



Plant Systematics



Michael G. Simpson

PLANT SYSTEMATICS

This Page Intentionally Left Blank

PLANT SYSTEMATICS

Michael G. Simpson



ELSEVIER
ACADEMIC
PRESS

Amsterdam Boston Heidelberg London
New York Oxford Paris San Diego
San Francisco Singapore Sydney Tokyo

Associate Acquisitions Editor: Kelly Sonnack
Publisher: Dana Dreibelbis
Project Manager: Sarah Hajduk
Marketing Manager: Linda Beattie
Cover Design: Eric DeCicco
Composition: Cepha Imaging Pvt Ltd
Cover and Interior Printer: Transcontinental Interglobe

Elsevier Academic Press
30 Corporate Drive, Suite 400, Burlington, MA 01803, USA
525 B Street, Suite 1900, San Diego, California 92101-4495, USA
84 Theobalds Road, London WC1X 8RR, UK

Cover Images (from left to right): *Magnolia grandiflora*, flowering magnolia (Magnoliaceae); *Graptopetalum paraguayense* (Crassulaceae); *Ferocactus* sp., barrel cactus (Cactaceae); *Faucaria tigrina*, tiger's jaw (Aizoaceae); *Nelumbo nucifera*, water-lotus (Nelumbonaceae); *Chorizanthe fimbriata*, fringed spineflower (Polygonaceae); *Swertia parryi*, deer's ears (Gentianaceae); *Stanhopea tigrina* (Orchidaceae).

This book is printed on acid-free paper. ☺

Copyright © 2006, Elsevier Inc. All rights reserved.

No part of this publication may be reproduced or transmitted in any form or by any means, electronic or mechanical, including photocopy, recording, or any information storage and retrieval system, without permission in writing from the publisher.

Permissions may be sought directly from Elsevier's Science & Technology Rights Department in Oxford, UK: phone: (+44) 1865 843830, fax: (+44) 1865 853333, e-mail: permissions@elsevier.com.uk. You may also complete your request on-line via the Elsevier homepage (<http://elsevier.com>), by selecting Customer Support and then Obtaining Permissions.

Library of Congress Cataloging-in-Publication Data

Simpson, Michael G. (Michael George), 1953-
Plant systematics / Michael G. Simpson.
p. cm.

Includes bibliographical references and index.

ISBN 0-12-644460-9 (casebound : alk. paper)

1. Plants—Classification. I. Title.

QK95.S566 2006

580'.12 dc22

2005014932

British Library Cataloguing in Publication Data

A catalogue record for this book is available from the British Library

ISBN 13: 978-0-12-644460-5

ISBN 10: 0-12-644460-9

For all information on all Elsevier Academic Press Publications
visit our Web site at www.books.elsevier.com

Printed in Canada

06 07 08 09 9 8 7 6 5 4 3 2 1



I wish to dedicate this book to three mentors I was very fortunate to know: Albert Radford, who taught critical thinking; P. Barry Tomlinson, who taught the fine art of careful observation; and Rolf Dahlgren, whose magnetic personality was inspirational. I also wish to thank my many students who have provided useful suggestions over the years, plus three writers who captured my interest in science and the wonder of it all: Isaac Asimov, Richard Feynman, and Carl Sagan.

CONTENTS

PREFACE.....	IX
ACKNOWLEDGMENTS	XI

UNIT I SYSTEMATICS

CHAPTER 1 PLANT SYSTEMATICS: AN OVERVIEW.....	3
CHAPTER 2 PHYLOGENETIC SYSTEMATICS.....	17

UNIT II EVOLUTION AND DIVERSITY OF PLANTS

CHAPTER 3 EVOLUTION AND DIVERSITY OF GREEN AND LAND PLANTS.....	51
CHAPTER 4 EVOLUTION AND DIVERSITY OF VASCULAR PLANTS.....	69
CHAPTER 5 EVOLUTION AND DIVERSITY OF WOODY AND SEED PLANTS	97
CHAPTER 6 EVOLUTION OF FLOWERING PLANTS	121
CHAPTER 7 DIVERSITY AND CLASSIFICATION OF FLOWERING PLANTS: AMBORELLALES, NYMPHAEALES, AUSTROBAILEYALES, MAGNOLIIDS, CERATOPHYLLALES, AND MONOCOTS	137
CHAPTER 8 DIVERSITY AND CLASSIFICATION OF FLOWERING PLANTS: EUDICOTS.....	227

UNIT III SYSTEMATIC EVIDENCE AND DESCRIPTIVE TERMINOLOGY

CHAPTER 9 PLANT MORPHOLOGY	347
CHAPTER 10 PLANT ANATOMY AND PHYSIOLOGY	409
CHAPTER 11 PLANT EMBRYOLOGY.....	437
CHAPTER 12 PALYNOLOGY	453
CHAPTER 13 PLANT REPRODUCTIVE BIOLOGY.....	465
CHAPTER 14 PLANT MOLECULAR SYSTEMATICS.....	477

UNIT IV RESOURCES IN PLANT SYSTEMATICS

CHAPTER 15 PLANT IDENTIFICATION	495
---------------------------------------	-----

CHAPTER 16	PLANT NOMENCLATURE	501
CHAPTER 17	PLANT COLLECTING AND DOCUMENTATION	517
CHAPTER 18	HERBARIA AND DATA INFORMATION SYSTEMS	525
APPENDIX 1	PLANT DESCRIPTION	535
APPENDIX 2	BOTANICAL ILLUSTRATIONS.....	541
APPENDIX 3	SCIENTIFIC JOURNALS IN PLANT SYSTEMATICS.....	545
GLOSSARY OF TERMS	547
INDEX.....	579

This Page Intentionally Left Blank

PREFACE

Plant Systematics is an introduction to the morphology, evolution, and classification of land plants. My objective is to present a foundation of the approach, methods, research goals, evidence, and terminology of plant systematics and to summarize information on the most recent knowledge of evolutionary relationships of plants as well as practical information vital to the field. I have tried to present the material in a condensed, clear manner, such that the beginning student can better digest the more important parts of the voluminous information in the field and acquire more detailed information from the literature.

The book is meant to serve students at the college graduate and upper undergraduate levels in plant systematics or taxonomy courses, although portions of the book may be used in flora courses and much of the book could be used in general courses in plant morphology, diversity, or general botany.

Each chapter has an expanded Table of Contents on the first page, a feature that my students recommended as very useful. Numerous line drawings and color photographs are used throughout. A key feature is that illustrated plant material is often dissected and labeled to show important diagnostic features. At the end of each chapter are (1) Review Questions, which go over the chapter material; (2) Exercises, whereby a student may apply the material; and (3) References for Further Study, listing some of the basic and recent references. Literature cited in the references is not exhaustive, so the student is encouraged to do literature searches on his/her own (see Appendix 3).

The book is classified into units, which consist of two or more chapters logically grouped together. Of course, a given instructor may choose to vary the sequence of these units or the chapters within, depending on personal preference and the availability of plant material. There is a slight amount of repetition between chapters of different units, but this was done so that chapters could be used independently of one another.

Unit 1, Systematics, gives a general overview of the concepts and methods of the field of systematics. Chapter 1 serves as an introduction to the definition, relationships, classification, and importance of plants and summarizes the basic concepts and principles of systematics, taxonomy, evolution, and phylogeny. Chapter 2 covers the details of phylogenetic systematics, and the theory and methodology for inferring phylogenetic trees or cladograms.

Unit 2, Evolution and Diversity of Plants, describes in detail the characteristics and classification of plants. The six chapters of this unit are intended to give the beginning student a basic understanding of the evolution of Green and Land Plants (Chapter 3), Vascular Plants (Chapter 4), Woody and Seed Plants (Chapter 5), and Flowering Plants (Chapters 6-8). Chapters 3-5 are formatted into two major sections. The first section presents cladograms (phylogenetic trees), which portray the evolutionary history of the group. Each of the major derived evolutionary features (apomorphies) from that cladogram is described and illustrated, with emphasis on the possible adaptive significance of these features. This evolutionary approach to plant systematics makes learning the major plant groups and their features conceptually easier than simply memorizing a static list of characteristics. Treating these features as the products of unique evolutionary events brings them to life, especially when their possible adaptive significance is pondered. The second section of Chapters 3 through 5 presents a brief survey of the diversity of the group in question. Exemplars within major groups are described and illustrated, such that the student may learn to recognize and know the basic features of the major lineages of plants.

Because they constitute the great majority of plants, the flowering plants, or angiosperms, are covered in three chapters. Chapter 6 deals with the evolution of flowering plants, describing the apomorphies for that group and presenting a brief coverage of their origin. Chapters 7 and 8 describe specific groups of flowering plants. In Chapter 7 the non-eudicot groups are treated, including basal angiosperms and the monocotyledons. Chapter 8 covers the eudicots, which make up the great majority of angiosperms. Numerous flowering plant families are described in detail, accompanied by photographs and illustrations. Reference to Chapter 9 and occasionally to Chapters 10-14 (or use of the comprehensive Glossary) may be needed with regard to the technical terms. Because of their great number, only a limited number of families are included, being those that are commonly encountered or for which material is usually available to the beginning student. I have tried to emphasize diagnostic features that a student might use to recognize a plant family, and have included some economically important uses of family members. The Angiosperm Phylogeny Group II system of classification is

used throughout (with few exceptions). This system uses orders as the major taxonomic rank in grouping families of close relationship and has proven extremely useful in dealing with the tremendous diversity of the flowering plants.

Unit 3, Systematic Evidence and Descriptive Terminology, begins with a chapter on plant morphology (Chapter 9). Explanatory text, numerous diagrammatic illustrations, and photographs are used to train beginning students to precisely and thoroughly describe a plant morphologically. Appendices 1 and 2 (see below) are designed to be used along with Chapter 9. The other chapters in this unit cover the basic descriptive terminology of plant anatomy (Chapter 10), plant embryology (Chapter 11), palynology (Chapter 12), plant reproductive biology (Chapter 13), and plant molecular systematics (Chapter 14). The rationale for including these in a textbook on plant systematics is that features from these various fields are described in systematic research and are commonly utilized in phylogenetic reconstruction and taxonomic delimitation. In particular, the last chapter on plant molecular systematics reviews the basic techniques and the types of data acquired in what has perhaps become in recent years the most fruitful of endeavors in phylogenetic reconstruction.

Unit 4, Resources in Plant Systematics, discusses some basics that are essential in everyday systematic research. Plant identification (Chapter 15) contains a summary of both standard dichotomous keys and computerized polythetic keys and reviews practical identification methods. The chapter on nomenclature (Chapter 16) summarizes the basic rules of the most recent International Code of Botanical Nomenclature, including the steps needed in the valid publication of a new species and a review of botanical names. A chapter on plant collecting and documentation (Chapter 17) emphasizes both correct techniques for collecting plants and thorough data acquisition, the latter of which has become increasingly important today in biodiversity studies and conservation biology. Finally, the chapter on herbaria and data information systems (Chapter 18) reviews the basics of herbarium management, emphasizing the role of computerized database systems in plant collections for analyzing and synthesizing morphological, ecological, and biogeographic data.

Lastly, three **Appendices** and a **Glossary** are included. I have personally found each of these addenda to be of value in

my own plant systematics courses. Appendix 1 is a list of characters used for detailed plant descriptions. This list is useful in training students to write descriptions suitable for publication. Appendix 2 is a brief discussion of botanical illustration. I feel that students need to learn to draw, in order to develop their observational skills. Appendix 3 is a listing of scientific journals in plant systematics, with literature exercises. The Glossary defines all terms used in the book and indicates synonyms, adjectival forms, plurals, abbreviations, and terms to compare.

By the time of publication, two Web sites will be available to be used in conjunction with the textbook: (1) a Student Resources site (<http://books.elsevier.com/companion/0126444609>), with material that is universally available; and (2) an Instructor Resources site (<http://books.elsevier.com/manualsprotected/0126444609>), with material that is password protected. Please contact your sales representative at <textbooks@elsevier.com> for access to the Instructor Resources site.

Throughout the book, I have attempted to adhere to **W-H-Y**, **What-How-Why**, in organizing and clarifying chapter topics: (1) What is it? What is the topic, the basic definition? (I am repeatedly amazed that many scientific arguments could have been resolved at the start by a clear statement or definition of terms.) (2) How is it done? What are the materials and methods, the techniques of data acquisition, the types of data analysis? (3) Why is it done? What is the purpose, objective, or goal; What is the overriding paradigm involved? How does the current study or topic relate to others? This simple W-H-Y method, first presented to me by one of my mentors, A. E. Radford, is useful to follow in any intellectual endeavor. It is a good lesson to teach one's students, and helps both in developing good writing skills and in critically evaluating any topic.

Finally, I would like to propose that each of us, instructors and students, pause occasionally to evaluate why it is that we do what we do. Over the years I have refined my ideas and offer these suggestions as possible goals: 1) to realize and explore the beauty, grandeur, and intricacy of nature; 2) to engage in the excitement of scientific discovery; 3) to experience and share the joy of learning. It is in this spirit that I sincerely hope the book may be of use to others.

ACKNOWLEDGMENTS

I sincerely thank Andy Bohonak, Bruce Baldwin, Lisa Campbell, Travis Columbus, Bruce Kirchoff, Lucinda McDade, Kathleen Pryer (and her lab group), Jon Rebman, and several anonymous reviewers for their comments on various chapters of the book and Peter Stevens for up-to-date information on higher level classification of angiosperms. As always, they bear no responsibility for any mistakes, omissions, incongruities, misinterpretations, or general stupidities.

Almost all of the illustrations and photographs are the product of the author. I thank the following for additions to these (in order of appearance in text):

The Jepson Herbarium (University of California Press) gave special permission to reproduce the key to the Crassulaceae (Reid Moran, author) in Figure 1.7.

Rick Bizzoco contributed the images of *Chlamydomonas reinhardtii* in Figures 3.2C and 3.3A.

Linda Graham contributed the image of *Coleochaete* in Figure 3.6A.

Figure 4.11A was reproduced from Kidston, R. and W. H. Lang. 1921. Transactions of the Royal Society of Edinburgh. vol. 52(4):831 902.

Figure 5.10 was reproduced and modified from Swamy, B. G. L. 1948. American Journal of Botany 35: 77 88, by permission.

Figure 5.15A,B was reproduced from: Beck, C. B. 1962. American Journal of Botany 49: 373 382, by permission.

Figure 5.15C was reproduced from Stewart, W. N., and T. Delevoryas. 1956. Botanical Review 22: 45 80, by permission.

Figure 5.23B was reproduced from Esau, K. 1965. Plant Anatomy. J. Wiley and sons, New York, by permission.

Mark Olsen contributed the images of *Welwitschia mirabilis* in Figure 5.24E G.

Figure 6.5 was based upon Jack, T. 2001. Relearning our ABCs: new twists on an old model. Trends in Plant Science 6: 310 316.

Figure 6.18A C was redrawn from Thomas, H. H. 1925. Philosophical Transactions of the Royal Society of London 213: 299 363.

Figure 6.18D was contributed by K. Simons and David Dilcher (); Figure 6.18E was contributed by David Dilcher () and Ge Sun.

Stephen McCabe contributed the images of *Amborella* in Figures 7.3A,C.

The Arboretum at the University of California-Santa Cruz contributed the image of *Amborella* in Figure 7.3B.

Sandra Floyd provided the image of *Amborella* in Figure 7.3D.

Jeffrey M. Osborn and Mackenzie L. Taylor contributed the images of the Cabombaceae of Figure 7.5.

Jack Scheper contributed the image of *Illicium floridanum* in Figure 7.6A.

Figure 7.16 was reproduced from Behnke, H.-D. 1972. Botanical Review 38: 155 197, by permission.

Constance Gramlich contributed the image of *Amorphophallus* in Figure 7.23C.

Wayne Armstrong contributed the image of a flowering *Wolffia* in Figure 7.23G.

John Kress contributed the Zingiberales drawing of Figure 7.53.

Figure 8.11B was reproduced from Behnke, H.-D. 1972. Botanical Review 38: 155 197, by permission.

David G. Smith contributed the images of *Phryma leptostachya* in Figure 8.71.

Figure 9.12 was redrawn from Hickey, L. J. 1973. American Journal of Botany 60: 17 33, by permission.

Darren Burton prepared several illustrations in Chapter 9.

Figure 13.4A was redrawn from Weberling. 1989. Morphology of Flowers and Inflorescences. Cambridge University Press, Cambridge, New York, by permission.

Figure 13.4B was redrawn from Kohn et al. 1996. Evolution 50:1454 1469, by permission.

Jon Rebman contributed the images of Figure 13.7D,E.

Figure 14.4 was redrawn from Wakasugi, T., M. Sugita, T. Tsudzuki, and M. Sugiura. 1998. Plant Molecular Biology Reporter 16: 231 241, by permission.

The Herbarium at the San Diego Natural History Museum contributed the images of Figure 17.2.

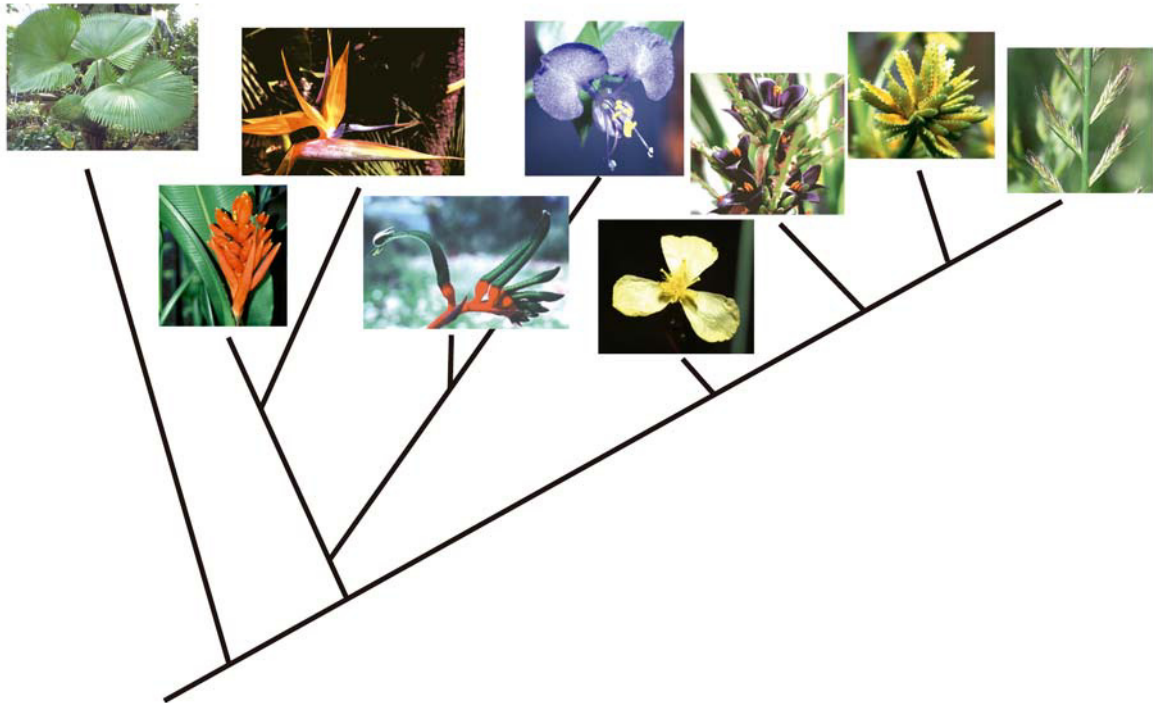
Jon Rebman contributed the image of the herbarium sheet in Figure 18.2.

Dirina Estrella contributed the stippled line drawing of Appendix 2.

This Page Intentionally Left Blank

I

SYSTEMATICS



This Page Intentionally Left Blank

1

PLANT SYSTEMATICS: AN OVERVIEW

PLANTS	3	Taxonomy	10
What Is a Plant?	3	Phylogeny	13
Plants and the Evolution of Life	3	Why Study Systematics?	13
Land Plants	5	REVIEW QUESTIONS	15
Why Study Plants?	5	EXERCISES	16
SYSTEMATICS	9	REFERENCES FOR FURTHER STUDY	16
What Is Systematics?	9		
Evolution	10		

This book is about a fascinating field of biology called plant systematics. The purpose of this chapter is to introduce the basics: what a plant is, what systematics is, and the reasons for studying plant systematics.

PLANTS

WHAT IS A PLANT?

This question can be answered in either of two conceptual ways. One way, the traditional way, is to define groups of organisms such as plants by the characteristics they possess. Thus, historically, plants included those organisms that possess photosynthesis, cell walls, spores, and a more or less sedentary behavior. This traditional grouping of plants contained a variety of microscopic organisms, all of the algae, and the more familiar plants that live on land. A second way to answer the question “What is a plant?” is to evaluate the evolutionary history of life and to use that history to delimit the groups of life. We now know from repeated research studies that some of the photosynthetic organisms evolved independently of one another and are not closely related.

Thus, the meaning or definition of the word *plant* can be ambiguous and can vary from person to person. Some still like to treat plants as an unnatural assemblage, defined by

the common (but independently evolved) characteristic of photosynthesis. However, delimiting organismal groups based on evolutionary history has gained almost universal acceptance. This latter type of classification directly reflects the patterns of that evolutionary history and can be used to explicitly test evolutionary hypotheses (discussed later; see Chapter 2).

An understanding of what plants are requires an explanation of the evolution of life in general.

PLANTS AND THE EVOLUTION OF LIFE

Life is currently classified as three major groups (sometimes called domains) of organisms: **Archaea** (also called **Archaeobacteria**), **Bacteria** (also called **Eubacteria**), and **Eukarya** or **eukaryotes** (also spelled eucaryotes). The evolutionary relationships of these groups are summarized in the simplified evolutionary tree or cladogram of Figure 1.1. The Archaea and Bacteria are small, mostly unicellular organisms that possess circular DNA, replicate by binary fission, and lack membrane-bound organelles. The two groups differ from one another in the chemical structure of certain cellular components. Eukaryotes are unicellular or multicellular organisms that possess linear DNA (organized as histone-bound chromosomes), replicate by mitotic and often meiotic division, and possess membrane-bound organelles such as nuclei, cytoskeletal structures, and (in almost all) mitochondria (Figure 1.1).

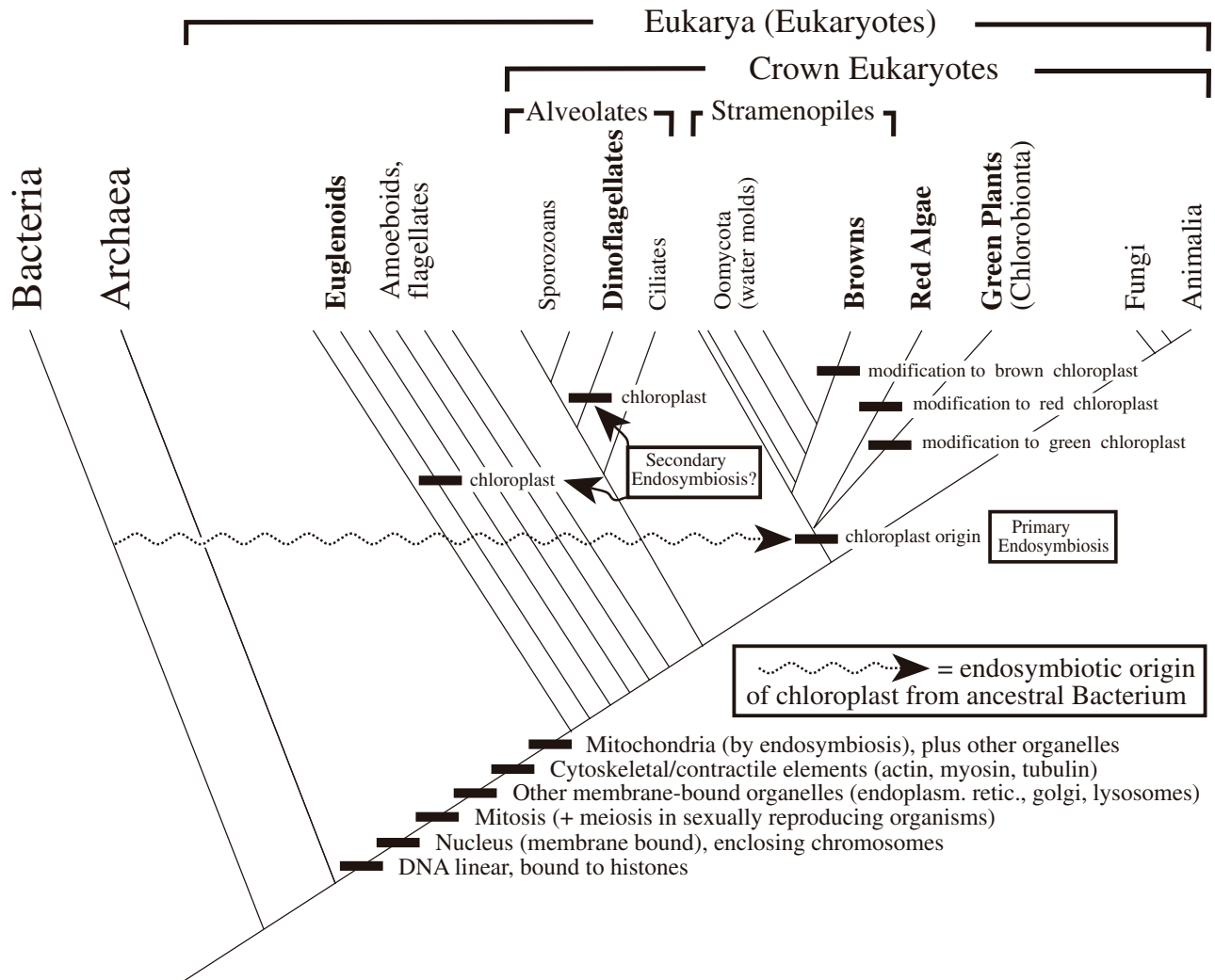


FIGURE 1.1 Simplified cladogram (evolutionary tree) of life (modified from Sogin 1994, Kumar & Rzhetsky 1996, and Yoon et al. 2002), illustrating the independent origin of chloroplasts via endosymbiosis (arrows) in the euglenoids, dinoflagellates, brown plants, red algae, and green plants. Eukaryotic groups containing photosynthetic, chloroplast-containing organisms in bold. The relative order of evolutionary events is unknown.

Some of the unicellular bacteria (including, e.g., the Cyanobacteria, or blue-greens) carry on photosynthesis, a biochemical system in which light energy is used to synthesize high-energy compounds from simpler starting compounds, carbon dioxide and water. These photosynthetic bacteria have a system of internal membranes called thylakoids, within which are embedded photosynthetic pigments, compounds that convert light energy to chemical energy. Of the several groups of eukaryotes that are photosynthetic, all have specialized photosynthetic organelles called **chloroplasts**, which resemble photosynthetic bacteria in having pigment-containing thylakoid membranes.

How did chloroplasts evolve? It is now largely accepted that the chloroplasts of eukaryotes originated by the engulfment of an ancestral photosynthetic bacterium (probably a

cyanobacterium) by an ancestral eukaryotic cell, such that the photosynthetic bacterium continued to live and ultimately multiply *inside* the eukaryotic cell (Figure 1.2). The evidence for this is the fact that chloroplasts, like bacteria today (a) have their own single-stranded, circular DNA; (b) have a smaller sized, 70S ribosome; and (c) replicate by binary fission. These engulfed photosynthetic bacteria provided high-energy products to the eukaryotic cell; the host eukaryotic cell provided a more beneficial environment for the photosynthetic bacteria. The condition of two species living together in close contact is termed symbiosis, and the process in which symbiosis results by the engulfment of one cell by another is termed **endosymbiosis**. Over time, these endosymbiotic, photosynthetic bacteria became transformed structurally and functionally, retaining their own DNA and

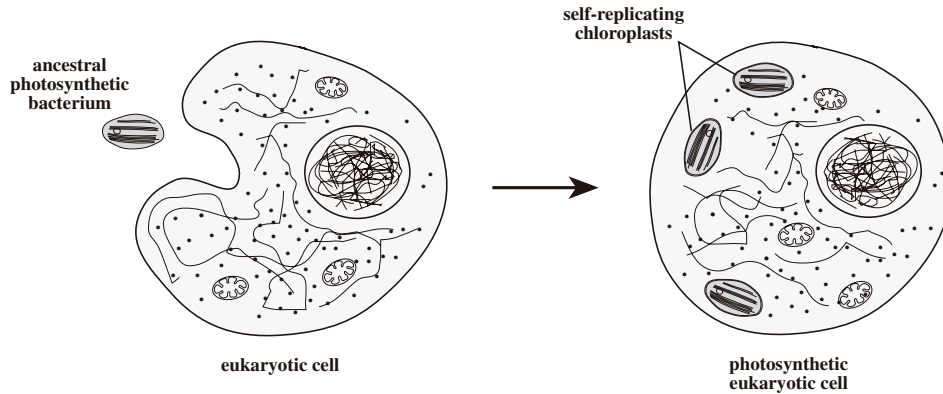


FIGURE 1.2 Diagrammatic illustration of the origin of chloroplasts by endosymbiosis of ancestral photosynthetic bacterium within ancestral eukaryotic cell.

the ability to replicate, but losing the ability to live independently of the host cell. In fact, over time there has been a transfer of some genes from the DNA of the chloroplast to the nuclear DNA of the eukaryotic host cell, making the two biochemically interdependent.

The most recent data from molecular systematic studies indicates that this so-called primary endosymbiosis of the chloroplast likely occurred one time, a shared evolutionary novelty of the red algae, green plants, and stramenopiles (which include the brown algae and relatives; Figure 1.1). This early chloroplast became modified with regard to photosynthetic pigments, thylakoid structure, and storage products into forms characteristic of the red algae, green plants, and browns (see Figure 1.1). In addition, chloroplasts may have been lost in some lineages, e.g., in the Oomycota (water molds) of the Stramenopiles. Some lineages of these groups may have acquired chloroplasts via secondary endosymbiosis, which occurred by the engulfment of an ancestral chloroplast-containing *eukaryote* by another eukaryotic cell. The euglenoids and the dinoflagellates, two other lineages of photosynthetic organisms, may have acquired chloroplasts by this process (Figure 1.1). The exact story is yet to be elucidated.

LAND PLANTS

Of the major groups of photosynthetic eukaryotes, the green plants (also called the Chlorobionta) are united primarily by distinctive characteristics of the green plant chloroplast with respect to photosynthetic pigments, thylakoid structure, and storage compounds (see Chapter 3 for details). Green plants include both the predominately aquatic green algae and a group known as embryophytes (formally, the Embryophyta), usually referred to as the land plants (Figure 1.3). The land plants are united by several evolutionary novelties that were adaptations to making the transition from an aquatic environment to living on land. These include (1) an outer cuticle,

which aids in protecting tissues from desiccation; (2) specialized gametangia (egg and sperm producing organs) that have an outer, protective layer of sterile cells; and (3) an intercalated diploid phase in the life cycle, the early, immature component of which is termed the embryo (hence, embryophytes; see Chapter 3 for details).

Just as the green plants include the land plants, the land plants are inclusive of the vascular plants (Figure 1.3), the latter being united by the evolution of an independent sporophyte and xylem and phloem vascular conductive tissue (see Chapter 4). The vascular plants are inclusive of the seed plants (Figure 1.3), which are united by the evolution of wood and seeds (see Chapter 5). Finally, seed plants include the angiosperms (Figure 1.3), united by the evolution of the flower, including carpels and stamens, and by a number of other specialized features (see Chapters 6–8).

For the remainder of this book, the term *plant* is treated as equivalent to the embryophytes, the land plants. The rationale for this is partly that land plants make up a so-called natural, monophyletic group, whereas the photosynthetic eukaryotes as a whole are an unnatural, paraphyletic group (see section on **Phylogeny**, Chapter 2). And, practically, it is land plants that most people are talking about when they refer to plants, including those in the field of plant systematics. However, as noted before, the word *plant* can be used by some to refer to other groupings; when in doubt, get a precise clarification.

WHY STUDY PLANTS?

The tremendous importance of plants cannot be overstated. Without them, we and most other species of animals (and so many other groups of organisms) wouldn't be here. Photosynthesis in plants and the other photosynthetic organisms changed the earth in two major ways. First, the fixation of carbon dioxide and the release of molecular oxygen in photosynthesis directly altered the earth's atmosphere over

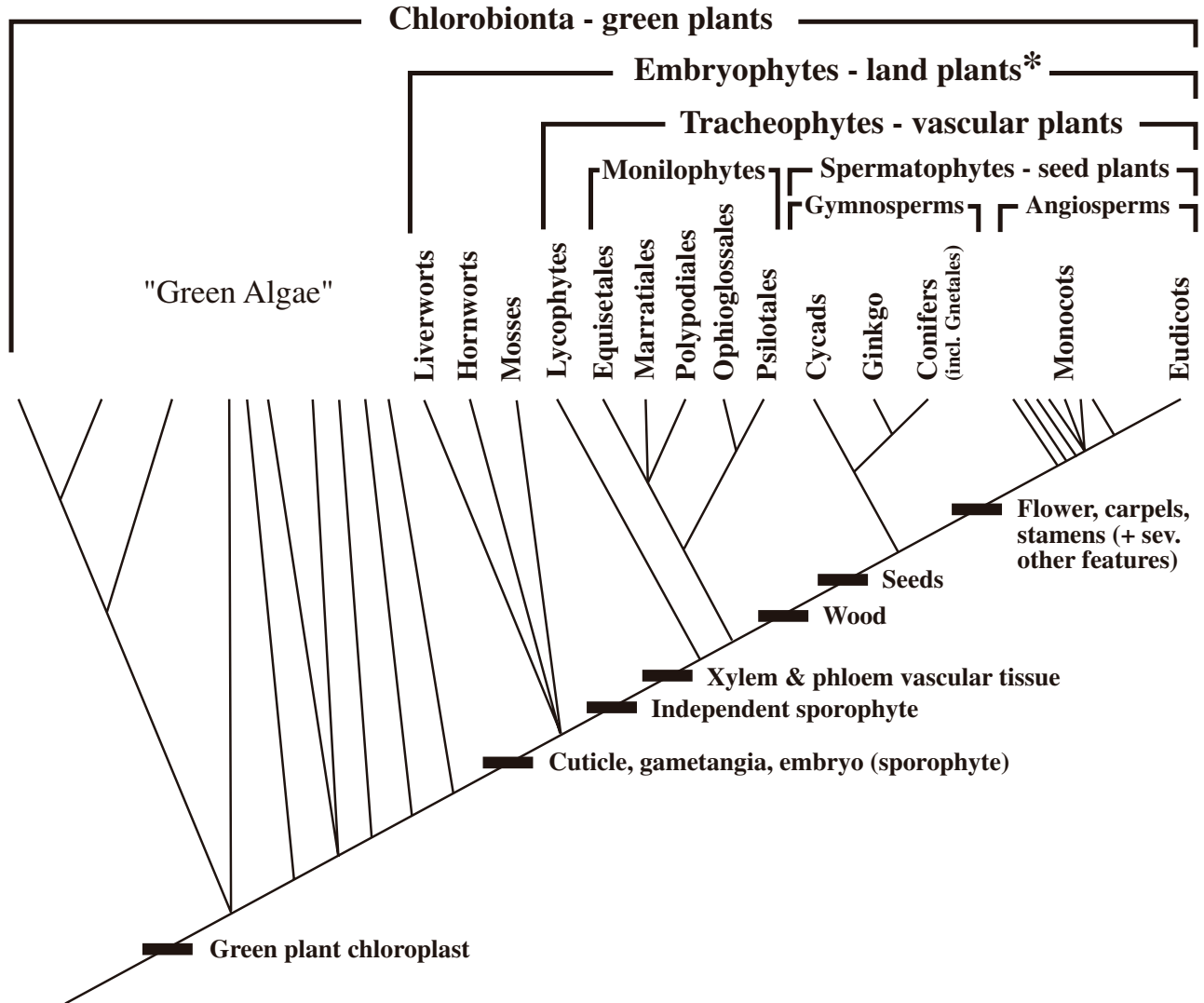


FIGURE 1.3 Simplified cladogram (evolutionary tree) of the green plants, illustrating major extant groups and evolutionary events (or apomorphies, hash marks). *Embryophytes are treated as plants in this book.

billions of years. What used to be an atmosphere deficient in oxygen underwent a gradual change. As a critical mass of oxygen accumulated in the atmosphere, selection for oxygen-dependent respiration occurred (via oxidative phosphorylation in mitochondria), which may have been a necessary precursor in the evolution of many multicellular organisms, including all animals. In addition, an oxygen-rich atmosphere permitted the establishment of an upper atmosphere ozone layer, which shielded life from excess UV radiation. This allowed organisms to inhabit more exposed niches that were previously inaccessible.

Second, the compounds that photosynthetic species produce are utilized, directly or indirectly, by nonphotosynthetic, heterotrophic organisms. For virtually all land creatures and

many aquatic ones as well, land plants make up the so-called primary producers in the food chain, the source of high-energy compounds such as carbohydrates, structural compounds such as certain amino acids, and other compounds essential to metabolism in some heterotrophs. Thus, most species on land today, including millions of species of animals, are absolutely dependent on plants for their survival. As primary producers, plants are the major components of many communities and ecosystems. The survival of plants is essential to maintaining the health of those ecosystems, the severe disruption of which could bring about rampant species extirpation or extinction and disastrous changes in erosion, water flow, and ultimately climate.

To humans, plants are also monumentally important in numerous, direct ways (Figures 1.4, 1.5). Agricultural plants,

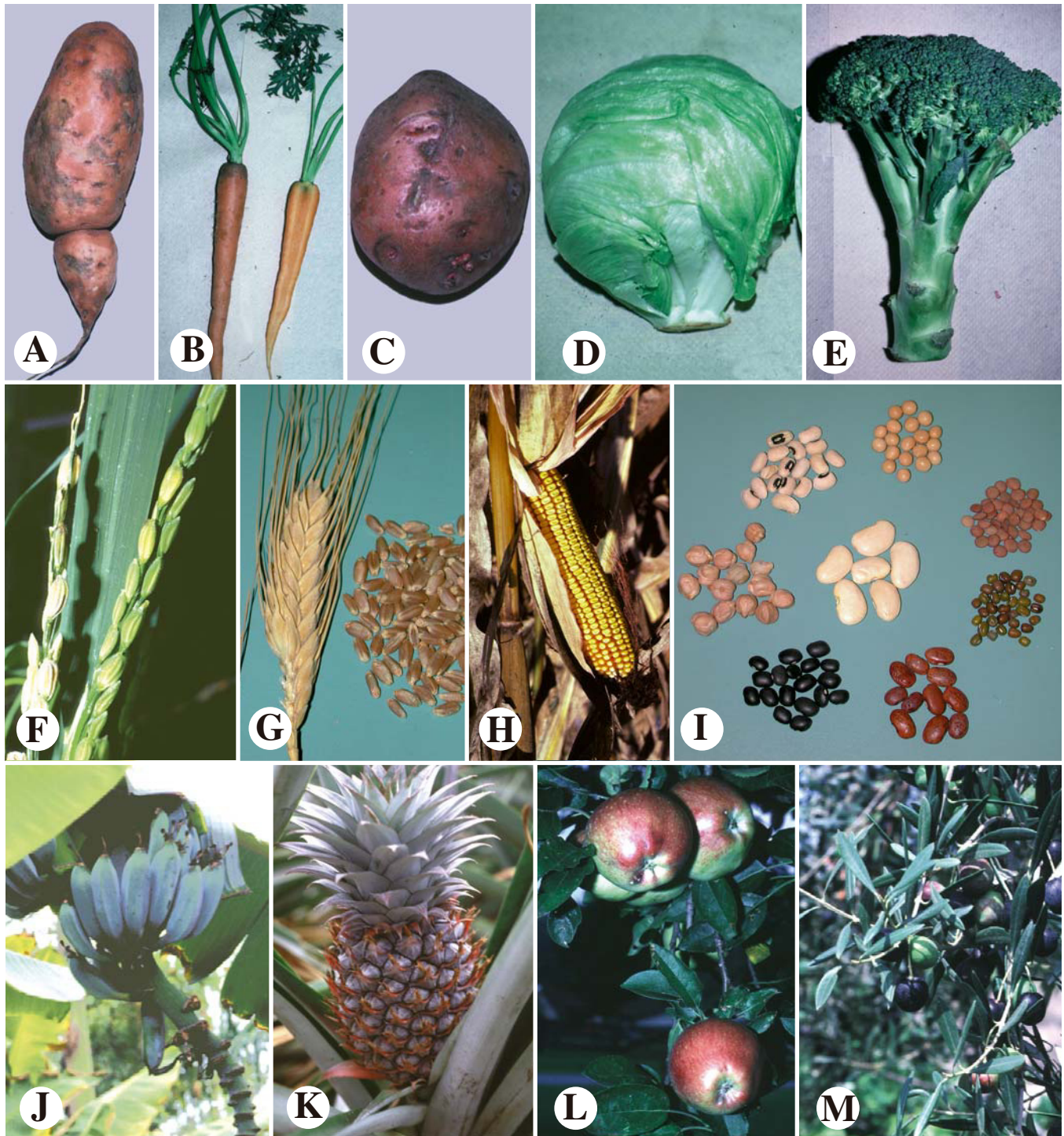


FIGURE 1.4 Examples of economically important plants. A–E. Vegetables. A. *Ipomoea batatas*, sweet potato (root). B. *Daucus carota*, carrot (root). C. *Solanum tuberosum*, potato (stem). D. *Lactuca sativa*, lettuce (leaves). E. *Brassica oleracea*, broccoli (flower buds). F–I. Fruits, dry (grains). F. *Oryza sativa*, rice. G. *Triticum aestivum*, bread wheat. H. *Zea mays*, corn. I. Seeds (pulse legumes), from top, clockwise to center: *Glycine max*, soybean; *Lens culinaris*, lentil; *Phaseolus aureus*, mung bean; *Phaseolus vulgaris*, pinto bean; *Phaseolus vulgaris*, black bean; *Cicer arietinum*, chick-pea/garbanzo bean; *Vigna unguiculata*, black-eyed pea; *Phaseolus lunatus*, lima bean. J–M. Fruits, fleshy. J. *Musa paradisiaca*, banana. K. *Ananas comosus*, pineapple. L. *Malus pumila*, apple. M. *Olea europaea*, olive.

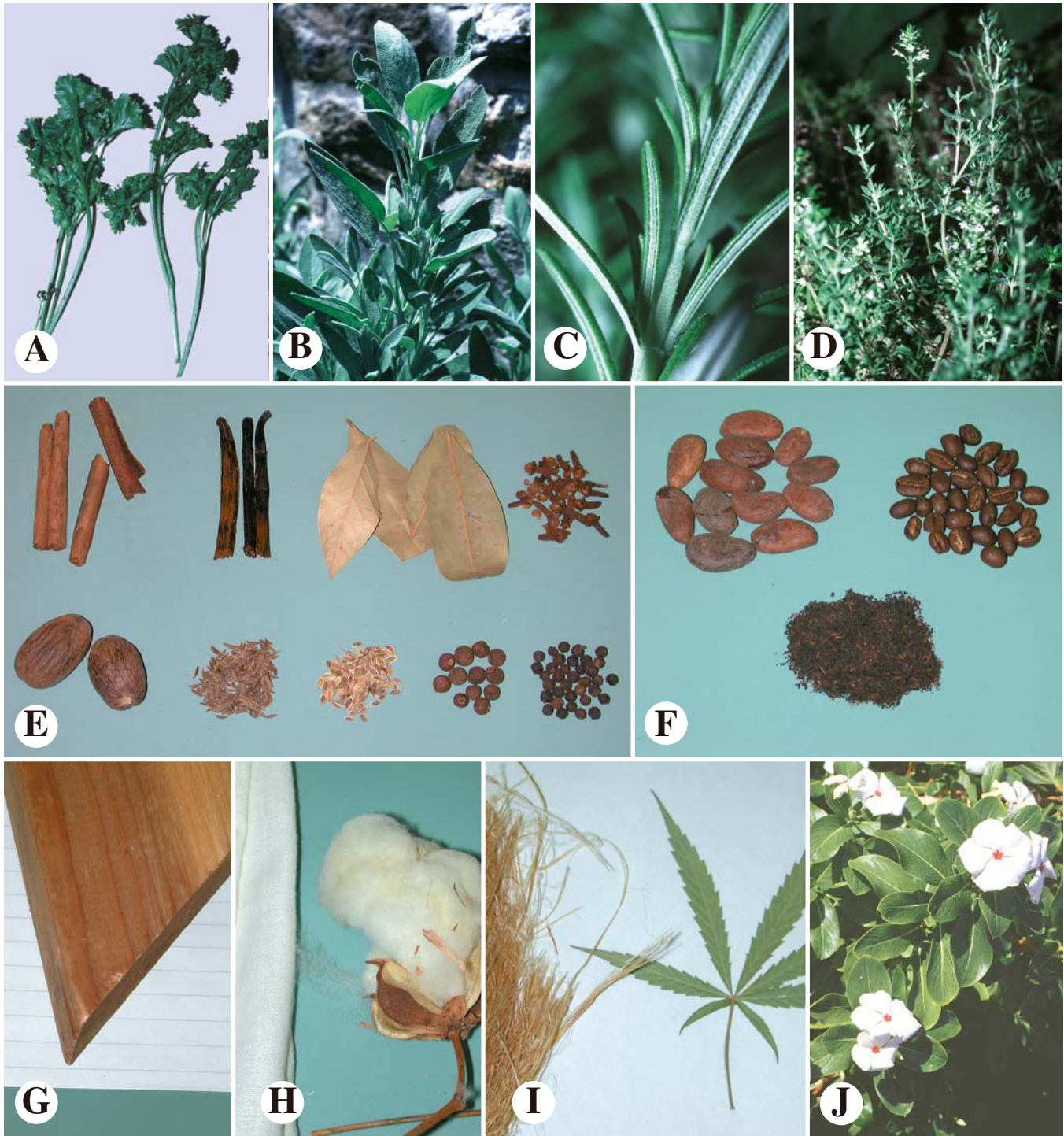


FIGURE 1.5 Further examples of economically important plants. **A–D.** Herbs. **A.** *Petroselinum crispum*, parsley. **B.** *Salvia officinalis*, sage. **C.** *Rosmarinus officinalis*, rosemary. **D.** *Thymus vulgaris*, thyme. **E.** Spices and herbs, from upper left: *Cinnamomum cassia/zeylanicum*, cinnamon (bark); *Vanilla planifolia*, vanilla (fruit); *Laurus nobilis*, laurel (leaf); *Syzygium aromaticum*, cloves (flower buds); *Myristica fragrans*, nutmeg (seed); *Carum carvi*, caraway (fruit); *Anethum graveolens*, dill (fruit); *Pimenta dioica*, allspice (seed); *Piper nigrum*, pepper (seed). **F.** Flavoring plants, from upper left, clockwise: *Theobroma cacao*, chocolate (seeds); *Coffea arabica*, coffee (seeds); *Thea sinensis*, tea (leaves). **G.** Wood products: lumber (*Sequoia sempervirens*, redwood), and paper derived from wood pulp. **H.** Fiber plant. *Gossypium* sp., cotton (seed trichomes), one of the most important natural fibers. **I.** Euphoric, medicinal, and fiber plant. *Cannabis sativa*, marijuana, hemp; stem fibers used in twine, rope, and cloth; resins contain the euphoric and medicinal compound tetrahydrocannabinol. **J.** Medicinal plant. *Catharanthus roseus*, Madagascar periwinkle, from which is derived vincristine and vinblastine, used to treat childhood leukemia.

most of which are flowering plants, are our major source of food. We utilize all plant parts as food products: roots (e.g., sweet potatoes and carrots; Figure 1.4A,B); stems (e.g., yams, cassava/manioc, potatoes; Figure 1.4C); leaves (e.g., cabbage, celery, lettuce; Figure 1.4D); flowers (e.g., cauliflower and broccoli; Figure 1.4E); and fruits and seeds, including grains such as rice (Figure 1.4F), wheat (Figure 1.4G), corn (Figure 1.4H), rye, barley, and oats, legumes such as beans and peas (Figure 1.4I), and a plethora of fruits such as bananas (Figure 1.4J), tomatoes, peppers, pineapples (Figure 1.4K), apples (Figure 1.4L), cherries, peaches, melons, kiwis, citrus, olives (Figure 1.4M), and others too numerous to mention. Other plants are used as flavoring agents, such as herbs (Figure 1.5A–D) and spices (Figure 1.5E), as stimulating beverages, such as chocolate, coffee, tea, and cola (Figure 1.5F), or as alcoholic drinks, such as beer, wine, distilled liquors, and sweet liqueurs. Woody trees of both conifers and flowering plants are used structurally for lumber and for pulp products such as paper (Figure 1.5G). In tropical regions, bamboos, palms, and a variety of other species serve in the construction of human dwellings. Plant fibers are used to make thread for cordage (such as sisal), for sacs (such as jute for burlap), and for textiles (most notably cotton, Figure 1.5H, but also linen and hemp, Figure 1.5I). In many cultures, plants or plant products are used as euphorics or hallucinogenics (whether legally or illegally), such as marijuana (Figure 1.5I), opium, cocaine, and a great variety of other species that have been used by indigenous peoples for centuries. Plants are important for their aesthetic beauty, and the cultivation of plants as ornamentals is an important industry. Finally, plants have great medicinal significance, to treat a variety of illnesses or to maintain good health. Plant products are very important in the pharmaceutical industry; their compounds are extracted, semisynthesized, or used as templates to synthesize new drugs. Many modern drugs, from aspirin (originally derived from the bark of willow trees) to vincristine and vinblastine (obtained from the Madagascar periwinkle, used to treat childhood leukemia; Figure 1.5J), are ultimately derived from plants. In addition, various plant parts of a great number of species are used whole or are processed as so-called herbal supplements, which have become tremendously popular recently.

The people, methods, and rationale concerned with the **plant sciences** (defined here as the study of land plants) are as diverse as are the uses and importance of plants. Some of the fields in the plant sciences are very practically oriented. Agriculture and horticulture deal with improving the yield or disease resistance of food crops or cultivated ornamental plants, e.g., through breeding studies and identifying new cultivars. Forestry is concerned with the cultivation and

harvesting of trees used for lumber and pulp. Pharmacognosy deals with crude natural drugs, often of plant origin. In contrast to these more practical fields of the plant sciences, the pure sciences have as their goal the advancement of scientific knowledge (understanding how nature works) through research, regardless of the practical implications. But many aspects of the pure sciences also have important practical applications, either directly by applicable discovery or indirectly by providing the foundation of knowledge used in the more practical sciences. Among these are plant anatomy, dealing with cell and tissue structure and development; plant chemistry and physiology, dealing with biochemical and biophysical processes and products; plant molecular biology, dealing with the structure and function of genetic material; plant ecology, dealing with interactions of plants with their environment; and, of course, plant systematics.

Note that a distinction should be made between botany and plant sciences. **Plant sciences** is the study of plants, treated as equivalent to land plants here. **Botany** is the study of most organisms traditionally treated as plants, including virtually all eukaryotic photosynthetic organisms (land plants and the several groups of algae) plus other eukaryotic organisms with cell walls and spores (true fungi and groups that were formerly treated as fungi, such as the Oomycota and slime molds). Thus, in this sense, botany is inclusive of but broader than the plant sciences. Recognition of both botany and plant sciences as fields of study can be useful, although how these fields are defined can vary and may require clarification.

SYSTEMATICS

WHAT IS SYSTEMATICS?

Systematics is defined in this book as a science that includes and encompasses traditional **taxonomy**, the description, identification, nomenclature, and classification of organisms, and that has as its primary goal the reconstruction of **phylogeny**, or evolutionary history, of life. This definition of *systematics* is not novel, but neither is it universal. Others in the field would treat taxonomy and systematics as separate but overlapping areas; still others argue that historical usage necessitates what is in essence a reversal of the definitions used here. But words, like organisms, evolve. The use of *systematics* to describe an all-encompassing field of endeavor is both most useful and represents the consensus of how most specialists in the field use the term, an example being the journal *Systematic Botany*, which contains articles both in traditional taxonomy and phylogenetic reconstruction. Plant systematics is studied by acquiring, analyzing, and synthesizing information about plants and

plant parts, the content and methodology of which is the topic for the remainder of this book.

Systematics is founded in the principles of **evolution**, its major premise being that there is one phylogeny of life. The goal of systematists is, in part, to discover that phylogeny.

EVOLUTION

Evolution, in the broadest sense, means change and can be viewed as the cumulative changes occurring since the origin of the universe some 15 billion years ago. Biological evolution, the evolution of life, may be defined (as it was by Charles Darwin) as descent with modification. **Descent** is the transfer of genetic material (enclosed within a cell, the unit of life) from parent(s) to offspring over time. This is a simple concept, but one that is important to grasp and ponder thoroughly. Since the time that life first originated some 3.8 billion years ago, *all life has been derived from preexisting life*. Organisms come to exist by the transfer of genetic material, within a surrounding cell, from one or more parents. Descent may occur by simple clonal reproduction, such as a single bacterial cell parent dividing by fission to form two offspring cells or a land plant giving rise to a vegetative propagule. It may also occur by complex sexual reproduction (Figure 1.6A), in which each of two parents produces specialized gametes (e.g., sperm and egg cells), each of which has half the complement of genetic material, the result of meiosis. Two of the gametes fuse together to form a new cell, the zygote, which may develop into a new individual or may itself divide by meiosis to form gametes. Descent through time results in the formation of a **lineage**, or **clade** (Figure 1.6B,C), a set of organisms interconnected through time and space by the transfer of genetic material from parents to offspring. So, in a very literal sense, we and all other forms of life on earth are connected in time and in space by descent, the transfer of DNA (actually the pattern of DNA) from parent to offspring (ancestor to descendant), generation after generation.

The **modification** component of evolution refers to a change in the genetic material that is transferred from parent(s) to offspring, such that the genetic material of the offspring is different from that of the parent(s). This modification may occur either by mutation, which is a direct alteration of DNA, or by genetic recombination, whereby existing genes are reshuffled in different combinations (during meiosis, by crossing over and independent assortment). Systematics is concerned with the identification of the unique modifications of evolution (see later discussion).

It should also be asked, *what* evolves? Although genetic modification may occur in offspring relative to their parents, individual organisms do not generally evolve. This is because a new individual begins when it receives its complement of

DNA from the parent(s); that individual's DNA does not change during its/his/her lifetime (with the exception of relatively rare, nonreproductive somatic mutations that cannot be transmitted to the next generation). The general units of evolution are populations and species. A **population** is a group of individuals of the same species that is usually geographically delimited and that typically have a significant amount of gene exchange. **Species** may be defined in a number of ways, one definition being a distinct lineage that, in sexually reproducing organisms, consists of a group of generally intergrading, interbreeding populations that are essentially reproductively isolated from other such groups. With changes in the genetic makeup of offspring (relative to parents), the genetic makeup of populations and species changes over time.

In summary, evolution is descent with modification occurring by a change in the genetic makeup (DNA) of populations or species over time. How does evolution occur? Evolutionary change may come about by two major mechanisms: (1) **genetic drift**, in which genetic modification is random; or (2) **natural selection**, in which genetic change is directed and nonrandom. Natural selection is the differential contribution of genetic material from one generation to the next, differential in the sense that genetic components of the population or species are contributed in different amounts to the next generation; those genetic combinations resulting in *increased survival or reproduction* are contributed to a greater degree. (A quantitative measure of this differential contribution is known as **fitness**.) Natural selection results in an **adaptation**, a structure or feature that performs a particular function and which itself brings about increased survival or reproduction. In a consideration of the evolution of any feature in systematics, the possible adaptive significance of that feature should be explored.

Finally, an ultimate result of evolution is **speciation**, the formation of new species from preexisting species. Speciation can follow lineage divergence, the splitting of one lineage into two, separate lineages (Figure 1.6D). Lineage divergence is itself a means of increasing evolutionary diversity. If two, divergent lineages remain relatively distinct, they may change independently of one another, into what may be designated as separate species.

TAXONOMY

Taxonomy is a major part of systematics that includes four components: **Description**, **Identification**, **Nomenclature**, and **Classification**. (Remember the mnemonic device: **DINC**.) The general subjects of study are **taxa** (singular, **taxon**), which are defined or delimited groups of organisms. Ideally, taxa should have a property known as **monophyly** (discussed

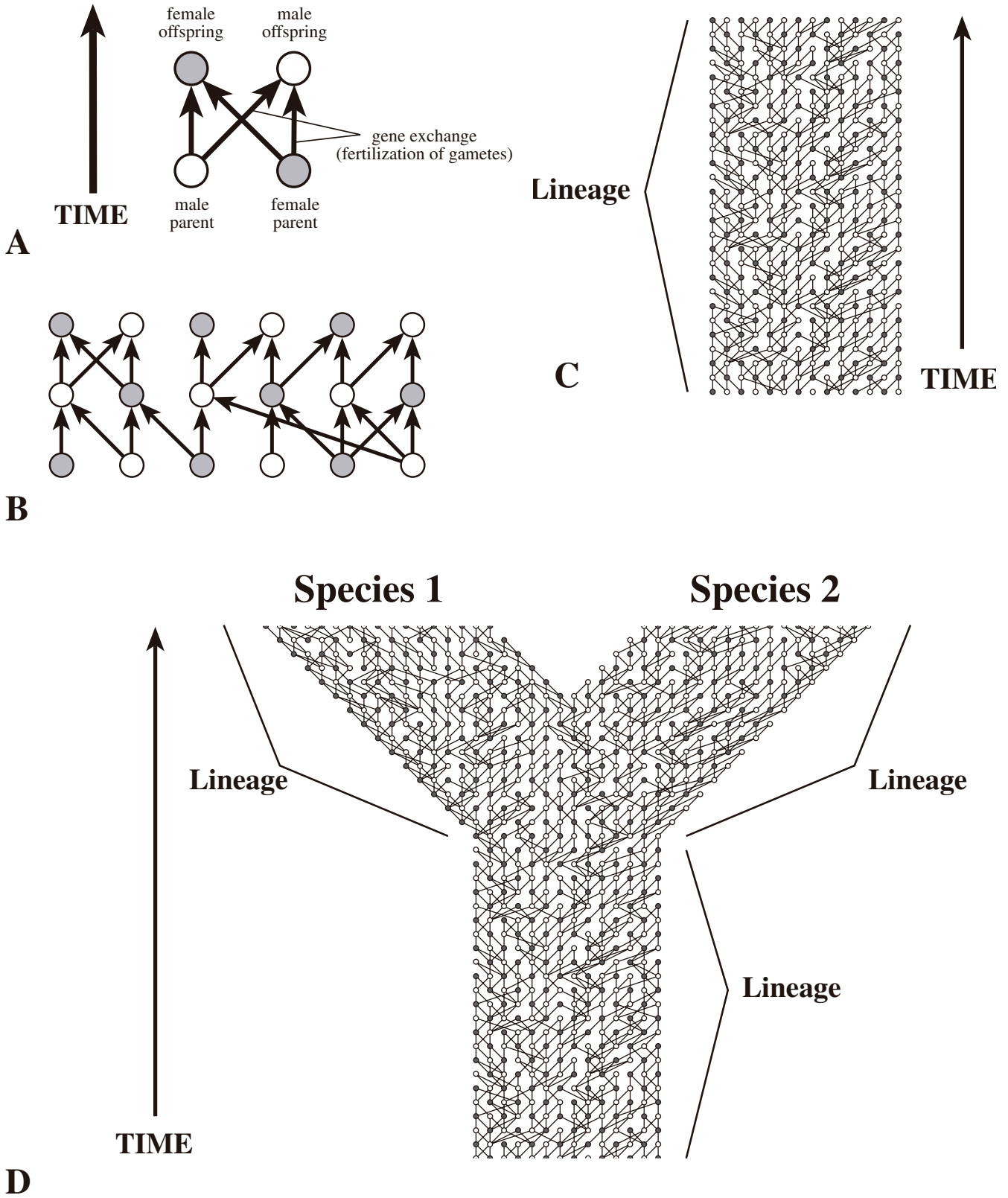


FIGURE 1.6 **A.** Diagram of descent in sexually reproducing species, in which two parents mate to form new offspring. **B.** Gene flow between individuals of a population. **C.** A lineage, the result of gene flow over time. **D.** Divergence of one lineage into two, which may result in speciation (illustrated here).

later; Chapter 2) and are traditionally treated at a particular rank (see later discussion). It should be pointed out that the four components of taxonomy are not limited to formal systematic studies but are the foundation of virtually all intellectual endeavors of all fields, in which conceptual entities are described, identified, named, and classified. In fact, the ability to describe, identify, name, and classify things undoubtedly has evolved by natural selection in humans and, in part, in other animals as well.

Description is the assignment of features or attributes to a taxon. The features are called **characters**. Two or more forms of a character are **character states**. One example of a character is petal color, for which two character states are yellow and blue. Another character is leaf shape, for which possible character states are elliptic, lanceolate, and ovate. Numerous character and character state terms are used in plant systematics, both for general plant morphology (see Chapter 9) and for specialized types of data (Chapters 10–14). The purpose of these descriptive character and character state terms is to use them as tools of communication, for concisely categorizing and delimiting the attributes of a taxon, an organism, or some part of the organism. An accurate and complete listing of these features is one of the major objectives and contributions of taxonomy.

Identification is the process of associating an unknown taxon with a known one, or recognizing that the unknown is new to science and warrants formal description and naming. One generally identifies an unknown by first noting its characteristics, that is, by describing it. Then, these features are compared with those of other taxa to see if they conform. Plant taxa can be identified in many ways (see Chapter 15). A taxonomic key is perhaps the most utilized of identification devices. Of the different types of taxonomic keys, the most common, used in virtually all florae, is a dichotomous key. A **dichotomous key** consists of a series of two contrasting statements. Each statement is a **lead**; the pair of leads

constitutes a **couplet** (Figure 1.7). That lead which best fits the specimen to be identified is selected; then all couplets hierarchically beneath that lead (by indentation and/or numbering) are sequentially checked for fit until an identification is reached (Figure 1.7).

Nomenclature is the formal naming of taxa according to some standardized system. For plants, algae, and fungi, the rules and regulations for the naming of taxa are provided by the International Code of Botanical Nomenclature (see Chapter 16). These formal names are known as **scientific names**, which by convention are translated into the Latin language. The fundamental principle of nomenclature is that all taxa may bear *only one scientific name*. Although they may seem difficult to learn at first, scientific names are much preferable to common (vernacular) names (Chapter 16).

The scientific name of a species traditionally consists of two parts (which are underlined or italicized): the genus name, which is always capitalized, e.g., *Quercus*, plus the specific epithet, which by recent consensus is not capitalized, e.g., *agrifolia*. Thus, the species name for what is commonly called California live oak is *Quercus agrifolia*. Species names are known as **binomials** (literally meaning two names) and this type of nomenclature is called binomial nomenclature, first formalized in the mid-18th century by Carolus Linnaeus.

Classification is the arrangement of entities (in this case, taxa) into some type of order. The purpose of classification is to provide a system for cataloguing and expressing relationships between these entities. Taxonomists have traditionally agreed upon a method for classifying organisms that utilizes categories called **ranks**. These taxonomic ranks are hierarchical, meaning that each rank is inclusive of all other ranks beneath it (Figure 1.8).

As defined earlier, a **taxon** is a group of organisms typically treated at a given rank. Thus, in the example of Figure 1.8, Magnoliophyta is a taxon placed at the rank of phylum; Liliopsida is a taxon placed at the rank of class; Arecaceae is a taxon

- | | | |
|-----------------|---|-------------------|
| | 1. Annual; leaves <<1 cm long; flowers 1–4 mm | |
| <i>Couplet:</i> | <i>Lead:</i> 2. Leaves opposite, pairs fused around stem; flowers axillary; petals <2 mm | <i>Crassula</i> |
| | <i>Lead:</i> 2. Leaves alternate above, free; flowers in terminal cyme; petals 1.5–4.5 mm | <i>Parvisedum</i> |
| | 1. Generally perennial herbs to shrubs; leaves >1 cm; flowers generally >10 mm (if annual, flowers >4 mm) | |
| | 3. Shrub or subshrub | |
| | 4. Leaves alternate, many in rosette, ciliate; sepals 6–16; petals ± free | <i>Aeonium</i> |
| | 4. Leaves opposite, few, not ciliate; sepals 5; petals fused, tube > sepals | <i>Cotyledon</i> |
| | 3. Perennial herb (annual or biennial in <i>Sedum radiatum</i>) | |
| | 5. Inflorescence axillary; cauline leaves different from rosette leaves | <i>Dudleya</i> |
| | 5. Inflorescence terminal; cauline leaves like rosettes, or basal leaves brown, scale-like | <i>Sedum</i> |

FIGURE 1.7 Dichotomous key to the genera of the Crassulaceae of California, by Reid Moran, ' *The Jepson Manual* (1993, Hickman, ed., University of California Press, Berkeley), reprinted by special permission.

Major Taxonomic Ranks	Taxa
Kingdom	Plantae
Phylum (Division also acceptable)	Magnoliophyta
Class	Liliopsida (Monocots)
Order	Arecales
Family	Areaceae
Genus (plural: genera)	<i>Cocos</i>
Species (plural: species)	<i>Cocos nucifera</i>

FIGURE 1.8 The primary taxonomic ranks accepted by the International Code of Botanical Nomenclature.

placed at the rank of family; etc. Note that taxa of a particular rank generally end in a particular suffix (Chapter 16). There is a trend among systematic biologists to eliminate the rank system of classification (see Chapter 16). In this book, ranks are used for naming groups but not emphasized as ranks.

There are two major means of arriving at a classification of life: phenetic and phylogenetic. **Phenetic** classification is that based on overall similarities. Most of our everyday classifications are phenetic. For efficiency of organization (e.g., storing and retrieving objects, like nuts and bolts in a hardware store) we group similar objects together and dissimilar objects apart. Many traditional classifications in plant systematics are phenetic, based on noted similarities between and among taxa. **Phylogenetic** classification is that which is based on evolutionary history, or pattern of descent, which may or may not correspond to overall similarity (see later discussion, Chapter 2).

PHYLOGENY

Phylogeny, the primary goal of systematics, refers to the evolutionary history of a group of organisms. Phylogeny is commonly represented in the form of a **cladogram** (or phylogenetic tree), a branching diagram that conceptually represents the evolutionary pattern of descent (see Figure 1.9). The lines of a cladogram represent **lineages** or **clades**, which (as discussed earlier) denote descent, the sequence of ancestral-descendant populations through time (Figure 1.9A). Thus, cladograms have an implied (relative) time scale. Any branching of the cladogram represents lineage **divergence**, the diversification of lineages from one **common ancestor**.

Changes in the genetic makeup of populations, i.e., evolution, may occur in lineages over time. Evolution may be recognized as a change from a preexisting, or **ancestral**, character state to a new, **derived** character state. The derived character state is an evolutionary novelty, also called an **apomorphy** (Figure 1.9A). **Phylogenetic systematics**, or **cladistics**, is a methodology for inferring the pattern of evolutionary

history of a group of organisms, utilizing these apomorphies (Chapter 2).

As cited earlier, cladograms serve as the basis for phylogenetic classification. A key component in this classification system is the recognition of what are termed monophyletic groups of taxa. A **monophyletic group** is one consisting of a common ancestor plus all (and only all) descendants of that common ancestor. For example, the monophyletic groups of the cladogram in Figure 1.9B are circled. A phylogenetic classification recognizes only monophyletic groups. Note that some monophyletic groups are included within others (e.g., in Figure 1.9B the group containing only taxa *E* and *F* is included within the group containing only taxa *D*, *E*, and *F*, which is included within the group containing only taxa *B*, *C*, *D*, *E*, and *F*, etc.). The sequential listing of monophyletic groups can serve as a phylogenetic classification scheme (see Chapter 2).

In contrast to a monophyletic group, a **paraphyletic group** is one consisting of a common ancestor but *not all* descendants of that common ancestor; a **polyphyletic group** is one in which there are two or more separate groups, each with a separate common ancestor. Paraphyletic and polyphyletic groups distort the accurate portrayal of evolutionary history and should be abandoned (see Chapter 2).

Knowing the phylogeny of a group, in the form of a cladogram, can be viewed as an important end in itself. As discussed earlier, the cladogram may be used to devise a system of classification, one of the primary goals of taxonomy. The cladogram also can be used as a tool for addressing several interesting biological questions, including biogeographic or ecological history, processes of speciation, and adaptive character evolution. A thorough discussion of the principles and methodology of phylogenetic systematics is discussed in Chapter 2.

WHY STUDY SYSTEMATICS?

The rationale and motives for engaging the field of systematics are worth examining. For one, systematics is important in

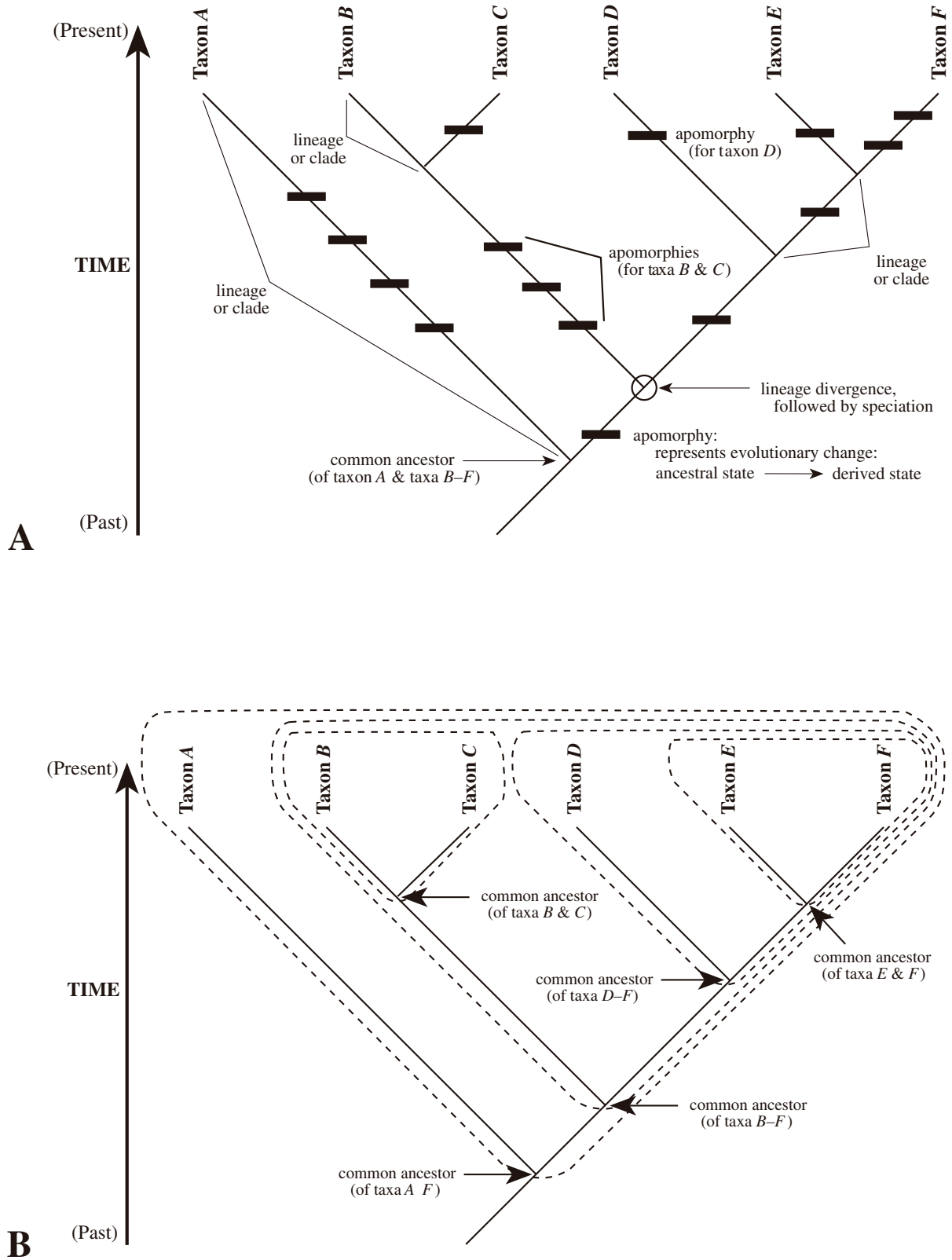


FIGURE 1.9 Example of a cladogram or phylogenetic tree for taxa A-F. **A.** Cladogram showing lineages and apomorphies, the latter indicated by thick hash marks. **B.** Cladogram with common ancestors shown and monophyletic groups circled.

providing a foundation of information about the tremendous diversity of life. Virtually all fields of biology are dependent on the correct taxonomic determination of a given study organism, which relies on formal description, identification, naming, and classification. Systematic research is the basis for acquiring, cataloguing, and retrieving information about life's diversity. Essential to this research is documentation, through collection (Chapter 17) and storage of reference specimens, e.g., for plants in an accredited herbarium (Chapter 18). Computerized data entry of this collection information is now vital to cataloguing and retrieving the vast amount of information dealing with biodiversity (Chapter 18).

Systematics is also an integrative and unifying science. One of the fun aspects of systematics is that it may utilize data from all fields of biology: morphology, anatomy, embryology/development, ultrastructure, paleontology, ecology, geography, chemistry, physiology, genetics, karyology, and cell/molecular biology. The systematist has an opportunity to understand all aspects of his/her group of interest in an overall synthesis of what is known from all biological specialties, with the goal being to understand the evolutionary history and relationships of the group.

Knowing the phylogeny of life can give insight into other fields and have significant practical value. For example, when a species of *Dioscorea*, wild yam, was discovered to possess steroid compounds (used first in birth control pills), examination of other closely related species revealed species that contained even greater quantities of these compounds. Other examples corroborate the practical importance of knowing phylogenetic relationships among plant species. The methodology of phylogenetics is now an important part of comparative biology, used by, for example, evolutionary

ecologists, functional biologists, and parasitologists, all of whom need to take history into account in formulating and testing hypotheses.

The study of systematics provides the scientific basis for defining or delimiting species and infraspecific taxa (subspecies or varieties) and for establishing that these are distinct from other, closely related and similar taxa. Such studies are especially important today in conservation biology. In order to determine whether a species or infraspecific taxon of plant is rare or endangered and warrants protection, one must first know the limits of that species or infraspecific taxon. In addition, understanding the history of evolution and geography may aid in conservation and management decisions, where priorities must be set as to which regions to preserve.

Finally, perhaps the primary motivation for many, if not most, in the field of systematics has been the joy of exploring the intricate complexity and incredible diversity of life. This sense of wonder and amazement about the natural world is worth cultivating (or occasionally rekindling). Systematics also can be a challenging intellectual activity, generally requiring acute and patient skills of observation. Reconstruction of phylogenetic relationships and ascertaining the significance of those relationships can be especially challenging and rewarding. But today we also face a moral issue: the tragic and irrevocable loss of species, particularly accelerated by rampant destruction of habitat, such as deforestation in the tropics. We can all try to help, both on a personal and professional level. Systematics, which has been called simply the study of biodiversity, is *the* major tool for documenting that biodiversity and can be a major tool for helping to save it. Perhaps we can all consider reassessing our own personal priorities in order to help conserve the life that we study.

REVIEW QUESTIONS

PLANTS

1. What is a plant? In what two conceptual ways can the answer to this question be approached?
2. What are the three major groups of life currently accepted?
3. Name and define the mechanism for the evolution of chloroplasts.
4. Name some chlorophyllous organismal groups that have traditionally been called plants but that evolved chloroplasts independently.
5. Draw a simplified cladogram showing the relative relationships among the green plants (Chlorobionta), land plants (embryophytes), vascular plants (tracheophytes), seed plants (spermatophytes), gymnosperms, and angiosperms (flowering plants).
6. Why are land plants treated as equivalent to plants in this book?
7. List the many ways that plants are important, both in the past evolution of life on earth and in terms of direct benefits to humans.

SYSTEMATICS

8. What is systematics and what is its primary emphasis?
9. Define biological evolution, describing what is meant both by descent and by modification.
10. What is a lineage (clade)?
11. Name and define the units that undergo evolutionary change.
12. What are the two major mechanisms for evolutionary change?
13. What is a functional feature that results in increased survival or reproduction called?
14. Name and define the four components of taxonomy.
15. Define character and character state.
16. Give one example of a character and character state from morphology or from some type of specialized data.
17. What is a dichotomous key? a couplet? a lead?
18. What is a scientific name?
19. Define binomial and indicate what each part of the binomial is called.
20. What is the difference between rank and taxon?
21. What is the plural of taxon?
22. Name the two main ways to classify organisms and describe how they differ.
23. Define phylogeny and give the name of the branching diagram that represents phylogeny.
24. What does a split, from one lineage to two, represent?
25. Name the term for both a preexisting feature and a new feature.
26. What is phylogenetic systematics (cladistics)?
27. What is a monophyletic group? a paraphyletic group? a polyphyletic group?
28. For what can phylogenetic methods be used?
29. How is systematics the foundation of the biological sciences?
30. How can systematics be viewed as unifying the biological sciences?
31. How is systematics of value in conservation biology?
32. Of what benefit is plant systematics to you?

EXERCISES

1. Obtain definitions of the word *plant* by asking various people (lay persons or biologists) or looking in reference sources, such as dictionaries or textbooks. Tabulate the various definitions into classes. What are the advantages and disadvantages of each?
2. Take a day to note and list the uses and importance of plants in your everyday life.
3. Pick a subject, such as history or astronomy, and cite how the principles of taxonomy are used in its study.
4. Do a Web search for a particular plant species (try common and scientific name) and note what aspect of plant biology each site covers.
5. Peruse relevant articles in a systematics journal and tabulate the different types of research questions that are addressed.

REFERENCES FOR FURTHER STUDY

- Kumar, S., and A. Rzhetsky. 1996. Evolutionary relationships of eukaryotic kingdoms. *Journal of Molecular Evolution* 42: 183-193.
- Reaka-Kudla, M. L., D. E. Wilson, and E. O. Wilson (eds.). 1997. *Biodiversity II: Understanding and Protecting Our Biological Resources*. Joseph Henry Press, Washington, DC.
- Simpson, B. B., and M. C. Ogorzaly. 2001. *Economic Botany: Plants in Our World*. McGraw-Hill, New York.
- Sogin, M. L. 1994. The origin of eukaryotes and evolution into major kingdoms. Bengtson, S. (ed.). *Nobel Symposium, No. 84. Early life on earth*; 84th Nobel Symposium, Karlskoga, Sweden, May 16, 1992. Columbia University Press, New York, pp. 181-192.
- Systematics Agenda 2000: Charting the Biosphere. 1994. Produced by Systematics Agenda 2000. [This is an excellent introduction to the goals and rationale of systematic studies, described as "a global initiative to discover, describe and classify the world's species. Available through SA2000, Herbarium, New York Botanical Garden, Bronx, New York 10458, USA]
- Wilson, E. O. (ed.), and F. M. Peter (assoc. ed.). 1988. *Biodiversity*. National Academy Press, Washington, DC.
- Yoon, H. S., J. D. Hackett, G. Pinto, and D. Bhattacharya. 2002. The single ancient origin of chromist plastids. *Proceedings of the National Academy of Sciences of the United States of America* 99: 15507-15512.

2

PHYLOGENETIC SYSTEMATICS

OVERVIEW AND GOALS	17	Reticulation	31
TAXON SELECTION	18	Taxon Selection and Polymorphic Characters	31
CHARACTER ANALYSIS	19	Polarity Determination: Outgroup Comparison	31
Description	19	Ancestral versus Derived Characters	34
Character Selection and Definition	19	Consensus Trees	35
Character State Discreteness	20	Long Branch Attraction	35
Character Correlation	21	Maximum Likelihood	36
Homology Assessment	21	Bayesian Analysis	36
Character State Transformation Series	22	Measures of Homoplasy	36
Character Weighting	22	Cladogram Robustness	37
Polarity	23	CLADOGRAM ANALYSIS	38
Character Step Matrix	24	Phylogenetic Classification	38
Character × Taxon Matrix	24	Character Evolution	40
CLADOGRAM CONSTRUCTION	24	Biogeography and Ecology	41
Apomorphy	24	Ontogeny and Heterochrony	41
Recency of Common Ancestry	26	A Perspective on Phylogenetic Systematics	43
Monophyly	26	REVIEW QUESTIONS	45
Parsimony Analysis	26	EXERCISES	47
Unrooted Trees	29	REFERENCES FOR FURTHER STUDY	48
Character Optimization	30	CLADISTIC COMPUTER PROGRAMS	48
Polytomy	30		

OVERVIEW AND GOALS

As introduced in the previous chapter, **phylogeny** refers to the evolutionary history or pattern of descent of a group of organisms and is one of the primary goals of systematics. **Phylogenetic systematics**, or **cladistics**, is that branch of systematics concerned with inferring phylogeny. Ever since Darwin laid down the fundamental principles of evolutionary theory, one of the major goals of the biological sciences has been the determination of life's history of descent. This phylogeny of organisms, visualized as a branching pattern, can be determined by an analysis of characters from living or fossil organisms, utilizing phylogenetic principles and methodology.

As reviewed in Chapter 1, a phylogeny is commonly represented in the form of a **cladogram**, or **phylogenetic tree**, a branching diagram that conceptually represents the best estimate of phylogeny (Figure 2.1). The lines of a cladogram are known as **lineages** or **clades**. Lineages represent the sequence of ancestral-descendant populations through time, ultimately denoting descent.

Thus, as previously reviewed, cladograms have an implied, but relative, time scale. Any branching of the cladogram represents lineage **divergence** or **diversification**, the formation of two separate lineages from one **common ancestor**. (The two lineages could diverge into what would be designated separate species, the process of forming two species from one termed **speciation**.) The *point* of divergence of one clade into

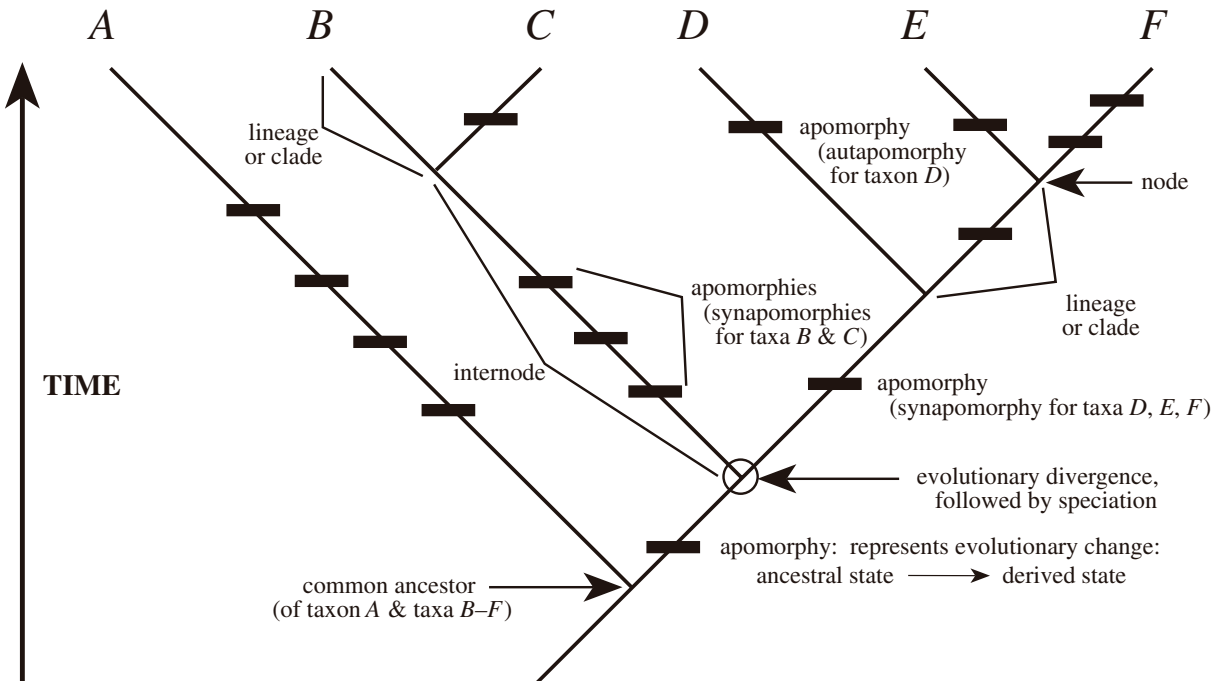


FIGURE 2.1 Example of a cladogram or phylogenetic tree for taxa A–F, with apomorphies indicated by thick hash marks; redrawn from Chapter 1. See text for explanation of terms.

two (where the most common ancestor of the two divergent clades is located) is termed a **node**; the region between two nodes is called an **internode** (Figure 2.1).

Evolution may occur within lineages over time and is recognized as a change from a preexisting **ancestral** (also called **plesiomorphic** or **primitive**) condition to a new, **derived** (also called **apomorphic** or **advanced**) condition. The derived condition, or **apomorphy**, represents an evolutionary novelty. As seen in Figure 2.1, an apomorphy that unites two or more lineages is known as a **synapomorphy** (*syn*, together); one that occurs within a single lineage is called an **autapomorphy** (*aut*, self). However, either may be referred to simply as an **apomorphy**, a convention used throughout this book. Cladograms may be represented in different ways. Figure 2.2 shows the same cladogram as in Figure 2.1, but shifted 90° clockwise and with the lineages drawn perpendicular to one another and of a length reflective of the number of apomorphic changes.

Why study phylogeny? Knowing the pattern of descent, in the form of a cladogram, can be viewed as an important end in itself. The branching pattern derived from a phylogenetic analysis may be used to infer the collective evolutionary changes that have occurred in ancestral/descendant populations through time. Thus, a knowledge of phylogenetic relationships may be invaluable in understanding structural

evolution as well as in gaining insight into the possible functional, adaptive significance of hypothesized evolutionary changes. The cladogram can also be used to classify life in a way that directly reflects evolutionary history. Cladistic analysis may also serve as a tool for inferring biogeographic and ecological history, assessing evolutionary processes, and making decisions in the conservation of threatened or endangered species.

The principles, methodology, and applications of phylogenetic analyses are described in the remainder of this chapter.

TAXON SELECTION

The study of phylogeny begins with the selection of **taxa** (taxonomic groups) to be analyzed, which may include living and/or fossil organisms. Taxon selection includes both the group as a whole, called the study group or **ingroup**, and the individual unit taxa, termed **Operational Taxonomic Units**, or **OTUs**. The rationale as to *which* taxa are selected from among many rests by necessity on previous classifications or phylogenetic hypotheses. The ingroup is often a traditionally defined taxon for which there are competing or uncertain classification schemes, the objective being to test the bases of those different classification systems or to provide a new

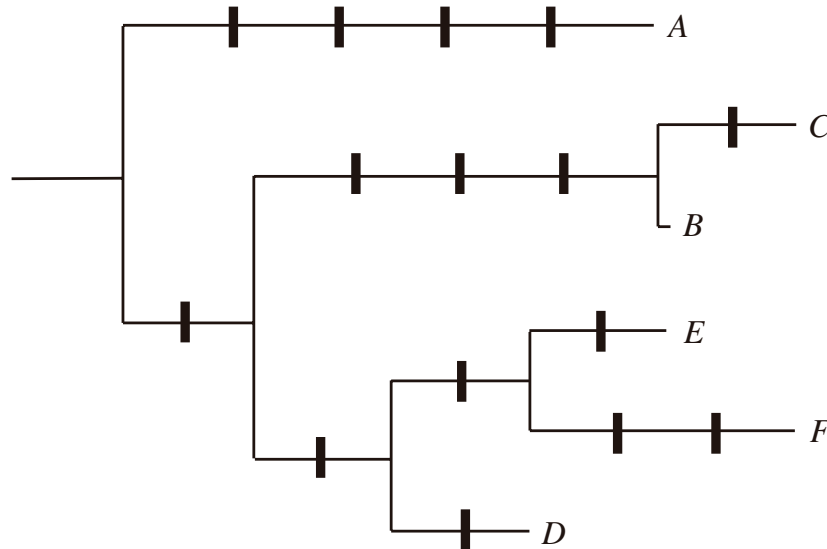


FIGURE 2.2 Cladogram of Fig. 2.1 drawn horizontally, with branch lengths scaled to number of apomorphic changes.

classification system derived from the phylogenetic analysis. The OTUs are previously classified members of the study group and may be species or taxa consisting of groups of species (e.g., traditional genera, families). Sometimes named subspecies or even populations, if distinctive and presumed to be on their own evolutionary track, can be used as OTUs in a cladistic analysis.

In addition, one or more outgroups OTUs are selected. An **outgroup** is a taxon that is closely related to but not a member of the ingroup (see **Polarity Determination: Outgroup Comparison**). Outgroups are used to root a tree (see later discussion).

Some caution should be taken in choosing which taxa to study. First, the OTUs must be well circumscribed and delimited from one another. Second, the study group itself should be large enough so that all probable closely related OTUs are included in the analysis. Stated strictly, both OTUs and the group as a whole must be assessed for **monophyly** before the analysis is begun (see below.) In summary, the initial selection of taxa in a cladistic analysis, both study group and OTUs, should be questioned beforehand to avoid the bias of blindly following past classification systems.

CHARACTER ANALYSIS

DESCRIPTION

Fundamental in any systematic study is description, the characterization of the attributes or features of taxa using any

number of types of evidence (see Chapters 9–14). A systematist may make original descriptions of a group of taxa or rely partly or entirely on previously published research data. In any case, it cannot be overemphasized that the ultimate validity of a phylogenetic study depends on the descriptive accuracy and completeness of the primary investigator. Thorough research and a comprehensive familiarity with the literature on the taxa and characters of concern are prerequisites to a phylogenetic study.

CHARACTER SELECTION AND DEFINITION

After taxa are selected and the basic research and literature survey are completed, the next step in a phylogenetic study is the actual selection and definition of **characters** and **character states** from the descriptive data. (Recall that a character is an attribute or feature; character states are two or more forms of a character.) Generally, those features that (1) are genetically determined and heritable (termed *intrinsic*), (2) are relatively invariable within an OTU, and (3) denote clear discontinuities from other similar characters and character states should be utilized. However, the selection of a finite number of characters from the virtually infinite number that could be used adds an element of subjectivity to the study. Thus, it is important to realize that any analysis is inherently biased simply by *which* characters are selected and *how* the characters and character states are defined. (In some cases, certain characters may be weighted over others; see later discussion.)

Because morphological features are generally the manifestation of numerous intercoordinated genes, and because

evolution occurs by a change in one or more of those genes, the precise definition of a feature in terms of characters and character states may be problematic. A structure may be defined broadly as a whole entity with several components. Alternatively, discrete features of a structure may be defined individually as separate characters and character states. For example, in comparing the evolution of fruit morphology within some study group, the character fruit type might be designated as two character states: berry versus capsule, *or* the characteristics of the fruit may be subdivided into a host of characters with their corresponding states, for example, fruit shape, fruit wall texture, fruit dehiscence, and seed number. (These characters may be correlated, however; see later discussion.) In practice, characters are divided only enough to communicate differences between two or more taxa. However, this type of terminological atomization may be misleading with reference to the effect of specific genetic changes in evolution, as genes do not normally correspond one for one with taxonomic characters. The morphology of a structure is the end product of development, involving a host of complex interactions of the entire genotype.

CHARACTER STATE DISCRETENESS

Because phylogenetic systematics entails the recognition of an evolutionary transformation from one state to another, an important requirement of character analysis is that character states be discrete or discontinuous from one another. Molecular characters and their states are usually discrete (see Chapter 14). For some nonmolecular, qualitative characters such as corolla color, the discontinuity of states is clear; e.g., the corolla is yellow in some taxa and red in others. But for other features, character states may not actually be clearly distinguishable from one another. This lack of discontinuity often limits the number of available characters and is often the result of variation of a feature either within a taxon or between taxa. Because character states must be clearly discrete from one another in order to be used in a cladistic analysis, they must be evaluated for discontinuity. A standard way to evaluate state discontinuity is to do a statistical analysis, e.g., by comparing the means, ranges, and standard deviations of each character for all taxa in the analysis (including outgroup taxa; see later discussion). Such a plot may reveal two or more classes of features that may be defined as discrete character

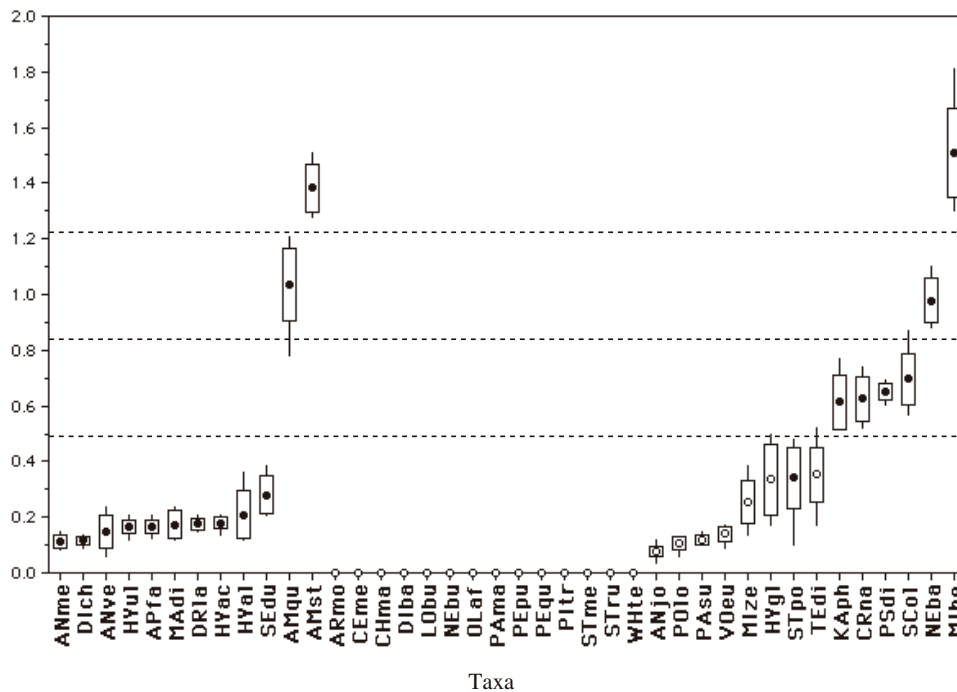


FIGURE 2.3 Example of a pollen character (exine wall foot-layer thickness) for which the character states are quantitatively analyzed for each taxon. The dashed horizontal lines represent breaks or discontinuities between states. Solid dots are means, vertical lines are ranges, and boxes are ± 1 standard deviation from the mean (used here as the measure of discreteness). Outgroup taxa are to the left, ingroup taxa to the right. (From Levin, G. A., and M. G. Simpson. 1994. *Annals of the Missouri Botanical Garden* 81: 203-238.)

states (Figure 2.3). The investigator must decide what constitutes discreteness, such as lack of overlap of ranges or lack of overlap of ± 1 standard deviation. Additional statistical tests, such as ANOVAS, *t*-tests, or multivariate statistics, may be used as other criteria for evaluating character state discontinuity.

CHARACTER CORRELATION

Another point to consider in character selection and definition is whether there is possible correlation of characters. Character correlation is an interaction between what are defined as separate characters, but which are actually components of a common structure, the manifestation of a single evolutionary novelty. Two or more characters are correlated if a change in one always accompanies a corresponding change in the other. When characters defined in a cladistic analysis are correlated, including them in the analysis (as two or more separate characters) may inadvertently weight what could otherwise be listed as a single character. In the example above, in which the original single character fruit type is subdivided into many characters (fruit shape, fruit wall texture, fruit dehiscence, and seed number), it is likely that these separate characters are correlated with an evolutionary shift from one fruit type (e.g., capsule) to another (e.g., berry). This is tested simply by determining if there is any variation in the character states of the subdivided characters between taxa. If characters appear to be correlated, they should either be combined into one character or *scaled*, such that each component character gets a reduced weight in a phylogenetic analysis (see **Character Weighting**).

HOMOLOGY ASSESSMENT

One concept critical to cladistics is that of **homology**, which can be defined as similarity resulting from common ancestry. Characters or character states of two or more taxa are homologous if those same features were present in the common ancestor of the taxa. For example, the flower of a daisy and the flower of an orchid are homologous as flowers because their common ancestor had flowers, which the two taxa share by continuity of descent. Taxa with homologous features are presumed to share, by common ancestry, the same or similar DNA sequences or gene assemblages that may, e.g., determine the development of a common structure such as a flower. (Unfortunately, molecular biologists often use the term *homology* to denote similarity in DNA sequence, even though the common ancestry of these sequences may not have been tested; using the term *sequence similarity* in this case is preferred.)

Homology may also be defined with reference to similar structures within the same individual; two or more structures are homologous if the DNA sequences that determine their

similarity share a common evolutionary history. For example, carpels of flowering plants are considered to be homologous with leaves because of a basic similarity between the two in form, anatomy, and development. Their similarity is thought to be the result of a sharing of common genes or of gene complexes of common origin that direct their development. The duplication and subsequent divergence of genes is a type of intraindividual or intraspecies homology; the genes are similar because of origin from a common ancestor, in this case the gene prior to duplication.

Similarity between taxa can arise not only by common ancestry, but also by independent evolutionary origin. Similarity that is *not* the result of homology is termed **homoplasy** (also sometimes termed *analogy*). Homoplasy may arise in two ways: **convergence** (equivalent to parallelism, here) or **reversal**. **Convergence** is the independent evolution of a similar feature in two or more lineages. Thus, liverwort gametophytic leaves and lycopod sporophytic leaves evolved independently as photosynthetic appendages; their similarity is homoplasious by convergent evolution. (However, although leaves in the two groups evolved independently, they could possibly be homologous in the sense of utilizing gene complexes of common origin that function in the development of bifacial organs. This is unknown at present.)

Reversal is the loss of a derived feature with the re-establishment of an ancestral feature. For example, the reduced flowers of many angiosperm taxa, such as *Lemna*, lack a perianth; comparative and phylogenetic studies have shown that flowers of these taxa lack the perianth by secondary loss, i.e., via a reversal, reverting to a condition prior to the evolution of a reproductive shoot having a perianth-like structure.

The determination of homology is one of the most challenging aspects of a phylogenetic study and may involve a variety of criteria. Generally, homology is hypothesized based on some evidence of similarity, either direct similarity (e.g., of structure, position, or development) or similarity via a gradation series (e.g., intermediate forms between character states). Homology should be assessed for each character of all taxa in a study, particularly of those taxa having *similarly termed* character states. For example, both the cacti and stem-succulent euphorbs have spines (Figure 2.4). Thus, for the character spine presence/absence, the character state spines present may be assigned to both of these two taxa in a broad cladistic analysis. Whether intended or not, this designation of the same character state for two or more taxa presupposes that these features are homologous in those taxa and arose by common evolutionary origin. Thus, a careful distinction should be made between terminological similarity and similarity by homology. In the above example, more detailed study demonstrates that the spines of cacti and

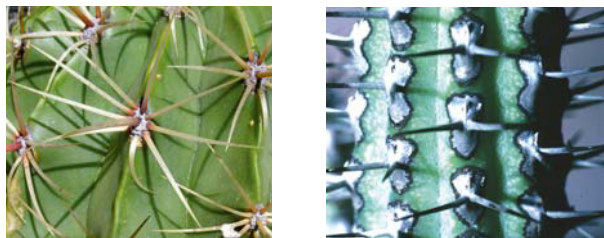


FIGURE 2.4 Comparison of spines in cacti (left) and stem-succulent euphorbs (right), which are not homologous as spines. See text for explanation.

euphorbs are quite different in origin, cacti having leaf spines arising from an areole (a type of short shoot), euphorbs having spines derived from modified stipules. Despite the similarity between spines of cacti and stem-succulent euphorbs, their structural and developmental *dissimilarity* indicates that they are homoplasious and had independent evolutionary origins (with similar selective pressures, i.e., protection from herbivores). This hypothesis necessitates a redefinition of the characters and character states, such that the two taxa are not coded the same.

Homology must be assessed for molecular data as well (Chapter 14). For DNA sequence data, alignment of the sequences is used to evaluate homology of individual base positions. In addition, gene duplication can confound comparison of homologous regions of DNA.

Hypotheses of homology are tested by means of the cladistic analysis. The *totality* of characters are used to infer the most likely evolutionary tree, and the original assessment of homology is checked by determining if convergences or reversals must be invoked to explain the distribution of character states on the final cladogram (see later discussion).

CHARACTER STATE TRANSFORMATION SERIES

After the characters and character states have been selected and defined and their homologies have been assessed, the character states for each character are arranged in a sequence, known as a **transformation series** or **morphocline**. Transformation series represent the hypothesized sequence of evolutionary change, from one character state to another, in terms of direction and probability. For a character with only two character states, known as a **binary character**, obviously only one transformation series exists. For example, for the character ovary position having the states inferior and superior, the implied transformation series is inferior \Leftrightarrow superior. This two-state transformation series represents (at least initially) a single, hypothesized evolutionary step, the direction of which is unspecified, being either inferior \Rightarrow superior or superior \Rightarrow inferior.

Characters having three or more character states, known as **multistate characters**, can be arranged in transformation series that are either ordered or unordered. An **unordered** transformation series allows for each character state to evolve into every other character state with equal probability, i.e., in a single evolutionary step. For example, an unordered transformation series for a three-state character is shown in Figure 2.5A; one for a four-state character is shown in Figure 2.5B and C. An **ordered** transformation series places the character states in a predetermined sequence that may be linear (Figure 2.5D) or branched (Figure 2.5E). Ordering a transformation series limits the direction of character state changes. For example, in Figure 2.5E, the evolution of 2 stamens from 5 stamens (or vice versa) takes two evolutionary steps and necessitates passing through the intermediate condition, 4 stamens; the comparable unordered series takes a single step between 2 stamens and 5 stamens (and between all other character states; Figure 2.5B).

The rationale for an ordered series is the assumption or hypothesis that evolutionary change proceeds gradually, such that going from one extreme to another most likely entails passing through some recognizable intermediate condition. Ordered transformation series are generally postulated *vis-à-vis* some obvious intergradation of character states or stages in the ontogeny of a character. A general suggestion in cladistic analyses is to code all characters as unordered unless there is compelling evidence for an ordered transformation, such as the presence of a vestigial feature in a derived structure. For example, a unifoliate leaf might logically be treated as being directly derived not from a simple leaf but from a compound leaf (in an ordered transformation series; see Figure 2.5D), evidence being the retention of a vestigial, ancestral petiolule (see **Polarity**).

CHARACTER WEIGHTING

As part of a phylogenetic analysis, the investigator may choose to weight characters. Character weighting is the assignment of greater or lesser taxonomic importance to certain characters over other characters in determining phylogenetic relationships. Assigning a character greater weight has the effect of listing it more than once in the character \times taxon matrix (see later section) in order to possibly override competing changes in unweighted characters. (Note that fractional weights can also be assigned using computer algorithms.)

In practice, character weighting is rarely done, in part because of the arbitrariness of determining the amount of weight a character should have. A frequent exception, however, is molecular data, for which empirical studies may justify the rationale for and degree of weighting.

Characters may be given greater weight in cases for which the designation of homology is considered relatively certain.

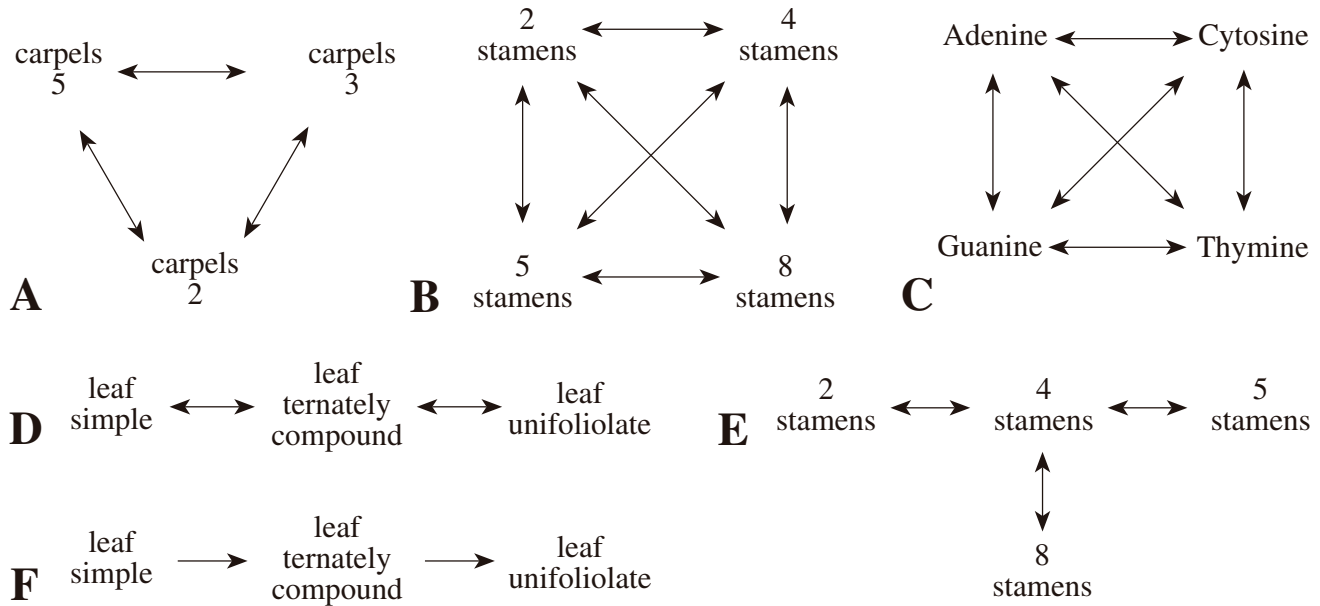


FIGURE 2.5 Examples of various character state transformations used in a cladistic analysis. **A.** Unordered, three-state character. **B,C.** Unordered, four-state character. **D.** Ordered, three-state character. **E.** Ordered, four-state character. **F.** Ordered and polarized, three-state character.

The expectation is that, by increasing the weight of characters for which homoplasy is deemed unlikely, taxa will be grouped by real, shared derived features. Such characters given greater weight may be hypothesized as having homologous states for various reasons. For example, a feature distinctive for two or more taxa may be structurally or developmentally complex, such that the independent evolution of the same character state would seem very unlikely. (It should be realized, however, that if a feature is most likely highly adaptive, convergence of similar complex features in two or more taxa may not necessarily be ruled out.)

Characters may be weighted unintentionally because they are correlated, i.e., the corresponding character state values of two or more characters are always present in all taxa and believed to be aspects of the same evolutionary novelty. In order to prevent excess weighting of correlated characters, they may be **scaled**, meaning that each character receives a weight that is the inverse of the number of characters (e.g., if there are three correlated characters, each receives a weight of 1/3).

Alternatively, weighting may be done *after* the first stage of a phylogenetic analysis. Those characters that exhibit reversals or parallelisms on the cladogram are recognized and given less weight over those that do not, sometimes as a direct function of the degree of homoplasy they exhibit. For example, if, after a cladistic analysis, a character exhibits two convergent changes, that character would be given a weight of

1/2 in a second cladistic analysis. This type of a posteriori analysis is called *successive weighting* (which relies on the assumption that the initial tree(s) are close to an accurate representation of phylogeny). Often, the rescaled consistency index (RC) value is used as a basis for successive weighting (see **Measures of Homoplasy**).

POLARITY

The final step of character analysis is the assignment of polarity. Polarity is the designation of relative ancestry to the character states of a morphocline. As summarized earlier, a change in character state represents a heritable evolutionary modification from a preexisting structure or feature (termed **plesiomorphic**, ancestral, or primitive) to a new structure or feature (**apomorphic**, derived, or advanced). For example, for the character ovary position, with character states superior and inferior, if a superior ovary is hypothesized as ancestral, the resultant polarized morphocline would be superior \Rightarrow inferior. For a multistate character (e.g., leaf type in Figure 2.5D), an example of a polarized, ordered transformation series is seen in Figure 2.5F. The designation of polarity is often one of the more difficult and uncertain aspects of a phylogenetic analysis, but also one of the most crucial. The primary procedure for determining polarity is outgroup comparison (see **Polarity Determination: Outgroup Comparison**).

		0	1	2	3
	0	0	1	2	3
	1	1	0	1	2
	2	2	1	0	1
A	3	3	2	1	0

		0	1	2	3
	0	0	1	1	1
	1	1	0	1	1
	2	1	1	0	1
B	3	1	1	1	0

		0	1	2	3
	0	0	1	2	3
	1	∞	0	1	2
	2	∞	∞	0	1
C	3	∞	∞	∞	0

		0	1	2	3
	0	0	1	5	5
	1	1	0	5	5
	2	5	5	0	1
D	3	5	5	1	0

FIGURE 2.6 Character step matrices for: **A.** Ordered character. **B.** Unordered character. **C.** Irreversible character. **D.** Differentially weighted character.

CHARACTER STEP MATRIX

As reviewed earlier assigning a character state transformation determines the number of steps that may occur when going from one character state to another. Computerized phylogeny reconstruction algorithms available today permit a more precise tabulation of the number of steps occurring between each pair of character states through a **character step matrix**. The matrix consists of a listing of character states in a top row and left column; intersecting numbers within the matrix indicate the number of steps required, going from states in the left column to states in the top row. For example, the character step matrix of Figure 2.6A illustrates an ordered character state transformation series, such that a single step is required when going from state 0 to state 1 (or state 1 to state 0), two steps are required when going from state 0 to state 2, etc. The character step matrix of Figure 2.6B shows an unordered transformation series, in which a single step is required when going from one state to any other (nonidentical) state. Character step matrices need not be symmetrical; that of Figure 2.6C illustrates an ordered transformation series but one that is irreversible, disallowing a change from a higher state number to a lower state number (e.g., from state 2 to state 1) by requiring a large number of step changes (symbolized by ∞). Character step matrices are most useful with specialized types of data. For example, the matrix of Figure 2.6D could represent DNA sequence data, where 0 and 1 are the states for the two purines (adenine and guanine) and 2 and 3 are the states for the two pyrimidines (cytosine and thymine; see Chapter 14). Note that in this matrix the change from one purine to another purine or one pyrimidine to another pyrimidine (each of these known as a *transition*) requires only one step, being biochemically more probable to occur, whereas a change from a purine to a pyrimidine or from a pyrimidine to a purine (termed a *transversion*) is given five steps, being more biochemically less likely. Thus, in a cladistic analysis, the latter change will be given substantially more weight.

CHARACTER X TAXON MATRIX

Prior to cladogram construction, characters and character states for each taxon are tabulated in a **character x taxon matrix**, as illustrated in Figure 2.7A. In order to analyze the

data using computer algorithms, the characters and character states must be assigned a numerical value. In doing so, character states are assigned nonnegative integer values, typically beginning with 0. Figure 2.7B shows the numerical coding of the matrix of Figure 2.7A. The states are numerically coded in sequence to correspond with the hypothesized transformation series for that character. For example, for the ordered transformation series *leaf type* of Figure 2.5D,F, the character states *simple*, *ternately compound*, and *unifoliolate* could be enumerated as 0, 1, and 2. In the character x taxon matrix, polarity is established by including one or more outgroup taxa as part of the character x taxon matrix (as in Figure 2.7A,B) and by subsequently rooting the tree by placing the outgroups at the extreme base of the final, most parsimonious cladogram (see later discussion). By convention, the ancestral character state (that possessed by the outgroup) is usually designated 0, even if intermediate in a morphocline (e.g., $1 \leftarrow 0 \Rightarrow 2$, in which state 0 is ancestral to both 1 and 2); however, *any* coded state may be designated as ancestral, including nonzero ones.

CLADOGRAM CONSTRUCTION

APOMORPHY

The primary tenet of phylogenetic systematics is that derived character states, or **apomorphies**, that are *shared* between two or more taxa (OTUs) constitute evidence that these taxa possess them because of common ancestry. These shared derived character states, or **synapomorphies**, represent the products of unique evolutionary events that may be used to link two or more taxa in a common evolutionary history. Thus, by sequentially linking taxa together based on their common possession of synapomorphies, the evolutionary history of the study group can be inferred.

The character x taxon matrix supplies the data for constructing a phylogenetic tree or cladogram. For example, Figure 2.7 illustrates construction of the cladogram for the five species of the hypothetical genus *Xid* from the character x taxon matrix at Figure 2.7A,B. First, the OTUs are grouped together as lineages arising from a single common ancestor

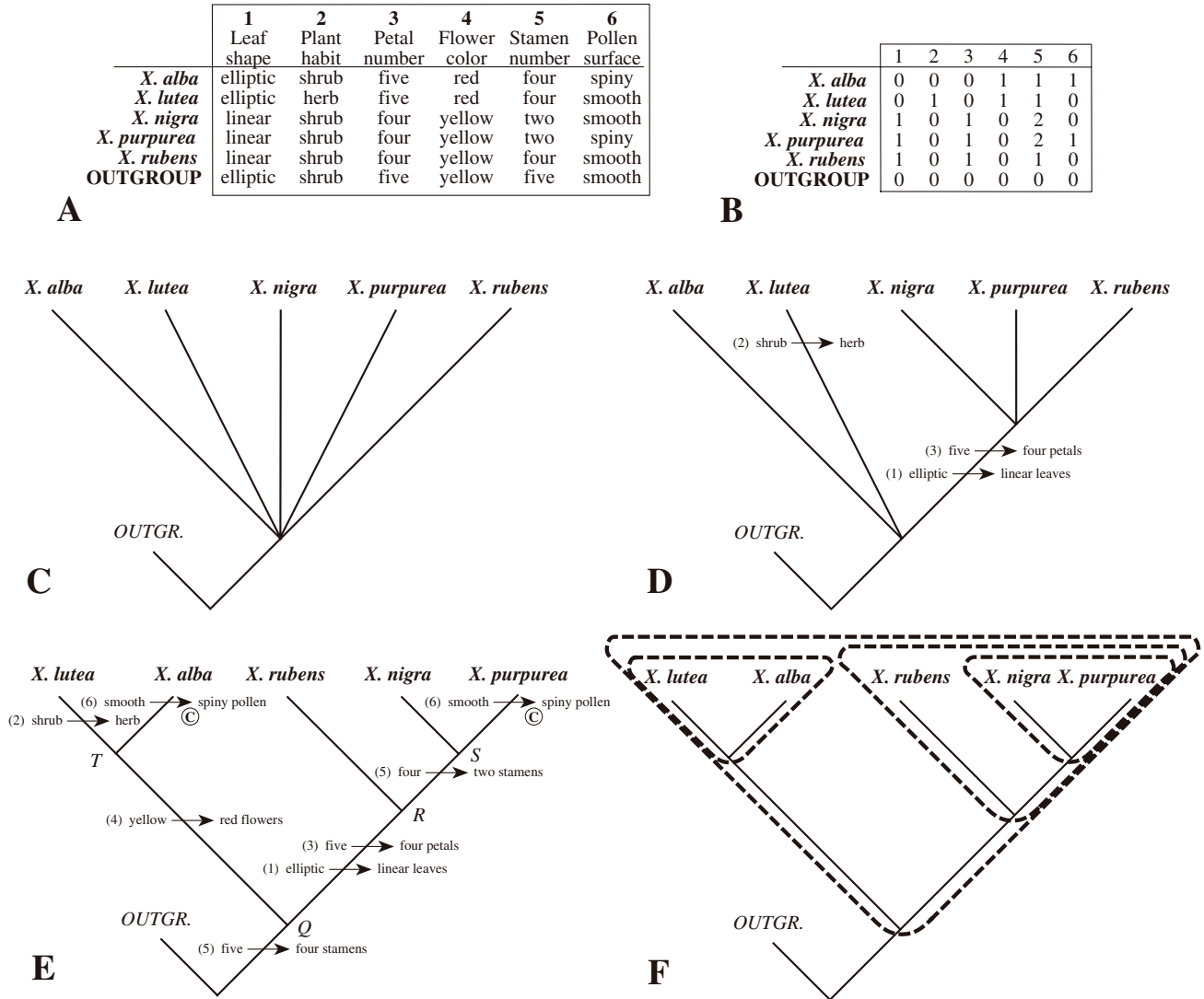


FIGURE 2.7 Character \times taxon matrix for five species of the hypothetical genus *Xid* plus an outgroup taxon (left column), showing six characters (top row) and their character states (inner columns). **A.** Character state names listed. **B.** Characters and character states converted to numerical values. **C.** Unresolved cladogram. **D.** Addition of characters 1-3. **E.** Most parsimonious cladogram, with addition of other characters. Note common ancestors *Q*, *R*, *S*, *T*, shown for illustrative purposes. **F.** Cladogram at **E**, with all monophyletic groups circled.

above the point of attachment of the outgroup (Figure 2.7C). This unresolved complex of lineages is known as a **polytomy** (see later discussion). Next, *derived* character states are identified and used to sequentially link sets of taxa (Figure 2.7D,E). In this example, synapomorphies include (1) the derived states of characters 1 and 3 that group together *X. nigra*, *X. purpurea*, and *X. rubens*; (2) the derived state of character 4 that groups together *X. alba* and *X. lutea*; (3) the derived state four stamens of character 5, which is found in all ingroup OTUs and constitutes a synapomorphy for the entire study group; and the derived state two stamens of character 5 that groups *X. nigra*

and *X. purpurea*. The derived state of character 2 is restricted to the taxon *X. lutea* and is therefore an **autapomorphy**. Autapomorphies occur within a single OTU and are not informative in cladogram construction. Finally, the derived state of character 6 evolved twice, in the lineages leading to both *X. alba* and *X. purpurea*; these independent evolutionary changes constitute homoplasies due to convergence.

One important principle is illustrated in Figure 2.7E for character 5, in which the derived state four stamens is an apomorphy for *all* species of the study group, including *X. nigra* and *X. purpurea*. Although the latter two species lack

the state four stamens for that character, they still *share the evolutionary event* in common with the other three species. The lineage terminating in *X. nigra* and *X. purpurea* has simply undergone additional evolutionary change in this character, transforming from four to two stamens (Figure 2.7E).

REGENCY OF COMMON ANCESTRY

Cladistic analysis allows for a precise definition of biological relationship. *Relationship* in phylogenetic systematics is a measure of **recency of common ancestry**. Two taxa are more closely related to one another if they share a common ancestor that is more recent in time than the common ancestor they share with other taxa. For example, in Figure 2.8A taxon *C* is more closely related to taxon *D* than it is to taxon *E* or *F*. This is true because the common ancestor of *C* and *D* is more recent in time (closer to the present) than is the common ancestor of *C*, *D*, *E*, and *F* (Figure 2.8A). In the earlier example of Figure 2.7E, it is evident that *X. nigra* and *X. purpurea* are more closely related to one another than either is to *X. rubens*. This is because the former two species together share a common ancestor (*S*) that is more recent in time than the common ancestor (*R*) that they share with *X. rubens*. Similarly *X. rubens* is more closely related to *X. nigra* and *X. purpurea* than it is to either *X. lutea* or *X. alba* because the former three taxa share a common ancestor (*R*) that is more recent in time than *Q*, the common ancestor shared by all five species.

Because descent is assessed by means of recency of common ancestry, the lineages of a given cladogram may be visually rotated around their junction point or node (at the common ancestor) with no change in phylogenetic relationships. For example, the cladogram portrayed in Figure 2.9A, *B*, and *C* are all the same as that in Figure 2.7E, differing only in that the lineages have been rotated about their common ancestors. The *topology* of all these cladograms is exactly the same; only the relative positioning of branches varies. (Again note that cladograms can be portrayed in different manners, with taxa at the top, bottom, or sides and with lineages drawn as vertical, horizontal, or angled lines; see Figure 2.7A C.)

MONOPHYLY

A very important concept in phylogenetic systematics is that of monophyly, or monophyletic groups. As introduced in Chapter one, a **monophyletic group** is one that consists of a common ancestor plus *all* descendants of that ancestor. The rationale for monophyly is based on the concept of recency of common ancestry. Members of a monophyletic group share one or more unique evolutionary events; otherwise, the group could not generally be identified as monophyletic. For example, four monophyletic groups can be delimited from the cladogram of

Figure 2.7E; these are circled in Figure 2.7F. In another example, the monophyletic groups of the cladogram of Figure 2.8A are shown in Figure 2.8B. Note that all monophyletic groups include the common ancestor plus all lineages derived from the common ancestor, with lineages terminating in an OTU.

Each of the two descendant lineages from one common ancestor is known as **sister groups** or **sister taxa**. For example, in Figure 2.7E and F, sister group pairs are: (1) *X. lutea* and *X. alba*; (2) *X. nigra* and *X. purpurea*; (3) *X. nigra* + *X. purpurea* and *X. rubens*; and (4) *X. lutea* + *X. alba* and *X. nigra* + *X. purpurea* + *X. rubens*.

The converse of monophyly is paraphyly. A **paraphyletic group** is one that includes a common ancestor and some, *but not all*, known descendants of that ancestor. For example, in Figure 2.7E, a group including ancestor *Q* and the lineages leading to *X. lutea*, *X. alba*, and *X. rubens* alone is paraphyletic because it has left out two taxa (*X. purpurea* and *X. nigra*), which are also descendants of common ancestor *Q*. Similarly, a **polyphyletic group** is one containing two or more common ancestors. For example, in Figure 2.7E, a group containing *X. lutea* and *X. purpurea* alone could be interpreted as polyphyletic as these two taxa do not have a single common ancestor that is part of the group. (*Paraphyletic* and *polyphyletic* may intergrade; the term *non-monophyletic* may be used to refer to either.)

Paraphyletic and polyphyletic groups are not natural evolutionary units and should be abandoned in formal classification systems. Their usage in comparative studies of character evolution, evolutionary processes, ecology, or biogeography will likely bias the results. In addition, paraphyletic groups cannot be used to reconstruct the evolutionary history of that group (see **Classification**). A good example of a paraphyletic group is the traditionally defined Dicots. Because most recent analyses show that some members of the Dicots are more closely related to Monocots than they are to other Dicots, the term *Dicot* should not be used in formal taxonomic nomenclature. (See Chapter 7.)

PARSIMONY ANALYSIS

In constructing a cladogram, a single branching pattern is selected from among many possibilities. The number of possible dichotomously branching cladograms increases dramatically with a corresponding increase in the number of taxa. For two taxa, there is only one cladogram (Figure 2.10A); for three taxa, three dichotomously branched cladograms can be constructed (Figure 2.10B); and for four taxa, 15 dichotomously branched cladograms are possible (Figure 2.10C). The formula for the number of trees is $\prod (2i-1)$, with \prod being the product of all the factors $(2i-1)$ from $i = 1$ to $i = n-1$,

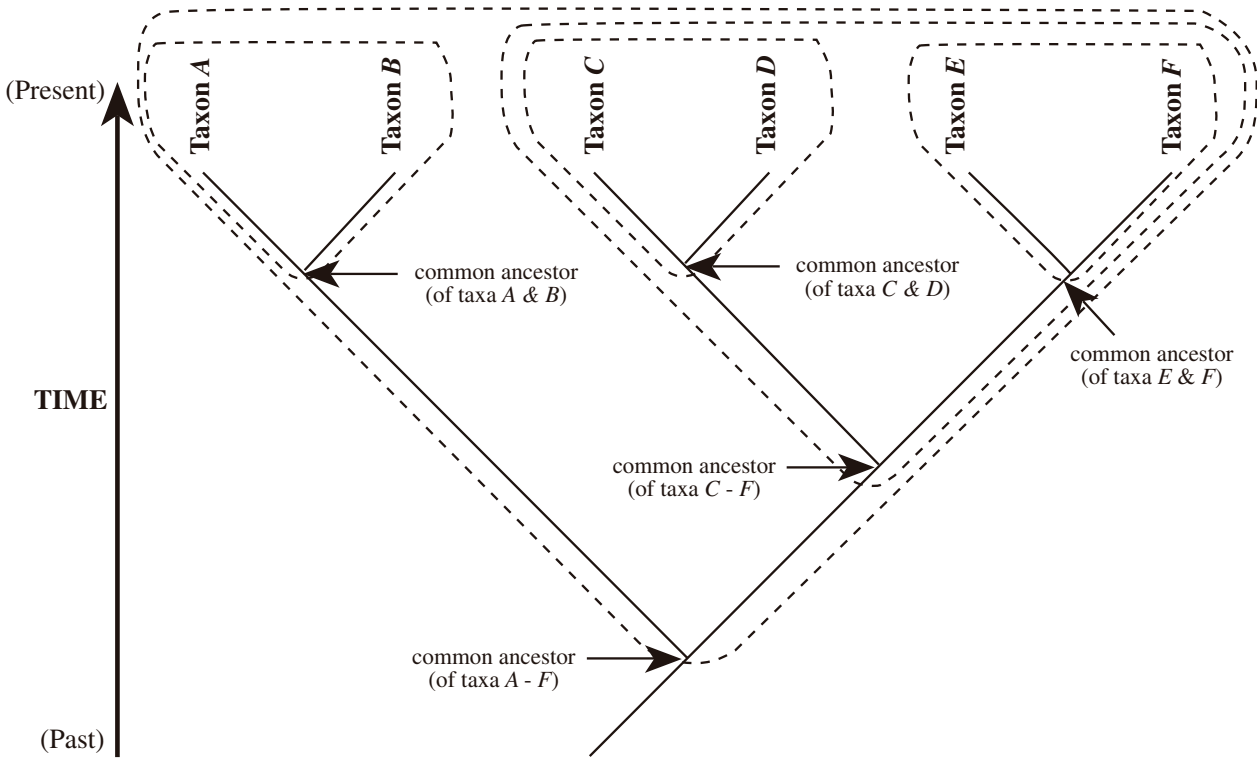
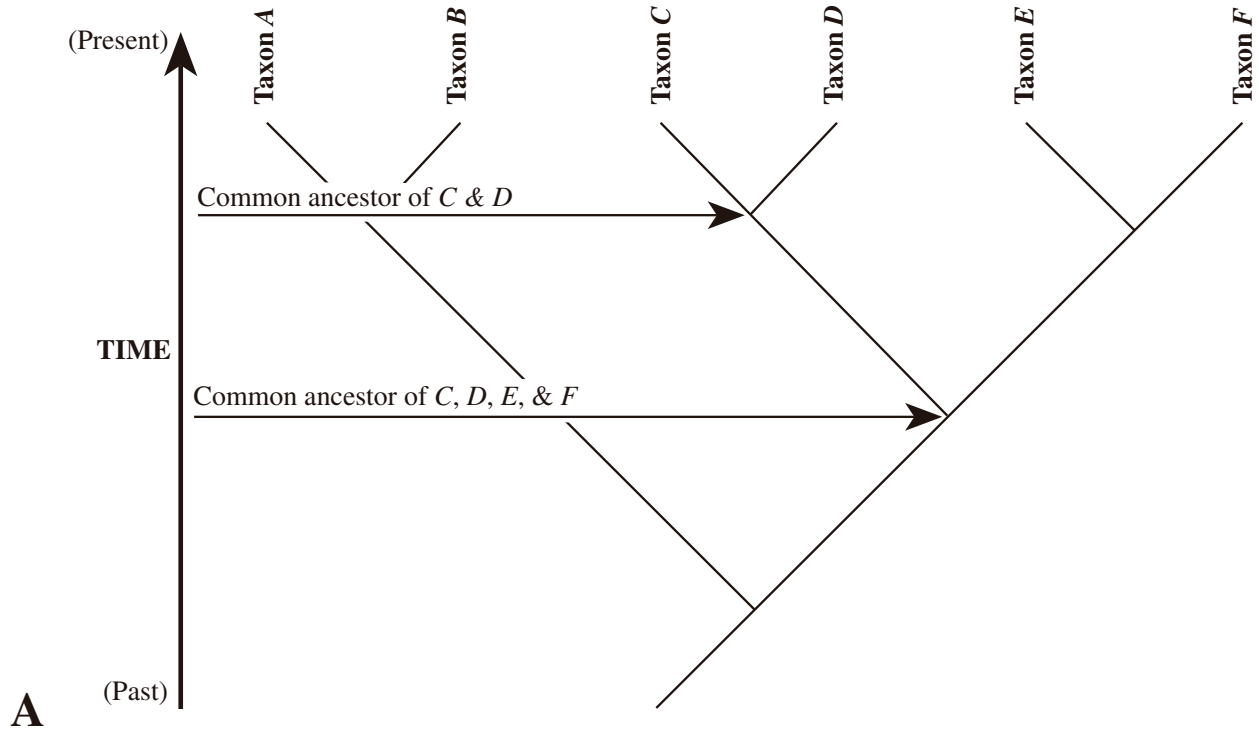


FIGURE 2.8 **A.** Hypothetical cladogram, illustrating recency of common ancestry. **B.** Cladogram of A with all monophyletic groups circled.

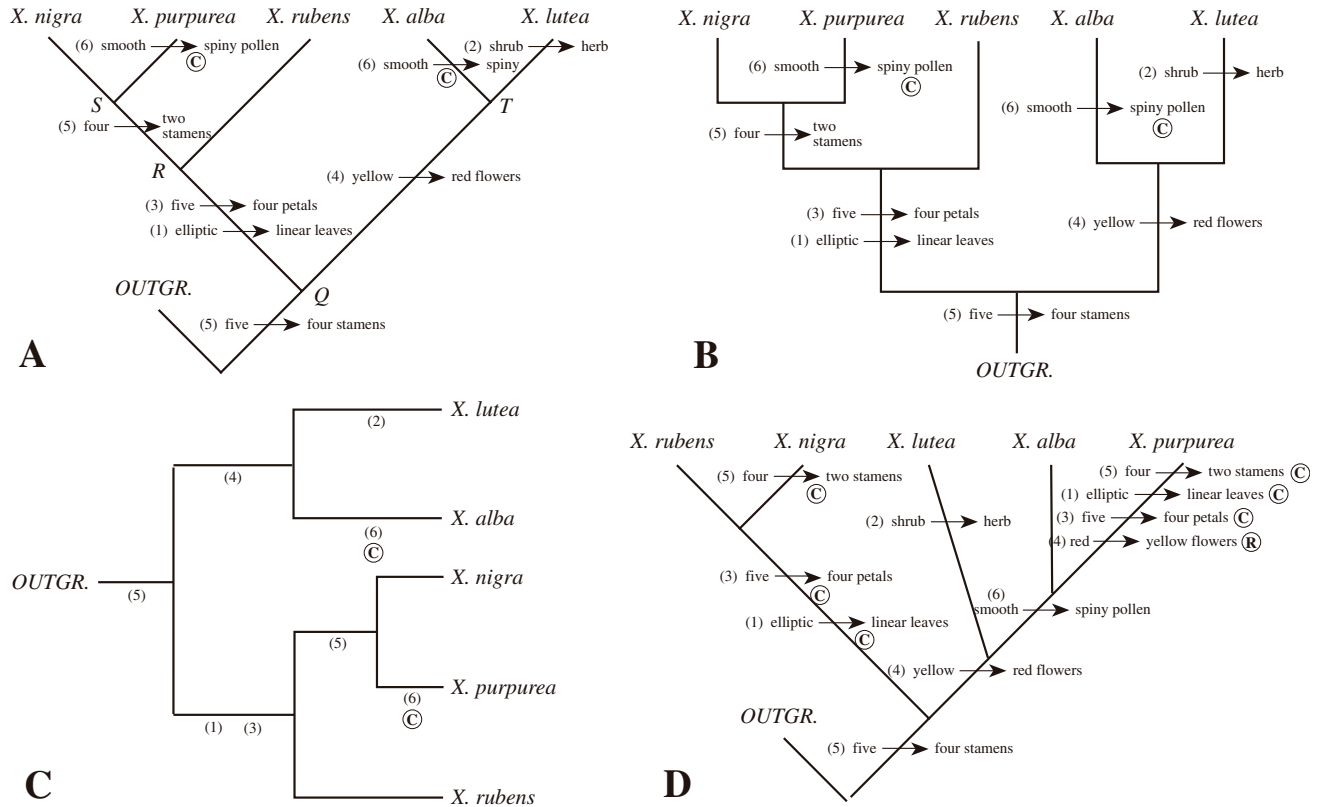


FIGURE 2.9 A–C. Most parsimonious cladogram of Fig. 2.7E. **A.** Cladogram with diagonal lines, but lineages rotated about common ancestor. **B.** Cladogram portrayed with perpendicular lines. **C.** As in B but rotated 90°. **D.** Alternative cladogram for the data set of Fig. 2.7A, showing a different relationship among the five species of genus *Xid*, requiring 11 character state changes, three more than the most parsimonious cladogram at Fig. 2.7E. Convergent and reversal homoplasies are denoted by circled **C** and **R**, respectively.

where n is the number of OTUs. For a cladistic analysis involving 54 OTUs, the number of possible dichotomously branching trees is 3×10^{84} (which is greater than the number of electrons in the universe!). The number of trees is even greater when the additional possibilities of reticulation or polytomies are taken into account (see later discussion).

Because there are generally many possible trees for any given data set, one of the major methods of reconstructing phylogenetic relationships is known as the **principle of parsimony** or **parsimony analysis**. The principle of parsimony states that of the numerous possible cladograms for a given group of OTUs, the one (or more) exhibiting the fewest number of evolutionary steps is accepted as being the best estimate of phylogeny. (Note that there may be two or more cladograms that are equally most parsimonious.) The principle of parsimony is actually a specific example of a general tenet of science known as Ockham's Razor (*Entia non sunt multiplicanda praeter necessitatem*), which states that given two or more competing hypotheses, each of which can explain the facts, the simplest one is accepted. The rationale for parsimony analysis is that the simplest explanation minimizes

the number of ad hoc hypotheses, i.e., hypotheses for which there is no direct evidence. In other words, of all possible cladograms for a given group of taxa, the one (or more) implying the fewest number of character state changes is accepted. A consequence of minimizing the total number of character state changes is to minimize the number of homoplasious reversals or convergences. The principle of parsimony is a valid working hypothesis because it minimizes uncorroborated hypotheses, thus assuming no additional evolutionary events for which there is no evidence.

Parsimony analysis can be illustrated as follows. For the example data set of Figure 2.7A, which includes five taxa (plus an outgroup), there are actually 105 possible dichotomously branching cladograms; the cladogram at Figure 2.7E (having total of eight character state changes) is only one of these. One of the other 104 alternative cladistic hypotheses is illustrated in Figure 2.9D. Note, however, that for this cladogram, there are a total of 11 character state changes (including three pairs of convergent evolutionary events and one reversal). Thus, of all the possible cladograms for the data set of Figure 2.7A, the one illustrated in Figure 2.7E is the

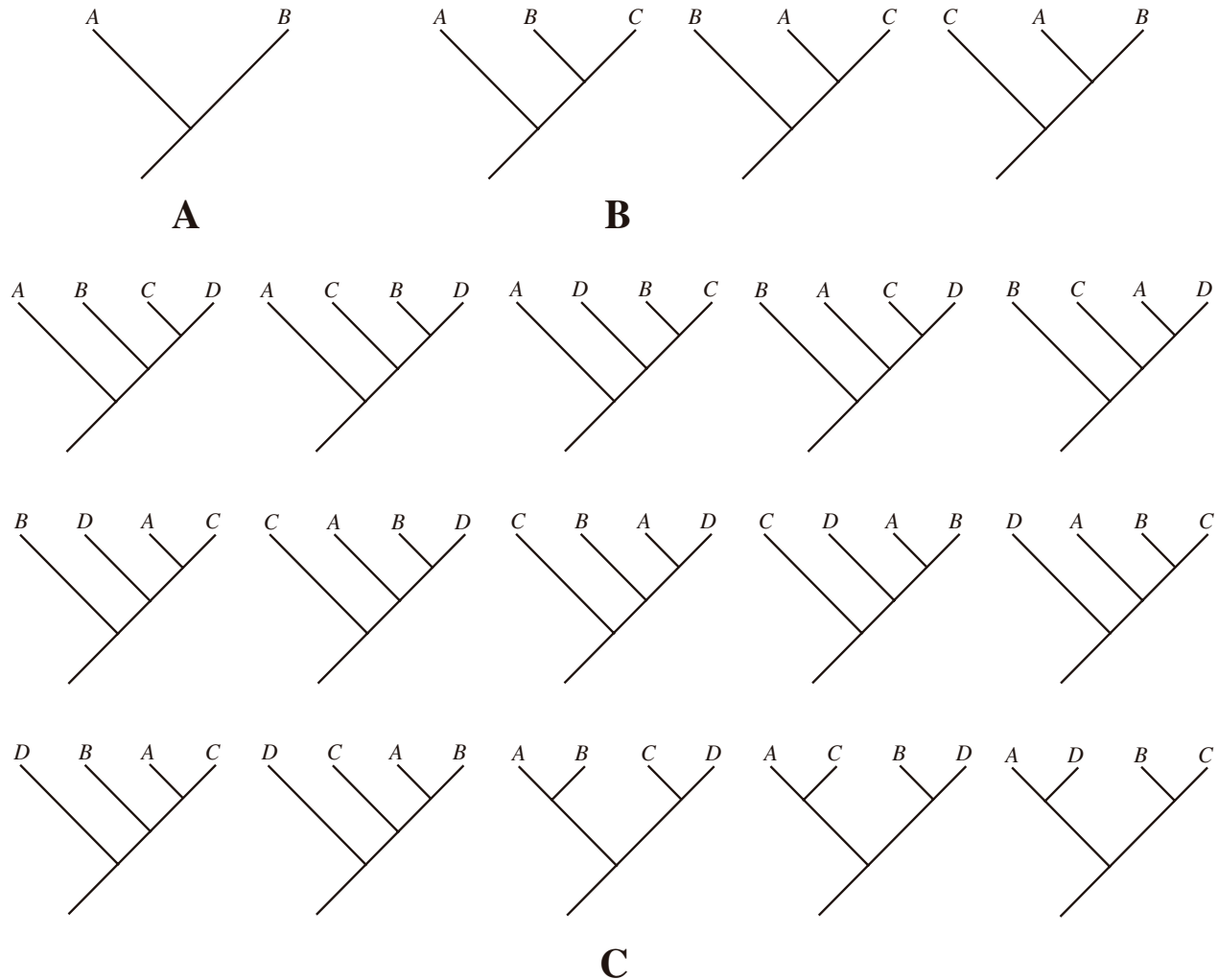


FIGURE 2.10 All possible dichotomously branched cladograms for a group consisting of the following. **A.** Two taxa (A and B). **B.** Three taxa (A, B, and C). **C.** Four taxa (A, B, C, and D).

shortest, containing the fewest number of evolutionary steps, and would be accepted as the best estimate of phylogeny.

Various computer programs (algorithms) are used to determine the most parsimonious cladogram from a given character \times taxon matrix. (See **Cladistic Computer Programs** at the end of this chapter.)

UNROOTED TREES

In contrast to a cladogram, a method for the representation of relative character state changes between taxa is the unrooted tree, sometimes called a network. An unrooted tree is a branching diagram that minimizes the total number of character state changes between all taxa. Unrooted trees are constructed by grouping taxa from a matrix in which polarity is not indicated (in which no hypothetical ancestor is designated), perhaps because the polarity of one or more

characters cannot be ascertained. Because no assumptions of polarity are made, no evolutionary hypotheses are implicit in an unrooted tree. Figure 2.11 illustrates the unrooted tree for the data set of Figure 2.7A,B. Note that monophyletic groups cannot be recognized in unrooted trees because relative ancestry (and therefore an outgroup) is not indicated. The character state changes noted on the unrooted tree simply denote evolutionary changes when going from one group of taxa to another, without reference to direction of change. After an unrooted tree is constructed, it may be rooted and portrayed as a cladogram. If the relative ancestry of one or more characters can be established, a point on the network may be designated as most ancestral, forming the root of the cladogram. For example, if the unrooted tree of Figure 2.11 is rooted at *, the result is the tree of Figure 2.7E. However, rooting is effectively done by simply including one or more

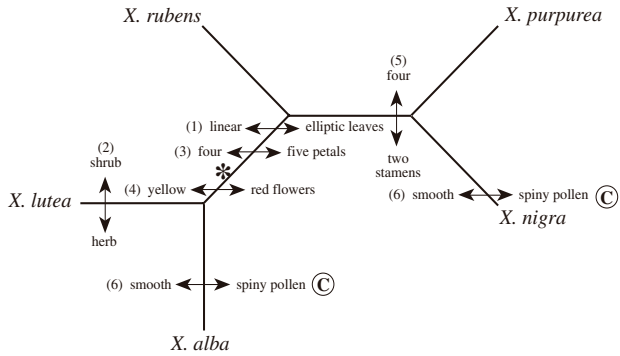


FIGURE 2.11 Unrooted tree for the data set of Fig. 2.7A (minus the *Outgroup* taxon). Direction of evolutionary change is not indicated and monophyletic groups cannot be defined. The * indicates the point of rooting that yields the tree of Fig. 2.7E.

outgroup(s) in the analysis and placing these outgroups at the base (the root) of the tree.

CHARACTER OPTIMIZATION

Optimization of characters refers to their representation (or plotting) in a cladogram in the most parsimonious way, such that the minimal number of character state changes

occur. Figure 2.12A,B shows a cladogram in which the evolution of a character is explained, but not in the most parsimonious way. In Figure 2.12C,D, the character is optimized, having the fewest number of state changes. In this example, character state evolution can be optimized in either of two equally parsimonious ways. *Acctran* (accelerated transformation) optimization hypothesizes an earlier initial state change with a later reversal of the same character (Figure 2.12C). *Deltran* (delayed transformation) optimization hypothesizes two later, convergent state changes (Figure 2.12D). Note that when alternative character optimization exists, there are nodes in the cladogram that are *equivocal*, i.e., for which the character state cannot be definitively determined.

Optimization is automatically performed by computer algorithms that trace characters and character states. (See end of chapter.)

POLYTOMY

Occasionally, the relationships among taxa cannot be resolved. A **polytomy** (also called a polychotomy) is a branching diagram in which the lineages of three or more taxa arise from a single hypothetical ancestor. Polytomies arise either because data are lacking or because three or more of the taxa were

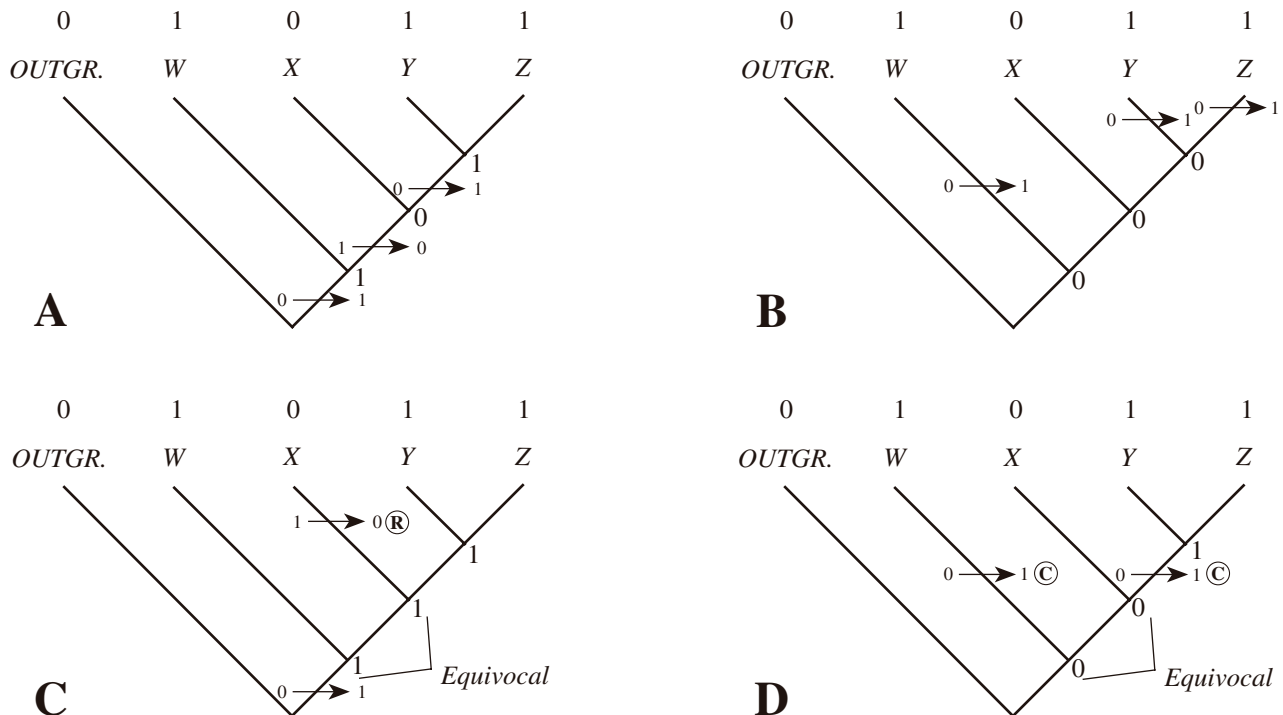


FIGURE 2.12 **A,B.** Cladograms for taxa W–Z and Outgroup, in which character states of a character (superposed above taxa) are accounted for by hypothesizing three state changes, not optimized. **C.** Optimization (*Acctran*) of character, hypothesizing two state changes, including one reversal. **D.** Optimization (*Deltran*) of character, hypothesizing two convergent state changes.

actually derived from a single ancestral species. (In addition, polytomies often are found in consensus trees; see later discussion.)

In the case of a polytomy arising via missing data, there are no derived character states identifying the monophyly of any two taxa among the group. For example, from the **character x taxon matrix** of Figure 2.13A, the relationships among taxa *W*, *X*, and *Y* cannot be resolved; synapomorphies link none of the taxon pairs. Thus, *W*, *X*, and *Y* are grouped as a polytomy in the most parsimonious cladogram (Figure 2.13B).

The other possible reason for the occurrence of a polytomy is that all of the taxa under consideration diverged independently from a single ancestral species. Thus, no synapomorphic evolutionary event links any two of the taxa as a monophyletic group. The occurrence of a polytomy in phylogenetic analysis should serve as a signal for the reinvestigation of taxa and characters, perhaps indicating the need for continued research.

RETICULATION

The methodology of phylogenetic systematics generally presumes the dichotomous or polytomous splitting of taxa, representing putative ancestral speciation events. However, another possibility in the evolution of plants is **reticulation**, the hybridization of two previously divergent taxa forming a new lineage. A reticulation event between two ancestral taxa (*E* and *F*) is exemplified in Figure 2.13D, resulting in the hybrid ancestral taxon *G*, which is the immediate ancestor of extant taxon *X*. Most standard phylogenetic analyses do not consider reticulation and would yield an incorrect cladogram if such a process had occurred. For example, the character x taxon matrix of Figure 2.13C is perfectly compatible with the reticulate cladogram of Figure 2.13D. However, the methods of phylogenetic systematics would construct the most parsimonious *dichotomously branching* cladogram of Figure 2.13E or 2.10F, which show homoplasy and require one additional character state change than Figure 2.13D.

Reticulation among a group of taxa should always be treated as a possibility. Data, such as chromosome analysis, may provide compelling evidence for past hybridization among the most recent common ancestors of extant taxa. A good example of this is the evolution of durum and bread wheat (*Triticum* spp.) via past hybridization and polyploidy (Figure 2.13G).

TAXON SELECTION AND POLYMORPHIC CHARACTERS

As alluded to earlier, the initial selection of taxa to be studied may introduce bias in a phylogenetic analysis. Prior to a phylogenetic analysis, each of the smallest unit taxa under study

(OTUs) *and* the group as a whole must be hypothesized to be monophyletic prior to the analysis. Monophyly is ascertained by the recognition of one or more unique, shared derived character states that argue for most recent common ancestry of all and only all members of the taxon in question. If such an apomorphy cannot be identified, any relationships denoted from the phylogenetic analysis may be in doubt. For example, in a cladistic analysis of several angiosperm genera (Figure 2.14A), only if each of the unit taxa (genera in this case) is monophyletic will the resultant cladogram be unbiased. If, however, genus *A* is not monophyletic, then it may be possible for some species of genus *A* to be more closely related to (i.e., have more recent common ancestry with) a species of another genus than to the other species of genus *A* (e.g., Figure 2.14B). Therefore, if any doubt exists as to the monophyly of component taxa to be analyzed, the taxa in question should be subdivided until the monophyly of these subtaxa is reasonably certain. If this is not possible, an exemplar species (selected as representative of a higher taxon and assumed to be monophyletic) may be chosen for a first approximation of relationships.

Related to the requirement of OTU monophyly is the problem of **polymorphic** characters, i.e., those that have variable character state values *within* an OTU. If an OTU for which monophyly has been established is polymorphic for a given character, then it may be subdivided into smaller taxonomic groups until each of these groups is monomorphic (i.e., invariable) for the character. If an OTU at the level of species is polymorphic, it is generally listed as such in computer algorithms.

If the ingroup as a whole is not monophyletic, the effect is identical to excluding taxa from the analysis, which could give erroneous results under certain conditions. For example, the most parsimonious cladogram constructed from the data matrix of Figure 2.14C is that of Figure 2.14D. However, if taxon *W* is inadvertently omitted from the ingroup (which is now not monophyletic; Figure 2.14E), then a different, most parsimonious cladogram topology may result for taxa *X*, *Y*, and *Z* (Figure 2.14F). The question of monophyly may be a serious problem for traditionally recognized taxa that were generally not defined by demonstrable apomorphies.

POLARITY DETERMINATION: OUTGROUP COMPARISON

As mentioned earlier in the discussion on character analysis, knowledge of character polarity is necessary to recognize shared derived character states that define monophyletic taxa. The only valid criterion for ascertaining polarity is outgroup comparison. An **outgroup** is a taxon that is not a member of

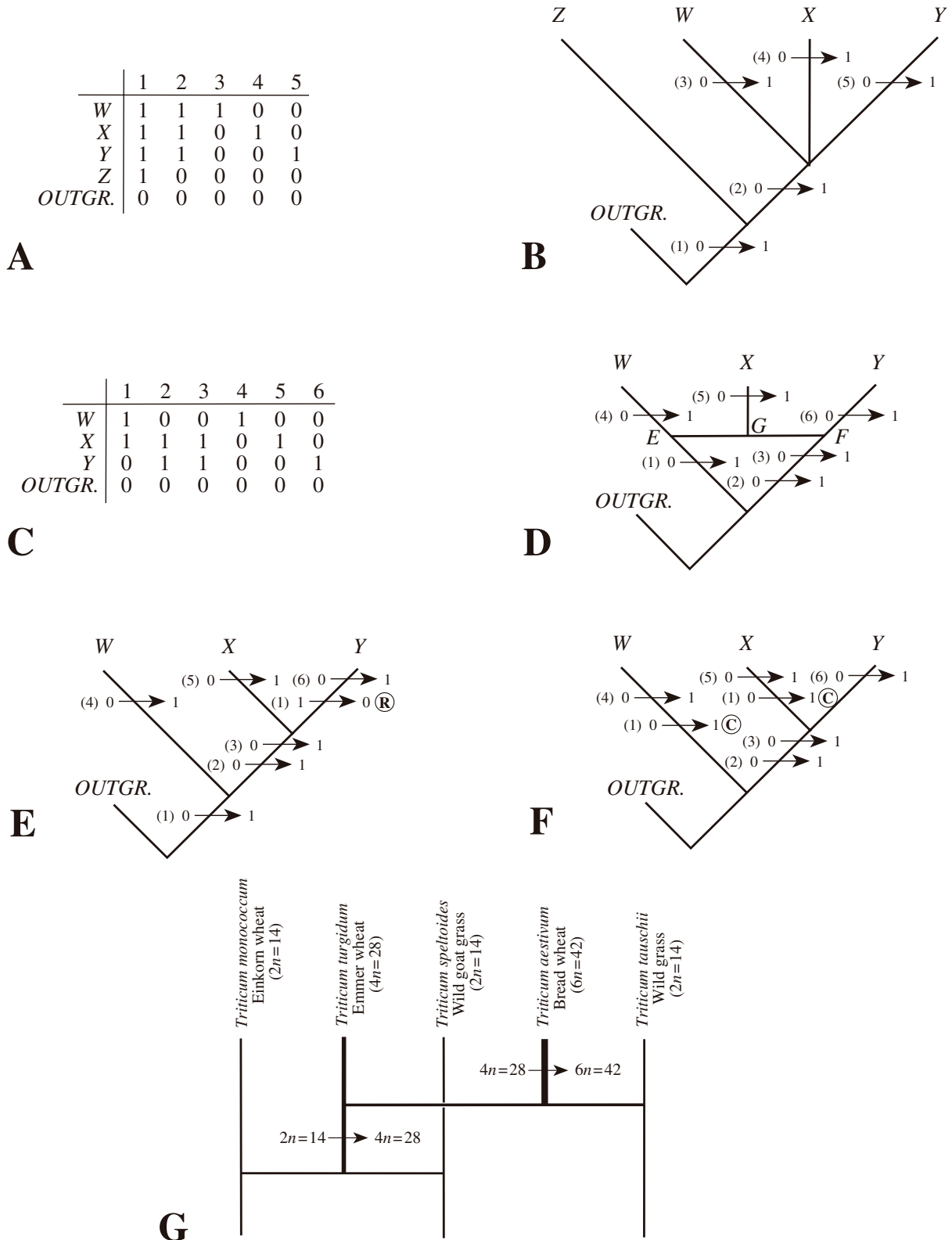


FIGURE 2.13 **A.** Hypothetical data set. **B.** Resultant tree from data set at A. Note polytomy of lineages to W, X, and Y. **C.** Hypothetical data set. **D.** Cladogram exhibiting reticulation that is compatible with data set at C. **E,F.** Dichotomously branching cladograms arising from data set at C, showing two alternative distributions of character state changes. **G.** Evolution of wheat via ancestral hybridization and polyploidy.

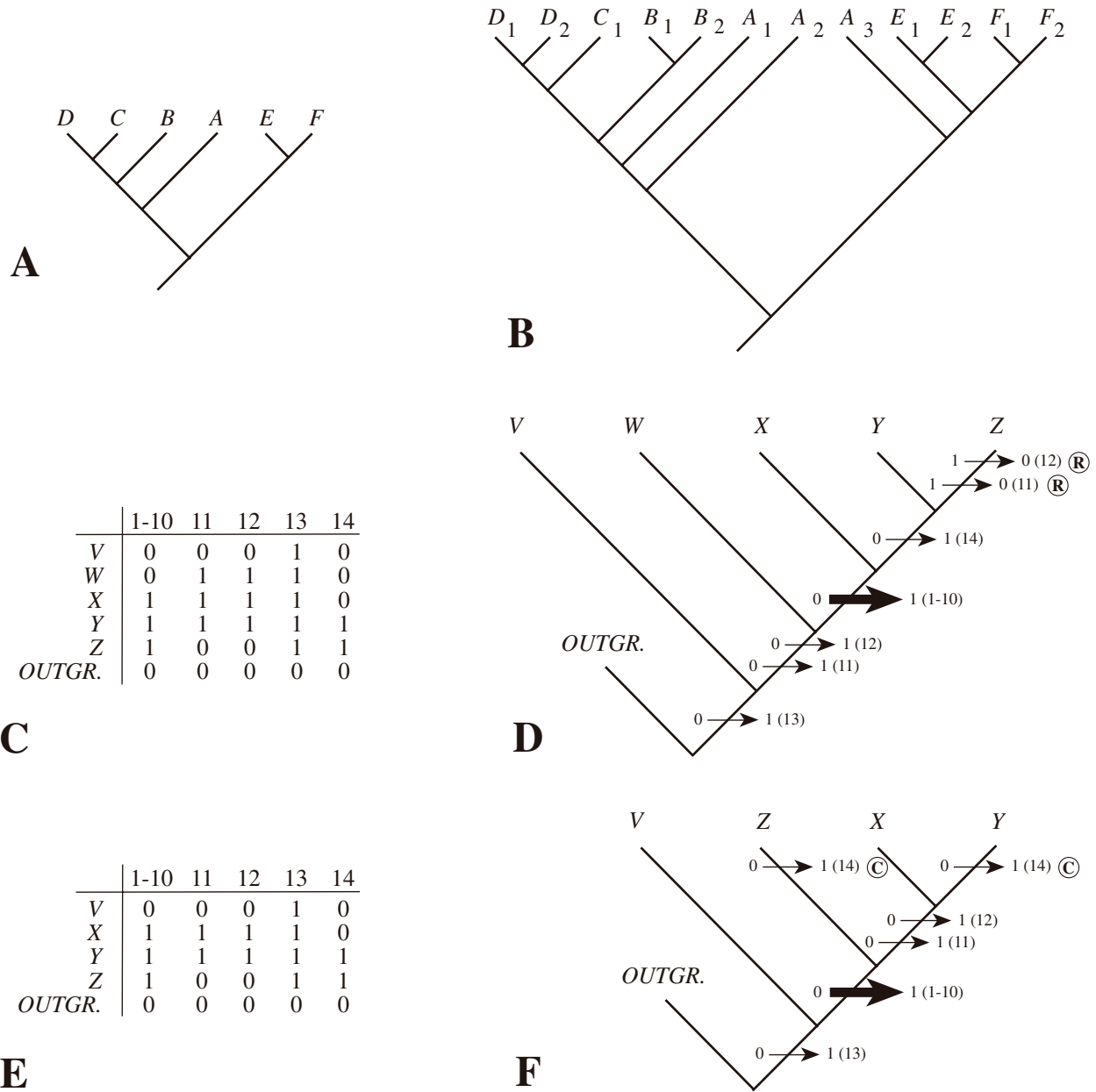


FIGURE 2.14 **A.** Cladogram of hypothetical genera A–E. **B.** Cladogram of the species of genera A–E. Note that genus A is paraphyletic. **C.** Character × taxon matrix for taxa V–Z plus *OUTGROUP*. **D.** Most parsimonious cladogram for taxa V, W, X, Y, and Z. **E.** Character × taxon matrix for same taxa, minus taxon W, omitted because it is not considered as part of the ingroup. **F.** Most parsimonious cladogram for taxa V, X, Y, and Z. Note different branching pattern for taxa X, Y, and Z.

the study group under investigation (the **ingroup**). Outgroup comparison entails character assessment of the *closest* outgroups to the ingroup. Those character states possessed by the closest outgroups (particularly by the sister group to the ingroup) are considered to be ancestral; states present in the ingroup, but not occurring in the nearest outgroups, are derived.

The rationale for outgroup comparison is founded in the principle of parsimony. For example, given some monophyletic ingroup X (Figure 2.15A), members of which possess either state 0 or 1 of a character, and given that taxon Y (nearest outgroup to X) possesses only character state 1, then the most parsimonious solution (requiring a single character change: 1 ⇒ 0) is that state 1 is ancestral and present in the

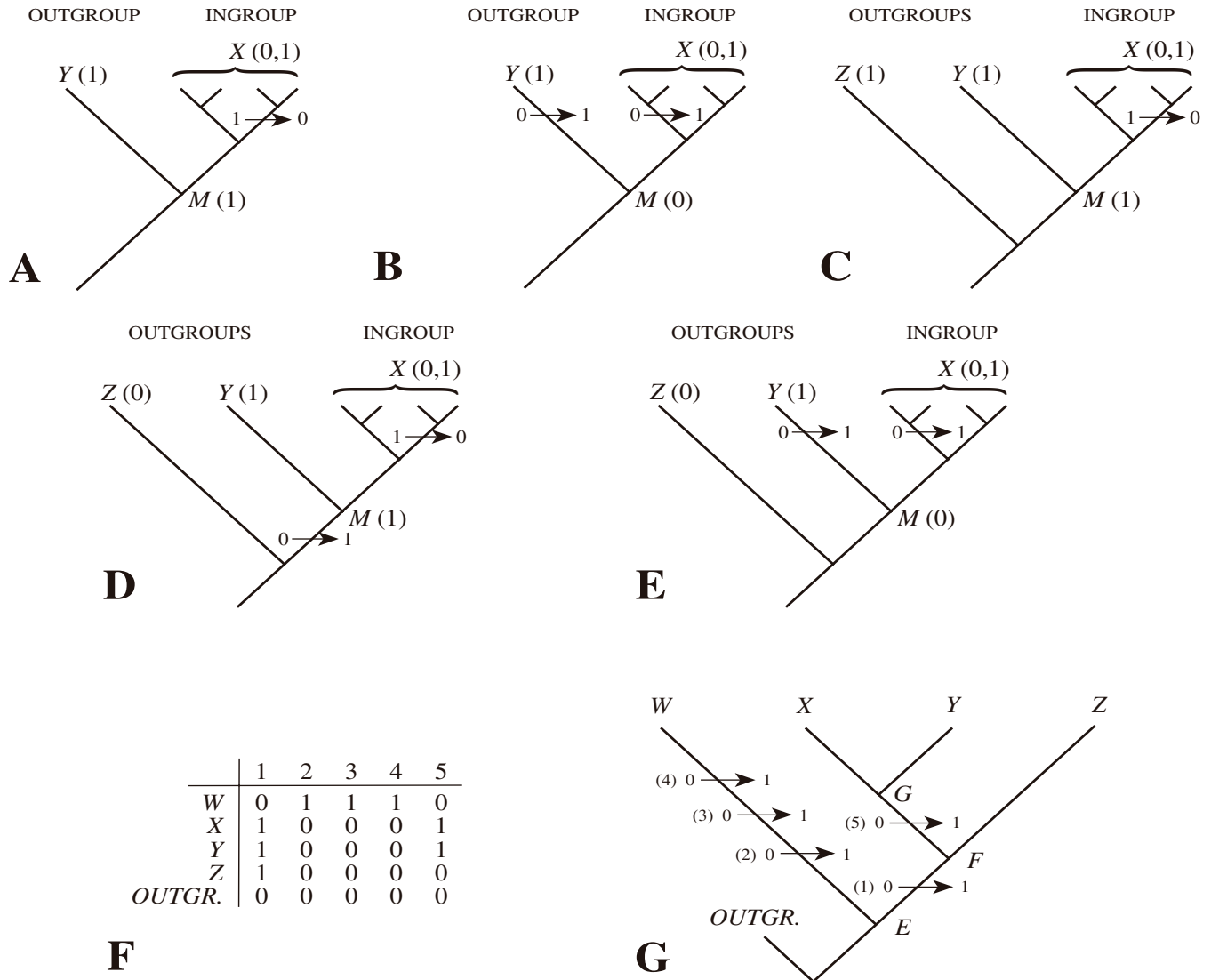


FIGURE 2.15 Determination of character state polarity using outgroup comparison. **A**. Most parsimonious assumption in which character state 1 is ancestral and present in ancestor *M*. **B**. Alternative, less parsimonious cladogram, in which state 0 is assumed to be ancestral. **C**. Verification of cladogram at A by addition of next outgroup Z, which also has state 1. **D, E**. Cladograms in which additional outgroup Z has state 0, showing that assumption of polarity is equivocal; ancestor *M* is equally likely to possess state 0 as opposed to state 1. **F**. Character x taxon matrix for taxa W Z plus *OUTGROUP*. **G**. Most parsimonious cladogram of taxa W, X, Y, and Z and ancestors E, F, and G.

common ancestor *M* (the outgroup node); character state 0 is derived within taxon *X* (Figure 2.15A). The alternative, that state 0 is ancestral, requires at least two character state changes (Figure 2.15B). Verification is made by considering an additional outgroup (e.g., taxon Z in Figure 2.15C). If this next outgroup possesses only character state 1, then the ancestral status of state 1 for taxon *Y* is substantiated (Figure 2.15C). If, however, outgroup Z contains only character state 0, then it is equally parsimonious to assume that state 1 is ancestral (Figure 2.15D) versus derived (Figure 2.15E). In this case, consideration of additional outgroups may resolve polarity.

The major problem with outgroup comparison is that the cladistic relationships of outgroup taxa may be unknown; in such a case, all possible outgroups (in all possible combinations) may be tested. In practice, prior studies at a higher taxonomic level are often used to establish near outgroups for a phylogenetic analysis.

ANCESTRAL VERSUS DERIVED CHARACTERS

A common point of confusion is seen in the use of the terms *ancestral* (plesiomorphic or primitive) and *derived* (apomorphic or advanced). It is advisable that these terms be limited

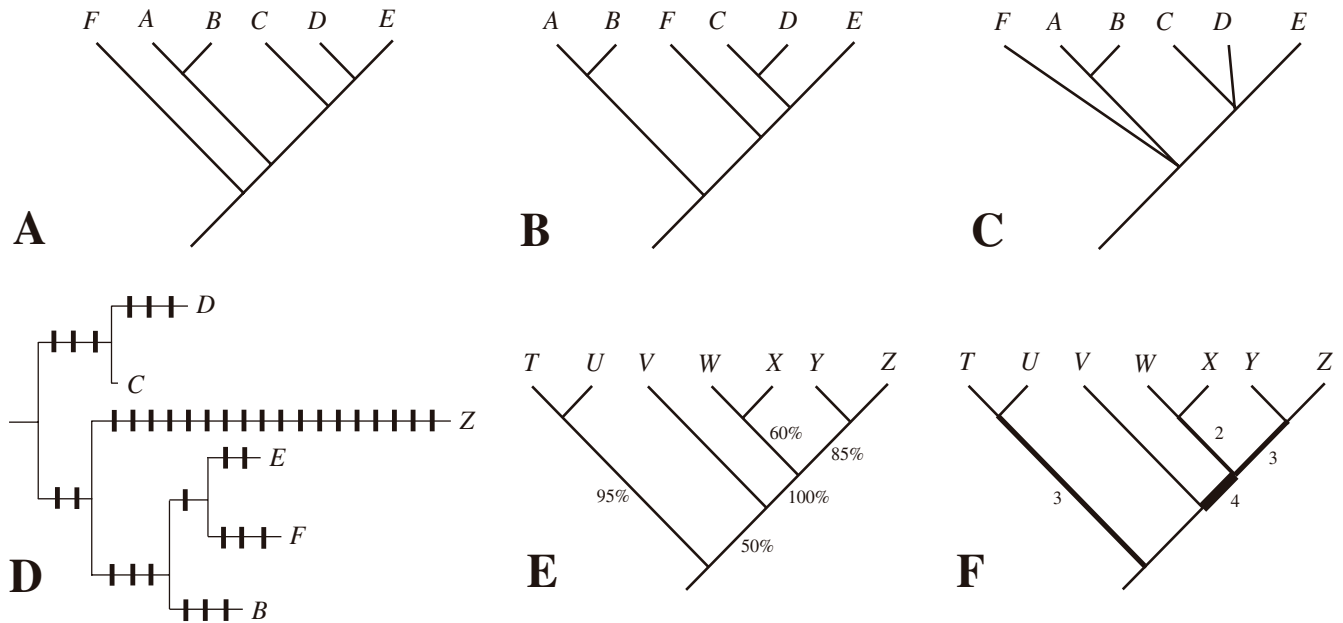


FIGURE 2.16 **A,B.** Two equally most parsimonious cladograms resulting from cladistic analysis. **C.** Strict consensus tree of cladograms at A and B. **D.** Cladogram illustrating taxon Z with very long branch. **E.** Cladogram showing bootstrap values at each internode. **F.** Same cladogram as in E showing decay index values. (Increasing numbers correspond to increasing line thickness; internodes not numbered have a decay index of 1.)

to the description of characters (not taxa) and then only relative to monophyletic groups. For example, in the cladogram of Figure 2.15G (constructed from the matrix of Figure 2.15F), state 1 of character 1 is derived *within* the group including W, X, Y, and Z (i.e., state 1 is absent in common ancestor E), but it is ancestral with regard to the monophyletic group X, Y, Z (i.e., state 1 is present in F, the common ancestor of X, Y, and Z). The use of the terms ancestral and derived to describe *taxa* should be avoided to prevent ambiguity. For example, from Figure 2.15G, it might be asked which taxon is most primitive? Confusion is avoided by describing, e.g., taxon W as phylogenetically most basal (or earliest diverging) and, e.g., taxon Z as possessing the fewest number of observed apomorphic states.

CONSENSUS TREES

In practice, most cladistic analyses yield numerous cladograms that are equally most parsimonious. Rather than view and discuss each of these cladograms, it is usually convenient to visualize the one tree that is compatible with all equally most parsimonious trees. A **consensus tree** is a cladogram derived by combining the features in common between two or more cladograms. There are several types of consensus trees. One of the most commonly portrayed is the **strict consensus tree**,

which collapses differences in branching pattern between two or more cladograms to a polytomy. Thus, the two equally parsimonious cladograms of Figure 2.16A,B are collapsible to the strict consensus tree of Figure 2.16C. Another type of consensus tree is the **50% majority consensus tree**, in which only those clades that occur in 50% or more of a given set of trees are retained. Consensus trees may be valuable for assessing those clades that are robust, i.e., that show up in all of the equally parsimonious trees. Greater confidence may be given to such clades in terms of recognition of accepted and named monophyletic groupings.

LONG BRANCH ATTRACTION

Sometimes, e.g., with molecular sequence data, one or more taxa will have a very long branch, meaning that these taxa have a large number of autapomorphies relative to other taxa in the analysis (e.g., taxon Z of Figure 2.16D). This can be caused by unequal rates of evolution among the taxa examined or can be the by-product of the particular data used. Such a situation can result in **long branch attraction**, in which taxa with relatively long branches tend to come out as close relatives of one another (or, if only one taxon has a long branch, its phylogenetic placement may easily shift from one analysis to another). Long branch attraction occurs because

when relatively numerous state changes occur along lineages, random changes can begin to outweigh nonrandom, phylogenetically informative ones. The phylogenetic placement of a taxon with a long branch can be uncertain and can unduly influence the placement of other taxa.

Taxa with long branches may need to be analyzed using a different data set. They are sometimes left out of an analysis to see what the effect is on cladogram robustness (see later discussion).

MAXIMUM LIKELIHOOD

The principle of parsimony can be viewed as evaluating all alternative trees (or as many subsets as feasible), calculating the length of those trees, and selecting those trees that are shortest, i.e., require the minimum number of character state changes under the set of conditions (character coding) specified. Another method of phylogenetic inference that deserves at least a brief mention is termed **maximum likelihood** (see references at end of chapter for a better understanding). Maximum likelihood, like parsimony methods, also evaluates alternative trees (hypotheses of relationship), but considers the *probability*, based on some selected model of evolution, that each tree explains the data. That tree which has the highest probability of explaining the data is preferred over trees having a lower probability. The appropriate model of evolution used is typically based on the data of the current analysis, but may be based on other data sets.

Maximum likelihood is used in practice for molecular sequence data, as illustrated in Figure 2.17A. In this simple example, there are three possible trees (shown as unrooted in Figure 2.17B and rooted arbitrarily at taxon Z in Figure 2.17C). Maximum likelihood evaluates each tree and calculates, for each character, the total probability that each node of the tree possesses a given nucleotide (Figure 2.17D). These individual probabilities (there are 16 in all; only three are illustrated in Figure 2.17D) are added together and the total probability for all characters is calculated. This total probability is compared with that for the other trees. That tree having the greatest overall probability is preferred over the others.

Maximum likelihood methods have an advantage over parsimony in that the estimation of the pattern of evolutionary history can take into account probabilities of a nucleotide substitution (e.g., purine to purine versus purine to pyrimidine; see Chapter 14) as well as varying rates of nucleotide substitution. The particular model used is typically evaluated from the data at hand, so in this sense it is empirical. Maximum likelihood methods may also eliminate the problem of long branch attraction (discussed earlier). Phylogenetic studies will very often present the results of both parsimony and maximum likelihood methods for comparison.

BAYESIAN ANALYSIS

Another more recent method of phylogenetic analysis is **Bayesian** inference (which is also worth mentioning briefly here, but see the references at the end of this chapter for a detailed understanding). This method is based upon *posterior probability*, utilizing a probability formula devised by T. Bayes in 1763.

Bayesian inference calculates the posterior probability of the phylogeny, branch lengths, and various parameters of the data. In practice, the posterior probability of phylogenies is approximated by sampling trees from the posterior probability distribution, using algorithms known as the *Markov chain Monte Carlo* (MCMC) or the *Metropolis-coupled Markov chain Monte Carlo* (MCMCMC). The results of a Bayesian analysis yield the probabilities for each of the branches of a given tree (derived from the 50% majority consensus tree of sampled trees). (Generally, a Bayesian probability of 95% or greater is considered robust for a particular clade: see **Cladogram Robustness**.)

MEASURES OF HOMOPLASY

If significant homoplasy occurs in a cladistic analysis, the data might be viewed as less than reliable for reconstructing phylogeny. One measure of the relative amount of homoplasy in the cladogram is the consistency index. **Consistency index (CI)** is equal to the ratio m/s , where m is the *minimum* number of character state changes that must occur and s is the *actual* number of changes that occur. The minimum number of changes is that needed to account for a single transformation between all character states of all characters. For example, a three-state character transformation, $0 \leftrightarrow 1 \leftrightarrow 2$, requires a minimum of two steps; e.g., one possibility (of several) is the change $0 \Rightarrow 1$ (first step) and then $1 \Rightarrow 2$ (second step).

A consistency index close to 1 indicates little to no homoplasy; a CI close to 0 is indicative of considerable homoplasy. As an example, the character x taxon matrix of Figure 2.7A,B necessitates a minimum of seven changes; i.e., there must be at least seven character state transformations to explain the distribution of states in the taxa. The actual number of changes in the most parsimonious cladogram is eight because of homoplasy (Figure 2.7E). Thus, the CI for this cladogram is $7/8 = 0.875$. The consistency index may be viewed as a gauge of confidence in the data to reconstruct phylogenetic relationships.

A consistency index may be calculated for individual characters as well. For example, relative to the most parsimonious cladogram of Figure 2.7E, the CI of all characters is equal to 1, except for character 6, which has a CI of 0.5 (because of two convergent character state changes).

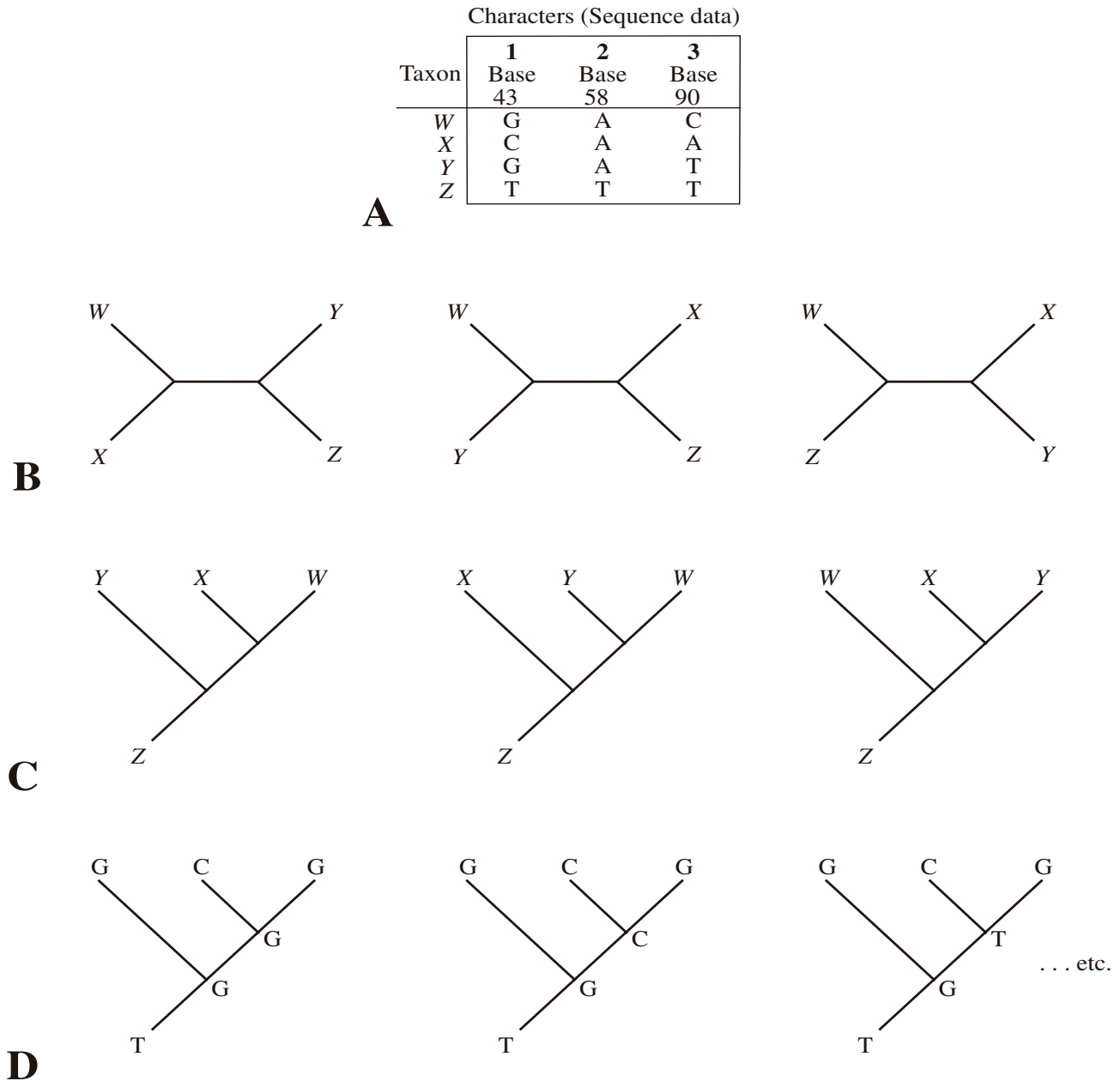


FIGURE 2.17 Maximum likelihood. **A.** Character \times taxon matrix (three nucleotide base sequences shown) for taxa *W*–*Z*. **B.** Three possible unrooted trees for taxa *W*–*Z*. **C.** Three possible rooted trees for taxa, with *Z* arbitrarily set as the root. **D.** Illustration of three of 16 possible nucleotide bases at nodes of one of the rooted trees.

Two other measures of homoplasy may be calculated: the **retention index (RI)** and the **rescaled consistency index (RC)**. The retention index is calculated as the ratio $(g - s) / (g - m)$, where g is the maximum possible number of state changes that could occur on any conceivable tree. Thus, the retention index is influenced by the number of taxa in the study. The rescaled consistency index (RC) is equal to the product of the CI and RI. The RC is used most often in

successive weighting; the rationale for its use is based on theoretical simulation studies.

CLADOGRAM ROBUSTNESS

It is very important to assess the confidence for which a tree actually denotes phylogenetic relationships. One way to evaluate cladogram robustness is the **bootstrap**. **Bootstrapping** is a method that reanalyzes the data of the original character \times

taxon matrix by selecting (resampling) characters *at random*, such that a given character can be selected more than once. The effect of this resampling is that some characters are given greater weight than others, but the total number of characters used is the same as that of the original matrix. This resampled data is then used to construct the most parsimonious cladogram(s). Many sequential bootstrapping analyses are generated (often 100 or more runs), and all most parsimonious cladograms are determined. From all of these most parsimonious trees, a 50% majority consensus tree is constructed; the percentages placed beside each internode of the cladogram represent the percentage of the time (from the bootstrap runs) that a particular clade is maintained (e.g., Figure 2.16E). A bootstrap value of 70% or more is generally considered a robustly supported node. The rationale for bootstrapping is that differential weighting by resampling of the original data will tend to produce the same clades if the data are good, i.e., reflect the actual phylogeny and exhibit little homoplasy. One problem with the bootstrapping method is that it technically requires a random distribution of the data, with no character correlation. These criteria are almost never verified in a cladistic analysis. However, bootstrapping is still the most used method to evaluate tree robustness. [Another method of measuring cladogram robustness is the so-called **jackknife** (or **jackknifing**), which is similar to the bootstrap but differs in that each randomly selected character may only be resampled once (not multiple times), and the resultant resampled data matrix is smaller than the original.]

A second way to evaluate clade confidence is by measuring clade decay. A **decay index** (also called Bremer support) is a measure of how many extra steps are needed (beyond the number in the most parsimonious cladograms) before the original clade is no longer retained. Thus, if a given cladogram internode has a decay index of 4, then the monophyletic group arising from it is maintained even in cladograms that are four steps longer than the most parsimonious (e.g., Figure 2.16F). The greater the decay index value, the greater the confidence in a given clade.

Finally, Bayesian analysis provides a measure of robustness in calculating posterior probabilities for each of the clades generated. Any branch with a posterior probability of 95% or greater is statistically well supported. (However, this method has come under some scrutiny because it often generates particularly high values of support.)

CLADOGRAM ANALYSIS

A typical cladistic analysis may involve the use of DNA sequence data from one or more genes plus the use of

morphological (i.e., nonmolecular) data. (Tests may be used to evaluate the homogeneity or compatibility of phylogenetic information from different types of molecular data, e.g., from chloroplast versus nuclear genes.) Often, separate analyses are done for (1) each of the gene sets individually; (2) all molecular data combined; (3) morphological data alone; and (4) a combined analysis utilizing all available data molecular and morphological. It has been demonstrated that utilizing the totality of data often results in the most robust cladogram. The strict consensus tree of this combined analysis generally represents the best estimate of phylogenetic relationships of the group studied.

From the most robust cladogram(s) derived from cladistic analyses, it is valuable to trace all character state changes. In addition, all monophyletic groupings should be evaluated in terms of their overall robustness (e.g., bootstrap support) and the specific apomorphies that link them together. Homoplasies (convergences or reversals) should also be noted. A homoplasy may represent an error in the initial analysis of that character that may warrant reconsideration of character state definition, intergradation, homology, or polarity. Thus, cladogram construction should be viewed not only as an end in itself, but as a means of pointing out those areas where additional research is needed to resolve satisfactorily the phylogeny of a group of organisms.

Cladograms represent an estimate of the pattern of evolutionary descent, both in terms of recency of common ancestry and in the distribution of derived (apomorphic) character states, which represent unique evolutionary events. Once a robust cladogram is derived, the pattern of relationships and evolutionary change may be used for a variety of purposes, discussed next.

PHYLOGENETIC CLASSIFICATION

One of the most important uses of cladograms is as a basis for classification. The pattern of evolutionary history portrayed in a cladogram may be used to classify taxa phylogenetically. A phylogenetic classification may be devised by naming and ordering monophyletic groups in a sequential, hierarchical classification, sometimes termed an **indented** method. The hierarchically arranged monophyletic groups may be assigned standard taxonomic ranks. For example, for the most parsimonious cladogram of Figure 2.18A, one possible classification of hypothetical genus *Xid* is seen in Figure 2.18B. Note that in this example, each named taxon corresponds to a monophyletic group (Figure 2.18A) and that these groups are sequentially nested such that the original cladogram may be directly reconstructed from this classification system. Two taxa of the same rank (e.g., Sections *Rubens* and *Nigropurpurea*) are automatically sister groups. Each higher taxon above

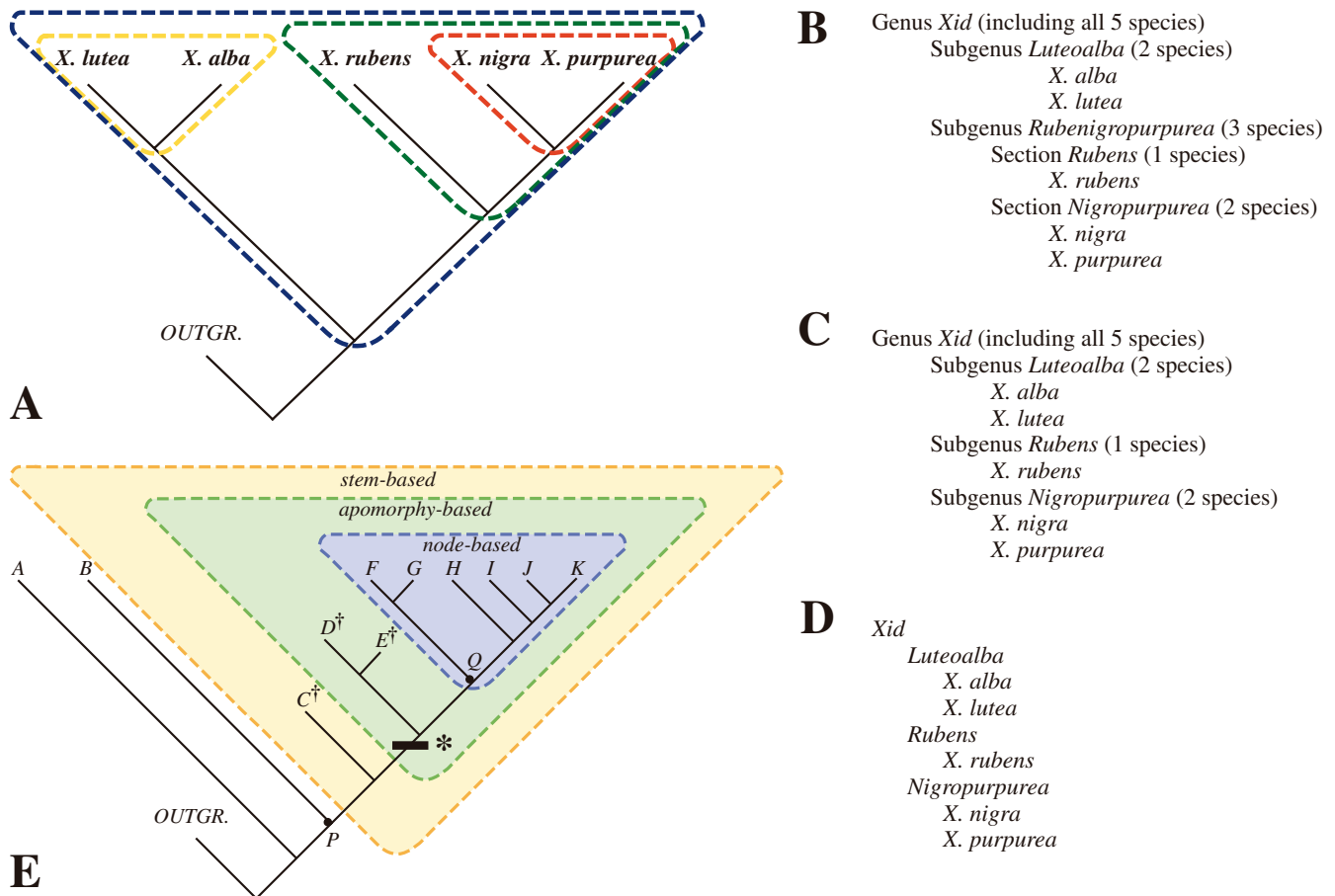


FIGURE 2.18 **A.** Cladogram from Fig. 2.7E. **B.** Indented classification scheme. **C.** Annotated classification scheme. **D.** Indented, but rankless, classification scheme. **E.** Cladogram illustrating node-based, apomorphy-based, and stem-based classification. †Extinct taxon. *Major evolutionary change, used as the basis for an apomorphy-based group.

(e.g., Subgenus *Luteoalba*) would also include automatically created lower taxa (e.g., species *Xid alba* and *Xid lutea* in this case).

An alternative, and often more practical, means of deriving a classification scheme from a cladogram is by annotation. **Annotation** is the sequential listing of derivative lineages from the base to the apex of the cladogram, each derivative lineage receiving the same hierarchical rank. The sequence of listing of taxa may be used to reconstruct their evolutionary relationships. For example, an annotated classification of the taxa from Figure 2.18A is seen in Figure 2.18C. In this case all named taxa are monophyletic, but taxa at the same rank are not necessarily sister groups.

The particular rank at which any given monophyletic group is given is arbitrary and is often done to conserve a past, traditional classification. A recent trend in systematics is to eliminate ranks altogether or, alternatively, to permit unranked names between the major rank names (see Chapter 16). In either case, the taxon

names, minus ranks, would still retain their hierarchical, evolutionary relationship (e.g., as in Figure 2.18D).

This most common type of phylogenetic classification is sometimes termed **node-based**, because it recognizes a node (common ancestor) of the cladogram and all descendants of that common ancestor as the basis for grouping (Figure 2.18E). In some cases, it may be valuable to recognize a group that is **stem-based**, i.e., one that includes the stem (internode) region just above a common ancestor plus all descendants of that stem (Figure 2.18E). A stem-based group might be useful, for example, in that it might include both a well-defined and corroborated node-based monophyletic group, plus one or more extinct, fossil lineages that contain some, but not all, of the apomorphies possessed by the node-based group. Yet a third general type of phylogenetic classification is **apomorphy-based**, in which all members of a monophyletic group that share a given, unique evolutionary event (illustrated by an * in Figure 2. 18E) are grouped together.

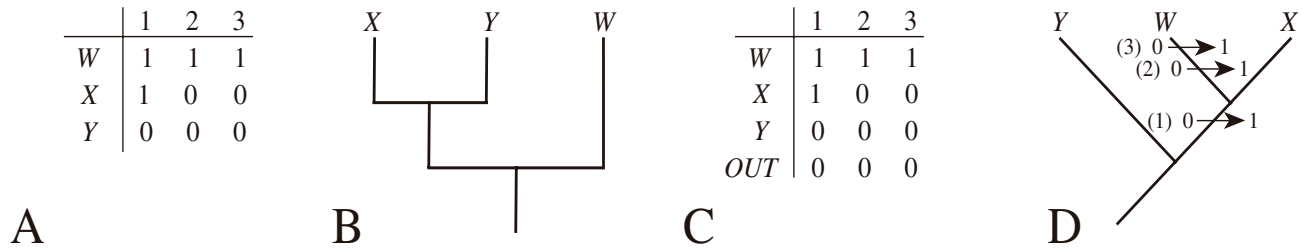


FIGURE 2.19 **A.** Character \times taxon matrix for taxa *W*–*Y*. **B.** Phenogram of taxa *W*, *X*, and *Y*. **C.** Character \times taxon matrix for taxa *W*–*Y* plus *OUTGROUP*. **D.** Most parsimonious cladogram of taxa *W*, *X*, and *Y*. Note different branching pattern.

Last, it should be mentioned that a monophyletic group can be recognized with a phylogenetic definition. For example, in Figure 2.18A, the monophyletic *Xid* might be defined as the least inclusive monophyletic group containing the common ancestor of *X. lutea* and *X. nigra*. The rationale is that this presents a more explicit and stable means of classification of taxa. However, any given phylogenetic definition is based on some cladistic analysis. If future cladistic analyses portray a somewhat different relationship of taxa, then the phylogenetically defined groups may contain taxa that were unintended, making them less useful and less stable than more standard classifications.

As mentioned in Chapter 1, a second major type of classification is **phenetic**, in which taxa are grouped by overall similarity. This phenetic grouping may be represented in the form of a branching diagram known as a **phenogram**. For example, for the data matrix of Figure 2.19A, the resultant phenogram is seen in Figure 2.19B. (Note that no outgroup is included in the matrix.) Phenetic classifications will often be quite different from phylogenetic ones because in a phenetic analysis, taxa may be grouped together by shared *ancestral* features (known as **symplesiomorphies**) as well as by shared derived character states (synapomorphies). For example, the data matrix of Figure 2.19C (identical to that of 2.16A except for the addition of an outgroup) yields the most parsimonious cladogram at Figure 2.19D, which has a different branching pattern from the phenogram of Figure 2.19B. Note that in the cladogram, taxa *W* and *X* are grouped as sister taxa because they share the derived state of character 1, which is a synapomorphy for *W* and *X*. In contrast, the phenogram of Figure 2.19B groups together taxa *X* and *Y* because they are more similar, having in common state 0 of characters 2 and 3; however, these are shared ancestral states (symplesiomorphies) and cannot be used to recognize monophyletic groups. Because many past classification systems have been based on overall phenetic similarity, great caution should be taken in evaluating relationship. Taxa that are most similar to one another may not, in fact, be particularly close relatives in a phylogenetic sense (i.e., by recency of common ancestry).

In summary, phylogenetic classification of taxa has the tremendous advantage of being based upon and of reflecting the

evolutionary history of the group in question. The International Code of Botanical Nomenclature (Chapter 16) has been used very successfully to assign taxonomic names based on the criterion of monophyly (although some problems persist that it is hoped will be addressed in future versions of the Code). Phylogenetic classifications have resulted in several name changes in some groups, but these are gradually beginning to stabilize, particularly with additional, robust molecular studies. In practice, assigning a name to every monophyletic group, whether ranked or not, is unwieldy, impractical, and unnecessary. Generally, only monophyletic groups that are well supported (and ideally that have a well-recognized apomorphy) should be formally named, and every effort should be made to retain (or modify) former classification systems, where possible.

CHARACTER EVOLUTION

Cladograms can be used as an analytical device to evaluate evolutionary change within a given character. Examination of the cladogram with reference to the distribution of the states of this character reveals one or more optimized (most parsimonious) explanations for the evolution of that character. Such an analysis can be used to verify the original coding of characters. For example, cladistic analyses of angiosperms (based on numerous characters) verify that the spines of cacti and those of euphorbs have evolved independently of one another. Had these features been coded as the same state of the character spine presence/absence, convergent evolution would be evident in the cladogram, indicative of an initially faulty assessment of homology.

In addition, certain characters may have been omitted from the original character \times taxon matrix because of incomplete data or uncertainty with regard to homology, polarity, or intergradation of character states. However, the sequence of evolutionary changes of these characters may be ascertained by superposing the states of the characters not included on the terminal taxa of the cladogram. If it is assumed that the cladogram is correct, then the most parsimonious explanation for the distribution of character states may be obtained by optimization. For example, Figure 2.20 illustrates the most

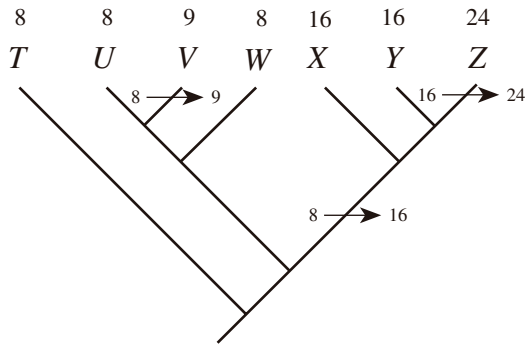


FIGURE 2.20 Analysis of character data that were not included in an original cladistic analysis. Cladogram in which chromosome data are superposed on taxa names. Note optimization of evolutionary change in this feature.

parsimonious cladogram for taxa *T*–*Z*, in which the character chromosome number was not originally included. A superposition of known haploid chromosome numbers may be used to hypothesize the most parsimonious (or optimum) evolutionary pattern (Figure 2.20).

BIOGEOGRAPHY AND ECOLOGY

A phylogenetic analysis can be used to evaluate past changes in biogeographic distribution and ecological habitat. Both distribution and habitat data are considered to be extrinsic in nature, i.e., not determined by the genetic makeup (genome) of a taxon, and, therefore, not subject to biological evolution. Thus, data on distribution and habitat cannot be included in the data matrix of a cladistic analysis. (Note that ecological data in the simple sense of the habitat a taxon occupies, such as desert or salt marsh, is extrinsic. However, the propensity or capability to survive in a particular habitat, e.g., physiological or morphological adaptations that allow survival in the desert, are intrinsic and may be used directly as characters in an analysis.) A historical analysis of extrinsic data may be accomplished by superposing the data onto an existing cladogram and optimizing the changes that would be needed, using the principle of parsimony (see later discussion).

Analysis of biogeographic data can give insight into the direction of change in biogeographic distribution. A change from one distribution to another can occur by either of two means: dispersal or vicariance. **Dispersal** is the movement of an organism or propagule from one region to another, such as the transport of a seed or fruit (by wind, water, or bird) from a continent to an island (Figure 2.21A). **Vicariance**, in contrast, is the splitting of one ancestral population into two (or more) populations, e.g., by continental drift or the formation of a new waterway or mountain range, resulting in a barrier between the split populations; this barrier prevents gene flow

between these populations, allowing them to diverge independently (Figure 2.21B).

Determining vicariance versus dispersal as an explanation for biogeographic change cannot always be made, and requires additional knowledge of geologic history. For example, Figure 2.21C illustrates a cladogram of taxa endemic to the Hawaiian archipelago, in which the ranges (by island) are superposed. A simple optimization shows the changes in geographic ranges that would be needed to explain the data. In this case, a shift from the island of Kauai to Maui and one from Maui to the island of Hawaii constitutes the simplest explanation needed to account for the current distribution of taxa. Because geologic data firmly suggests that the Hawaiian islands arose from sequential hot-spot volcanic activity and that the major islands were never connected, vicariance as an explanation is ruled out, leaving dispersal as the mechanism for biogeographic change. The hypothetical example of Figure 2.21D shows another cladogram in which both biogeographic distributions are superposed. A likely explanation for change in biogeographic distributions in this example is the splitting of the three continents from an ancestral Gondwana (Figure 2.21D). Although dispersal across oceans cannot be ruled out, vicariance might be more likely because the changes in distribution correspond to a hypothesis of continental drift. (Note that the continentally delimited groups need not be monophyletic.)

An example of tracing extrinsic ecological data is seen in Figure 2.21E, in which habitat types are superposed on the taxa from a cladistic analysis. Note in this example the shift from a terrestrial to an aquatic habitat. Analyses such as this may yield insight into the adaptive significance of evolutionary changes in anatomy, morphology, or physiology relative to differing habitat requirements.

ONTOGENY AND HETEROCHRONY

Phylogeny and character evolution are normally studied only with regard to the mature features of adult individuals. However, a mature structure, whether organ, tissue, or cell, is the end product of **ontogeny**, the developmental sequence under the control of a number of genes. Ontogeny may be visualized in either of two ways. First, a study of the developmental pattern may reveal a series of discrete structural stages or entities, one transforming into the next until the end point (the mature adult structure) is obtained. These discrete stages are identified and named and the transformation in **ontogenetic sequence**, from one stage to the next, is compared in different taxa (Figure 2.22A). Second, some feature of the developmental change of a structure may be measured quantitatively as a function of real time. This plot of morphology as a function of time is called an **ontogenetic trajectory** (Figure 2.22B). Ontogenetic trajectories may be compared

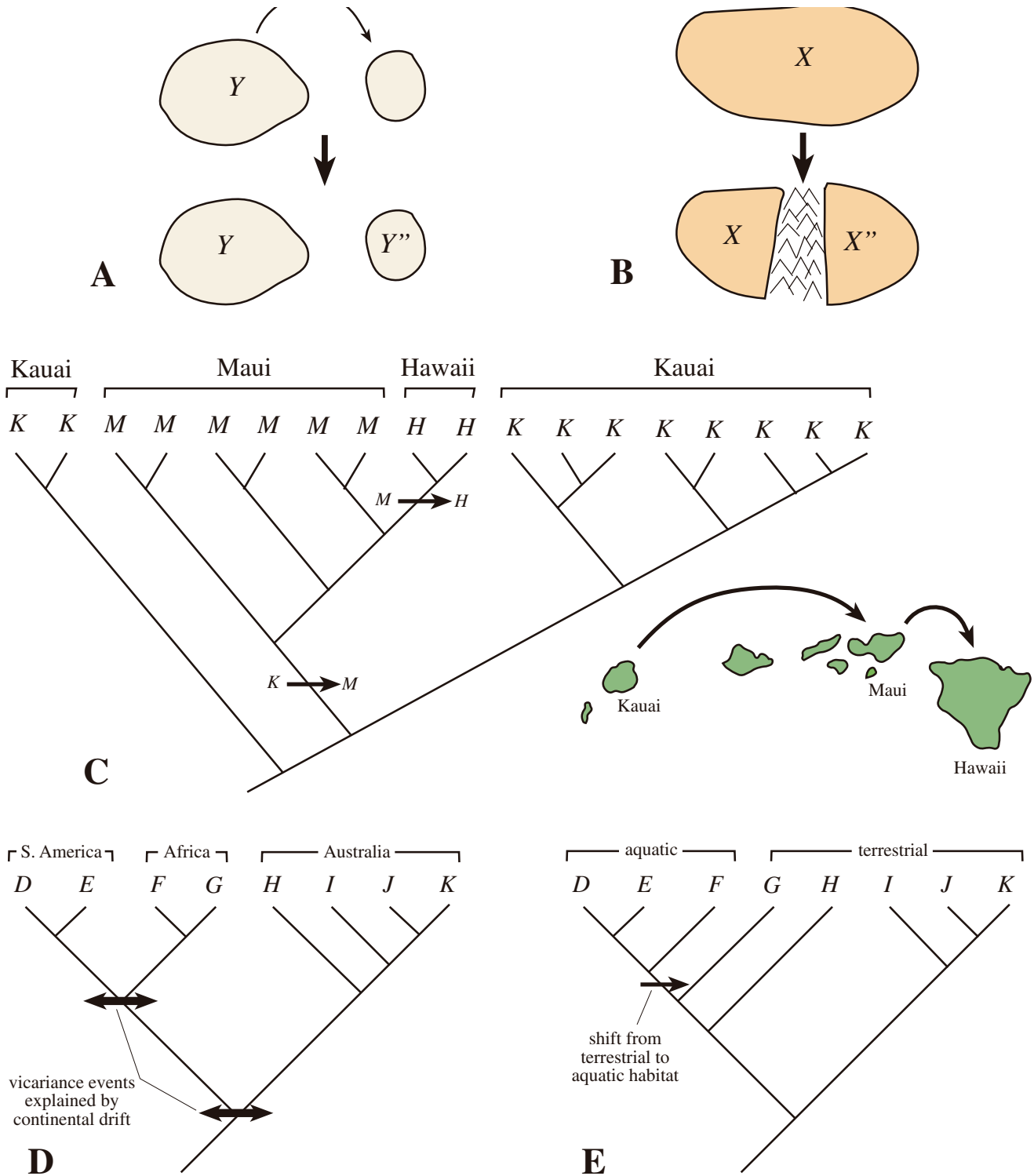


FIGURE 2.21 Cladistic analysis of biogeographic data. **A.** Hypothesis of dispersal, in which a propagule of species *Y* lands on another island. The isolated population subsequently diverges into *Y'*. **B.** Hypothesis of vicariance, in which ancestral population *X* is divided into two populations by a mountain range. The two populations, now isolated, can subsequently diverge, one becoming *X'*. **C.** Cladogram in which geographic distributions are superposed atop lineages, illustrating dispersal in Hawaiian archipelago. Optimized explanation is dispersal of ancestral taxa from Kauai to Maui and then from Maui to Hawaii. **D.** Cladogram in which geographic distributions are superposed atop lineages, illustrating vicariance. Continental drift explains current distribution of taxa *D*–*K*. **E.** Superposition of ecological habitat data, illustrating use of cladogram to deduce history of ecological change.

between different taxa. Note, e.g., in Figure 2.22B that taxon *Z* and taxa *W* and *Y* have the same adult structures but differing ontogenetic trajectories.

Ontogenetic data may be used in a cladistic analysis like any other character. Thus, two or more discrete ontogenetic sequences (Figure 2.22A) or ontogenetic trajectories (Figure 2.22B) may be defined as separate character states of a developmental character. The polarity of ontogenetic character states may be assessed by outgroup comparison as can be done for any other character.

Evolution may often be manifested by a change in ontogeny. An evolutionary change in the rate or timing of development is known as **heterochrony**. Heterochrony has apparently been an important evolutionary mechanism in many groups, in which the relatively simple evolutionary alteration of a regulatory gene results in often profound changes in the morphology of a descendant. Heterochrony can be assessed by performing a cladistic analysis and determining from this the ancestral versus the derived condition of an ontogenetic sequence or trajectory. The two major categories of heterochrony are peramorphosis and paedomorphosis. **Peramorphosis** is a derived type of heterochrony in which ontogeny passes through *and goes beyond* the stages or trajectory of the ancestral condition. Peramorphosis can result in the addition of a new stage or an ontogenetic trajectory that continues beyond that of the ancestral trajectory. For example, in Figure 2.22C, the derived ontogenetic sequence of taxa *A* and *D* ($s^1 \Rightarrow s^2 \Rightarrow S^3$) is the result of peramorphosis via the terminal addition of stage S^3 to the ancestral sequence ($s^1 \Rightarrow S^2$). (Note that s represents a juvenile developmental stage; S is a mature, adult feature.) Thus, the adult condition (S^2) in the ancestral ontogeny is homologous with a juvenile condition (s^2) in the derived ontogeny of taxa *A* and *D*. This principle is termed *terminal addition* or *Haeckelian recapitulation* and is often summarized by the expression ontogeny recapitulates phylogeny.

Paedomorphosis is a type of heterochrony in which the mature or adult stage of the derived ontogenetic sequence resembles a juvenile ontogenetic stage of the ancestral condition. (**Neotony** is one type of paedomorphosis that is caused by a *decrease* in the rate of development of a structure.) For example, in Figure 2.22D, the derived ontogenetic sequence of taxon *Z* ($s^1 \Rightarrow S^2$) is the result of paedomorphosis by the terminal loss of stage S^3 in the ancestral sequence ($s^1 \Rightarrow s^2 \Rightarrow S^3$). Thus, the adult condition (S^2) in the derived ontogeny of taxon *Z* is homologous with a juvenile condition (s^2) in the ancestral ontogeny. In a cladistic analysis paedomorphosis is portrayed as the *reversal* of a character state and can only be detected via the utilization of other characters in the analysis.

Evolutionary change may result in the modification of mature structures by affecting early developmental stages. For example, if the ontogeny of structure S^3 occurs in two discrete stages ($s^1 \Rightarrow s^2$) and ($s^2 \Rightarrow S^3$), then a single alteration of the regulatory pathway controlling the first developmental sequence (represented by $*$ in Figure 2.22 E) may cause a change in both the final structure and the intermediate stage (e.g., to $s^1 \Rightarrow s^4 \Rightarrow S^5$; Figure 2.22E). Thus, structural evolution may occur by modification at any developmental stage, and mature ancestral structures need not be preserved as extant juvenile developmental stages.

A PERSPECTIVE ON PHYLOGENETIC SYSTEMATICS

The careful researcher, in constructing cladograms or critically reading cladistic analyses in the literature, should be aware of several potential pitfalls in phylogenetic studies. Lack of consideration of any of the following renders the study questionable at best and useless at worst. Are unit taxa (OTUs) and the group as a whole monophyletic? If evidence for monophyly is not presented, the study may be faulty from the start. What are the sources of the data? The validity of a phylogenetic study is based on the comprehensiveness and accuracy of the original descriptive data. *Which* characters are selected and how are they defined? It is important to question the basis for the selection of these characters and not others. Are character states assessed for discreteness? Is homology assessed? Has an effort been made to determine whether similar characters and character states presumably have a common evolutionary origin? Or is observed similarity more one of traditional and imprecise terminology and possibly homoplasious? Have any characters been weighted? If so, what is the rationale behind it? How are polarities determined? The evidence for selection and relative placement of outgroups should be thoroughly investigated. Finally, is the resultant cladogram analyzed in terms of monophyletic groupings, character state changes, assessment of convergences and reversals, testing of homology, and possible reevaluation of characters and character states? The thorough phylogenetic study critically reviews each step of cladogram construction, considers all alternatives, and evaluates and reevaluates the significance of the phylogenetic analysis in terms of future research that might clarify our understanding of plant evolutionary relationships. Although the determination of phylogeny using the methodology of phylogenetic systematics may be problematic, it has the significant advantage of being repeatable and explicit. Each step of the analysis can be duplicated, evaluated, and critiqued in subsequent investigations.

Taxon *T*: $s^1 \longrightarrow S^2$
 Taxon *U*: $s^1 \longrightarrow s^2 \longrightarrow S^3$
 Taxon *V*: $s^1 \longrightarrow s^2 \longrightarrow s^3 \longrightarrow S^4$

A

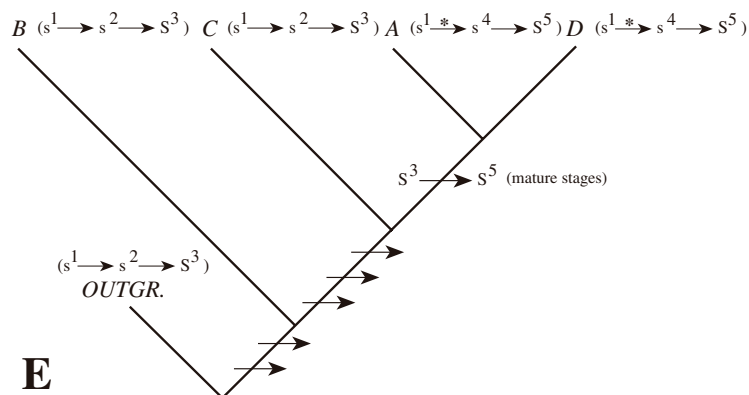
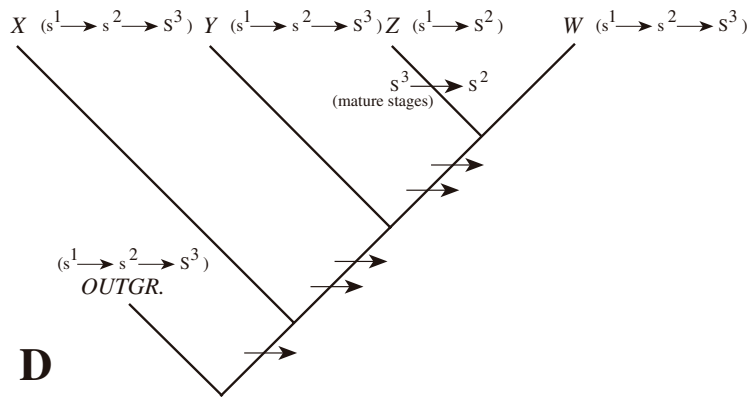
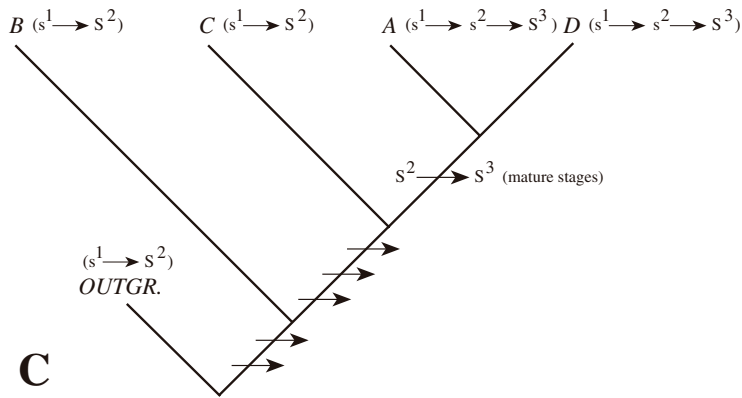
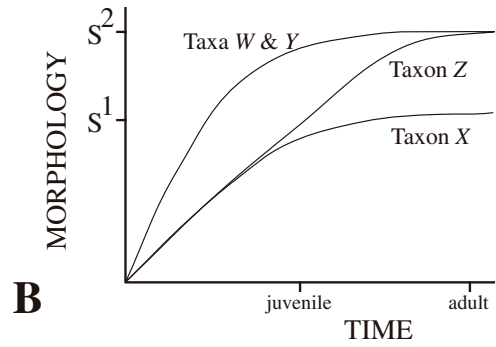


FIGURE 2.22 **A.** Representation of an ontogenetic sequence, a change from one discrete stage to another in various taxa. **B.** Ontogenetic trajectories of various taxa. Note juvenile and adult stages. **C–E.** Cladograms, with ontogenetic data (in parentheses next to taxa) and character state changes of mature structures (along lineage internodes). Note that s represents a juvenile developmental stage; S is a mature, adult feature. See text for further explanation.

REVIEW QUESTIONS

OVERVIEW, TAXON SELECTION, AND CHARACTER ANALYSIS

1. Define phylogeny and give the name of the branching diagram that represents phylogeny.
2. What is phylogenetic systematics and what are its goals?
3. What are the lines of a cladogram called and what do they represent?
4. What does a split, from one lineage to two, represent?
5. Name the term for both a preexisting feature and a new feature.
6. What is the difference between an autapomorphy and a synapomorphy?
7. What names are given to both the group as a whole and the individual component taxa in a cladistic analysis?
8. What precautions must be taken in taxon selection?
9. What criteria are used in the selection and definition of characters and character states?
10. Why and how are characters assessed for character state discreteness?
11. How might characters be correlated, and what should be done in a cladistic analysis if they are?
12. What is homology and how may it be assessed?
13. What is homoplasy?
14. Name and define the two types of homoplasy and give an example of each.
15. What is a transformation series or morphocline?
16. Name, define, and discuss the rationale for the two basic types of transformation series.
17. What is character weighting? Scaling? Why is either done?
18. What is polarity?
19. What is a character step matrix? A character \times taxon matrix?

CLADOGRAM CONSTRUCTION

20. What is meant by recency of common ancestry?
21. What is a monophyletic group?
22. What is the rationale for using synapomorphies in recognizing monophyletic groups?
23. What are sister groups?
24. What is a paraphyletic group?
25. Name a traditional taxonomic plant group that is paraphyletic (refer to Chapters 3–6).
26. What is the principle of parsimony and what is the rationale of this principle?
27. From the data set of Figure 2.7, construct five trees that are different from the one in Figure 2.7E, draw in all character state changes, and calculate the total length of these trees.
28. What is an unrooted tree and what can it not represent?
29. What is a polytomy and how may polytomies arise in cladistic analyses?
30. What is reticulation? How might it be detected?
31. Why do the OTUs of a study need to be verified for monophyly?
32. Why does the whole study group (ingroup) need to be verified for monophyly?
33. What is outgroup comparison and what is the rationale for using it to determine character state polarity?
34. Why should the terms ancestral/plesiomorphic and derived/apomorphic not be applied to taxa?
35. What is a consensus tree?
36. What is long branch attraction and why is it a problem in phylogenetic analysis?
37. Briefly describe the methods of maximum likelihood and Bayesian analysis.
38. What is a consistency index and what does it measure?
39. What are bootstrapping and the decay index and what do they assess?

CLADOGRAM ANALYSIS

40. Describe two ways in which a classification system may be derived from a cladistic analysis.
41. What are the differences between a node-based, apomorphy-based, and stem-based classification system?

42. Give an example as to how a cladistic analysis can be used to assess (a) character evolution; (b) change in ecological habitat; (c) biogeographic history.
43. Name the two major explanations for changes in distribution and indicate how they differ.
44. What is ontogeny and how may ontogeny be measured?
45. Define heterochrony, peramorphosis, paedomorphosis, and neotony.
46. Review the precautions to be taken in a cladistic analysis.
47. For the following data sets: (a) draw the three possible (dichotomously branching) cladograms; (b) for *each* of the three cladograms indicate (with arrows and corresponding characters and states) the minimum character state changes that are needed to explain the data; (c) indicate which of the three trees would be accepted by a cladist as the best estimate of phylogeny and why.

1

	1	2	3	4	5
A	1	1	1	1	1
B	1	0	0	0	0
C	0	0	1	1	1
OUTGROUP	0	0	0	0	0

2

	1	2	3	4	5
A	0	1	1	1	0
B	0	0	1	0	0
C	1	0	1	1	1
OUTGROUP	0	0	0	0	0

48. For each of the following data sets: (a) draw the most parsimonious cladogram; (b) indicate all character state changes; (c) circle all monophyletic groups; (d) derive a hypothetical classification scheme. Assume an ordered transformation series where more than two character states per character occur.

1

GENERA:	1	2	3	4	5	6
	Flower symmetry	Perianth tube	Perianth aestivation	Stamen number	Anther shape	Pollen exine
<i>Aahh</i>	bilateral	present	valvate	6	oblong	homogeneous
<i>Batahr</i>	bilateral	present	valvate	6	oblong	homogeneous
<i>Conarus</i>	radial	present	valvate	6	oblong	homogeneous
<i>Phlebus</i>	radial	absent	imbricate	6	oblong	tectate
<i>Tribus</i>	radial	present	imbricate	6	fringed	homogeneous
OUTGROUP	radial	absent	imbricate	3	oblong	tectate

2

SPECIES:	1	2	3	4	5	6
	Stem type	Carpel number	Pollen sculpturing	Perianth type	Staminode +/-	Leaf vestiture
<i>C. cordatus</i>	rhizome	2	psilate	rotate	-	tomentose
<i>C. ellipticus</i>	corm	2	psilate	rotate	-	glabrous
<i>C. lanecolatus</i>	rhizome	5	spinulose	rotate	-	glabrous
<i>C. ovatus</i>	rhizome	5	psilate	salverform	+	glabrous
<i>C. rhomboideus</i>	rhizome	5	psilate	salverform	+	glabrous
OUTGROUP	rhizome	5	spinulose	rotate	-	glabrous

3

GENERA:	1	2	3	4	5	6	7
	Glu-Ph. allozyme	Pollen aperture no.	Anther dehiscence	Ovary position	Chromosome number	Leaf shape	Calyx merosity
<i>Queesus</i>	B + C	7 8	latrorse	inferior	28	ovate	5
<i>Racamupa</i>	B + C	7 8	latrorse	inferior	14	ovate	4
<i>Shoota</i>	B + C	3	poricidal	inferior	14	lanceolate	5
<i>Tumblus</i>	B + C	3	poricidal	inferior	14	lanceolate	5
<i>Uvulus</i>	A + B	3	latrorse	inferior	7	lanceolate	5
<i>Vertex</i>	A + B	1	latrorse	superior	7	lanceolate	4
OUTGROUP	A	1	latrorse	superior	7	lanceolate	5

EXERCISES

1. **Computer phylogeny applications.**

If computers are available, you may wish to explore one of the commonly used phylogeny software applications, such as MacClade (Maddison and Maddison, 2000; see others cited hereafter). These programs allow the user to input data, including taxa names and their characters and character states, and enable both the phylogenetic relationships of taxa and specific character state changes to be visualized.

With the help of your instructor, enter a data file using MacClade or some other phylogeny application for a given taxonomic group. You may use the data matrix below for the families of the Zingiberales.

Examine the optimal (most parsimonious) tree. Engage the function that displays characters and visualize several, noting the distribution of their states. You may also swap branches on the cladogram, exploring alternative evolutionary hypotheses and noting the change in tree length.

If time allows, choose a volunteer to re-draw the cladogram from MacClade onto the chalkboard. List each apomorphy illustrated on MacClade by placing the derived character state (apomorphy) beside a hatch-mark on the cladogram. Circle and tentatively name all monophyletic groups.

Review as a class the following terms: cladogram, lineage/clade, common ancestor, lineage divergence/diversification, apomorphy, synapomorphy, autapomorphy, monophyletic, paraphyletic.

Example data set of the families of the Zingiberales.

	LEAF ARRANGEMENT	SEED ARIL	POLYARC ROOT	INNER MED. STAMEN	RAPHIDES	SILICA CRYSTALS
Cannaceae	distichous	+	+	+	-	+
Costaceae	distichous	+	+	+	-	+
Heliconiaceae	distichous	+	+	+	+	-
Lowiaceae	distichous	+	-	-	+	-
Marantaceae	distichous	+	+	+	-	+
Musaceae	spiral	-	-	-	+	-
Strelitziaceae	distichous	+	+	-	+	-
Zingiberaceae	distichous	+	+	+	-	+

	STAMEN NUMBER	STAMINODE PETALOID	PERISPERM	OUT. TEPALS FUSED	ANTHER TYPE
Cannaceae	1	+	+	-	monotheical
Costaceae	1	+	+	+	bithecal
Heliconiaceae	5	-	-	-	bithecal
Lowiaceae	5	-	-	-	bithecal
Marantaceae	1	+	+	-	monotheical
Musaceae	5	-	-	-	bithecal
Strelitziaceae	5	-	-	-	bithecal
Zingiberaceae	1	+	+	+	bithecal

2. **Web trees.**

If you have available a computer with a Web browser Internet connection, log onto The Tree of Life (<http://tolweb.org>), TreeBASE (<http://www.treebase.org>), Angiosperm Phylogeny Website (<http://www.mobot.org/MOBOT/research/Apweb>), or a similar Web page. These Web pages contain up-to-date information on the relationships of organismal groups and plants, respectively. Browse through the trees illustrated on the sites and note the source of the data. Examine the apomorphies denoted at the nodes for these trees.

REFERENCES FOR FURTHER STUDY

- Brooks, D. R., and McLennan, D. A. 1991. *Phylogeny, Ecology, and Behavior: A Research Program in Comparative Biology*. Univ. Chicago Press, Chicago.
- Felsenstein, J. 2003. *Inferring Phylogenies*. Sinauer Associates, Sunderland, Massachusetts.
- Gould, S. J. 1977. *Ontogeny and Phylogeny*. Belknap Press of Harvard University, Cambridge, Massachusetts.
- Hennig, W. 1966. *Phylogenetic Systematics*. University of Illinois Press, Urbana.
- Hillis, D. M., C. Moritz, and B. Mable (eds.). 1996. *Molecular Systematics*. Second edition. Sinauer, Sunderland, Massachusetts.
- Huelsenbeck, J. P., and J. P. Bollback. 2001. Empirical and hierarchical Bayesian estimation of ancestral states. *Syst Biol* 50: 351-366.
- Kitching, I. J. 1998. *Cladistics: The Theory and Practice of Parsimony Analysis*, 2nd ed. Oxford University Press, Oxford.
- Li, W. 1997. *Molecular Evolution*. Sinauer Associates, Sunderland, Massachusetts.
- Maddison, W. P., and D. R. Maddison. 2000. *MacClade 4: Analysis of Phylogeny and Character Evolution*. Sinauer Associates, Sunderland, Massachusetts.
- Nei, M. and S. Kumar. 2000. *Molecular Evolution and Phylogenetics*. Oxford University Press, New York.
- Page, R. D., and E. C. Holmes. 1998. *Molecular Evolution: A Phylogenetic Approach*. Blackwell Science, Oxford.
- Semple, C., and M. A. Steel. 2003. *Phylogenetics*. Oxford University Press, Oxford.
- Wiley, E. O., D. Siegel-Causey, D. R. Brooks, and V. A. Funk. 1991. *The Compleat Cladist: A Primer of Phylogenetic Procedures*. Univ. Kansas Museum Nat. History Sp. Publ. no. 19.

CLADISTIC COMPUTER PROGRAMS

- Felsenstein, J. 1993. PHYLIP (Phylogeny Inference Package) version 3.5c. Distributed by the author. Department of Genetics, University of Washington, Seattle. [Mac & Windows OS]
- Goloboff, P. 1993. Nona. Software and documentation by the author, Tucuman, Argentina. [Windows OS]
- Huelsenbeck, J. P., and F. Ronquist. 2001. MR-BAYES: Bayesian inference of phylogeny. *Bioinformatics* 17: 754-755. Version 3. <http://morphbank.ebc.uu.se/mrbayes3> [Mac, Unix, and Windows OS]
- Maddison, W. P., and D. R. Maddison. 2000. *MacClade 4: Analysis of phylogeny and character evolution*. Sinauer, Sunderland, Massachusetts. [Mac OS; Windows OS compatible with an emulator]
- Nixon, K. C. 1999. WinClada. Software and documentation by the author. Cornell University, Ithaca, New York. [Windows OS]
- Swofford, D. L. 2003. PAUP*. *Phylogenetic Analysis Using Parsimony (*and Other Methods)*. Version 4. Sinauer Associates, Sunderland, Massachusetts. [Mac, Unix/VMS, DOS, and Windows OS]

II

EVOLUTION AND DIVERSITY OF PLANTS



This Page Intentionally Left Blank

3

EVOLUTION AND DIVERSITY OF GREEN AND LAND PLANTS

THE GREEN PLANTS	51	Mosses	63
THE LAND PLANTS	54	Polysporangiophytes	66
DIVERSITY OF NONVASCULAR LAND PLANTS	59	REVIEW QUESTIONS	66
Liverworts	59	EXERCISES	67
Hornworts	59	REFERENCES FOR FURTHER STUDY	67

THE GREEN PLANTS

The green plants, or Chlorobionta, are a monophyletic group of eukaryotic organisms that includes what have traditionally been called green algae plus the land plants or embryophytes (Figure 3.1). Like all eukaryotes, the Chlorobionta have cells with membrane-bound organelles, including a nucleus (containing chromosomes composed of linear chains of DNA bound to proteins, that are sorted during cell division by mitosis), microtubules, mitochondria, an endoplasmic reticulum, vesicles, and golgi bodies. Although the interrelationships of the non land plant Chlorobionta will not be covered in detail here, it is important to realize that some of the evolutionary innovations, or apomorphies, that we normally associate with land plants actually arose before plants colonized the land.

Several apomorphies unite the Chlorobionta (Figure 3.1). One possible novelty for this group is a **cellulosic cell wall** (Figure 3.2A). Cellulose, like starch, is a polysaccharide, but one in which the glucose sugar units are bonded in the beta-1,4 position (=β-1,4-glucopyranoside). This slight change in chemical bond position results in a very different molecule. Cellulose is secreted outside the plasma membrane as microscopic fiber-like units called **microfibrils** that are further intertwined into larger fibril units, forming a supportive meshwork. The function of cellulose is to impart rigidity to

the cells, acting as a sort of cellular exoskeleton. The evolution of a cellulosic cell wall was a preamble to the further evolution of more complex types of growth, particularly of self-supporting shoot systems. It is not clear if a cellulosic cell wall constitutes an apomorphy for the Chlorobionta alone, as it may have evolved much earlier, constituting an apomorphy for the Chlorobionta plus one or more other groups; in any case, its adaptive significance seems clear.

Perhaps the primary apomorphy for the Chlorobionta is a specialized type of **chloroplast** (Figure 3.2). As discussed in Chapter 1, chloroplasts are one of the major defining characteristics of traditionally defined plants; their adaptive significance as organelles functioning in photosynthesis, the conversion of light energy to chemical energy, is unquestioned. Chloroplasts in the Chlorobionta, the green plants, differ from those of most other organisms, such as the red and brown algae, in (1) containing **chlorophyll b** in addition to chlorophyll a, the former of which acts as an accessory pigment in light capture; (2) having **thylakoids**, the chlorophyll-containing membranes that are **stacked into grana**, which are pancake-like aggregations (see Figure 3.2B); and (3) manufacturing as a storage product true **starch**, a polymer of glucose sugar units (= *polysaccharide*) in which the glucose molecules are chemically bonded in the alpha-1,4 position (α-1,4-glucopyranoside). Thus, all green plants, from filamentous green algae in a pond or tide pool to giant sequoia or *Eucalyptus* trees

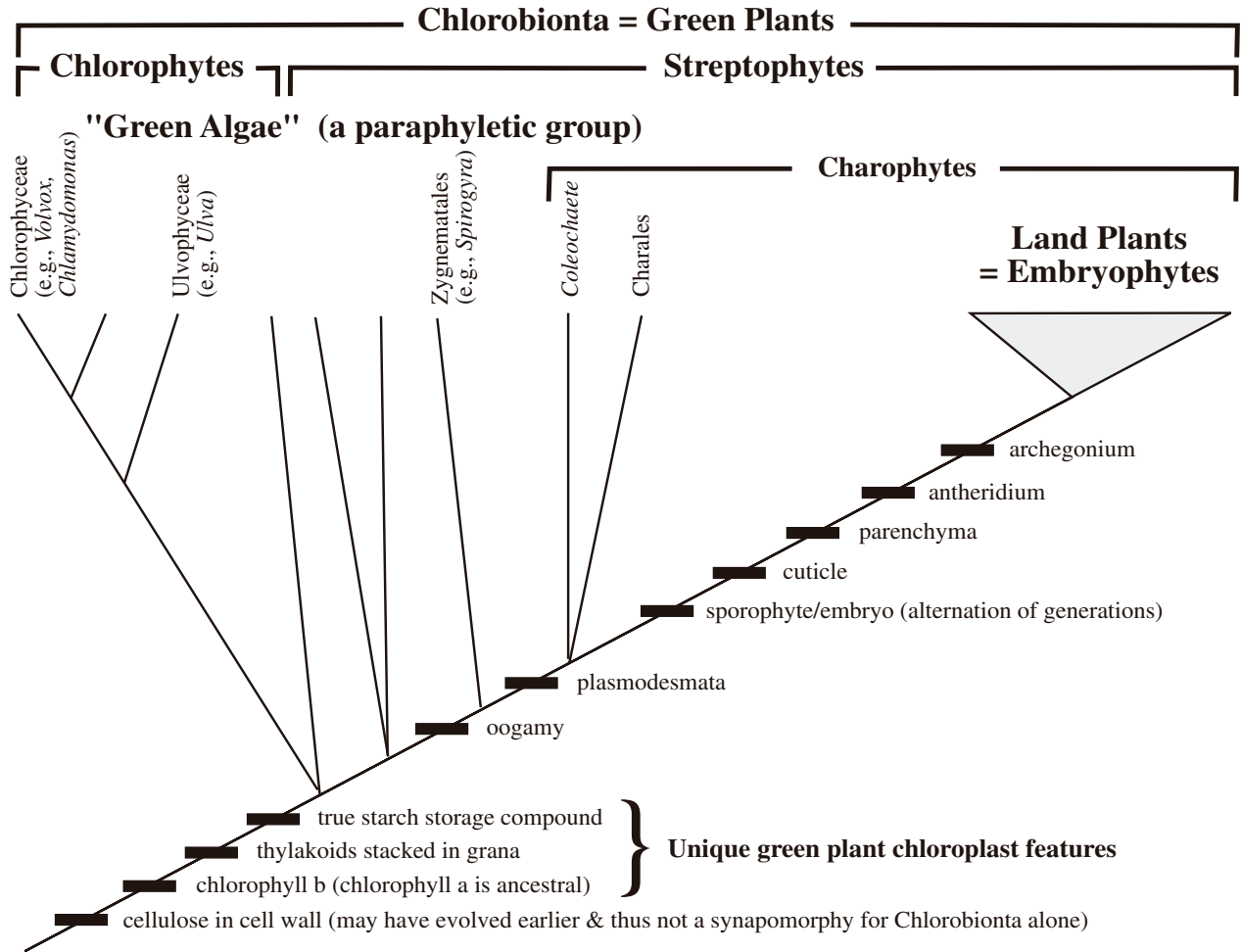


FIGURE 3.1 Cladogram of the green plants (Chlorobionta), modified from Bremer (1985), Mishler and Churchill (1985), and Mishler et al. (1994). Important apomorphies discussed in the text are listed beside thick hash marks.

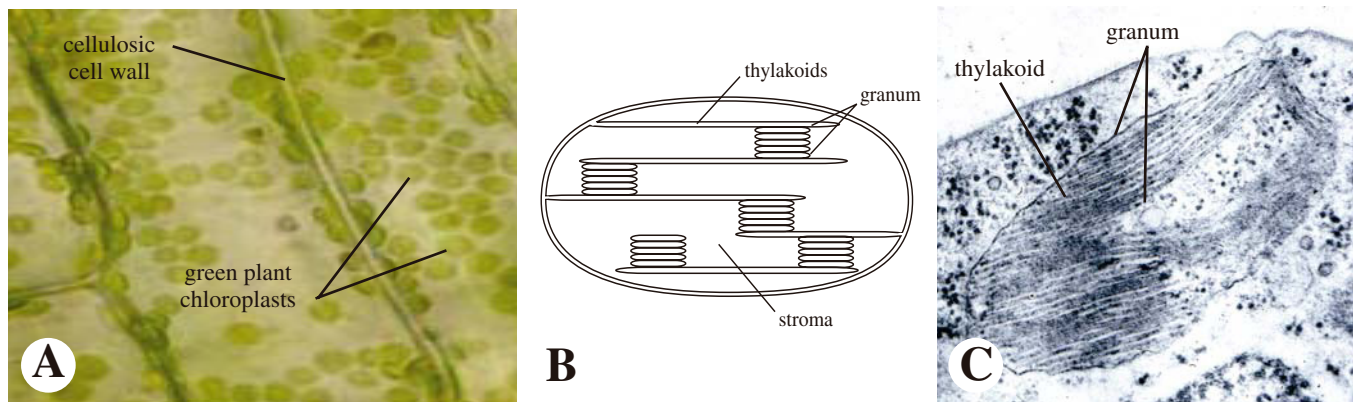


FIGURE 3.2 **A.** *Elodea*, whole leaf in face view, showing apomorphies of the Chlorobionta: a cellulosic cell wall and green plant chloroplasts. **B.** Diagram of chloroplast structure of green plants, showing thylakoids and grana. **C.** Electron micrograph of *Chlamydomonas reinhardtii*, a unicellular green alga, showing granum of chloroplast. (Photo courtesy of Rick Bizzoco.)

have this same type of chloroplast. Recent data imply that chloroplasts found in the green plants today were modified from those that evolved via **endosymbiosis**, the intracellular cohabitation of an independently living, unicellular prokaryote inside a eukaryotic cell (see Chapter 1).

The Chlorobionta as a whole are classified as two sister groups: chlorophytes, or Chlorophyceae, and streptophytes, or Streptophyceae (Figure 3.1). The traditional green algae are a paraphyletic group (which is why the name is placed in quotation marks) and are defined as the primarily aquatic Chlorobionta, consisting of all chlorophytes and the non land plant streptophytes. Green algae occur in a tremendous variety of morphological forms. These include flagellated unicells (Figure 3.3A) with or without flagella, thalloid forms (Figure 3.3B), motile and nonmotile colonies (Figure 3.3C), and nonmotile filaments (Figure 3.3D). Many have flagellated motile cells in at least one phase of their life history. Green algae inhabit fresh and marine waters and some live in or on soil (or even on snow!) or in other terrestrial but moist habitats.

The primitive type of green plant sexual reproduction seems to have been the production of flagellate, haploid (n) gametes that are isomorphic, that is, that look identical. Fertilization occurs by union of two of these gametes, resulting in a diploid ($2n$) **zygote** (Figure 3.4A). The zygote, which is free-living, then divides by meiosis to form four haploid **spores**, each of which may germinate and develop into a new haploid individual, which produces more gametes, completing what is termed a **haplontic** (or haplobiontic) life cycle (Figure 3.4A).

Within the streptophyte lineage that gave rise to the land plants, a few innovations evolved that may have been preadaptations to survival on land. First of these was the

evolution of **oogamy**, a type of sexual reproduction in which one gamete, the **egg**, becomes larger and nonflagellate; the other gamete is, by default, called a **sperm cell** (Figure 3.4B). Oogamy is found in all land plants and independently evolved in many other groups, including many other algae and in the animals. Two other evolutionary novelties that occurred prior to the evolution of land plants were retention of the egg and retention of the zygote on the parent body (Figure 3.1). Retention of the egg and zygote was adaptive, at least in part, by making possible the future *nutritional dependence* of the zygote upon the haploid plant, ultimately leading to the sporophyte (see later discussion).

Several other apomorphies of and within the Chlorobionta include ultrastructural specializations of flagella and some features of biochemistry. Although these have been valuable in elucidating phylogenetic relationships, their adaptive significance is unclear, and they will not be considered further here.

An apomorphy for the Charophytes, a clade within the streptophytes that includes the *Coleochaete* (Figure 3.6A), Charales (Figure 3.6B–D), and the land plants (Figure 3.1), are **plasmodesmata**. Plasmodesmata are essentially pores in the primary (1°) cell wall through which membranes traverse between cells, allowing for transfer of compounds between cells (Figure 3.5). Plasmodesmata may function in more efficient or rapid transport of solutes, including regulatory and growth-mediating compounds, such as hormones.

Members of the Charales, such as the genera *Chara* and *Nitella*, are perhaps the closest living relatives to the land plants. These aquatic organisms form whorls of lateral branches (Figure 3.6B) and grow by means of a single apical cell, resembling that of some land plants (but differing in lacking true **parenchyma**; see later discussion). The Charales

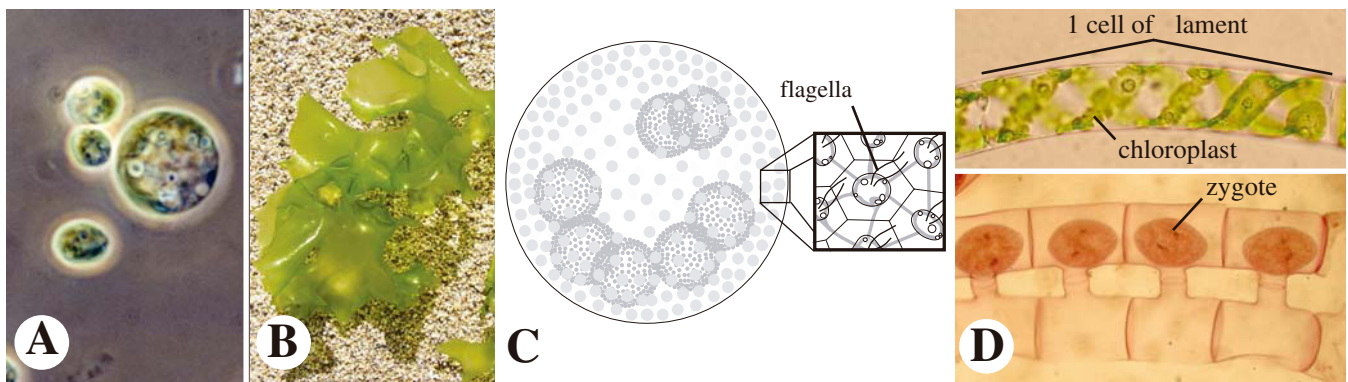


FIGURE 3.3 Examples of non land plant Chlorobionta. **A.** *Chlamydomonas reinhardtii*, a unicellular form. (Photo courtesy of Rick Bizzoco.) **B.** *Ulva*, a thalloid form. **C.** *Volvox*, a colonial form. **D.** *Spirogyra*, a filamentous form. *Above:* vegetative form, with large, spiral chloroplasts. *Below:* reproductive conjugation stage, showing + and – mating strains and nonmotile zygotes.

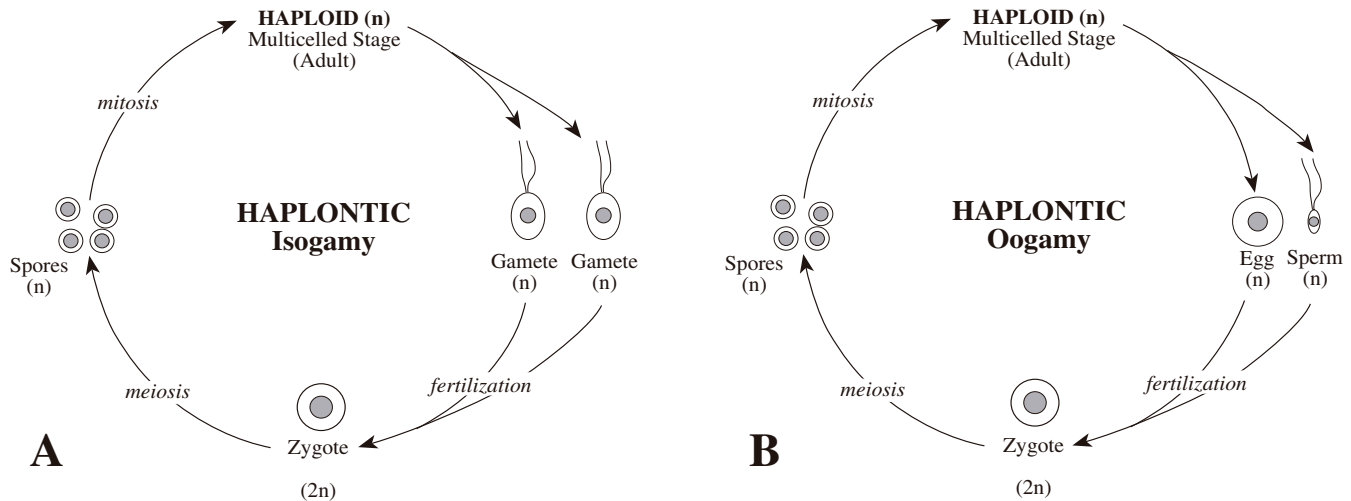


FIGURE 3.4 Haplontic life cycles in some of the green plants. **A.** Isogamy. **B.** Oogamy.

have specialized male and female gametangia, termed antheridia and oogonia (Figure 3.6C,D). The oogonia are distinctive in having a spirally arranged group of outer cells (Figure 3.6D); these have been found in the fossil record. Oogonia and antheridia of the Charales resemble the archegonia and antheridia of land plants (see later discussion) in having an outer layer of protective cells, but have been thought not to be directly homologous because of differences in structure. However, these specialized gametangia, as well as the apical cell growth, may represent a transition to what is seen in the land plants.

THE LAND PLANTS

The land plants, or embryophytes (also known as Embryophyta), are a monophyletic assemblage within the green plants

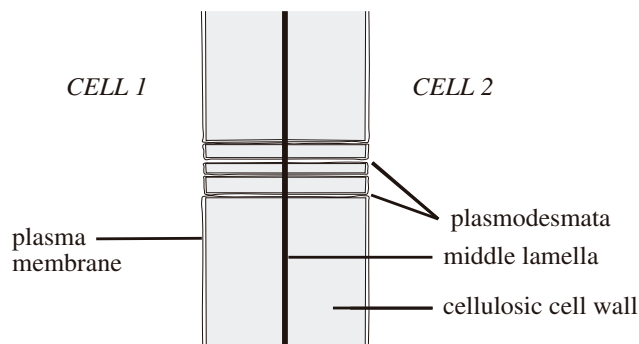


FIGURE 3.5 Diagram of plasmodesmata in cellulose cell wall, an apomorphy of some green plants, including the land plants.

(Figures 3.1, 3.7). The first colonization of plants on land during the Silurian period, ca. 400 million years ago, was concomitant with the evolution of several important features. These shared, evolutionary novelties (Figure 3.7A,B) constituted major adaptations that enabled formerly aquatic green plants to survive and reproduce in the absence of a surrounding water medium.

One major innovation of land plants was the evolution of the **embryo** and **sporophyte** (Figure 3.8). The sporophyte is a separate diploid (2n) phase in the life cycle of all land plants. The corresponding haploid, gamete-producing part of the life cycle is the **gametophyte**. The life cycle of land plants, having both a haploid gametophyte and a diploid sporophyte, is an example of a **haplodiplontic** (also called **diplobiontic**) life cycle, commonly called **alternation of generations** (Figure 3.8). Note that alternation of generations does not necessarily mean that the two phases occur at different points in time; at any given time, *both* phases may occur in a population.

The sporophyte can be viewed as forming from the zygote by the delay of meiosis and spore production. Instead of meiosis, the zygote undergoes numerous *mitotic* divisions, which result in the development of a separate entity. The **embryo** is defined as an immature sporophyte that is attached to or surrounded by the gametophyte. In many land plants, such as the seed plants, the embryo will remain dormant for a period of time and will begin growth only after the proper environmental conditions are met. As the embryo grows into a mature sporophyte, a portion of the sporophyte differentiates as the spore-producing region. This spore-producing region of the sporophyte is called the **sporangium**. A sporangium contains

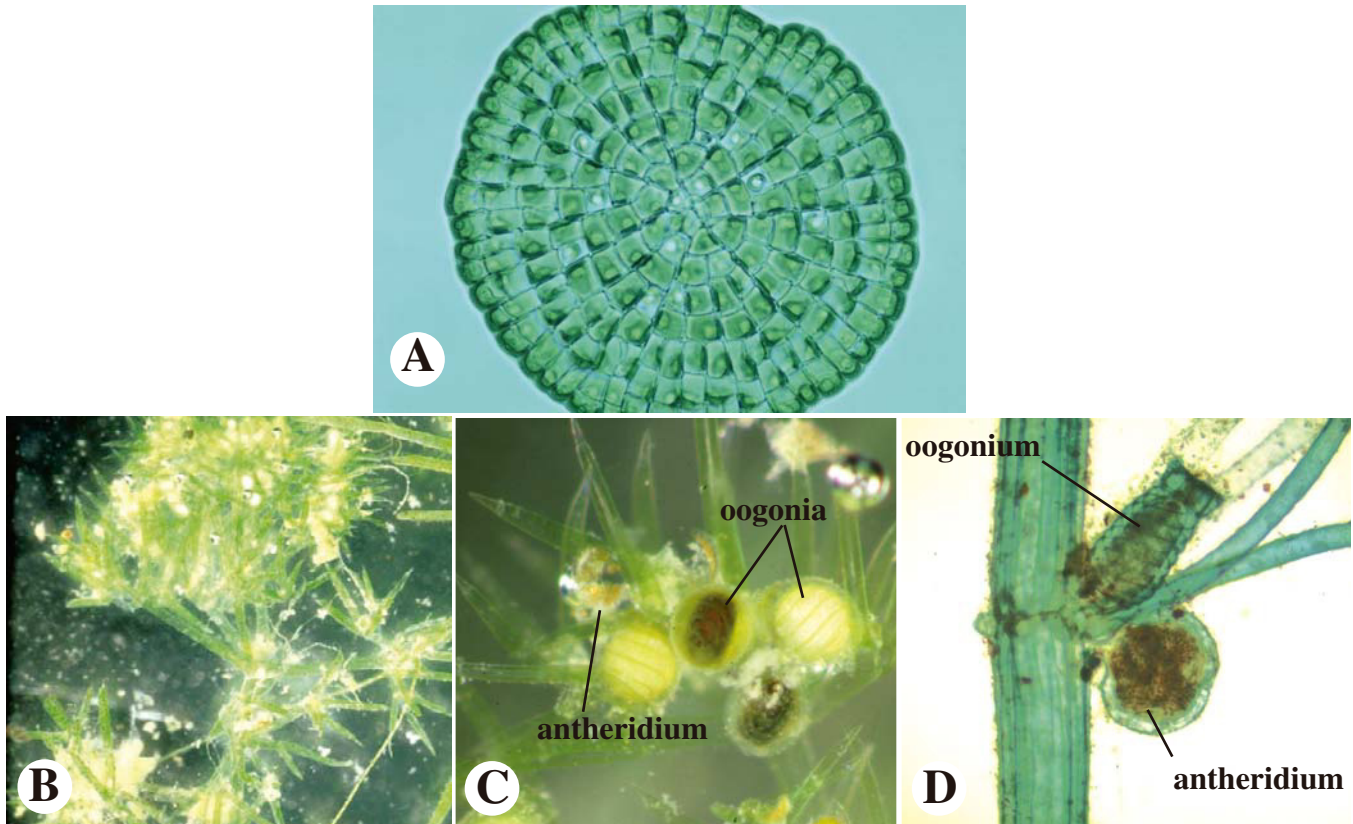


FIGURE 3.6 A. *Coleochaete* sp., a close relative to the embryophytes. (Photo courtesy of Linda Graham.) B–D. Charales. B,C. *Nitella* sp. B. Whole plant. C. Oogonia and antheridia. D. *Chara* sp., oogonium and antheridium. Note spiral wall of oogonia.

sporogenous tissue, which matures into **sporocytes**, the cells that undergo meiosis. Each sporocyte produces, by meiosis, four haploid **spores**. The sporangium is enveloped by a **sporangial wall**, which consists of one or more layers of sterile, non-spore-producing cells.

One adaptive advantage of a sporophyte generation as a separate phase of the life cycle is the large increase in spore production. In the absence of a sporophyte, a single zygote (the result of fertilization of egg and sperm) will produce four spores. The elaboration of the zygote into a sporophyte and sporangium can result in the production of literally millions of spores, a potentially tremendous advantage in reproductive output and increased genetic variation.

Another possible adaptive value of the sporophyte is associated with its diploid ploidy level. The fact that a sporophyte has two copies of each gene may give this diploid phase an increased fitness in either of two ways: (1) by potentially preventing the expression of recessive, deleterious alleles (which, in the sporophyte, may be shielded by dominant alleles, but which, in the gametophyte, would always be expressed); and (2) by permitting increased genetic variability in the

sporophyte generation (via genetic recombination from two parents) upon which natural selection acts, thus increasing the potential for evolutionary change.

A second innovation in land plant was the evolution of **cutin** and the **cuticle** (Figure 3.9). A **cuticle** is a protective layer that is secreted to the outside of the cells of the **epidermis** (Gr. *epi*, upon + *derma*, skin), the outermost layer of land plant organs. The epidermis functions to provide mechanical protection of inner tissue and to inhibit water loss. The cuticle consists of a thin, homogeneous, transparent layer of **cutin**, a polymer of fatty acids, and functions as a sealant, preventing excess water loss. Cutin also impregnates the outer cellulosic cell walls of epidermal cells; these are known as a cutinized cell wall. The adaptive advantage of cutin and the cuticle is obvious: prevention of desiccation outside the ancestral water medium. In fact, plants that are adapted to very dry environments will often have a particularly thick cuticle (as in Figure 3.9) to inhibit water loss.

A third apomorphy for the land plants was the evolution of parenchyma tissue (Figure 3.10). All land plants grow by means of rapid cell divisions at the apex of the stem, shoot,

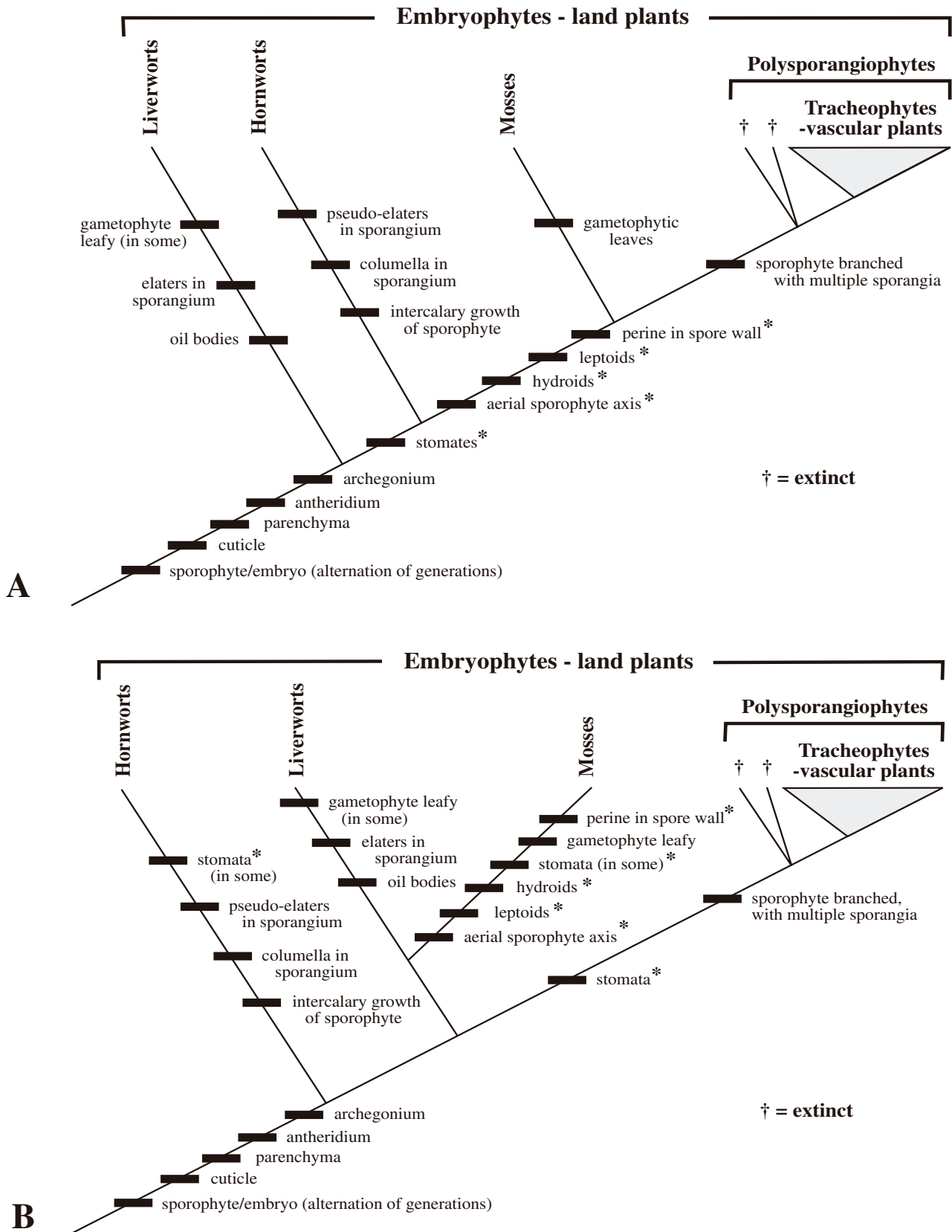


FIGURE 3.7 Two alternative cladograms of the land plants (Embryophyta), with major apomorphies indicated, those with an * varying between the two cladograms. **A.** Traditional topology, based on morphology (e.g., Bremer, 1985; Mishler and Churchill, 1985; Mishler et al., 1994). **B.** Recent molecular study (e.g., Nickrent et al., 2000; Renzaglia et al., 2000).

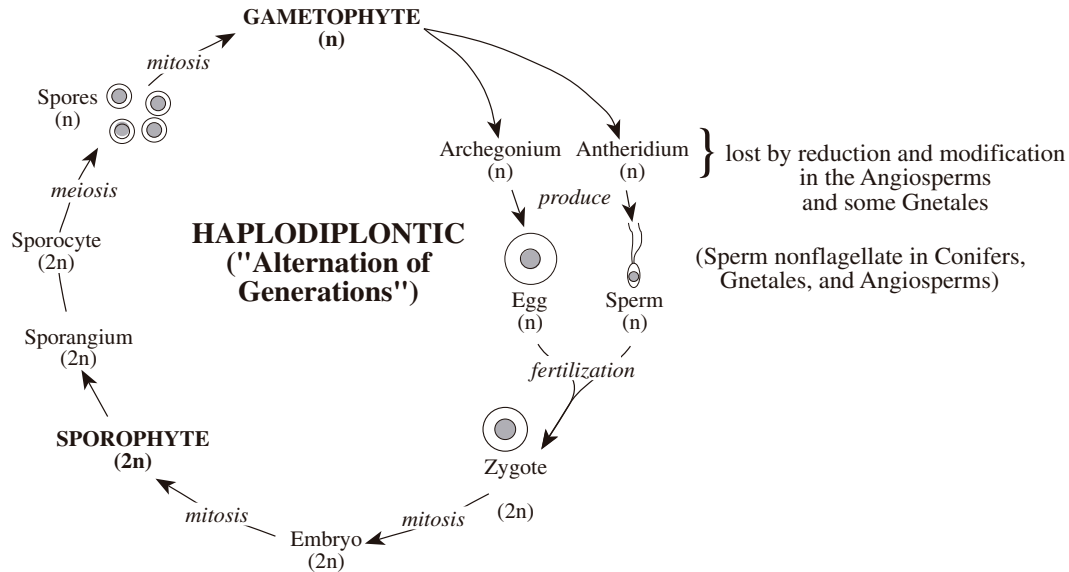


FIGURE 3.8 Haplodiplontic alternation of generations in the land plants (Embryophytes).

and thallus or (in most vascular plants) of the root. This region of actively dividing cells is the **apical meristem**. The apical meristem of liverworts, hornworts, and mosses (discussed later), and of the monilophytes (see Chapter 4) have a single apical cell (Figure 3.10), the ancestral condition for the land plants. In all land plants the cells derived from the apical meristem region form a solid mass of tissue known as **parenchyma** (Gr. *para*, beside + *enchyma*, an infusion; in reference to a concept that parenchyma infuses or fills up space beside and between the other cells). Parenchyma tissue consists of cells that most resemble the unspecialized, undifferentiated cells of actively dividing meristematic tissue. Structurally, parenchyma cells are (1) elongate to isodiametric; (2) have a primary (1°) cell wall only (rarely a secondary wall); and (3) are living at maturity and potentially capable of

continued cell divisions. Parenchyma cells function in metabolic activities such as respiration, photosynthesis, lateral transport, storage, and regeneration/wound healing. Parenchyma cells may further differentiate into other specialized cell types. It is not clear if the evolution of both apical growth and true parenchyma is an apomorphy for the land plants alone, as shown here (Figure 3.7). Both may be interpreted to occur

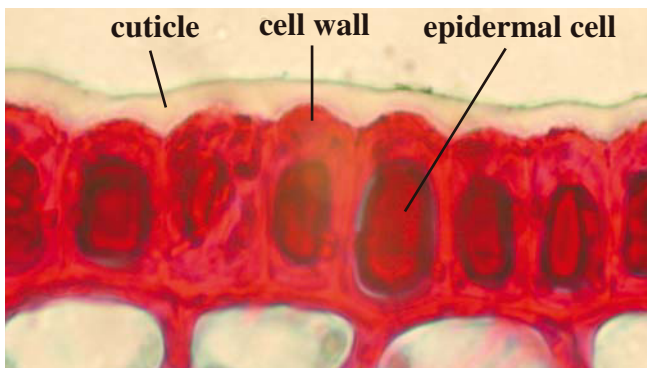


FIGURE 3.9 The cuticle, an apomorphy for the land plants.

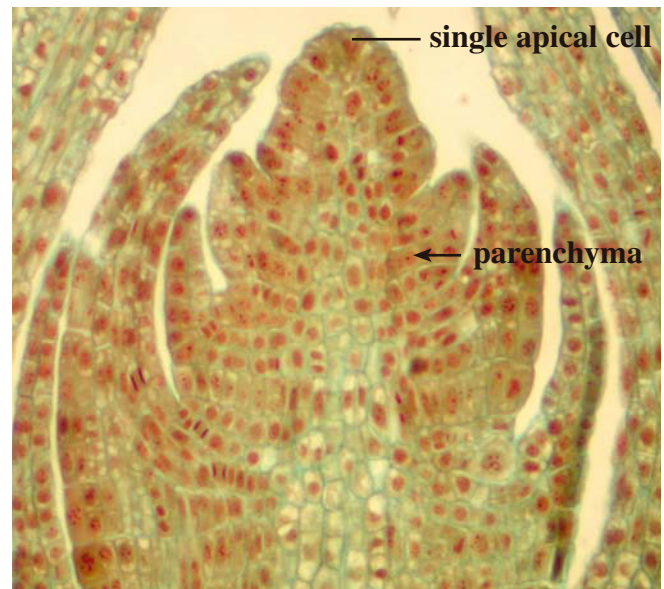


FIGURE 3.10 *Equisetum* shoot apex, showing parenchymatous growth form, from an apical meristem.

in certain closely related green plants, including the Charales.

Correlated with the evolution of parenchyma may have been the evolution of a middle lamella in land plants. The middle lamella is a pectic-rich layer that develops between the primary cell walls of adjacent cells (Figure 3.5). Its function is to bind adjacent cells together, perhaps a prerequisite to the evolution of solid masses of parenchyma tissue.

Another evolutionary innovation for the land plants was the **antheridium** (Figure 3.11A). The antheridium is a type of specialized gametangium of the haploid (n) gametophyte, one that contains the sperm-producing cells. It is distinguished from similar structures in the Chlorobionta in being surrounded by a layer of sterile cells, the antheridial wall. The evolution of the surrounding layer of sterile wall cells, which is often called a sterile jacket layer, was probably adaptive in protecting the developing sperm cells from desiccation. In all of the nonseed land plants, the sperm cells are released from the antheridium into the external environment and must swim to the egg in a thin film of water. Thus, a wet environment is needed for fertilization to be effected in the nonseed plants, a vestige of their aquatic ancestry. Members of the Charales also have a structure termed an antheridium, which has an outer layer of sterile cells (Figure 3.6C,D). However, because of its differing anatomy, the Charales antheridium may not be homologous with that of the land plants, and thus may have evolved independently.

Another land plant innovation was the evolution of the **archegonium**, a specialized female gametangium (Figure 3.11B). The archegonium consists of an outer layer of sterile cells, termed the **venter**, that immediately surround the **egg** plus others that extend outward as a tube-like **neck**. The archegonium is stalked in some taxa; in others the egg is rather deeply embedded in the parent gametophyte. The egg cell is located inside and at the base of the archegonium. Immediately above the egg is a second cell, called the **ventral canal cell**, and above this and within the neck region, there may be several **neck canal cells**. The archegonium may have several adaptive functions. It may serve to protect the developing egg. It may also function in fertilization. Before fertilization occurs, the neck canal cells and ventral canal cell break down and are secreted from the terminal pore of the neck itself; the chemical compounds released function as an attractant, acting as a homing device for the swimming sperm. Sperm cells enter the neck of the archegonium and fertilize the egg cell to form a diploid ($2n$) zygote. In addition to effecting fertilization, the archegonium serves as a site for embryo/sporophyte development and the establishment of a nutritional dependence of the sporophyte upon gametophytic tissue.

The land plants share other possible apomorphies: the presence of various ultrastructural modifications of the sperm cells, flavonoid chemical compounds, and a proliferation of heat shock proteins. These are not discussed here.

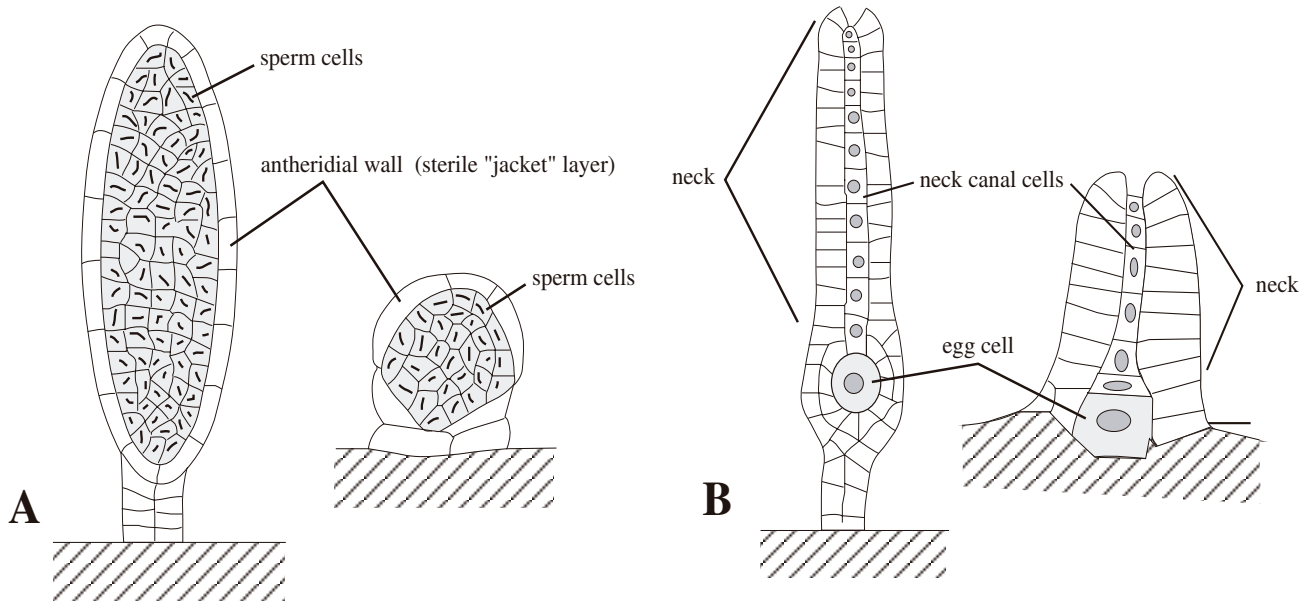


FIGURE 3.11 A. Antheridia. B. Archegonia. Both are apomorphies of land plants.

DIVERSITY OF NONVASCULAR LAND PLANTS

During the early evolution of land plants, three major, monophyletic lineages diverged *before* the vascular plants (Chapter 4). These lineages may collectively be called the nonvascular land plants or bryophytes and include the liverworts, hornworts, and mosses. Bryophytes are a paraphyletic group, defined by the absence of derived features; the name, placed in quotation marks, is no longer formally recognized.

Liverworts, hornworts, and mosses differ from the vascular plants in lacking true vascular tissue and in having the gametophyte as the dominant, photosynthetic, persistent, and free-living phase of the life cycle; it is likely that the ancestral gametophyte of the land plants was thalloid in nature, similar to that of the hornworts and many liverworts. The sporophyte of the liverworts, hornworts, and mosses is relatively small, ephemeral, and attached to and nutritionally dependent upon the gametophyte (see later discussion).

The relationships of the liverworts, hornworts, and mosses to one another and to the vascular plants remain unclear. Many different relationships among the three lineages have been proposed, two of which are seen in Figure 3.7A and B. Note, when considering differing phylogenetic relationships, that the position of apomorphies may shift.

LIVERWORTS

Liverworts, also traditionally called the **Hepaticae**, are one of the monophyletic groups that are descendents of some of the first land plants. Today, liverworts are relatively minor components of the land plant flora, growing mostly in moist, shaded areas (although some are adapted to periodically dry, hot habitats). Among the apomorphies of liverworts are (1) distinctive **oil bodies** and (2) specialized structures called **elaters**, elongate, nonsporogenous cells with spiral wall thickenings, found inside the sporangium. Elaters are hygroscopic, meaning that they change shape and move in response to changes in moisture content. Elaters function in spore dispersal; as the sporangium dries out, the elaters twist out of the capsule, carrying spores with them (Figures 3.12, 3.13H).

There are two basic morphological types of liverwort gametophytes: thalloid and leafy (Figures 3.12, 3.13). **Thalloid** liverworts consist of a **thallus**, a flattened mass of tissue; this is likely the ancestral form, based on cladistic studies. As in hornworts and mosses, the gametophyte bears **rhizoids**, uniseriate, filamentous processes that function in anchorage and absorption. **Pores** in the upper surface of the thallus function in gas exchange (Figure 3.13I). These pores are not true

stomata (discussed later), as they have no regulating guard cells. Some liverworts, like the hornworts (discussed next), have a symbiotic relationship with Cyanobacteria. On the upper surface of the gametophytes of some thalloid liverworts, such as *Marchantia*, are specialized structures called **gemma cups**, which contain propagules called **gemmae**. These structures function in vegetative (asexual) reproduction; when a droplet of water falls into the gemma cup, the gemmae themselves may be dispersed some distance away, growing into a haploid genetic clone of the parent.

Leafy liverworts have gametophytes consisting of a stem axis bearing three rows of thin leaves. In most leafy liverworts, the stem is prostrate and the leaves are modified such that the upper two rows of leaves are larger and the lowermost row (on the stem underside) are reduced (Figures 3.12, 3.13K). Other leafy liverworts are more erect, with the three rows of leaves similar. The leaves of leafy liverworts evolved independently from those of mosses (discussed later) or vascular plants (Chapter 4).

As in all of the early diverging land plant lineages, liverworts have antheridia and archegonia that develop on the gametophyte. In some liverwort taxa (e.g., *Marchantia*), the gametangia form as part of stalked, peltate structures: **antheridiophores** bearing antheridia and **archegoniophores** bearing archegonia (Figures 3.12, 3.13). Sperm released from an antheridium of the antheridiophore swims in a film of water to the archegonia of the archegoniophore, effecting fertilization.

After fertilization the zygote divides mitotically and eventually differentiates into a diploid (2n) embryo, which matures into the diploid (2n) sporophyte. This sporophyte is relatively small, nonphotosynthetic, and short lived. It consists almost entirely of a **sporangium** or **capsule** (Figure 3.13G). At a certain stage, the internal cells of the capsule divide meiotically, forming haploid (n) spores (see Figure 3.8). In liverworts the spores are released by a splitting of the capsule into four valves. The spores may land on a substrate, germinate (under the right conditions), and grow into a new gametophyte, completing the life cycle.

HORNWORTS

The hornworts, or **Anthocerotae**, are a monophyletic group comprising a second major lineage of land plants. Some hornworts have **stomates** (also termed **stomata**), specialized epidermal cells generally found on leaves, but sometimes on stems. Stomata consist of two chloroplast-containing cells, the **guard cells**, which, by changes in turgor pressure, can increase or decrease the size of the opening between them, the **stoma** (Figure 3.14). Each guard cell has one or more ridge-like deposits on the side facing the stoma (Figure 3.14).

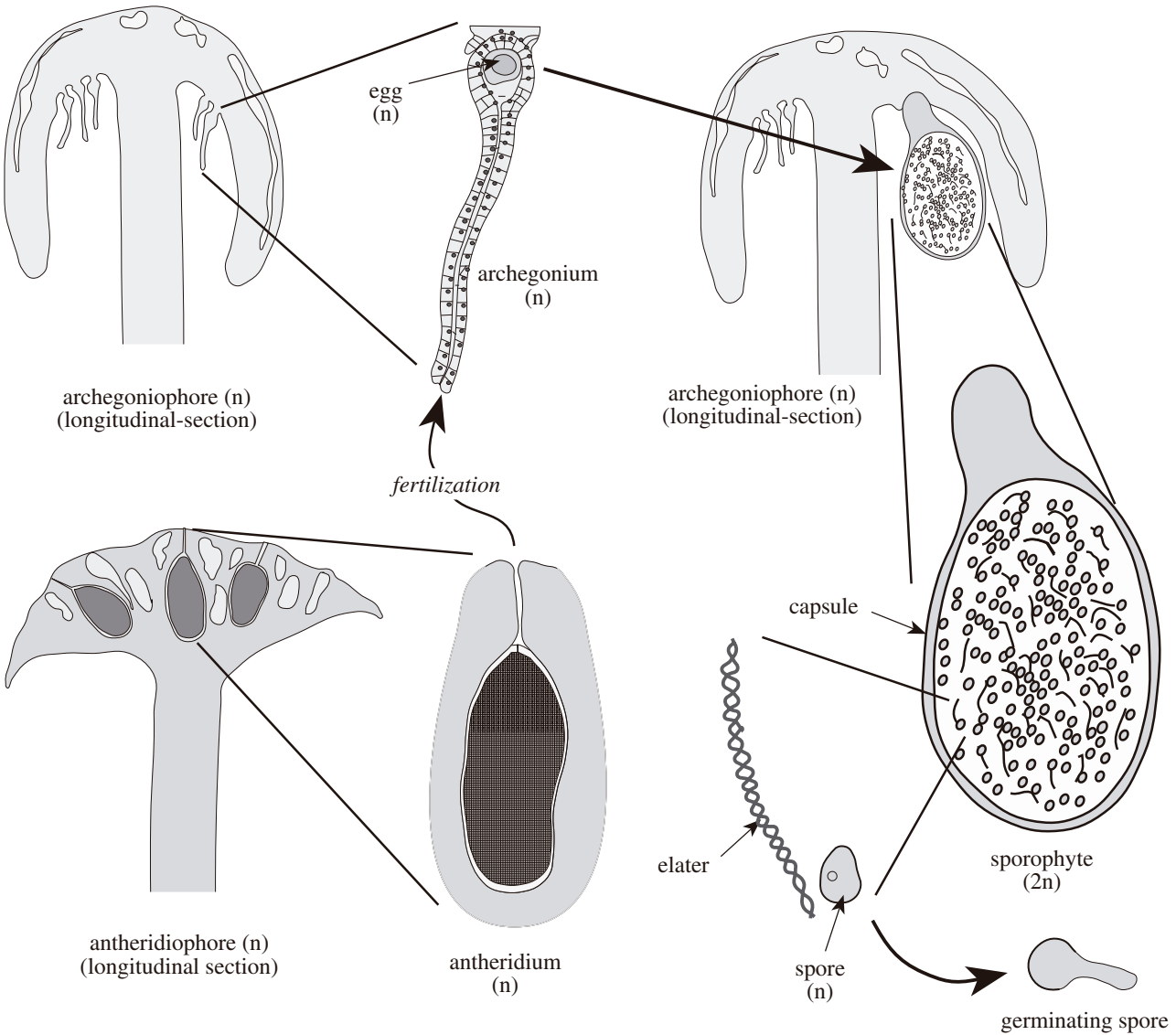
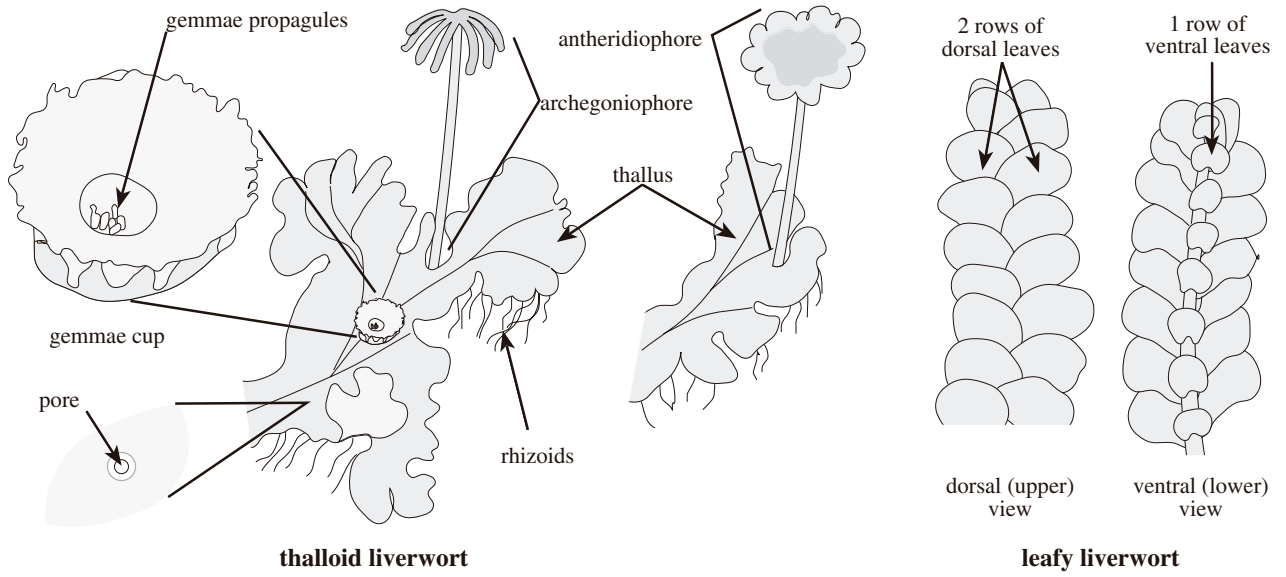


FIGURE 3.12 Liverwort morphology and life cycle.

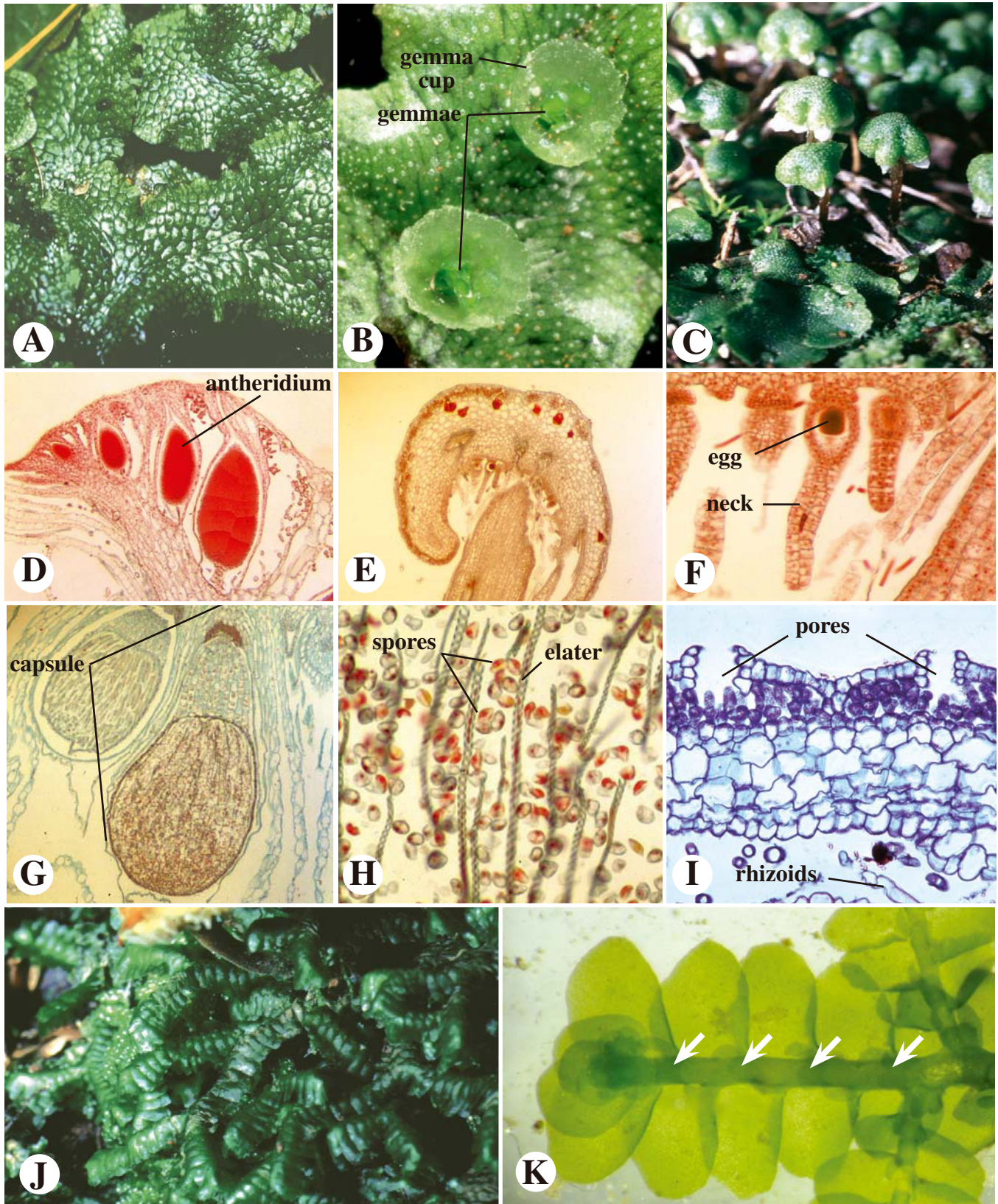


FIGURE 3.13 Liverworts. **A.** *Conocephalum* sp., a thalloid liverwort. **B.** *Marchantia*, thallus with gemma cups and gemmae. Note whitish pores. **C.** *Asterella*, a thalloid liverwort with archegoniophores. **D–I.** *Marchantia*. **D.** Anteridiophore, longitudinal section. **E.** Archegoniophore, longitudinal section. **F.** Archegonium. **G.** Capsule, longitudinal section, showing sporogenous tissue. **H.** Close-up, sporogenous tissue, showing spores and elaters. **I.** Cross-section of thallus, showing rhizoids and upper pores. **J.** *Bazania trilobata*, a leafy liverwort. **K.** *Porella*, a leafy liverwort, showing third row of reduced leaves at arrows (lower side facing).

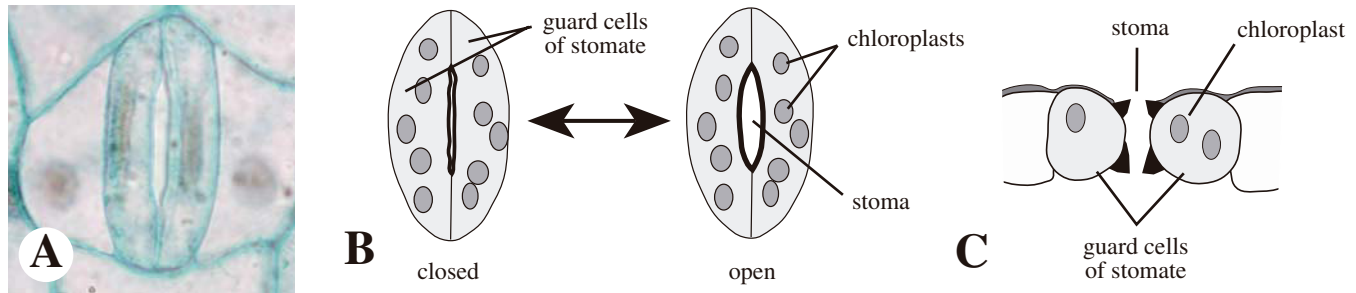


FIGURE 3.14 The stomate, an innovation for mosses, hornworts, and vascular plants. **A.** Face view, slightly open. **B.** Diagram, face view, open and closed. **C.** Diagram, cross-section.

This material, which is rich in suberin, a waxy, water-resistant substance, functions to better seal the stomata. Stomata function in regulation of gas exchange, in terms of both photosynthesis and water uptake. Carbon dioxide passing through the stoma diffuses to the chloroplasts of photosynthetic cells within and is used in the dark reactions of photosynthesis. Oxygen, a by-product of photosynthesis, exits via the stoma. Stomata also allow water vapor to escape from the leaf. In most plants stomata open during the day when photosynthesis takes place; thus, heat from the sun may cause considerable water loss through stomata. In some plants, loss of water via stomata is simply a by-product, a price to be paid for entry of carbon dioxide, which is essential for photosynthesis.

However, in other plants, such as tall trees, stomatal water loss may actually be adaptive and functional, as a large quantity of water must flow through the leaves in order to supply sufficient quantities of mineral nutrients absorbed via the roots.

By one hypothesis, stomates represent an apomorphy for all land plants except for the liverworts (Figure 3.7A); by another hypothesis, stomata evolved independently in the vascular plants and in lineages within the hornworts and mosses (Figure 3.7B).

Hornworts are similar to the thalloid liverworts in gametophyte morphology (Figure 3.15) and are found in similar habitats. Hornworts differ from liverworts, however, in lacking

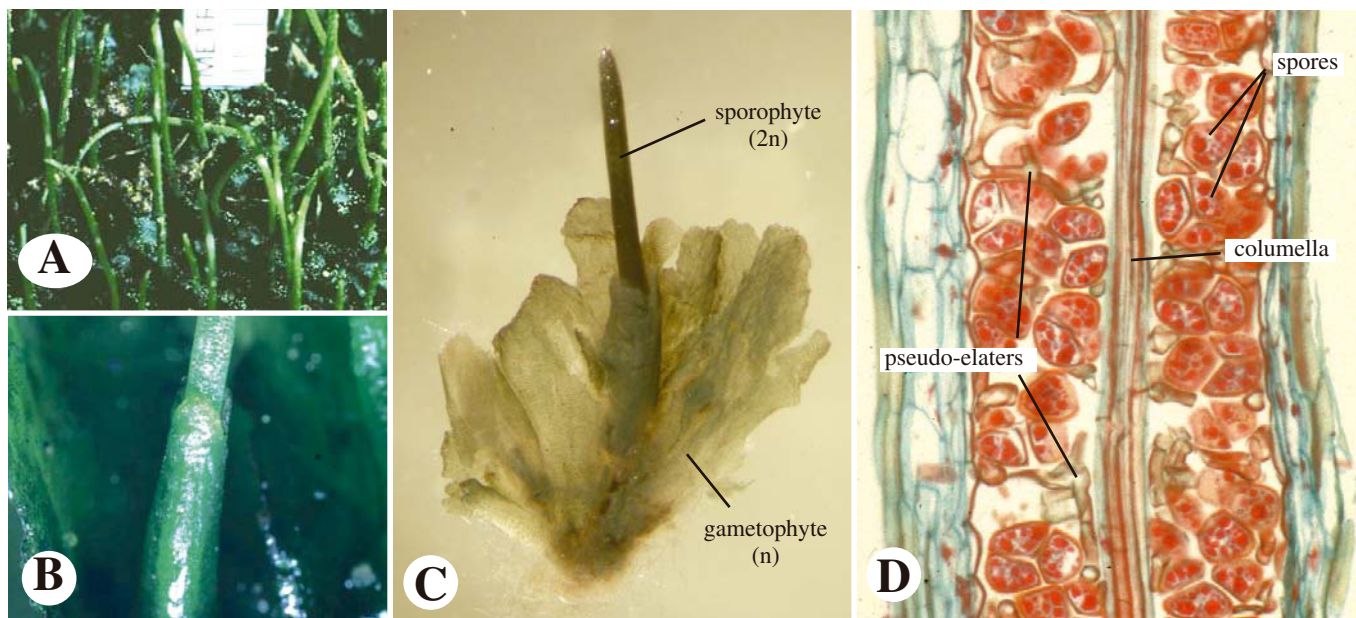


FIGURE 3.15 Hornworts, *Anthoceros* sp. **A.** Population of gametophytes with attached sporophytes. **B.** Close-up of sporophyte base, showing ensheathing collar of gametophytic tissue surrounding intercalary meristem of sporophyte. **C.** Gametophyte with attached, cylindrical sporophyte. **D.** Sporophyte longitudinal section, showing columella, spores, and pseudo-elaters.

pores (having stomates in some species, as discussed above). All hornworts have a symbiotic relationship with Cyanobacteria (blue-greens), which live inside cavities of the thallus. This relationship is found in a few thalloid liverworts as well (probably evolving independently), but not in mosses. Interestingly, hornworts and liverworts may also have a symbiotic association between the gametophytes and a fungus, similar to the mycorrhizal association with the roots of vascular plants.

The basic life cycle of hornworts is similar to that of liverworts and mosses. However, the sporophyte of hornworts is unique in being elongate, cylindrical, and photosynthetic (Figure 3.15A,C). This cylindrical sporophyte has indeterminate (potentially continuous) growth, via a basal, **intercalary meristem** (Figure 3.15B). The intercalary meristem is a region of actively dividing cells near the base of the sporophyte (just above the point of attachment to the gametophyte), constituting an apomorphy for the hornworts. Other apomorphies include a unique central column of sterile (non-spore-producing) tissue called a **columella** and the production of specialized structures in the sporangium called **pseudo-elaters**, groups of cohering, nonsporogenous, elongate, generally hygroscopic cells, which are nonhomologous with but have a similar function to the elaters of liverworts (Figure 3.15D).

MOSESSES

The mosses, or **Musci**, are by far the most speciose and diverse of the three major groups of nonvascular land plants and inhabit a number of ecological niches. Mosses may share some apomorphies with the vascular plants. One of these is an elongate, **aerial sporophyte axis**, an apomorphy for the mosses alone (Figure 3.7B) or a possible precursor to the evolution of the sporophytic stem in vascular plants (Figure 3.7A). Some mosses have specialized conductive cells called **hydroids**, which function in water conduction, and **leptoids**, which function in sugar conduction. These cells resemble typical xylem tracheary elements and phloem sieve elements (Chapter 4), but lack the specializations of the latter cell types. They may represent intermediates in the evolution of true vascular tissue (Figure 3.7A) or evolved independently of vascular tissue (Figure 3.7B). The spores of mosses have a thick outer layer called a **perine layer** (Figure 3.16), which may be apomorphic for the mosses alone (Figure 3.7B) or for the mosses and vascular plants combined (Figure 3.7A). The perine layer may function in preventing excess desiccation and provide additional mechanical protection of the spore cytoplasm. As with liverworts and hornworts, a three-lined structure, called a **trilete mark**, develops on the spore wall; the trilete mark is the scar of attachment of the

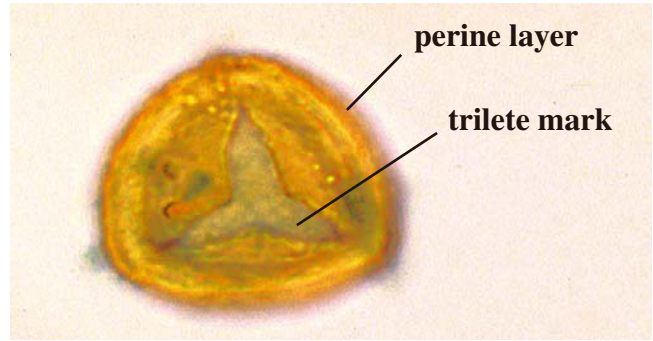


FIGURE 3.16 Moss spore. Note protective perine layer and trilete mark.

adjacent three spores of the four spores produced at meiosis (Figure 3.16).

Moss gametophytes are always leafy, with a variable number of leaf ranks or rows (Figures 3.17, 3.18B). The leaves of mosses are thought to have evolved independently from those in liverworts and, thus, constitute an apomorphy for the mosses alone. Moss leaves are mostly quite small and thin, but may have a central **costa**, composed of conductive cells, that resembles a true vein (Figure 3.17). Antheridia and archegonia in mosses are usually produced at the apex of gametophytic stems (Figures 3.17, 3.18C,D,E). After fertilization, the sporophyte grows upward (Figures 3.17, 3.18F) and often carries the apical portion of the original archegonium, which continues to grow. This apical archegonial tissue, known as a **calyptra** (Figures 3.17, 3.18G), may function in protecting the young sporophyte apex. The sporophyte generally develops a long stalk, known as a **stipe**, at the apex of which is born the sporangium or capsule (Figures 3.17, 3.18F,G). The capsule of most mosses has a specialized mechanism of dehiscence. At the time of spore release, a lid known as an **operculum** falls off the capsule apex, revealing a whorl of **peristome teeth**. The peristome teeth, like the elaters of liverworts, are hygroscopic. As the capsule dries up, the peristome teeth retract, effecting release of the spores (Figures 3.17, 3.18G,H).

Under the right environmental conditions, moss spores will germinate and begin to grow into a new gametophyte. The initial development of the gametophyte results in the formation of filamentous structure, known as a **protonema** (Figure 3.18A). The protonema probably represents an ancestral vestige, resembling a filamentous green alga. After a period of growth, the protonema grows into a parenchymatous gametophyte.

One economically important moss worth mentioning is the genus *Sphagnum*, or peat moss, containing numerous species. *Sphagnum* grow in wet bogs and chemically modifies its

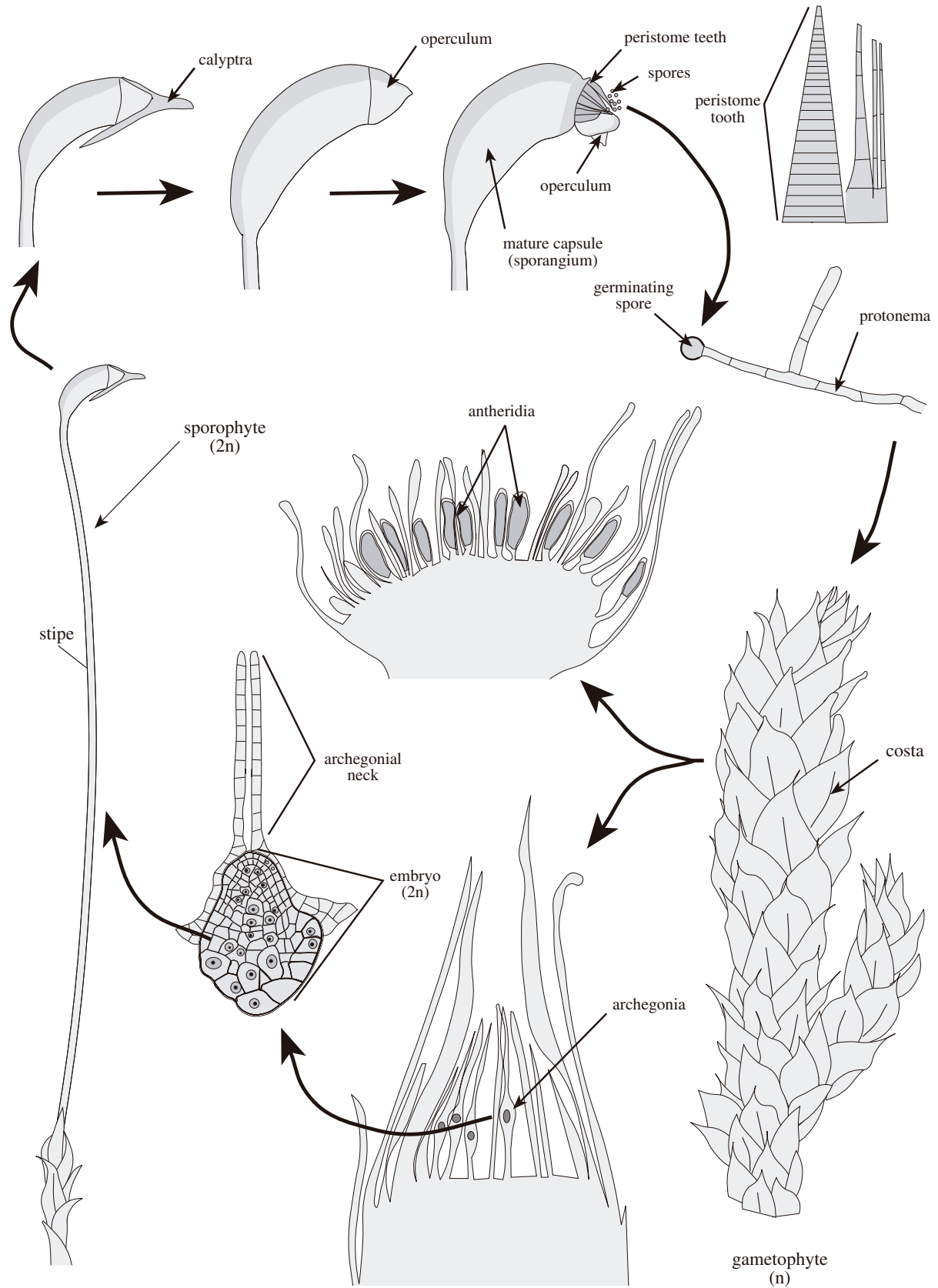


FIGURE 3.17 Moss morphology and life cycle.

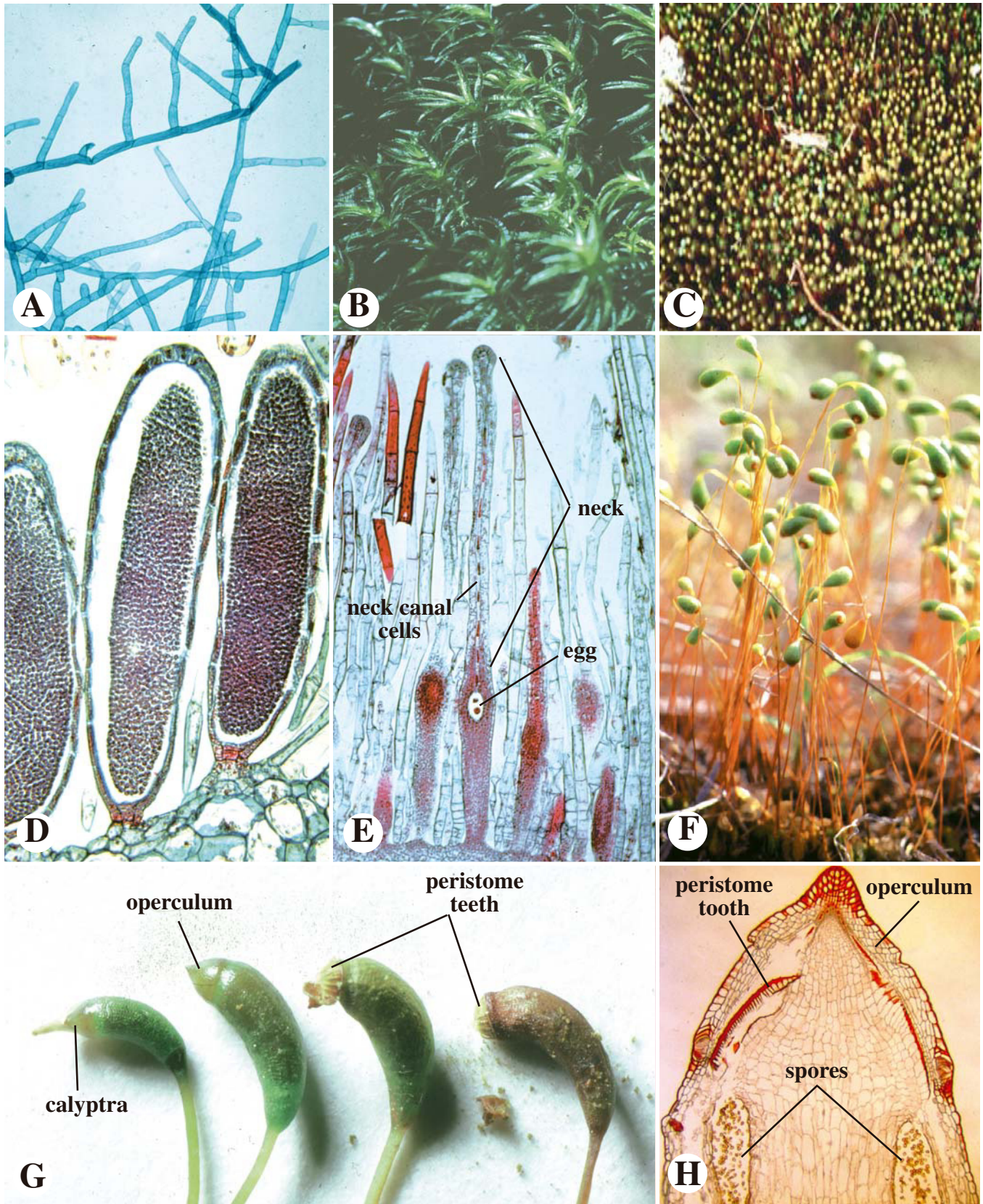


FIGURE 3.18 Mosses. **A.** Protonema of *Sphagnum*. **B.** *Atrichum* sp. gametophyte. **C.** *Polytrichum* sp. gametophyte, face view, showing antheridia at tips of branches. **D,E.** *Mnium* sp. **D.** Antheridia, longitudinal section, showing external capsule wall (sterile layer) and internal sporogenous tissue. **E.** Archegonia, showing stalk, egg cell, neck, and neck canal cells. **F.** Sporophytes of moss, showing capsules. **G.** Moss sporophyte close-up, showing developmental series (left to right). **H.** *Mnium*, capsule (sporangium) longitudinal section, showing operculum, one of several peristome teeth, and spores within sporangium.

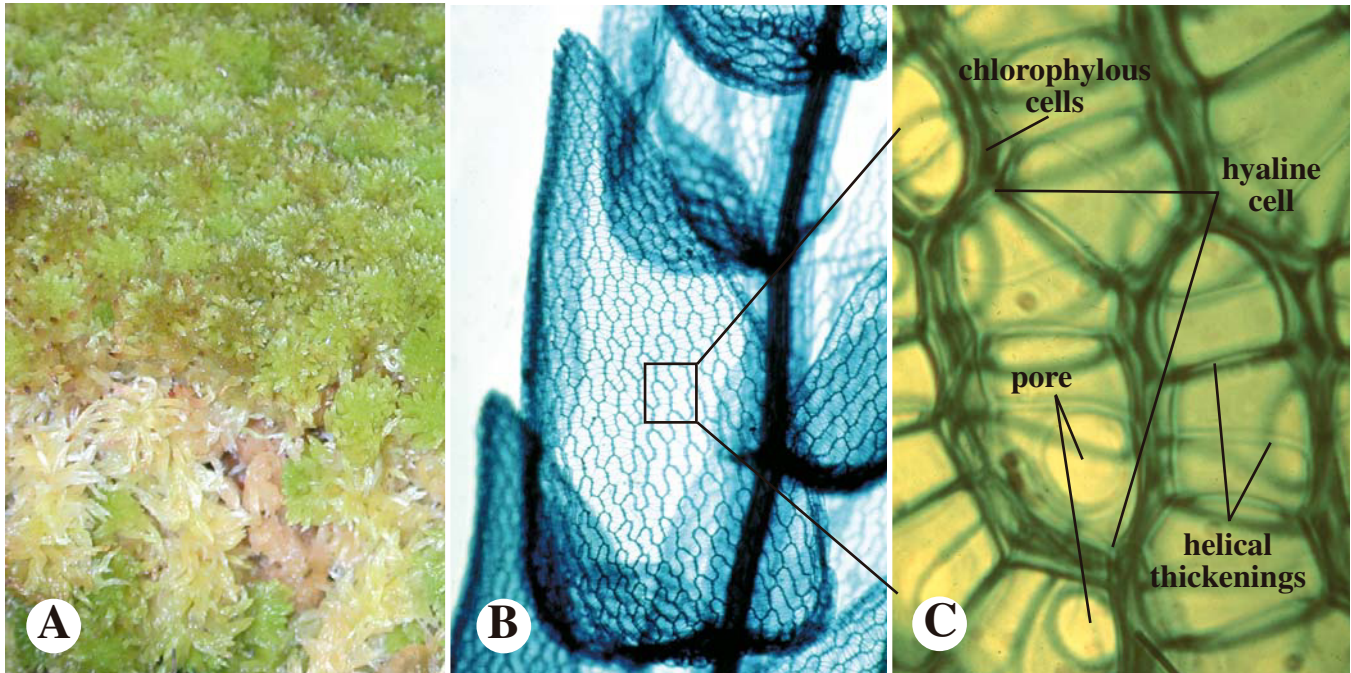


FIGURE 3.19 *Sphagnum*, or peat moss. **A.** Clonal population. **B.** Individual leaf at center, showing the specialized chlorophyllous and hyaline cells. **C.** Leaf close-up, showing chlorophyllous cells, hyaline cells, pores, and spiral wall thickenings of hyaline cells.

environment by making the surrounding water acidic. The leaves of *Sphagnum* are unusual in having two cell types: **chlorophyllous cells**, which form a network, and large, clear **hyaline cells**, having characteristic pores and helical thickenings (Figure 3.19). The pores of the hyaline cells give *Sphagnum* remarkable properties of water absorption and retention, making it quite valuable horticulturally in potting mixtures. **Peat** is fossilized and partially decomposed *Sphagnum* and is mined for use in potting mixtures and as an important fuel source in parts of the world.

POLYSPORANGIOPHYTES

This group is inclusive of a few, basal fossil taxa plus all of the true vascular plants, or tracheophytes (Chapter 4). The basal (first-evolving) polysporangiophytes, such as the genus *Horneophyton* (not illustrated), were similar to hornworts, liverworts, and mosses in lacking vascular tissue. However, they are different from bryophytes, and linked to the vascular plants, in having branched stems with multiple sporangia. Thus, the polysporangiophytes include taxa that were transitional to the evolution of tracheophytes.

REVIEW QUESTIONS

GREEN PLANTS

1. What is a formal name for the green plants?
2. What are the unique features of green plant chloroplasts?
3. How are chloroplasts thought to have originated?
4. The bulk of the primary cell wall of green plants is composed of what substance? (Give the common name and chemical name.)
5. Is the cell wall synthesized inside or outside the plasma membrane?
6. What are plasmodesmata?
7. What is a haplontic life cycle?
8. What is oogamy?

LAND PLANTS

9. What is the formal name for the land plants?
10. Name the major apomorphies of the land plants.
11. Draw and label the basic haplodiplontic life cycle (alternation of generations) of all land plants, illustrating all structures, processes, and ploidy levels.
12. What is an embryo?
13. What is a sporangium?
14. Name the possible adaptive features of the sporophyte.
15. What are cutin and cuticle and what are their adaptive significance?
16. Define apical growth and parenchyma.
17. In land plants what is the name of the pectic-rich layer between adjacent cell walls that functions to bind them together?
18. What is an antheridium? Draw.
19. What is an archegonium? Draw.

HORNWORTS, LIVERWORTS, AND MOSSES

20. Draw two, different phylogenetic trees denoting relationships of the mosses, liverworts, hornworts, and vascular plants.
21. Name two apomorphies of the liverworts.
22. What are the two major morphological forms of liverworts? Which is likely ancestral?
23. What are gemmae and gemma cups?
24. What is an antheridiophore? an archegoniophore?
25. What land plant groups possess stomates?
26. Describe the structural makeup and function of a stomate.
27. How do the hornworts differ from the liverworts?
28. Name major apomorphies either shared by the mosses alone or possibly shared by the mosses plus vascular plants.
29. What is a calyptra, stipe, operculum, peristome tooth?
30. What is the scientific name of peat moss?
31. What feature of the leaf anatomy of peat moss enables the leaves to absorb and retain water?
32. How is peat moss of economic importance?
33. What apomorphy links the Polysporangiates with the vascular plants?

EXERCISES

1. Peruse the most recent literature on phylogenetic relationships of the green algae relative to the land plants. Are there any differences relative to Figure 3.1?
2. Peruse the recent literature on phylogenetic relationships of the hornworts, liverworts, and mosses. Do any show relationships different from that of Figure 3.7?
3. Peruse botanical journals and find a systematic article on a moss, liverwort, or hornwort. What is the objective of the article and what techniques were used to address it?
4. Collect and identify local liverworts, hornworts, and mosses. What features are used to distinguish among families, genera, and species?

REFERENCES FOR FURTHER STUDY

- Bremer, K. re. 1985. Summary of green plant phylogeny and classification. *Cladistics* 1(4): 369-385.
- Graham, Linda. 1985. The origin of the life cycle of land plants. *American Scientist* 73: 178-186.
- Kernick, P., and P. R. Crane. 1997. The origin and early diversification of land plants: a cladistic study. Smithsonian Institution Press, Washington, DC.
- Mishler, Brent D., and Steven P. Churchill. 1984. A cladistic approach to the phylogeny of the Bryophytes. *Brittonia* 36(4): 406-424.
- Mishler, Brent D., and Steven P. Churchill. 1985. Transition to a land flora: phylogenetic relationships of the green algae and bryophytes. *Cladistics* 1(4): 305-328.

- Mishler, Brent D., Louise A. Lewis, Mark A. Buchheim, Karen S. Renzaglia, David J. Garbary, Charles F. Delwiche, Frederick W. Zechman, Thomas S. Kantz, and Russell L. Chapman. 1994. Phylogenetic relationships of the Green Algae and Bryophytes. *Annals of the Missouri Botanical Garden* 81: 451-483.
- Nickrent, D. L., C. L. Parkinson, J. D. Palmer, and R. J. Duff. 2000. Multigene phylogeny of land plants with special reference to bryophytes and the earliest land plants. *Molecular Biology and Evolution* 17: 1885-1895.
- Renzaglia, K. S., R. J. Duff, D. L. Nickrent, and D. J. Garbary. 2000. Vegetative and reproductive innovations of early land plants: implications for a unified phylogeny. *Philosophical Transactions: Biological Sciences* 355: 769-793.

4

EVOLUTION AND DIVERSITY OF VASCULAR PLANTS

VASCULAR PLANT APOMORPHIES	69	Lycopodiophyta Lycophytes	78
Lignified Secondary Cell Walls	69	Euphyllophytes	82
Sclerenchyma	70	Monilophytes Ferns	82
Tracheary Elements (of Xylem)	71	Ophioglossales Ophioglossoid Ferns	83
Sieve Elements (of Phloem)	72	Psilotales Whisk Ferns	84
Endodermis	73	Equisetales Horsetails	85
Independent, Long-Lived Sporophyte	74	Marattiales Marattioid Ferns	87
Sporophytic Leaves and Shoot	75	Polypodiales Leptosporangiate Ferns	87
Roots	76	REVIEW QUESTIONS	92
VASCULAR PLANT DIVERSITY	77	EXERCISES	94
Rhyniophyta Rhyniophytes	78	REFERENCES FOR FURTHER STUDY	94

VASCULAR PLANT APOMORPHIES

The vascular plants, or Tracheophyta (also called tracheophytes), are a monophyletic subgroup of the land plants. The major lineages of tracheophytes (excluding many fossil groups) are seen in Figure 4.1. Vascular plants together share a number of apomorphies, including (1) **lignified secondary walls**, with pits, in certain specialized cells; (2) **sclerenchyma**, specialized cells that function in structural support; (3) **tracheary elements**, cells of **xylem** tissue; (4) **sieve elements**, cells of **phloem** tissue (the xylem and phloem comprising the vascular tissue); (5) an **endodermis**; and (6) an **independent, long-lived sporophyte**. In addition, all extant vascular plants, and all except for the earliest fossil lineages such as rhyniophytes (discussed later), possess two other apomorphies: (7) **sporophytic leaves**, which are associated with the stem in a **shoot system**; and (8) **roots** (secondarily lost in the psilophytes; see later discussion).

LIGNIFIED SECONDARY CELL WALLS

Vascular plants have evolved a chemical known as **lignin**, which is a complex polymer of phenolic compounds. Lignin is

incorporated into an additional cell wall layer, known as the **secondary (2°) wall** (Figure 4.2), which is found in certain, specialized cells of vascular plants. Secondary walls are secreted to the outside of the plasma membrane (between the plasma membrane and the primary cell wall) *after* the primary wall has been secreted, which is also after the cell ceases to elongate. Secondary cell walls are usually much thicker than primary walls and, like primary walls, contain cellulose. However, in secondary walls, lignin is secreted into the space between the cellulose microfibrils, forming a sort of interbinding cement. Thus, lignin imparts significant strength and rigidity to the cell wall.

In virtually all plant cells with secondary, lignified cell walls, there are holes in the secondary wall called **pits** (Figure 4.2). Pits commonly occur in pairs opposite the sites of numerous plasmodesmata in the primary cell wall. This group of plasmodesmata is called a **primary pit field**. Pits function in allowing communication, via the plasmodesmata of the primary pit field, between cells during their development and differentiation. They may also have specialized functions in water conducting cells (discussed later). Plant cells with secondary walls include **sclerenchyma** and **tracheary elements** (see later discussion).

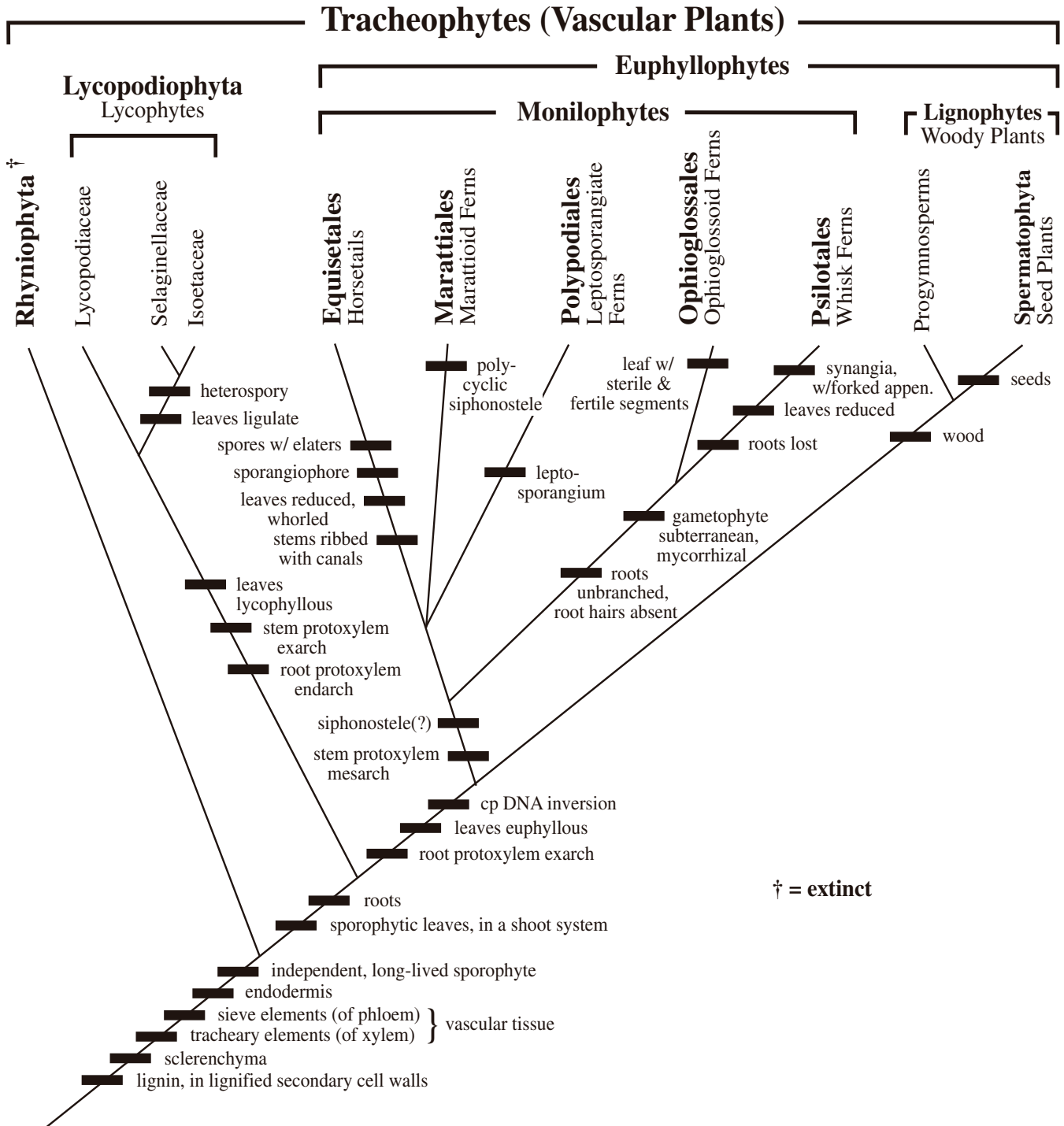


FIGURE 4.1 Phylogeny of the tracheophytes, the vascular plants, modified from Pryer et al. (2001, 2004a).

SCLERENCHYMA

Sclerenchyma (Gr. *scleros*, hard + *enchyma*, infusion, in reference to the infusion of lignin in the secondary cell walls) are nonconductive cells that have a thick, lignified secondary cell wall, typically with pits, and that are dead at maturity.

There are two types of sclerenchyma (Figure 4.3): (1) **fibers**, which are long, very narrow cells with sharply tapering end walls; and (2) **scleireids**, which are isodiametric to irregular or branched in shape. Fibers function in mechanical support in various organs and tissues, sometimes making up the bulk

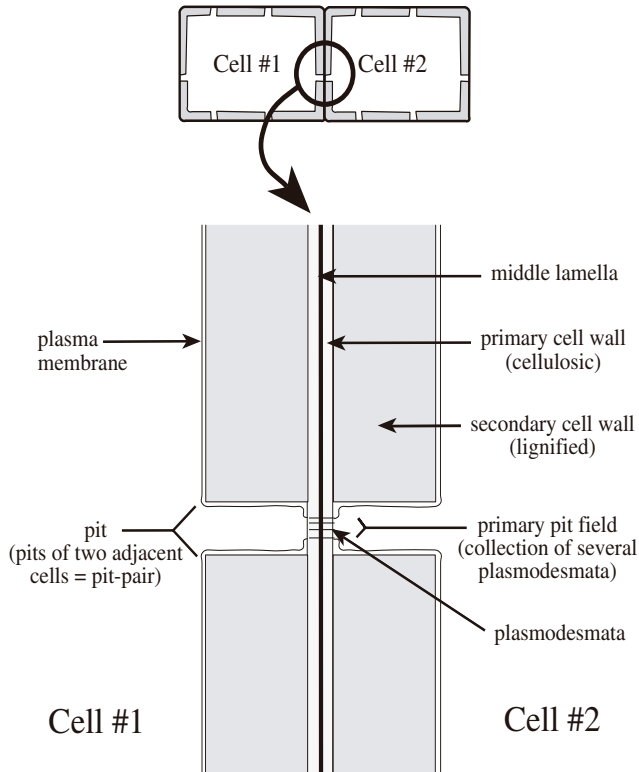


FIGURE 4.2 Lignified secondary cell wall of specialized cells of vascular plants. Note pit-pair, adjacent to primary pit field.

of the tissue. Fibers often occur in groups or bundles. They may be components of the xylem and/or phloem or may occur independently of vascular tissue. Sclereids may also function in structural support, but their role in some plant organs is unclear; they may possibly aid to deter herbivory in some plants. The evolution of sclerenchyma (especially fibers), with lignified secondary cell walls, constituted a major plant adaptation, permitting the structural support needed to attain greater stem height.

Another tissue type that functions in structural support is **collenchyma**, consisting of live cells with unevenly thickened, pectic-rich, primary cell walls (see Chapter 10). Collenchyma is found in many vascular plants, but is probably not an apomorphy for the group.

TRACHEARY ELEMENTS (OF XYLEM)

The vascular plants, as the name states, have true vascular tissue, consisting of cells that have become highly specialized for conduction of fluids. (A **tissue** consists of two or more cell types that have a common function and often a common developmental history; see Chapter 10.) Vascular tissue was a major adaptive breakthrough in plant evolution;

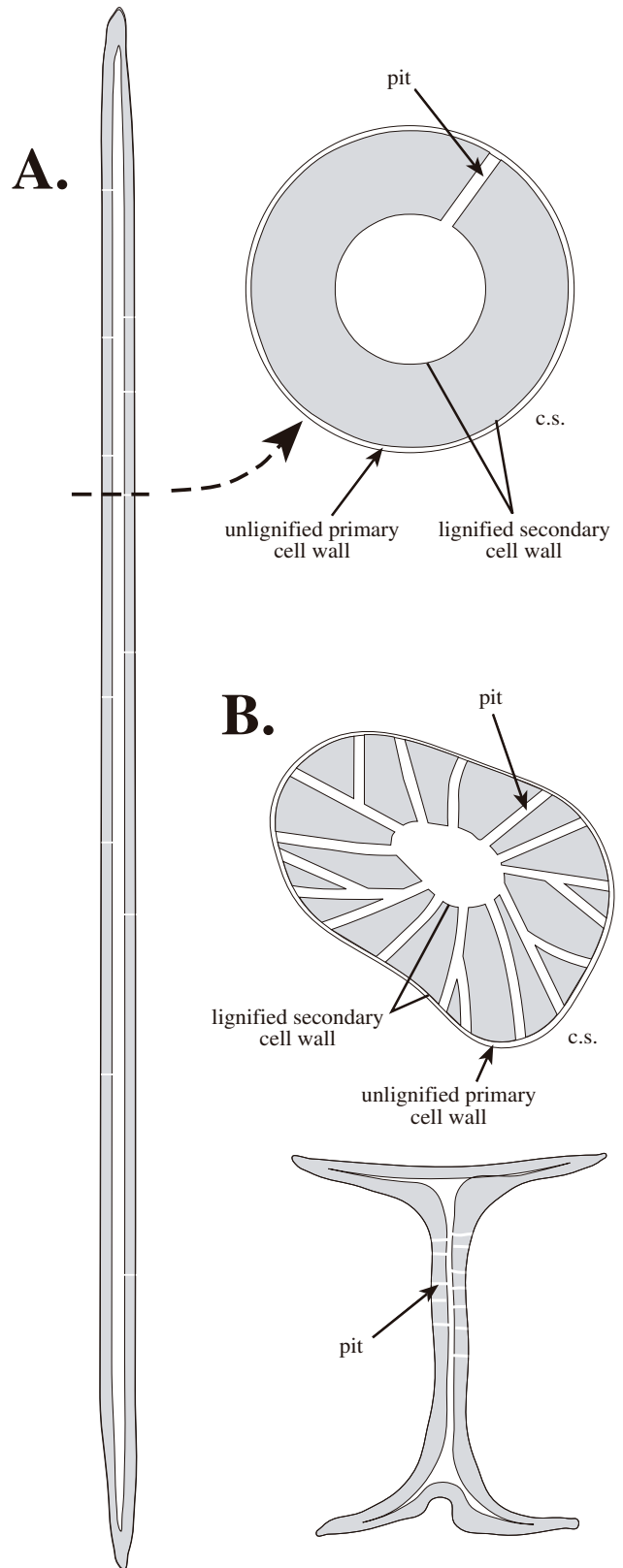


FIGURE 4.3 Sclerenchyma. **A.** Fiber cell. **B.** Sclereid cells. Cross-section = c.s.

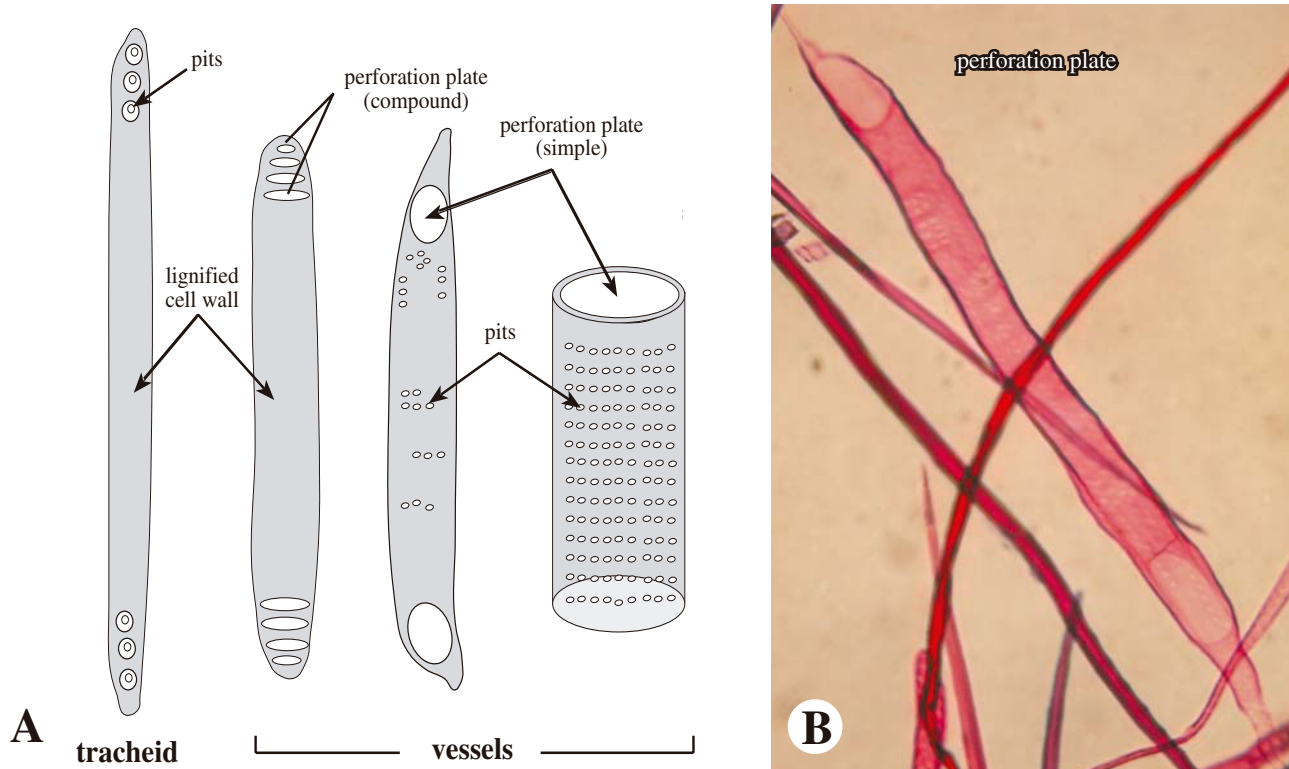


FIGURE 4.4 Conductive cells of vascular plants: tracheary elements. **A.** Types of tracheary elements. **B.** Vessel.

more efficient conductivity allowed for the evolution of much greater plant height and diversity of form.

Tracheary elements are specialized cells that function in water and mineral conduction. Tracheary elements are generally elongate cells, are dead at maturity, and have lignified 2° cell walls (Figure 4.4A,B). They are joined end-to-end, forming a tubelike continuum. Tracheary elements are typically associated with parenchyma and often some sclerenchyma in a common tissue known as **xylem** (Gr. *xylo*, wood, after the fact that wood is composed of secondary xylem). The function of tracheary elements is to conduct water and dissolved essential mineral nutrients, generally from the roots to other parts of the plant.

There are two types of tracheary elements: **tracheids** and **vessel members** (Figure 4.4A). These differ with regard to the junction between adjacent end-to-end cells, whether *imperforate* or *perforate*. Tracheids are imperforate, meaning that water and mineral nutrients flow between adjacent cells through the primary cell walls at pit-pairs, which are adjacent holes in the lignified 2° cell wall. Vessel members are perforate, meaning that there are one or more continuous holes or perforations, with no intervening 1° or 2° wall between adjacent cells through which water and minerals may pass.

The contact area of two adjacent vessel members is called the **perforation plate**. The perforation plate may be **compound**, if composed of several perforations, or **simple**, if composed of a single opening (see Chapter 10). Vessels may differ considerably in length, width, angle of the end walls, and degree of perforation.

Tracheids are the primitive type of tracheary element. Vessels are thought to have evolved from preexisting tracheids *independently* in several different groups, including in a few species of *Equisetum*, a few leptosporangiate ferns, all Gnetales (Chapter 5), and almost all angiosperms (Chapter 6).

SIEVE ELEMENTS (OF PHLOEM)

Sieve elements are specialized cells that function in conduction of sugars. They are typically associated with parenchyma and often some sclerenchyma in a common tissue known as **phloem** (Gr. *phloe*, bark, after the location of secondary phloem in the inner bark). Sieve elements are elongate cells having only a primary (1°) wall, with no lignified 2° cell wall. This primary wall has specialized pores (Figure 4.5C), which are aggregated together into **sieve areas** (Figure 4.5A). Each pore of the sieve area is a continuous hole in the 1° cell wall that is lined with a substance called

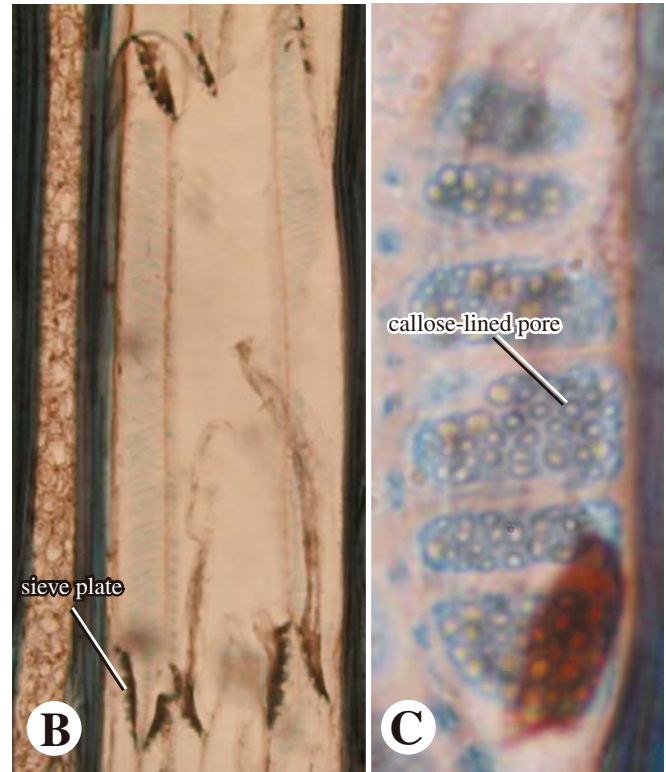
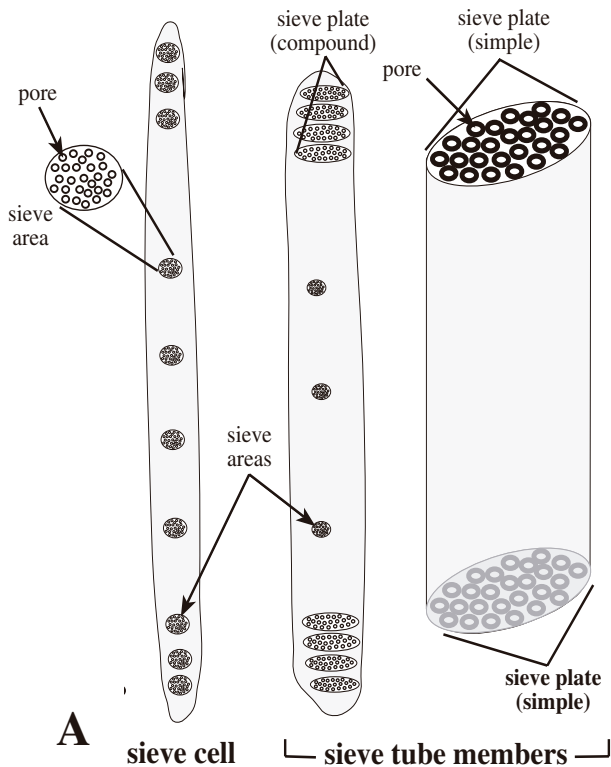


FIGURE 4.5 Conductive cells of vascular plants: sieve elements. **A.** Types of sieve elements. **B,C.** Sieve tube members.

callose, a polysaccharide composed of β -1,3-glucose units. (Note the difference in chemical linkage from cellulose, which is a polymer of β -1,4-glucose.) Sieve elements are semi-alive at maturity. They lose their nucleus and other organelles but retain the endoplasmic reticulum, mitochondria, and plastids. Like tracheary elements, sieve elements are oriented end-to-end, forming a tubelike continuum. Sieve elements function by conducting dissolved sugars from a sugar-rich source to a sugar-poor sink region of the plant. Source regions include the leaves, where sugars are synthesized during photosynthesis, or mature storage organs, where sugars may be released by the hydrolysis of starch. Sinks can include actively dividing cells, developing storage organs, or reproductive organs such as flowers or fruits.

There are two types of sieve elements: **sieve cells** and **sieve tube members** (Figure 4.5A). **Sieve cells** have only sieve areas on both end and side walls. **Sieve tube members** have both sieve areas and sieve plates (Figure 4.5B). **Sieve plates** consist of one or more sieve areas at the end wall junction of two sieve tube members; the pores of a sieve plate, however, are significantly larger than are those of sieve areas located on the side wall (Figure 4.5C). Both sieve cells and sieve tube members have parenchyma cells associated with them.

Parenchyma cells associated with sieve cells are called **albuminous cells**; those associated with sieve tube members are called **companion cells**. The two differ in that companion cells are derived from the same parent cell as are sieve tube members, whereas albuminous cells and sieve cells are usually derived from different parent cells. Both albuminous cells and companion cells function to load and unload sugars into the cavity of the sieve cells or sieve tube members. Sieve cells (and associated albuminous cells) are the ancestral sugar-conducting cells and are found in all nonflowering vascular plants. Sieve tube members were derived from sieve cells and are found only in flowering plants (angiosperms; see Chapter 6).

ENDODERMIS

Another apparent apomorphy for the vascular plants is the occurrence in some (especially underground) stems and all roots of a special cylinder of cells, known as the **endodermis** (Figure 4.6A,B). Each cell of the endodermis possesses a **Casparian strip**, which is a band or ring of lignin and **suberin** (chemically similar to lignin) that infiltrates the cell wall, oriented tangentially (along the two transverse walls) and axially (along the two radial walls; Figure 4.6C). The Casparian

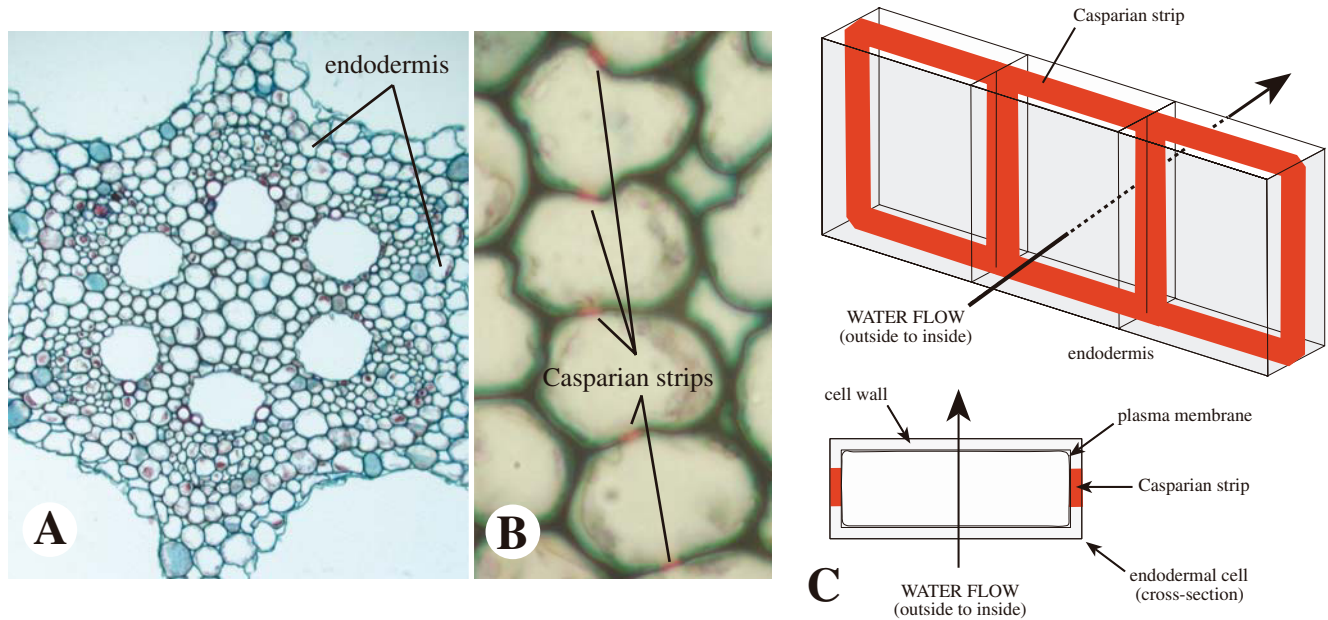


FIGURE 4.6 Endodermis of vascular plants. **A,B.** *Equisetum* rhizome. **A.** Rhizome cross-section, showing single layer of endodermal cells. **B.** Close-up of endodermal cells (in cross-section), showing Casparian strip thickenings. **C.** Diagram of Casparian strip, indicating function.

strip acts as a water-impermeable material that binds to the plasma membrane of the endodermal cells. Because of the presence of the Casparian strip, absorbed water and minerals that flow from the outside environment to the central vascular tissue must flow through the plasma membrane of the endodermal cells (as opposed to flowing through the intercellular spaces, i.e., between the cells or through the cell wall). Because the plasma membrane may differentially control solute transfer, the endodermis (with Casparian strips) selectively controls *which* mineral nutrients are or are not absorbed by the plant; thus, toxic or unneeded minerals may be differentially excluded.

INDEPENDENT, LONG-LIVED SPOROPHYTE

Like all land plants, the vascular plants have a haplodiplontic alternation of generations, with a haploid gametophyte and a diploid sporophyte. Unlike the liverworts, hornworts, and mosses, however, vascular plants have a dominant, free-living, photosynthetic, relatively persistent *sporophyte* generation. In the vascular plants, the gametophyte generation is also (ancestrally) free-living and may be photosynthetic, but it is smaller (often much more so) and much shorter-lived than the sporophyte generation. In all land plants, the sporophyte is initially attached to and nutritionally dependent upon the gametophyte. However, in the vascular plants, the sporophyte soon grows larger and becomes nutritionally independent, usually with the subsequent death of the gametophyte. (In seed

plants the female gametophyte is attached to and nutritionally dependent upon the sporophyte; see Chapter 5.)

The sporophytic axes, or **stems**, of vascular plants are different from those of liverworts, hornworts, and mosses in being branched and bearing multiple sporangia. Vascular plants share this feature with some fossil plants that are transitional between the bryophytes and the tracheophytes. This more inclusive group, containing these basal, fossil taxa (having branched sporophytic stems and multiple sporangia) plus the tracheophytes, is referred to as the **polysporangiophytes** (see Chapter 3).

Stems function as supportive organs, bearing and usually elevating leaves and reproductive organs; they also function as conductive organs, via vascular tissue, of water/minerals and sugars between roots, leaves, and reproductive organs. Structurally, stems can be distinguished from roots by several anatomical features (below).

Stems of the vascular plants typically have a consistent and characteristic spatial arrangement of xylem and phloem. This organization of xylem and phloem in the stem is known as a **stele**. In several groups of early vascular plant lineages, the stelar type is a **protostele**, in which there is a central solid cylinder of xylem and phloem (Figure 4.7). The largely parenchymatous tissue between the epidermis and vascular tissue defines the **cortex**. Protosteles are thought to be the most ancestral type of stem vasculature, found, e.g., in the rhyniophytes (below).

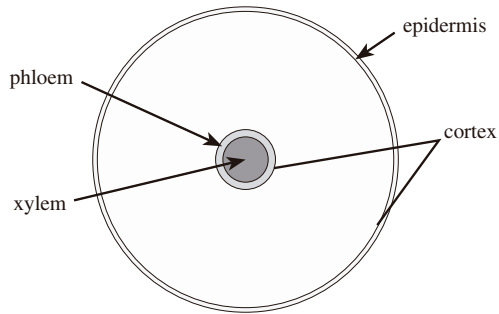


FIGURE 4.7 Example of a protostele, an ancestral vasculature of vascular plants.

SPOROPHYTIC LEAVES AND SHOOT

Another apomorphy of all extant vascular plants is the **sporophytic leaf**. Sporophytic leaves are dorsiventrally flattened organs that generally function as the primary organ of photosynthesis. Although some liverworts and all mosses have leaves, these occur on gametophytes only and are not strictly homologous with the sporophytic leaves of vascular plants. The evolution of sporophytic leaves (usually just called leaves) constituted a major adaptive innovation for extant vascular plants by greatly increasing the tissue area available for photosynthesis. This paved the way for the evolution of various ecological adaptive strategies, enabling some vascular plants to survive in previously inaccessible habitats. In addition, leaves or leaflike homologues have become evolutionarily modified for numerous other functions in plants, to be discussed later.

Leaves have a characteristic anatomy (Figure 4.8). Because they are usually dorsiventrally flattened organs (with some exceptions), both an upper and lower epidermis can be defined. As with all land plants, a cuticle covers the outer cell wall of the epidermal cells. One or more **vascular bundles**, or **veins**, contain xylem and phloem tissue and conduct water and sugars to and from the chloroplast-containing **mesophyll** cells. The mesophyll of some leaves is specialized into upper, columnar palisade mesophyll cells and lower, irregularly shaped spongy mesophyll cells, the latter with large intercellular spaces (Figure 4.8). Stomata, which function in gas exchange (see Chapter 3), are typically found only in the lower epidermis of leaves (Figure 4.8).

Sporophytic leaves originate developmentally as part of an integral association of stem plus leaves known as a **shoot** (Figure 4.9). The tip of a shoot contains one or more actively dividing cells of the **apical meristem**. These cells undergo continuous mitotic divisions. [The ancestral apical meristem consisted of a single, apical cell; in seed plants (see later discussion; Chapter 5), the apical meristem is complex, consisting of a number of continuously dividing cells.] Vertically down from the apical meristem, the cells undergo considerable elongation, literally pushing the cells of the apical meristem upward or forward. Even further down from the shoot tip, the fully grown cells differentiate into their mature, specialized form. To the sides of the apical meristem region, certain regions of the outermost cell layers of a shoot undergo cell division and elongation. Further growth and differentiation in these regions result in the formation of a **leaf** (Figure 4.9A,D,E). The point of attachment of a leaf to the stem is known as the **node**; the region between two nodes is

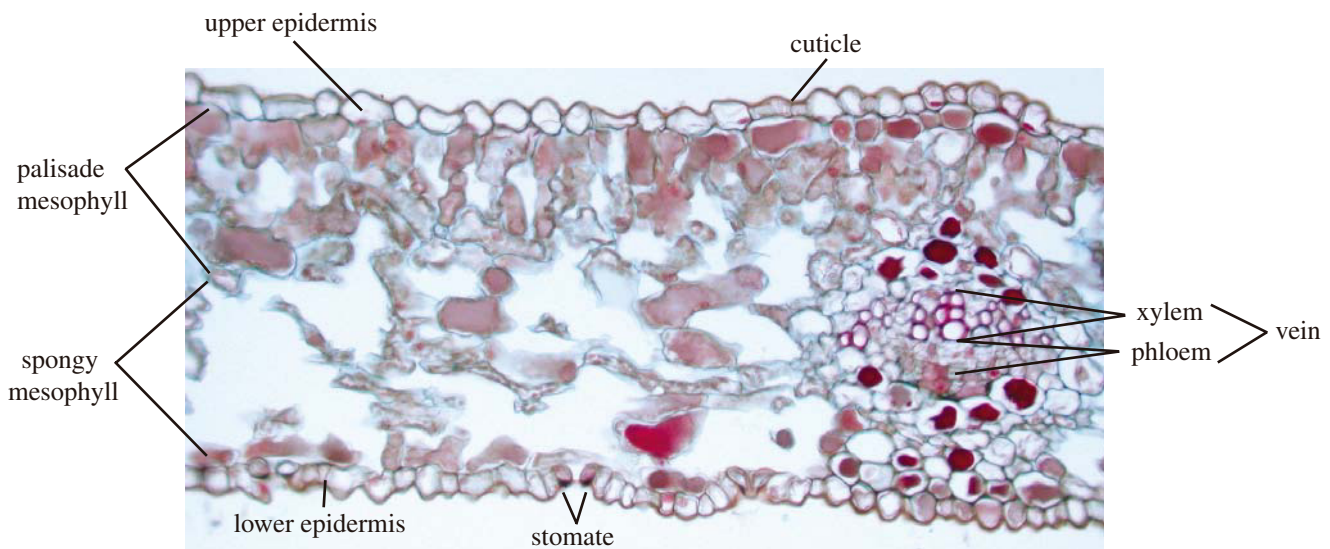


FIGURE 4.8 Cross-section of a vascular plant leaf.

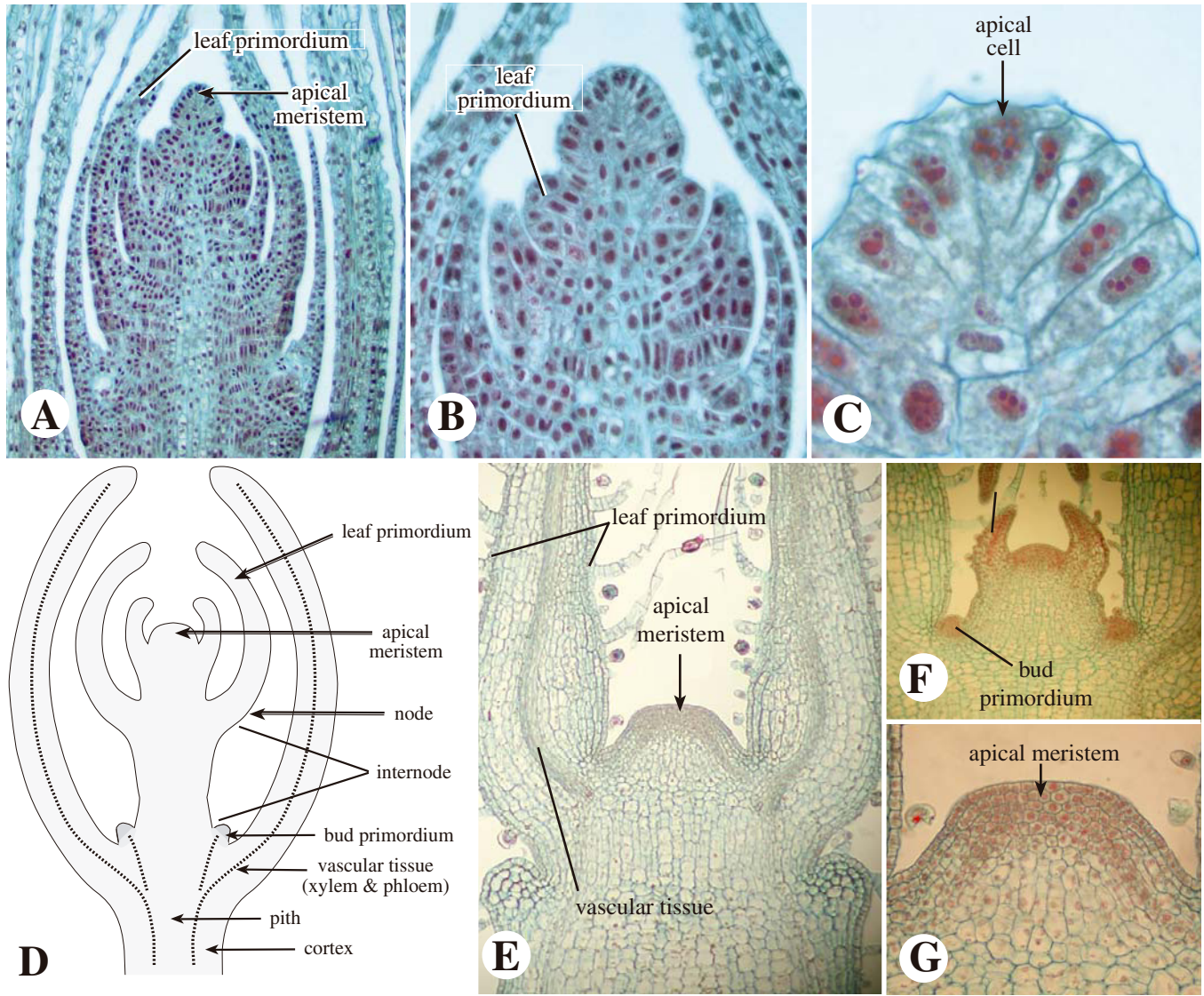


FIGURE 4.9 Shoot, longitudinal section. **A–C.** *Equisetum*, showing single, apical meristem cell. **D.** Diagram. **E–G.** *Plectranthus* [*Coleus*] shoot. Note complex apical meristem, leaf and bud primordia, and vasculature.

called an **internode** (Figure 4.9D). As the shoot matures, the leaves fully differentiate into an amazing variety of forms (see Chapter 9), and the stem differentiates a vascular system. Vascular strands run between stem and leaf, providing a connection for fluid transport.

Later in shoot development, the tissue at the region of the junction of stem and upper leaf, termed the **axil**, may begin to divide and differentiate into a **bud** (Figure 4.9F), defined as an immature shoot system. Buds have an architectural form identical to that of the original shoot. They may develop into a lateral branch or may terminate by developing into a reproductive structure. It is growth of new shoots from buds that result in branching of sporophytes in the vascular plants.

ROOTS

Roots are specialized plant organs that function in anchorage and absorption of water and minerals. Roots are found in all vascular plants except for the (extinct) rhyniophytes and the psilophytes (discussed later). Other fossil groups of vascular plants may have lacked roots; plants lacking roots generally have uniseriate (one cell thick), filamentous **rhizoids** (similar to those of bryophytes), which assume a similar absorptive function. Although roots are apparently not a strict apomorphy for all vascular plants, they constituted a major adaptive advance in enabling much more efficient water and mineral acquisition and conduction, permitting the evolution of plants in more extreme habitats.

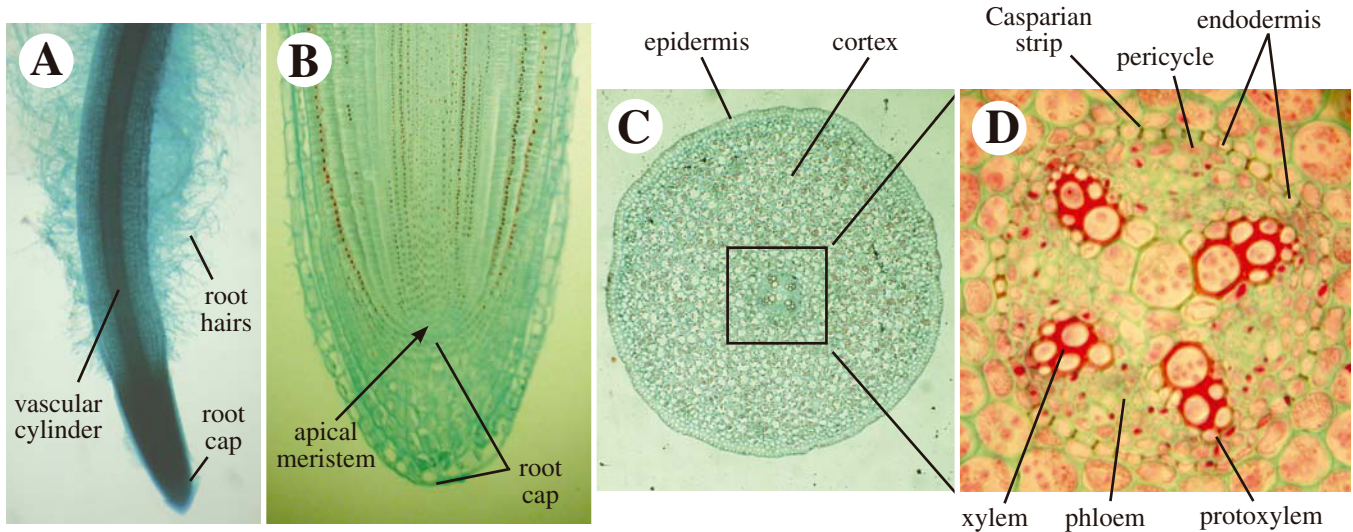


FIGURE 4.10 Anatomy of the root, an apomorphy of all but the earliest of the vascular plants. **A.** Root, whole mount. **B.** Root longitudinal section. **C.** Whole root cross-section. **D.** Close-up of root cross-section, showing vascular cylinder.

Roots, like shoots, develop by the formation of new cells within the actively growing **apical meristem** of the root tip, a region of continuous mitotic divisions (Figure 4.10B). At a later age and further up the root, these cell derivatives elongate significantly. This cell growth, which occurs by considerable expansion both horizontally and vertically, pushes the apical meristem tissue downward. Even later in age and further up the root, the fully-grown cells differentiate into specialized cells. [As with shoots, the ancestral apical meristem of roots consisted of a single, apical cell; in seed plants (see later discussion; Chapter 5), the apical meristem is complex, consisting of a number of continuously dividing cells.]

Roots are characterized by several anatomical features. First, the apical meristem is covered on the outside by a **rootcap** (Figure 4.10B); stems lack such a cell layer. The rootcap functions both to protect the root apical meristem from mechanical damage as the root grows into the soil and to provide lubrication as the outer cells slough off. Second, the epidermal cells away from the root tip develop hairlike extensions called **root hairs** (Figure 4.10A); these are absent from stems. Root hairs function to *greatly* increase the surface area available for water and mineral absorption. Third, roots have a central vascular cylinder, in which ridges of xylem *alternate* with cylinders of phloem; i.e., xylem and phloem are on alternate radii (Figure 4.10C,D). As in stems, the mostly parenchymatous region between the vasculature and epidermis is called the **cortex** (Figure 4.10C); the center of the vascular cylinder, if vascular tissue is lacking, is called a **pith**. Fourth, the vascular cylinder of roots is surrounded by an **endodermis**

with **Casparian strips** (Figure 4.10D). As with some stems, the endodermis in roots selectively controls which chemicals are and are not absorbed by the plant, functioning in selective absorption. Fifth, roots have no exogenous (externally developing) organs like leaf primordia; all **secondary roots arise endogenously** from the internal tissues of the root. Secondary roots develop by cell divisions within either the **endodermis** or the **pericycle**; the latter is a cylindrical layer of parenchyma cells located just inside the endodermis itself. Secondary roots must actually penetrate the surrounding tissue of the cortex and epidermis during growth.

Numerous modifications of roots have evolved, most of these restricted to the flowering plants (see Chapter 9). Roots of many, if not most, vascular plants have an interesting symbiotic interaction with various species of fungi, this association between the two known as **mycorrhizae**. The fungal component of mycorrhizae appears to aid the plant in both increasing overall surface area for water and mineral absorption and increasing the efficiency of selective mineral absorption, such as of phosphorus.

VASCULAR PLANT DIVERSITY

Of the tremendous diversity of vascular plants that have arisen since their first appearance some 400 million years ago, only the major lineages will be described here. These include the rhyniophytes, known only from fossils, plus clades that have modern-day descendants: the Lycopodiophyta,

Equisetales, Marattiales, Polypodiales (leptosporangiate ferns), Ophioglossales, Psilotales, and seed plants (Figure 4.1). The evolution of seed plants will be discussed in Chapter 5.

RHYNIOPHYTA — RHYNIOPHYTES

The Rhyniophyta, or rhyniophytes, were among the first vascular land plants. They include only extinct, fossil plants and may constitute a paraphyletic group. Rhyniophytes include the genus *Rhynia* (Figure 4.11A,B), a well-known vascular plant from the early Devonian, ca. 410–360 million years ago. Rhyniophyte sporophytes consisted of dichotomously branching axes that bore terminal sporangia.

Rhyniophytes ancestrally lacked both roots and a leaf-bearing shoot system; these two features evolved later, prior to or within the Lycophyte lineage (discussed next). The stems of Rhyniophytes were protostelic (Figure 4.7) in which the first-formed xylem (known as protoxylem) was centrarch (positioned at the center).

LYCOPODIOPHYTA — LYCOPHYTES

The Lycopodiophyta, or lycophytes (also commonly called lycopods), are a lineage of plants that diverged after the rhyniophytes. An extinct, fossil group, known as the zosterophylls [Zosterophyllophytina], are either immediately basal to or sister to the lycophytes. Zosterophylls had no leaves, but possessed lateral sporangia, similar to those of the lycophytes (see later discussion). Within the lycophytes, the now extinct

Lepidodendron, *Sigillaria*, and relatives (Figure 4.11C–E) were woody trees that comprised a large portion of the primary biomass of forests during the Carboniferous, approximately 300 million years ago. Fossil remains of these plants today make up much of the Earth's coal deposits.

A number of apomorphies characterize the lycophytes, three of which are mentioned here. First, the roots of lycophytes have an **endarch protoxylem**. **Protoxylem** refers to the first tracheary cells that develop within a patch of xylem and that are typically smaller and have thinner cell walls than the later formed **metaxylem**. In the roots of lycophytes, the protoxylem forms in a position *interior* to the metaxylem (i.e., toward the stem center). Second, the stems of lycophytes have an **exarch protoxylem** (just the reverse of the roots). In the stems of lycophytes, the protoxylem forms in a position *exterior* to the metaxylem (i.e., away from the stem center; Figure 4.12A,B). Third, lycophytes have a sporophytic leaf structural type known as a **lycophyll** (essentially synonymous with microphyll). **Lycophylls** are characterized as having an intercalary meristem (at the proximal side of the leaf base) and lacking a gap in the vasculature of the stem (Figure 4.12C). Lycophylls also have a single, unbranched (very rarely branched) vein. Lycophylls may have evolved from small appendages called **enations** (found in rhyniophytes and some lycophyte relatives), which may resemble lycophylls but which lack vascular tissue. Thus, lycophylls may have formed by the innervation of vasculature tissue from the stem into the

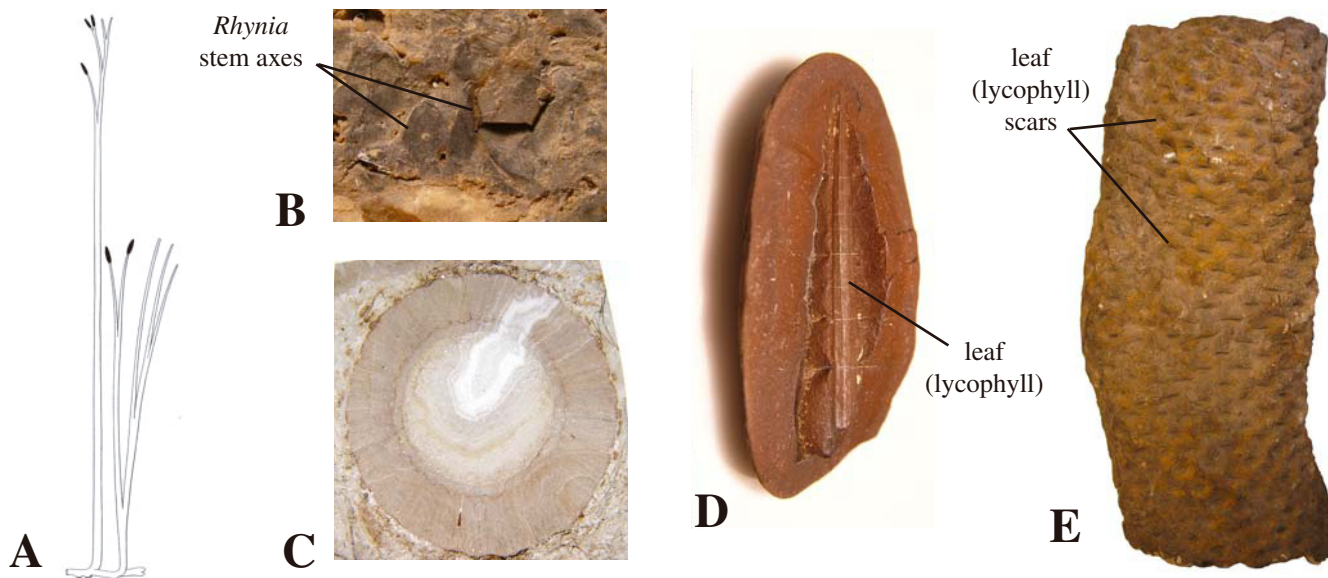


FIGURE 4.11 A–B. Rhyniophytes. A. Reconstruction of *Rhynia major*, an early, extinct vascular plant. Note erect, branched stem (without leaves) bearing terminal sporangia. (Reproduced from Kidston, R. and W. H. Lang. 1921. Transactions of the Royal Society of Edinburgh. vol. 52(4):831–902.) B. *Rhynia* stem axes, embedded in Rhynie chert. C–E. Lycophytes. C–D. *Sigillaria*, an extinct, woody lycophyte. C. Stem cross-section, showing outer wood. D. Fossil impression of lycophyll leaf, showing single vein. E. Fossil cast of *Lepidodendron*, an extinct, woody, tree-sized lycophyte. Note lycophyll scars.

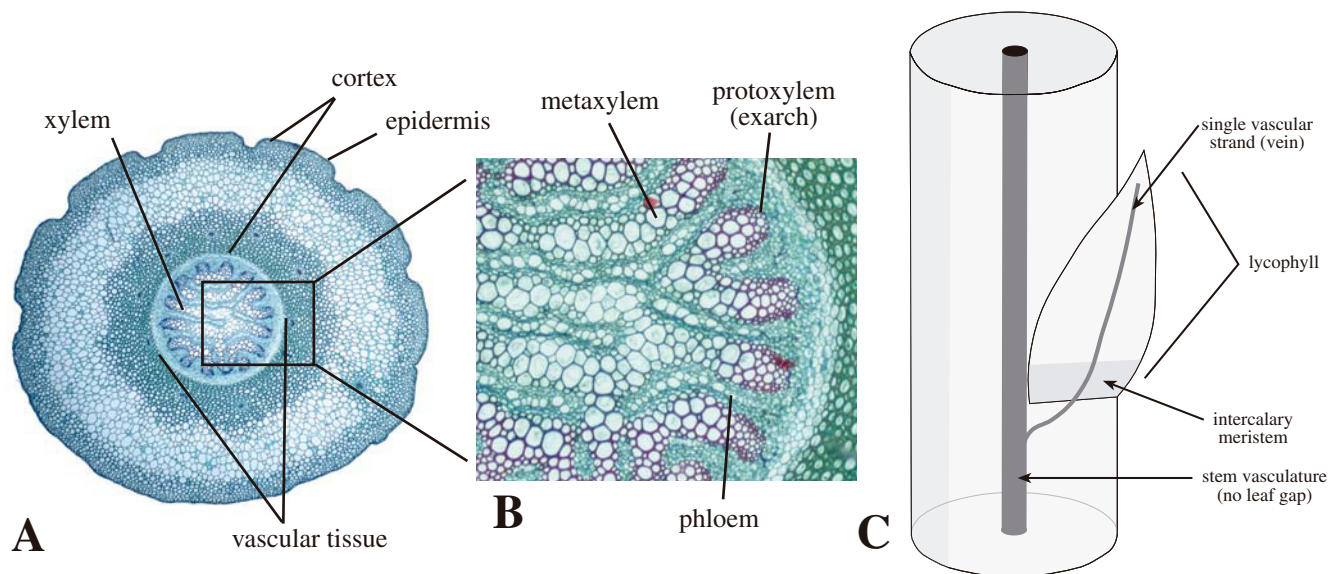


FIGURE 4.12 **A,B.** *Lycopodium* stem cross-section, showing protoxylem that is exarch (to periphery of stem). **C.** Lycophyll structure.

enation and flattening of this structure into a dorsiventral, planar posture; such a gradation, from enation to lycophyll, may be seen in some fossil plants.

The only lycophytes that survived to the present are small, nonwoody, herbaceous plants, typically grouped into three families: Lycopodiaceae, Sellaginellaceae, and Isoetaceae. The Lycopodiaceae (ca. 380 species; Figure 4.13), which are often commonly called club-mosses, are distinguished in having one type of spore, a condition known as **homospory**. The Lycopodiaceae contain about 300 species in five genera: *Diphasiastrum*, *Huperzia* (Figure 4.13A,C), *Lycopodiella*, *Lycopodium* (Figure 4.13B,D G), and *Phylloglossum* (Figure 4.13H). Some family members may in fact resemble a large moss (e.g., Figure 4.13A), but they are true vascular plants, the persistent, long-lived phase being sporophytic. Sporangia of the Lycopodiaceae, like those of all lycophytes, develop laterally (relative to the stem) in the axils of specialized leaves termed **sporophylls** (Figure 4.13E). In some members of the family, the sporophylls are similar to the vegetative leaves (Figure 4.13C) and co-occur with them on shoots that are indeterminate, i.e., with continuous growth. In other family members, the sporophylls differ in size or shape from vegetative leaves and are aggregated into a terminal shoot system that is determinate, meaning that it terminates growth after formation. This determinate reproductive shoot, consisting of a terminal aggregate of sporophylls with associated sporangia, is known as a **strobilus** or **cone** (Figure 4.13B,D,G,H).

The two other extant lycophyte families are the Selaginellaceae and Isoetaceae. The Selaginellaceae (Figure 4.14A G) contain approximately 700 species in the single

genus *Selaginella*, commonly called spike-moss. Species of *Selaginella* occur in two vegetative forms. Some have spirally arranged vegetative leaves that are isomorphic, of only one size and shape (Figure 4.14A). Other *Selaginella* species, which are generally prostrate, have leaves that are dimorphic, of two forms, arranged in four rows: two lateral rows of larger leaves and two upper, or dorsal, rows of smaller leaves (Figure 4.14B,C). The Isoetaceae (Figure 4.14H J) consist of approximately 150 species in the single genus *Isoetes*, commonly called quillwort or Merlin's-grass. *Isoetes* plants consist of a cormose (rarely rhizomatous) stem bearing numerous acicular (needle-like) leaves (Figure 4.14J). Species of *Isoetes* are aquatics found in shallow, sometimes periodically inundated, pools.

The Selaginellaceae and Isoetaceae differ from the Lycopodiaceae in having leaf **ligules** and in being **heterosporous**, both of which are apomorphies within the lycophytes (Figure 4.1). Ligules are tiny appendages on the upper (adaxial) side of the leaf (both vegetative and reproductive), near the leaf base (Figures 4.14D, 4.15). The function of ligules is not clear; one proposal is that they act as glands, providing hydration for young, developing lycophylls. **Heterospory** refers to the production of *two* types of spores: **microspores** and **megaspores**, which form within specialized sporangia: **microsporangia** and **megasporangia** (Figure 4.14E). Microspores are relatively small (Figure 4.14F) and are produced in large numbers. Megaspores (Figure 4.14G) are much larger in size and are produced in fewer numbers (typically four) per sporangium. Megasporangia and microsporangia may be produced together in the same shoot or in different shoots.

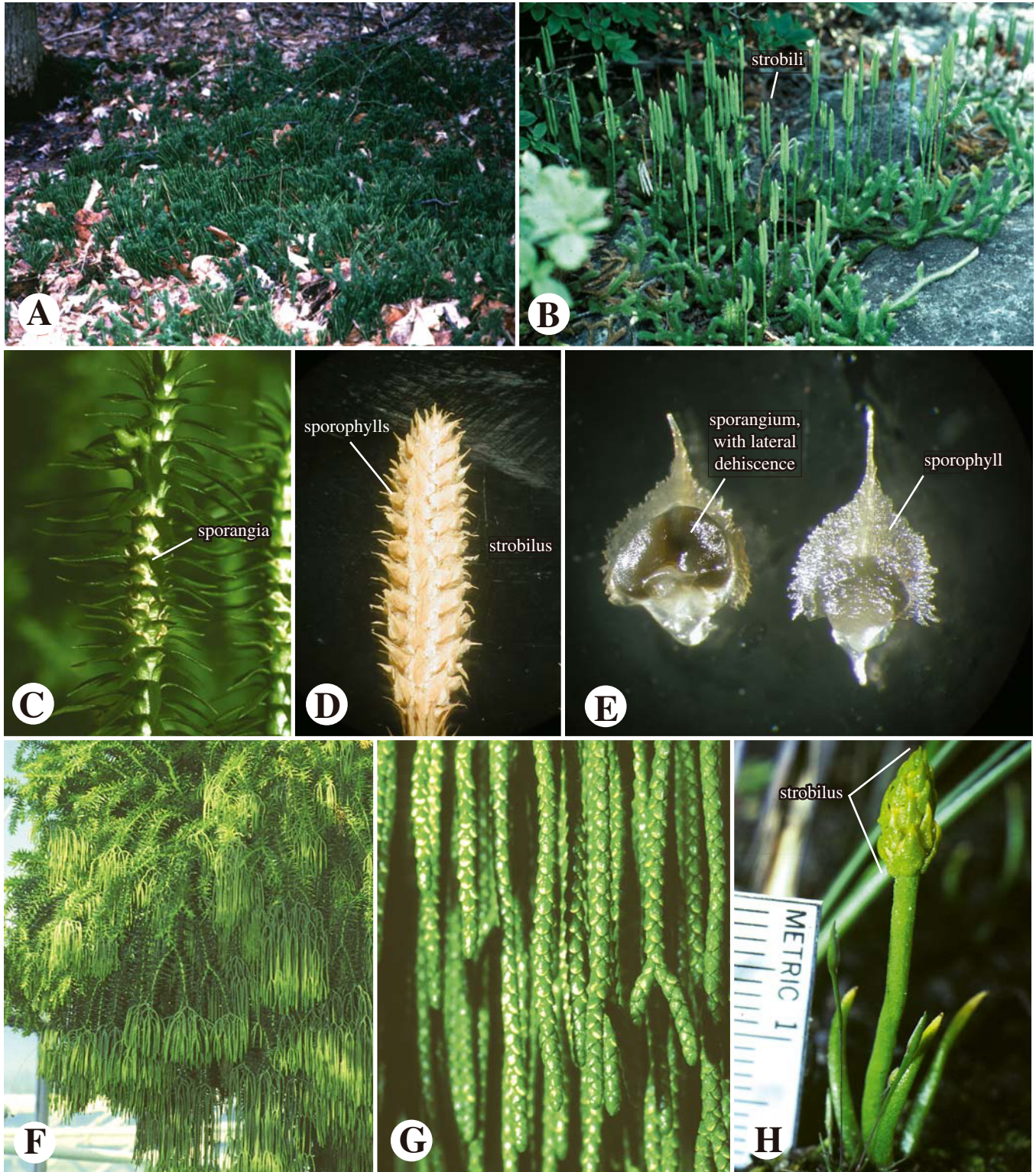


FIGURE 4.13 Lycopodiophyta homosporic taxa. **A.** *Huperzia lucidula*, a species with unspecialized reproductive organs. **B.** *Lycopodium clavatum*, a species with strobili. **C.** *Huperzia lucidula*, showing sporangia in axils of leaves, with no specialized cones. **D.** *Lycopodium annotinum*, strobilus close-up, showing sporophylls. **E.** *Lycopodium clavatum*, sporophylls removed from strobilus, showing sporangia having lateral dehiscence, adaxial view (left) and abaxial view (right). **F–G.** *Lycopodium squarrosum*, a large epiphytic lycopod with pendant branches and specialized strobili (at **G**). **H.** *Phylloglossum drummondii*, a small, cormose lycopod.

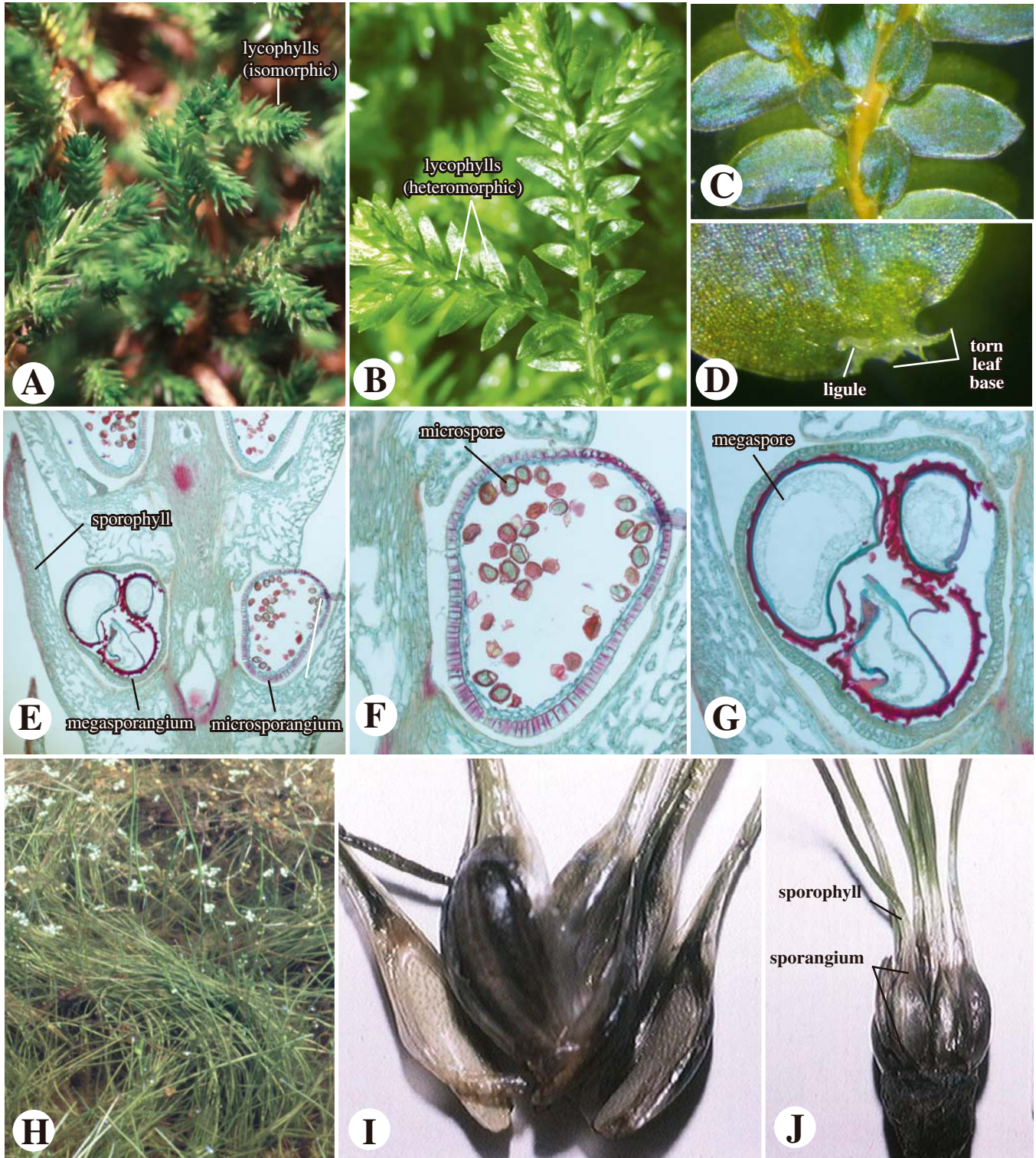


FIGURE 4.14 Lycopodiophyta heterosporic taxa. **A.** *Selaginella bigelovii*, with isomorphic leaves. **B.** *Selaginella apoda*, with dimorphic leaves. **C,D.** *Selaginella* sp. **C.** Close-up of dimorphic leaves. **D.** Close-up of ligule, adaxial side of leaf base. **E–G.** *Selaginella* sp., reproductive. **E.** Strobilus longitudinal section, showing sporophylls, megasporangia, and microsporangia. **F.** Close-up of microsporangium, containing microspores. **G.** Close-up of megasporangium, containing megaspores. **H,I.** *Isoetes howellii*. **H.** Plants growing in vernal pool. **I.** Close-up of male and female sporangia, containing microspores and megaspores, respectively. **J.** *Isoetes orcutii*, showing sporophylls with basal microsporangia or megasporangia.

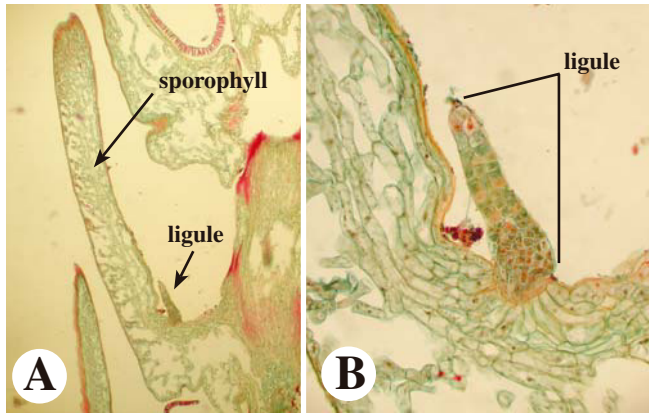


FIGURE 4.15 Lycopodiophyta heterosporic taxa. **A.** Longitudinal section of *Selaginella* strobilus, showing sporophyll and ligule. **B.** Ligule, close-up.

Some species of *Selaginella* have strobili, with specialized sporophylls subtending the sporangia on a determinate shoot (Figure 4.14E). In *Isoetes*, the sporophylls bear enlarged microsporangia or megasporangia on the upper (adaxial) side of the sheathing base (Figure 4.14I,J); male sporophylls (**microsporophylls**) are usually located inner to the female sporophylls (**megasporophylls**). The size and sculpturing pattern of the spores can be an important feature in identifying different species of *Isoetes*. In both *Selaginella* and *Isoetes*, the megaspore develops into a **female gametophyte**, which contains only archegonia, housing the egg cell. Each microspore germinates to form a **male gametophyte**, which produces only antheridia, the sperm-manufacturing organs. The gametophytes of *Selaginella* and *Isoetes* are **endosporic**, meaning that the gametophytes develop entirely *within* the original spore wall. Heterospory and endospory also evolved independently in the seed plants (see Chapter 5).

Interestingly, the fossil tree *Lepidodendron* belongs to the ligulate lycophytes, being most closely related to *Isoetes* among the extant lycophytes. *Lepidodendron* possessed leaf ligules and was heterosporous.

EUPHYLLOPHYTES

The sister group of the lycophytes are the euphyllophytes, including all the other vascular plants (Figure 4.1). Two major apomorphies that unite the euphyllophytes are mentioned here. First, the roots have an exarch protoxylem, in which the protoxylem is placed outer to the metaxylem (Figure 4.10D). Second, the leaves are **euphyllous**, meaning that they grow by means of either marginal or apical meristems and have an associated **leaf gap**, a region of nonvascular, parenchyma tissue interrupting the vasculature of the stem (Figure 4.16). **Euphylls** typically have more than one vein and generally

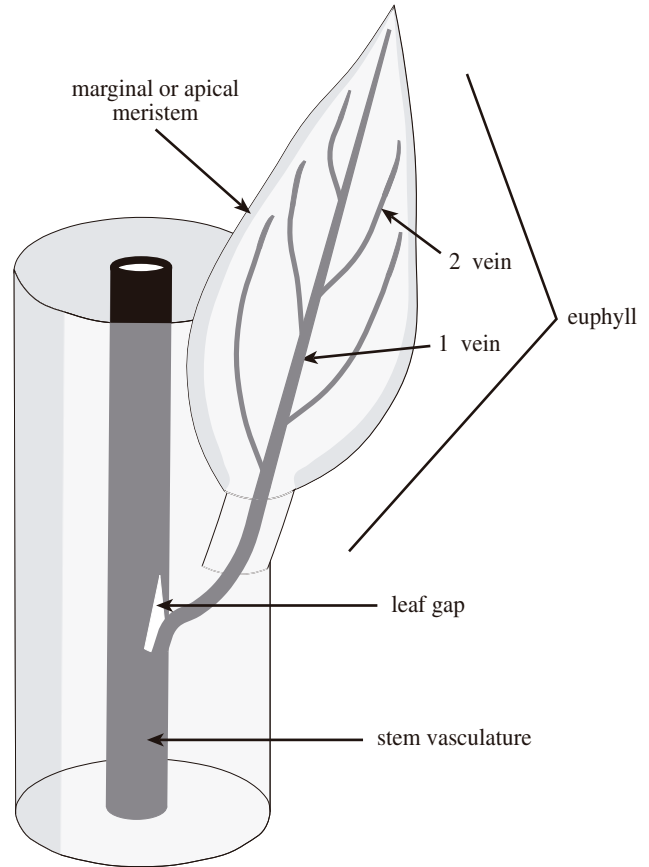


FIGURE 4.16 Euphyll structure.

have a highly branched system of veins, although in a few euphyllous taxa, the veins have become secondarily reduced again to a single mid-vein. (Note that *euphyll* is essentially synonymous with *megaphyll*, a more traditional term.) Fossil evidence suggests that euphylls evolved from a planar branch system, different from that of lycophylls. Third, euphylls have a molecular apomorphy, a 30-kilobase **inversion** located in the large single-copy region of **chloroplast DNA** (see Figure 14.4 of Chapter 14).

Euphyllophytes are composed of two major groups, which are sister to one another: monilophytes (ferns, in the broad sense) and spermatophytes (seed plants), the latter to be discussed in Chapter 5.

MONILOPHYTES — FERNS

Recent morphological and molecular phylogenetic studies (e.g., Kenrick and Crane, 1997; Pryer et al., 2001) support the recognition of a monophyletic group of vascular plants that are inclusive of five major lineages: Equisetales (horsetails), Marattiales (marattioid ferns), Ophioglossales (ophioglossoid ferns), Psilotales (whisk ferns), and Polypodiales

(leptosporangiate ferns). This monophyletic group has been termed the monilophytes (or moniliformopses); the common name is often now termed ferns, in the broad sense of the word. One recognized anatomical apomorphy for the monilophytes is that the **stem protoxylem is mesarch** in position (Figure 4.17E), meaning that tracheary elements first mature near the middle of a patch of xylem; this protoxylem (unlike that of some related fossil taxa) is restricted to the lobes of the xylem. The derivation of monilophyte (*L. monilo*, necklace or string of beads + Gr. *phyt*, plant) is in reference to this anatomy.

Lastly, the ancestral stem vasculature of the monilophytes, found in most (but not all) extant members, is the **siphonostele**. A **siphonostele** (Figure 4.17A–D) is a type of stem vasculature in which a ring of xylem is surrounded by an outer layer of phloem (ectophloic siphonostele, Figure 4.17A)

or by an outer and inner layer of phloem (amphiphloic siphonostele, Figure 4.17B; if dissected, called a dictyostele, Figure 4.17C); siphonosteles have a central, parenchymatous pith (Figure 4.17). Siphonosteles have evidently become secondarily modified in some monilophytes.

OPHIOGLOSSALES — OPHIOGLOSSOID FERNS

The Ophioglossales (=Ophioglossidae), or ophioglossoid ferns, consist of a few genera of fernlike plants. The ophioglossoid ferns are unique in that each leaf (or frond) consists of a **sterile segment**, which contains the photosynthetic blade or lamina, and a **fertile segment**. The underground rhizome gives rise to unbranched roots that lack root hairs. The most common genera of the Ophioglossales are *Botrychium*, commonly called grape fern or moonwort, and *Ophioglossum*, commonly called adder's tongue. *Botrychium* species have

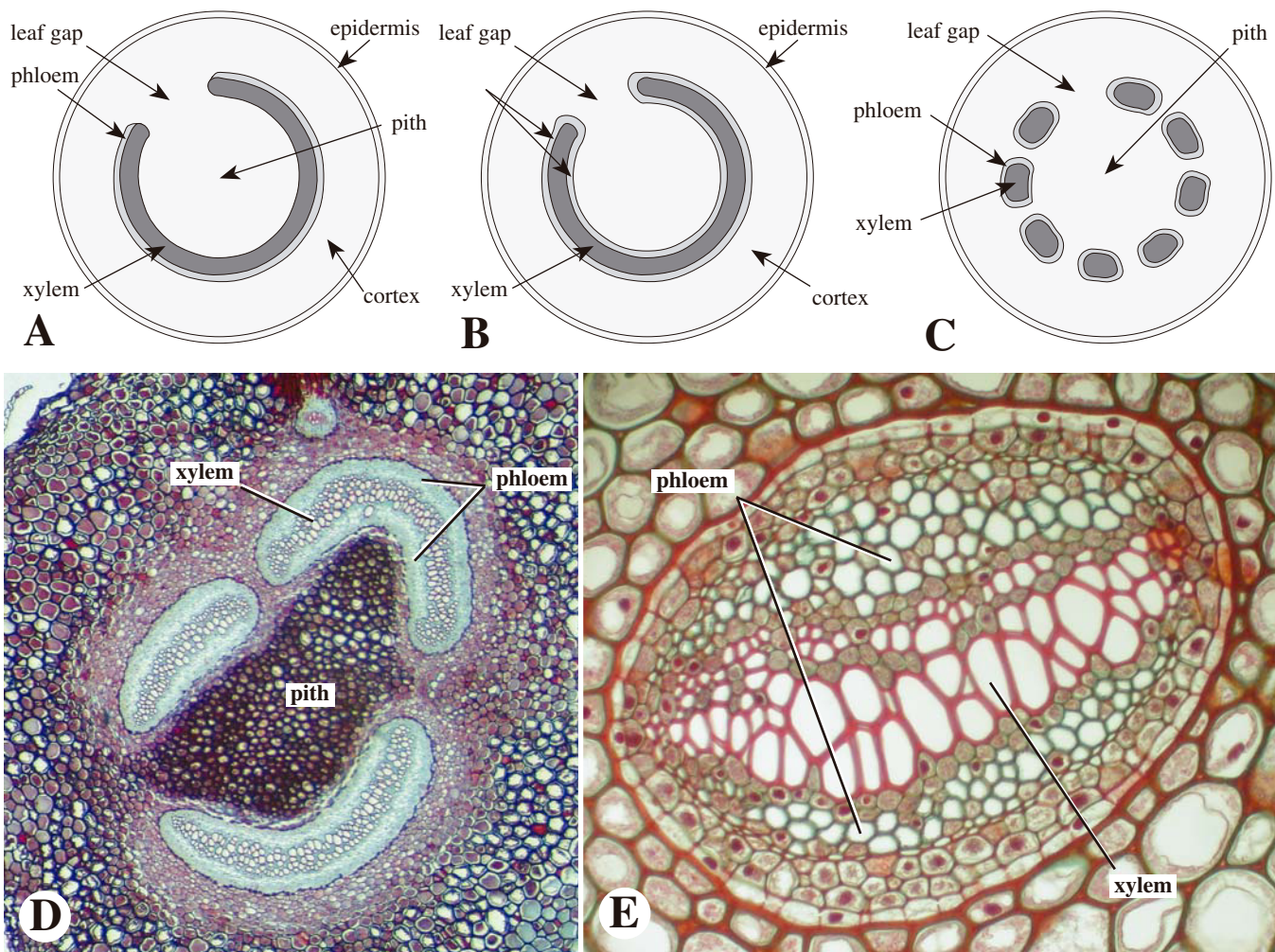


FIGURE 4.17 A–C. Siphonostele types. A. Ectophloic siphonostele, with phloem to outside of xylem. B. Amphiphloic siphonostele, with phloem to outside and inside. C. Dictyostele, a dissected amphiphloic siphonostele. D. *Adiantum* rhizome, an amphiphloic siphonostele. E. *Polypodium* rhizome, close-up of vasculature showing mesarch protoxylem, an apomorphy for the monilophytes (ferns).

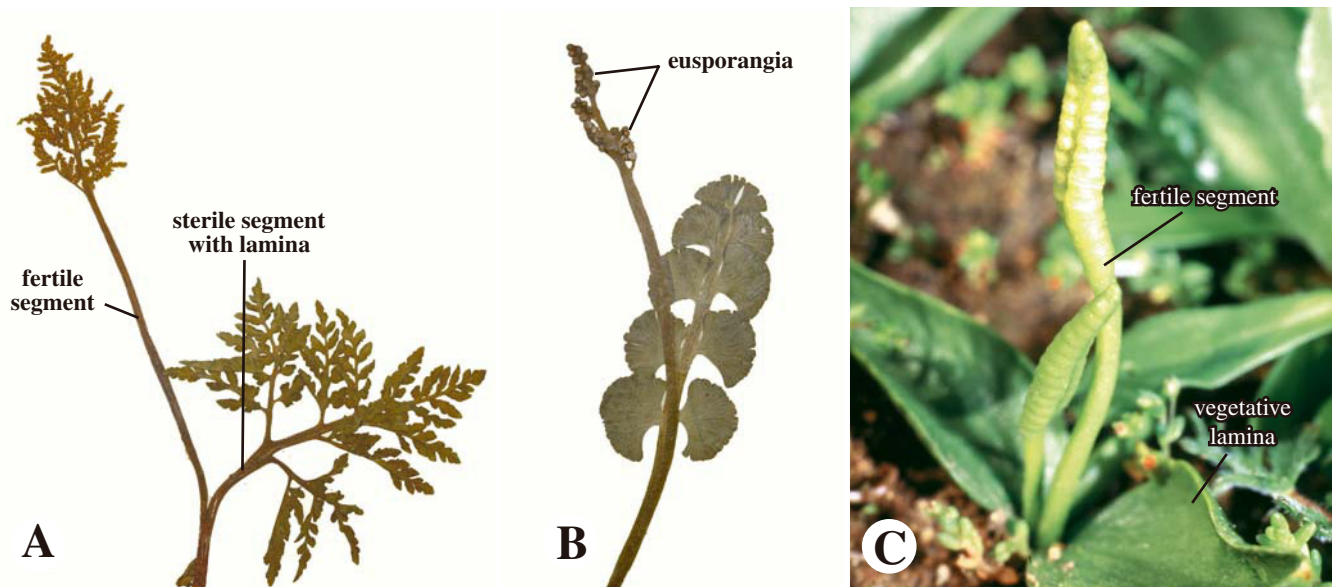


FIGURE 4.18 Ophioglossales. **A,B.** *Botrychium* species, showing vegetative lamina and fertile segment. **A.** *B. multi dum*. **B.** *B. lunaria*. **C.** *Ophioglossum californicum*, adder s tongue. Note elongate fertile segment bearing eusporangia.

a divided to compound lamina and a branched fertile segment (Figure 14.18A,B), whereas species of *Ophioglossum* have a simple, undivided lamina and an unbranched fertile segment (Figure 14.18C).

The sporangia of the Ophioglossales, and all other monilophytes except for the leptosporangiate ferns, are often termed **eusporangia** (or eusporangiate sporangia) to contrast them with leptosporangia of the leptosporangiate ferns (see later discussion). A **eusporangium** is relatively large, is derived from several epidermal cells, and has a sporangial wall comprised of more than one cell layer (Figure 4.18B,C). Eusporangia are the ancestral condition of the land plants.

Two features may constitute apomorphies, linking the ophioglossoid ferns with the Psilotales, the whisk ferns (discussed later). First, the roots of ophioglossoid ferns are unusual in lacking both root branches and root hairs. This may represent a transitional stage to the total loss of roots in the whisk ferns. Second, the gametophytes of the Ophioglossales and Psilotales are nonphotosynthetic (heterotrophic), contain mycorrhizal fungi, and are often subterranean (Figure 4.1).

PSILOTALES — WHISK FERNS

The Psilotales, or psilophytes (commonly called whisk ferns), consist of only two genera of plants, *Psilotum* (two species) and *Tmesipteris* (ca. 10 species). Like all vascular plants, the whisk ferns have an independent, dominant, free-living sporophyte; the haploid gametophyte is small, obscure,

and free-living in or on the soil. The sporophyte consists of a horizontal rhizome that gives rise to aerial, photosynthetic, generally dichotomously branching stems (Figure 4.19A,B). Plants are often epiphytic, with rhizomes having mycorrhizal symbiotic associations. All psilophytes **lack true roots**, an apomorphy for the group; only absorptive rhizoids arise from the rhizome. The absence of roots in the psilophytes has often been considered to be a primitive retention, the psilophytes having been viewed as direct descendants of the rhyniophytes. However, molecular studies clearly indicate that psilophytes are sister to the Ophioglossales (Figure 4.1) and likely lost roots secondarily.

The leaves of psilophytes are very reduced and peglike (Figure 4.19C) and may lack a vascular strand, in which case they are termed **enations**. The sporangia (which, like the Ophioglossales, could be termed eusporangia) are two- or three-lobed, which is interpreted as a **synangium**, a fusion product of two or three sporangia (Figure 4.19D). The synangia are yellowish at maturity and are subtended by a forked appendage, an apomorphy for the group. As in the Ophioglossales, the gametophytes of the Psilotales are nonphotosynthetic (subterranean or surface-dwelling) and may contain mycorrhizal fungi.

Psilotum nudum, the whisk broom, is the most widespread species of the psilophytes, one that commonly serves as an exemplar for the group (Figure 4.19). *Psilotum nudum* is native to tropical regions and is cultivated in greenhouses and naturalized in warm climates worldwide.



FIGURE 4.19 Psilotophyta. *Psilotum nudum*. **A.** Whole plant, showing dichotomous branching. **B.** Close-up of plant. **C.** Vegetative stem close-up, showing reduced leaves or enations. **D.** Close-up of synangia, subtended by forked appendage.

EQUISETALES — HORSETAILS

The Equisetales, also called the equisetophytes, sphenophytes, or sphenopsids, are a monophyletic group that diverged early in the evolution of vascular plants. As with the lycophytes, some equisetophytes in the Carboniferous period, approximately 300 million years ago, were large woody trees. Among these was *Calamites* (Figure 4.20), another contributor to coal deposits. Current molecular systematic studies (Figure 4.1) place the equisetophytes near the Marattiales and within a group containing the Polypodiales (leptosporangiate ferns; see later discussion). However, this may contradict fossil interpretations, so the position of this group needs further investigation.

Equisetales are united by several apomorphies, four of which are cited here (Figure 4.1): (1) **ribbed stems** (Figure 4.21A,J), these often associated with internal **hollow canals** (Figure 4.21C); (2) **reduced, whorled leaves** that are usually marginally fused (Figure 4.21A,J); (3) **sporangiophores**, each of which consists of a peltate axis bearing pendant longitudinally dehiscent sporangia (Figure 4.21F,L); and (4) photosynthetic spores with **elaters** (Figure 4.21G,H; see later discussion).

Today, the only remaining equisetophytes are species of the genus *Equisetum*. *Equisetum* species generally have an extensive underground rhizome system with adventitious roots; the rhizome gives rise to erect, aerial shoots. The ribbed stems contain epidermal cells that are impregnated with silica. Thus, the stems are rather tough, laying claim to having

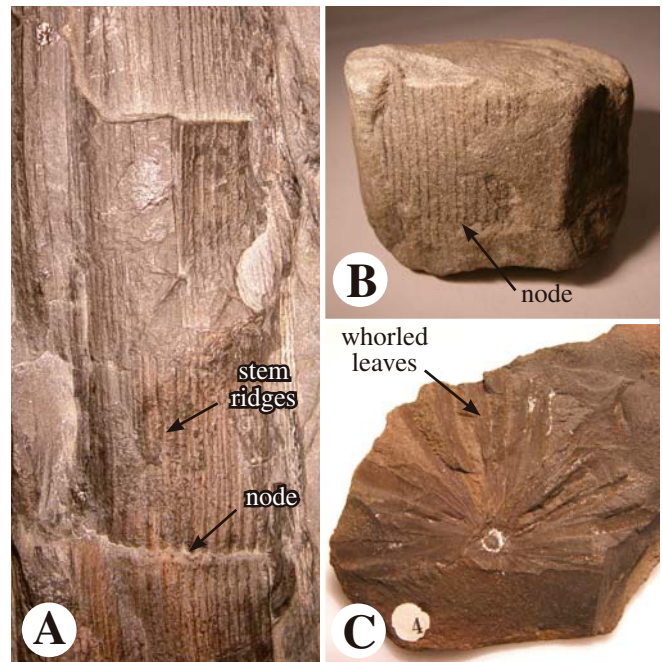


FIGURE 4.20 *Calamites*, an extinct, tree-sized equisetophyte. **A.** Fossil impression, showing nodes and stem ridges. **B.** Fossil cast of stem. **C.** Fossil impression showing whorled leaves of branch.

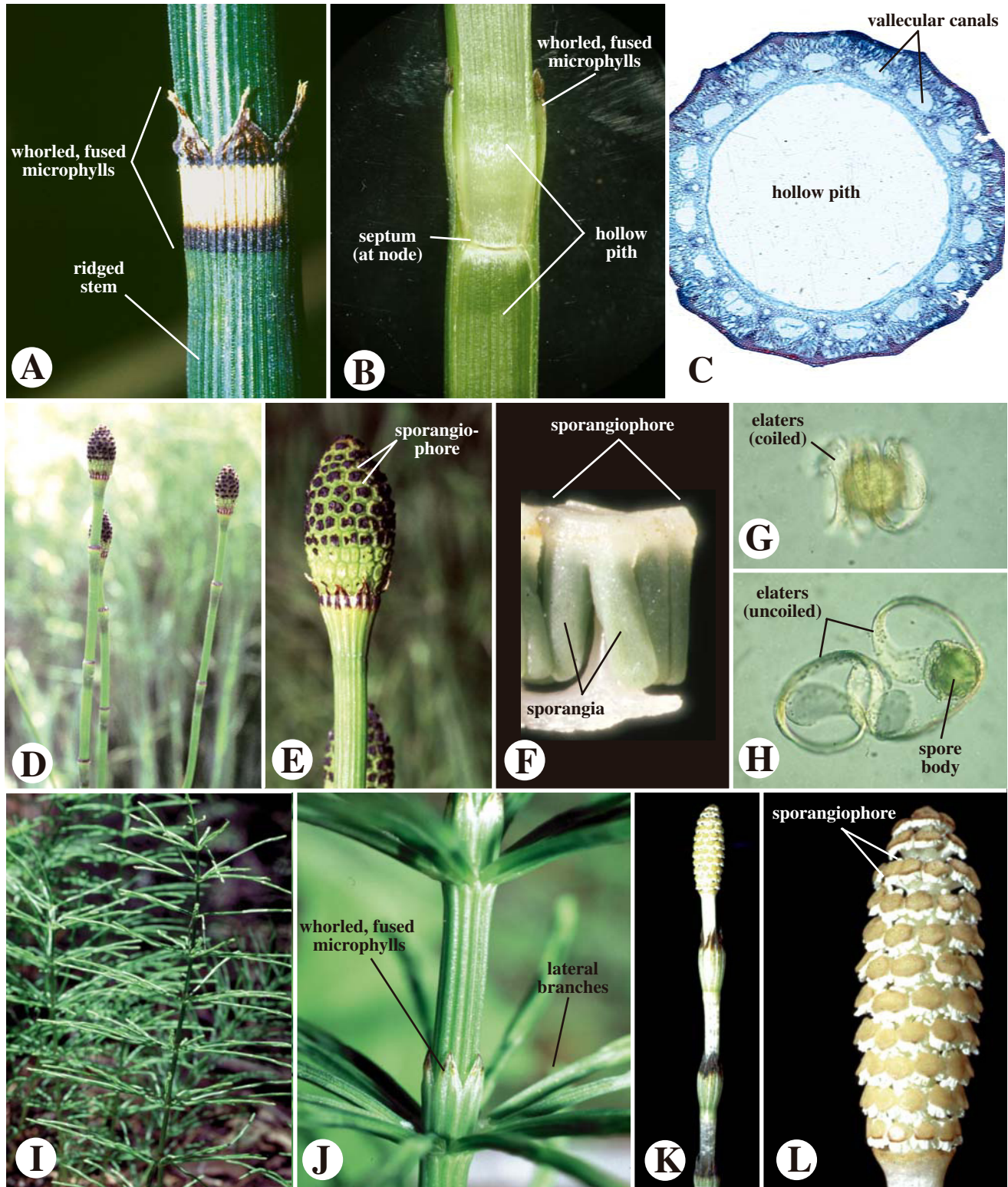


FIGURE 4.21 Equisetales. *Equisetum*, the only extant genus of the equisetophytes. **A,B.** *Equisetum hyemale*. **A.** Vegetative stem. Note ridged stem and whorled microphylls. **B.** Stem longitudinal section, showing central hollow pith and septum at nodes. **C.** Stem cross-section of *Equisetum* sp., showing central, hollow pith and peripheral, vallicular canals. **D,E.** *Equisetum laevigatum*, a scouring rush, having photosynthetic, generally unbranched aerial stems. **F.** Sporangio- phore, with several pendant sporangia. **G,H.** Spores, each with four elaters. **G.** Elaters coiled. **H.** Elaters uncoiled. **I–L.** *Equisetum arvense*, a horsetail, with dimorphic aerial stems. **I,J.** Sterile, photosynthetic stems with whorls of lateral branches. **K.** Reproductive, nonphotosynthetic aerial stem, lacking whorls of branches and terminating in a strobilus. **L.** Strobilus close-up, showing sporangiophores.

been used in the past for cleaning cooking utensils, hence the common name scouring rush. The stems are hollow (have a hollow pith), with cross walls called **septa** at each node (Figure 4.21B) and peripheral canals (termed vallicular canals; Figure 4.21C). The leaves are whorled and laterally fused, forming a sheathlike structure at the nodes (Figure 4.21A,B,J).

Equisetum species are classified in part based on their aerial branching pattern. In some species, whorls of lateral branches arise at the node from the axils of the leaves, actually penetrating the marginally fused leaves; because of their appearance, these species are called horsetails (Figure 4.21I,J) and are classified as the subgenus *Equisetum*. The other species, which lack extensive branching at the nodes, are classified as subgenus *Hippochaete* (Figure 4.21A,D). The two subgenera differ in stomate anatomy as well, those of subgenus *Hippochaete* being sunken, and those of subgenus *Equisetum* occurring at the (stem) surface.

At the tip of some aerial stems are **strobili** or **cones** (Figure 4.21E,L) containing the **sporangia**, which are pendant from a stalked, peltate structure called the **sporangio-phore** (Figure 4.21F,L). The sporangiophore is thought to represent an evolutionary fusion product of an aggregate of ancestrally distinct, recurved sporangia. Some species of *Equisetum*, e.g., *E. arvense*, are unusual in having two types of aerial stems: photosynthetic vegetative stems (Figure 4.21I,J) and nonphotosynthetic reproductive stems that terminate in strobili (Figure 4.21K,L). The spores of

Equisetum are unique among vascular plants in containing chloroplasts and unique among land plants in having four or more unusual appendages called **elaters** (Figure 4.21G,H). The elaters of *Equisetum* spores (which are not homologous with elaters in the sporangia of liverworts) are hygroscopic and uncurl from the spore body upon drying, aiding in spore dispersal.

MARATTIALES — MARATTIROID FERNS

The Marattiales are a group of about six genera and have traditionally been called ferns. They are very similar to the Polypodiales or leptosporangiate ferns (discussed later) in general form, having large pinnate or bipinnate leaves (Figure 4.22A,D) with circinate vernation, sporangia located on the abaxial surface of leaflet blades, and a photosynthetic gametophyte (see later discussion). However, the sporangia of the Marattiales are **eusporangiate**, like those of all vascular plants except for the leptosporangiate ferns. In some taxa of the Marattiales, the sporangia are fused into a common structure, a **synangium** (Figure 4.22B,C). A distinctive apomorphy of the Marattiales is the occurrence of a polycyclic siphonostele (Figure 4.1), which appears as concentric rings of siphonosteles in cross-section (the vasculature of which is, however, connected at a lower level).

POLYPODIALES — LEPTOSPORANGIATE FERNS

The Polypodiales (also known as Filicales or Pteridales) correspond to what are commonly known as the leptosporangiate ferns. Of the five major monilophyte groups, the

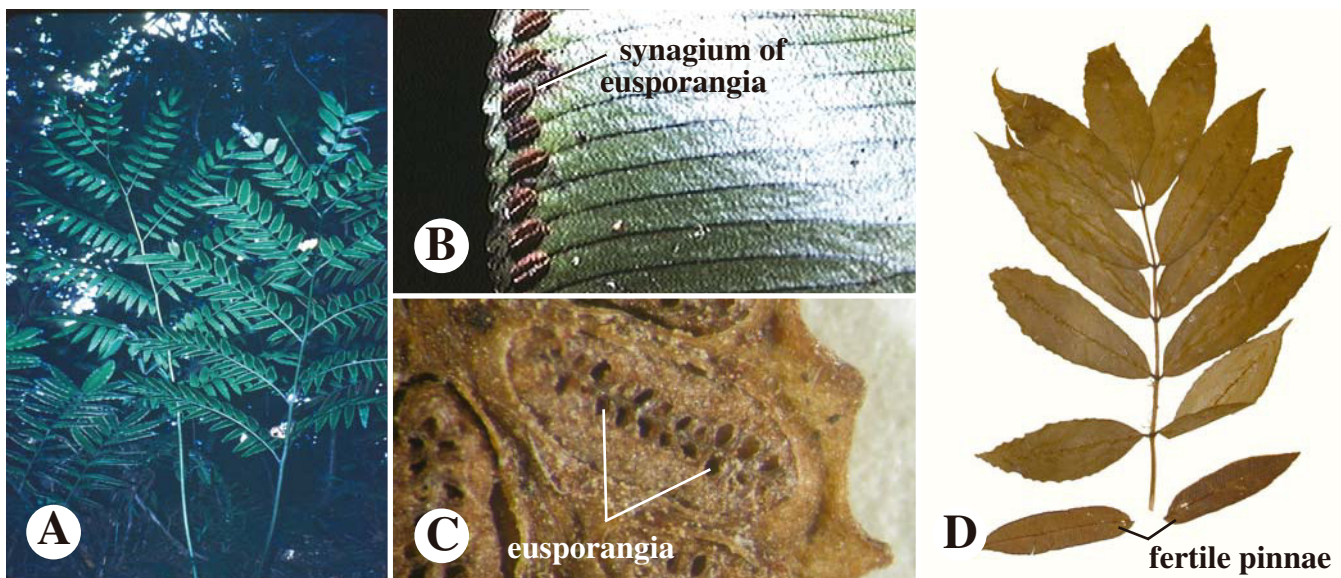


FIGURE 4.22 A–D. Marattiales. A,B. *Marattia* sp. A. Whole leaf, bipinnately compound. B. Close-up of synangium containing eusporangiate sporangia near margin on abaxial leaf surface. C,D. *Danaea* sp. C. Synangium containing several eusporangia. D. Pinnately compound leaf, with lower, fertile pinnae having numerous synangia on abaxial surface.

leptosporangiate ferns contain by far the greatest diversity, with more than 11,000 species.

Most leptosporangiate ferns have a horizontal stem, the **rhizome**, which is usually underground but may sprawl at ground level. Some leptosporangiate ferns have erect aerial stems, which in the so-called tree ferns (Figure 4.23G) can attain heights approaching 100 feet. A few ferns are vines (Figure 4.23E,F), with weak stems that sprawl on the ground or upon another plant. The leaves of ferns come in a great variety of forms (Figure 4.23, 4.24).

Like those of the Marattiales, the immature leaves of Polypodiales are coiled and known as **fiddleheads** or **croziers** (Figure 4.23A). This type of developmental morphology is called **circinate vernation**. Leptosporangiate ferns often have trichomes or scales on the rhizome or leaves, which are a valuable taxonomic character (Figure 4.23B). Circinate vernation with crozier formation may constitute an apomorphy for the Polypodiales and Marattiales together; however, this feature is also shared with the cyads of the seed plants (see Chapter 5).

Leptosporangiate fern leaves have a terminology slightly different from that of other vascular plants (see Chapter 9). The leaf itself is called a **frond**; the petiole is called a **stipe**; the first discrete leaflets or blade divisions of a fern leaf are called **pinnae** (singular **pinna**). If there is more than one division, the terms **1° pinna**, **2° pinna**, and so forth may be used. The ultimate leaflets or blade divisions are called **pinnules** (Figure 4.23C,D; see also Chapter 9).

The primary apomorphy of the Polypodiales is the **leptosporangium** (Figure 4.1, 4.24I). Leptosporangia are unique among vascular plants in (1) developing from a single cell, and (2) having a single layer of cells making up the sporangium wall. Leptosporangia are often aggregated into clusters, known as **sori** (singular **sorus**; Figure 4.24A,B,D), which may or may not be covered by a flap of tissue, the **indusium** (Figure 4.24E). Some species have an extension of the pinnule margin called a **false indusium** that overlaps the sorus (Figure 4.24F,G). In addition to general frond morphology, the position and shape of the sorus and indusium are useful taxonomic characters in delimiting the ferns. For example, the family Polypodiaceae are largely distinguished in being **exindusiate** (sori lacking an indusium), whereas other families, such as the Pteridaceae, are **indusiate** (sori having an indusium).

The leptosporangium may have been an important adaptation in the ferns because of a unique mechanism of spore dispersal. On the outer rim of the leptosporangium is a single row of specialized cells, collectively known as an **annulus**, in which the cell walls are differentially thickened on the inner cell face and on the cell faces between adjacent annular cells (Figure 4.24I, 4.25). As the leptosporangium matures and begins to dry, water evaporates from the cells of the annulus.

The force of capillarity causes the cells to buckle on the *outer* faces, as these are regions in which the cell wall is not thickened and therefore structurally weakest. This buckling provides a force resulting in splitting, or dehiscence, of the leptosporangium, followed by a backward retraction of the annulus (Figure 4.25). A short time after the annular cells fully retract, total evaporation of water within the cells causes the release of the capillarity tensile strength, which catapults the annulus forward, ejecting the spores in the process (Figure 4.25).

Leptosporangiate ferns, like all nonseed tracheophytes, have a haploid gametophyte phase that is free-living from the dominant sporophyte phase. The gametophytes are quite small and generally consist of a thin flat sheet of photosynthetic cells, which is variable (but often cordate) in shape. These bear several rootlike **rhizoids** as well as sperm-producing **antheridia** and egg-producing **archegonia** (Figure 4.25). As in all the nonflowering land plants, a sperm cell fertilizes an egg cell of the archegonium. The resultant zygote divides and differentiates into a new sporophyte, which initially remains attached to the gametophyte (Figure 4.25). Soon, however, the sporophyte attains independence of the gametophyte (which subsequently dies), the sporophyte becoming the persistent, dominant phase of the life cycle, a characteristic of all vascular plants (Figure 4.1).

The economic importance of leptosporangiate ferns is mostly as important ornamental cultivars in the horticultural trade. These include, among many others, species of *Adiantum* (maiden hair fern), *Asplenium* (e.g., *A. nidus*, bird's nest fern), *Cyathea* (a tree fern), and *Nephrolepis* (Boston fern, sword fern). Ostrich fern (*Matteuccia struthiopteris*) has edible croziers. *Pteris vittata* has recently been used to remove arsenic from toxic landfills. The family circumscription of the leptosporangiate ferns is still in flux and awaits further studies. See Pryer et al. (2004a) for a recent phylogenetic analysis of the group.

One group of leptosporangiate ferns is unusual in being aquatic, with members sometimes cultivated in small ponds or aquaria. These so-called water ferns comprise two major subgroups. One group is composed of three genera, *Pilularia* (Figure 4.26D,E), *Regnellidium*, and *Marsilea*, the water clover (Figure 4.26A), the latter considered by some to be the true four-leaf clover. Another subgroup of water ferns contains two genera, *Salvinia*, the water spangles (Figure 4.26A,B), and *Azolla*, the mosquito fern; Figure 4.26A C). *Azolla* is interesting in having a symbiotic, nitrogen-fixing cyanobacterium living inside leaf clefts of the plant. Because the cyanobacterium ultimately adds nitrogen to the environment, rice farmers in parts of the world have learned to seed the paddies with *Azolla*, ensuring a better crop.

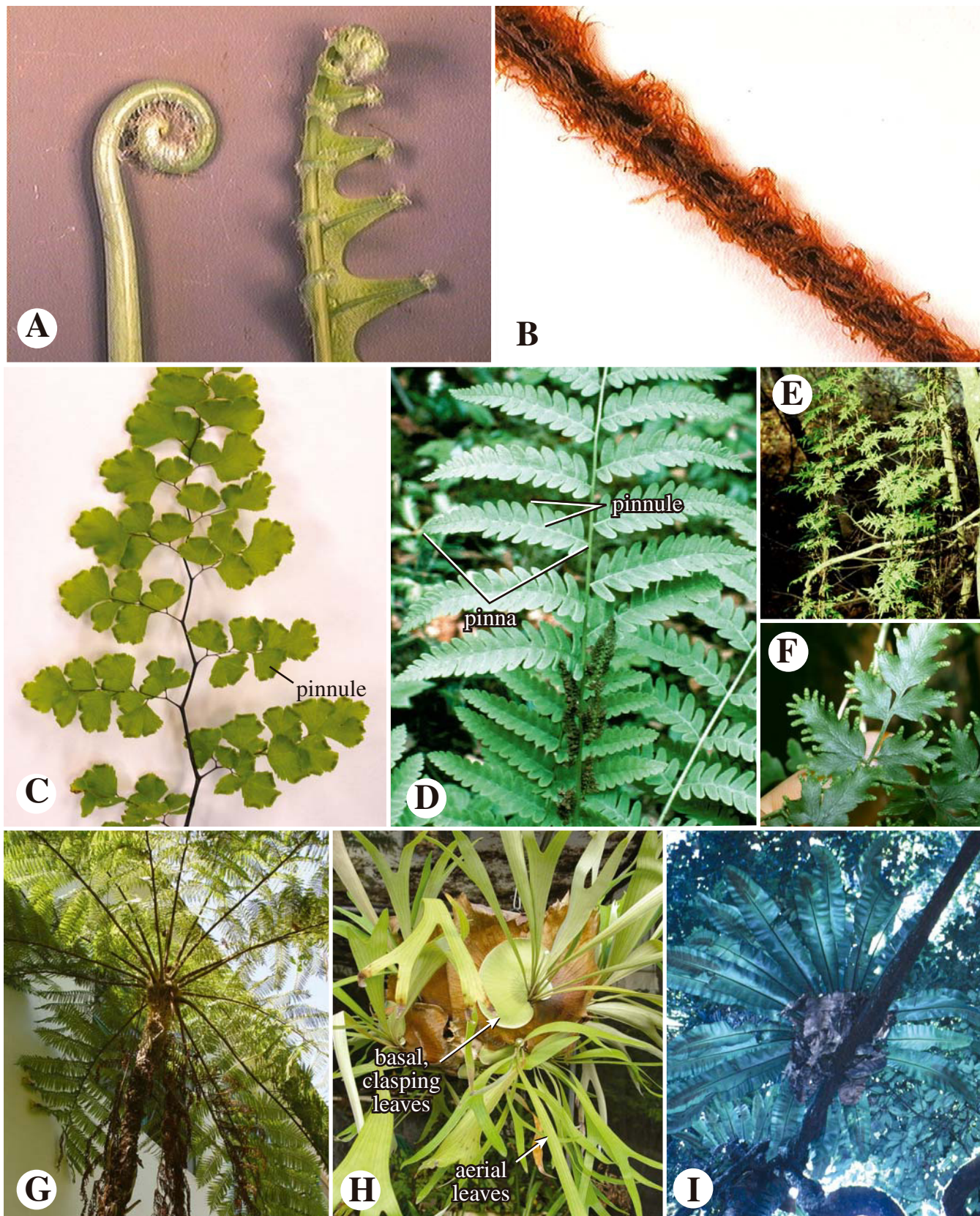


FIGURE 4.23 Polypodiales leptosporangiate ferns. **A.** *Polypodium aureum*, showing croziers or fiddleheads in early (left) and later (right) stages. **B.** *Nephrolepis cordifolia*, showing scales at base of stipe. **C.** *Adiantum capillus-veneris*, with bipinnate leaf, having ultimate pinnules. **D.** *Osmunda claytoniana*, with bipinnate leaves bearing pinnules, divided into pinnules. **E,F.** *Lygodium japonicum*, an epiphytic vine. **G.** *Cyathea cooperi*, a tree fern with erect, elongate stems. **H.** *Platycerium* sp., staghorn fern, epiphytic with dimorphic leaves. **I.** *Asplenium nidus*, bird's nest fern, epiphytic with simple leaves.

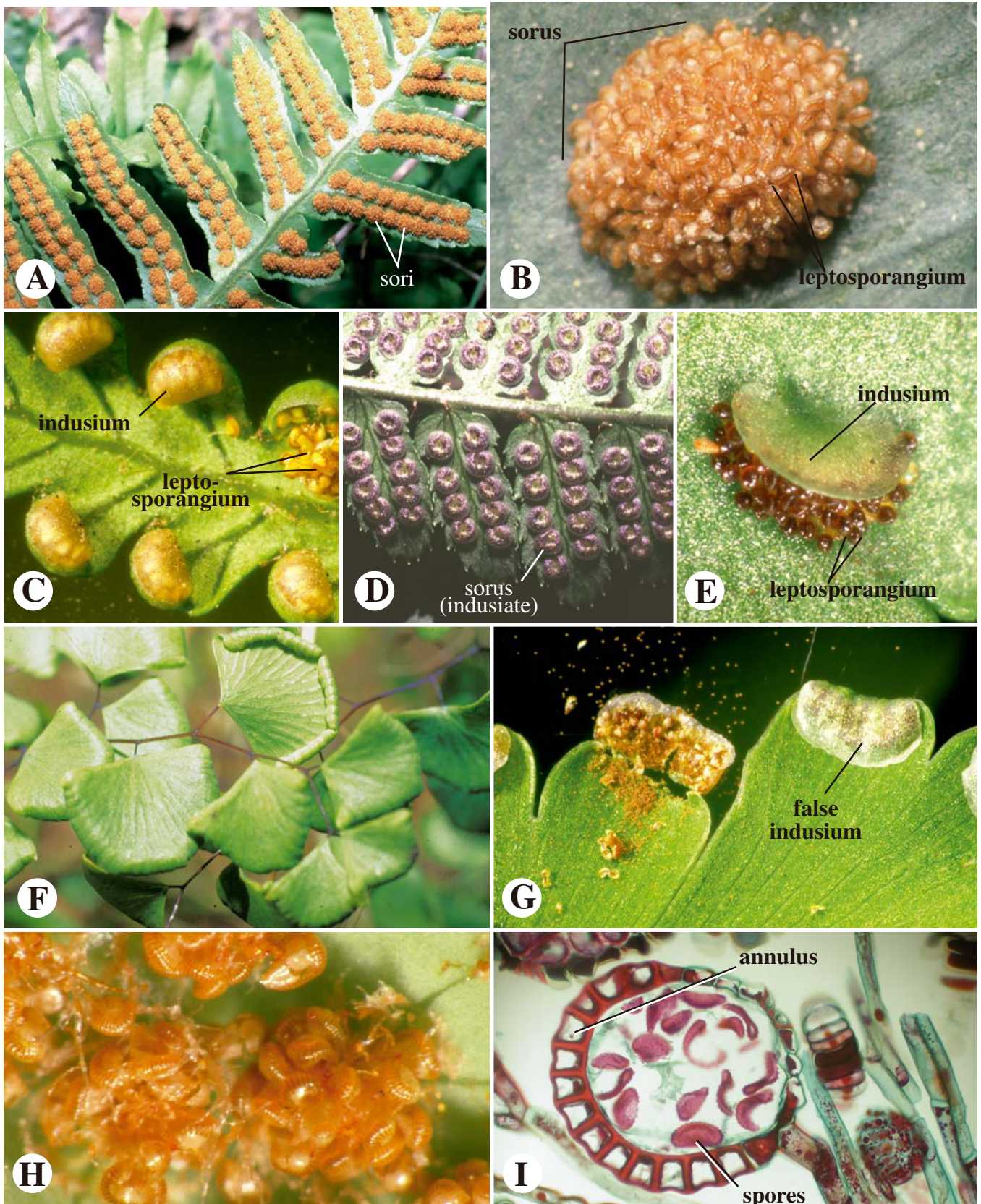


FIGURE 4.24 Polypodiales Leptosporangiate ferns. **A.** *Polypodium californicum*, an indusiate species. **B.** *Polypodium aureum*, sorus close-up. **C.** *Cibotium* sp., a tree fern, showing indusia at margin of pinnules. **D.** *Dryopteris arguta*, with orbicular-reniform indusiate sori on leaf surface. **E.** *Nephrolepis cordifolia*, close-up of indusium and sorus of leptosporangia. **F.** *Adiantum jordani*, with false indusia. **G.** *Adiantum capillus-veneris*, close-up of false indusia. **H.** Close-up of leptosporangia. **I.** Leptosporangium in sagittal section, showing annulus and internal spores. Note single cell layer of wall.

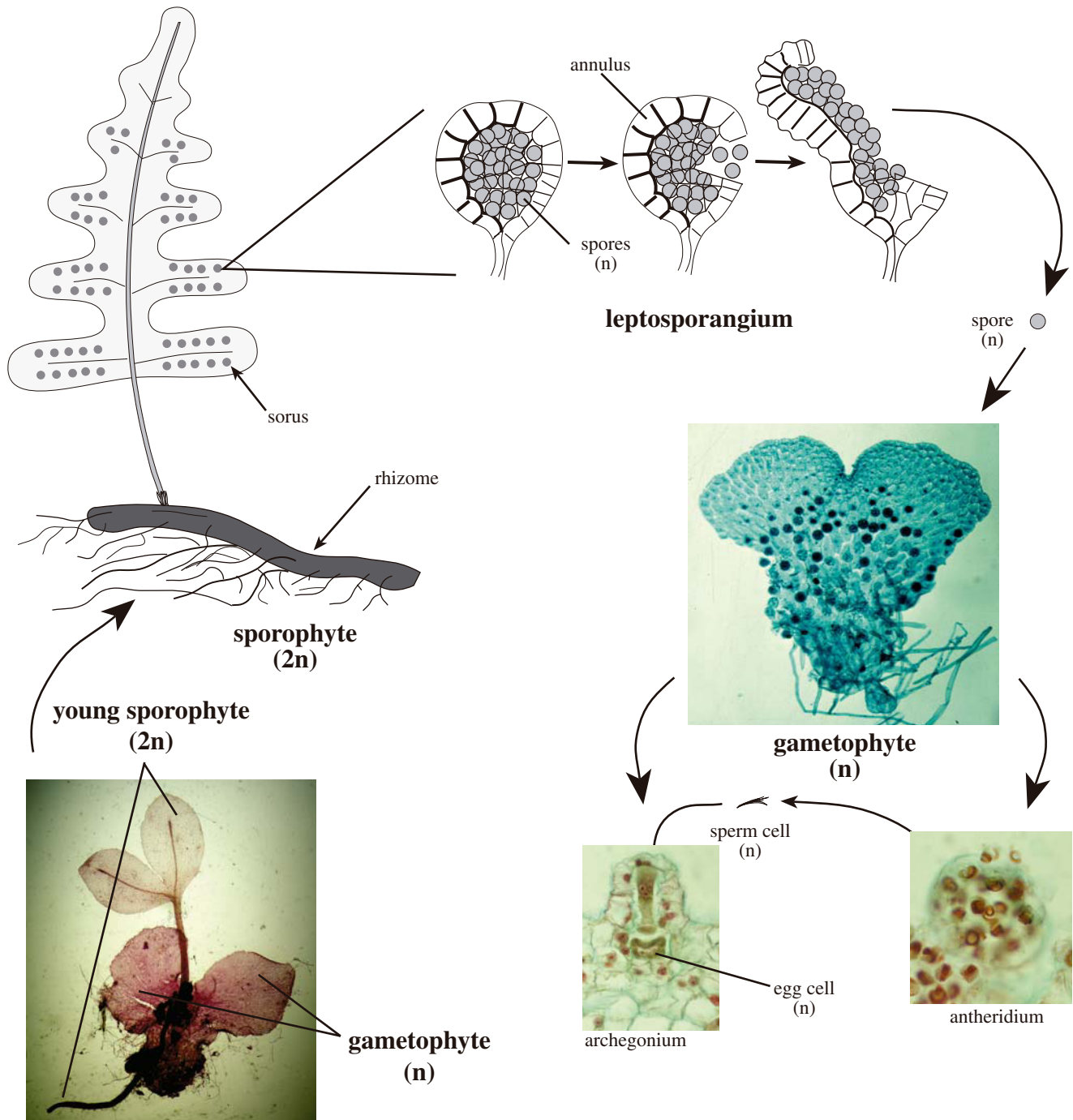


FIGURE 4.25 Life cycle of leptosporangiate ferns. Note mechanism of spore dispersal, gametophyte development, fertilization, and sporophyte development.

All of the aquatic ferns are virtually unique among the leptosporangiate ferns in being **heterosporous**. Recall that **heterospory** is the development of two types of spores, male and female. From these spores develop the male and female gametophytes, which are **endosporic**, similar to *Selaginella* and *Isoetes* of the lycophytes. The reproductive structures of

these aquatic ferns are complicated and are organized into generally spherical **sporocarps** (Figure 4.26E). The sporocarps allow the sporangia and spores to remain dormant for long periods of time, an adaptation that enables them to survive and persist when the ponds or pools where these plants are found dry up.

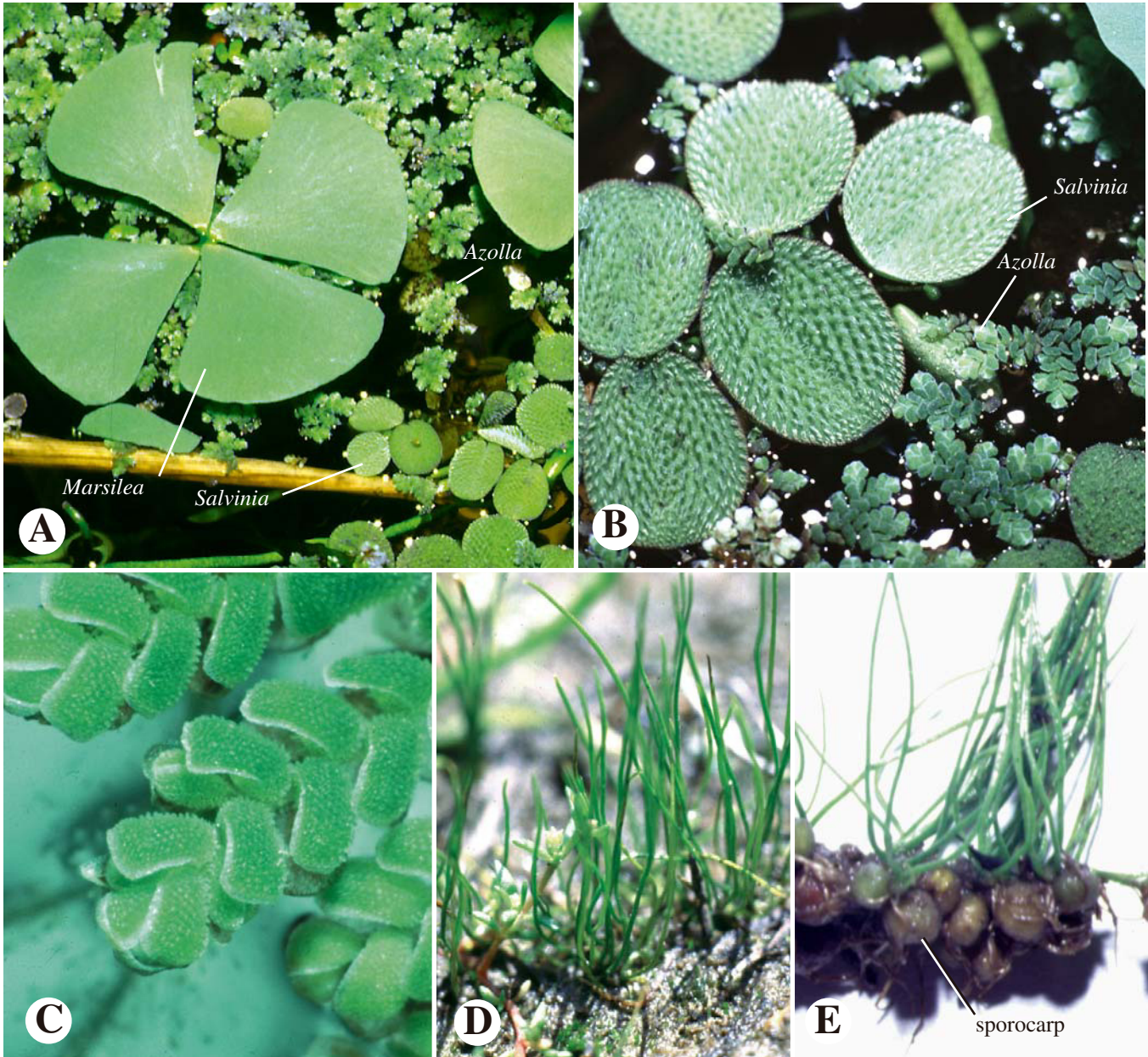


FIGURE 4.26 Polydiales aquatic leptosporangiate ferns. **A.** *Azolla*, *Marsilea*, and *Salvinia*, growing together. **B.** *Salvinia* and *Azolla*, close-up. **C.** Close-up of *Azolla*, with lower leaf cavities housing a blue-green alga. **D,E.** *Pilularia americana*. **D.** Habit. **E.** Close-up of rhizome bearing spherical sporocarps.

REVIEW QUESTIONS

VASCULAR PLANT APOMORPHIES

1. What is the formal, scientific name for the vascular plants?
2. Name the major apomorphies of the vascular plants.
3. How was the evolution of lignin a major adaptive feature of the vascular plants?
4. What is the difference between a primary and secondary cell wall in terms of time of deposition and chemistry?
5. What is a pit? a primary pit field?

6. Is the secondary cell wall formed inside or outside the plasma membrane? inside or outside the primary cell wall?
7. What are the general characteristics of sclerenchyma cells?
8. Name the two types of sclerenchyma and state how they differ.
9. How are sclerenchyma and tracheary elements similar? How do they differ?
10. What is the function of tracheary elements?
11. What is xylem?
12. Name the two types of tracheary elements and cite how they differ structurally.
13. What is a perforation plate?
14. In what taxa are vessels found?
15. What is the function of sieve elements?
16. What is phloem?
17. What is a sieve area and what compound is associated with them?
18. What is the difference, in morphology and taxonomic group found, between a sieve cell and a sieve tube member?
19. What is the endodermis and Casparian strip, and what is the function of these?
20. How are sporophytes of the vascular plants different from those of the liverworts, hornworts, and mosses?
21. What is the definition and function of a stem?
22. What is a stele?
23. What is the ancestral stelar type in the vascular plants and what is its structural anatomy?
24. What is the general morphology and function of leaves?
25. What are the internal, chlorophyllous cells of a leaf called? Into what two layers are these cells typically formed?
26. What is a vein?
27. What is a shoot?
28. What is the name of the region of actively dividing cells in the shoot?
29. What is the definition of a bud?
30. Where are buds typically located?
31. Define node; internode.
32. What is the function of roots?
33. What is the name of the region of actively dividing cells in the root?
34. What is the function of: (a) rootcap; (b) root hairs; (c) endodermis/Casparian strips?
35. What are the major differences between roots and stems?
36. What are mycorrhizae?

VASCULAR PLANT DIVERSITY

37. What is the most basal (earliest diverging) lineage of the vascular plants, now extinct?
38. What are the major apomorphies of the lycophytes?
39. What fossil lycophyte was a large tree in the Carboniferous and now makes up a large percentage of coal deposits?
40. What is a lycophyll (microphyll)? an enation?
41. What is the position of the sporangia in lycophytes?
42. What is a sporophyll? a strobilus?
43. What is homosporous? Name two genera of lycophytes that have this condition.
44. Name and define the two types of leaf morphology in *Selaginella* species.
45. Name two genera of extant lycophytes that are heterosporous. What structure is associated with their leaves?
46. Define the terms heterosporous and endosporic.
47. Name the apomorphies of the euphyllophytes, and list the two major, vascular plant groups included.
48. Name the putative apomorphies of the monilophytes, and list the five major groups contained within it.
49. What is distinctive about the leaves of the ophioglossid ferns?
50. What is a eusporangium?
51. What two features may link the ophioglossid ferns with the psilophytes (whisk ferns)?
52. What is distinctive (and apomorphic) about the roots, leaves, and sporangia of the Psilotales?
53. What is the most commonly cultivated species of whisk fern?

54. What is a fossil member of Equisetales, making up a component of coal deposits.
55. Name the major apomorphies of the Equisetales.
56. What is the only extant genus of this group?
57. What do equisetophytes have as a component of the cell wall?
58. What is the difference between a scouring rush and a horsetail? Into what two subgenera are these classified?
59. Describe the morphology of the strobilus (cone), sporangiophore, and sporangia of *Equisetum*.
60. What is unique about the spores of *Equisetum*? What is the function of this novelty?
61. Describe the diagnostic features and a putative apomorphy of the Marattiales.
62. How do the gametophytes and leaf development of the Marattiales resemble the Polypodiales (below)?
63. What type of sporangium is found in the Marattiales?
64. What is the major evolutionary novelty of the Polypodiales? Describe its development and morphology.
65. Name three stem types/habits that occur in the Polypodiales.
66. What is circinate vernation? What terms are used for immature fern leaves that exhibit this?
67. Define frond, stipe, pinna, pinnule.
68. Define sorus, indusium, false indusium, annulus.
69. In a fern gametophyte, what is the name of the male gametangium? the female gametangium? What do they look like?
70. Name three or more genera of aquatic ferns.
71. What reproductive features unite the aquatic ferns?

EXERCISES

1. Peruse the most recent literature on phylogenetic relationships of the vascular plants. Are there any differences relative to Figure 4.1?
2. Peruse botanical journals and find a systematic article on a nonseed vascular plant (e.g., a leptosporangiate fern or fern group). What is the objective of the article and what techniques were used to address it? What types of morphological characters are discussed by the author(s)?
3. Collect and identify local lycophytes, equisetophytes, psilophytes, ophioglossoid ferns, or leptosporangiate ferns. What diagnostic features are used to distinguish between species?

REFERENCES FOR FURTHER STUDY

- Cracraft, J., and M. J. Donoghue. 2004. *Assembling the Tree of Life*. Oxford University Press, New York.
- Des Marais, D. L., A. R. Smith, D. M. Britton, and K. M. Pryer. 2003. Systematics phylogenetic relationships and evolution of extant horsetails, *Equisetum*, based on chloroplast DNA sequence data (*rbcL* and *trnL-F*). *International Journal of Plant Sciences* 164: 737-751.
- Foster, A. S., and E. M. Gifford. 1974. *Comparative morphology of vascular plants*, 2nd edition. W. H. Freeman, San Francisco.
- Flora of North America Editorial Committee. 1993+. *Pteridophytes and Gymnosperms. Volume 2, in Flora of North America North of Mexico*. 7+ vols. New York and Oxford.
- Gensel, P. G., and C. M. Berry. 2001. Early lycophyte evolution. *American Fern Journal* 91: 74-98.
- Gifford, E. M., and A. S. Foster. 1989. *Morphology and evolution of vascular plants*, 3rd edition. W. H. Freeman and Co., New York.
- Kenrick, P., and P. R. Crane. 1997. *The Origin and Early Diversification of Land Plants: a Cladistic Study*. Smithsonian Institution Press, Washington, DC.
- Lellinger, D. B. 1985. *A Field Manual of the Ferns and Fern-Allies of the United States and Canada*. Smithsonian Institution Press, Washington, DC.
- Mickel, John T. 1979. *How to Know the Ferns and Fern Allies*. Wm. C. Brown, Dubuque, IA.
- Pryer, K. M., A. R. Smith, and J. E. Skog. 1995. Phylogenetic relationships of extant ferns based on evidence from morphology and *rbcL* sequences. *American Fern Journal* 85: 205-282.
- Pryer, K. M., H. Schneider, A. R. Smith, R. Cranfill, P. G. Wolf, J. S. Hunt, and S. D. Sipes. 2001. Horsetails and ferns are a monophyletic group and the closest living relatives to seed plants. *Nature* 409: 618-622.
- Pryer, K. M., E. Schuettpelz, P. G. Wolf, H. Schneider, A. R. Smith, and R. Cranfill. 2004a. Phylogeny and evolution of ferns (monilophytes) with a focus on the early leptosporangiate divergences. *American Journal of Botany* 91: 1582-1598.
- Pryer, K. M., H. Schneider, and S. Magallon. 2004b. The radiation of vascular plants. In J. Cracraft and M. J. Donoghue (eds.). *Assembling the Tree of Life*, pp. 138-153. Oxford University Press, London.

- Schneider, H., E. Schuettpelz, K. M. Pryer, R. Cranfill, S. Magallon, and R. Lupia. 2004. Ferns diversified in the shadow of angiosperms. *Nature* 428: 553-557.
- Schneider, H., K. M. Pryer, R. Cranfill, A. R. Smith, and P. G. Wolf. 2002. Evolution of vascular plant body plans: a phylogenetic perspective. *Systematics Association special volume* 65: 330-364.
- Stewart, W. N., and G. W. Rothwell. 1993. *Paleobotany and the Evolution of Plants*, 2nd edition. Cambridge University Press, Cambridge, UK.
- Wikström, N. 2001. Diversification and relationships of extant homosporous lycopods. *American Fern Journal* 91:150-165.
- Wolf, P. G., K. M. Pryer, A. R. Smith, and M. Hasebe. 1998. Phylogenetic studies of extant pteridophytes. In: *Molecular Systematics of Plants*, 2nd ed. P. S. Soltis, D. E. Soltis, and J. J. Doyle (eds.). Chapman and Hall, New York.

This Page Intentionally Left Blank

5

EVOLUTION AND DIVERSITY OF WOODY AND SEED PLANTS

LIGNOPHYTES—WOODY PLANTS.....	97	Pteridosperms Seed Ferns	107
SPERMATOPHYTES—SEED PLANTS.....	98	Gymnosperms.....	107
Seed Evolution	98	Cycadophyta Cycads	108
Pollen Grains	101	Ginkgophyta <i>Ginkgo</i>	109
Pollen Tube.....	101	Coniferophyta Conifers	109
Pollination Droplet.....	101	Gnetales	114
Ovule and Seed Development	104	REVIEW QUESTIONS	118
Seed Adaptations	105	EXERCISES.....	119
Eustele	105	REFERENCES FOR FURTHER STUDY.....	119
DIVERSITY OF WOODY AND SEED PLANTS.....	107		
<i>Archeopteris</i>	107		

LIGNOPHYTES—WOODY PLANTS

The lignophytes, or woody plants (also called Lignophyta), are a monophyletic lineage of the vascular plants that share the derived features of a **vascular cambium**, which gives rise to **wood**, and a **cork cambium**, which produces **cork** (Figures 5.1, 5.2). These features also occurred in now extinct lineages within the lycophytes (e.g., *Lepidodendron*) and equisetophytes (e.g., *Calamites*), but are thought to have been derived independently in these taxa. A vascular cambium is a sheath, or hollow cylinder, of cells that develops within the stems and roots as a continuous layer, between the xylem and phloem in extant, eustelic spermatophyte (see later discussion). The cells of the vascular cambium divide mostly in a tangential plane, resulting initially in two layers of cells (Figure 5.3). One of these layers remains as the vascular cambium and continues to divide indefinitely; the other layer eventually differentiates into either **secondary xylem = wood**, if produced to the inside of the cambium, or **secondary phloem**, if produced to the outside of the cambium (Figure 5.3, 5.4). Generally, much more secondary xylem is

produced than secondary phloem. As secondary tissue is formed, the inner cylinder of wood expands (Figures 5.4, 5.5). Many woody plants have regular growth periods, e.g., forming annual rings of wood (Figure 5.5). A cork cambium is similar to a vascular cambium, only it differentiates near the periphery of the stem or root axis. The cork cambium and its derivatives constitute the **periderm** (referred to as the outer bark). The outermost layer of the periderm is **cork** (Figure 5.4). Cork cells contain a waxy polymer called **suberin** (similar to cutin) that is quite resistant to water loss (see Chapter 10).

The vascular cambium and cork cambium were a major evolutionary novelty. Secondary xylem, or wood, functions in structural support, enabling the plant to grow tall and acquire massive systems of lateral branches. Thus, the vascular cambium was a precursor to the formation of intricately branched shrubs or trees with tall overstory canopies (e.g., Figure 5.2), a significant ecological adaptation. Cork produced by the cork cambium functions as a thick layer of cells that protects the delicate vascular cambium and secondary phloem from mechanical damage, predation, and desiccation.

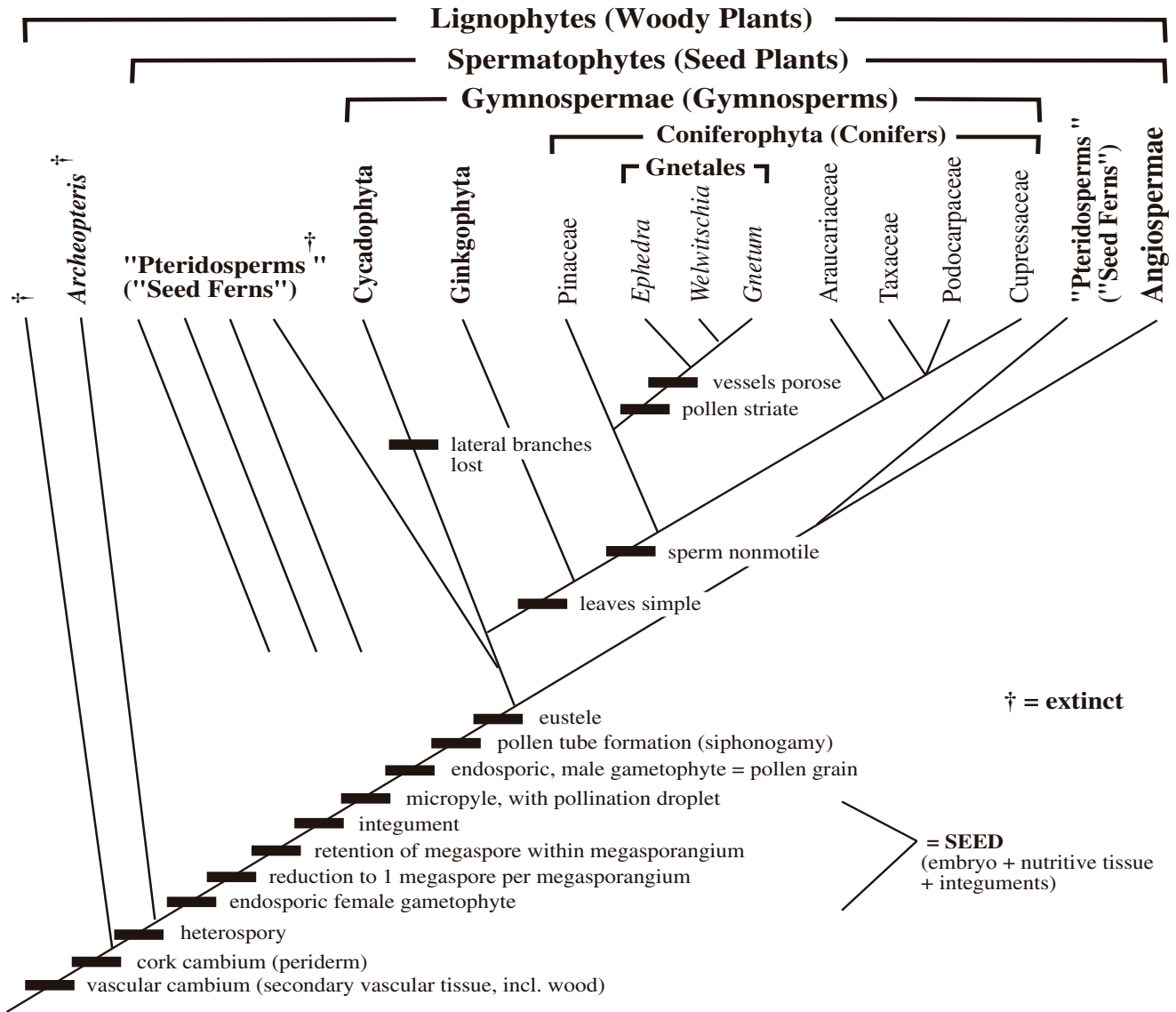


FIGURE 5.1 Cladogram of the woody and seed plants. Major apomorphies are indicated beside a thick hash mark. Modified from Bowe et al., 2000; Chaw et al., 2000; Frohlich et al., 2000; and Samigullin et al., 1999.

Wood anatomy can be quite complex. The details of cellular structure are important characters used in the classification and identification of woody plants. Wood anatomical features may also be used to study the past, a specialty known as **dendrochronology** (see Chapter 10).

SPERMATOPHYTES—SEED PLANTS

The spermatophytes, or seed plants (also called Spermato-phyta), are a monophyletic lineage within the lignophytes (Figure 5.1). The major evolutionary novelty that unites this group is the **seed**. A seed is defined as an **embryo**, which is an

immature diploid sporophyte developing from the zygote, surrounded by **nutritive tissue** and enveloped by a **seed coat** (Figure 5.6). The embryo generally consists of an immature root called the **radicle**, a shoot apical meristem called the **epicotyl**, and one or more young seed leaves, the **cotyledons**; the transition region between root and stem is called the **hypocotyl** (Figures 5.6, 5.12). An immature seed, prior to fertilization, is known as an **ovule**.

SEED EVOLUTION

The evolution of the seed involved several steps. The exact sequence of these is not certain, and two or more steps in seed evolution may have occurred concomitantly.



FIGURE 5.2 Composite photograph of *Sequoiadendron giganteum*, a woody conifer that is the most massive organism on Earth, and among the tallest of trees.

The probable steps in seed evolution are as follows (Figure 5.7):

1. **Heterospory.** Heterospory is the formation of *two* types of haploid spores within *two* types of sporangia: large, fewer-numbered **megaspores**, which develop via meiosis in the **megasporangium**, and small, more numerous **microspores**, the products of meiosis in the **microsporangium** (Figures 5.7, 5.8). The ancestral condition, in which a single spore type forms, is called **homospory**. Each megaspore develops into a **female gametophyte** that bears only archegonia; a microspore develops into a **male gametophyte**, bearing only antheridia. Although heterospory was prerequisite to seed evolution, there are fossil plants that were heterosporous but had not evolved seeds, among these being species of *Archeopteris* (Figure 5.1, 5.15A; see later discussion). Note that heterospory has evolved *independently* in other, nonseed plants, e.g., in the extant lycophytes *Selaginella* and *Isoetes* and in the water ferns (Chapter 4).
2. **Endospory.** Endospory is the complete development of, in this case, the female gametophyte *within the original spore wall* (Figure 5.7). The ancestral condition, in which the spore germinates and grows as an external gametophyte, is called **exospory**.
3. **Reduction of megaspore number to one.** Reduction of megaspore number occurred in two ways. First, there evolved a reduction in the number of cells within the megasporangium that undergo meiosis (each termed a **megasporocyte** or **megaspore mother cell**) was reduced to one (Figure 5.7). After meiosis, the single diploid megasporocyte gives rise to four haploid megaspores. Second, of the four haploid megaspores produced by meiosis, *three consistently abort*, leaving only one functional megaspore. This single megaspore also undergoes a great increase in size, correlated with the increased availability of space and resources in the megasporangium.
4. **Retention of the megaspore.** Instead of the megaspore being released from the sporangium (the ancestral condition, as occurs in all homosporous nonseed plants), in seed plants it is retained within the megasporangium (Figure 5.7). This was accompanied by a reduction in thickness of the megaspore wall.
5. **Evolution of the integument.** Most likely, the final event in seed evolution was the envelopment of the megasporangium by tissue, called the **integument** (Figure 5.7). The integument grows from the base of the **megasporangium** (which is often called a **nucellus** when surrounded by an integument) and surrounds it, except at the distal end. Fossil evidence suggests that integuments may have evolved first as separate lobes. In all extant seed plants, however, the integument develops as a continuous sheath

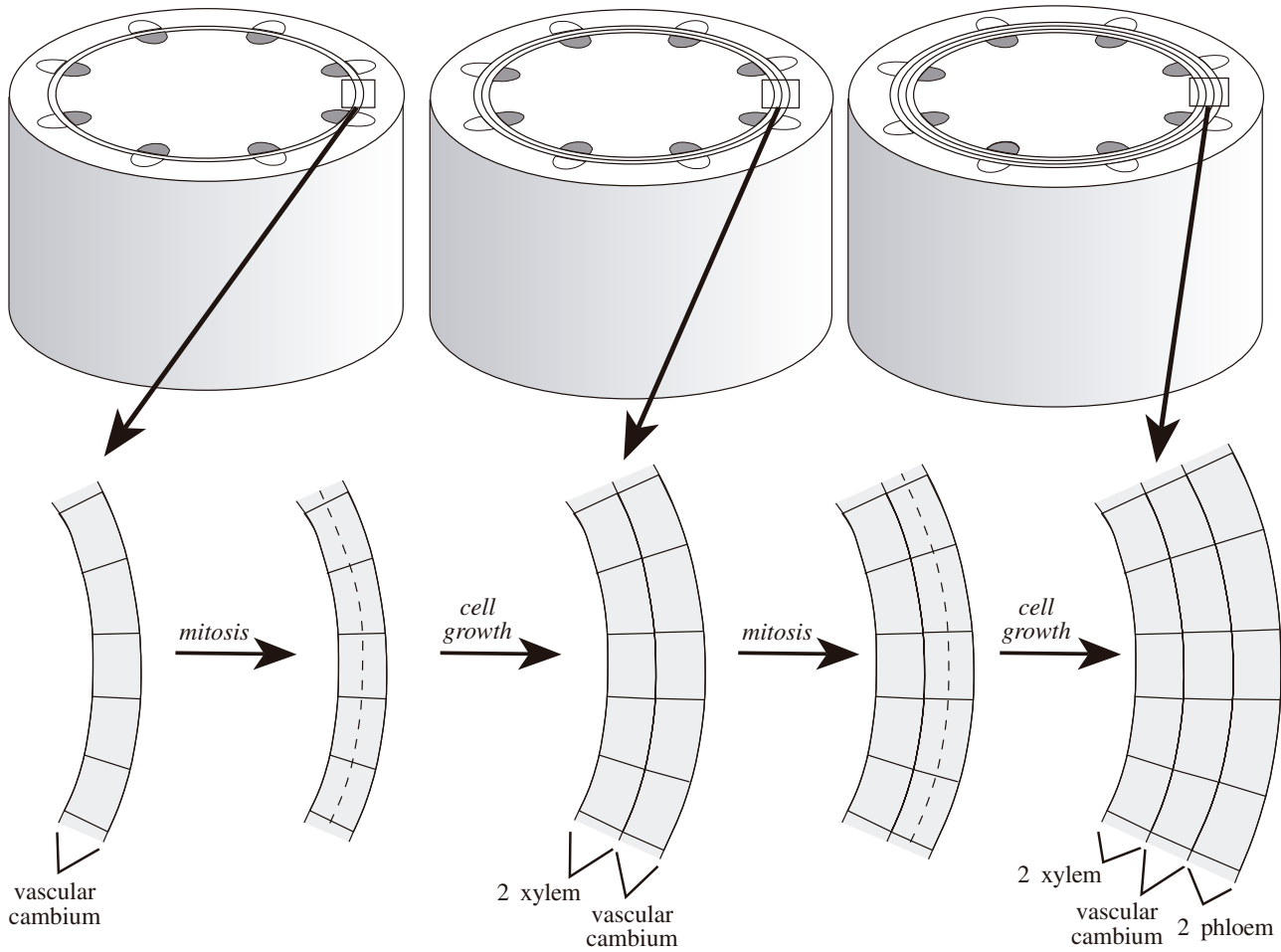


FIGURE 5.3 Development of the vascular cambium.

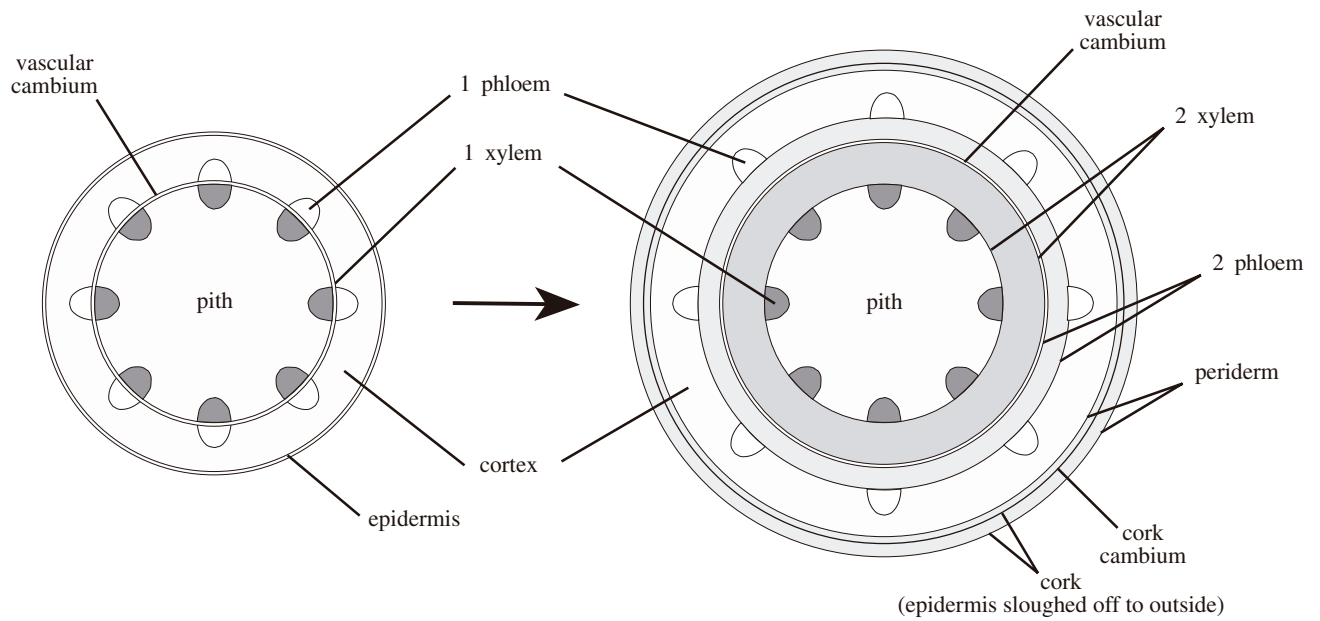


FIGURE 5.4 Development of secondary vascular tissue in the stem, illustrated here for a eustelic stem.

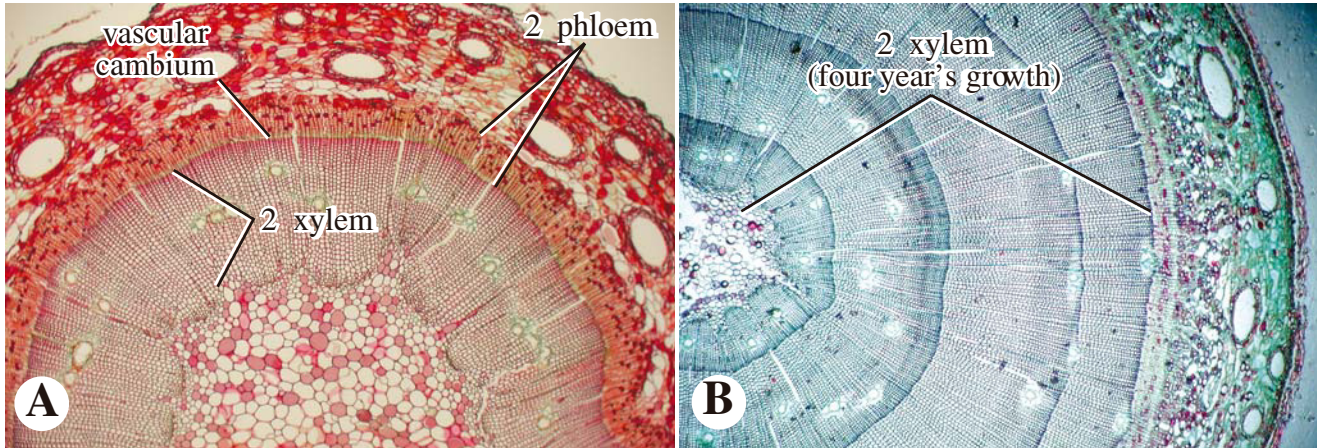


FIGURE 5.5 Woody stem cross-section, *Pinus* sp. **A.** One year's growth. **B.** Four years' growth.

and completely surrounds the nucellus except for a small pore at the distal end called the **micropyle**. The micropyle functions as the site of entry of pollen grains (or in angiosperms, of pollen tubes), which effect fertilization of the egg (see later discussion). The micropyle also functions in the mechanics of pollination droplet formation and resorption (see later discussion). Note that a single integument represents the ancestral condition of spermatophytes; in angiosperms a second integument layer evolved (Chapter 6).

POLLEN GRAINS

Concomitant with the evolution of the seed was the evolution of **pollen grains** (Figure 5.9). A pollen grain is, technically, an immature, endosporic male gametophyte. **Endospory** in pollen grain evolution was similar to the same process in seed evolution, involving the development of the male gametophyte within the original spore wall. Pollen grains of seed plants are extremely reduced male gametophytes, consisting of only

a few cells. They are termed immature male gametophytes because, at the time of their release, they have not fully differentiated.

After being released from the microsporangium, pollen must be transported to the micropyle of the ovule (or, in angiosperms, to the stigmatic tissue of the carpel; see Chapter 6) in order to ultimately effect fertilization. Wind dispersal, in combination with an ovule pollination droplet (see later subsection), was probably the ancestral means of pollen transport. After being transported to the ovule (or stigmatic tissue), the male gametophyte completes development by undergoing additional mitotic divisions and differentiation. The male gametophyte grows an exosporic **pollen tube**, which functions as a haustorial organ, obtaining nutrition by absorption from the surrounding sporophytic tissue (Figure 5.10; see **Pollen Tube**).

POLLEN TUBE

The male gametophytes of all extant seed plants form a pollen tube (Figure 5.10) soon after the pollen grains make contact with the megasporangial (nucellar) tissue of the ovule. The formation of pollen tubes is termed **siphonogamy** (*siphono*, tube + *gamos*, marriage). The pollen tubes, which may become branched in some taxa, function as a haustorial organ, growing into and feeding from the megasporangial (nucellar) tissue. Pollen tubes also function to deliver the sperm cells, directly or indirectly, to the egg of the ovule (see later discussion).

POLLINATION DROPLET

One possible evolutionary novelty associated with seed evolution is the **pollination droplet**. This is a droplet of liquid that is secreted by the young ovule through the micropyle

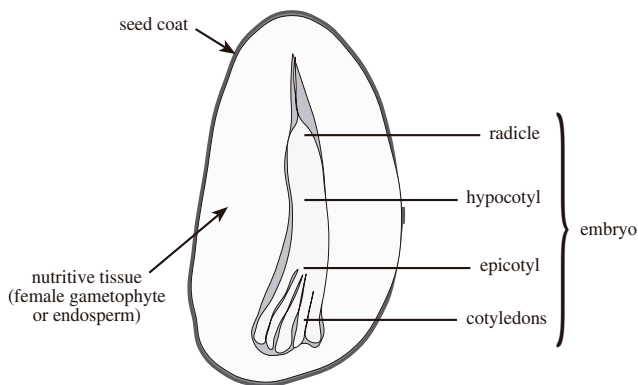
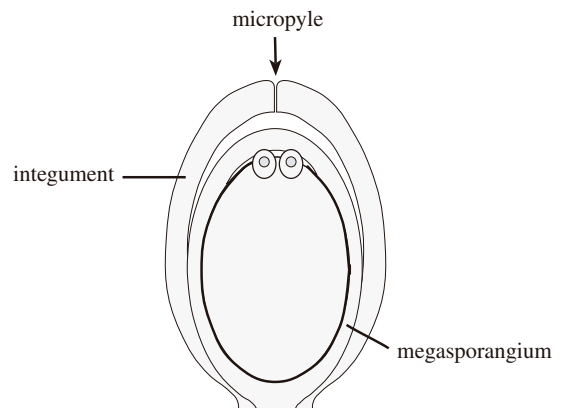
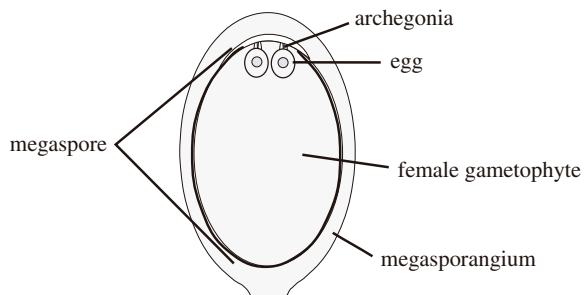
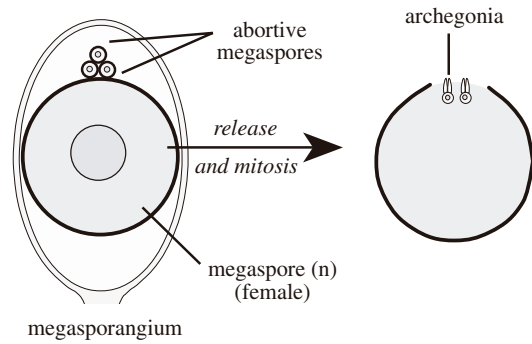
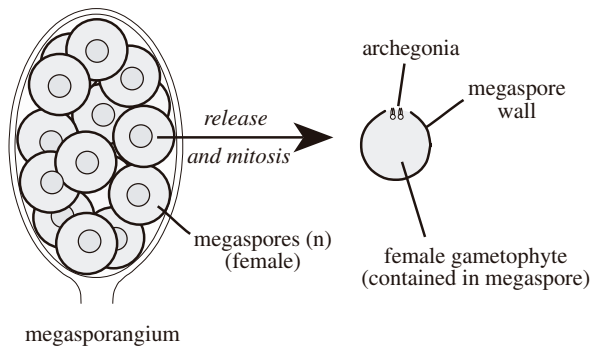
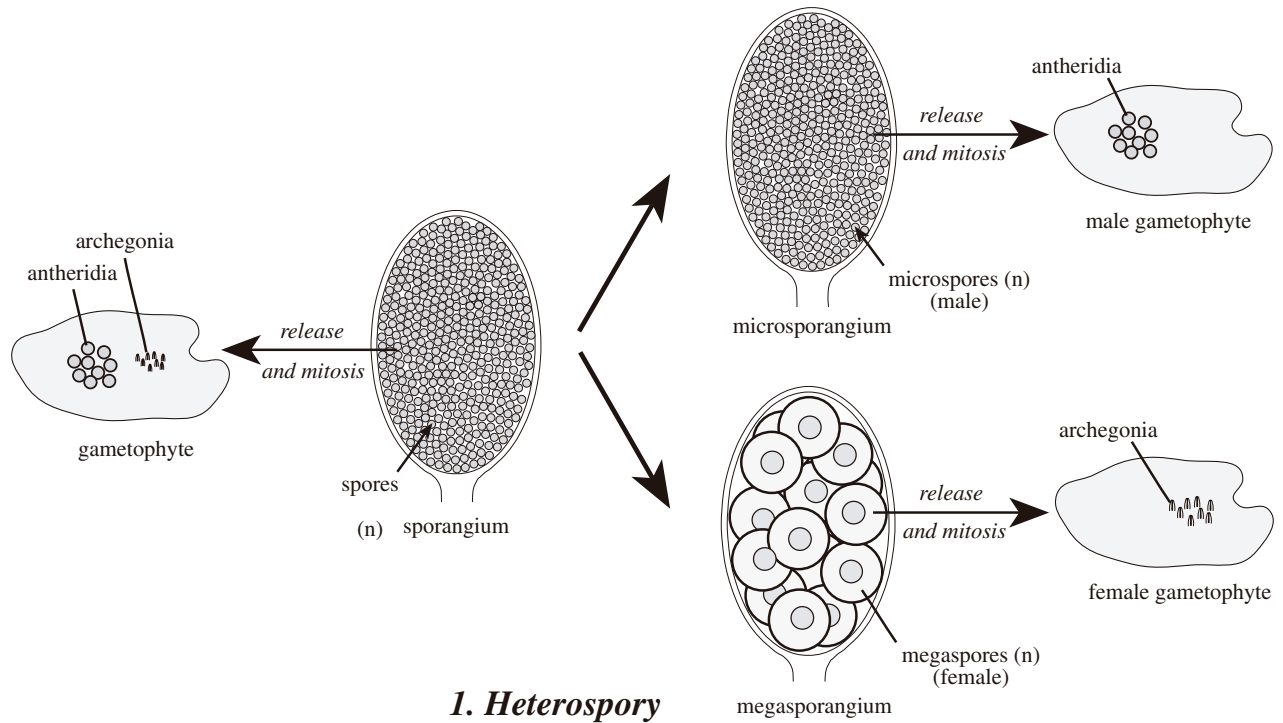


FIGURE 5.6 Morphology of a seed. *Pinus* sp. illustrated here.



4. Retention of megaspore

5. Evolution of Integument

FIGURE 5.7 Ovule and seed evolution in the spermatophytes (hypothetical, for purpose of illustration).

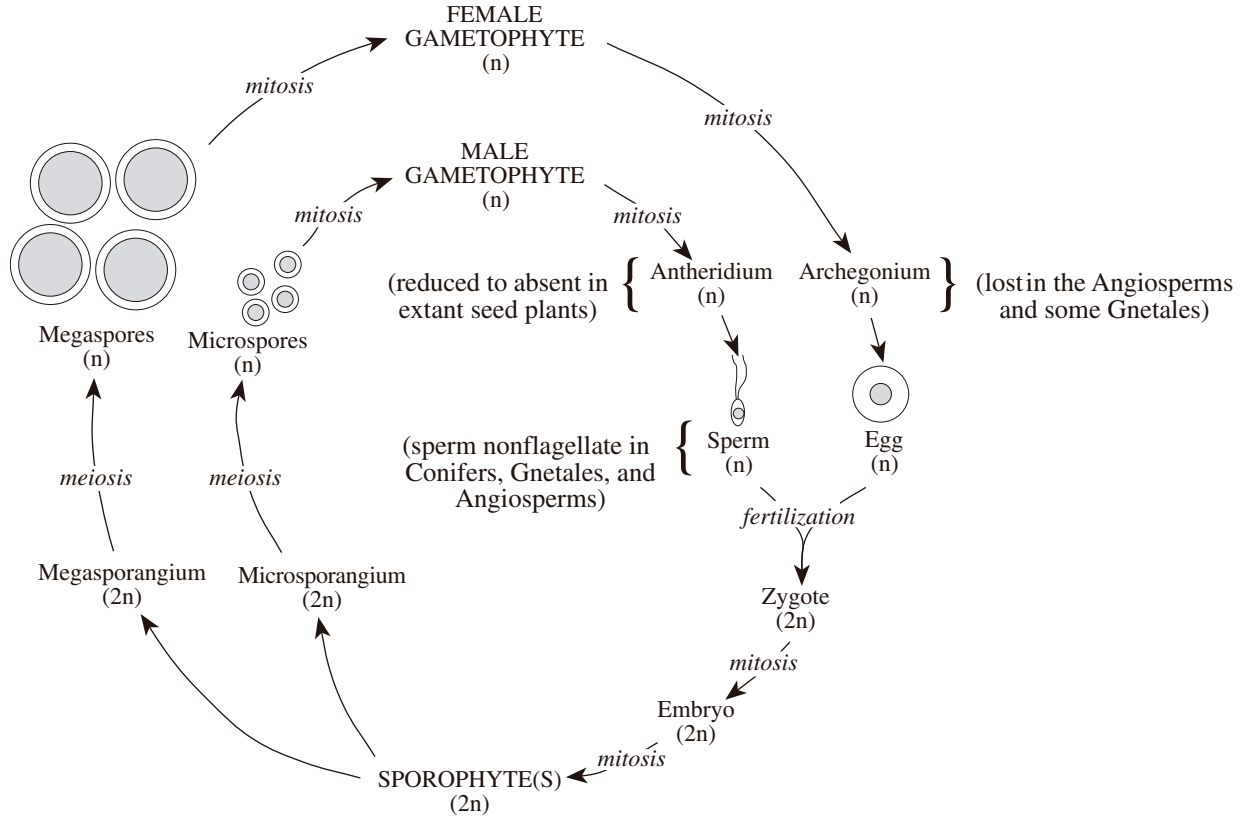


FIGURE 5.8 Life cycle of heterosporous plants.

(Figure 5.11). This droplet is mostly water plus some sugars or amino acids and is formed by the breakdown of cells at the distal end of the megasporangium (nucellus). The cavity formed by this breakdown of cells is called the **pollination chamber** (Figure 5.11). The pollination droplet functions in transporting pollen grains through the micropyle. This occurs

by resorption of the droplet, which pulls pollen grains that have contacted the droplet into the pollination chamber. It is unknown whether a pollination droplet was present in the earliest seed plants. However, the presence of a pollination droplet in many nonflowering seed plants suggests that its occurrence may be ancestral for at least extant seed

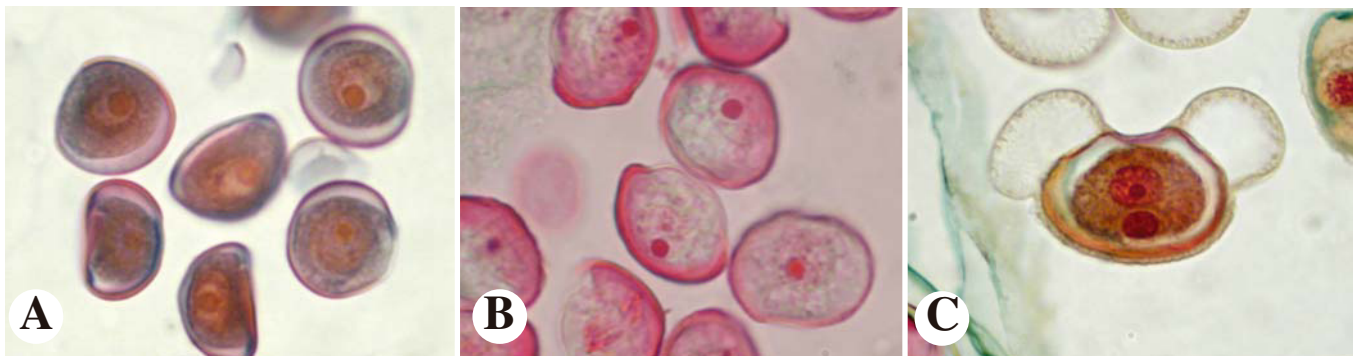


FIGURE 5.9 Pollen grains. A. *Zamia* sp., a cycad. B. *Ginkgo biloba*. C. *Pinus* sp., a conifer.

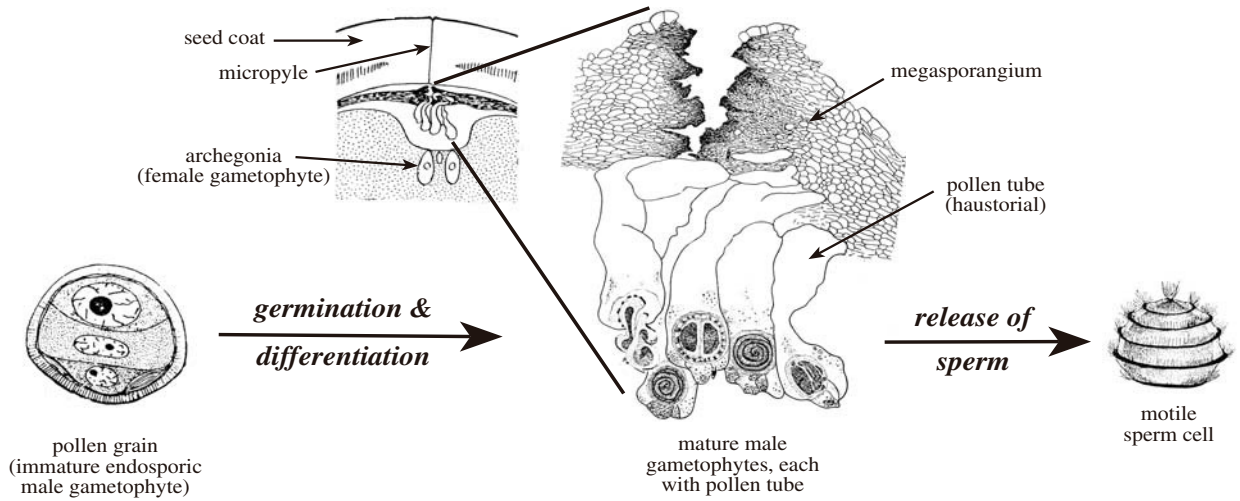


FIGURE 5.10 Male gametophyte morphology and development in the non flowering Spermatophytes; *Cycas* sp., illustrated. (Reproduced and modified from Swamy, B. G. L. 1948. American Journal of Botany 35: 77-88, by permission.)

plant lineages. Note that the ovules of angiosperms lack pollination droplets or pollination chambers, as flowering plants have evolved a different mechanism of pollen grain transfer (see Chapter 6).

OVULE AND SEED DEVELOPMENT

After pollination, the **megasporocyte** develops within the megasporangium of the ovule (Figures 5.11, 5.13A). The megasporocyte is a single cell that undergoes meiosis, producing a tetrad of four haploid megaspores, which in most extant seed plants are arranged in a straight line, or linearly (Figure 5.11). The three megaspores that are distal (away from the ovule base)

abort; only the proximal megaspore (near the ovule base) continues to develop. In the pollination chamber, the resorbed pollen grains (Figures 5.11, 5.13A) develop into mature male gametophytes and form pollen tubes, which grow into the tissue of the megasporangium (Figures 5.11, 5.13B). In gymnosperms these male gametophytes may live in the megasporangial tissue for some time, generally several months to a year.

The functional megaspore greatly expands, accompanied by numerous mitotic divisions, to form the endosporic female gametophyte (Figures 5.11, 5.13B,C). In the seeds of gymnosperms, archegonia differentiate at the apex of the female gametophyte (Figure 5.13C,D). As in the nonseed

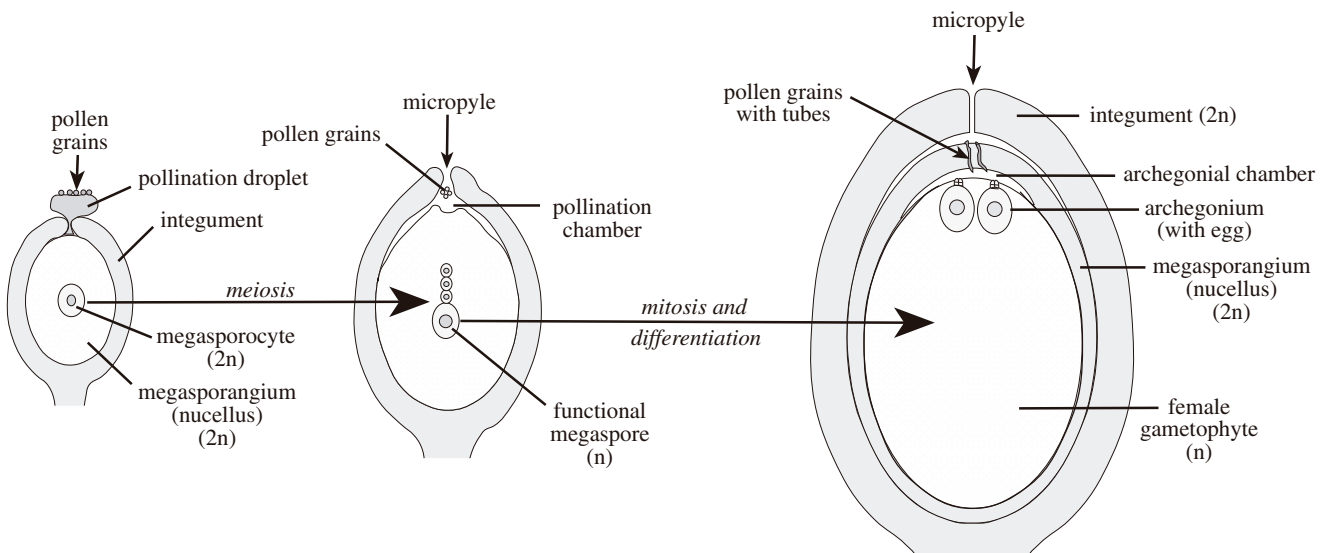


FIGURE 5.11 Ovule development in the non flowering Spermatophytes.

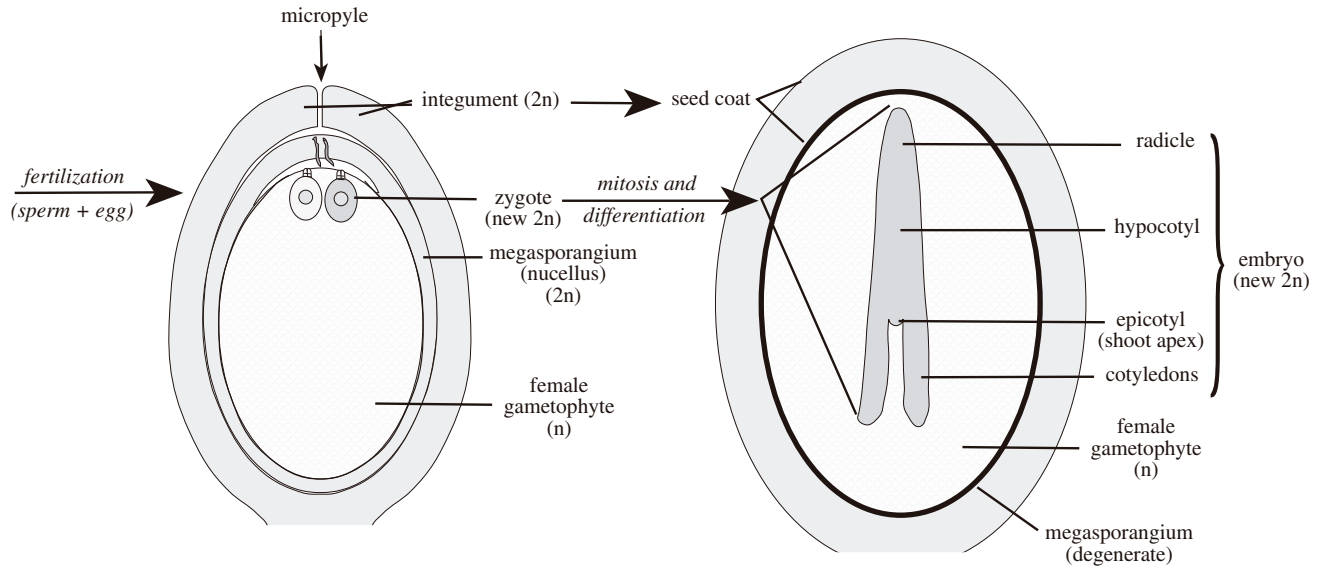


FIGURE 5.12 Seed development in the non-flowering Spermatophytes.

land plants, each archegonium has a large egg cell and a short line of neck cells (plus typically a ventral canal cell or nucleus). Eventually, the male gametophytes release or transport sperm cells (motile or nonmotile) into a cavity between the megasporangium and female gametophyte known as the **archegonial chamber** (Figure 5.11). (Note that the ovules of angiosperms lack archegonia and an archegonial chamber.) Here, the sperm cells either swim to (in cycads and *Ginkgo*) or are released in close proximity to (in conifers and Gnetales) an archegonium of the female gametophyte. A sperm cell entering the archegonium then fertilizes the egg. A long time (perhaps a year or more) may ensue between pollination, which is delivery of the pollen grains to the ovule, and fertilization, actual union of sperm and egg. **Note:** This is not true for the flowering plants, in which fertilization occurs very soon after pollination (see Chapter 6).

The resulting diploid zygote, once formed, undergoes considerable mitotic divisions and differentiation, eventually maturing into the **embryo**, the immature sporophyte (Figures 5.12, 5.13E). The tissue of the female gametophyte continues to surround the embryo (Figure 5.13E) and serves as nutritive tissue for the embryo upon seed germination (except in the flowering plants; see Chapter 6). The megasporangium (nucellus) eventually degenerates. The integument matures into a peripheral **seed coat**, which may differentiate into various hard and/or fleshy layers.

SEED ADAPTATIONS

The adaptive significance of the seed is unquestioned. First, seeds provide *protection*, mostly by means of the seed coat,

from mechanical damage, desiccation, and often predation. Second, seeds function as the *dispersal unit* of sexual reproduction. In many plants the seed has become specially modified for dispersal. For example, a fleshy outer seed coat layer may function to aid in animal dispersal. In fact, in some plants the seeds are eaten by animals, the outer fleshy layer is digested, and the remainder of the seed (including the embryo protected by an inner, hard seed coat layer) passes harmlessly through the gut of the animal, ready to germinate with a built-in supply of fertilizer. In other plants, differentiation of the seed coat into one or more wings functions in seed dispersal by wind. Third, the seed coat may have *dormancy mechanisms* that ensure germination of the seed only under ideal conditions of temperature, sunlight, or moisture. Fourth, upon germination, the *nutritive tissue* surrounding the embryo provides *energy* for the young seedling, aiding in successful establishment.

Interestingly, in seed plants the female gametophyte (which develops within the megaspore) remains attached to and nutritionally dependent upon the sporophyte. This is exactly the reverse condition as is found in the liverworts, hornworts, and mosses (Chapter 3).

EUSTELE

In addition to the seed, an apomorphy for *most* spermatophytes, including all *extant* spermatophytes (Figure 5.1), is the **eustele** (Figure 5.14). A eustele is a primary stem vasculature (primary meaning prior to any secondary growth) that consists of a single ring of discrete vascular bundles. Each vascular bundle contains an internal strand of xylem

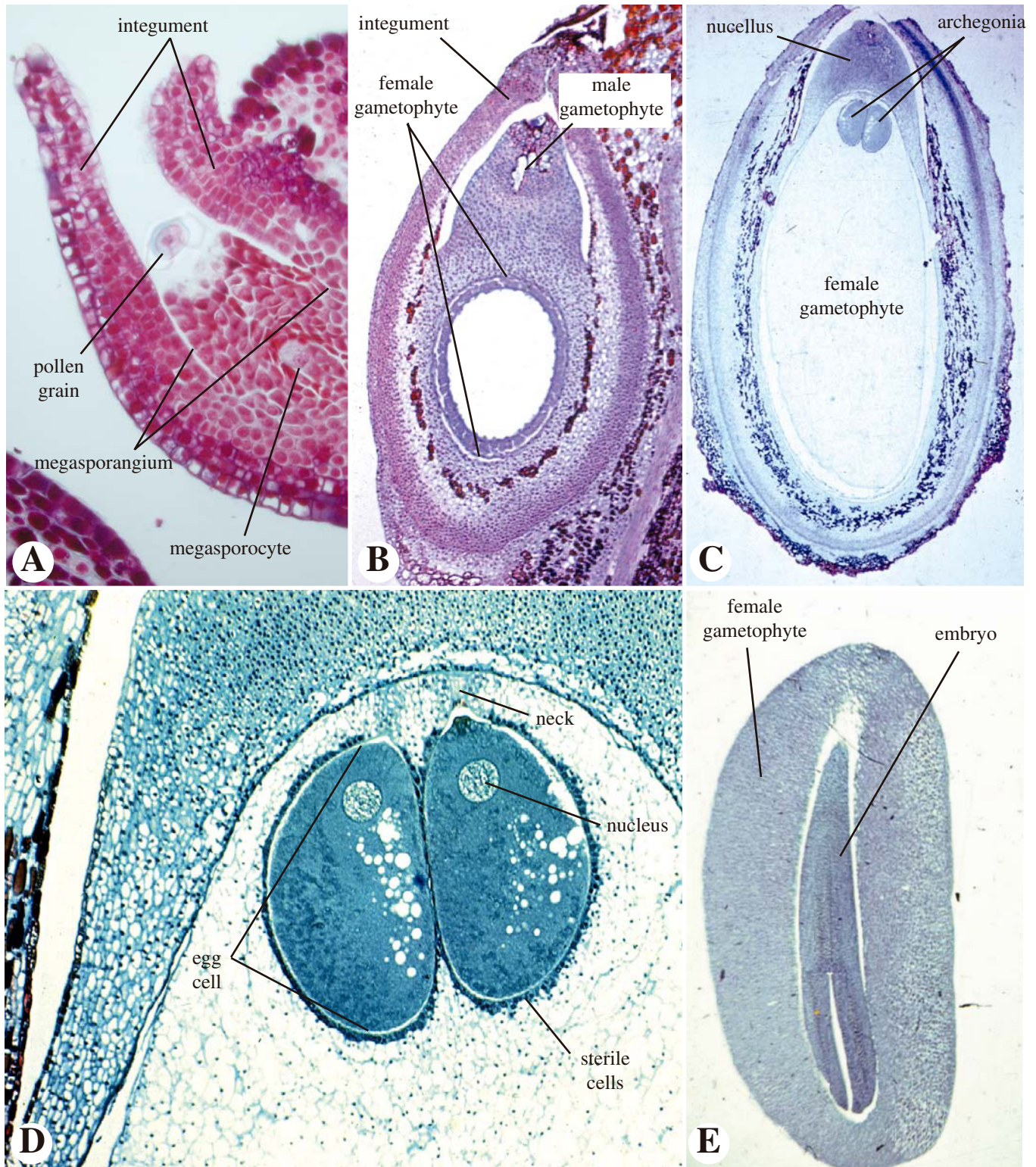


FIGURE 5.13 Ovule and seed development, illustrated by *Pinus* sp. **A.** Young ovule, longitudinal section, at time of pollination. Pollen grains are pulled into micropyle by resorption of pollination droplet. Meiosis of the megasporocyte has yet to occur. **B.** Postpollination, showing development of the female gametophyte and haustorial pollen tube growth of the male gametophytes within tissue of megasporangium (nucellus). **C.** Mature ovule, showing two functional archegonia within female gametophyte. **D.** Close-up of archegonia, each containing a large egg cell with a surrounding layer of sterile cells and apical neck. **E.** Seed longitudinal section, seed coat removed, showing embryo and surrounding nutritive layer of female gametophytic tissue.

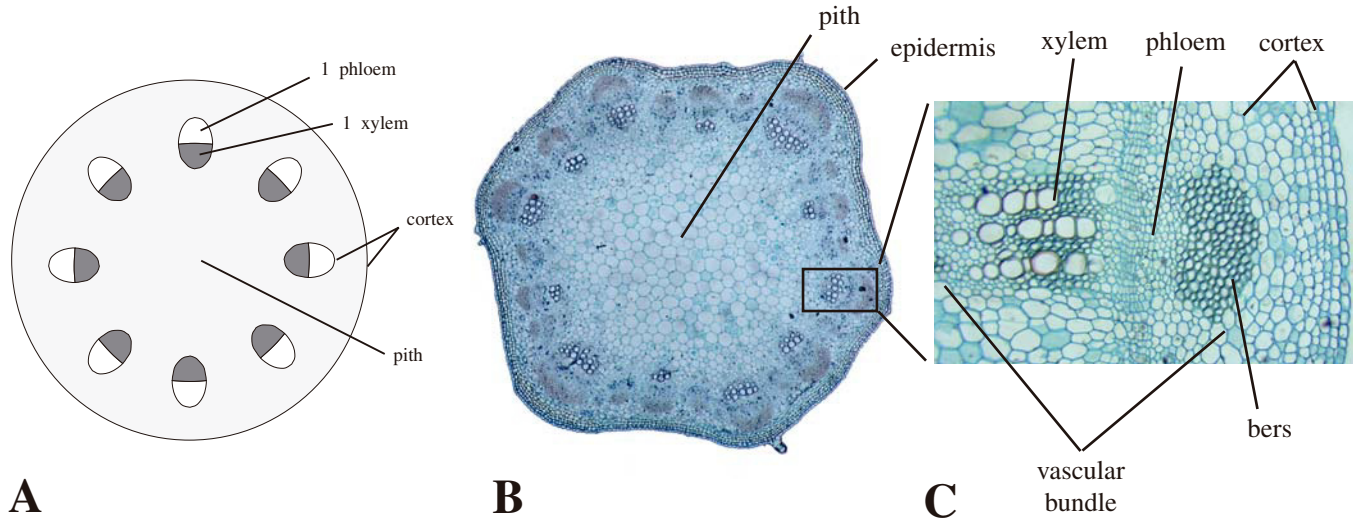


FIGURE 5.14 Eustele. **A.** Diagram of eustele. Note single ring of vascular bundles, with xylem inside, phloem outside. **B.** *Helianthus* stem cross-section, an example of a eustele. **C.** Close-up of vascular bundle, showing xylem, phloem, and associated fibers.

and an external strand of phloem that are radially oriented, i.e., positioned along a radius (Figure 5.14).

The protoxylem of the vascular bundles of a eustele is endarch in position, i.e., toward the center of the stem. This is distinct from the exarch protoxylem of the lycophytes and the mesarch protoxylem of the monilophytes (Chapter 4).

DIVERSITY OF WOODY AND SEED PLANTS

ARCHEOPTERIS

A well-known lignophyte that lacked seeds was the fossil plant *Archeopteris* (not to be confused with the very famous fossil, reptilian bird *Archeopteryx*). *Archeopteris* was a large tree, with wood like a conifer but leaves like a fern (Figure 5.15A,B). Sporangia, producing spores, were born on fertile branch systems. Some species of *Archeopteris* were heterosporous.

“PTERIDOSPERMS” — “SEED FERNS”

The pteridosperms, or seed ferns, are almost certainly a nonnatural, paraphyletic group of fossil plants that had fernlike foliage, yet bore seeds. *Medullosa* is a well-known example of a seed fern (Figure 5.15C E). As in many fossil plants, different organs of *Medullosa* are placed in separate form genera. For example, the fernlike leaves of *Medullosa* are in the form genera *Alethopteris* and *Neuropteris*. *Dolerotheca*, which had huge pollen grains, refers to the

pollen-bearing organs of *Medullosa*, and seeds of *Medullosa* are placed in the genus *Pachytesta*.

The relationships of various pteridosperms to extant seed plants are unclear. Some are basal to the extant seed plants; others may be more closely related to the gymnosperms and others to the angiosperms (Figure 5.1).

GYMNOSPERMS

The extant, nonangiospermous seed plants are included within a group known as the Gymnospermae, or gymnosperms (after *gymnos*, naked + *sperm*, seed). In the past decade or so, based on morphological and limited molecular studies, the gymnosperms were largely accepted to be an unnatural, paraphyletic taxon, grouped together based more on what they lacked (flowers) than on any definitive apomorphy. In addition, the Gnetales (discussed later) were considered to be the closest living relative of the angiosperms, together comprising a group termed the Anthophytes. However, these results were never viewed as particularly robust.

Very recently, more intensive cladistic analyses using multiple gene sequences have provided quite strong evidence that the gymnosperms are in fact a monophyletic group and are sister to the angiosperms (Figure 5.1). Relationships within the gymnosperms are somewhat unclear, but many results show the cycads (or Cycadophyta) as the most basal lineage, followed by the *Ginkgo* group (Ginkgophyta), then the conifers (Coniferophyta). Interestingly, the Gnetales are most frequently placed within the conifers (often as the sister group to

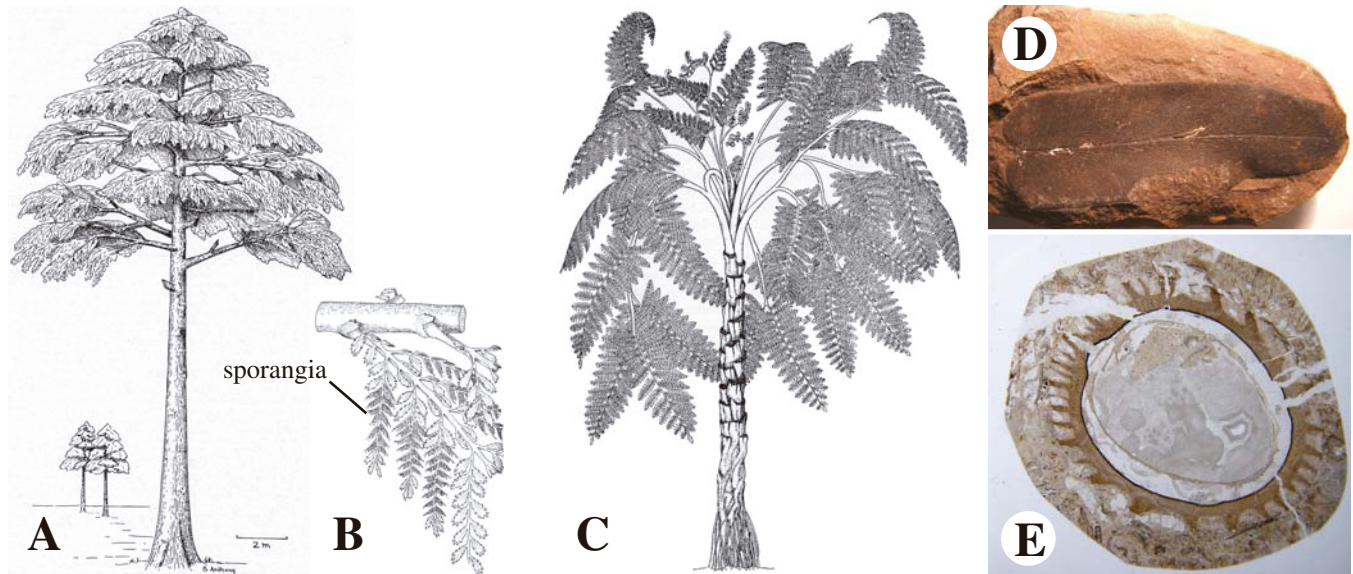


FIGURE 5.15 A–B. *Archeopteris*, an extinct lignophyte. A. Reconstruction of plant. B. Branch system, showing leaves and sporangia. (Reproduced from: Beck, C. B. 1962. *American Journal of Botany* 49: 373–382, by permission.) C–E. *Medullosa*, an extinct seed fern. C. Reconstruction of plant. (Reproduced from: Stewart, W. N., and T. Delevoryas. 1956. *Botanical Review* 22: 45–80, by permission.) D. Fossil leaf impression (*Neuropteris*). E. Seed longitudinal section (*Pachytosta*).

the Pinaceae. Thus, the Anthophytes are no longer recognized as a natural group.

CYCADOPHYTA—CYCADS. The Cycadophyta, or cycads, are a relatively ancient group of plants that were once much more common than today and served as fodder for plant-eating nonavian dinosaurs. Extant cycads are now fairly restricted in distribution, consisting of approximately 185 species in 11 or so genera. Cycads are found in southeastern North America, Mexico, Central America, some Caribbean islands, South America, eastern and southeastern Asia, Australia, and parts of Africa. Many cycads throughout the world are of economic importance in being used as a source of food starch (sometimes termed sago), typically collected from the apex of the trunk just prior to a flush of leaves or reproductive structures. Some cycads, especially *Cycas revoluta*, the sago palm, are planted horticulturally.

Cycads are an apparently monophyletic lineage consisting of plants with a mostly short, erect stem or trunk, rarely tall and palmlike (as in the misnamed genus *Microcycas*). The trunk bears spirally arranged, mostly pinnately compound leaves (Figure 5.16A,C,E). Only the genus *Bowenia* of Australia has bipinnately compound leaves (Figure 5.16B). The trunk of cycads does not usually exhibit lateral (axillary) branching; thus, the **loss of axillary branching** on the aerial trunk may be an apomorphy for the cycads (Figure 5.1). Interestingly, the leaves of cycads have **circinate vernation**

(Figure 5.17B) as in ferns, perhaps a primitive retention that was lost in other seed plants. Reproductively, all cycad individuals are either male or female; this plant sex is termed **dioecious** (see Chapter 9).

The classification of cycads varies, but recent evidence suggests they are best grouped as two families: Cycadaceae and Zamiaceae. The Cycadaceae, which consists solely of the genus *Cycas*, is distinguished by not forming female cones. In species of *Cycas*, seeds are produced on the lower margins of numerous **female sporophylls** (also called megasporophylls) that are congregated at the trunk apex in dense masses (Figure 5.17E,G). *Cycas* species do have male cones (Figure 5.17A,C), which are found in all cycads (see later discussion).

The family Zamiaceae differs from the Cycadaceae in having both male and female **cones**, also called **strobili**. Recall that cones are determinate shoot systems, consisting of a single axis that bears **sporophylls**, modified leaves with attached sporangia. Male cones (Figures 5.17A,B, 5.18A,B,G) have **male sporophylls** (also called microsporophylls), each of which bears numerous male sporangia (Figure 5.17D). The male sporangia, also called **microsporangia**, produce haploid microspores that develop into pollen grains. Female cones (Figures 5.16C,E, 5.18C,D,F,G) have **female sporophylls** (also called megasporophylls), each of which bears two seeds (Figure 5.18E,H,I).

Interestingly, the pollen of all cycads release motile sperm cells (Figure 5.10) into the ovule of a female cone, a vestige of an ancestrally aquatic condition.



FIGURE 5.16 Cycad diversity. Zamiaceae. **A.** *Macrozamia moorei* (Australia). **B.** *Bowenia spectabilis* (Australia), showing a single, bipinnately compound leaf. **C.** *Lepidozamia peroffskyana* (eastern Australia). **D.** *Stangeria eriopus* (Africa). **E.** *Encephalartos lebomboensis* (Africa).

GINKGOPHYTA—GINKGO. The Ginkgophyta, or ginkgophytes, have an extensive fossil record but contain only one extant species, *Ginkgo biloba*. This species is native only to certain remote regions of China but has now been planted worldwide as a popular street tree. *Ginkgo biloba*, unlike the cycads (and similar to conifers, discussed next), is a highly branched, woody tree. It can be recognized by the fact that it has short shoots in addition to long shoots, and by the distinctive obtriangular (fan-shaped), often two-lobed leaves with dichotomous venation (Figure 5.19A C). *Ginkgo*, like the cycads, is **dioecious** and has ancestrally motile sperm.

Male *Ginkgo* trees bear reproductive structures that are called cones but that do not bear structures that resemble sporophylls. These male cones consist of a central axis with lateral stalks (Figure 5.19D E), each of which bears two microsporangia (Figure 5.19F,G). The microsporangia dehisce longitudinally, releasing pollen grains. Female *Ginkgo* trees do not bear cones. The female reproductive structures each consist of an axis having two terminal ovules (Figure 5.19H,I).

CONIFEROPHYTA—CONIFERS. The Coniferophyta, or conifers, are another ancient group of land plants that were

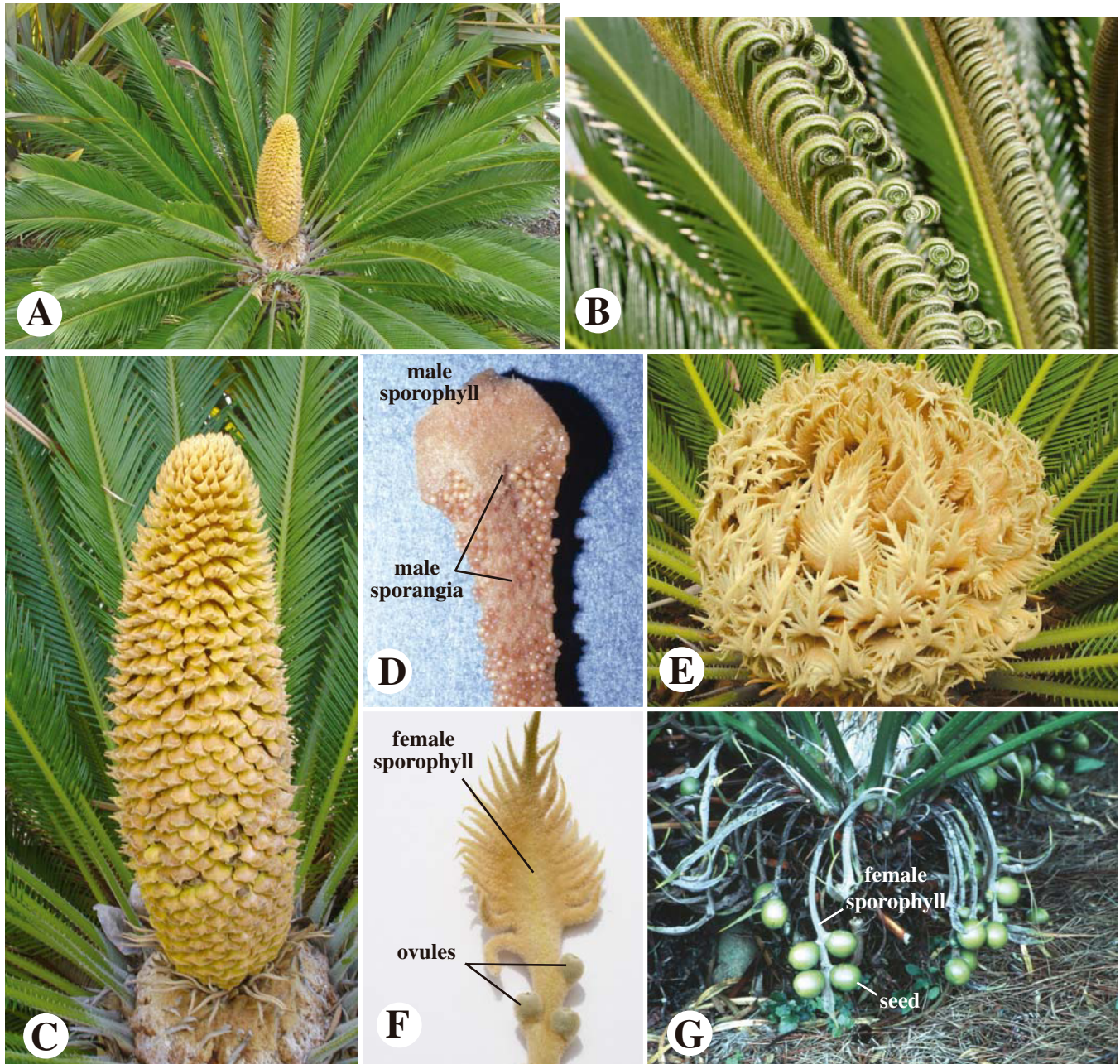


FIGURE 5.17 Cycad reproduction. Cycadaceae (*Cycas*). A–F. *Cycas revoluta*, sago palm. A. Male individual with male cone. B. Leaves with circinnate vernation. C. Close-up of male cone. D. Male sporophyll with sporangia. E. Female individual, showing aggregate of female sporophylls (cones lacking). F. Female sporophyll with marginal, immature seeds. G. *Cycas circinalis*, female, showing mature female sporophylls with seeds (cones lacking).

once dominant in most plant communities worldwide. Today, they have largely been replaced by angiosperms, but still constitute the dominant species in various coniferous forests.

Conifers comprise a monophyletic group of highly branched trees or shrubs with **simple leaves**, the latter a possible

apomorphy for the group. Leaves of conifers are linear, acicular (needle-like), or subulate (awl-shaped; see Chapter 9). In some conifers the leaves are clustered into **short shoots**, in which adjacent internodes are very short in length. An extreme of this is the **fascicle**, e.g., in species of *Pinus*, the pines.

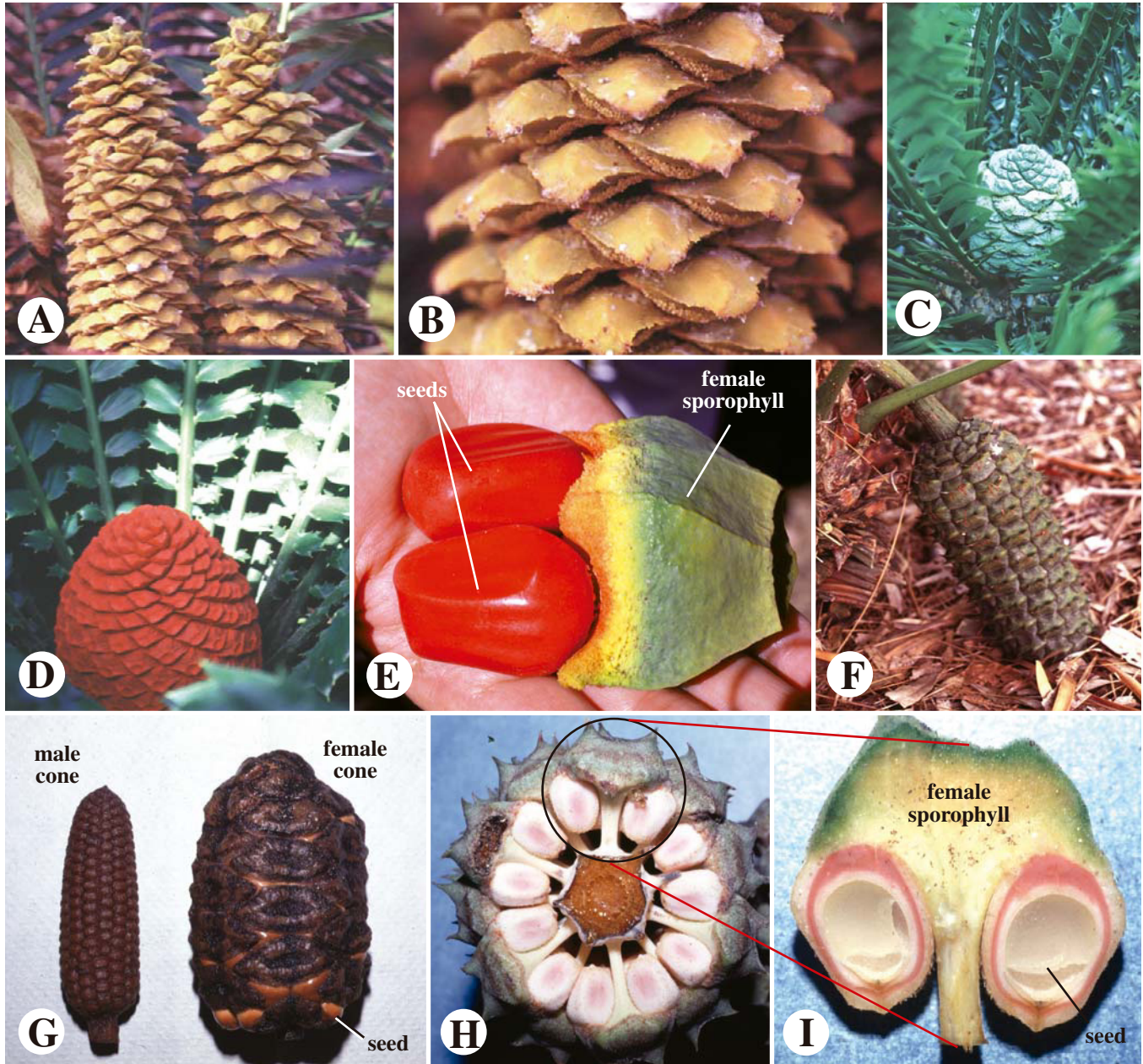


FIGURE 5.18 Cycad reproduction. Zamiaceae. **A,B.** *Encephalartos altensteinii*, male cones. **C.** *Encephalartos arenarius*, female with cone. **D.** *Encephalartos ferox*, female, with bright red cone. **E.** *Encephalartos maikensis*, female sporophyll with two attached seeds. **F.** *Ceratozamia mexicana*, female cone. **G.** *Zamia* sp., male and female cones. **H,I.** *Ceratozamia* sp., female cone and sporophyll with ovules.

A fascicle is a specialized short shoot consisting of stem tissue, one or more needle-shaped leaves, and persistent basal bud scales (Figure 5.20A,B; Chapter 9).

A second, apparent apomorphy of the conifers, including the Gnetales (discussed next), is the loss of sperm cell motility (Figure 5.1). This distinguishes the conifers from the other

gymnosperms, which have flagellated sperm cells. Conifers, like all extant seed plants, are **siphonogamous**, i.e., the male gametophytes develop pollen tubes. As in cycads and *Ginkgo*, these pollen tubes are haustorial, consuming the tissues of the nucellus (megasporangial tissue) for a year or so after pollination. One difference, however, (likely correlated with

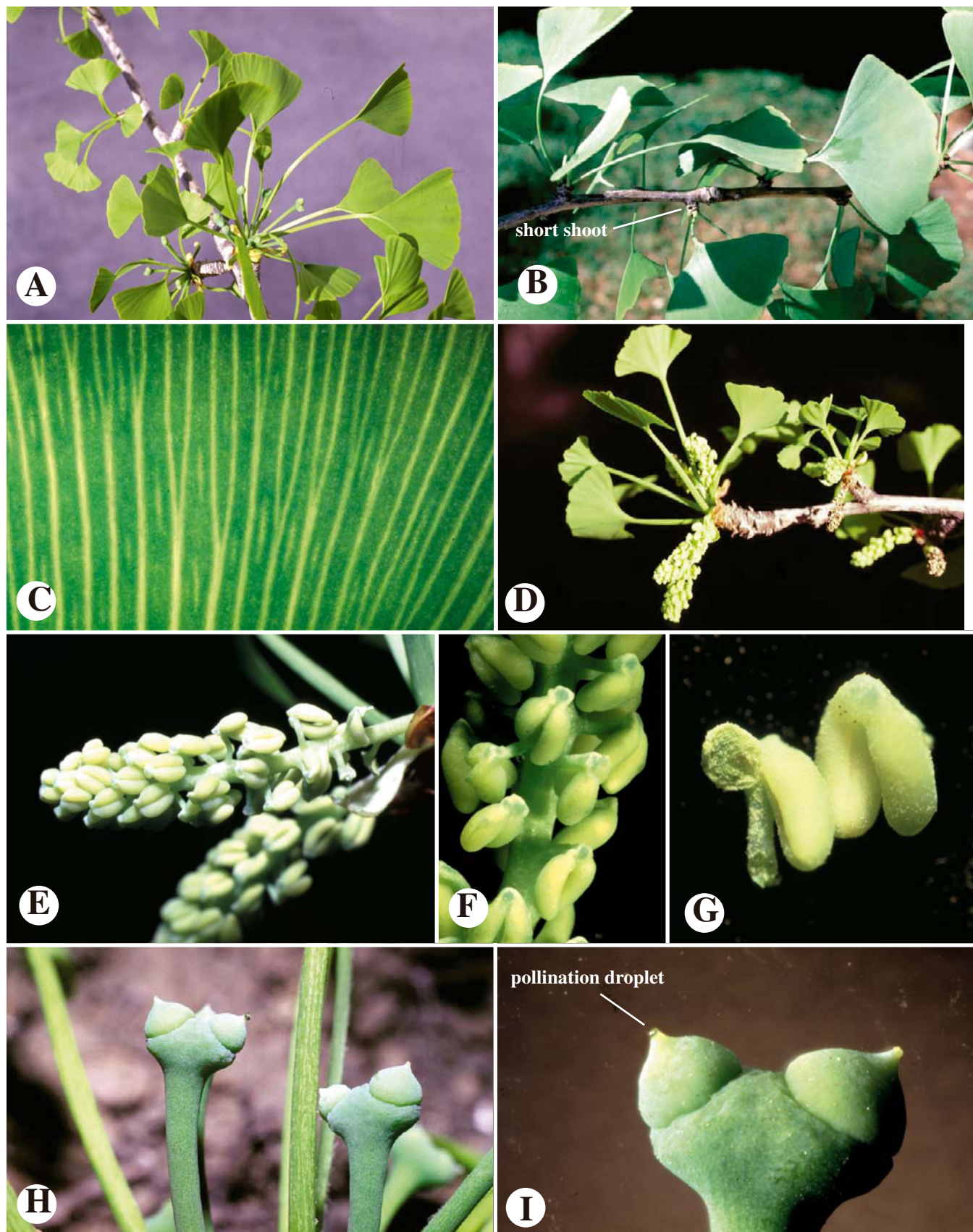


FIGURE 5.19 *Ginkgo biloba*. A,B. Vegetative growth. Note fan-shaped leaves, clustered into short shoots. C. Leaf close-up, showing dichotomous venation. D. Male tree bearing male cones. E. Male cone. F,G. Close-up of male sporangia, born in pairs on stalk arising from central axis of male cone. H. Female plant bearing stalk with pair of ovules. I. Close-up of ovule pair. Note pollination droplet from micropyle.

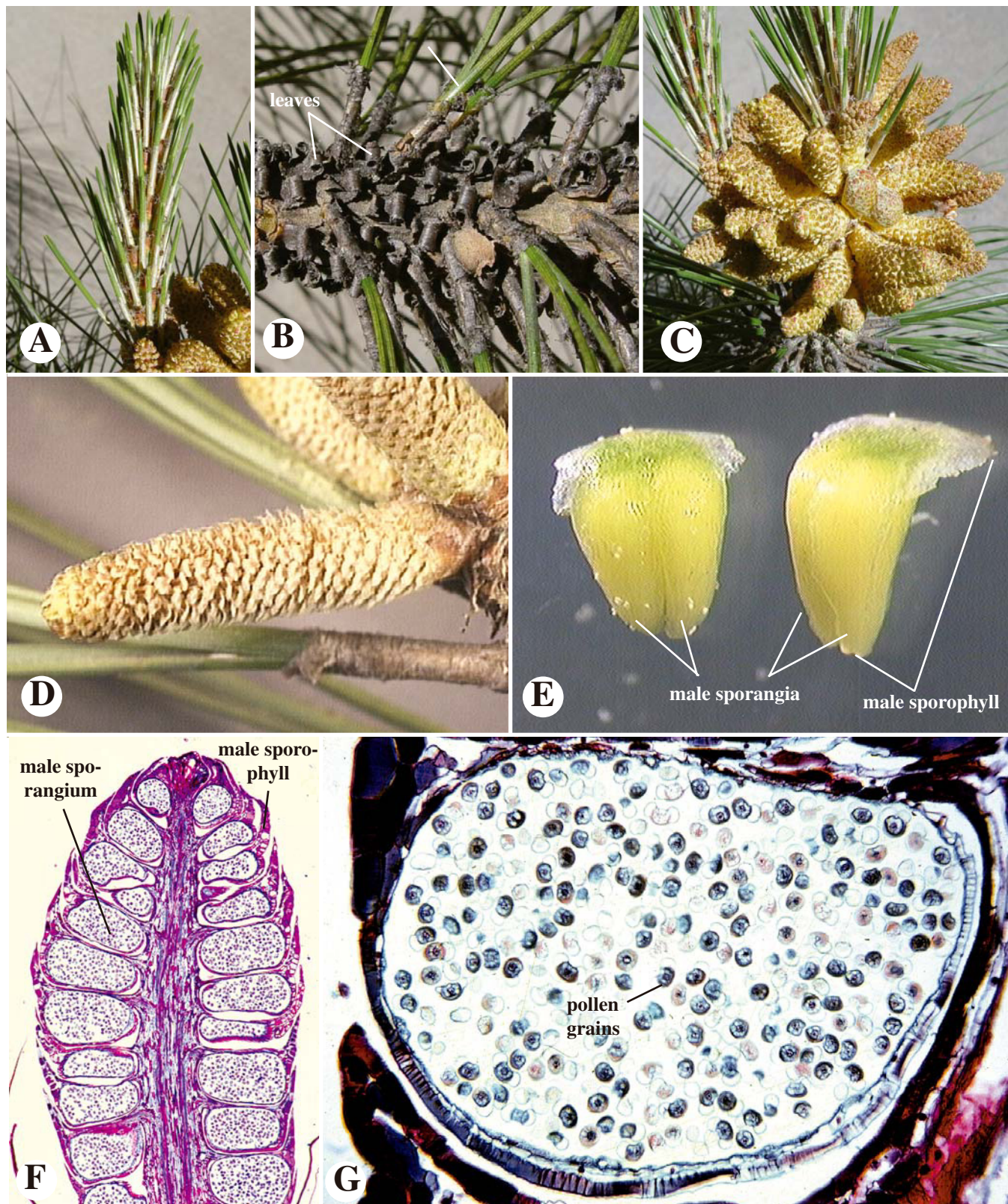


FIGURE 5.20 Conifers. A–G. *Pinus* spp. A. Shoot with young fascicles. B. Branch, showing scale leaves and fascicles. C. Apex of branch with fascicles and male cones. D. Male cones, close-up. E. Male sporophylls of male cones, each with two male sporangia. F. Male strobilus, longitudinal section, showing microsporangia and subtending microsporophylls. G. Close-up of microsporangium, full of mature pollen grains.

sperm nonmotility) is that the male gametophyte of conifers delivers the sperm cells more directly to the egg by the growth of the pollen tube into the archegonial chamber, where it makes contact with the female gametophyte at or near the archegonia. The nonswimming sperm cells are then released from the pollen tube, make contact with the archegonial egg cell, and fertilize the egg nucleus. Because there is more than one archegonium per seed, multiple fertilization events may occur, resulting in multiple young embryos, but usually only one survives in the mature seed.

Reproductively, conifers produce male cones and female cones, either on the same individual (**monoecy**) or, less commonly, on different individuals (**dioecy**). As with all vascular plants, cones consist of an axis that bears sporophylls. As in cycads, male strobili (Figure 5.20C,D) have **male sporophylls** (microsporophylls; Figure 5.20E,F). These male sporophylls bear the **male sporangia** (microsporangia) that produce **pollen grains** (Figure 5.20E G). The pollen grains of conifers are interesting in mostly being bi-saccate, in which two bladders develop from the pollen grain wall (Figure 5.9C). These saccate structures, like air bladders, may function to transport the pollen more efficiently by wind. They may also function as flotation devices, to aid in the capture and transport of pollen grains by a pollination droplet formed in the nonflowering seed plants.

Female cones of most conifers are different from those of other seed plants. Conifer female cones are a **compound** structure. They consist of an axis that bears modified leaves called **bracts**, each of which subtends the seed-bearing structure, called an **ovuliferous scale** (Figure 5.21). The ovuliferous scale is actually a modified lateral branch system. The evidence for this is the inverted vasculature orientation and fossil intermediates between extant conifers and fossil conifers plus another fossil group called the Cordaitales. In most conifer female cones, the ovuliferous scales are much bigger than the small bracts (Figure 5.21D F). In a few conifers, e.g., *Pseudotsuga*, or Douglas-fir, the bracts are elongated and can be seen on the outside of the ovuliferous scales (Figure 5.21G). The female cones of most conifers have two seeds on the upper surface of each ovuliferous scale (Figure 5.21H). Mature seeds are typically winged (Figure 5.21H,I), an adaptation for seed dispersal by wind.

Important families of conifers include the **Araucariaceae** (e.g., buya bunya, monkey puzzle, and Norfolk Island-pine, Figure 5.22A); **Cupressaceae**, or cypress family (e.g., cypress, junipers, incense cedar, bald cypress, redwood, and giant sequoia, inclusive of the Taxodiaceae; Figure 5.22B G); **Podocarpaceae** (including the yew pine; Figure 5.22H); **Taxaceae** (yews; Figure 5.22I,J); and **Pinaceae**, or Pine family (including cedars, pines, spruces, firs, Douglas-fir, larches, and hemlock; Figures 5.20, 5.21, 5.22K,L).

GNETALES. The Gnetales, also referred to as the Gnetopsida or sometimes Gnetophyta, are an interesting group containing three extant families: Ephedraceae (consisting of *Ephedra*, with about 65 species), Gnetaceae (consisting of *Gnetum*, with 28 species, plus the monotypic genus *Vinkarella*), and Welwitschiaceae (consisting of the sole species *Welwitschia mirabilis*). The Gnetales has often been thought to be the sister group to the angiosperms, the two groups united by some obscure features, possibly including whorled, somewhat perianth-like microsporophylls in structures that may resemble flowers (see Chapter 6). However, as reviewed earlier, recent molecular studies have placed the Gnetales within the conifers (Figure 5.1).

The Gnetales are united by (among other things) the occurrence of (1) striate pollen (Figure 5.23A); and (2) vessels with porose (porelike) perforation plates (Figure 5.23B), as opposed to scalariform (barlike) perforation plates in basal angiosperms (see Chapter 6). The vessels of Gnetales were derived independently from those of angiosperms. The reproductive structures in various Gnetales show some parallels to the flowers of angiosperms.

Ephedra of the Ephedraceae is a rather common desert shrub (Figure 5.24A C) and can be recognized by the photosynthetic, striate stems and the very reduced scale-like leaves, only two or three per node. Male or female cones may be found in the axils of the leaves (Figure 5.24B,C). The Gnetaceae are tropical vines (rarely trees or shrubs) with opposite (decussate), simple leaves (Figure 5.24D), looking for all the world like an angiosperm but, of course, lacking true flowers. *Welwitschia mirabilis* of the Welwitschiaceae is a strange plant native to deserts of Namibia in southwestern Africa. An underground caudex bears only two leaves (Figure 5.24E,F), these becoming quite long and lacerated in old individuals. Male and female cones are born on axes arising from the apex of the caudex (Figure 5.24G J).

Recently, the occurrence of a type of **double fertilization** was verified in species of the Gnetales. Double fertilization in *Ephedra* entails the fusion of each of two sperm cells from a male gametophyte with nuclei in the archegonium of the female gametophyte. One sperm fuses with the egg nucleus and the other fuses with the ventral canal nucleus. In fact, the fusion product of sperm and ventral canal cell may even divide a few times mitotically, resembling angiospermous endosperm (Chapter 6), but this does not persist. Thus, double fertilization, which has long been viewed as a defining characteristic of the angiosperms alone, was recently interpreted as a possible apomorphy of the Gnetales and angiosperms together (the Anthophytes). This notion is rejected with the current acceptance of seed plant relationships as seen in Figure 5.1, in which the Gnetales are nested within the conifers. Thus, double fertilization in the Gnetales and angiosperms presumably evolved independently.

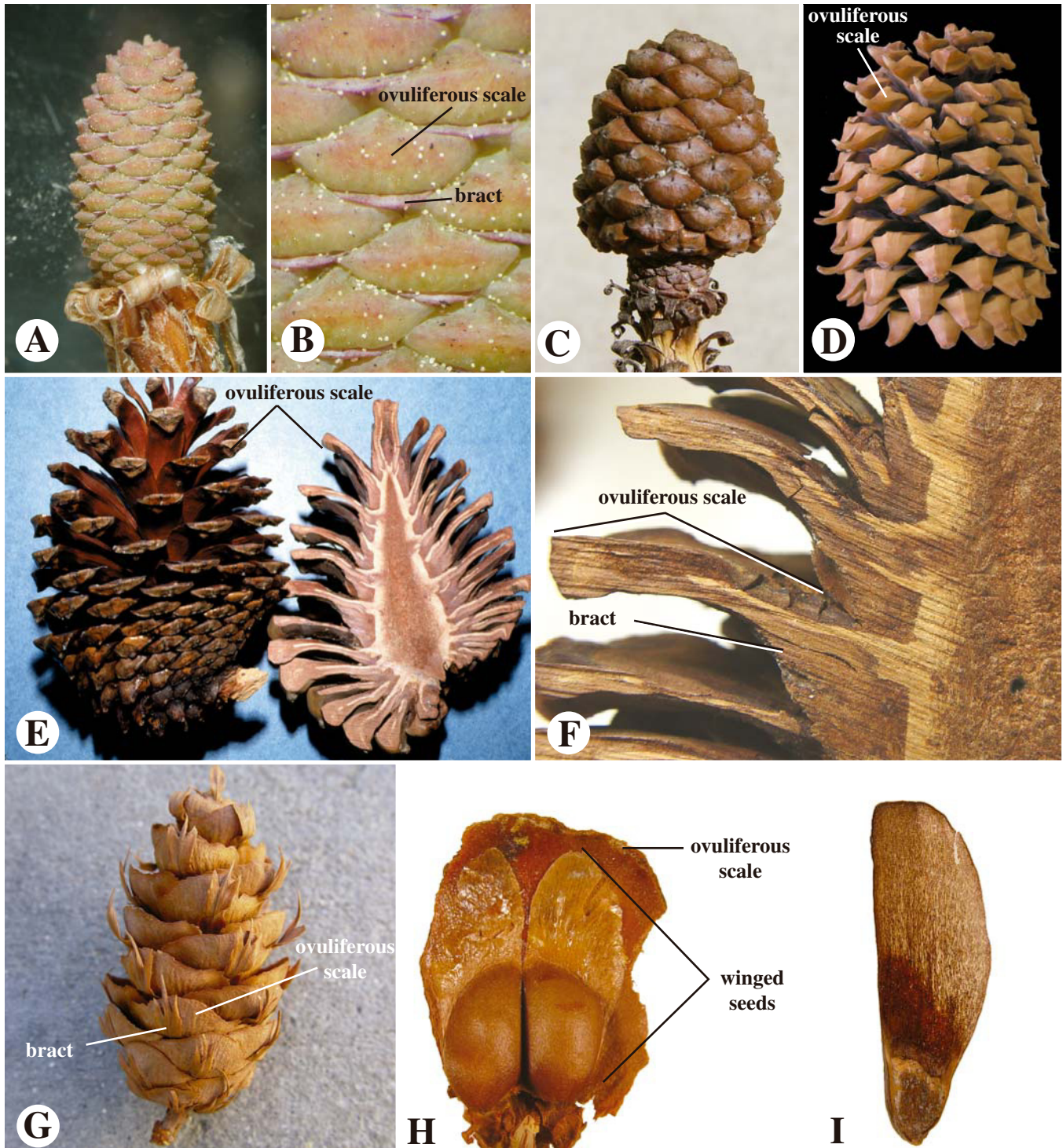


FIGURE 5.21 Conifers. A–F. *Pinus* spp. A. Young female cone, at time of pollination. B. Close-up, showing ovuliferous scales and bracts. Note pollen grains. C. One-year-old female cone. D. *Pinus coulteri*, coulter pine, mature female cone (most massive of any species). E. Female pine cones, right in section. F. Close-up of longitudinal section, showing bract and ovuliferous scale. G. *Pseudotsuga* sp. (Douglas- r) female cone. Note elongate bracts and wide ovuliferous scales. H. Immature ovuliferous scale, top view, showing two winged seeds. I. *Pinus*, mature winged seed.

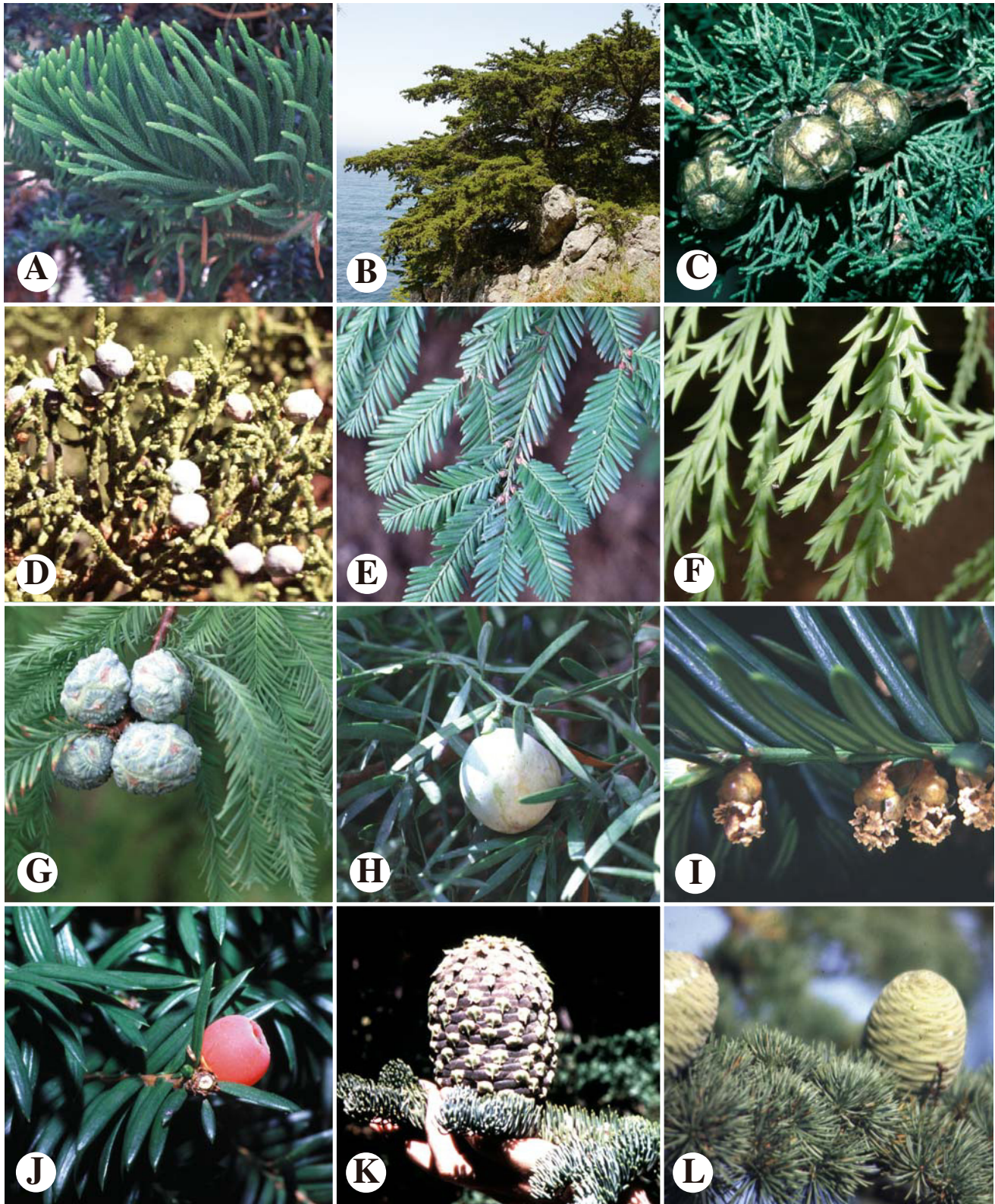


FIGURE 5.22 Conifer diversity. **A.** Araucariaceae. *Araucaria heterophylla*, Norfolk Island-pine. **B–G.** Cupressaceae. **B.** *Cupressus macrocarpa*, Monterey cypress. **C.** *Cupressus sempervirens*, with female cones. **D.** *Juniperus californica*, California juniper, bearing fleshy female cones. **E.** *Sequoia sempervirens*, redwood, with flattened branch system having linear leaves. **F.** *Sequoiadendron giganteum*, giant sequoia, branches with awl-like leaves. **G.** *Taxodium distichum*, bald cypress, with leaf-like branches and female cones. **H.** Podocarpaceae. *Podocarpus gracilior*. Note seeds not aggregated in female cones. **I, J.** Taxaceae, *Taxus* sp., yew. **I.** Branch with bearing male cones. **J.** Branch with female cone of single seed surrounded by red, fleshy aril. **K–L.** Pinaceae. **K.** *Abies magnifica*, spruce, female cone. **L.** *Cedrus* sp., cedar, with female cones.

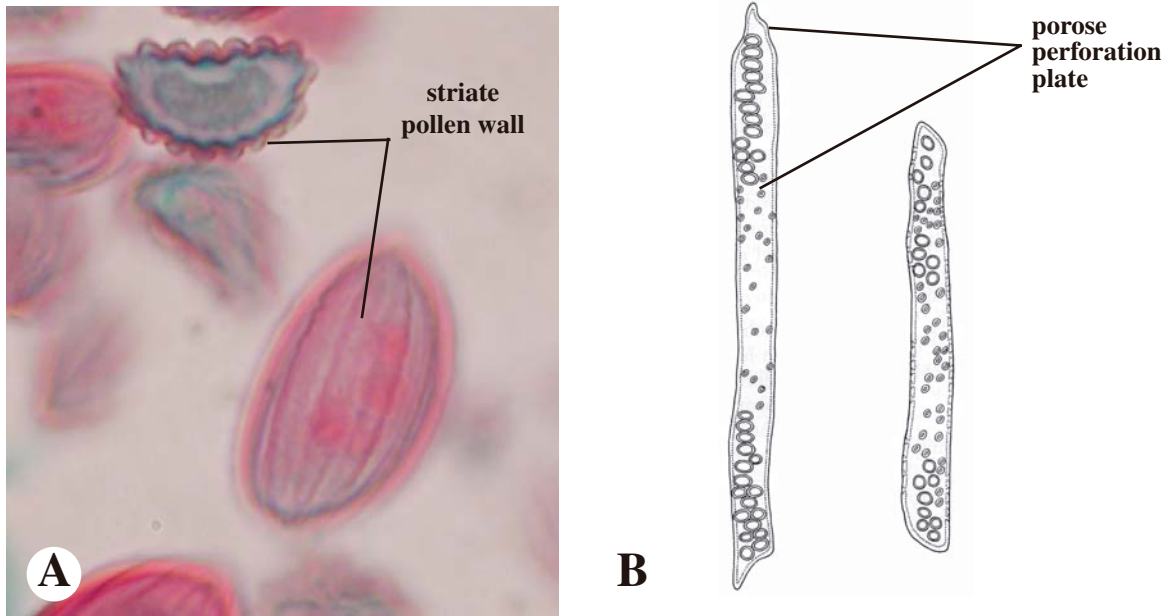


FIGURE 5.23 Gnetales apomorphies, illustrated by *Ephedra*. **A.** Striate pollen grains, face view below, cross-section above. **B.** Vessels with porose perforation plates. (**B** reproduced from Esau, K. 1965. *Plant Anatomy*. J. Wiley and Sons, New York, by permission.)

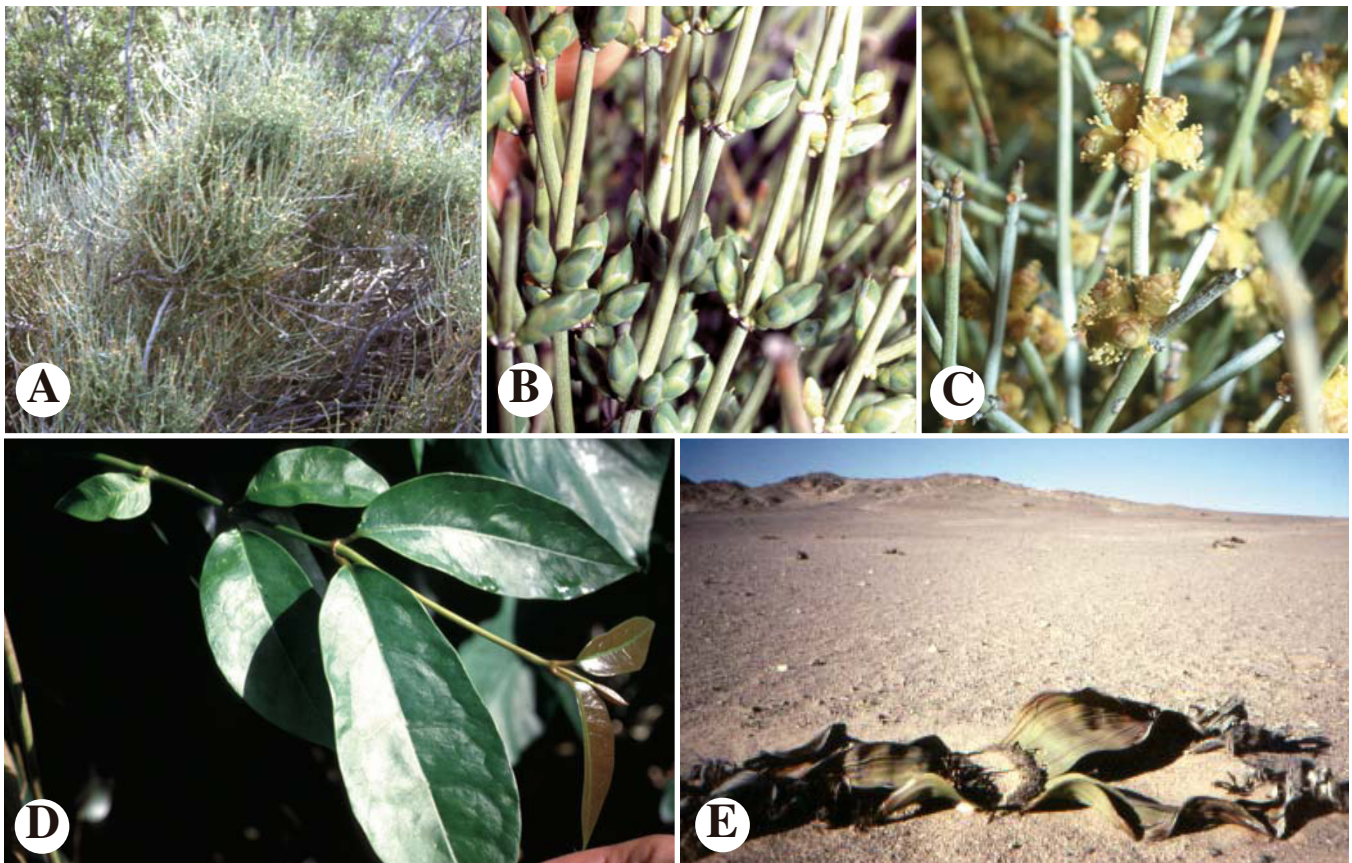


FIGURE 5.24 Gnetales exemplars. **A–C.** *Ephedra* sp. **A.** Whole plant. **B.** Female plant with cones. **C.** Male plant with cones. **D.** *Gnetum* sp. Note opposite leaves. **E–J.** *Welwitschia mirabilis*. **E.** Whole plant, showing two elongate opposite leaves. **F.** Close-up of central region of plant. **G.** Male cones. **H–J.** Female cones. [**E–G.** contributed by Mark Olsen] (Continued)

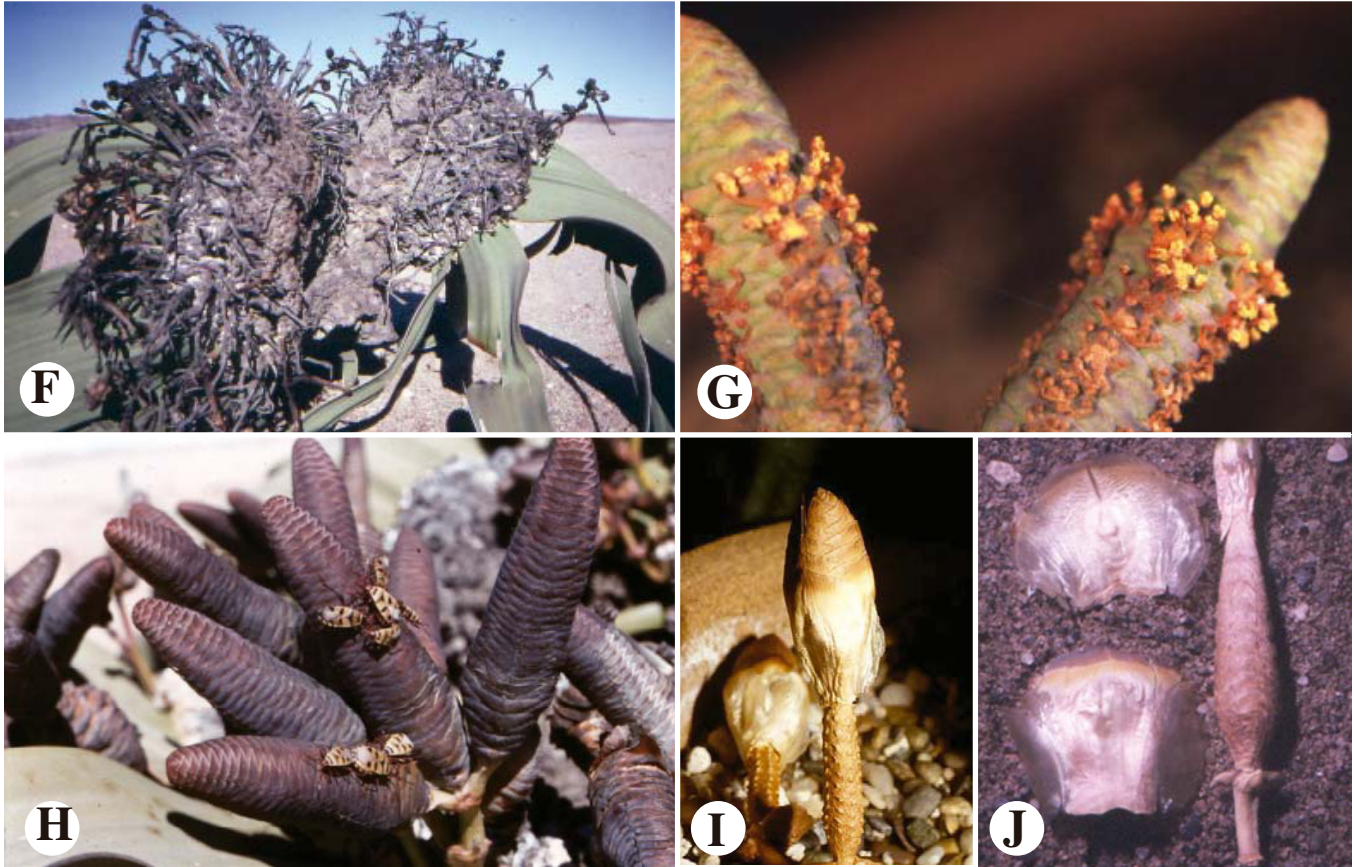


FIGURE 5.24 Continued

REVIEW QUESTIONS

WOODY PLANT APOMORPHIES

1. What are the major evolutionary novelties for the lignophytes?
2. Describe how a cambium undergoes secondary growth.
3. What are the products of secondary growth of the vascular cambium? the cork cambium?

SEED PLANT APOMORPHIES

4. Define seed and ovule.
5. Including heterospory, name and describe the steps that were involved in the evolution of the seed.
6. What is the definition of a pollen grain? From what does it develop?
7. What is a pollen tube and how does it function?
8. Define and state the significance of the pollination droplet.
9. Review the stages of ovule and seed development, and describe how a lag period can occur between pollination and fertilization.
10. Name four ways that seeds are adaptive.
11. Name and describe the stem stelar type that is an apomorphy for all extant seed plants.

SEED PLANT DIVERSITY

12. What were the basic features of *Archeopteris*?
13. What is a pteridosperm (seed fern)? Name a genus of the seed ferns.

14. What group of seed plants is characterized by generally short trunks, pinnate, coriaceous leaves (with circinate vernation) and dioecy, bearing either male or female cones?
15. What are the conifers and what are some families of conifers?
16. What is the definition of a cone (strobilus)? What are the parts of a female cone? a male cone?
17. What group/species is a tree having short shoots and obtriangular leaves with dichotomous venation?
18. What is the name of the structure in a pine cone that directly bears the ovules/seeds? What was it derived from? What subtends this structure?
19. What is a pine fascicle?
20. What is the morphology of a conifer pollen grain? What is the possible function of this morphology?
21. Name two apomorphies for the Gnetales.
22. Name the three families and four genera of the Gnetales. What do they look like and where do they occur?

EXERCISES

1. Peruse the most recent literature on phylogenetic relationships of the seed plants. Are there any differences relative to Figure 5.1?
2. Peruse botanical journals and find a systematic article on a conifer, ginkgo, a Gnetales, or a cycad. What is the objective of the article and what techniques were used to address it?
3. Collect and identify several local conifers. What diagnostic features are used to distinguish between species?

REFERENCES FOR FURTHER STUDY

- Bowe, L. Michelle, GwØnañle Coat, and Claude W. dePamphilis. 2000. Phylogeny of seed plants based on all three genomic compartments: Extant gymnosperms are monophyletic and Gnetales closest relatives are conifers. *Proceedings of the National Academy of Sciences of the United States of America* 97: 4092-4097.
- Chaw, Shu-Miaw, Christopher L. Parkinson, Yuchang Cheng, Thomas M. Vincent, and Jeffrey D. Palmer. 2000. Seed plant phylogeny inferred from all three plant genomes: Monophyly of extant gymnosperms and origin of Gnetales from conifers. *Proceedings of the National Academy of Sciences of the United States of America* 97: 4086-4091.
- Crane, Peter. 1985. Phylogenetic relationships in seed plants. *Cladistics* 1(4): 329-348.
- Doyle, James A., and Michael J. Donoghue. 1986. Seed plant phylogeny and the origin of angiosperms: an experimental cladistic approach. *The Botanical Review* 52(4): 321-431.
- Frohlich, Michael W., and David S. Parker. 2000. The mostly male theory of flower evolutionary origins: from genes to fossils. *Syst. Bot.* 25(2): 155-170.
- Gifford, E. M., and A. S. Foster. 1989. *Morphology and evolution of vascular plants*, 3rd edition. W.H. Freeman and Co., New York.
- Gugerli, F., C. Sperisen, U. Buchler, I. Brunner, S. Brodbeck, J. D. Palmer, and Y.-L. Qiu. 2001. The evolutionary split of Pinaceae from other conifers: evidence from an intron loss and a multigene phylogeny. *Molecular Phylogenetics and Evolution* 21: 167-175.
- Hill, K. D., M. W. Chase, D. W. Stevenson, H. G. Hills, and B. Schutzman. 2003. The families and genera of cycads: a molecular phylogenetic analysis of Cycadophyta based on nuclear and plastid DNA sequences. *International Journal of Plant Sciences* 164: 933-948.
- Jones, David L. 1993. *Cycads of the World: Ancient Plants in Today's Landscape*. Smithsonian Institution Press, Washington, DC.
- Nixon, Kevin C., William L. Crepet, Dennis Stevenson, and Else Marie Friis. 1994. A reevaluation of seed plant phylogeny. *Annals of the Missouri Botanical Garden* 81: 484-533.
- Rai, H. S., H. E. O'Brien, P. A. Reeves, R. G. Olmstead, and S. W. Graham. 2003. Inference of higher-order relationships in the cycads from a large chloroplast data set. *Molecular Phylogenetics and Evolution* 29: 350-359.
- Rothwell, Gar W., and Rudolph Serbet. 1994. Lignophyte phylogeny and the evolution of spermatophytes: a numerical cladistic analysis. *Systematic Botany* 19: 443-482.
- Samigullin, Tagir Kh., William F. Martin, Aleksey V. Troitsky, Andrey S. Antonov. 1999. Molecular data from the chloroplast *rpoC1* gene suggest a deep and distinct dichotomy of contemporary spermatophytes into two monophyla: gymnosperms (including Gnetales) and angiosperms. *Journal of Molecular Evolution* 49: 310-315.

This Page Intentionally Left Blank

6

EVOLUTION OF FLOWERING PLANTS

ANGIOSPERM APOMORPHIES	121	Sieve Tube Members	132
Flower	121	Angiosperm Specializations	132
Stamens	124	Vessels	133
Reduced Male Gametophyte	126	ORIGIN OF ANGIOSPERMS	133
Carpel	127	REVIEW QUESTIONS	135
Two Integuments	130	EXERCISES	136
Reduced Female Gametophyte	130	REFERENCES FOR FURTHER STUDY	136
Endosperm Formation	130		

The flowering plants, or angiosperms (also called Angiospermae, Magnoliophyta, or Anthophyta), are a monophyletic group currently thought to be the sister group to the gymnosperms (Chapter 5). Angiosperms are by far the most numerous, diverse, and successful extant plant group, containing well over 95% of all land plant species alive today. Flowering plants grow in virtually every habitable region and are dominant in some aquatic and most terrestrial ecosystems, the notable exception to the latter being coniferous forests. Angiosperms comprise the great bulk of our economically important plants, including our most valuable food crops (Chapter 1).

Several apomorphies distinguish the angiosperms from all other land plants (Figure 6.1): (1) the flower, usually with an associated perianth; (2) stamens with two lateral thecae, each composed of two microsporangia; (3) a reduced, 3-nucleate male gametophyte; (4) carpels and fruit formation; (5) ovules with two integuments; (6) a reduced, 8-nucleate female gametophyte; (7) endosperm formation; and (8) sieve tube members. Some of these apomorphic features, which represent the product of a unique evolutionary event, have become further modified in particular lineages of angiosperms (see Chapters 7, 8).

Figure 6.1 shows a simplified cladogram of the major groups of angiosperms. The diversity and classification of these groups are discussed in Chapter 7 (Amborellales, Nymphaeales, Austrobaileyales, Magnoliids, Ceratophyllales, and Monocots) and Chapter 8 (Eudicots). The following is

a review of flowering plant apomorphies and general evolutionary history.

ANGIOSPERM APOMORPHIES

FLOWER

Perhaps the most obvious distinguishing feature of angiosperms is the **flower** (Figure 6.2; see Chapter 9 for detailed terminology of flower parts). A flower can be defined as a *modified, determinate shoot* system bearing one or more **stamens**, collectively called the **androecium**, and/or one or more **carpels** (making up one or more **pistils**), collectively called the **gynoecium** (see later discussion). Most angiosperm flowers are **bisexual (perfect)**, containing both stamens and carpels, but some are **unisexual (imperfect)**, having only stamens or carpels. In addition, most (but not all) flowers have a **perianth**, consisting of modified leaves at the base of the shoot system.

The perianth of a flower both protects the other floral parts during floral development and functions as an attractant for pollination (see later discussion and Chapter 13). Most flowers have a perianth of *two* discrete whorls or series of parts: an outer **calyx** and an inner **corolla** (Figure 6.3A). The calyx is generally green and photosynthetic, composed of leaf-like **sepals** or (if these are fused) of **calyx lobes**. The corolla is typically colorful, showy, and odoriferous and is composed

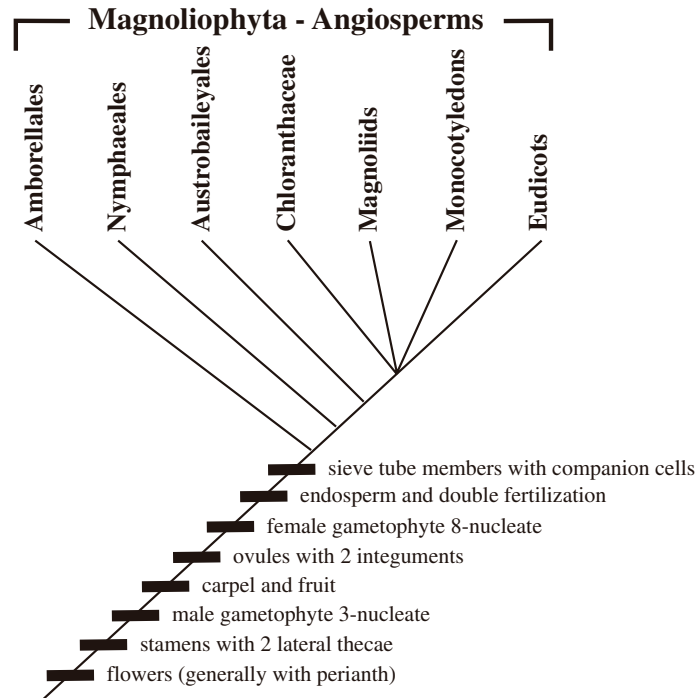


FIGURE 6.1 Cladogram of the angiosperms, showing apomorphies and major taxonomic groups, the latter after APG II (2003).

of individual **petals** or (if these are fused) of **corolla lobes**. However, in some flowering plants, there are two whorls of parts, but the outer and inner whorl of perianth parts are not otherwise differentiated, resembling one another in color and texture. The term **tepals** is often used for such similar perianth parts, and one may refer to **outer tepals** and **inner tepals** for the two whorls (Figure 6.3B). More rarely, the perianth may consist of a single whorl (this usually called the calyx, by tradition) or of three or more discrete whorls (see Chapter 9). Finally, the perianth of some flowers consists of spirally arranged units that grade from sepal-like structures on the outside to petal-like structures on the inside, but with no clear point of differentiation between them; in this case, the units may be termed **tepals**, **perianth parts**, or **perianth segments** (Figure 6.3C).

The components of a flower develop in a manner very similar to leaves. In early floral development actively dividing regions of cells grow, forming bumpy mounds of tissue, the primordia. Typically, the primordia develop in whorls from outside to inside, in sequence as sepal (or outer tepal) primordia first, petal (or inner tepal) primordia second, stamen primordia third (often in two or more whorls), and carpel primordia last (Figure 6.4A–C). Each primordium typically becomes innervated by one or more vascular bundles (veins); primordia may also transform into a flattened, or dorsiventral (having a dorsal and ventral side) shape,

resembling leaves. Fusion of floral parts may occur after they form, termed postgenital fusion. Alternatively, floral parts may appear to be fused at maturity but may actually develop as a single structure. For example, the basal tube of a corolla in which the petals are fused (known as a sympetalous corolla; see Chapter 9) may form by vertical expansion of a ring of actively dividing tissue that develops beneath discrete primordia; only the upper corolla lobes may develop from discrete primordia. Overall, the resemblance of floral organs to leaves in terms of initiating like leaf primordia of a vegetative shoot, being innervated by veins, and often having a dorsiventral shape is why these organs sepals, petals, stamens, and carpels are thought to be homologous to leaves (Chapter 2).

Ongoing studies of the molecular basis of development in plants, especially those using the species *Arabidopsis thaliana* (termed the *Drosophila* of the plant world), have helped to elucidate the genetic basis of floral development and the nature of these presumed homologies. Research in this field is summarized in the ABC model of floral development, in which gene products of the so-called A, B, and C classes combine to produce the four major floral organs: sepals, petals, stamens, and carpels (Figure 6.5). In this model, sepals are expressed by A activity alone; petals by a combination of A and B activities, stamens by a combination of B and C activities, and carpels by C activity alone (Figure 6.5).

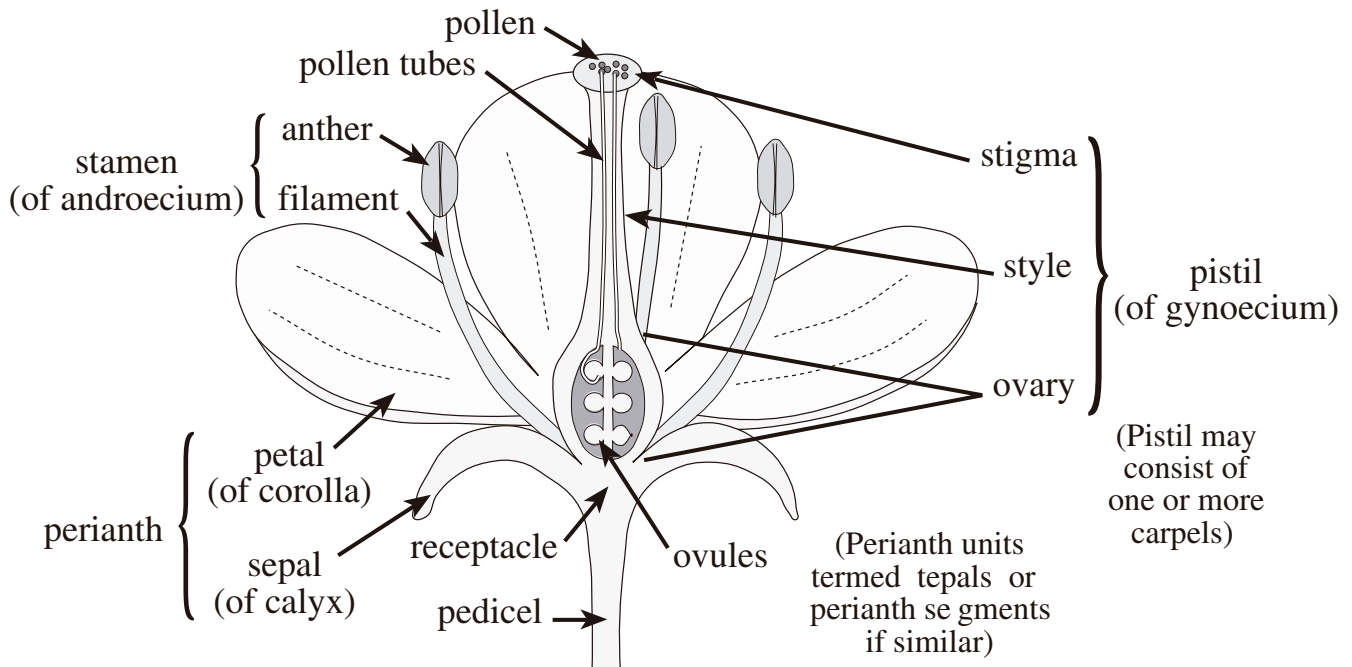


FIGURE 6.2 A typical (diagrammatic) flower, illustrating the parts.

In addition, genes of the so-called SEPALLATA class are needed in combination with those of the A, B, and C classes to effect proper floral organ identity (Figure 6.5). All of these floral organ identity genes work by producing transcription factors in the proper location of the flower, i.e., in the outermost, second, third, and innermost floral whorls. The transcription factors induce the expression of other genes that bring about the development of the four floral organs. Developmental studies like these, in a wide range of species, will help to understand both the molecular basis of homology and

the mechanisms of evolution that have given rise to the rich diversity of floral forms.

The flower, with its typically showy and often scented perianth, evidently evolved in response to selective pressure for the transfer of pollen by animals. Animal pollination appears to be the primitive condition in the angiosperms, separating them from the predominantly wind-pollinated gymnosperms (Chapter 5). Numerous, intricate pollination mechanisms have evolved in various angiosperm lineages. These pollination mechanisms have largely driven the evolution of innumerable

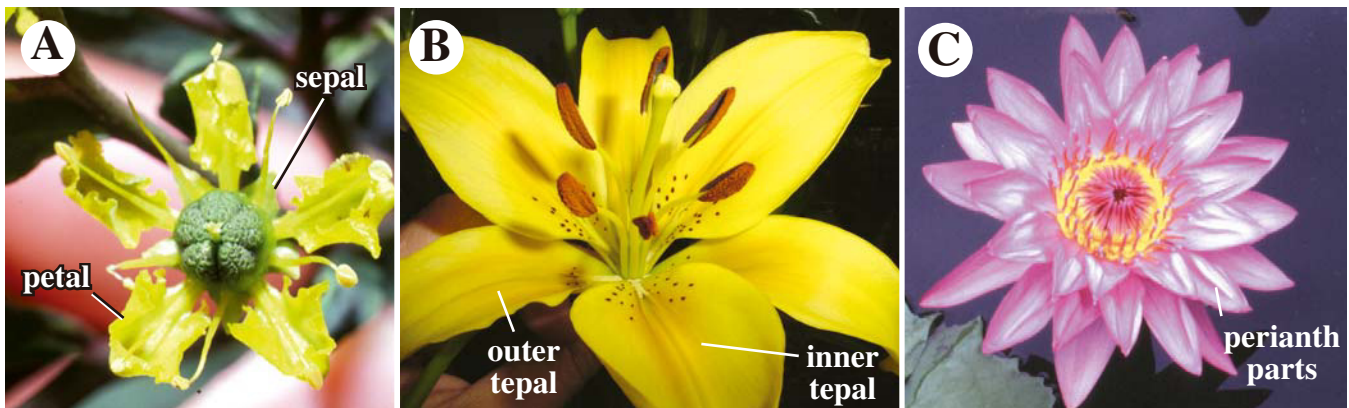


FIGURE 6.3 Various perianth types in flowers. **A.** Perianth of two whorls, dissimilar in appearance: a calyx of sepals and a corolla of petals (*Ruta*). **B.** Perianth of two whorls, similar in appearance: outer tepals and inner tepals (*Lilium*). **C.** Perianth undifferentiated, spiral (*Nymphaea*).

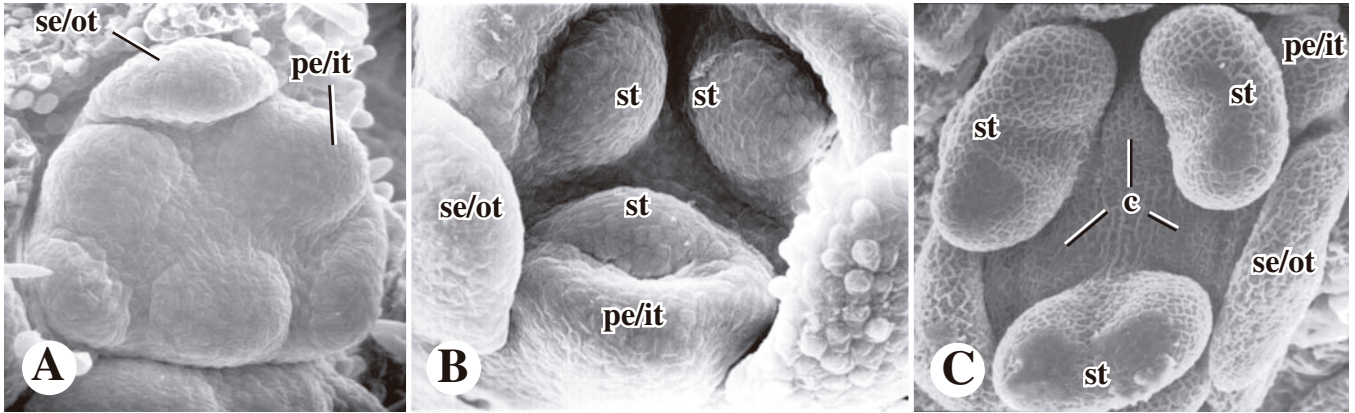


FIGURE 6.4 Flower development. **A.** Early development of sepal/outer tepal (se/ot) primordia and petal/inner tepal (pe/it) primordia. **B.** Later formation of stamen (st) primordia. **C.** More mature stamens and early initiation of carpel (c) primordia.

oral forms, accounting in large part for the distinctiveness of many angiosperm families (see Chapter 13 for floral syndromes related to pollination biology). Animal pollinators may include bees (Figure 6.6A), butterflies and moths (Figure 6.6B), flies (Figure 6.6C), bats (Figure 6.6D,E), and birds (Figure 6.6F). However, flowers of many groups are quite reduced in size or structural complexity, often lacking a perianth altogether; these may be water pollinated (Figure 6.6G) or wind pollinated (Figure 6.6H).

STAMENS

A distinctive apomorphy for the angiosperms is the **stamen**, the male reproductive organ of a flower. Stamens are interpreted as modified microsporophylls, modified leaves that bears microsporangia (see Chapter 5). Microsporangia produce microspores, which develop into pollen grains (Chapter 5; see later discussion). Some stamens have a laminar (leaf-like) structure, to which the anther is attached or embedded (Figure 6.7A). However, the stamens of most flowering plants have two parts: a stalk, known as a **filament**, and the pollen bearing part, known as the **anther** (Figure 6.7B). Some stamens lack a filament (or lamina), in which case the anther is **sessile**, directly attached to the rest of the flower.

The angiosperm anther is a type of synangium, a fusion product of sporangia. Anthers are unique in (ancestrally) containing two pairs of microsporangia arranged in a bilateral symmetry (i.e., having two mirror image halves). Each pair of microsporangia is typically located within a discrete half of the anther called a **theca** (plural, **thecae**; Figure 6.7C). Thus, such an anther consists of two thecae (termed **bithecal**), each theca having two microsporangia for a total of four (termed **tetrasporangiate**; Figure 6.7D). At maturity, the two microsporangia of a theca typically coalesce into a single, contiguous chamber, called the **anther locule**; each theca

then opens to the outside by a specific dehiscence mechanism, releasing the pollen (Figure 6.7E). (Note that anthers of some angiosperms are secondarily reduced to a single theca, known as **monothecal** or **bisporangiate**, a distinctive systematic character; see Chapters 7-9.)

The adaptive value of the stamens of angiosperms over the microsporophylls of gymnosperms is likely connected with selective pressures for the flower itself. Stamens are generally smaller and lighter than gymnosperm microsporophylls, and stamens generally occur in bisexual flowers, rather than in more massive, unisexual cones. Modifications of the stamen

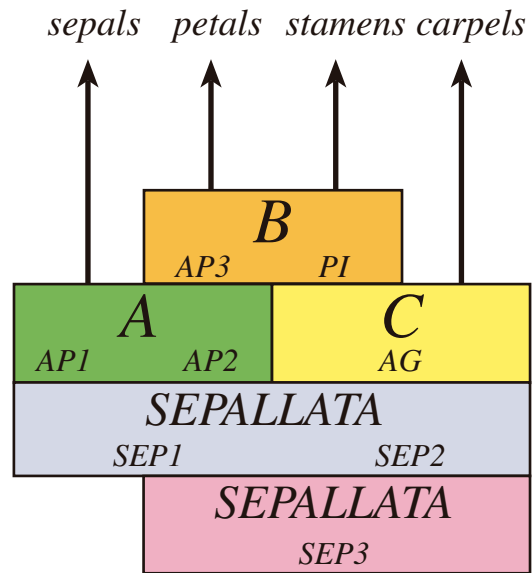


FIGURE 6.5 The ABC model of floral development. Within each gene class are specific genes (AP1, AP2, AP3, AG, PI, SEP1, SEP2, SEP3), identified in mutant forms in *Arabidopsis thaliana*. (Diagram after Jack, 2001.)

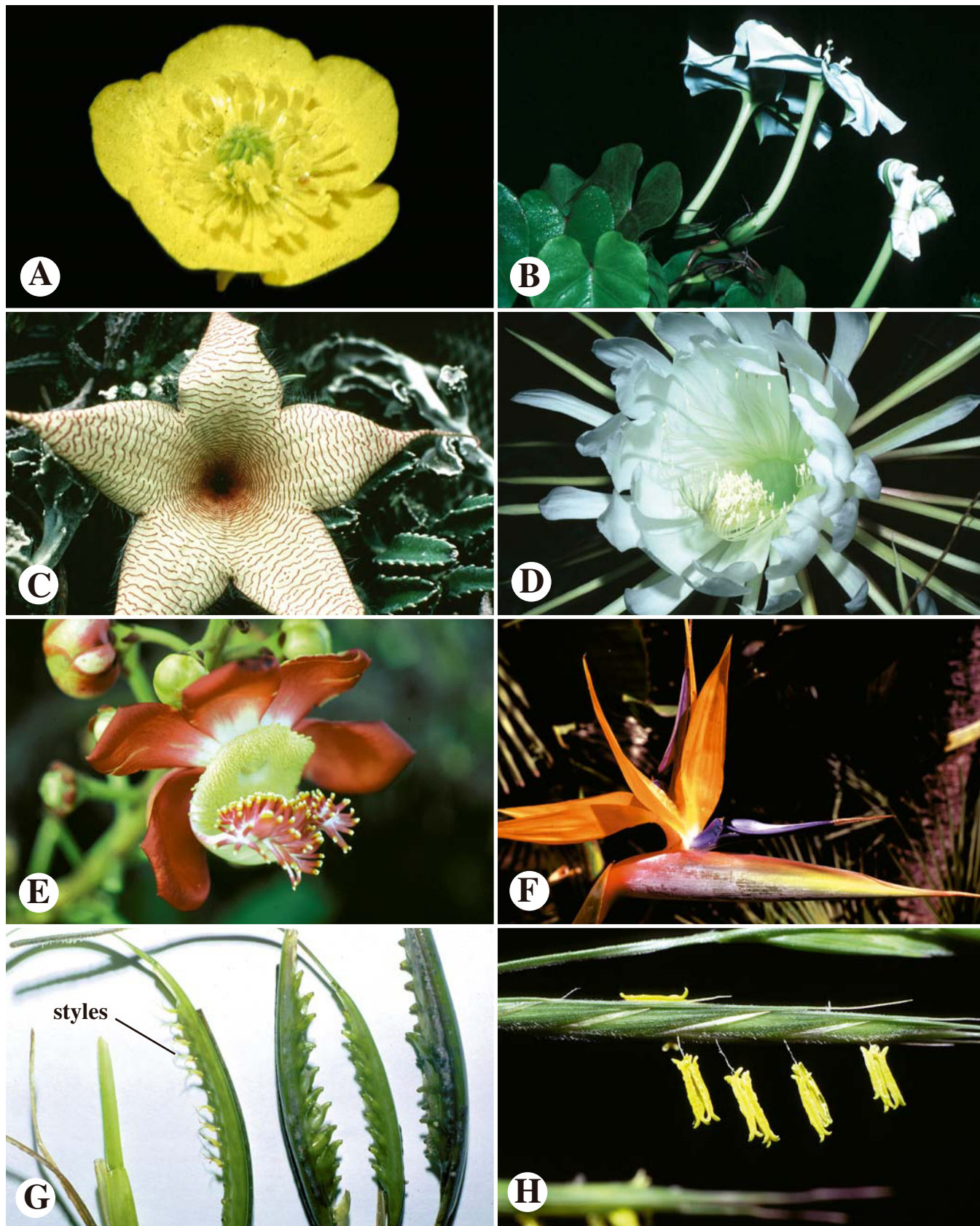


FIGURE 6.6 Flower modifications. **A.** *Ranunculus* sp., buttercup, insect-pollinated. **B.** *Calonyction* sp., moonflower, moth-pollinated. **C.** *Stapelia* sp., starflower, fly-pollinated. **D.** *Selenicereus*, night-blooming cactus, bat-pollinated. **E.** *Couroupida guianensis*, cannonball tree, bat-pollinated. **F.** *Strelitzia reginae*, bird of paradise, bird-pollinated. **G.** *Phyllospadix torreyi*, surf-grass, water-pollinated. **H.** Grass, wind-pollinated.

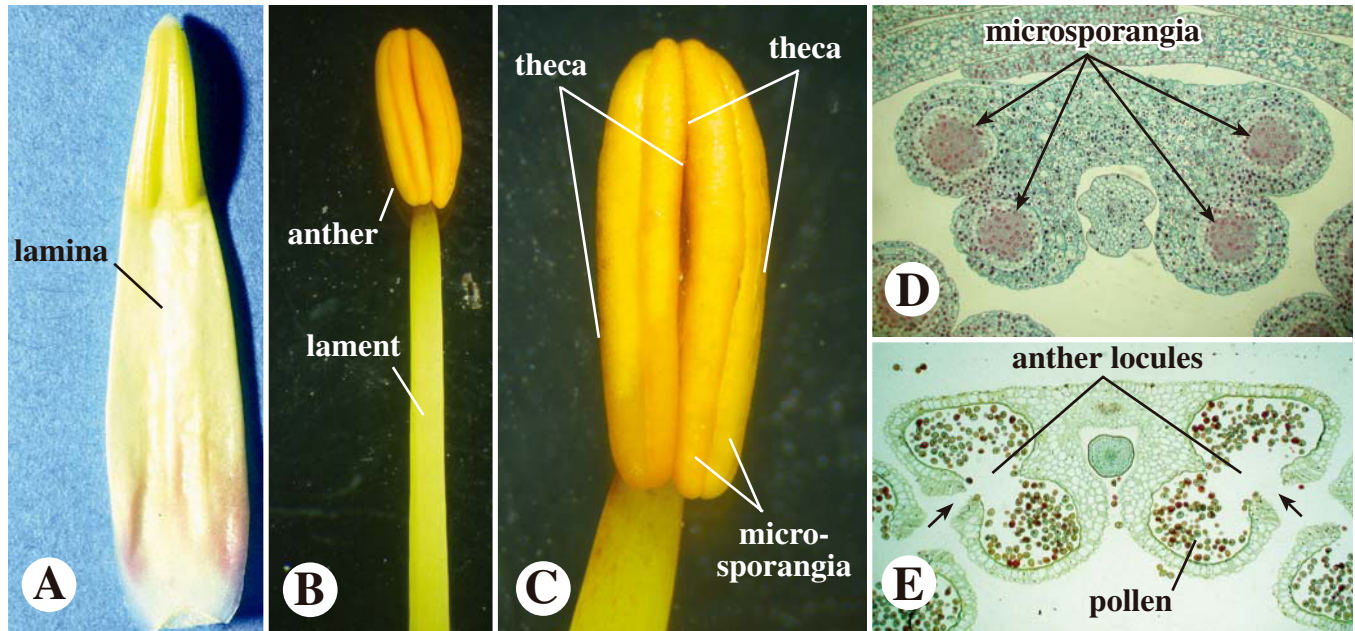


FIGURE 6.7 Stamen morphology. **A.** Laminar stamen, *Nymphaea*. **B,C.** Filamentous stamen, *Aloe*. Note anther composed of two thecae, each with two microsporangia. **D.** Young anther in cross-section, showing four microsporangia. **E.** Cross-section of older anther at time of dehiscence. Note that walls between adjacent microsporangia of each theca have broken down. Dehiscence line indicated by arrows.

have enabled the evolution of specialized pollination mechanisms, such as those involving stamens of the proper length or orientation to transfer pollen to a specific pollinator, flower heteromorphism (associated with stamens at different levels in the flower relative to differing style/stigma lengths), trigger devices, and very modified stamens such as pollinia (see Chapters 12 and 13 for more details).

REDUCED MALE GAMETOPHYTE

Another apomorphy for the angiosperms is a **reduced, three-celled male gametophyte** (Figure 6.8). No other plant group has a male gametophyte so reduced in cell number. After each **microspore** is formed by meiosis within the microsporangium, its single nucleus divides mitotically to form two cells: a **tube cell** and a **generative cell** (Figure 6.8A,B). When this happens, the microspore is transformed to an immature, endosporic male gametophyte or **pollen grain** (Chapter 5). The generative cell divides one time, producing **two sperm cells** (Figure 6.8A). Pollen grains are shed in either a two- or three-celled condition, depending on whether the generative cell division occurs before or after the pollen grains are released. If pollen is released as two-celled, then the generative cell divides within the pollen tube as it travels down the style (Figure 6.8A). Whether pollen grains are 2- or 3-nucleate at release can be an important taxonomic character (Chapter 11).

The pollen grains of angiosperms, like those of gymnosperms, germinate during development, meaning that an

elongate **pollen tube** grows out of the pollen grain wall, a condition known as **siphonogamy** (Figure 6.8A,C,D). In gymnosperms the pollen tube develops after the pollen grains enter the micropyle of the ovule and functions as a haustorial device (feeding from the tissues of the nucellus) for a long period of time (see Chapter 5). In contrast, the pollen tube of angiosperms forms immediately after transfer of pollen to the stigma. The pollen tube of angiosperms elongates through (and feeds upon) the tissues of the stigma and style of the carpel and soon reaches the ovule, where it penetrates the micropyle and transports the two sperm cells directly to the female gametophyte (see later discussion). The sperm cells of angiosperms lack flagella or cilia and are thus nonmotile, a derived condition among the land plants. The loss of motility may be a function of the direct transport of the sperm cells to the micropyle of the ovule. The only other land plants with nonmotile sperm cells are the gymnospermous conifers (including the Gnetales), which lost sperm motility independently of flowering plants.

The adaptive significance of the reduced male gametophytes of angiosperms is probably correlated with the evolution of a reduced female gametophyte and relatively rapid seed development (discussed later). In gymnosperms fertilization of sperm and egg occurs long after pollination, sometimes as long as a year; the male gametophytes must persist during this long period, feeding off the tissues of the nucellus. In angiosperms, however, fertilization occurs very soon

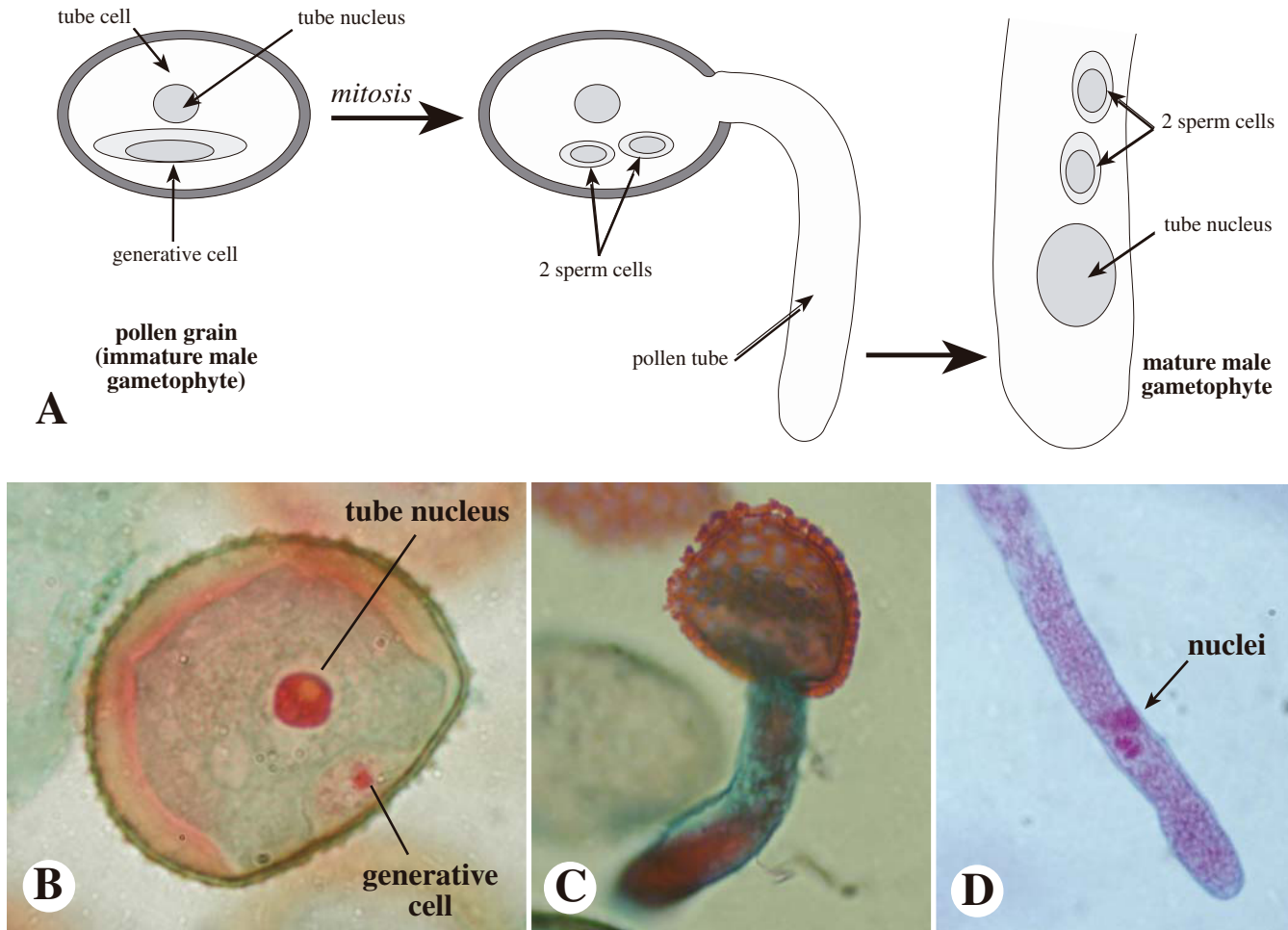


FIGURE 6.8 Angiosperm male gametophyte. **A.** Development of reduced 3-nucleate male gametophyte from pollen grain. **B.** Mature binucleate pollen grain, with tube nucleus and generative cell. **C.** Germinating pollen grain, forming pollen tube. **D.** Tip of pollen tube, housing nuclear material (nuclei types unclear in this image).

after pollination. Thus, angiospermous male gametophytes are lean, apparently requiring a minimum number of cells and nuclei; they function to deliver sperm cells to the female gametophyte and effect fertilization very rapidly compared with gymnosperms.

CARPEL

A major apomorphy of angiosperms is the **carpel**. According to the most widely accepted hypothesis, the carpel constitutes a modified, conduplicate megasporophyll bearing two, adaxial rows of ovules (Figure 6.9D). (Recall that a megasporophyll is a modified leaf that bears megasporangia, which in the seed plants are components of the ovules and seeds; see Chapter 5. Conduplicate means inwardly folded longitudinally and along the central margin; see Chapter 9.) This megasporophyll is modified in that the margins by virtue of the conduplicate folding come together and fuse

(Figures 6.9A D, 6.10A), with certain parts differentiating into tissue for pollen reception and pollen tube growth, typically forming an apical stigma and style (Figure 6.9D). At maturity the carpel body completely encloses the ovules and seeds, accounting for the name angiosperm (Gr. *angio*, vessel + *sperm*, seed).

The sporophyll-like nature of the carpel is evident in that (1) it may develop like a leaf, having an initially attenuated, dorsiventral shape, with an adaxial (toward the top-center of the flower) and abaxial (away from the top-center of the flower) surface; and (2) it has veins, typically one in the middle termed the **dorsal** (median) vein or bundle, corresponding to the midvein of a leaf, and two others near the two carpel margins termed the **ventral** (lateral or placental) veins/bundles (Figures 6.9D, 6.10A). Additional veins often occur between the dorsal and ventral bundles (e.g., Figure 6.10B), and veins will sometimes fuse together. The veins of a carpel are

typically collateral (see Chapter 10), with xylem on the adaxial side and phloem on the abaxial side. The ventral veins become inverted in orientation after carpel formation, with the xylem and phloem disposed 180° from their original orientation, i.e., prior to conduplicate folding (Figure 6.9D).

The carpels of some angiosperm taxa show no evidence of a conduplicate, leaf-like nature during development. It is generally accepted that these have become secondarily modified

or specialized, particularly in compound pistils (see later discussion). One type, known as an **ascidiate** carpel, develops from a ring of tissue that grows upward, sometimes assuming a somewhat peltate form. However, taxa that lack a conduplicate carpel development usually still have inverted ventral veins, evidence of the ancestral condition.

A given flower can have one to many carpels. If two or more carpels are present, they may be separate from one

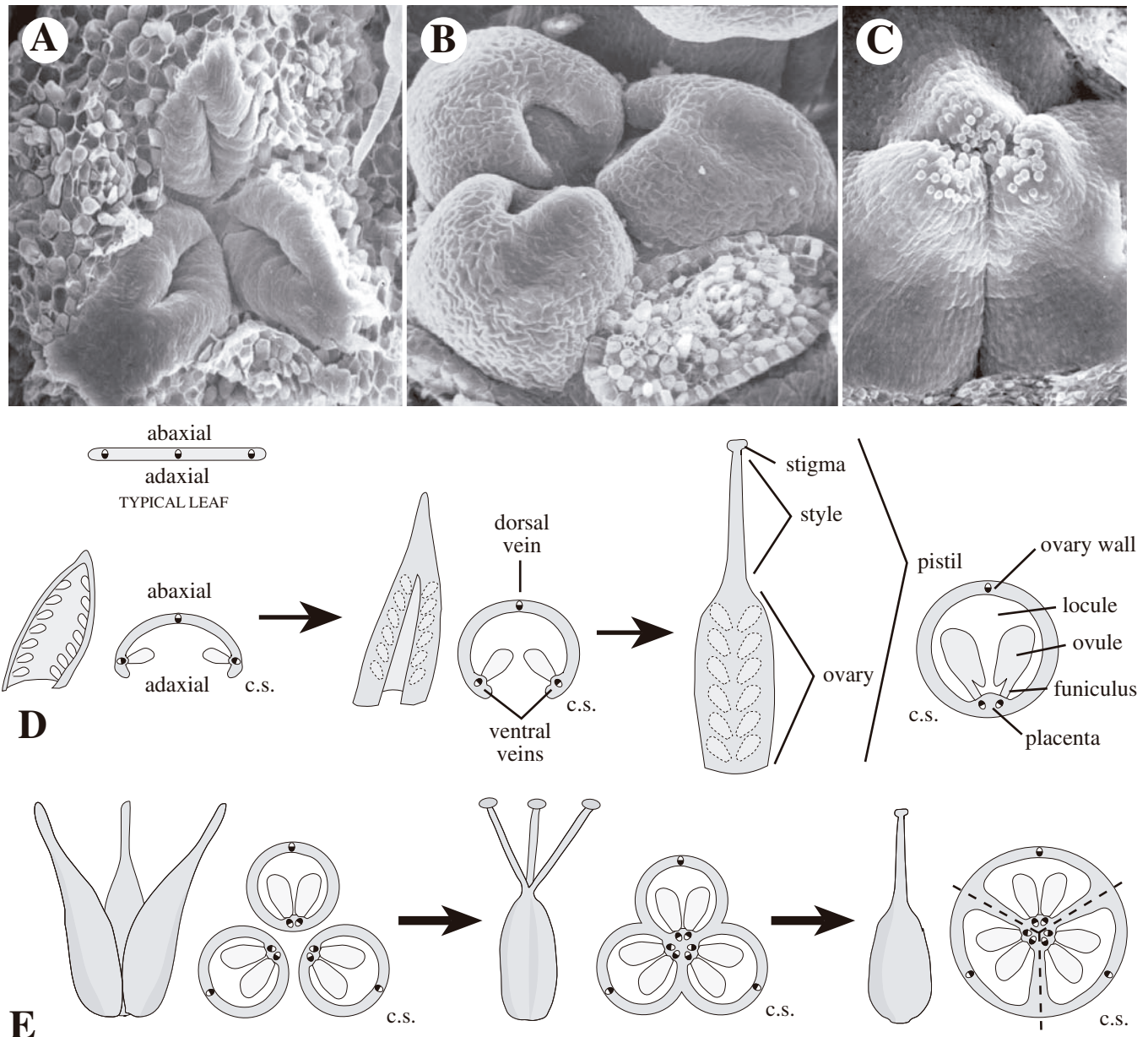


FIGURE 6.9 The carpel, an apomorphy of the angiosperms. **A–C.** Scanning electron micrographs of carpel development. **A.** Early formation of three carpels, showing conduplicate formation. **B.** Intermediate developmental stage. Note lateral contact of the three carpels. **C.** Mature stage, in which carpel margins have closed in and adjacent carpels have fused into a syncarpous gynoecium (compound pistil). **D.** Diagram of carpel development from early stages to mature ovary, adaxial side below. Note dorsal and ventral veins (black=xylem; white=phloem), the latter becoming inverted. **E.** Diagram illustrating evolutionary sequence of carpel fusion (dashed lines=carpel boundaries).

another (distinct), termed **apocarpous**, or fused together (connate), termed **syncarpous**. Because of the frequent fusion of carpels, additional terms are useful in describing the female parts of a flower. The term **gynoecium** is the totality of female reproductive structures in a flower, regardless of their structure. Thus, a carpel may be alternatively defined as a unit of the gynoecium. The gynoecium is composed of one or more **pistils**. Each pistil consists of a basal **ovary**, an apical **style** (or styles), which may be absent, and one or more **stigmas**, the tissue receptive to pollen grains (Figure 6.9D). A pistil may be equivalent to one carpel (in which case, it may be termed a **simple pistil**) or composed of two or more, fused carpels (termed a **compound pistil**; Figures 6.9E, 6.10B). (The position of one or more ovules and the fusion of one or more carpels determine various placentation types; see Chapter 9 for complete terminology.)

The evolution of the carpel had considerable adaptive significance. First, because carpels are the receivers of pollen, they may function to selectively control fertilization. The transfer of pollen to the carpels is followed by germination of the pollen grain to form a pollen tube, which grows through the tissue of the stigma and style to the micropyle of the ovule. However, chemicals that are present in the stigma and style may inhibit either pollen germination or pollen tube growth; this is known as an **incompatibility reaction**, mediated by incompatibility genes (see Chapter 13). This type of chemical incompatibility often occurs between the pollen and stigmatic regions of *different* species. However, it may

also occur between individuals of the *same* species, notably between individuals that are genetically similar and possess the same incompatibility alleles. Thus, incompatibility reactions may inhibit inbreeding, allowing for reproduction only between genetically dissimilar individuals of the species (i.e., promoting out-crossing; see Chapter 13 for more details). Thus, the carpel may ultimately provide some selective control as to which pollen grains contribute the sperm cells that fertilize the egg.

A second major adaptive function of the carpel pertains to fruit formation and seed dispersal. A **fruit** is the mature ovary or ovaries (made up of one or more carpels) plus any accessory tissue that might be present (see Chapter 9). Fruits generally do not mature from ovaries if fertilization of the seed(s) does not occur. The mature ovary wall, termed the **pericarp**, may be highly modified. These modifications generally function in a tremendous variety of dispersal mechanisms (Chapter 9). In general, if the pericarp is fleshy, fruits are dispersed by animals. In these fleshy, animal-dispersed fruits, the seeds are transported either by passing through the gut of the animal unharmed (with only the pericarp being digested) or by being spilled during a sloppy eating session. Dry fruits may also be dispersed by animals, but typically via external barbs or prickles that catch on skin, fur, or feathers. Last, fruits may be dispersed by wind (aided by the development of wings or trichomes), water (via various flotation devices), or mechanically (by various explosive, hygroscopic, or catapulting methods).

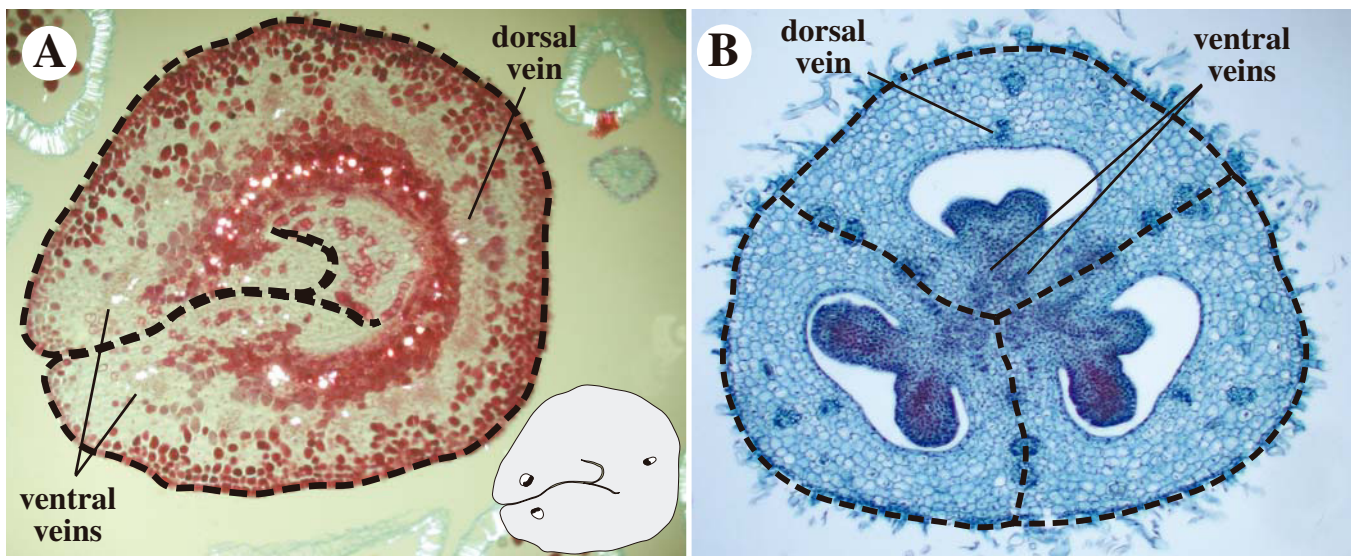


FIGURE 6.10 **A.** Ovary cross-section of a taxon with a single carpel per flower (unicarpellate gynoecium). Note outline of carpel boundary (dashed line). Inset diagram: note orientation of xylem (black) and phloem (white) of veins. **B.** Ovary cross-section of a taxon with a 3-carpellate, syncarpous pistil (carpels outlined by dashed lines), showing dorsal and ventral veins. (Note: vascular bundle outside dorsal vein supplies perianth and stamens, this ovary being inferior.)

TWO INTEGUMENTS

A unique apomorphy of angiosperms is the growth of **two integuments** during ovule development, the ovules known as **bitegmic** (Figure 6.11). All non flowering seed plants have ovules with a single integument, termed **unitegmic**. The two integuments of angiosperms usually completely surround the nucellus, forming a small pore at the distal end; this opening, the **micropyle**, is the site of pollen tube entrance. Both of the integuments of angiosperm ovules contribute to the seed coat. The two integuments typically coalesce during seed coat development, but may form anatomically different layers.

The possible adaptive significance of two integuments, if any, is not clear, but may have enabled the evolution of specialized seed coat layers, although differential seed coat layers are found in several gymnosperm taxa as well. Interestingly, several angiosperm lineages have secondarily lost an integument, and are thus unitegmic. Notable unitegmic groups are many Poales of the Monocots (Chapter 7) and most of the Asterids of the Eudicots (Chapter 8).

REDUCED FEMALE GAMETOPHYTE

Several novelties of the angiosperms have to do with the evolution of a specialized type of ovule and seed. A major apomorphy of angiosperms is a **reduced 8-nucleate female gametophyte**. As in other seed plants, a single megasporocyte within the megasporangium (nucellus) divides meiotically to form four haploid megaspores (Figure 6.12). The female gametophyte typically generates from only one of these megaspores (Figure 6.12), with a few exceptions in which others may contribute (see Chapter 11). Typically, the megaspore divides in a sequence of three mitotic divisions,

resulting in a total of eight haploid nuclei. Further differentiation usually results in an arrangement of these eight nuclei into seven cells (Figures 6.12, 6.14A). In the micropylar region three cells develop: an **egg cell** flanked by two **synergid cells**. Egg plus synergids is sometimes called the **egg apparatus**. In the **chalazal** region, which is opposite the micropyle, three **antipodal cells** form. The remaining volume of the female gametophyte is technically a single cell, called the **central cell**, which contains two **polar nuclei**. Archegonia do not form within the female gametophyte of angiosperms as they do in virtually all other seed plants. The female gametophyte in various angiospermous taxa may become further modified from the ancestral type described here by variations in cell divisions, nuclear fusions, and cell formations (see Chapter 11). (**Note:** The female gametophyte of angiosperms is often called an **embryo sac**; this terminology, although often used, is to be avoided, as it fails to denote the homology with the female gametophyte of other seed plants.)

The significance of a reduced female gametophyte in flowering plants is likely correlated with developmental timing. Fertilization in angiosperms occurs very shortly after pollination, unlike that of the gymnosperms, in which a long period of time may ensue between the two events. Thus, angiosperms have the capacity to more quickly generate seeds. This feature may be of tremendous adaptive value, enabling, for example, the evolution of rapidly spreading annual herbs.

ENDOSPERM FORMATION

Another major apomorphy of the angiosperms is the presence of **endosperm**. Endosperm is the product of **double fertilization**. When the pollen tube enters the micropyle of the ovule,

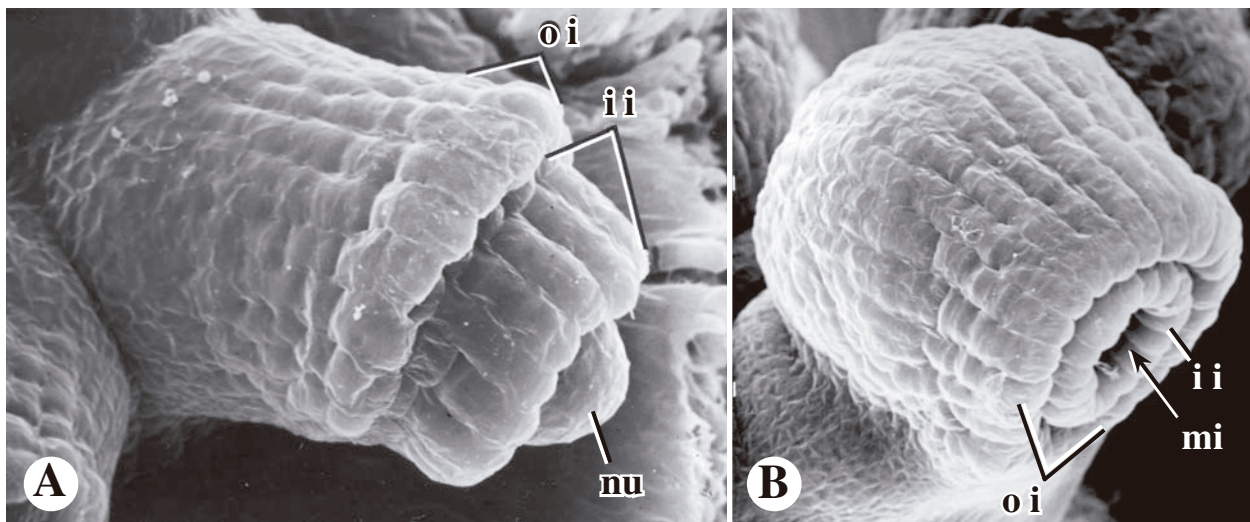


FIGURE 6.11 Bitegmic ovule, the ancestral condition of the angiosperms. **A.** Young ovule, showing initiation of inner integument (ii) and outer integument (oi), both growing around the nucellus (nu). **B.** Older ovule, in which inner and outer integuments have enveloped the nucellus, forming a micropyle (mi).

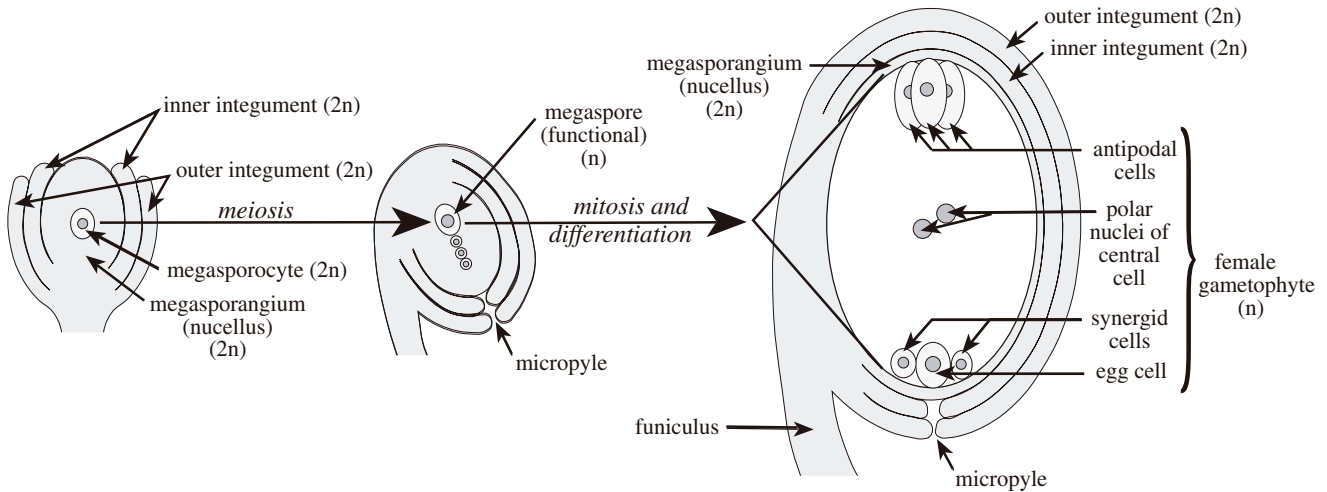


FIGURE 6.12 Angiosperm ovule development and morphology. Note meiosis of megasporocyte, producing four haploid megaspores, one of which undergoes mitotic divisions and differentiation, resulting in an 8-nucleate female gametophyte.

it penetrates one of the synergid cells and releases the two sperm cells into the central cell of the female gametophyte (Figure 6.13). One sperm cell migrates toward and fuses with the egg cell to produce a diploid **zygote**. As in other land plants, the zygote matures into an embryo, with structures similar to those in other seed plants (Figure 6.13). The other sperm cell fuses with the two polar nuclei to produce a triploid, or $3n$, **endosperm cell**. This endosperm cell then repeatedly divides by mitosis, eventually forming the **endosperm**, a mass of tissue that generally envelops the embryo of the seed (Figures 6.13, 6.14B,C). Endosperm replaces the female

gametophyte as the primary nutritive tissue for the embryo in virtually all angiosperms, containing cells rich in carbohydrates, oil, or protein.

The adaptive significance of endosperm is, like that of the reduced female gametophyte, possibly correlated with developmental timing. The endospermous nutritive tissue of angiosperms does not begin to develop until *after* fertilization is achieved. This is in contrast with gymnospermous seed plants, in which considerable female gametophytic nutritive tissue is deposited after pollination, even if the ovules are never ultimately fertilized. Thus, a major selective pressure

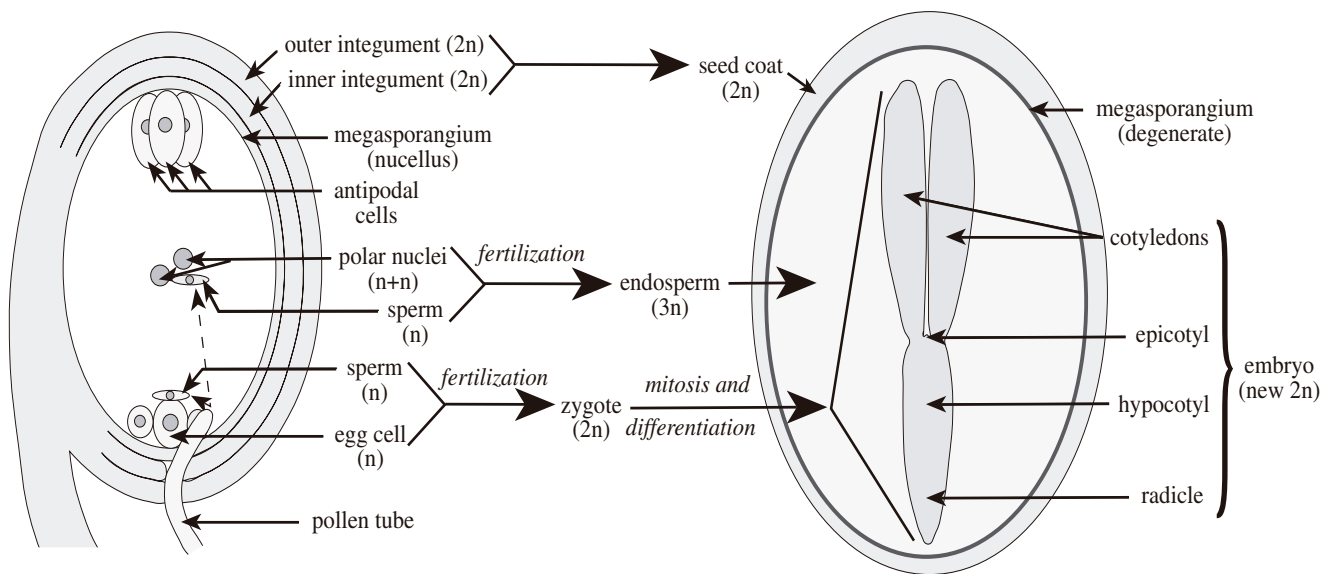


FIGURE 6.13 Angiosperm seed development and morphology. Note fertilization of egg, forming zygote and embryo, and fertilization of polar nuclei, forming triploid endosperm.

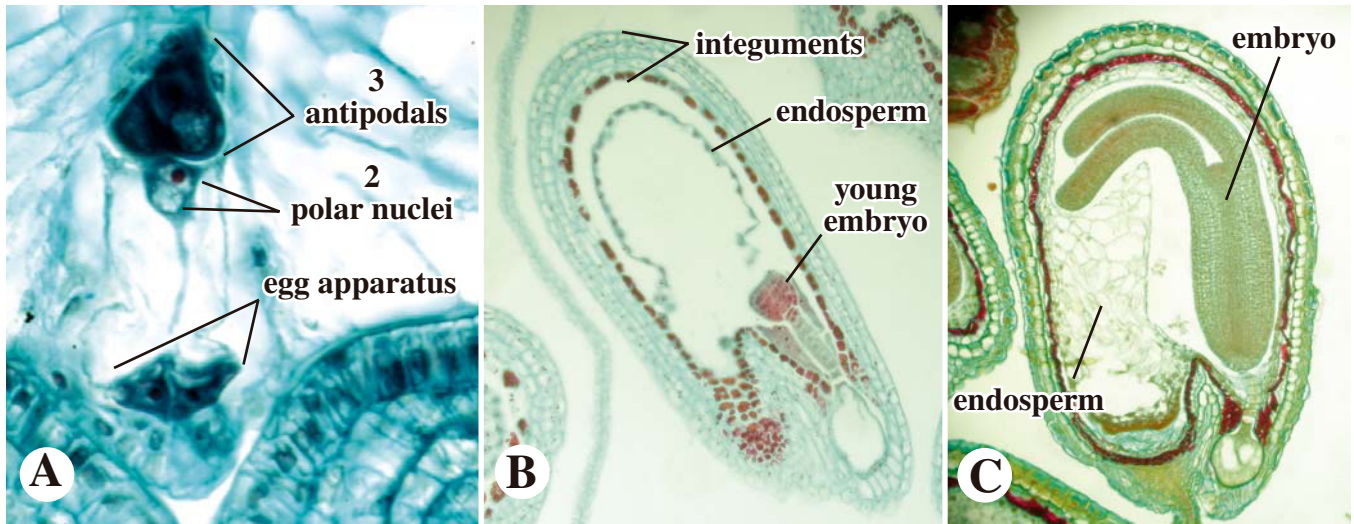


FIGURE 6.14 A. Reduced, 8-nucleate female gametophyte (*Lachnanthes*), showing egg apparatus (egg + synergid cells), polar nuclei, and antipodals. B,C. Endosperm formation (*Capsella*). B. Early stage. C. Later stage, forming seed.

for the evolution of endosperm may have been conservation of resources, such that seed storage compounds are not formed unless fertilization is assured. An additional, functional feature of endosperm derives from the tissue being triploid. Having three sets of chromosomes (one from the male and two from the female) may enable the endosperm to develop more rapidly (correlated with rapid overall seed development) and may also provide greater potential for chemical variation in nutritive contents.

SIEVE TUBE MEMBERS

Angiosperms are unique (with minor exceptions) in having **sieve tube members** as the specialized sugar-conducting cells (Figure 6.15). Sieve cells (and associated albuminous cells) are the primitive sugar-conducting cells and are found in all non-flowering vascular plants (see Chapter 4). Sieve tube members (and associated companion cells) were evolutionarily modified from sieve cells and are found only in flowering plants. Sieve tube members differ from the ancestral sieve cells in that the pores at the end walls are differentiated, being much larger than those on the side walls. These collections of differentiated pores at the end walls are called **sieve plates**. Sieve plates may be either compound (composed of two or more aggregations of pores) or simple (composed of one pore region). Parenchyma cells associated with sieve tube members are called **companion cells**. Companion cells function to load and unload sugars into the cavity of sieve tube members. Unlike the similar albuminous cells of gymnosperms, companion cells are derived from the same parent cell as the conductive sieve tube members.

The adaptive significance of sieve tube members over sieve cells is not clear, though they may provide more efficient sugar conduction.

ANGIOSPERM SPECIALIZATIONS

Angiosperms are a tremendously diverse group of seed plants and have evolved a great number of novel structural features. Various lineages of angiosperms have acquired an amazing

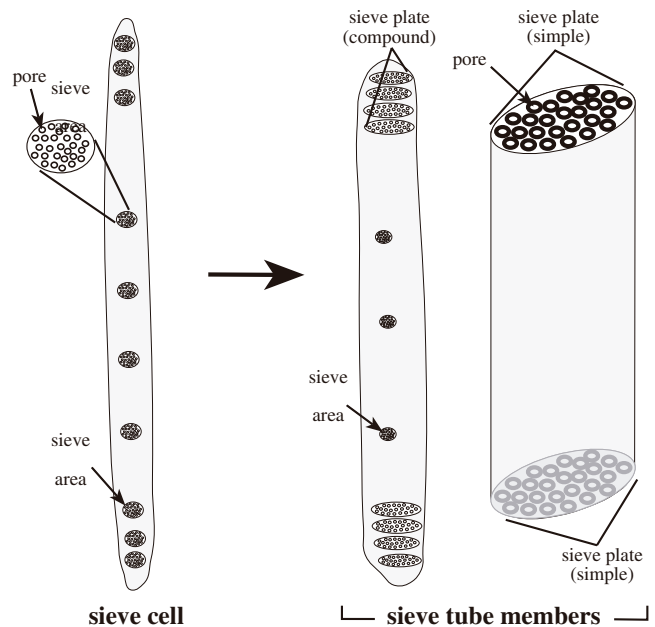


FIGURE 6.15 Evolutionary change from sieve cells (left) to sieve tube members, the latter an apomorphy of the angiosperms.

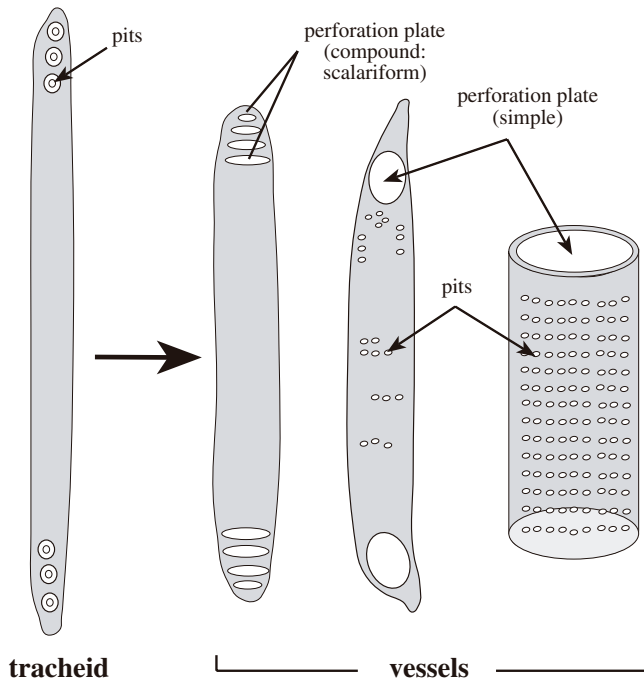


FIGURE 6.16 Evolution of vessels in the angiosperms. Note transformation from imperforate tracheid to vessels with perforation plates. Trends within the angiosperms include change from elongate vessels with scalariform perforation plate to short vessels with simple perforation plates.

variety of specialized roots, stems, and leaf types not found in any other land plant taxa (see Chapters 7-9). And, as mentioned earlier, angiosperms have a number of specialized pollination systems and fruit/seed dispersal mechanisms, by-products of the evolution of flowers and fruits (see Chapter 13).

VESSELS

One angiosperm specialization concerns water and mineral conductive cells. The great majority of angiosperms have **vessels**, in which the two ends of the cells have openings, termed perforation plates (Figure 6.16; see Chapters 4, 10). Vessels constituted a major evolutionary innovation within the angiosperms, presumably providing for more efficient solute conduction. Not all angiosperms have vessels, however, and some basal flowering plant groups (e.g., Amborellales, some Nymphaeales; see Figure 6.1, Chapter 7) are vessel-less, having only tracheids (which lack perforation plates). Thus, vessels may not constitute an apomorphy for the flowering plants as a whole, and likely arose independently in more than one angiosperm lineage.

The tracheids of basal, vessel-less angiosperms characteristically have numerous transversely elongated pits (called scalariform pitting), especially at the tapering end walls where they join other tracheid cells. Tracheids with scalariform

pitting may be the ancestral tracheary element for the angiosperms. In general, primitive vessels resemble tracheids in having scalariform perforation plates (Figure 6.17A) in which the openings consist of numerous, transversely oriented pits. Specializations of vessels (Figure 6.16) include (1) modification of the perforation plate from scalariform to one with fewer, less transversely oriented openings, to a simple perforation plate (having a single opening; e.g., Figure 6.17B,C); (2) modification from tapering end walls to perpendicular ones; and (3) modification from long, narrow cells to short, wide cells (Figure 6.17D).

ORIGIN OF ANGIOSPERMS

As is often stated, Charles Darwin described the relatively rapid diversification of the higher plants (presumed to mean angiosperms) as an abominable mystery. The earliest definitive fossils of flowering plants are dispersed pollen grains from the earliest Cretaceous period, approximately 140 million years ago. The earliest definitive flowers occur slightly later in the fossil record, as early as 130 million years ago. These early flowering plant fossils can largely be assigned to recognizable, extant groups. Once angiosperms arose, they radiated rapidly into several, distinct lineages and gradually replaced gymnosperms as the dominant plant life form on the earth.

However, the details of angiosperm evolution from a gymnosperm precursor are not clear. One problem is what to call an angiosperm. Many angiosperm features cited earlier, such as a reduced male gametophyte, reduced female gametophyte, and double fertilization with triploid endosperm, are microscopic and cytological and would be unlikely to be preserved in the fossil record. Cladistic analyses of extant angiosperms may help elucidate the features possessed by the common ancestor of the flowering plants. Given this, we might expect to find at least some of these features in the closest fossil relatives of the angiosperms. Based on recent cladistic studies, *Amborella trichopoda* of the Amborellales (Figure 6.1) is accepted as the best hypothesis for the most basal angiosperm lineage (see Chapter 7). *Amborella* lacks vessels and has unisexual flowers with a spiral perianth, laminar stamens, and separate carpels. However, other, near-basal lineages of flowering plants vary in these features, making an assessment of the characteristics of the common ancestor of the angiosperms unclear.

An ongoing hypothesis on the origin of angiosperms is that they were derived by modification of some member of the group known as pteridosperms (mentioned in Chapter 5), a paraphyletic assemblage of extinct plants that possessed

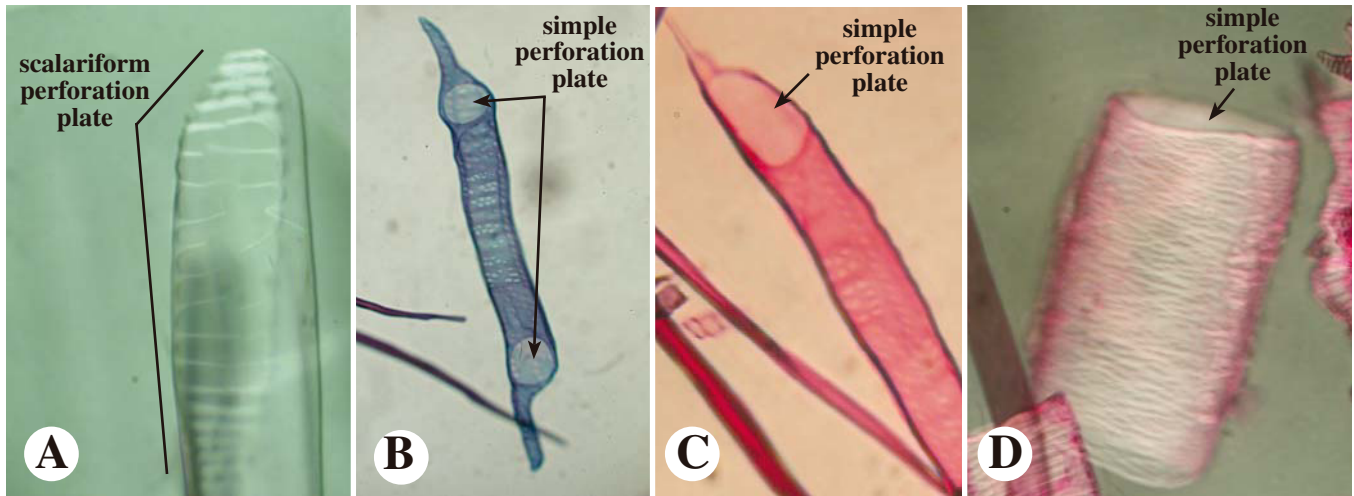


FIGURE 6.17 Variation in vessel anatomy in the angiosperms. **A.** *Liriodendron tulipifera*, with scalariform perforation plates. **B,C.** *Quercus* sp., elongate with simple perforation plates. **D.** *Cucurbita* sp., short and cylindrical, with simple perforation plates.

seeds and had generally fernlike foliage. Some Pteridosperms may represent possible angiosperm progenitors. One fossil taxon that exemplifies a putative transition to angiosperms is *Caytonia* of the Caytoniales (Figure 6.18A–C). *Caytonia* possessed reproductive structures similar to those of the angiosperms. The male reproductive structures resemble anthers in consisting of a fusion product (synangium) of three or four microsporangia; however, these differ from angiosperm anthers in being radially (not bilaterally) symmetric (Figure 6.18A). The female reproductive structures of *Caytonia* consist of a spikelike arrangement of units that have been termed **cupules** (Figure 6.18B,C). Each cupule encloses a cluster of unitegmic ovules/seeds, with a small opening in the cupule near the proximal end (Figure 6.18C). The cupule has been hypothesized as being homologous with the angiosperm carpel. However, the cupule of *Caytonia* is different from what is presumed to be the ancestral carpel morphology, a conduplicate megasporophyll bearing ovules along two margins. In addition, (monosulcate) pollen grains have been discovered at the micropyle of *Caytonia* ovules, evidence that the pollen grains were transported directly to the ovules (perhaps by means of a pollination droplet, as occurs in extant gymnosperms), rather than to a stigmatic region where pollen tubes formed. Thus, the cupule apparently did not function as a carpel in terms of a site for pollen germination. Another interpretation of the cupule of *Caytonia* is that it is the homologue of the second integument apomorphic of all angiosperms, evolving by the reduction of the number of ovules within the cupule to one. In summary, the homology of the reproductive structure in *Caytonia* is difficult to decipher, and no other pteridosperm is clearly an angiosperm progenitor.

However, some pteridosperms, like *Caytonia*, may still be more closely related to the angiosperms than to the gymnosperms (see Figure 5.1).

An example of a fossil that may help elucidate early angiosperm evolution is the genus *Archaeoфраuctus*, fairly recently collected from China, and evidently now dated to no earlier than 130 million years ago of the early Cretaceous. *Archaeoфраuctus* (with two described species) was apparently an aquatic plant, having dissected leaves and elongate reproductive axes, each of the latter with paired stamens below and several-seeded carpels above (Figure 6.18D,E). Although *Archaeoфраuctus* appears to have bona fide carpels, its relationship to extant angiosperms is debatable. By one hypothesis the reproductive axis is interpreted as an entire, perianth-less flower (with stamens below and carpels above), the axis perhaps homologous to an elongate receptacle reminiscent of some Magnoliaceae (see Chapter 7). By this interpretation, this reproductive structure might represent an ancestral flower (or flower precursor), and *Archaeoфраuctus* might be sister to the extant angiosperms. An alternative hypothesis views the reproductive axis of *Archaeoфраuctus* not as a single, atypical flower, but as an inflorescence of individual, reduced male and female flowers, as seen in some aquatic angiosperms today. By this viewpoint, *Archaeoфраuctus* may just as likely represent an extinct off-shoot of an extant lineage within the angiosperms (such as the Nymphaeales).

In summary, it seems that more fossils may need to be discovered and described (or reinvestigated with new techniques) before the abominable mystery can be satisfactorily solved. Cladistic analyses help, but there is always the problem of

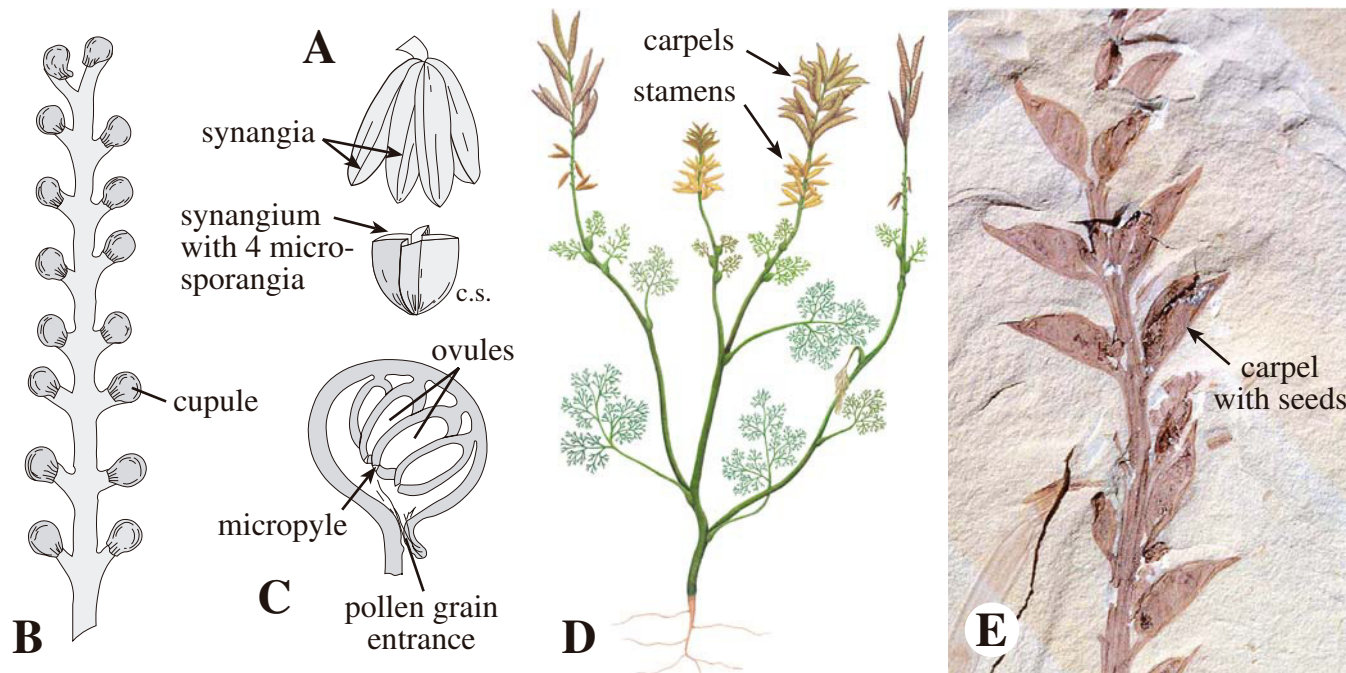


FIGURE 6.18 **A–C.** *Caytonia*, diagram redrawn from Thomas (1925). **A.** Cluster of male reproductive units, each a radially symmetrical synangium of 4 microsporangia. (Cross section=c.s.) **B.** Reproductive axis, bearing two rows of cupules. **C.** Cupule, in sagittal section, showing four ovules and opening at base. **D,E.** *Archaeoфраuctus*. **D.** Reconstruction of *Archaeoфраuctus sinensis*, showing reproductive axis bearing stamens proximally and carpels distally. (Contributed by K. Simons and David Dilcher (').) **E.** Fossil impression of carpel units of *Archaeoфраuctus lianogensis*. (Contributed by David Dilcher (') and Ge Sun.)

homology assessment with structures that are vastly different from contemporary forms. Despite the fact that the relationships among extant flowering plants are much better known with advanced molecular techniques (see Chapter 7), fossils

will be key to understanding their origin. Paleobotanical work should be continuously emphasized as of the utmost importance in understanding plant relationships.

REVIEW QUESTIONS

ANGIOSPERM APOMORPHIES

1. What is another name for the flowering plants?
2. Name the apomorphies of the flowering plants.
3. What is the definition of a flower?
4. Name the major components of a typical flower.
5. Describe the morphology and adaptive significance of the perianth.
6. What is the ABC model of floral development, and what species served as the original exemplar for this?
7. What was a major selective pressure that resulted in the evolution of specialized types of flowers?
8. What is unique about the angiosperm stamen, and what are the types and parts of a stamen?
9. What is a theca and of what is it composed?
10. What about the male gametophyte of flowering plants is unique?
11. Describe the structure and function of a mature male gametophyte in the flowering plants.
12. What is the definition of a carpel?
13. What is the difference between carpel, pistil, and gynoecium?
14. Name and describe two major adaptive features of the carpel.

15. Contrast integument number in gymnosperms versus that in angiosperms.
16. Draw and label a mature female gametophyte in the flowering plants.
17. How many cells and nuclei are present in a typical, mature, female gametophyte of the flowering plants?
18. How might the reduced angiospermous female gametophyte be adaptive?
19. What is endosperm and what is its function?
20. What is the difference between a sieve cell and a sieve tube member? In what groups are each found?
21. What type of tracheary element do most angiosperms have, and what is its adaptive significance?

ANGIOSPERM ORIGINS

22. When are the earliest definitive angiosperm fossils found?
23. Describe the example of *Caytonia* as a putative angiosperm progenitor, citing evidence for and against this idea.
24. Describe the reproductive structure of *Archaeofructus* and indicate two competing hypotheses for its homology.

EXERCISES

1. Collect and observe a flowering plant. Looking at specific parts of the plant, go over in your mind the apomorphies (both macroscopic and microscopic) that have enabled the angiosperms to dominate the world's vegetation. Especially review all parts of a flower, citing the adaptive significance of each component.
2. Place various angiospermous pollen grains on a microscope slide, stain (e.g., with toluidine blue), and observe these reduced male gametophytes under a microscope. Look for the cells and nuclei inside. Are the pollen grains two-celled or three-celled at maturity?
3. Observe an angiosperm ovule in sagittal-section under the microscope. Look for the two integuments and the (typically) eight nuclei and seven cells of the female gametophyte.
4. Contrast popcorn (an angiosperm) with pine nuts (a gymnosperm) in terms of the ploidy level and development of the nutritive tissue. Cite the selective advantage that flowering plant seeds might have in this regard.

REFERENCES FOR FURTHER STUDY

- Andrews, H. N. 1961. *Studies in Paleobotany*. Wiley, New York.
- Angiosperm Phylogeny Group. 2003. An update of the Angiosperm Phylogeny Group classification for the orders and families of flowering plants: APG II. *Botanical Journal of the Linnean Society* 141: 399–436.
- Crane, P. R., E. M. Friis, and K. Pedersen. 1995. The origin and early diversification of angiosperms. *Nature* 374: 27.
- Crepet, W. L. 1998. The abominable mystery. *Science* 282: 1653–1654.
- Cronquist, A. 1981. *An integrated system of classification of flowering plants*. Columbia University Press, New York.
- Davies, T. J., T. G. Barraclough, M. W. Chase, P. S. Soltis, D. E. Soltis, and V. Savolainen. 2004. Darwin's abominable mystery: insights from a supertree of the angiosperms. *Proceedings of the National Academy of Sciences of the United States of America* 101: 1904–1909.
- Friis, E. M., J. A. Doyle, P. K. Endress, and Q. Leng. 2003. *Archaeofructus*: Angiosperm precursor or specialized early angiosperm? *Trends in Plant Science* 8: 369–373.
- Friis, E. M., K. R. Pedersen, and P. R. Crane. 2000. Reproductive structure and organization of basal angiosperms from the Early Cretaceous (Barremian or Aptian) of western Portugal. *International Journal of Plant Sciences* 161: S169–S182.
- Jack, T. 2001. Relearning our ABCs: new twists on an old model. *Trends in Plant Science* 6: 310–316.
- Jenik, P. D., and V. F. Irish. 2000. Regulation of cell proliferation patterns by homeotic genes during *Arabidopsis* floral development. *Development* 127: 1267–1276.
- Stebbins, G. L. 1974. *Flowering Plants: Evolution above the Species Level*. Belknap Press of Harvard University Press, Cambridge, MA.
- Sun, G., D. L. Dilcher, S. Zheng, and Z. Zhou. 1998. In search of the first flower: a Jurassic angiosperm, *Archaeofructus*, from Northeast China. *Science* 282: 1692–1695.
- Sun, G., Q. Ji, D. L. Dilcher, S. Zheng, K. C. Nixon, and X. Wang. 2002. *Archaeofractaceae*, a new basal angiosperm family. *Science* 296: 899–904.
- Takhtajan, A. L. 1991. *Evolutionary Trends in Flowering Plants*. Columbia University Press, New York.
- Thomas, H. H. 1925. The Caytoniales, a new group of angiospermous plants from the Jurassic rocks of Yorkshire. *Philosophical Transactions of the Royal Society of London* 213: 299–363.
- Veit, B., R. J. Schmidt, S. Hake, and M. F. Yanofsky. 1993. Maize floral development: new genes and old mutants. *The Plant Cell* 5: 1205–1215.
- Zanis, M. J., P. S. Soltis, Y. L. Qiu, E. Zimmer, and D. E. Soltis. 2003. Phylogenetic analyses and perianth evolution in basal angiosperms. *Annals of the Missouri Botanical Garden* 90: 129–150.

7

DIVERSITY AND CLASSIFICATION OF FLOWERING PLANTS: AMBORELLALES, NYMPHAEALES, AUSTROBAILEYALES, MAGNOLIIDS, CERATOPHYLLALES, AND MONOCOTS

INTRODUCTION	138	MONOCOTYLEDONS	153
MAJOR ANGIOSPERM CLADES	138	Monocot Apomorphies	155
FAMILY DESCRIPTIONS	140	Classification of the Monocotyledons	156
AMBORELLALES	141	ACORALES	157
Amborellaceae	142	Acoraceae	157
NYMPHAEALES	143	ALISMATALES	160
Nymphaeaceae	143	Araceae	160
Cabombaceae	143	Alismataceae	160
AUSTROBAILEYALES	143	ASPARAGALES	163
Illiciaceae	145	Agavaceae	165
MAGNOLIIDS	146	Alliaceae	165
LAURALES	146	Amaryllidaceae	169
Lauraceae	146	Asphodelaceae	171
MAGNOLIALES	146	Iridaceae	171
Annonaceae	146	Orchidaceae	171
Magnoliaceae	149	Themidaceae	177
PIPERALES	149	DIOSCOREALES	178
Aristolochiaceae	149	Dioscoreaceae	178
Piperaceae	149	LILIALES	180
Saururaceae	153	Liliaceae	180
CERATOPHYLLALES	153	PANDANALES	180
Ceratophyllaceae	153	Pandanaaceae	180
		COMMELINIDS	184

(Continued)

ARECALES	185	POALES	203
Arecaceae (Palmae)	185	Bromeliaceae	203
COMMELINALES, ZINGIBERALES, AND POALES	188	Typhaceae	205
COMMELINALES	188	Sparganiaceae	206
Commelinaceae	188	Juncaceae	206
Haemodoraceae	190	Cyperaceae	206
Pontederiaceae	190	Eriocaulaceae	211
ZINGIBERALES	192	Xyridaceae	212
Musaceae	192	Restionaceae	212
Strelitziaceae	195	Poaceae (Gramineae)	213
Zingiberaceae	197	REVIEW QUESTIONS	219
Cannaceae	198	EXERCISES	220
		REFERENCES FOR FURTHER STUDY	221

INTRODUCTION

The phylogenetic relationships within the angiosperms has been and continues to be a field of active research in plant systematics. Much progress has been made with the use of cladistic methodology and the incorporation of morphological, anatomical, embryological, palynological, karyological, chemical, and molecular data (see Chapters 9–14). The more recent use of multiple gene sequence data has been particularly useful in assessing higher-level angiosperm relationships. However, the phylogenetic relationships and classification presented in this chapter can be viewed as somewhat preliminary, to be further refined with continued research. For a more precise understanding of relationships within a particular group, there is no substitute for consulting the most recent, primary scientific literature.

MAJOR ANGIOSPERM CLADES

Portrayal of the relationships of major angiosperm groups is modeled (with very few exceptions) after the system of the Angiosperm Phylogeny Group, 2003 (referred to as APG II, 2003), which supersedes Angiosperm Phylogeny Group, 1998. The APG II system is based on published cladistic analyses primarily utilizing molecular data (e.g., Chase et al. 1993, 2000; Graham and Olmstead 2000b; Soltis et al. 1997, 2000; Qui et al. 2000; Zanis et al. 2002) or a combination of morphological and molecular data (e.g., Nandi et al. 1998). In the APG II system, an attempt was made to recognize only those angiosperm families that are monophyletic. In many cases, angiosperm families have been redefined from their past, traditional circumscription, either being split into separate groups (e.g., the traditional Liliaceae and Scrophulariaceae)

or united into one family (e.g., the Bombaceae, Malvaceae, Sterculiaceae, and Tiliaceae united into one family, Malvaceae, s.l.). The APG II system classifies one to several families into orders (thus, each group having the ending *-ales*), where strong evidence suggests that the order is monophyletic. It must be understood, however, that the designated orders are not comparable evolutionary units and are not indicative of a hierarchical classification system (see Chapter 2). For example, a single order may be sister to a monophyletic group containing several orders. The orders can be viewed simply as convenient placeholders for one or more families that appear to comprise a monophyletic group with relatively high certainty. Some monophyletic groups containing several orders are given informal names, such as Magnoliids, Monocots, Eudicots, Rosids, Eurosids I and II, Asterids, and Euasterids I and II.

The precise interrelationships of the major groups of angiosperms still show some uncertainty, but recent results have begun to converge. Figure 7.1 illustrates higher-level phylogenetic relationships from various analyses that are summarized in APG II, 2003. Note that some polytomies occur; further research may, in time, resolve many of these. In particular, the elucidation of the most basal branches of the flowering plants may yield insight into early angiosperm evolution and radiation.

As seen in Figure 7.1, the angiosperms can be broadly delimited into several groups: the Amborellales, Nymphaeales, Austrobaileyales, Chloranthaceae, Magnoliids (consisting of Laurales, Magnoliales, Canellales, and Piperales), monocotyledons (the Monocotyledonae or monocots), Ceratophyllales, and the eudicots. Of these major groups, the current chapter deals with all but the eudicots, which are covered in Chapter 8. Those angiosperm groups other than the eudicots are sometimes referred to as basal flowering plants because

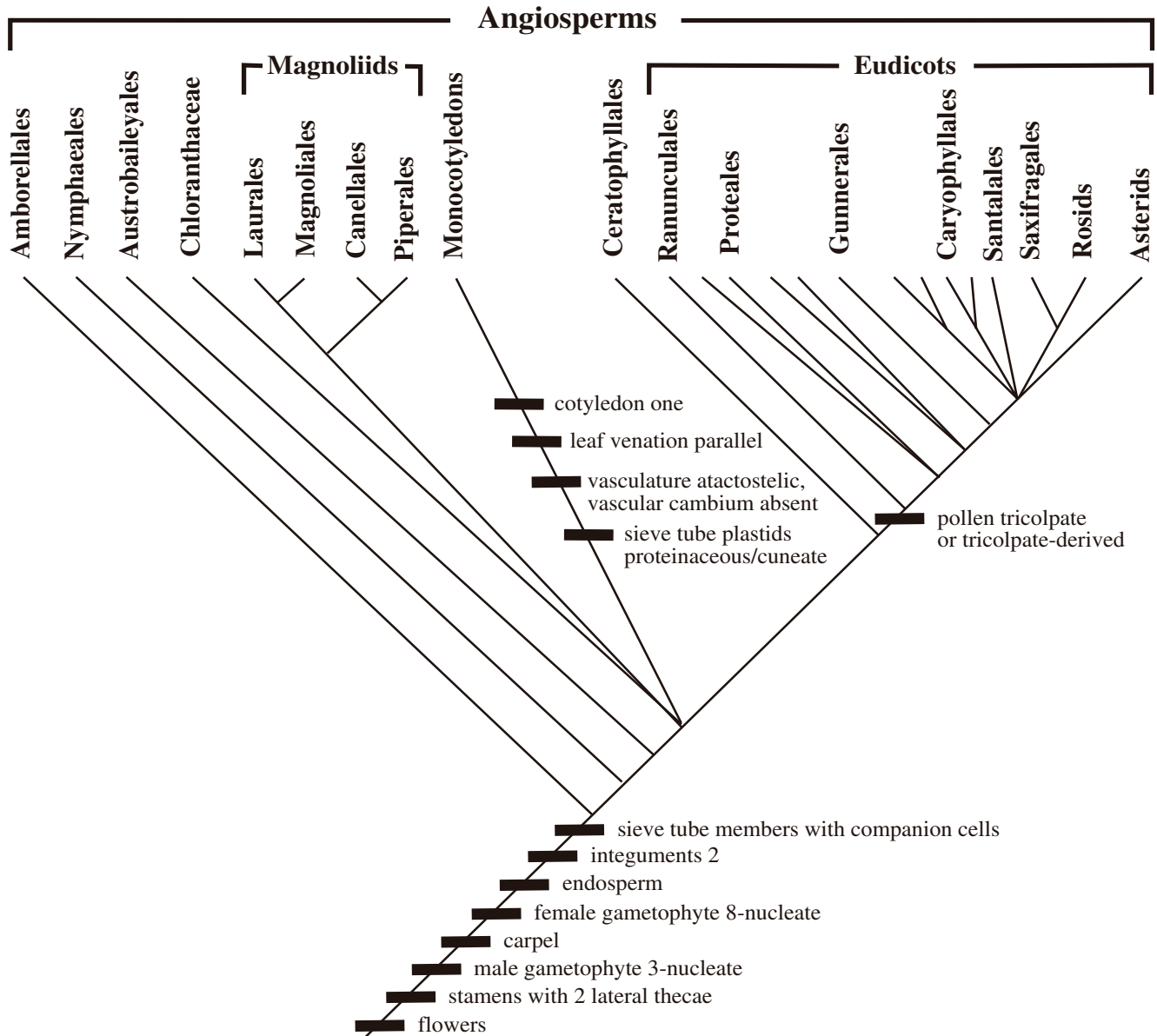


FIGURE 7.1 Phylogenetic relationships of major angiosperm clades, from APG II, 2003, with selected apomorphies shown (see Chapter 6).

they include the first lineages that diverged from the common ancestor of the angiosperms. However, as portrayed in Figure 7.1, it is evident that this is an arbitrary designation, in that some of these groups are no more basal than the eudicots. The families within the orders are listed in Table 7.1 (all except the monocots), Tables 7.2 (basal monocots) and 7.3 (commelinid monocots); eudicot families are listed in Tables 8.1–8.3 of Chapter 8.

The great bulk of the angiosperms in terms of species diversity are contained within the monocots and eudicots. The monocotyledons are a large group, containing approximately 22% of all angiosperms (see later discussion). The eudicots comprise a very large group, including approximately 75% of all angiosperms, and will be treated separately in Chapter 8.

The traditionally defined group Dicotyledonae, the dicotyledons or dicots, have been defined in the past by their possession of embryos with two cotyledons. It is now thought that the possession of two cotyledons is an ancestral feature for the taxa of the flowering plants and not an apomorphy for any group within. Thus, dicots as traditionally delimited (all angiosperms other than monocots), are paraphyletic and must be abandoned as a formal taxonomic unit.

In the descriptions in this chapter and in Chapter 8, exemplars are used for each order or other major group. The choice of these exemplars is very limited in the context of the huge diversity of the angiosperms. These treatments are not designed as a substitute for the many fine references on flowering plant family characteristics (see the references at the end of

TABLE 7.1 Major groups of the angiosperms, listing orders and included families (after APG II, 2003) for groups other than monocots (see Tables 7.2, 7.3) and eudicots (see Chapter 8). Families in **bold** are described in detail. An asterisk denotes an acceptable deviation from APG II, with brackets indicating the more inclusive family recommended by APG II.

ANGIOSPERMS		
AMBORELLALES*	MAGNOLIIDS	MAGNOLIIDS (continued)
Amborellaceae	LAURALES	CANELLALES
Nymphaeales*	Atherospermataceae	Canellaceae
Nymphaeaceae*	Calycanthaceae	Winteraceae
Cabombaceae* [Nymphaeaceae]	Gomortegaceae	PIPERALES
AUSTROBAILEYALES	Hernandiaceae	Aristolochiaceae
Austrobaileyaceae	Lauraceae	Hydnoraceae
Illiciaceae* [Schisandraceae]	Monimiaceae	Lactoridaceae
Schisandraceae	Siparunaceae	Piperaceae
Trimeniaceae	MAGNOLIALES	Saururaceae
Chloranthaceae	Annonaceae	MONOCOTS (see Table 7.2, p. 159; Table 7.3, p. 185)
	Degeneriaceae	Ceratophyllales
	Eupomatiaceae	Ceratophyllaceae
	Himantandraceae	EUDICOTS (see Chapter 8)
	Magnoliaceae	
	Myristicaceae	

this chapter), but are intended as an introduction to some of the common or important groups for the beginning student.

Taxa at the traditional rank of family are utilized as exemplar units; in a few cases subfamilies or tribes are described. Only major, general features of commonly encountered plant families are presented, with examples cited to show diagnostic features. More thorough descriptions and illustrations of angiosperm families may be obtained from references cited in the family descriptions and listed at the end of the chapter.

FAMILY DESCRIPTIONS

The family descriptions that follow use technical terms that are defined and illustrated in Chapter 9 and listed in the Glossary; some embryological or anatomical terms are defined in Chapters 10 and 11. The descriptions begin with a heading that lists the family name (scientific and common), the etymology of the family name where known (Gr. = Greek; L. = Latin), and the number of genera and species. The first paragraph is a description of plant characteristics of the family members, starting with plant habit and vegetative features, in the order of **root, stem, and leaf**. This is followed by reproductive features, in the order of **inflorescence, flower, perianth** (if undifferentiated) or **calyx and corolla** (if differentiated), **androecium, gynoecium, fruit, and seed**. Important anatomical or chemical characteristics are occasionally listed as well. The second paragraph lists infra-familial classification (where pertinent), distribution, and

economically important members of the family. The third paragraph lists the diagnostic features of the family, i.e., how the family can be distinguished from other, related families. This is to aid the beginning student in recognizing the family at a glance; the most important diagnostic features are shown in **boldface-italics**. Features thought to represent apomorphies for the family or groups within the family are cited as such. Finally, the family descriptions end with a floral formula.

The **floral formulas** are used to summarize the number and fusion of floral parts. In these formulas, **P** refers to perianth parts and is used where the perianth is undifferentiated into a typical outer calyx and inner corolla (e.g., being homochlamydeous, or having outer, calyx-like series and inner corolla-like series that intergrade). If the perianth is differentiated into a distinct calyx and corolla, **K** represents the number of sepals or calyx lobes and **C** the number of petals or corolla lobes. The androecium is denoted by **A** and represents the number of stamens; staminodes may also be tabulated, but are indicated as such in the formula. The gynoecium is denoted by **G**, showing the number of carpels in the gynoecium, followed by superior or inferior to denote ovary position. Connation, the fusion of similar parts, is illustrated with parentheses () that enclose the number . Separate, discrete whorls of parts are separated by the + sign, delimiting the number of parts per whorl; the outermost whorl is indicated by the first number, the innermost whorl by the last number. Numbers that are enclosed by brackets [] represent a less common or rare condition. If there are more than about 10–12 parts, the ∞ sign is used for numerous.

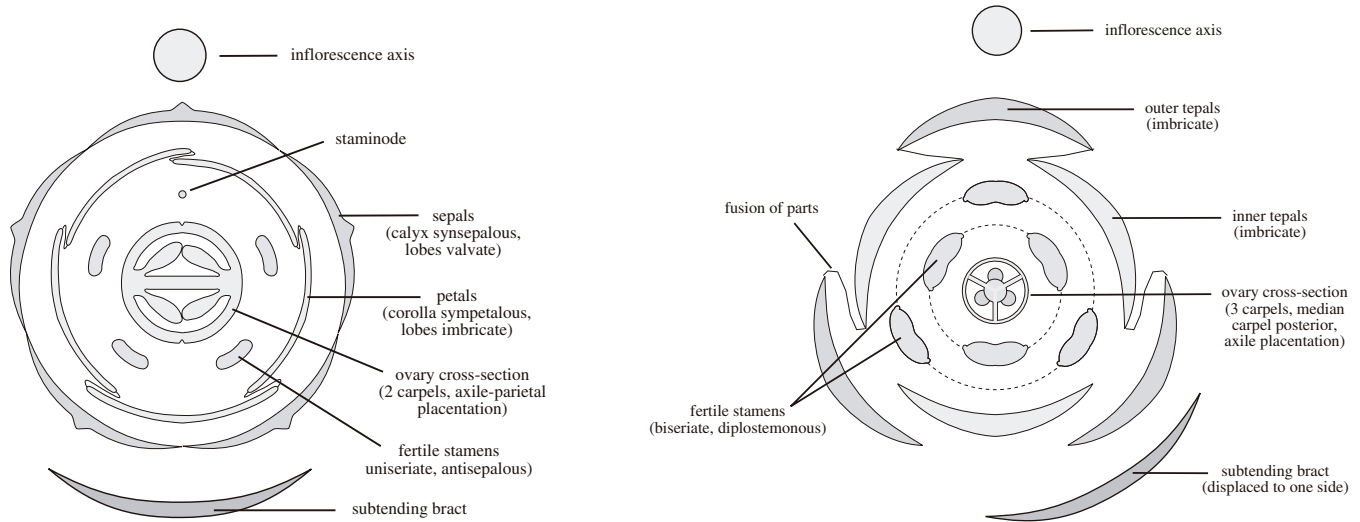


FIGURE 7.2 Floral diagrams, illustrating relative relationships among the components of the perianth, androecium, and gynoecium.

The floral formulas used here summarize the variation that occurs *within the family as a whole*, not necessarily that for a single species. However, floral formulas certainly may also be used to summarize the floral characteristics of a single species. Some hypothetical examples of floral formulas are:

K (5) [(4)] **C** 5 [4] **A** 5+5 [4+4] **G** 5 [4], superior: represents a flower having a synsepalous calyx with five [rarely four] lobes, an apopetalous corolla of five [rarely four] petals, an androecium with ten distinct (not fused to one another) stamens in two whorls of five each [rarely eight stamens in two whorls of four], and an apocarpous gynoecium with five [rarely four], superior-ovary carpels.

P (3+3) **A** 3+3 **G** (3), inferior: represents a flower with a homochlamydeous perianth (i.e., one not delimited into calyx and corolla) having connate, outer and inner whorls of three tepals each, six distinct stamens in two whorls of three each, and a syncarpous, inferior-ovary gynoecium with three carpels.

Family descriptions are accompanied by figures of photographs and line drawings of exemplars. An effort is made to illustrate both diagnostic and apomorphic features. **Floral diagrams** are sometimes illustrated. These represent a diagrammatic cross-sectional view of a flower bud, showing the relative relationship of perianth, androecial, and gynoecial components (examples in Figure 7.2). Floral diagrams may show fusion of floral parts as well as things such as stamen position, placentation, and perianth, calyx, or corolla aestivation (see chapter 9). They are very useful in visualizing floral structure, and, along with floral formulas, are a succinct summary of the characteristics of the group.

The following are detailed descriptions of selected families (shown in **bold** in Table 7.1) from these major groups.

Those selected families were done so largely because live material is more likely to be available for classroom examination and dissection or because of their tremendous importance ecologically or with respect to biodiversity. An attempt was made to describe only information that can be generally seen by the student, unless the characters are of significant diagnostic significance. The source of data for family descriptions was largely taken from *The Plant-Book* (Mabberley 1997), an excellent compendium of descriptions of vascular plant families and genera and highly recommended as a general reference. Another major reference used was Cronquist (1981). Very good recent family descriptions are found in the ongoing series *The Families and Genera of Flowering Plants*: Kubitzki et al. (1993, 1998a,b); Kubitzki and Bayer (2002); and Kubitzki (2004). The family descriptions that follow were often difficult to do, since many families have undergone vastly different circumscriptions in the APG II system. Refer to the references cited earlier and at the end of the chapter for additional information and for descriptions of families not treated here. The *Angiosperm Phylogeny Website* (Stevens, 2001 onward) is an excellent, up-to-date resource for cladograms, classification, references, and apomorphies.

AMBORELLALES

This order comprises one family and one species (below). The Amborellaceae is purported in most molecular studies to be the most basal angiosperm group, although some studies suggest other possibilities (notably that *Amborella* + Nymphaeaceae together are sister to the rest of the

angiosperms). See Mathews and Donoghue (1999, 2000); Qui et al. (1999, 2000); Graham and Olmstead (2000a,b); Parkinson et al. (1999); Barkman et al. (2000); Zanis et al. (2002); and Borsch et al. (2003) for studies on relationships of *Amborella* within the angiosperms. See Doyle and Endress (2000) and Zanis et al. (2003) for a discussion of character evolution in the basal angiosperms.

The absence of vessels in the order, which is rare in angiosperms, is possibly an ancestral condition, and the absence of aromatic (ethereal) oil cells is significant in light of other basal groups that have them.

Amborellaceae Amborella family. (L. for around a little mouth, perhaps in reference to the flower). 1 genus and species. (Figure 7.3)

The Amborellaceae comprises the single species *Amborella trichopoda*, a dioecious, tropical shrub. The **leaves** are alternate, spiral to distichous, undivided, exstipulate, evergreen, and simple. The **inflorescence** is an axillary cyme. The **flowers** are unisexual, actinomorphic, and hypogynous to perigynous. The **perianth** consists of 5–8, spiral, distinct to basally connate perianth parts (termed sepals by default). The **stamens** of male flowers are ∞ , and somewhat laminar. **Anthers** are longitudinal in dehiscence. The **gynoecium** of female flowers is apocarpous, comprising 5–6 superior-ovary pistils that are apically open. **Placentation** is marginal; the **ovule** is solitary in each pistil. The **fruit** is a drupe. Vessels and ethereal oil cells are lacking.

Amborella trichopoda, the single species of the Amborellaceae, is native only to New Caledonia. There are no

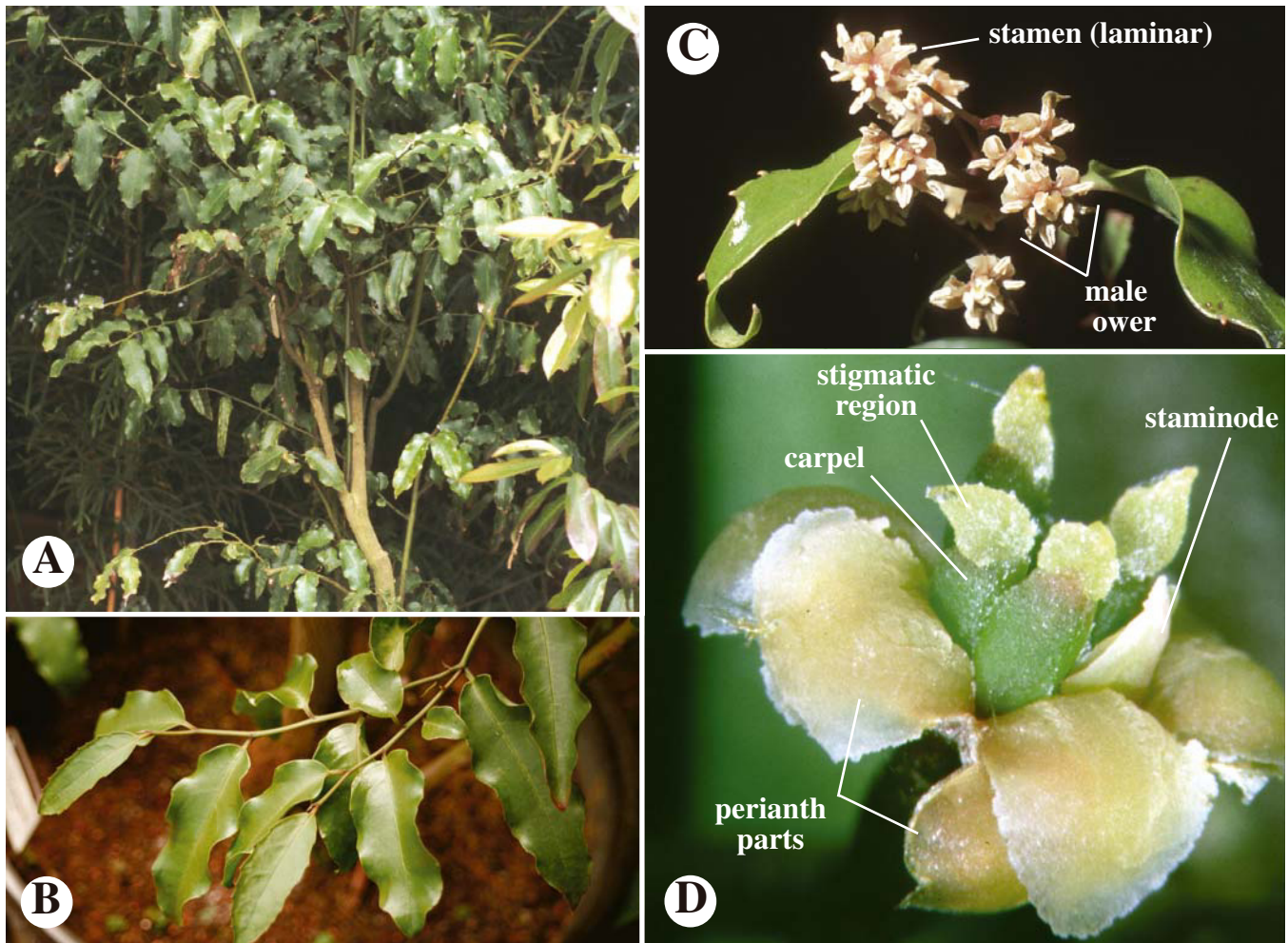


FIGURE 7.3 AMBORELLALES. Amborellaceae, *Amborella trichopoda*. **A.** Whole plant, in cultivation. **B.** Close-up of leaves. **C.** Male flowers, showing laminar stamens. **D.** Female flower close-up, showing spiral perianth and apocarpous gynoecium. (A and C, courtesy of Stephen McCabe; B, courtesy of the Arboretum at University of California-Santa Cruz; D, courtesy of Sandra K. Floyd.)

economic uses, other than being a cultivar sought because of its distinctive, basal position in the angiosperms. See Thien et al. (2003) for a study of the population structure and floral biology of *Amborella*.

The Amborellaceae are distinctive in being **vessel-less**, **evergreen** shrubs with **unisexual** flowers having an **undifferentiated**, **spiral** perianth, **numerous**, **laminar** stamens, and an **apocarpous**, **apically-open** gynoecium, with **1-ovuled** carpels.

Male flowers: P 5-8 A ∞.

Female flowers: P 5-8 G 5-6, superior.

NYMPHAEALES

This order consists of two families, Nymphaeaceae and Cabombaceae, which are sometimes treated together (e.g., as subfamilies) in a broader Nymphaeaceae, s.l. See Les et al. (1999) for recent information on the phylogeny and classification of the order.

Nymphaeaceae Water-Lily family (*Nymphae*, a water nymph). 6 genera / 60 species. (Figure 7.4)

The Nymphaeaceae consist of aquatic, annual or perennial herbs, with a milky latex often present. The underground **stems** are rhizomatous or tuberous. The stem vasculature is an atactostele or eustele. The **leaves** are simple, often peltate, stipulate or extipulate, floating, spiral, usually orbicular in shape. The **inflorescence** consists of a solitary, floating or emergent flower. **Flowers** are bisexual, actinomorphic, and hypogynous or epigynous, with long peduncles arising from the underground stem. The **perianth** is usually differentiated into calyx and corolla, the parts spirally arranged. The **calyx** consists of 4-6 [up to 14], aposepalous sepals. The **corolla** consists of 8 many [0], apopetalous petals, the inner of which grade into laminar stamens. **Stamens** are numerous, spiral, apostemonous; the filaments are laminar to the outside, grading morphologically into petals, to terete toward the flower center; anthers are longitudinal in dehiscence, dithecal, with thecae and connective often extending beyond the anther. The **gynoecium** is syncarpous, with a superior or inferior ovary, and 3 many carpels; placentation is lamellate or parietal; ovules are anatropous, bitegmic, and numerous per carpel. The **fruit** is a berry.

The Nymphaeaceae has in the past included the subfamilies Cambomboideae and Nelumboideae, but these are treated here as separate families: Cabombaceae and Nelumboaceae (the latter distantly grouped within the eudicots; see Chapter 8). Members of the Nymphaeaceae are distributed worldwide. Economic uses include species with edible rhizomes and seeds; many species are used as ornamental

cultivars, especially *Nuphar* (cow-lily), *Nymphaea* (water-lily), and *Victoria* (giant water-lily), the last having huge, peltate, floating leaves with upturned, ridged margins. See Scheinder and Williamson (1993) for more information on the family.

The Nymphaeaceae are distinguished from related families in being **aquatic** herbs with **floating** leaves and **solitary**, **floating to emergent** flowers with mostly **spiral** floral parts and petals **grading into usually laminar** stamens.

K 4-6 [-14] C 8-∞ [0] A ∞ G (3-∞), superior or inferior.

Cabombaceae F anwort family (*Cabomba*, Spanish for a S. Am. aquatic plant). 2 genera (*Brasenia* and *Cabomba*) / 6 species. (Figure 7.5)

The Cabombaceae consist of aquatic herbs. The underground **stems** are rhizomatous, which give rise to elongate leafy shoots. The stem vasculature is atactostelic. The **leaves** are dimorphic, floating or submersed, extipulate, spiral, opposite, or whorled, simple and undivided or highly divided into numerous segments. The **inflorescence** consists of a solitary, emergent flower. **Flowers** are bisexual, actinomorphic, and hypogynous. The **perianth** is dichlamydeous (differentiated into calyx and corolla), the parts whorled. The **calyx** consists of 3 [2 or 4] aposepalous sepals. The **corolla** consists of 3 [2 or 4] apopetalous petals. **Stamens** are 3 or 6 (in *Cabomba*) or 12 many (in *Brasenia*); the filaments are somewhat laminar. The **gynoecium** is apocarpous, with a superior ovary, and 2-18 [1] carpels; placentation is parietal; ovules are anatropous, bitegmic, and 2-3 [1] per carpel; styles are terminal or decurrent along the carpel. The **fruit** unit is a coriaceous follicle.

The Cabombaceae are distributed in tropical to temperate areas. *Cabomba* is found in the tropical Americas, whereas the monotypic *Brasenia* (*B. schreberi*) is distributed in tropical to temperate regions of the Americas, Africa, and Australasia. The Cabombaceae are sometimes treated as a subfamily (Cabomboideae) of the Nymphaeaceae, being different from the latter in having a trimerous [2 or 4] number of non spirally arranged sepals and petals. See Williamson and Schneider (1993) for more information on the family.

The Cabombaceae are distinguished in being **aquatic herbs** with **atactostelic** stems (resembling those of monocots); dimorphic, floating or submersed, **undivided or highly divided** leaves; a perianth with **3 [2,4] sepals and petals**, and an **apocarpous** gynoecium.

K 3 [2,4] C 3 [2,4] A 3, 6 or 12-∞ G 2-18 [1], superior.

AUSTROBAILEYALES

The Austrobaileyales, sensu APG II (2003) consist of four families (Table 7.1). Only the Illiciaceae are described here.

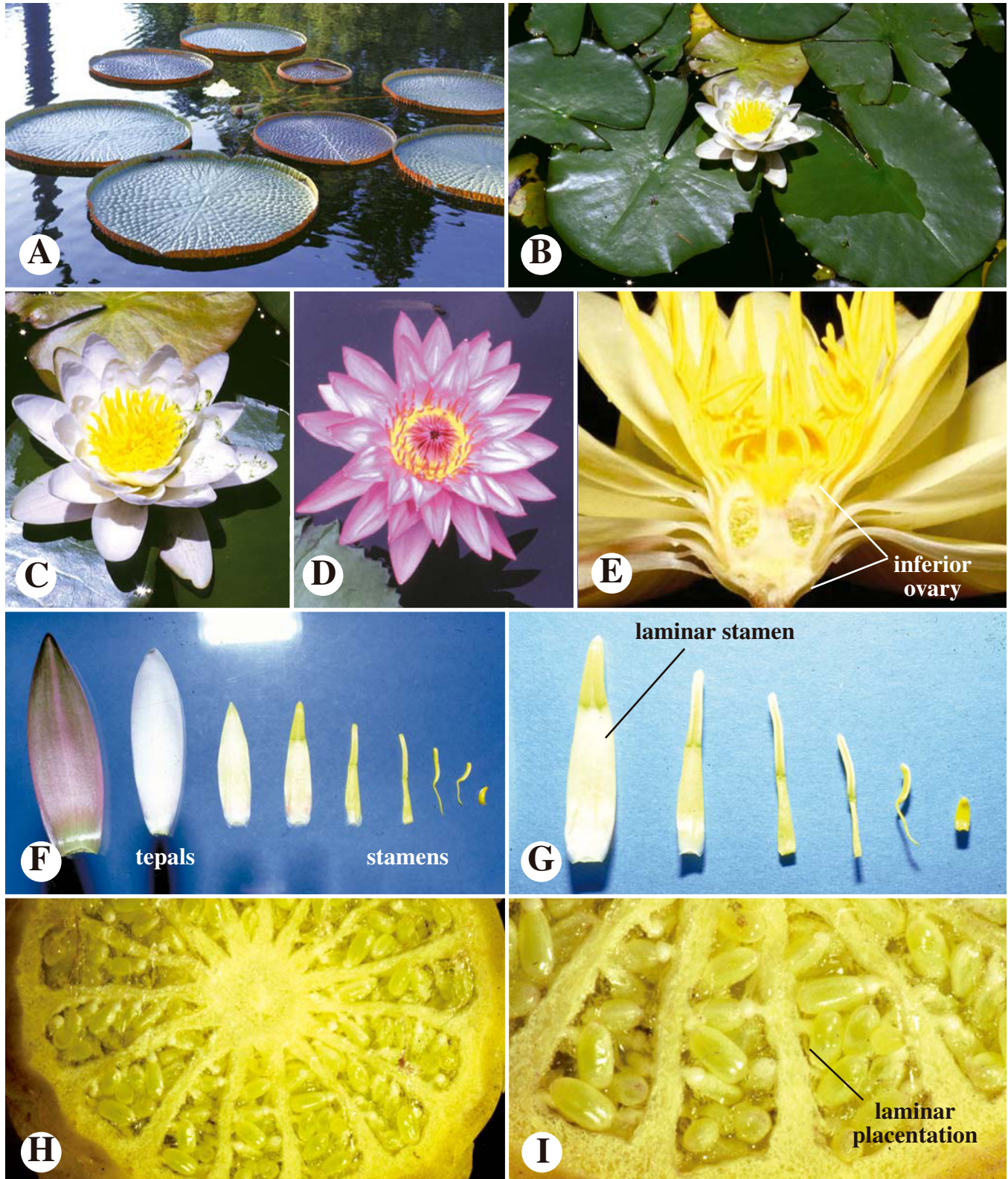


FIGURE 7.4 NYMPHAEALES. Nymphaeaceae. **A.** *Victoria amazonica*, with large, floating leaves having upturned, rimlike margins. **B–I.** *Nymphaea* spp. **B.** Whole plant, showing floating leaves and solitary flower. **C, D.** Close-up of flower. Note numerous, spiral perianth parts and stamens. **E.** Flower in longitudinal section, showing perianth series, inferior ovary, and numerous stamens. **F.** Removed floral parts (outer to inner = left to right), showing gradation from sepal-like structures (left) to petal-like structures (second and third from left) to stamens (right). **G.** Close-up of stamens, showing gradation from outer, laminar stamens (left) to filamentous stamens (middle) to sub-sessile stamens (right). **H.** Ovary cross-section. **I.** Close-up of ovary cross-section, showing laminar placentation, i.e., attachment of ovules to inner surface of septae.



FIGURE 7.5 NYMPHAEALES. Cabombaceae. *Brasenia schreberi*. **A.** Floating leaves and emergent flowers. **B,C.** Flower (protogynous) close-up. **B.** Pistils mature. **C.** Stamens mature. All photos courtesy of Jeffrey M. Osborn and Mackenzie L. Taylor.

Illiciaceae Star-Anise family (L., for alluring, enticing). 1 genus/42 species. (Figure 7.6)

The Illiciaceae consist of trees and shrubs with aromatic (ethereal) oil cells. The **leaves** are simple, spiral (often appearing whorled), pellucid-punctate, exstipulate, evergreen, and glabrous. The **inflorescence** is an axillary or supra-axillary, solitary flower or group of 2 or 3 flowers. The **flowers** are small, bisexual, actinomorphic, and hypogynous. The **perianth** consists of numerous (7-33), distinct tepals, typically spirally arranged, the outer sepal-like parts grading into inner petal-like parts, which grade into central anther-like parts. The **stamens** are few to numerous (4 ca.50), in one or more spiral series, and apostemonous; filaments are short and thick. **Anthers** are longitudinal in dehiscence, with an extended connective. The **gynoecium** is apocarpous, with numerous (5-21), superior, unilocular carpels in a single whorl. The **style** is open.

Placentation is ventrally sub-basal; **ovules** are anatropous, 1 per carpel. The **fruit** is an aggregate of follicles (follicetum). The **seeds** are endospermous, the endosperm oil-rich. Flowers are beetle-pollinated.

The Illiciaceae have distributions in S.E. Asia and S.E. U.S. to the Caribbean. Economic importance includes *Illicium anisatum*, Japanese anise, used to kill fish and used medicinally and in religious rites, and *Illicium verum*, star anise, used as a spice, e.g., in liqueurs (Figure 7.7).

The Illiciaceae are distinctive in being **evergreen** trees or shrubs having **aromatic oil cells**, with glabrous, spiral, **pellucid-punctate**, **exstipulate** leaves, the flowers with **numerous, spiral** tepals (outer sepal-like, inner petal-like), few-numerous stamens, and few-numerous, **one-seeded, apocarpous** pistils **in a single whorl**, the fruit a **follicetum**. $P \infty [7-33]$ $A \infty [4-50]$ $G \infty [5-21]$, superior.

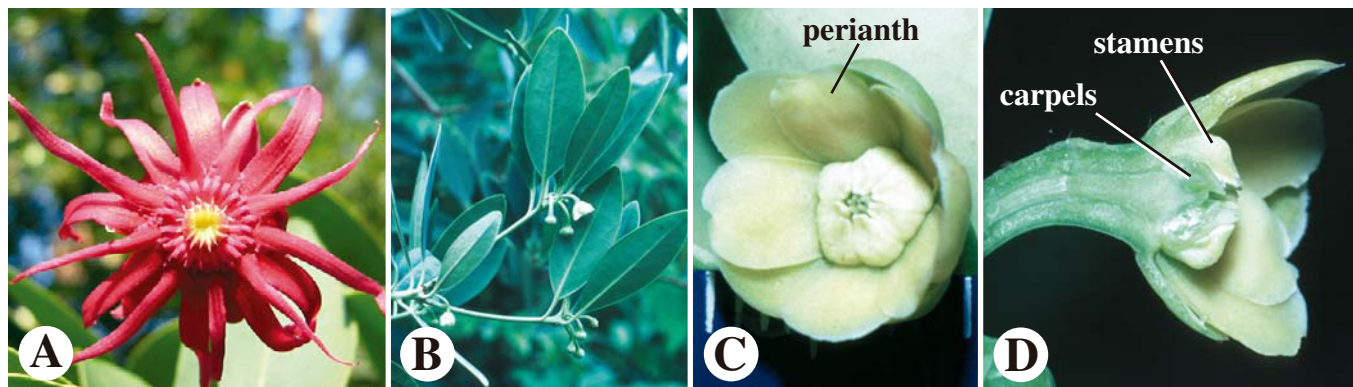


FIGURE 7.6 AUSTROBAILEYALES. Illiciaceae. **A.** *Illicium oridanum*, flower face view. (Photo courtesy of Jack Scheper, Floridata.com LC) **B-D.** *Illicium parviflorum*. **B.** Branch, showing simple, glabrous, evergreen leaves and one- to few-flowered inflorescences. **C.** Flower, face-view, showing grading perianth. **D.** Flower, longitudinal-section, showing encircling stamens and apocarpous gynoecium.



FIGURE 7.7 AUSTROBAILEYALES. Illiciaceae. *Illicium verum*. Fruits of star anise.

MAGNOLIIDS

This group, recognized by APG II (2003), contains the four orders Laurales, Magnoliales, Canellales, and Piperales. See Kim et al. (2004).

LAURALES

The Laurales, sensu APG II (2003), contain seven families (Table 7.1). Only the Lauraceae are described here. See Endress and Igersheim (1997), Renner (1999), and Renner and Chanderbali (2000) for further information.

Lauraceae Laurel family (*L. laurus*, laurel or bay). 45 genera / 2200 species. (Figure 7.8)

The Lauraceae consist of mostly trees or shrubs (except *Cassytha*, a parasitic vine) with aromatic oil glands. The **leaves** are evergreen, simple, exstipulate, spiral, rarely whorled or opposite, undivided or lobed, pinnate-netted, usually punctate. The **inflorescence** is an axillary cyme or raceme, rarely a solitary flower. **Flowers** are small, bisexual or unisexual, actinomorphic, perigynous or epiperigynous, the subtending receptacle often enlarging in fruit. The **perianth** is 1 3-whorled, usu. 3+3 [6, 2+2, or 3+3+3], apotepalous, hypanthium present. **Stamens** are 3 12 or more, with staminodes often present as an inner whorl; filaments often have a pair of basal, nectar-bearing appendages; anthers are valvular, with 2 4 [1] valves per anther opening from the base, introrse or extrorse in dehiscence, dithecal [monothechal], tetrasporangiate [bi- or monosporangiate]. The **gynoecium** consists of a single superior, rarely inferior, ovary, unicarpellous or syncarpous, consisting of 1 [up to 3] carpel, 1 locule, 1 terminal style, and 1 3 stigmas; placentation

is apical; ovules are anatropous, bitegmic, 1 per carpel. The **fruit** is a berry, drupe, or is dry and indehiscent, often with an enlarged receptacle and accrescent calyx; seeds are exalbuminous.

The Lauraceae are distributed in tropical to warm temperate regions, esp. S.E. Asia and tropical America. Economic importance includes several timber trees, spice and other flavoring plants (including the bark of *Cinnamomum cassia*, cassia, and *C. zeylanicum*, cinnamon; oils derived from *C. camphora*, camphor; and the leaves of *Laurus nobilis*, laurel or bay), and food plants, especially avocado, *Persea americana*. See Rohwer (1993) for more information on the family.

The Lauraceae are distinguished in being **perennial trees or shrubs** [rarely vines] with **aromatic oil glands**, **evergreen** leaves, an **undifferentiated** perianth, **valvular** anther dehiscence, and a **single, superior** ovary having one ovule per carpel with **apical** placentation, seeds lacking endosperm.

P 3+3 [6, 2+2, or 3+3+3] **A** 3-12+ **G** 1 [- (3)], superior, rarely inferior, hypanthium present.

MAGNOLIALES

The Magnoliales, sensu APG II (2003), contain six families (Table 7.1), of which two are described here. Notable among the others are the Myristicaceae, containing *Myristica fragrans*, from which are derived nutmeg and mace (from the seeds and aril, respectively). See Sauquet et al. (2003) and general references for angiosperm phylogeny.

Annonaceae Custard-Apple family (*Anona*, a Haitian name). 112 genera / 2150 species. (Figure 7.9)

The Annonaceae consist of trees, shrubs, or woody vines (lianas). The **leaves** are usually distichous, simple, and exstipulate. The **inflorescence** is a solitary flower or cyme. The **flowers** are bisexual [unisexual] and hypogynous. The **perianth** is triseriate, usu. 3+3+3, hypanthium absent. The **stamens** are numerous, usually spiral, apostemonous, rarely basally connate. **Anthers** are longitudinally dehiscent. The **pollen** is released as monads, tetrads, or polyads. The **gynoecium** consists of numerous carpels with superior ovaries, either apocarpous with usually spiral carpels, or rarely syncarpous with whorled carpels. **Placentation** is variable; **ovules** are anatropous or campylotropous, bitegmic or rarely tritegmic, 1-numerous per carpel. The **fruit** is an aggregate of berries or dry and indehiscent units, or a syncarp in which the unit berries fuse to a fleshy receptacular axis. The **seeds** are endospermous, the endosperm ruminant (having an uneven, coarsely wrinkled texture), oily, sometimes starchy. Resin canals and a septate pith are usually present.

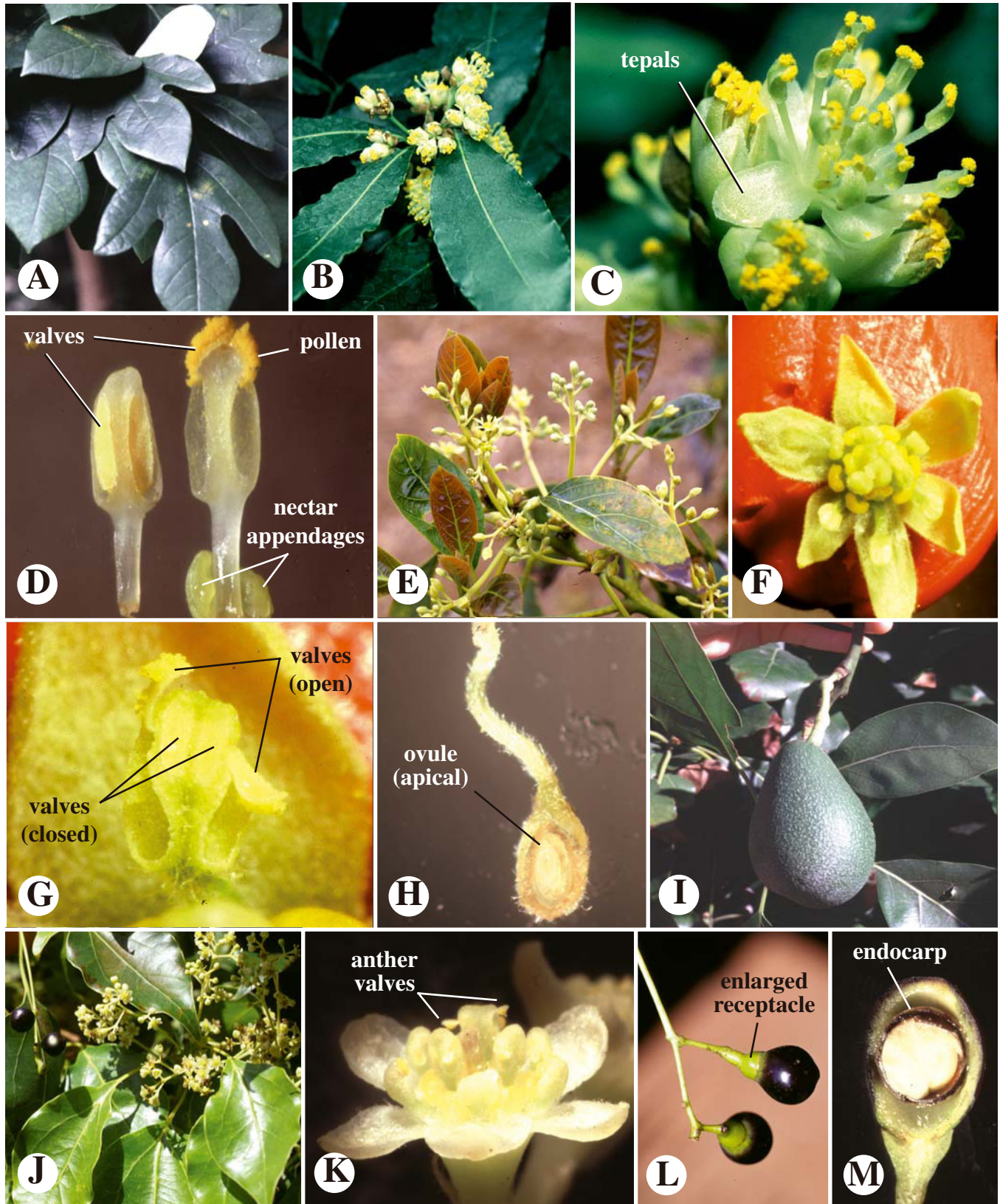


FIGURE 7.8 LAURALES. Lauraceae. **A.** *Sassafras albidum*, sassafras. **B–D.** *Laurus nobilis*, laurel. **B.** Branch, in flower. **C.** Whole flower, showing tepals and multiple whorls of stamens. **D.** Anther, showing valvular dehiscence from base of anther, one valve per theca. **E–L.** *Persea americana*, avocado. **E.** Shoot and inflorescence. **F.** Flower. **G.** Anther, showing valvular dehiscence from base of anther, two valves per theca. **H.** Pistil, ovary in longitudinal section, showing single, apical ovule. **I.** Mature fruit, a drupe. **J–M.** *Cinnamomum camphora*, camphor tree. **J.** Shoot, with inflorescence. **K.** Flower. **L.** Drupes, with enlarged receptacle at base. **M.** Drupe longitudinal section, showing hard endocarp.

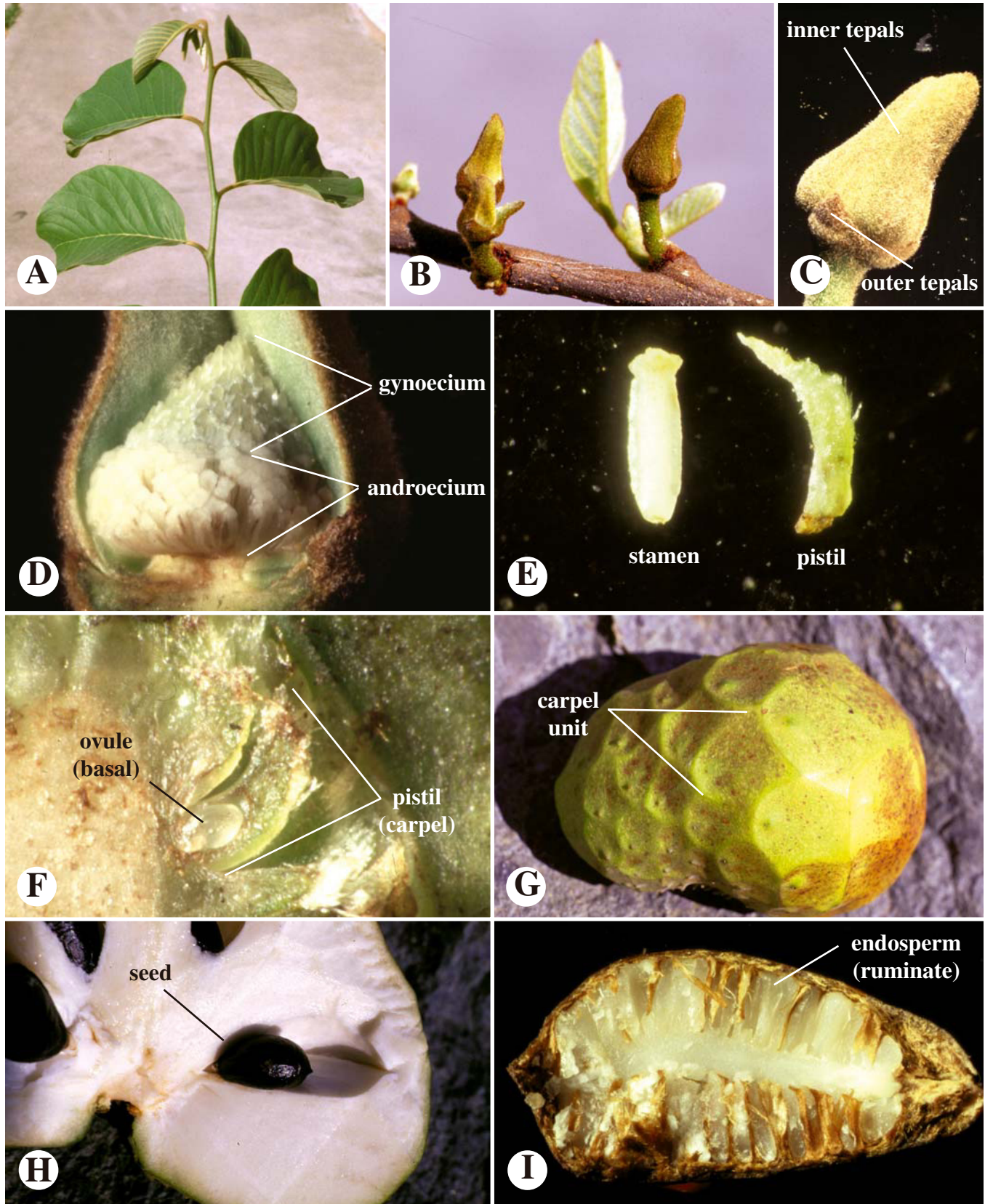


FIGURE 7.9 MAGNOLIALES. Annonaceae. *Annona cherimola*. **A.** Shoot, showing distichous leaves. **B.** Close-up of shoot, with young leaves and flowers. **C.** Flower close-up, showing undifferentiated perianth. **D.** Flower, perianth removed. Note basal androecium of numerous stamens and apical, apocarpous gynoecium of numerous pistils. **E.** Close-up of removed stamen and pistil. **F.** Pistil, longitudinal section, showing single ovule with basal placentation. **G.** Fruit, a syncarp of laterally fused carpel units. **H.** Fruit in section, showing dark seeds and surrounding fleshy tissue. **I.** Seed in longitudinal section, showing characteristic ruminant endosperm.

The Annonaceae have a mainly tropical distribution. Economic importance includes *Annona* spp. (e.g., *Annona cherimola*, cherimoya/custard-apple) grown for their edible fruits, species used for scent or timber, and some cultivated ornamentals, e.g., *Polyalthia*. See Doyle and le Thomas (1994, 1996) for analyses with emphasis on pollen evolution.

The Annonaceae are distinctive in being trees, shrubs, or woody vines with **simple**, usually **distichous** leaves, a **trimerous** perianth, **numerous, usually spiral** stamens and pistils (apocarpous or syncarpous), and seeds with **ruminate endosperm**.

P 3+3+3 **A** ∞ **G** ∞, superior.

Magnoliaceae Magnolia family (after Pierre *Magnol* of Montpellier, 1638 1715). 7 genera/200 species. (Figure 7.10)

The Magnoliaceae consist of species of trees or shrubs. The **leaves** are simple, spiral, pinnate-netted, and stipulate, with caducous stipules enclosing the buds. The **inflorescence** is a terminal solitary flower. **Flowers** are large, bisexual (rarely unisexual), actinomorphic, hypogynous; the receptacle grows into an elongate axis (called a torus or androgynophore), which bears the androecium and gynoecium. The **perianth** is multiwhorled or spiral, and apotepalous. **Stamens** are numerous, spiral, apostemonous; filaments are thickened to laminar; anthers are longitudinal in dehiscence (variable in direction), tetrasporangiate, dithecal, the paired sporangia sometimes appearing embedded, with a connective often extending beyond thecae. The **gynoecium** is apocarpous, with [2-] numerous, superior, spirally arranged ovaries/carpels, each unilocular, with one terminal style, and one stigma; placentation is marginal; ovules are anatropous and bitegmic, 2 numerous per carpel. The **fruit** is an aggregate of follicles, berries, or samaras; seeds are endospermous, rich in oils and protein with a sarcotesta (fleshy seed coat resembling an aril) usually present.

The Magnoliaceae are distributed in tropical to warm temperate regions, especially in the northern hemisphere. Economic importance includes ornamental cultivars and some important timber trees, e.g., *Liriodendron*, *Magnolia*, and *Michelia*. See Kim et al. (2001) for a more detailed treatment of the family.

The Magnoliaceae are distinguished in being trees and shrubs with **stipulate** leaves, **solitary** flowers, a usually **undifferentiated petaloid** perianth with **numerous** tepals, and **numerous, spiral** stamens and an **apocarpous gynoecium** of **numerous, spiral** pistils born on **elongate receptacular axis (torus or androgynophore)**, the fruit an **aggregate** of follicles, berries, or samaras, seeds usually with a **sarcotesta**.

P ∞ **A** ∞ **G** ∞ [2-∞], superior.

PIPERALES

The Piperales, sensu APG II (2003), contain five families (Table 7.1), of which three are described here. Notable among the others are the achlorphyllous, parasitic Hydnoraceae. See Nickrent et al. (2002) and general angiosperm phylogeny references.

Aristolochiaceae Birthwort family (Gr. *aristos*, best + *lochia*, childbirth, from resemblance of a species of *Aristolochia* to the correct fetal position). 7 genera/410 species. (Figure 7.11)

The Aristolochiaceae consist of shrubs, vines, or rhizomatous herbs, usually climbing. The **leaves** are simple, petiolate, spiral, and usually exstipulate. The **inflorescence** consists of a solitary flower or of terminal or lateral racemes or cymes. **Flowers** are bisexual, actinomorphic or zygomorphic (in *Aristolochia*), generally epigynous. The **perianth** consists of a three-lobed, synsepalous, petaloid calyx. The **corolla** is absent or reduced to three minute petal-like structures (in *Asarum*). **Stamens** are 6 ca.40, free or fused with the style forming a gynostemium (also called a column or androgynophore); filaments, when present, are short and thick; anthers are longitudinal and extrorse [introrse] in dehiscence, dithecal. The **gynoecium** is syncarpous, with a mostly inferior [half-inferior] ovary, with 4-6 carpels, 4-6 locules, one style, and 4-6 stigmas; placentation is axile; ovules are usually anatropous, bitegmic, many per carpel. The **fruit** is usually a capsule, less commonly a schizocarp of follicles or indehiscent; seeds are oily to starchy endospermous.

Members of the family have distributions in tropical and warm temperate regions, esp. in the Americas. Economic importance includes cultivated ornamentals, e.g., *Aristolochia* (Dutchman's-pipe, pelican flower, birthwort) and *Asarum* (wild ginger), with some species used medicinally (*Aristolochia*, *Thottea*), some to cure snakebites. See Kelly and Gonzalez (2003) for a recent analysis of the family.

The Aristolochiaceae are distinguished in being usually **climbing** plants, having an **enlarged, petaloid** calyx, an **absent to reduced** corolla, often adnate stamens (forming a **gynostemium**), and an **inferior to half-inferior, 4-6-carpeled and loculed** ovary.

K (3) **C** 0 [3] **A** 6-∞, usu. adnate to style **G** (4-6), inferior (half-inferior).

Piperaceae Pepper family (*piper*, Indian name for pepper). 14 genera/1940 species. (Figure 7.12)

The Piperaceae consist of herbs, shrubs, vines, or trees. The **leaves** are spiral, simple, stipulate (the stipules adnate to the petiole) or exstipulate. The **inflorescence** is a spadix. The **flowers** are very small, bisexual or unisexual, actinomorphic,

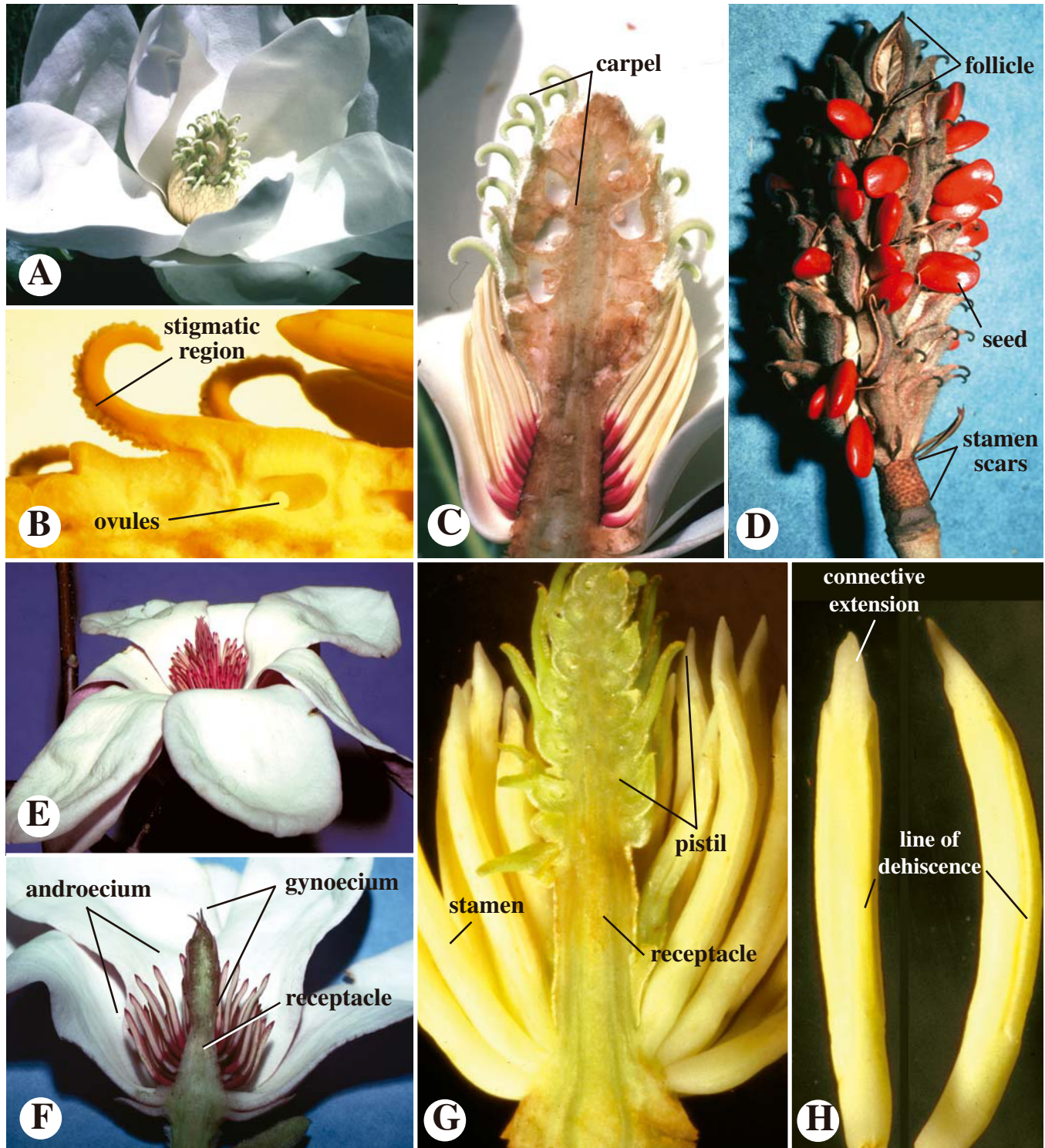


FIGURE 7.10 MAGNOLIALES. Magnoliaceae. **A–D.** *Magnolia grandiflora*. **A.** Whole flower, showing numerous tepals. **B.** Close-up of pistil. Note marginal placentation. **C.** Flower l.s., showing pistils. **D.** Fruit, an aggregate of follicles. Note seeds, having fleshy (red) sarcotesta. **E,F.** *Magnolia stellata*. **E.** Whole flower. **F.** Flower l.s. Note elongate, central receptacle (torus, androgynophore). **G,H.** *Michelia doltsopa*. **G.** Flower l.s., close-up, showing androecium (below) and receptacle bearing pistils. **H.** Stamens, adaxial (left) and side (right) views. Note lack of differentiation between filament and anther.

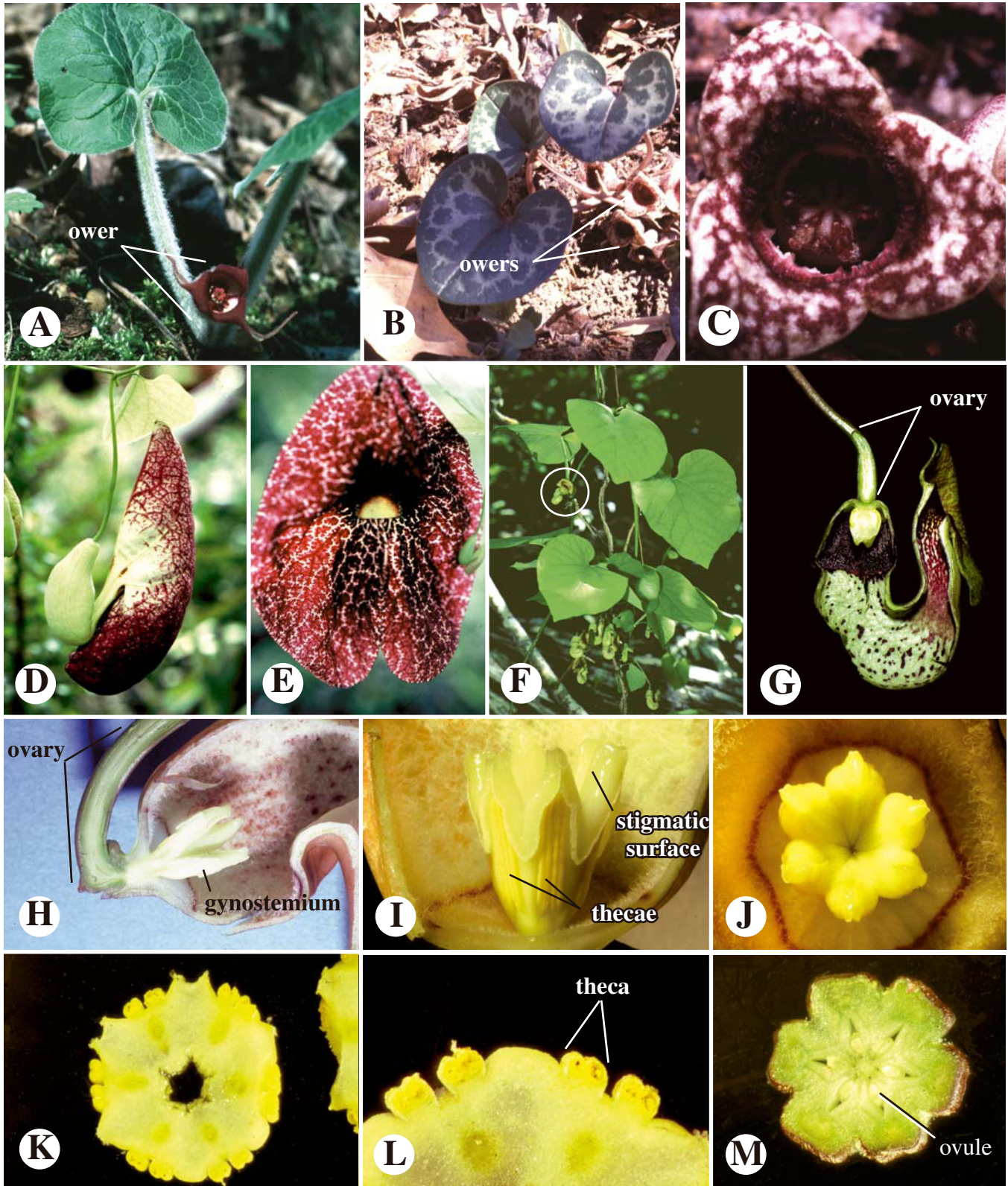


FIGURE 7.11 PIPERALES. Aristolochiaceae. **A.** *Asarum canadense*, wild ginger. **A.** Whole plant, showing ovary with trimerous calyx. **B,C.** *Hexastylis minor*. **B.** Whole plant. **C.** Close-up of ovary, showing trimerous, petaloid calyx. **D,E.** *Aristolochia elegans*. **D.** Flower bud, just prior to opening. **E.** Mature ovary, face view. **F,G.** *Aristolochia macrophylla*. **F.** Vine, with large, cordate leaves and ovary (circled). **G.** Flower longitudinal section. Note synsepalous perianth and inferior ovary. **H-M.** *Aristolochia trilobata*. **H.** Flower base longitudinal section, showing inferior ovary and gynostemium (column or androgynophore). **I.** Gynostemium close-up. Note anther thecae and stigmatic surface. **J.** Gynostemium, upper view, showing six lobes, corresponding to carpels. **K.** Gynostemium cross-section. **L.** Close-up of K, showing thecae of anthers. **M.** Ovary cross-section. Note six carpels and axile placentation.

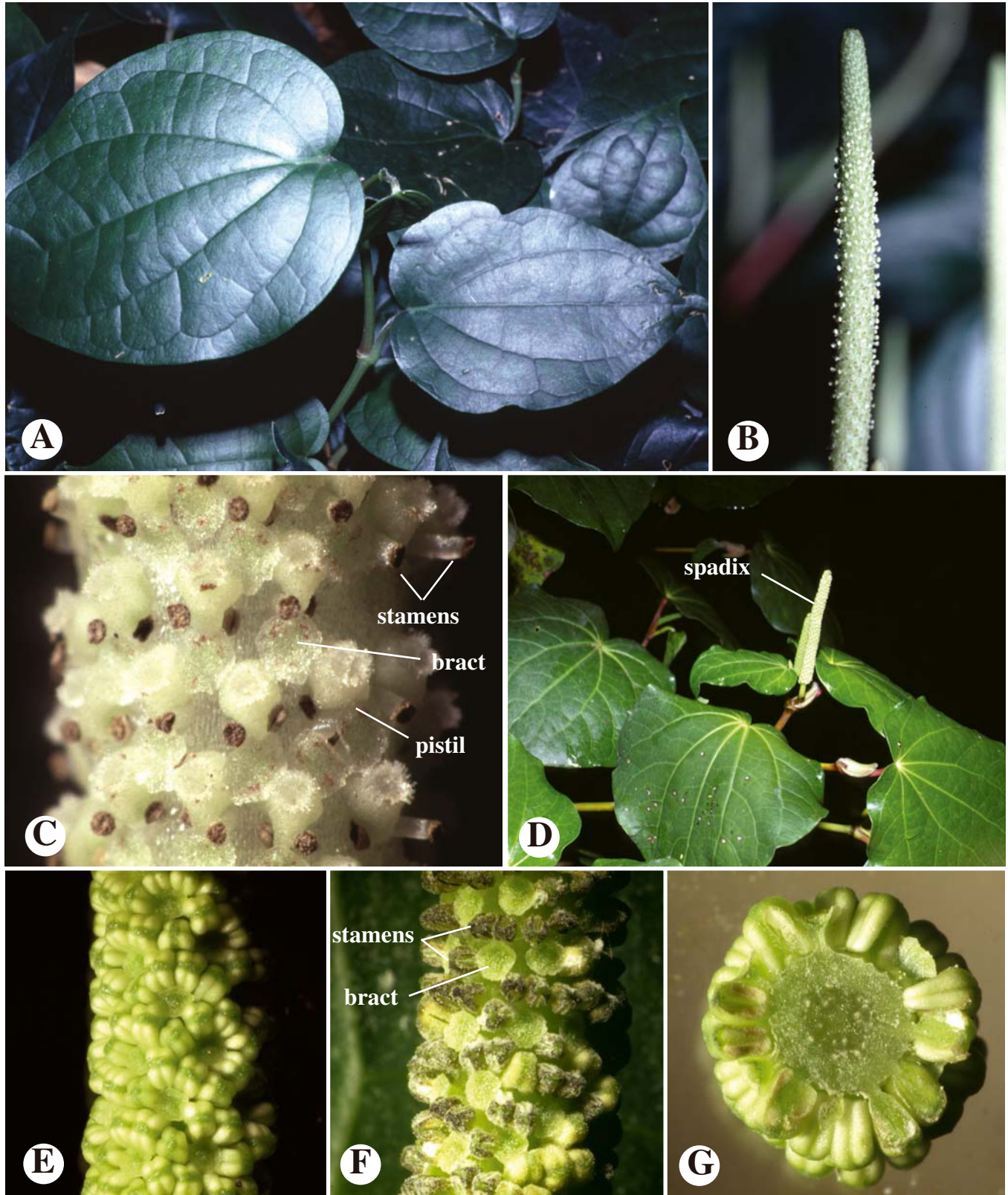


FIGURE 7.12 PIPERALES. Piperaceae. **A.** *Piper nigrum*, pepper. Vegetative morphology. **B.** *Peperomia argyreia*, watermelon peperomia. Spadix in orescence. **C.** *Peperomia* sp. Close-up of in orescence, showing numerous small, bracteate flowers. Note absence of perianth. **D–G.** *Macropiper excelsum*. **D.** Whole plant with spadix. **E.** Immature male flowers. **F.** Mature male flowers, anthers dehiscing. **G.** In orescence cross-section, showing thick, fleshy axis.

bracteate, with bracts peltate, and hypogynous. The **perianth** is absent. The **stamens** are 3+3 [1 10]. **Anthers** are longitudinally dehiscent, dithecal (sometimes appearing monothechal by fusion of thecae). The **gynoecium** consists of a single pistil with a superior ovary, having 1 or 3 4 carpels, and one locule. The **style** is absent or solitary; **stigma(s)** are 1 or 3 4, being brushlike and lateral in *Peperomia*. **Placentation** is basal; **ovules** are orthotropous, bitegmic or (in *Peperomia*) unitegmic, one per ovary. The **fruit** is a 1-seeded berry or drupe. The **seeds** have a starchy perisperm (the endosperm scanty). Plants have spherical, aromatic (ethereal) oil cells in the parenchyma and an atactostele-like vasculature (but with an outer cambium).

Members of the family have distributions in tropical regions. Economic importance includes *Piper nigrum*, the source of black and white pepper; other species are used for flavoring, medicinal plants, euphoric plants (e.g., *Piper methysticum*, kava), and cultivated ornamentals, e.g., *Peperomia* spp.

The Piperaceae are distinctive in having an *atactostelic* stem, a *spadix* with *numerous, very small, unisexual or bisexual* flowers *lacking* a perianth, the ovary solitary, 1-ovulate, the fruit a *1-seeded berry or drupe*.

P 0 **A** 3+3 [1-10] **G** 1 or (3,4), superior.

Saururaceae Lizard s-Tail family (Gr. *saur*, lizard + *our*, tail, in reference to the tail-shaped inflorescence of *Saururus cernuus*). 4 genera/6 species. (Figure 7.13)

The Saururaceae consist of perennial herbs. The **leaves** are spiral, simple, and stipulate, the stipules adnate to the petiole. The **inflorescence** is a bracteate spike or raceme, with involucrate bracts enlarged and petal like in some taxa. The **flowers** are bisexual, hypogynous. The **perianth** is absent. The **stamens** are 3, 3+3, or 4+4, apostemonous, adnate to base of the gynoecium in some taxa. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous or apically apocarpous, with a superior ovary, 3 5 carpels, and one locule. The **styles** are 3 5. **Placentation** is parietal (to marginal in *Saururus*); **ovules** are orthotropous to hemitropous, bitegmic, 1 10 per ovary. The **fruit** is an apically dehiscent capsule. The **seeds** are perispermous. Stems have 1 or 2 vascular bundle rings.

Members of the family have distributions in eastern Asia and N. America. Economic importance includes some cultivated ornamentals. See Meng et al. (2003) for a recent phylogenetic analysis of the family.

The Saururaceae are distinctive in being *perennial herbs* with a *bracteate spike or raceme* and with flowers *lacking* a perianth, the ovary solitary, many-ovulate, the fruit a *capsule*.

P 0 **A** 3, 3+3, or 4+4 **G** (3-5), superior.

CERATOPHYLLALES

This order, containing one family and genus (APG II 2003; Table 7.1), has been placed in different positions in various phylogenetic analyses, presumably because of long-branch attraction. Here it is placed as the sister group to the Eudicots, but some studies place it sister to the monocots.

Ceratophyllaceae Hornwort family (Gr. *cerato*, horn + *phyllum*, leaf, from the forked leaves resembling horns). 1 genus/ 2 (30, depending on treatment) species. (Figure 7.14)

The Ceratophyllaceae consist of monoecious, floating or submerged, aquatic, perennial herbs with rootlike anchoring branches. The **leaves** are exstipulate, whorled, 3 10 per node, 1 4 dichotomously divided, and marginally serrulate. The **inflorescence** consists of solitary and axillary flowers, male and female usually on alternate nodes. **Flowers** are unisexual. The **perianth** is uniseriate and consists of 8 12, basally fused, linear tepals. **Stamens** are generally numerous (5 27), spirally arranged on a flat receptacle; filaments are not clearly distinct from anthers, the thecae and connective apically two-pointed. The **gynoecium** is unicarpellous, with a superior ovary, 1 carpel, and 1 locule; placentation is marginal with a solitary anatropous or orthotropous, unitegmic ovule. The **fruit** is an achene, with a persistent, spiny style; seeds are exalbuminous.

Members of the family are worldwide in distribution. Economically, *Ceratophyllum demersum* is used as an aquarium plant and as a protective cover in fisheries. See Les (1993) for more information on the family.

The Ceratophyllaceae are distinguished from related families in being *monoecious, aquatic herbs* with *whorled, dichotomously branched, serrulate* leaves, and *solitary, unisexual* flowers.

P (8-12) **A** 5-27 **G** 1, superior.

MONOCOTYLEDONS

The monocotyledons, or monocots (also known as the Monocotyledonae or Liliidae), have long been recognized as a major and distinct group, comprising roughly 56,000 species, 22% of all angiosperms. All recent studies, including several molecular ones, agree with the notion that monocots are monophyletic (Figure 7.1). Monocots include the well-known aroids, arrowleaf, lilies, ginger, orchids, irises, palms, and grasses. Grasses are perhaps the most economically important of all plants, as they include grain crops such as rice, wheat, corn, barley, and rye.

Traditionally, monocots have been defined in part by the occurrence of floral parts in multiples of three. However, this

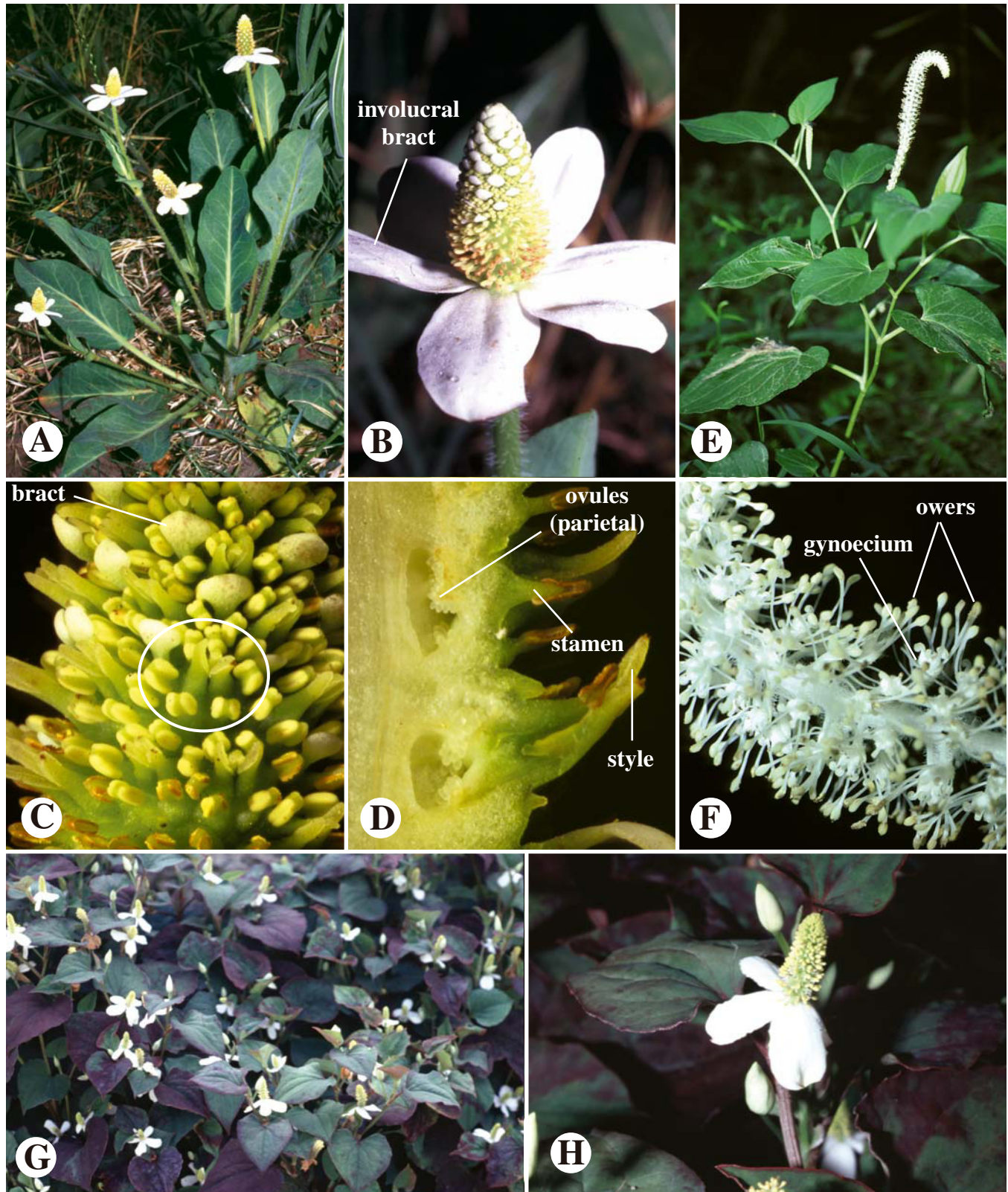


FIGURE 7.13 PIPERALES. Saururaceae. A–D. *Anemopsis californica*, yerba santa. A. Whole plant, showing basal leaves and scapose spikes. B. In orescence, close-up, with showy bracts. C. Close-up of in orescence. Note tightly clustered owers (one ower circled). D. In orescence in longitudinal section, showing partially embedded gynoecium with parietal placentation. E,F. *Saururus cernuus*, lizard s tail. E. Whole plant, showing cauline leaves and elongate raceme. F. Close-up of in orescence. G,H. *Houttuynia cordata*.



FIGURE 7.14 CERATOPHYLLALES. Ceratophyllaceae. *Ceratophyllum demersum*. A. Whole plant. B. Close-up, showing whorled, dichotomously forked leaves.

feature is now thought to represent an ancestral condition, one present or common in several basal, non-monocot lineages of flowering plants such as the Laurales, Magnoliales, and Piperales.

The phylogenetic relationships of the major groups of monocots, as summarized from recent studies, are seen in Figure 7.15. The monophyly of monocots is supported by several major morphological, anatomical, and ultrastructural apomorphies. These apomorphies will be discussed first, followed by a treatment of the major groups and exemplar families.

MONOCOT APOMORPHIES

First, all monocots have **sieve tube plastids** with **cuneate** (wedge-shaped) **proteinaceous inclusions** (Figure 7.16) of the P2 type; see Behnke (2000). This sieve tube plastid type (which can only be resolved with transmission electron microscopy) is found in all investigated monocotyledons, with some variation in form (Behnke 2000). Thus, it is likely that the cuneate, proteinaceous plastid type constitutes an apomorphy for the monocots (Figures 7.1, 7.15). The adaptive significance of this plastid type in monocots (if any) is unknown.

Second, all monocots have an **atactostele** stem vasculature, an apparent apomorphy for the group. An **atactostele** (Figure 7.17) consists of numerous discrete vascular bundles that, in cross-section, consist of two or more rings or (more commonly) appear to be rather randomly organized (but which actually have a high complexity of organization). In addition, no monocot has a true vascular cambium that produces true wood (Chapter 5); this feature is likely correlated with the evolution of the atactostele. Thus, for example,

tall palm trees have no wood, relying on the deposition and expansion of cells during primary growth for support. Some monocots (e.g., members of the Agavaceae and Asphodelaceae) do have secondary growth by means of so-called anomalous cambia, but these do not develop as a single continuous cylinder that deposit rings of secondary tissue, as in plants that produce true wood. A few eudicots (e.g., some Nelumbonaceae) have evolved an atactostele, but this was most likely a secondary innovation. Atactosteles may have evolved in response to selective pressure for adaptation to an aquatic habitat, but this is not clear.

Third, *most* monocots have **parallel leaf venation** (Figure 7.18), another apomorphy for the group. In leaves with parallel venation, the veins are either strictly parallel (as in most grasses), curved and approximately parallel, or penni-parallel (= pinnate-parallel). A penni-parallel leaf has a central midrib with secondary veins that are essentially parallel to one another (Figure 7.18). In all types of parallel venation, the ultimate veinlets connecting the major parallel veins are transverse and do not form a netlike **reticulate venation** (see Chapter 9) as found in almost all nonmonocotyledonous flowering plants. Parallel leaf venation is not a characteristic of all monocots. Numerous monocot taxa, for example some Araceae, the Dioscoreaceae (yam family), Smilacaceae (green briar family), and many others, have a reticulate leaf venation similar to that found in nonmonocots. However, the evidence supports the notion that a reticulate venation evolved in these monocot taxa secondarily, after the common evolution of parallel veins.

Fourth, all monocots have a **single cotyledon** (Figure 7.19), the feature responsible for the name *monocot*. A single cotyledon appears to be a valid apomorphy for all monocots.

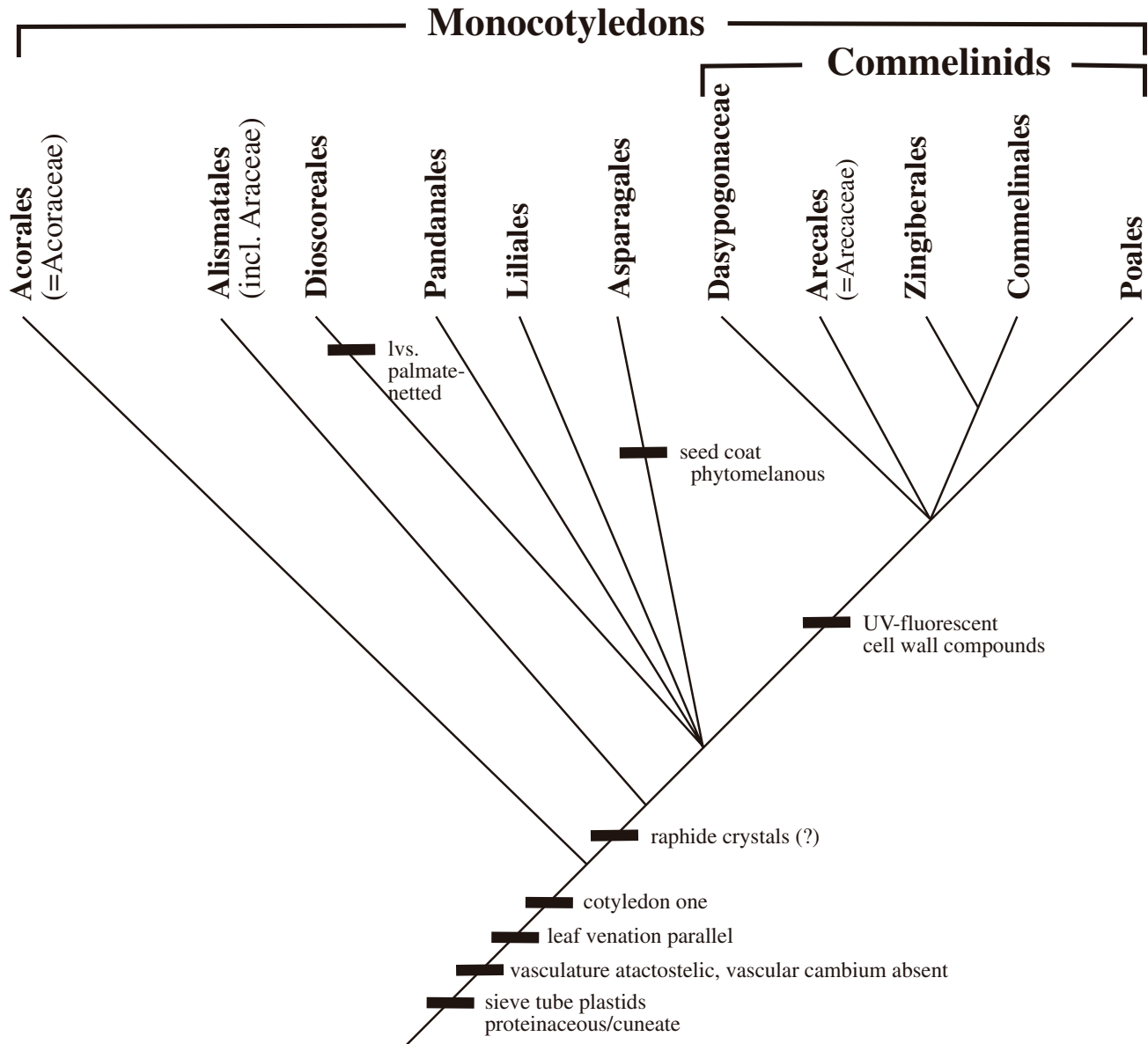


FIGURE 7.15 Major clades of the monocotyledon, modified from APG II (2003), with selected apomorphies shown.

Its adaptive significance, if any, is unknown. Some of the angiosperm lineages closely related to monocots may have a reduced second cotyledon, a possible precursor to the single cotyledon.

CLASSIFICATION OF THE MONOCOTYLEDONS

The orders of monocots and their included families are listed in Tables 7.2 and (for the Commelinid monocots) 7.3. The Acorales, which consists of the single family Acoraceae and the single genus *Acorus*, is the most basal monocot lineage as determined by numerous molecular analyses. The Alismatales, as treated here, includes the family Araceae,

which is often treated in a separate order, the Arales. Note that the Asparagales, Dioscoreales, Liliales, Pandanales, and Commelinids form an unresolved polytomy. Finally, the Commelinid monocots form a well-resolved clade that consists of the Dasygongonaceae, the Arecales (the sole member being the Arecaceae, or Palmae, the palms), the Commelinales, the Zingiberales (ginger group), and the Poales (grasses and their close relatives). See Chase et al. (2000a), Stevenson et al. (2000), and general references on angiosperm phylogeny for recent analyses of the monocots. See Rudall et al. (1995), Wilson and Morrison (2000), and Columbus et al. (2005) for collections of papers from monocot symposia.

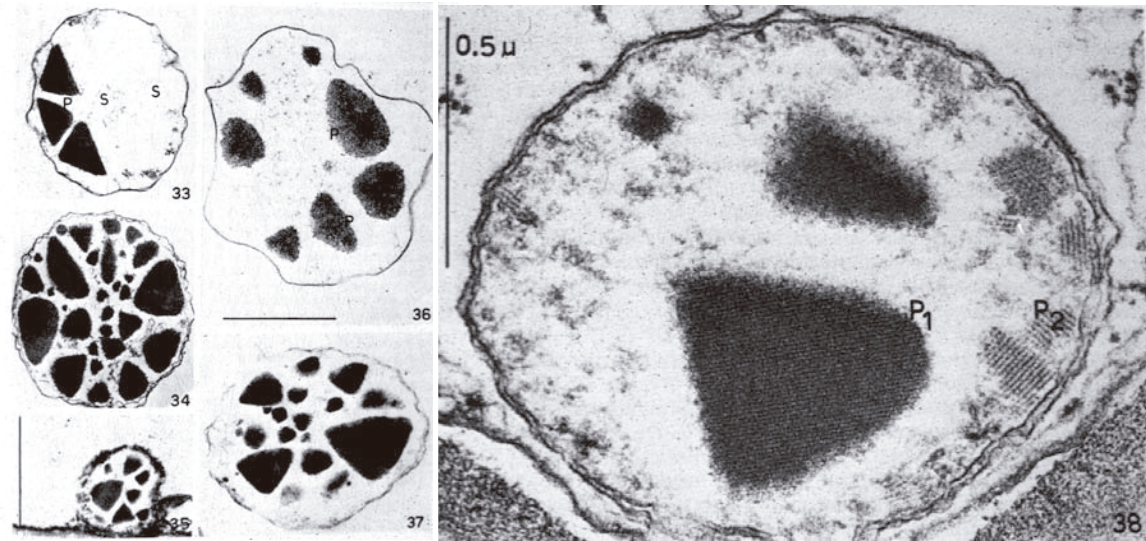


FIGURE 7.16 Sieve tube plastids with cuneate proteinaceous inclusions, an apomorphy of the monocotyledons. (Reproduced from Behnke, H.-D., 1972, by permission.)

ACORALES

The Acorales contain only one family, one genus, and 23 species. In molecular analyses, it usually comes out as the most basal lineage of the monocots (see general angiosperm phylogeny references; Duvall et al., 1993; Chase et al., 2000a; and Chen et al., 2002; however, see Stevenson et al., 2000).

Acoraceae Sweet Flag family (*Acorus*, meaning without pupil, originally in reference to a species of *Iris* used to treat cataracts). 1 genus/23 species. (Figure 7.20)

The Acoraceae consist of perennial herbs found in marshy habitats. The **stems** are rhizomatous. The **leaves** are ensiform, unifacial, distichous, sheathing, simple, undivided, exstipulate, and parallel veined, with intravaginal (axillary) squamules present. The **inflorescence** is a terminal spadix borne on a leaf-like peduncle and subtended by a long, linear spathe. The **flowers** are bisexual, actinomorphic, ebracteate, sessile, and hypogynous. The **perianth** is biseriata, of 3+3 distinct tepals. The **stamens** are biseriata, 3+3, apostemonous, with flattened filaments. **Anthers** are longitudinal and introrse in dehiscence. The **gynoecium** is syncarpous, with a superior ovary,

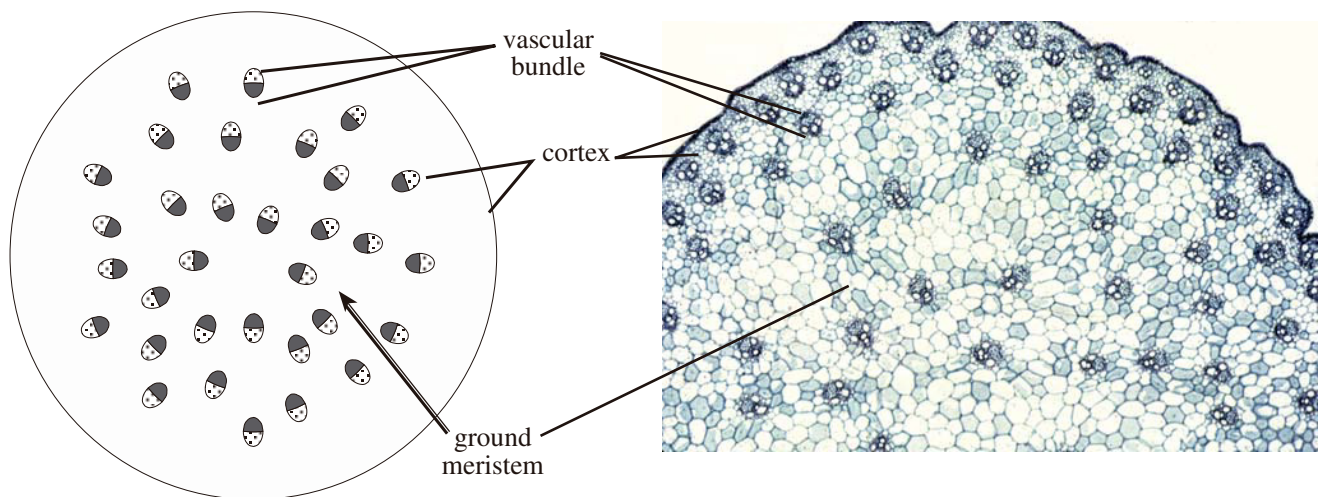


FIGURE 7.17 The atactostele, an apomorphy of the monocotyledons. Note numerous vascular bundles; at left: xylem = dark; phloem = stippled.

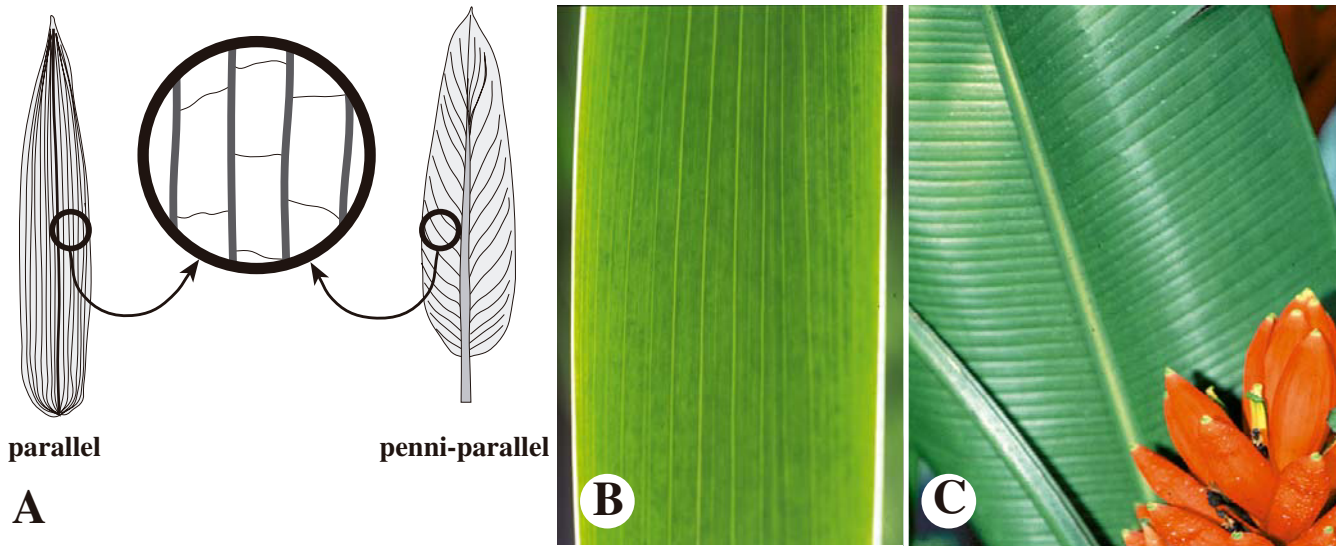


FIGURE 7.18 Parallel venation, an apomorphy of the monocotyledons. **A.** Parallel venation (left) and penni-parallel venation (right). **B.** *Leymus condensatus* (Poaceae), an example of parallel venation. **C.** *Musa coccinea* (Musaceae), an example of penni-parallel venation.

2-3 carpels, 2-3 locules, and a minute stigma. **Placentation** is apical-axile; **ovules** are ∞ per carpel, pendent. The **fruit** is a 1-5 [9] seeded berry, with a persistent perianth. The **seeds** are perispermous and endospermous. Aromatic ethereal oil cells are present. Raphide crystals are absent.

The Acoraceae are similar to the family Araceae (discussed later) in having a spadix and spathe, but is clearly separated from that family (within which it used to be placed) based on morphology and analyses of DNA sequence data. The Acoraceae differs from the Araceae in having ensiform, unifacial leaves, perispermous/endospermous seeds, and aromatic (ethereal) oil cells, and in lacking raphide crystals.

Members of the Acoraceae are distributed in the Old World and North America. Economic importance includes *Acorus calamus* used medicinally (e.g., as calamus oil), in religious rituals, as an insecticide, and as a perfume and flavoring plant (e.g., in liqueurs). See Grayum (1990) and Bogner and Mayo (1998) for more information on the family.

The Acoraceae are distinctive in being *marsh* plants with a **spadix** and **spathe** (resembling Araceae) but having **distichous, ensiform, unifacial** leaves, **perispermous and endospermous** seeds, and **ethereal oil cells**, and in **lacking raphide crystals**.

P 3+3 **A** 3+3 **G** (2-3) superior.

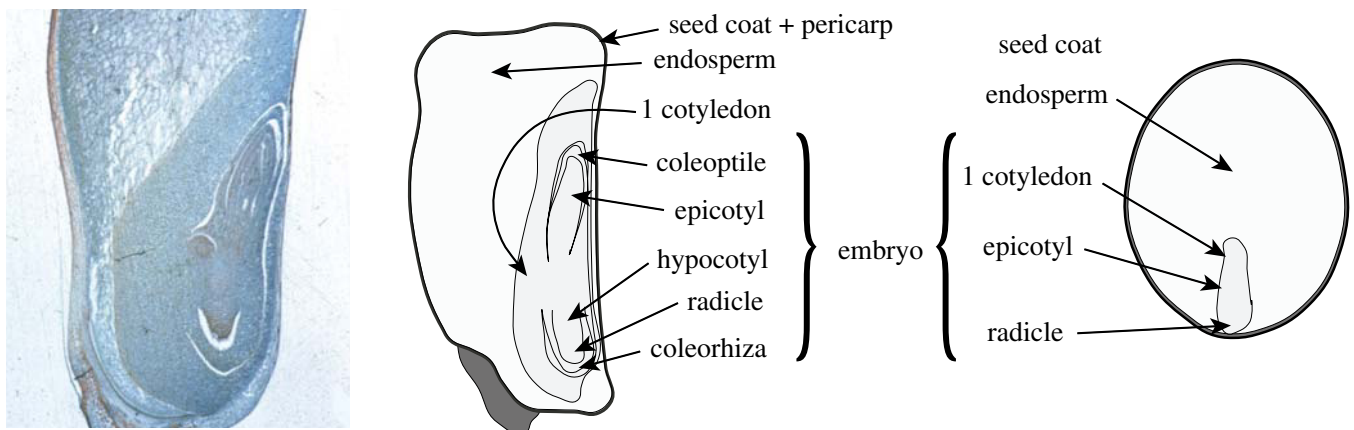


FIGURE 7.19 A single cotyledon, an apomorphy of the monocotyledons. Left, *Zea mays* (Poaceae). Right, *Xiphidium caeruleum* (Haemodoraceae).

TABLE 7.2 Orders and included families of Monocotyledons (excluding Commelinids, see Table 7.3), after APG II, 2003. Families in **bold** are described in detail. An asterisk denotes an acceptable deviation from APG II, with brackets indicating the more inclusive family recommended by APG II. A double asterisk indicates a change suggested by Angiosperm phylogeny Web site (Stevens, 2001 onwards). Including = incl.

MONOCOTYLEDONS		
<p>PETROSAVIALES** Petrosaviaceae</p> <p>ACORALES Acoraceae</p> <p>ALISMATALES Alismataceae Aponogetonaceae Araceae (incl. Lemnaceae) Butomaceae Cymodoceaceae Hydrocharitaceae Juncaginaceae Limnocharitaceae Posidoniaceae Potamogetonaceae Ruppiceae Scheuchzeriaceae To eldiaceae Zosteraceae</p> <p>ASPARAGALES Agapanthaceae* [Alliaceae] Agavaceae* [Asparagaceae] (incl. Hesperocallidaceae) Alliaceae* Amaryllidaceae* [Alliaceae] Aphyllanthaceae* [Asparagaceae]</p>	<p>ASPARAGALES (continued) Asparagaceae* Asphodelaceae* [Xanthorrhoeaceae] Asteliaceae Blandfordiaceae Boryaceae Doryanthaceae Hemerocallidaceae* [Xanthorrhoeaceae] Hyacinthaceae* [Asparagaceae] Hypoxidaceae Iridaceae Ixioliriaceae Lanariaceae Laxmanniaceae* [Asparagaceae] Orchidaceae Ruscaceae* [Asparagaceae] (incl. Convallariaceae) Tecophilaeaceae Themidaceae* [Asparagaceae] Xanthorrhoeaceae* Xeronemataceae</p>	<p>DIOSCOREALES Burmanniaceae Dioscoreaceae Nartheciaceae</p> <p>LILIALES Alstroemeriaceae Campynemaceae Colchicaceae Corsiaceae Liliaceae Luzuriagaceae Melanthiaceae Petermanniaceae** Philesiaceae Rhipogonaceae Smilacaceae</p> <p>PANDANALES Cyclanthaceae Pandanaceae Stemonaceae Triuridaceae Velloziaceae</p> <p>COMMELINIDS (see Table 7.3)</p>

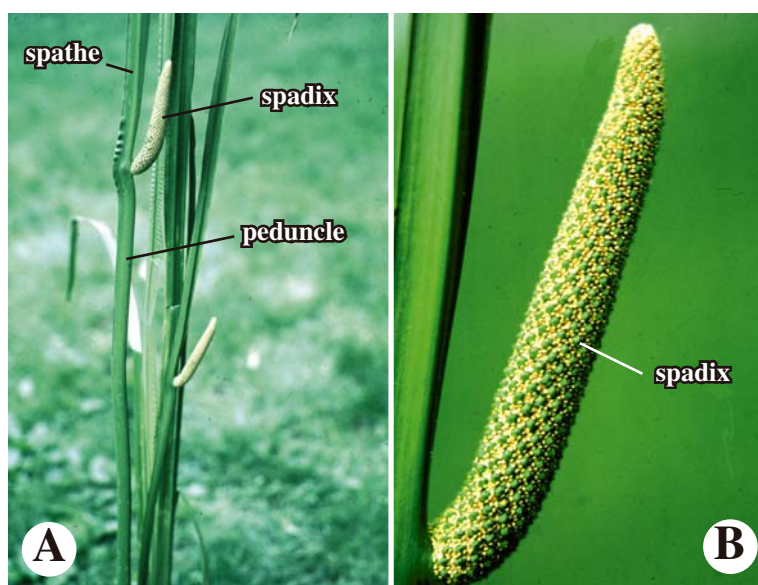


FIGURE 7.20 ACORALES. Acoraceae. *Acorus calamus*. **A.** Whole plant. **B.** Close-up of spadix in orescence.

ALISMATALES

The Alismatales, sensu APG II (2003), contain 14 families, only two of which are described here. The order has often been split into the Arales (containing only the Araceae) and the Alismatales, s.s. (largely equivalent to the Alismatidae, sensu Cronquist, 1981, and Takhtajan, 1997), but some recent molecular studies unite these two groups. Notable among the families of the order that are not described here (see Figure 7.21) are the terrestrial **Tofieldiaceae** and a number of aquatic groups, including the **Aponogetonaceae** (e.g., *Aponogeton distachyon*), **Cymodoceaceae** (several marine sea-grasses), **Hydrocharitaceae** (Figure 7.21A,B, including marine sea-grasses such as *Halophila* and *Thalassia*, fresh water aquatics such as the aquarium plants *Elodea* and *Valisneria*, and problematic weedy species of *Elodea*, *Hydrilla*, and *Lagarosiphon*), **Juncaginaceae** (Figure 7.21C,D), **Posidoniaceae** (*Posidonia* spp., marine sea-grasses), **Potamogetonaceae** (freshwater aquatics, Figure 7.21E,F), **Ruppiceae** (*Ruppia* spp., fresh to brackish water plants), and **Zosteraceae** (including deep, marine sea-grass species such as *Phyllospadix*, Figure 7.21G J). See Les and Haynes (1995) and Les et al. (1997) for more information on the order.

Trichomes located in the axils of sheathing leaves, known as **intravaginal squamules** (see Chapter 9), are common in many Alismatales (also found in the Acorales). The evolution of raphide crystals (see Chapter 10) may constitute an apomorphy for the monocots after the Acorales lineage (Figure 7.15). However, if so, they have been secondarily lost in a number of monocot lineages, including many Poales, Zingiberales, and most of the Alismatales themselves (except for the Araceae).

Araceae Arum family (Arum, a name used by Theophrastus). 104 genera/ca. 3300 species. (Figures 7.22, 7.23)

The Araceae consist of terrestrial or aquatic shrubs, vines, or herbs (the vegetative body reduced and globose to thalloid in the Lemnoideae). The **roots** are often mycorrhizal, without root hairs. The **stems** are rhizomatous, cormose, tuberous, or reduced. The **leaves** are simple, bifacial, spiral, or distichous, sometimes highly divided or fenestrate (often exhibiting heteroblasty), with parallel, penni-parallel, or netted venation. The **inflorescence** is a terminal, many-flowered spadix (with a sterile apical portion in some), usually subtended by a prominent, often colored spathe, or reduced to 1–4 flowers in a small pouch in the Lemnoideae. **Flowers** are small, bisexual or unisexual (female flowers often proximal, and the male distal on a spadix), actinomorphic, sessile, ebracteate, hypogynous, sometimes foul-smelling. The **perianth** is biseriate and 2+2 or 3+3 [4+4] or absent, apotepalous or basally

syntepalous, a hypanthium absent. **Stamens** are 4, 6, or 8 [1–12], distinct or connate, antitepalous in bisexual flowers; anthers are poricidal, longitudinal, or transverse in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 [1 ca.50] carpels, usu. as many locules as carpels, style and stigma one and short or absent; placentation is variable; ovules are usu. anatropous and bitegmic, 1–∞ per carpel. The **fruit** is typically a multiple of berries, less often dry, e.g., of utracles. **Seeds** are oily (sometimes also starchy) endospermous (rarely endosperm absent) with a sometimes fleshy seed coat. Some have cyanogenic compounds or alkaloids. Raphides are present and laticifers are common.

The Araceae are traditionally divided into several subfamilies; the traditional Lemnaceae (small, thalloid to globose aquatics with very reduced flowers; Figure 7.23E–G) are now known to be nested within the Araceae and may be classified as subfamily Lemnoideae. Members of the family have distributions in tropical and subtropical regions. Economic importance includes many taxa that are important food sources (from rootstocks, leaves, seeds, or fruits) in the tropics, e.g., *Alocasia*, *Amorphophallus*, *Colocasia esculenta* (taro), *Monstera*, *Xanthosoma sagittifolium*; indigenous medicinal, fiber (from roots), or arrow-poison plants; and numerous cultivated ornamentals, such as *Aglaonema*, *Anthurium*, *Caladium* (elephant's ear), *Dieffenbachia* (dumb cane), *Epipremnum*, *Monstera*, *Philodendron*, *Spathiphyllum*, *Syngonium*, and *Zantedeschia* (calla lily). *Amorphophallus titanum* (Figure 7.23C) is unique in having among the most massive inflorescences of any flowering plant; *Wolffia* spp. (Figure 7.23F,G) are unique in having the smallest flowers. See Grayum (1990), French et al. (1995), and Mayo et al. (1998) for more information and detailed phylogenetic studies.

The Araceae are distinguished from related families in having **bifacial** leaves with **parallel or netted** venation, usually a **spadix** of numerous, small flowers with a subtending **spathe**, **endospermous** seeds, and **raphide crystals**.

P 2+2, 3+3, (2+2), (3+3) or 0 [4+4, (4+4)] **A** 4, 6, 8 or (4, 6, 8) [1–12] **G** (3) [1–(∞)] superior.

Alismataceae Water-Plantain family (Alisma, a name used by Dioscorides for a plantain-leaved aquatic plant). 11 genera/ca. 80 species. (Figure 7.24)

The Alismataceae consist of perennial [or annual], monoecious, dioecious, or polygamous, floating to emergent, aquatic or marsh herbs. The **stem** is a corm or rhizome, the latter sometimes bearing tubers. The **leaves** are basal, simple, petiolate [rarely sessile], sheathing, spiral, and often dimorphic (the juvenile linear, adult leaves linear to ovate to triangular sagittate or hastate), parallel, or reticulate in venation. The **inflorescence** is a scapose raceme or panicle [sometimes

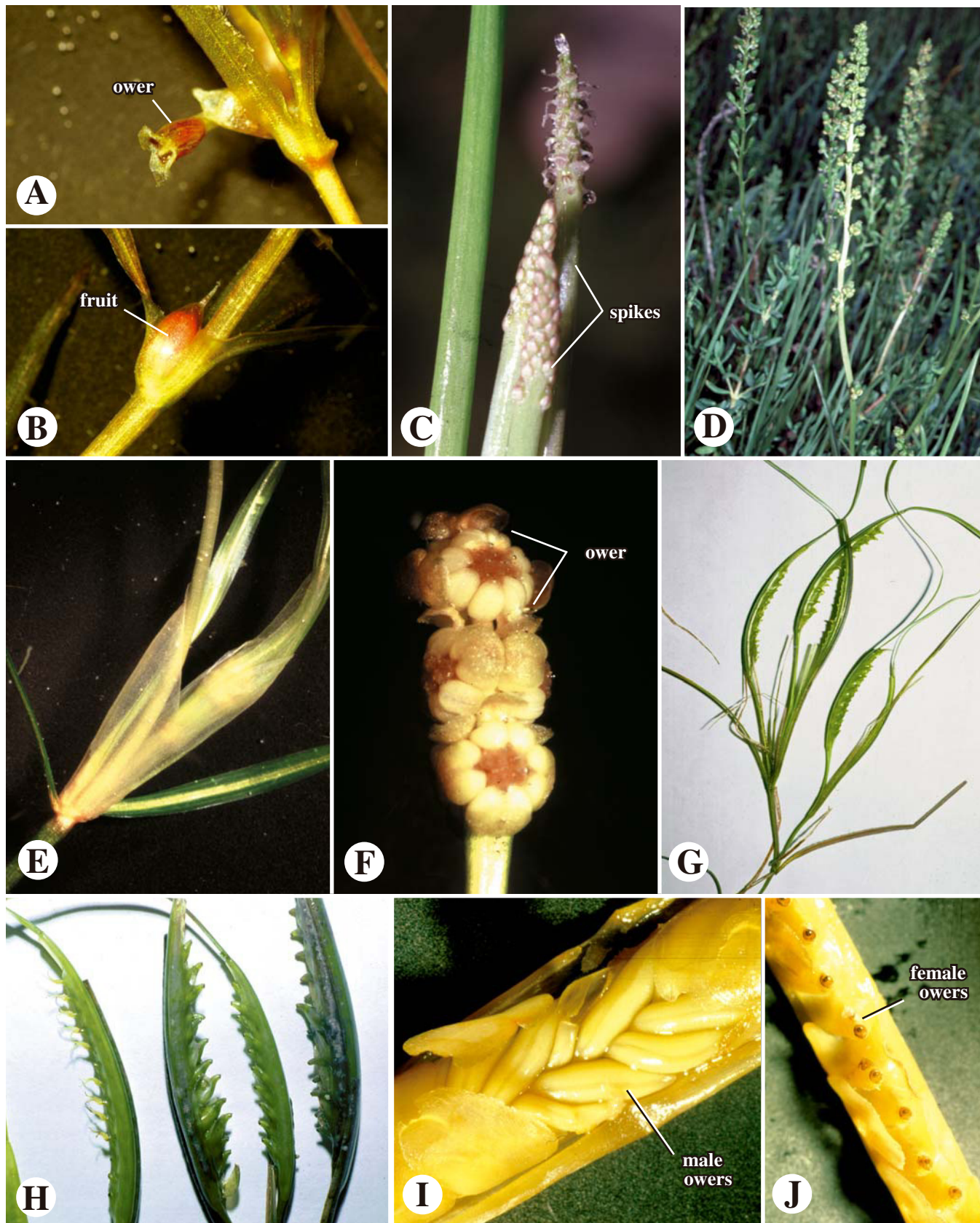


FIGURE 7.21 ALISMATALES exemplars. **A,B.** *Najas guadalupensis*, Hydrocharitaceae. **C.** *Lilaea scilloides*, Juncaginaceae. **D.** *Triglochin concinnus*, Juncaginaceae. **E,F.** *Potamogeton foliosus*, Potamogetonaceae. **G–J.** *Phyllospadix torreyi*, Zosteraceae.

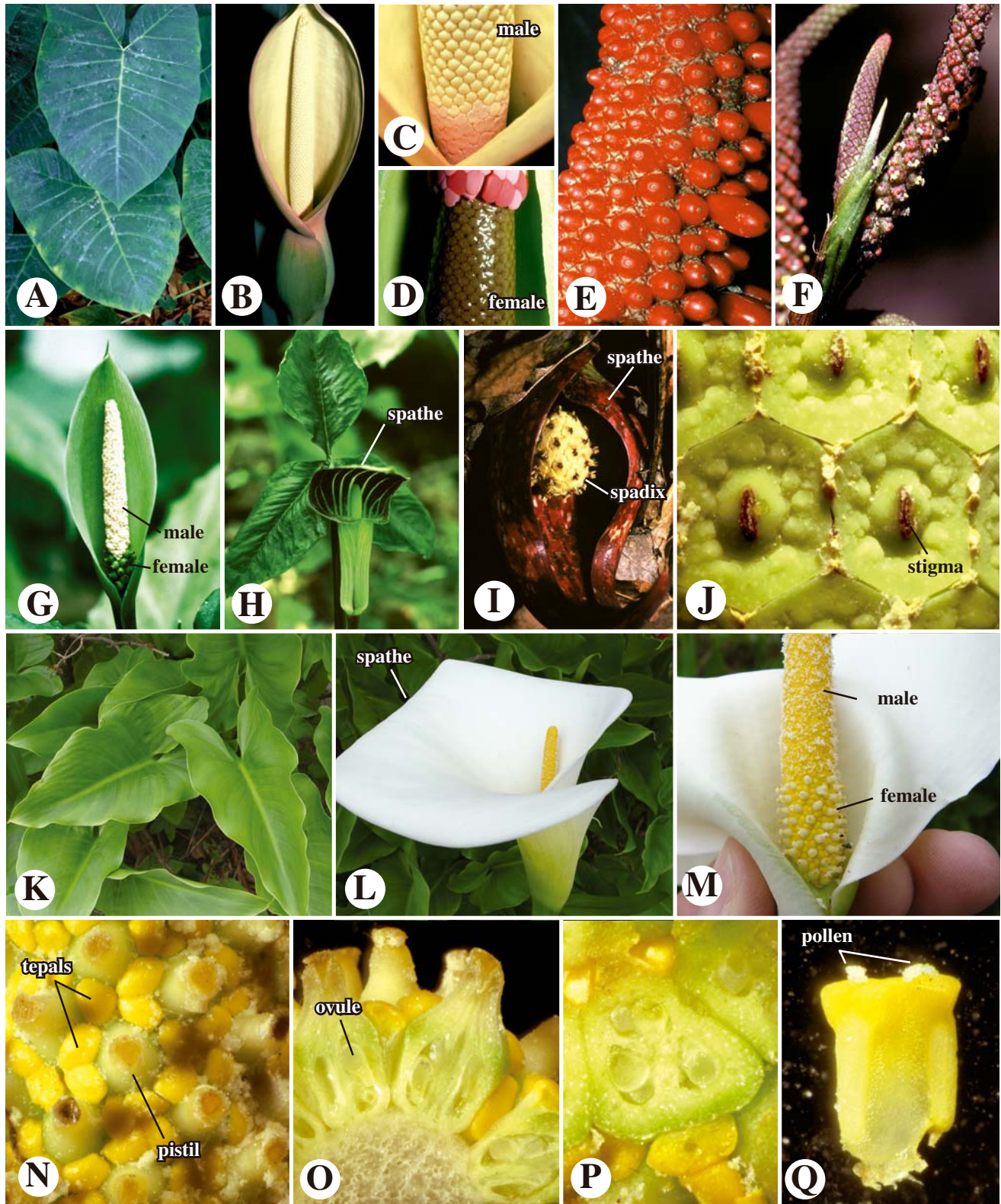


FIGURE 7.22 ALISMATALES. Araceae. A–D. *Xanthosoma sagittifolium*. A. Whole plant, with large, sagittate leaves. B. Inflorescence, a spadix and surrounding spathe. C. Close-up of distal male flowers. D. Close-up of proximal female flowers. E. *Anthurium* sp., multiple fruit of berries. F. *Gymnostachys anceps*, inflorescence. G. *Aglaonema modestum*, inflorescence. H. *Arisaema triphyllum* (jack-in-the-pulpit), inflorescence and leaf. I. *Symplocarpus foetidus* (skunk weed), inflorescence. J. *Monstera deliciosa*, flowers (bisexual), showing outer face of hexagonal pistil and peripheral stamens. K–Q. *Zantedeschia aethiopica* (calla lily). K. Sagittate leaves. L, M. Inflorescence. N. Female flowers, face view. O. Female flowers, pistil longitudinal section, showing basal placentation. P. Ovary cross-section, showing three carpels and locules. Q. Anther, with poricidal dehiscence.



FIGURE 7.23 ALISMATALES. Araceae diversity. **A.** *Pothos* sp., a vine. **B.** *Philodendron selloum*, a rhizomatous shrub. **C.** *Amorphophallus titanum*, having among the largest inflorescences of any flowering plant. **D.** *Pistia stratiotes*, water-lettuce, a floating aquatic. **E.** *Lemna* sp., duckweed, a floating aquatic. **F.** *Wolfia* sp., water-meal, a floating aquatic, having the smallest flower of any flowering plant. **G.** *Wolfia borealis* in flower, with no perianth, one anther, and one pistil (with circular stigma) inside dorsal pouch. Note tip of sewing needle for size. (C, courtesy of Constance Gramlich; G, courtesy of Wayne Armstrong.)

umbel-like], with flowers or flower axes whorled and spathe absent. **Flowers** are bisexual or unisexual, actinomorphic, subsessile to pedicellate, bracteate, hypogynous; the receptacle is flat or expanded and convex. The **perianth** is biseriata and dichlamydeous, trimerous, hypanthium absent. The **calyx** consists of 3, aposepalous sepals. The **corolla** consists of 3, apopetalous, caducous petals. **Stamens** are 6, 9, or ∞ [3], whorled, distinct, free, uniseriate or biseriata (often in pairs); anthers are longitudinal, and extrorse or latrorse in dehiscence. The **gynoecium** is apocarpous, with a superior ovary, 3, 6, or ∞ carpels, and 1 terminal style and stigma; placentation is basal [rarely marginal]; ovules are anatropous, bitegmic, 1 [∞] per carpel. The **fruit** is an aggregate of achenes or of basally dehiscent follicles. **Seeds** are exalbuminous.

The Alismataceae have a worldwide distribution, esp. in N. temperate regions. Economic importance includes taxa used

as food by indigenous people, others used as aquatic, cultivated ornamentals. See Haynes et al. (1998) for more information on the family.

The Alismataceae are distinguished from related families in consisting of *aquatic or marsh herbs* with *basal* leaves, usually *whorled* flowers or flower axes, and *dichlamydeous* flowers with an *apocarpous* gynoecium having *basal placentation*.

K 3 **C** 3 **A** 6,9- ∞ [3] **G** 3,6- ∞ superior.

ASPARAGALES

The Asparagales, sensu APG II (2003), contain approximately 24 families of monocotyledons, including a large and diverse number of taxa (Table 7.2, although note that many

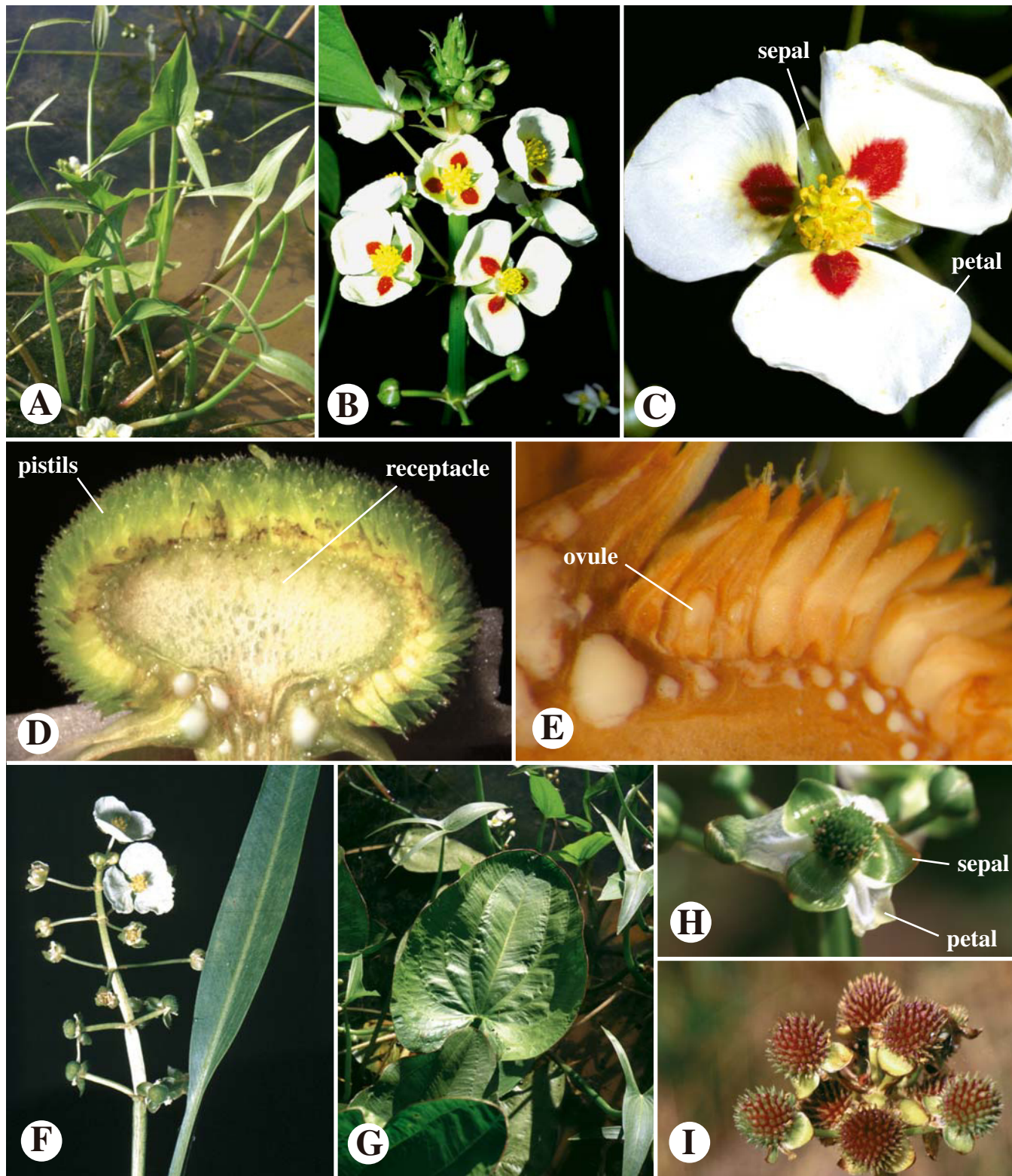


FIGURE 7.24 ALISMATALES. Alismataceae. **A–C.** *Sagittaria montevidensis*, arrowhead. **A.** Emergent, aquatic plant, with sagittate leaves. **B.** In inflorescence. **C.** Male flower close-up, showing dichlamydeous perianth. **D–F.** *Sagittaria* spp. **D.** Female flower longitudinal section, showing expanded receptacle and numerous pistils. **E.** Close-up of pistils, each with a single ovule having basal placentation. **F.** Leaf and inflorescence, the latter a raceme of whorled flowers. **G–I.** *Echinodorus berteroi*, burhead. **G.** Leaf. **H.** Flower (bisexual) close-up. **I.** Maturing fruit, an achenecetum, with persistent sepals.



FIGURE 7.25 ASPARAGALES. Seeds of *Agapanthus* (left) and *Yucca* (right), both with black, phytomelan-encrusted seed coat, a general apomorphy for the order.

families could alternatively be united). A possible apomorphy uniting the order (aside from molecular sequence data) is the presence of seeds having a seed coat containing a black substance called **phytomelan** (Figure 7.25). The phytomelaniferous seeds of the Asparagales have apparently been lost in some taxa, particularly those that have evolved fleshy fruits.

Family delimitations of the Asparagales have undergone a number of changes in recent years and more work is needed before these stabilize. Seven families are described here. Notable among the others are the **Agapanthaceae**, with *Agapanthus* spp. being common cultivars (Figure 7.26A,B); **Asparagaceae**, including the vegetable, *Asparagus officinalis*, and several ornamental species, such as *A. setaceus*, asparagus fern; **Blandfordiaceae** (Figure 7.26C); **Doryanthaceae** (Figure 7.26J,K); **Hemerocallidaceae** (Figure 7.26E G), including *Hemerocallis fulva*, day-lily; **Hyacinthaceae**, including several ornamental cultivars; **Hypoxidaceae** (Figure 7.26H); **Laxmanniaceae** (Figure 7.26D,I); and **Xanthorrhoeaceae**, the grass trees (Figure 7.26L,M). See Fay et al. (2000) and Rudall (2003) for recent phylogenetic and morphological studies of the Asparagales.

Agavaceae Agave family (after Agave, meaning admired one). Ca. 8 (-12+) genera/300+ species. (Figures 7.27, 7.28)

The Agavaceae consist of perennial subshrubs, shrubs, trees, or possibly herbs. The **stems** are a caulescent caudex, rhizome, bulb, or are arborescent, sympodial in taxa with branched stems, some species with anomalous secondary growth. The **leaves** are parallel veined, often large, xeromorphic, fibrous or rarely succulent, basal and rosulate or acrocaulis, spiral, simple, undivided, the apex or margin sometimes toothed or spined. The **inflorescence** is a panicle, raceme, or spike in

some producing vegetative plantlets. The **flowers** are bisexual, actinomorphic or zygomorphic, bracteate, hypogynous or epigynous. The **perianth** is biseriata, homochlamydeous of 3+3 tepals, apotepalous or syntepalous, a hypanthium present in some. The **stamens** are 6, distinct, the filaments long and thin to short and thick. **Anthers** are dorsifixed, versatile, longitudinal and introrse in dehiscence, tetrasporangiate, dithecal. The **gynoecium** is syncarpous, with a superior or inferior ovary and 3 carpels and locules. The **style** is solitary; **stigmas** are solitary or 3-lobed. **Placentation** is axile; **ovules** are anatropous, bitegmic, ∞ and in 2 rows per carpel. **Septal nectaries** are present. The **fruit** is a loculicidal or septicidal capsule or indehiscent (dry or fleshy). The **seeds** are black, phytomelanous, and flattened. Flowers are pollinated by bats, bees, hummingbirds, or moths; *Tegiticula* moths have a symbiotic relationship with *Yucca* species, the female moths transferring pollen and ovipositing the ovaries (the developing larvae feeding on some of the seeds). The chromosomes are dimorphic in size, characteristically 5 long and 25 short.

Members of the Agavaceae occur in xeric to mesic habitats, with many found in dry areas, and often have CAM photosynthesis. The family is distributed in the New World, ranging from the central U.S. to Panama, Caribbean islands, and northern South America. Economic importance includes use by indigenous cultures as a source of fiber, food, beverages, soap, and medicinals. The leaves of *Agave sisalana* are the source of sisal fiber and *A. fourcroydes* of henequen. The fermented and distilled young flowering shoots of *Agave tequilana* are the primary source of tequila.

A recent study by Bogler et al. (2005) suggests that the Agavaceae could be expanded (as Agavaceae s.l.) to include at least four other basal genera, *Camassia*, *Chlorogalum*, *Hesperocallis*, and *Hosta*, with additional genera likely to be added. Many of these are herbaceous, and all seem to have dimorphic chromosomes as occur in traditional family members. See also Bogler and Simpson (1995, 1996) for phylogenetic studies within the family and Verhoek (1998) for a recent family treatment.

The Agavaceae are distinctive in being **perennial** subshrubs to branched trees with spiral, **xeromorphic**, **generally fibrous** leaves, trimerous **hypogynous to epigynous** flowers, and characteristic **dimorphic chromosomes** (base number with 5 long and 25 short chromosomes), the latter a possible apomorphy.

P 3+3 **A** 6 **G** (3), superior or inferior, hypanthium present in some.

Alliaceae Onion family (Latin name for garlic). 13 genera/ca. 600 species. (Figure 7.29)

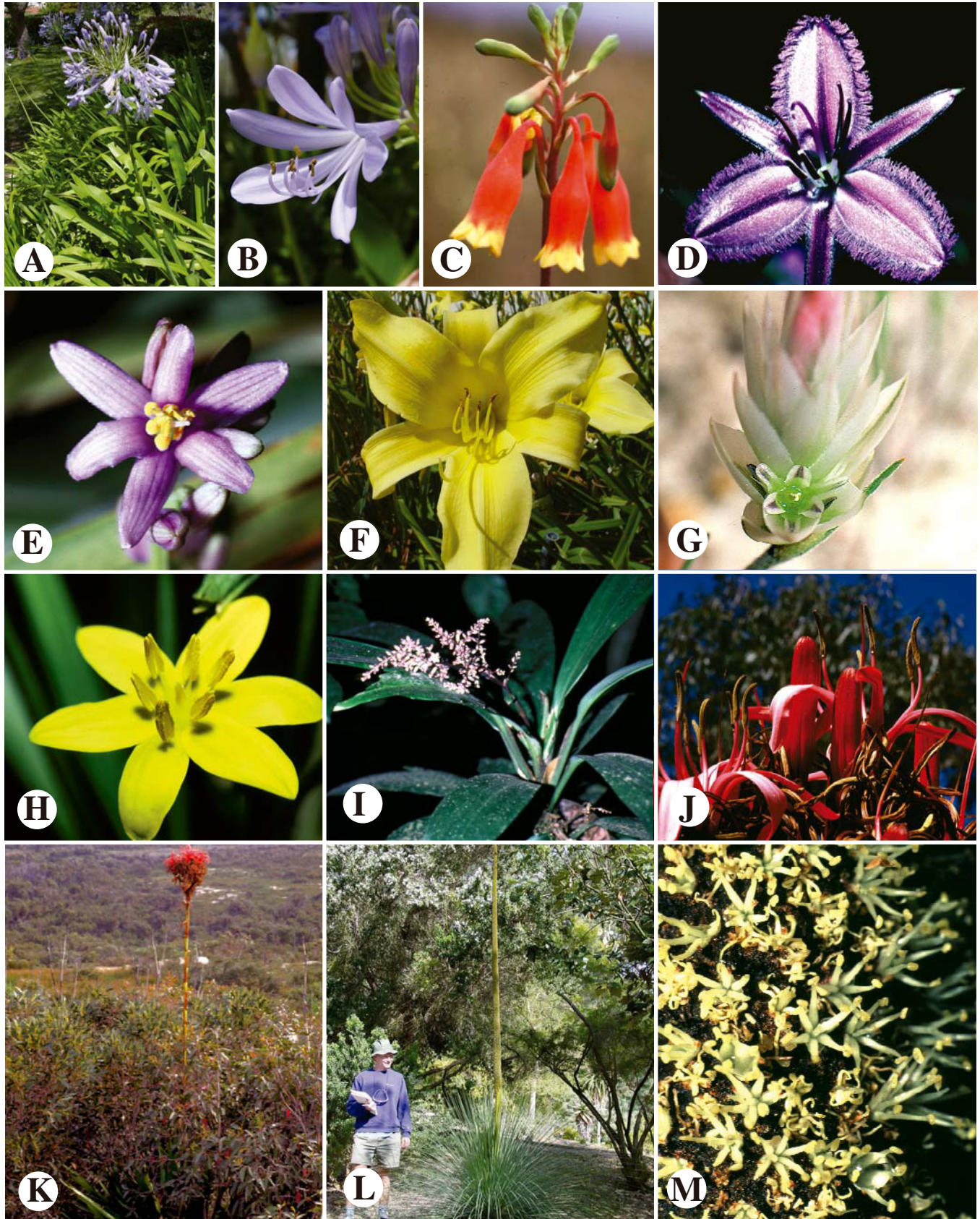


FIGURE 7.26 ASPARAGALES exemplars. A–B. *Agapanthus orientalis*, Agapanthaceae. C. *Blandfordia nobilis*, Blandfordiaceae. D. *Thysanotus* sp., Laxmanniaceae. E–G. Hemerocallidaceae. E. *Dianella laevis*. F. *Hemerocallis fulva*, day-lily. G. *Johnsonia* sp. H. *Hypoxis* sp., Hypoxidaceae. I. *Cordyline* sp., Laxmanniaceae. J–K. *Doryanthes excelsa*, Doryanthaceae. L–M. *Xanthorrhoea* spp., Xanthorrhoeaceae.

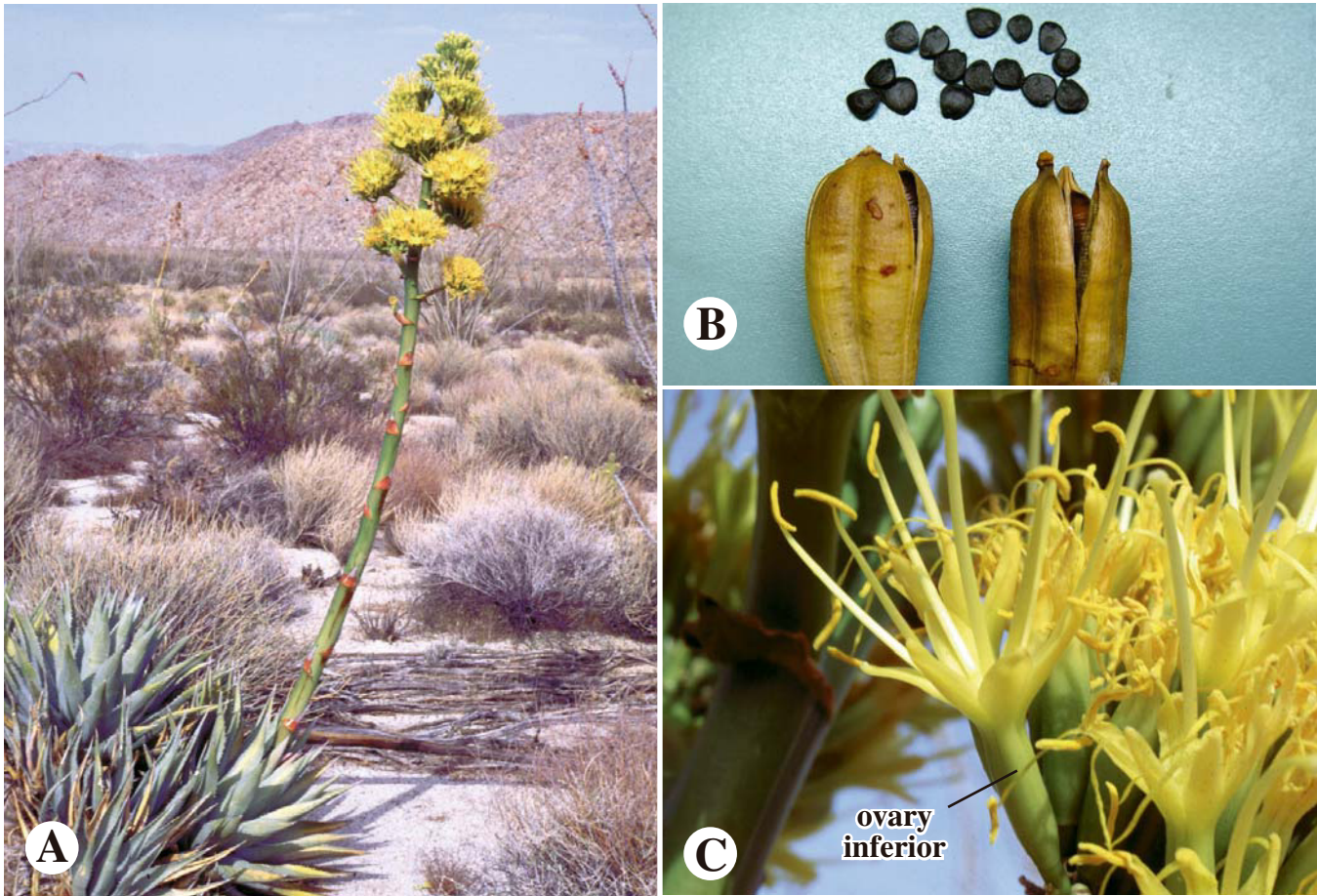


FIGURE 7.27 ASPARAGALES. Agavaceae. *Agave deserti*. **A.** Whole plant, with basal, brood leaves and terminal panicle. **B.** Fruit, a loculicidal capsule; note phytomelan-encrusted seeds. **C.** Flower close-up, showing homochlamydeous perianth and inferior ovary.

The Alliaceae consist of biennial or perennial herbs, usually with a distinctive onion-like (alliaceous) odor. The **stems** are acaulescent and usually a bulb, rarely a short rhizome or corm, typically enveloped by membranous scale leaves or leaf bases. The **leaves** are simple, basal, spiral, closed-sheathing, acicular, linear, or lanceolate [rarely ovate], parallel veined. The **inflorescence** is a terminal, scapose umbel (derived from condensed, monochasial cymes, sometimes termed a pseudo-umbel), rarely a spike or of solitary flowers, with membranous and spathe-like bracts. The **flowers** are bisexual, actinomorphic, pedicellate (pedicels sometimes apically articulate), membranous-bracteate, and hypogynous. The **perianth** is biseriate, homochlamydeous, campanulate to tubular, hypanthium absent, with 3 outer and 3 inner, distinct to connate tepals, a corona sometimes present. The **stamens** are 3+3 [rarely 3 or 2 with staminodes], whorled, diplostemonous, biseriate, unfused or epitepalous; the filaments are generally flat. **Anthers** are versatile, longitudinal and introrse

in dehiscence. The **gynoecium** is syncarpous, with a superior [rarely half-inferior] ovary, 3 carpels, and 3 locules. The **style** is solitary, terminal or gynobasic; the **stigma** is solitary, trilobed to capitate, dry to wet. **Placentation** is axile; **ovules** are campylotropous to anatropous, 2∞ per carpel. **Septal nectaries** are present. The **fruit** is a loculicidal capsule. The **seeds** are black, phytomelanous, ovoid, ellipsoid or subglobose, endospermous, the endosperm rich in oils and aleurone. Family members contain alliin, which is enzymatically converted by wounding to allyl sulfide compounds, the latter imparting the distinctive onion-like odor and taste.

The Alliaceae have a mostly worldwide distribution, mainly northern hemisphere, S. American, and S. African. Economic importance includes important food and flavoring plants, including onion (*Allium cepa*), garlic (*A. sativum*), leek (*A. ampeloprasum*), chive (*A. schoenoprasum*), and other *Allium* species. Garlic also has documented medicinal properties. Several taxa are used as ornamental cultivars,

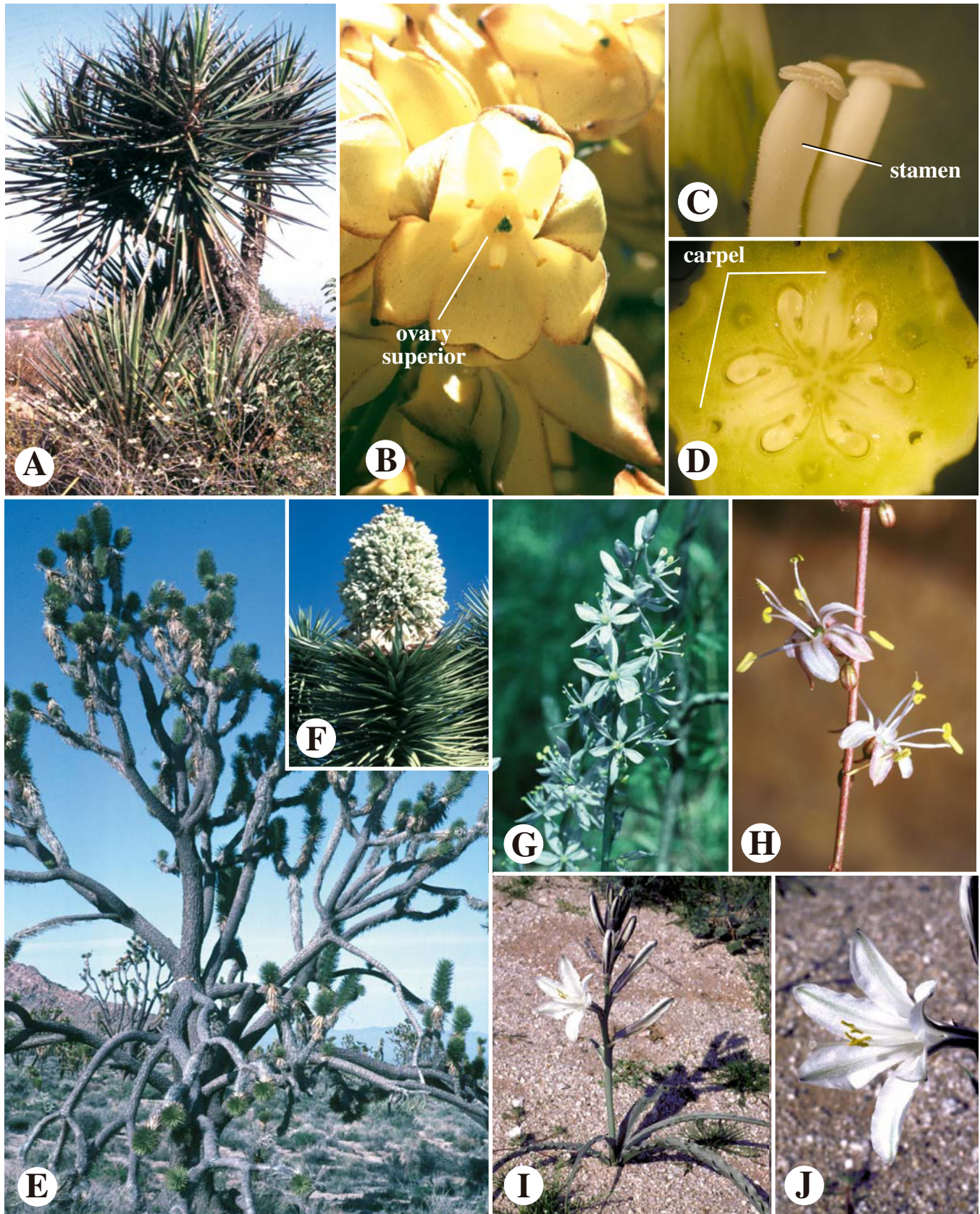


FIGURE 7.28 ASPARAGALES. Agavaceae. **A.** *Yucca schidigera*, showing arborescent habit with acrocaulis leaves. **B.** *Hesperoyucca whipplei*. Note trimerous, homochlamydeous flowers with superior ovaries. **C.** *Yucca* sp. anther. **D.** *Yucca* sp. ovary. **E,F.** *Yucca brevifolia*, Joshua tree. **E.** Whole plant, arborescent with acrocaulis leaves and terminal panicles. **F.** Close-up of leaves and panicle. **G–J.** Basal members of expanded Agavaceae s.l. **G.** *Camassia scilloides*. **H.** *Chlorogalum parviflorum*. **I,J.** *Hesperocallis undulata*, desert-lily, a bulbous, perennial herb.

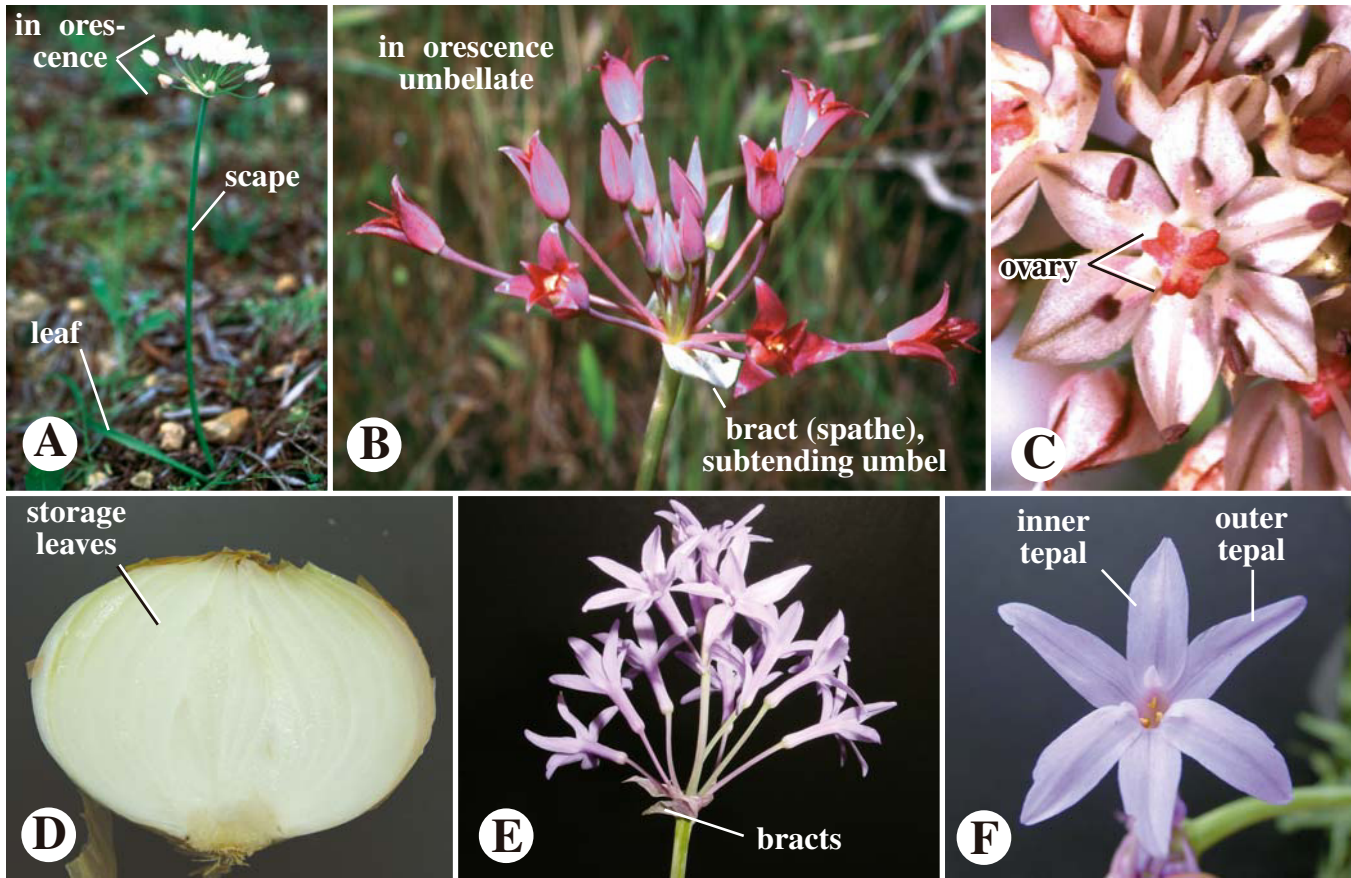


FIGURE 7.29 ASPARAGALES. Alliaceae. **A.** *Allium praecox*, showing basal leaves, scape, and simple umbel. **B.** *Allium peninsulare*, showing close-up of umbel with subtending spathe-like bract. **C.** *Allium praecox*, flower close-up, showing biseriate, homochlamydeous perianth, six stamens, and superior ovary. **D.** *Allium cepa*, onion, bulb longitudinal section. **E,F.** *Tulbaghia violacea*.

e.g., *Ipheion*, *Leucocoryne*, and *Tulbaghia* spp. See Rahn (1998a) for a recent family treatment of the Alliaceae.

The Alliaceae are distinctive in being generally **bulbous herbs**, with **basal**, usually **narrow** leaves, an **umbellate** inflorescence, and a usually **superior ovary**.

P 3+3 **A** 3+3 [3,2] **G** (3), superior [rarely half-inferior].

Amaryllidaceae Amaryllis family (Latin name for a country girl). 59 genera / 850 species. (Figure 7.30)

The Amaryllidaceae consist of terrestrial, rarely aquatic or epiphytic, perennial herbs. The **stems** are bulbs, covered by membranous leaf bases, the **tunica**. The **leaves** are simple, undivided, spiral or distichous, sheathing or not, sessile or petiolate, and parallel veined. The **inflorescence** is a terminal, scapose umbel (derived from condensed, monochasial cymes, sometimes termed a pseudo-umbel), rarely of solitary flowers, with bracts present, enclosing the flower buds. The **flowers** are bisexual, actinomorphic or zygomorphic,

pedicellate or sessile, bracteate, epigynous to epiperigynous. The **perianth** is biseriate, homochlamydeous, trimerous, apotepalous or syntepalous, and forming a short to long hypanthial tube, sometimes with a perianth corona (e.g., *Narcissus*). The **stamens** are generally biseriate, 3+3 [3 18], distinct or connate, forming a staminal corona in some (e.g., *Hymenocallis*). **Anthers** are usually dorsifixed, longitudinal [rarely poricidal], and introrse in dehiscence. The **gynoecium** is syncarpous, with an inferior ovary, 3 carpels, and 3 [1] locules. **Placentation** is axile or basal; ovules are anatropous, bitegmic, unitegmic, or ategmic. The **fruit** is a loculicidal capsule or rarely a berry. The **seeds** are phytomelaniferous.

The Amaryllidaceae have a worldwide distribution, being especially concentrated in South America and South Africa. Economic importance is primarily as innumerable cultivated ornamentals, such as *Amaryllis* (belladonna-lily), *Crinum*, *Galanthus* (snowdrop), *Hippeastrum* (amaryllis), *Leucojum* (snowflake), *Lycoris* (spider-lily), and *Narcissus* (daffodil);

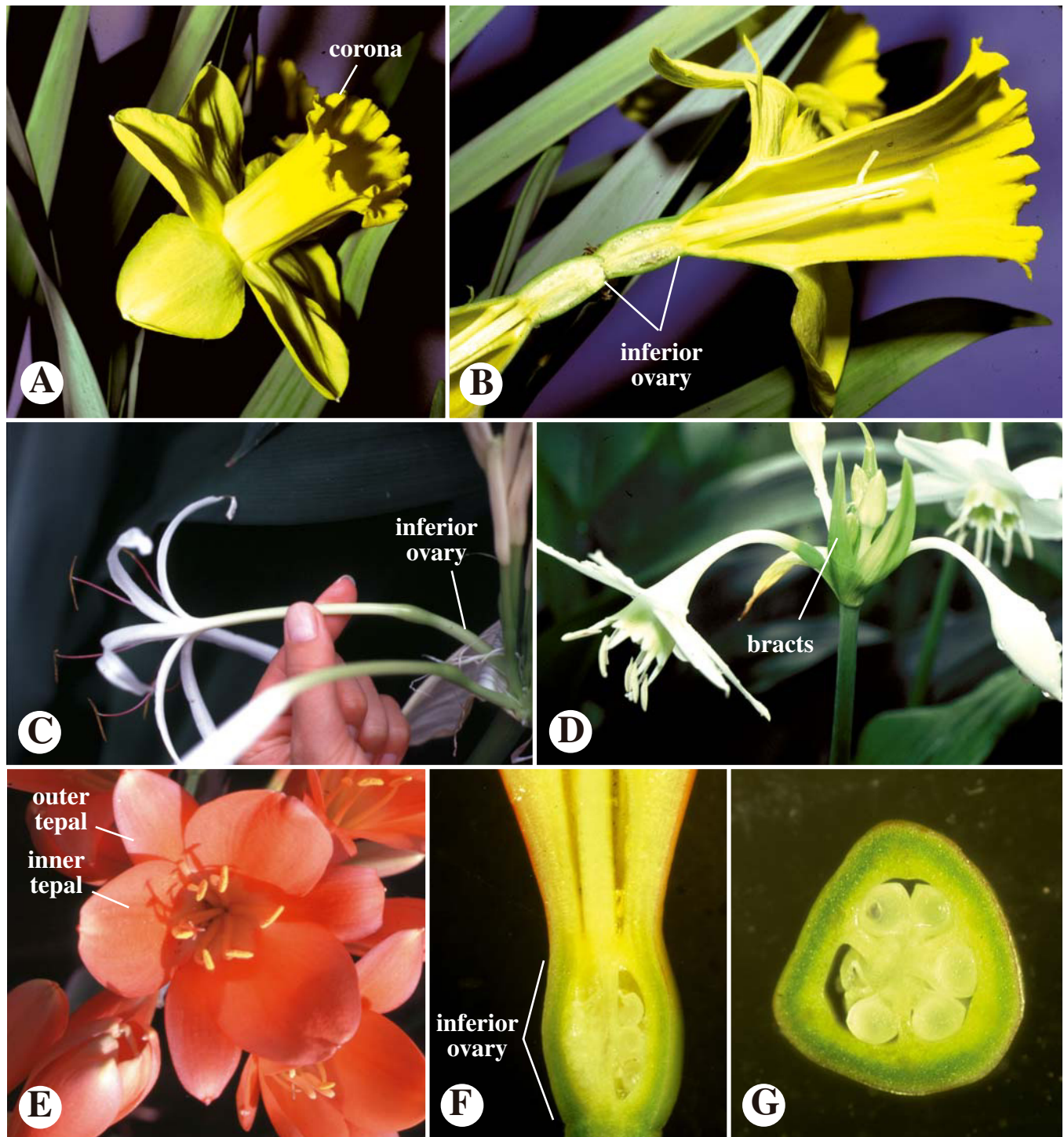


FIGURE 7.30 ASPARAGALES. Amaryllidaceae. **A,B.** *Narcissus pseudonarcissus*. **A.** Flower, showing elongate, tubular corona. **B.** Flower, longitudinal section. Note inferior ovary. **C.** *Crinum* sp., showing inferior ovary. **D.** *Eucharis grandiflora*. Note spatheous bracts subtending ovaries. **E–G.** *Clivia miniata*. **E.** Flower, face view. **F.** Ovary longitudinal section. **G.** Ovary cross-section. Note three carpels and locules with axile placentation.

several taxa are used by indigenous peoples for medicinal, flavoring, psychotropic, or other purposes. See Meerow and Snijman (1998) and Meerow et al. (1999, 2000) for phylogenetic studies of the family.

The Amaryllidaceae are distinctive in being perennial, **bulbous herbs** with an **umbellate** inflorescence and an **inferior ovary**.

P 3+3 or (3+3) **A** 3+3 or (3+3) [3-18] **G** (3), inferior, hypanthium present.

Asphodelaceae Asphodel or Aloe family. 15 genera / 780 species. (Figure 7.31)

The Asphodelaceae consist of herbs to [rarely] pachycaulous trees. **Roots** are often succulent, with a velamen in some taxa. The **stems** exhibit anomalous secondary growth in some taxa, as in *Aloe*. The **leaves** are usually succulent, simple, spiral to distichous, undivided, parallel-veined, and dorsiventral to terete, the margins entire to toothed or spinose. The **inflorescence** is a raceme or panicle. The **flowers** are bisexual, actinomorphic or zygomorphic, pedicellate, bracteate or not, hypogynous. The **perianth** is biseriate, homochlamydeous, 3+3, apotepalous or syntepalous. The **stamens** are 3+3, distinct. **Anthers** are dorsifixed to basifixed, longitudinal and introrse in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 carpels, and 3 locules. **Placentation** is axile; **ovules** are 2 ∞ per carpel. Septal nectaries are present. The **fruit** is a loculicidal capsule or (rarely) berry. The **seeds** have an aril present.

Members of the Asphodelaceae grow in temperate and subtropical Africa, particularly southern Africa. Economic importance includes *Aloe* spp. (esp. *A. vera* and *A. ferox*, from which aloin is derived), which have important uses medicinally (e.g., as laxatives and treatment of burns) as well as in skin, hair, and health products; many family members are important as cultivated ornamentals, e.g., *Aloe*, *Asphodelus*, *Gasteria*, *Haworthia*, *Kniphofia*. See Smith and v. Wyk (1998) for a recent family treatment and Chase et al. (2000b) for a phylogenetic analysis of the family.

The Asphodelaceae are distinguished from related taxa in being **herbs** or **pachycaulous trees** with leaves usually **succulent**, flowers trimerous with a superior ovary, and the seeds **arillate**.

P 3+3 or (3+3) **A** 3+3 **G** (3), superior.

Iridaceae Iris family (after Iris, mythical goddess of the rainbow). 70 genera/1750 species. (Figure 7.32)

The Iridaceae consist of perennial [rarely annual] herbs or shrubs with anomalous secondary growth, achlorophyllous and saprophytic in *Geosiris*. The **stems** are rhizomatous, cormose, bulbous, or a woody caudex. The **leaves** are unifacial

(with leaf plane parallel to stem) or terete, simple, narrow and generally ensiform, sheathing, often equitant, distichous, and parallel-veined [scalelike and achlorophyllous in *Geosiris*]. The **inflorescence** is a terminal spike, solitary flower, or a spike or panicle of clusters of 1 many monochasial cymes (often rhipidia), typically subtended by two spathe-like bracts; inflorescence subterranean in *Geosiris*. **Flowers** are bisexual, actinomorphic or zygomorphic, pedicellate or sessile, bracteate, epigynous or rarely hypogynous (*Isophysis*). The **perianth** is biseriate, homochlamydeous, 3+3, apotepalous or syntepalous (forming a prominent tube in Ixioideae), a hypanthium present or absent. **Stamens** are 3, opposite the outer tepals, distinct or monadelphous; anthers are longitudinally extrorse or poricidal in dehiscence. The **gynoecium** is syncarpous, with an inferior (superior in *Isophysis* only) ovary, 3 carpels and locules, style(s) terminal, petaloid in many Iridoideae; placentation is axile (rarely parietal); ovules are anatropous, bitegmic, 1 ∞ per carpel. The **fruit** is a loculicidal capsule; seeds are endospermous with a dry or fleshy seed coat.

The Iridaceae has been classified into two subfamilies, Isophysidoideae (one genus, *Isophysis*, having a superior ovary) and Iridoideae (all other genera, with an inferior ovary). Within the latter subfamily are three commonly recognized tribes: Iridoideae and Nivenioideae with radial, pedicellate flowers and rhipidia enclosed by large, spathe-like bracts (Nivenioideae differing in having paired rhipidia) and Ixioideae with radial or bilateral, sessile flowers (with two bracts at base) usually with a long perianth tube and arranged on a spike or flowers solitary. Members of the family have a worldwide distribution, being especially diverse in southern Africa. Economic importance includes extensive use as ornamental cultivars, e.g., as cut flowers, especially species of *Iris*, *Gladiolus*, *Freesia*, and *Crocus*; the styles and stigmas of *Crocus sativus* are the source of the spice saffron; corms of some species are eaten by indigenous people. See Goldblatt et al. (1998) and Reeves et al. (2001) for phylogenetic studies in the Iridaceae.

The Iridaceae are distinguished from related families in being usually **perennial herbs** with generally **ensiform, unifacial** leaves, a bracteate **spike or panicle** of **solitary flowers** or **monochasial cyme (rhipidia) clusters**, and flowers with **three stamens opposite outer tepals**.

P 3+3 or (3+3) **A** 3 or (3) **G** (3), inferior (superior in *Isophysis*).

Orchidaceae Orchid family (*orchis*, testicle, from the shape of the root tubers). 700-800 genera / ca. 20,000 species. (Figures 7.33-7.36)

The Orchidaceae consist of terrestrial or epiphytic, perennial [rarely annual] herbs [rarely vines]. The **roots** are often

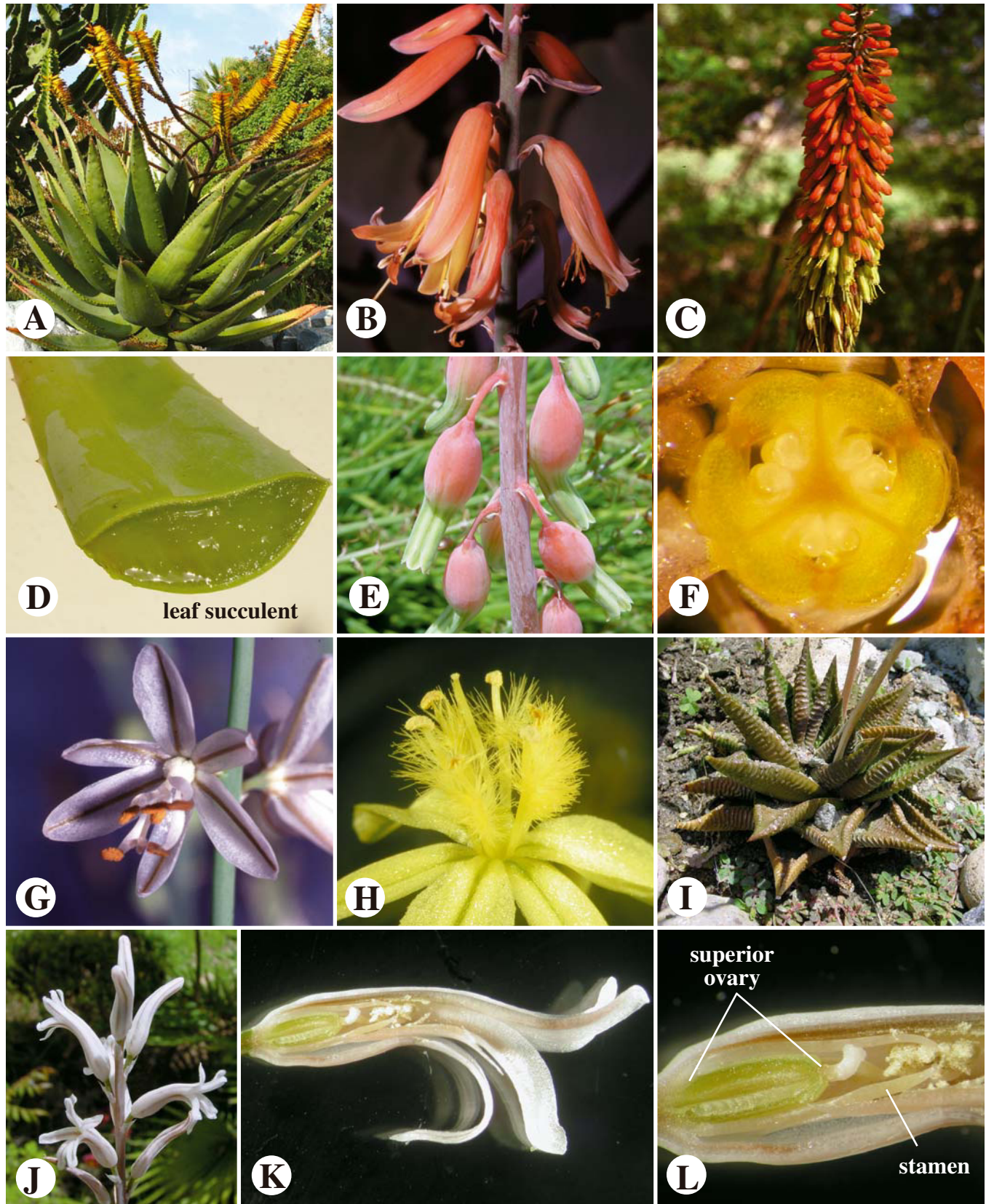


FIGURE 7.31 ASPARAGALES. Asphodelaceae. **A.** *Aloe marlothii*, rosette of succulent, spinose leaves. **B.** *Aloe* sp., showing zygomorphic flowers with tubular perianth. **C.** *Kniphofia* sp., red-hot poker. **D.** *Aloe* sp., cut succulent leaf. **E,F.** *Gasteria trigona*. **E.** Flowers. **F.** Ovary cross-section. **G.** *Asphodelus stulosus*, flower. **H.** *Bulbine* sp., showing actinomorphic flower with apotepalous perianth and pilose stamen filaments. **I.** *Haworthia linifolia*, basal rosette of leaves. **J–L.** *Haworthia cooperi*. **J.** Inflorescence, a raceme. **K.** Flower longitudinal section, showing syntepalous perianth. **L.** Close-up of ovary longitudinal section and epitepalous stamens.

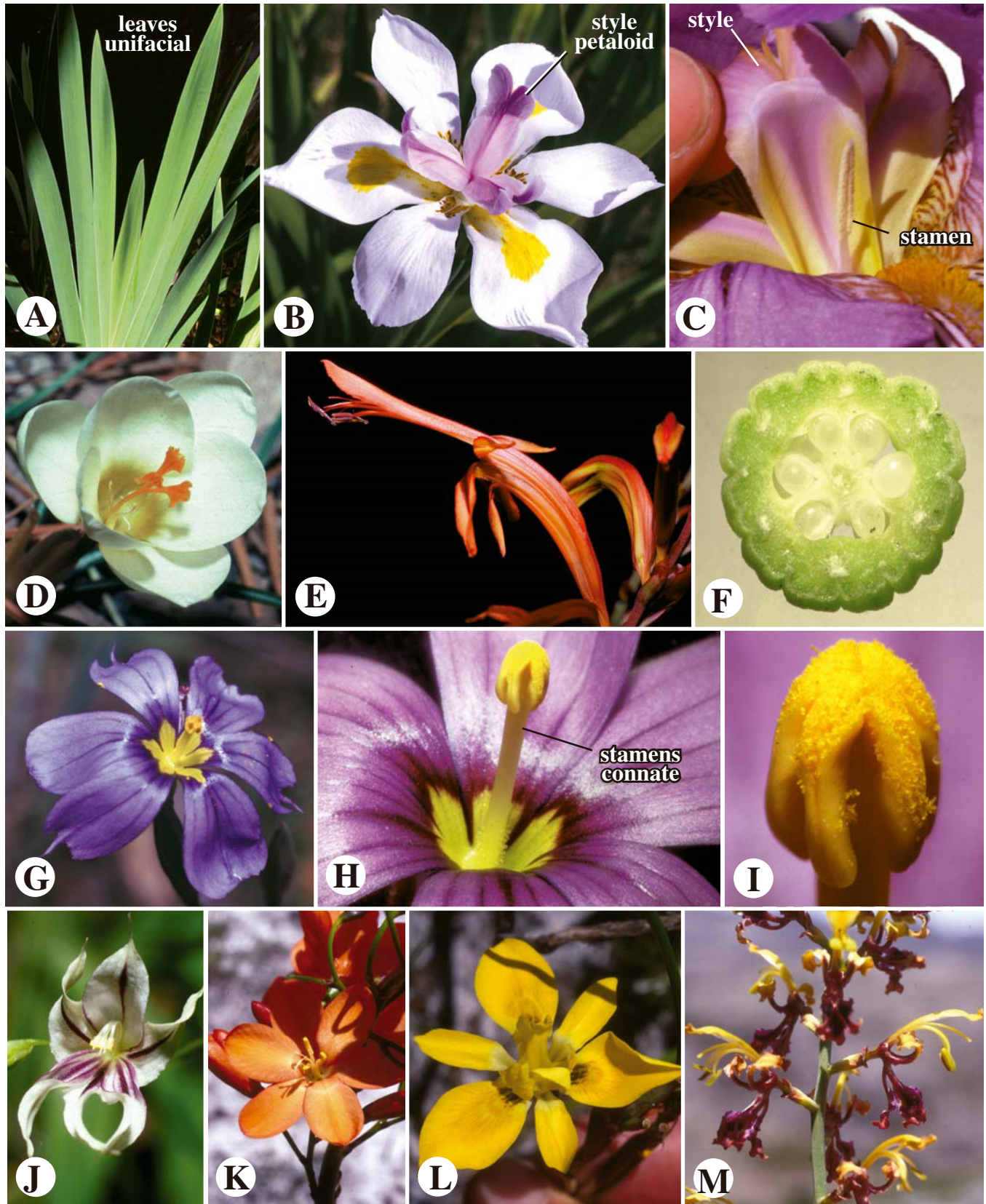


FIGURE 7.32 ASPARAGALES. Iridaceae. **A.** *Iris* sp., showing unifacial leaves that are equitant and distichous. **B.** *Dietes* sp., showing the three outer tepals, three inner tepals, and petaloid styles (corresponding in position to the three carpels). **C.** *Iris* sp., with petaloid style pulled back to show stamen opposite outer tepal. **D.** *Crocus* sp. **E.** *Chasmanthe aethiopica*, an example of a zygomorphic member of the family. **F.** *Iris* sp., inferior ovary cross-section, showing axile placentation. **G,H.** *Sisyrinchium bellum*. **G.** Whole flower. **H.** Close-up of flower, showing central connate stamens. **I.** Close-up of connate anthers. **J.** *Melasphaerula ramosa*. **K.** *Pillansia templemannii*. **L.** *Moraea fugax*. **M.** *Tritoniopsis*.

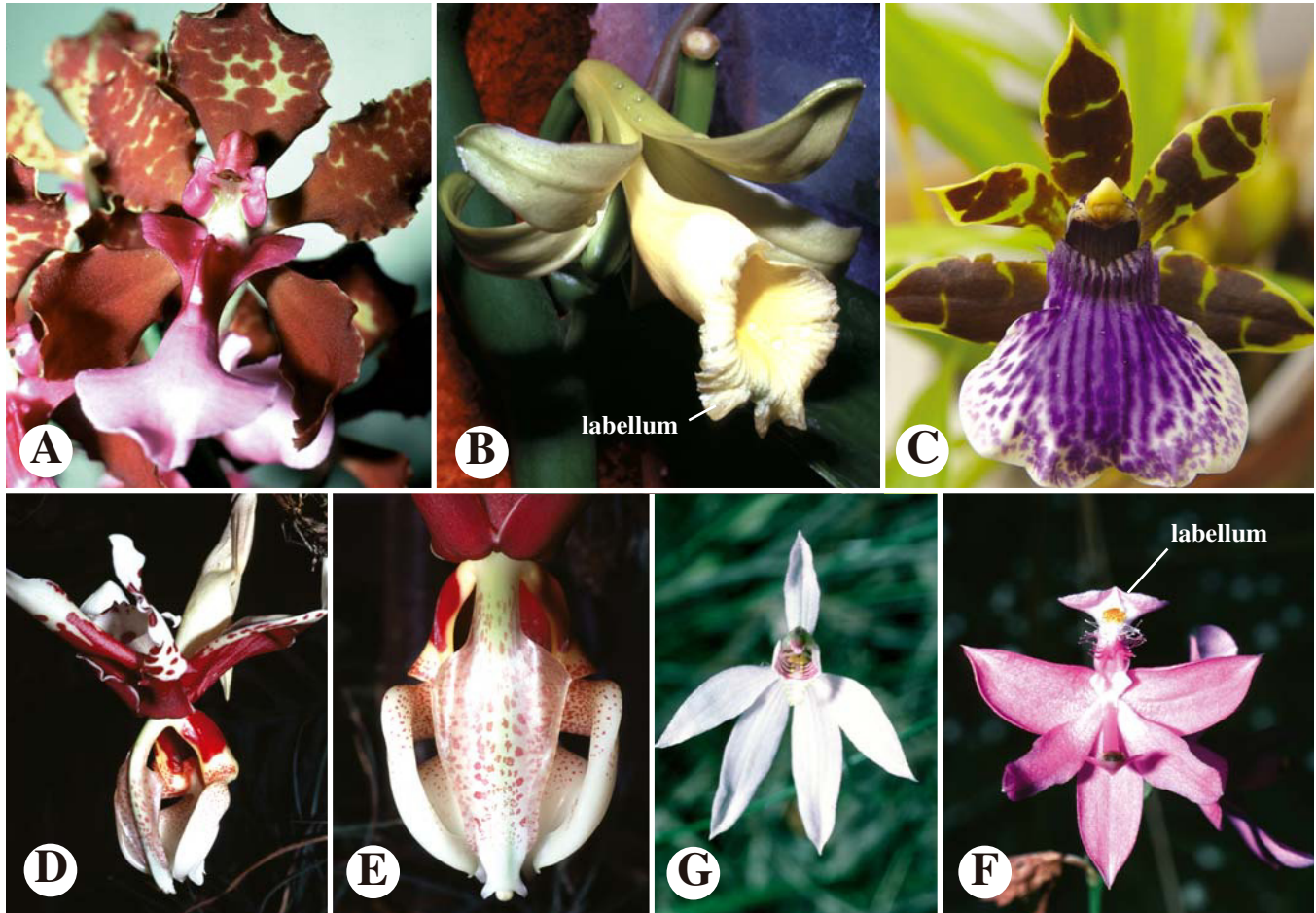


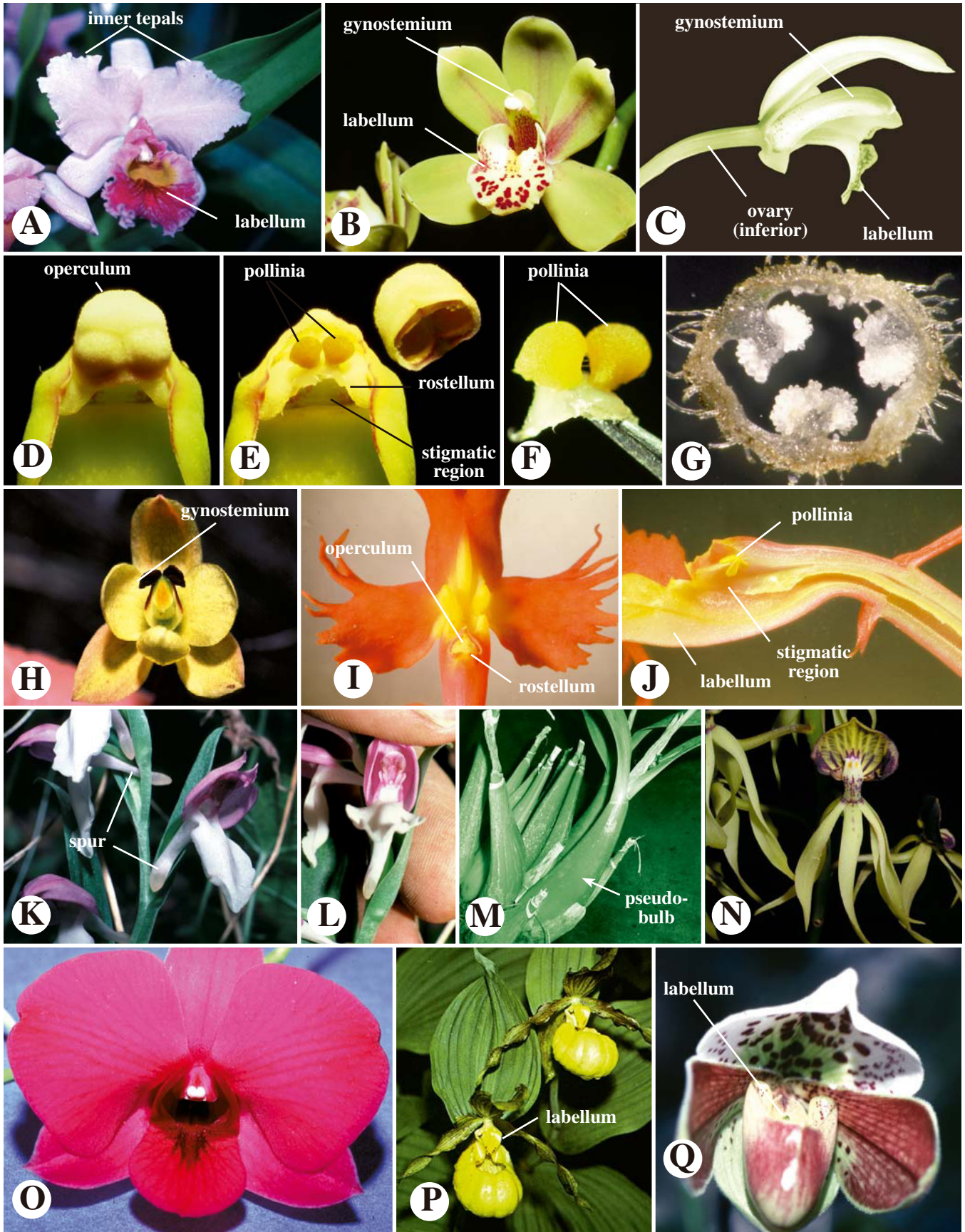
FIGURE 7.33 ASPARAGALES. Orchidaceae. **A.** *Oncidium lanceanum*. **B.** *Vanilla planifolia*, vanilla. **C.** *Zygopetalum* sp. **D,E.** *Stanhopea tigrina*, with pendant flowers. **F.** *Caladenia fuscata*. **G.** *Calopogon* sp., a nonresupinate species.

tuberous (in terrestrial species) or aerial (in epiphytic species), typically with a multilayered velamen. The **stems** are rhizomatous or cormose in terrestrial species, the epiphytic species often with pseudobulbs. The **leaves** are spiral, distichous, or whorled, usually sheathing, simple, and parallel veined. The **inflorescence** is a raceme, panicle, spike, or a solitary flower. The **flowers** are bisexual, rarely unisexual,

zygomorphic, usually resupinate, resulting in a 180° shift of floral parts (Figure 7.36), epigynous. The **perianth** is biseriate, homochlamydeous (although outer and inner whorls are often differentiated), 3+3, apotepalous or basally syntepalous, extremely variable in shape and color, sometimes spurred or with enlarged saclike tepal. The inner median, anterior tepal (when resupinate; actually posterior early in development)

FIGURE 7.34 ASPARAGALES. Orchidaceae (following page). **A.** *Cattleya* sp., showing basic structure of a resupinate orchid flower. Note enlarged and colorful inner, median tepal, the labellum. **B–F.** *Cymbidium* sp. **B.** Whole flower, illustrating prominent gynostemium. **C.** Flower longitudinal section. **D.** Close-up of gynostemium apex. Note operculum covering anther. **E.** Gynostemium apex with operculum removed. **F.** Close-up of pollinarium with two pollinia. **G.** *Ludisia* sp., cross-section of inferior ovary, showing parietal placentation. **H.** *Thelymitra antennifera*, an orchid mimicking an insect. **I,J.** *Epidendrum* sp. **I.** Close-up of gynostemium, which is adnate to the labellum. **J.** Flower longitudinal section. **K,L.** *Orchis spectabilis*. **K.** Whole flower. **L.** Close-up view of gynostemium. **M,N.** *Encyclia cochleata*. **M.** Pseudobulb, found in many epiphytic orchids. **N.** Flower, showing rare nonresupinate orientation. **O.** *Dendrobium* sp. **P.** *Cypripedium* sp., lady's slippers. Note enlarged, swollen labellum. **Q.** *Paphiopedilum* sp.

(See Figure in opposite page)



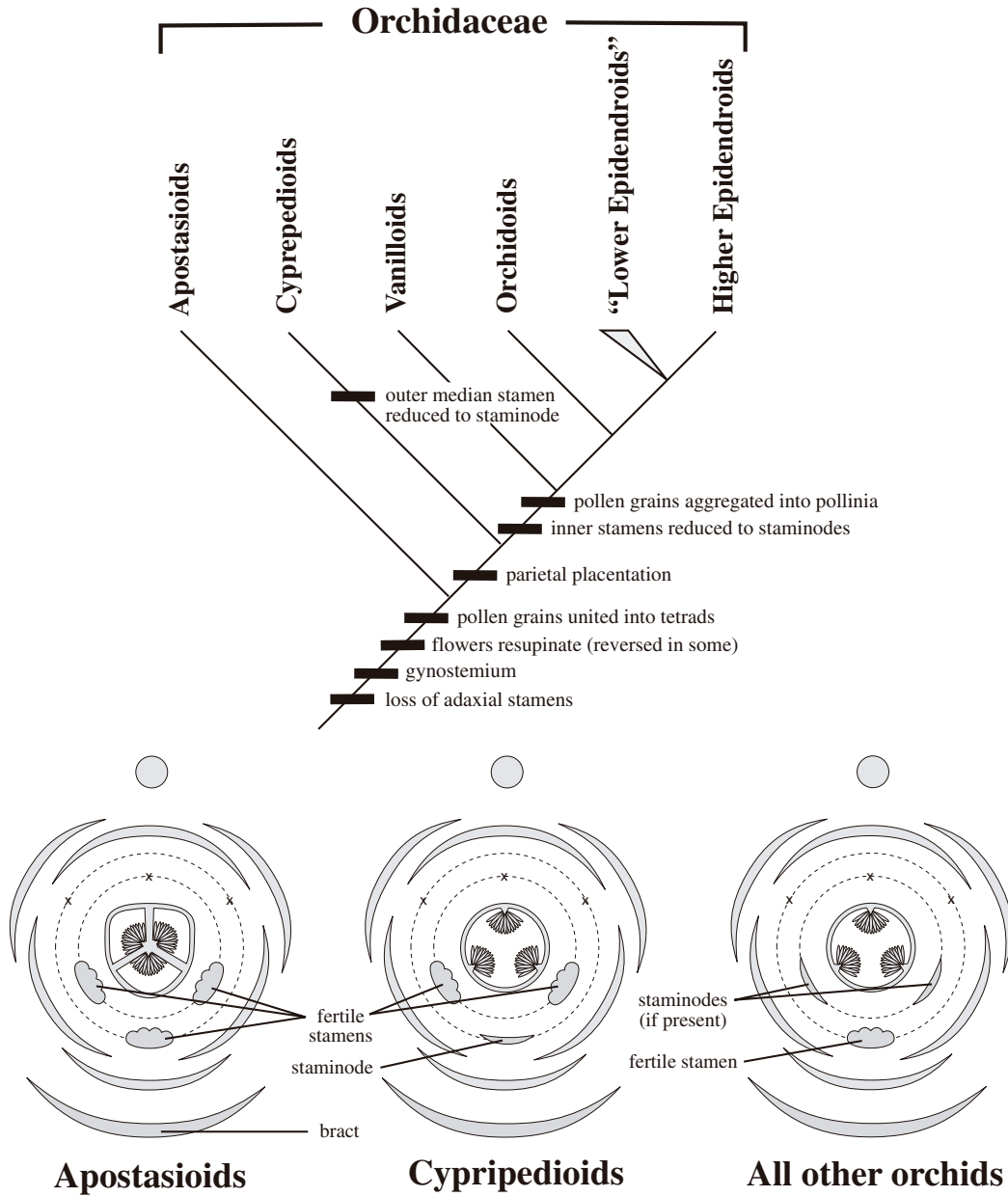


FIGURE 7.35 ASPARAGALES. Orchidaceae. Cladogram of major orchid groups, after Cameron et al. (1999), with putative apomorphies; oral diagrams of Apostasioids, Cyripedioids, and all other orchids (lower), after Dahlgren et al. (1985).

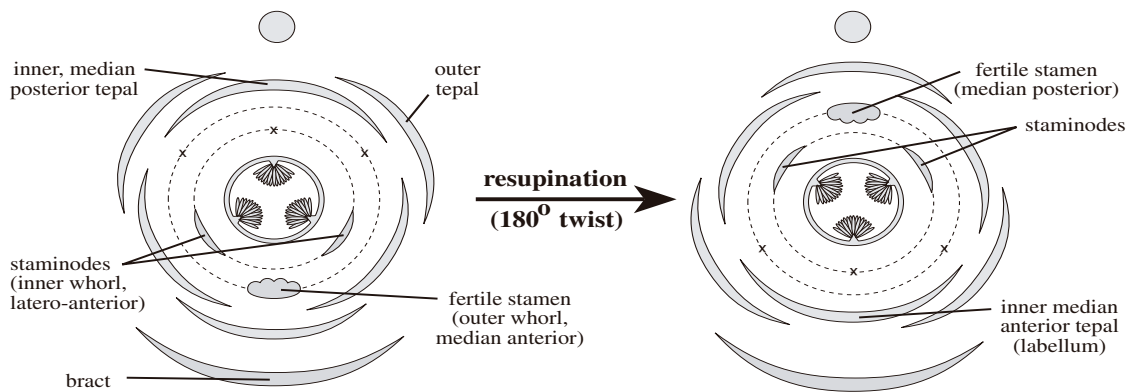


FIGURE 7.36 ASPARAGALES. Orchidaceae. Floral diagram before (left) and after (right) resupination.

is termed the **labellum**, which is typically enlarged, sculptured, or colorful and often functions as a landing platform for pollinators. The **stamen** in most species is solitary, derived from the median stamen of the ancestral outer whorl, often with two vestigial staminodes derived from the lateral stamens of an ancestral inner whorl; in Apostasioideae or Cyripedioideae, there are two or three fertile stamens, when two, derived from the two lateral stamens of the ancestral inner whorl, when three, derived from these plus the median stamen of the outer whorl; the androecium is fused with the style and stigma to form the **gynostemium** (also called the **column** or **gynostegium**). **Anthers** are longitudinally or modified in dehiscence, bisporangiate, dithecal; in all but the Apostasioideae and most Cyripedioideae, the pollen is agglutinated into 1–12 (typically 2 or 4) discrete masses, each termed a **pollinium** (derived from individual anther microsporangia or from fusion products or subdivisions of the microsporangia); the pollinia plus a sticky stalk (derived from either the anther or stigma) are together termed a **pollinarium**, the unit of transport during pollination, the anther connective often modified into an **operculum** (anther cap) that covers the anther(s) prior to pollination. The **pollen** consists of tetrads in most family members, massulae in some, monads in Apostasioideae and Cyripedioideae. The **gynoecium** is syncarpous, with an inferior ovary, 3 carpels, and 1–3 locules. The style is solitary and terminal and is the major component of the gynostemium; a single, enlarged lobe, termed the **rostellum** and interpreted as part of the stigma(s), is positioned above the stigmatic region; the **rostellum** typically is adherent to the pollinarium stalk, the tip of which derives a sticky substance from the surface of the **rostellum** (this sticky region termed the **viscidium**). **Placentation** is parietal or axile; **ovules** are anatropous, usually bitegmic, very many per carpel (sometimes on the order of a million). Nectaries are typically present, variable in position and type. The **fruit** is a loculicidal capsule or rarely a berry. The **seeds** are often membranous-winged, possibly functioning in wind dispersal, and exalbuminous, the endosperm abortive early in development. **Pollination** is effected by various insects (often one species having a specific association with one orchid species), birds, bats, or frogs. The transfer of pollen grains together within the pollinia is an apparent adaptation for ensuring fertilization of many of the tremendous number of ovules. Some species have remarkable adaptations for pollination. Among the more remarkable are several species with visual and chemical mimicry, fooling a male insect into perceiving the flower as a potential mate. The bucket orchid, *Coryanthes*, has an pouchlike labellum that fills with a fluid secreted from the gynostemium; a bee, falling into this fluid, must travel

through a tunnel, forcing deposition of the pollinarium on its body.

The Orchidaceae consist of the basal Apostasioideae or Apostasioids (2–3 stamens, axile placentation, lacking pollinia), Cyripedioideae or Cyripedioids (2 stamens, parietal placentation, lacking pollinia), and the remainder of the orchids (1 stamen, parietal placentation, pollinia), the latter grouped by Cameron et al. (1999) into the Vanilloids, Orchidoids, a paraphyletic Lower Epidendroids, and Higher Epidendroids (Figure 7.35). Members of the family are distributed worldwide. Economic importance is largely as cultivated ornamentals, including some quite monetarily valuable in the horticultural trade; the fermented capsules of *Vanilla planifolia* (Figure 7.33B) are the source of vanilla food flavoring. *Angraecum sesquipedale* Thouars (Madagascar) is known for its long spur (up to 45 cm long); this orchid is pollinated by a moth with a proboscis of that spur length, a fact that Charles Darwin predicted prior to the discovery of the moth. See Cameron et al. (1999), Cameron and Chase (2000), and Cameron (2004) for recent phylogenetic analyses of the orchids.

The Orchidaceae are distinctive in consisting of *mycorrhizal*, mostly perennial, *terrestrial or epiphytic* herbs having trimerous, often *resupinate* flowers with a showy **labellum**, the **androecium and gynoecium adnate** (termed a **column**, **gynostegium**, or **gynostemium**), the pollen grains often fused into 1–several masses (**pollinia**), bearing a sticky-tipped stalk, pollinia and stalk termed a **pollinarium**, which is the unit of pollen dispersal during pollination.

P (3+3) **A** 1–3, when 1 a pollinarium **G** (3), inferior, with gynostemium.

Themidaceae The Brodiaea family. ca. 12 genera / ca. ~60 species. (Figure 7.37)

The Themidaceae consist of perennial herbs. The **stems** are corms, typically with a membranous to fibrous covering from previous leaf bases, termed a **tunica**. **Leaves** are simple, closed-sheathing, flat, terete, or fistulose, acicular, linear, or lanceolate in outline. The **inflorescence** consists of a terminal scapose umbel. **Flowers** are bisexual, actinomorphic, and hypogynous. The **perianth** is biseriate and homochlamydeous, tepals 3+3, connate below or distinct. **Stamens** are 6 (3+3) or 3 (3 outer staminodes + 3 fertile, or 3 fertile in the position of the inner whorl), whorled, diplostemonous or antipetalous, usually distinct. The **gynoecium** is syncarpous; the ovary is superior, with 3 carpels, 3 locules, and 1 terminal style. **Placentation** is axile with 2 many ovules per carpel. The **fruit** is a loculicidal capsule. **Seeds** are ovoid, ellipsoid, or subglobose, endospermous, rich in oils and aleurone. An onionlike (alliaceous) odor is absent.

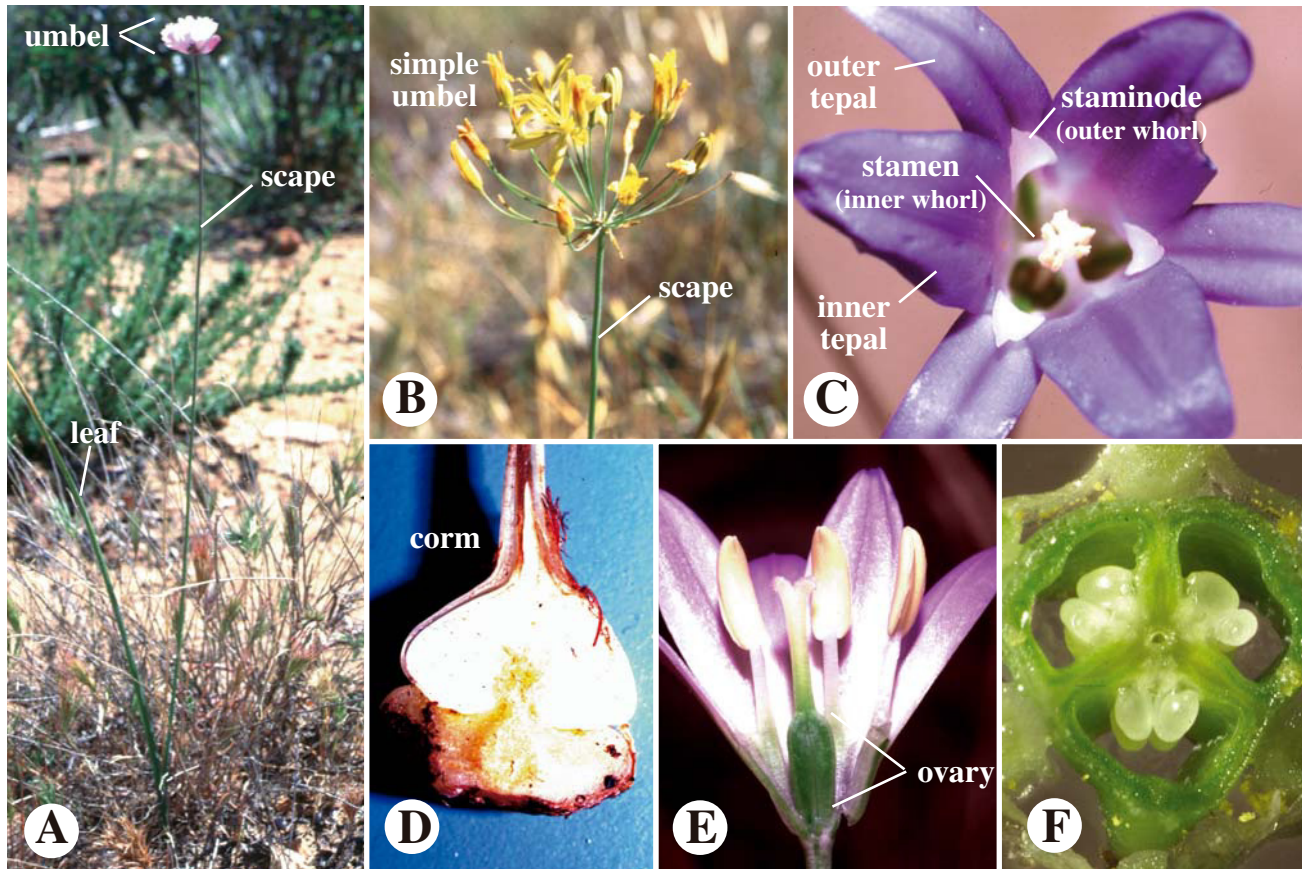


FIGURE 7.37 ASPARAGALES. Themidaceae. **A.** *Dichelostemma capitatum*, showing basal leaves, scape, and simple umbel. **B.** *Bloomeria crocea*, showing close-up of umbel with scape. **C.** *Brodiaea elegans*, flower close-up, showing biseriate, homochlamydeous perianth, three central, fertile stamens, and three staminodes. **D.** *Dichelostemma capitatum*, corm in longitudinal section. **E,F.** *Brodiaea orcutii*. **E.** Flower close-up in longitudinal section, showing three fertile stamens and superior ovary. **F.** Ovary cross-section, showing axile placentation.

Members of the Themidaceae are distributed in North America from S.W. Canada to Central America. There are no economic uses other than a few being used in cultivation. See Fay and Chase (1996) regarding the resurrection of the Themidaceae, Rahn (1998b) for detailed information on the family, and Pires and Sytsma (2002) for a phylogenetic analysis.

The Themidaceae are distinctive in being perennial, *cormose herbs, lacking an onionlike odor*, and having an *umbellate* inflorescence.

P 3+3 **A** 3+3, 3+3 staminodes, or 0+3 **G** (3), superior.

DIOSCOREALES

This order contains three families in APG II (2003): Burmanniaceae, Dioscoreaceae, and Nartheciaceae (Table 7.2). Only the Dioscoreaceae (united in APG II with the Taccaceae

and Trichopodaceae) are described here. See Caddick et al. (2002a,b) for a recent cladistic analysis of the group.

Dioscoreaceae Yam family (after Dioscorides, Greek herbalist and physician of 1st century A.D.). 4 genera / 300+ species. (Figure 7.38)

The Dioscoreaceae consist of dioecious or hermaphroditic, perennial herbs. The **stems** are rhizomatous or tuberous, often with climbing aerial stems, secondary growth present in some taxa. The **leaves** are spiral, opposite, or whorled, petiolate (typically with a pulvinus at proximal and distal ends), simple to palmate, undivided to palmately lobed, stipulate or not, with parallel or often net (reticulate) venation, the primary veins arising from the leaf base. The **inflorescence** is an axillary panicle, raceme, umbel, or spike of monochasial units (reduced to single flowers), with prominent involucre bracts in *Tacca*. The **flowers** are bisexual or unisexual, actinomorphic, pedicellate, bracteate or not, and epigynous.

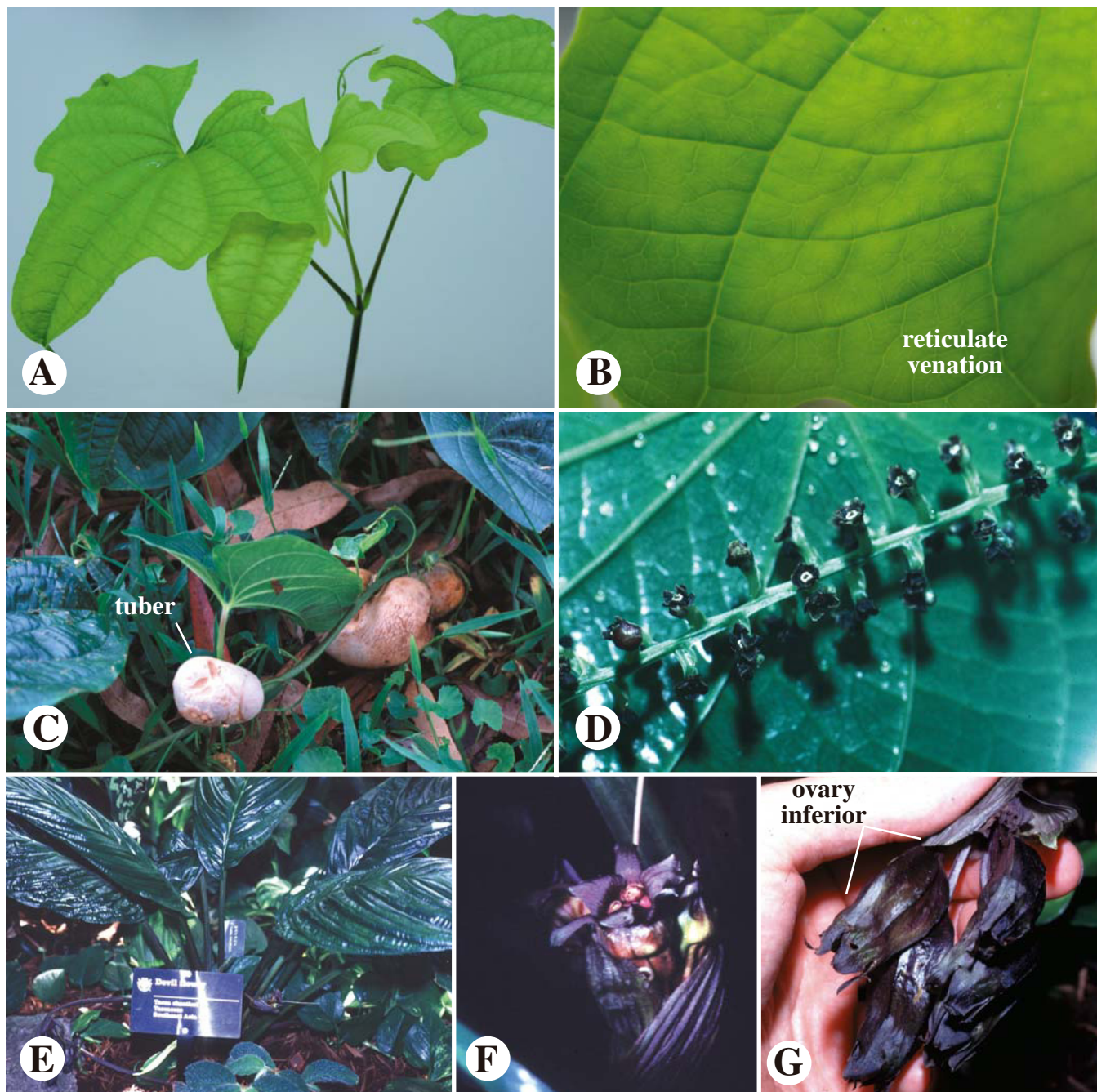


FIGURE 7.38 DIOSCORALES. **A,B.** Dioscoreaceae, yam family. *Dioscorea* spp. **A.** Shoot. **B.** Leaf surface close-up, showing reticulate venation. **C.** Tubers. **D.** Raceme of male flowers. **E–G.** *Tacca chantieri*. **E.** Whole plant; note net-veined leaves. **F.** Flower close-up. **G.** Fruits.

The **perianth** is biseriata, homochlamydeous, 3+3, a hypanthium absent or present. The **stamens** are 3+3 or 3+0, whorled, diplostemonous or antisepalous, distinct or monadelphous, free or epitepalous. **Anthers** are longitudinal and introrse or extrorse in dehiscence, tetrasporangiate, dithecal. The **gynoecium** is syncarpous, with a inferior ovary, 3 carpels,

and 3 locules. The **style(s)** are 3 or 1 and terminal; **stigmas** are 3. **Placentation** is axile or parietal; **ovules** are 1 2 [∞] per carpel. The **fruit** is a capsule or berry, often winged, 1 3 locular at maturity. **Seeds** are exalbuminous.

Members of the Dioscoreaceae have a mostly pantropical distribution. The family as most recently circumscribed

contains 4 genera: *Dioscorea*, *Stenomeris*, *Tacca* (previously classified in Taccaceae), and *Trichopus* (sometimes classified in Trichopodaceae). Several segregate genera have been merged into *Dioscorea* (Caddick et al. 2002). Economic importance includes various species of *Dioscorea*, the true yam, which are very important food sources in many tropical regions and which are also a source of steroidal saponins, used pharmaceutically in semisynthetic corticosteroid and sex hormones (especially birth control products) and used indigenously as a poison or soap. See Caddick et al. (2002) and Huber (1998a,b).

The Dioscoreaceae are distinctive in being perennial, **hermaphroditic or dioecious**, **rhizomatous or tuberous** herbs with **simple to palmate leaves** having **net venation** and **epigynous**, trimerous flowers.

P 3+3 **A** 3+3 or 3+0 **G** (3), inferior, hypanthium absent or present.

LILIALES

The Liliales is a fairly large group of monocotyledons that include 10 families (Table 7.2; Figure 7.39). As with the Asparagales, family delimitations of the Liliales have undergone a number of changes in recent years. Only the Liliaceae is described here. Notable among the other families are the **Alstroemeriaceae** (Figure 7.39A–D), *Alstroemeria* being a commonly cultivated ornamental, having interesting resupinate leaves; **Colchicaceae** (Figure 7.39E), containing *Colchicum autumnale*, autumn-crocus, source of colchicine used medicinally (e.g., formerly to treat gout) and in plant breeding (inducing chromosome doubling); **Melanthiaceae** (Figure 7.39G–I); **Philesiaceae** (Figure 7.39F); and **Smilacaceae** (Figure 7.39J), including *Smilax*, the green-briers, species of which are of economic importance as the source of sarsaparilla. See Rudall et al. (2000) for a phylogenetic analysis of the order.

Liliaceae [including Calochortaceae] Lily family (after *Lilium*, a name used in Virgil's writings). ca. 16 genera / ca. 600 species. (Figure 7.40)

The Liliaceae consist of perennial herbs. The **roots** are typically contractile. The **stems** are usually bulbous, rhizomatous in some. The **leaves** are basal or cauline, spiral or (in *Lilium* and *Fritillaria* spp.) whorled, usually sheathing, rarely petiolate, simple, and parallel veined [rarely net-veined]. The **inflorescence** is a terminal raceme, of a solitary flower, or rarely an umbel. The **flowers** are bisexual, actinomorphic or zygomorphic, pedicellate, bracteate or not, hypogynous. The **perianth** is biseriate and 3+3, homochlamydeous or

dichlamydeous, apotepalous, perianth parts sometimes spotted or striate. The **stamens** are 3+3, whorled, diplostemonous, distinct and free. **Anthers** are peltately attached to the filament or pseudo-basifixed (the filament tip surrounded by but not adnate to connective tissue), and longitudinally dehiscent. The **gynoecium** is syncarpous, with a superior ovary, 3 carpels, and 3 locules. The **style** is solitary; **stigmas** are 3, trilobed or with 3 crests. **Placentation** is axile.

Perigonal nectaries are present, at the tepal bases. The female gametophyte is the tetrasporic, *Fritillaria* type. The **fruit** is a loculicidal capsule. The **seeds** are flat and discoid or ellipsoid, the endosperm with aleurone and fatty oils, but no starch. Raphide crystals and chelidonic acid are lacking. Allyl sulfide compounds are absent.

The Liliaceae in the past has been treated as a large assemblage (Liliaceae sensu lato), which has more recently been broken up into numerous segregate families. Members of the family grow in mostly steppes and mountain meadows of the northern hemisphere, with the center of diversity in S.W. Asia to China. Economic importance includes several taxa of value as ornamental cultivars, including lilies, *Lilium*, and tulips, *Tulipa*. See Hayashi and Kawano (2000), Patterson and Givnish (2002), and Tamura (1998a,b).

The Liliaceae are characterized in being **perennial**, **usually bulbous herbs**, **lacking an onion-like odor**, with basal or cauline leaves, the inflorescence a **raceme**, **umbel or of solitary flowers** with a **superior** ovary.

P 3+3 **A** 3 **G** (3), superior.

PANDANALES

This order contains five families in APG II (2003), only one of which is described here. Notable among the other four is the Cyclanthaceae, containing *Carludovica palmata*, source of fiber, e.g., for Panama hats. See general references for more information on the order.

Pandanaceae Screw-Pine family (after Pandanus, a Malayan name for screw-pines). 3 genera / ca. 900 species. (Figure 7.41)

The Pandanaceae consist of perennial, dioecious, woody trees, shrubs, or vines. The adventitious **roots** are often branched, prop roots. The **stems** are sympodially branched, with prominent, encircling leaf scars. The **leaves** are acrocaulis, 3- or 4-ranked, appearing spiral because of twisting of the stem, sheathing, simple, undivided, linear to ensiform, parallel veined, the margin and adaxial midrib typically with prickles. The **inflorescence** is a terminal, rarely axillary, panicle, spike, or raceme or a pseudo-umbel of spikes or spadices subtended

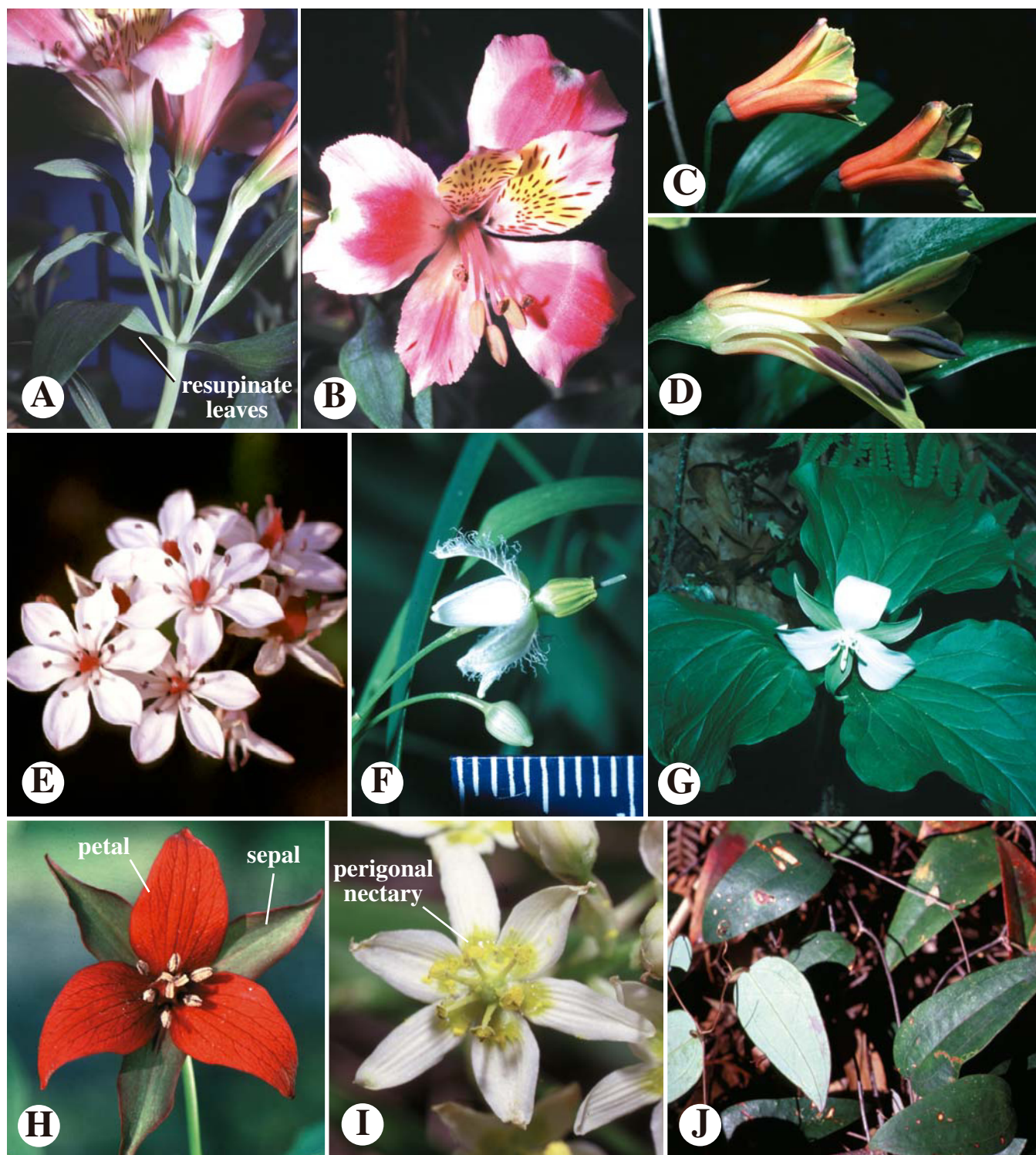


FIGURE 7.39 LILIALES, exemplars. A–D. Alstroemeriaceae. A,B. *Alstroemeria* sp. C,D. *Bomarea* sp. E. Colchiaceae, *Burchardia umbellata*. F. Philesiaceae, *Geitonoplesium* sp. G–I. Melanthiaceae. G. *Trillium grandiflorum*. H. *Trillium erectum*. I. *Zigadenus fremontii*. J. Smilacaceae, *Smilax glyciphylla*.

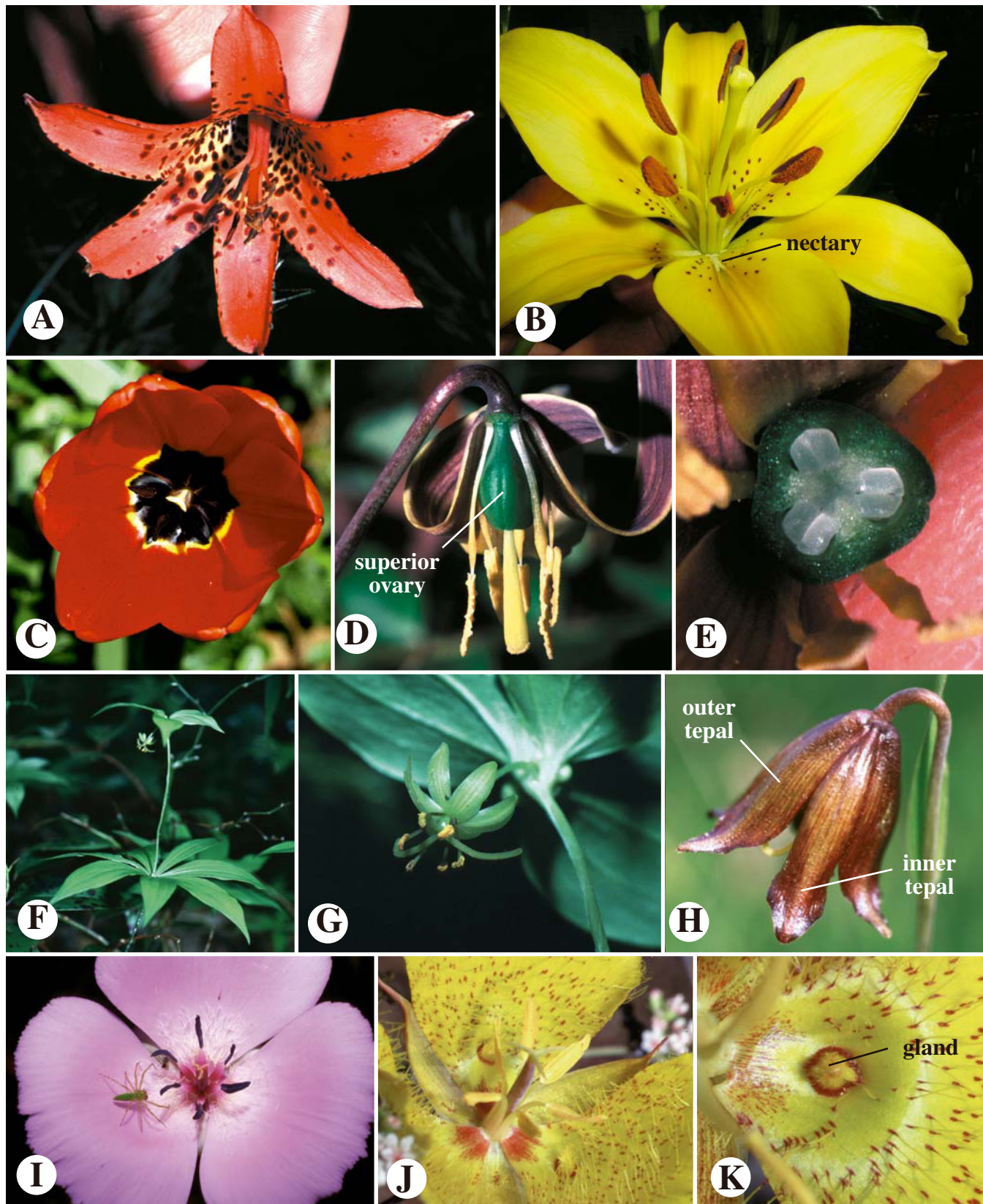


FIGURE 7.40 LILIALES. Liliaceae, Lily family. **A.** *Lilium canadense*, showing pendant ovary. **B.** *Lilium* sp., with erect ovary. Note nectary at base of tepal. **C.** *Tulipa* sp., tulip. **D,E.** *Erythronium americanum*, trout lily. **D.** Flower close-up, some tepals removed to show parts. **E.** Ovary cross-section, showing three carpels and locules. **F,G.** *Medeola virginiana*, Indian cucumber-root. **F.** Whole plant, with whorled leaves. **G.** Close-up of ovary. **H.** *Fritillaria bi-ora*, chocolate lily, ovary. **I.** *Calochortus splendens*. **J,K.** *Calochortus weedii*. **J.** Whole ovary. **K.** Close-up of perigonal gland (at base of inner tepal).

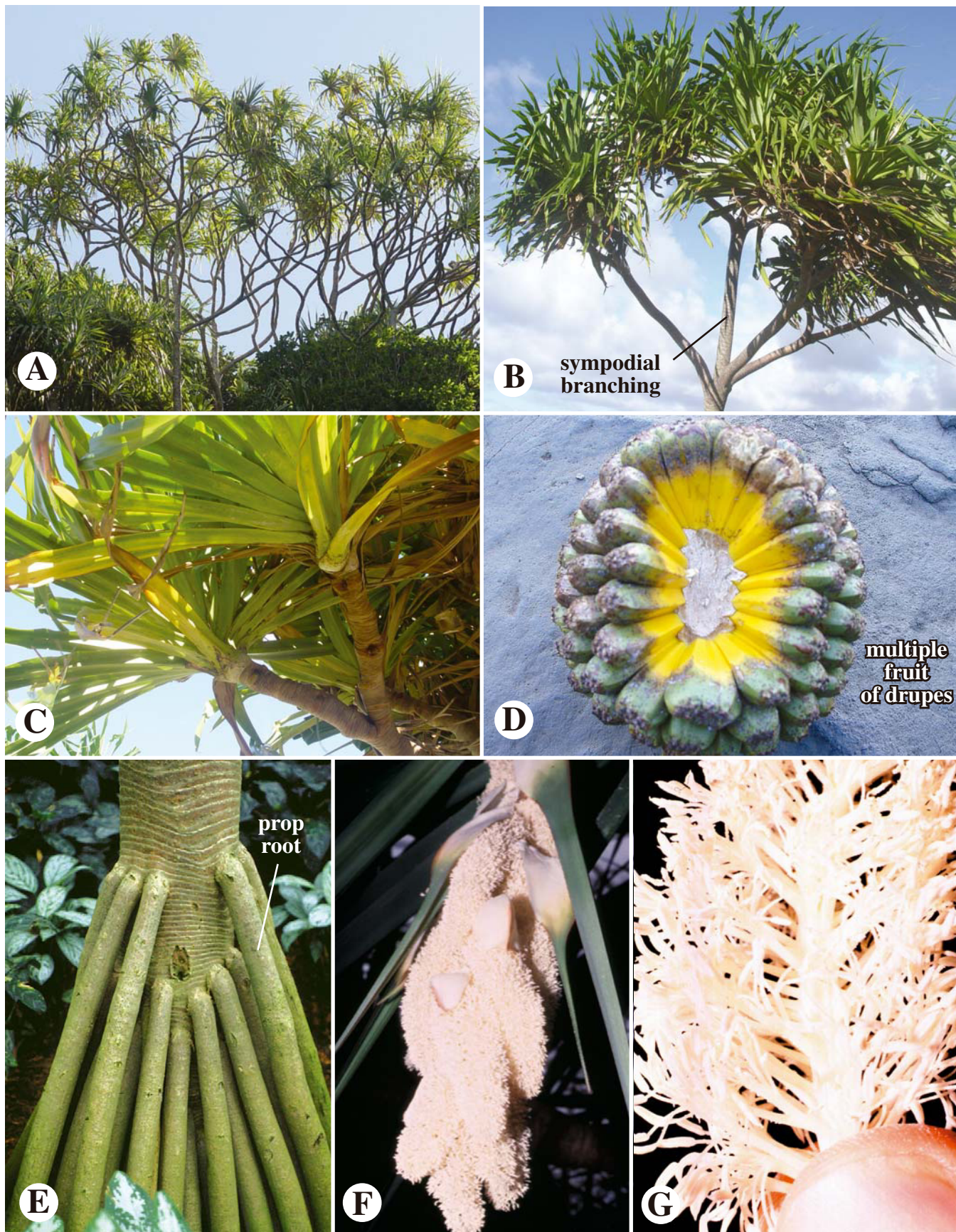


FIGURE 7.41 PANDANALES. Pandanceae, Screw-pine family. *Pandanus* sp. A–C. Whole plant, showing acrocaulis, narrow leaves. D. Fruit, a multiple fruit of berries. E. Base of stem with prop roots. F. Male inflorescence. G. Male flowers, close-up.

by spathes. The **flowers** are minute, usually unisexual, often with pistillodes or staminodes present, pedicellate, bracteate, hypogynous. The **perianth** is absent or an obscure 3-4-lobed, cuplike structure. The **stamens** are ∞ ; filaments are fleshy. The **gynoecium** is syncarpous, with a superior ovary and 1 ∞ carpels and locules. **Ovules** are anatropous, bitegmic, 1 ∞ . The **fruit** is a berry or drupe, forming multiple fruits in some taxa.

Members of the Pandanaceae are distributed from western Africa east to the Pacific islands. Economic importance includes use as ornamentals in some taxa and uses by indigenous people for thatch (for roofing), weaving, fiber, food (fruits and stems), spices, and perfumes. See Cox et al. (1995) and Stone et al. (1998) for more information on the family.

The Pandanaceae are distinctive in being mostly *dioecious*, *sympodially branched*, woody plants with *prop roots*, *3- or 4-ranked*, simple, *acrocaulis*, linear to ensiform leaves (appearing spiral), and small, usually *unisexual* flowers of variable morphology, the fruit a *berry or drupe*, multiple in some. **P** (3-4) or 0 **A** ∞ (male) **G** 1(- ∞) (female), superior.

COMMELINIDS

The Commelinids (also called the Commelinoids) are a monophyletic assemblage of monocots, as evidenced by morphological and molecular data (Figure 7.42). The Commelinids are characterized by an apparent chemical apomorphy, the presence of a class of organic acids (including coumaric, diferulic, and ferulic acid) that impregnate the cell walls. These acids can be identified microscopically in being UV-fluorescent (Figure 7.43). The orders and families of the Commelinids (after APG II, 2003) are listed in Table 7.3).

The Commelinids include a number of economically important plants, including the palms (Arecaceae), gingers and bananas (Zingiberales), and grasses (Poaceae). The grass family in particular is perhaps the most important family of plants, as grasses include the grain crops. As can be seen from Figure 7.42, the Dasygongonaceae and Arecaceae (palm family) are likely the most basal members of the Commelinid monocots. See Givnish et al. (1999), Chase et al. (2000a), and Davis et al. (2004) for recent analyses.

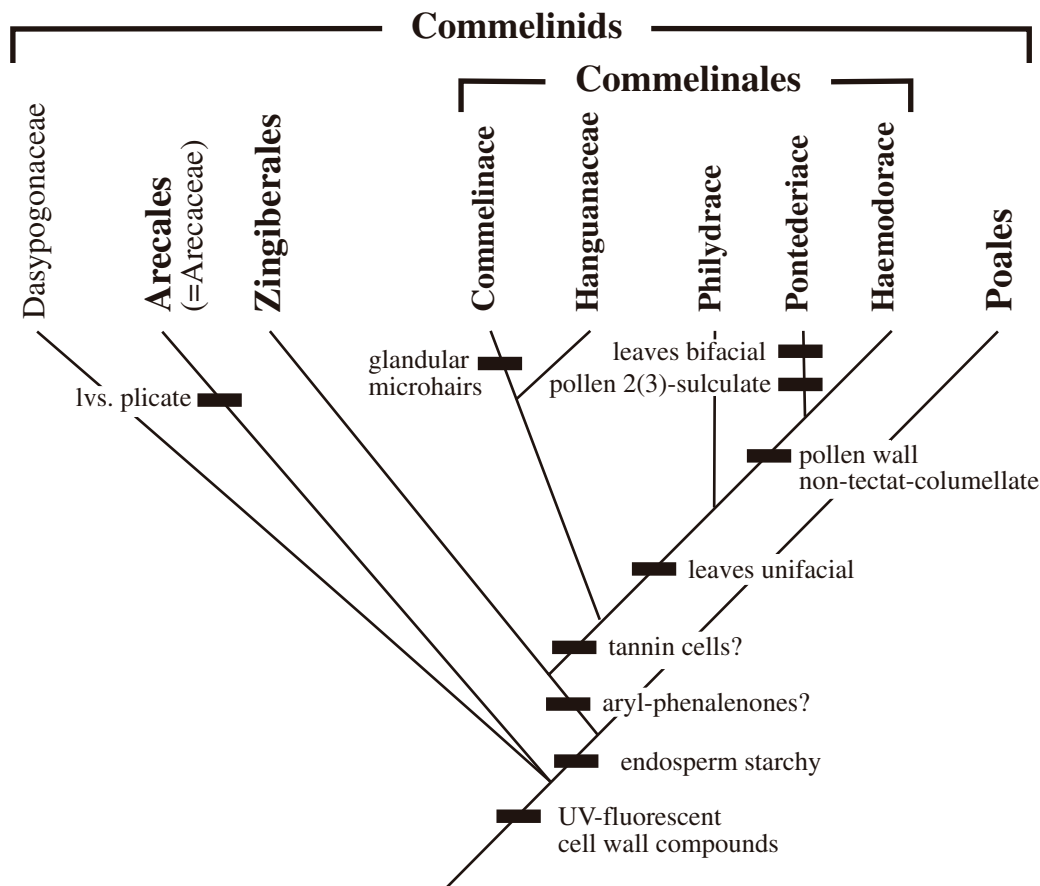


FIGURE 7.42 Major clades of the commelinid monocotyledons, modified from APG II (2003) with selected apomorphies shown.

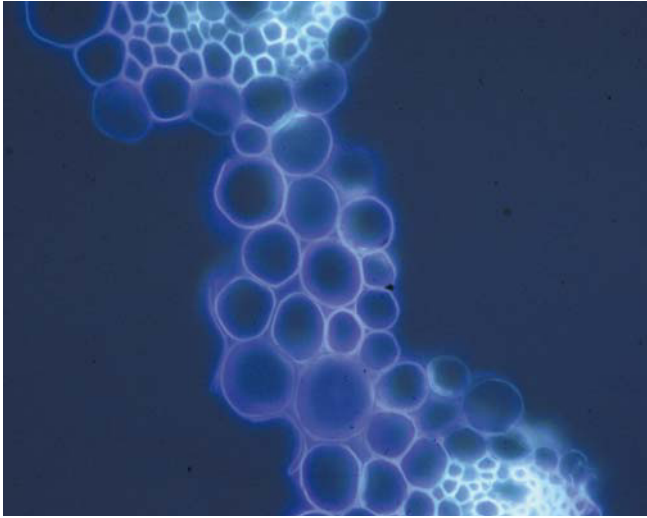


FIGURE 7.43 Leaf cross-section of *Lachnanthes caroliniana* (Haemodoraceae), showing the UV fluorescence of nonlignified cell walls (center). This fluorescence is indicative of the presence of certain organic acids, apomorphic for the Commelinid monocots.

ARECALES

This order contains the single family Arecaceae. See Dransfield and Uhl (1998), Asmussen et al. (2000), Hahn (2002), and Lewis and Doyle (2001) for information and phylogenetic analyses of the palms.

Arecaceae (Palmae) Palm family (from *areca*, Portuguese for the betel palm). ca. 190 genera / ca. 2000 species. (Figures 7.44, 7.45)

The Arecaceae consist of perennial trees, large rhizomatous herbs, or lianas. Plant sex is variable, and secondary growth is absent. The **roots** are mycorrhizal, lacking root hairs.

The **stem** is usually arborescent, consisting of a single, unbranched trunk [dichotomously branched in *Hyphaene*], or a caespitose cluster of erect stems, or a stout, dichotomously branched rhizome (*Nypa*), or an elongate liana with long internodes (rattan palms). The **leaves** are typically quite large, generally terminal (acrocaulis), spiral [rarely distichous or tristichous], with a sheathing base and an elongate, stout petiole (sometimes referred to as pseudopetiole) between the sheath apex and blade. In arborescent taxa the sheathing bases of adjacent leaves may overlap one another, forming a distinctive crownshaft at the trunk apex. Leaves are simple, pinnate, bipinnate, costapalmate, or palmate; if simple, the leaves are often pinnately or palmately divided, sometimes bifid, with leaflet spines present in some taxa. Leaves are typically ligulate (with an appendage, the ligule, at the inner junction of blade and petiole); in taxa with palmate leaves, another distinctive process, called the hastula, may be present at the junction of the petiole and blade. The leaf blade is characteristically plicate (pleated), with the leaflets or blade divisions in cross-section either induplicate (V-shaped, with the point of the fold below, or abaxial) or reduplicate (Λ-shaped, with the point of the fold above, or adaxial). Venation is pinnate- or palmate-parallel. The **inflorescence** is typically an axillary, bracteate panicle or spike of solitary flowers or of cyme units, the inflorescence arising either below (infrafoliar) or among (interfoliar) or above (suprafoliar) the leaves of the crownshaft. The peduncle is subtended by an often large prophyll and 1 ∞ spathes. The **flowers** are unisexual or bisexual, actinomorphic, sessile, and hypogynous. The **perianth** is usually biseriate and homochlamydeous, 3+3 [0, 2+2, or ∞], apotepalous. The **stamens** are 3+3 [3 or ∞], distinct or connate, epitepalous in some spp., staminodes present in some spp. **Anthers** are longitudinal, rarely poricidal, in dehiscence. The **gynoecium** is syncarpous or

TABLE 7.3 Orders and included families of the Commelinid Monocotyledons, after APG II (2003). Families in **bold** are described in detail.

COMMELINIDS			
Dasygongonaceae	ZINGIBERALES	POALES	POALES (continued)
ARECALES	Cannaceae	Anarthriaceae	Poaceae
Arecaceae (Palmae)	Costaceae	Bromeliaceae	(Gramineae)
COMMELINALES	Heliconiaceae	Centrolepidaceae	Rapateaceae
Commelinaceae	Lowiaceae	Cyperaceae	Restionaceae
Haemodoraceae	Marantaceae	Ecdeiocoleaceae	Sparganiaceae
Hanguanaceae	Musaceae	Eriocaulaceae	Thurniaceae
Philydraceae	Strelitziaceae	Flagellariaceae	(including Prioniaceae)
Pontederiaceae	Zingiberaceae	Hydatellaceae	Typhaceae
		Joinvilleaceae	Xyridaceae
		Juncaceae	
		Mayacaceae	

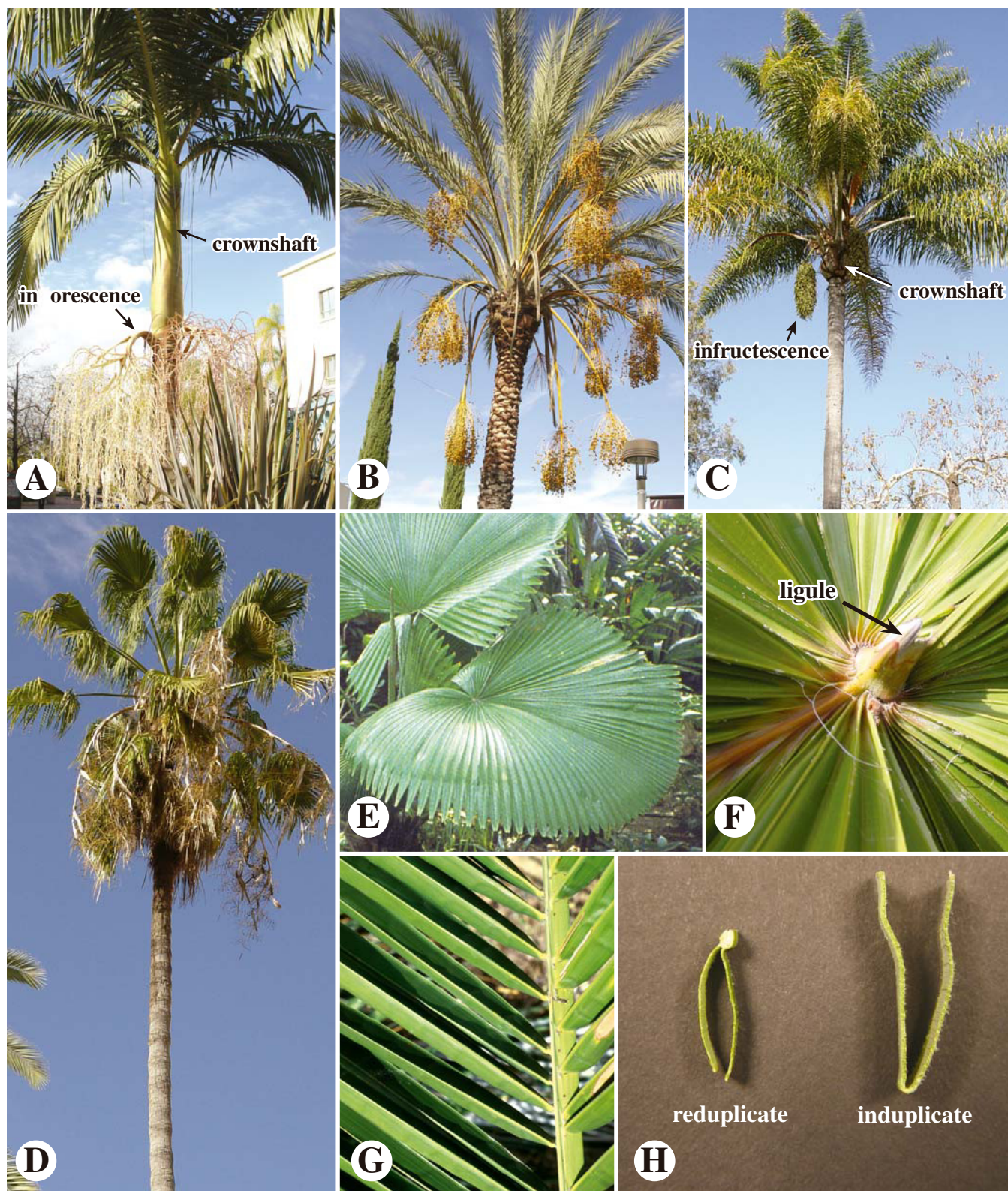


FIGURE 7.44 ARECALES. Areaceae. **A.** *Archontophoenix cunninghamiana*, king palm, showing single, unbranched trunk with acrocaulis crown of pinnately compound leaves and lateral in orescences below crownshaft (intrafoliar). **B.** *Phoenix dactylifera*, date palm, with several in orescences arising within crownshaft (interfoliar). **C.** *Syagrus romanzof ana*, queen palm, with pinnate leaves. **D.** *Washingtonia robusta*, with palmately divided leaves. **E.** *Licuala peltata*, with palmately lobed leaves. **F.** *Livistona drudei* leaf close-up, showing ligula at junction of petiole and blade. **G.** *Jubaea chilensis* leaf close-up, showing plicate posture of pinnate leaves. **H.** Reduplicate (*Syagrus romanzof ana*) and induplicate (*Phoenix dactylifera*) leaf posture. Adaxial side of leaf blade is at top.

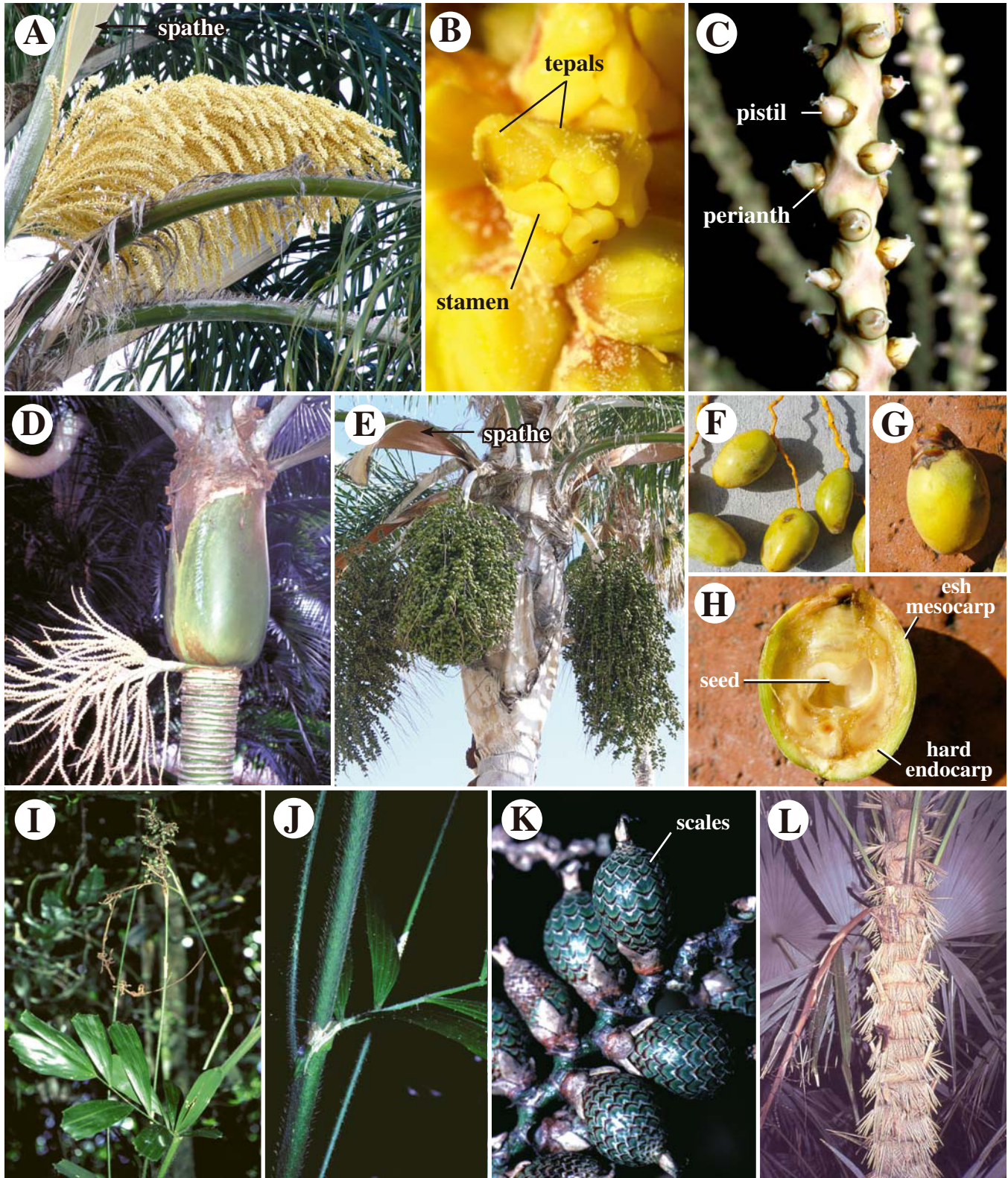


FIGURE 7.45 ARECALES. Arcaceae. **A.** *Syagrus romanzof ana*, queen palm, in orescence, with spathe. **B.** *Chamaerops humilis*, Mediterranean palm. Close-up of flowers, showing trimerous perianth and androecium. **C, D.** *Rhopalostylis sapida*. **C.** Close-up of female flowers, with reduced, scalelike perianth and superior, 3-carpeled ovary. **D.** Close-up of sheathing leaf bases forming crownshaft; note lateral, infrapetiolar in orescence. **E.** *Syagrus romanzof ana*, infructescences, with spathes. **F.** *Phoenix dactylifera*, date palm, drupes. **G, H.** *Syagrus romanzof ana*, drupes. **G.** Whole fruit. **H.** Drupe longitudinal section, showing pericarp layers (hard endocarp and fleshy mesocarp). **I–K.** *Calamus* sp. (rattan palm). **I.** Whole plant, showing pinnate leaf. **J.** Leaf base close-up; note sheath and long internodes. **K.** Fruit (drupe) close-up, showing retrorse scales typical of the rattan palms. **L.** *Zombia antillarum*, a palm with numerous spines.

apocarpous, with a superior ovary, usually 3 [1, 2, 4 ∞] carpels, and 3 or 1 [∞] locules. The styles, if present, are distinct or connate; stigmas are sessile or at tip of styles. **Placentation** is variable; ovules are variable in type, bitegmic, and 1 per locule. Septal nectaries are present in some taxa. The **fruit** is fleshy or fibrous, usually a drupe, rarely dehiscent, some with outer scales (Calamoideae), hairs, prickles, or other processes. **Seeds** are usually 1 per fruit and have an oil or hemicellulose-rich, sometimes ruminant, endosperm; starch is absent.

The Arecaceae have distributions mostly in warm, tropical regions and are often ecologically important where they occur. The family has recently classified into several subfamilies, including the Nypoideae, Calamoideae, Arecoideae, Coryphoideae, Phytelphantoideae, and Ceroxyloideae, although some of these are likely paraphyletic. The rhizomatous Nypoideae (*Nypa*) are thought to be the most basal palm lineage, with the lianous Calamoideae (*Calamus* and relatives) possibly next basal. Thus, it is likely that the arborescent habit that we typically associate with palms evolved from a rhizomatous habit within the family. The plicate leaf is a probable apomorphy for the family as a whole, as is the drupaceous fruit. (The family Cyclanthaceae and a few other scattered monocot taxa also have plicate leaves, but these are thought to have evolved independently.) The palms are of great economic importance, including uses as fruits (e.g., *Cocos nucifera*, coconut palm, *Phoenix dactylifera*, date palm), furniture/canes (rattan palms), fibers (e.g., coir from the mesocarp of *Cocos nucifera*), oils (e.g., *Elaeis oleifera*, oil palm), starch (e.g., *Metroxylon* spp., sago palms), waxes (e.g., *Copernicia cerifera*, wax palm), and many species used indigenously as timber or in building construction; fruits of *Areca catechu*, betel palm, are chewed in India (with *Piper betle* leaves and lime) as a stimulant.

The Arecaceae are distinctive in having a rhizomatous, lianous, or usually **arborescent** stem, with large, sheathing, **plicate** leaves, a **fleshy, usually drupaceous** fruit, and seeds lacking starch. The plicate leaf posture and drupaceous fruit are likely apomorphies for the family.

P 3+3 [0,2+2,∞] **A** 3+3 or (3+3) [3,∞; 0 in female fls.] **G** 3 or (3) [1,2,4-∞; 0 in male fls.], superior.

COMMELINALES, ZINGIBERALES, AND POALES

The taxa of the Commelinid monocots other than the Dasypogonaceae and Arecaceae are classified into the three orders Zingiberales, Commelinales, and Poales. All of these have seeds that contain endosperm rich in starch, an apparent apomorphy for the three orders (Figure 7.42). In contrast, the

palms have seeds rich in oils and hemicellulose and lacking in starch.

The Commelinales and Zingiberales are sister taxa according to some phylogenetic analyses (see general angiosperm phylogeny studies; Chase et al. 2000a; Davis et al. 2004). A possible apomorphy uniting them is the presence of arylphenalenone chemical compounds (Figure 7.42), which are common in the Haemodoraceae and have been discovered also in some Pontederiaceae and Zingiberales.

COMMELINALES

The Commelinales, sensu APG II (2003) consist of five families, three of which are described here. One hypothesis of interrelationships and putative apomorphies is portrayed in Figure 7.42. The family Hanguanaceae has only recently been placed here. The Commelinales is not well defined morphologically, although floral tannin cells may constitute an apomorphy (Figure 7.42). The Haemodoraceae and Philydraceae have unifacial leaves, a likely apomorphy, but this would necessitate the reversal to bifacial leaves in the aquatic Pontederiaceae (Figure 7.42). The Haemodoraceae and Pontederiaceae may be united by the apomorphy of non-tectate-columellate pollen wall structure (Figure 7.42). See Givnish et al. (1999) and Davis et al. (2004) for recent phylogenetic analyses of the order.

Commelinaceae Spiderwort family (for Caspar Commelijn, Dutch botanist, 1667–1731). 39 genera/640 species. (Figure 7.46)

The Commelinaceae consist of mostly perennial herbs. The **stems** typically have swollen nodes. The **leaves** are spiral, sheathing (sheath closed), simple, undivided, with each half of the blade rolled adaxially toward the midrib early in development. The **inflorescence** is a cyme, rarely a raceme or of solitary flowers, the flowers often piercing the subtending bract. The **flowers** are usually bisexual, actinomorphic or zygomorphic, and hypogynous. The **perianth** is biseriate, usually dichlamydeous. The **calyx** consists of 3 distinct or basally fused sepals or lobes. The **corolla** contains 3, equal or unequal (anterior petal smaller), distinct or basally connate [sometimes clawed] petals or lobes, which are characteristically ephemeral. The **stamens** are usually 3+3, sometimes with 3 fertile and 3 staminodes [rarely of 1 fertile stamen], apostemonous, the filaments often with pilose trichomes, fertile stamens sometimes dimorphic. **Anthers** are basifixed, versatile, longitudinally dehiscent [rarely poricidal apically and basally], with the connective often extended; prominent, antherodes (sterile anthers) present on staminodes.

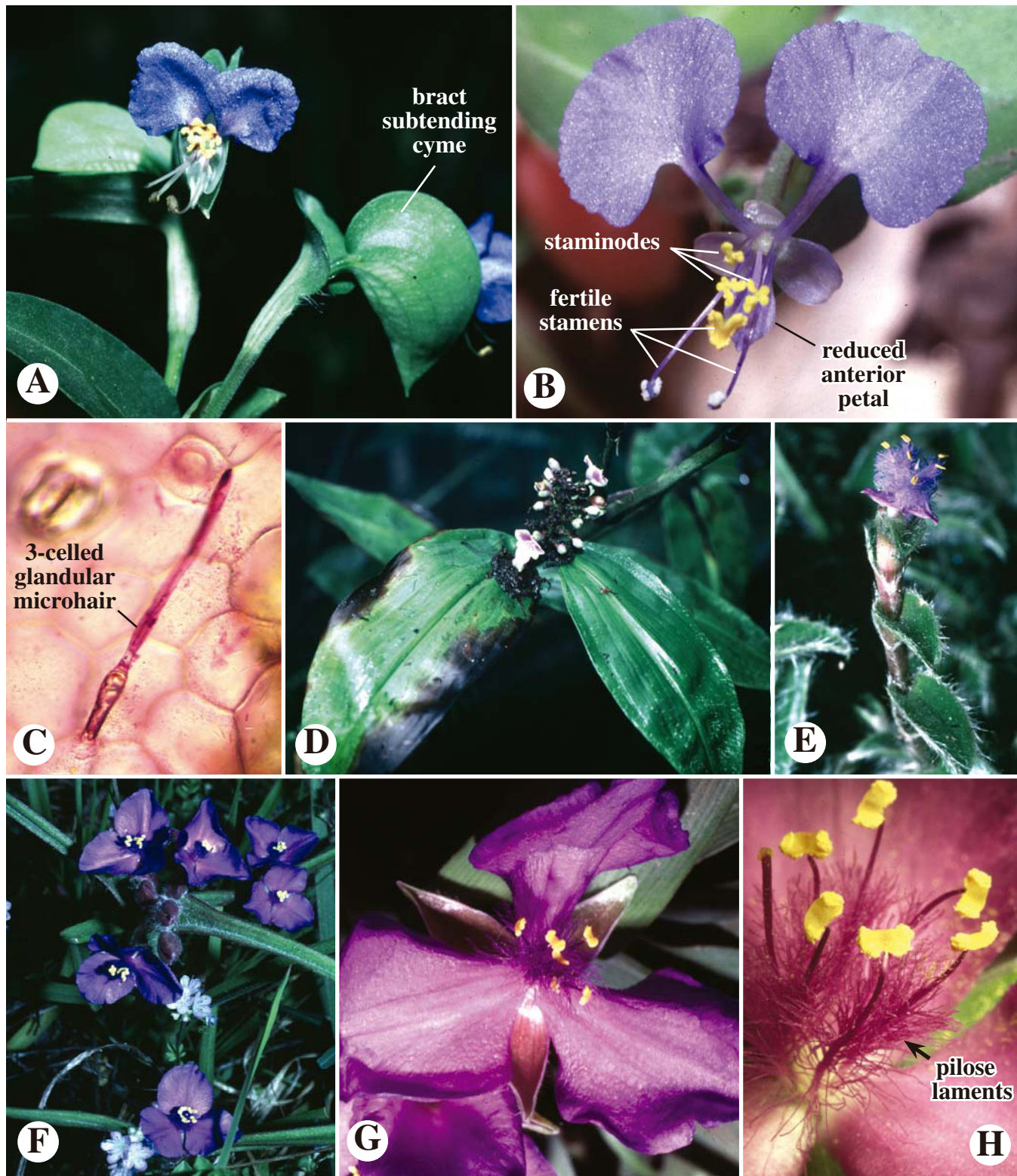


FIGURE 7.46 COMMELINALES. Commelinaceae. A–C. *Commelina* sp. A. In rosette apex, showing closed sheath of leaves and spathe-like bracts. B. Flower, face view. Note staminodes and fertile stamen dimorphism. C. Tricellular microhair, a putative apomorphy of the Commelinaceae. D. *Dichorisandra reginae*. E. *Cyanotis somaliensis*. F–H. *Tradescantia*. F. *T. hirsuta*. G. *T. virginiana*. H. *Tradescantia* sp. Close-up of androecium, showing pilose filaments.

The **gynoecium** is syncarpous, with a superior ovary, 3 carpels (the median carpel anterior), locules 3 or 1 at the apex only or 1 2 (the other locule(s) undeveloped or absent). **Placentation** is axile; **ovules** are orthotropous to anatropous, bitegmic, 1 ∞ in number. The **fruit** is a loculicidal capsule, rarely an indehiscent capsule or berry. The **seeds** are rarely winged or arillate, having a starchy endosperm. Plant surfaces typically bear 3-celled, glandular microhairs, a putative apomorphy for the family (Figure 7.42), and tissues often have raphide-containing mucilage cells.

Members of the Commelinaceae have distributions in most tropical to subtropical regions worldwide. Economic importance includes ornamental cultivars, such as *Rhoeo*, *Tradescantia*, and *Zebrina*, and some local medicinal and edible species. See Faden (1998) and Evans et al. (2003) for recent treatments and phylogenetic analyses of the family.

The Commelinaceae are distinctive in being mostly perennial **herbs** with **closed sheathed** leaves and a trimerous, hypogynous flower with an **ephermal** corolla, staminodia in some, most species with characteristic **3-celled glandular microhairs**, the latter a probable apomorphy for the family (Figure 7.42).

K 3 or (3) **C** 3 or (3) **A** 3 or 3 + 3 staminodes or 1 **G** (3), superior

Haemodoraceae Bloodwort family (Gr. *haimo*, blood, in reference to red pigmentation in roots and rootstocks of some members). 13 genera / 100 species. (Figures 7.47, 7.48)

The Haemodoraceae consist of perennial herbs. The **stems** are rhizomatous, stoloniferous or cormose. The **leaves** are simple, unifacial, mostly basal, distichous, sheathing and often equitant, undivided, narrow, flat or terete, and parallel veined. The **inflorescence** is a terminal thyrse or corymb of single or 2 3 branched monochasial cyme units, a simple raceme, or rarely reduced to a single flower. The **flowers** are bisexual, actinomorphic or zygomorphic, pedicellate, bracteate, hypogynous, epigynous, or epiperigynous, glabrous to densely tomentose, trichomes tapering, pilate-glandular, or branched, often brightly colored; the receptacle is extended proximally in some taxa (e.g., *Wachendorfia*). The **perianth** is apotepalous or syntepalous, biseriate, imbricate, and homochlamydeous with 3 outer and 3 inner tepals (median outer tepal posterior in zygomorphic flowers) or syntepalous and uniseriate with 6 valvate tepal lobes; tepals are red, red-orange, yellow-orange, yellow, to green, white, or black in color, hypanthium present or absent. The **stamens** are 6, 3, or 1, whorled, diplostemonous or antitropalous in taxa with 6 stamens, or antitropalous in taxa with 3 or 1 stamens, unfused or epitepalous, with staminodes present in some taxa. **Anthers** are basifixed, longitudinal and introrse in dehiscence, tetrasporangiate,

dithecal, with thecae and connective having appendages in *Tribonanthes*. The **pollen** is monosulcate or 2 many porate and binucleate at release. The **gynoecium** is syncarpous, with an inferior or superior ovary, 3 carpels (the median carpel posterior), and 3 (rarely 1 at ovary apex) locules. The **style** and **stigma** are solitary, the latter often 3-lobed. **Placentation** is axile; **ovules** are anatropous, bitegmic, 1, 2, 5 7, or ∞ per carpel. Septal nectaries occur in most taxa. The **fruit** is a capsule or rarely a schizocarp. The **seeds** are globose, ellipsoid and ridged, or flattened and marginally winged, with starchy endosperm. Distinctive arylphenalenone chemicals are found in all investigated family members, comprising a reddish pigmentation in the roots and rhizomes of some taxa (hence the name Bloodwort).

The Haemodoraceae contain two monophyletic groups: Haemodoroideae [Haemodoreae], with unbranched, pilate or tapering trichomes, 3 (1) stamens, and monosulcate pollen, and Conostylidoideae [Conostylideae], with branched to dendritic trichomes, 6 stamens, and porate pollen. Members of the family grow in seasonally wet habitats with distributions in S.W. and E. Australia, New Guinea, S. South Africa, N. South America, Central America and S. Mexico, Cuba, or E. to S.E. North America. Economic importance includes ornamental cultivars, especially *Anigozanthos* spp. (kangaroo-paws), and historical uses by native people for food and as euphorics. See Simpson (1990, 1998) and Hopper et al. (1999) for recent treatments of the family.

The Haemodoraceae are distinctive in being perennial herbs with **arylphenalenone** compounds (imparting a reddish coloration to stems and roots in almost all Haemodoroideae), **unifacial** leaves, and variable flowers.

P 3+3 or (3+3) or (6) **A** 1,3,6 **G** (3), inferior or superior, hypanthium present or absent.

Pontederiaceae Pickerel-Weed family (for Buillo Pontedera, former Professor of Botany at Padua, 1688 1757). 7 genera / 31 species. (Figures 7.49, 7.50)

The Pontederiaceae consist of perennial or rarely annual, emergent or free-floating, aquatic herbs. The **stems** are rhizomatous or stoloniferous. The **leaves** are bifacial, ligulate, mostly basal, distichous or spiral, basally sheathing and petiolate (the petiole swollen in *Eichhornia*), simple, undivided, narrow to broad, flat, and parallel curved-convergent veined (filiform in *Hydrothrix*). The **inflorescence** is a terminal or axillary raceme, spike, thyrse, or of solitary flowers, with spathe-like bract present. The **flowers** are bisexual, zygomorphic or actinomorphic, hypogynous, glabrous or with scattered pilate-glandular trichomes on the outer perianth, filaments, or style. The **perianth** is biseriate and homochlamydeous with 3 [4] outer and 3 [4] inner imbricate tepals,

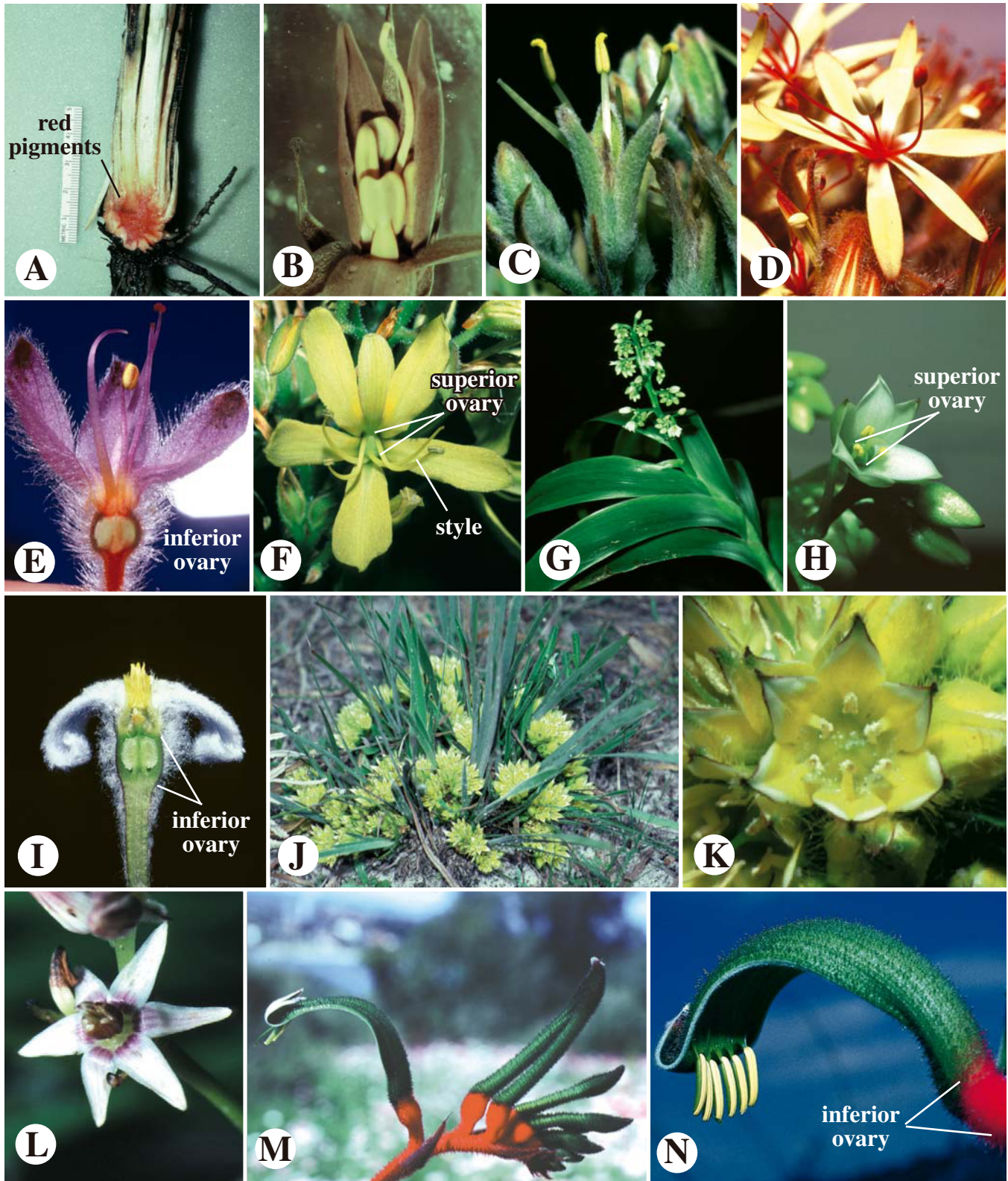


FIGURE 7.47 COMMELINALES. Haemodoraceae. **A.** *Haemodorum laxum*, showing cormlike rootstock with red pigmentation. **B.** *Haemodorum spicatum*, opened flower. **C.** *Lachnanthes caroliniana*. **D.** *Dilatris viscosa*, showing three, dimorphic stamens. **E.** *Dilatris corymbosa*, flower longitudinal section. **F.** *Wachendor a thyrsi* flower, showing zygomorphic, enantiostylous flower. **G.** *Xiphidium caeruleum*, showing unifacial, ensiform leaves. **H.** *Xiphidium caeruleum*, actinomorphic flower with three stamens and superior ovary. **I.** *Tribonanthes uni* flower, with inferior ovary. **J,K.** *Conostylis aurea*. **J.** Whole plant. **K.** Flower close-up, showing six stamens. **L.** *Phlebocarya ciliata*, with actinomorphic flowers. **M,N.** *Anigozanthos manglesii*. **M.** Inflorescence, the unit a monochasium. **N.** Zygomorphic flower with six stamens and inferior ovary.

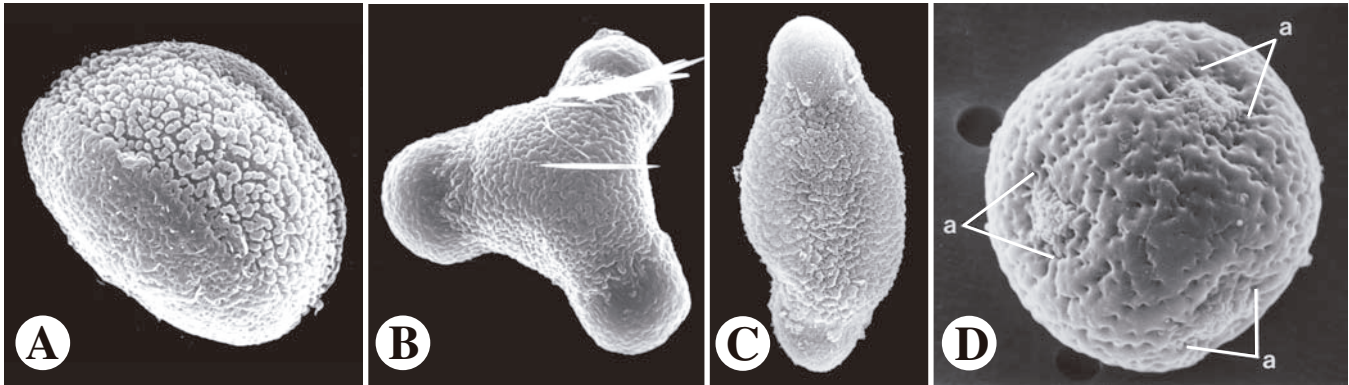


FIGURE 7.48 COMMELINALES. Haemodoraceae. Pollen diversity. **A.** Ellipsoid and monosulcate, *Schiekia orinocensis*. **B.** Triangular and triplicate, *Conostylis aurea*. **C.** Fusiform and diporate, *Macropidia fuliginosa*. **D.** Spherical and oligoporate, *Tribonanthes australis*.

with median inner tepal posterior in zygomorphic flowers; tepals basally connate, and blue, lilac, white, or yellow, with nectar guide on median tepal in zygomorphic flowers, hypanthium present. The **stamens** are six (3+3) or three (+ staminodes) or 1 (+ 2 staminodes), whorled, of different lengths in some taxa (often associated with tristylly), diplostemonous (in taxa with 6 stamens) or antipetalous (in taxa with 3 or 1 stamens), epitepalous, filaments with appendages in some taxa. **Anthers** are basifixed, introrse, longitudinal or (in *Monochoria*) poricidal in dehiscence, tetrasporangiate and dithecal. The **pollen** is di- (tri-) sulcate and trinucleate at release. The **gynoecium** is syncarpous, with a superior ovary, 3 carpels (2 carpels reduced and abortive in *Pontederia* and *Reussia*), and 1 to 3 locules. The **styles** are heteromorphic or enantiostylous in some taxa; the **stigma** is solitary, often 3-lobed. **Placentation** is apical, axile, parietal, or axile below and parietal above; **ovules** are anatropous, bitegmic, 1 ∞ per carpel. Septal nectaries are present or absent. The **fruit** is a loculicidal capsule or nut/utricle (e.g., *Pontederia*). The **seeds** are longitudinally ribbed, with a starchy endosperm.

Members of the Pontederiaceae have distributions in tropical to north temperate regions in Africa, Asia, and esp. the Americas. Economic importance includes species that are serious weeds (especially *Eichhornia crassipes*, water hyacinth, which clogs waterways), some species with edible parts, and cultivated ornamentals (e.g., *Eichhornia*, *Heteranthera*, and *Pontederia*). See Barrett and Graham (1997) and Cook (1998) for recent phylogenetic analyses and treatments of the family.

The Pontederiaceae are distinctive in being **emergent to free-floating aquatic herbs** with simple, sheathing, **bifacial** leaves, actinomorphic or zygomorphic flowers, and **di-(tri-)sulcate** pollen. The bifacial leaves and sulcate pollen are probable apomorphies for the family (Figure 7.42).

P (3+3) or (4+4) **A** 3+3 or 3+staminodes or 1+2 staminodes **G** (3), superior, hypanthium present.

ZINGIBERALES

The Zingiberales, commonly called the gingers and bananas, are a well-defined, monophyletic clade of eight families (Figure 7.51, 7.53), four of which are described here. (See Figure 7.59 for images of three of the other families.) Several apomorphies unite the ginger group. One obvious apomorphy is the occurrence of leaves with **penni-parallel venation** (Figures 7.52A, 7.54A). In addition, virtually all members of the Zingiberales have a **ptyxis** (the posture of immature leaves or leaf parts; see chapter 9) that is **supervolute**, in which the two opposing (left and right) halves of the blade are rolled along a longitudinal axis, one half being rolled completely within the other (Figure 7.52A,B). Leaves and stems of all members of the order have **diaphragmed air chambers** (Figure 7.52C) and possess **silica cells** (although the latter is not apomorphic for this order alone). Lastly, all Zingiberales have an **inferior ovary** (Figures 7.51, 7.56D).

The four terminal families of the Zingiberales make up a monophyletic assemblage, commonly known as the ginger group; their phylogenetic interrelationships are well accepted (Figure 7.51). The other four families, which are referred to as the bananas, constitute a paraphyletic assemblage whose precise interrelationships are not clear; the phylogeny shown in Figure 7.51 is one possibility. See Kress et al. (2001) for a recent treatment of this group.

Musaceae Banana family (after Antonia Musa, physician to Emperor Augustus 63–14 BC). 3 genera (*Ensete*, *Musa*, and *Musella*) / ca. 40 species. (Figure 7.54)

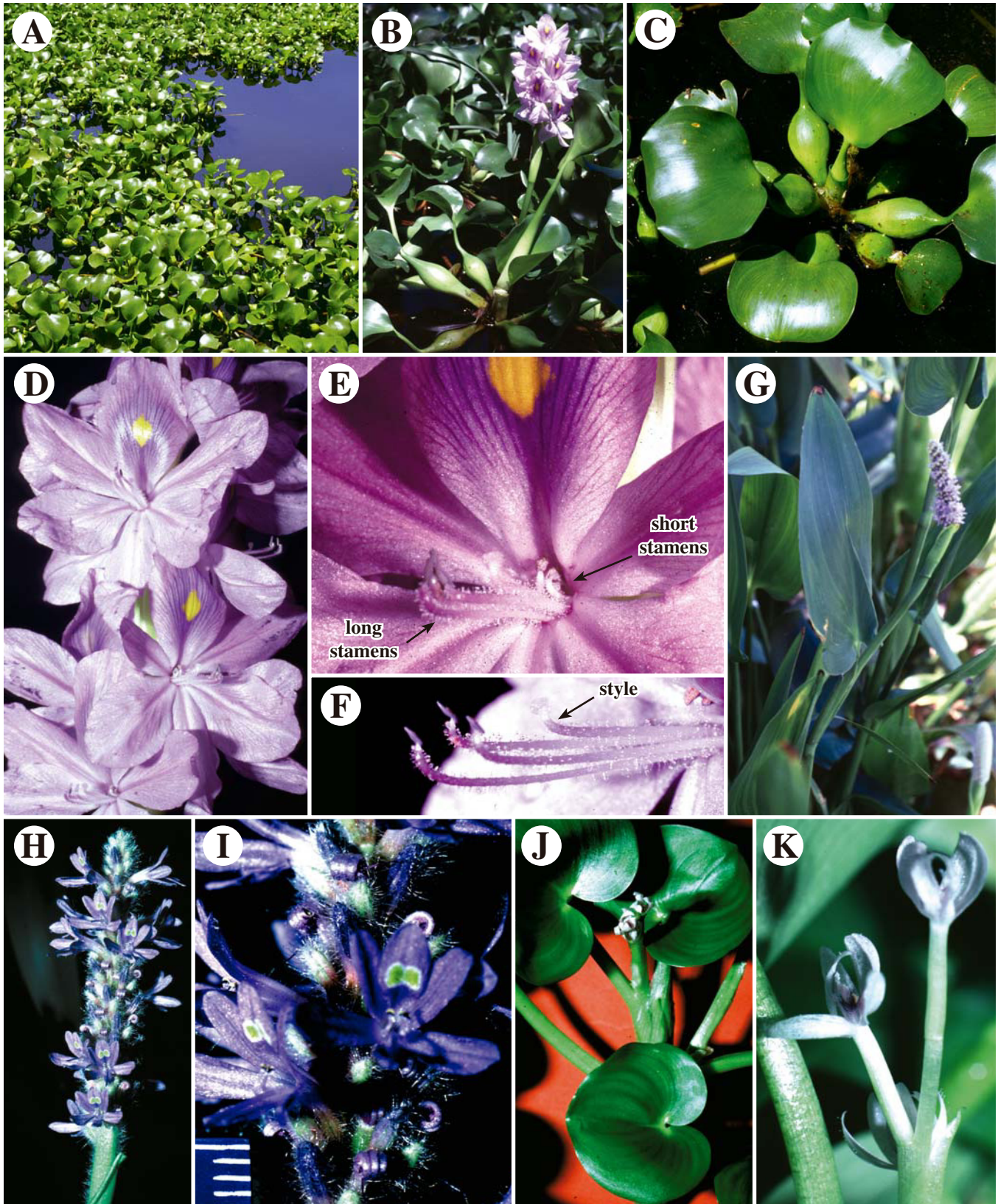


FIGURE 7.49 COMMELINALES. Pontederiaceae. A–F. *Eichhornia crassipes*. A. Habit, showing masses of floating clonal plants clogging lake. B. Plant with inflorescence. C. Close-up of leaves with swollen petioles. D. Inflorescence. E. Close-up of zygomorphic flower, showing median posterior inner tepal with nectar guide. Note dimorphic stamens: three long and three short. F. Flower close-up, showing three long stamens and style (above). G–I. *Pontederia cordata*. G. Whole plant, in flower. H. Inflorescence. I. Flower close-up, showing zygomorphic perianth with nectar guides. J, K. *Heteranthera reniformis*. J. Whole plant, with inflorescence. K. Close-up of tubular, actinomorphic flowers.

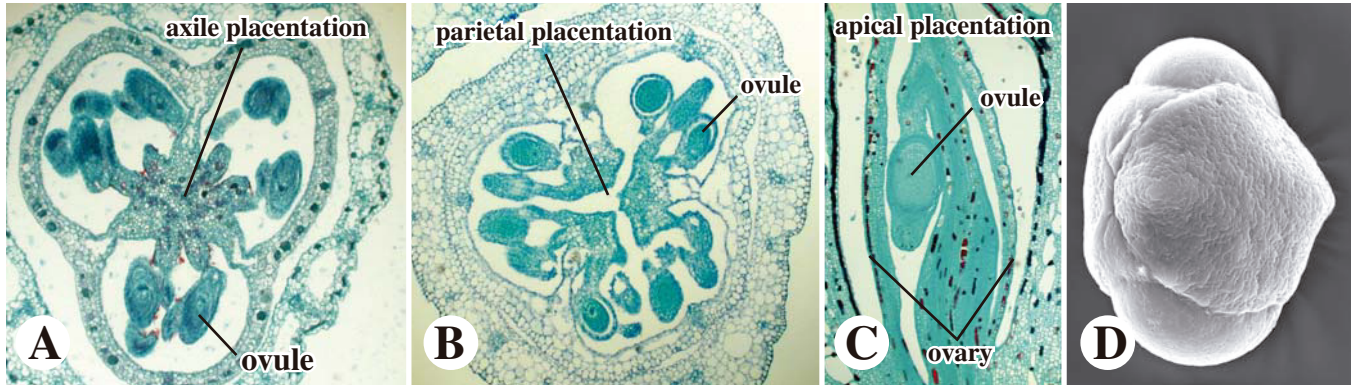


FIGURE 7.50 COMMELINALES. Pontederiaceae. **A.** *Eichornia diversifolia*, ovary cross-section, showing axile placentation. **B.** *Heteranthera reniformis*, ovary cross-section, with parietal placentation. **C.** *Pontederia cordata*, ovary longitudinal section, showing apical placentation. **D.** Pollen of *Pontederia cordata*, showing disulcate apertures.

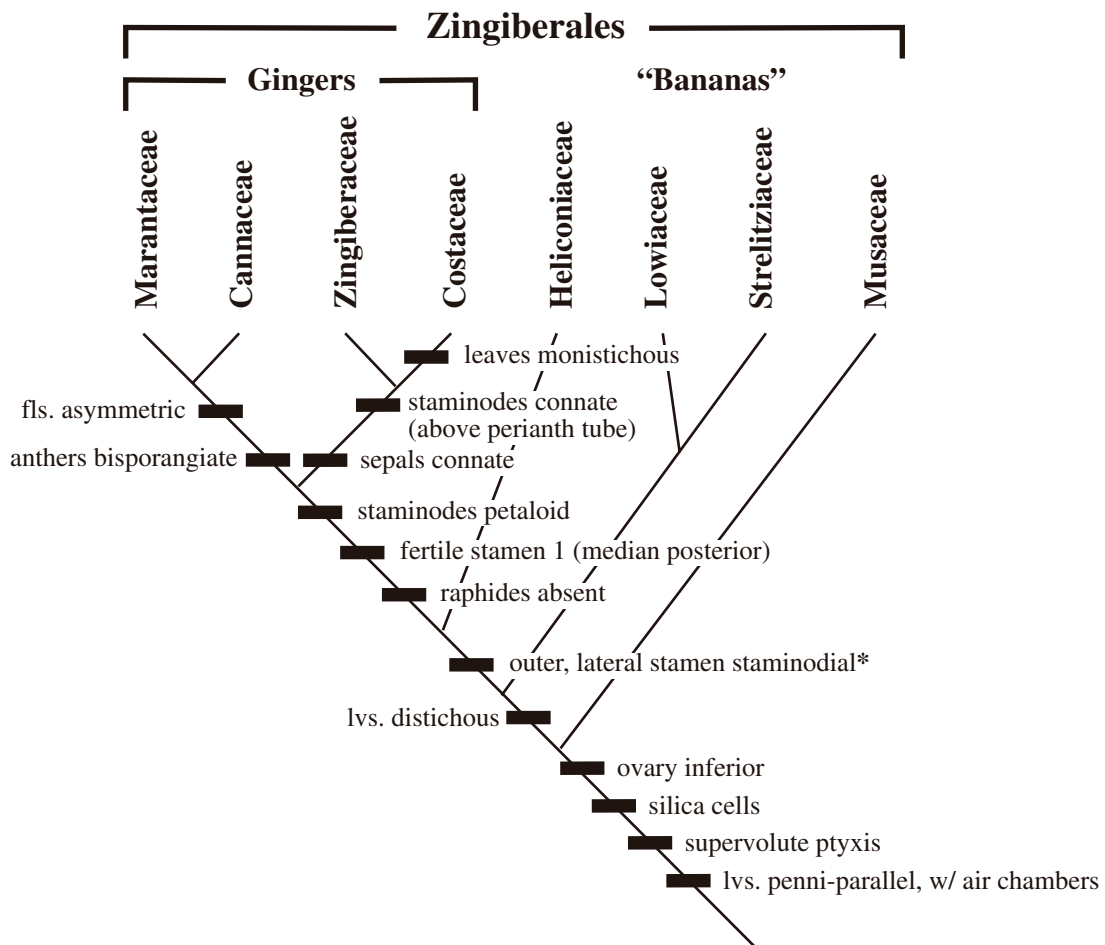


FIGURE 7.51 Cladogram of the Zingiberales (after Kress et al. 2001), with selected apomorphies, that at * after Kirchoff (2003).

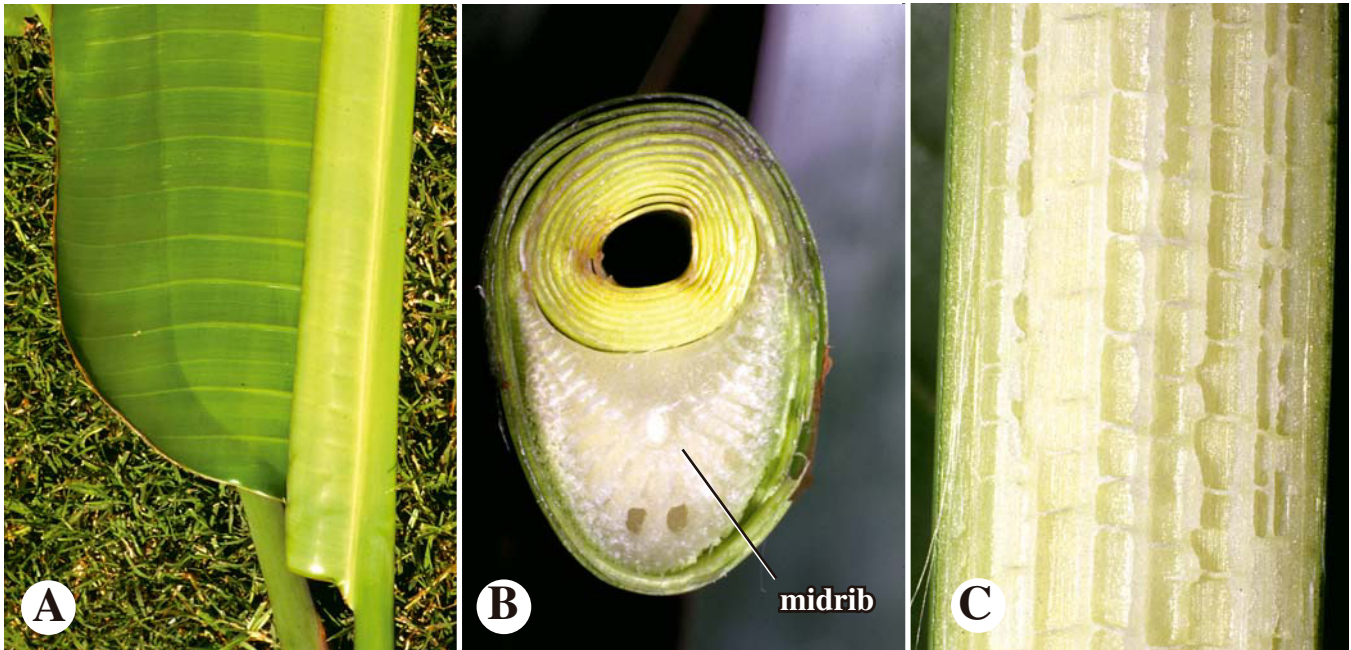


FIGURE 7.52 Apomorphies of the Zingiberales. **A–B.** Supervolute ptyxis. **A.** *Strelitzia reginae* (bird-of-paradise) young leaf, showing right half still partially coiled. **B.** *Musa acuminata* (banana), immature leaf cross-section, showing rolled left and right leaf blade halves. **C.** Diaphragmed air chambers, seen here in a leaf petiole longitudinal section of *Musa acuminata* (banana).

The Musaceae consist of monoecious, perennial herbs. The **stems** are subterranean, sympodial, rhizomatous to cormose, and hapaxanthic. The **leaves** are large, basal, spiral, sheathing (with the long, sheathing leaf bases overlapping, forming a pseudo-stem), a petiole (sometimes termed a pseudo-petiole) present in *Musa*, lacking in *Ensete*, simple (often tearing in several places perpendicular to the midrib), and penni-parallel-veined. The **inflorescence** (which arises from the apical meristem of the corm and grows inside the rolled leaf sheaths) is a terminal thyrse, equivalent to a raceme of spirally arranged, fasciculate, monochasial cymes (commonly called banana hands), bracteate, the bracts large, coriaceous, each enclosing a fasciculate unit cyme, female cymes proximal, male cymes distal. **Flowers** are ebracteate, unisexual, zygomorphic, epigynous. The **perianth** is biseriate and homochlamydeous, 3+3, syntepalous (the inner, adaxial tepal usually distinct). **Stamens** are apostemonous, 5 or 6, the missing stamen or staminode opposite the inner, median, adaxial tepal; anthers are longitudinal in dehiscence, dithecal. The **gynoecium** is syncarpous, with an inferior ovary, 3 carpels (the median carpel anterior), and 3 locules; the styles are terminal; placentation is axile; ovules are anatropous, bitegmic, ∞ per carpel. Septal nectaries are present and occur above the locules. The **fruit** is a berry; seeds are endospermous, with rudimentary arils. Pollinated by bats or birds in the wild.

Members of the family have Old World distributions in tropical Africa and southeast Asia to northern Australia. Economic importance includes use of fruits of *Musa* spp. as a food source (esp. triploid forms of *Musa acuminata* and the triploid hybrid *Musa* \times *paradisica*); *Musa textilis* (Manila-hemp, abac \ddot{E}) and *Musa basjoo* are used as a fiber source for twine, textiles, and building materials. See Andersson (1998a) for a recent treatment of the family.

The Musaceae are distinguished from related families of the Zingiberales in having a **spiral** leaf arrangement and **monoecious** plant sex.

P (3+3) **A** 5-6 **G** (3), inferior.

Strelitziaceae Bird-of-paradise family (after Charlotte of Mecklenburg-Strelitz, wife of King George III). 3 genera (*Phenakospermum*, *Ravenala*, *Strelitzia*)/7 species. (Figures 7.55A, 7.56)

The Strelitziaceae consist of perennial herbs or trees. The underground **stems** are rhizomatous (dichotomously branching in at least some), the aerial stems decumbent and herbaceous or arborescent and woody-textured. The **leaves** are distichous, sheathing, petiolate, simple, and penni-parallel-veined (veins marginally fused). The **inflorescence** is a terminal or axillary thyrse of 1 many monochasial cymes, each cyme subtended by a large, spatheaceous bract. **Flowers** are bisexual, zygomorphic,



The Zingiberales

FIGURE 7.53 Cladogram of the Zingiberales, after Kress et al. 2001. (Artwork by Ida Lopez, by permission of W. J. Kress.)

bracteate, epigynous. The **perianth** is biseriate and homochlamydeous, 3+3, syntepalous, the median inner tepal smaller than the lateral, sometimes connivent inner tepals. **Stamens** are 5 or 6; anthers are basifixed, longitudinal in dehiscence, and bithecal. The **gynoeceum** is syncarpous, with a inferior ovary, 3 carpels (the median carpel anterior), and 3 locules; the style is terminal and filiform; placentation is axile; ovules are anatropous, bitegmic, and ∞ per carpel. Septal nectaries are present. The **fruit** is a loculicidal capsule; seeds are arilate, with a starch-rich endosperm and starch-less perisperm. Pollinated by insects or birds.

Members of the family have distributions in tropical South America, Southern Africa, and Madagascar. Economic importance includes some species used as ornamental cultivars, e.g., *Strelitzia reginae* (bird-of-paradise) and *S. nicolai* (tree bird-of-paradise). See Andersson (1998b) for a recent treatment of the family.

The Strelitziaceae are distinguished from related families of the Zingiberales in having **rhizomatous and decumbent or erect, arborescent** stems with **distichous** leaves and flowers having **5–6 stamens**.

P (3+3) **A** 5 or 6 **G** (3), inferior.

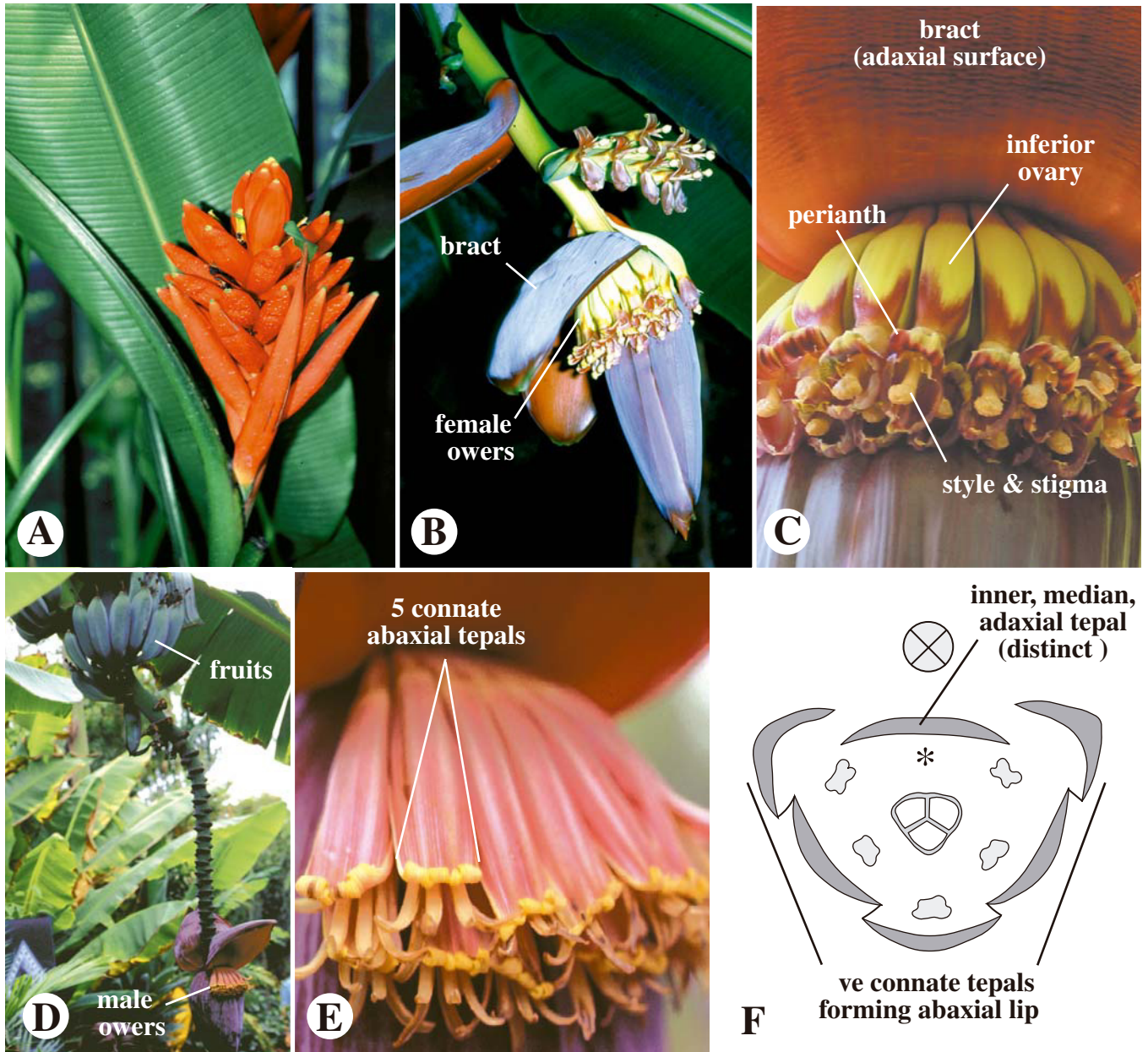


FIGURE 7.54 ZINGIBERALES. Musaceae. **A.** *Musa coccinea*, showing leaf with penni-parallel venation and terminal inflorescence (of bright red flowers and bracts). **B–F.** *Musa acuminata*, cultivated banana. **B.** Young inflorescence, with proximal cyme unit (banana hand) of female flowers, subtended by large bract. **C.** Cyme of female flowers. **D.** Inflorescence, which grew through pseudostem, having proximal female flowers (in fruit) and distal male flowers (below). **E.** Close-up of male flowers. **F.** Floral diagram (combining male and female flowers); * = missing stamen.

Zingiberaceae Ginger family (from a pre-Gr. name, possibly from India). 50 genera / ca. 1300 species. (Figures 7.55B, 7.57)

The Zingiberaceae consist of perennial herbs. The stems are rhizomatous and sympodial. The leaves are distichous, simple, sheathing (sheaths forming a pseudo-stem in some), petiolate, usually ligulate, penni-parallel-veined, a pulvinus

present in *Zingiber*. The **inflorescence** is a bracteate spike, raceme, thyrse, or of solitary flowers. **Flowers** are bisexual, zygomorphic, bracteate, and epigynous. The **perianth** is biseriata and homochlamydeous, 3+3, syntepalous, each whorl 3-lobed. **Stamens** are 1 fertile (median posterior in position); the anther is longitudinal or poricidal in dehiscence, dithecal.

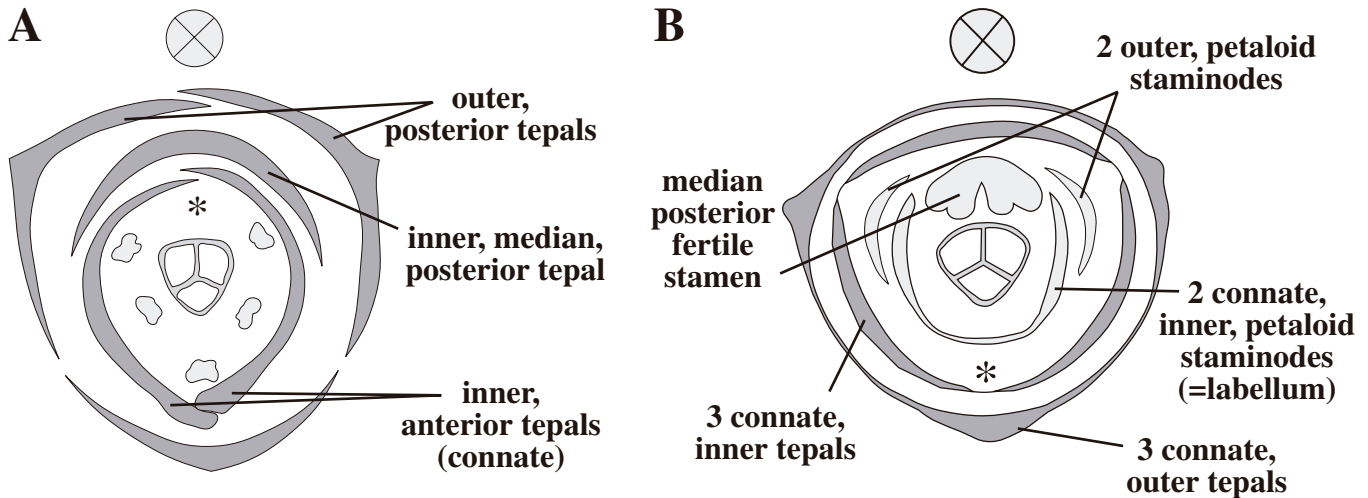


FIGURE 7.55 Floral diagrams. **A.** Strelitziaceae. **B.** Zingiberaceae. * = missing stamen.

Staminodes are 4, petaloid, the two in the inner whorl connate, forming an anterior labellum, the two in the outer whorl distinct above the floral tube or fused to labellum (the third member of the outer whorl absent). The **gynoecium** is syncarpous, with an inferior ovary, 3 carpels (the median carpel anterior), and 1 or 3 locules; the style is terminal and positioned in the furrow of the filament and between the anther thecae; placentation is axile or parietal; ovules are anatropous, bitegmic, and ∞ per carpel. Septal nectaries are absent and replaced by two epigynous nectaries. The **fruit** is a dry or fleshy loculidial or indehiscent capsule; seeds are arillate, with a starch-rich endosperm and perisperm. Plants are insect-pollinated.

The Zingiberaceae are a large family, usually classified into four tribes: Hedychieae (leaves parallel to rhizome, lateral staminodes petaloid, not fused to labellum), Zingibereae (style exerted past anther and enveloped by elongate anther crest), Alpinieae (leaves perpendicular to rhizome, lateral staminodes absent or small and fused to labellum), and Globbeae (filament long-exserted and arched, gynoecium 1-locular). Members of the family have distributions in the tropics of south and southeastern Asia, especially Indomalaysia. Economic importance includes the source of important spice plants, e.g., *Curcuma* spp., including *C. domestica* (turmeric), *Elettaria cardamomum* (cardamom), and *Zingiber* spp., including *Z. officinale* (ginger); some species are grown as cultivated ornamentals, e.g., *Alpinia* and *Hedychium*. See Larsen et al. (1998) and Kress et al. (2002) for recent treatments of the family.

The Zingiberaceae are distinguished from related families of the Zingiberales in having **distichous, usually ligulate**

leaves with a **single, dithecal** stamen and a **petaloid labellum derived from two staminodes**.

P (3+3) **A** 1 fertile + 2 + (2) petaloid staminodes **G** (3), inferior.

Cannaceae—Canna-Lily family (Gr. *canna*, a reed). 1 genus (*Canna*) / 10 25 species. (Figure 7.58)

The Cannaceae consist of perennial herbs. The **stems** are rhizomatous and sympodial. The **leaves** are distichous [to spiral], sheathing, petiolate, simple, and penni-parallel-veined. The **inflorescence** is a bracteate thyrses consisting of a spike or raceme of 2-flowered cymes (or reduced to a raceme). **Flowers** are bisexual, asymmetric, and epigynous. The **perianth** is biseriata and homochlamydeous, 3+3, and apotepalous. **Stamens** are 1 fertile, (median posterior in position), the fertile stamen petaloid. Staminodes are 1 4[5], large, petaloid, resembling the fertile stamen; the anther is laterally subapically positioned on the petaloid stamen, longitudinal in dehiscence, bisporangiate, and monotheical. The **gynoecium** is syncarpous, with an inferior ovary, 3 carpels (the median carpel anterior), and 3 locules; the style is terminal and laminar; placentation is axile; ovules are anatropous, bitegmic, ∞ per carpel. The **fruit** is a usually a capsule; seeds are exarillate, with a starch-rich endosperm and perisperm.

Members of the Cannaceae have distributions in the warm American tropics. Economic importance includes ornamental cultivars of *Canna* spp. (canna lily) and a source of starch (from rhizome of *Canna edulis*). See Kubitzki (1998c) for a recent family treatment.

The Cannaceae are distinguished from related families of the Zingiberales in having usually **distichous** leaves and



FIGURE 7.56 ZINGIBERALES. Strelitziaceae. A–E. *Strelitzia reginae*, bird-of-paradise. A. Whole plant, showing (basal) leaves (arising from rhizome) and lateral, erect inflorescence. B. Close-up of inflorescence. Note large, subtending spathe and two visible ovaries. C. Inflorescence cross-section, showing spathe and monochasium of ovaries (ovaries seen in cross-section), each ovary subtended by a bract. D. Flower close-up, showing inner and outer tepals and inferior ovary. E. Inner anterior tepals pulled back, exposing the enclosed stamens and central style/stigma. F–H. *Strelitzia nicolai*, giant bird-of-paradise. F. Whole plant. Note distichous, sheathing, cauline leaves. G. Inflorescence. H. Loculicidal capsular fruits. Note black seeds covered at base with (orange) arils.

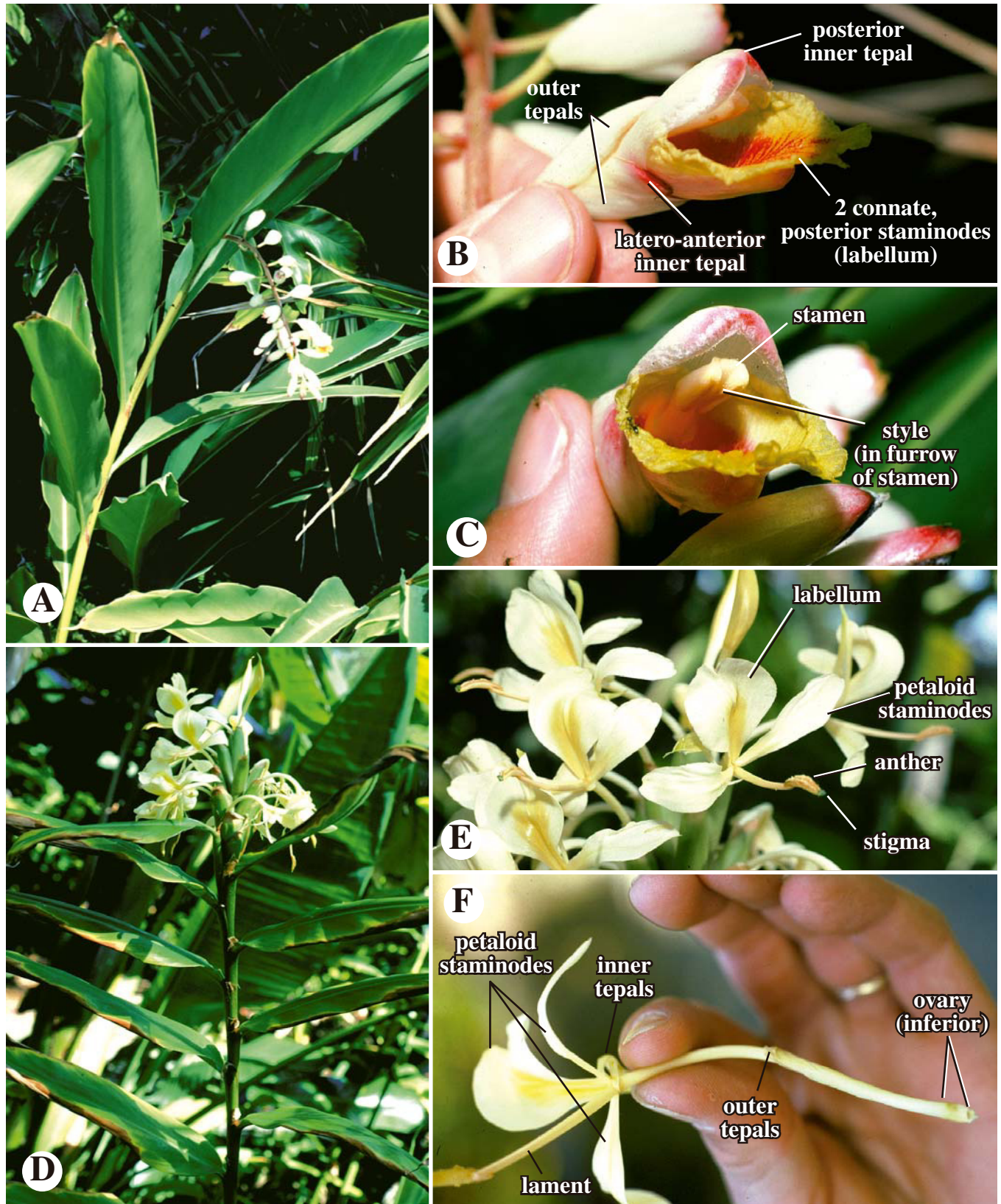


FIGURE 7.57 ZINGIBERALES. Zingiberaceae. A–C. *Alpinia* sp. (shell ginger). A. Whole plant, erect aerial stem with inflorescence. B. Flower close-up, showing tepals and petaloid staminodes, forming an anterior labellum. C. Flower close-up. Note single anther of stamen posterior to and partially enclosing style. D–F. *Hedychium* sp. (ginger-lily). D. Whole plant. Note distichous leaves. E. Inflorescence, showing single stamen of flower. F. Flower close-up (removed), showing inferior ovary, outer and inner tepals, and showy, petaloid staminodes.

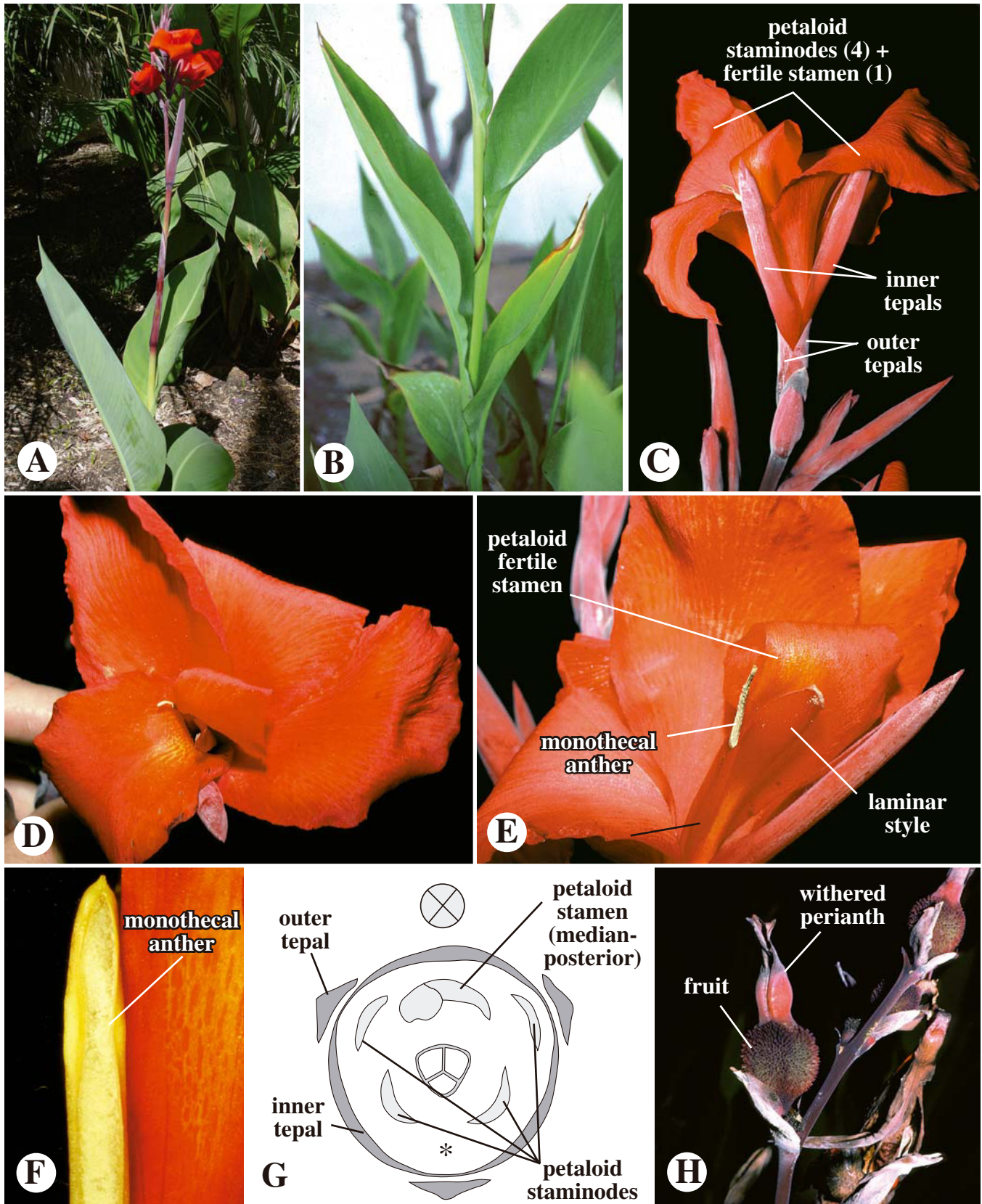


FIGURE 7.58 ZINGIBERALES. Cannaceae. A–F. *Canna x generalis* (canna-lily). A. Whole plant, aerial shoot bearing terminal inflorescence. B. Close-up of shoot, showing sheathing leaves. C. Flower. Note reduced outer tepals, narrow, showy inner tepals, and large, showy, petaloid staminodes and stamen. D. Top view of flower, showing petaloid staminodes. E. Flower close-up, showing petaloid fertile stamen with laterally adnate, monotheical anther. Note laminar style. F. Monotheical anther, close-up. G. Floral diagram. * = missing stamen. H. Close-up of fruit, a capsule.

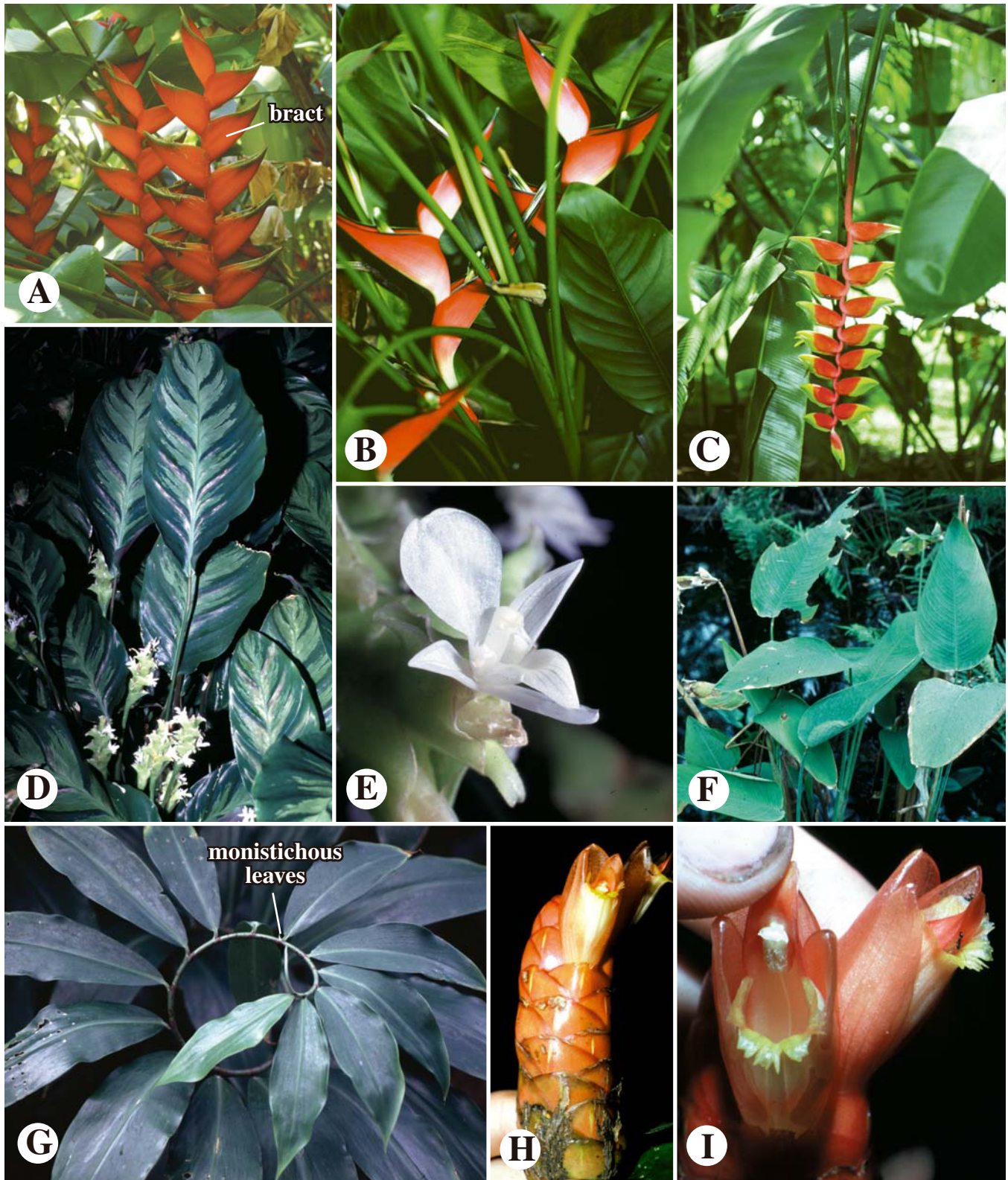


FIGURE 7.59 ZINGIBERALES, exemplars of three other families, not described. A–C. Heliconiaceae, *Heliconia* spp., all with showy bracts. A,B. Taxa with erect inflorescences. C. Taxa with pendant inflorescences. D–F. Marantaceae. D,E. *Calathea louisae*. F. *Thalia geniculata*. G–I. Costaceae, *Costus* spp. G. Leaves spiral, monostichous (borne along one row). H. Inflorescence. I. Flowers.

flowers with *one petaloid, monothechal* stamen associated with *1–4[5] petaloid staminodes*.

P 3+3 **A** 1, petaloid & monothechal + 1-4 petaloid staminodes **G** (3), inferior.

POALES

The Poales, as circumscribed by APG II (2003), is a large group of 18 families, of which nine are described here. One hypothesis of relationships of the families is seen in Figure 7.60. The order includes several basal groups with showy, insect-pollinated flowers. A distinctive, large sub-group of the Poales is characterized by monoulcerate pollen grains (Figure 7.60); these have small, reduced, typically wind-pollinated flowers. See general studies of the angiosperms and monocots (e.g., Chase et al., 2000a, Davis et al., 2004),

plus Bremer (2002) and Michelangeli et al. (2003) for recent studies of the order.

Bromeliaceae Bromeliad family (after Swedish medical doctor and botanist O. Bromell, 1639 1705). 59 genera / 2400 species. (Figure 7.61)

The Bromeliaceae consist of terrestrial or epiphytic, perennial herbs to rosette trees. **Roots** are absorbing or function as holdfasts or rarely absent. The **stem** is a caudex or rarely arborescent, often sympodially branched. The **leaves** are spiral, simple, often adaxially concave, sheathing, in some (the tank bromeliads) tightly overlapping and channeling rain water and runoff to storage in a central cavity, the margins entire or serrate-spinose, the surface usually bearing (at least when young) absorptive, usually peltate, scale-like trichomes, functioning in water and mineral uptake. The **inflorescence** is a terminal, bracteate, spike, raceme, or head,

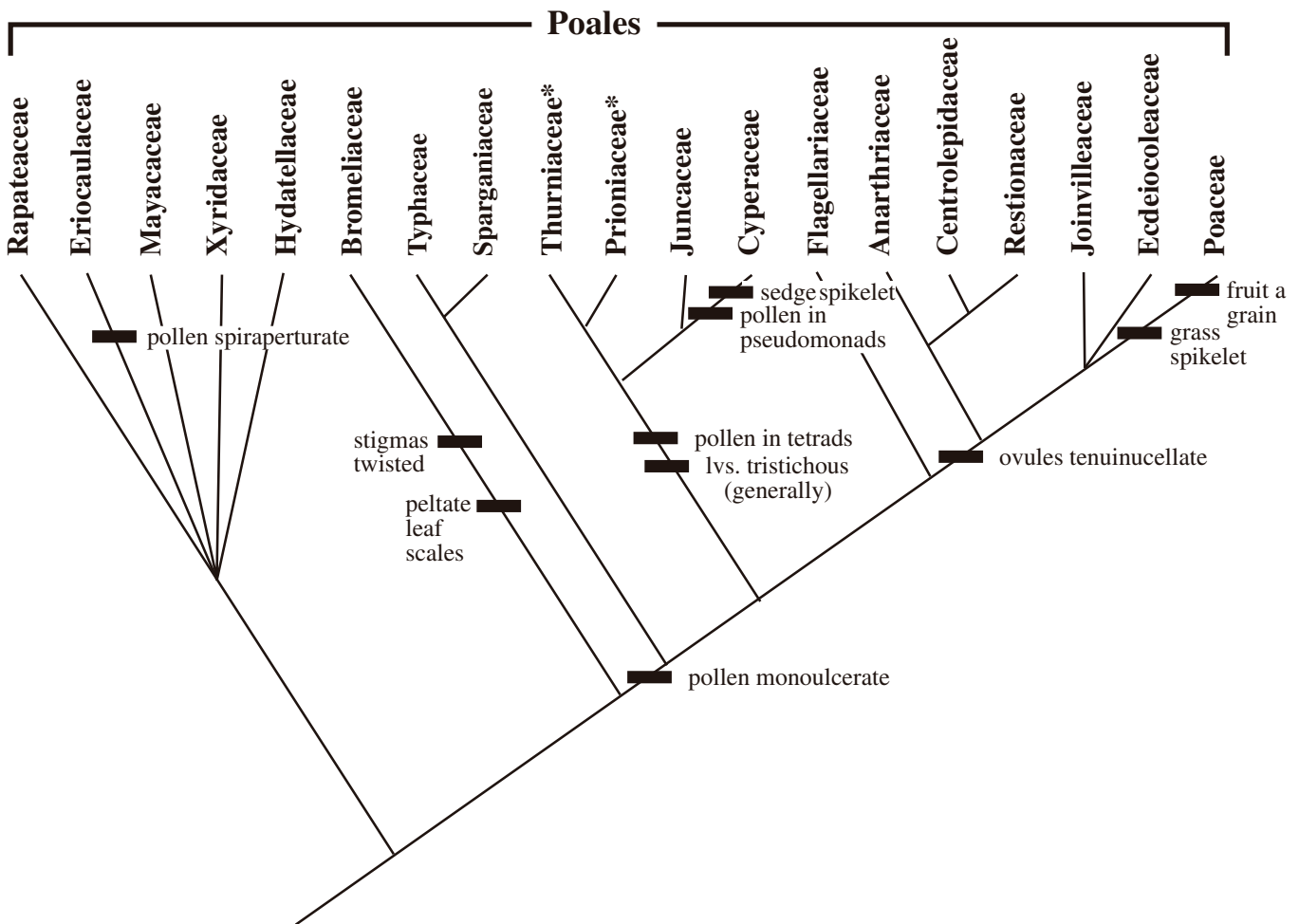


FIGURE 7.60 Cladogram of the families of the Poales, after Michelangeli et al. (2003), with selected apomorphies shown. *Note that the Prioniaceae is united into the Thurniaceae sensu APG II (2003).

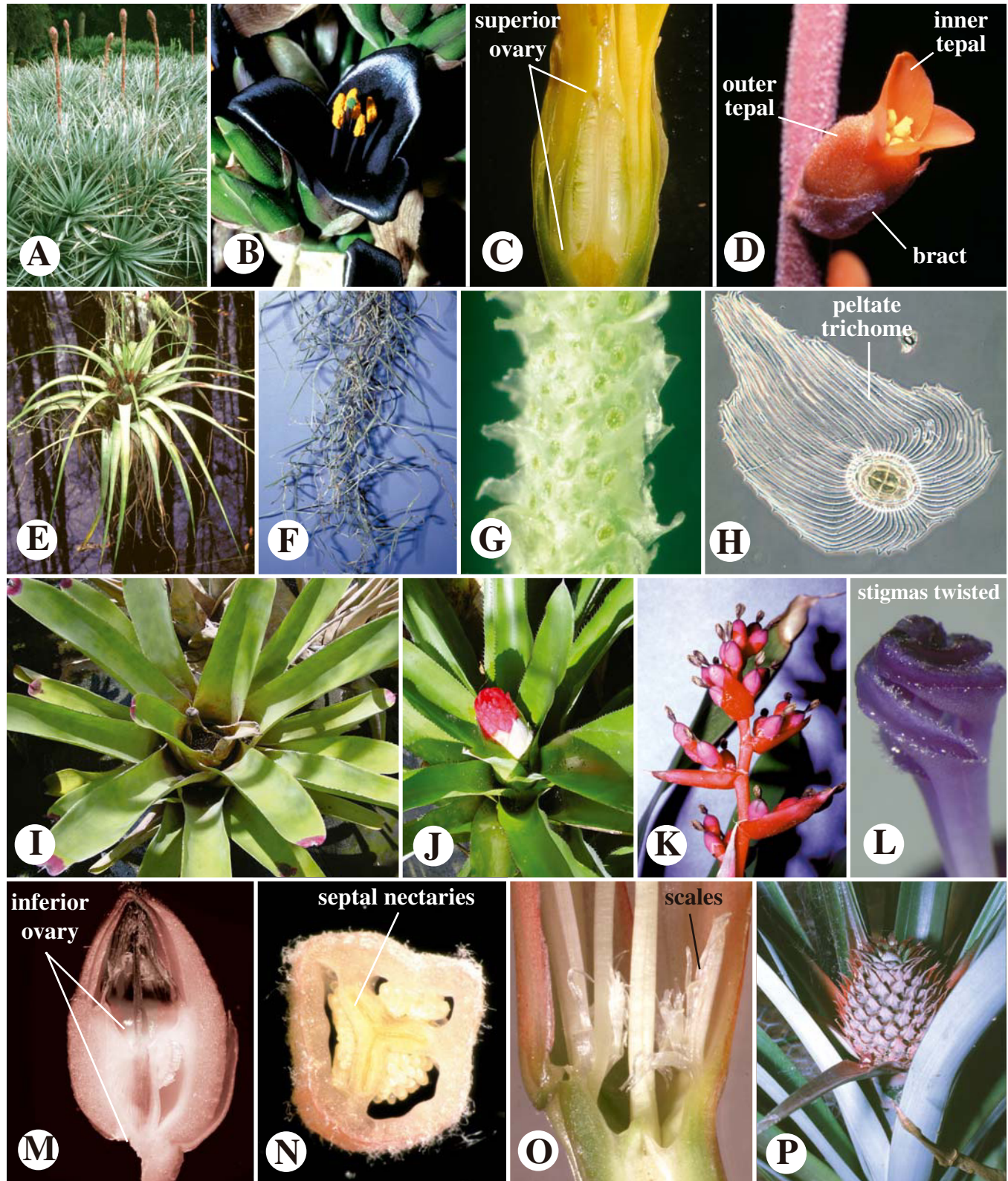


FIGURE 7.61 POALES. Bromeliaceae. A–D. Puyoideae. A. *Puya venusta*, whole plants. B. *Puya alpestris*, flower. C. *Puya* sp., showing superior ovary. D. Pitcairnioideae. *Dyckia dawsonii*, flower. E–H. Tillandsioideae. E. *Tillandsia fasciculata*. F–H. *Tillandsia usneoides*. F. Whole plant. G. Close-up of stem, covered with absorptive, scale-like trichomes. H. Close-up of absorptive, peltate trichome, an apomorphy for the family. I–P. Bromelioideae. I, J. *Neoregalia* sp. I. Whole plant, a tank bromeliad. J. Inflorescence emerging from apex. K–N. *Aechmea* sp. K. Inflorescence. L. Style/stigma close-up. M. Flower, longitudinal section, showing inferior ovary. N. Ovary cross-section. Note septal nectaries. O. *Bilbergia* sp., flower longitudinal section. Note inferior ovary and scales at base of perianth. P. *Ananas comosus*, pineapple. Classification of subfamilies after Givnish et al. (2005).

the bracts often brightly colored. The **flowers** are bisexual [rarely unisexual], actinomorphic or slightly zygomorphic, bracteate, hypogynous or epigynous. The **perianth** is biseriata and dichlamydeous to homochlamydeous, 3+3, the perianth parts distinct to basally connate, the petals/outer tepals often with a basal scale-like or hardened appendage. The **stamens** are diplostemonous, 3+3, distinct or connate, often epipetalous. **Anthers** are longitudinally and introrsely dehiscent. The **gynoecium** is syncarpous, with a superior or inferior ovary, 3 carpels, and 3 locules. The **style** is solitary, with 3 typically twisted **stigmas**. **Placentation** is axile; **ovules** are mostly anatropous, bitegmic, few ∞ per carpel. Septal nectaries are present. The **fruit** is a septicidal capsule or berry, rarely a sorosis (*Ananas*). The **seeds** are winged, plumose, or glabrous. Pollinated by birds, insects, bats, wind [rarely], or flowers cleistogamous.

The Bromeliaceae are traditionally classified into three subfamilies: Pitcairnioideae, with superior (to half-inferior) ovaries, forming capsules with winged seeds; Tillandsioideae, with superior ovaries forming capsules with plumose seeds; and Bromelioideae, with inferior ovaries forming berries with unappendaged seeds. Givnish et al. (2005) proposed a classification into eight monophyletic subfamilies: Brocchinioideae, Lindmanioideae, Hechtioideae, Tillandsioideae, Navioideae, Pitcairnioideae, Puyoideae, and Bromelioideae. Their Tillandsioideae and Bromelioideae correspond to the traditional subfamilies; the other six were formerly classified together (as a now paraphyletic Pitcairnioideae, s.l.). Members of the

Bromeliaceae are distributed almost entirely in the American tropics. Economic importance includes uses as fruit plants (e.g., *Ananas comosus*, pineapple), fiber plants, and cultivated ornamentals. See Smith and Till (1998) for a general description and Givnish et al. (2005) for a phylogenetic analysis of the family.

The Bromeliaceae are distinctive in being perennial **terrestrial or epiphytic** herbs or shrubs with **absorptive, peltate trichomes**, often **colorful** bracts, and trimerous flowers, the petals/outer tepals often with **basal scales or appendages**, stigmas typically **twisted**.

P 3+3 or (3)+(3) **A** 3+3 **G** (3), superior or inferior.

Typhaceae Cattail family (Gr. for various plants). 1 genus (*Typha*) / 8 13 species. (Figure 7.62)

The Typhaceae consist of emergent, aquatic, monoecious, perennial herbs. The **stems** are rhizomatous. The **leaves** are bifacial, mostly basal, distichous, sheathing, simple, undivided, flat, elongate and narrow, and parallel veined, with spongy parenchyma. The **inflorescence** is a terminal, cylindrical spike of very dense flowers, male above and female below. The **flowers** are very small, unisexual, actinomorphic, female flowers hypogynous. The **perianth** consists of 0 3 [8] bristle-like tepals in male flowers, ∞ bristle or scale-like tepals (in 1 4 whorls) in female flowers. The **stamens** are 3 [1 8], apostemonous. **Anthers** are basifixed, with connective broad, extended beyond thecae. The **pollen** is released as tetrads or monads. The **gynoecium** is unicarpellous, with

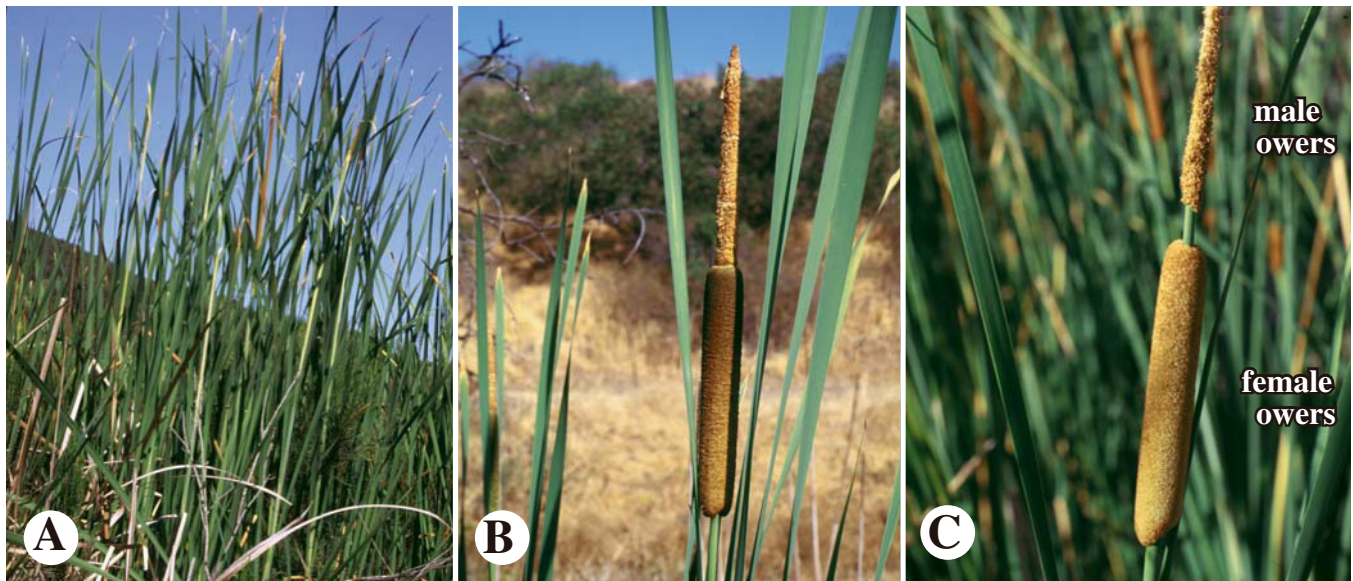


FIGURE 7.62 POALES. Typhaceae. **A,B.** *Typha latifolia*. **A.** Whole plants, emergent, rhizomatous herbs. **B.** Inflorescence. **C.** *Typha domingensis*, close-up of inflorescence, female flowers below, male above.

a superior ovary. The **style** is accrescent. **Placentation** is apical; the **ovule** is solitary, anatropous, bitegmic. **Nectaries** are absent. The **fruit** is a dehiscent, achenelike fruit, with an accrescent gynophore (stipe) and style and persistent perianth parts, aiding in wind dispersal. The **seeds** are starchy endospermous. Flowers are wind pollinated.

Members of the Typhaceae grow as emergents in ponds, ditches, and marshes with worldwide distributions. Economic importance includes local uses as food (pollen or starchy rhizome), matting (leaves), paper, or as ornamental cultivars. See Kubitzki (1998d) for a description of the Typhaceae.

The Typhaceae are distinctive in being *perennial, rhizomatous, monoecious, emergent aquatics* with *distichous, bifacial* leaves, a *spike* of numerous, minute, *wind-pollinated* flowers (male above and female below) having a *bristlelike or scalelike* perianth, and an *achenelike, dehiscent* fruit with an *accrescent stipe and style*.

Male flowers: **P** 0-3 [-8] **A** 3 [1-8].

Female flowers: **P** ∞ **G** 1, superior.

Sparganiaceae Bur-Reed family (Gr. for band used to wrap or bind, after the long, narrow leaves). 1 genus (*Sparganium*) / 14 species. (Figure 7.63)

The Sparganiaceae consist of emergent, aquatic, monoecious, perennial herbs. The **stems** are rhizomatous. The **leaves** are bifacial, distichous, sheathing, simple, undivided, flat, elongate and narrow, and parallel veined. The **inflorescence** is compound, of globose, bracteate, unisexual heads, male heads above, female below. The **flowers** are small, unisexual, actinomorphic, sessile, the female flowers hypogynous. The **perianth** is apparently bracteate in female flowers, the scale-like tepals 1-6 in males, 3-4 [2-5] in females. The **stamens** are 1-8, antitepalous, distinct or basally connate. The **gynoecium** is unicarpellous or syncarpous, with a superior ovary, 1 [2-3] carpel(s), and 1 [2-3] locule(s). **Placentation** is apical; **ovules** are anatropous, bitegmic, 1 per carpel. **Nectaries** are absent. The **fruit** is dry and drupelike with a persistent perianth and style. The **seeds** are endospermous. Flowers are wind pollinated.

Members of the Sparganiaceae have a worldwide distribution. Taxa are of no significant economic importance. The family is sometimes united in the Typhaceae; see Kubitzki (1998d).

The Sparganiaceae are distinctive in being *perennial, rhizomatous, monoecious, emergent aquatics* with *distichous, bifacial* leaves and *unisexual, globose heads* (male heads above, female below) of numerous, minute, wind-pollinated flowers having a *scalelike* perianth, and a *drupelike* fruit with *persistent style*.

Male flowers: **P** 1-6 **A** 1-8 or (1-8).

Female flowers: **P** 3-4 [2-5] in female **G** 1 [(2-3)], superior.

Juncaceae Rush family (L. for binder, in reference to use in weaving and basketry). 7 genera/ca. 350 species. (Figure 7.64)

The Juncaceae consist of perennial, rarely annual, herbs. The **stems** of perennials are usually rhizomatous. The **leaves** are simple, parallel veined, undivided, bifacial or unifacial, mostly basal, spiral, usually tristichous [rarely distichous], sheathing, usually with auricles and ligulate, flat or terete. The **inflorescence** is of solitary flowers or compound of 1 many cymes, glomerules, or heads. The **flowers** are bisexual, rarely unisexual, actinomorphic, bracteate, hypogynous. The **perianth** is usually scarious, biseriate, homochlamydeous, rarely uniseriate, 3+3 [2+2 or 3], apotepalous, with hypanthium absent. The **outer** and **inner tepals** are distinct, each whorl of 3 [2] parts. The **stamens** are 3+3 [3+0 or 2+2], whorled, diplostemonous when biseriate, unfused. **Anthers** are basifixed, longitudinally dehiscent. The **pollen** is released as tetrads. The **gynoecium** is syncarpous, with a superior ovary, 3 carpels, and 3 or 1 locules. The **style** is usually 3-branched, **stigmas** sometimes twisted. **Placentation** is axile, basal, or parietal; **ovules** are anatropous, bitegmic, 1 ∞ per carpel. The **fruit** is a loculicidal capsule, rarely indehiscent. **Seeds** are starchy endospermous. Flowers are wind or insect pollinated.

The Juncaceae, Cyperaceae, and another family (Thurniaceae, including Prioniaceae) probably share two major apomorphies: tristichous leaves and pollen in tetrads (Figure 7.60). Members of the Juncaceae have a worldwide distribution, generally in temperate and cool regions. Economic importance is limited, some used as ornamental cultivars, *Juncus* spp. used indigenously to make matting, bowls, or other products; cushion-forming *Distichia* is used as fuel in Peru. See Balslev (1998) and Drabkova et al. (2003) for recent treatments and phylogenetic analyses of the Juncaceae.

The Juncaceae are distinctive in being usually *perennial herbs* with *spiral, sheathing*, bifacial or unifacial leaves, trimerous, actinomorphic flowers with a typically *scarious* perianth and a *loculicidal capsule*.

P 3+3 [2+2 or 3] **A** 3+3 [3+0 or 2+2] **G** (3), superior.

Cyperaceae Sedge family (Gr. for several species of the genus *Cyperus*). 104 genera / ca. 5000 species. (Figures 7.65, 7.66)

The Cyperaceae consist of perennial or annual herbs, rarely shrubs or lianas. The **stems** of perennials are rhizomes, stolons, bulbs, or caudices bearing aerial culms that are often tufted (cespitose), usually 3-sided, with a solid pith. The **leaves** are bifacial, spiral, and usually tristichous [rarely distichous], sheathing (sheath usually closed), simple, undivided, narrow, flat, and parallel veined, a ligule present or absent;

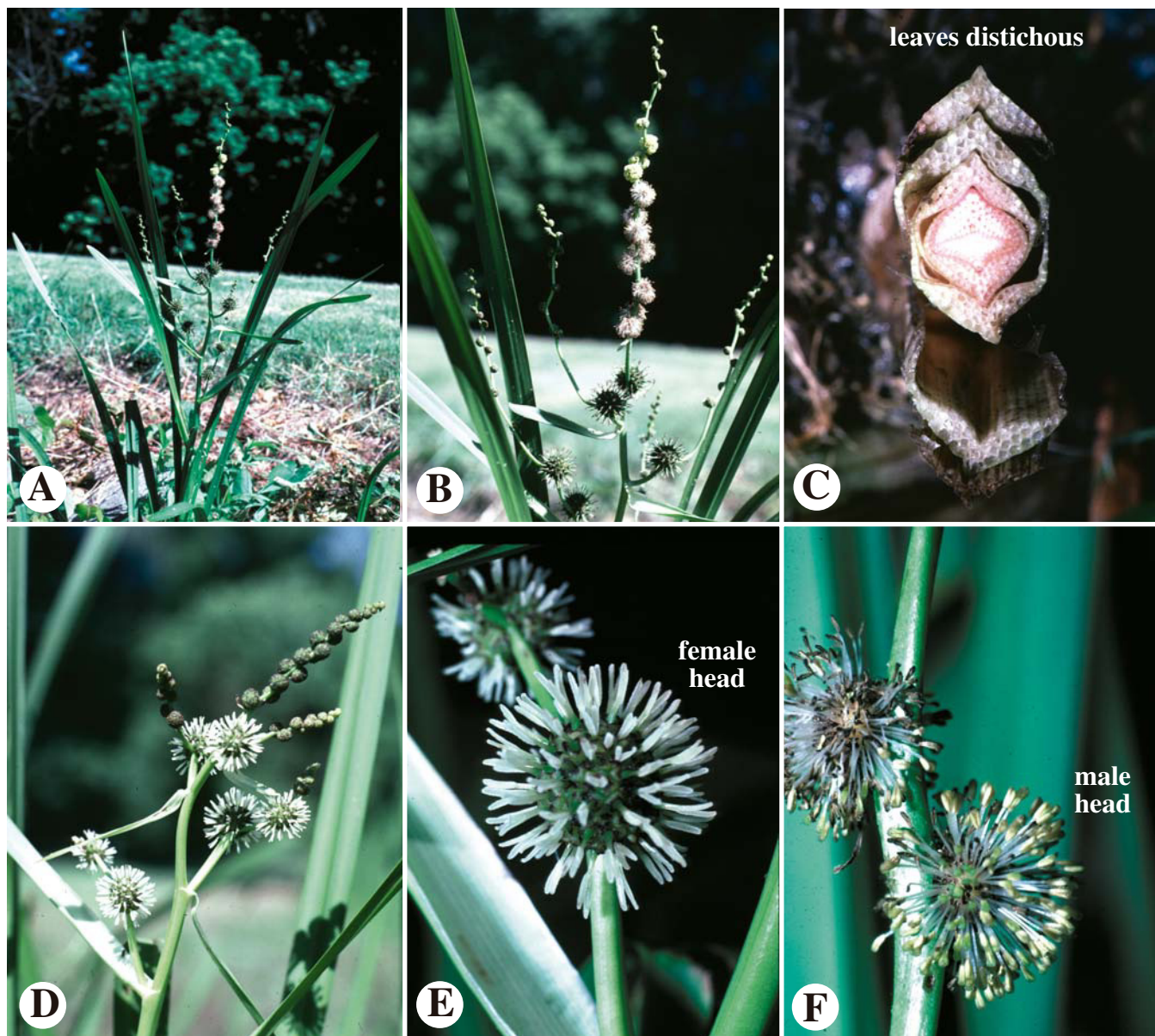


FIGURE 7.63 POALES. Sparganiaceae. *Sparganium* sp. A. Whole plant. B. Inflorescence, a panicle of globose heads. C. Shoot cross-section, showing distichous leaf arrangement. D. Inflorescence, female heads below, male above. E. Female heads. F. Male heads.

lower leaves (or in some taxa all leaves) reduced to sheaths. The **inflorescence** consists of one or more bisexual or unisexual sedge spikelets (either solitary or in various types of secondary inflorescences), each spikelet consisting of a central axis (the rachilla), bearing spiral or distichous bracts (also called scales or glumes), each (except sometimes the lower) subtending a single flower. The **flowers** are small, unisexual or bisexual, actinomorphic, hypogynous. The **perianth** is absent or 6-merous [1∞], of reduced, distinct bristles or scalelike tepals. **Stamens** are 3 [$1-6+$], anthers

introse and longitudinal in dehiscence, filaments elongating during anthesis. The **pollen** is released as pseudomonads, in which 3 of the 4 nuclei of the microspore tetrad degenerate after microsporogenesis. The **gynoecium** is syncarpous, with a superior ovary, 2 or 3 [rarely 4] carpels, and 1 locule; the gynoecium of *Carex* and relatives is surrounded by an inflated bract, known as the perigynium, at the apex of which the style protrudes. The **styles** are usually 2 or 3. **Placentation** is basal; **ovules** are anatropous, bitegmic, 1 per ovary. **Nectaries** are absent. The **fruit** is a lenticular (2-sided)

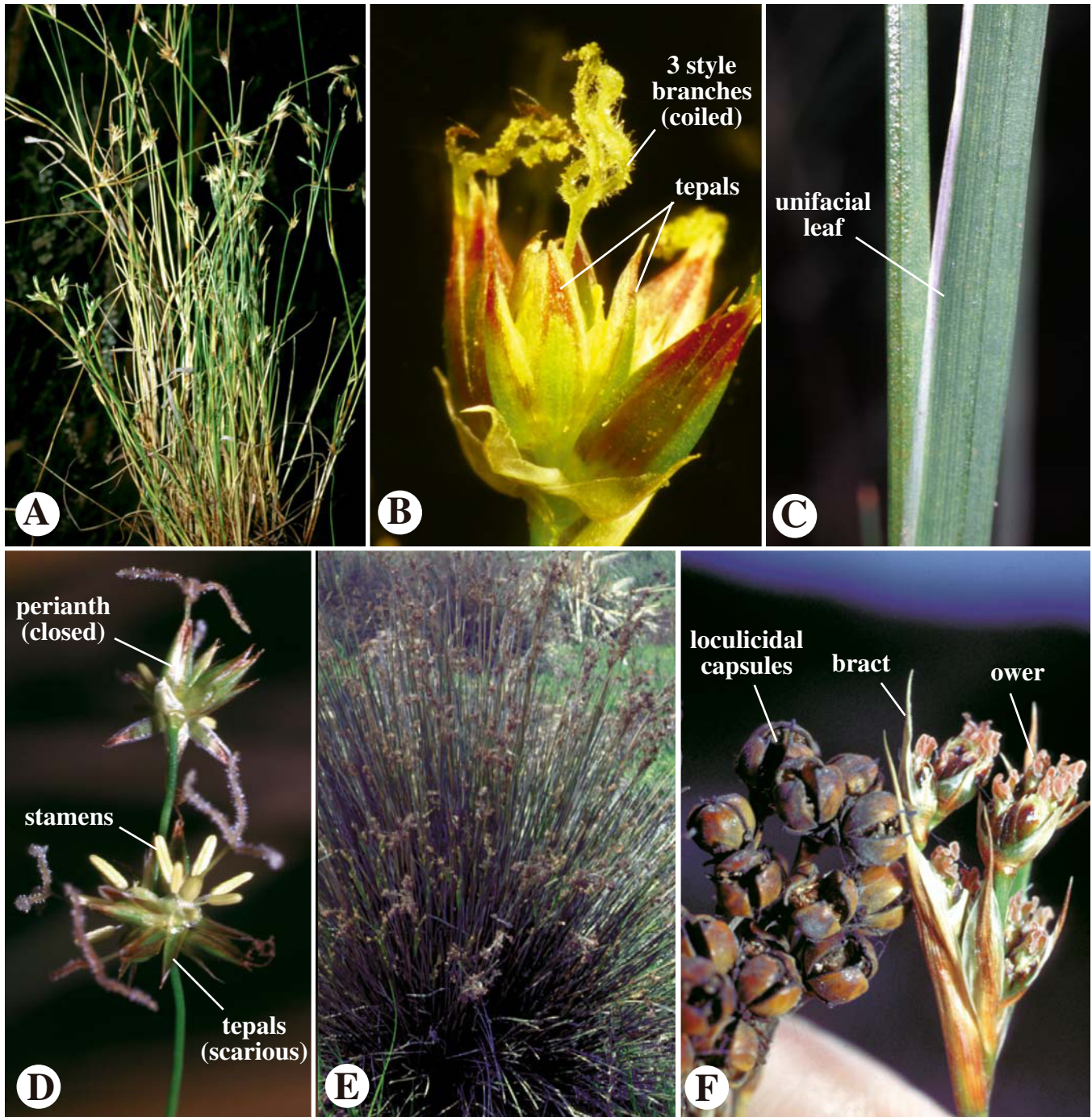


FIGURE 7.64 POALES. Juncaceae. **A.** *Juncus bufonius*, whole plant. **B.** *Juncus dubius*, flower close-up. Note six, imbricate, scarious tepals. **C,D.** *Juncus phaeocephalus*. **C.** Close-up of unifacial leaves. **D.** Flowers, showing exerted styles. **E.** *Juncus acutus*. **E.** Whole plant. **F.** Close-up of infructescence of capsules (left) and inflorescence (right).

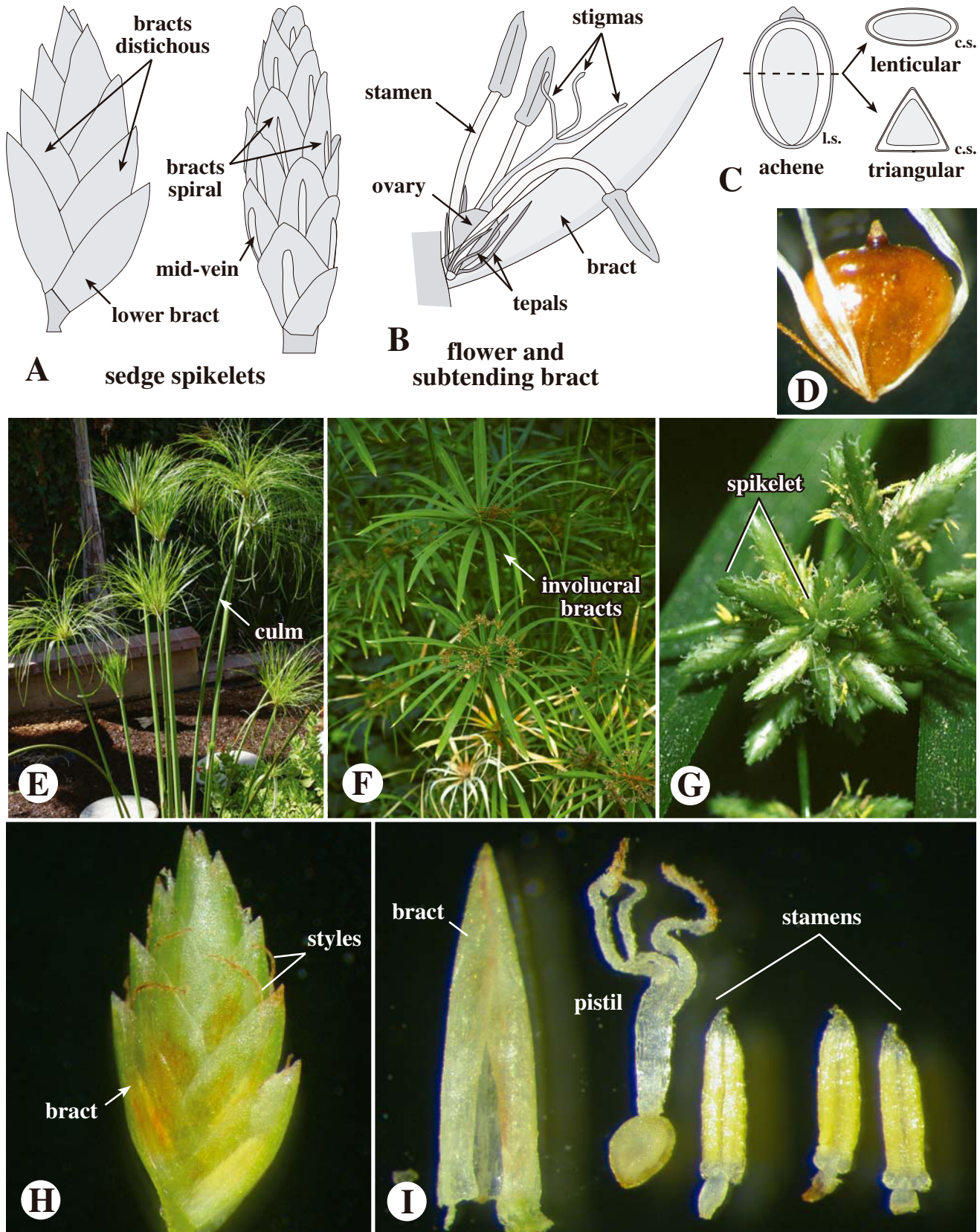


FIGURE 7.65 POALES. Cyperaceae. **A.** Diagram of sedge spikelets with distichous (left) and spiral (right) bracts. **B.** Diagram of sedge flower. **C.** Diagram of achene, illustrating two shape types; l.s. = longitudinal-section; c.s. = cross-section. **D.** *Scirpus* sp., mature achene. **E.** *Cyperus papyrus*, papyrus. **F–I.** *Cyperus involucreatus*. **F.** Whole plant showing prominent involucre bracts. **G.** Close-up of involucre, a glomerule of spikelets. **H.** Spikelet, close-up. **I.** Subtending bract (left) and dissected flower components (right).

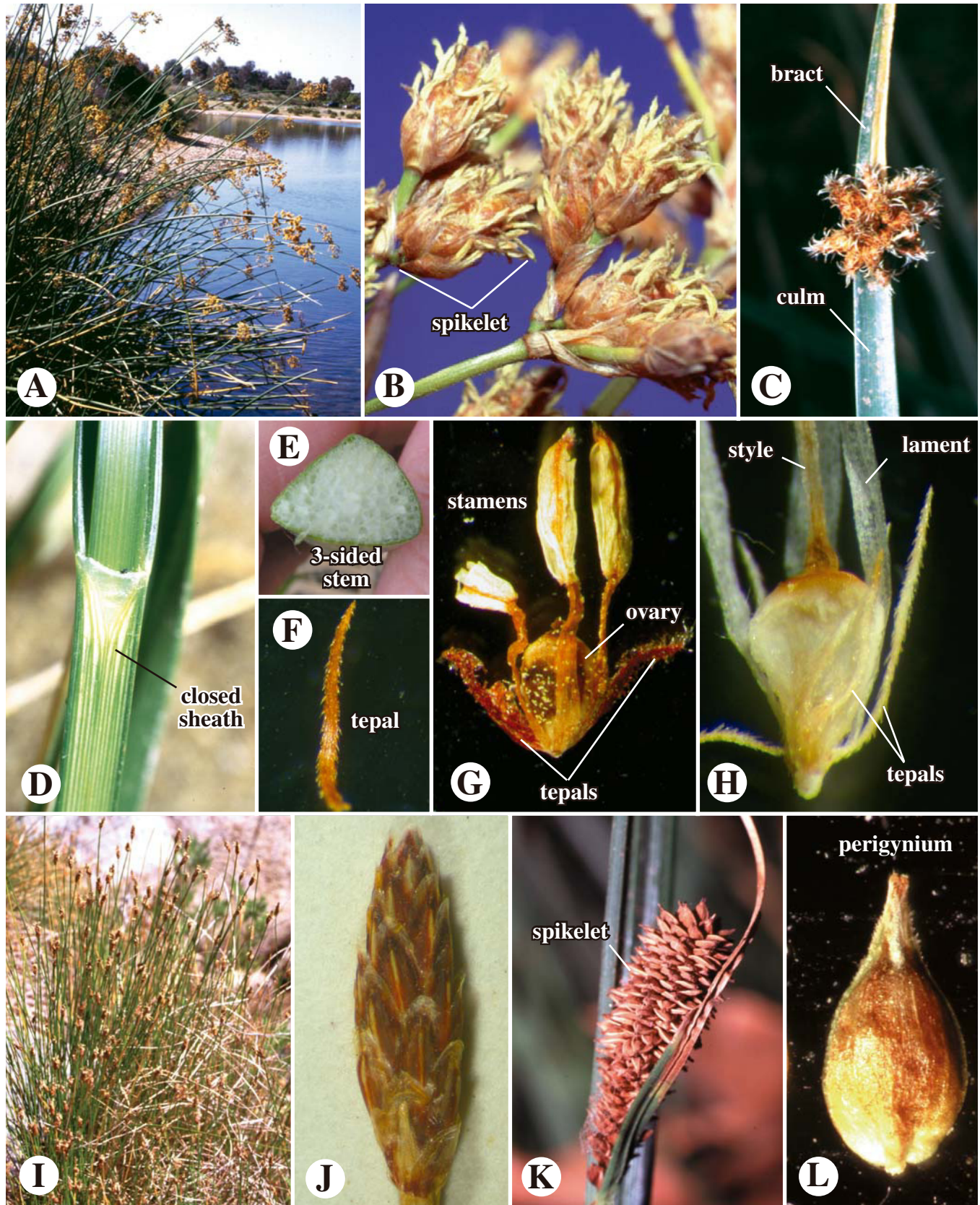


FIGURE 7.66 POALES. Cyperaceae. **A,B.** *Scirpus californicus*. **A.** Plants growing along pond edge. **B.** Close-up of inflorescence, a dense panicle of spikelets. **C.** *Scirpus americanus*, inflorescence arising from culm. Note stemlike bract at apex. **D,E.** *Scirpus maritimus*. **D.** Closed leaf sheath. **E.** Stem cross-section, showing three-sided shape. **F–H.** *Scirpus* spp. **F.** Tepal, detached. **G.** Flower. **H.** Base of flower, showing tepals. **I,J.** *Eleocharis montevidensis*. **I.** Whole plant; note cespitose culms. **J.** Spikelet, close-up. **K.** *Carex barbarae*, spike of female spikelets. **L.** *Carex praegracilis*, perigynium, covering pistil (inside).

or trigonous (3-sided) achene (also called a nutlet). Plants are generally wind pollinated.

Members of the Cyperaceae have a worldwide distribution, especially in temperate regions. The genus *Carex* is especially diverse with ca. 2000 species, important in a number of ecosystems. Economic importance is limited, with some species used as mats, thatch, weaving material, or writing material (*Cyperus papyrus*, papyrus, the culm pith of which was historically used to make paperlike scrolls), a few used as ornamental cultivars (e.g., *Cyperus involucratus*, umbrella plant), and some species, such as the nutsedges, being noxious weeds. See Goetghebeur (1998) and Simpson et al. (2003) for recent descriptions and phylogenetic analyses of the family.

The Cyperaceae are distinctive in being herbs with usually **3-sided, solid-pithed stems, closed-sheathed, often tristichous**

leaves, the inflorescence a **sedge spikelet**, consisting of a central axis bearing many sessile, **distichous or spiral bracts**, each subtending a single, reduced unisexual or bisexual flower, with perianth **absent or reduced to bristles or scales**, usually 3 stamens, and a 2-3-carpellate ovary, the fruit a 2- or 3-sided **achene**.

P 6 or 0 [1-∞] **A** 3 [1-6+] **G** (2-3)[(4)], superior.

Eriocaulaceae Pipewort family (Gr., woolly stem). 10 genera/700-1400 species. (Figure 7.67)

The Eriocaulaceae consist of monoecious [rarely dioecious], perennial or annual herbs. The **stems** are rhizomatous, basal shoots often tufted (cespitose). The **leaves** are basal, often rosulate, spiral [rarely distichous], basally sheathing, simple, usually narrow (flat or terete, canaliculate in some), and parallel veined. The **inflorescence** is a scapose head with

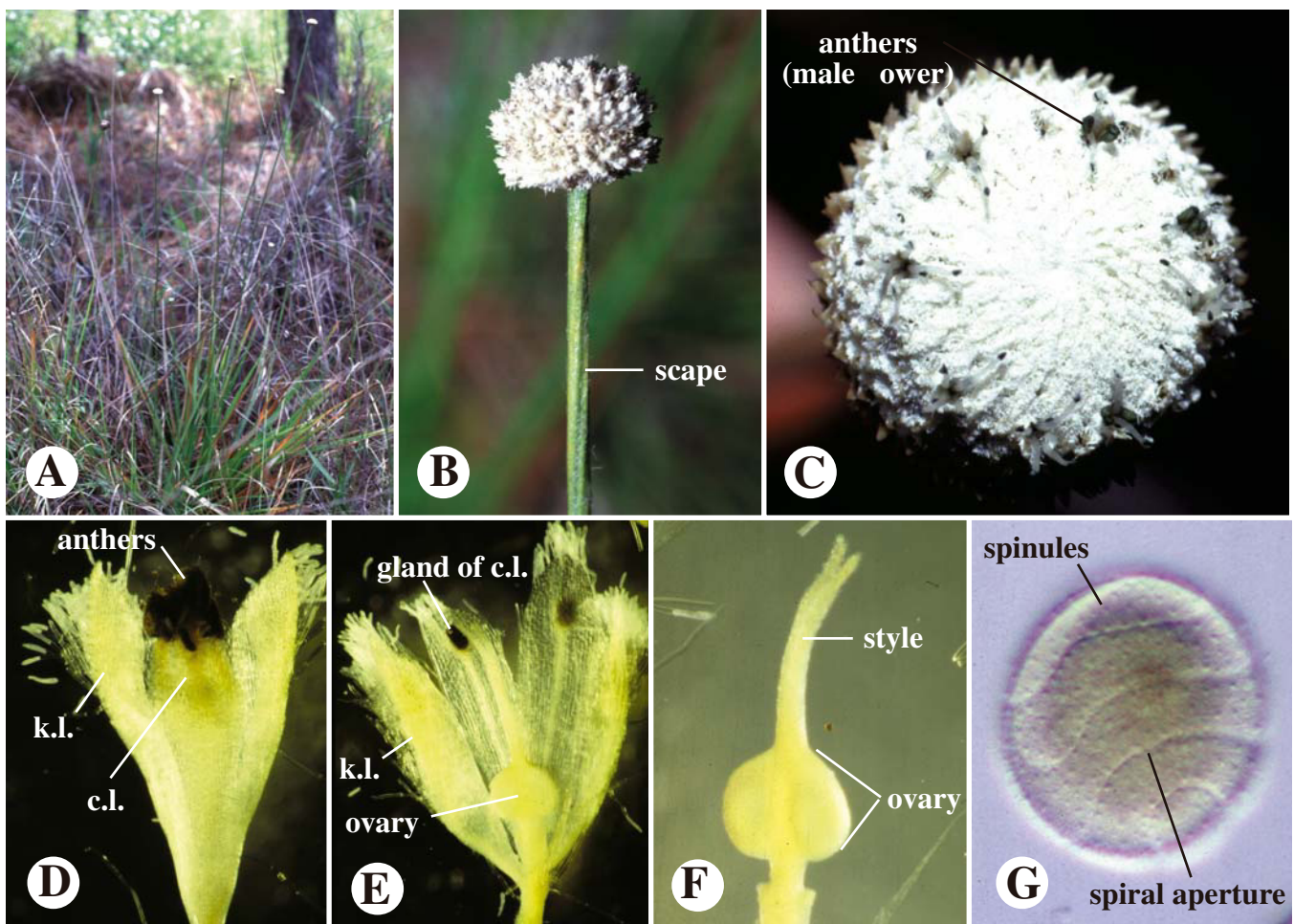


FIGURE 7.67 POALES. Eriocaulaceae. *Eriocaulon* sp. **A**. Whole plant. **B**. Inflorescence, a scapose head. **C**. Head, face view, showing dark anthers of male flowers. **D**. Male flower. **E**. Female flower. **F**. Ovary of female flower, tepals removed. **G**. Spiraperturate pollen. Abbreviations: c.l. = corolla lobe; k.l. = calyx lobe.

subtending imbricate bracts (phyllaries), the compound receptacle often with trichomes or chaffy bracts, in monoecious species the male and female flowers mixed or females marginal. The **flowers** are small, whitish, unisexual, actinomorphic or zygomorphic, sessile or short-pedicellate, subtended by chaffy bracts or bractless. The **perianth** is biseriate, dichlamydeous, transparent, whitish, or variously colored. The **calyx** is distinct or basally fused into a tube with 2 or 3 parts. The **corolla** is also distinct or basally fused into a tube and with 2 or 3 parts [corolla rarely absent]. The **stamens** are 2 or 4 in dimerous flowers, 3, 6, or 1 in trimerous flowers, antipetalous (when 2 or 3), epipetalous or arising from an androphore. **Anthers** are longitudinal and introrse in dehiscence, bi- or tetrasporangiate. The **pollen** is spheroidal, usually spiraperturate-spinulose. The **gynoecium** is syncarpous, with a superior ovary, 2 or 3 carpels, and 2 or 3 locules. The **style** is solitary, sometimes style-like appendages also present; **stigmas** are 2-3, dry. **Placentation** is apical, ventral-pendulous; **ovules** are orthotropous, bitegmic, 1 per carpel. **Nectaries** are absent except for glands at tepal tips in some taxa. The **fruit** is a loculicidal capsule. The **seeds** are ellipsoidal, endospermous, starchy. Flowers are wind or insect pollinated.

The Eriocaulaceae consist of the principal genera *Eriocaulon*, *Leiothrix*, *Paepalanthus*, and *Syngonanthus*. Members of the family grow in wet areas with distributions in tropical to subtropical warm regions, especially the Americas, a few northern temperate. Economic importance includes inflorescences of *Syngonanthus* used in the floral trade as everlasting. See St tzel (1998) for a detailed family description.

The Eriocaulaceae are distinctive in being perennial or annual herbs with **basal, often rosulate** leaves and a **scapose head** of very small, **unisexual** usually **white** flowers.

K 2-3 or (2-3) **C** 2-3 or (2-3) **A** 2, 3, 2+2 or 3+3 **G** (2-3), superior.

Xyridaceae Yellow-eyed-grass family (Gr. name for plant with razorlike leaves). 5 genera / ca. 385 species. (Figure 7.68)

The Xyridaceae consist of perennial or annual herbs. The **stem** of perennials is a caudex, less commonly a rhizome or corm. The **leaves** are bifacial or unifacial-ensiform, usually basal and rosulate, alternate distichous or spiral, sheathing with sheaths often persistent, simple, ligulate in some, narrow, flat or terete, and parallel veined. The **inflorescence** is a terminal, scapose, usually solitary spike or head with bracts subtending single or (in *Achlyphila*) 2 or 3 flowers. The **flowers** are bisexual, hypogynous, actinomorphic or slightly zygomorphic, sessile or pedicellate, subtended by imbricate, indurate bracts. The **perianth** is biseriate, dichlamydeous. The **calyx** is aposepalous with 3 sepals, the anterior one

reduced to absent. The **corolla** is ephemeral, of three, distinct or basally connate petals/corolla lobes, usually yellow, rarely white, blue, or magenta. The **stamens** are 3 [6], whorled, biseriate or uniseriate, or with 3 staminodes and 3 fertile stamens. **Anthers** are longitudinal in dehiscence. The **pollen** is sulcate or inaperturate. The **gynoecium** is syncarpous, with a superior ovary, 3 carpels and 1 or 3 locules. The **style** is solitary; **stigmas** are 3 [1]. **Placentation** is axile, basal, free-central, or parietal; **ovules** are orthotropous or anatropous, bitegmic, few ∞ per carpel. The **fruit** is a loculicidal or irregularly dehiscent capsule, sometimes enclosed by persistent sepals and bracts. The **seeds** are small, endospermous (starchy and proteinaceous, sometimes oily).

Three genera of the family, *Abolboda*, *Aratitiopea*, and *Orectanthe* (sometimes classified in the family Abolbodaceae) have spiral, bifacial leaves, spinose pollen, highly connate petals, and asymmetric, appendaged styles. The other two genera, *Achlyphila* and *Xyris*, have distichous, unifacial leaves, nonspinose pollen, slightly connate or distinct petals, and symmetric, unappendaged styles. Members of the Xyridaceae grow in wet areas, such as marshy savannas, and have a worldwide distribution in tropical and warm and some temperate regions; three genera are restricted to northern South America. Economic importance includes *Xyris* spp. used occasionally ornamentally and medicinally. See Kral (1998) for a detailed family treatment.

The Xyridaceae are distinctive in being perennial or annual herbs with a terminal, **scapose bracteate head or dense spike**, bracts subtending showy flowers with **ephemeral, usually yellow petals**.

K 3 **C** 3 or (3) **A** 3 or 3+3 or 3+3 staminodes **G** (3), superior.

Restionaceae Restio family (*restio*, L. for rope, cord, in reference to the cordlike stems). 55 genera / 490 species. (Figure 7.69)

The Restionaceae consist of dioecious [rarely monoecious or hermaphroditic], evergreen, perennial herbs. The underground **stems** are rhizomatous or stoloniferous, the erect culms photosynthetic, hollow or solid. The **leaves** are simple, unifacial, spiral, with a usually open sheath, usually eligulate, often reduced to sheaths in mature plants, sometimes caducous. The **inflorescence** is a solitary flower or an aggregate of spikelets, in variously branched groups, each group sometimes subtended by bracts (spathes). The spikelets consist of an axis bearing 1 ∞ flowers, each flower subtended by 1 [2] bract, lowermost bracts often sterile; male and female spikelets may be similar or dimorphic. The **flowers** are small, unisexual, actinomorphic, hypogynous. The **perianth** is biseriate, homochlamydeous, 3+3 [0 2+0 2], apotepalous. The **tepals**

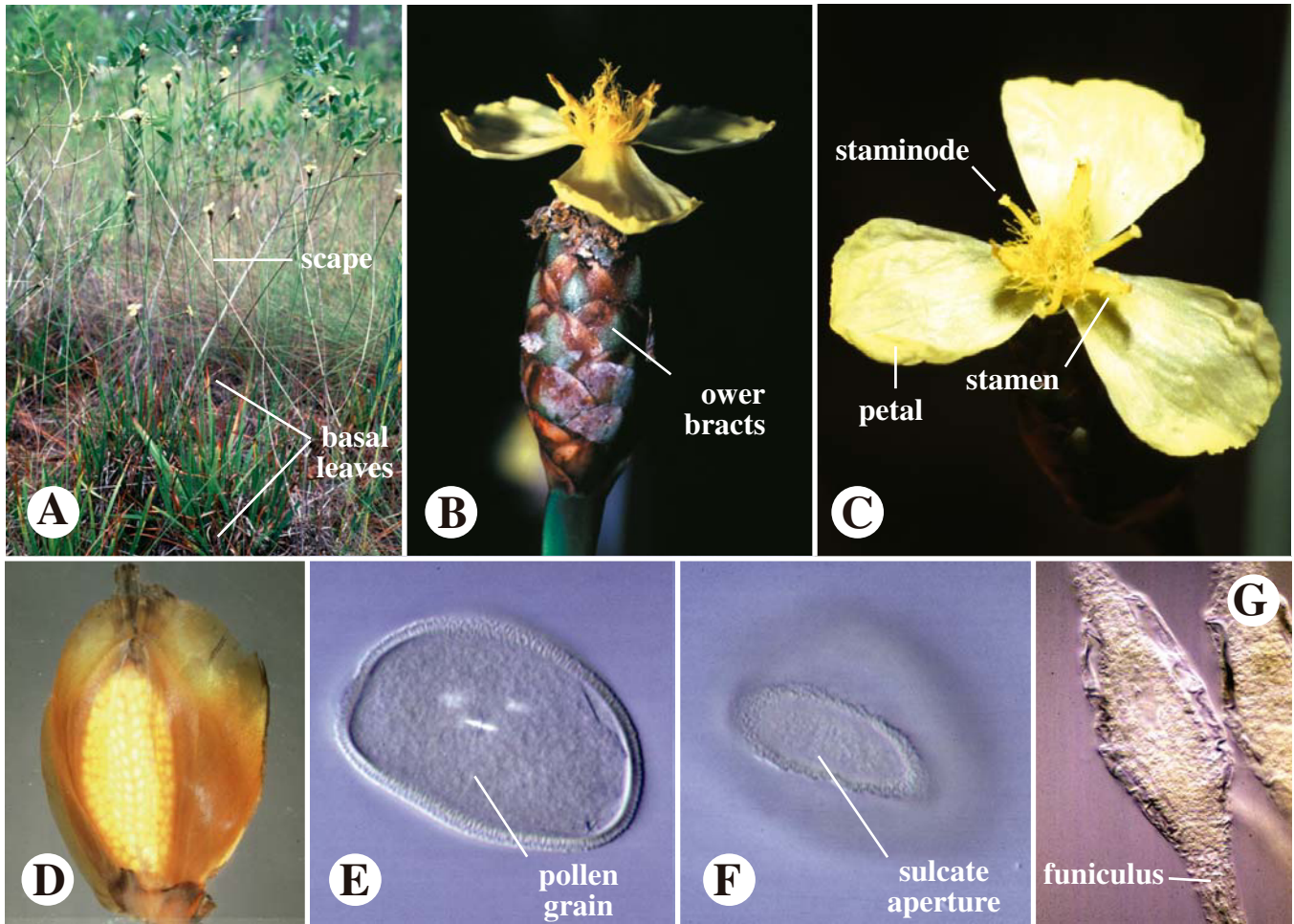


FIGURE 7.68 POALES. Xyridaceae. *Xyris* sp. **A.** Whole plant. **B.** In-orecence, a scapose, bracteate spike. **C.** Flower, face view; note three stamens and three staminodes. **D.** Fruit, with persistent sepals and bract. **E,F.** Pollen grain in midsagittal (**E**) and surface (**F**) views, the latter showing sulcate aperture type. **G.** Orthotropous ovules.

are membranous to indurate. The **stamens** are 3 [1-4], antipetalous, distinct [rarely connate]. **Anthers** are usually unilocular, bisporangiate, and monothechal, longitudinally and usually introrse in dehiscence. The **pollen** is monoulcerate at release. The **gynoecium** is syncarpous or unicarpelous, with a superior ovary, 3 [1 or 2] carpels, and 3 [1 or 2] locules. The **style(s)** are 1-3. **Placentation** is apical-axile; **ovules** are orthotropous, bitegmic, solitary. **Nectaries** are absent. The **fruit** is an achene, nut, or capsule. The **seeds** are endospermous and sometimes have an elaiosome, functioning in ant dispersal. Flowers are wind-pollinated.

Members of the Restionaceae are a major component of fynbos or heath vegetation with distributions in the southern hemisphere, especially South Africa and Australia. Economic importance includes local use as thatching and brooms. See Linder et al. (1998, 2000) for recent phylogenetic analyses and treatments of the family.

The Restionaceae are distinctive in being *perennial, rhizomatous*, mostly *dioecious herbs* with photosynthetic erect stems, *leaves reduced to sheaths*, an inflorescence of solitary flowers or variously branched *spikelets*, and *small, unisexual, wind-pollinated* flowers with usually *monothechal, bisporangiate* anthers.

P 3+3 [0-2+0-2] **A** 3 [1-4] **G** (3) [1-(2)], superior.

Poaceae (Gramineae) Grass family (from *poa*, Greek name for a grass). 668 genera / 9500 species. (Figures 7.70-7.73)

The Poaceae consist of perennial or annual, hermaphroditic, monoecious, or dioecious herbs or (in the bamboos) trees. The **roots** are adventitious, often endomycorrhizal. The underground **stems** of perennials are rhizomes or stolons, the erect stems (termed **culms**) are hollow (solid at the nodes), often cespitose, woody-textured in some (e.g., bamboos). The **leaves** are simple, basal or cauline, distichous, rarely

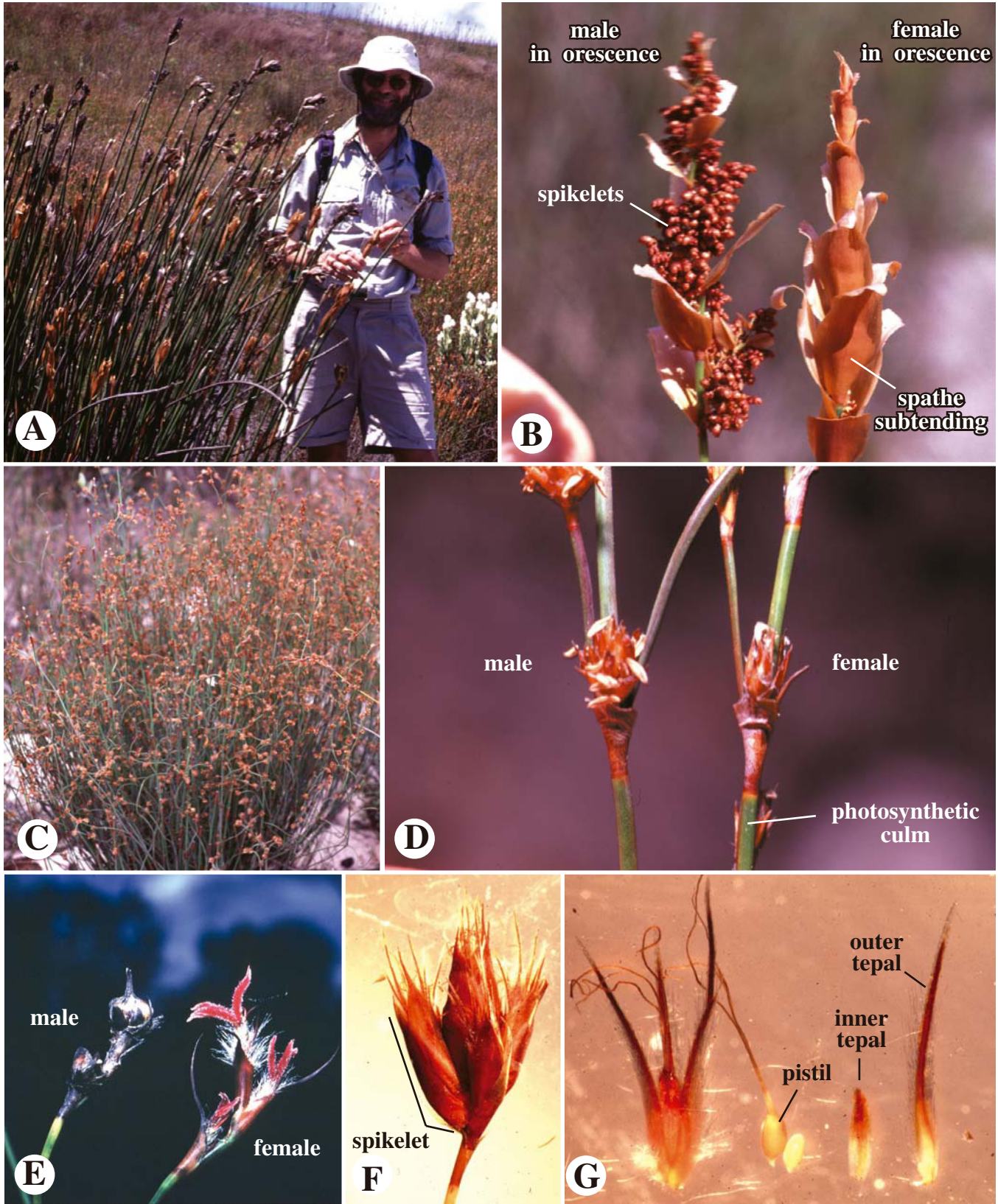


FIGURE 7.69 POALES. Restionaceae. A. *Chondropetalum mucronatum*, whole plant. B. *Elegia* sp., close-up of male and female in orescences. C,D. *Empodisma minus*. C. Whole plants. D. Close-up of male and female in orescences. E. *Leptocarpus* sp. male and female in orescences. F,G. *Leptocarpus aristatus*. G. In orescence. H. Female flowers, whole flower (left) and dissected gynoecium and tepals (right).

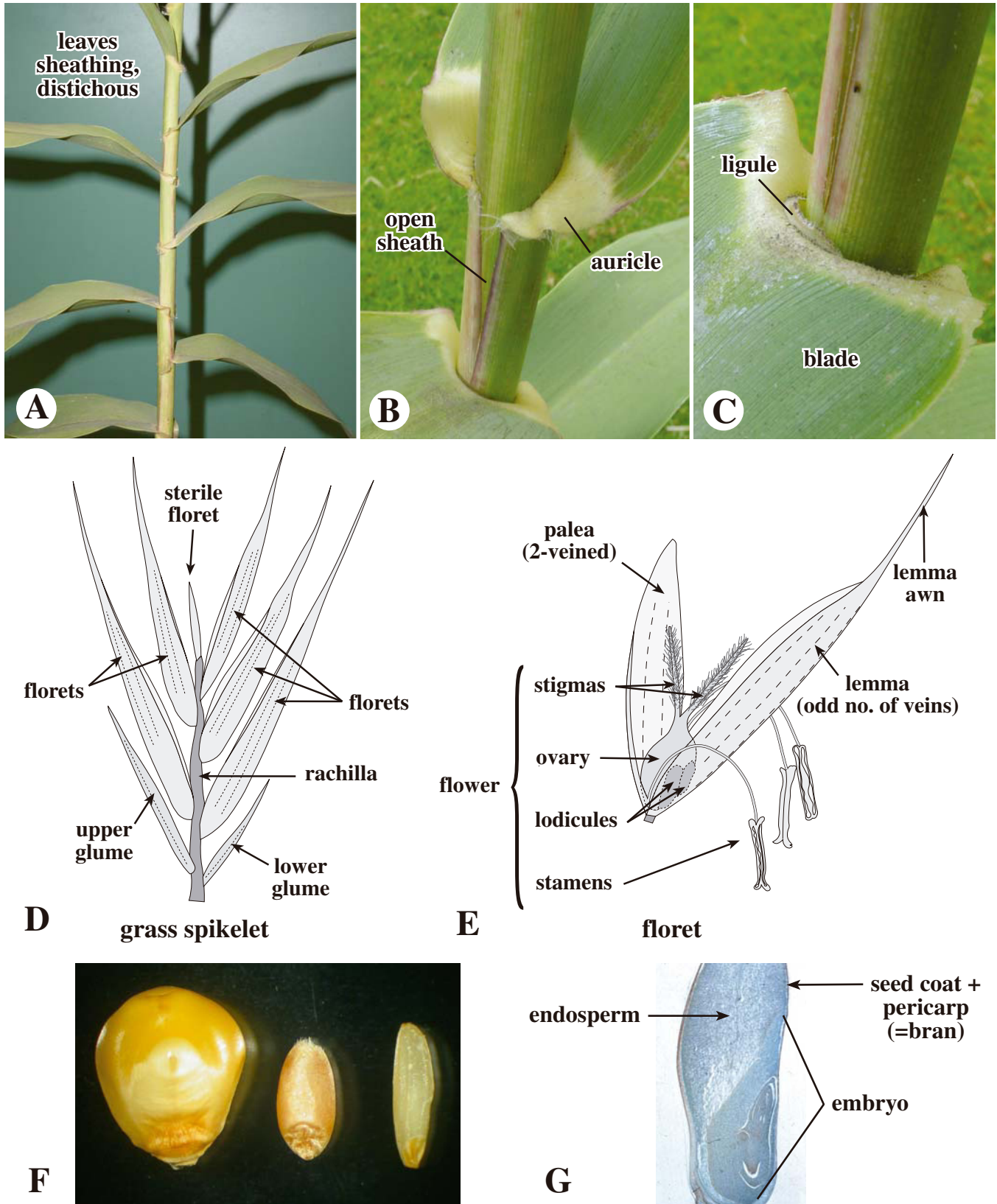


FIGURE 7.70 POALES. Poaceae. **A.** Aerial shoot, showing distichous leaf arrangement. **B.** Close-up of leaf, showing open leaf sheath and basal auricles of blade. **C.** Ligule, at adaxial junction of sheath and blade. **D.** Diagram of grass spikelet, consisting of an axis (rachilla) bearing two, basal glumes (one or both absent or modified in some taxa) plus 1 ∞ florets. **E.** Floret, consisting of a short, lateral branch bearing two bracts, lemma and palea, plus a flower; a given floret may be sterile or unisexual in some taxa. **F.** Grains of (left to right) *Zea mays* (corn), *Triticum aestivum* (bread wheat), and *Oryza sativa* (rice). **G.** Grain of *Zea mays* in longitudinal section, showing embryo, endosperm, and fusion product of seed coat and pericarp.

spiral, with a usually open, basal sheath; the leaf blade is bifacial, parallel-veined, often auriculate at base, and typically ligulate, with a ligule at junction of sheath and blade (resembling a sheathlike structure or tuft of trichomes); in the bamboos the first leaves are scalelike and sheathing, followed by branches that bear photosynthetic leaves; in the bamboos and other taxa, a stalklike pseudo-petiole is present between the sheath and blade. The **inflorescence** consists of terminal or axillary spikelets (more properly termed grass spikelets), these aggregated in secondary inflorescences of spikes, racemes, panicles, or glomerules; the spikelets are sessile or stalked (the spikelet stalk termed a pedicel), and are whorled, opposite, or distichous (on 1 or 2 sides) on the inflorescence axes; the grass spikelet itself consists of an axis (termed the rachilla) bearing distichous parts: two basal bracts (termed glumes, the lower one called the first glume, the upper the second glume, sometimes modified or absent) and one or more florets; each floret consists of a minute lateral axis with two additional bracts (termed the lemma and palea) and a flower; the lemma is the lower and larger bract, typically with an odd number of veins (nerves); the palea is the upper, smaller bract, which has 2 veins and is partially enveloped or enclosed by the lemma. A bristlelike **awn** may

be present at the apex of glumes or lemmas. The **flowers** are bisexual or unisexual, sessile, and hypogynous. The **perianth** is absent or modified into 2 or 3 **lodicules** (located on the lower side, toward the lemma), which upon swelling function to open the floret by separating the lemma from palea. The **stamens** are 2 or 3. **Anthers** are basifixed-versatile, usually sagittate at the base, generally pendulous on elongate filaments, dithecal, and longitudinal in dehiscence. The **pollen** is monoporate. The **gynoecium** is syncarpous, with a superior ovary, 2-3 carpels, and 1 locule. The **stigmas** are 2 or 3, usually plumose. **Placentation** is basal; **ovules** are orthotropous to anatropous, usually bitegmic, 1 per ovary. **Nectaries** are absent. The **fruit** is a caryopsis (grain). The **seeds** are endospermous. Plants are wind pollinated.

The Poaceae are worldwide in distribution. The grasses are perhaps the most economically important group of plants, containing the agricultural grains (vital food and alcoholic beverage sources), including barley (*Hordeum*), corn (*Zea*), oats (*Avena*), rice (*Oryza*), rye (*Secale*), wheat (*Triticum*), and others, as well as important forage and grazing plants. Members of the family are also important components of many ecosystems, such as grasslands and savannahs. See the Grass Phylogeny Working Group (2001) for information on

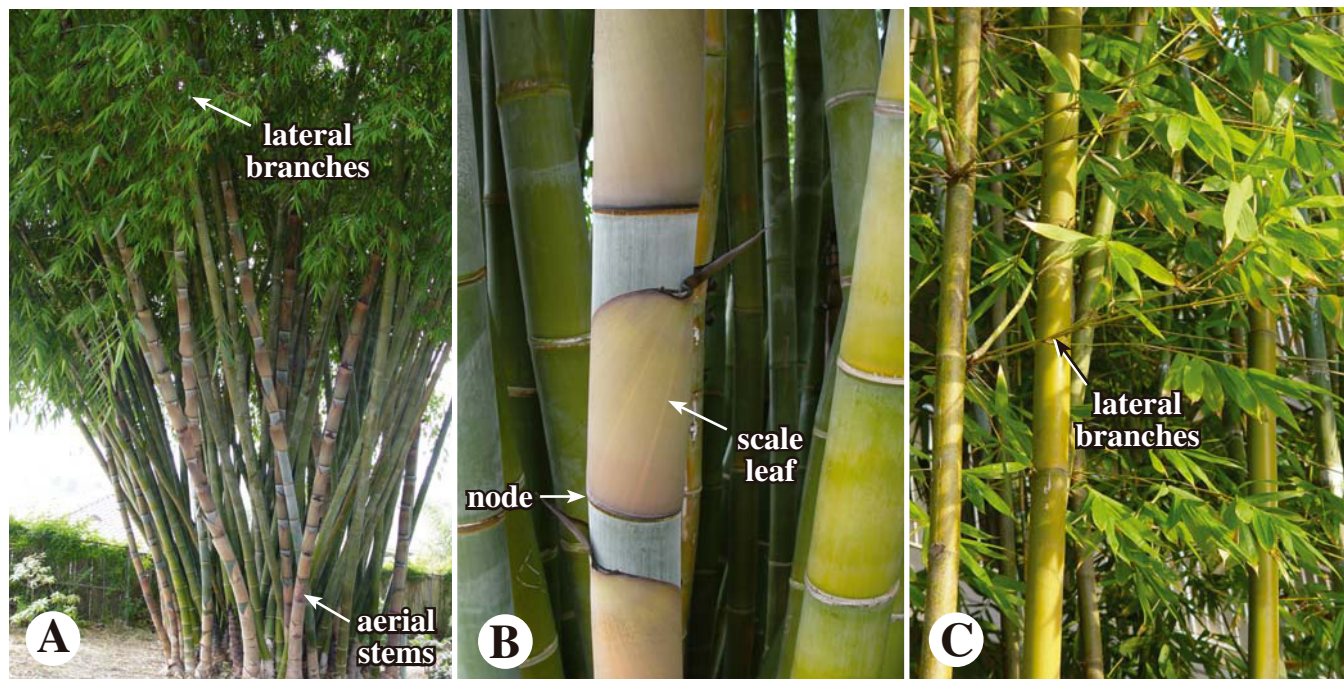


FIGURE 7.71 POALES. Poaceae. Bamboos, having stout, woody, aerial stems that bear large, nonphotosynthetic, scale leaves and upper, lateral branches with photosynthetic leaves. **A,B.** *Dendrocalamus giganteus*. **C.** Bamboo showing lateral branches with photosynthetic leaves.

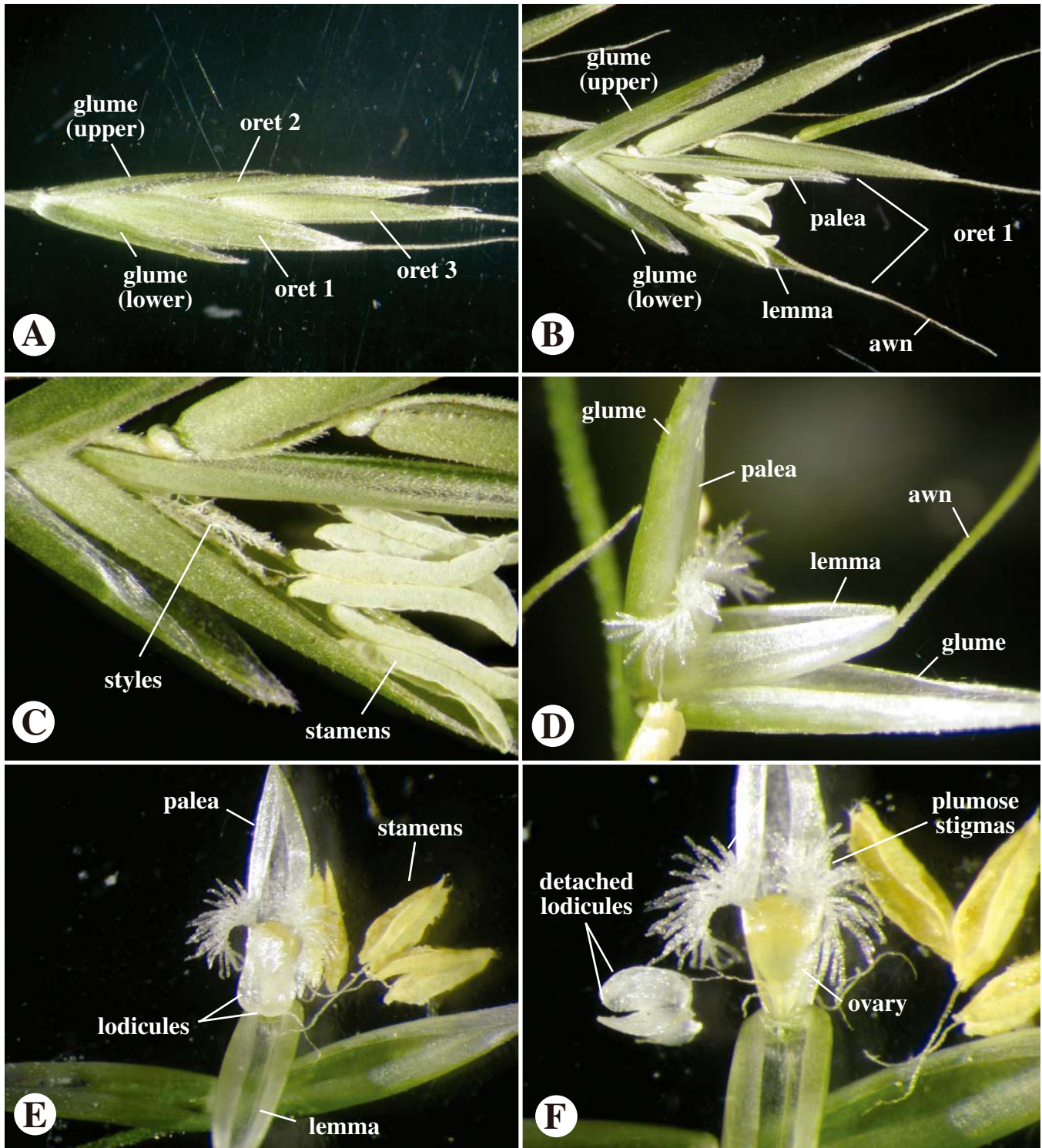


FIGURE 7.72 POALES. Poaceae. Spikelet morphology. A–C. *Elymus glaucus*. A. Spikelet, immature and closed, with two glumes and three orets. B. Mature spikelet, showing palea and awned lemma of open oret. C. Close-up of lower, showing three stamens and styles of ovary. D–F. *Agrostis stolonifera*. D. Open, mature spikelet, showing two glumes and palea and lemma of single oret. E. Floret dissected open, showing three stamens and two lodicules on lemma side of ovary. F. Lodicules removed; note ovary with two plumose styles.

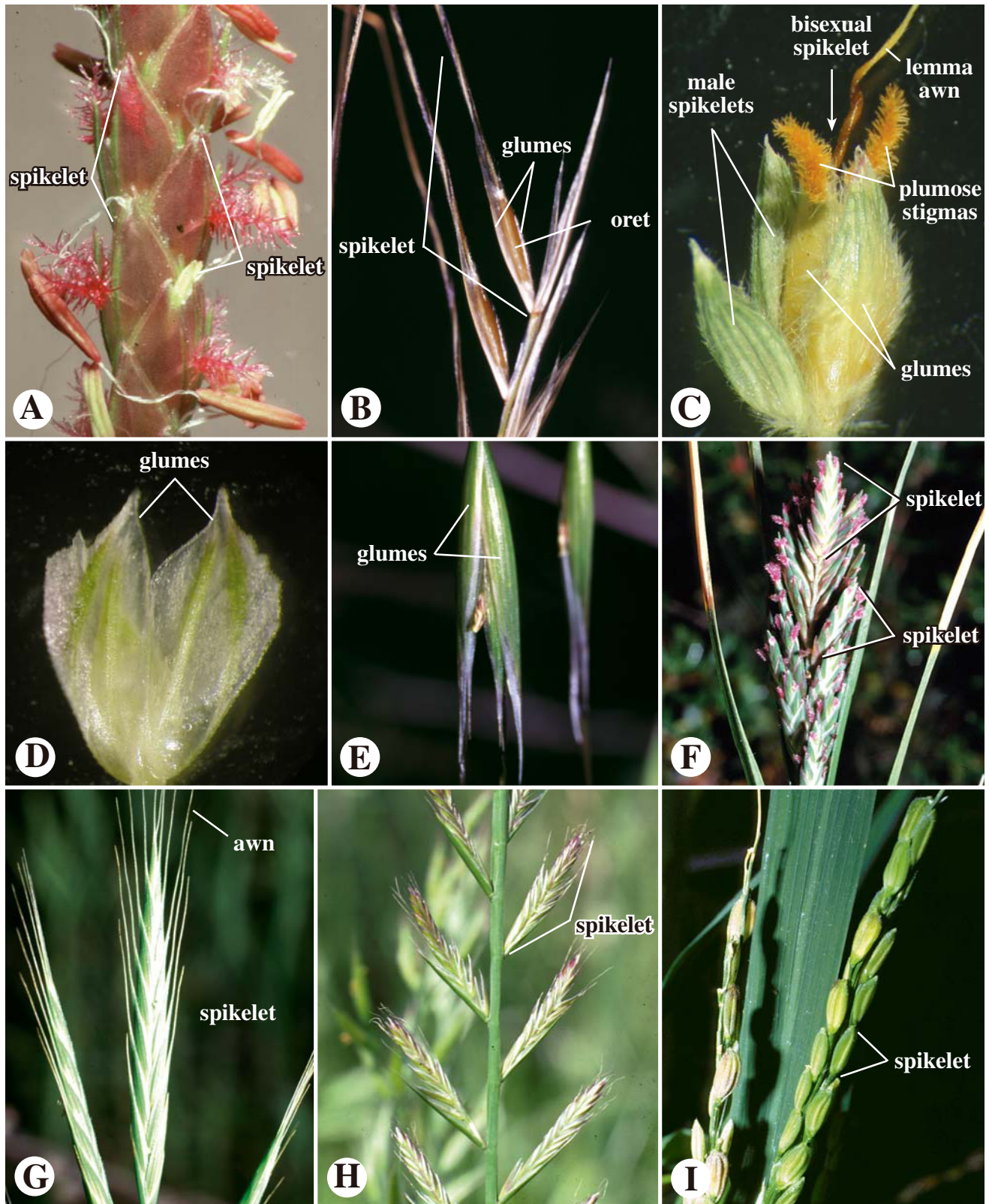


FIGURE 7.73 POALES. Poaceae. Spikelet diversity. **A.** *Cynodon dactylon*, bermuda grass, close-up of spikelets in two rows, each bearing exerted, pendulous anthers and red, bifurcate styles. **B.** *Nasella pulchra*, needle grass, having one oret per spikelet. **C.** *Sorghum bicolor*, in which two, reduced, male spikelets are grouped with a single, bisexual spikelet. **D.** *Phalaris minor*; spikelet with prominent glumes and one oret. **E.** *Avena barbata*, pendulous spikelet. **F.** *Distichlis spicata*, in orescence of female plant, a condensed panicle of spikelets. **G.** *Brachypodium distachyon*, spikelet with numerous, awned orets. **H.** *Lolium multi orum*, in orescence a spike of distichously arranged spikelets. **I.** *Oryza sativa*, rice.

character evolution in the Poaceae plus recommendations regarding infrafamilial classification.

The Poaceae are distinctive in being herbs (trees in the bamboos) with *hollow-pithed stems* and *open-sheathed, distichous leaves* with a *ligule* at inner junction with blade; the inflorescence is a *grass spikelet*, typically with 2 basal bracts (*glumes*) on a central axis and 1 ∞ *florets*, each

consisting of a short lateral axis with 2 bracts (a lower, odd-veined *lemma* and an upper, 2-veined *palea*) and a flower, the flower with perianth reduced to usually 2-3 *lodicules*, usually 2-3 pendulous stamens, and a single 2-3-carpellate, 1-ovuled ovary with 2-3 *plumose stigmas*, the fruit a *caryopsis (grain)*.

P 2-3 [-6+] lodicules **A** 2-3 [1] **G** (2-3), superior.

REVIEW QUESTIONS

GENERAL

1. What is the Angiosperm Phylogeny Group system of classification and what ranking does it utilize?
2. What are the major groups of basal angiosperms?
3. Why have the traditional dicots been abandoned as a taxonomic group?
4. What is a floral formula? What are the symbols used in floral formulas?
5. What is a floral diagram and what does it represent?

NONMONOCOT GROUPS

6. Name the family and species of what is currently thought to represent the most basal lineage of angiosperms.
7. Name the diagnostic characteristics of the Amborellaceae. Do these necessarily represent ancestral angiosperm features?
8. How does the Nymphaeales compare with the Amborellaceae in: plant habit, flower sex, perianth arrangement, stamen number and type, gynoecial fusion type, and ovary position?
9. What anatomical feature is characteristic of the family Illiciaceae? How is this family different from and similar to the Amborellaceae and Nymphaeales?
10. For the Ceratophyllaceae name the plant habitat, plant habit, leaf arrangement and morphology, and economic importance.
11. What distinctive anther dehiscence occurs in the Lauraceae?
12. Name two economically important members of the Lauraceae.
13. Name at least two families of the Magnoliales.
14. For the Annonaceae, what is distinctive about the leaf arrangement and endosperm structure?
15. What is distinctive about the receptacle and gynoecial fusion of the Magnoliaceae?
16. What is the fruit type of the Magnoliaceae?
17. Name at least three families of the Piperales.
18. What is the etymology of *Aristolochia*?
19. What are the diagnostic features of the Aristolochiaceae?
20. How does the Piperaceae differ from the Aristolochiaceae?
21. What is an economically important member of the Piperaceae?
22. How does the Saururaceae differ from the Piperaceae?

MONOCOTS: BASAL LINEAGES

23. Name and describe the major apomorphies of the monocots.
24. Name the order, family, and genus of the most basal lineage of monocots.
25. How does *Acorus* differ from the Araceae in: leaf structure; seed nutritive tissue; crystal type?
26. What is the leaf venation of members of the Araceae?
27. What is the inflorescence type of the Araceae?
28. Name an economically important member of the Araceae.
29. What is a major apomorphy of the Asparagales?
30. What is a cytological apomorphy of the Agavaceae?
31. What is the ovary position of the Agavaceae?
32. What is a chemical apomorphy of the Alliaceae?

33. Name two economically important members of the Alliaceae.
34. How are members of the Asphodelaceae distinguished? What is their distribution?
35. Name and define the leaf structure of the Iridaceae.
36. What is the range of inflorescence morphology of the Iridaceae?
37. What is the floral formula of the Iridaceae?
38. How many species occur in the orchid family?
39. For the Orchidaceae, name the ovary position, placentation, and name for specialized androecium.
40. What is a gynostemium? What are other names for this structure?
41. What orchid is used as a food flavoring and what part of the plant is utilized?

MONOCOTS: COMMELINIDS

42. Name and describe the major chemical apomorphy of the Commelinid monocots.
43. Name an apomorphy of the palms.
44. What are the two acceptable scientific names of the palm family?
45. For the Arecaceae, what is the: flower sex, ovary position, fruit type?
46. What is the seed nutrition of the Commelinid monocots, minus the Arecales and Dasypogonaceae?
47. Name three apomorphies of the Zingiberales.
48. What is the leaf arrangement and plant sex of the the Musaceae?
49. What is the scientific name of banana?
50. What leaf arrangement apomorphy unites all of the Zingiberales, minus the Musaceae?
51. Name two apomorphies that unite the clade Cannaceae + Costaceae + Marantaceae + Zingiberaceae.
52. What, in reality, are the showy petaloid structures of this group?
53. Name two economically important members of the Zingiberaceae.
54. What is the specialized anther type of the Cannaceae and Marantaceae?
55. Name three families of the Commelinales.
56. Name one or more apomorphies for these families.
57. Name two apomorphies of the Bromeliaceae.
58. What is the ovary position of the Bromeliaceae?
59. What special ecological adaptations do various bromeliads have?
60. Name an economically important bromeliad.
61. What two apomorphies that unite the Poales, minus the Bromeliaceae, Mayacaceae, and Rapateaceae?
62. What is the flower sex of the Sparganiaceae and Typhaceae?
63. How do these two families differ?
64. What pollen and leaf apomorphies may unite the Cyperaceae, Thurniaceae (including Prionaceae) and Juncaceae?
65. How do the Cyperaceae and Juncaceae families differ in inflorescence, perianth, and fruit type?
66. How do the Eriocaulaceae and Xyridaceae differ in flower sex? pollen aperture type?
67. Name the characteristics and distribution of the Restionaceae.
68. What are the two scientific names for the grass family?
69. How does the leaf arrangement and attachment of grasses differ from that of sedges?
70. What is the structural difference between a sedge spikelet and a grass spikelet?
71. What is a: grass spikelet? floret? glume? lemma? palea? lodicule?
72. What is the fruit type of the grasses?
73. Name the generic and common names of five economically important grain crops.

EXERCISES

1. Select a family of angiosperms treated in this chapter and learn everything you can about it. Perform a literature search (e.g., family name + systematics) on journal articles published in the last 5 years. Consult family descriptions, recent data on phylogenetic relationships, and information on intrafamilial groupings.

2. From this same family, collect living material of an exemplar. Describe this species in detail, using the character list of Appendix 1 as a guide (see Chapter 9). Illustrate the vegetative and reproductive parts (see Appendix 2).
3. Assimilate all of your information in a written report and computerized slide presentation to present to an audience.

REFERENCES FOR FURTHER STUDY

GENERAL REFERENCES ON ANGIOSPERM RELATIONSHIPS AND EVOLUTION

- Angiosperm Phylogeny Group. 1998. An ordinal classification for the families of flowering plants. *Annals of the Missouri Botanical Garden* 85: 531 553.
- Angiosperm Phylogeny Group. 2003. An update of the Angiosperm Phylogeny Group classification for the orders and families of flowering plants: APG II. *Botanical Journal of the Linnean Society* 141: 399 436.
- Barkman, T. J., G. Chenery, J. R. McNeal, J. Lyons-Weiler, W. J. Ellisens, G. Moore, A. D. Wolfe, and C. W. dePamphilis. 2000. Independent and combined analyses of sequences from all three genomic compartments converge on the root of flowering plant phylogeny. *Proceedings of the National Academy of Sciences of the United States of America* 97: 13166 13171.
- Behnke, H.-D. 1972. Sieve-tube plastids in relation to Angiosperm systematics an attempt towards a classification by ultrastructural analysis. *Botanical Review* 38: 155 197.
- Borsch, T., K. W. Hilu, D. Quandt, V. Wilde, C. Neinhuis, and W. Barthlott. 2003. Noncoding plastid trnT-trnF sequences reveal a well resolved phylogeny of basal angiosperms. *Journal of Evolutionary Biology* 16: 558 576.
- Chase, M. W., D. E. Soltis, R. G. Olmstead, D. Morgan, D. H. Les, B. D. Mishler, M. R. Duvall, R. A. Price, H. G. Hills, Y.-L. Qiu, K. A. Kron, J. H. Rettig, E. Conti, J. D. Palmer, J. R. Manhart, K. J. Sytsma, H. J. Michaels, W. J. Kress, K. G. Karol, W. D. Clark, M. Hedrøn, B. S. Gaut, R. K. Jansen, K.-J. Kim, C. F. Wimpee, J. F. Smith, G. R. Furnier, S. H. Strauss, Q.-Y. Xiang, G. M. Plunkett, P. S. Soltis, S. M. Swensen, S. E. Williams, P. A. Gadek, C. J. Quinn, L. E. Eguiarte, E. Golenberg, G. H. Learn Jr., S. W. Graham, S. C. H. Barrett, S. Dayanandan, and V. A. Albert. 1993. Phylogenetics of seed plants: an analysis of nucleotide sequences from the plastid gene *rbcL*. *Annals of the Missouri Botanical Garden* 80: 528 580.
- Chase, M. W., M. F. Fay, and V. Savolainen. 2000. Plant Systematics: A half-century of progress (1950 2000) and future challenges Higher-level classification in the angiosperms: New insights from the perspective of DNA sequence data. *Taxon* 49: 685 704.
- Doyle, J. A., and P. K. Endress. 2000. Morphological phylogenetic analysis of basal angiosperms: Comparison and combination with molecular data. *International Journal of Plant Sciences* 161: S121 S153.
- Graham, S. W. and R. G. Olmstead. 2000a. Evolutionary significance of an unusual chloroplast DNA inversion found in two basal angiosperm lineages. *Current Genetics* 37: 183 188.
- Graham, S. W., and R. G. Olmstead. 2000b. Utility of 17 chloroplast genes for inferring the phylogeny of the basal angiosperms. *American Journal of Botany* 87: 1712 1730.
- Hilu, K. W., T. Borsch, K. Muller, D. E. Soltis, P. S. Soltis, V. Savolainen, M. W. Chase, M. P. Powell, L. A. Alice, R. Evans, H. Sauquet, C. Neinhuis, T. A. B. Slotta, J. G. Rohwer, C. S. Campbell, and L. W. Chatrou. 2003. Angiosperm phylogeny based on *matK* sequence information. *American Journal of Botany* 90: 1758 1766.
- Hutchinson, J. 1973. *The Families of Flowering Plants; Arranged According to a New System Based on their Probable Phylogeny*, 3rd edition. Clarendon Press, Oxford.
- Kubitzki, K., J. G. Rohwer, and V. Bittrich (eds). 1993. *The Families and Genera of Vascular Plants. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families*. Springer, Berlin.
- Kubitzki, K., H. Huber, P. J. Rudall, P. S. Stevens, and T. Stuetzel (eds.). 1998a. *The Families and Genera of Vascular Plants III: Flowering Plants Monocotyledons Liliaceae (except Orchidaceae)*. Springer-Verlag, Berlin.
- Kubitzki, K., H. Huber, P. J. Rudall, P. S. Stevens, and T. Stuetzel (eds.). 1998b. *The Families and Genera of Vascular Plants IV: Flowering Plants Monocotyledons Alismatanae and Commelinanae (except Gramineae)*. Springer-Verlag, Berlin.
- Kubitzki, K., and C. Bayer. 2002. *The Families and Genera of Vascular Plants. V. Flowering Plants. Dicotyledons: Malvales, Capparales, and Non-betalain Caryophyllales*. Springer, Berlin.
- Kubitzki, K. 2004. *The Families and Genera of Vascular Plants. VI. Flowering Plants. Dicotyledons. Ceslastrales, Oxalidales, Rosales, Cornales, Ericales*. Springer, Berlin.
- Mathews, S., and M. J. Donoghue. 1999. The root of angiosperm phylogeny inferred from duplicate phytochrome genes. *Science* 286: 947 950.
- Mathews, S., and M. J. Donoghue. 2000. Basal angiosperm phylogeny inferred from duplicate phytochromes A and C. *International Journal of Plant Sciences* 161: S41 S55.
- Nandi, O. I., M. W. Chase, and P. K. Endress. 1998. A combined cladistic analysis of angiosperms using *rbcL* and non-molecular data sets. *Annals of the Missouri Botanical Garden* 85: 137 212.
- Parkinson, C. L., K. L. Adams, and J. D. Palmer. 1999. Multigene analyses identify the three earliest lineages of extant flowering plants. *Current Biology* 9: 1485 1488.
- Qui, Y.-L., M. W. Chase, S. B. Hoot, E. Conti, P. R. Crane, K. J. Sytsma, C. R. Parks. 1998. Phylogenetics of Hamamelidae and their allies: parsimony analyses of nucleotide sequences of the plastid gene *rbcL*. *International Journal of Plant Sciences* 159: 891 905.

- Qiu, Y.-L., J. Lee, F. Bernasconi-Quadroni, D. E. Soltis, P. S. Soltis, M. Zanis, E. A. Zimmer, Z. Chen, V. Savolainen, and M. W. Chase. 1999. The earliest angiosperms: evidence from mitochondrial, plastid and nuclear genomes. *Nature* 402: 404-407.
- Qiu, Y.-L., J. Lee, F. Bernasconi-Quadroni, D. E. Soltis, P. S. Soltis, M. Zanis, E. A. Zimmer, Z. Chen, V. Savolainen, and M. W. Chase. 2000. Phylogeny of basal angiosperms: analyses of five genes from three genomes. *International Journal of Plant Sciences* 161: S3-S27.
- Ronse Decraene, L.-P., P. S. Soltis, and D. E. Soltis. 2003. Evolution of floral structures in basal angiosperms. *International Journal of Plant Sciences* 164: S329-S363.
- Savolainen, V., M. W. Chase, S. B. Hoot, C. M. Morton, D. E. Soltis, C. Bayer, M. F. Fay, A. Y. de Bruijn, S. Sullivan, and Y.-L. Qiu. 2000. Phylogenetics of flowering plants based on combined analysis of plastid *atpB* and *rbcL* sequences. *Systematic Biology* 49: 306-362.
- Soltis, D. E., P. S. Soltis, D. L. Nickrent, L. A. Johnson, W. J. Hahn, S. B. Hoot, J. A. Sweere, R. K. Kuzoff, K. A. Kron, M. W. Chase, S. M. Swensen, E. A. Zimmer, S.-M. Chaw, L. J. Gillespie, W. J. Kress, and K. J. Sytsma. 1997. Angiosperm phylogeny inferred from 18S ribosomal DNA sequences. *Annals of the Missouri Botanical Garden* 84: 1-49.
- Soltis, D. E., P. S. Soltis, M. W. Chase, M. E. Mort, D. C. Albach, M. Zanis, V. Savolainen, W. H. Hahn, S. B. Hoot, M. F. Fay, M. Axtell, S. M. Swensen, L. M. Prince, W. J. Kress, K. C. Nixon, J. S. Farris. 2000. Angiosperm phylogeny inferred from 18S rDNA, *rbcL*, and *atpB* sequences. *Botanical Journal of the Linnean Society* 133(4): 381-461.
- Takhtajan, A. L. 1991. *Evolutionary Trends in Flowering Plants*. Columbia University Press, New York.
- Takhtajan, A. L. 1997. *Diversity and Classification of Flowering Plants*. Columbia University Press, New York.
- Thorne, Robert F. 2000. The classification and geography of the flowering plants: Dicotyledons of the class Angiospermae: (Subclasses Magnoliidae, Ranunculidae, Caryophyllidae, Dilleniidae, Rosidae, Asteridae, and Lamiidae). *Botanical Review* 66(4): 441-647.
- Zanis, M. J., D. E. Soltis, P. S. Soltis, S. Mathews, and M. J. Donoghue. 2002. The root of the angiosperms revisited. *Proceedings of the National Academy of Sciences of the United States of America* 99: 6848-6853.
- Zanis, M. J., P. S. Soltis, Y. L. Qiu, E. Zimmer, and D. E. Soltis. 2003. Phylogenetic analyses and perianth evolution in basal angiosperms. *Annals of the Missouri Botanical Garden* 90: 129-150.

WEB PAGE SITES

- Reveal, J. L. (2001 onwards). *Indices Nominum Supragenericorum Plantarum Vascularium*. Alphabetical Listing by Family of Validly Published Suprageneric Names.
<http://www.inform.umd.edu/PBIO/fam/revfam.html>
 An index of accepted family names.
- Royal Botanic Garden Vascular Plant Families and Genera Database
<http://www.rbghkew.org.uk/data/vascplnt.html>
 A searchable database taken from Vascular Plant Families and Genera, compiled by R. K. Brummitt and published by the Royal Botanic Gardens, Kew, in 1992.
- Stevens, P. F. (2001 onwards). Angiosperm Phylogeny Website. Version 5, May 2004.
<http://www.mobot.org/MOBOT/research/APweb>
 An excellent graphic representation of the Angiosperm Phylogeny Group classification, up-to-date cladograms, family characteristics, references, and apomorphies.

ANGIOSPERM FAMILY DESCRIPTIONS AND ILLUSTRATIONS

- Cronquist, A. 1981. *An Integrated System of Classification of Flowering Plants*. Columbia University Press, New York.
- Heywood, V. H. 1985. *Flowering Plants of the World*. Prentice Hall, Englewood Cliffs, NJ.
- Mabberly, D. J. 1997. *The Plant-Book*, 2nd ed. Cambridge University Press, Cambridge, UK.
- Takhtajan, A. 1997. *Diversity and Classification of Flowering Plants*. Columbia University Press, New York.
- Zomlefer, W. B. 1994. *Guide to Flowering Plant Families*. University of North Carolina Press, Chapel Hill.

NONMONOCOTYLEDON REFERENCES

- Doyle, J. A., and A. le Thomas. 1994. Cladistic analysis and pollen evolution in Annonaceae. *Acta Botanica Gallica* 141: 149-170.
- Doyle, J. A., and A. le Thomas. 1996. Phylogenetic analysis and character evolution in Annonaceae. *Bulletin of the Museum of Natural History, Paris*, 4e s^{er}e [sect. B, Adansonia] 18: 279-334.
- Endress, P. K., and A. Igersheim. 1997. Gynoecium diversity and systematics of the Laurales. *Botanical Journal of the Linnean Society* 125: 93-168.
- Kelly, L. M., and F. Gonzalez. 2003. Phylogenetic relationships in Aristolochiaceae. *Systematic Botany* 28: 236-249.
- Kim, S., C.-W. Park, Y.-D. Kim, and Y. Suh. 2001. Phylogenetic relationships in family Magnoliaceae inferred from *ndhf* sequences. *American Journal of Botany* 88: 717-728.
- Kim, S., D. E. Soltis, P. S. Soltis, M. J. Zanis, and Y. Suh. 2004. Phylogenetic relationships among early-diverging eudicots based on four genes: Were the eudicots ancestrally woody? *Molecular Phylogenetics and Evolution* 31: 16-30.
- Les, D. H. 1993. Ceratophyllaceae. In: Kubitzki, K., J. G. Rohwer, and V. Bittrich (eds.), *The Families and Genera of Vascular Plants*. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families. Pp. 246-249. Springer, Berlin.

- Les, D. H., E. L. Schneider, D. J. Padgett, P. S. Soltis, D. E. Soltis, and M. Zanis. 1999. Phylogeny, classification and floral evolution of water lilies (Nymphaeaceae; Nymphaeales): a synthesis of non-molecular, *rbcL*, *matK*, and 18s rDNA data. *Systematic Botany* 24: 28-46.
- Meng, S.-W., A. W. Douglas, D.-Z. Li, Z.-D. Chen, H.-X. Liang, and J.-B. Yang. 2003. Phylogeny of Saururaceae based on morphology and five regions from three plant genomes. *Annals of the Missouri Botanical Garden* 90: 592-602.
- Nickrent, D. L., A. Blarer, Y.-L. Qiu, D. E. Soltis, P. S. Soltis, and M. Zanis. 2002. Molecular data place Hydnoraceae with Aristolochiaceae. *American Journal of Botany* 89: 1809-1817.
- Renner, S. S. 1999. Circumscription and phylogeny of the Laurales: Evidence from molecular and morphological data. *American Journal of Botany* 86: 1301-1315.
- Renner, S. S., and A. S. Chanderbali. 2000. What is the relationship among Hernandiaceae, Lauraceae, and Monimiaceae, and why is this question so difficult to answer? *International Journal of Plant Sciences* 161: S109-S119.
- Rohwer, J. G. 1993. Lauraceae. In: Kubitzki, K., J. G. Rohwer, and V. Bittrich (eds.), *The Families and Genera of Vascular Plants. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families*. Pp. 366-390. Springer, Berlin.
- Sauquet, H., J. A. Doyle, T. Scharaschkin, T. Borsch, K. W. Hilu, L. Chatrou, and A. Le Thomas. 2003. Phylogenetic analysis of Magnoliaceae and Myristicaceae based on multiple data sets: implications for character evolution. *Botanical Journal of the Linnean Society* 142: 125-186.
- Schneider, E. L. and P. S. Williamson. 1993. Nymphaeaceae. In: Kubitzki, K., J. G. Rohwer, and V. Bittrich (eds.), *The Families and Genera of Vascular Plants. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families*. Pp. 486-493. Springer, Berlin.
- Thien, L. B., T. L. Sage, T. Jaffre, P. Bernhardt, V. Pontieri, P. H. Weston, D. Malloch, H. Azuma, S. W. Graham, M. A. McPherson, H. S. Rai, R. F. Sage, and J.-L. Dupre. 2003. The population structure and floral biology of *Amborella trichopoda* (Amiborellaceae). *Annals of the Missouri Botanical Garden* 90: 466-490.
- Williamson, P. S. and E. L. Schneider. 1993. Cabombaceae. In: Kubitzki, K., J. G. Rohwer, and V. Bittrich (eds.), *The Families and Genera of Vascular Plants. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families*. Pp. 157-160. Springer, Berlin.

MONOCOTYLEDON REFERENCES

- Andersson, L. 1998a. Musaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 296-300. Springer, Berlin.
- Andersson, L. 1998b. Strelitziaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 451-454. Springer, Berlin.
- Arber, A. 1925. *Monocotyledons; A Morphological Study*. The University Press, Cambridge, England.
- Asmussen, C. B., Baker, W. J., and Dransfield, J. 2000. Phylogeny of the palm family (Arecaceae) based on *rps16* intron and *trnL-trnF* plastid DNA sequences. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 525-535. CSIRO, Collingwood.
- Balslev, H. 1998. Juncaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 252-259. Springer, Berlin.
- Barrett, S. C. H., and S. W. Graham. 1997. Adaptive radiation in the aquatic plant family Pontederiaceae: Insights from phylogenetic analysis. In: Givnish, T. J., and K. J. Sytsma (eds.), *Molecular Evolution and Adaptive Radiation*. Pp. 225-258. Cambridge University Press, Cambridge, UK.
- Behnke, H.-D. 2000. Forms and sizes of sieve-element plastids and evolution of the monocotyledons. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 163-188. CSIRO, Collingwood.
- Bogler, D. J., and B. B. Simpson. 1995. A Chloroplast DNA Study of the Agavaceae. *Systematic Botany* 20: 191.
- Bogler, D. J., and B. B. Simpson. 1996. Phylogeny of Agavaceae based on ITS rDNA sequence variation. *American Journal of Botany* 83: 1225.
- Bogler, D. J., C. Pires, and J. Francisco-Ortega. 2005 (in press). Phylogeny of Agavaceae based on *ndhF*, *rbcL*, and ITS sequences: implications of molecular data for classification. In: Columbus, J. T., E. A. Friar, J. M. Porter, L. M. Prince, and M. G. Simpson (eds.), *Monocots: Comparative Biology and Evolution*, Rancho Santa Ana Botanic Garden, Claremont, California.
- Bogner, J., and S. J. Mayo. 1998. Acoraceae. In: Kubitzki, K., H. Huber, P. J. Rudall, P. S. Stevens, and T. Stuetzel (eds.), *The Families and Genera of Vascular Plants IV: Flowering Plants Monocotyledons Alismatanae and Commelinanae (except Gramineae)*. Springer-Verlag, New York.
- Bremer, K. 2002. Gondwanan evolution of the grass alliance of families (Poales). *Evolution* 56: 1374-1387.
- Caddick, L. R., P. J. Rudall, P. Wilkin, T. A. J. Hedderson, and M. W. Chase. 2002a. Phylogenetics of Dioscoreales based on combined analyses of morphological and molecular data. *Botanical Journal of the Linnean Society* 138: 123-144.
- Caddick, L. R., P. Wilkin, P. J. Rudall, T. A. J. Hedderson, and M. W. Chase. 2002b. Yams reclassified: A recircumscription of Dioscoreaceae and Dioscoreales. *Taxon* 51: 103-114.
- Cameron, K. M. 2004. Utility of plastid *psaB* gene sequences for investigating intrafamilial relationships within Orchidaceae. *Molecular Phylogenetics and Evolution* 31: 1157-1180.
- Cameron, K. M., and M. W. Chase. 2000. Nuclear 18S rDNA sequences of Orchidaceae confirm the subfamilial status and circumscription of Vaniilloideae. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 457-464. CSIRO, Collingwood.
- Cameron, K. M., M. W. Chase, W. M. Whitten, P. J. Kores, D. C. Jarrell, V. A. Albert, T. Yukawa, H. G. Hills, and D. H. Goldman. 1999. A phylogenetic analysis of the Orchidaceae: evidence from *rbcL* nucleotide. *American Journal of Botany* 86: 208-224.

- Chase, M. W., D. E. Soltis, P. S. Soltis, P. J. Rudall, M. F. Fay, W. J. Hahn, S. Sullivan, J. Joseph, M. Molvray, P. J. Kores, T. J. Givnish, K. J. Sytsma, and J. C. Pires. 2000a. Higher-level systematics of the monocotyledons: an assessment of current knowledge and a new classification. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. CSIRO, Collingwood.
- Chase, M. W., A. Y. De Bruijn, A. V. Cox, G. Reeves, P. J. Rudall, M. A. T. Johnson, and L. E. Eguiarte. 2000b. Phylogenetics of Asphodelaceae (Asparagales): an analysis of plastid rbcL and trnL-F DNA sequences. *Annals of Botany* 86: 935-952.
- Chen, Y.-Y., D.-Z. Li, and H. Wang. 2002. Infrageneric phylogeny and systematic position of the Acoraceae inferred from ITS, 18S and rbcL sequences. *Acta Botanica Yunnanica* 24: 699-706.
- Columbus, J. T., E. A. Friar, J. M. Porter, L. M. Prince, and M. G. Simpson (eds.). 2005 (in press). *Monocots: Comparative Biology and Evolution*. 2 vols. Rancho Santa Ana Botanic Garden, Claremont, CA.
- Cook, C. D. K. 1998. Pontederiaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 395-403. Springer, Berlin.
- Cox, P. A., K.-L. Huynh, and B. C. Stone. 1995. Evolution and systematics of Pandanaceae. In: Rudall, P. J., P. J. Cribb, D. F. Cutler, and C. J. Humphries (eds.), *Monocotyledons: Systematics and Evolution*. Pp. 663-684. Royal Botanic Gardens, Kew.
- Dahlgren, R., H. T. Clifford, and P. F. Yeo. 1985. *The Families of the Monocotyledons: Structure, Evolution, and Taxonomy*. Springer-Verlag, New York.
- Davis, J. I., D. W. Stevenson, G. Petersen, O. Seberg, L. M. Campbell, J. V. Freudenstein, D. H. Goldman, C. R. Hardy, F. A. Michelangeli, and M. P. Simmons. 2004. A phylogeny of the monocots, as inferred from rbcL and atpA sequence variation, and a comparison of methods for calculating jackknife and bootstrap values. *Systematic Botany* 29: 467-510.
- Drabkova, L., J. Kirschner, O. Seberg, G. Petersen, and C. Vlcek. 2003. Phylogeny of the Juncaceae based on rbcL sequences, with special emphasis on *Luzula* DC. and *Juncus* L. *Plant systematics and evolution = Entwicklungsgeschichte und Systematik der Pflanzen* 240: 133-148.
- Dransfield, J. and N. W. Uhl. 1998. Palmae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 306-388. Springer, Berlin.
- Duvall, M. R., M. T. Clegg, M. W. Chase, W. D. Clark, W. J. Kress, H. G. Hills, L. E. Eguiarte, J. F. Smith, B. S. Gaut, E. A. Zimmer, and G. H. Learn Jr. 1993. Phylogenetic hypotheses for the monocotyledons constructed from rbcL sequence data. *Annals of the Missouri Botanical Garden* 80: 607-619.
- Evans, T. M., K. J. Sytsma, R. B. Faden, and T. J. Givnish. 2003. Phylogenetic relationships in the Commelinaceae: II. A cladistic analysis of rbcL sequences and morphology. *Systematic Botany* 28: 270-292.
- Faden, R. B. 1998. Commelinaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 109-127. Springer, Berlin.
- Fay, M. F., and M. W. Chase. 1996. Resurrection of Themidaceae for the *Brodiaea* alliance, and recircumscription of Alliaceae, Amaryllidaceae and Agapanthoideae. *Taxon* 45: 441-451.
- Fay, M. F., P. J. Rudall, S. Sullivan, K. L. Stobart, A. Y. de Bruijn, G. Reeves, F. Qamaruz-Zaman, W.-P. Hong, J. Joseph, W. J. Hahn, J. G. Conran, and M. W. Chase. 2000. Phylogenetic studies of Asparagales based on four plastid DNA regions. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 360-371. CSIRO, Collingwood.
- French, J. C., M. Chung, and Y. Hur. 1995. Chloroplast DNA phylogeny of Ariflorae. In: Rudall, P. J., P. J. Cribb, D. F. Cutler, and C. J. Humphries (eds.), *Monocotyledons: Systematics and Evolution*. Pp. 255-275. Royal Botanic Gardens, Kew.
- Givnish, T. J., T. M. Evans, J. C. Pires, and K. J. Sytsma. 1999. Polyphyly and convergent morphological evolution in Commelinales and Commelinidae: Evidence from rbcL sequence data. *Molecular Phylogenetics and Evolution* 12: 360-385.
- Givnish, T. J., K. C. Millam, P. E. Berry, and K. J. Sytsma. 2005. Phylogeny, adaptive radiation, and historical biogeography of Bromeliaceae inferred from ndhF sequence data. In: Columbus, J. T., E. A. Friar, C. W. Hamilton, J. M. Porter, L. M. Prince, and M. G. Simpson (eds.), *Monocots: Comparative Biology and Evolution*. 2 vols. Rancho Santa Ana Botanic Garden, Claremont, CA.
- Goetghebeur, P. 1998. Cyperaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 141-189. Springer, Berlin.
- Goldblatt, P., J. C. Manning, and P. Rudall. 1998. Iridaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae)*. Pp. 295-333. Springer, Berlin.
- Grass Phylogeny Working Group. 2001. Phylogeny and subfamilial classification of the grasses (Poaceae). *Annals of the Missouri Botanical Garden* 88: 373-457.
- Grayum, M. H. 1990. Evolution and phylogeny of the Araceae. *Annals of the Missouri Botanical Garden* 77: 628-697.
- Hahn, W. J. 2002. A molecular phylogenetic study of the Palmae (Arecaceae) based on atpB, rbcL and 18S nrDNA sequences. *Systematic Botany* 51: 92-112.
- Hayashi, K., and Kawano, S. 2000. Molecular systematics of *Lilium* and allied genera (Liliaceae): Phylogenetic relationships among *Lilium* and related genera based on the rbcL and matK gene sequence data. *Plant Species Biology* 15: 73-93.
- Haynes, R. R., D. H. Les, and L. B. Holm-Nielsen. 1998. Alismataceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae)*. Pp. 11-18. Springer, Berlin.
- Hopper, S. D., M. G. Fay, M. Rossetto, and M. W. Chase. 1999. A molecular phylogenetic analysis of the bloodroot and kangaroo paw family, Haemodoraceae: taxonomic, biogeographic and conservation implications. *Botanical Journal of the Linnean Society* 131: 285-299.
- Huber, H. 1998a. Dioscoreaceae. Pp. 216-235. In: *The Families and Genera of Flowering Plants. IV. Flowering Plants, Monocotyledons: Liliaceae (except Orchidaceae)* (Kubitzki, K. (ed.), Springer-Verlag, Berlin, Heidelberg, New York.

- Huber, H. 1998b. Trichopodaceae. Pp. 441 444. In: The Families and Genera of Flowering Plants. IV. Flowering Plants, Monocotyledons: Liliaceae (except Orchidaceae) (Kubitzki, K. (ed.), Springer-Verlag, Berlin, Heidelberg, New York.
- Kirchoff, B. K. 2003. Shape matters: Hofmeister's rule, primordium shape, and flower orientation. *International Journal of Plant Sciences* 164: 505 517.
- Kral, R. 1998. Xyridaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 461 469. Springer, Berlin.
- Kress, W. J., L. M. Prince, W. J. Hahn, and E. A. Zimmer. 2001. Unraveling the evolutionary radiation of the families of the Zingiberales using morphological and molecular evidence. *Systematic Biology* 50: 926 944.
- Kress, W. J., L. M. Prince, and K. J. Williams. 2002. The phylogeny and a new classification of gingers: evidence from molecular data. *American Journal of Botany* 89: 1682 1696.
- Kubitzki, K. 1998c. Cannaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 103 105. Springer, Berlin.
- Kubitzki, K. 1998d. Typhaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 457 460. Springer, Berlin.
- Larsen, K., J. M. Lock, H. Maas, and P. J. M. Maas. 1998. Zingiberaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 474 495. Springer, Berlin.
- Les, D. H., and Haynes, R. R. 1995. Systematics of subclass Alismatidae: a synthesis of approaches. In: Rudall, P. J., P. J. Cribb, D. F. Cutler, and C. J. Humphries (eds.), *Monocotyledons: Systematics and Evolution*, Vol. 2. Pp. 353 377. Royal Botanic Gardens, Kew.
- Les, D. H., M. A. Cleland, and M. Waycott. 1997. Phylogenetic studies in Alismatidae, II: evolution of marine angiosperms (seagrasses) and hydrophyly. *Systematic Botany* 22: 443 463.
- Lewis, C. E., and J. J. Doyle. 2001. Phylogenetic utility of the nuclear gene malate synthase in the palm family (Arecaceae). *Molecular Phylogenetics and Evolution* 19: 409 420.
- Linder, H. P., B. G. Briggs, and L. A. S. Johnson. 1998. Restionaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 425 444. Springer, Berlin.
- Linder, H. P., B. G. Briggs, and L. A. S. Johnson. 2000. Restionaceae: A morphological phylogeny. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 653 660. CSIRO, Collingwood.
- Mayo, S. J., J. Bogner, and P. C. Boyce. 1998. Araceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 26 73. Springer, Berlin.
- Meerow, A. W., and D. A. Snijman. 1998. Amaryllidaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 83 110. Springer, Berlin.
- Meerow, A. W., M. F. Fay, C. L. Guy, Q.-B. Li, F. Q. Zaman, and M. W. Chase. 1999. Systematics of Amaryllidaceae based on cladistic analysis of plastid *rbcL* and *trnL-F* sequence data. *American Journal of Botany* 86: 1325 1345.
- Meerow, A. W., M. F. Fay, M. W. Chase, C. L. Guy, Q.-B. Li, D. A. Snijman, and S. L. Yang. 2000. Phylogeny of Amaryllidaceae: Molecules and morphology. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 372 386. CSIRO, Collingwood.
- Michelangeli, F. A., J. I. Davis, and D. W. Stevenson. 2003. Phylogenetic relationships among Poaceae and related families as inferred from morphology, inversions in the plastid genome, and sequence data from the mitochondrial and plastid genomes. *American Journal of Botany* 90: 93 106.
- Patterson, T. B. and T. J. Givnish. 2002. Phylogeny, concerted convergence, and phylogenetic niche conservatism in the core Liliales: Insights from *rbcL* and *ndhF* sequence data. *Evolution* 56: 233 252.
- Pires, J. C., and K. J. Sytsma. 2002. A phylogenetic evaluation of a biosystematic framework: *Brodiaea* and related petaloid monocots (Themidaceae). *American Journal of Botany* 89: 1342 1359.
- Rahn, K. 1998a. Alliaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 70 78. Springer, Berlin.
- Rahn, K. 1998b. Themidaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 436 440. Springer, Berlin.
- Reeves, G., M. W. Chase, P. Goldblatt, P. Rudall, M. F. Fay, A. V. Cox, B. Lejeune, and T. Souza-Chies. 2001. Molecular systematics of Iridaceae: evidence from four plastid DNA regions. *American Journal of Botany* 88: 2074 2087.
- Rudall, P. 2003. Unique floral structures and iterative evolutionary themes in Asparagales: insights from a morphological cladistic analysis. *Botanical Review* 68: 488 509.
- Rudall, P., K. L. Stobart, W.-P. Hong, J. G. Conran, C. A. Furness, G. C. Kite, and M. W. Chase. 2000. Consider the lilies: Systematics of Liliales. In: Wilson, K. L., and D. A. Morrison (eds.), *Monocots: Systematics and Evolution*. Pp. 347 359. CSIRO, Collingwood.
- Rudall, P., P. J. Cribb, D. F. Cutler, and C. Humphries (eds). 1995. *Monocotyledons: Systematics and Evolution*. Royal Botanic Gardens Kew [London].
- Simpson, M. G. 1990. Phylogeny and classification of the Haemodoraceae. *Annals of the Missouri Botanical Garden* 77: 722 784.
- Simpson, M. G. 1998. Haemodoraceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 212 222. Springer, Berlin.
- Simpson, D. A., C. A. Furness, T. R. Hodkinson, A. M. Muasya, and M. W. Chase. 2003. Phylogenetic relationships in Cyperaceae subfamily Mapanioideae inferred from pollen and plastid DNA sequence data. *American Journal of Botany* 90: 1071 1086.

- Smith, G. F., and B.-E. v. Wyk. 1998. Asphodelaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 130-140. Springer, Berlin.
- Smith, L. B., and W. Till. 1998. Bromeliaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 74-99. Springer, Berlin.
- Stevenson, D. W., J. I. Davis, J. V. Freudenstein, C. R. Hardy, M. P. Simmons, and C. D. Specht. 2000. A phylogenetic analysis of the monocotyledons based on morphological and molecular character sets, with comments on the placement of *Acorus* and Hydatellaceae. In: Wilson, K. L., and D. A. Morrison (eds.), Monocots: Systematics and Evolution, CSIRO, Collingwood.
- Stone, B. C., K.-L. Huynh, and H.-H. Poppendieck. 1998. Pandanaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 397-403. Springer, Berlin.
- St tzel, T. 1998. Eriocaulaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. IV. Flowering Plants. Monocotyledons. Alismatanae and Commelinanae (except Gramineae). Pp. 197-207. Springer, Berlin.
- Tamura, M. N. 1998a. Calochortaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 164-172. Springer, Berlin.
- Tamura, M. N. 1998b. Liliaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 343-353. Springer, Berlin.
- Verhoek, S. 1998. Agavaceae. In: Kubitzki, K. (ed.), The Families and Genera of Vascular Plants. III. Flowering Plants. Monocotyledons. Liliaceae (except Orchidaceae). Pp. 60-70. Springer, Berlin.
- Wilson, K. L., and D. A. Morrison (eds). 2000. Monocots: Systematics and Evolution. CSIRO, Collingwood, VIC, Australia.

8

DIVERSITY AND CLASSIFICATION OF FLOWERING PLANTS: EUDICOTS

EUDICOTS	228	Cucurbitaceae	259
RANUNCULALES	228	FABALES	262
Berberidaceae	228	Fabaceae (Leguminosae)	262
Papaveraceae	230	Polygalaceae	265
Ranunculaceae	233	FAGALES	265
PROTEALES	233	Betulaceae	265
Nelumbonaceae	233	Fagaceae	265
Platanaceae	235	MALPIGHIALES	269
Proteaceae	235	Euphorbiaceae	269
CORE EUDICOTS	238	Passifloraceae	271
CARYOPHYLLALES	238	Violaceae	272
Aizoaceae	238	OXALIDALES	272
Amaranthaceae	241	Oxalidaceae	272
Cactaceae	244	ROSALES	275
Caryophyllaceae	244	Moraceae	275
Polygonaceae	249	Rhamnaceae	275
SANTALALES	249	Rosaceae	275
Santalaceae	249	EUROSIDS II	278
SAXIFRAGALES	250	BRASSICALES	278
Crassulaceae	250	Brassicaceae (Cruciferae)	280
Saxifragaceae	252	MALVALES	280
ROSIDS	252	Malvaceae s.l.	280
GERANIALES	252	SAPINDALES	286
Geraniaceae	252	Anacardiaceae	286
MYRTALES	254	Rutaceae	286
Melastomataceae	254	Sapindaceae	289
Myrtaceae	256	ASTERIDS	289
Onagraceae	259	CORNALES	291
EUROSIDS I	259	Cornaceae	291
CUCURBITALES	259		

(Continued)

ERICALES	293	SOLANALES	319
Ericaceae	293	Convolvulaceae	319
Fouquieriaceae	295	Solanaceae	319
Polemoniaceae	298		
EUASTERIDS I.	298	EUASTERIDS II.	319
Boraginaceae s.s.	298	APIALES	322
Hydrophyllaceae	301	Apiaceae (Umbelliferae)	322
		Araliaceae	322
GENTIANALES	301	ASTERALES	322
Apocynaceae	301	Asteraceae (Compositae)	326
Gentianaceae	305	Goodeniaceae	331
Rubiaceae	305		
LAMIALES	305	DIPSACALES	331
Acanthaceae	308	Caprifoliaceae s.s.	331
Bignoniaceae	308	Dipsacaceae	335
Lamiaceae (Labiatae)	308	Valerianaceae	335
Scrophulariaceae s.l.	310		
Scrophulariaceae s.s.	310	REVIEW QUESTIONS	335
Orobanchaceae	315	EXERCISES.	339
Phrymaceae	315	REFERENCES FOR FURTHER STUDY.	339
Plantaginaceae s.l.	315		

EUDICOTS

The **eudicots** are a large, monophyletic assemblage of angiosperms, comprising roughly 190,000 described species, or 75% of all angiosperms. The monophyly of eudicots is well supported from molecular data and delimited by at least one palynological apomorphy: a **tricolpate or tricolpate-derived** pollen grain (Figure 8.1). A tricolpate pollen grain is one that has three apertures, equally spaced and approximately parallel to the polar axis of the grain (Figure 8.2; see Chapter 12). Apertures are differentiated regions of the pollen grain wall that may function as the site of pollen tube exitus as well as to allow for expansion and contraction of the pollen grain with changes in humidity (Chapter 12). Tricolpate pollen grains evolved from a **monosulcate** type (having a single distal aperture; Figure 8.2), which is considered to be ancestral in the angiosperms, as well as for many seed plant clades. Many eudicots have pollen grains with more than three apertures, of a great variety of numbers, shapes, and position (constituting important taxonomic characters; see Chapter 12). These are all thought to have been derived from a tricolpate type.

The orders of the eudicots and their included families (after APG II 2003) are listed in Tables 8.1–8.3. Table 8.1 lists the most basal groups, including the families of the Ranunculales, Proteales, Caryophyllales, Santalales, and Saxifragales. Table 8.2 lists orders and families within a large group termed the Rosids. Table 8.3 lists orders and families of another large

group termed the Asterids. See FAMILY DESCRIPTIONS, Chapter 7, for a summary of the format for these. For detailed information on eudicot relationships, see the references on general angiosperm relationships (e.g., Chase et al. 1993, 2000; Graham and Olmstead 2000; Nandi et al. 1998; Soltis et al. 1997, 2000; Hilu et al. 2003; Savolainen et al. 2000a). Also see Hoot and Crane (1995) and Hoot et al. (1999) for an analysis of the basal eudicots and Savolainen et al. (2000b) for a treatment of the eudicots alone.

RANUNCULALES

The Ranunculales, sensu APG II (2003), contain seven families, of which three are described here. See Hoot et al. (1999) for a phylogenetic analysis that includes this order.

Berberidaceae Barberry family (after Barbary, an ancient Arabic name for N. Africa). 15 genera / 670 species. (Figure 8.3)

The Berberidaceae consist of perennial trees, shrubs, or herbs. The **leaves** are spiral [rarely opposite], petiolate with the petiole often flared basally, and either pinnate, ternate, simple, or unifoliolate. The **inflorescence** is a raceme, spike, panicle, cyme, or a solitary, axillary flower. The **flowers** are bisexual, actinomorphic, and hypogynous. The **perianth** is 6–7-seriate with 3 [2,4] parts per whorl, the outer 2 whorls sepaloid, the inner 4–5 whorls petaloid, with the innermost

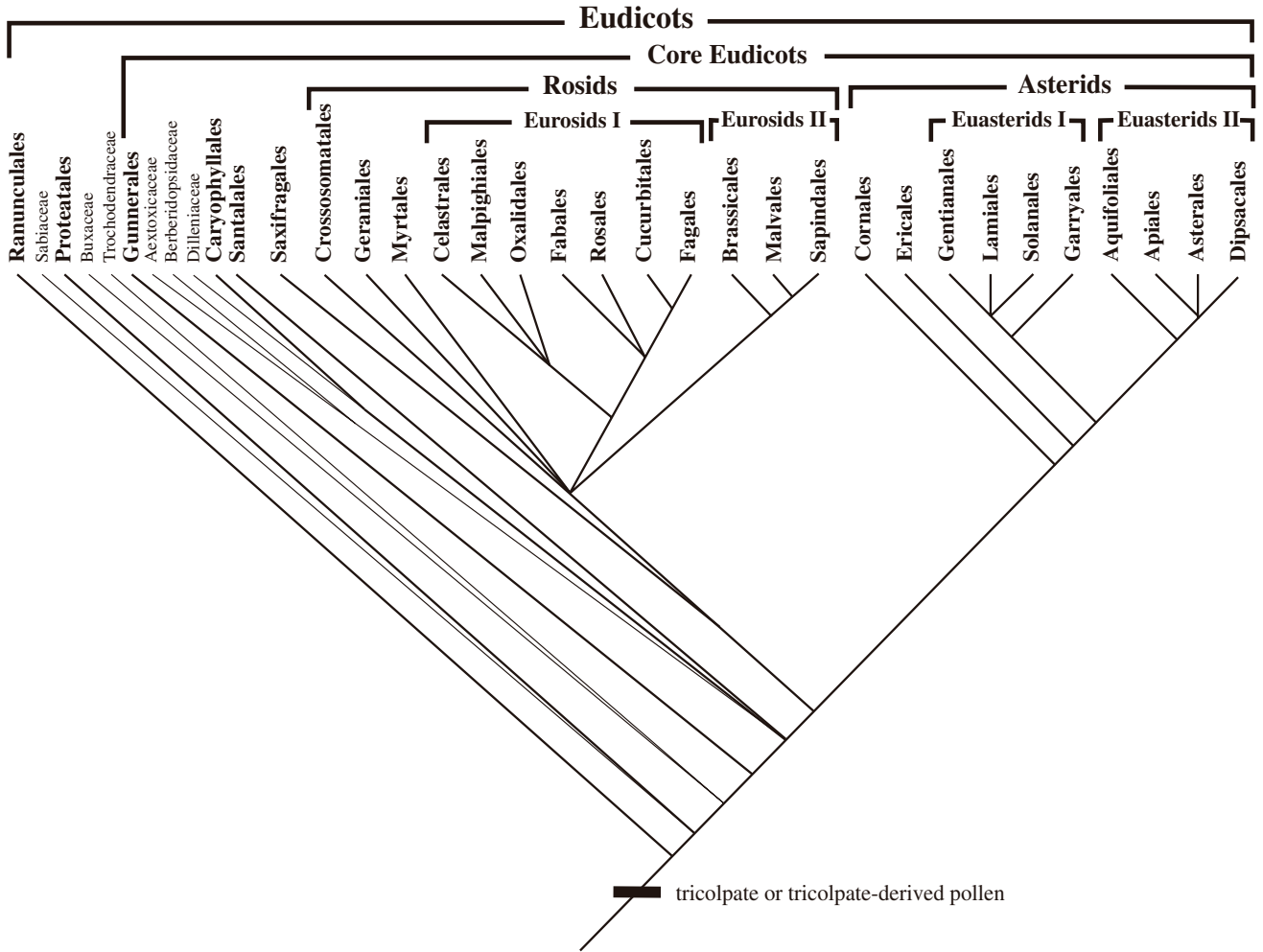


FIGURE 8.1 Cladogram of the orders of the Eudicots (modified from APG II, 2003).

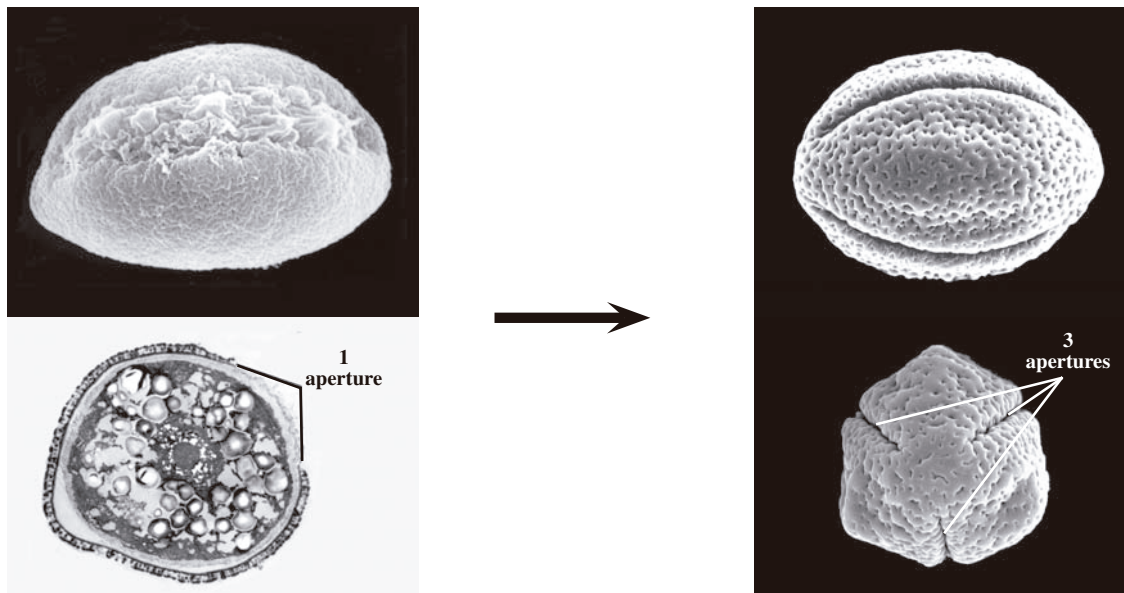


FIGURE 8.2 Transformation from monosulcate to tricolpate pollen grain, the latter an apomorphy of the Eudicots.

TABLE 8.1 Orders and families of the Eudicots, after APG II (2003). Families in bold are described in detail. An asterisk denotes an acceptable deviation from APG II, with brackets indicating the more inclusive family recommended by APG II. A double asterisk indicates a change suggested by Angiosperm Phylogeny Website (Stevens 2001 onwards). See Tables 8.2 and 8.3 for listings in the Rosids and Asterids.

EUDICOTS		
UNPLACED** Apodanthaceae, Balanophoraceae, Cynomoriaceae, Cytinaceae, Hoplestigmataceae, Medusandraceae, Metteniusaceae, Pottingeriaceae BUXALES** Buxaceae Didymelaceae** Sabiaceae TROCHODENDRALES** Trochodendraceae RANUNCULALES Berberidaceae Circaeasteraceae Eupteleaceae Lardizabalaceae Menispermaceae Papaveraceae Ranunculaceae PROTEALES Nelumbonaceae Platanaceae* [Proteaceae] Proteaceae CORE EUDICOTS BERBERIDOPSISIDALES** Aextoxicaceae Berberidopsidaceae DILLENIALES** Dilleniaceae GUNNERALES Gunneraceae Myrothamnaceae**	CARYOPHYLLALES Achatocarpaceae Aizoaceae Amaranthaceae Ancistrocladaceae Asteropeiaceae Barbeuiaceae Basellaceae Cactaceae Caryophyllaceae Didiereaceae Dioncophyllaceae Droseraceae Drosophyllaceae Frankeniaceae Gisekiaceae Halophytaceae** Hectorellaceae** Limeaceae** Molluginaceae Nepenthaceae Nyctaginaceae Phyteneaceae Phytolaccaceae Plumbaginaceae Polygonaceae Portulacaceae Rhabdodendraceae Sarcobataceae Simmondsiaceae Stegnospermataceae Tamaricaceae	SANTALALES Erythralaceae** Olacaceae Opiliaceae Loranthaceae Misodendraceae Santalaceae Schoep aceae** SAXIFRAGALES Altingiaceae Aphanopetalaceae** Cercidiphyllaceae Crassulaceae Daphniphyllaceae Grossulariaceae Haloragaceae Hamamelidaceae Iteaceae Paeoniaceae Penthoraceae** Peridiscaceae** Pterostemonaceae** Saxifragaceae Tetracarpaceae** ROSIDS (see Table 8.2, p. 255) ASTERIDS (see Table 8.3, p. 292)

2-3 of these nectariferous (sometimes interpreted as staminodes). The **stamens** are 6 [4-18], mostly in two whorls, opposite the inner most whorls of petals. **Anthers** are valvular (opening from the base) or longitudinal in dehiscence, and are tetrasporangiate and dithecal. The **gynoecium** is unicarpellous (sometimes interpreted as derived from 2-3 carpels), with a superior ovary, and 1 [2] locules. **Placentation** is marginal; **ovules** are ∞ [1,2] per ovary. The **fruit** is a berry [rarely dry].

Members of the family have a worldwide distribution, especially in north-temperate regions. Economic importance includes cultivated ornamentals, such as *Berberis* and *Mahonia*; *Podophyllum* is reported to be used against testicular cancer; *Berberis vulgaris*, the common barberry, is the alternate host of stem rust of wheat.

The Berberidaceae are distinctive in having flowers with a **multiseriate** perianth (possibly apomorphic for the family) differentiated into **outer sepaloid** and **inner petaloid** parts

(the innermost nectariferous), a **biseriate androecium**, and a **single**, apparently **unicarpellate** pistil.

P ∞ , 6-7-seriate in whorls of 3 [2,4] each **A** 6 [4-18] **G** 1, superior.

Papaveraceae (including Fumariaceae) Popp y family (Latin for poppy). 40 genera / 760 species. (Figure 8.4)

The Papaveraceae consist of annual or perennial herbs, shrubs, or small (sometimes pachycaulis) trees, with milky latex from articulated laticifers in some taxa. The **leaves** are spiral to subopposite, usually lobed to divided or dissected, exstipulate. The **inflorescence** is a solitary flower or cyme. The **flowers** are bisexual, actinomorphic, zygomorphic, or biradial, hypogynous [rarely perigynous]. The **perianth** is dichlamydeous, in 3 [2,4] series, a hypanthium absent [rarely present]. The **calyx** is uniseriate and aposepalous to basally synsepalous, with 2 [3], usually caducous sepals. The **corolla** is biseriata, apopetalous, of 2+2 or 3+3 [16] petals, sometimes

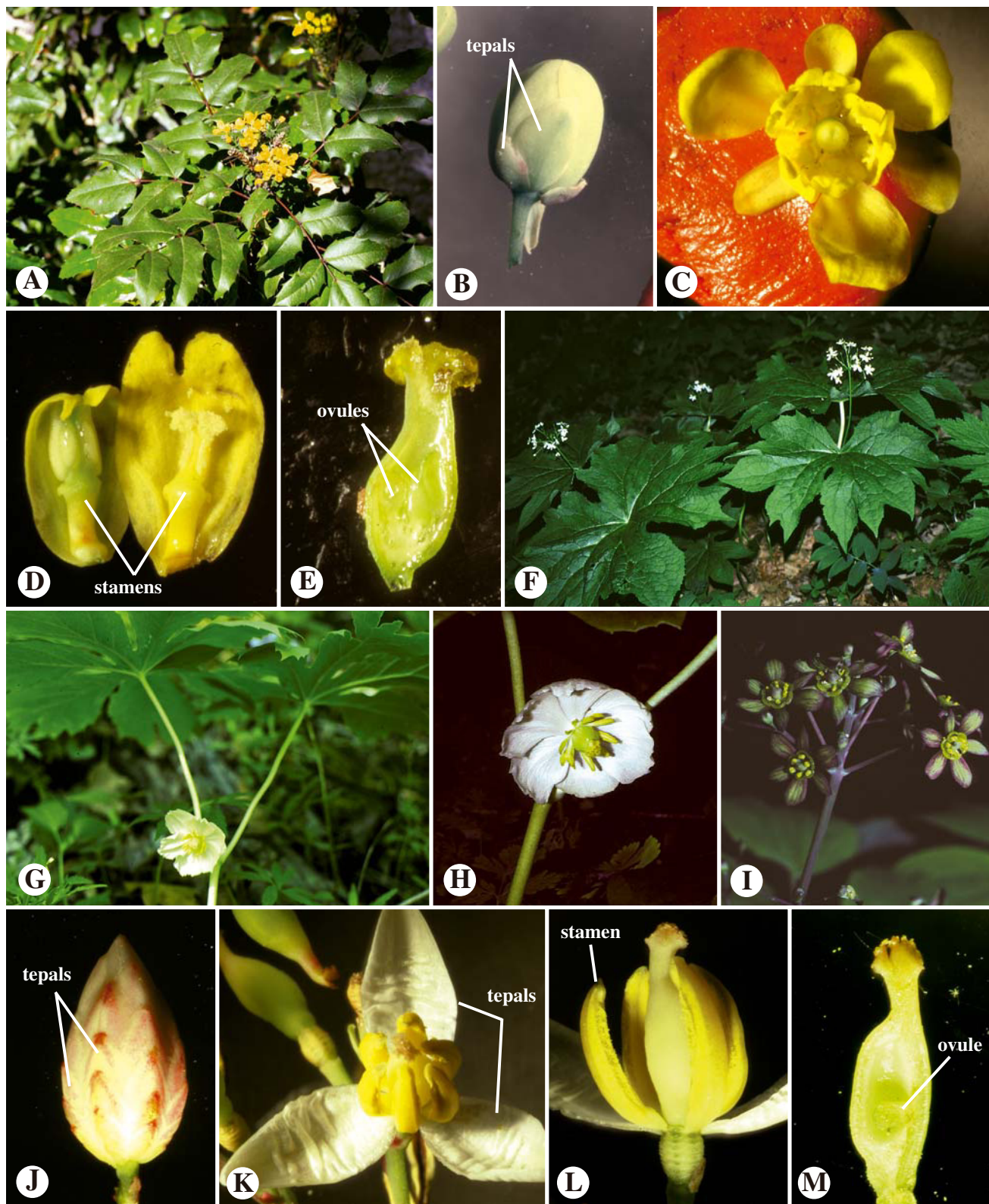


FIGURE 8.3 RANUNCULALES. Berberidaceae. A–E. *Berberis* sp., barberry. A. Whole plant, with pinnately compound leaves. B. Flower bud, showing numerous tepals. C. Open flower. D. Tepals with opposing stamens. E. Pistil, longitudinal section, showing marginal placentation. F. *Diphylleia cymosa*, umbrella-leaf; whole plant with peltate, palmately cleft leaves. G, H. *Podophyllum peltatum*, mayflower. G. Plant, showing peltate leaves. H. Solitary flower. I. *Caulophyllum thalictroides*, blue cohosh; cymose inflorescence. J–M. *Nandina domestica*. J. Flower bud, showing multiserial perianth. K. Open flower. L. Flower close-up, showing pistil and stamens. M. Ovary longitudinal section, showing marginal placentation.

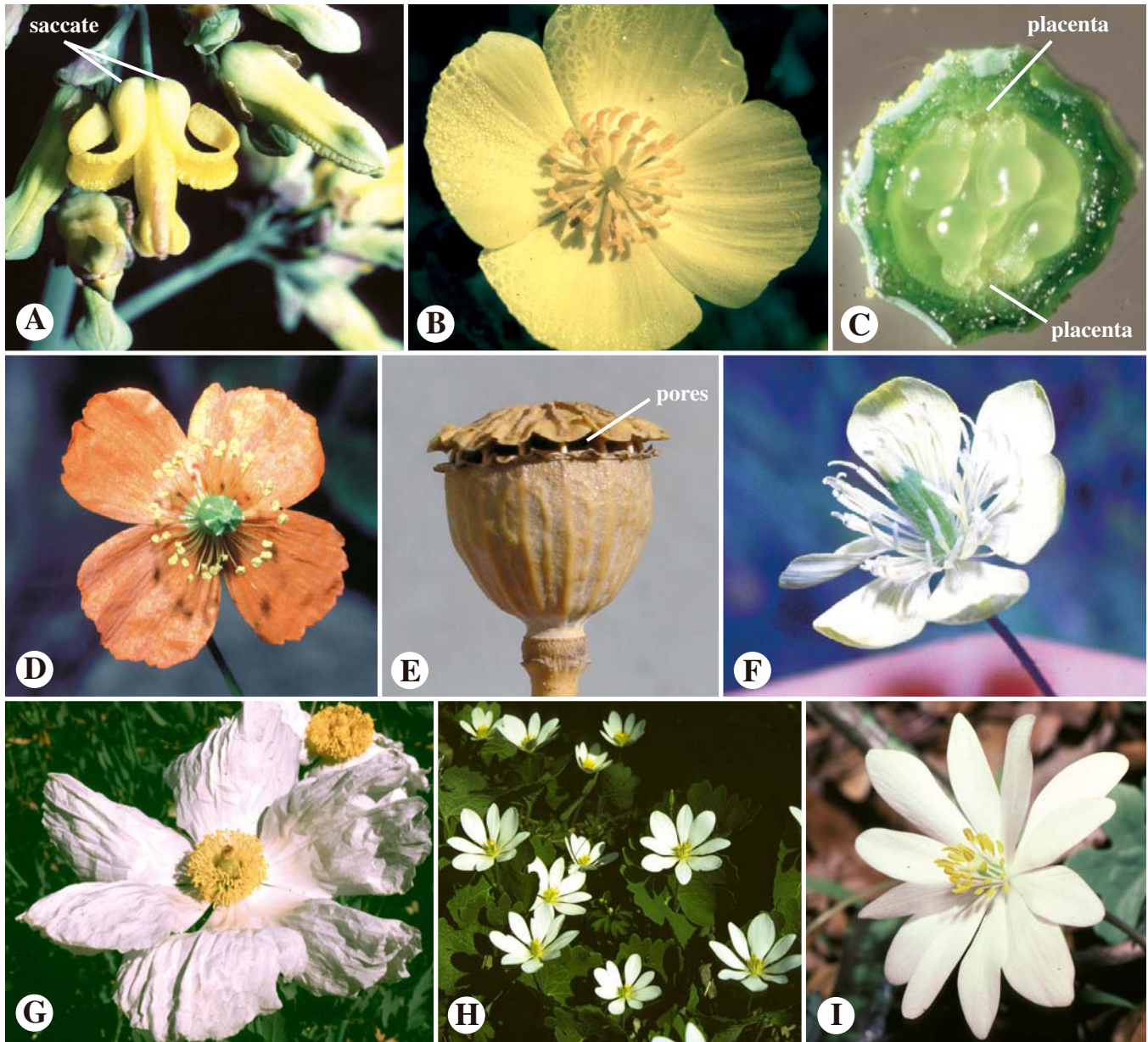


FIGURE 8.4 RANUNCULALES. Papaveraceae. **A.** *Dicentra chrysantha* (Fumarioideae), with biradial, saccate flowers. **B.** *Dendromecon rigida*, tree poppy, showing biseriata (2+2) corolla (sepals caducous). **C.** *Eschscholzia californica*, California poppy, ovary cross-section showing parietal placentation. **D.** *Papaver californicum*, ree poppy. **E.** *Papaver somniferum*, mature poricidal capsule. **F.** *Platystemon californicus*, cream cups. **G.** *Romneya trichocalyx*, Matilija poppy. **H,I.** *Sanguinaria canadensis*, bloodroot, having several petals per flower.

imbricate and crumpled in bud, the outer petals with a spur or sac in some taxa. The **stamens** are usually numerous [4–6], centripetal, sometimes in two or three bundles. **Anthers** are tetrasporangiate or bisporangiate, dithecal or monotheical (sometimes in the same flower). The **gynoecium** is syncarpous, with a superior ovary, 2 [to several] carpels, and 1 [2 several] locule(s). The **style** and **stigma** are usually solitary, the latter sometimes connate to form a disclike

structure. **Placentation** is parietal [rarely axile]; **ovules** are anatropous to campylotropous, bitegmic, 1∞ per carpel. **Nectaries** are sometimes present at base of stamens. The **fruit** is a longitudinally dehiscent or poricidal capsule, sometimes a schizocarp or nut. The **seeds** are oily endospermous, arillate in some taxa.

The Papaveraceae are treated here as encompassing the Fumariaceae (often as subfamily Fumarioideae), the latter

differing in having biradial or bilateral flowers, the outer whorl of petals usually with a spur or sac. Members of the family are distributed in mostly north temperate regions. Economic importance includes many cultivated ornamentals and taxa used as oil seeds. *Papaver somniferum*, opium poppy, is an addictive narcotic plant, the source of heroin (which has shaped human history) and very important medicinally, e.g., as the source of the analgesic morphine and other alkaloids. See Hoot et al. (1997) and Kadereit et al. (1994, 1995) for detailed phylogenetic studies of the family.

The Papaveraceae are distinctive in being herbs, shrubs, or small trees (some with milky sap), with a **dichlamydeous, triseriate** perianth (the corolla **biseriate**), usually **numerous** stamens, and a superior, compound ovary usually with **parietal** placentation, the fruit usually a **loculicidal or poricidal capsule**. **K** 2 [3] or (2 [3]) **C** 2+2 or 3+3 [-16] **A** ∞ [4-6] **G** (2) [-several], superior.

Ranunculaceae Buttercup family (meaning little frog, after the amphibious habit of many species). 62 genera / 2,450 species. (Figure 8.5)

The Ranunculaceae consist of terrestrial or aquatic, perennial or annual shrubs, herbs, or lianas. The **leaves** are spiral, simple to compound, stipulate or exstipulate. The **inflorescence** is a cyme or a solitary flower. The **flowers** are bisexual, rarely unisexual, actinomorphic or zygomorphic, hypogynous; the receptacle is somewhat elongate. The **perianth** is dichlamydeous, a hypanthium absent. The **calyx** is aposepalous with 5-8 [3], often petaloid sepals. The **corolla** is apopetalous with few ∞ [rarely 0] petals. The **stamens** are apostemonous, usually ∞ , spiral and usually centripetal, sometimes multiwhorled. **Anthers** are longitudinal in dehiscence, tetrasporangiate and dithecal. The **gynoecium** is apocarpous [rarely syncarpous], usually of ∞ [1 few] pistils/carpels, each unilocular, ovaries superior. **Placentation** is marginal, apical, or basal, axile in the syncarpous taxa; **ovules** are anatropous or hemitropous, bitegmic or unitegmic, several ∞ per carpel. **Nectaries** are often perigonal (at the base of staminode-like petals). The **fruit** is an aggregate of follicles, achenes, or berries. Flowers are insect or wind-pollinated.

Members of the Ranunculaceae have distributions mainly in temperate and boreal regions. Economic importance includes cultivated ornamentals, medicinal plants, poisonous plants, and weeds. See Hoot (1991, 1995) for phylogenetic analyses of the family.

The Ranunculaceae are distinctive in being herbs, shrubs, or lianas having flowers with **spirally arranged** perianth parts, usually in two series, **numerous** stamens, and an **apocarpous** gynoecium.

K 5-8 [3] **C** few- ∞ [0] **A** ∞ **G** ∞ [1-few], superior.

PROTEALES

The Proteales contain three very different families, classified together only rather recently. All three are similar in having mostly 1 ovule per ovary. The Nelumbonaceae and Platanaceae are similar in having apical placentation. The Platanaceae and Proteaceae have similarities in wood anatomy.

Nelumbonaceae Water Lotus family (*Nelumbo*, a Sinhalese name). 1 genus (*Nelumbo*) / 2 species. (Figure 8.6)

The Nelumbonaceae consist of aquatic, perennial herbs, with milky latex present from articulated laticifers. The **stems** are rhizomatous. The **leaves** are spiral, peltate, petiolate (petiole emergent), simple, undivided, orbicular, concave above, and net-veined. The **inflorescence** is an axillary (from a scale leaf of the rhizome), emergent, solitary flower. The **flowers** are large, bisexual, actinomorphic, long-pedunculate, ebracteate, hypogynous; the receptacle is enlarged and spongy, with numerous sunken cavities containing individual pistils. The **perianth** is approximately 3-seriate, with an outermost whorl of 2 green, sepaloid tepals, and two inner whorls of numerous, yellow or red, petaloid tepals, all tepals distinct. The **stamens** are numerous, spiral, apostemonous; filaments are narrow. **Anthers** are latrorse to introrse and longitudinal in dehiscence, tetrasporangiate, with a laminar connective. The **gynoecium** is apocarpous, with 12-40 one-loculed superior ovaries. The **style** and **stigma** are solitary and terminal. **Placentation** is apical; **ovules** are anatropous and bitegmic, 1 per ovary. The **fruit** is an aggregate of nuts, each sunken in an accrescent receptacle. The **seeds** are exalbuminous. Flowers are beetle-pollinated. The stems have a eustele or atactostele-like vasculature.

The Nelumbonaceae were formerly treated as a subfamily (Nelumboideae) of the Nymphaeaceae. The two species that make up the Nelumbonaceae are distributed from eastern North America to northern South America (*N. lutea*) and Asia to northern Australia (*N. nucifera*). Economic importance includes use as cultivated ornamentals, edible rhizomes and seeds, medicine, and in religious rites (the sacred lotus); *N. nucifera* is famous for having long-lived seeds, some discovered 3000 years old. See Williamson and Schneider (1993) for more detailed information.

The Nelumbonaceae are distinctive in being **aquatic herbs** with often atactostelic stems, **emergent concave-peltate** leaves, and **emergent, solitary** flowers with numerous tepals, numerous stamens, and an **apocarpous** gynoecium having pistils **partially embedded within an expanded receptacle**; the fruit is an **aggregate of nuts** within an **accrescent receptacle**.

P 2+ ∞ + ∞ **A** ∞ **G** ∞ , superior.

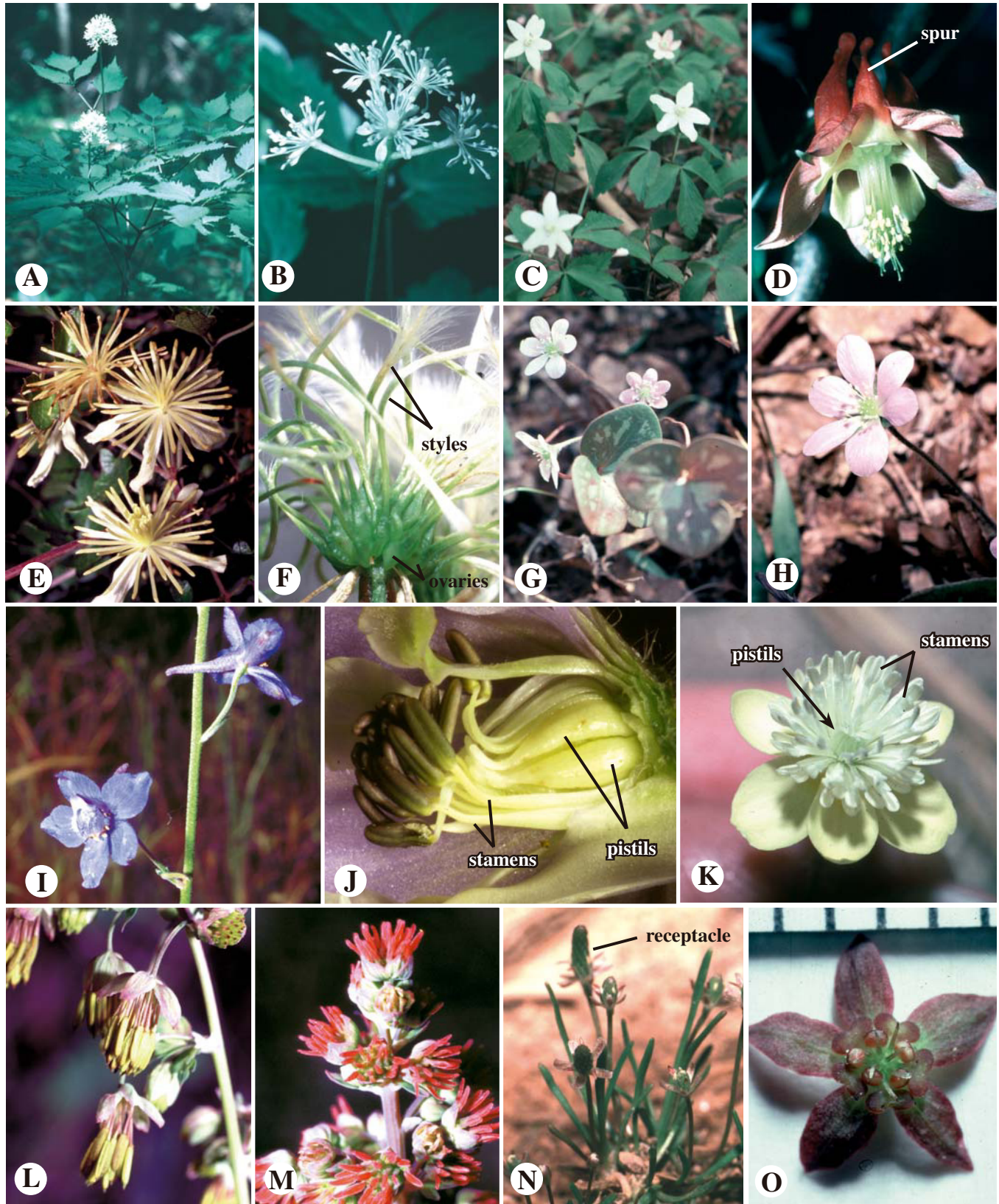


FIGURE 8.5 RANUNCULALES. Ranunculaceae. **A,B.** *Actaea alba*, baneberry. **C.** *Anemone lancifolia*, wind-owen. **D.** *Aquilegia canadensis*, columbine, perianth with prominent spurs. **E,F.** *Clematis pauciflora*, virgin's bower. **G,H.** *Hepatica americana*, liverleaf. **I.** *Delphinium parryi*, larkspur. **J.** *Delphinium* sp., larkspur, perianth opened to show numerous stamens and pistils. **K.** *Ranunculus californica*, buttercup. **L,M.** *Thalictrum fendleri*, meadow-rue. **L.** Male flowers, pendant. **M.** Female flowers, erect, each with several pistils. **N.** *Myosurus minimus*, mouse-tail, with elongate receptacle. **O.** *Xanthorhiza simplicissima*, yellow-root.

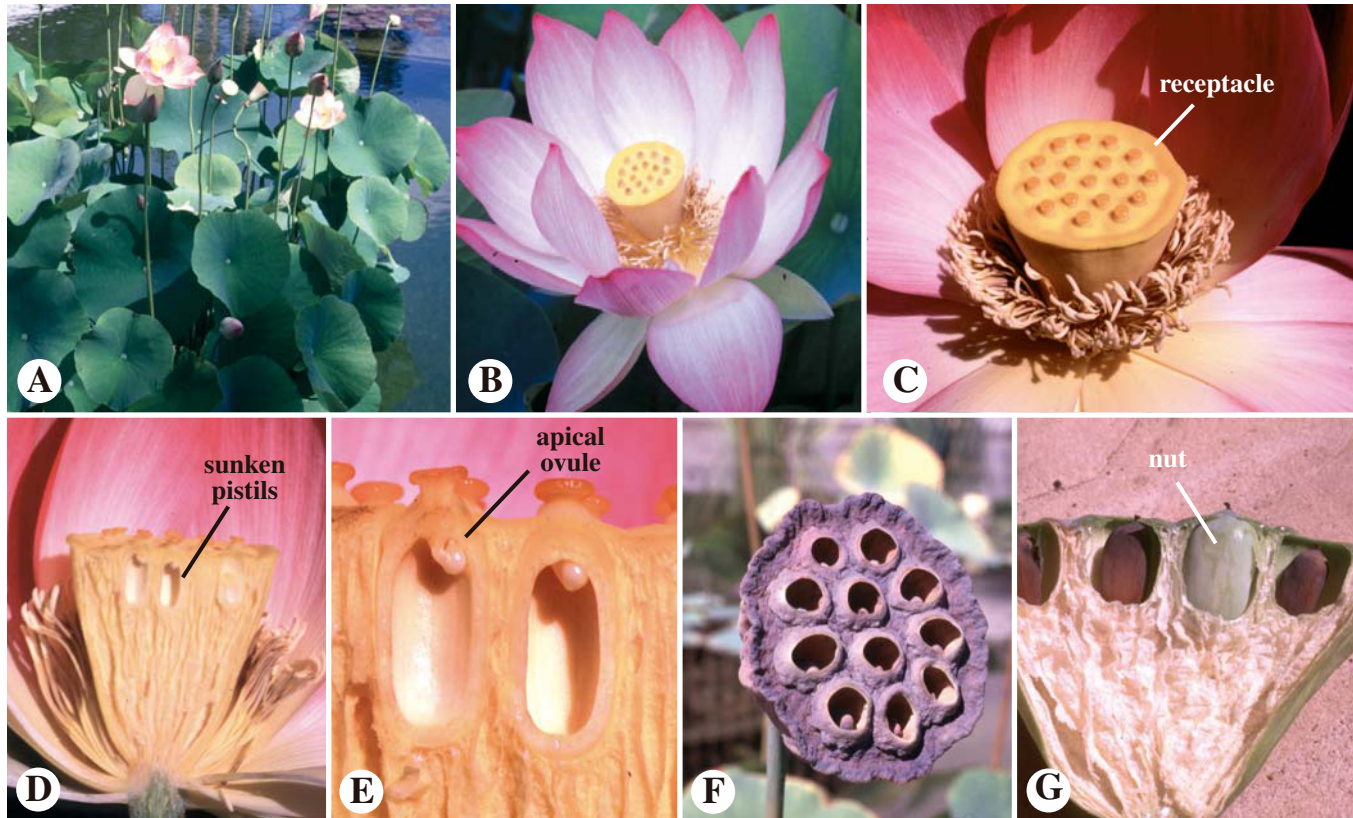


FIGURE 8.6 PROTEALES. Nelumbonaceae. *Nelumbo nucifera*, Indian/Chinese lotus. **A.** Whole plants, showing emergent, peltate/concave leaves and flowers. **B.** Whole flower, showing numerous tepals. **C.** Flower close-up; note numerous stamens and enlarged receptacle. **D.** Receptacle longitudinal-section, with sunken pistils at apex. **E.** Close-up of pistils in longitudinal section, showing apical placentation. **F.** Fruit, an aggregate of nuts, sunken in accrescent receptacle. **G.** Close-up of fruit, longitudinal section, showing nuts.

Platanaceae Plane Tree or Sycamore family (Greek *Platanus*, broad leaf). 1 genus (*Platanus*) / 7 species. (Figure 8.7)

The Platanaceae consist of monoecious trees with exfoliating bark. The **leaves** are alternate (the petiole enclosing an infrapetiolar bud), simple, palmately lobed, stipulate (stipules usually encircling twig), deciduous, with stellate trichomes, and usually palmately netted venation. The **inflorescence** is a terminal, pendant spike of unisexual heads. The **flowers** are small, unisexual. The **perianth** is biseriate, hypanthium absent. The **calyx** is aposepalous to basally synsepalous, of 3-4 [7] sepals. The **corolla** is apopetalous, of 3-4 [7] (usually 0 in females) petals. The **stamens** are 3-4 [7], antisepalous, apostemonous, staminodes present in some female flowers. **Anthers** are sessile, longitudinal in dehiscence, the connective with an apical, peltate appendage. The **gynoecium** is apocarpous, with a superior ovary and 5-8 [3-9] carpels and locules. **Placentation** is apical; **ovules** are orthotropous, bitegmic, 1 [2] per ovary. **Nectaries** are absent.

The **fruit** is a multiple of achenes, with an accrescent, bristly perianth (functioning in wind dispersal) and persistent style. The **seeds** are endospermous. Flowers are wind-pollinated.

Members of the Platanaceae are distributed in the northern hemisphere. Economic importance includes timber and use as cultivated ornamentals. See Kubitzki et al. (1993) for a treatment of the family.

The Platanaceae are distinctive in being in being *monoecious trees* with *encircling stipules*, *infrapetiolar buds*, usually *palmately lobed and veined leaves*, and a *pendant spike of heads* bearing *unisexual* flowers, with a *multiple fruit of bristly achenes*.

Male: K 3-4 [7] or (3-4 [7]) C 3-4 [7] A 3-4 [7].

Female: K 3-4 [7] or (3-4 [7]) C 0 G 5-8 [3-9], superior.

Proteaceae Protea family (after Proteus, the sea god, for his versatility in changing form). 75 genera / 1350 species. (Figure 8.8)

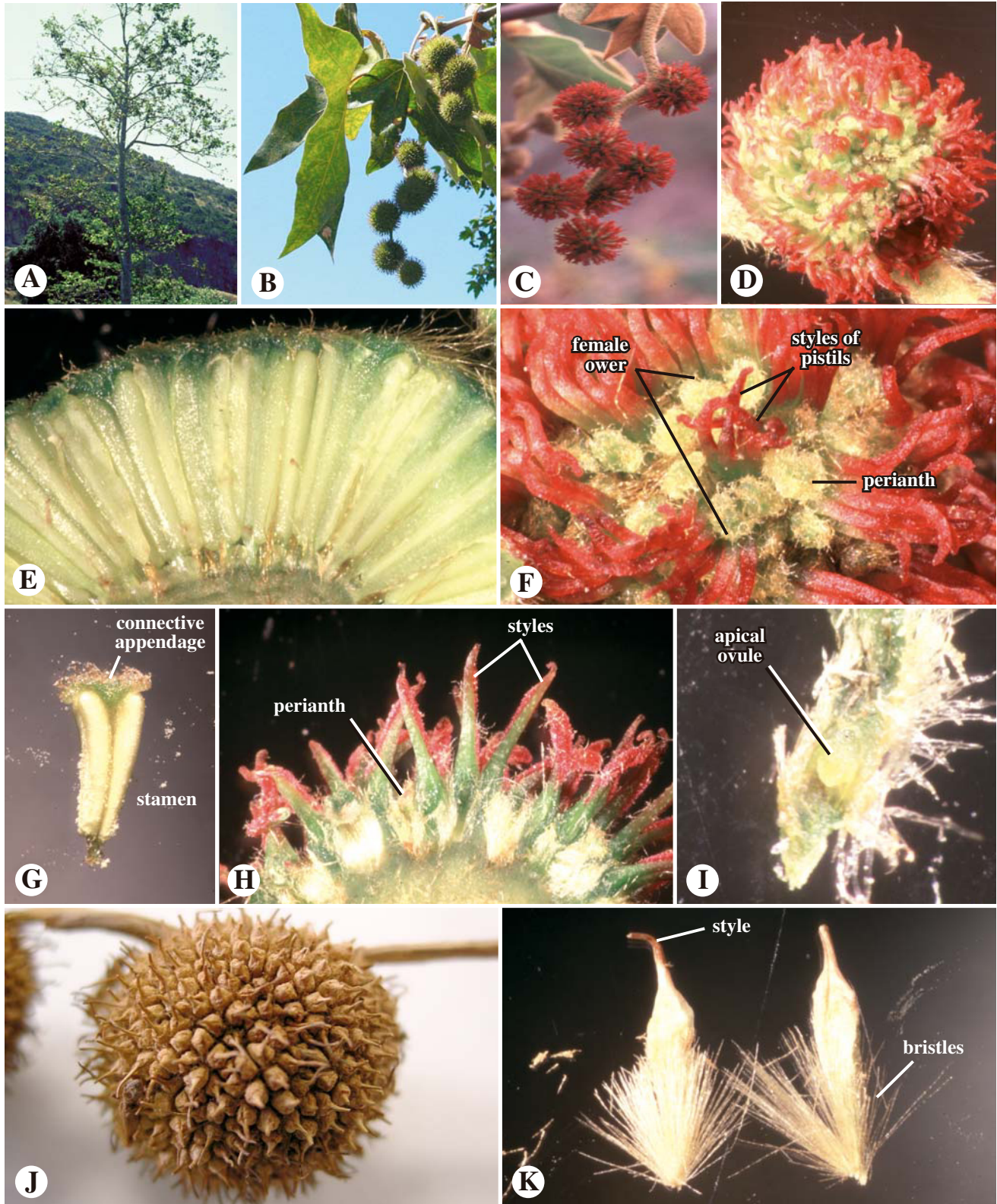


FIGURE 8.7 PROTEALES. Platanaceae, *Platanus racemosa*. **A.** Tree, early in spring. **B.** In orencence (maturing to fruit), a spike of heads. **C.** Close-up of in-orencence. **D.** Close-up of head. **E.** Male head, longitudinal section. **F.** Close-up of female head, showing pistils and perianth of ill-de ned owers. **G.** Stamen of male ower. Note apical extension of connective. **H.** Female head, longitudinal section. Note distinct pistils (apocarpous gynoecium) and bristlelike perianth. **I.** Ovary, longitudinal section, showing single, apical ovule. **J.** Fruit, a multiple of achenes. Note persistent styles. **K.** Individual achenes, with accrescent, bristly perianth.

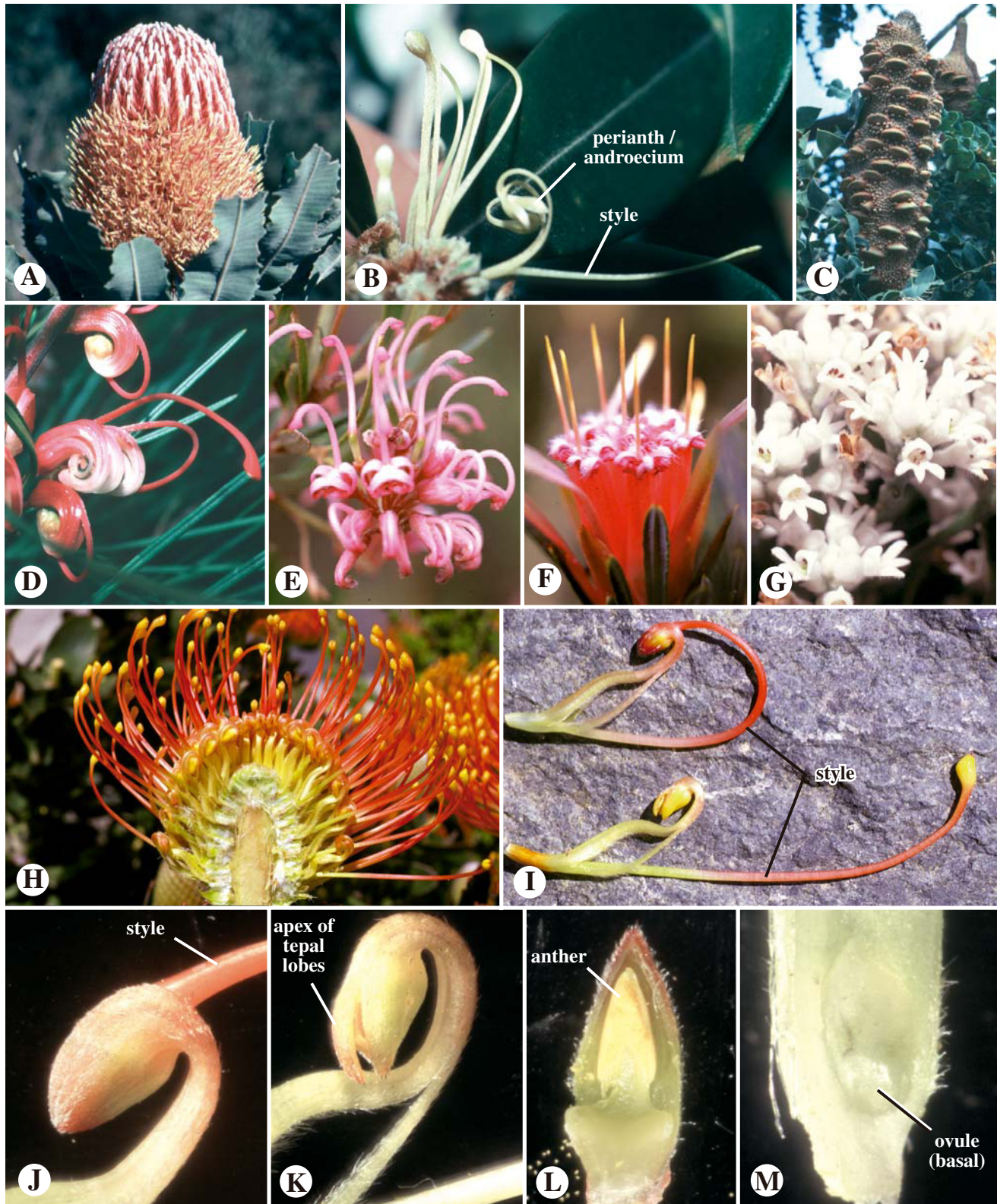


FIGURE 8.8 PROTEALES. Proteaceae. A–C. *Banksia* spp. A. *Banksia menziesii*, headlike in orescence. B. *Banksia* sp., showing partially opened (above) and open (below) owers. C. *Banksia grandis*, multiple fruit of follicles. D. *Grevillea longistyla*, owers. E. *Grevillea sericea*, in orescence. F. *Lambertia formosa*, in orescence. G. *Conospermum taxifolium*, owers. H–M. *Leucospermum* sp., pincushions. H. Headlike in orescence in longitudinal section. I. Flowers removed, before (above) and after (below) anthesis. J. Close-up of ower tip before anthesis, showing four coherent tepal lobe apices enclosing style tip. K. Flower tip, after release of style and stigma. L. Close-up of tepal lobe apex, abaxial surface, showing sessile anther. M. Ovary longitudinal section, showing single, basal ovule.

The Proteaceae consist of shrubs and trees. The **roots** are without mycorrhizae, often with short, lateral proteoid roots. The **leaves** are usually spiral and simple, pinnate or bipinnate, evergreen, and coriaceous. The **inflorescence** is a bracteate raceme, umbel, involucrate head, or of solitary or paired flowers. The **flowers** are bisexual or unisexual, actinomorphic or zygomorphic. The **perianth** is uniseriate. The **calyx** is valvate, consisting of 4 distinct or connate sepals. The **corolla** is absent (or interpreted as modified into 4-lobed nectariferous disk or minute scales). The **stamens** are 4, anti-sepalous, usually episepalous. **Anthers** are longitudinal in dehiscence, tetrasporangiate or bisporangiate, dithecal or monothecal, with thecae and connective often extended as an appendage. The **gynoecium** is unicarpellous, with a superior ovary and 1 locule. **Placentation** is marginal or appearing basal; **ovules** are variable in type, bitegmic, 1 2 [∞] per ovary. The **fruit** is a follicle, nut, achene, or drupe. **Seeds** are exalbuminous.

Members of the Proteaceae occur in rain forest to xeric habitats with distributions in tropical and subtropical regions, especially Australia and South Africa. Economic importance includes numerous cultivated ornamentals (e.g., *Banksia*, *Grevillea*, *Protea*), important timber trees, and species with edible seeds (e.g., *Macadamia*). See Hoot and Douglas (1998) for a recent phylogenetic study of the Proteaceae.

The Proteaceae are distinctive in having flowers with a **uniseriate perianth** of **4 sepals**, **4 antisepalous stamens**, and a **unicarpellous**, superior ovary.

K 4 or (4) **C** 0 **A** 4 **G** 1, superior.

CORE EUDICOTS

The so-called Core Eudicots of the APG II system include (among others) the orders Caryophyllales, Santalales, and Saxifragales, plus two large groups, the Rosids and Asterids (Figure 8.1, Table 8.1).

CARYOPHYLLALES

The Caryophyllales, sensu APG II (2003), contain 28 families (plus three more, after Stevens, 2001 onwards), of which five are treated here (Table 8.1). Notable among the families not described are the **Droseraceae** (Figure 8.9A E), including the spectacular carnivorous plants *Dionaea muscipula*, Venus fly-trap with trap leaves, and *Drosera* spp., the sundews with tentacular leaves; **Frankeniaceae** (Figure 8.9F H), herbaceous or shrubby halophytes; **Nepenthaceae** (Figure 8.9I), carnivorous plants with pitchers at the ends of photosynthetic leaves; **Nyctaginaceae** (Figure 8.9J,K), four-o'clock family,

including *Bougainvillea*; **Phytolaccaceae** (Figure 8.9L), the pokeberry family; **Plumbaginaceae** (Figure 8.9M), including the cultivars *Armeria*, *Limonium* (statice) and *Plumbago*; **Portulacaceae** (Figure 8.9N,O), the purslane family, mostly succulents, including a number of cultivated ornamentals; **Simmondsiaceae** (Figure 8.9P,Q), a monotypic family consisting of *Simmondsia chinensis*, jojoba, the seed a source of oil-like wax with many uses, including cosmetics and skin/hair products; and **Tamaricaceae** (Figure 8.9R), including the cultivar and weed *Tamarix*.

The Caryophyllales encompass a traditional group formerly known as the Centrospermae (or Caryophyllidae, after Cronquist 1981). The traditional Centrospermae or Caryophyllidae are largely equivalent to the Core Caryophyllales, a complex of approximately 15 families (Figure 8.10), although the Polygonaceae, Plumbaginaceae, Frankeniaceae, and Tamaricaceae were often included. Knowledge of the interrelationships within the Core Caryophyllales is still in flux, with several additional families having been proposed by various workers.

Many, though not all, members of the Core Caryophyllales possess pollen that is trinucleate upon being released from the anther, a relatively rare feature in angiosperms (most being binucleate at release). Also, many members of the Core Caryophyllales have either free-central (hence the name Centrospermae, from the seeds arising from a central column) or basal placentation. More clear-cut apomorphies for the Core Caryophyllales are an ultrastructural feature **sieve tube plastids with protein crystalloid inclusions surrounded by proteinaceous filaments** (Figure 8.11B) and ovule/seed apomorphies **ovules campylotropous** (Figure 8.11A) and **perispermous seeds** (see Chapter 11).

A subgroup of approximately 10 families of the Core Caryophyllales, termed the Higher Caryophyllales, are largely united by the presence of **betalains** (Figure 8.11C), reddish, purplish, or yellowish pigmented compounds that functionally replace the anthocyanins found in other angiosperms.

The Caryophyllales of the APG II system also include a number of basal families previously unsuspected of close relationships to the Core Caryophyllales prior to detailed molecular studies. These families include the Simmondsiaceae and the carnivorous groups Droseraceae, Drosophyllaceae, and Nepenthaceae. The relationships of these families portrayed in Figure 8.9 shows one possibility of evolution in this group, necessitating the reversal of carnivory in two families. See Meimberg et al. (2000) and Soltis et al. (2000) for more information.

Aizoaceae Mesembryanthemum or Vygie family (meaning always alive). 128 genera / 1850 species. (Figure 8.12)

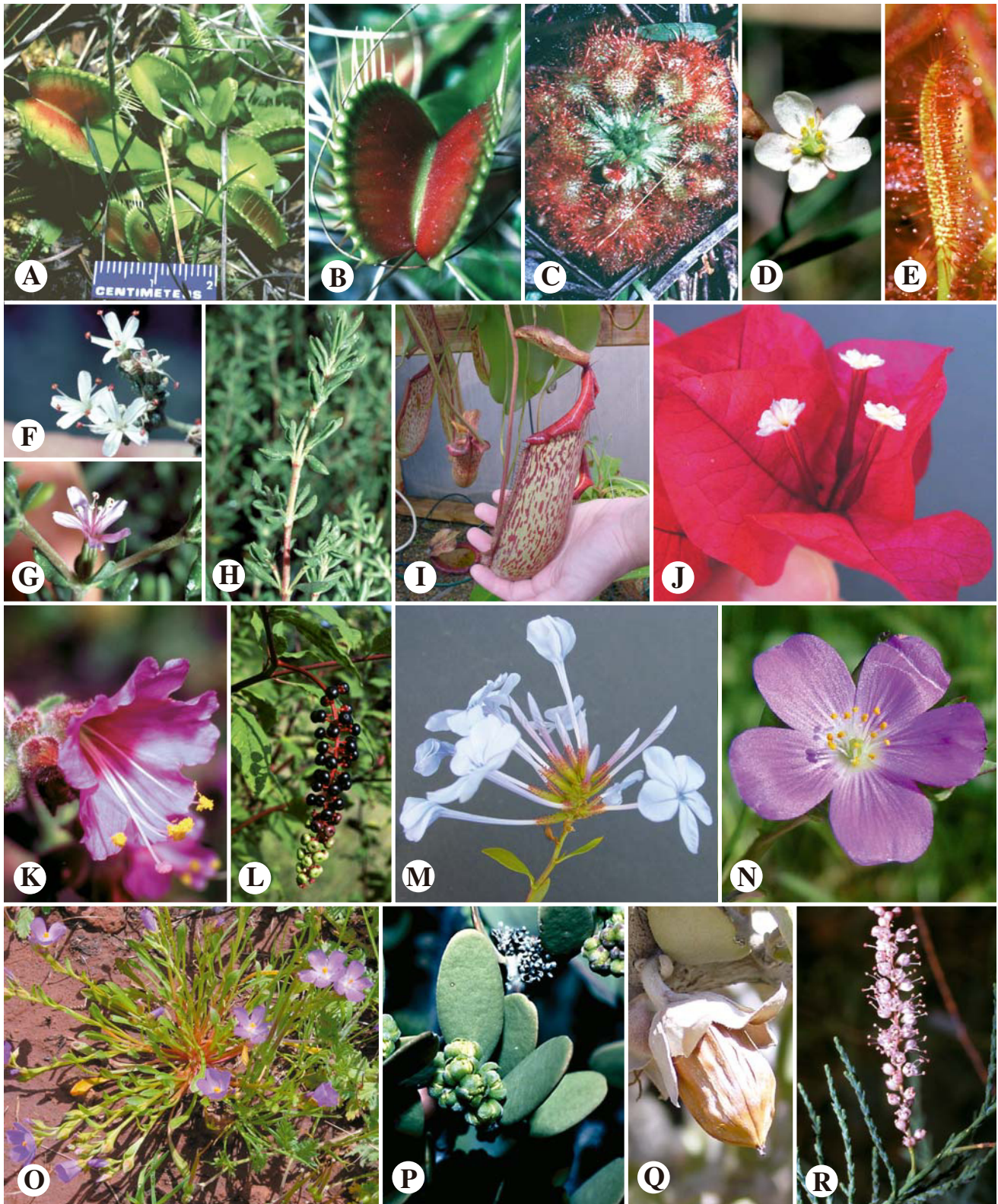


FIGURE 8.9 CARYOPHYLLALES. A–E. Droseraceae. A,B. *Dionaea muscipula*, Venus fly trap, with trap leaves. C–E. *Drosera* spp., sundews, with tentacular leaves. C,D. *Drosera capillaris*. E. *Drosera capensis*. F–H. Frankeniaceae. F. *Frankenia palmeri*. G,H. *Frankenia salina*. I. Nepenthaceae, *Nepenthes rafflesiana*, with pitchers from end of leaves. J,K. Nyctaginaceae. J. *Bougainvillea spectabilis*. K. *Mirabilis laevis*. L. Phytolaccaceae, *Phytolacca americana*, pokeberry. M. Plumbaginaceae, *Plumbago auriculata*. N,O. Portulacaceae, *Calandrinia* sp. P,Q. Simmondsiaceae, *Simmondsia chinensis*. jojoba. P. Flowers of male individual. Q. Fruit of female individual. R. Tamaricaceae, *Tamarix ramosissima*, tamarisk.

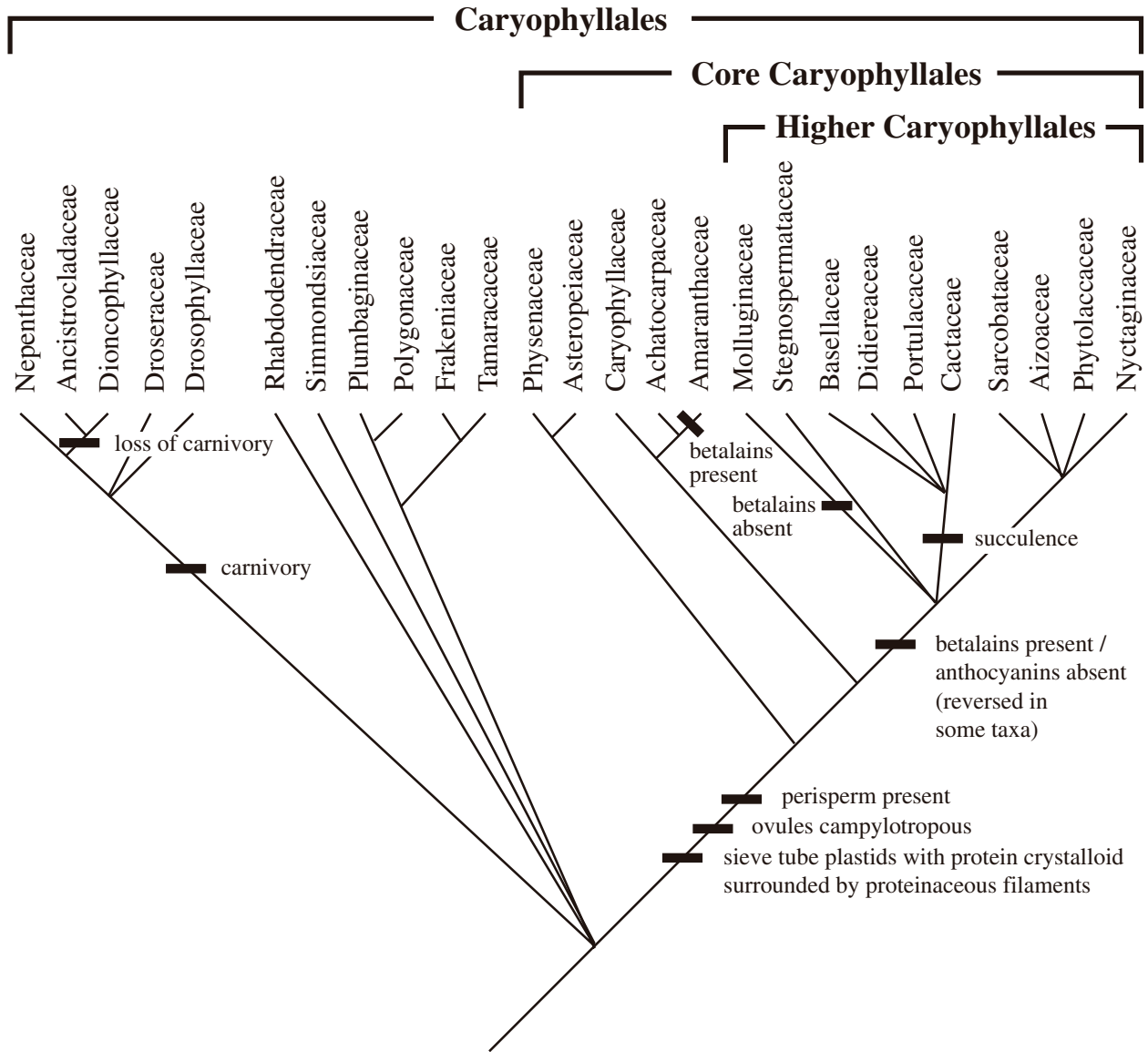


FIGURE 8.10 Cladogram of the Caryophyllales, with selected apomorphies. Simplified and modified from Meimberg et al. (2000) and Soltis et al. (2000). Two families (Barbueviaceae and Gisekiaceae) recognized by APG II (2003) and three families (Halophytaceae, Hectorellaceae, and Limeaceae) recognized by Stevens (2001 onwards) were not considered in these studies.

The Aizoaceae consist of annual or perennial herbs, rarely shrubs or trees, rarely spiny. The **leaves** are often centric, without a bifacial structure (some with apical lens through which light enters), opposite or whorled (rarely alternate), simple, undivided, usually exstipulate, succulent, often terete or angled. The **inflorescence** is terminal or axillary, of solitary flowers or cymes. The **flowers** are bisexual (rarely unisexual), actinomorphic, epiperigynous or perigynous. The **perianth** is uniseriate, a hypanthium present. The **calyx** is distinct with 5 [3–8] sepals. The **corolla** is absent, the petaloid structures of some taxa interpreted as petaloid staminodes. The **stamens**

are [4] 5 ∞, apostemonous or basally connate into bundles or monadelphous, with an outer whorl(s) of petaloid staminodes in some taxa. The **gynoecium** is syncarpous, with a superior or inferior ovary, 2 ∞ carpels, and 2 ∞ (rarely 1) locules. The **styles** are generally as many as carpels. **Placentation** is axile, parietal with septa, or basal; **ovules** are campylotropous to anatropous, bitegmic, [1] ∞. **Nectaries** are present, inner to the insertion of the androecium. The **fruit** is a loculicidal capsule or berry. Betalain pigments are present, anthocyanins absent. Photosynthesis is often C4 or CAM (see Chapter 10).

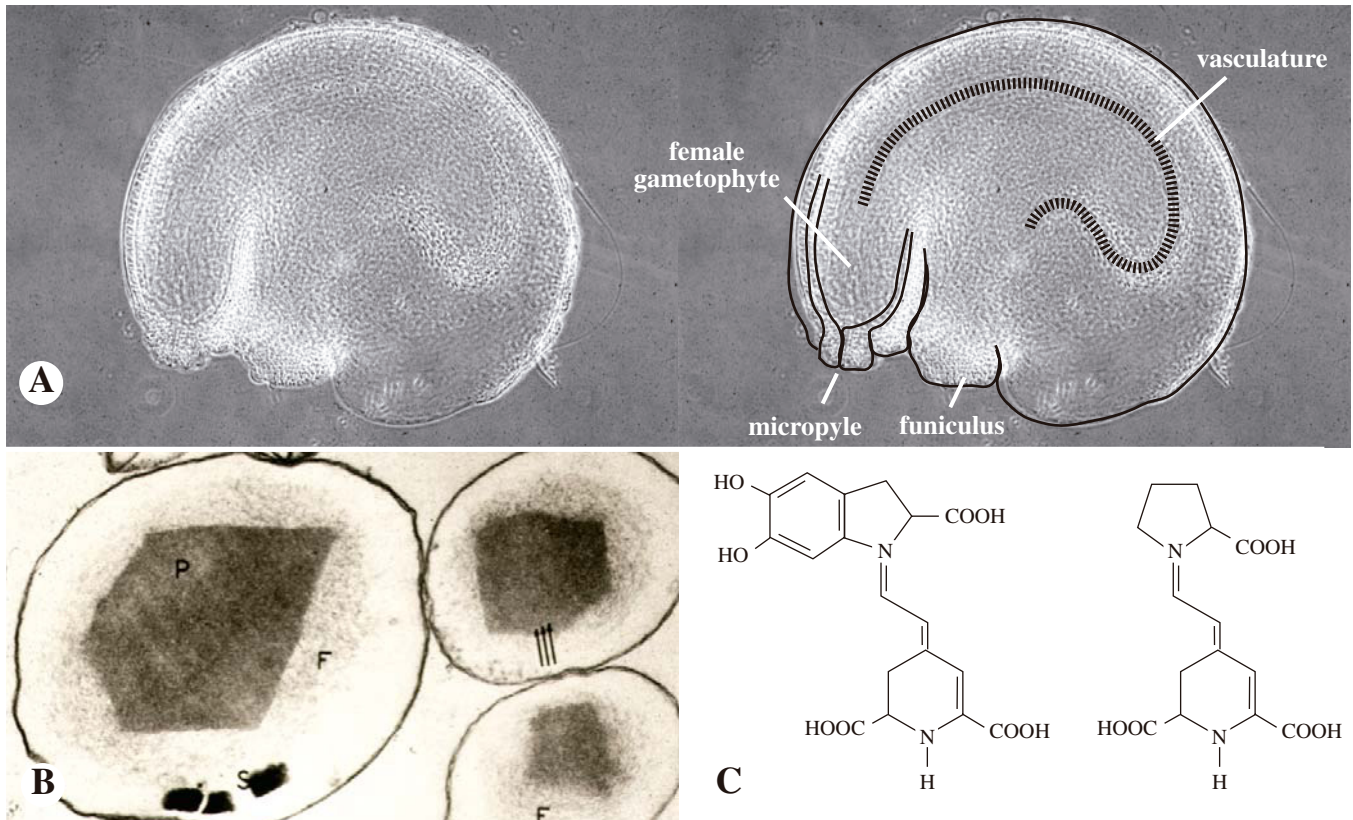


FIGURE 8.11 Caryophyllales apomorphies. **A.** Campylotropous ovule. **B.** Sieve tube plastid with proteinaceous lamellae (F) encircling central crystalloid protein (P); from Benhke (1972), by permission. **C.** Betanidin, a betacyanin (left) and indicoxanthin, a betaxanthin, both examples of betalain pigments.

Members of the Aizoaceae grow in tropical and subtropical regions, primarily in South Africa, less so in Australia. Economic importance includes mostly numerous ornamental cultivars, some (e.g., *Sceletium*) with medicinal properties, *Tetragonia* (New Zealand spinach) used as table greens. See Klak et al. (2003) for a recent treatment of the Aizoaceae.

The Aizoaceae are distinctive in being herbs, rarely shrubs or trees, with generally *opposite, succulent leaves* (often with C4 or CAM photosynthesis) and solitary or cymose flowers with a *uniserial perianth (outer petaloid staminodes present in many)*, usually *numerous stamens*, and usually *numerous ovules*, betalain pigments only present.

P 5 [3-8] **A** [4] 5-∞ **G** (2-∞), superior or inferior, hypanthium present.

Amaranthaceae (including Chenopodiaceae) Amaranth family (Greek unfading, from the persistent bracts and perianth). ca. 174 genera / 2050 species. (Figure 8.13)

The Amaranthaceae consist of annual or perennial, hermaphroditic, dioecious, monoecious, or polygamous herbs,

vines, shrubs, or rarely trees. The **stems** are sometimes jointed or succulent. The **leaves** are simple, spiral or opposite, exstipulate, succulent or reduced in some taxa. The **inflorescence** is of solitary flowers or a spike, panicle, cyme, or thyrse, with bracts and bracteoles bristle-like and pigmented in some taxa. The **flowers** are small, bisexual or unisexual, usually actinomorphic, hypogynous or rarely epihypogynous. The **perianth** is uniseriate (usually termed a calyx, by default), consisting of [0 2] 3 5 [6 8] distinct or rarely basally connate sepals. The **stamens** are [1 2] 3 5 [6 8], generally the same number as sepals and antisepalous, distinct or basally connate and forming a tube. **Anthers** are longitudinal in dehiscence, dithecal or monothecal. The **gynoecium** is unicarpellous or syncarpous, with a superior, rarely half-inferior ovary, 1 3 [5] carpels, and 1 locule. The **style(s)** are 1 several. **Placentation** is basal; **ovules** are campylotropous or amphitropous, bitegmic, 1 [∞]. **Nectaries** are present in some, typically an annular disk. The **fruit** is a nutlet, berry, irregularly dehiscent capsule, or rarely a circumscissile capsule or multiple fruit. The **seeds** are mostly starchy-perispermous with curved embryo.

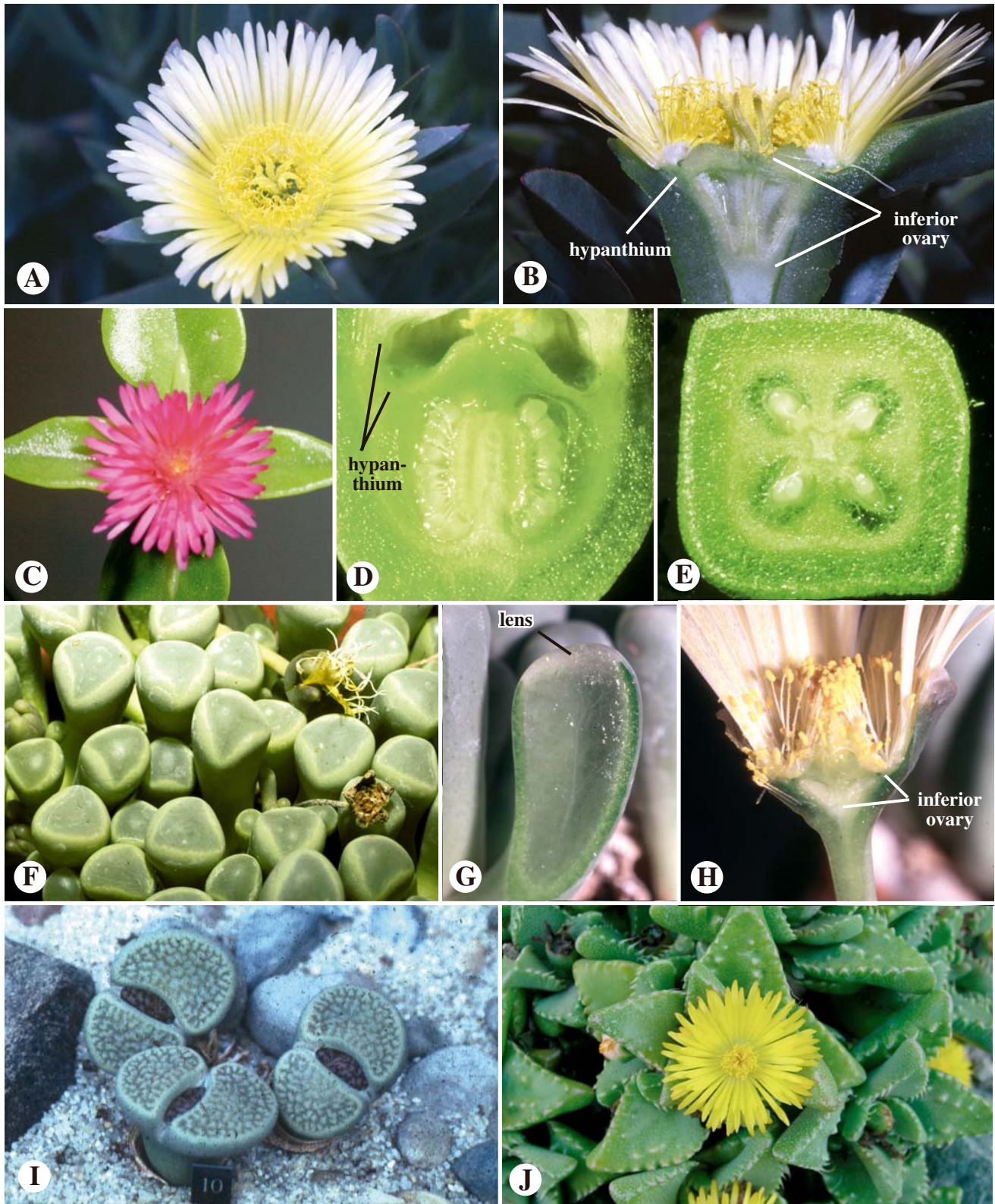


FIGURE 8.12 CARYOPHYLLALES. Aizoaceae. **A,B.** *Carpobrotus edulis*. **A.** Flower, top view, showing numerous, petaloid staminodes and numerous fertile stamens. **B.** Flower, longitudinal section. Note inferior ovary and hypanthium. **C–E.** *Aptenia cordifolia*. **C.** Flowering shoot. Note decussate leaf arrangement. **D.** Flower, longitudinal section, showing inferior ovary and hypanthium. **E.** Flower, cross-section, showing axile placentation (four carpels and locules in this species). **F–H.** *Fenestera aurantiaca*. **F.** Leaves, with apical lens. **G.** Leaf longitudinal section. **H.** Flower, longitudinal section, showing numerous petals and stamens and inferior ovary. **I.** *Lithops* sp., one of the stone plants, the leaves camouflaged as pebbles. **J.** *Faucaria tigrina*, tiger's jaw, flower and shoots.

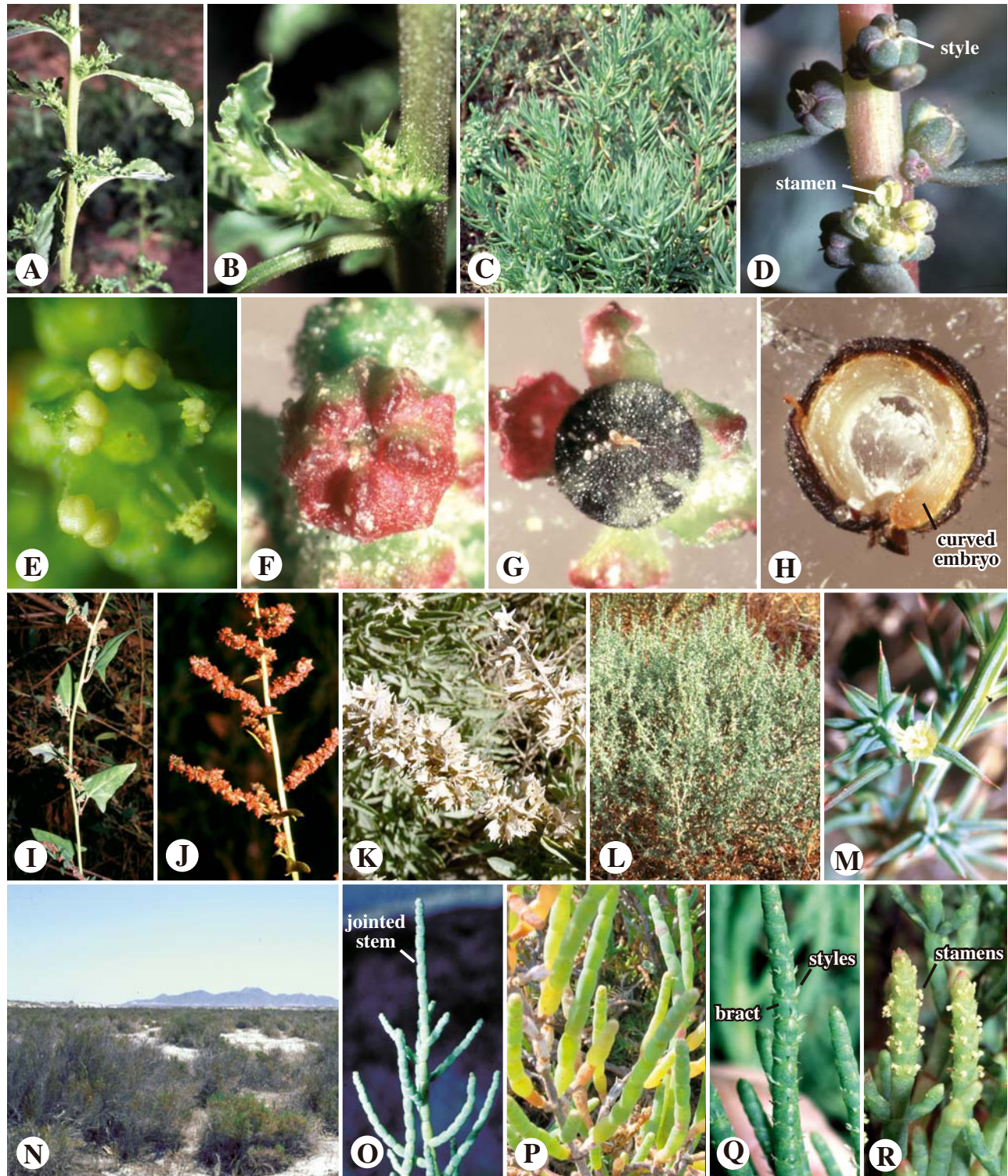


FIGURE 8.13 CARYOPHYLLALES. Amaranthaceae. **A,B.** *Amaranthus* sp. **A.** Whole plant. **B.** Inflorescence, showing bristlelike bracts subtending flower clusters. **C,D.** *Suaeda esteroa*. **C.** Whole plant, a succulent halophyte. **D.** Solitary flowers in axils of leaves; note antiseipalous stamens. **E–H.** *Chenopodium* spp. **E.** Open bisexual flower, with five sepals, five antiseipalous stamens, and single pistil. **F.** Fruit, surrounded by persistent calyx. **G.** Calyx removed to show fruit. **H.** Fruit sectioned, showing single seed with curved embryo. **I,J.** *Atriplex triangularis*. **I.** Whole plant. **J.** Inflorescence, a compound spike of cymes. **K.** *Atriplex canescens*, saltbush/shadscale. Note accrescent bracts surrounding fruits. **L,M.** *Salsola tragus*. **L.** Whole plant, a tumbleweed. **M.** Flower close-up, showing spine-tipped bracts and prominent sepals. **N,O.** *Allenrolfea occidentalis*, iodine bush. **N.** Estuarine community dominated by plant. **O.** Stem close-up, showing jointed, succulent stems with reduced leaves. **P,Q.** *Salicornia virginica*, pickleweed. **P.** Shoot, with succulent, jointed stems and reduced leaves. **Q.** Inflorescence, flowers inside bracts, styles exserted. **R.** *Salicornia subterminalis*, in inflorescence with stamens exserted from bracts.

Betalain pigments are present, anthocyanins absent. Plants have anomalous secondary growth, forming concentric rings of vascular bundles or alternating concentric rings of xylem and phloem, often with C4 or CAM photosynthesis.

The Amaranthaceae has a largely worldwide distribution, members common in some deserts, estuarine or alkaline regions, tropical areas, and some temperate regions. Economic importance includes vegetable crops such as beet (*Beta vulgaris*) and spinach (*Spinacea oleracea*), pseudograin crops such as *Amaranthus* and *Chenopodium* spp. (e.g., *C. quinoa*), some cultivated ornamentals such as *Celosia argentea*, cockscomb, several detrimental weeds, and some local firewood and medicinal plants.

The Amaranthaceae are distinctive in being herbs to trees with **anomalous secondary growth**, simple leaves (succulent to reduced in some), bristlelike, pigmented bracts in some, a **uniseriate perianth** of mostly 3-5 [0-2, 6-8] sepals, basally connate stamens of same number and opposite perianth parts, a **1-loculed, mostly 1-ovuled** ovary with **basal** placentation, seeds with **curved embryo**, and betalain pigments only present. **K** [0-2] **3-5** [6-8] **C** 0 **A** [1-2] **3-5** [6-8] **G** 1(-3) [(5)], superior, rarely half-inferior.

Cactaceae Cactus family (Greek for a spiny plant). 97 genera / 1400 species. (Figures 8.14, 8.15)

The Cactaceae consist of perennial shrubs or trees, with leaf spines arising from specialized axillary meristems termed **areoles**, which in some taxa may also bear small, trichome-like leaves with retorse barbs called **glochidia**. The **stems** are typically succulent, and may be cladodes (e.g., prickly-pears) or radially plicate (e.g., barrel cacti). The **leaves** are simple, often succulent, spiral, usually caducous (persistent and well-developed in Pereskioideae). The **inflorescence** is of axillary, solitary flowers or rarely terminal cymes. The **flowers** are often large, bisexual (rarely unisexual), acinomorphic, epiperigynous. The **perianth** consists of numerous, distinct, spirally arranged tepals, grading from outer bractlike to inner petal-like structures, a hypanthium present. The **stamens** are numerous, spiral or in whorled clusters, and apostemonous. **Anthers** are longitudinal in dehiscence, tetrasporangiate and dithecal. The **gynoecium** is syncarpous, with an inferior ovary, 3 many carpels, and 1 locule. The **style(s)** are single at the base, branched above. **Placentation** is parietal [basal]; **ovules** are campylotropous [anatropous], bitegmic, numerous per carpel. **Nectaries** are present within the hypanthium. The **fruit** is a berry, rarely dry and indehiscent. The **seeds** are arillate in some, exalbuminous, perispermous in some, embryos straight to curved. Flowers are pollinated by bees, moths, hummingbirds, or bats. Betalain pigments are present, anthocyanins absent. Photosynthesis is CAM (Crassulacean

Acid Metabolism), in which stomata are opened at night (when carbon dioxide is fixed and stored), closed during the day to conserve water (see Chapter 10).

The Cactaceae are typically classified into three subfamilies: Pereskioideae (thought to be basal in the family), with persistent, broad vegetative leaves, glochidia absent, and seeds exarillate; Opuntioideae, with cylindrical, caducous leaves, specialized glochidia, and arillate seeds; and Cactoideae, with leaves and glochidia absent and seeds exarillate. Only the Opuntioideae are likely monophyletic. Members of the Cactaceae grow mostly in desert regions with distributions in the New World (except for *Rhipsalis* in Africa). Economic importance includes numerous cultivated ornamentals; *Opuntia* spp. are eaten for their fruits (prickly-pears) and stems (nopales); *Lophophora williamsii* (peyote) is used as a hallucinogen and in religious ceremonies (e.g., Religion of the Native American Church). See Nyffeler (2002) and Wallace and Gibson (2002) for recent studies of Cactaceae systematics.

The Cactaceae are distinctive in being typically **stem-succulent**, CAM shrubs or trees, with **leaves usually reduced or absent**, the axillary meristems modified into specialized **areoles** bearing **leaf spines**, the flowers **epiperigynous** with **spiral perianth parts** intergrading from outer bractlike to inner petal-like parts, having **numerous stamens** and an **inferior ovary** with numerous ovules and **parietal** placentation, betalain pigments only present.

P ∞ **A** ∞ **G** (3-∞), inferior, hypanthium present.

Caryophyllaceae Carnation family (meaning clove-leaved). 87 genera / 2300 species. (Figure 8.16)

The Caryophyllaceae consist of annual or perennial herbs, rarely shrubs, lianas, or trees. The **stems** often have swollen nodes. The **leaves** are opposite (rarely spiral), simple, usually exstipulate. The **inflorescence** is of dichasial cymes or solitary flowers. The **flowers** are bisexual or unisexual, actinomorphic, hypogynous, rarely perigynous. The **perianth** is biseriate, dichlamydeous, hypanthium absent [rarely present]. The **calyx** is synsepalous [rarely aposepalous] with 5 [4] sepals. The **corolla** is apopetalous and often unguiculate (clawed), with 5 [4] petals. The **stamens** are 5-10 [1-4], uniseriate or biseriate, apostemonous, epipetalous, or epise-palous, basally epipetalous and forming a tube in some species. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary (often with a stipe/gynophore), 2-5+ carpels, and 1 locule, often with basal septa. The **style(s)** are terminal, single below, often branched above. **Placentation** is free-central at least above, often axile below; **ovules** are campylotropous to hemitropous, bitegmic, usually numerous per ovary. **Nectaries** occur as a nectariferous disk in some. The **fruit** is a capsule, with valves or teeth 1-2 × the

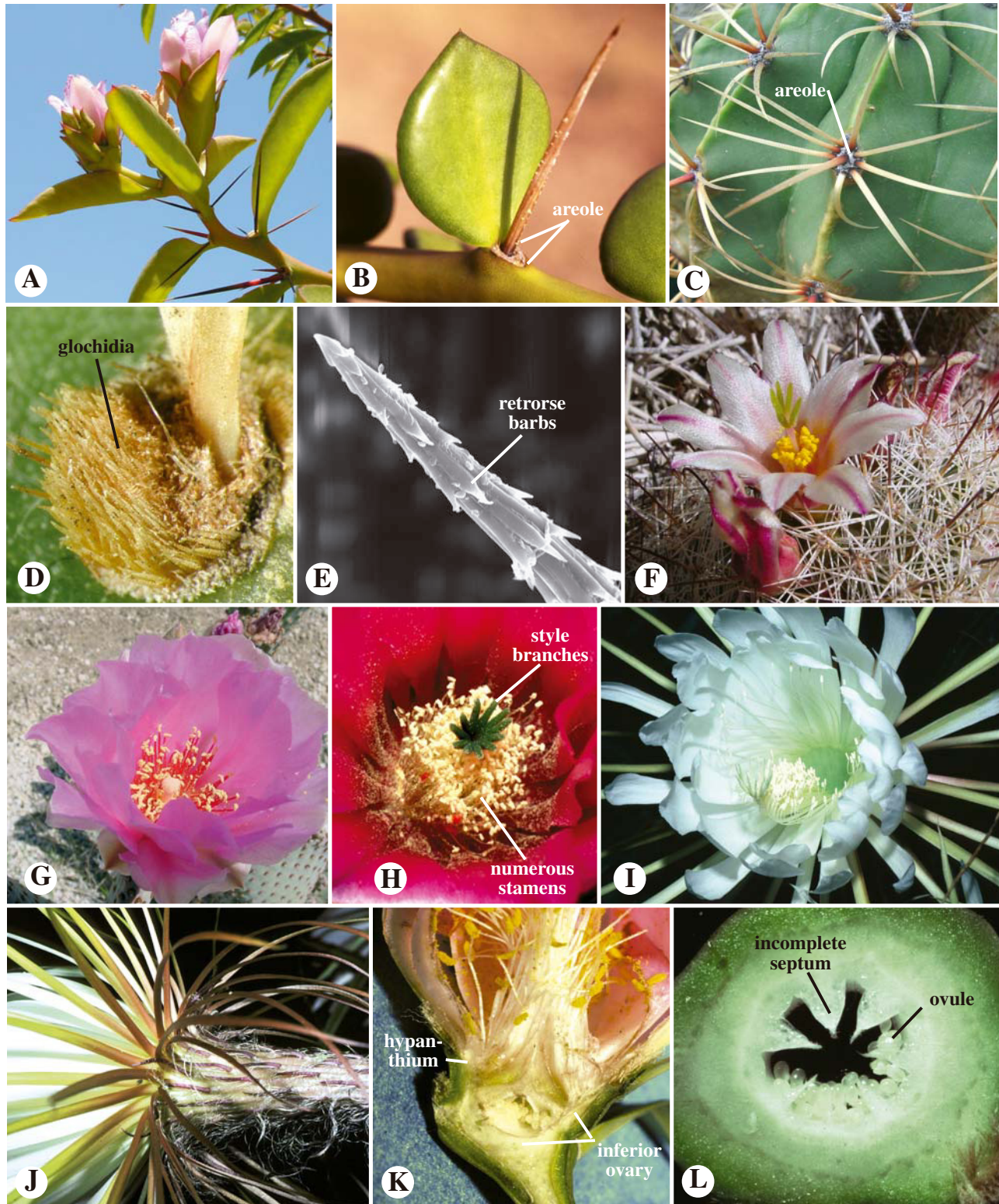


FIGURE 8.14 CARYOPHYLLALES. Cactaceae. **A,B.** Pereskioideae, having persistent, photosynthetic leaves. **A.** *Pereskia grandiflora*. **B.** *Quiabentia zehntneri*. **C.** Cactoideae, typical, leafless stem with areole bearing leaf spines. **D,E.** Opuntioideae, glochidia. **D.** Mass of glochidia. **E.** Single glochidium with retrorse barbs (high magnification, scanning electron microscopy). **F.** *Mammillaria dioica*, with uncinata spines. **G.** *Opuntia basilaris*, flower. **H.** *Echinocereus engelmannii*, close-up of stamens and style branches. **I,J.** *Selenicereus* sp., night-blooming cactus. **I.** Open, zygomorphic flower, with numerous tepals and stamens. **J.** Side view, with transition between bracts and inner petaloid tepals. **K,L.** *Pereskia grandiflora*. **K.** Flower longitudinal section, showing inferior ovary and hypanthium. **L.** Ovary cross-section, showing parietal placentation.

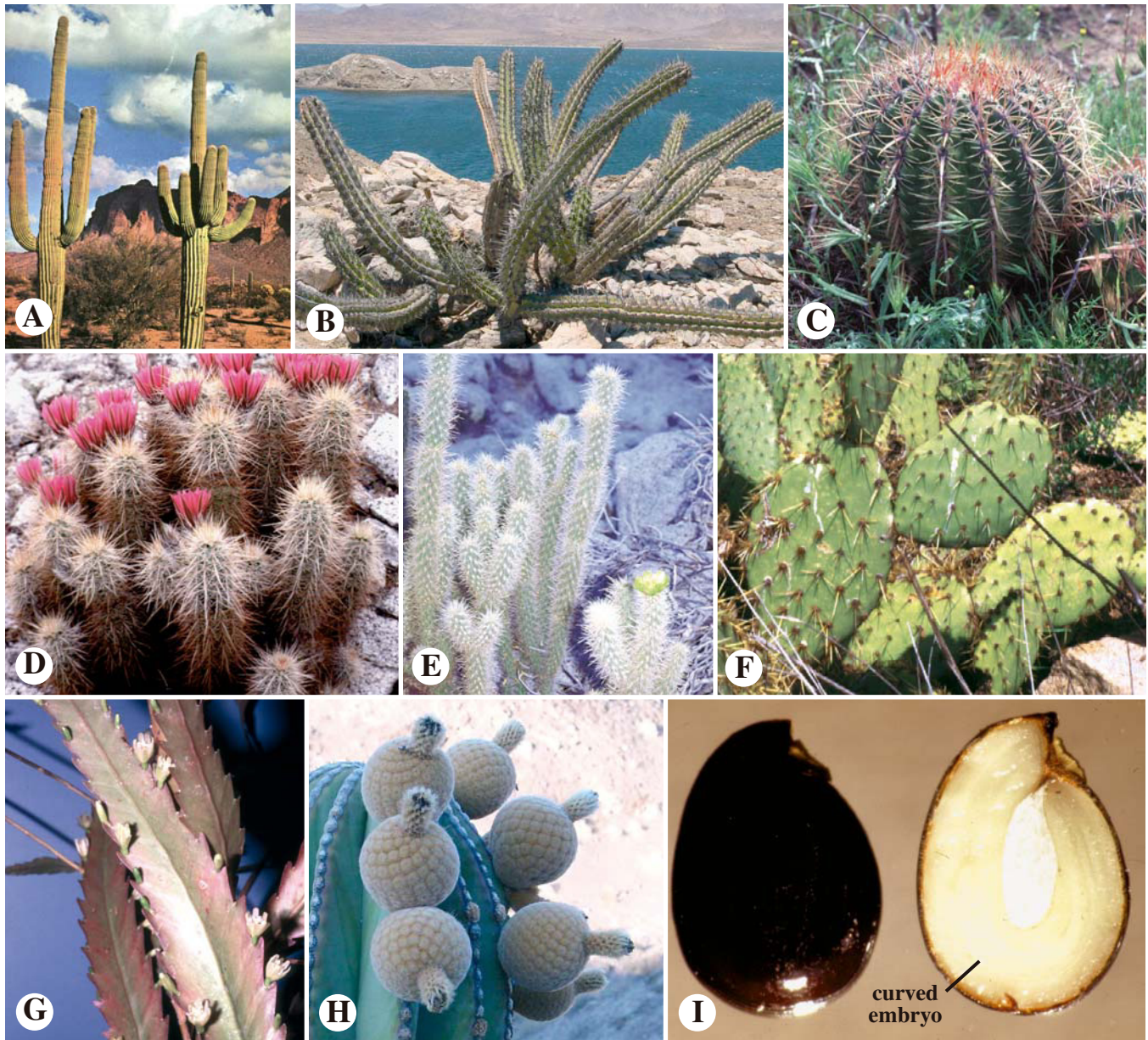


FIGURE 8.15 CARYOPHYLLALES. Cactaceae. A–G. Diversity of growth forms in cacti. A. *Carnegiea gigantea*, saguaro cactus. B. *Stenocereus gummosus*, pitaya agria. C. *Ferocactus viridescens*, a barrel cactus with pleated stems. D. *Echinocereus engelmannii*, hedgehog cactus, with numerous, erect, clumped stems. E. *Cylindropuntia ganderi*, a cholla. F. *Opuntia littoralis*, a prickly pear with flattened cladodes. G. *Zygocactus* sp., a cactus with cladodes bearing marginal flowers. H, I. Propagules. H. Fruits, *Pachycereus pringlei*, cardon. I. *Pereskia grandiflora* seed, longitudinal section.

carpel number. The seeds are often with sculptured seed coat and are perispermous. Anthocyanin pigments are present, betalains absent. Anomalous secondary growth with concentric rings of vascular tissue occurs in some taxa.

The Caryophyllaceae have a worldwide distribution, especially in the northern hemisphere. Economic importance includes several ornamental cultivars, such as *Dianthus*, carnation.

See Smitsen et al. (2002) for a recent phylogenetic analysis of the Caryophyllaceae.

The Caryophyllaceae are distinctive in having *nodes often swollen*, with *simple, opposite leaves*, an inflorescence of *solitary flowers or dichasial cymes*, and biserial, actinomorphic, usually pentamerous flowers with *distinct, clawed petals*, a superior ovary with distally *free-central*



FIGURE 8.16 CARYOPHYLLALES. Caryophyllaceae. **A,B.** *Arenaria caroliniana*. **A.** Whole plants, an annual herb. **B.** Flower close-up. **C.** *Cerastium glomeratum*, capsules, with apical lobes (teeth). **D–H.** *Lychnis coronaria*. **D.** Flower, face view. **E.** Calyx, side view, showing synsepalous fusion. **F.** Single petal, showing basal claw. **G.** Ovary, with five styles. **H.** Ovary cross-section, showing free-central placentation (lacking septa). **I.** *Silene gallica*, side view of flower and immature fruits. **J.** *Silene laciniata*, flower with strongly lobed petals. **K.** *Spergularia villosa*, flower close-up. **L,M.** *Spergula arvensis*. **L.** Swollen node of stem, bearing lobed leaves. **M.** Fruit, a five-valved capsule. **N,O.** *Stellaria media*. **N.** Shoot, showing swollen node and opposite leaves. **O.** Flower, with five bifid petals.

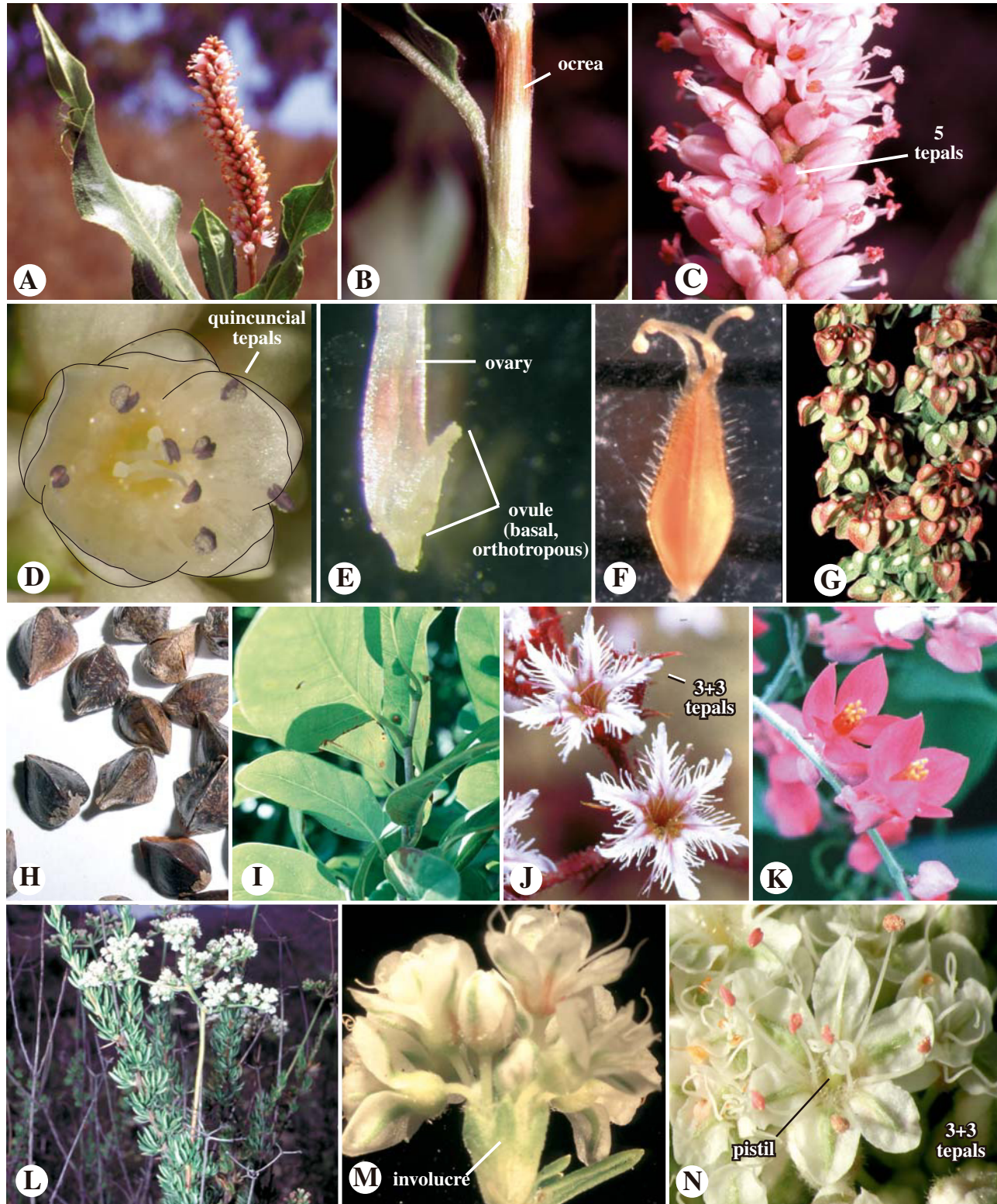


FIGURE 8.17 CARYOPHYLLALES. Polygonaceae. **A,B.** *Polygonum amphibium*. **A.** Shoot with spikelike inflorescence. **B.** Close-up of node, showing ocrea. **C.** Close-up of flowers. **D.** *Polygonum* sp., flower close-up, showing quincuncial aestivation (outlined). **E.** *Eriogonum fasciculatum*, ovary longitudinal section, showing single basal, orthotropous ovule. **F.** *Dedekera eurekensis*, ovary with three styles. **G.** *Rumex crispus*, fruits, surrounded by accrescent calyx. **H.** *Fagopyrum esculentum*, buckwheat, trigonous achenes. **I.** *Cocoloba diversifolia*, sea-grape. **J.** *Chorizanthe mbriata*, spine-flower, showing 3+3 tepals. **K.** *Antigonon leptopus*, coral vine. **L,M.** *Eriogonum fasciculatum*, California buckwheat. **L.** Whole plant. **M.** Fascicle with involucre. **N.** Flower close-up, showing 3+3 tepals, androecium, and superior ovary.

placentation, and a **capsular fruit**, anthocyanin pigments only present.

K 5[4] or (5[4]) **C** 5[4] **A** 5 or 5+5 [1-4] **G** (2-5+), superior, hypanthium usually absent.

Polygonaceae Buckwheat family (meaning many knees, from swollen nodes found in some species). 46 genera / 1100 species. (Figure 8.17)

The Polygonaceae consist of annual or perennial herbs, shrubs, lianas, vines, or trees. The **stems** often have swollen nodes. The **leaves** are usually spiral, simple, stipulate or exstipulate, when stipulate, stipules are typically connate into a scarious, appressed sheath extending above the node, termed an **ocrea**. The **inflorescence** consists of involucrate fasciculate units, the fascicles arranged in various branched or unbranched secondary inflorescences. The **flowers** are hypogynous, small, bisexual or unisexual, actinomorphic, pedicellate, the pedicels often articulated (jointed) above, an **ocreola** often subtending individual flowers. The **perianth** is uniseriate or appearing biseriate (actually spiral), homochlamydeous, usually 3+3, or 5 and quincuncial (rarely 2+2), the tepals (perianth parts) basally connate, hypanthium absent or present. The **stamens** are 3+3 or 8 [2,9+], often of two lengths, generally antitepalous, apostemonous to basally connate. **Anthers** are versatile, longitudinal and introrse in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 [2,4] carpels, and 1 locule. The **styles** are distinct or basally connate. **Placentation** is basal; **ovules** are orthotropous, bi- or unitegmic, solitary. **Nectaries** are often present, consisting of a nectariferous disk or nectary pads at base of stamens. The **fruit** is usually a 3-sided achene or nutlet, sometimes with an accrescent perianth or hypanthium. The **seeds** are endospermous, oily and starchy. Anthocyanin pigments are present, betalains absent. The vasculature is often anomalous.

The Polygonaceae are typically classified into two sub-families: Polygonoideae, with **ocrea** present, and Eriogonoideae, with **ocrea** absent, although the **ocrea**-containing taxa are likely paraphyletic. Members of the family have a worldwide distribution, especially in the northern temperate hemisphere. Economic importance includes edible plants, such as *Fagopyrum esculentum*, buckwheat, and *Rheum × hybridum*, rhubarb; medicinal plants; timber, charcoal, and tanning plants; and a number of cultivated ornamentals, such as *Antigonon leptopus*, coral vine, *Muelenbeckia*, and *Polygonum*. See Lamb Frye and Kron (2003) for a recent phylogenetic analysis of the family.

The Polygonaceae are distinctive in having simple, spiral leaves, with or without a stipular **ocrea**, an inflorescence of **fasciculate units**, small actinomorphic flowers usually with **3+3 or 5 (quincuncial) connate tepals**, a 3 [2,4] carpellate ovary with a **single, basal, mostly orthotropous ovule**,

and a usually **3-sided achene or nutlet**, anthocyanin pigments only present.

P (3+3) or (5) [(2+2)] **A** 3+3, 8 [2,9+] **G** (3) [(2,4)], superior, hypanthium absent or present.

SANTALALES

The Santalales, sensu APG II (2003), contain five families (seven in Stevens, 2001 onwards), of which one is described here. See Nickrent and MalØcot (2001) for more details.

Santalaceae [incl. Viscaceae] Sandal wood family (from Persian name for sandalwood). 41 genera / 925 species. (Figure 8.18)

The Santalaceae (sensu APG II 2003) consist of herbs, shrubs, or trees that are hemiparasites (parasites that carry on photosynthesis). The **roots** are haustorial either on the roots (sandalwoods) or stems (mistletoes) of a host plant. The **stems** are bifurcating (pseudo-dichotomous) in some (e.g., mistletoes), thorns sometimes present. The **leaves** are simple, exstipulate, usually opposite, either typical (bifacial and photosynthetic) or reduced and scale-like. The **inflorescence** is usually of axillary dichasia in a compound inflorescence. The **flowers** are small, bisexual or unisexual, actinomorphic, bracteate, hypogynous to epigynous. The **perianth** is uniseriate, of 3 5 [6 8], usually valvate tepals, apotepalous or syntepalous with a fleshy tube. The **stamens** are 4 5 [3 8], opposite the tepals, apostemonous, free or epitepalous. **Anthers** are longitudinal, porcidial, or transverse in dehiscence, and either tetra-, bi-, or unisporangiate. The **gynoecium** is syncarpous, with a superior to inferior ovary, 3 4 [2,5] carpels, and 1 locule. The **style** is solitary and terminal. **Placentation** is free-central, apical/pendulous, or with a large, basal placenta; **ovules** are 1 4 per ovary. An ovary disk is present in some. The **fruit** is a drupe, nut, berry, or explosive capsule. **Seeds** with seed coat absent, endospermous, the endosperm chlorophyllous in some.

The Santalaceae in the APG II (2003) treatment encompass the families Santalaceae, s.s. (sandal woods) and Viscaceae (mistletoes). Members of the family have a worldwide distribution. Economic importance includes uses as timber, oil, incense (e.g., *Santalum album*, Indian sandalwood), plus some used for edible fruits or tubers and for tanning; mistletoes, esp. *Phoradendron* spp., are used in decorative displays, some are pests on crop trees (e.g., *Arceuthobium*). See Nickrent and MalØcot (2001) for a recent treatment on the family and its relationships.

The Santalaceae are distinctive in being photosynthetic **hemiparasites** with **haustorial** roots attached to roots or



FIGURE 8.18 SANTALALES. Santalaceae [including Viscaceae]. **A.** *Arceuthobium campylopodum*, parasitic on pine. **B,C.** *Phoradendron californicum*, desert mistletoe. **D.** *Comandra umbellata*, close-up of flowers; note uniseriate, valvate perianth. **E.** *Pyrrularia pubera*, buffalo nut. **F.** *Thecium* sp., flower close-ups.

branches of a host plant, having *small*, often *unisexual flowers* with a *uniseriate, valvate perianth*, antitepalous stamens, and free-central/apical or basal placentation, *seeds lacking a seed coat*, the fruit a drupe, nut, berry, or explosively dehiscent.

P 3-5 [6-8] or (3-5) [(6-8)] **A** 3-5 [6-8] **G** (3-4) [(2,5)], superior to inferior.

SAXIFRAGALES

The Saxifragales, sensu APG II (2003) include 10 families (15 sensu Stevens, 2001 onwards), of which two are described here. Notable among the families not treated are the **Altingiaceae** (including *Liquidambar*, sweetgum), **Cercidiphyllaceae** (only 2 species, used as timber trees, much more widespread in the past), **Grossulariaceae** (including *Ribes*, currants/gooseberries, with edible fruits), **Haloragaceae** (including aquarium aquatics such as *Myriophyllum*), **Hamamelidaceae** (including *Hamamelis*, witch-hazel), and **Paeoniaceae** (peony family). See Soltis and Soltis (1997) and Fishbein et al. (2001) for information on the order.

Crassulaceae Stonecrop family (meaning thick or succulent little plant). 33 genera / ca. 1500 species. (Figure 8.19)

The Crassulaceae consist of herbs, shrubs, or rarely trees. The **leaves** are spiral, opposite, or whorled, simple, exstipulate, and characteristically succulent. The **inflorescence** is a branched cyme or of solitary flowers. The **flowers** are usually bisexual, actinomorphic, pedicellate or sessile, hypogynous or slightly perigynous. The **perianth** is biseriate and dichlamydeous. The **calyx** is usually aposepalous, with 5 [rarely 3 6 +] sepals. The **corolla** is apopetalous to basally sympetalous, with 5 [rarely 3 6+] petals. The **stamens** are 1 2× the sepal or petal number, biseriate and obdiplostemonous or uniseriate and antisepalous, free or epipetalous. **Anthers** are longitudinal in dehiscence, tetrasporangiate, dithecal. The **gynoecium** is apocarpous, with a superior ovary and 5 [3 6 or more] carpels. **Placentation** is marginal; **ovules** are anatropous, bitegmic, and numerous [rarely few or 1]. **Nectaries** are present, consisting of scale-like structures at the base of and opposite the carpels. The **fruit** is a follicetum, rarely a capsule. The **seeds** are endospermous (oily and proteinaceous). The stem xylem is usually in a continuous cylinder;

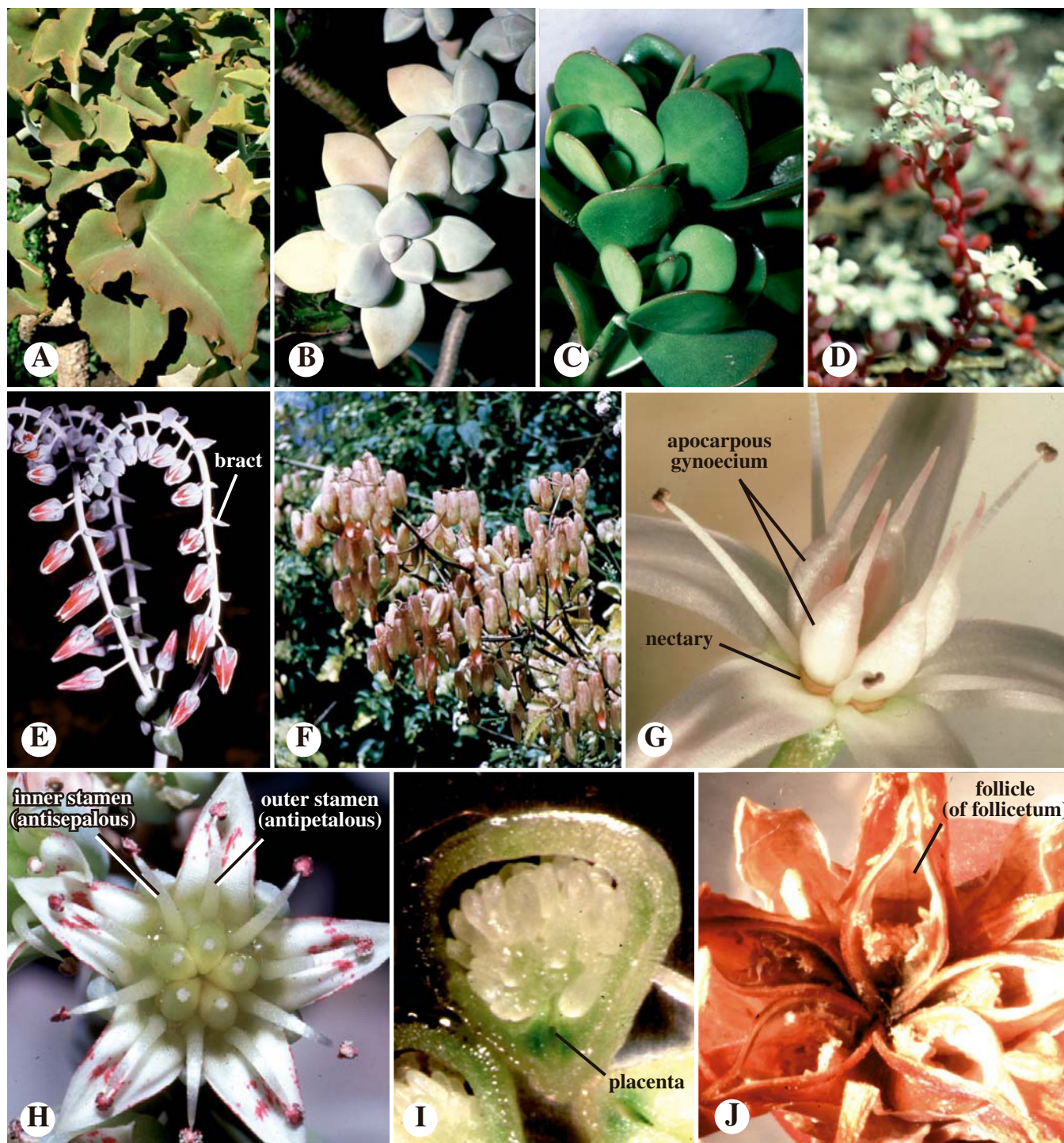


FIGURE 8.19 SAXIFRAGALES. Crassulaceae. A–D. Vegetative morphology. A. *Kalanchoë beharensis*, large shrub with thick, tomentose leaves. B. *Graptopetalum paraguayense*, shrub with succulent, glaucous leaves. C. *Crassula argentea*, shrub with succulent, glabrous leaves. D. *Sedum smallii*, small annual, with minute, succulent leaves. E–J. Reproductive morphology. E. *Dudleya pulverulenta*, showing branched, scorpioid cyme; note bracts displaced ca. 90° from pedicel. F. *Bryophyllum pinnatum*, with pedulous, tubular flowers on large, paniculate inflorescence. G. *Crassula argentea*. Note flower with apocarpous gynoecium and scalelike nectaries at base of and opposite carpels. H, I. *Graptopetalum paraguayense*. H. View of biserial, obdiplostemonous androecium. I. Carpel (ovary) cross-section, showing marginal placentation. J. *Pachyphytum hookeri*, mature follicetum fruit.

leaves often have Kranz anatomy, with Crassulacean Acid Metabolism (CAM) photosynthesis (see Chapter 10).

The Crassulaceae is traditionally treated in six subfamilies, but recent studies discount the monophyly of most of these. Members of the family often grow in arid environments, but also occur in mesic or moist habitats with distributions worldwide, except Australia and Pacific islands; species are most diverse in S. Africa and mtns. of Mexico and Asia. Economic importance includes cultivated ornamentals, especially *Aeonium*, *Crassula*, *Echeveria*, *Kalanchoë*, and *Sedum*. See Ham and Hart (1998) and Mort et al. (2001) for recent phylogenetic analyses.

The Crassulaceae are distinctive in being herbs, shrubs, or rarely trees, with simple, **succulent leaves** having **CAM photosynthesis**, a cymose inflorescence with bisexual, actinomorphic, dichlamydeous flowers, obdiplostemonous or uniseriate stamens, and an **apocarpous gynoecium** with opposed, **scale-like nectaries**, the fruit a **follicetum**.

K 5 [3-6+] **C** 5 [3-6+] **A** 5+5 or 5 [3-6+] **G** 5 [3-6+], superior.

Saxifragaceae Saxifrage family (Latin for rock breaking). 35 genera / 660 species. (Figure 8.20)

The Saxifragaceae consist of perennial herbs or subshrubs. The **leaves** are usually spiral, often rosulate, simple, pinnate, or palmate, usually exstipulate, sometimes succulent. The **inflorescence** is a cyme, raceme, or of solitary flowers. The **flowers** are bisexual, actinomorphic, and hypogynous to epigynous. The **perianth** is biseriate, dichlamydeous. The **calyx** is aposepalous with 5 [3-10] sepals; the **corolla** is sympetalous with 5 [3-10] petals. The **stamens** are variable in number, uniseriate or biseriate, with staminodes present in some. The **gynoecium** is syncarpous and often lobed, rarely apocarpous, with a superior to inferior ovary, and 2-4 [7] carpels. **Placentation** is marginal, axile, or parietal; **ovules** are anatropous, uni- or bitegmic, usually numerous. A **nectariferous** ovary disk is often present. The **fruit** is a (usually) septicidal capsule. The **seeds** are (oily) endospermous.

The Saxifragaceae are widely distributed, especially in northern temperate and cold regions. Economic importance is primarily as cultivated ornamentals. See Soltis and Soltis (1997) and Soltis et al. (2001).

The Saxifragaceae are distinctive in being **perennial herbs, rarely subshrubs**, with spiral, sometimes succulent leaves (often in rosettes), the flowers generally with 5 distinct sepals, 5 connate petals, and 1-2 whorls of stamens, the **gynoecium usually syncarpous and lobed**, the **ovary superior to inferior**, with numerous ovules, the fruit a **septicidal capsule**.

K 5 [3-10] **C** (5) [(3-10)] **A** 5 or 5+5 [variable] **G** (2-4) [(-7)], superior to inferior.

ROSIDS

The Rosids, as delimited by the APG II (2003) system, comprise a very large group of eudicots. Recent molecular studies, upon which the APG II system is based, verify the monophyly of this group. Rosids are largely equivalent to the subclass Rosidae of Cronquist (1981), but also contain several taxa that various authors placed in other groups (particularly within the subclass Dilleniidae of Cronquist).

No clear nonmolecular apomorphies unite the Rosids. Members tend to have perianths with unfused parts and a stamen merosity greater than that of the calyx or corolla, but there are many exceptions. Generally, Rosids have bitegmic, crassinucellate ovules, distinguishing them from the Asterids, which largely have unitegmic, tenuinucellate ovules.

The Rosids are currently delimited into 13-16 orders, many of which are members of the subgroups Eurosids I and Eurosids II (Figures 8.1, 8.21). A listing of families placed within these orders is seen in Table 8.2. Some families are of uncertain placement (insertae sedis); these are listed at the beginning of the Rosids and Eurosids I (Table 8.2).

GERANIALES

The Geraniales, sensu APG II, 2003, contain four families (Table 8.2), only one of which, the Geraniaceae, is described here.

Geraniaceae — Geranium family (Gr. for crane, from accrescent styles resembling a long bird's beak). 11 genera / 700 species. (Figure 8.22)

The Geraniaceae consist of herbs or shrubs. The **stems** are a pachycaul in some taxa. The **leaves** are simple or compound, if simple, usually pinnately or palmately lobed to divided, spiral, rarely opposite, usually stipulate. The **inflorescence** is a cyme or a solitary, axillary flower. The **flowers** are bisexual, actinomorphic (zygomorphic in *Pelargonium*), hypogynous, often bracteate, an epicalyx present in some. The **perianth** is biseriate and dichlamydeous. The **calyx** is aposepalous or synsepalous with 5 [4], imbricate or valvate sepals, the adaxial sepal a nectariferous spur in *Pelargonium*. The **corolla** is apopetalous with 5 [0,4,8] imbricate, rarely convolute petals. The **stamens** are usually 10 [rarely 8 or 15], in two whorls, basally connate, with staminodes present in the outer whorl of some. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 5 [rarely 2, 3, or 8] carpels, and as many locules as carpels. The **style** is usually solitary; **stigmas** are as many as carpels. **Placentation** is apical-axile; **ovules** are anatropous to campylotropous, bitegmic, usually

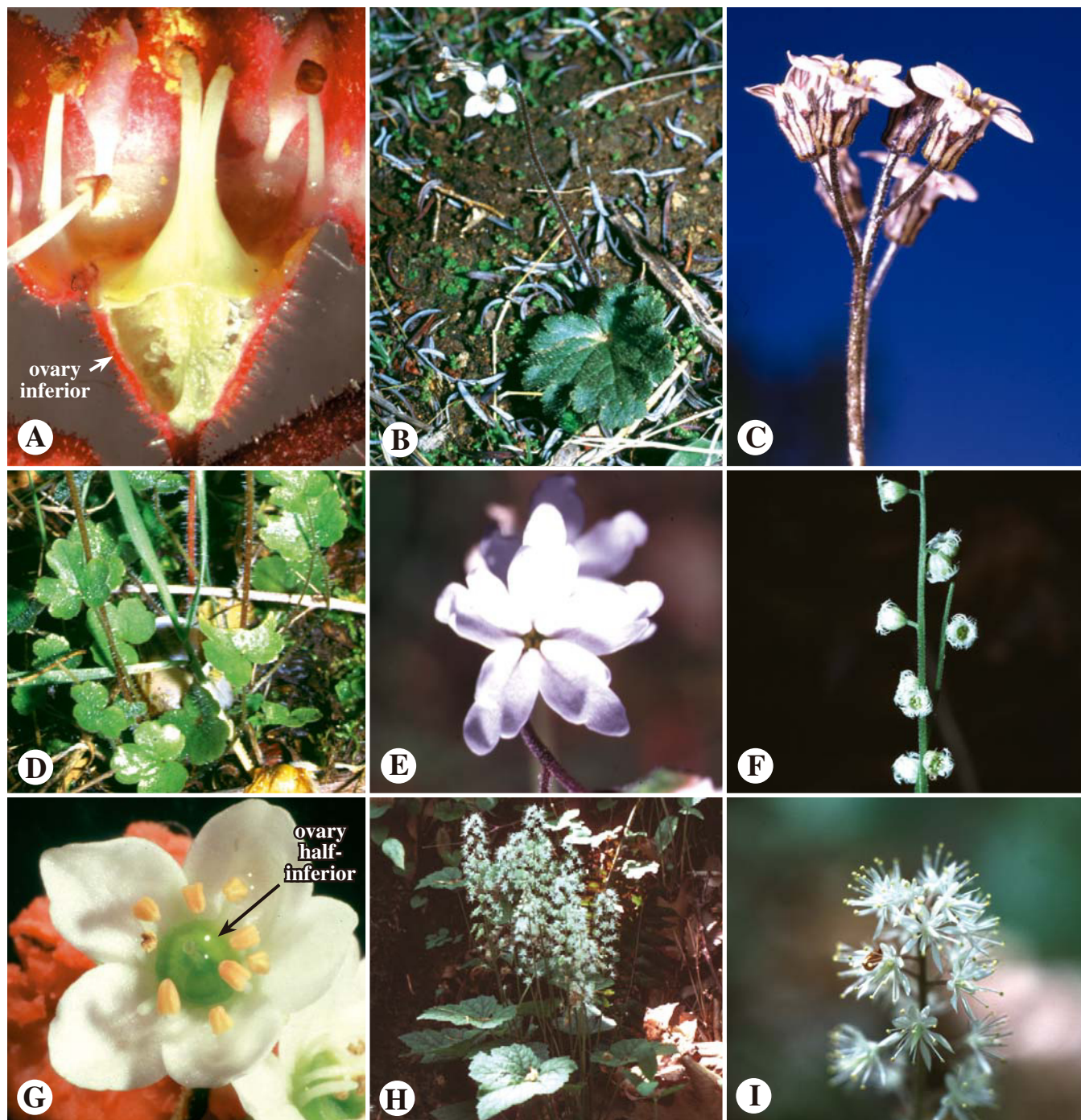


FIGURE 8.20 SAXIFRAGALES. Saxifragaceae. **A.** *Heuchera sanguinea*, showing inferior ovary. **B,C.** *Jepsonia aphylla*. **B.** Whole plant, with single leaf. **C.** Inflorescence. **D,E.** *Lithophragma affine*, woodland star. **D.** Basal leaves. **E.** Flower. **F.** *Mitella diphylla*. **G.** *Saxifraga californica*, flower close-up, with half-inferior ovary. **H,I.** *Tiarella cordifolia*, foam flower.

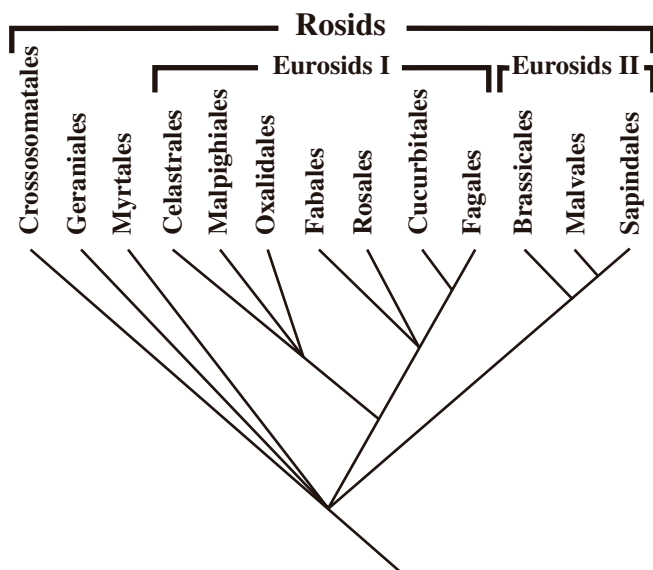


FIGURE 8.21 Cladogram of the orders of the Rosids, after Angiosperm Phylogeny Group II (APG II, 2003).

two per carpel. **Nectaries** are present, between petals and stamens (except in *Pelargonium*). The **fruit** is a loculicidal capsule or a schizocarp of mericarps or follicles, usually separating from a persistent beak arising from an accrescent style. Mucicellular, capitate, glandular trichomes are often present, usually with aromatic oils in trichome glands.

The Geraniaceae contain the tribe Geranieae, including *Geranium*, *Pelargonium*, and *Erodium* (stork's bill, crane's bill, or filaree), distinctive in having a schizocarpic fruit with an elongate, persistent beak. Members of the family are distributed in mostly temperate, some tropical, regions. Economic importance includes the use of taxa as cultivated ornamentals (such as *Geranium* and *Pelargonium*), forage plants (such as *Alfilaria* in w. U.S.), and in essential oil extraction (e.g., *Pelargonium*). See Lis-Balchin (2002) for more information on members of the family.

The Geraniaceae are distinctive in being herbs or shrubs with generally pentamerous, dichlamydeous flowers usually having *nectariferous glands alternating with the petals* and generally *two or more whorls of stamens, staminodes* often present; the tribe Geranieae is distinctive in having *beaked, shizocarpic fruits*.

K 5 [4] **C** 5 [0,4,8] **A** 5+5 [8,15] **G** (5) [(2,3,8)], superior.

MYRTALES

The Myrtales, sensu APG II, 2003, contain 13 families (Table 8.2), of which three are described here. A possible

apomorphy for the order may be an inferior ovary with a hypanthium (epiperigynous perianth/androecial position). Notable among the families not described here are the **Combretaceae** (including savannah and mangrove species, timber and dye plants, and ornamental cultivars) and the **Lythraceae** (the loosestrife family, including dye and timber plants, ornamental cultivars such as *Cuphea* and *Lagerstroemia*, and the food plant *Punica granatum*, pomegranate [formerly classified in the Punicaceae]). See Conti et al. (1996, 1998) for detailed studies of the Myrtales.

Melastomataceae Melastome family (Greek for black mouth, from fruits that stain). 188 genera / 4,950 species. (Figure 8.23)

The Melastomataceae consist of shrubs, herbs, rarely trees or lianas. The **stems** are often 4-sided. The **leaves** are simple, opposite, rarely whorled, usually exstipulate, often with 3–9 subparallel, major veins. The **inflorescence** is a cyme. The **flowers** are bisexual, epiperigynous, with perianth mostly actinomorphic, the androecium zygomorphic. The **perianth** has a hypanthium usually present. The **calyx** consists of 4–5 [3–10], valvate or calyptrate sepals. The **corolla** is apopetalous, with 4–5 [3–10] connate, volute petals. The **stamens** are usually 8 or 10, biseriate, often dimorphic; filaments are often twisted during anthesis, positioning the anthers to one side of flower. **Anthers** are poricidal with 1–2 pores per anther, rarely longitudinal in dehiscence, the anther connective often appendaged. The **gynoecium** is syncarpous, with an inferior ovary, 3–5 [2–15] carpels, and locules, or unilocular by formation of incomplete septa. **Placentation** is usually axile; **ovules** are anatropous to campylotropous, bitegmic, with zig-zag micropyle, ∞ [1] per carpel. The **fruit** is a loculicidal or septifragal capsule or a berry. The **seeds** are exalbuminous.

The Melastomataceae have distributions in tropical regions, especially South America. Economic importance includes timber trees, edible fruit plants, dye plants, and several ornamental cultivars, such as *Tibouchina*. See Clausing and Renner (2001) for more detailed studies of the family.

The Melastomataceae are distinctive in being shrubs, herbs, rarely trees or lianas, the **stems often 4-sided**, with *simple, opposite* (rarely whorled) **leaves**, usually with *3–9 subparallel major veins*, the inflorescence a cyme, *flowers epiperigynous*, the perianth usually 4–5-merous, the **stamens biseriate and dimorphic**, anthers oriented to one side of flower by filament twisting, *connective often appendaged and dehiscence usually poricidal*, the inferior ovary with usually axile placentation, and fruit a capsule or berry.

K 4–5 [3–10] **C** 4–5 [3–10] **A** 4+4 or 5+5 **G** (3–5) [(2–15)], inferior, hypanthium usually present.

TABLE 8.2 Orders and families of the Rosids, after APG II (2003). Families in **bold** are described in detail. An asterisk denotes a deviation from APG II, with brackets indicating the more inclusive family recommended by APG II. A double asterisk indicates a change suggested by Angiosperm Phylogeny Website (Stevens 2001 onwards). See Table 8.3 for listing of orders and families in the Asterids.

ROSIDS		
Picramniaceae	FAGALES	ROSALES
VITALES**	Betulaceae	Barbeyaceae
Vitaceae	Casuarinaceae	Cannabaceae
CROSSOSOMATALES	Fagaceae	Dirachmaceae
Aphloiaceae**	Juglandaceae	Elaeagnaceae
Crossosomataceae	(incl. Rhoipteleaceae)	Moraceae
Geissolomataceae**	Myricaceae	Rhamnaceae
Ixerbaceae**	Nothofagaceae	Rosaceae
Stachyuraceae	Ticodendraceae	Ulmaceae
Staphyleaceae	MALPIGHIALES	Urticaceae
Strasburgeriaceae**	Achariaceae	EUROSIDS II
GERANIALES	Balanopaceae	HUERTEALES**
Geraniaceae	Bonnetiaceae	Dipentodontaceae**
Ledocarpaceae	Caryocaraceae	Tapisciaceae
Melianthaceae	Centropiaceae**	BRASSICALES
Vivianiaceae	Chrysobalanaceae	Akaniaceae
MYRTALES	Clusiaceae (Guttiferae)	Bataceae
Alzateaceae	Ctenolophonaceae	Brassicaceae (Cruciferae)*
Combretaceae	Elatinaceae	Capparaceae* [Brassicaceae]
Cryteriaceae	Erythroxylaceae* [Rhizophoraceae]	Caricaceae
Heteropyxidaceae	Euphorbiaceae s.s.	Cleomaceae* [Brassicaceae]
Lythraceae	Goupiaceae	Emblingiaceae
Melastomataceae	Humiriaceae	Gyrostemonaceae
Myrtaceae	Hypericaceae	Koeberliniaceae
Oliniaceae	Irvingiaceae	Limnanthaceae
Onagraceae	Ixonanthaceae	Moringaceae
Penaecaceae	Lacistemataceae	Pentadiplandraceae
Psiloxylaceae	Linaceae	Resedaceae
Rhynchocalycaceae	Lophopyxidaceae	Salvadoraceae
Vochysiaceae	Malpighiaceae	Setchellanthaceae
EUROSIDS I	Ochnaceae	Tovariaceae
Huaceae	Pandaceae	Tropaeolaceae
ZYGOPHYLLALES**	Passi oraceae	MALVALES
Krameriaceae* [Zygophyllaceae]	Phyllanthaceae	Bixaceae
Zygophyllaceae*	Picrodendraceae	Cistaceae
CELASTRALES	Podostemaceae	Dipterocarpaceae
Celastraceae	Putranjivaceae	Malvaceae
Lepidobotryaceae	Rafesiaceae**	Mutingiaceae
Parnassiaceae	Rhizophoraceae*	Neuradaceae
CUCURBITALES	Salicaceae	Sarcolaenaceae
Anisophylleaceae	Violaceae	Sphaerosepalaceae
Begoniaceae	OXALIDALES	Thymelaeaceae
Coriariaceae	Brunelliaceae	SAPINDALES
Corynocarpaceae	Cephalotaceae	Anacardiaceae
Cucurbitaceae	Connaraceae	Biebersteiniaceae
Datiscaceae	Cunoniaceae	Burseraceae
Tetramelaceae	Elaeocarpaceae	Kirkiaceae
FABALES	Oxalidaceae	Meliaceae
Fabaceae (Leguminosae)		Nitriaceae
Polygalaceae		Rutaceae
Quillajaceae		Sapindaceae
Surianaceae		Simaroubaceae

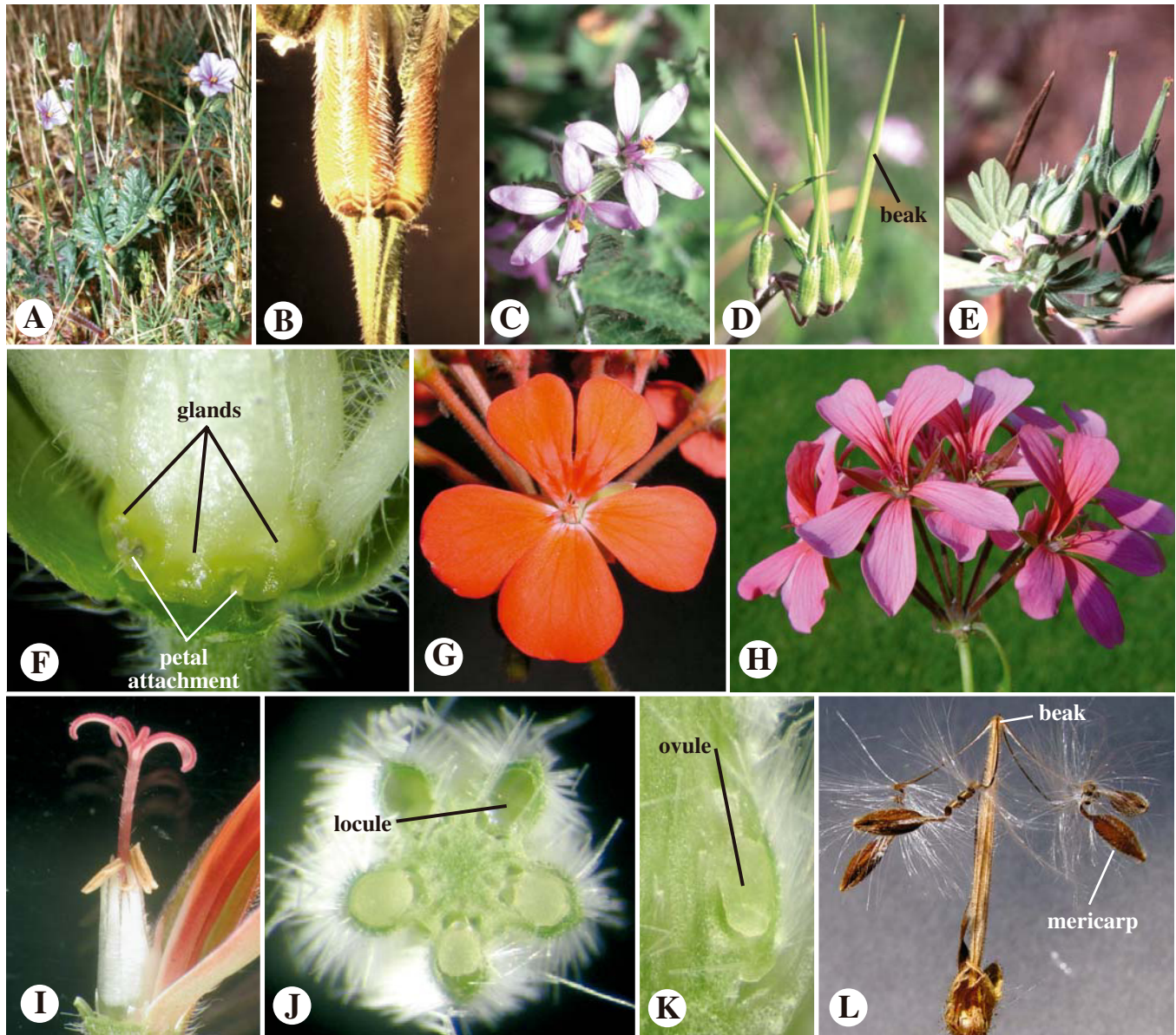


FIGURE 8.22 GERANIALES. Geraniaceae. **A,B.** *Erodium botrys*, storksbill. **A.** Whole plant in flower. **B.** Base of mature schizocarpic fruit. **C,D.** *Erodium moschatum*. **C.** Flowers. **D.** Young fruits, showing accrescent styles, forming beak. **E.** *Geranium carolinianum*, in flower and immature fruit. **F.** *Geranium* sp., base of flower with sepals and petals removed, showing alternipetalous glands. **G–L.** *Pelargonium* spp. **G.** Flower, face view, slightly zygomorphic. **H.** Inflorescence. **I.** Androecium and gynoecium, showing style branches. **J.** Cross-section of ovary, strongly 5-lobed. **K.** Ovary lobe longitudinal section, showing apical-axile placentation. **L.** Fruit, a beaked schizocarp of mericarps.

Myrtaceae Myrtle family (*myrtus*, Gr. name for myrtle). 120 genera / 3850 species. (Figure 8.24)

The Myrtaceae consist of trees and shrubs. The **roots** possess ectotrophic mycorrhizae. The **stems** have secretory cavities and internal phloem in the pith. **Leaves** are opposite (usually) or spiral, rarely whorled, simple, glandular-punctate or pellucid, and often coriaceous, with stipules present and small or absent. The **inflorescence** is variable. The **flowers**

are bisexual, actinomorphic, bracteate, epiperigynous, rarely perigynous. The **perianth** is biseriate, perianth segments distinct or connate, fused into a lidlike calyptra (operculum) in some (e.g., *Eucalyptus*). The **calyx** consists of 4–5 [3,6] sepals. The **corolla** consists of 4–5 [3,6] petals. The **stamens** are ∞ and are distinct or connate into 4 or 5 groups. **Anthers** are loculicidal or poricidal in dehiscence. The **gynoecium** is syncarpous, with an inferior [rarely half-inferior or superior]

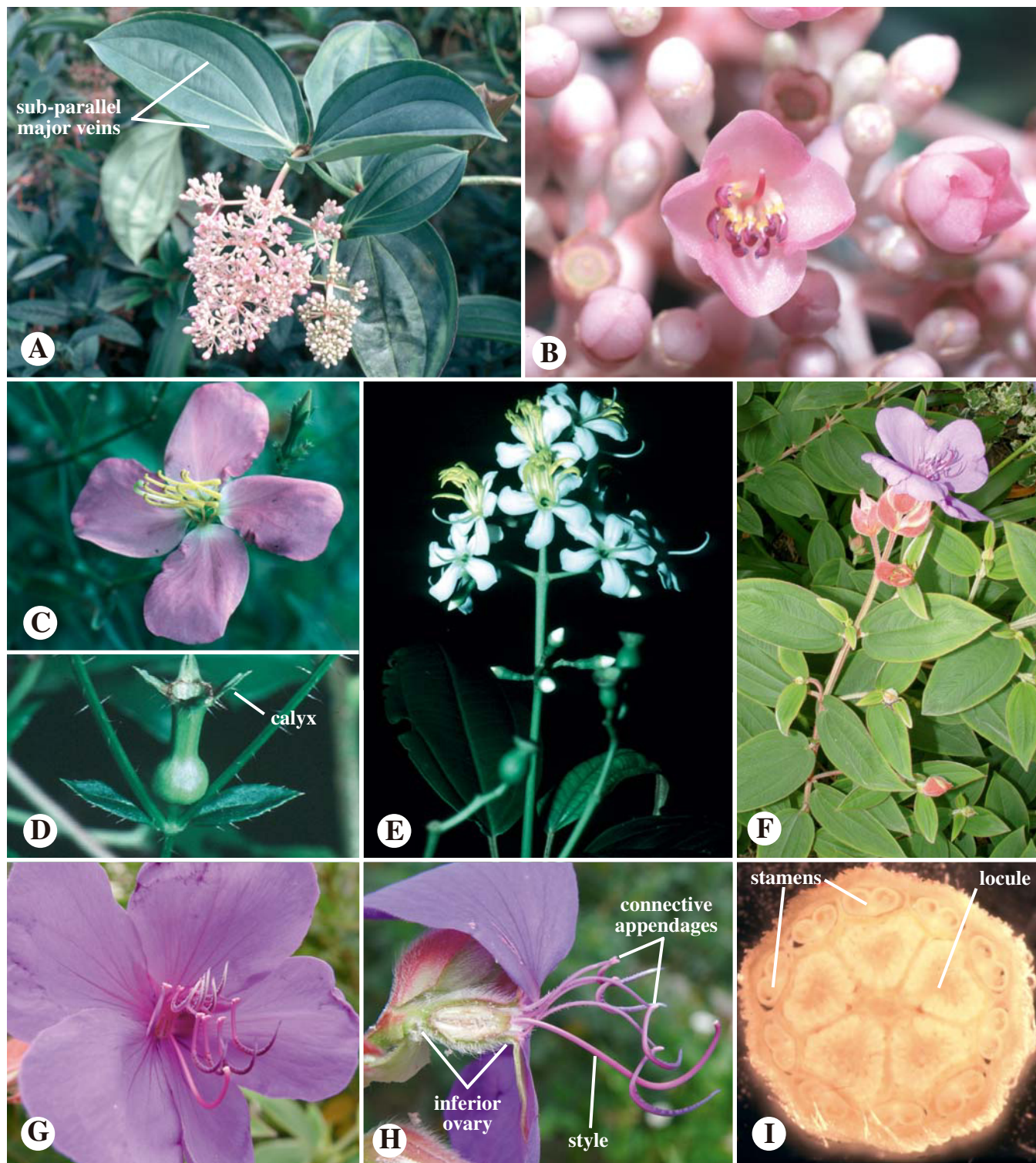


FIGURE 8.23 MYRTALES. Melastomataceae. **A,B.** *Medinilla* sp. **A.** Flowering shoot, showing subparallel major veins. **B.** Flower close-up. **C,D.** *Rhexia mariana*. **C.** Flower. **D.** Urn-shaped fruit; note inferior ovary. **E.** *Tetrazygia bicolor*, in orescence. **F-H.** *Tibouchina urvilleana*. **F.** Flowering shoot. **G.** Flower, face view. **H.** Flower, longitudinal section, showing inferior ovary and stamen connective appendages. **I.** *Tibouchina multijuga*, bud cross-section, showing ovary with axile placentation and recessed anthers of stamens.



FIGURE 8.24 MYRTALES. Myrtaceae. **A.** *Actinodium cunninghamii*, having a capitulum resembling that of the Asteraceae. **B,C.** *Callistemon citrinus*. **B.** Note capsules and flower spike, which continues apical vegetative growth. **C.** Close-up of flower, showing reduced perianth and numerous, showy stamen filaments. **D.** *Calothamnus sanguineus*, with connate, zygomorphic stamens (in bundles). **E.** *Darwinia fascicularis*, having heads and flowers with elongate styles. **F.** *Eucalyptus obtusiflora*, showing capsules and flowers. **G.** *Eucalyptus torquata*. Note calyptra and incurved stamens (prior to unfurling). **H,I.** *Eucalyptus sideroxylon*. **H.** Flower longitudinal section, showing inferior ovary and hypanthium. **I.** Style and stigma close-up. **J.** Ovary cross-section, showing axile placentation. **K.** *Eucalyptus calophylla*, a very tall tree species. **L.** *Leptospermum laevigatum*. **M.** *Myrtus communis*, leaf close-up showing internal, pellucid glands. **N.** *Tristania conferta*, Brisbane box, the flower with fascicles of stamens. **O.** *Verticordia grandiflora*, with fringed petals.

ovary, 2 5 [16] carpels, and 2 5 [16, 1] locules. The **style** is terminal; the **stigma** is capitate or lobed. **Placentation** is axile (being basal-axile to apical-axile) [parietal if unilocular]; **ovules** are anatropous or campylotropous, bitegmic or unitegmic, 2 ∞ per locule. **Nectaries** are present, as a disk atop the ovary or on the inner hypanthium. The **fruit** is a berry or loculicidal capsule [rarely a drupe or nut].

The Myrtaceae are typically classified into two subfamilies (with several tribes): Leptospermoideae, with fruits usually a capsule and leaves spiral or opposite, and Myrtoideae, with fruits fleshy and leaves always opposite. Members of the family have distributions in warm tropics and temperate Australia. Economic importance includes important timber trees, especially *Eucalyptus* spp., edible fruits (e.g., *Psidium guajava*, guava), spices (e.g., *Syzygium aromaticum*, cloves, *Pimenta dioica*, allspice), oils (e.g., *Eucalyptus* spp.), and cultivated ornamentals such as *Callistemon* (bottlebrush), *Chamelaucium* (wax-flower), *Eucalyptus* spp., *Leptospermum* (tea tree), and *Myrtus* (myrtle). See Wilson et al. (2001) for a detailed study of the family.

The Myrtaceae are distinctive in being trees and shrubs with **glandular-punctate or pellucid leaves** and usually **epiperigynous flowers** with **numerous stamens**.

K 4-5 [3,6] **C** 4-5 [3,6] **A** ∞ **G** (2-5) [(-16)], inferior [rarely half-inferior or superior], with hypanthium.

Onagraceae Evening-Primrose family (after *onagra*, Gr. for oleander, an unrelated plant). 24 genera / 650 species. (Figure 8.25)

The Onagraceae consist of terrestrial or aquatic herbs and shrubs, rarely trees. The **stems** have internal phloem, often with epidermal oil cells. The **leaves** are spiral, opposite, or whorled, simple, undivided to pinnatifid, and exstipulate or stipulate. The **inflorescence** is a spike, panicle, or of solitary flowers. The **flowers** are bisexual (usually), actinomorphic, and epiperigynous, the hypanthium elongate in some taxa. The **perianth** is biseriate and dichlamydeous. The **calyx** is valvate, aposepalous, consisting of usually 4 [rarely 2 6] sepals. The **corolla** is valvate or imbricate, apopetalous, consisting of usually 4 [rarely 2 6 or 0] petals. The **stamens** are 4+4 [rarely 2 6]. **Anthers** are longitudinal in dehiscence, tetrasporangiate and dithecal, with cross-partitions in some species. The **pollen** is shed in monads or tetrads, often with viscin threads, which function to adhere grains together. The **gynoecium** is syncarpous, with an inferior ovary, and usually 4 [rarely 2 6] carpels and locules. **Placentation** is axile or parietal; **ovules** are anatropous, bitegmic, usually ∞ per locule, with a monosporic, 4-nucleate (*Oenothera* type) female gametophyte. The **fruit** is a capsule, berry, or nut. **Seeds** are (oily) endospermous.

The Onagraceae have a worldwide distribution. Economic importance includes several cultivated ornamentals, such as species of *Clarkia*, *Fuchsia*, and *Oenothera*. See Levin et al. (2003) for a detailed phylogenetic study of the family.

The Onagraceae are distinctive in being herbs and shrubs (rarely trees) with usually **4-merous [2-6-merous], epiperigynous flowers** with usually **4+4 stamens** and a **monosporic, 4-nucleate female gametophyte**, the latter a possible apomorphy for the family.

K 4 [2-6] **C** 4 [2-6,0] **A** 4+4 [2-6] **G** (4) [(2-6)], inferior, with hypanthium.

EUROSIDS I

This subgroup of the Rosids contains 7 8 orders plus one unplaced family (Table 8.2, Figure 8.21). Some of the orders are quite large, both in terms of number of families and number of species. Among the Eurosids I group are taxa of great agricultural importance, such as members of the Cucurbitaceae (squash family), Fabaceae (bean/pea family), Rosaceae (rose family), and Euphorbiaceae (spurge family). Others are of great ecological or industrial significance, such as the oaks (Fagaceae).

CUCURBITALES

This order, sensu APG II, 2003, contains seven families (Table 8.2), one of which is described here. Well known among the other families are the **Begoniaceae**, with *Begonia* a prominent ornamental cultivar. See Zhang and Renner (2003) for a recent study of the order.

Cucurbitaceae Cucumber / Gourd family (L. for gourd). 120 genera / 775 species. (Figure 8.26)

The Cucurbitaceae consist of monoecious or dioecious [rarely hermaphroditic] vines [rarely tree-like], usually with one tendril per node. The **leaves** are simple, palmately veined and often palmately lobed, spiral, and exstipulate. The **inflorescence** is axillary, variable in type or with flowers solitary. The **flowers** are usually unisexual, actinomorphic, the female flowers epiperigynous. The **perianth** is biseriate and dichlamydeous, with hypanthium present. The **calyx** is aposepalous with 5 [3 6] imbricate sepals. The **corolla** is apopetalous or sympetalous with 5 [3 6] valvate petals. The **stamens** are 3 5, alternipetalous, distinct or connate. **Anthers** are longitudinal in dehiscence, dithecal or monotheical. The **gynoecium** is syncarpous, with an inferior ovary, 3 [1 5] carpels, and 1 locule [locules rarely as many as carpels]. The **styles** are

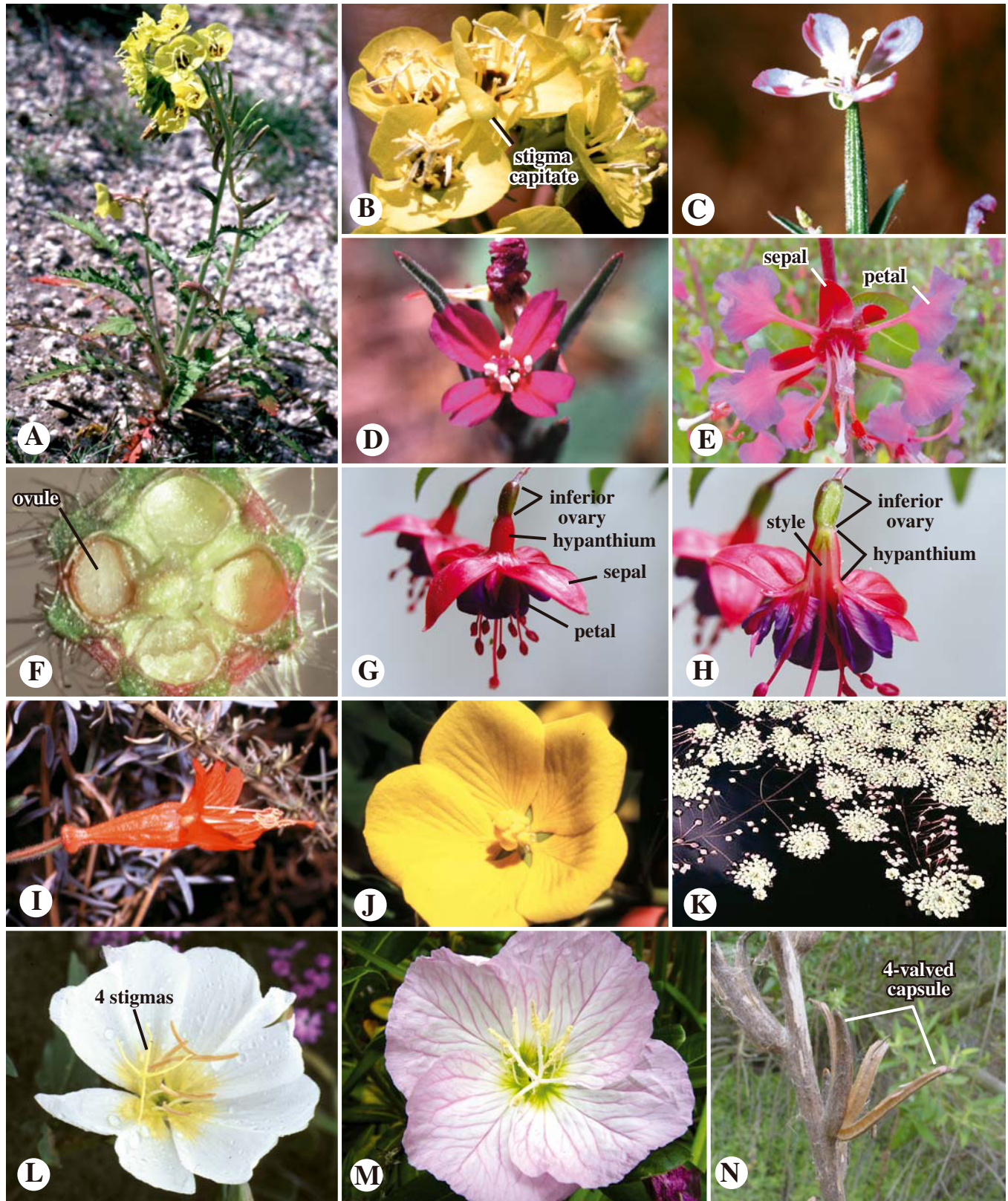


FIGURE 8.25 MYRTALES. Onagraceae. **A,B.** *Camissonia claviformis*, with capitulate stigma. **C.** *Clarkia delicata*. **D.** *Clarkia purpurea*. **E,F.** *Clarkia unguiculata*. **E.** Flower. **F.** Ovary cross-section, with axile placentation. **G,H.** *Fuchsia xhybrida*. **G.** Pendant flower. **H.** Flower in longitudinal section, showing epiperigynous structure. **I.** *Epilobium canum*, with zygomorphic flowers. **J.** *Ludwigia hexapetala*, with pentamerous flowers. **K.** *Ludwigia sedoides*, a floating aquatic. **L.** *Oenothera deltoides*, with four linear stigmas. **M.** *Oenothera speciosa*. **N.** *Oenothera elata*, showing four-valved capsule.

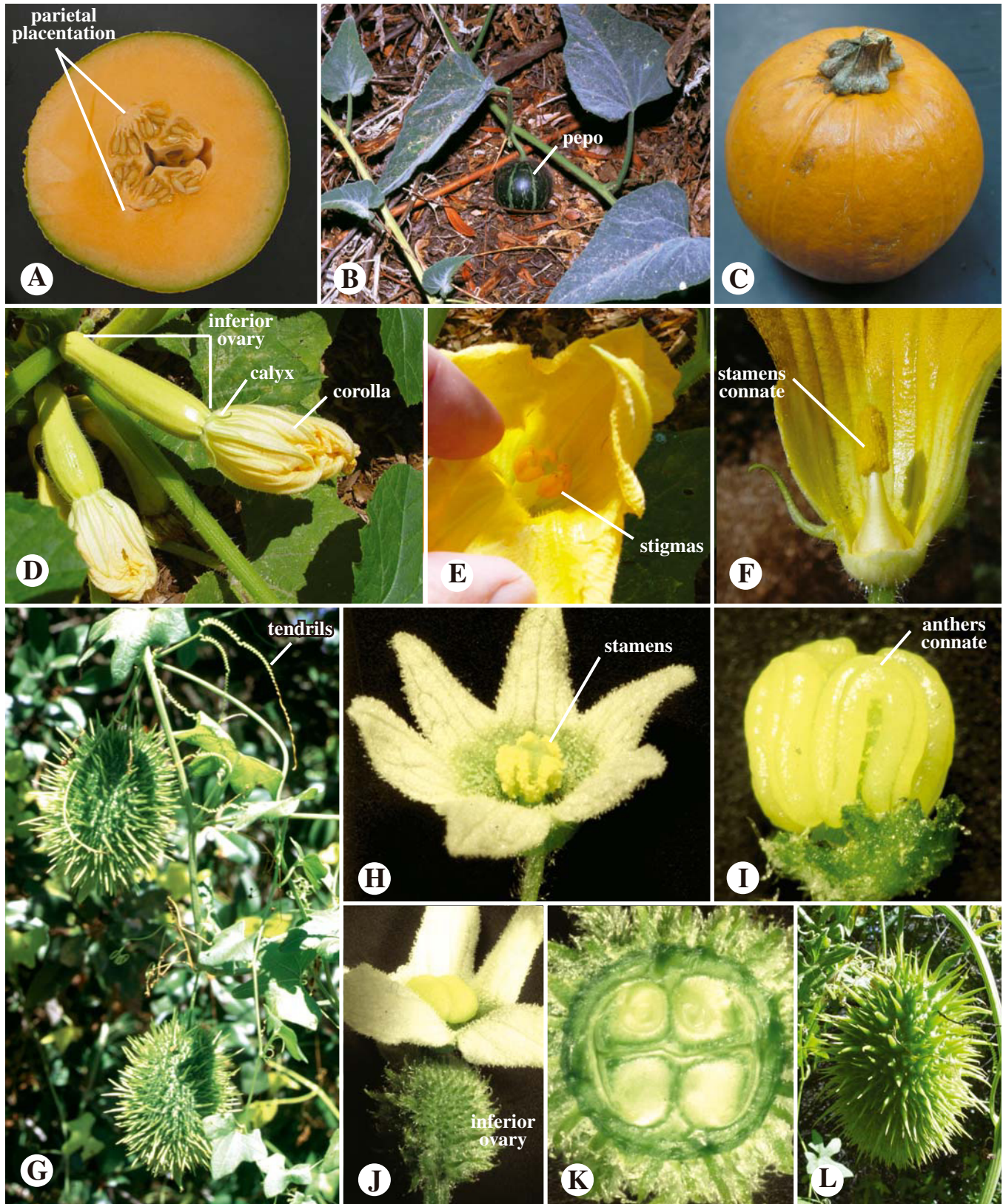


FIGURE 8.26 CUCURBITALES. Cucurbitaceae. **A.** *Cucumis melo*, cantaloupe. **B.** *Cucurbita foetidissima*, calabazilla. **C.** *Cucurbita pepo*, a pumpkin. **D–F.** *Cucurbita pepo*, a squash. **D,E.** Female flower. **F.** Male flower. **G–L.** *Marah macrocarpus*, manroot. **G.** Whole plant. Note vine habit, simple, palmately lobed leaves, and tendrils. **H.** Male flower. **I.** Close-up of connate anthers of male flower. **J.** Female flower, showing inferior ovary. **K.** Ovary cross-section, showing parietal placentation (only two carpels). **L.** Immature, prickly fruit.

1 3; **stigmas** are 1 2. **Placentation** is parietal, rarely axile; **ovules** are anatropous, bitegmic, generally ∞ [rarely 1 fe w]. Extrafloral **nectaries** are often present. The **fruit** is a berry, pepo, capsule, or samara. The **seeds** are exalbuminous. Stem anatomy is typically bicyclic, with bicollateral vascular bundles.

The Cucurbitaceae have largely worldwide distributions, but occur mostly in tropical regions. Economic importance includes important food crops such as *Citrullus lanatus* (watermelon), *Cucumis melo* (melons), *Cucumis sativa* (cucumber), *Cucurbita pepo* and other spp. (squashes, pumpkins) and a number of other taxa; the dried fruits of several species are used as gourds, those of *Luffa* (luffa) are used as a sponge; some taxa have medicinal or horticultural uses. See Renner et al. (2002) for a recent study of the family.

The Cucurbitaceae are distinctive in being mostly *monoecious or dioecious vines* with *simple, palmately veined and/or lobed leaves*, usually with *tendrils*, the *female flowers epiperigynous*, with usually *parietal placentation and three carpels*, the fruit a berry, pepo, capsule, or samara. **K** 5 [3-6] **C** 5 [3-6] or (5) [(3-6)] **A** 3-5 or (3-5) **G** (3) [(2-5)], inferior, hypanthium present.

FABALES

The Fabales, sensu APG II, 2003, contain four families (Table 8.2), the two most common of which are described here. See Forest et al. (2002) for a recent study of the order.

Fabaceae (Leguminosae) — Bean/Pea family (after *faba*, Latin name for broad bean). 643 genera / 18,000 species. (Figures 8.27, 8.28)

The Fabaceae consist of herbs, shrubs, trees, or vines, with spines sometimes present. The **roots** of many members have a symbiotic association with nitrogen-fixing bacteria (*Rhizobium* spp.), which induce formation of root nodules (this especially common in the Faboideae). The **leaves** are usually compound (pinnate, bipinnate, trifoliolate, rarely palmate), sometimes simple or unifoliolate, usually spiral, basal pulvini often present, sometimes functioning in tactile (thigmonastic), leaflet folding responses (e.g., *Mimosa* spp.), generally stipulate, sometimes stipellate, stipules spinose in some. The **inflorescence** is variable, typically bracteate. The **flowers** are usually bisexual, sometimes unisexual, actinomorphic or zygomorphic, pedicellate or sessile, hypogynous or perigynous. The **perianth** is biseriate, dichlamydeous, with a hypanthium sometimes present. The **calyx** is aposepalous or synsepalous with 5 [3 6] sepals. The **corolla** is apopetalous or sympetalous, with 5 [rarely 0] valvate or imbricate petals. The **stamens** are 5 or 10 ∞ , distinct

or connate. **Anthers** are longitudinal, rarely poricidal in dehiscence. The **gynoecium** is unicarpellous, with a superior ovary, 1 [rarely 2 or more in some Mimosoideae] carpel, and 1 locule. The **style** and **stigma** are solitary. **Placentation** is marginal; **ovules** are anatropous or campylotropous, bitegmic, 2 ∞ [1] per carpel. **Nectaries** are often present as a ring at the base of the ovary. The **fruit** is generally a legume, sometimes indehiscent (e.g., *Arachis*, peanut), winged (a samara), drupe-like, or divided into transverse partitions (a loment).

The Fabaceae are traditionally classified into three subfamilies (sometimes treated as separate families): Caesalpinioideae, Mimosoideae, and Faboideae (=Papilionoideae).

The **Caesalpinioideae** are distinctive in having generally *zygomorphic flowers* with usually **5 or 10** [1 ∞] *distinct stamens* (staminodes present in some) and a corolla (imbricate in bud) with typically *five distinct petals* (sometimes reduced or lacking), the *posterior, median petal inner* to (overlapped by) the two lateral petals. The Caesalpinioideae is likely paraphyletic and, hence, placed in quotations here.

The **Mimosoideae** are distinctive in having *actinomorphic flowers* with a corolla of typically *five, distinct or basally fused petals* (valvate in bud), a hypanthium sometimes present, and with usually *numerous, distinct or basally fused stamens*.

The **Faboideae (Papilionoideae)** are distinctive in having “*papilionaceous*” flowers, which are *zygomorphic*, with *connate stamens* (either *10 monadelphous* or *9+1 diadelphous*) and a corolla (imbricate in bud) with five petals consisting of a *large, median, usually posterior petal* (the *banner* or *standard*), which is *outer* to (overlapping) the adjacent petals, *two lateral “wing” petals* (overlapped by the banner), and *two anterior, distally fused “keel” petals*. The flowers are resupinate in some species, e.g., *Clitoria*, in which the banner is anterior in position.

The Fabaceae is a very large group with a worldwide distribution. Members of the family are dominant species in some ecosystems (e.g., *Acacia* spp. in parts of Africa and Australia) and ecologically important for containing nitrogen-fixing rhizobial nodules. Economically, legumes are one of the important plant groups, being the source of numerous pulses (such as *Arachis hypogaea*; peanut; *Glycine max*, soybeans; *Lens culinaris*, lentil; *Phaseolus* spp., beans; *Pisum sativum*, peas); flavoring plants (such as *Ceratonia siliqua*, carob), fodder and soil rotation plants (such as *Medicago sativa*, alfalfa, or *Trifolium* spp., clovers) oils, timber trees, gums, dyes, and insecticides. See Doyle et al. (2000), Bruneau et al. (2001), and references therein for information on the phylogeny of the Fabaceae.

The Fabaceae are distinctive in being trees, shrubs, vines, or herbs, with *stipulate*, often *compound* leaves and typically pentamerous flowers, usually with a *single, unicarpellous pistil* with *marginal placentation*, the fruit a *legume* (or modified legume).

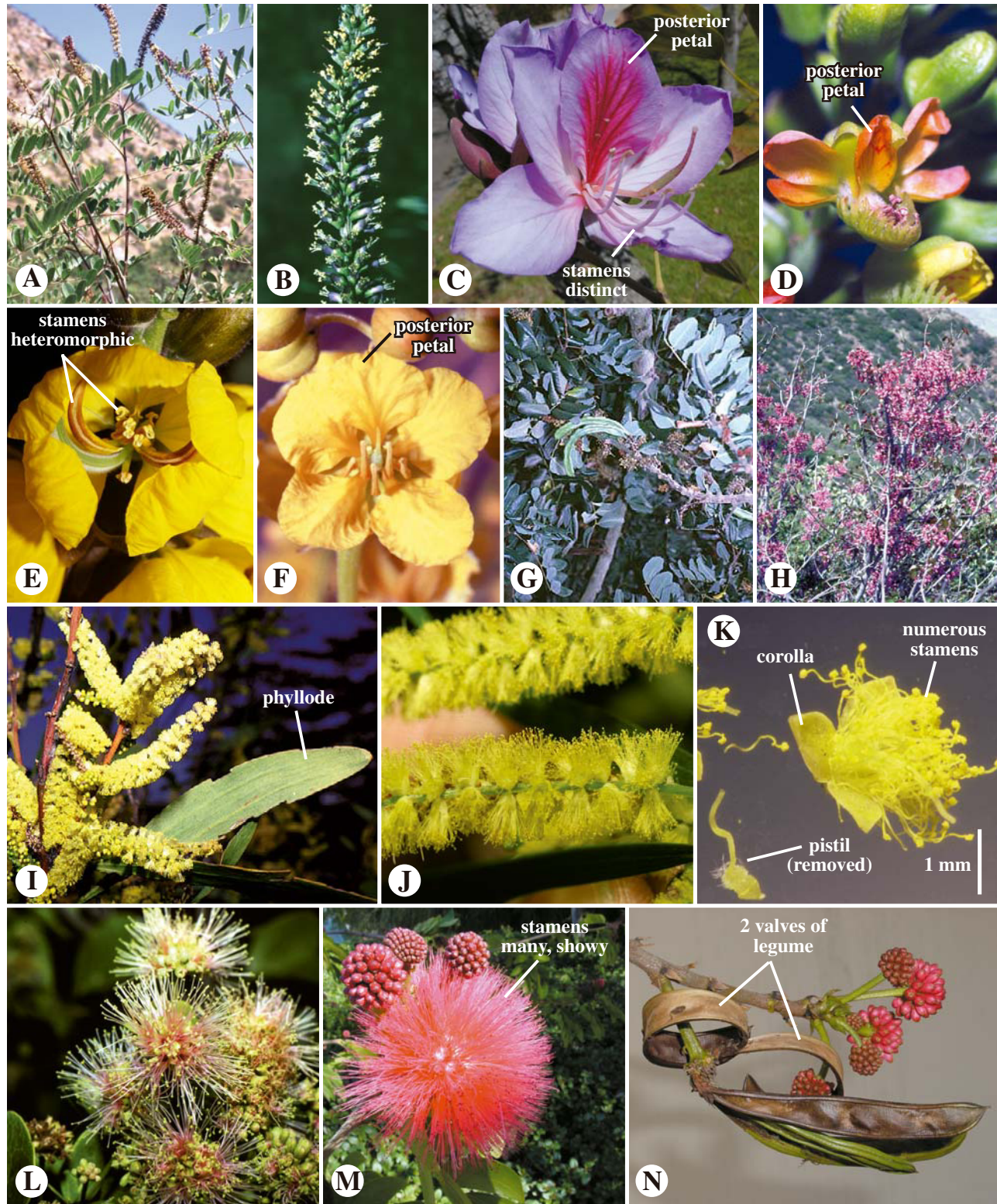


FIGURE 8.27 FABALES. Fabaceae. A–H. "Caesalpinoideae." A,B. *Amorpha fruticosa*. A. Whole plant. B. Close-up of flowers of spike. C. *Bauhinia variegata*. Note medial posterior petal inner to laterals and distinct stamens. D. *Caesalpinia spinosa*. Note unusual pectinate anterior sepal. E. *Cassia didymobotrya*, with heteromorphic (trimorphic) stamens. F. *Senna armata*, a desert species. G. *Ceratonia siliqua*, carob, showing legumes. H. *Cercis occidentales*, red bud. I–N. Mimosoideae. I–K. *Acacia longifolia*. I. Leaves (phyllodes) and spicate inflorescences. J. Close-up of flowers of spike. K. Individual flower. Note inconspicuous perianth, numerous stamens, and ovary (removed from another flower). L. *Pithecellobium unguis-cati*, cat-claw, showing heads of flowers. M,N. *Calliandra haematocephala*. M. Inflorescence heads, the flowers with very long, showy stamens. N. Shoot with fruits (legumes) and immature heads.

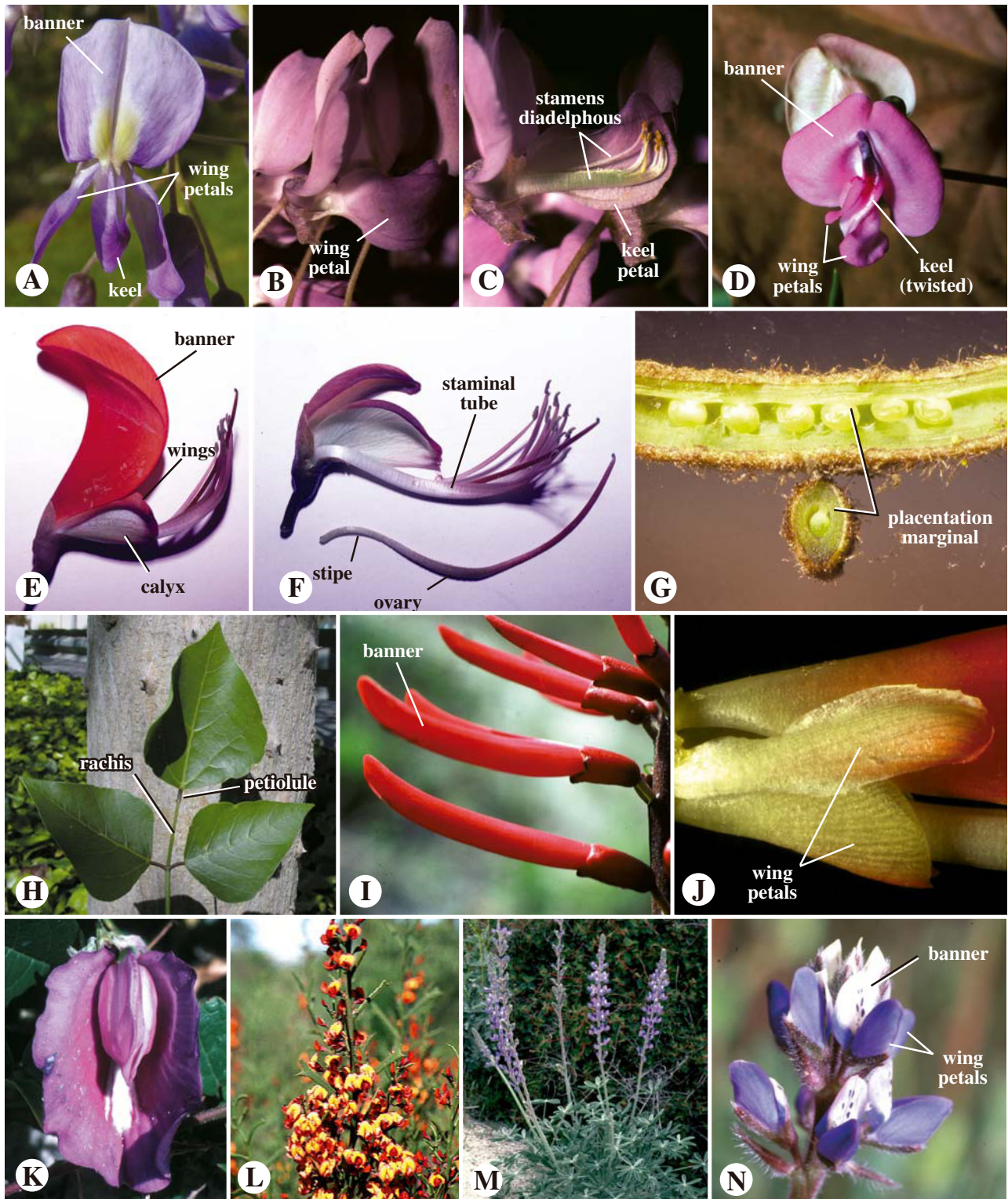


FIGURE 8.28 FABALES. Fabaceae, Faboideae (Papilionoideae). A–C. *Wisteria sinensis*. A. Papilionaceous lower, face view. Note large banner, outer to wing petals. B. Side view, showing banner and one of two wing petals. C. Side view, one wing and keel petal removed, showing diadelphous stamen fusion. D. *Strophostyles umbellata*, with assymmetric owers. E,G. *Erythrina caffra*. E. Flower, side view, showing banner and reduced wing and keel petals. F. Dissected ower, showing staminal tube (monadelphous stamen fusion) and removed stipitate ovary. G. Immature legume in longitudinal (above) and cross- (below) section, showing marginal placentation. H–J. *Erythrina corraloides*, coral bean tree. H. Pinnate-ternate leaf. I. Flowers, side view, with elongate, semitubular banner petals. J. Close-up of reduced wing petals. K. *Clitoria mariana*, a resupinate papilionoid, with banner below, keel above. L. *Daviesia* sp., a papilionoid. M. *Lupinus excubitus*, with palmate leaves. N. *Lupinus bicolor*, ower close-ups, with contrasting, spotted banner.

K 5 or (5) [(3-6)] **C** 5 or (5) [0,1-6, or (1-6)] **A** 5, 10, or (∞) [1- ∞] **G** 1 [2-16], superior, hypanthium sometimes present.

Polygalaceae—Milkwort family (Greek for much milk, in the belief that some species consumed by cows increase milk flow). 17 genera / 950 species. (Figure 8.29)

The Polygalaceae consist of trees, shrubs, lianas, or herbs. The **leaves** are simple, spiral, usually exstipulate (modified as a pair of glands or spines in some). The **inflorescence** is a spike, raceme, or panicle. The **flowers** are bisexual, zygomorphic, hypogynous to perigynous, and subtended by a pair of bracteoles. The **perianth** is biseriate, a hypanthium present in some. The **calyx** is usually aposepalous, sepals 5, the two inner, latero-posterior sepals often petaloid (resembling wing petals), rarely all or the two anterior sepals basally connate. The **corolla** is often adnate to the androecium forming a tube, petals 5 or 3 (the latter by suppression or loss of two lateral petals), when 3, the median-anterior (lower) petal apically fringed and boat-shaped. The **stamens** are 4+4, 10, or 3-7, usually basally connate forming a staminal tube. **Anthers** are longitudinal or apically poricidal in dehiscence. The **gynoecium** is syncarpous with a superior ovary, 2-5 [8] carpels, and locules [locule rarely 1]. The **style** is often curved, often 2-lobed, one lobe stigmatic, the other sterile and comose. **Placentation** is usually apical-axile; **ovules** are pendulous, epitropous, anatropous to hemitropous, bitegmic, 1 per carpel [rarely 1 per ovary]. **Nectaries** consist of a nectariferous disk surrounding the base of the ovary; extrafloral nectaries are present in many species. The **fruit** is a loculicidal capsule, nut, samara, or drupe. The **seeds** are arillate (with caruncle) and endospermous (proteinaceous).

Members of the Polygalaceae have a mostly worldwide distribution. Economic importance includes some ornamental cultivars and plants of local medicinal value. See Persson (2001) for a study of the family.

The Polygalaceae are distinctive in being trees, shrubs, lianas, or herbs, with **simple, spiral**, usually exstipulate leaves, the flowers bisexual, the perianth biseriate, with the **2 inner (of 5) sepals often petaloid** (resembling wing petals), petals 3-5, when 3, the anterior petal often **apically fringed and boat-shaped**, the anthers **poricidal or longitudinally dehiscent**, the style **often 2-lobed with one lobe stigmatic, the other sterile, ovule 1 per carpel**, the seeds **arillate** (with caruncle). **K** 5 or (5) or (2)+3 **C** 3 or 5 **A** 4+4, 10, 3-7 **G** (2-5) [(-8)], superior.

FAGALES

The Fagales, sensu APG II, 2003, contain seven families (Table 8.2), members of which are largely monoecious and

wind pollinated. Two families are described here. Notable among the others are the **Casuarinaceae** (including *Casuarina*, Australian-pine), **Juglandaceae** (walnut family, including *Carya*, pecan, and *Juglans*, walnut), **Myricaceae** (wax-myrtle family), and **Nothofagaceae** (southern-beech family, which include important timber trees). See Li et al. (2002) and Manos and Steele (1997) for detailed treatments of the order.

Betulaceae—Birch family (Latin name for birch or for pitch, derived from bark). 6 genera / ca. 110 species. (Figure 8.30)

The Betulaceae consist of monoecious trees or shrubs. The **leaves** are simple, deciduous, usually spiral, caducous-stipulate, with the margin usually toothed. The **inflorescences** are unisexual, the male inflorescence is a pendulous catkin, the female inflorescence a short, pendulous or erect catkin, both bearing numerous, 1-3-flowered, bracteate, simple dichasia. The **flowers** are unisexual, hypogynous or epigynous. The **perianth** is uniseriate (by default termed a **calyx**), of 1-6 [0], scale-like sepals/lobes. A **corolla** is absent. The **stamens** are 1- ∞ , generally the same number as the perianth parts, antisepalous. **Anthers** have thecae either divided along the connective or connate. The **gynoecium** is syncarpous, with an inferior or superior ovary (the latter sometimes termed **nude** because of lacking a perianth and therefore a point of reference for ovary position), and 2-3 carpels; locules are 2-3 below, 1 above. The **styles** are 2-3, distinct. **Placentation** is apical-axile, the ovules pendulous from the apex of the septa; **ovules** are anatropous, unitegmic or bitegmic, 1-2 per locule. The **fruit** is a nut or 2-winged samara, subtended by woody bracts of a cone-like infructescence or partially enclosed by leafy bracts. The **seeds** are with or without endosperm. Plants are wind pollinated.

The Betulaceae are distributed in northern temperate and mountainous, tropical regions. The family is usually divided into two subfamilies: Betuloideae (including *Alnus*, alders, and *Betula*, birches) with male flowers in groups of 3 (a full, simple dichasium), and the Coryloideae (including *Corylus*, hazels/filberts, *Carpinus*, ironwood, and *Ostrya*, hornbeam) with male flower units reduced to 1. Economic importance of the family includes lumber trees (some woods very dense), chemical derivatives, nuts (e.g., *Corylus*, hazels/filberts), cultivated ornamentals, and numerous uses by aboriginal people.

The Betulaceae are distinctive in being **monoecious trees or shrubs** with simple, toothed leaves, and bearing **pendulous, elongate male catkins** and **pendulous to erect female catkins**, each with numerous, **bracteate dichasia**, the fruit a **nut or 2-winged samara**.

P 1-6 [0] **A** 1- ∞ **G** (2-3), superior (nude) or inferior.

Fagaceae Oak family (Latin for the beech tree). 8 genera / 700 species. (Figure 8.31)

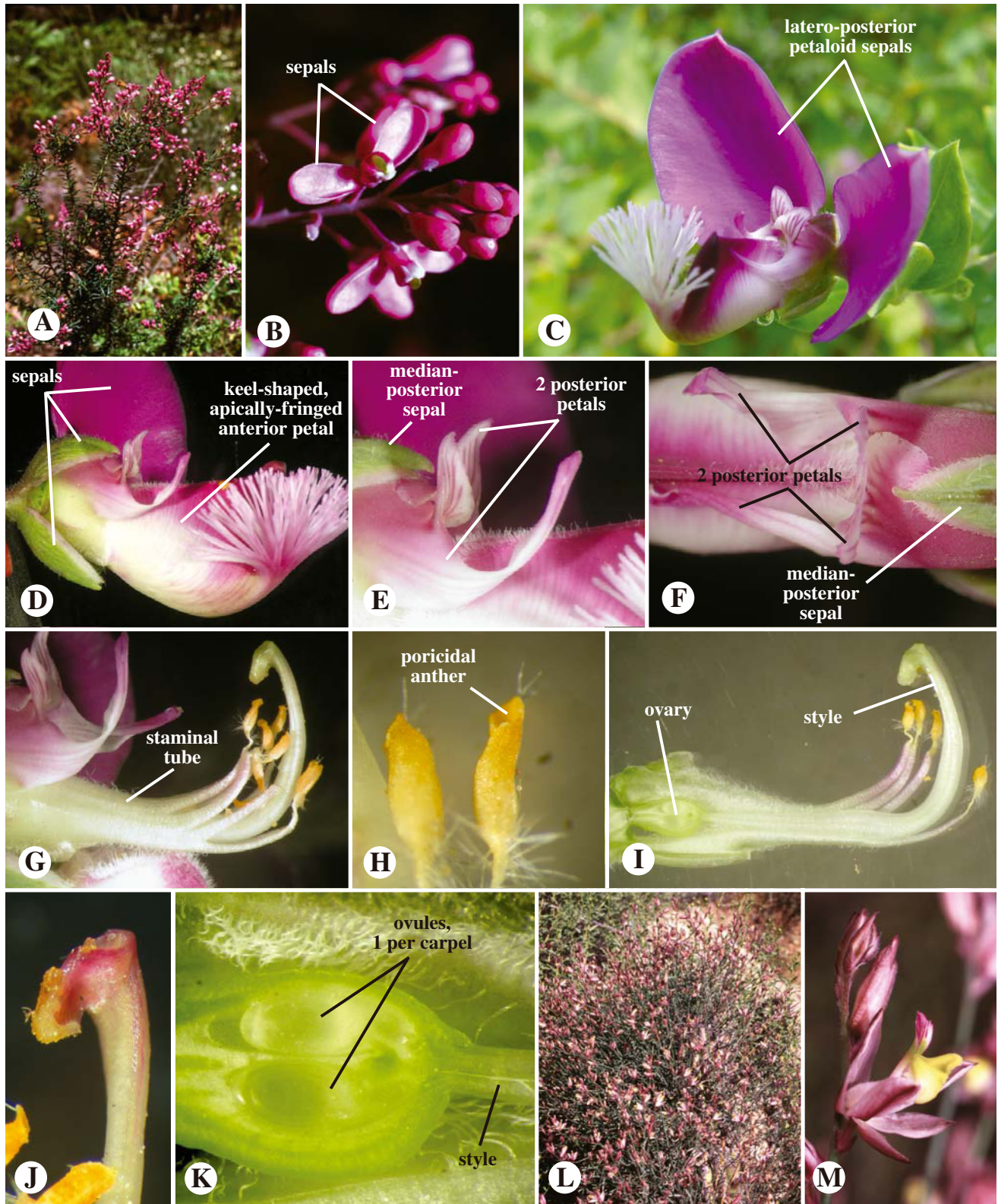


FIGURE 8.29 FABALES. Polygalaceae. **A,B.** *Comesperma ericinum*. **A.** Shrub. **B.** Flowers, showing winglike, petaloid sepals. **C–J.** *Polygala dalmaisiana*. **C.** Flower, showing two latero-posterior, petaloid sepals. **D.** Flower close-up, showing three of five sepals (one petaloid sepal removed) and petals of perianth, the anterior petal keel-shaped and apically fringed. **E.** Side view of two posterior petals, each two-lobed. **F.** Top view of two posterior petals. **G.** Androecium, the filaments basally fused into a staminal tube. **H.** Poricidal anthers. **I.** View of pistil, with style extending through middle of staminal tube. **J.** Style tip, showing two-lobed stigma. **K.** Ovary longitudinal-section, showing ovules with apical-axile placentation. **M,N.** *Polygala desertorum*.

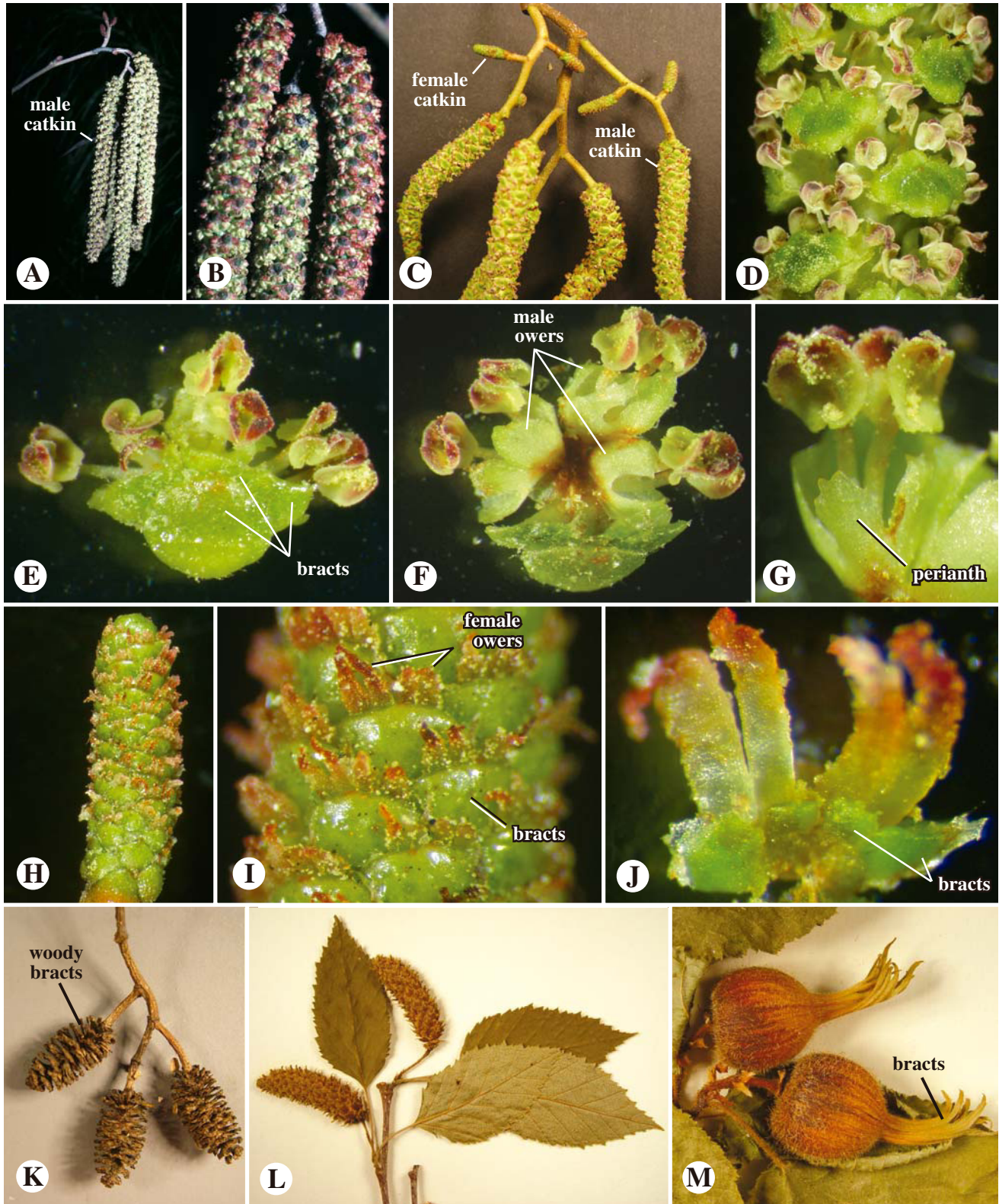


FIGURE 8.30 FAGALES. Betulaceae. **A,B.** *Alnus serrulata*, showing male catkins. **C–K.** *Alnus rhombifolia*, white alder. **C.** Male and female catkins. **D.** Close-up of male catkin. **E,F.** Male simple dichasium unit, three owers with subtending bracts. **G.** Male ower, with perianth. **H.** Female catkin. **I.** Catkin, close-up, showing styles exerted from bracts. **J.** Female owers with subtending bracts. **K.** Inflorescence with woody-textured bracts. **L.** *Betula papyrifera*, paper birch, with inflorescence. **M.** Fruits of *Corylus cornuta*, beaked hazelnut, with subtending bracts.

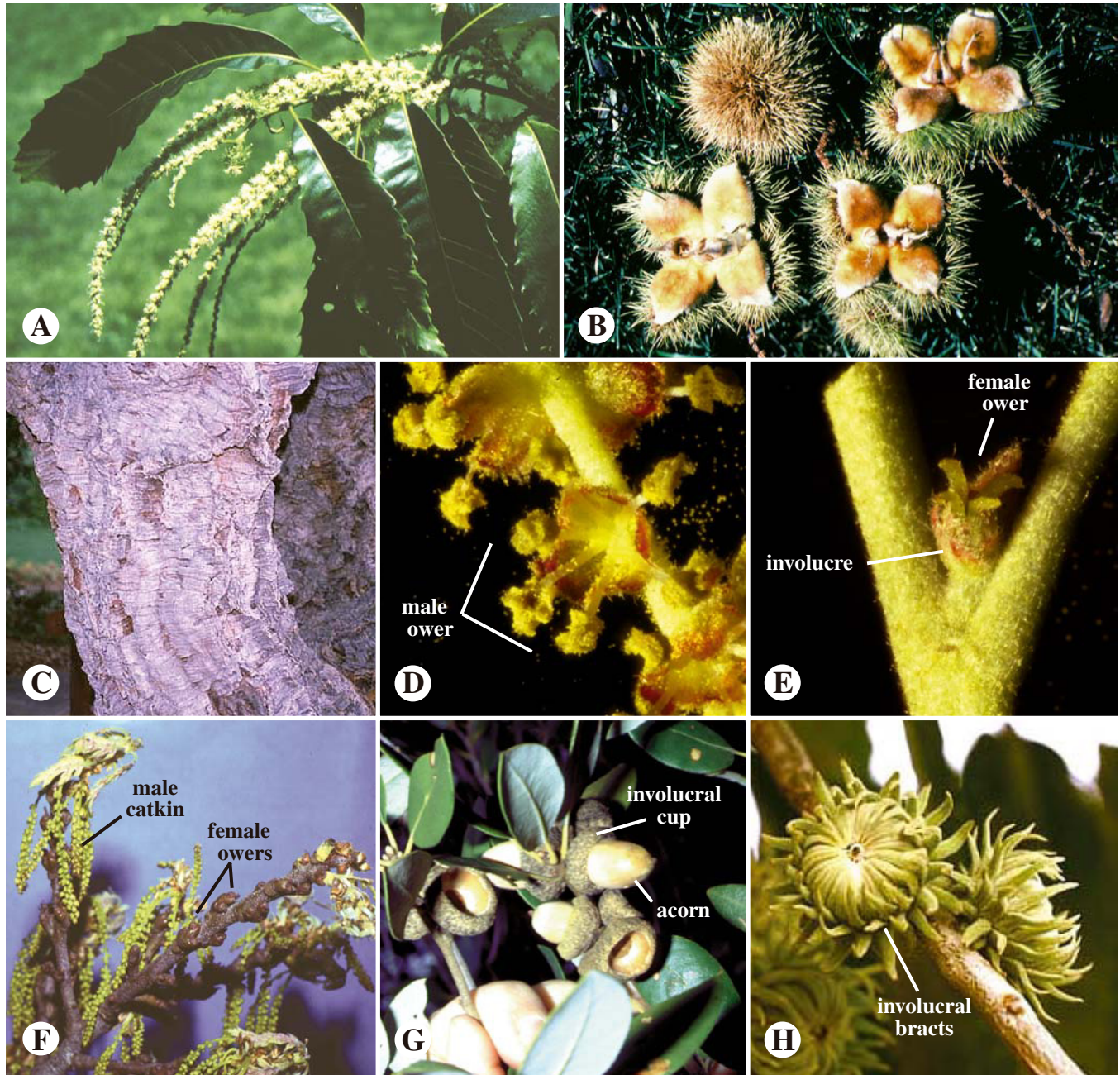


FIGURE 8.31 FAGALES. Fagaceae. **A,B.** *Castanea dentata*, chestnut. **A.** Male spikes. **B.** Fruits, showing prickly involucre bracts. **C-E.** *Quercus suber*, cork oak. **C.** Outer bark, from which commercial cork is derived. **D.** Male flowers of catkin. **E.** Female flower, showing styles and young involucre bracts. **F.** *Quercus* sp., showing catkins and female flowers. **G.** *Quercus chrysolepis*, with mature acorns in which involucre bracts are fused into acorn cup. **H.** *Quercus acutissima*, with prominent, relatively distinct involucre bracts.

The Fagaceae consist of monoecious, rarely dioecious, trees or shrubs. The **leaves** are simple, undivided to divided, usually spiral, rarely opposite or whorled, stipulate, the stipules deciduous. The **inflorescence** is usually unisexual, the male inflorescence a catkin or head of reduced dichasia, the female flowers located at the base of male inflorescences or solitary.

The **flowers** are small, unisexual, actinomorphic, the female flowers epigynous and involucre, the involucre bracts often fused forming a cupule (e.g., acorn cup). The **perianth** is composed of 6 [4–9] tepals. The **stamens** are 6–12 [4–90], distinct. **Anthems** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with an inferior ovary, and usually

3 or 6 [2,7-12] carpels, the locules as many as carpels basally, opening to one locule apically. The **styles** are as many in number as carpels. **Placentation** is basally axile; **ovules** are anatropous, bitegmic, 2 per carpel. The **fruit** is a nut (sometimes termed a glans), with a usually hard pericarp and subtended by accrescent involucre bracts, forming the cupule. The **seeds** are exalbuminous. Plants are usually wind pollinated, although they are insect pollinated in *Castanea*.

The Fagaceae have a mostly worldwide distribution in nontropical regions. Economic importance includes important lumber trees, such as *Quercus* (oak), *Fagus* (beech), and *Castanea* (chestnut); the outer bark of *Quercus suber* is the source of commercial cork; the seeds of various species have been a traditionally important source of food for humans and other animals. See Manos et al. (2001) for a recent study of the Fagaceae.

The Fagaceae are distinctive in being *monoecious* (rarely dioecious) *trees or shrubs* with *simple leaves* (sometimes divided), the *flowers unisexual and small*, the male flowers in *catkins or heads of reduced dichasia*, the female at base of male inflorescences or solitary, with an inferior, multicarpellate ovary, the fruit a *nut with subtending cupule of connate involucre bracts*.

Male: **P** 6 [4-9] **A** 6-12 [4-90]

Female: **P** 6 [4-9] **G** (3,6) [(2,7-12)], inferior.

MALPIGHIALES

The Malpighiales, sensu APG II, 2003, contain ca. 29 (-31) families (Table 8.2), of which three are described here. Notable among the other members of the order are **Erythroxylaceae** (containing *Erythroxylum* spp., the source of the alkaloid cocaine), **Linaceae** (containing *Linum usitatissimum*, flax, the source of linen cloth), **Malpighiaceae** (containing edible fruit plants and hallucinogens such as *Banisteriopsis*), **Rhizophoraceae** (the mangrove family), and **Salicaceae** (*Salix*, the willows, being the original source of salicin, from which aspirin was derived). See Chase et al. (2002) for a recent phylogenetic analysis of families of this order.

Euphorbiaceae Spurge family (after Euphorbus, physician to the king of Mauritania, 1st century). ca. 300 genera / ca. 8,000 species. (Figure 8.32)

The Euphorbiaceae consist of monoecious or dioecious, herbs, shrubs, vines, or trees, latex present in some major groups (Crotonoideae and Euphorbioideae). The **stems** are succulent and cactuslike in some (e.g., some Euphorbias). The **leaves** are simple, rarely trifoliate or palmate, spiral, opposite, or whorled, stipules generally present, these sometimes modified

as glands or spines (e.g., many succulent Euphorbias). The **inflorescence** is generally a cyme, modified as a cyathium in the Euphorbioideae. The **flowers** are unisexual, actinomorphic, rarely zygomorphic, bracteate in some, hypogynous. The **perianth** is biseriate, uniseriate, or absent, generally 5-merous. The **calyx** is usually aposepalous with 5 [rarely 0] sepals. The **corolla** is usually apopetalous with 5 [rarely 0] valvate or imbricate petals. The **stamens** are 1 ∞ , distinct or connate. **Anthers** are longitudinal, poricidal, or transverse in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 [2 ∞] carpels and locules. The **styles** are as many as carpels, each style sometimes 2-branched. **Placentation** is apical-axile and pendulous, with an obturator (a protuberance arising from the funiculus or placenta at the base of the ovule); **ovules** are anatropous or hemitropous, bitegmic, 1 per carpel. **Nectaries** are often present. The **fruit** is a schizocarp, drupe, berry, or samara.

The Euphorbiaceae have traditionally been classified into five subfamilies: Phyllanthoideae, Oldfieldioideae, Acalyphoideae, Crotonoideae, and Euphorbioideae. The former two subfamilies have two ovules per carpel (bioovulate); the latter three subfamilies have one ovule per carpel (unioovulate). In the APG II (2003) classification, only the latter three subfamilies are included within the Euphorbiaceae, as is designated here. The bioovulate Phyllanthoideae and Oldfieldioideae are elevated to family rank, Phyllanthaceae and Picrodendraceae, respectively (in the Malpighiales). Of the three remaining subfamilies of the Euphorbiaceae, the Crotonoideae and Euphorbioideae are similar in having a usually milky (white) latex (red or yellow in some Crotonoideae), and the Euphorbioideae have a characteristic cyathium as an inflorescence. However, a recent phylogenetic study has suggested further subdivision of the family (see Wurdack and Chase, 2002).

Members of the Euphorbiaceae have worldwide distributions. Economic importance includes *Ricinus communis*, the source of castor bean oil and the deadly poison ricin; *Hevea brasiliensis*, the major source of natural rubber; *Manihot esculentus*, cassava/manioc, a very important food crop and the source of tapioca; and various oil, timber, medicinal, dye, and ornamental plants. Succulent *Euphorbia* species are major components of plant communities in southern Africa, as well as important ornamental cultivars.

The Euphorbiaceae are distinctive in having *unisexual* flowers with a superior, *usually 3-carpellate* ovary with *1 ovule per carpel, apical-axile* in placentation; Crotonoideae and Euphorbioideae have a *red, yellow, or usually white* ("milky") *latex* and the Euphorbioideae alone have a characteristic *cyathium* inflorescence.

K 5 [0] **C** 5 [0] **A** 1- ∞ **G** (3) [(2 ∞)], superior.

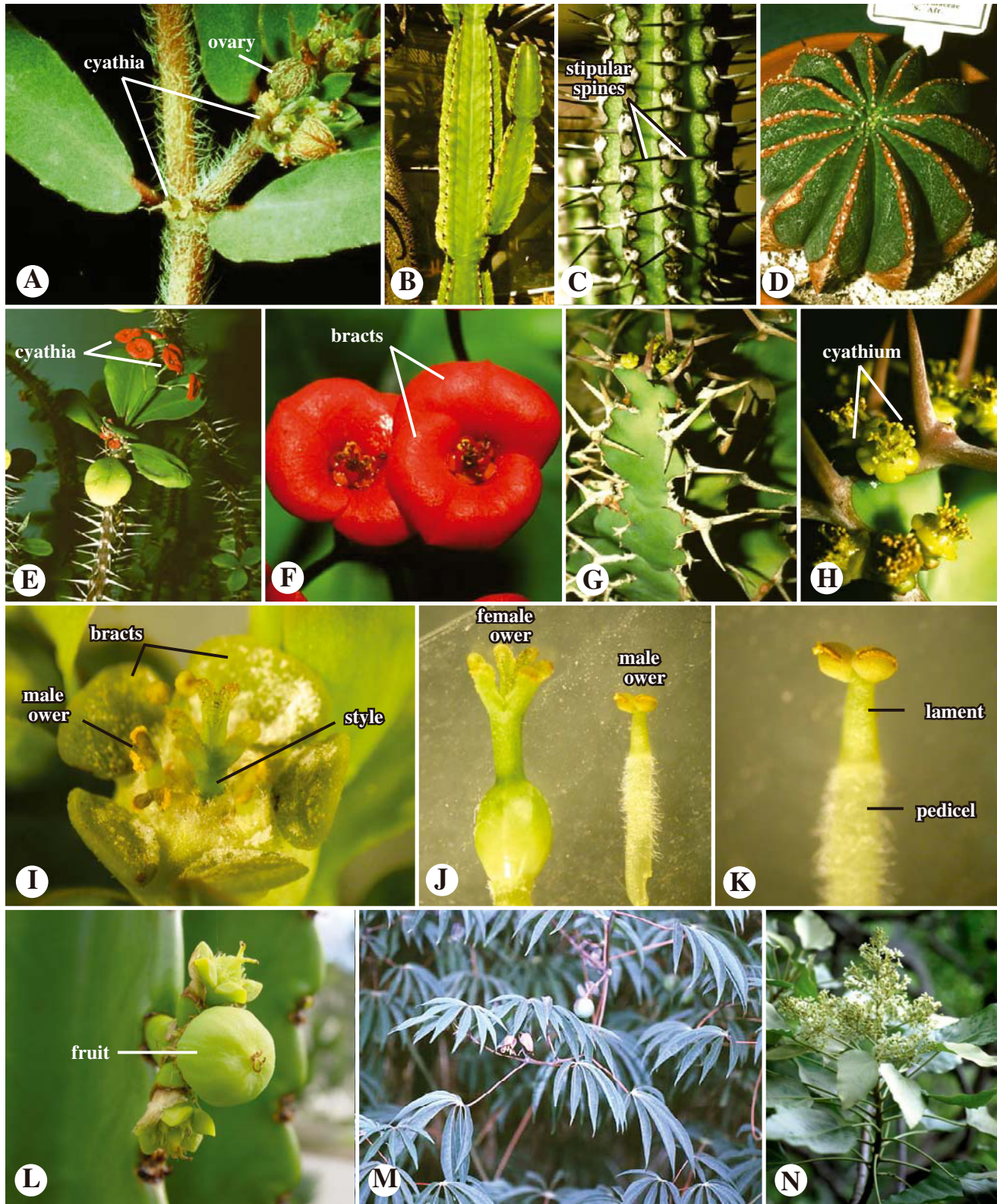


FIGURE 8.32 MALPIGHIALES. Euphorbiaceae, s.l. **A.** *Chamaesyce maculata*, close-up of leaves and cyathia. **B.** *Euphorbia ingens*, a stem-succulent plant. **C.** *Euphorbia bairoensis*, showing paired stipular spines. **D.** *Euphorbia obesa*. **E.** *Euphorbia milii*, crown-of-thorns. **E.** Aerial shoot of plant. **F.** Close-up of cyathia, having red bracts. **G,H.** *Euphorbia grandicornis*. **G.** Whole plant, showing enlarged stipular spines. **H.** Cyathia. **I–K.** *Euphorbia* sp. **I.** Cyathium, showing bracts and male and female owers. **J.** Female and male ower, removed. **K.** Close-up of male ower, showing junction between lamenet and pedicel. **L.** *Euphorbia ingens*, simple dichasium of cyathia, the central one in fruit. **M.** *Manihot esculenta*, cassava. **N.** *Aleurites moluccana*, candle-nut tree.

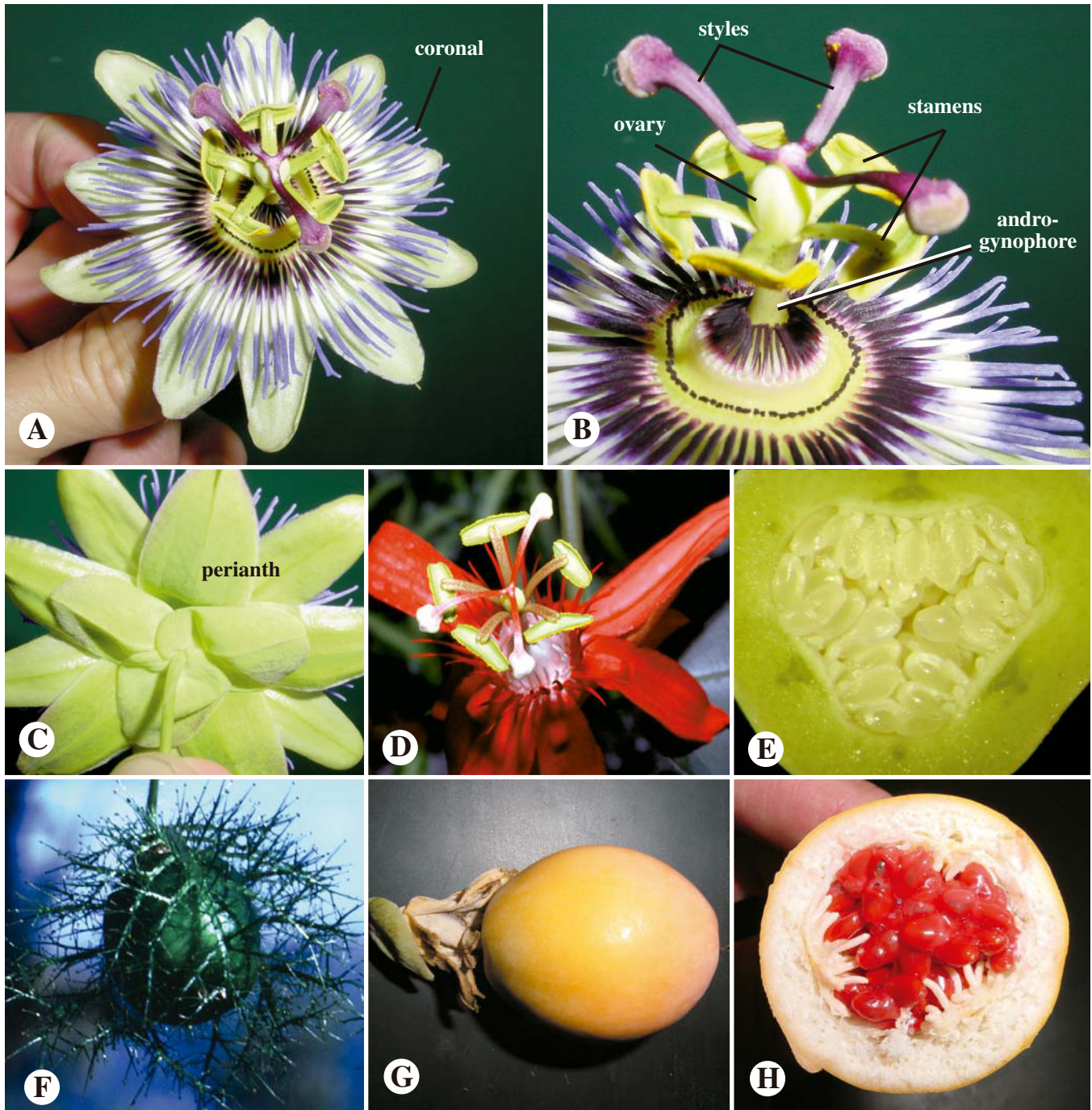


FIGURE 8.33 MALPIGHIALES. Passifloraceae. A–C. *Passiflora* sp. A. Flower. B. Close-up, showing androgynophore. Note five stamens and three-branched style. C. Perianth, back view, showing three bracts, five sepals, and five petals. D. *Passiflora* sp., flower. E. *Passiflora* sp., ovary cross-section, showing parietal placentation with three carpels. F. *Passiflora foetida*, fruit. G, H. *Passiflora edulis*, fruit, a berry.

Passifloraceae — Passion Flower family (Latin for passion flower, after events of the Christian Passion, signified in floral parts). 17 genera / 575 species. (Figure 8.33)

The Passifloraceae consist of lianas, shrubs, or trees. The **stems** have axillary tendrils in lianous species. The **leaves** are

simple, rarely palmately compound, often palmately lobed, spirally arranged, stipulate or exstipulate, the petioles often with extrafloral nectaries. The **inflorescence** is a cyme or a solitary flower. The **flowers** are bisexual or unisexual, bracteate, actinomorphic, usually perigynous, rarely hypogynous.

The **calyx** is aposepalous or basally synsepalous, with 5 [3-8] sepals or calyx lobes. The **corolla** is apopetalous or basally sympetalous, with 5 [3-8] petals or corolla lobes, and a corona of 1 or more whorls of filamentous or scalelike structures between the perianth and androecium. The **stamens** are 5 [4-∞], alternipetalous, whorled, uniseriate, distinct or connate, free or adnate to an androgynophore. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 [2-5] carpels, and 1 locule, usually arising from a prominent androgynophore, rarely from a gynophore or sessile. The **styles** and **stigma** are as many as carpels and usually basally connate. **Placentation** is parietal; **ovules** are anatropous, bitegmic, and numerous per carpel. The **fruit** is a berry or capsule. The **seeds** are endospermous.

The Passifloraceae have a worldwide distribution in tropical and subtropical regions. Economic importance includes use of many species as cultivated ornamentals, and some, e.g., *Passiflora edulis* (passion fruit) as edible fruits.

The Passifloraceae are distinctive in being lianas, shrubs, or trees with actinomorphic, usually bisexual flowers, having **one or more whorls of coronal appendages** between perianth and androecium, an **androgynophore** usually present, and typically **5 stamens and 3 carpels**, with **parietal placentation**, the fruit a **berry or capsule**.

K 5 [3-8] or (5) [(3-8)] **C** 5 [3-8] or (5) [(3-8)] **A** 5 [4-∞] or (5) [(4-∞)] **G** (3) [(2-5)], superior, hypanthium present in most species.

Violaceae Violet family (Latin for fragrant plants). 20 genera / 800 species. (Figure 8.34)

The Violaceae consist of herbs, shrubs, trees, or lianas. The **leaves** are simple, undivided to divided, usually spiral, and stipulate. The **inflorescence** is of solitary, axillary flowers or in heads, panicles, or racemes. The **flowers** are usually bisexual (cleistogamous in some, e.g., *Viola* spp.), actinomorphic or zygomorphic. The **calyx** is aposepalous with 5 imbricate, often persistent sepals. The **corolla** is apopetalous with 5 imbricate or convolute petals, the anterior petal in zygomorphic flowers often with a nectar-storing spur. The **stamens** are 5 [3], apotemonous or monadelphous, usually connivent around the ovary. The **gynoecium** is syncarpous, with a superior ovary, 3 [2-5] carpels, and 1 locule. The **style** is solitary. **Placentation** is parietal; **ovules** are anatropous, bitegmic, 1 ∞ per carpel. **Nectaries** are present on the dorsal side of stamens. The **fruit** is a berry or loculicidal capsule, rarely a nut. The **seeds** are endospermous and often with an aril or caruncle.

The Violaceae have a mostly worldwide distribution. Economic importance includes some plants of medicinal value, cultivated ornamentals, e.g., *Viola xwittrockiana* (pansies), and use for extraction of volatile oils.

The Violaceae are distinctive in being **herbs, shrubs, trees, or lianas** with simple, undivided or divided leaves, actinomorphic or zygomorphic flowers with a 5-merous perianth, usually **5, connivent stamens**, and a superior, usually **3-carpellate ovary** with **parietal placentation**, the seeds often with a **caruncle or aril**.

K 5 **C** 5 **A** 5 [3] or (5) [(3)] **G** (3) [(2-5)], superior.

OXALIDALES

The Oxalidales, sensu APG II, 2003, contain six families (Table 8.2), only one of which is described here. See Matthews and Endress (2002) and Kubitzki (2004) for recent information about this order.

Oxalidaceae — Oxalis family (Greek for sour, from the accumulation of oxalic acid in the tissues). 6 genera / 775 species. (Figure 8.35)

The Oxalidaceae consist of herbs, shrubs, or small trees. The **stems** are bulbs or tubers in some herbaceous taxa. The **leaves** are pinnate, palmate, often trifoliolate, rarely unifoliolate or of phyllodes, spirally arranged, usually exstipulate, the leaflets often folded at night. The **inflorescence** is a cyme. The **flowers** are bisexual, actinomorphic, and hypogynous. The **calyx** is aposepalous with 5 imbricate sepals. The **corolla** is apopetalous or basally sympetalous with 5 [0], convolute to imbricate petals. The **stamens** are biseriata, 5+5, whorled, the outer fertile stamens shorter, filaments basally connate, with outer whorl of staminodes sometimes present. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 5 [3] carpels, and 5 [3] locules. The **styles** are 5 [3,1]. **Placentation** is axile; **ovules** are anatropous or hemitropous, bitegmic, 2 ∞ [1] per carpel. **Nectaries** are often present at base of outer stamens. The **fruit** is a loculicidal capsule or berry. The **seeds** are endospermous and often with a basal aril.

The Oxalidaceae are mostly worldwide in distribution. Economic importance includes fruit trees, e.g., *Averrhoa carambola* (star-fruit), tuber plants, e.g., *Oxalis tuberosa*, and ornamental cultivars, e.g., *Oxalis* spp.

The Oxalidaceae are distinctive in being herbs, shrubs, or small trees with usually **pinnate or palmate (often trifoliolate)** leaves (**leaflets often folding** at night), bisexual, actinomorphic pentamerous flowers, **stamens usually biseriata**, with **outer, basal nectaries**, the fruit a loculicidal capsule or berry.

K 5 **C** 5 or (5) [0] **A** 5+5 [+5 staminodes] **G** (5) [(3)], superior.



FIGURE 8.34 MALPIGHIALES. Violaceae. **A.** *Hybanthus calycinus*, flower. Note large anterior petal. **B.** *Hybanthus concolor*, with reduced flowers. **C.** *Viola pedata*. **D.** *Viola purpurea*, a perennial herb. **E,F.** *Viola papilionacea*. **E.** Flower, face view. **F.** Flower longitudinal section. Note connivent stamens and nectary inside corolla spur. **G–K.** *Viola Xwittrockiana*, pansy. **G.** Flower, face view. **H–J.** Flower, longitudinal sections. **H.** Corolla spur with recessed nectaries. **I.** Connivent stamens appressed to pistil. **J.** Ovary, style, and stigma. **K.** Ovary cross-section, showing three carpels with parietal placentation, placental regions at arrows.

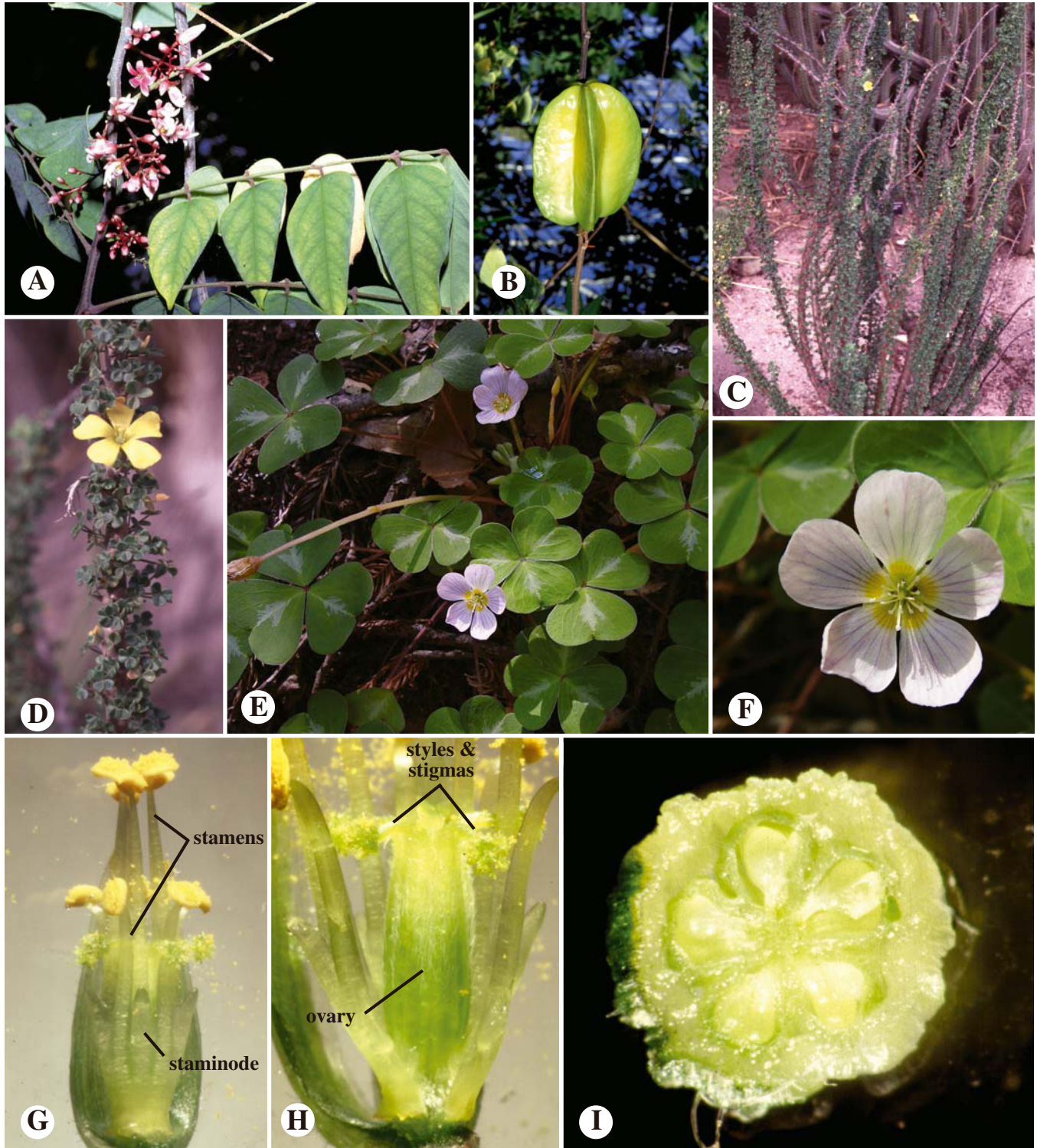


FIGURE 8.35 OXALIDALES. Oxalidaceae. **A,B.** *Averrhoa carambola*. **A.** Plant in flower. **B.** Winged fruit. **C,D.** *Oxalis arborescens*, a shrub. **E,F.** *Oxalis oregana*, redwood sorrel, a perennial herb. **G–I.** *Oxalis pes-caprae*. **G.** Flower, dissected, showing biseriate androecium, plus an outer whorl of staminodes. **H.** Central ovary, with apical, divergent styles and stigmas. **I.** Ovary cross-section, showing five carpels with axile placentation.

ROSALES

The Rosales, sensu APG II, 2003, contain nine families (Table 8.2), three of which are described here. Of these, the large family Rosaceae is of particular economic importance. Notable among the other families are the **Cannabaceae** (containing the euphoric and fiber plant *Cannabis sativa*, marijuana/hemp, and the beer-flavoring agent *Humulus lupulus*, hops), **Ulmaceae** (the elm family, source of important cultivars), and **Urticaceae** (stinging nettles and relatives). See Kubitzki (2004) for recent information on the order.

Moraceae Mulberry family (Latin name for mulberry). ca. 40 genera / 1100 species (Figure 8.36)

The Moraceae consist of monoecious or dioecious trees, shrubs, lianas, and herbs, often with laticifers bearing a milky latex. The **roots** are prop or buttress in some taxa. The **leaves** are simple [rarely compound], spiral or opposite, stipulate. The **inflorescence** is axillary and variable in morphology, consisting of a spike (catkinlike in *Morus*), raceme, head (in some taxa with flowers borne upon the surface of an invaginated compound receptacle), or in *Ficus* an enclosed hypanthodium. The **flowers** are unisexual, small, actinomorphic, hypogynous or epigynous. The **perianth** is uniseriate [rarely biseriate], 0–10, the perianth parts (often termed a calyx) connate, at least basally. The **stamens** are 1–6, opposite and usually as many as the perianth parts; anthers are dithecal or (in *Ficus* spp.) monothechal. The **gynoecium** is syncarpous, with a superior or inferior ovary, 2–3 carpels, and 1–2 locules. The **styles** are typically 2. **Placentation** is apical (to subapical); **ovules** are solitary, anatropous to campylotropous, and bitegmic. The **fruit** is a multiple of achenes, each unit achene often surrounded by the accrescent, fleshy perianth (thus resembling a drupe) or borne on a fleshy compound receptacle, in *Ficus* borne on the inner surface of an enclosed syconium. The **seeds** are 1 per unit fruit, albuminous or exalbuminous. Plants are wind or insect pollinated, in *Ficus* spp. having an intricate pollination mechanism with wasps.

The Moraceae have a worldwide distribution, from tropical to temperate regions. Economic importance includes fruit trees, such as *Artocarpus altilis* (breadfruit), *Ficus carica* (edible fig), and *Morus* spp. (mulberry); paper, rubber, and timber trees; and some cultivated ornamentals, especially *Ficus* spp., figs; the leaves of *Morus alba* are the food source of silkworm moth larvae.

The Moraceae are distinctive in being *monoecious or dioecious* trees, shrubs, lianas, or herbs with a *milky latex*, *stipulate*, simple leaves, and *unisexual flowers*, the female with a usually *2-carpellate (2 styled) pistil* and a *single*,

apical to subapical ovule, the fruit a *multiple of achenes*, in some taxa with an enlarged compound receptacle or syconium. **P** (0–10) **A** 1–6 **G** (2) [(3)], superior or inferior.

Rhamnaceae — Buckthorn family (Greek name for buckthorn or other thorny shrubs). 49 genera / 900 species. (Figure 8.37)

The Rhamnaceae consist of trees, shrubs, lianas, or rarely herbs. The **roots** of some taxa are associated with nitrogen-fixing Actinomycetes bacteria. The **stems** are sometimes modified as thorns, tendrils, or hooks. The **leaves** are simple, sometimes rudimentary, pinnately or palmately veined, spiral or opposite, stipulate or exstipulate, with stipular spines present in some taxa. The **inflorescence** is a cyme, thyrse, fascicle, or rarely a solitary flower. The **flowers** are unisexual or bisexual, actinomorphic, perigynous to epiperigynous. The **perianth** typically has a hypanthium, sometimes adnate to staminal disk. The **calyx** is aposepalous with 4–5 sepals. The **corolla** is apopetalous with 4–5 petals [absent in some taxa], these often clawed, concave, and cucullate (hooded). The **stamens** are 4–5, whorled, alternisepalous, and apostemonous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior to inferior ovary and 2–3 [5] carpels and locules. **Placentation** is apical-axile; **ovules** are anatropous, bitegmic, 1–2 [2] per carpel. **Nectaries** consist of a staminal disk, often fused to the hypanthium and/or (inferior) ovary. The **fruit** is a drupe with 1 many endocarps, a circumscissile capsule, or a schizocarp of mericarps. The **seeds** are exalbuminous.

The Rhamnaceae have a mostly worldwide distribution, especially in tropics. Economic importance includes edible fruits (e.g., *Ziziphus jujuba*, jujube, or *Z. lotus*, lotus fruit), ornamental cultivars, and dye, medicinal, soap, timber and varnish plants; *Ziziphus spina-christi* is purported to be the true Christ's crown of thorns. See Richardson et al. (2000) for a recent study of infrafamilial relationships in the Rhamnaceae.

The Rhamnaceae are distinctive in being trees, shrubs, lianas, or rarely herbs with simple, spiral or opposite leaves, unisexual or bisexual, *perigynous to epiperigynous flowers*, the perianth/androecium *4–5-merous*, petals sometimes absent, *stamens alternisepalous*, a *nectariferous disk usually adnate to hypanthium*, the fruit a *drupe, circumscissile capsule, or schizocarp*.

K 4–5 **C** 4–5 [0] **A** 4–5 **G** (2–3) [(-5)], superior to inferior, hypanthium usually present.

Rosaceae Rose family (Latin for various roses). 95 genera / 2800 species. (Figures 8.38, 8.39)

The Rosaceae consist of trees, shrubs, or herbs. The **leaves** are spiral (rarely opposite), simple or compound, undivided to divided, usually stipulate, the stipules often adnate to the

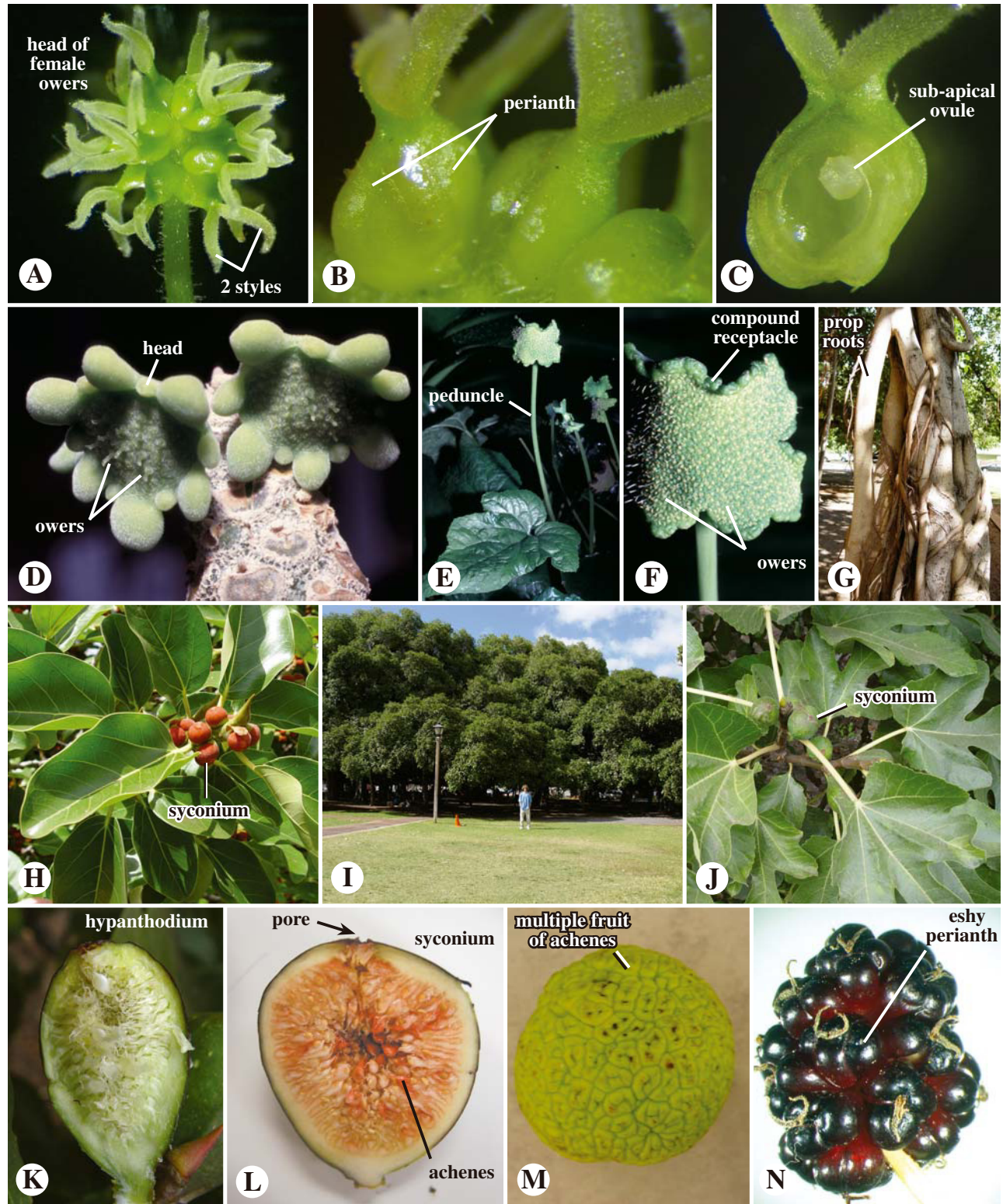


FIGURE 8.36 ROSALES. Moraceae. A–C. *Morus* sp., mulberry. A. Female inflorescence, a catkinlike spike of flowers. B. Flower close-up, showing outer perianth, which forms a fleshy layer in fruit. C. Ovary longitudinal section, showing subapical placentation. D. *Dorstenia gigas*, showing headlike inflorescences. E,F. *Dorstenia* sp., with headlike, compound receptacle bearing minute flowers. G–I. *Ficus benghalensis*. G. Prop roots. H. Shoot close-up, showing syconia. I. Habit, a tree, here covering more than an acre of ground. J–L. *Ficus carica*, edible fig. J. Shoot. K. Young syconium, longitudinal section. L. Mature syconium, longitudinal section. M. *Maclura pomifera*, osage-orange, showing globose, multiple fruit of drupelike achenes. N. *Morus* sp., fruit, a multiple of achenes, each surrounded by a fleshy perianth.

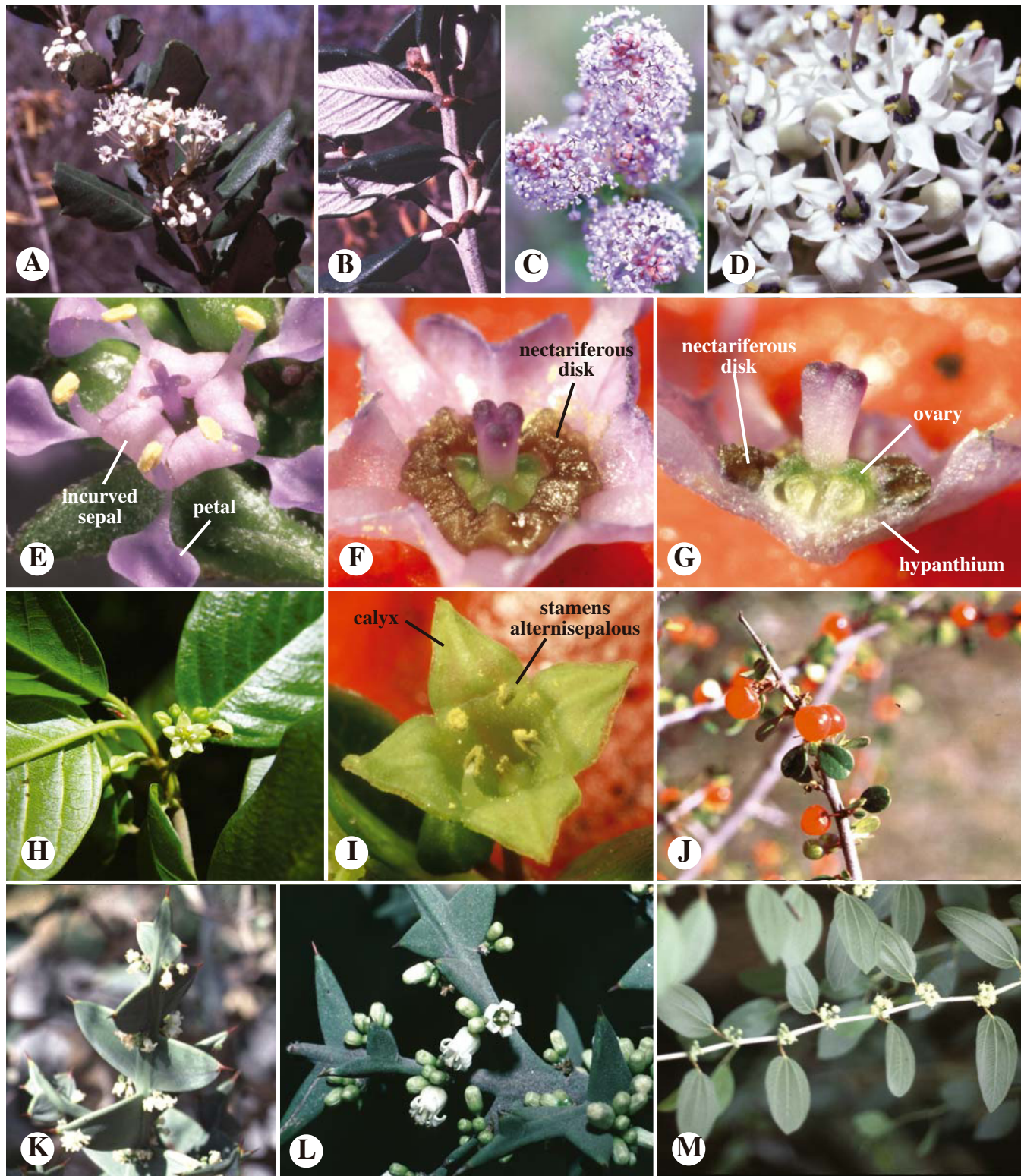


FIGURE 8.37 ROSALES. Rhamnaceae. **A,B.** *Ceanothus crassifolius*. **C.** *Ceanothus tomentosus*, in orescence. **D.** *Ceanothus verrucosus*, flower close-up. **E.** *Ceanothus griseus*, flower close-up. Note antipetalous stamens. **F,G.** *Ceanothus tomentosus*. Flowers, showing nectariferous disk, ovary, and hypanthium. **H.** *Rhamnus californica*. **I,J.** *Rhamnus crocea*. **I.** Flower. Note absence of petals and alternisepalous stamens. **J.** Shoot bearing fruits. **K.** *Colletia cruciata*, with thick, spinose leaves. **L.** *Colletia paradoxa*. **M.** *Zizyphus* sp.



FIGURE 8.38 ROSALES. Rosaceae. **A.** *Rosa* sp., one of many forms of cultivated roses. **B.** *The Rose Family*, by Robert Frost. (Frost, R., and J. J. Lankes. 1928. West-Running Brook. H. Holt, New York.)

The Rose Family

The rose is a rose,
And was always a rose.
But the theory now goes
That the apple s a rose,
And the pear is, and so s
The plum, I suppose.
The dear only knows
What will next prove a rose.
You, of course, are a rose
But were always a rose.

Robert Frost (1874-1963)

petiole base. The **inflorescence** is variable. The **flowers** are bisexual (usually), actinomorphic, perigynous or epiperigynous; the receptacle is sometimes expanded or sunken. The **perianth** is biseriate and dichlamydeous, usually pentamerous, imbricate, a hypanthium present. The **calyx** is aposepalous with 5 [3 10] sepals. The **corolla** is apopetalous with 5 [0, 3 10] petals. The **stamens** are 20∞ [1,5], whorled, arising centripetally, usu. apostemonous. **Anthers** are longitudinal or rarely poricidal in dehiscence and dithecal. The **gynoecium** is syncarpous or apocarpous, with a superior or inferior ovary, 1∞ carpels, and 1∞ locules. The **style(s)** are terminal or lateral. **Placentation** is axile, basal, or marginal; **ovules** are 1∞ . **Nectaries** are often present on the hypanthium. The **fruit** is a drupe, pome, hip, follicetum, achenecectum, or capsule. The **seeds** are usually without endosperm.

The Rosaceae is traditionally classified into four subfamilies (some of which are likely paraphyletic; see Potter et al. 2002): Spiraeoideae, with an apocarpous gynoecium forming a follicetum; Rosoideae, with an apocarpous gynoecium forming an achenecectum or drupecectum, the receptacle varying from expanded and fleshy (e.g., *Fragaria*) to sunken (e.g., the hips of *Rosa*); Prunoideae, with a single, superior-ovary pistil bearing one ovule, the fruit a drupe; and Maloideae, with an inferior ovary, forming a pome. Members of the family have mostly worldwide distributions, but are more concentrated in north temperate regions. The family is very economically important as the source of many cultivated fruits, including *Fragaria* (strawberry), *Malus* (apples), *Prunus* (almond, apricot, cherry, peach, plum), *Pyrus* (pear), and *Rubus* (blackberry, raspberry), as well as essential oils (e.g., *Rosa*), and numerous ornamental cultivars, such as

Cotoneaster, *Photinia*, *Prunus* (cherries), *Pyracantha*, *Rosa* (roses), and *Spiraea*. See Potter et al. (2002) for a recent study of the Rosaceae.

The Rosaceae are distinctive in having usually **stipulate** leaves (often adnate to petiole) and an actinomorphic, generally **pentamerous** flower with **hypanthium present**, variable in gynoeceal fusion, ovary position, and fruit type.

K 5[3-10] **C** 5[0,3-10] **A** $20-\infty$ [1,5] **G** $1(-\infty)$, superior or inferior, hypanthium present.

EUROSIDS II

This second major subgroup of the Rosids contains 3 4 orders (Table 8.2). Some of the orders are quite large, in terms of both number of families and number of species. Among the Eurosids II group are taxa of great agricultural importance, such as members of the Anacardiaceae (cashew family), Brassicaceae (mustard family), and Rutaceae (citrus family). Of course, many other groups are quite important more locally.

BRASSICALES

The Brassicales are composed of ca. 15 17 families. Only the largest and most economically important family, the Brassicaceae, is described here. The order is generally united in having **glucosinolate** secondary compounds (Figure 8.40). Glucosinolates function to deter herbivory and parasitism and also serve as flavoring agents in the commercially important

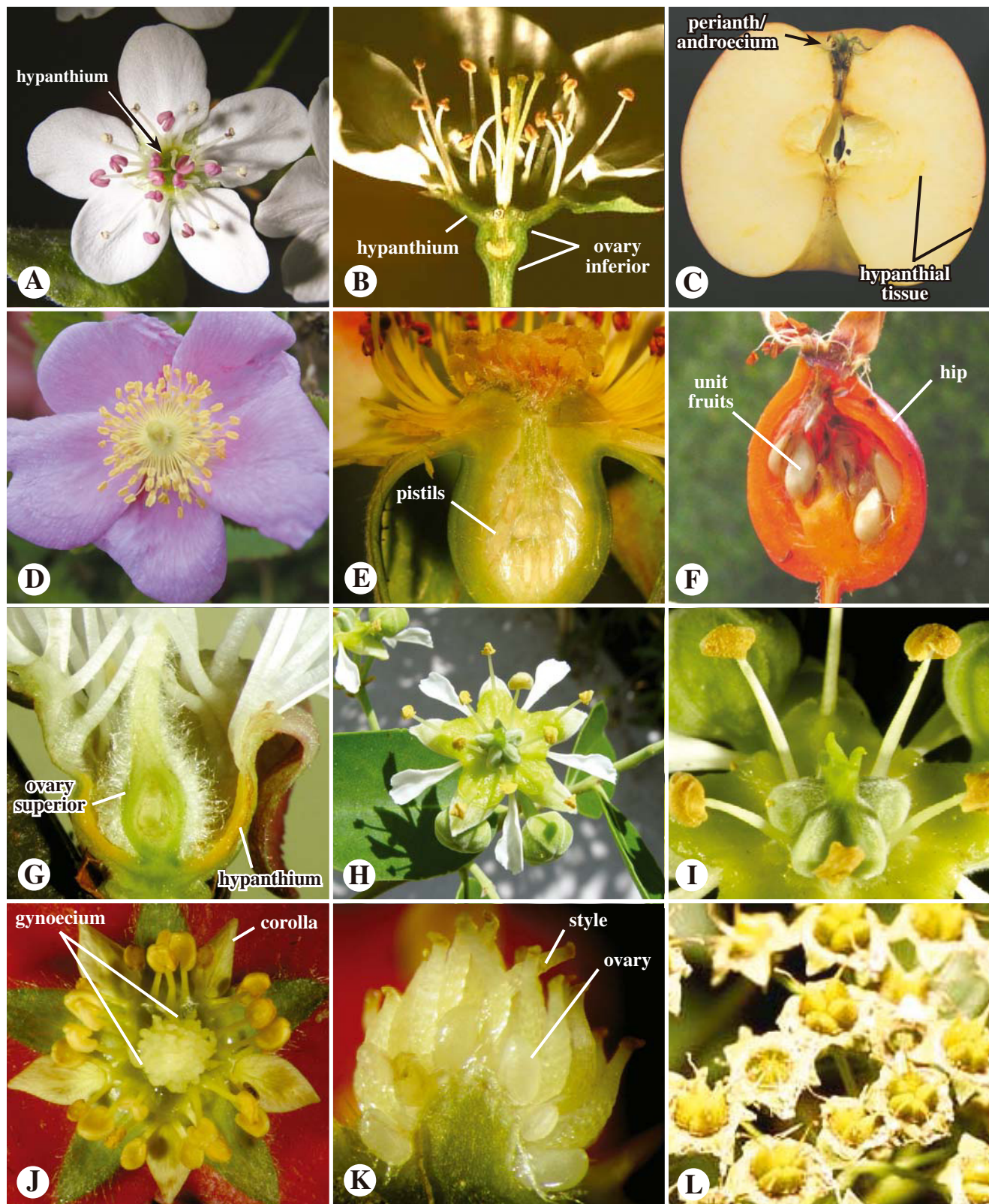


FIGURE 8.39 ROSALES. Rosaceae. A–C. Maloideae. A. *Pyrus kawakamii*, flower, face view. B,C. *Malus domestica*, apple. B. Flower longitudinal section, showing inferior ovary, characteristic of the subfamily. C. Apple fruit, a pome, comprised mostly of hypanthial tissue. D,F. Rosoidae, *Rosa californica*. D. Flower. E. Sunken receptacle of flower. F. Fruit, a hip. G. Prunoideae, *Prunus* sp., cherry, with a superior ovary. H,I. *Quillaja saponaria*. J,K. *Potentilla glandulosa*, having numerous pistils of an apocarpous gynoecium. L. Spiraeoideae. *Spiraea japonica* immature fruits, each an aggregate of follicles (follicetum).

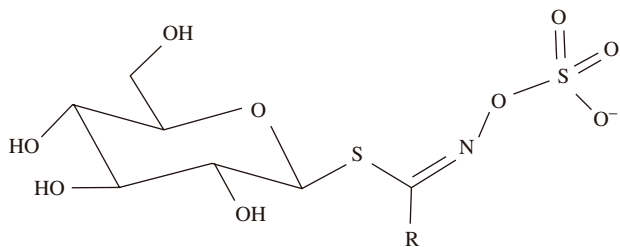


FIGURE 8.40 General structure of glucosinolates, a common feature of the Brassicales.

members of the Brassicaceae, such as broccoli, cauliflower, and mustard. Some well-known families of the order not described here include the **Capparaceae** (including capers, *Capparis spinosa*), **Caricaceae** (including the fruit tree *Carica papaya*, papaya) and **Tropaeolaceae** (including *Tropaeolum majus*, the ornamental nasturtium). See Rodman et al. (1997, 1998) and Kubitzki and Bayer (2002) for more information on the order.

Brassicaceae (Cruciferae) — Mustard family (name used by Pliny for cabbagelike plants). 365 genera / 3250 species. (Figures 8.41, 8.42)

The Brassicaceae consist of usually hermaphroditic herbs, rarely shrubs (pachycaulis in some). The **leaves** are simple [rarely compound], often lobed to divided, spiral [rarely opposite], exstipulate. The **inflorescence** is usually a raceme, rarely of solitary, axillary flowers. The **flowers** are bisexual, rarely unisexual, usually actinomorphic, pedicellate, ebracteate, hypogynous; the receptacle is rarely elongate into a gynophore. The **perianth** is dichlamydeous, cruciate. The **calyx** is aposepalous [rarely synsepalous] with 2+2, decussate outer sepals, often basally gibbous. The **corolla** is apopetalous, rarely basally connate, with 4 [rarely absent] petals, which are often clawed. The **stamens** are apostemonous, biseriate, 2+4 tetradynamous [rarely 2 or 4 or up to 16], the outer 2 shorter, antisealous, the inner 4 longer, of two pairs, each pair (from a single primordium) flanking adjacent petals. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 2 locules. The **style** is 1 or absent. **Placentation** is axile-parietal, each carpel with two rows of ovules, the placentae at junction of septum and ovary wall; **ovules** are anatropous or campylotropous, 1 ∞ per carpel. **Nectaries** are discrete or ringlike around stamens or pistil. The **fruit** is a silique or silicle, with a persistent septum termed a replum, rarely transversely dehiscent. The **seeds** are usually exalbuminous.

The Brassicaceae, sensu APG II (2003), include the traditional families Capparaceae and the Cleomaceae (these two

in the past treated as subfamilies of the Capparaceae). Each of these three groups appear to be monophyletic. Thus, an alternative, acceptable classification is to treat them as three separate families, as done here: Brassicaceae, Capparaceae, and Cleomaceae (Table 8.2). The Capparaceae (Figure 8.42A,B) differ from the Brassicaceae largely in having a woody habit, an elongate gynophore or androgynophore, a generally greater number of stamens, a unilocular ovary with parietal placentation, and an indehiscent fruit type lacking a replum. The Cleomaceae (Figure 8.42C E) resemble the Capparaceae but are largely herbaceous and have a dehiscent fruit with a replum (but lacking a complete partition, thus the ovary unilocular). See Hall et al. (2002) for a recent phylogenetic analysis of the group.

The Brassicaceae have a worldwide distribution. Economic importance includes numerous vegetable plants (notably the crucifers or mustard plants), including broccoli, brussels sprouts, cauliflower, cabbage, collards, kale (all cultivars of *Brassica oleracea*), rutabaga and canola oil (*B. napus*), mustard (*B. nigra*), turnip (*B. rapa*), and many more, plus numerous cultivated ornamentals, dye plants (*Isatis tinctoria*, woad), and some noxious weeds; *Arabidopsis thaliana* is noted as a model for detailed molecular studies.

The Brassicaceae as treated here are distinctive in being herbs, rarely shrubs, with **glucosinolates** (mustard oil glucosides), the **perianth cruciate** (petals usually clawed), the androecium with usually **2+4, tetradynamous stamens**, the gynoecium with a superior, **2-carpellate/loculate ovary**, with **axile-parietal placentation** and a usually **2-valved, dehiscent fruit with a replum (silique or silicle)**.

K 2+2 **C** 4 **A** 2+4 [2,4-16] **G** (2), superior.

MALVALES

The Malvales, sensu APG II (2003), include nine families (Table 8.2). Of these nine families, only the Malvaceae (s.l.) will be covered here. More well-known among the others are the **Bixaceae** (containing *Bixa orellana*, anatto, commonly used as a natural food coloring), **Cistaceae** (the rock-rose family), **Dipterocarpaceae** (the dipterocarps of s.e. Asia, source of important hardwood timber trees and gum/resin plants), and **Thymelaeaceae**. The order as a whole may have chemical and anatomical apomorphies, including the presence of lysigenous mucilage canals in most members. See Kubitzki and Chase (2002) for more information.

Malvaceae [including Bombaceae, Sterculiaceae, and Tiliaceae] — Mallow family (name used by Pliny, meaning soft). ca. 250 genera / 4230 species. (Figures 8.43 8.46)

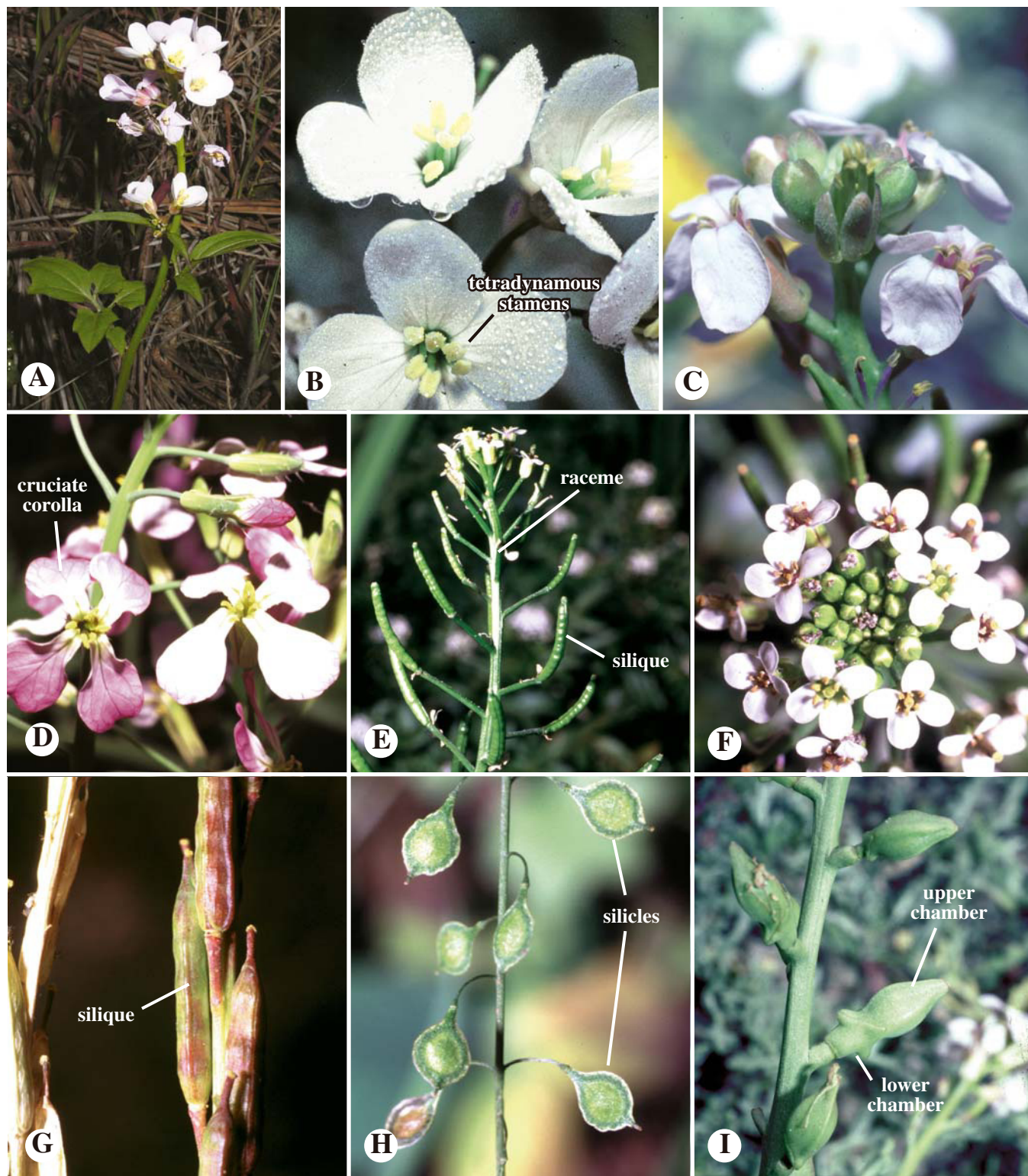


FIGURE 8.41 BRASSICALES. Brassicaceae. **A,B.** *Cardamine californica*, showing raceme and cruciate, tetradynamous flowers. **C.** *Cakile maritima*, flowers. **D.** *Raphanus sativus*, radish, flowers. **E,F.** *Rorippa nasturtium-aquaticum*, flowers and fruits. **G.** Silique of *Brassica nigra*, mustard. **H.** Silicle of *Thysanocarpus laciniatus*. **I.** Transversely dehiscent capsule of *Cakile maritima*.

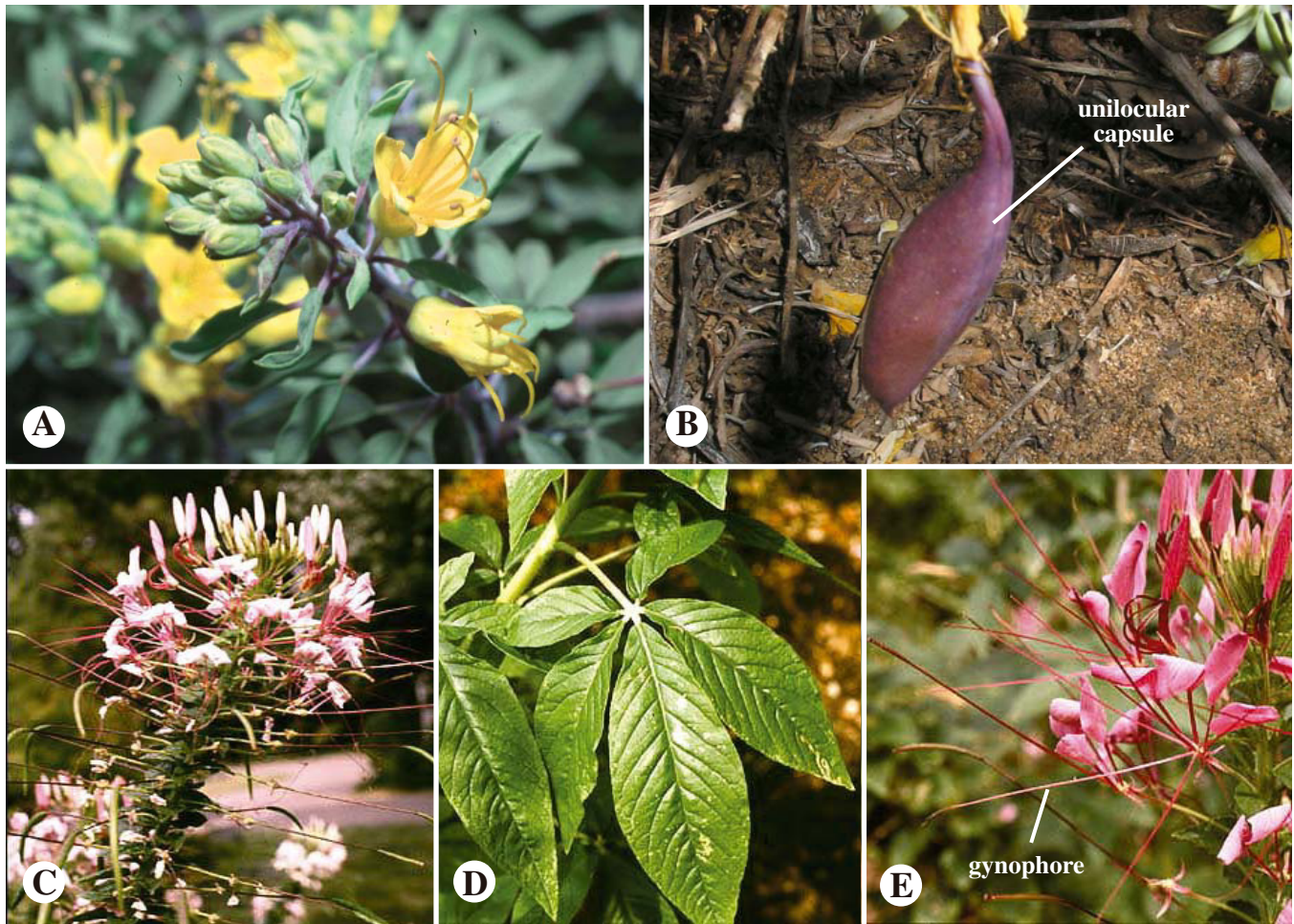


FIGURE 8.42 BRASSICALES. **A,B.** Capparaceae, *Isomeris arborea*. **A.** Flowers. **B.** Fruit. **C–E.** Cleomaceae, *Cleome bassleriana*. **C.** Inflorescence. **D.** Palmate leaves. **E.** Flower, showing elongate gynophores.

The Malvaceae, sensu APG II (2003), consist of usually hermaphroditic, rarely monoecious or polygamous trees, shrubs, or herbs, often with either stellate trichomes or peltate scales. The **leaves** are simple or palmately compound, sometimes lobed to divided, palmately or pinnately veined, usually spiral and stipulate, the stipules often caducous. The **inflorescence** is of solitary or paired flowers or cymelike, sometimes complex. The **flowers** are bisexual [rarely unisexual], mostly actinomorphic, an epicalyx typically present, hypogynous, rarely perigynous. The **perianth** is biseriate, the petals alternating with the sepals. The **calyx** is aposepalous or basally synsepalous with 5 [less often 3–4], valvate sepals. The **corolla** is apopetalous [sometimes adnate to the base of an androecium tube; rarely absent], when present of 5 [3–4], sometimes clawed, convolute, valvate, or imbricate petals. The **stamens** are 5 ∞ , the filaments usually connate, either as

a tube surrounding the ovary, or as 5–15 bundles of stamens or a tube bearing bundles. **Anthers** are longitudinal or poricidal in dehiscence. The **pollen** is spinulose or smooth. The **gynoecium** is syncarpous, rarely apocarpous or with carpels fused only apically, with a superior [rarely inferior] ovary, 2 ∞ carpels, and 2 ∞ [1] locules. The **style** is unlobed, lobed, or branched at the apex. **Placentation** is usually axile, rarely marginal; **ovules** are 2 ∞ [1] per carpel. **Nectaries** consist of glandular trichomes typically present at the adaxial base of the calyx. The **fruit** is a loculicidal, septicidal, or indehiscent capsule, a schizocarp of mericarps, or rarely a berry or samara. The **seeds** are exalbuminous or endospermous (oily or starchy).

The Malvaceae s.l. as treated here were formerly (and still commonly) divided into four families: Malvaceae s.s., Bombacaceae, the Bombax family, Sterculiaceae, the chocolate family, and Tiliaceae, the Linden family. Recent morphological

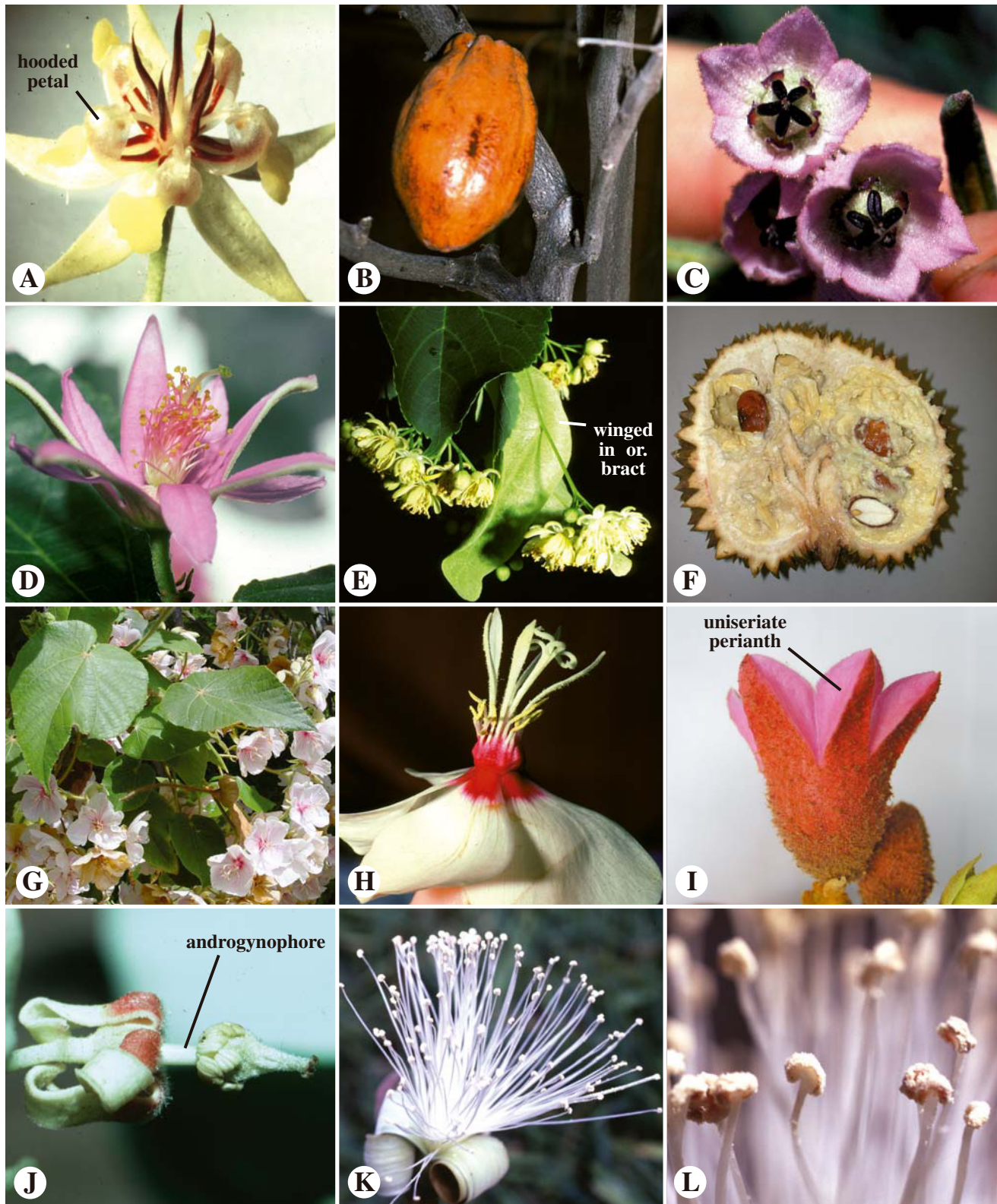


FIGURE 8.43 MALVALES. Malvaceae. **A,B.** *Theobroma cacao*, cacao (Byttnerioideae). **A.** Flower, showing cucullate (hooded) petals. **B.** Fruit, a berry, source of chocolate. **C.** *Guichenotia ledifolia* (Byttnerioideae). **D.** *Grewia occidentalis* (Grewioideae). **E.** *Tilia americana* (Tilioideae), showing prominent in orscence bract. **F.** *Durio zebethinus*, durian (Helicteroideae?), fruit in longitudinal section. **G,H.** *Dombeya* spp. (Dombeyoideae). **I.** *Brachychiton discolor* (Sterculioideae), with uniseriate perianth (corolla absent). **J.** *Heritiera littoralis* (Sterculioideae), showing androgynophore. **K,L.** *Pseudobombax ellipticum* (Bombacoideae). **K.** Flower with numerous stamens. **L.** Close-up of monothechal anthers. Subfamilies after Bayer et al. (1999).

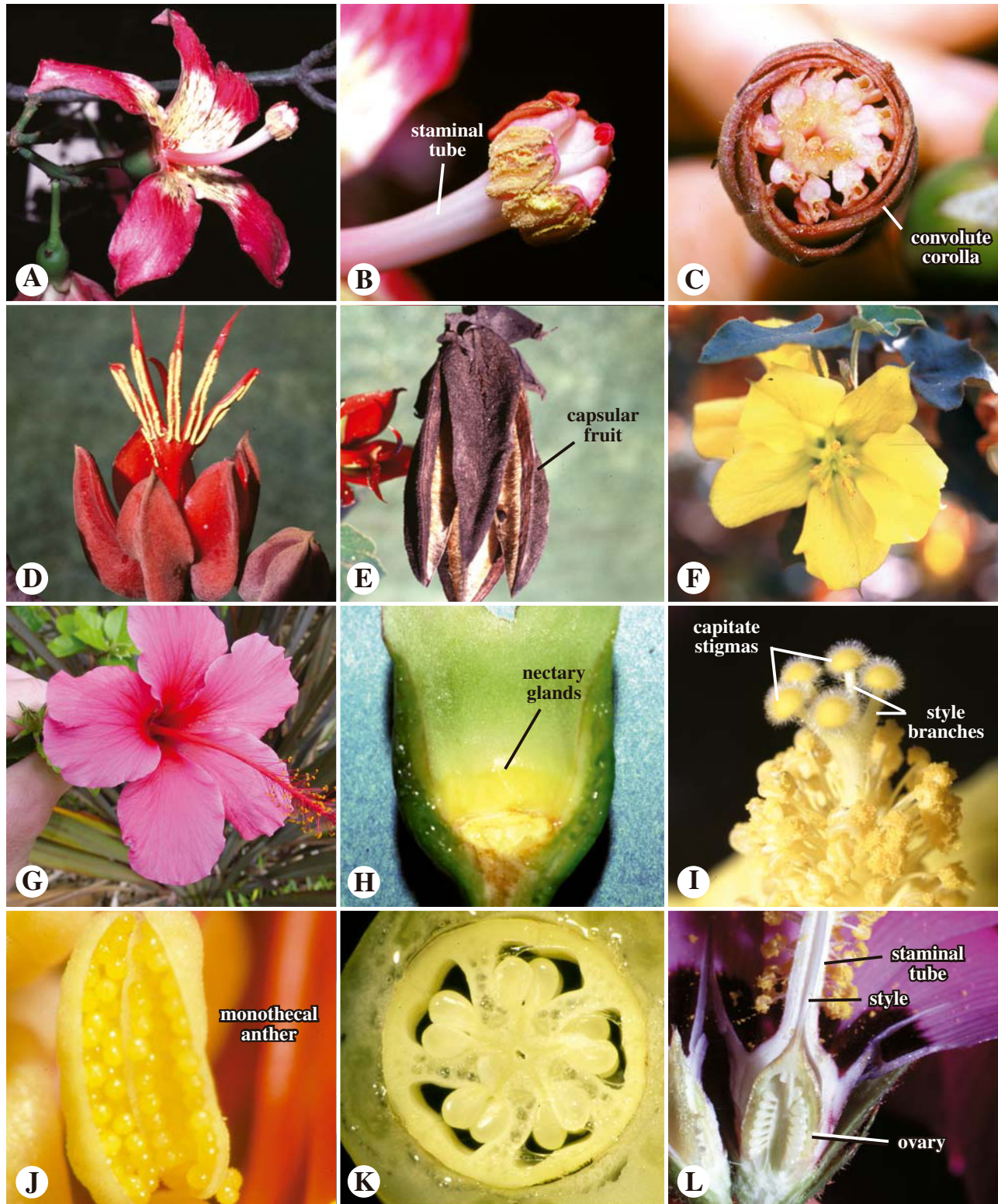


FIGURE 8.44 MALVALES. Malvaceae (continued). A–C. *Chorisia speciosa* (Bombacoideae). A. Whole flower, showing staminal tube. B. Close-up of staminal tube bearing monothechal anthers. C. Bud cross-section, showing convolute corolla. D,E. *Chiranthodendron pentadactylon* (Bombacoideae). D. Flower, abaxial view, showing five connate stamens. E. Fruit, a capsule. F. *Fremontodendron mexicanum* (Bombacoideae). G–K. *Hibiscus* sp. (Malvoideae). G. Whole flower, showing convolute corolla and staminal tube. H. Nectariferous adaxial surface of calyx. I. Close-up of style branches and anthers of surrounding staminal tube. J. Close-up of monothechal anther. K. Ovary cross-section, showing axile placentation. L. *Alyogyne huegelii* (Malvoideae), flower longitudinal section, showing corolla connate to androecium, staminal tube, and superior ovary. Subfamilies after Bayer et al. (1999).

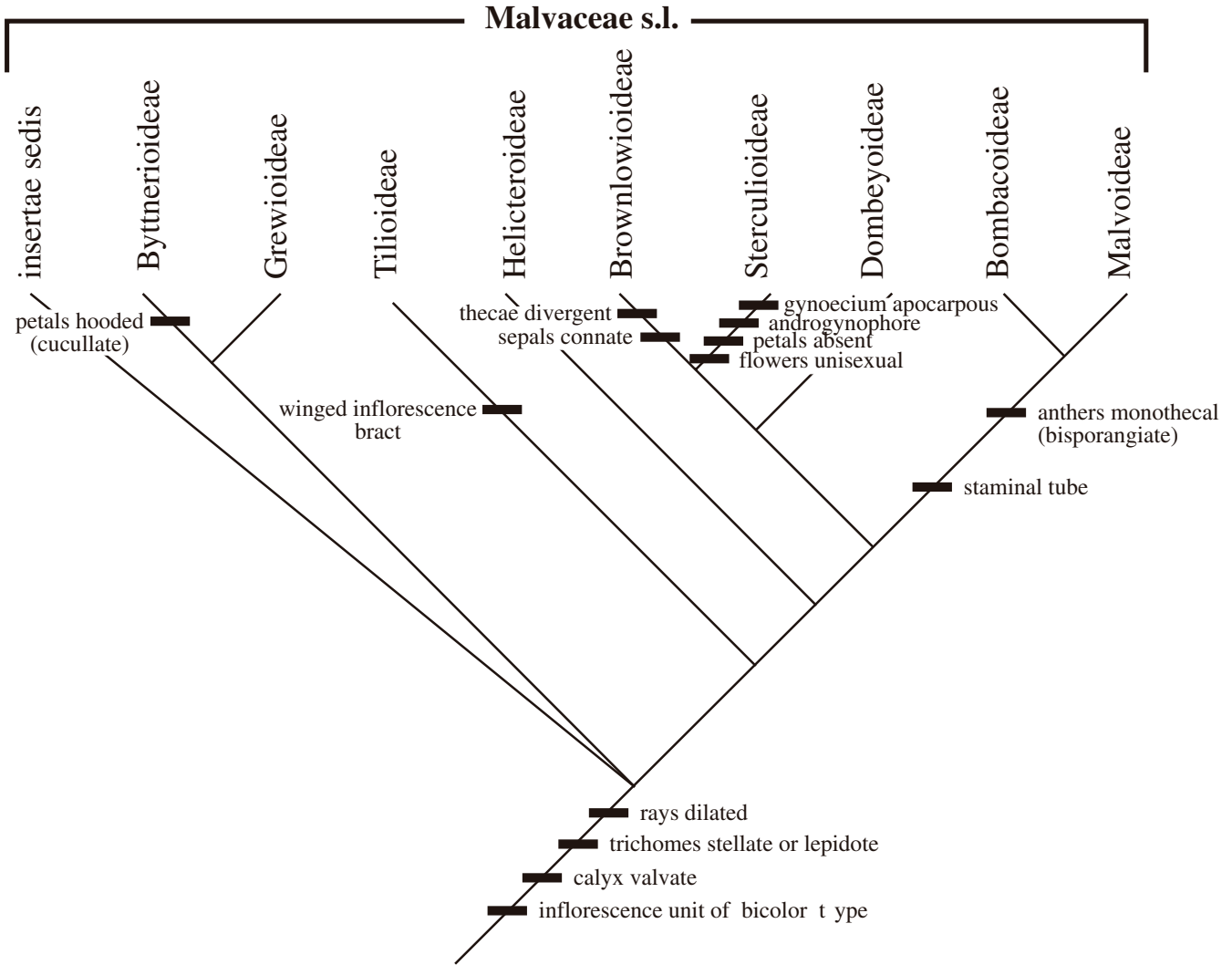


FIGURE 8.45 MALVALES. Cladogram of the Malvaceae s.l., after Bayer et al. (1999). Apomorphies listed are suggestive and may need further study for verification.

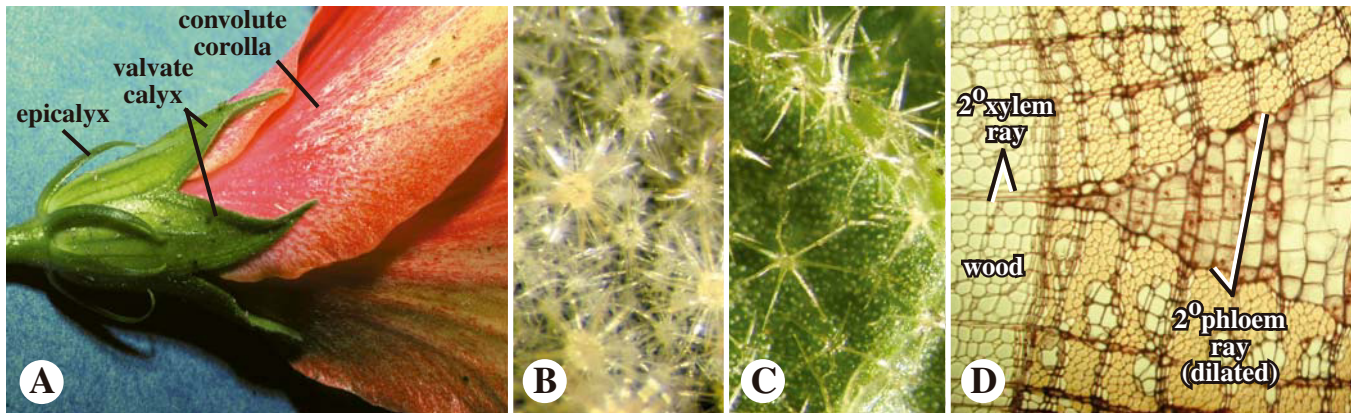


FIGURE 8.46 MALVALES. Possible apomorphies of the Malvaceae s.l. **A.** Valvate calyx (*Hibiscus*); also note epicalyx and convolute corolla. **B,C.** Stellate trichomes (*Alyogyne* and *Fremontodendron*, respectively). **D.** Dilated wood rays (*Tilia*).

and molecular analyses indicate that these groups are largely nonmonophyletic and best classified together. Bayer et al. (1999) tentatively recognized nine subfamilies, some putative apomorphies shown in Figure 8.45. The Malvaceae s.l. as a whole may be united by an inflorescence apomorphy, the occurrence of a bicolor unit (the term derived from *Theobroma bicolor*, where it was first observed), consisting of a modified, 3-bracted cyme, the trimerous epicalyx of family members possibly derived from these 3 bracts. Other apomorphies of the family may be a valvate calyx, stellate or lepidote trichomes, and dilated secondary tissue rays (Figures 8.45, 8.46). Members of the family are distributed worldwide, especially in tropical regions. Economic importance includes medicinal plants; several fiber plants, especially *Gossypium* spp. (cotton, the world's most important fiber plant) and *Ceiba pentandra* (kapok), in both of which the seed trichomes are utilized, and *Corchorus* spp. (jute), a bast fiber plant and source of burlap; food and flavoring plants, such as *Theobroma cacao* (cacao, the source of chocolate), *Cola nitida* (cola), *Abelmoschus* (okra), and *Durio zibethinus* (durian); wood, such as *Ochroma pyramidale* (balsa) and *Pachira aquatica*; and numerous ornamental cultivars, such as *Brachychiton*, *Chorisia* (floss-silk tree), *Dombeya*, *Fremontodendron*, *Hibiscus* (mallows), and *Tilia* (linden tree). Many others, such as *Adansonia digitata* (baobab, tropical Africa) are of great local economic or ecological importance. See Bayer (1999), Bayer et al. (1999), and references therein.

The Malvaceae are distinctive in being herbs, shrubs, or trees, often with *stellate trichomes*, typically with an *epicalyx*, the *calyx valvate*, the *corolla often convolute* [sometimes valvate or imbricate] the *stamens connate* as a tube or 5-∞ bundles, with *monotheical or ditheical anthers*, gynoecium syncarpous [rarely apocarpous], ovary superior [rarely inferior], ovules axile or marginal, the fruit a capsule, schizocarp of mericarps, berry, or samara.

K 3-5 or (3-5) **C** 3-5 [0] **A** 5-∞ **G** 2-∞ [1], superior [rarely inferior].

SAPINDALES

This order contains nine families, three of which are described here. Among the others, the **Burseraceae** is notable as the source of frankincense (*Boswellia* spp., esp. *B. sacra*) and myrrh (*Commiphora* spp., esp. *C. myrrha*). See Gadek et al. (1996) for a phylogenetic analysis of the order.

Anacardiaceae Cashew family (Gr. for heart-shaped, after swollen, red pedicel in cashew fruit). 70 genera / 875 species. (Figure 8.47)

The Anacardiaceae consist of trees, shrubs, lianas, or rarely perennial herbs, tissues of plant organs with resin ducts or laticifers, the resin allergenic in some taxa. The **leaves** are pinnate, trifoliolate, or simple, spiral, rarely opposite or whorled, exstipulate or stipules vestigial. The **inflorescence** is a terminal or axillary thyrses. The **flowers** are bisexual or unisexual, actinomorphic, usually hypogynous; the receptacle is swollen and fleshy in some taxa (e.g., *Anacardium*). The **perianth** is biseriate and dichlamydeous, parts valvate or imbricate. The **calyx** is usually basally synsepalous with usually 5 sepals or lobes. The **corolla** is apopetalous with usually 5 [0] petals. The **stamens** are 5-10 [1, ∞], apostemonous or rarely basally connate. The **gynoecium** is syncarpous [rarely apocarpous], with a superior [rarely inferior] ovary, 1-3, or 5 [rarely 12] carpels, and usually 1 [sometimes as many as carpels] locule. **Placentation** is apical/pendulous or basal; **ovules** are anatropous, bitegmic or unitegmic, 1 per carpel. **Nectaries** are present as a staminal, intrastaminal, or extrastaminal nectariferous disk. The **fruit** is a drupe, with the mesocarp usually resinous. The **seeds** have endosperm absent or scanty.

The Anacardiaceae have a broad distribution in tropical to temperate regions. Economic importance includes ornamental cultivars (e.g., *Schinus* spp.), fruit and seed trees, such as *Pistacia vera* (pistachio), *Rhus* spp. (sumacs), *Anacardium occidentale* (cashew), and *Mangifera indica* (mango), plus several dye, timber, and lacquer trees. *Toxicodendron* spp. (poison oak, poison ivy) and related taxa cause contact dermatitis, and fruits/seeds can be allergenic in sensitive individuals. See Pell and Urbatsch (2001) for a recent analysis of the family.

The Anacardiaceae are distinctive in being trees, shrubs, lianas, or perennial herbs with *resin ducts or laticifers* (some species causing allergenic responses), flowers generally 5-merous, with a *nectariferous disk* and *single ovule per carpel*, the fruit a *drupe* with a resinous mesocarp.

K usu. 5 or (5) **C** usu. 5 [0] **A** 5-10 [1, ∞] **G** (1-3,5) [(12)], superior, rarely inferior.

Rutaceae Rue / Citrus family (Latin for rue). 153 genera / 1800 species. (Figure 8.48)

The Rutaceae consist of trees, shrubs, lianas, or rarely herbs. The **stems** of some taxa have thorns. The **leaves** are simple, trifoliolate, or pinnate, sometimes pinnatifid, exstipulate, usually with pellucid glands. The **inflorescence** is a cyme or raceme, rarely of solitary flowers. The **flowers** are usually bisexual and actinomorphic, hypogynous, rarely epigynous. The **calyx** is aposepalous or synsepalous with 4-5 [2-3] sepals or lobes. The **corolla** is apopetalous or sympetalous with 4-5 [0, 2-3], imbricate or valvate petals or lobes.

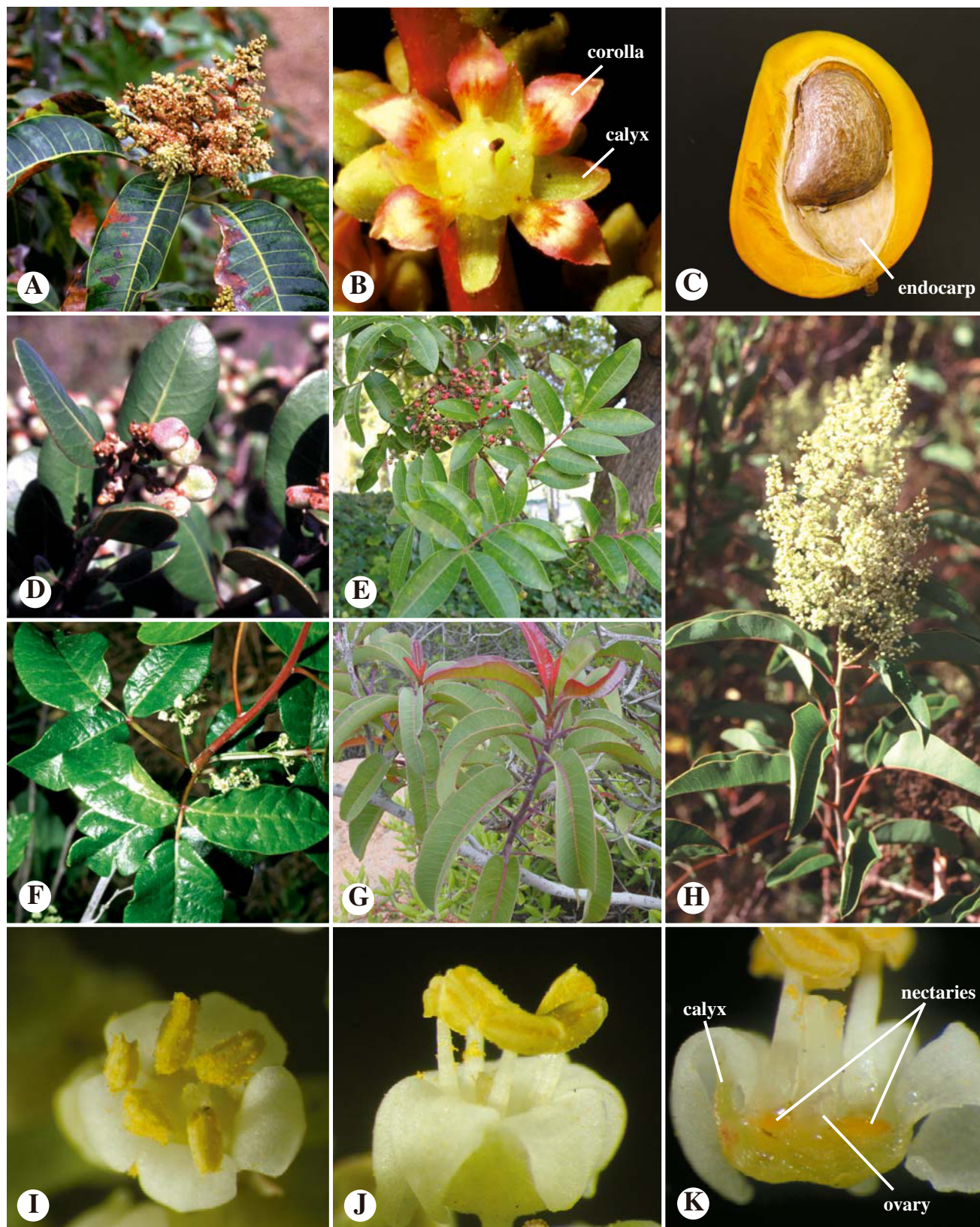


FIGURE 8.47 SAPINDALES. Anacardiaceae. A–C. *Mangifera indica*, mango. A. In florescence and leaves. B. Flower close-up. C. Fruit (drupe) longitudinal section, showing single, apical seed and endocarp. D. *Rhus integrifolia*, shoot with drupes. E. *Schinus terebinthifolius*, Brazilian pepper, shoot with drupes. F. *Toxicodendron diversilobum*, poison oak. G, K. *Malosma laurina*, laurel sumac. G. Vegetative shoot. H. In florescence. I, J. Flower close-ups. K. Flower longitudinal section, showing nectaries and ovary.

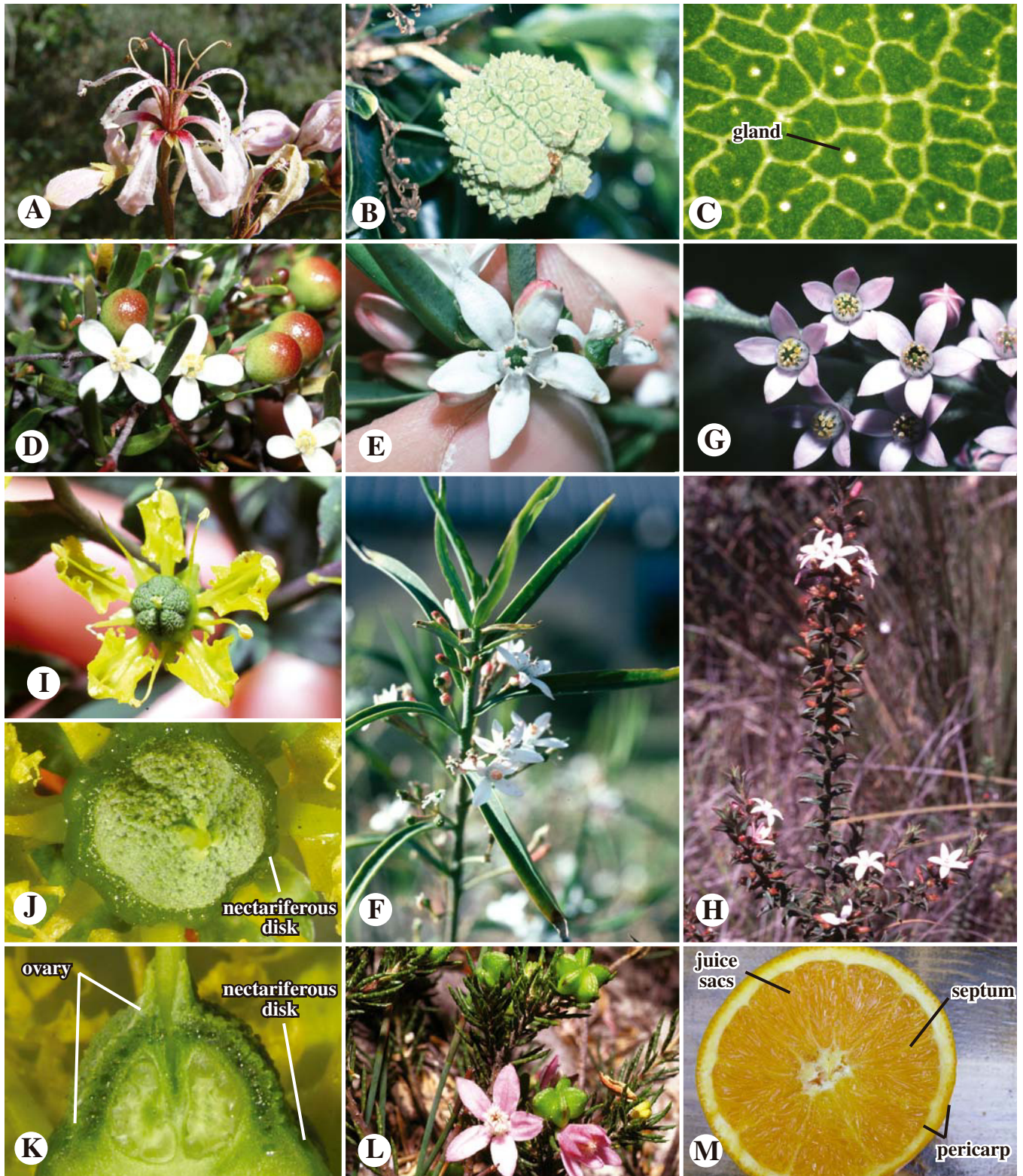


FIGURE 8.48 SAPINDALES. Rutaceae. A–C. *Calodendrum capense*. A. Flower. B. Fruit. C. Leaf close-up, showing internal, pellucid glands. D. *Cneoridium dumosum*. Note glands on fruits. E, F. *Eriostemon myoporoides*. G. *Eriostemon spicatum*. H. *Eriostemon buxifolius*. I–K. *Ruta graveolens*, rue. I. Flower, face view. J. Ovary close-up, showing glandular surface. K. Flower longitudinal section. L. *Philotheca salsolifolia*. M. *Citrus aurantiacus*, navel orange, fruit (seedless cultivar), a hesperidium.

The **stamens** are 8–10 ∞, usually diplostemonous, in 2 [1–4] whorls, with staminodes present in some taxa; filaments are often basally connate. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, rarely apocarpous, with a superior ovary, 4–5 [1–∞] carpels, and 4–5 [1–∞] locules. **Placentation** is axile; **ovules** are anatropous or hemitropous, bitegmic, 2 [1–∞] per carpel. **Nectaries** are usually present as an annular disk at the base of the ovary. The **fruit** is a schizocarp, berry, drupe, or hesperidium (the last with internal, swollen trichomes termed juice sacs). Secretory cavities containing ethereal oils are present in many tissues, including the leaves and pericarp.

Members of the Rutaceae have a worldwide distribution, especially in tropical regions. Economic importance includes many important fruits, among them *Citrus* spp. (oranges, grapefruits, lemons, limes, etc.), herbs such as *Ruta graveolens* (rue), timber trees, medicinal plants, and a number of ornamental cultivars. See Chase et al. (1999) for a recent phylogenetic analysis of the family.

The Rutaceae are distinctive in being trees, shrubs, lianas, or herbs, with simple to compound leaves and usually bisexual, actinomorphic, hypogynous, 4–5-merous flowers, typically with an **annular, nectariferous disk**, the fruit a schizocarp, berry, hesperidium, or drupe; **secretory glands containing ethereal oils** occur in many tissues, appearing as **pellucid glands** in the leaves and pericarp.

K 4–5 [2–3] **C** 4–5 [0, 2–3] **A** 8–10–∞ **G** (4–5) [(1–∞)], superior.

Sapindaceae [including Aceraceae] Soapberry / Maple family (name meaning Indian soap, from the use of soapberry). 133 genera / 1560 species. (Figures 8.49, 8.50)

The Sapindaceae consist of trees, shrubs, lianas, or herbaceous vines, tendrils present in some viney species. The **leaves** are simple, palmate, trifoliolate, pinnate, or bipinnate, usually spiral, but opposite in *Acer* and relatives, usually exstipulate. The **inflorescence** is a cyme or thyrs, rarely of solitary, axillary flowers, sometimes umbel- or corymblike. The **flowers** are usually unisexual, actinomorphic or zygomorphic, hypogynous, rarely perigynous. The **perianth** is with hypanthium present or absent. The **calyx** is aposepalous or basally synsepalous with 4–5 sepals or lobes. The **corolla** is apopetalous with 4–5 [3, 6+, 0] petals or lobes. The **stamens** are 8 [4–10+], with filaments often bearing trichomes. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary and 2–3 [–6] carpels and locules. The **styles** are 1–2. **Placentation** is apical-axile; **ovules** are anatropous, orthotropous, to campylotropous, 1 ∞ per carpel. **Nectaries** consist usually of an extrastaminal or intrastaminal annular disk or pad. The **fruit** is variable, fleshy or dry at maturity, being

a schizocarp of samaras in *Acer*. The **seeds** are exalbuminous, often with an aril or fleshy integuments. Tissues of many taxa contain soaplike saponins.

The Sapindaceae are distributed in tropical and temperate regions worldwide. Economic importance includes edible fruits/seeds, such as *Blighia* (akee), *Dimocarpus* (longan), and *Litchi* (litchi nut); timber trees; oil seeds; medicinal plants; stimulating (caffeine-containing) beverages, especially *Paullinia cupana* (guarana, prepared in a soft drink, especially popular in parts of South America); ornamental cultivars, such as *Koelreuteria* (golden-rain tree) and *Acer* (maple) spp.; arrow or fish poisons, e.g., *Jagera* and *Paullinia* spp.; and various taxa (e.g., *Sapindus saponaria*) used locally as a soap.

The Sapindaceae sensu APG II (2003) are distinctive in being trees, shrubs, lianas, or herbaceous vines with **simple, palmate, trifoliolate, or pinnate leaves**, the flowers typically with a **4–5-merous perianth, extrastaminal or intrastaminal nectariferous disk**, and **superior, 2–3-carpellate ovary**, the seeds often with an **aril or fleshy seed coat** (except in *Acer* and relatives); many taxa with soaplike **saponins** in tissues.

K 4–5 **C** 4–5 [3, 6+, 0] **A** 8 [4–10+] **G** (2–3) [(–6)], superior, hypanthium sometimes present.

ASTERIDS

The Asterids are a major group of eudicots, comprising a large percentage of angiosperms in total. According to the APG II (2003) system of classification, Asterids are divided into 10 orders, plus several families unplaced in orders (Figure 8.51, Table 8.3). The Asterids include such well known plants as dogwoods, hydrangeas, blueberries, phlox, tea, borage, gentians, mints, snapdragons, tomatoes/potatoes, carrots, scheffleras, hollies, bluebells, daisies, and a host of others.

Asterids are generally united by three major characters. One of these common features is chemical: the presence of **iridoid compounds** (Figure 8.52A). In addition, most Asterids have a **sympetalous corolla** (Figure 8.52B) and **unitegmic, tenuinucellate ovules** (Figure 8.52C; see Chapter 11 for an explanation). It isn't clear if any of these features are apomorphic for the entire group, as there is some variation and certainly homoplasy within.

The following are representative family descriptions of Asterids from most of the orders listed in Table 8.3. See Albach et al. (2001), Bremer et al. (2001, 2002), and references therein for more information about relationships within the Asterids.

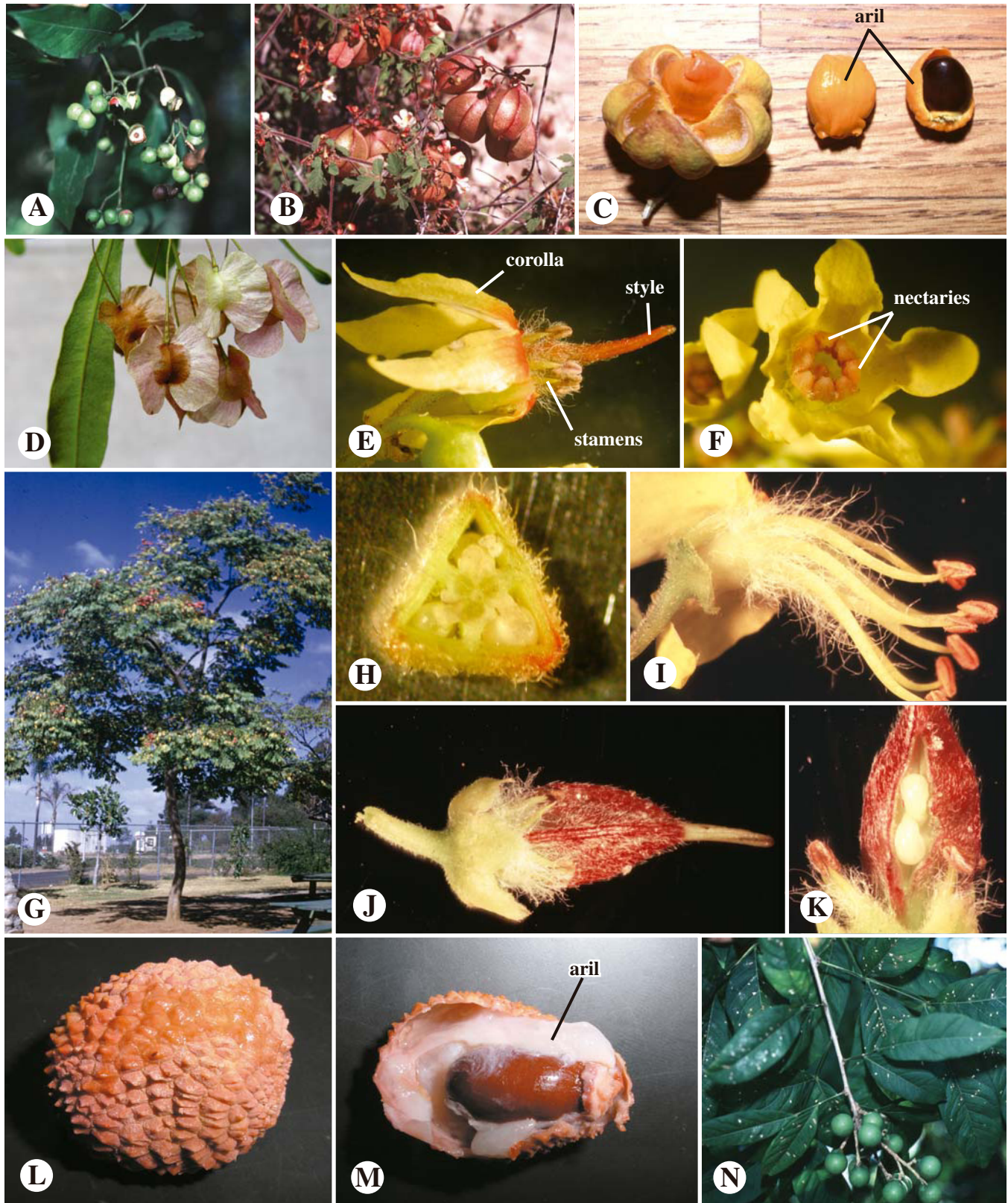


FIGURE 8.49 SAPINDALES. Sapindaceae. **A.** *Alectryon subcinereus*. **B.** *Cardiospermum corindum*. **C.** *Cupaniopsis anacardioides*, fruit with arillate seeds. **D.** *Dodonaea triquetra*, winged fruits. **E–K.** *Koelreuteria* spp. **E, F.** Flower close-ups. **G.** Tree. **H.** Ovary cross-section, showing axile placentation and three carpels. **I.** Androecium. **J, K.** Immature fruit. **L, M.** *Litchi chinensis*, litchi, the fruit a nut with a single, arillate seed (the aril edible). **N.** *Sapindus saponaria*, soap tree.

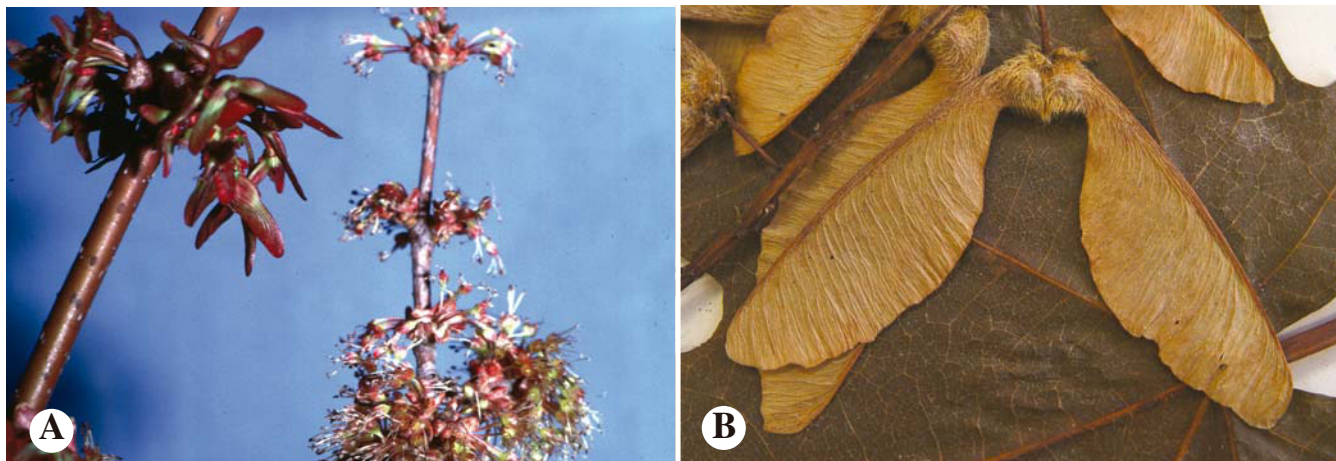


FIGURE 8.50 SAPINDALES. Sapindaceae, maples. **A.** *Acer rubrum*, red maple, in inflorescence (right) and young fruits (samaras, at left). **B.** *Acer macrophyllum*, big-leaf maple, the fruit a schizocarp of samaras.

CORNALES

The Cornales, sensu APG II, 2003, consists of six families (Table 8.3) and is the most basal order of the Asterids. Only the Cornaceae is treated here. Notable among the other families is the **Hydrangeaceae**, including important cultivated ornamentals, such as *Hydrangea*. See Xiang et al. (2002), Kubitzki (2004), and references therein for information about relationships and families within the order.

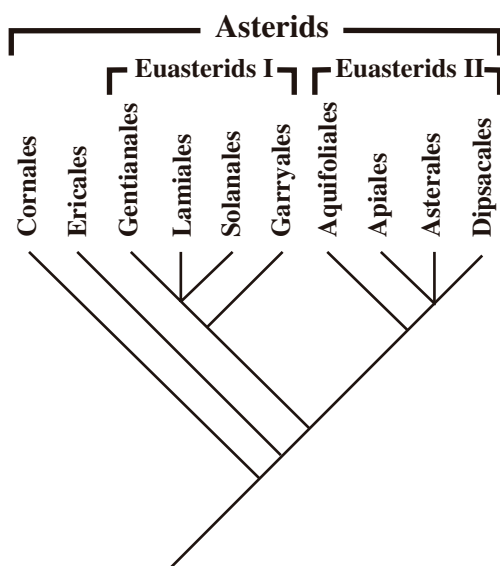


FIGURE 8.51 Cladogram of the orders of Asterids, from APG II (2003) and Albach et al. (2001).

Cornaceae [including Nyssaceae] Dogwood family (Latin for horn, after the hard wood). ca. 13 genera / 120 species. (Figure 8.53)

The Cornaceae consist of trees, shrubs, or rarely perennial, rhizomatous herbs. The **leaves** are simple, usually undivided [rarely pinnatifid], usually opposite [rarely spiral], and usually exstipulate. The **inflorescence** is a cyme or head of cymes, rarely a raceme, with showy, petaloid inflorescence bracts in some taxa. The **flowers** are bisexual [rarely unisexual], actinomorphic, and epigynous. The **perianth** is biseri-ate. The **calyx** is aposepalous, synsepalous and tubular in some unisexual male flowers, with 4 [5 7,0] sepals or calyx lobes. The **corolla** is apopetalous with 4 5 [10, 0 in unisexual female flowers] valvate or imbricate petals. The **stamens** are 4 5 [10], usually alternipetalous, uniseriate, rarely biseri-ate, and apostemonous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with an inferior ovary, 2 4 [9] carpels, and as many locules as carpels [rarely 1]. The **style** is solitary or as many as there are carpels. **Placentation** is apical and pendulous; **ovules** are anatropous, bitegmic [rarely unitegmic], 1 per locule. **Nectaries** usually consist of an infrastaminal annular disk. The **fruit** is usually a drupe, the endocarp grooved, 1 5-locular. The **seeds** are endospermous.

The Cornaceae are generally distributed in northern temperate regions. Economic importance includes cultivated ornamentals, such as *Cornus* (e.g., *C. florida*, flowering dogwood) and some timber and edible fruit trees. See Xiang et al. (1997, 1998) for more detailed information about relationships of the family.

TABLE 8.3 Orders and families of the Asterids, after APG II (2003). Families in bold are described in detail. An asterisk denotes a deviation from APG II, with brackets indicating the more inclusive family recommended by APG II. A double asterisk indicates a change suggested by Angiosperm Phylogeny Website (Stevens 2001 onwards). Including = incl.

ASTERIDS		
<p>CORNALES</p> <p>Cornaceae (incl. Nyssaceae)</p> <p>Curtisiaceae Grubbiaceae Hydrangeaceae Hydrostachyaceae Loasaceae</p> <p>ERICALES</p> <p>Actinidiaceae Balsaminaceae Clethraceae Cyrillaceae Diapensiaceae Ebenaceae Ericaceae (incl. Empetraceae, Epacridaceae, Monotropaceae, Pyrolaceae, etc.)</p> <p>Fouquieriaceae Lecythidaceae Maesaceae Marcgraviaceae Mitrastemonaceae** Myrsinaceae Pentaphylacaceae**</p> <p>Polemoniaceae Primulaceae Roridulaceae Sapotaceae Sarraceniaceae Sladeniaceae** Styracaceae Symplocaceae Tetrameristaceae (incl. Pellicieraceae) Theaceae Theophrastaceae</p>	<p>EUASTERIDS I</p> <p>Boraginaceae* Cordiaceae* [Boraginaceae] Ehretiaceae* [Boraginaceae] Heliotropaceae* [Boraginaceae] Hydrophyllaceae* [Boraginaceae] Lennoaceae* [Boraginaceae] Icacinaceae Oncothecaceae Vahliaceae</p> <p>GARRYALES Eucommiaceae Garryaceae</p> <p>GENTIANALES</p> <p>Apocynaceae (incl. Asclepiadaceae) Gelsmiaceae Gentianaceae Loganiaceae Rubiaceae</p> <p>LAMIALES</p> <p>Acanthaceae Bignoniaceae Byblidaceae Calceolariaceae (incl. some Scrophulariaceae, s.l.) Carlemanniaceae Gesneriaceae Gratiolaceae**</p> <p>Lamiaceae (Labiatae) Lentibulariaceae Linderniaceae** Martyniaceae Oleaceae Orobanchaceae (incl. some Scrophulariaceae, s.l.) Paulowniaceae Pedaliaceae Phymaceae (incl. some Scrophulariaceae, s.l.) Plantaginaceae (incl. Callitrichaceae, Plantaginaceae, s.s., and some Scrophulariaceae, s.l.) Plocospermataceae Schlegeliaceae Scrophulariaceae s.s. (incl. Myoporaceae) Stilbaceae (incl. some Scrophulariaceae, s.l.) Tetrachondraceae Verbenaceae</p> <p>SOLANALES</p> <p>Convolvulaceae Hydroleaceae Montiniaceae Solanaceae Sphenocleaceae</p>	<p>EUASTERIDS II</p> <p>Bruniaceae Columelliaceae Eremosynaceae Escalloniaceae Paracryphiaceae Polyosmaceae Sphenostemonaceae Tribelaceae</p> <p>APIALES</p> <p>Apiaceae (Umbelliferae)** (incl. Mackinlayaceae)</p> <p>Araliaceae Aralidiaceae** (incl. Torricelliaceae) Griselinaceae Melanophyllaceae Myodocarpaceae Pennantiaceae Pittosporaceae</p> <p>AQUIFOLIALES</p> <p>Aquifoliaceae Cardiopteridaceae Helwingiaceae Phyllonomaceae Stemonuraceae</p> <p>ASTERALES</p> <p>Alseuosmiaceae Argophyllaceae Asteraceae (Compositae) Calyceraceae Campanulaceae (incl. Lobeliaceae) Goodeniaceae Menyanthaceae Pentaphragmataceae Phellinaceae Rousseaceae Stylidiaceae (incl. Donatiaceae)</p> <p>DIPSACALES</p> <p>Adoxaceae Caprifoliaceae* Diervillaceae* [Caprifoliaceae] Dipsacaceae* [Caprifoliaceae] Linnaeaceae* [Caprifoliaceae] Morinaceae* [Caprifoliaceae] Valerianaceae* [Caprifoliaceae]</p>

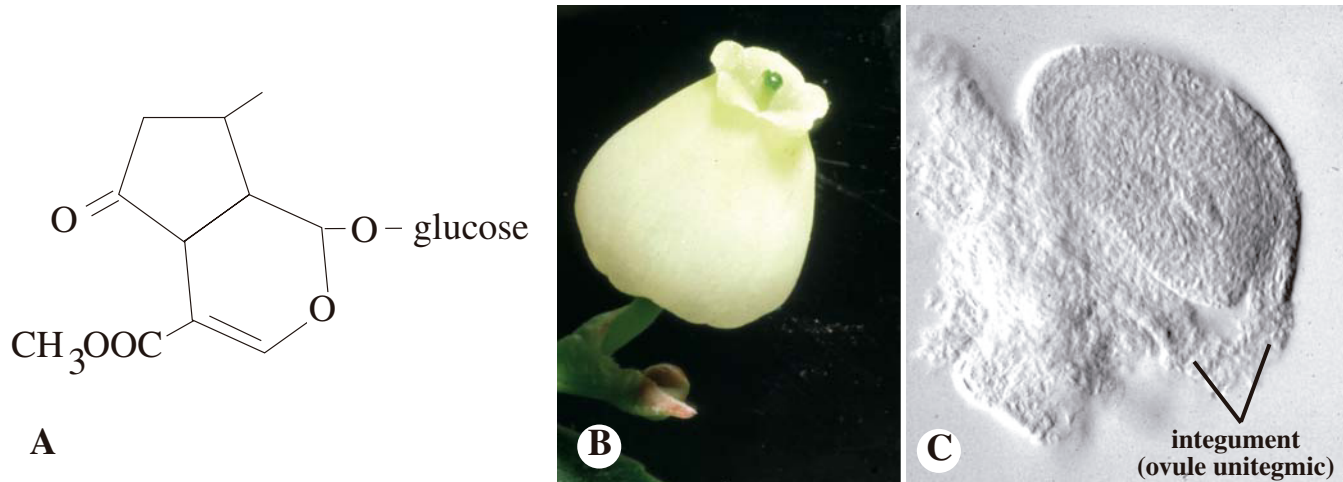


FIGURE 8.52 Putative apomorphies of the Asterids. **A.** Cornin, an iridoid compound. **B.** Sympetalous corolla. **C.** Unitemic, tenuinucellate ovule.

The Cornaceae are distinctive in being trees, shrubs, or rhizomatous herbs, with *simple, usually opposite and undivided leaves*, a usually *cymose inflorescence*, sometimes in *heads*, with *showy, petaloid bracts* in some taxa, the flowers usually biseriate and bisexual, *epigynous*, generally 4-5-merous, with an infrastaminal annual disk and *inferior ovary*, the fruit usually a multi-locular *drupe*.

K 4 [5-7, 0, connate in male flowers] **C** 4-5 [10, 0 in female flowers] **A** 4-5 [10] **G** (2-4) [(-9)], inferior.

ERICALES

The Ericales, sensu APG II (2003), is a large group of 23 (25) families (Table 8.3), only three of which are treated here. Notable among the families not described are the **Actinidiaceae** (containing *Actinidium chinensis*, kiwi fruit), **Ebenaceae** (including *Diospyros* spp., persimmon, and *Euclea pseudebenus*, black ebony), **Primulaceae** (the primrose family), **Sapotaceae** (source of important fruit and timber trees, as well as of *Palaquium gutta*, gutta-percha, and *Manilkara zapote*, chicle, the original chewing gum), **Sarraceniaceae** (pitcher plants), and **Theaceae** (including *Camellia sinensis*, tea). See Geuten et al. (2004) and Kubitzki (2004) for information about relationships within the order.

Ericaceae [including Empetraceae, Epacridaceae, Monotropaceae, Pyrolaceae] Heath family (from *Erica*, a name used by Pliny, adapted from Theophrastus). 106 genera / ca. 3355 species. (Figures 8.54, 8.55)

The Ericaceae consist of perennial, hermaphroditic or dioecious, shrubs and small trees, rarely lianas. Some taxa

are achlorophyllous and mycotrophic (fungus feeding, i.e., obtaining nutrition from mycorrhizal fungi in the soil, which in turn obtain nutrition from roots of vascular plants). The **roots** have endotrophic mycorrhizae. The **leaves** are simple, exstipulate, spiral, opposite, or whorled, sometimes with a basal pulvinus, evergreen, generally coriaceous, variable in shape, ranging from linear and strongly abaxially concave-revolute (ericoid) to broad and relatively flat, with pinnate-netted venation. The **inflorescence** is a raceme, fascicle, headlike cluster, or of solitary flowers. The **flowers** are bisexual [rarely unisexual], actinomorphic, pedicellate, bracteate (with two bracteoles), hypogynous or epigynous. The **perianth** is biseriate and dichlamydeous or uniseriate, urceolate, tubular, or campanulate. The **calyx** is aposepalous with 5 [2-7], valvate or imbricate sepals. The **corolla** is sympetalous, rarely apopetalous, with 5 [2-7 or 0 in some], convolute or imbricate lobes or petals. The **stamens** are 5+5 [rarely >10, 2-4], whorled, usually obdiplostemonous and biseriate [rarely uniseriate], usually distinct and epipetalous. **Anthers** of all but basal members are developmentally inverted such that the anther base assumes an apical position, dehiscence poricidal or longitudinal and introrse at maturity by inversion, connective appendages present in some taxa. The **pollen** is typically shed in tetrahedral tetrads [monads in basal members]. The **gynoecium** is syncarpous, with a superior or less often inferior (e.g., *Vaccinium*) ovary, 5 [2-10] carpels, and 5 [2-10] locules. The **style** is solitary, terminal, hollow, fluted. **Placentation** is axile, apical-axile, basal-axile, or parietal; **ovules** are anatropous to campylotropous, unitemic, 1 ∞ per carpel. **Nectaries** are present, as an intrastaminal disk. The **fruit** is a capsule, berry, or drupe. The **seeds** are endospermous (oily and proteinaceous).

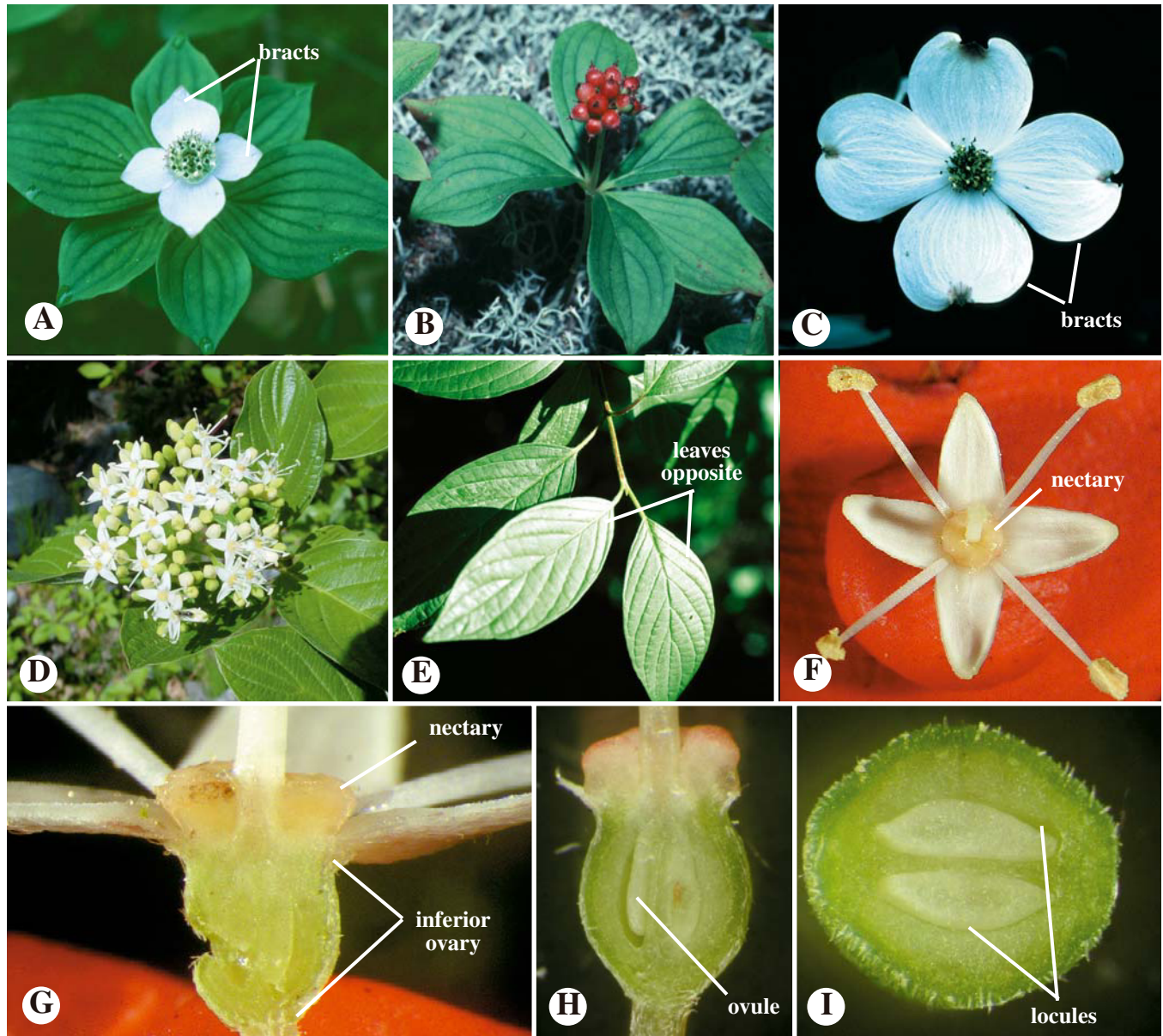


FIGURE 8.53 Cornaceae. **A,B.** *Cornus canadensis*, a subshrub. **A.** Inflorescence with showy bracts subtending flowers. **B.** Inflorescence, of berries. **C.** *Cornus florida*, a tree with large showy inflorescence bracts. **D–I.** *Cornus sericea*, a small tree. **D.** Inflorescence, a corymb lacking showy bracts. **E.** Leaves, opposite in arrangement. **F.** Flower, face view, showing petals, alternipetalous stamens, nectariferous disk, and style/stigma. **G.** Flower, longitudinal section, showing inferior ovary and nectary. **H.** Ovary, longitudinal section, showing apical-axile placentation. **I.** Ovary, cross-section, 2-carpellate and 2-loculate.

The Ericaceae are now circumscribed to include the former families Empetraceae, Epacridaceae, Monotropaceae, Pyrolaceae, and others. The Ericaceae has recently been classified into 8 subfamilies: Enkianthoideae, Monotropoideae (including some achlorophyllous and parasitic members), Arbutoideae (including *Arbutus* and *Arctostaphylos*, the manzanitas), Cassiopoideae, Ericoideae (including *Erica*,

Rhododendron, and the former Empetraceae), Harrimanelloideae, Styphelioideae (including the former Epacridaceae), and Vaccinioideae (including *Vaccinium*, blueberries, and relatives; see Kron et al., 2002.) Members of the family grow in acid soils typically; various species (e.g., *Erica*) are dominants in bog, moorland or heathland communities with distributions worldwide in temperate and tropical

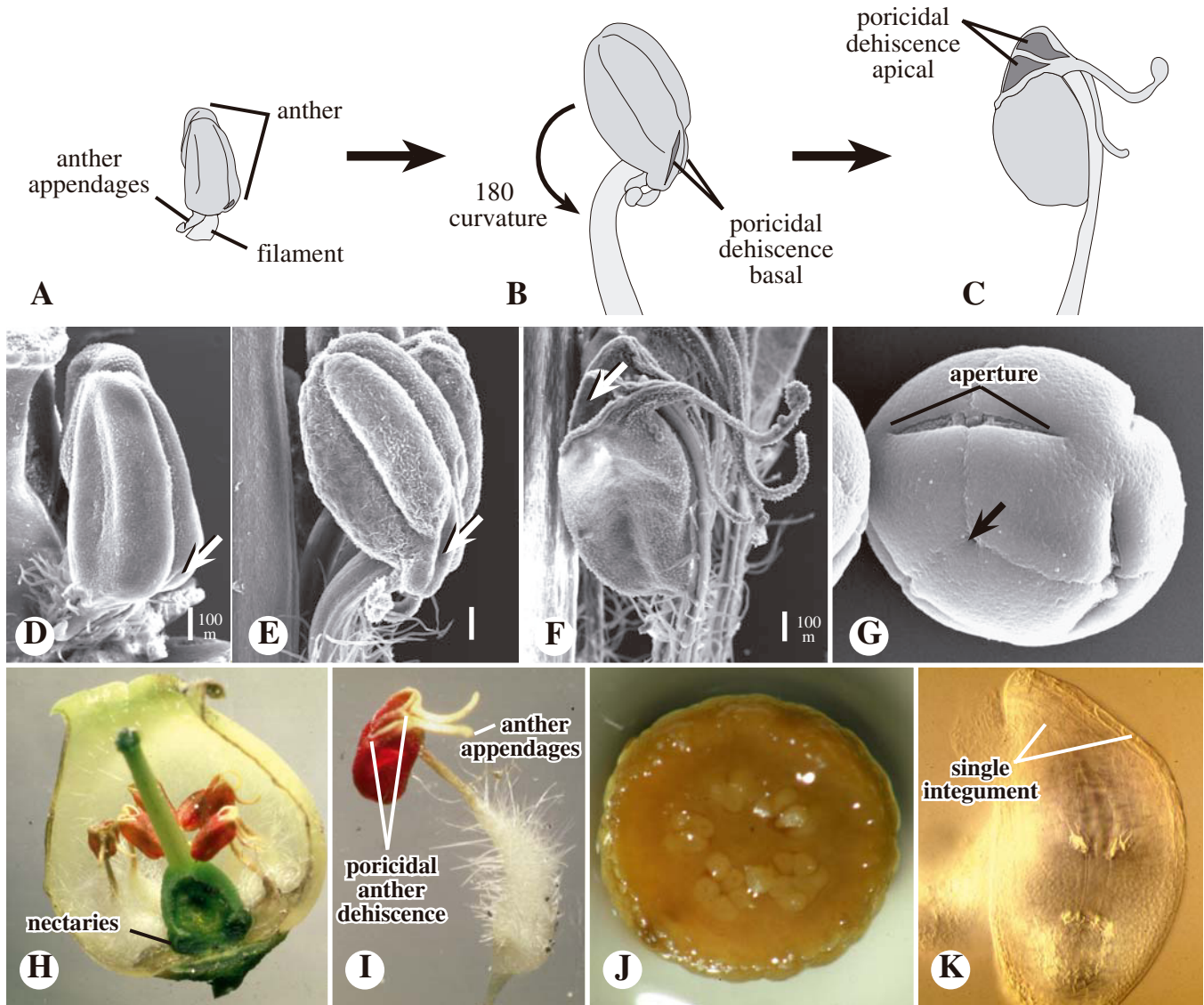


FIGURE 8.54 ERICALES. Ericaceae. A–C. Diagrammatic development of inverted anthers, characteristic of the family; note 180° inversion of anther base, in this case associated with anther appendages. D–F. *Xylococcus bicolor*, development of inverted anthers, showing inversion of anther base (arrow). G–K. *Arbutus unedo*. G. Pollen grain tetrad, showing colporate apertures; note junction of three (of four) grains (arrow). H. Flower in longitudinal section; note urceolate corolla, superior ovary, and basal nectaries. I. Stamen, showing poricidal dehiscence and anther appendages. J. Ovary cross-section, showing 5 carpels and locules and axile placentation. K. Unitegmatic ovule, typical of the Ericaceae and other Asterids.

(mostly montane) regions. Economic importance includes cultivated ornamentals, especially *Rhododendron* [*Azalea*] and *Erica*; *Vaccinium* species (including blueberry and cranberry) are important fruit plants.

The Ericaceae are distinctive in being usually *evergreen shrubs* (some achlorophyllous and mycotrophic), with *coriaceous, linear-revolute to broad-flat* leaves, a sympetalous corolla, stamens usually developing by *anther inversion*,

dehiscence often *poricidal* and an intrastaminal disk. See Kron et al. (2002) and references therein for information on phylogenetic relationships within the Ericaceae.

K 5 [2-7] **C** (5) [(2-7), 0] **A** 5+5 [2-4, ∞] **G** (5) [(2-10)], superior (usually) or inferior.

Fouquieriaceae Ocotillo family (after the Frenchman P. E. Fouquier). 1 genera / 11 species. (Figure 8.56)

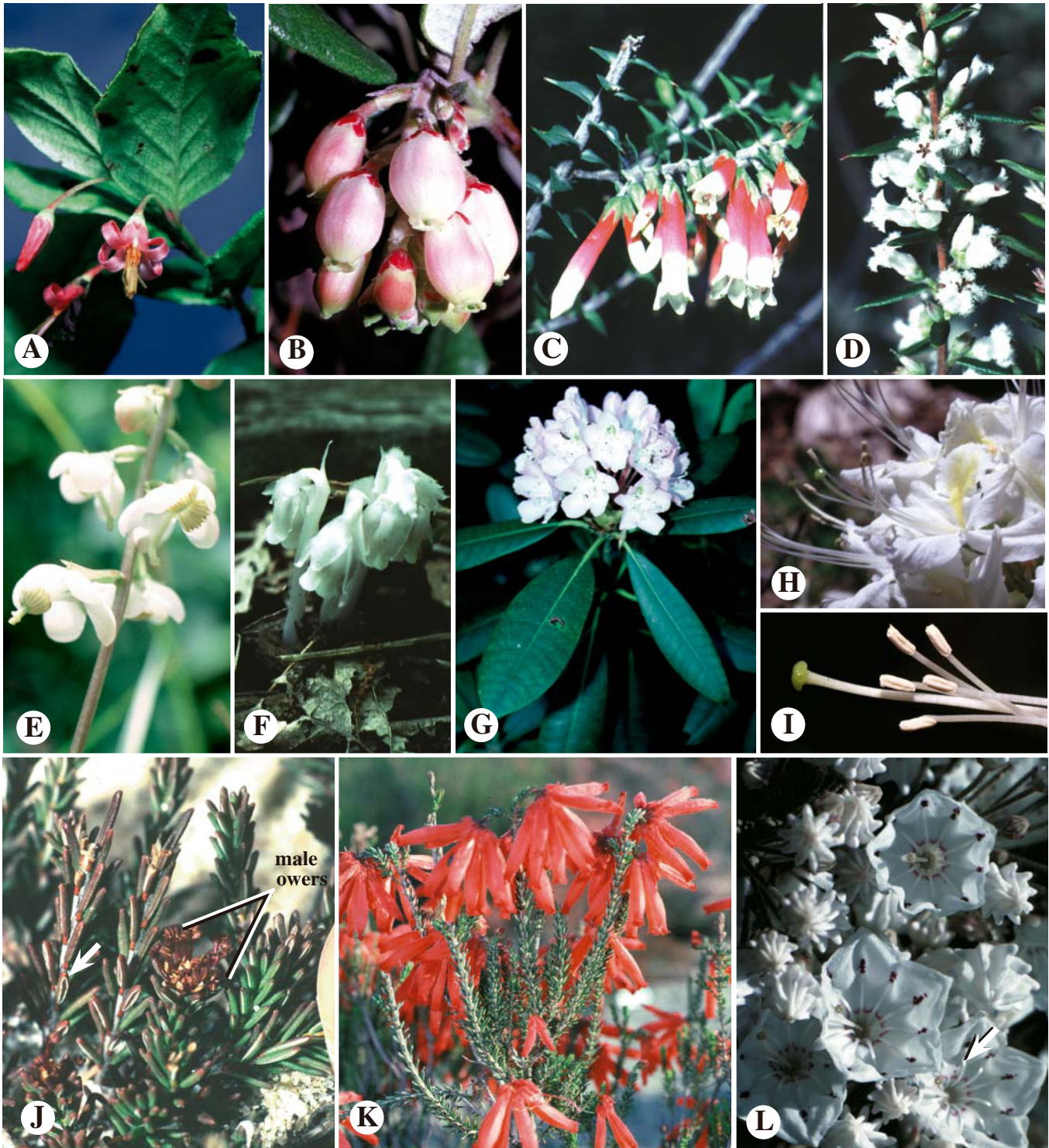


FIGURE 8.55 ERICALES. Ericaceae. **A.** *Vaccinium erythrocarpum*. **B.** *Xylococcus bicolor*, with urceolate corollas, typical of many Ericaceae. **C.** *Epacris longiora*, with long tubular, reddish flowers and short, coriaceous leaves. **D.** *Leucopogon ericoides*. **E.** *Pyrola rotundifolia*, an achlorophyllous, mycotrophic species. **F.** *Monotropa uniflora*, another achlorophyllous, mycotrophic species. **G.** *Rhododendron maximum*, with zygomorphic flowers and large, evergreen, coriaceous leaves. **H,I.** *Rhododendron occidentale*. **H.** Zygomorphic flowers with long, exserted stamens and style. **I.** Close-up of style and stamens, the latter with poricidal anthers. **J.** *Corema conradii*, close-up of low, decumbent shrub; note reduced, wind-pollinated flowers and "ericoid" leaves with abaxial cleft (arrow). **K.** *Erica centranthoides*, with red, tubular corollas and small, linear, "ericoid" leaves. **L.** *Kalmia latifolia*, in which stamens are bent (arrow) into pockets that catapult pollen with release of anthers.

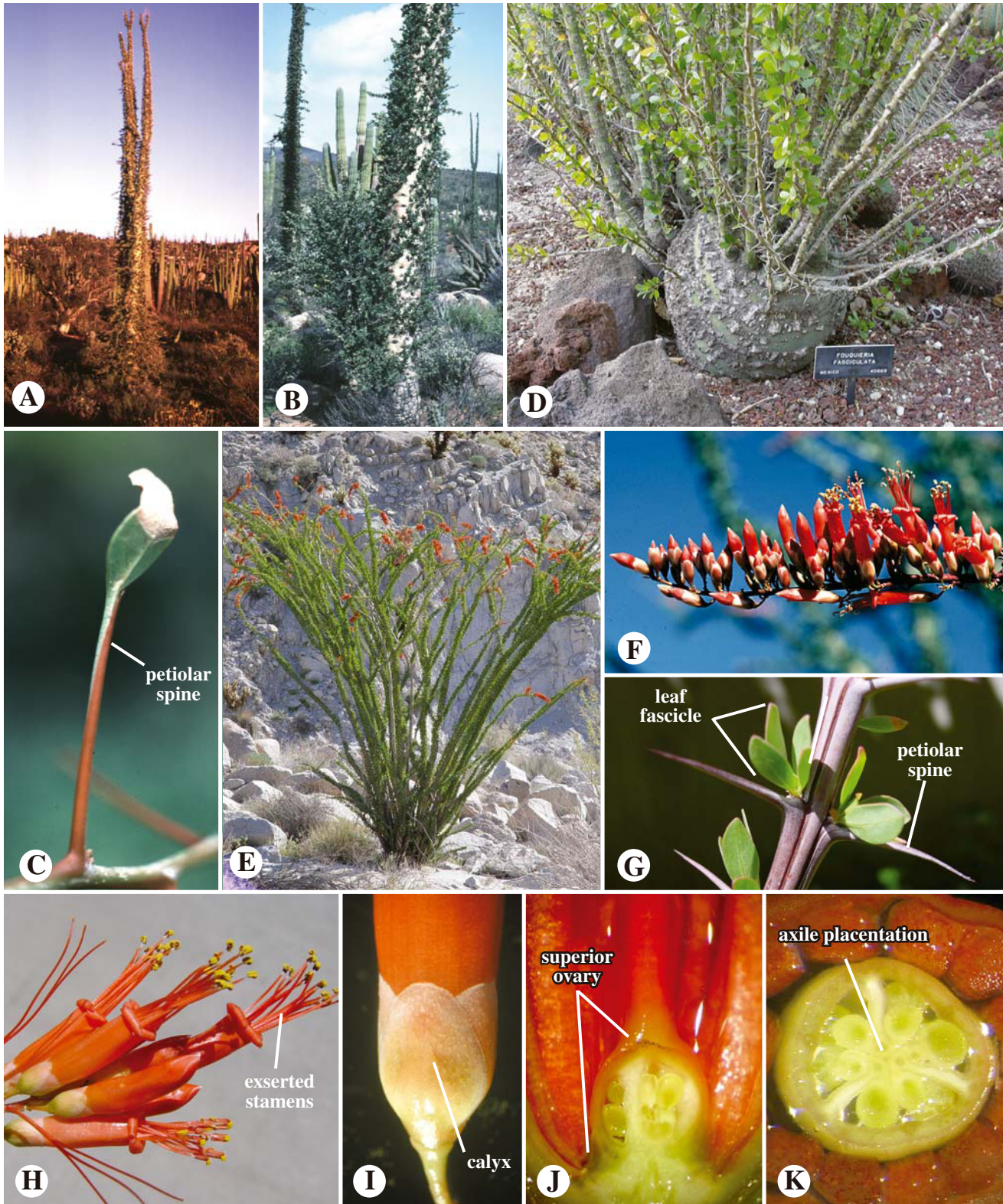


FIGURE 8.56 ERICALES. Fouquieriaceae. **A–C.** *Fouquieria columnaris*, boojum or cirio, a pachycaulis plant native to deserts of Baja California, Mexico. **A,B.** Plant habit. **C.** Close-up of leaf of long shoot, the blade senescing, leaving a petiolar spine. **D.** *Fouquieria fasciculata*, a caudiciform species. **E–K.** *Fouquieria splendens*, ocotillo. **D.** Whole plant, a tall shrub ca. 4 m tall. **F.** Inflorescence. **G.** Close-up of shoot, showing petiolar spines (derived from original leaves of long shoot) plus fascicles (short shoots), having drought-deciduous leaves. **H.** Flowers, with exserted stamens. **I.** Flower base, showing calyx. **J.** Ovary longitudinal section. **K.** Ovary cross-section; note three carpels with axile placentation.

The Fouquieriaceae consist of xeromorphic shrubs or trees. The **stems** are woody to succulent, ridged. The **leaves** are simple, undivided, spiral, those of long shoots forming petiolar spines, these with axillary fascicles of drought-deciduous nonspiny leaves. The **inflorescence** is a terminal spike, raceme, or panicle. The **flowers** are bisexual, actinomorphic, and hypogynous. The **calyx** is aposepalous with 5 imbricate sepals. The **corolla** is sympetalous and with 5, tubular to salverform, imbricate lobes. The **stamens** are 10–18 [23], uniseriate, apostemonous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 carpels, with 3 locules at base, 1 at apex. The **style** is solitary and 3-branched. **Placentation** is axile below, parietal above; **ovules** are ∞ , anatropous, bitegmic. The **fruit** is a loculicidal capsule. The **seeds** are endospermous.

The Fouquieriaceae are distributed in southwestern North America. Economic importance is limited; *Fouquieria splendens* (ocotillo) is planted as a fence or hedge; *Fouquieria columnaris* (boojum, cirio) is a spectacular pachycaul of Mexican deserts.

The Fouquieriaceae are distinctive in being *xeromorphic, sometimes succulent* shrubs or trees, bearing *long shoot leaves with petiolar spines*, in axils of which develop *fascicles of drought-deciduous, nonspiny leaves*, the flowers mostly pentamerous, with a sympetalous corolla and superior, *tricarpellate ovary* having axile-parietal placentation. See Schultheis and Baldwin (1999) for more information about family systematics.

K 5 **C** (5) **A** 10–18 [23] **G** (3), superior.

Polemoniaceae Phlox family (Greek, possibly from a King Polemon, name used by Pliny). 20 genera / 290 species. (Figure 8.57)

The Polemoniaceae consist of annual or perennial herbs, shrubs, lianas, or small trees. The **leaves** are simple or pinnate, divided in some, spiral, opposite, or whorled, exstipulate. The **inflorescence** is a head, cyme, or of solitary flowers. The **flowers** are bisexual, actinomorphic or zygomorphic, hypogynous. The **perianth** is biseriate and dichlamydeous. The **calyx** is usually synsepalous with five [4–6] lobes. The **corolla** is sympetalous and rotate, salverform, or bilabiate, with five [4–6] lobes, convolute in bud. The **stamens** are five [4–6], whorled, alternipetalous, uniseriate, epipetalous (inserted at different levels on corolla tube in some taxa). **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 3 [2,4] carpels, and 3 [2,4] locules. The **style** branches and **stigmas** are 3 [2,4]. **Placentation** is axile; **ovules** are anatropous to hemitropous, unitegmic, 1 ∞ . **Nectaries** consist of an annular nectariferous disk around ovary base. The **fruit** is a capsule, usually longitudinally dehiscent. The **seeds**

are endospermous (oily). The stem xylem typically occurs in a continuous ring.

Members of the Polemoniaceae have distributions in the Americas (especially western North America) and Eurasia. Economic importance includes numerous cultivated ornamentals, such as *Cobaea*, *Gilia*, *Ipomopsis*, *Plox*, and *Polemonium*. See Wilken (2004) for a general description and Porter (1997), Porter and Johnson (1998), and Prather et al. (2000) for data on infrafamilial relationships.

The Polemoniaceae are distinctive in being herbs, shrubs, or small trees with simple (divided in some) or pinnate leaves, a typically 5- [4,6] *merous, synsepalous, sympetalous perianth*, 5 [4,6] *epipetalous stamens*, and a typically 3- [2,4] *carpellate*, superior ovary, the fruit a *loculicidal capsule*.

K (5) [4,6] **C** (5) [4,6] **A** 5 [4,6] **G** (3) [(2),(4)], superior.

EUASTERIDS I

Euasterids I is an informal name used to denote a monophyletic grouping including four orders Garryales, Gentianales, Lamiales, and Solanales plus some families unplaced in orders (Table 8.3). Two, unplaced families are described first.

Boraginaceae s.s. — Borage family (possibly meaning shaggy coat, in reference to the leaves). 130 genera / 2300 species. (Figure 8.58)

The Boraginaceae s.s. consist of usually hermaphroditic herbs, often with hirsute or hispid vestiture. The **leaves** are simple, spiral, and exstipulate. The **inflorescence** unit is a monochasial (e.g., scorpioid or helicoid) cyme, rarely of solitary, axillary flowers. The **flowers** are usually bisexual, actinomorphic, hypogynous. The **perianth** is biseriate and dichlamydeous, often salverform or rotate, with hypanthium absent. The **calyx** is apo- or synsepalous with five [4–8] imbricate, rarely valvate sepals or lobes. The **corolla** is sympetalous and with five [4–6], convolute or imbricate lobes. The **stamens** are 5 [4–6], whorled, and epipetalous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 4 locules. The **style** is solitary and gynobasic, with the ovary deeply 4-lobed by formation of false septa dividing each carpel; **stigmas** are 1–2. **Placentation** is basal; **ovules** are anatropous to hemitropous, unitegmic, 2 per carpel. **Nectaries** are present in some taxa as a ring around the ovary base. The **fruit** is a schizocarp of usually four nutlets.

The Boraginaceae s.s., as treated here, are essentially equivalent to the subfamily Boraginoideae of an expanded

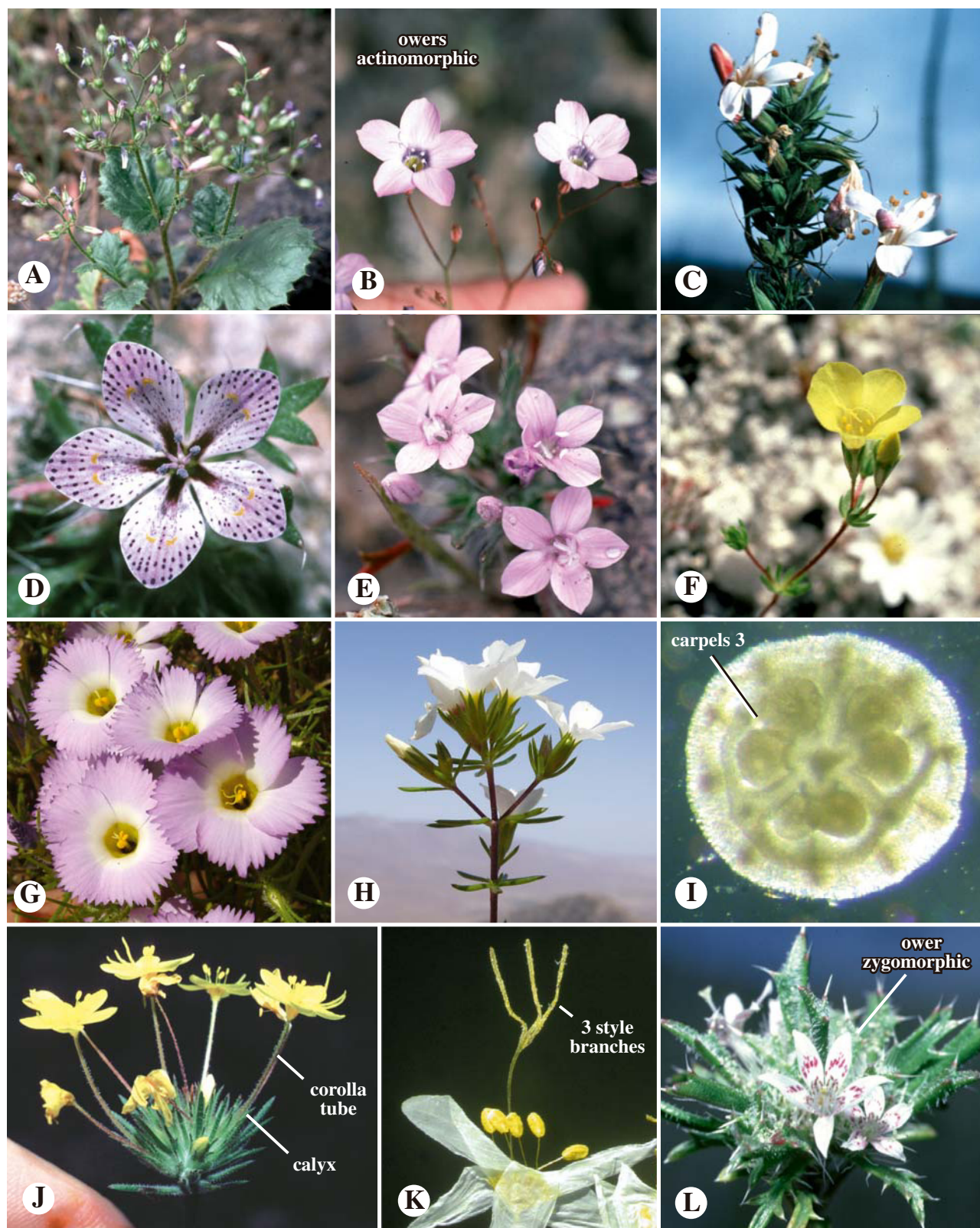


FIGURE 8.57 ERICALES. Polemoniaceae. **A.** *Gilia latifolia*, whole plant. **B.** *Gilia* sp., flower close-up, showing actinomorphic, sympetalous corolla. **C.** *Acanthogilia gloriosa*, a spiny shrubby species. **D.** *Langloisia punctata*, corolla with prominent nectar guides. **E.** *Langloisia setosissima*. **F.** *Leptosiphon aureus*. **G.** *Linanthus dianthiflorus*, with fringed corolla lobes. **H.** *Leptosiphon oribundus*, showing cyme in inflorescence. **I–K.** *Leptosiphon parviflorus*. **I.** Ovary cross-section, showing three carpels/locules and axile placentation. **J.** Whole plant, showing long corolla tube. **K.** Style, apically three-branched. **L.** *Loeseliastrum schottii*, with zygomorphic flowers.



FIGURE 8.58 Boraginaceae. **A.** *Amsinckia intermedia*, showing salverform corolla and helicoid in inflorescence. **B–H.** *Borago officinalis*. **B.** In inflorescence, a monochasial cyme. **C.** Flower close-up. **D.** Close-up of corona and staminal appendages. **E,F.** Gynoecium, showing four-lobed ovary and gynobasic style. **G.** Fruit close-up. **H.** Single nutlet, with caruncle. **I.** *Cryptantha intermedia*, in inflorescence unit, a monochasial cyme. **J.** *Plagiobothrys acanthocarpus*, having uncinately nutlets. **K,L.** *Echium fastuosum*, with monochasial (helicoid) cyme unit. **M.** *Mertensia virginica*. **N–Q.** Nutlet variation in *Cryptantha* spp. Scale bar = 1 mm. **N.** *C. abata*. **O.** *C. pterocarya*. **P.** *C. cf. nivalis*. **Q.** *C. dumetorum*.

Boraginaceae (as classified by APG II 2003). Subfamilies Cortioideae, Ehretioideae, and Heliotropoideae of Boraginaceae s.l. are treated here as separate families, after Gottschling et al. (2001) (Table 8.3). Members of the Boraginaceae s.s. have a worldwide distribution. Economic importance is limited, some used as herbs (e.g., *Borago officinalis*, borage), dyes, or cultivated ornamentals (e.g., *Myosotis*, forget-me-not).

The Boraginaceae s.s. are distinctive in being mostly herbs with simple, spiral leaves, a inflorescence of *monochasial (helicoid or scorpioid) cymes*, and *actinomorphic, sympetalous* flowers with a 4-ovuled, *deeply 4-lobed ovary* (by development of false septa) with a *gynobasic style*, the fruit a *schizocarp of usually 4 nutlets*. See Gottschling et al. (2001) for more information about relationships of the Boraginaceae to other groups.

K 5 or (5) [4-8,(4-8)] **C** (5) [(4,6)] **A** 5 [4,6] **G** (2), superior, hypanthium absent.

Hydrophyllaceae — Waterleaf family (*hydro*, water, + *phyllum*, leaf). 18 genera / 270 species. (Figure 8.59)

The Hydrophyllaceae consist of herbs or shrubs, often odiferous with glandular trichomes or hirsute/hispid. The **leaves** are simple or pinnate, rarely palmate, pinnatisect in some, spiral to opposite, exstipulate. The **inflorescence** is a cyme, often helicoid, or of solitary flowers. The **flowers** are bisexual, actinomorphic to slightly zygomorphic. The **perianth** is biserate and dichlamydeous, hypanthium absent. The **calyx** is usually synsepalous (only basally connate in some taxa) with five [4,10 12], imbricate lobes or sepals. The **corolla** is sympetalous with five [4,10 12], convolute lobes. The **stamens** are five [4,10 12], whorled, alternipetalous, epipetalous, usually with scalelike structures at either side of the filament at the junction with the corolla tube. The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 1 or 2 locules. The **styles** are 2 or 2-branched. **Placentation** is parietal or axile; **ovules** are anatropous to amphitropous, unitegmic, 2 ∞ per carpel. **Nectaries** are usually absent. The **fruit** is a capsule, sometimes irregularly or not dehiscent. The **seeds** are oily endospermous.

Members of the Hydrophyllaceae have a mostly worldwide distribution (except in Australia), with many in western North America. Economic importance includes some cultivated ornamentals. See Gottschling et al. (2001) for more information about relationships of the Hydrophyllaceae to other groups.

The Hydrophyllaceae are distinctive in being herbs, shrubs, or rarely trees with simple, pinnatisect, or pinnate, *spiral leaves*, an inflorescence *often of helicoid cymes*, and *actinomorphic, sympetalous* flowers, often with a *pair of scales* at junction of stamen filament with corolla tube, a 2 ∞ ovuled, *unlobed, 2-carpellate, superior ovary*, the fruit a *capsule*.

K 5 or (5) [4-∞, (4-∞)] **C** (5) [(4-∞)] **A** 5 [4-∞] **G** (2), superior.

GENTIANALES

The Gentianales, sensu APG II, 2003, contain five families (Table 8.3), three of which are treated here.

Apocynaceae [incl. Asclepiadaceae] — Dogbane/Milkweed family (Greek for away from dog, in reference to some taxa used as dog poison). 411 genera / 4650 species. (Figures 8.60, 8.61)

The Apocynaceae consist of lianas, trees, shrubs, or herbs, with latex present in tissues. The **stems** are succulent in some taxa, e.g., the stapelioids. The **leaves** are simple, undivided, sometimes reduced, opposite, whorled, or rarely spiral, usually exstipulate. The **inflorescence** is a cyme (often umbelliform in Asclepiadoids), raceme, or of solitary flowers. The **flowers** are usually bisexual, actinomorphic, and hypogynous. The **calyx** is usually synsepalous (at least basally) with 5 imbricate or valvate lobes. The **corolla** is sympetalous with 5 convolute (rarely valvate or imbricate) lobes. The **stamens** are 5, alternipetalous, epipetalous, apostemonous to monadelphous; in Asclepiadoids the stamens are connate to the stigma to form a gynostegium, often elaborate with appendages (hoods and horns). The **pollen** in Asclepiadoids is connate into pollinia, each consisting of a 2-lobed, glandlike corpusculum attached to two armlike retinacula, each bearing connate pollen grains from one of the thecae of separate, adjacent anthers. The **gynoecium** is syncarpous, often only apically (with ovaries distinct), with a superior, rarely half-inferior ovary, 2 [8] carpels, and 1 2 locules. The **stigma(s)** are capitate, a single, broad fusion product in Asclepiadoids. **Placentation** is apical and pendulous or marginal; **ovules** are anatropous, unitegmic, ∞ [1] per carpel. **Nectaries** are sometimes with 5 nectar glands or a disk at ovary base. The **fruit** is a variable and can be a berry, drupe, or follicle; in Asclepiadoids the fruit is a schizocarp of two follicles (one often not developing). The **seeds** are endospermous. Plants typically contain various glycosides and alkaloids. In Asclepiadoids, the glandlike corpusculum of the pollinia becomes attached to the legs of insect pollinators, the entire apparatus pulled through grooves in the side of the gynostegium.

The Apocynaceae is often treated as two families, Apocynaceae (Dogbane family) and Asclepiadaceae (Milkweed family), the latter now usually classified as subfamily Asclepiadoideae (Asclepiadoids). Members of the family have a mostly worldwide distribution, mostly in tropical regions. Economic importance includes uses as cultivated ornamentals, such as *Nerium* (oleander), *Plumeria*, *Stapelia*, and *Vinca*

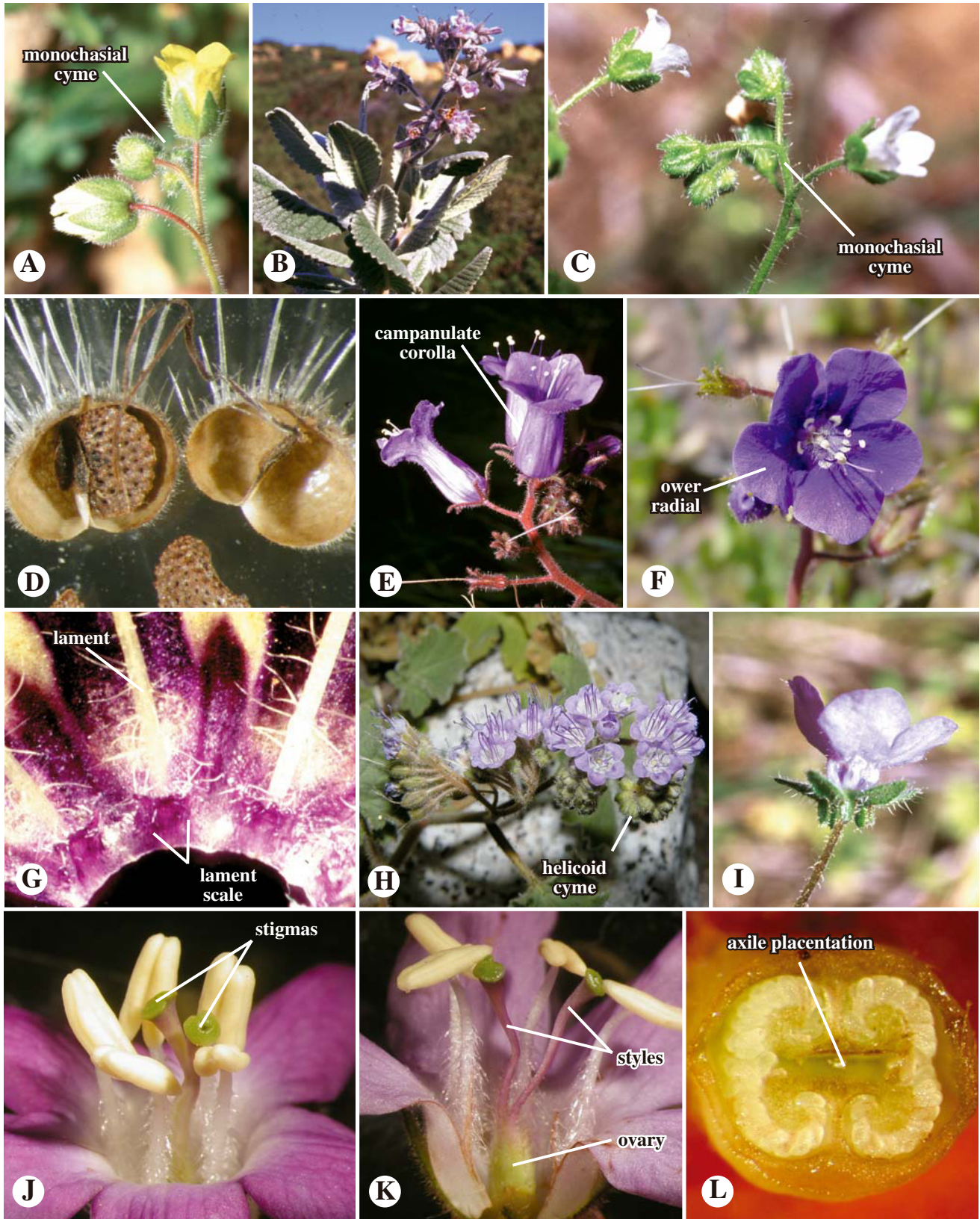


FIGURE 8.59 Hydrophyllaceae. **A.** *Emmenanthe penduli ora*, showing monochasial cyme. **B.** *Eriodictyon crassifolium*, yerba santa. **C.** *Eucrypta chrysanthemifolia*, in orescence unit. **D.** *Phacelia cicutaria*, two valves of capsule. **E.** *Phacelia minor*, owers. **F,G.** *Phacelia parryi*. **F.** Flower. **G.** Corolla, basal adaxial side, showing scales at base of stamen laments. **H.** *Phacelia pedicellata*, with helicoid cyme units. **I.** *Pholistoma auritum*, ower, side view, showing sympetalous, rotate corolla. **J–L.** *Wigandia caracasana*. **J.** Whole ower. **K.** Flower longitudinal section. **L.** Ovary cross-section, showing two carpels and axile placentation.

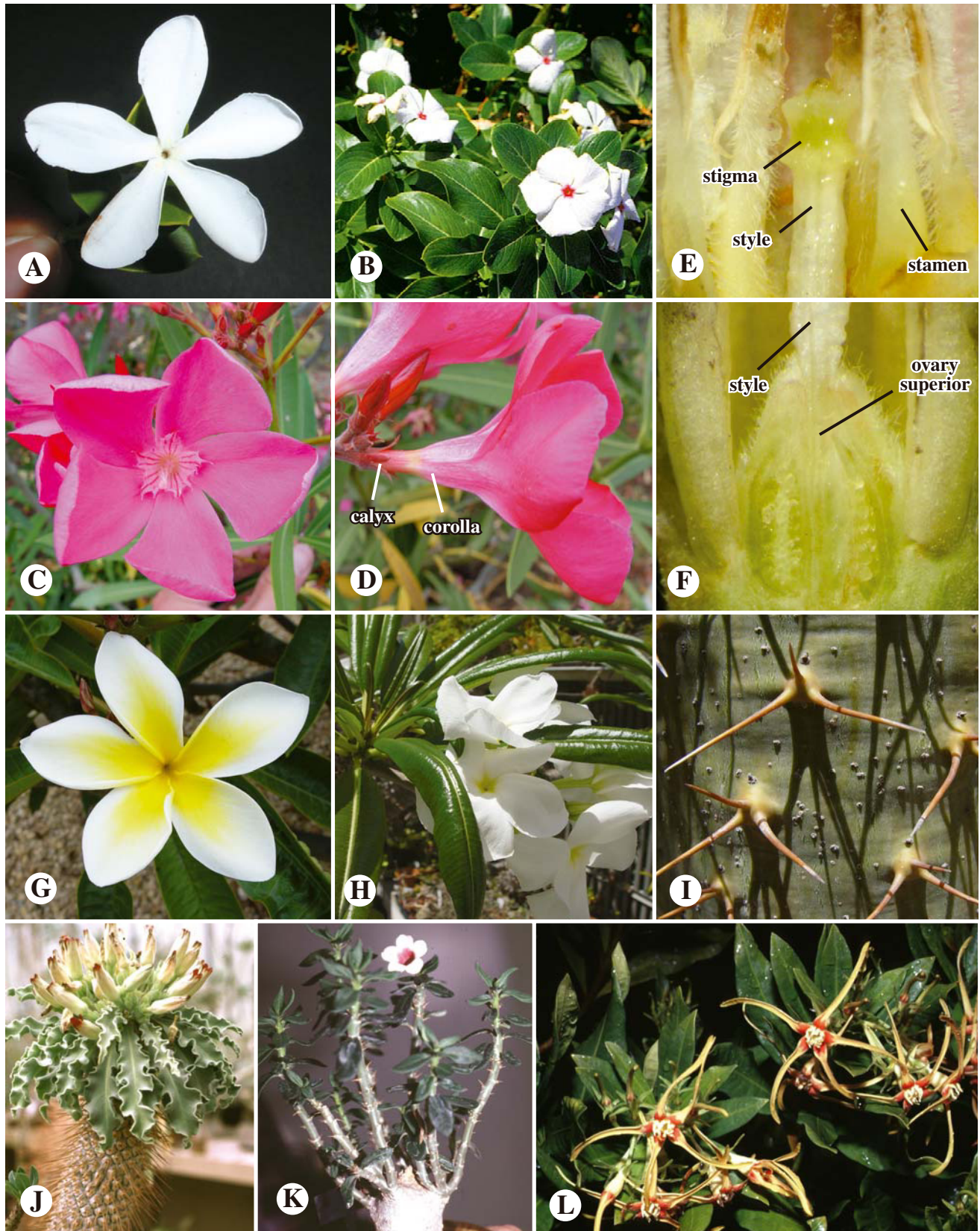


FIGURE 8.60 GENTIANALES. Apocynaceae. **A.** *Carissa grandiflora*, Natal plum; flower, face view. **B.** *Catharanthus roseus*, Madagascar periwinkle. **C–F.** *Nerium oleander*, oleander. **C.** Flower, face view. **D.** Flower, side view. **E,F.** Flower longitudinal section, showing style, stigma (**E**), and ovary (**F**). **G.** *Plumeria* sp., flower in face view. **H,I.** *Pachypodium lamerei*. **H.** Upper shoot. **I.** Spines of trunk. **J.** *Pachypodium namaquanum*. **K.** *Pachypodium bispinosum*. **L.** *Strophanthus speciosus*.

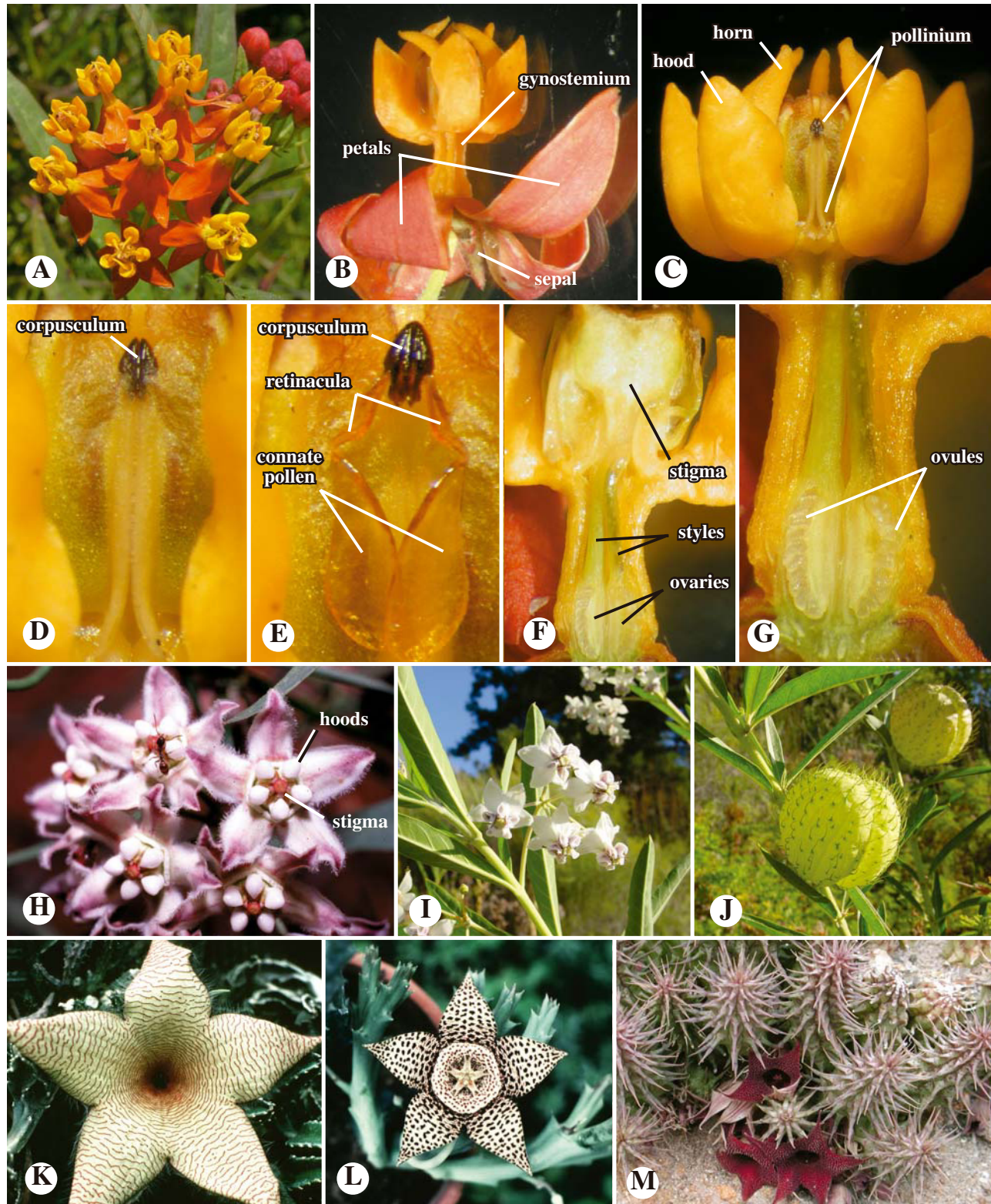


FIGURE 8.61 GENTIANALES. Apocynaceae, Asclepiadoids. **A–G.** *Asclepias tuberosa*. **A.** In orescence, an umbelliform cyme. **B.** Flower, side view. **C.** Gynostemium close-up, showing hood and horn appendages and one pollinium (of ve). **D.** Pollinium, inside sheath of tissue, with protruding, glandular corpusculum. **E.** Pollinium removed, showing retinacula (arms) and connate pollen grains. **F.** Flower longitudinal section, showing two distinct ovaries and styles and a single stigma. **G.** Close-up of ovaries in longitudinal section. **H.** *Sarcostemma cynanchoides*, owers. **I, J.** *Gomphocarpus physocarpus*. **I.** Umbelliform cyme. **J.** Fruits, bladderlike schizocarpic follicles. **K–M.** Succulent taxa. **K.** *Stapelia gigantea*, producing a foul odor and y-pollinated. **L.** *Stapelia variegata*. **M.** *Huernia* sp.

(periwinkle); medicinal uses, such as *Catharanthus roseus* (Madagascar periwinkle), from which vincristine/vinblastine used to treat childhood leukemia, and *Rauvolfia serpentina*, from which the drug reserpine is derived; and uses as timber, fiber, rubber, dye, and poison plants. See Endress and Bruyns (2000) for an updated classification of the family.

The Apocynaceae are distinctive in being lianas, trees, shrubs, or herbs with a 5-merous perianth/androecium, the gynoecium usually with 2 carpels, the *ovaries distinct in some taxa with stigmas connate* (in Asclepiadoids *androecium adnate to single stigma forming a gynostegium* and pollen fused to form *pollinia*, each half derived from an adjacent anther), the fruits variable, but a *schizocarp of follicles* in the Asclepiadoids.

K (5) **C** (5) **A** 5 or (5) **G** (2) [(-8)], superior, rarely half-inferior.

Gentianaceae Gentian family (after Gentius, king of Illyria). 78 genera / 1225 species. (Figure 8.62)

The Gentianaceae consist of trees, shrubs, or usually herbs. The **stems** of tree species are sometimes pachycaulous, and some taxa are achlorophyllous and mycotrophic (obtaining nutrition from mycorrhizal fungi in the soil). The **leaves** are simple (leaves scalelike in mycotrophic species), opposite (rarely whorled or spiral), and exstipulate. The **inflorescence** is a cyme, raceme, or of solitary flowers. The **flowers** are usually bisexual and actinomorphic, and hypogynous. The **calyx** is usually synsepalous with 4-5 [12], imbricate or valvate lobes. The **corolla** is sympetalous, rotate to salverform, with 4-5 [12], usually convolute lobes. The **stamens** are 4-5 [12], alternipetalous, epipetalous, with staminodes present in some. **Anthers** are usually longitudinal (rarely poricidal) in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 1, rarely 2 locules. The **style** is solitary and terminal; the **stigma** is solitary, 2-lobed, or decurrent. **Placentation** is parietal with placentae sometimes protruding and branched, rarely axile or free-central; **ovules** are anatropous, unitegmic, numerous. **Nectaries** are often present as pits on corolla lobes, with a nectariferous disk or glands usually at the ovary base. The **fruit** is a septicidal capsule, rarely a berry. The **seeds** are oily endospermous.

The Gentianaceae have a mostly worldwide distribution. Economic importance consists primarily of ornamental cultivars such as *Eustoma*, *Exacum*, *Gentiana* (gentian), and *Sabatia*; some species are used medicinally or as timber.

The Gentianaceae are distinctive in being trees, shrubs, or *usually herbs* (sometimes achlorophyllous and mycotrophic), with a usually **4-5 [-12]-merous perianth/androecium**, often with *nectary pits on corolla lobes* and a *disk or glands at ovary base*, the ovary superior, 2-carpellate, with *usually parietal* (sometimes axile or free-central) placentation, the fruit a *septicidal capsule or berry*.

K (4-5) [(-12)] **C** (4-5) [(-12)] **A** 4-5 [-12] **G** (2), superior.

Rubiaceae Coffee family (after *rubia*, name used by Pliny for madder). 630 genera / 10,200 species. (Figure 8.63)

The Rubiaceae consist of terrestrial (rarely epiphytic or aquatic) trees, shrubs, lianas, or herbs. The **leaves** are simple, undivided and entire, usually decussate, rarely whorled or spiral by suppression, stipulate, stipules of opposite leaves connate, often bearing structures termed colleters, which produce mucilaginous compounds protecting the young shoot. The **inflorescence** is a cyme, rarely of solitary flowers. The **flowers** are usually bisexual and usually epigynous. The **perianth** is usually biseriolate, although the calyx is lost in some taxa. The **calyx** is synsepalous with 4-5 or 0 lobes. The **corolla** is sympetalous with 4-5 [rarely 3 or 8-10], actinomorphic or bilabiate lobes. The **stamens** are 4-5 [rarely 3 or 8-10], alternipetalous and epipetalous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with a usually inferior [rarely superior] ovary, 2 [3-5+] carpels, and 1-2 [3-5+] locules. **Placentation** is axile, rarely parietal; **ovules** are anatropous to hemitropous, unitegmic, with a funicular obturator, 1 ∞ per carpel. **Nectaries** are often present as a nectariferous disk atop ovary. The **fruit** is a berry, capsule, drupe, or schizocarp. The **seeds** are usually endospermous.

The Rubiaceae have a mostly worldwide distribution, more concentrated in tropical regions. Economic importance includes *Cinchona*, the source of quinine used to treat malaria, *Coffea arabica* and other species, the source of coffee, *Pausinystalia johimbe*, the source of the sexual stimulant yohimbine, some timber trees, fruiting plants, dye plants (such as *Rubia*, madder), and ornamental cultivars (e.g., *Pentas*, among others).

The Rubiaceae are distinctive in being trees, shrubs, lianas, or herbs with *simple, entire, usually decussate leaves* and *connate stipules*, the stipules often with mucilage-secreting *colleters*, the *inflorescence usually a cyme*, flowers usually bisexual, the perianth dichlamydeous, perianth and androecium often 4-5-merous (calyx absent in some), the *ovary usually inferior* (rarely superior), often with an apical *nectariferous disk*, ovules with a *funicular obturator*, the fruit a berry, capsule, drupe, or schizocarp.

K (4-5) [0] **C** (4-5) [(3,8-10)] **A** 4-5 [3,8-10] **G** (2) [(3-5+)], usually inferior, rarely superior.

LAMIALES

The Lamiales, sensu APG II (2003), contain approximately 21 (23) families (Table 8.3), many of which have undergone considerable changes in classification (e.g., see Scrophulariaceae s.l., discussed later). Seven families are described here.

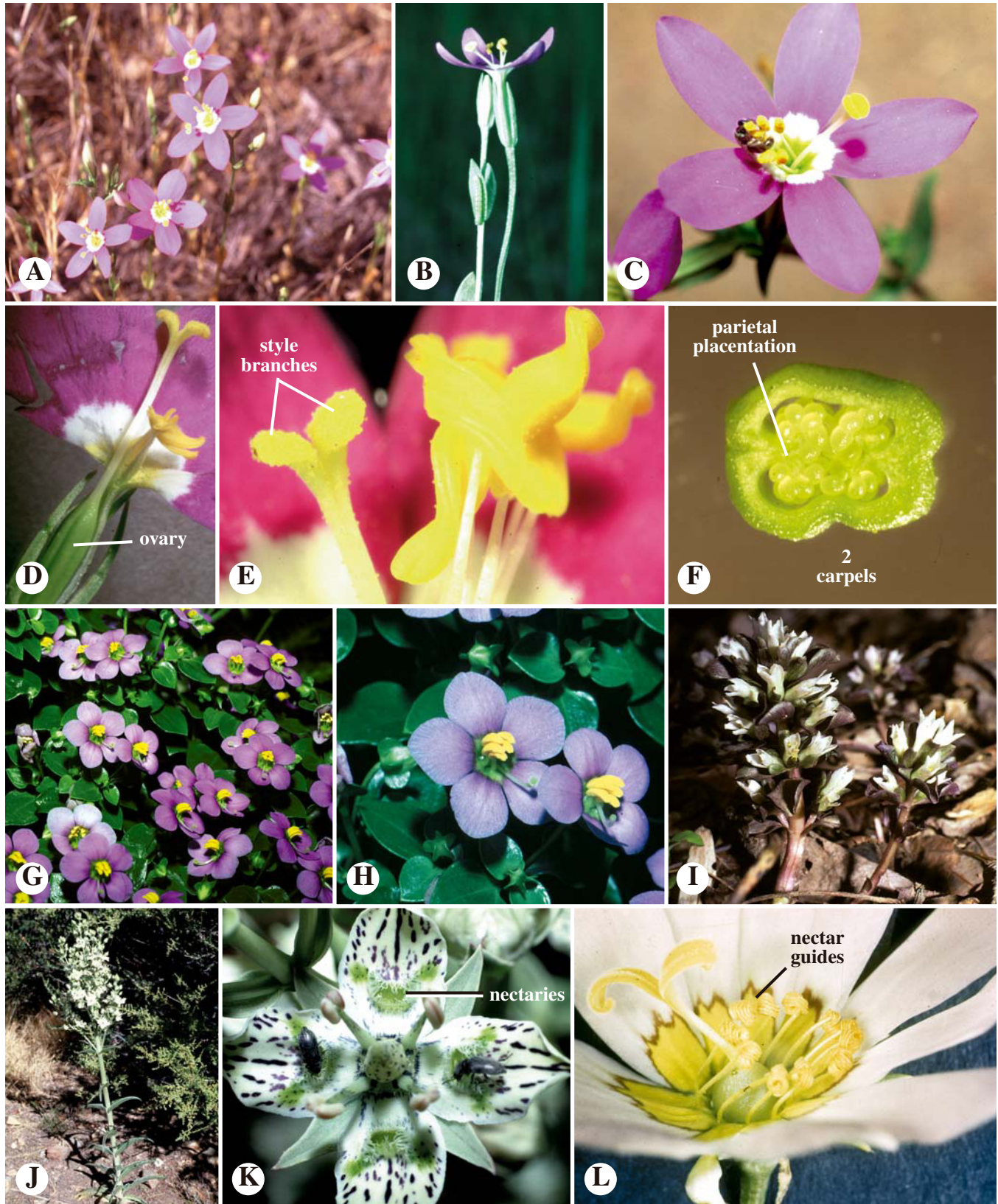


FIGURE 8.62 GENTIANALES. Gentianaceae. A–F. *Centaurium venustum*. A, B. Whole plants. C. Flower close-up. D. Flower in longitudinal section. E. Close-up of coiled stamens and style. F. Ovary cross-section, showing parietal placentation. G, H. *Exacum af ne.* I. *Obolaria virginica*, an herb. J, K. *Swertia parryi*. J. Whole plant. K. Flower close-up, showing corolla nectaries. L. *Sabatia dodecandra*, a species with multiple perianth and androecial parts.

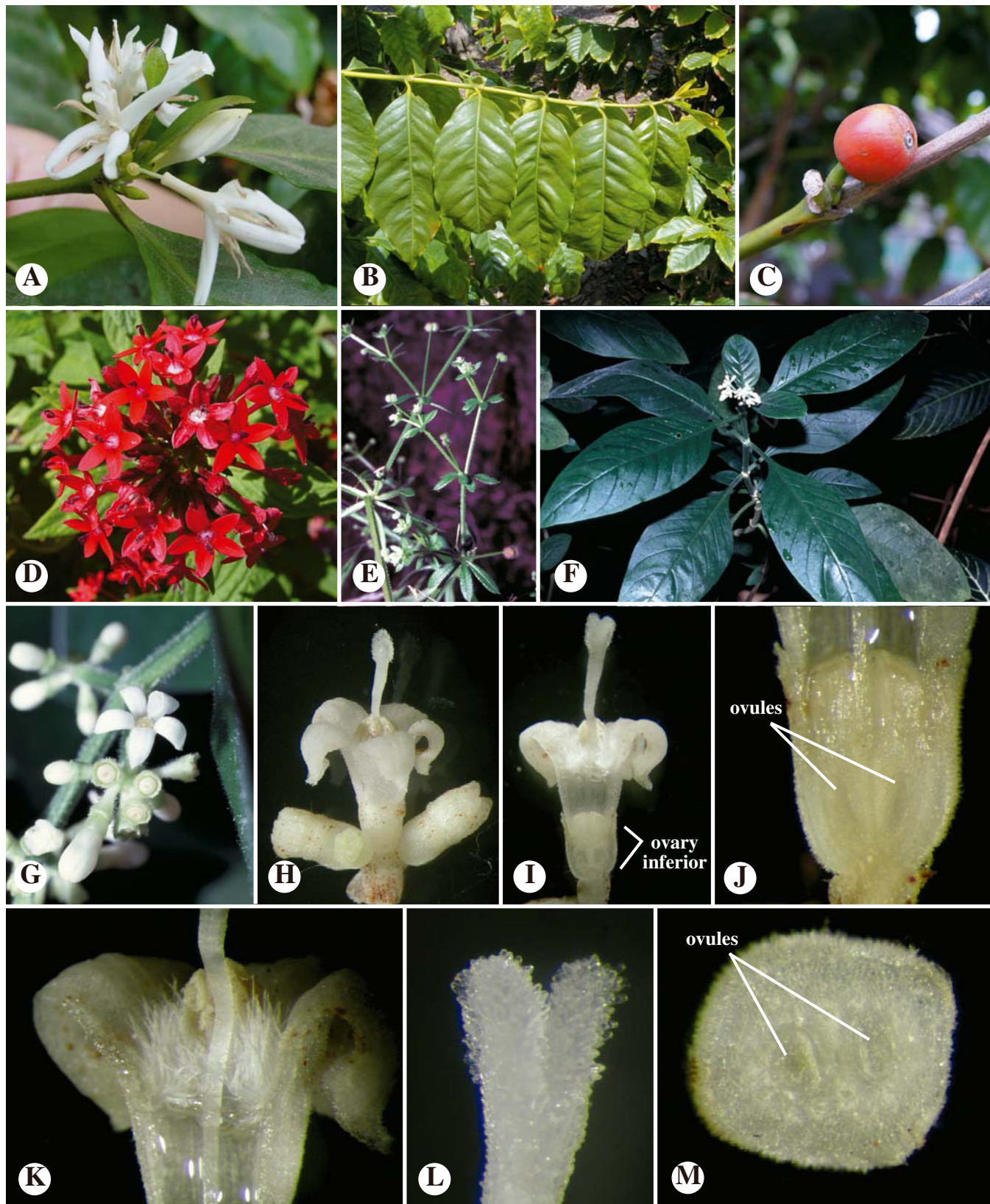


FIGURE 8.63 GENTIANALES. Rubiaceae. A–C. *Coffea arabica*, coffee. A. Flowers. B. Leaves; note opposite arrangement. C. Fruit, a drupe. D. *Pentas* sp. E. *Galium aparine*, bedstraw. F, G. *Psychotria kirkii*. H–M. *Psychotria viridi ora*, ower morphology. H. Simple dichasium. I. Flower longitudinal section, showing inferior ovary. J. Ovary, showing two ovules, with axile placentation. K. Apex of corolla tube, showing style. L. Stigmas. M. Ovary cross-section; note two carpels and locules.

Among those not described are the **Gesneriaceae** (a large family including many cultivated ornamentals, such as *Saintpaulia*, African violets, and *Sinningia*, gloxinia), **Lentibulariaceae** (containing the interesting carnivorous plants *Pinguicula*, butterwort, and *Utricularia*, bladderwort), **Oleaceae** (the olive family, containing olive, *Olea europaea*), **Pedaliaceae** (source of *Sesamum*, sesame), and **Verbenaceae** (the verbena family). See Kadereit (2004) for more information on classification in the order.

Acanthaceae Acanthus family (from Acanthus, prickly-one). 229 genera / 3450 species. (Figure 8.64)

The Acanthaceae consist of terrestrial or aquatic herbs, shrubs, or rarely trees. The **leaves** are opposite (usually) and simple. The **inflorescence** is a cyme, raceme, or of solitary flowers. The **flowers** are bisexual, zygomorphic, bracteate and bracteolate (the bracts often colored), and hypogynous. The **perianth** is biseriate and dichlamydeous, with hypanthium absent. The **calyx** is synsepalous with 5 [4,6], imbricate or valvate lobes. The **corolla** is sympetalous and usually bilabiate (the upper lip suppressed in some species) with 4–5 imbricate or convolute lobes. The **stamens** are 2, 4, or rarely 5, with staminodes present in some. **Anthers** are tetrasporangiate or bisporangiate, dithecal or monothechal, with parallel or divergent thecae. The **pollen** is tricolpate, triporate, diporate, pantoporate, or inaperturate. The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 2 locules. The **style** is solitary and terminal. **Placentation** is axile; **ovules** are variable in type, 2 ∞ per carpel. Nectaries are usually present as a disk at the ovary base. The **fruit** is usually an explosively dehiscent, loculicidal capsule. The seeds have funiculi that are modified into rigid, often hook-shaped structures that function to catapult the seeds; these are termed funicular retinacula or jaculators. Cystoliths are characteristic of some taxa, appearing as streaks in the leaves.

Members of the Acanthaceae are distributed from the tropics to temperate regions. Economic importance includes several cultivated ornamentals, such as *Acanthus mollis*, *Aphelandra*, and *Justicia* [including *Beloperone*]. For information about intrafamilial relationships, see McDade et al. (2000).

The Acanthaceae are distinctive in having simple, **opposite leaves** with **zygomorphic, bracteate**, usually **bilabiate flowers**, the fruit an explosively dehiscent, loculicidal capsule with distinctive **funicular retinacula** (jaculators) that function in seed dispersal, the funicular retinacula a presumed apomorphy of the family.

K (5) [(4,6)] **C** (4–5) **A** 2,4, or 5 **G** (2), superior.

Bignoniaceae Bignonia family (after AbbØ Jean-Paul Bignon, 1662–1743, court librarian at Paris, friend of Tournefort). 109 genera / 750 species. (Figures 8.65, 8.66)

The Bignoniaceae consist of trees, shrubs, and lianas, rarely herbs. The **leaves** are usually pinnate or ternate, less often simple or palmate, usually opposite, sometimes whorled, rarely simple, exstipulate, the terminal leaflets modified as tendrils in some taxa. The **inflorescence** is a cyme, raceme, or of solitary flowers. The **flowers** are bisexual, zygomorphic [corolla rarely actinomorphic], hypogynous. The **calyx** is synsepalous with 5 zygomorphic, often bilabiate lobes [sometimes unlobed or spathaceous, rarely calyprate]. The **corolla** is sympetalous with 5, usually bilabiate lobes [rarely actinomorphic]. The **stamens** are alternipetalous, whorled, usually didynamous, 2+2 or 2+2 + 1 staminode [rarely 2 + 3 staminodes]. The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 1, 2, or 4 locules. The **style** is solitary and terminal, with two **stigmas**. **Placentation** is axile or parietal with intruding septae [false septa dividing each carpel into two locules in some]; **ovules** are anatropous or hemitropous, unitegmic, numerous. **Nectaries** are usually present as a ring or cup-shaped structure around ovary base. The **fruit** is a two-valved capsule, rarely fleshy or fibrous and indehiscent. The **seeds** are usually flat and winged in taxa with capsules, exalbuminous. The stem anatomy of members of the family with lianas is unique.

Members of the Bignoniaceae are distributed primarily in tropical [some temperate] regions. Economic importance includes important timber trees and many ornamental cultivars (e.g., *Jacaranda*, *Spathodea*). See Spangler and Olmstead (1999) and Olmstead et al. (2002) for recent phylogenetic studies of the family.

The Bignoniaceae are distinctive in being trees, shrubs, or vines with **opposite leaves** and usually **zygomorphic**, often **bilabiate**, flowers with **didynamous** stamens, a superior, 2-carpellate ovary having **axile or parietal placentation** with numerous ovules, the fruit a **capsule** [rarely indehiscent] with usually **flat, winged, exalbuminous** seeds.

K (5) **C** (5) **A** 2+2 [+1 staminode in some; rarely 2 fertile +3 staminodes] **G** (2), superior.

Lamiaceae (Labiatae) — Mint family (*lamium*, gullet, after the shape of the corolla tube or old Latin name used by Pliny). 251 genera / 6700 species. (Figures 8.67, 8.68)

The Lamiaceae consist of hermaphroditic, sometimes gynodioecious, herbs, shrubs, or rarely trees, often with short-stalked glandular trichomes producing aromatic etheral oils. The **stems** are usually 4-sided (square in cross-section), at least when young. The **leaves** are simple [rarely pinnate], opposite, sometimes whorled [rarely spiral], exstipulate. The **inflorescence** consists of lateral cyme units in a verticillaster or thyrses, or of solitary, axillary flowers. The **flowers** are bisexual [rarely unisexual], mostly zygomorphic, bracteate and bracteolate, hypogynous. The **perianth** is biseriate and

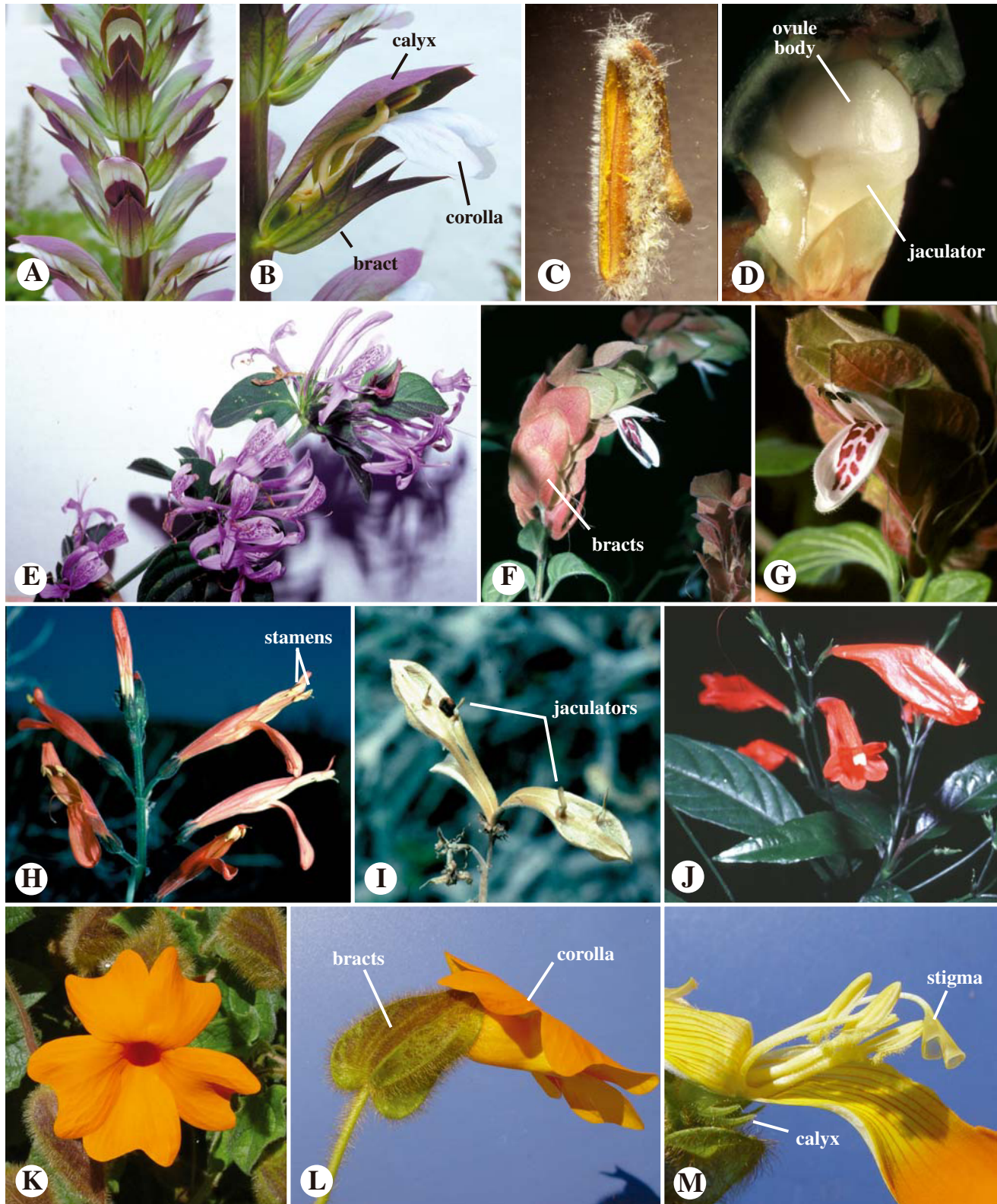


FIGURE 8.64 LAMIALES. Acanthaceae. A–D. *Acanthus mollis*. A. In inflorescence. B. Flower close-up, showing reduced corolla. C. Anther close-up. D. Ovule and jaculator. E. *Hypoestes aristata*, in inflorescence. F, G. *Justicia brandegeana*, shrimp plant. F. In inflorescence, with showy bracts. G. Flower close-up, showing bilabiate corolla. H, I. *Justicia californica*. H. In inflorescence. I. Open fruit, showing peglike jaculators. J. *Ruellia graecizans*. K–M. *Thunbergia gregorii*. K. Flower, face view. L. Flower, side view, showing basal, spathe-like bracts. M. Open flower. Note calyx, stamens, and style/stigma.

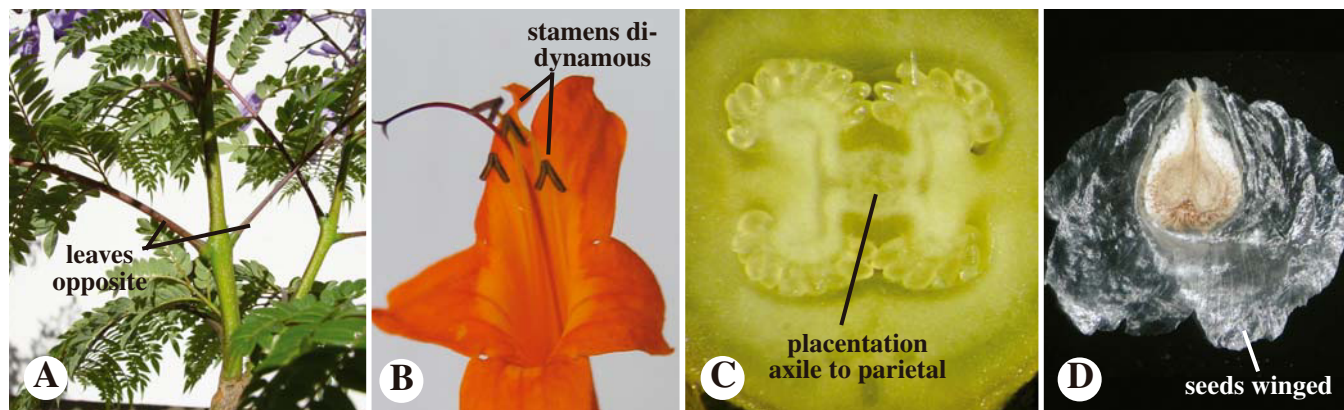


FIGURE 8.65 LAMIALES. Bignoniaceae. Diagnostic characteristics. **A.** Opposite leaf arrangement (*Jacaranda mimosifolia*). **B.** Didynamous stamen arrangement (*Tecomania capensis*). **C.** Axile to parietal placentation (*Spathodea campanulata*). **D.** Winged seeds (*S. campanulata*).

dichlamydeous, the corolla usually bilabiate, sometimes actinomorphic, hypanthium absent. The **calyx** is synsepalous of 5 zygomorphic, sometimes bilabiate, lobes. The **corolla** is sympetalous, of 4 or 5 corolla lobes (if 4, by fusion of two lobes). The **stamens** are 2, 4, or 2 fertile + 2 staminodes, whorled, epipetalous (adnate to corolla tube). **Anthers** are longitudinal in dehiscence, with connective split in *Salvia* and relatives, separating the thecae of anthers (one theca lost in some taxa). The **gynoecium** is syncarpous, with a superior ovary, 2 carpels, and 4 locules. The **style** is solitary, often apically 2-branched, terminal or gynobasic with the ovary deeply 4-lobed by formation of false septa dividing each carpel; **stigmas** are usually 2. **Placentation** is basal; **ovules** are anatropous to hemitropous, unitegmic, 2 per carpel, 1 per locule. **Nectaries** are usually present as a disk or pad of tissue at base of ovary. The **fruit** is a schizocarp of usually four [1 3] nutlets, a drupe, or a berry. Plants often have ethereal oils and stachyose carbohydrate.

The Lamiaceae have a mostly worldwide distribution. Economic importance includes medicinal plants, culinary herbs (e.g., *Mentha*, mint; *Ocimum*, basil; *Rosmarinus*, rosemary; *Salvia*, sage; *Thymus*, thyme), fragrance plants (e.g., *Lavandula*, lavender; *Pogostemon*, patchouli), and a number of cultivated ornamentals. See Harley et al. (2004) and references therein for treatments of the family.

The Lamiaceae are distinctive in being herbs or shrubs, **often aromatic with ethereal oils**, with usually **4-sided stems**, **opposite** [or whorled] leaves, a **verticillaster or thyrs**e inflorescence [flowers solitary and axillary in some], and zygomorphic [rarely actinomorphic], **usually bilabiate** flowers having a superior ovary, often **deeply 4-lobed** (by formation of false septa) with a **gynobasic style**, the fruit a **schizocarp of usually 4 nutlets** or a berry or drupe.

K (5) **C** (5) [(4)] **A** 4 or 2 [+2 staminodes] **G** (2), superior.

SCROPHULARIACEAE S.L.

The Scrophulariaceae s.l. has traditionally been characterized as shrubs or herbs with usually opposite leaves and mostly pentamerous, zygomorphic flowers, the stamens usually 2 or 4 (sometimes 5 or with staminodes), the ovary superior and usually 2-carpellate with axile or parietal placentation with numerous ovules, the fruit a capsule, berry, or schizocarp. The floral formula for this family as traditionally defined can be summarized as:

K (4-5) [(2,3)] **C** (5) [0,(4-8)] **A** 2,4,5 [some having 2 or 4 fertile stamens with one or more staminodes] **G** (2) [(-3)], superior.

Recent molecular studies demonstrate that the Scrophulariaceae as traditionally defined is not monophyletic. Several major clades that include at least some taxa that were formerly placed in the Scrophulariaceae s.l., are now recognized at the family rank (Olmstead et al. 2001; Oxelman et al. 2005; see Stevens, 2001 onwards): Calceolariaceae, Gratiolaceae, Linderniaceae, Orobanchaceae (Figure 8.70), Pawloniaceae, Phrymaceae (Figure 8.72), Plantaginaceae (Figure 8.71), Scrophulariaceae s.s. (Figure 8.69), and Stilbaceae. These families are difficult to diagnose and differentiate from one another morphologically. Their phylogenetic boundaries are still somewhat tentative, and they have yet to be characterized and thoroughly circumscribed in terms of number of genera and species. Four of these newly defined families are summarized briefly here, but only tentatively and superficially. See Fischer (2004) for a general treatment of the complex.

Scrophulariaceae s.s. [including Loganiaceae and Myoporaceae] Figwort family (after Scrophula, from glands on the corolla). (Figure 8.69)



FIGURE 8.66 LAMIALES. Bignoniaceae. **A.** *Campsis radicans*, a vine. **B.** *Chilopsis linearis*, desert-willow, flowers. **C,D.** *Distictis buccinatoria*, a vine. **C.** Flowers. **D.** Corolla, opened to reveal didynamous stamens and style/stigmas. **E–G.** *Kigelia africana*, African sausage tree. **E.** Flower. **F.** Flower with corolla removed and calyx dissected, showing large nectaries. **G.** Fruit, a rare indehiscent capsule. **H,I.** *Jacaranda mimosifolia*, a tree. **H.** Opened fruit, showing winged seeds. **I.** Flower, close-up, with bilabiate corolla. **J–L.** *Markhamia* sp., a vine. **J.** Flower. **K.** Dehiscent fruit, with persistent septum (replum). **L.** Winged seeds. **M,N.** *Spathodea campanulata*, a tree. **M.** Inflorescence. **N.** Flower close-up, showing spathe-like calyx and large corolla. **O.** *Pyrostegia venusta*, a vine.



FIGURE 8.67 LAMIALES. Lamiaceae, diagnostic features. **A.** *Salvia leucantha*, showing opposite leaf arrangement. **B.** *Marrubium vulgare*, unit in orescence of verticillaster. **C.** *Orthosiphon*, showing thyrse. **D.** *Rosmarinus of cinalis*, flower with zygomorphic, bilabiate corolla. **E.** *Hemiantra pungens*, showing four-lobed ovary and gynobasic style. **F,G.** *Rosmarinus of cinalis*. **F.** Style, 2-branched at apex. **G.** Fruit, showing two of four schizocarpic nutlets. **H.** *Acanthomintha ilicifolia*, an endangered species, with solitary, axillary flowers. **I.** *Hemiantra pungens*, corolla. **J.** *Hyptis emoryi*, in orescence. **K,L.** *Monarda stulosa*, congested headlike verticillaster, with showy bracts. **M.** *Monarda didyma*, bee-balm.

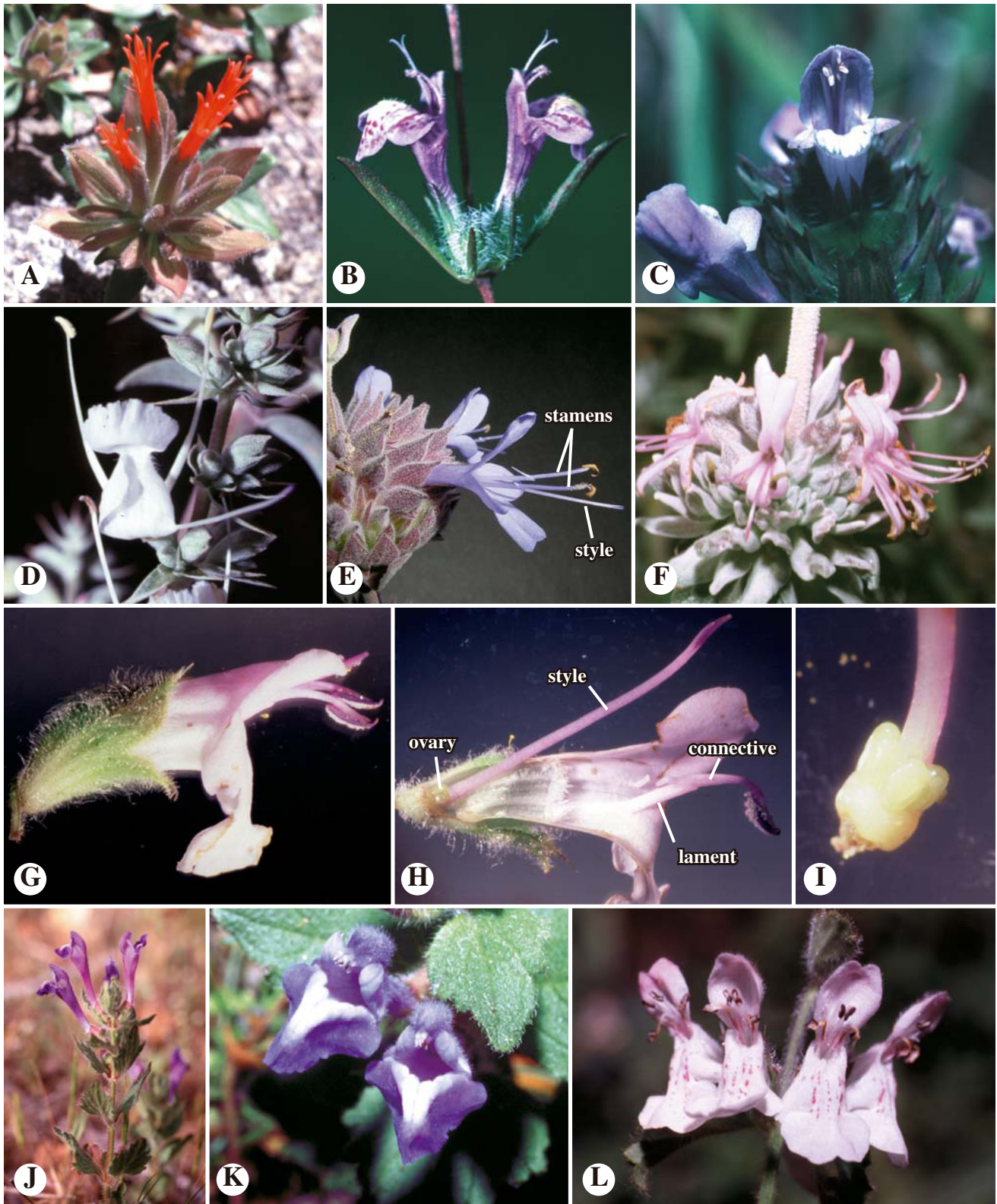


FIGURE 8.68 LAMIALES. Lamiaceae, diversity. **A.** *Monardella macrantha*. **B.** *Pogogyne abramsii*, with thyrse reduced to solitary, axillary flowers. **C.** *Prunella vulgaris*. **D.** *Salvia apiana*, having enantiostyly. **E.** *Salvia clevelandii*. **F.** *Salvia leucophylla*. **G–I.** *Salvia mellifera*. **G.** Flower, side view. **H.** Flower, longitudinal section, showing gynoeceium and epipetalous stamens, with unusual, extended connective and monotheal anthers. **I.** Ovary, four-lobed with gynobasic style. **J,K.** *Scutellaria tuberosa*. **L.** *Stachys ajugoides*, with strongly bilabiate corolla.

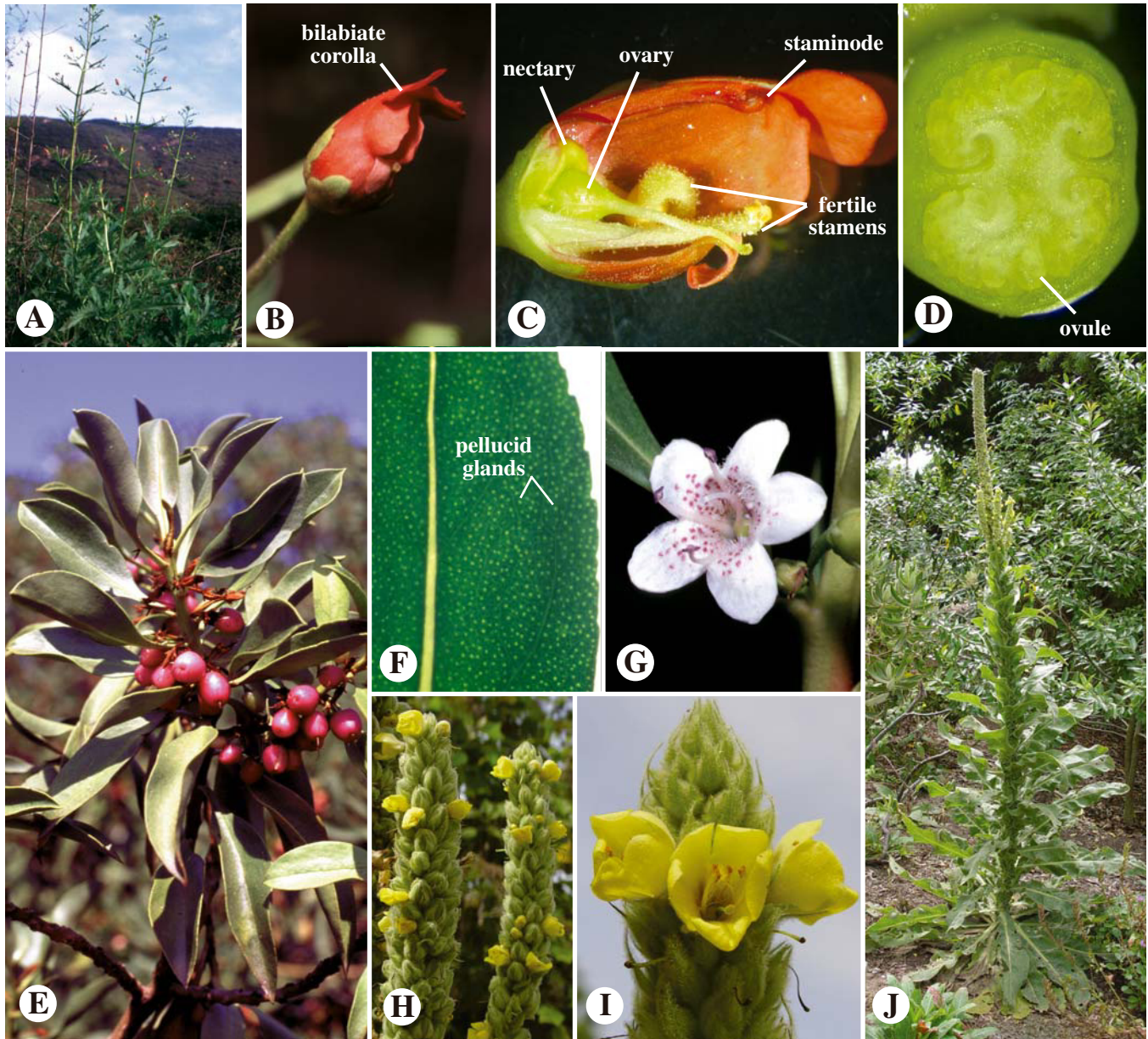


FIGURE 8.69 LAMIALES. Scrophulariaceae s.s. **A–D.** *Scrophularia californica*. **A.** Whole plant. **B.** Flower close-up, showing bilabiate corolla. **C.** Flower longitudinal section; note superior ovary. **D.** Ovary cross-section, with axile placentation. **E–G.** Myoporoids. *Myoporum laetum*. **E.** Branch of tree, in fruit (a berry). **F.** Leaf close-up showing glands. **G.** Flower close-up. **H–J.** *Verbascum thapsus*. **H.** Inflorescence, a spike. **I.** Flower close-up. **J.** Whole plant.

This clade, containing the type genus *Scrophularia*, also includes members of several subgroups of the traditionally defined Scrophulariaceae s.l., such as *Verbascum* and *Selago*. No morphological characteristics clearly differentiate this newly circumscribed family from other groups within the traditional, larger one. The Scrophulariaceae s.s. subsumes at least some members of other former families, including

the Buddlejaceae (when recognized), Loganiaceae, and Myoporaceae. These latter two families might logically be treated as subfamilies [Loganioideae and Myoporoideae, respectively] within the Scrophulariaceae s.s., when further systematic studies are done.

The **Loganioids** [Loganiaceae] as traditionally defined contain approximately 29 genera and 570 species of trees,

shrubs, herbs, or lianas, with usually opposite, simple leaves. The flowers are rather typical of the Scrophulariaceae s.l. in having a 4-5-merous perianth and androecium, a mostly superior ovary with 2 [1-4] carpels, mostly axile placentation with 2-∞ ovules, the fruit a capsule, berry, or drupe. This group is rather disparate with a varying history of classification, and much more additional work will be necessary to firmly place its members.

The **Myoporoids** [Myoporaceae] contain 3 genera and ca. 235 species. They can be distinguished from other members of the Scrophulariaceae s.s. in having simple, **gland-dotted, usually spirally arranged leaves**. This group, which is fairly well delimited, is largely Australian, with other taxa occurring primarily in lands bordering the Pacific or Indian oceans.

Orobanchaceae Broom-Rape family (Greek for the gummy strangler, after the parasitic habit of most). (Figure 8.70)

The Orobanchaceae consist almost entirely of root parasites (Olmstead et al. 2001). Included within this group are the more traditionally delimited Orobanchaceae s.s., which contains achlorophyllous root parasites. Also included are members of the traditional Scrophulariaceae s.l. such as *Castilleja*, which are chlorophyllous root hemiparasites. Economic importance of the Orobanchaceae is minimal, with some species being a local source of food.

The traditionally defined Orobanchaceae s.s. might logically be treated as a subfamily of this expanded treatment of the family. The former are termed here as Orobanchoids and are briefly described next.

The **Orobanchoids** [Orobanchaceae s.s.] contain approximately 15 genera and 210 species with a mostly northern hemisphere distribution. They are distinctive in being **achlorophyllous root-parasites**, with often fleshy stems and scale-like, usually spiral leaves. The flowers are mostly pentamerous, with a **sympetalous, bilabiate corolla**, typically with 4 stamens, a posterior staminode present in some, the ovary **superior, 2-carpellate, often with parietal placentation**.

Phrymaceae s.l. — Hopseed family (after an early genus name). ca. 10 genera / ca. 190 species. (Figure 8.71)

The Phrymaceae s.l. consist of annual or perennial herbs or shrubs. The **leaves** are simple, opposite, exstipulate. The **inflorescence** is variable, including a spike or of solitary, axillary flowers. The **flowers** are bisexual, zygomorphic or actinomorphic, ebracteate, hypogynous. The **perianth** is dichlamydeous. The **calyx** is synsepalous with (5) [(3,4)] teeth-like lobes. The **corolla** is sympetalous with 5, usually bilabiate lobes, the upper lip 2-lobed. The **stamens** are 4, alternipetalous and epipetalous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is unicarpellous or syncarpous,

with a superior ovary, 2 carpels, (appearing as 1 in *Phryma*, but stigma 2-lobed) and 1 or 2 locules. The **style** is solitary, terminal; **stigma(s)** are 2 or 2-lobed, bilamellate and thigmognastic (functioning in pollination) in some taxa. **Placentation** is axile, parietal, or basal; **ovules** are anatropous, unitegmic, 1-∞ per carpel. The **fruit** is a dehiscent capsule (opening late in some taxa), an achene with persistent calyx (*Phryma*), or a berry.

The Phrymaceae s.l. are recently expanded from a single species (*Phryma leptostachya*) to approximately 190 species, including the large genus *Mimulus*. See Beardsley and Olmstead (2002) and Cantino (2004). Members have a mostly worldwide distribution, especially in western North America. Economic importance includes some cultivars in *Mimulus*.

The Phrymaceae s.l. are distinctive in being herbs or shrubs with **simple, opposite** leaves, zygomorphic or actinomorphic **synsepalous and sympetalous** flowers, a **superior, 1- or 2-carpellate** ovary, the fruit a **capsule, achene, or berry**.

K (5) [(3,4)] **C** (5) **A** 4 **G** (2), superior.

Plantaginaceae s.l. (including Callitrichaceae, Globulariaceae, Halorigidaceae, Plantaginaceae s.s., and many members of the traditional Scrophulariaceae s.l.) Plantain/Speedwell family (L. for sole of the foot, after the leaves of some species that lie flat on the ground) (Figure 8.72)

The Plantaginaceae, as delimited by Olmstead et al. (2001) (which they tentatively termed the Veronicaceae, a name that is illegitimate, however) contains many taxa that have traditionally been placed in the Scrophulariaceae s.l., including *Antirrhinum* (snapdragons), *Chelone* (turtleheads), *Collinsia* (Chinese houses), and *Digitalis* (foxglove). Many members of the Plantaginaceae are important in the horticultural trade, e.g., *Antirrhinum*. *Digitalis* is both horticulturally and medically important, being the source of the cardiac glycoside digitoxin and others, used to treat heart ailments.

The Plantaginaceae as a whole cannot be readily delimited from other Scrophulariaceae s.l. based on morphological characteristics. Within the Plantaginaceae are clades that have previously been treated as the separate families Callitrichaceae, Globulariaceae, Halorigidaceae, and Plantaginaceae s.s. It might be logical to treat some of these as subfamilies within the Plantaginaceae, but this awaits much more detailed studies. Two of these former families, the Callitrichaceae and Plantaginaceae s.s., are briefly described here, denoted simply as Callitrichoids and Plantaginoids.

The **Callitrichoids** [Callitrichaceae], commonly known as Starworts, contain 1 genus, *Callitriche*, and approximately 17 species. The Callitrichoids are distinctive in being aquatic, submerged to floating herbs with decussate leaves and unisexual flowers lacking a perianth, the stamens usually solitary,

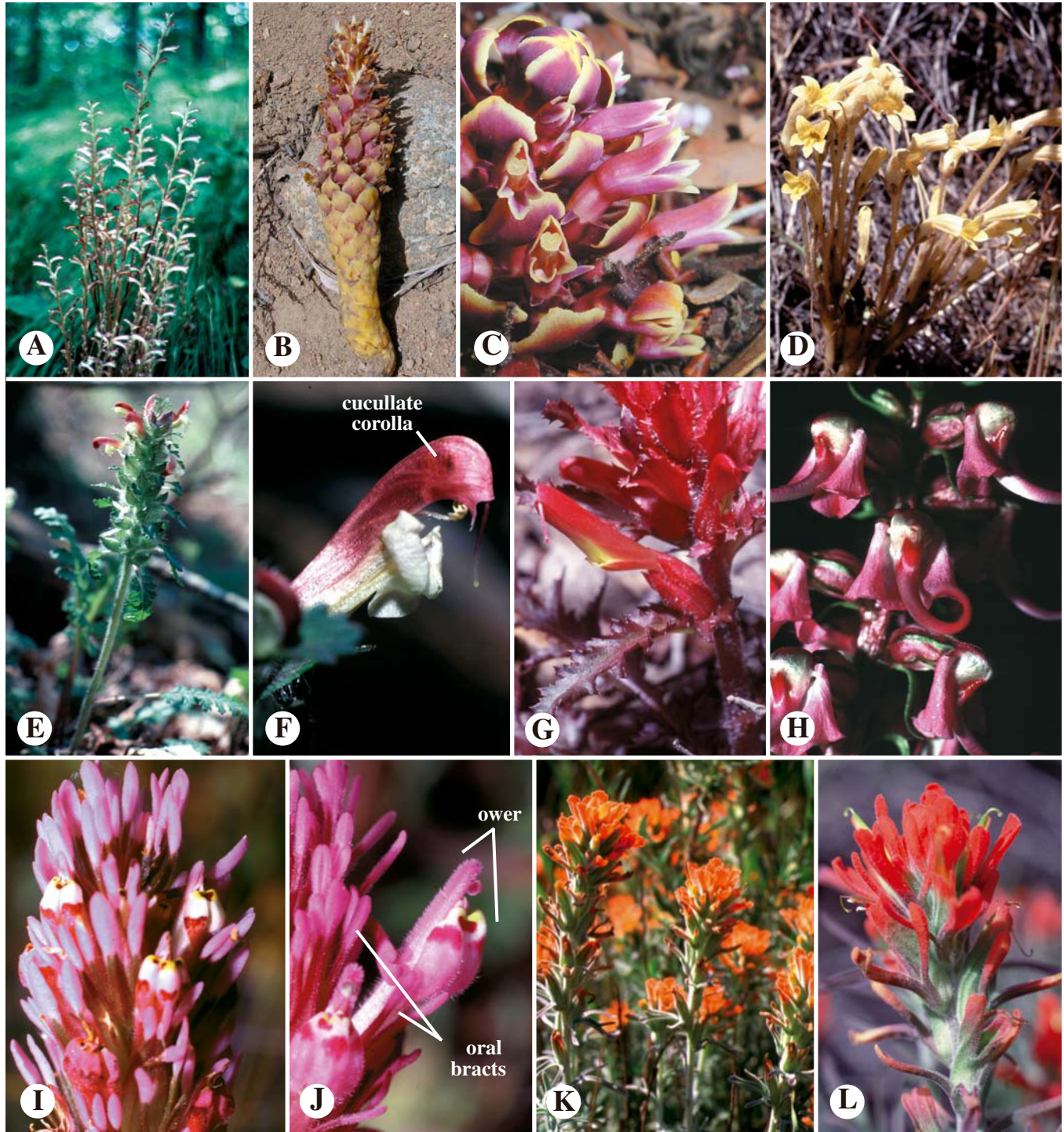


FIGURE 8.70 LAMIALES. Orobanchaceae. **A.** *Epifagus virginiana*, beech-drops. **B,C.** *Boschniackia strobilacea*. **B.** Whole plant (dug up). **C.** Flower close-ups. **D.** *Orobanche fasciculata*. **E,F.** *Pedicularis canadensis*. **E.** Whole plant. **F.** Flower close-up, showing bilabiate corolla. **G.** *Pedicularis densi ora*, owers. **H.** *Pedicularis groenlandica*, elephant heads. **I–L.** *Castilleja* spp., Indian paintbrush, hemiparasites with showy bracts. **I.** *C. densi ora*. **J.** *C. exserta*. **K,L.** *C. foliolosus*.

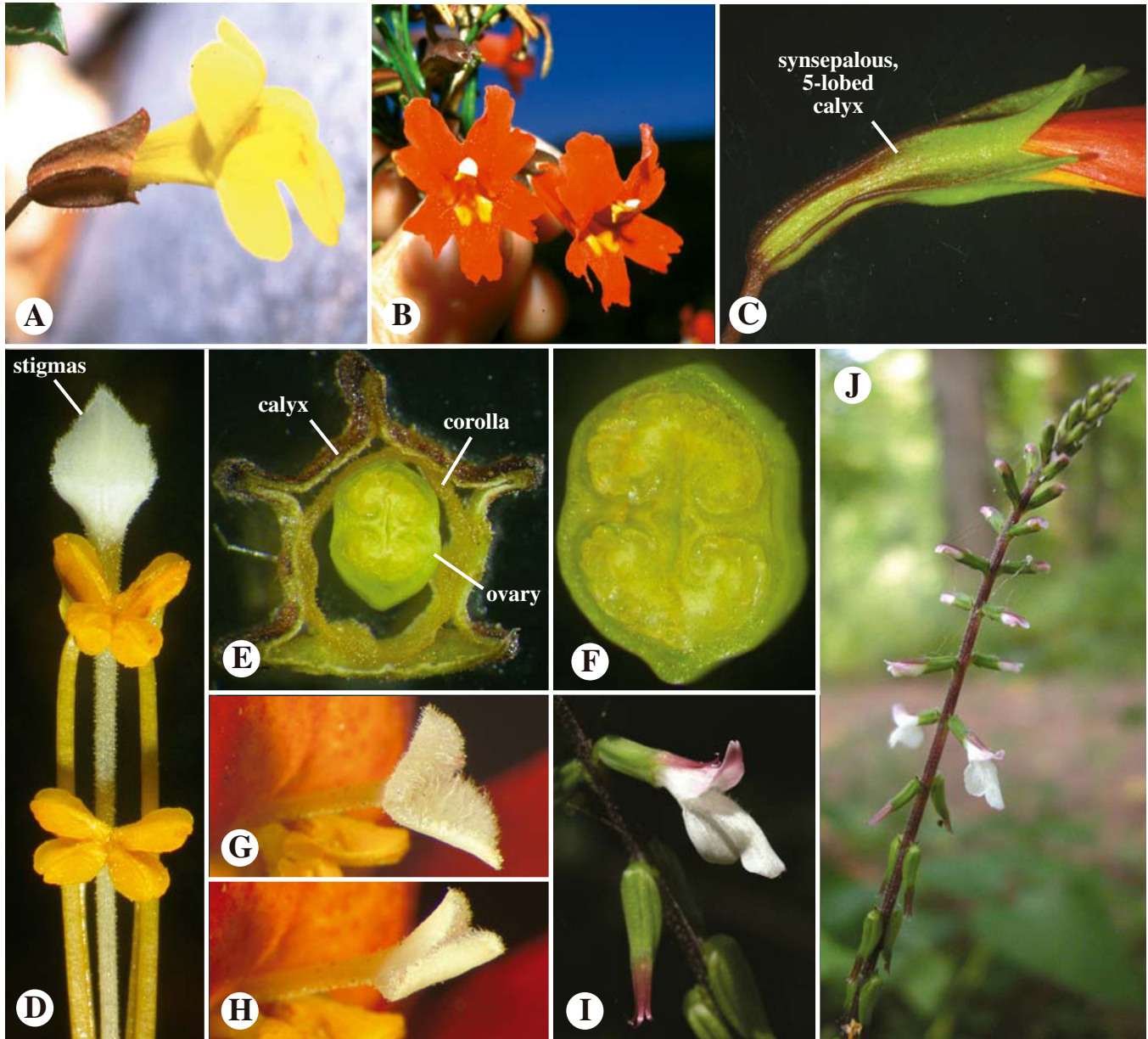


FIGURE 8.71 LAMIALES. Phrymaceae. **A.** *Mimulus guttatus*, seep monkey flower. **B–H.** *Mimulus aurantiacus*, coast monkey flower. **B.** Axillary, opposite flowers, face view. **C.** Winged calyx, side view. **D.** Androecium (didynamous) and style/stigma. **E.** Cross-section at base of flower, showing perianth and ovary. **F.** Ovary cross-section, with parietal placentation (septum incomplete). **G.** Bilamellate stigmas, open. **H.** Stigmas closed, 5 seconds after physical contact. **I, J.** *Phryma leptostachya*, hopseed; images courtesy of David G. Smith. **I.** Flower close-up, showing bilabiate corolla; note depressed orientation after flowering (below). **J.** Inflorescence, a spike.

the gynoecium of 2 carpels, each with 2 locules, and the fruit a schizocarp of 4 nutlets. The floral formula is: $\text{P } 0 \text{ A } 1 [2,3] \text{ G } (2)$, superior. Members of the Callitrichoids have a mostly worldwide distribution. Economic importance includes some species grown as aquarium plants.

The **Plantaginoids** [Plantaginaceae s.s.], commonly known as Plantains (not to be confused with the bananas

known as plantains) contain 3 genera and approximately 275 species, almost all of these in the genus *Plantago*. The Plantaginoids are distinctive in being herbs, rarely shrubs, the leaves spiral with parallel major veins, the flowers bisexual or unisexual, with 4 [3] connate, membranous to scarious sepals and petals, usually 4 [1–3] stamens, and a superior, 2-carpellate ovary, the fruit a circumscissile capsule or achene.

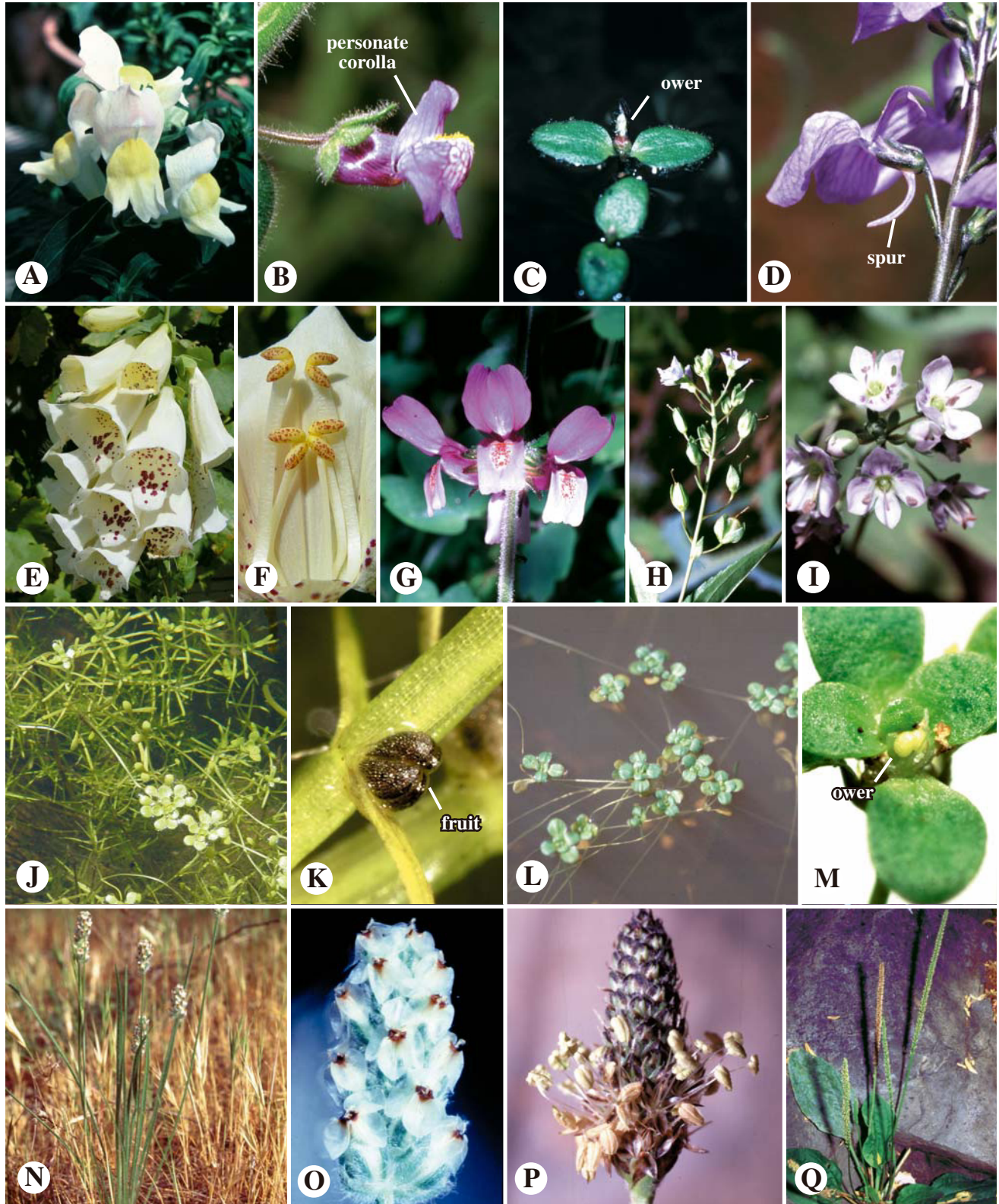


FIGURE 8.72 LAMIALES. Plantaginaceae. **A.** *Antirrhinum majus*, snapdragons. **B.** *Antirrhinum nuttallianum*. **C.** *Amphianthus pusillus*, an aquatic species. **D.** *Linaria canadensis*. **E,F.** *Digitalis purpurea*, foxglove, source of digitoxin. **G.** *Collinsia heterophylla*, Chinese houses. **H,I.** *Veronica anagallis-aquatica*. **J–M.** Callitrichoids. **J,K.** *Callitriche heterophylla*. **J.** Note dimorphic leaves, floating and submerged. **K.** Fruit close-up. **L,M.** *Callitriche marginata*. **L.** Floating shoots. **M.** Flower close-up. **N–Q.** Plantaginoids. **N,O.** *Plantago erecta*. **N.** Whole plant. **O.** Inflorescence. **P.** *Plantago lanceolata*, inflorescence, with exserted stamens. **Q.** *Plantago major*, whole plant.

The floral formula is **K** (4) [(3)] **C** (4) [(3)] **A** 4 [1-3] **G** (2), superior. Members of the Plantaginoids are worldwide in distribution. Economic importance includes weedy species, medicinal plants, and food plants. See Schwarzbach (2004).

SOLANALES

The Solanales, sensu APG II (2003), contain 5 families (Table 8.3). Of these, two families are described here.

Convolvulaceae — Morning Glory / Bindweed family (Latin for interwoven). 56 genera / 1600 species. (Figure 8.73)

The Convolvulaceae consist of herbaceous to woody vines, less commonly herbs, shrubs, or rarely trees. Some family members are achlorophyllous and parasitic (e.g., *Cuscuta*). The **roots** are haustorial in parasitic taxa. The **stems** of viney members are dextrorse (twining clockwise when moving away, like the grooves of a typical right-handed screw). The **leaves** are simple, undivided to divided, spiral, exstipulate, reduced and scalelike in *Cuscuta*. The **inflorescence** is a head, dichasium, or of solitary flowers, bracteate, of usually two, often accrescent bracts. The **flowers** are bisexual, actinomorphic, and hypogynous. The **perianth** is dichlamydeous. The **calyx** has 5 [3,4], sepals or lobes. The **corolla** is sympetalous, often infundibular, with 5 [3,4] lobes, with usually involute (plicate) aestivation (imbricate in *Cuscuta*). The **stamens** are 5 [3,4], filaments often unequal in length, the stamens epipetalous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous with a superior ovary, 2 [3 5] carpels, and 1 several locules (as many as carpels). The **style(s)** are solitary to as many as carpels. **Placentation** is basal; **ovules** are anatropous, unitegmic, 2 per carpel [rarely ∞]. **Nectaries** are present, consisting of an annular disk around base of ovary. The **fruit** is a capsule (loculicidal, circumscissile, or irregularly dehiscent), berry, drupe, or nut. The **seeds** are endospermous.

The Convolvulaceae have a mostly worldwide distribution. Economic importance includes cultivated ornamentals such as *Convolvulus*, *Ipomoea*, and *Jacquemontia*; *Ipomoea batatas* is the sweet potato, source of the starchy storage root.

The Convolvulaceae are distinctive in being often **dextrorse-twining vines**, less commonly shrubs or trees, with simple, spiral leaves, and **actinomorphic, sympetalous** flowers, corollas typically with **involute aestivation** and often **infundibular**. **K** (5) [(3,4)] **C** (5) [(3,4)] **A** (5) [3,4] **G** (2) [(3-5)], superior.

Solanaceae Nightshade family (Latin for sleeping or comforter, after narcotic properties of some). 94 genera/2950 species. (Figure 8.74)

The Solanaceae consist of herbs, shrubs, trees, or lianas, with prickles present in some taxa, many with stellate trichomes. The **leaves** are simple, pinnate, or ternate, usually spiral and exstipulate. The **inflorescence** is of solitary flowers or cyme units. The **flowers** are bisexual, actinomorphic, rarely zygomorphic. The **perianth** is biseriate, dichlamydeous, usually tubular, rotate, or salverform, hypanthium absent. The **calyx** is synsepalous, persistent, sometimes accrescent, with 5 calyx lobes. The **corolla** is sympetalous and with 5 [4,6] convolute, imbricate, or valvate lobes, with usually involute (plicate) aestivation. The **stamens** are 5 [rarely 4 or 2 + 2 staminodes], antisepalous and epipetalous, the anthers often connivent, with staminodes rarely present. **Anthers** are longitudinal or poricidal in dehiscence. The **gynoecium** is syncarpous, with a superior ovary, 2 [rarely 3 5] carpels, and 2 [rarely 1 or 4 5] locules. **Placentation** is axile, rarely basal; **ovules** are variable in type, unitegmic, ∞ [rarely 1 few] per carpel. The **fruit** is a berry, drupe, or capsule (often septicidal). The **seeds** are endospermous. Alkaloids and internal phloem (inner to the xylem, surrounding pith) are present in many family members.

The Solanaceae are currently classified into seven subfamilies (Olmstead et al. 1999). Members of the family have mostly worldwide distributions, concentrated in South America. Economic importance includes many edible plants, such as *Capsicum* (peppers), *Lycopersicon esculentum* (tomato), *Physalis philadelphica* (tomatillo), and *Solanum tuberosum* (potato), and the infamous fumatory *Nicotiana tabacum* (tobacco). Alkaloids from various taxa have medicinal properties (e.g., atropine from *Atropa belladonna*), hallucinogenic properties (e.g., *Datura*, Jimson weed), or are deadly poisons (e.g., *Datura*, *Solanum* spp.) or known carcinogens (e.g., *Nicotiana tabacum*); some are used as ornamental cultivars, others are noxious weeds.

The Solanaceae are distinctive in being herbs, shrubs, trees, or lianas with **internal phloem**, spiral leaves, a usually **actinomorphic, 5-merous** perianth and androecium (**corolla involute in aestivation**), a usually **bicarpellate, syncarpous** gynoecium, and usually **numerous ovules** per carpel, the fruit a berry, drupe, or capsule.

K (5) **C** (5) [(4),(6)] **A** 5 [4 or 2+2 staminodes] **G** (2) [(3-5)], superior.

EUASTERIDS II

Euasterids II is an informal name used to denote a monophyletic grouping of four orders: Apiales, Aquifoliales, Asterales, and Dipsacales, plus several unplaced families (Table 8.3).

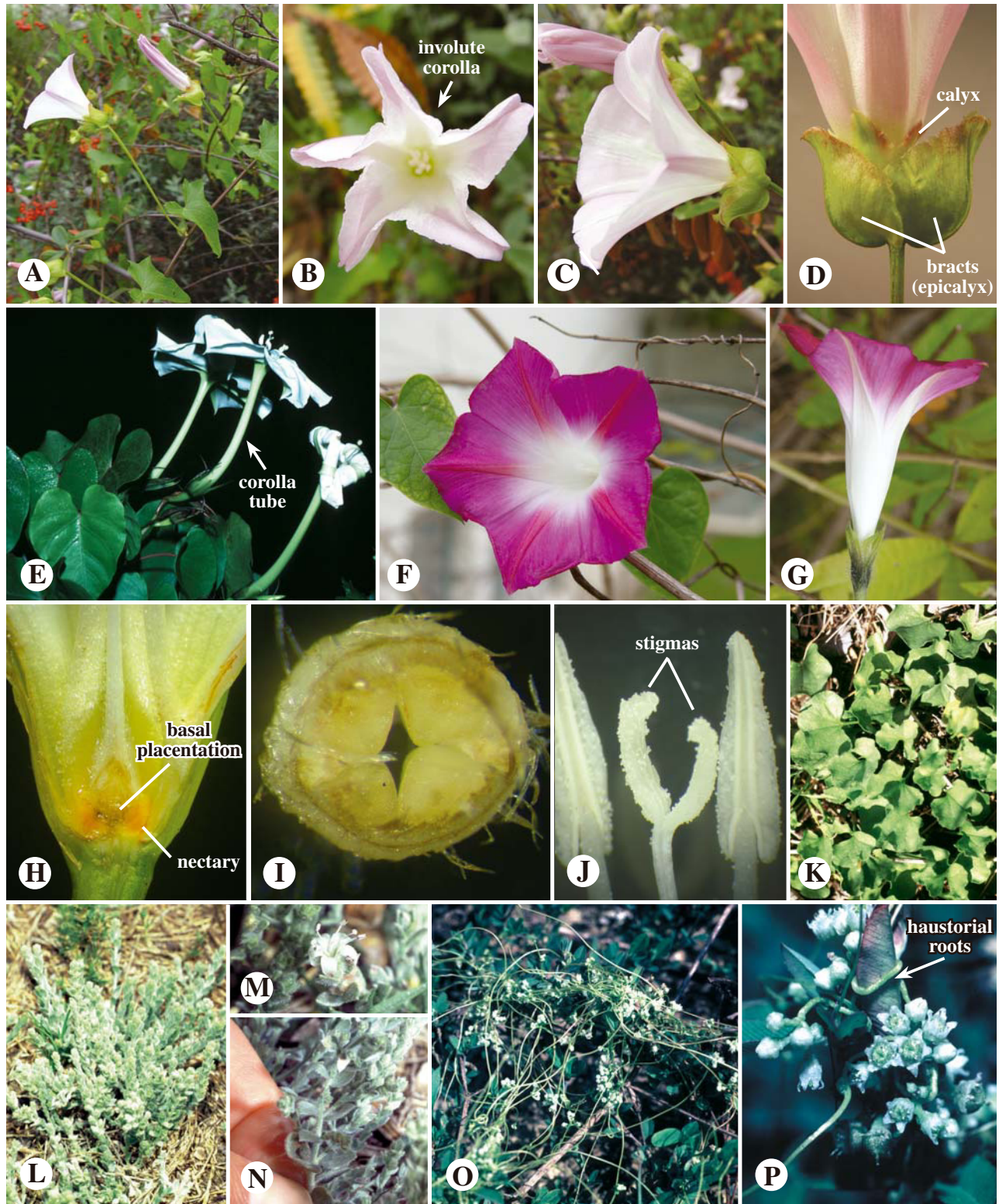


FIGURE 8.73 SOLANALES. Convolvulaceae. **A–D.** *Calystegia* sp. **A.** Whole vine, in flower. **B.** Young flower, showing involute aestivation. **C.** Flower, side view. **D.** Bracts (epicalyx) subtending flower. **E.** *Calonyction* sp., moonflower, with long, narrow corolla tube (moth pollinated). **F–J.** *Ipomoea* sp., morning glory. **F–G.** Open flowers, with infundibular corolla. **H.** Flower base longitudinal section, showing basal placentation. **I.** Ovary cross-section, showing four ovules. **J.** Style, with two style branches, and adjacent anthers. **K.** *Dichondra occidentalis*. **L–N.** *Cressa truxillensis*. **O, P.** *Cuscuta gronovii*, an achlorophyllous, parasitic vine with haustorial roots.

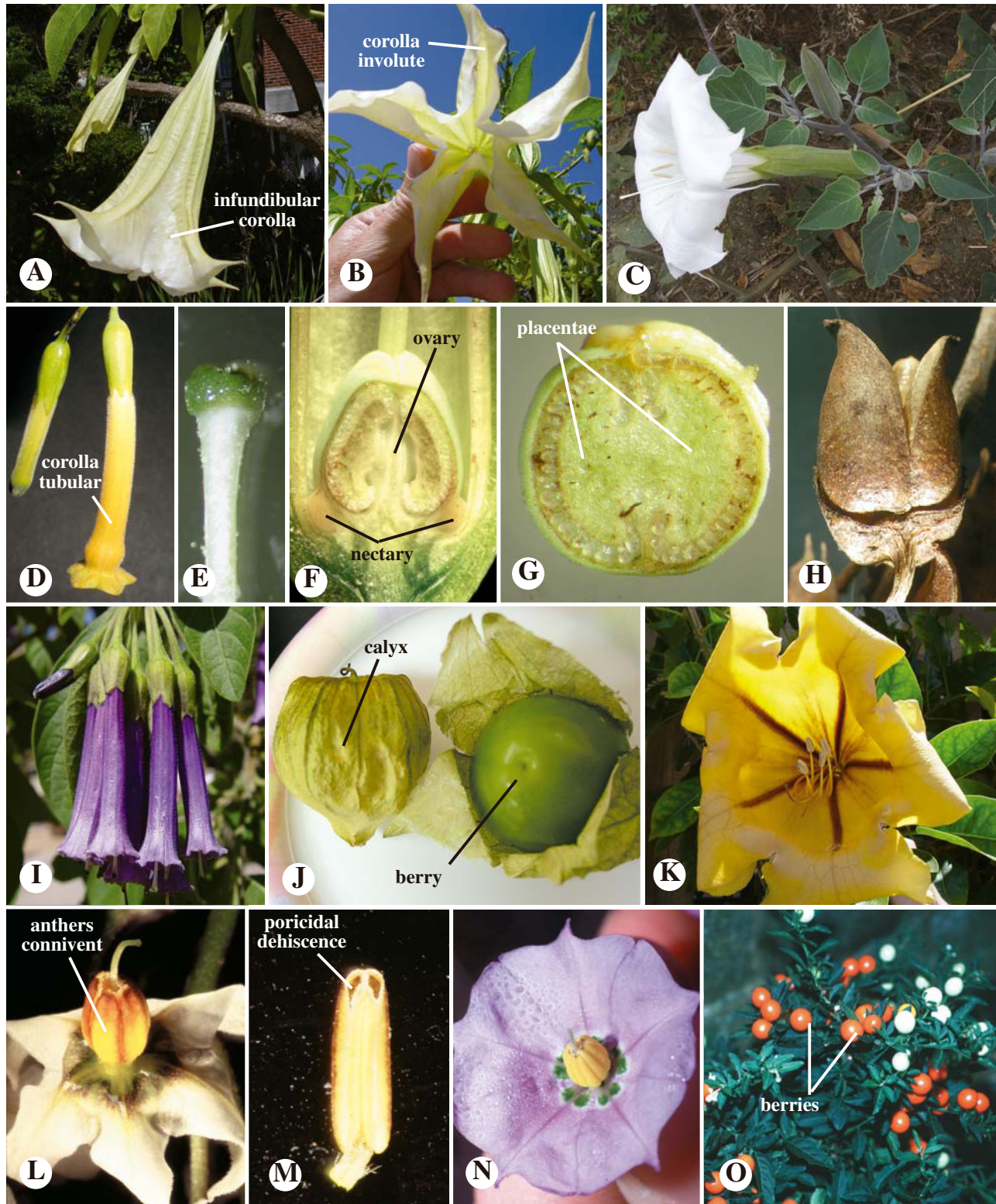


FIGURE 8.74 SOLANALES. Solanaceae. **A,B.** *Brugmansia* sp. Note actinomorphic, plicate corolla. **C.** *Datura wrightii*, jimson weed, a poisonous/hallucinogenic plant. **D–H.** *Nicotiana glauca*, tree tobacco. **D.** Corolla. **E.** Style and stigma. **F.** Ovary in longitudinal section. **G.** Ovary cross-section, showing bicarpellate axile placentation with large placentae. **H.** Fruit, a capsule. **I.** *Iochroma cyaneum*, with tubular corollas. **J.** *Physalis ixocarpa*, tomatillo, having an accrescent calyx. **K.** *Solandra maxima*. **L,M.** *Solanum douglasii*, one of the nightshades. **L.** Flower, with connivent stamens. **M.** Stamen, showing poricidal dehiscence. **N.** *Solanum parishii*, with prominent nectar guides. **O.** *Solanum pseudocapsicum*, an ornamental with berry fruits.

APIALES

The Apiales contain 8–10 families (Table 8.3), of which the two most common are described here. See Chandler and Plunkett (2004) for more detailed information on the order.

Apiaceae (Umbelliferae) Carrot family (*apium*, used by Pliny for a celerylike plant). 446 genera/3540 species. (Figures 8.75, 8.76)

The Apiaceae consist of herbs, less often shrubs or trees. The **leaves** are usually pinnate, ternate, or decomposed [rarely simple, palmate, or phyllodinous], spiral, with a broad sheathing base, stipular flanges sometimes present. The **inflorescence** is usually a compound umbel often with subtending involucre bracts, sometimes a head or simple umbel or reduced to a single flower or dichasium. The **flowers** are small, bisexual [marginal flowers sometimes sterile], actinomorphic, epigynous. The **perianth** is biseriate and dichlamydeous or uniseriate by loss of the calyx. The **calyx** is aposepalous with 5 lobes, which may be reduced or absent. The **corolla** is apopetalous and with 5 [rarely 0], valvate petals. The **stamens** are 5, whorled, alternipetalous, and apostemonous. The **gynoecium** is syncarpous, with an inferior ovary, 2 carpels, and 2 [rarely 1] locules, often with a stylopodium at apex of ovary. **Placentation** is apical-axile; **ovules** are anatropous, pendulous, unitegmic, 1 per carpel. The **fruit** is a schizocarp of mericarps, supported by carpophores upon splitting. The **seeds** are endospermous, endosperm oily. Some taxa have anomalous secondary thickening.

The Apiaceae have a worldwide distribution. Economically important members include a number of food, herb, and spice plants, such as *Anethum*, dill; *Apium*, celery; *Carum*, caraway; *Coriandrum*, coriander; *Cuminum*, cumin; *Daucus*, carrot; *Foeniculum*, fennel; and *Petroselinum*, parsely; some species are poisonous, such as *Conium maculatum*, poison-hemlock (an extract of which Socrates drank in execution); others are used as ornamental cultivars. See Plunkett et al. (1997) for a more detailed study of the Apiaceae.

The Apiaceae are distinctive in being **herbs**, with **sheathing leaves** (compound or simple, often decomposed), the inflorescence usually an **involucre compound umbel** [rarely a head, simple umbel, or reduced] with actinomorphic flowers having a 2-carpellate and 2-loculate, **inferior ovary**, each carpel with **one, axile-apical, pendulous ovule**, the fruit a **schizocarp of mericarps**.

K 5 or 0 **C** 5 [0] **A** 5 **G** (2), inferior.

Araliaceae — Ginseng family (possibly from French Canadian Aralie). 47 genera / 1325 species. (Figure 8.77)

The Araliaceae consist of trees, shrubs, lianas, or herbs. The **stems** of trees are often pachycaulis. The **leaves** are palmate, pinnate, or simple (these often divided), usually spiral, rarely opposite or whorled, usually stipulate. The **inflorescence** is a usually terminal umbel, head, or secondary inflorescence (e.g., panicle) of umbels, rarely of solitary flowers. The **flowers** are usually bisexual, actinomorphic, epigynous (rarely hypogynous). The **calyx** is aposepalous with 5 [3–∞] sepals, often reduced or absent. The **corolla** is apopetalous, rarely basally sympetalous or calyptrate, with 5 [rarely 3–12], valvate or imbricate petals or lobes. The **stamens** are 5–10 [rarely 3–∞]. **Anthems** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with an inferior (rarely superior and secondarily derived) ovary, 2–5 [∞] carpels, and 1–∞ locules. The **styles** are 1–∞. **Placentation** is apical-axile; **ovules** are anatropous, unitegmic, 1 [2] per carpel. **Nectaries** are sometimes present as an epigynous disk. The **fruit** is a drupe with multiple endocarps, berry, or schizocarp with carpophore. The **seeds** are oily endospermous. Plant tissues usually have secretory canals; leaf nodes are multilacunar.

The Araliaceae have a mostly worldwide distribution, mainly in tropical regions. Economic importance includes medicinally important plants, especially *Panax* (ginseng); *Tetrapanax papyrifer*, used as Chinese rice paper; some timber plants; and several cultivated ornamentals such as *Fatsia*, *Hedera* (ivy), and *Schefflera*. See Wen et al. (2000) for a study of the Araliaceae.

The Araliaceae are distinctive in being **mostly tropical** trees, shrubs, lianas, or herbs with **palmate or pinnate** (rarely simple, then usually divided) leaves, an **inflorescence of heads, umbels, or with umbel units**, the flowers with often reduced calyx, apopetalous to sympetalous corolla, and a **1–∞-carpellate inferior ovary** with usually **apical-axile placentation**, the fruit a berry, drupe, or schizocarp.

K 5 [0, 3–∞] **C** 5 [3–12] **A** 5–10 [3–∞] **G** (2–5) [(–∞)], inferior, rarely superior.

ASTERALES

The Asterales, sensu APG II (2003), contain 11 families (Table 8.3), two of which are described here. Notable among the other families are the **Campanulaceae** (the bluebell/lobelia family, a large group containing a number of species used as ornamentals) and **Stylidiaceae** (including the trigger plants, with an interesting pollination mechanism; see Chapter 13). See Lundberg and Bremer (2003) for a detailed study of the Asterales.

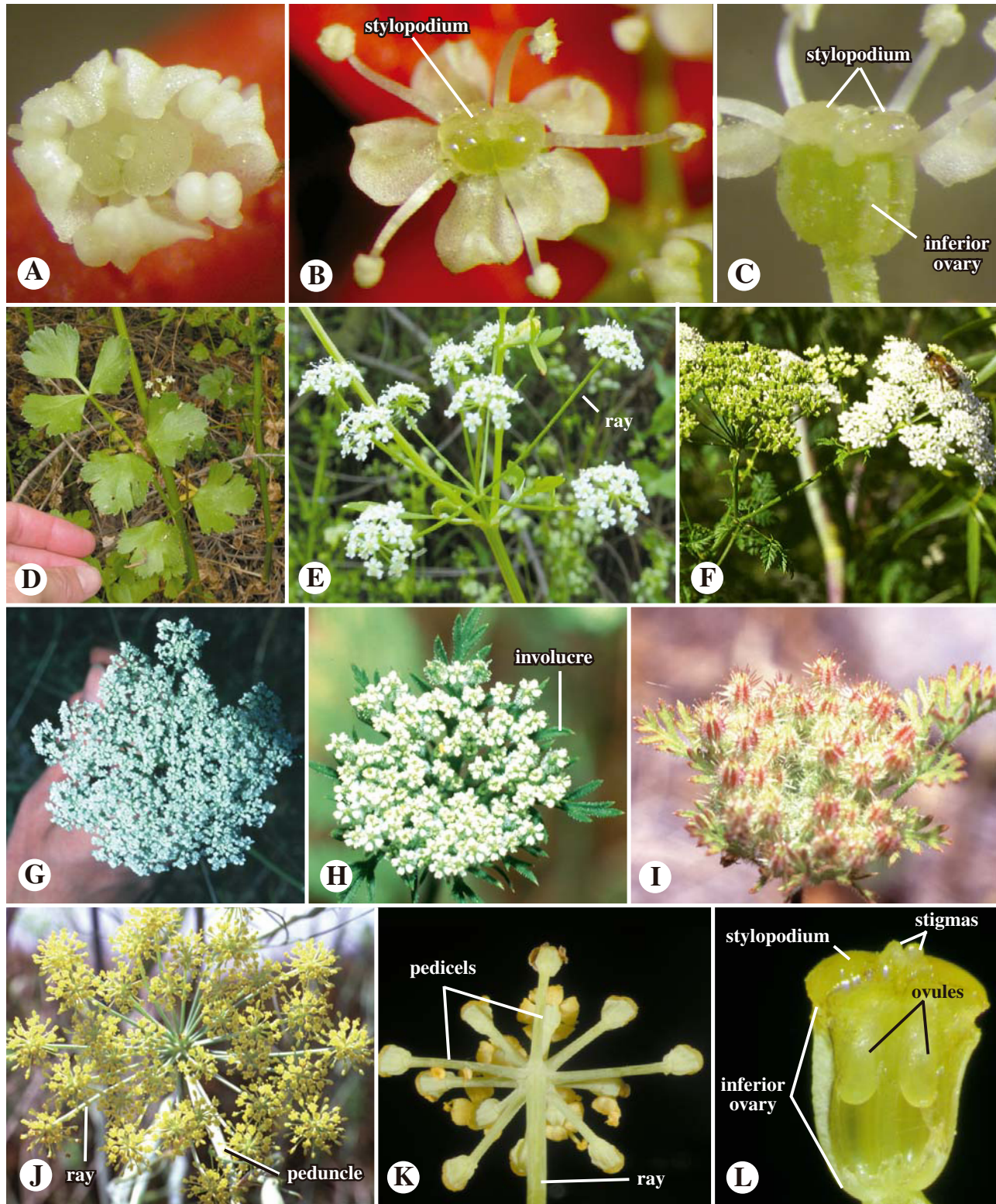


FIGURE 8.75 APIALES. Apiaceae. A–E. *Apium graveolens*, wild celery. A. Flower, immature, with incurved petals. B. Flower at anthesis, showing stylopodium and alternipetalous stamens. C. Flower in side view; note inferior ovary. D. Leaf. E. Inflorescence, a compound umbel. F. *Conium maculatum*, poison-hemlock, of Socrates fame. G. *Daucus carota*, wild carrot or Queen Anne’s lace, in inflorescence. H, I. *Daucus pusillus*. H. Inflorescence, showing subtending involucre. I. Inflorescence, with bristly fruits. J–L. *Foeniculum vulgare*, wild fennel. J. Compound umbel, showing rays. K. Close-up of simple umbel unit. L. Ovary longitudinal section, showing two carpels and apical-axile placentation.

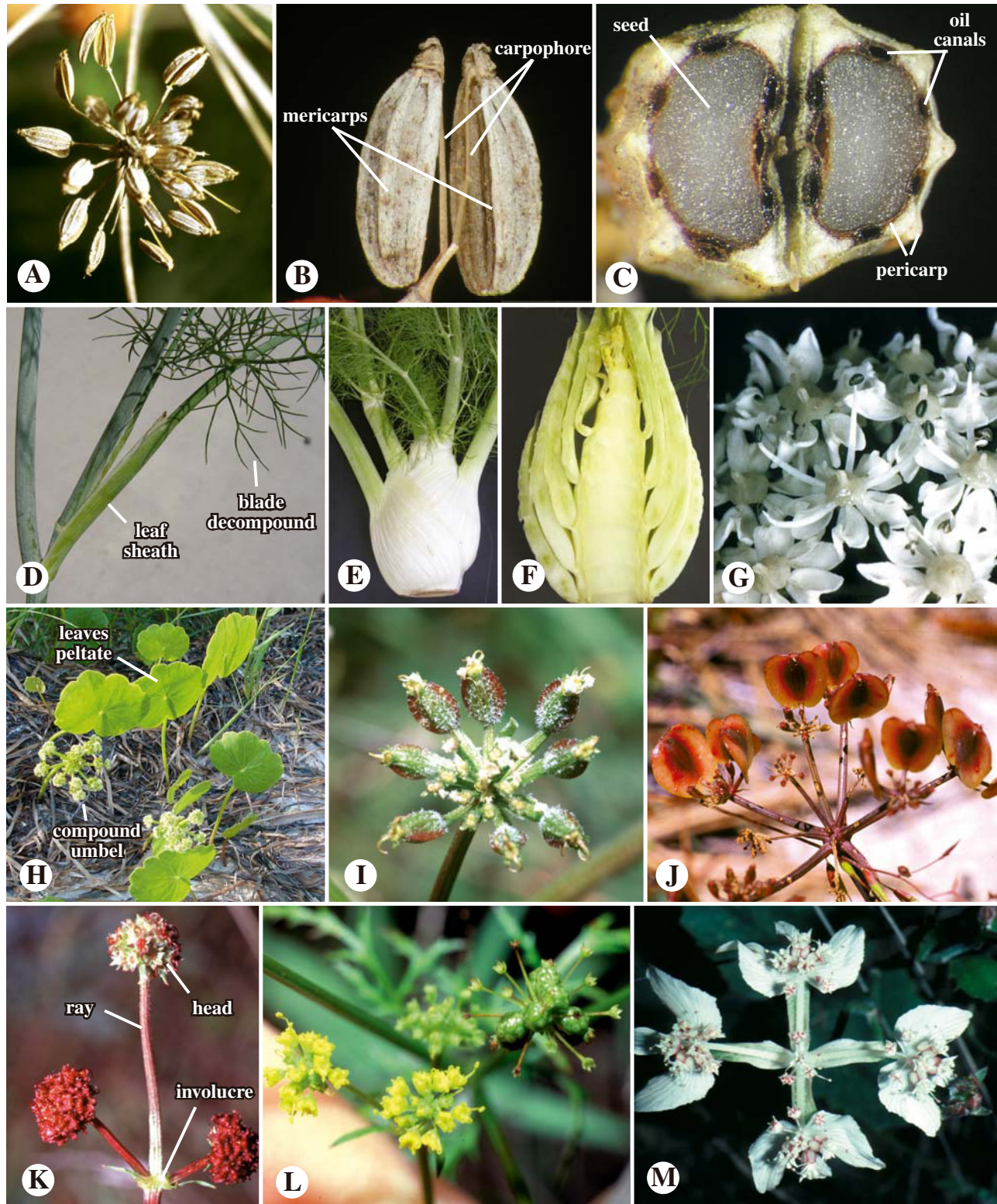


FIGURE 8.76 APIALES. Apiaceae. A–F. *Foeniculum vulgare*. A. Umbel unit of infructescence. B. Fruit, a schizocarp of mericarps, showing supporting carpophores. C. Fruit in cross-section, showing two carpels, oil canals, and seeds. D. Leaf, with prominent, adnate stipules. E, F. Florence fennel (var. *dulce*), bred for succulent stems and petioles, used as a vegetable. G. *Heracleum lanatum*, showing flowers of umbel unit. H. *Hydrocotyle* sp., pennywort. I. *Lomatium dasycarpum*, with immature fruits. J. *Lomatium lucidum*, having winged fruits. K. *Sanicula bipinnatifida*, with an umbel of headlike units. L. *Sanicula tuberosa*, showing short, bisexual flowers and long-pedicellate male flowers. M. *Xanthoxia* sp., southern cross, with unusual bracts and winged rays.

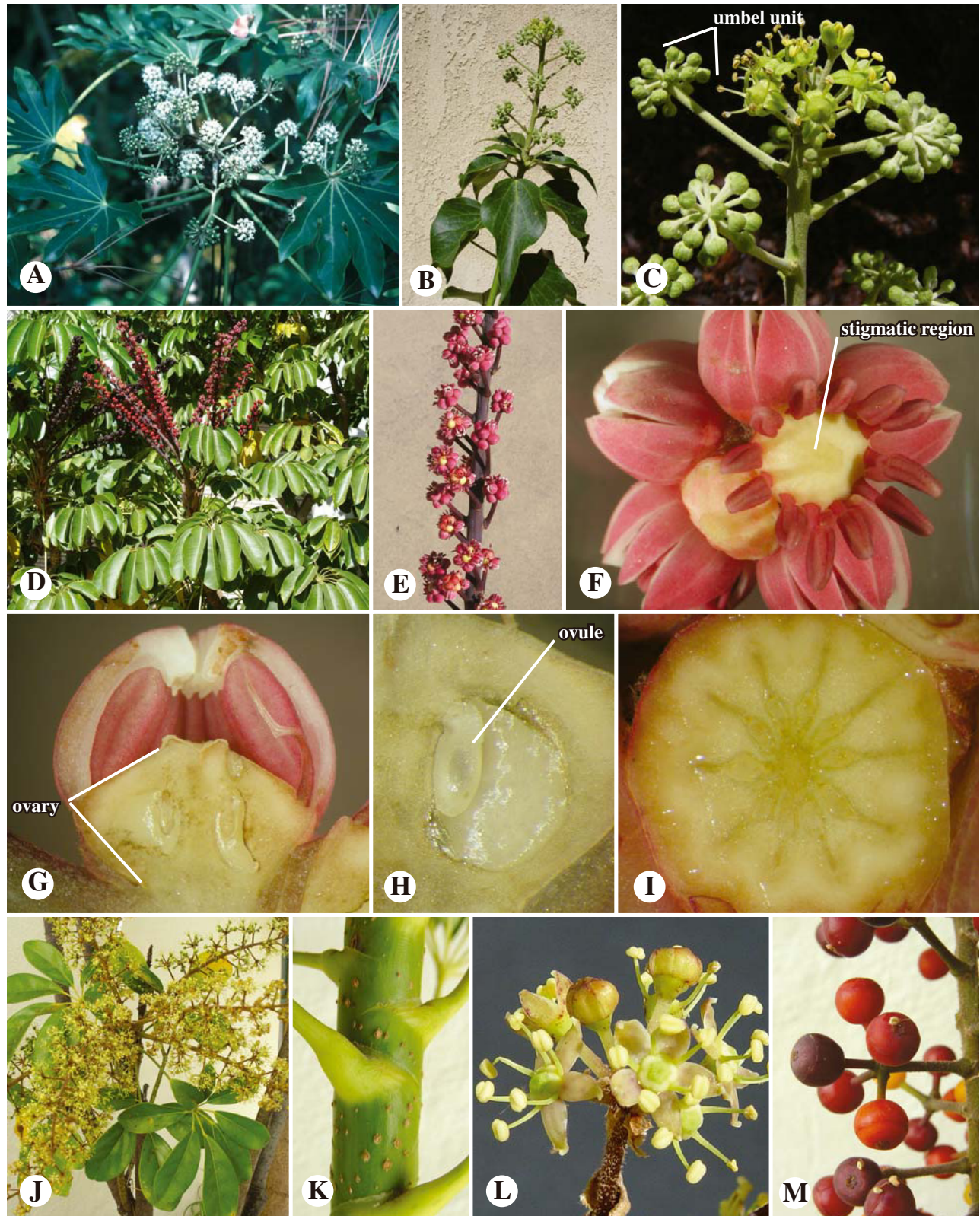


FIGURE 8.77 APIALES. Araliaceae. **A.** *Fatsia japonica*, the inflorescence a panicle of umbels. **B,C.** *Hedera canariensis*, Algerian ivy. **B.** Reproductive shoot, with a raceme of umbels. **C.** Close-up of unit umbels. **D–I.** *Brassia actinophylla*, octopus tree. **D.** Whole plant, showing palmate leaves and large, terminal panicle of umbels. **E.** Close-up of inflorescence axis. **F.** Unit umbel. Note open flower, petals caducous. **G.** Mature bud in longitudinal section. **H.** Carpel close-up, showing apical-axile placentation. **I.** Ovary cross-section, 10-carpellate. **J–M.** *Scheffera* sp. **J.** Whole plant. **K.** Close-up of stem nodes, showing broad point of attachment. **L.** Inflorescence umbel unit. **M.** Fruits, drupes.

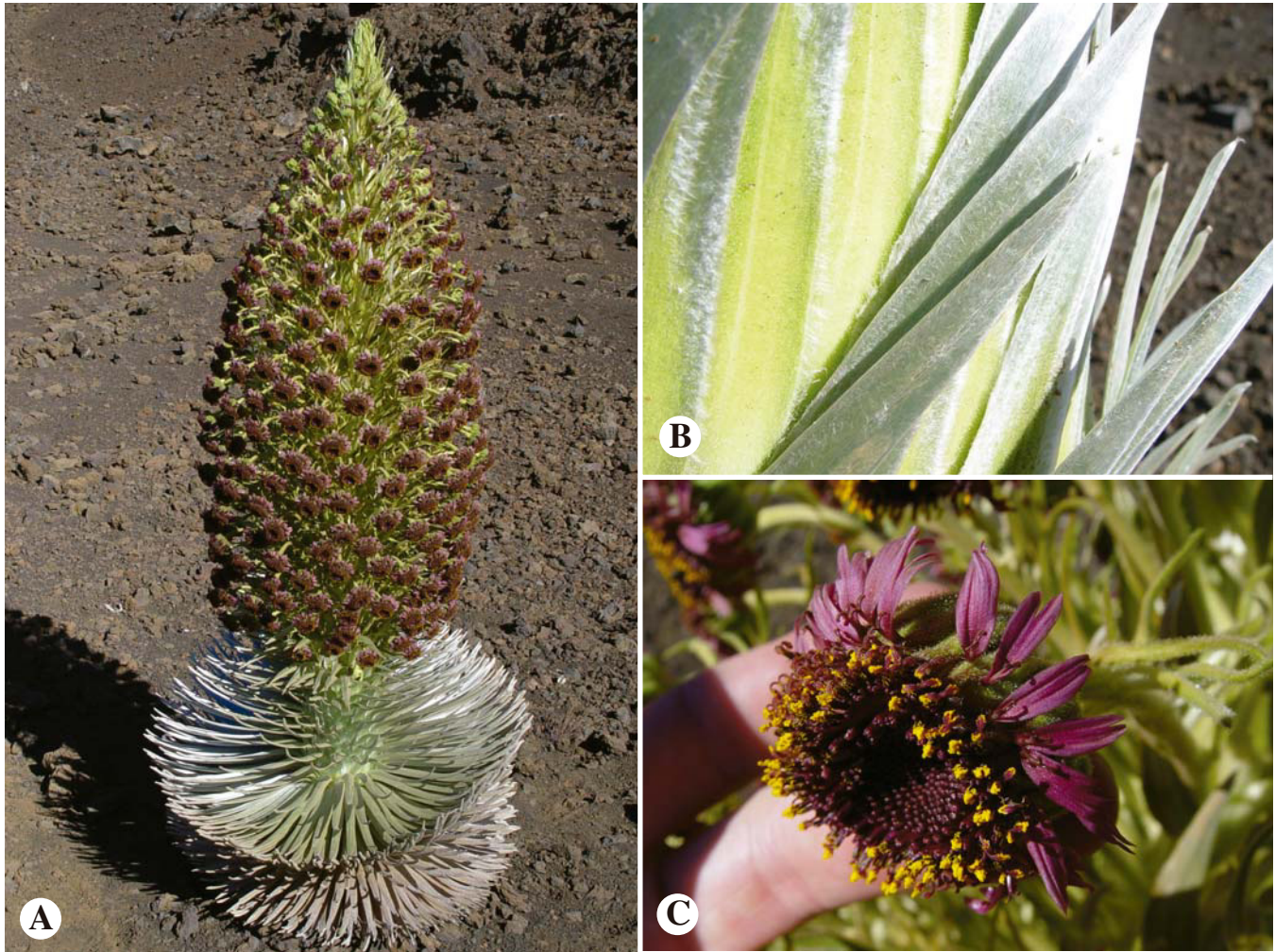


FIGURE 8.78 ASTERALES. Asteraceae. *Argyroxiphium sandwicense*, silversword. **A.** Whole plant at time of flowering. **B.** Close-up of basal leaves, showing silver, UV-reflectant, sericeous trichome layer. **C.** Close-up of head (capitulum), characteristic of the Asteraceae.

Asteraceae (Compositae) Sunflower family (after Aster, meaning star). 1528 genera / 22,750 species. (Figures 8.78–8.82)

The Asteraceae consist of herbs, shrubs, trees, or vines, with laticifers or resin ducts present in some taxa. The **leaves** are simple or compound, spiral or opposite [rarely whorled], exstipulate. The **inflorescence** consists of one or more heads (capitula) arranged in various secondary inflorescences, each head consisting of a flat to conical compound receptacle that bears one to many flowers (developing centripetally) and is subtended by one or more series of bracts, the phyllaries (collectively termed the involucre); heads of five general types: (1) discoid, with only disk flowers, all bisexual; (2) disciform, with only disk flowers, a mixture of pistillate and sterile with bisexual and staminate, in the same or different heads; (3) radiate, with central (bisexual or male) disk

flowers and peripheral (female or sterile) ray flowers; (4) ligulate, with all ray flowers (typically with 5-toothed corolla apices); and (5) bilabiate, with all bilabiate flowers. The **flowers** are epigynous, bisexual or unisexual, subtended in some taxa by bracts, known as chaff, or bristles (as in the thistles). The **perianth** is biseriate or uniseriate with hypanthium absent. The **calyx**, known as the **pappus**, is modified as 2∞ (sometimes connate) awns, scales, or capillary bristles (typically barbed or plumose), pappus absent in some. The **corolla** is sympetalous with 5 [rarely 4] lobes (reduced to 3 marginal teeth in some), of three structural types (also called flower types): (1) **bilabiate**, corolla zygomorphic with a short tube having upper and lower lips; (2) **disk**, corolla actinomorphic with short to elongate tube bearing 5 [4] teethlike or elongate lobes; or (3) **ray** or **ligulate**, corolla zygomorphic

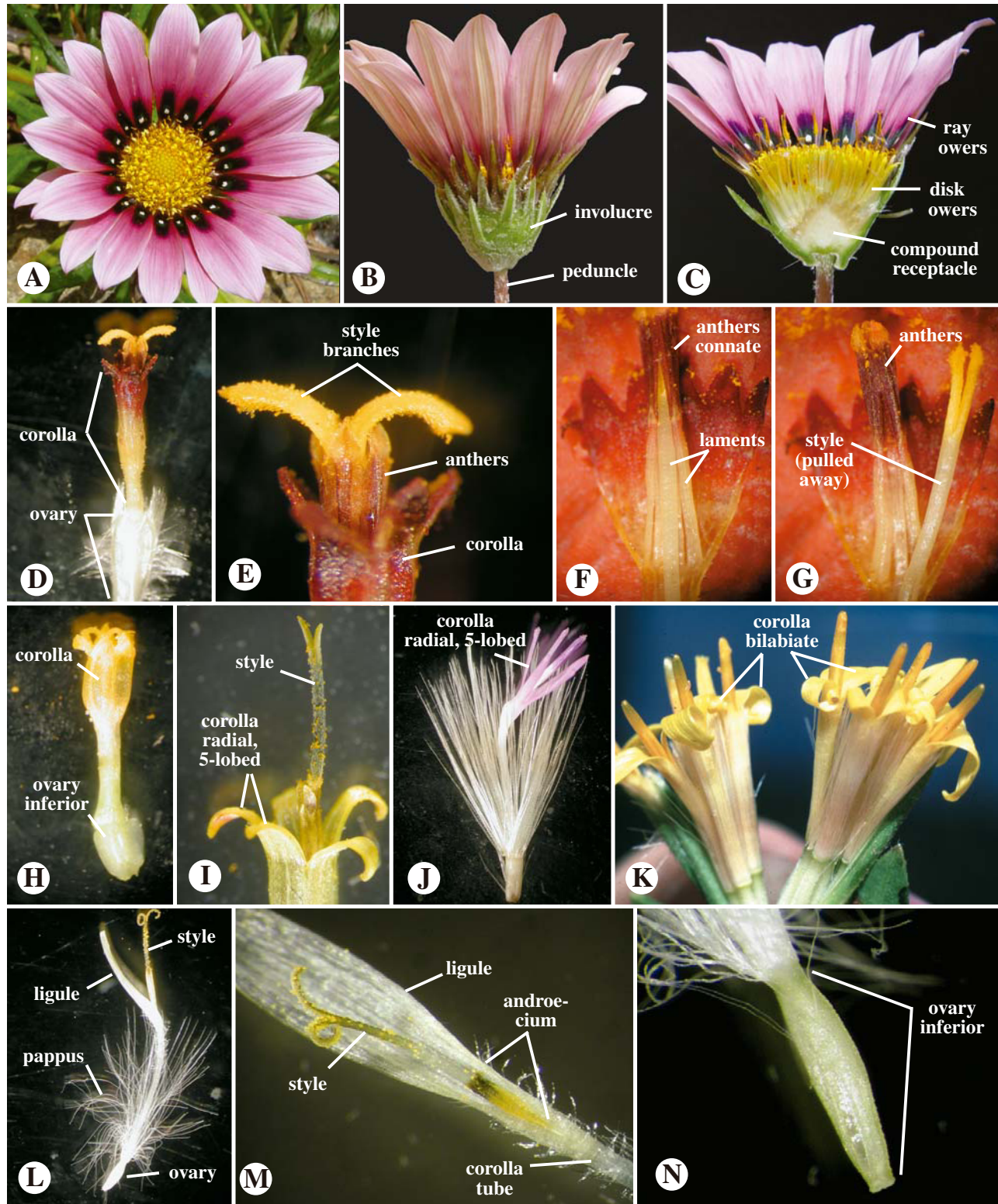


FIGURE 8.79 ASTERALES. Asteraceae. A–C. *Gazania* sp., head morphology. A. Radiate head, face view. B. Head, side view, showing involucre. C. Head, longitudinal section, showing compound receptacle, central disk flowers, and peripheral ray flowers. D–G. *Encelia californica*. D. Disk flower, showing inferior ovary and corolla. E. Flower apex. Note two-branched style and subtending anthers. F, G. Dissected flower, showing syngenesious androecium, with central style. H–J. Disk flower morphology. H. *Chrysanthemum coronarium*. I. *Vigiera laciniata*. J. *Carduus pycnocephalus*. K. Bilabiate flower morphology, *Trixis californica*. L–N. Ray flower morphology, *Sonchus oleraceus*.



FIGURE 8.80 ASTERALES. Asteraceae. A–D. Involucre morphology. A. Imbricate, multiseriata, isomorphic phyllaries in *Encelia californica*. B. Imbricate, multiseriata, dimorphic phyllaries in *Ursinia* sp. C. Mostly uniseriate phyllaries (but with tiny outer bracts below) in *Senecio vulgaris*. D. Decussate phyllaries in *Jaumea carnosa*. E–G. Secondary inflorescence types. E. Glomerule of heads in *Carduus pycnocephalus* (also having spine-tipped phyllaries). F. Corymb of heads in *Achillea* sp. G. Secund, narrow panicle of heads in *Solidago pinetorum*. H–J. Radiate heads, having inner disk flowers and peripheral ray flowers. H. *Aster* sp. I. *Layia platyglossa*. J. *Tithonia rotundifolia*. K–M. Ligulate heads, with all ray flowers. K. *Malacothrix californica*. L. *Cichorium intybus*, chicory. M. *Ranunculus acris*.



FIGURE 8.81 ASTERALES. Asteraceae. **A,B.** Bilabiate heads, i.e., having bilabiate owers; *Acurtia microcephala*. **C–F.** Discoid heads, with all disk owers. **C.** *Chaenactis artemisiifolia*. **D.** *Chaenactis gabriuscula*. **E.** *Cirsium vulgare*, a thistle. **F.** *Carduus pycnocephalus*. **G,H.** Disciform heads, with inner staminate or bisexual and outer pistillate owers. *Cotula coronopifolia*, brass buttons. Note reduced, 4-merous owers. **I,J.** Discaid heads. **I.** *Palafoxia arida*. **J.** *Psathyrotes ramosissima*. **K–M.** Disciform heads, having unisexual owers. **K.** *Ambrosia chamissonis*, male (staminate) heads above, female (pistillate) below. **L,M.** *Ambrosia [Hymenoclea] salsola*. **L.** Male heads. **M.** Female heads.

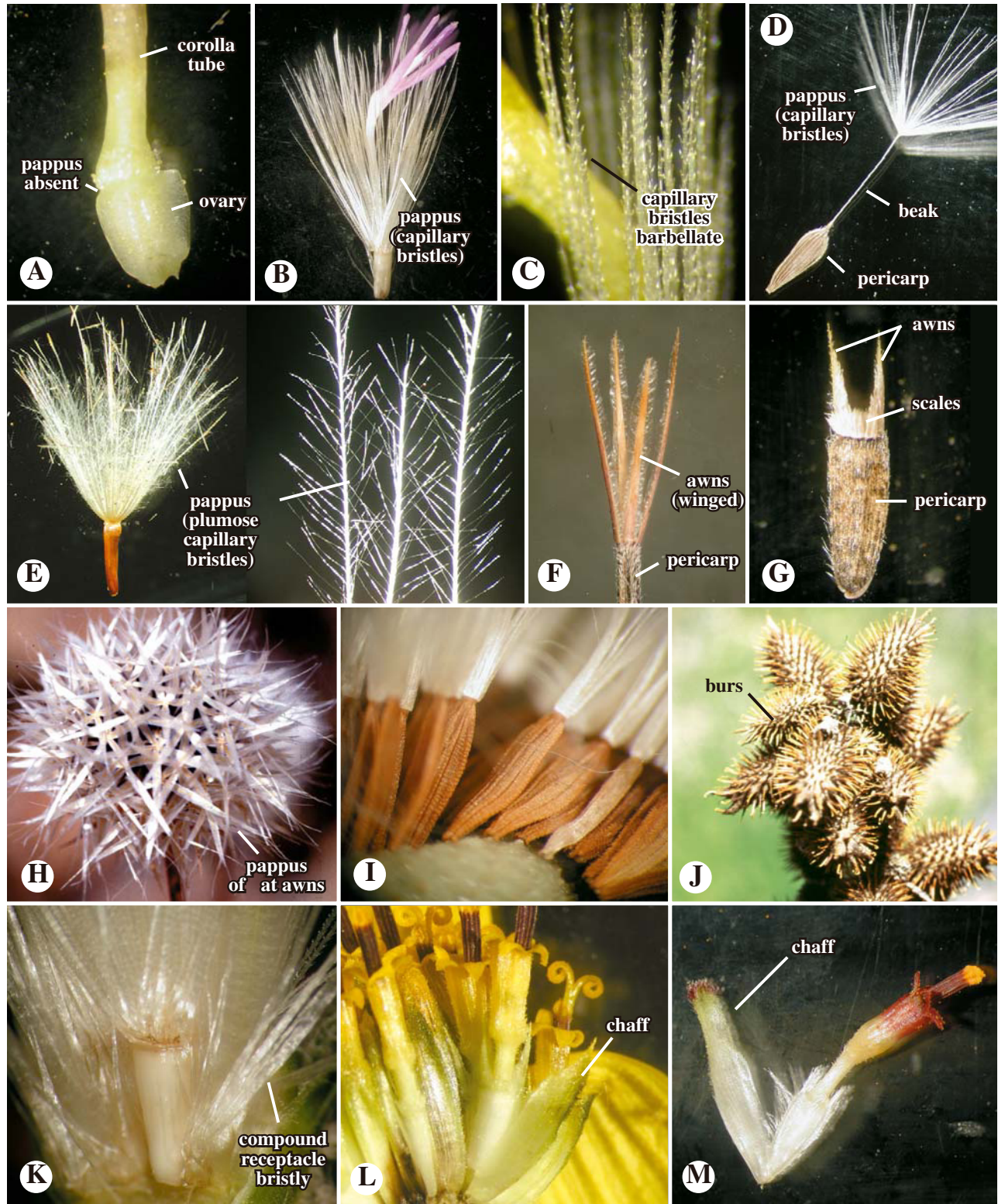


FIGURE 8.82 ASTERALES. Asteraceae. A–H. Pappus morphology. A. Pappus absent, *Chrysanthemum coronarium*. B. Pappus of capillary bristles, *Carduus pycnocephalus*. C. Capillary bristles antrorsely barbellate, *Isocoma menziesii*. D. Fruit, with beak and distal pappus of capillary bristles, *Lactuca serriola*. E. Pappus of plumose capillary bristles, *Cirsium mohavense*. F. Pappus of awns, *Palafoxia arida*. G. Pappus of two awns and several scales, *Viguiera laciniata*. H. Pappus of at awns, *Uropappus lindleyi*. I. Fruits (achenes), *Sonchus oleraceus*. J. Bur fruits, achenes with a spiny involucre as accessory tissue, *Xanthium strumarium*. K. Compound receptacle bearing numerous bristles, *Carduus pycnocephalus*. L, M. Heads with chaff subtending flowers. L. *Viguiera laciniata*. M. *Encelia californica*.

with generally short tube having elongate, flat, extension bearing 3–5 apical teeth. The **stamens** are 5 [4], whorled, alternipetalous, usually syngenesious, the anthers fused into a tube through which the style grows. **Anthers** are basifixed, with apical extensions and sometimes basal lobes, longitudinal and introrse in dehiscence. The **gynoecium** is syncarpous, with an inferior ovary, 2 carpels, and 1 locule. The **style** is solitary and apically two-branched; **stigmas** are two, occurring as stigmatic lines on the adaxial surface of style branches. **Placentation** is basal; **ovules** are anatropous, unitegmic, 1 per ovary. **Nectaries** are usually present at apex of ovary. The **fruit** is an achene (or cypsela, an achene derived from an inferior ovary), typically a multiple fruit of achenes, an elongate beak forming between fruit and pappus in some taxa. The **seeds** are exalbuminous.

The Asteraceae has recently been classified into at least ten subfamilies (Panero and Funk, 2002). Members of the family have a worldwide distribution. Economic importance includes some food (e.g., *Cynara scolymus*, artichoke, and *Helianthus annuus*, sunflower), a number of ornamental cultivars, and various species used locally or industrially. See Bremer (1994, 1996) and Panero and Funk (2002) for more detailed information on relationships in the Asteraceae.

The Asteraceae are distinctive in being herbs, shrubs, vines, or trees, the inflorescence a **head (capitulum)** subtended by an **involucre** of **phyllaries**, flowers either **bilabiate**, **disk**, or **ray/ligulate** (heads of many taxa a mixture of central disk flowers and peripheral ray flowers), with the calyx, termed a **pappus**, modified as scales, awns, or capillary bristles (or absent), the androecium **syngenesious**, and with an **inferior ovary** with a **single, basal ovule**, the fruit a multiple of **achenes**.

K 0–∞ (pappus) **C** (5) [(4)] or (3) in some ray flowers **A** (5) [(4)] **G** (2), inferior.

Goodeniaceae — Goodenia family. 12 genera / 400 species. (Figure 8.83)

The Goodeniaceae consist of shrubs, herbs, or rarely trees. The **leaves** are simple, alternate, spiral [rarely opposite or whorled], exstipulate. The **inflorescence** is of solitary flowers or a head, raceme, or cyme. **Flowers** are bisexual, zygomorphic, epigynous [to hypogynous]. The **perianth** is dichlamydeous. The **calyx** is synsepalous with 5 [3] lobes, reduced in some taxa. The **corolla** is valvate, sympetalous with 5 lobes, bilabiate or unilabiate with the 5 lobes anterior. The **stamens** are 5, alternipetalous, epipetalous or free. **Anthers** are connivent or connate, forming a tube, longitudinal introrse in dehiscence. The **gynoecium** is syncarpous, with an inferior [rarely half-inferior or superior] ovary, 2 carpels, and 2 [rarely 1 or 4] locules. The **style** is solitary, growing

through the tube formed by the anthers, the style having, below the stigma, a cupular indusium (having marginal hairs) that collects pollen and presents to a visiting insect. **Placentation** is axile; **ovules** are anatropous, unitegmic, 1–∞ per carpel. Intrastaminal **nectaries** are present in some taxa. The **fruit** is a capsule, rarely a nut or drupe. The **seeds** are flat, winged in some taxa, and oily endospermous.

The Goodeniaceae are mostly Australian in distribution. Economic importance includes some taxa used as ornamental cultivars.

The Goodeniaceae are distinctive in being herbs, shrubs, rarely trees, the flowers **5-merous**, with sympetalous, **bilabiate or often unilabiate (with the 5 lobes anterior)** corollas, **stamens forming a tube**, the style **growing through the connivent or connate anthers** with a **cupular indusium** that collects pollen, and a **usually inferior ovary**, the fruit a capsule, rarely a drupe or nut.

K (5) [(3)] **C** (5) **A** (5) or 5 **G** (2), inferior [rarely half-inferior or superior].

DIPSACALES

The Dipsacales, as treated by APG II (2003), contain two families: Adoxaceae and Caprifoliaceae. The Adoxaceae includes *Viburnum*, with several species used as ornamentals, and *Sambucus*, elderberry, used as fruit and wine plants (Figure 8.84A,B). The Caprifoliaceae in APG II is united with five previously recognized families: Diervillaceae, Dipsacaceae, Linnaeaceae, Morinaceae, and Valerianaceae. However, these families are here treated separately (an option in APG II) for convenience of circumscription and description, the Caprifoliaceae cited as *sensu stricto* (Table 8.3). Three families are treated here. See Donoghue et al. (2001) and Zhang et al. (2003) for recent phylogenetic analyses of the Dipsacales.

Caprifoliaceae s.s. — Honeysuckle family (Latin for goat leaf). 15 genera / 420 species. (Figure 8.84C–H)

The Caprifoliaceae consist of shrubs, trees, lianas, or herbs. The **leaves** are simple [rarely pinnate], opposite, exstipulate or reduced-stipulate. The **inflorescence** is usually a cyme. The **flowers** are bisexual, epigynous, rarely epihypogynous. The **calyx** is synsepalous with 5 [4] imbricate or open lobes, the calyx often constricted below lobes. The **corolla** is sympetalous with 5 [4], imbricate or valvate lobes. The **stamens** are 5 [2,4], alternipetalous, and epipetalous. **Anthers** are longitudinal in dehiscence. The **gynoecium** is syncarpous, with an inferior ovary, 2–5 [8] carpels (not all carpels fertile in some taxa), and 1–5 [8] locules. The **style** is solitary or absent. **Placentation** is axile or parietal; **ovules** are anatropous,

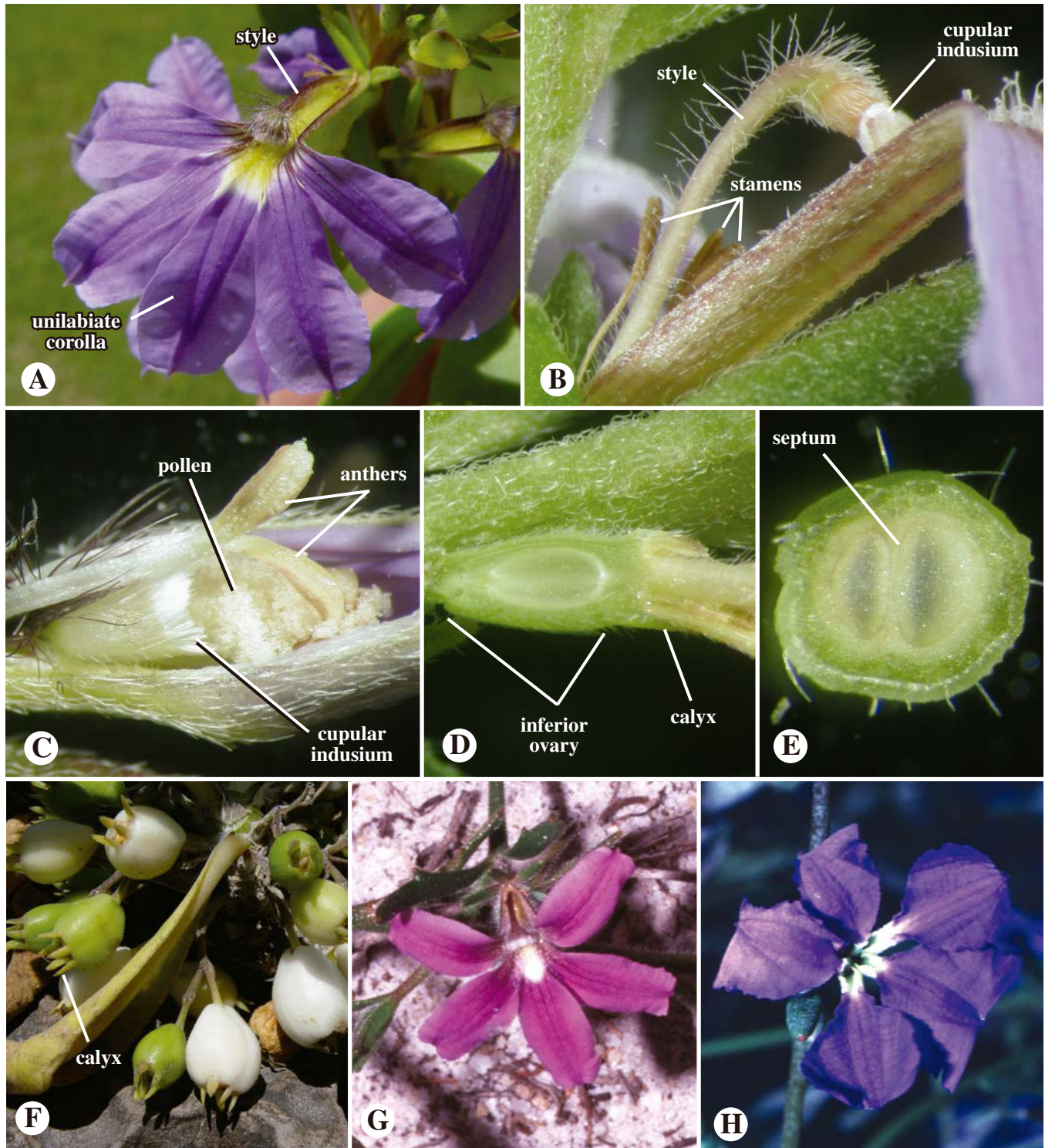


FIGURE 8.83 ASTERALES. Goodeniaceae. A–G. *Scaevola* spp. A. Flower, face view, showing unilabiate corolla and style along cleft of corolla tube. B. Close-up of style, with cupular indusium. Note lower stamens. C. Bud stage, showing style during elongation between stamens, the cupular indusium collecting pollen grains. D. Flower base in longitudinal section, showing inferior ovary. E. Ovary cross-section, showing two carpels and locules. F. Fruits (berries). G. Flower, face view. H. *Dampiera* sp.

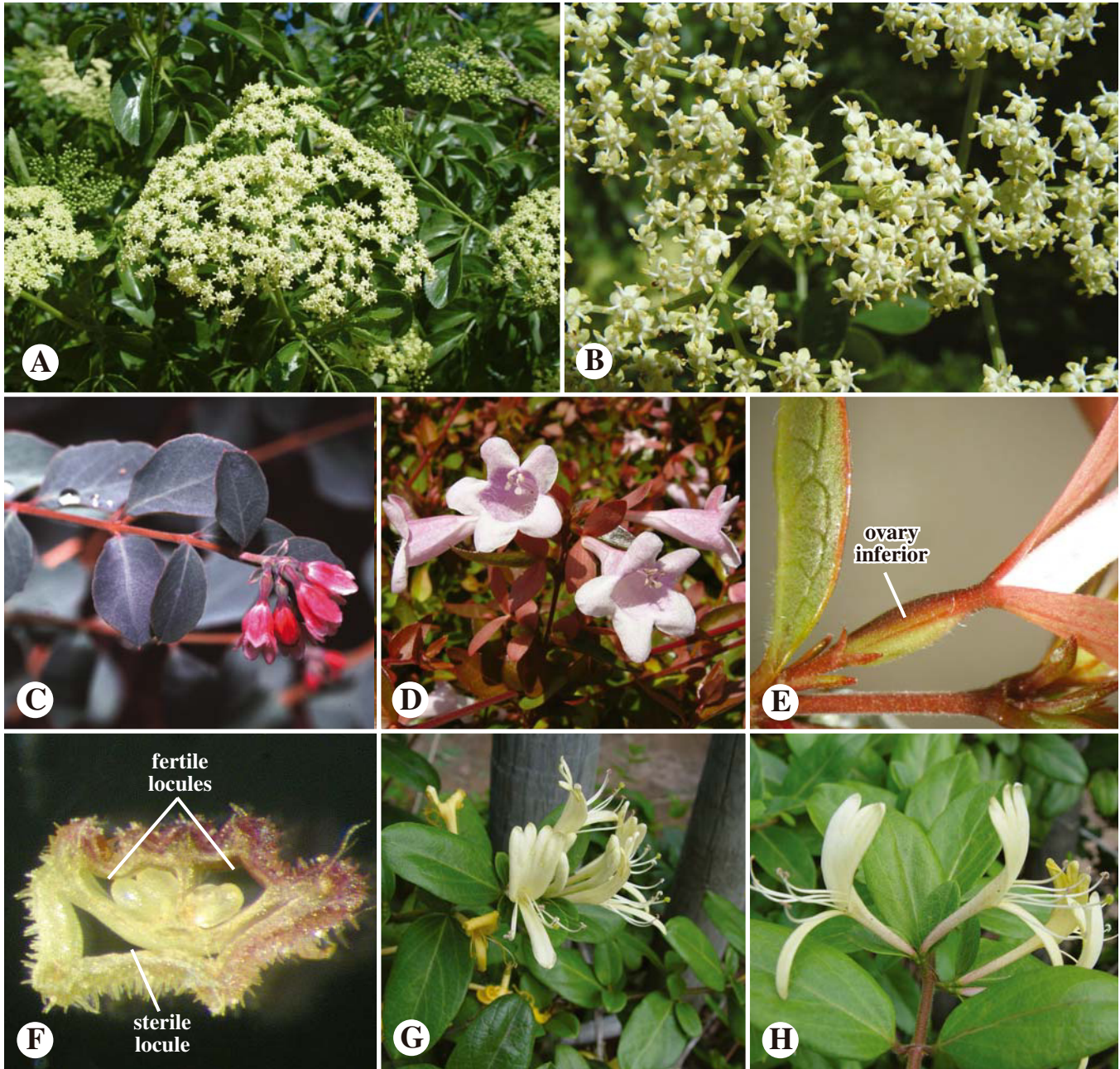


FIGURE 8.84 DIPSACALES. **A,B.** Adoxaceae, *Sambucus mexicana*, one of the elderberries. **C–H.** Caprifoliaceae s.s. **C.** *Symphoricarpos mollis*, snow berry. **D–F.** *Abelia x grandiflora*. **D.** Inflorescence and flowers. **E.** Ovary close-up. **F.** Ovary cross-section, showing three locules, lower one without ovules. **G,H.** *Lonicera japonica*, honeysuckle.

unitegmic, 1 ∞ per carpel. **Nectaries** are often present on inner corolla tube, with extrafloral nectaries present in some taxa. The **fruit** is a capsule, berry, or drupe. The **seeds** are oily endospermous.

The Caprifoliaceae have a mostly worldwide distribution. Economic importance includes several ornamental cultivars, such as *Abelia*, *Lonicera* (honeysuckles), and *Viburnum*.

The Caprifoliaceae are distinctive in being trees, shrubs, herbs, or lianas with *opposite, usually simple* (rarely pinnate) *leaves*, a *cymose inflorescence*, the flowers usually *epigynous*, with a *4–5-merous perianth*, 5 [2,4] stamens, and 2.5 [8] carpels, *not all fertile in some taxa*, the fruit a berry, capsule, or drupe.

K (5) [(4)] **C** (5) [(4)] **A** 5 [2,4] **G** (2-5) [(-8)], inferior.

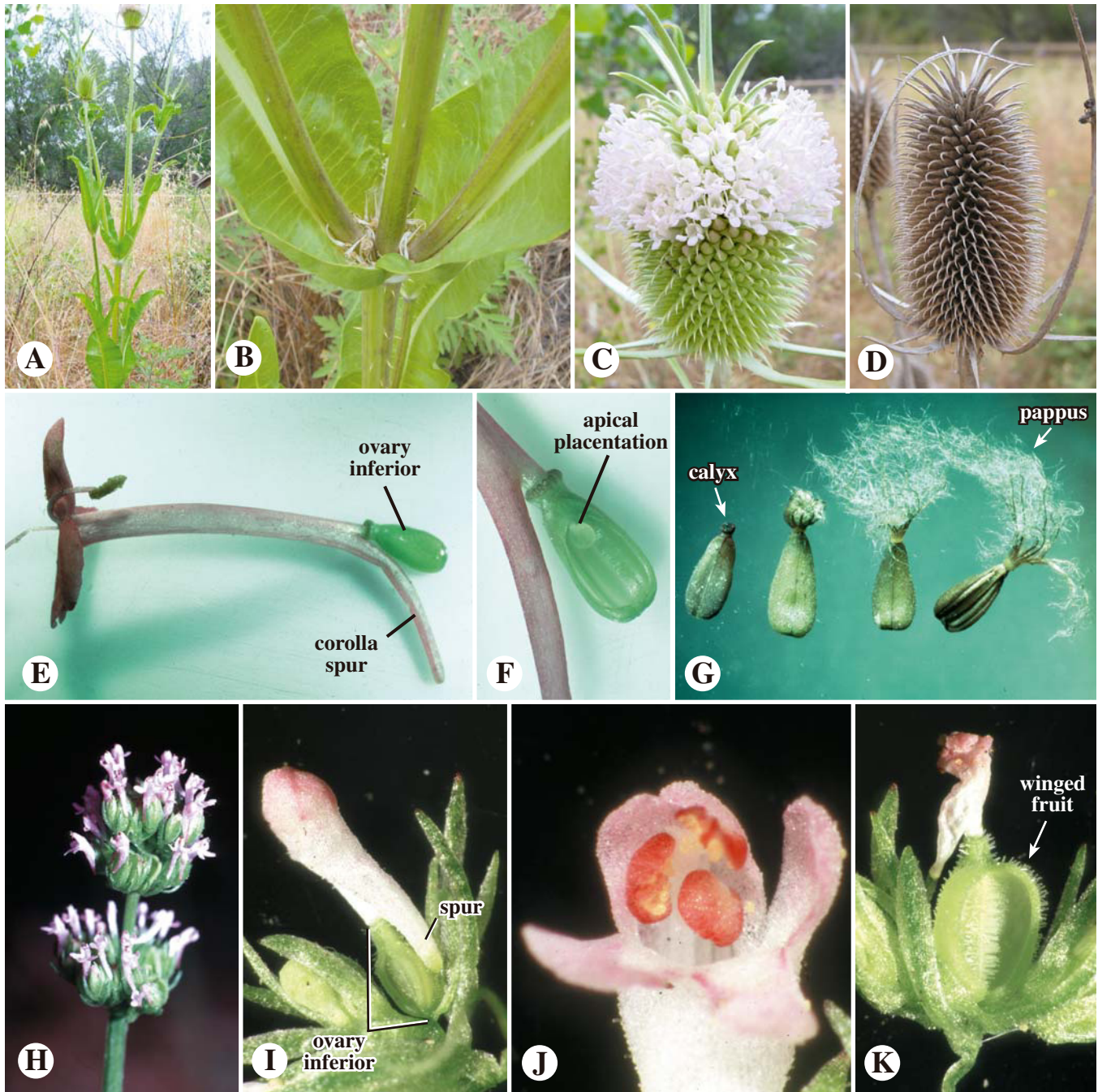


FIGURE 8.85 DIPSACALES. A–D. Dipsacaceae, *Dipsacus sativus*, teasel. A. Plant habit, an herb. B. Opposite, connate-perfoliate, water-collecting leaves, functioning to stop crawling insects. C. Inflorescence, an involucre head. D. Infructescence, with persistent, woody flower bracts. E–K. Valerianaceae. E–G. *Centranthus ruber*, red valerian. E. Whole flower, with sympetalous corolla and spur, solitary stamen, and inferior ovary. F. Ovary longitudinal section, showing apical placentation. G. Series of fruit development (left to right), showing maturation of pappus from calyx. H–K. *Plectritis ciliosa*. H. Inflorescence. I. Flower with inferior ovary and corolla spur. J. Apex of ve-lobed corolla, showing three epipetalous stamens. K. Winged fruit, beneath withering perianth/androecium.

Dipsacaceae — Teasel family (from *dipsa*, thirst, relative to water-collecting, in reference to the water collecting leaf bases). 11 genera / 290 species. (Figure 8.85A D)

The Dipsacaceae consist of herbs or shrubs. The **leaves** are simple, opposite or whorled, and exstipulate. The **inflorescence** is a head or raceme of cyme units, subtended by an involucre. The **flowers** are bisexual, usually bracteate, forming an epicalyx in some. The **calyx** is synsepalous or bristle-like with 4-5 [0,10] lobes or elements. The **corolla** is sympetalous with 4-5 imbricate lobes. The **stamens** are 4, alternipetalous, and epipetalous. **Anthers** are longitudinally dehiscent. The **gynoecium** is syncarpous, with an inferior ovary, 2 carpels, and 1 functional locule. The **style** is solitary. **Placentation** is apical; the solitary **ovule** is anatropous and unitegmic. A **nectary** is present, consisting of an annular disk at base of style. The **fruit** is an achene, usually enclosed by the epicalyx and with a terminal, persistent calyx. The **seeds** have an oily endosperm.

The Dipsacaceae are distributed in Eurasia and Africa. Economic importance includes some cultivated ornamentals.

The Dipsacaceae are distinctive in being herbs or shrubs with *opposite or whorled* leaves, an *involucrate head of cyme units*, the flowers usually *bracteate* with an *epicalyx*, the ovary *inferior with 2 carpels and 1 locule* (1 locule abortive) and a *single, apical ovule*, the fruit an *achene* usually with a *persistent epicalyx and calyx*.

K (4-5) [0, 10] **C** (4-5) **A** 4 **G** (2), inferior.

Valerianaceae — Valerian family (Latin *valere*, to be strong, or Valeria, a Roman province where plant found, or perhaps

after Valerianus, Roman emperor of 3rd century A.D.). 10 genera / 300 species. (Figure 8.85E K)

The Valerianaceae consist of herbs, rarely shrubs. The **leaves** are simple or pinnate, opposite, and exstipulate. The **inflorescence** is composed of cyme units. The **flowers** are usually bisexual, bracteate. The **perianth** is biseriolate or uniseriate. The **calyx** is absent, composed of teeth, or of 5 sepals, forming an accrescent pappus in some taxa. The **corolla** is sympetalous with 5 [3,4] imbricate lobes and a basal, nectariferous spur, actinomorphic or zygomorphic, bilabiate in some. The **stamens** are 1-4 (less than the number of corolla lobes), whorled, epipetalous. **Anthers** are versatile and longitudinally dehiscent. The **gynoecium** is syncarpous, with an inferior ovary, 3 carpels, and 1 functional locule (2 locules abortive). The **style** is solitary. **Placentation** is apical; the **ovule** is solitary, anatropous and unitegmic. The **fruit** is an achene, sometimes winged, with a plumose, pappuslike calyx in some. The **seeds** have an oily endosperm.

The Valerianaceae are mostly worldwide in distribution. Economic importance includes some cultivated ornamentals (e.g., *Centranthus*) and minor edible, medicinal, or essential oil plants.

The Valerianaceae are distinctive in being herbs, rarely shrubs, with *opposite* leaves, a *sympetalous, spurred* corolla, 1-4 stamens, and a *trilocellate, inferior* ovary with *1 functional locule* (2 locules abortive) and a *single, apical ovule*, the fruit an *achene*, with a *pappuslike calyx* in some members.

K 0-5 **C** (5) [(3,4)] **A** 1-4 **G** (3), inferior.

REVIEW QUESTIONS

GENERAL AND BASAL EUDICOTS: RANUNCULALES AND PROTEALES

1. Name and describe the major apomorphy of the Eudicots.
2. Name three families in the order Ranunculales.
3. What is distinctive about the perianth of the Berberidaceae?
4. What is the corolla cycle and placentation of the Papaveraceae?
5. What economically important member of the Papaveraceae has shaped human history?
6. What is the etymology of the root name for Ranunculaceae?
7. What is the gynoeceal fusion of the Ranunculaceae?
8. Name the three families of the Proteales.
9. For the Nelumbonaceae what is the family common name, plant habitat, leaf base/shape, floral formula, and placentation?
10. What is the fruit type of the Nelumbonaceae and what accessory tissue is part of this fruit?
11. What is the single genus and common name of the Platanaceae?
12. What is the stipule type, bud type, inflorescence type, flower sex, and fruit type of the Platanaceae?
13. How are the Nelumbonaceae and Platanaceae similar with regard to placentation?
14. For the Proteaceae, name the perianth cycle, perianth merosity, stamen number, ovary position, and placentation.
15. In what two regions of the world are most Proteaceae found?

CORE EUDICOTS: CARYOPHYLLALES, SANTALALES, AND SAXIFRAGALES

16. What three, basal orders and two major groups comprise the Core Eudicots?
17. What carnivorous plant families are found in the Caryophyllales?
18. Within the order Caryophyllales, name three apomorphies for the Core Caryophyllales.
19. Name and give the function of the major chemical apomorphy for the Higher Caryophyllales.
20. For the Aizoaceae, what is the leaf texture and typical leaf arrangement?
21. What is the perianth cycly of the Aizoaceae, and what structures appear like petals in many members?
22. What types of photosynthesis are found in many Aizoaceae?
23. What is distinctive about the perianth cycly, locule number, and placentation of the Amaranthaceae?
24. For the Cactaceae, what is the plant habit, stem texture, and geographic distribution?
25. What are the specialized axillary meristems of cacti termed, and what do these produce?
26. What is unusual about the perianth of the Cactaceae?
27. What type of photosynthesis is found in cacti and what is its physiological significance?
28. Give the floral formula, perianth/androecial position, and placentation for the Cactaceae.
29. What is the common name of the Caryophyllaceae, and what genus denotes this common name?
30. For the Caryophyllaceae, what is distinctive about the stem nodes, leaf arrangement, corolla (petal) type, and placentation?
31. What is the common name of the Polygonaceae, and what species denotes this common name?
32. What two perianth morphologies occur in the Polygonaceae?
33. What is the name of the distinctive stipular structures found in some Polygonaceae?
34. What is the ovule position and type of the Polygonaceae?
35. What former family is included within the Santalaceae, s.l. (sensu APG II 2003)?
36. What is the root type and plant nutrition of the Santalaceae, s.l.?
37. Name two economically important members of the Santalaceae.
38. What is the leaf arrangement and typical leaf texture of the Crassulaceae?
39. For the Crassulaceae, give the photosynthetic mechanism, inflorescence type, perianth cycly, gynoecium fusion, and fruit type.
40. Name a few common cultivars of the Crassulaceae.
41. For the Saxifragaceae, what is the leaf arrangement, and what is the variation of gynoecial fusion and ovary position?

ROSIDS: GERANIALES AND MYRTALES

42. What are some common features of members of the Rosids, sensu APG II, 2003?
43. Name five common families of the Rosids.
44. In the Geraniaceae, what is distinctive about the nectary position?
45. What is distinctive about the fruit type of the tribe Geranieae of the Geraniaceae?
46. What is distinctive about the leaf arrangement and venation in the Melastomataceae?
47. What is the ovary position, anther dehiscence, and anther connective form of the Melastomataceae?
48. How can the Myrtaceae often be recognized with respect to leaf structure?
49. What is the perianth/androecial position and stamen number of the Myrtaceae?
50. Name two spices, one fruit tree, and an important timber/pulp genus of the Myrtaceae.
51. What is the common name of the Onagraceae family?
52. Name the typical floral formula (including ovary position) of the Onagraceae.

EUROSIDS I: CUCURBITALES, FABALES, FAGALES, MALPIGHIALES, OXALIDALES, AND ROSALES

53. What is the typical plant habit, plant sex, and leaf morphology of the Cucurbitaceae.
54. Name the typical ovary position, perianth/androecial position, carpel number, and placentation of the Cucurbitaceae.
55. Name three economically important members of the Cucurbitaceae.
56. What is the alternate, traditional/classical name for the Fabaceae family?
57. What is distinctive about the typical gynoecial fusion, carpel number, placentation, and fruit type of the Fabaceae?

58. Prepare a table of the three subfamilies of the Fabaceae and for each listing the (a) flower symmetry; (b) position of the median posterior petal (where pertinent); (c) stamen number; and (d) stamen fusion.
59. Describe in detail a papilionaceous flower and indicate the subfamily (giving both acceptable names) having this type of flower.
60. Name several economically important members of the Fabaceae.
61. What is distinctive about the calyx of the Polygalaceae, and what other family and subfamily do the flowers superficially resemble?
62. What is the common name, plant sex, inflorescence type, and fruit type of the Betulaceae?
63. How does the Fagaceae differ from the Betulaceae?
64. For the Fagaceae what is distinctive about the (a) male inflorescence? (b) fruit accessory part?
65. What is the plant sex of members of the Euphorbiaceae?
66. What two groups (typically treated as subfamilies) of the Euphorbiaceae yield a latex?
67. What is the typical carpel and locule number in the Euphorbiaceae?
68. What is the inflorescence type of the Euphorbioideae?
69. Name three (scientific and common name) economically important members of the Euphorbiaceae.
70. Which Euphorbiaceae are important members of plant communities in southern Africa?
71. What is the common name of the Passifloraceae?
72. What two floral features are distinctive in the Passifloraceae?
73. What is the typical floral formula of the Violaceae?
74. Many members of the Violaceae with zygomorphic flowers have the anterior petal modified into what structure and for what function?
75. What is the anther orientation and placentation of the Violaceae?
76. What are the diagnostic features of the Oxalidaceae?
77. Name three diagnostic features of the Moraceae.
78. What are three economically important members of the Moraceae?
79. What is unusual about the roots in some members of the family Rhamnaceae?
80. What is the significance of *Ziziphus spina-christi*?
81. What is the characteristic perianth/androecial position and stamen position of the Rhamnaceae?
82. Name the traditional subfamilies of the Rosaceae and note how they differ with respect to gynoeceal fusion, ovary position, and fruit type.
83. Name several economically important members (scientific and common names) of the Rosaceae.

EUROSIDS II: BRASSICALES, MALVALES, AND SAPINDALES

84. What is an apomorphy for most members of the Brassicales?
85. What is the alternate, traditional/classical name for the Brassicaceae family? the common name?
86. What is the corolla type for the Brassicaceae?
87. What is the stamen arrangement of the Brassicaceae?
88. How do the Brassicaceae differ from the Capparaceae and Cleomaceae with respect to placentation and fruit morphology?
89. Name several economically important members of the Brassicaceae.
90. Name four putative apomorphies for the Malvaceae, s.l.
91. What former three families are now included as part of the family Malvaceae?
90. For the Malvoideae and Bombacoideae together, name the (a) stamen fusion; (b) anther type.
92. What is the name of the specialized bracts that subtend the calyx in many Malvaceae?
93. What are the common names of *Gossypium* spp., *Theobroma cacao*, and *Cola nitida*?
94. Name the common name of and several economically important members of the family Anacardiaceae.
95. How is the Anacardiaceae distinctive with regard to (a) nectaries; (b) anatomy?
96. Name several economically important members of the family Rutaceae.
97. How is the Rutaceae distinctive with regard to (a) nectaries; (b) glandular secretions?
98. How are many Sapindaceae distinctive with regard to (a) leaf morphology; (b) nectaries; (c) chemistry?
99. What former family is now included within the Sapindaceae?

ASTERIDS: CORNALES AND ERICALES

100. Name three potential apomorphies for the Asterids.
101. What is the common name of the Cornaceae, and how is the family distinctive with regard to (a) leaf arrangement; (b) inflorescence types; (c) ovary position; and (d) fruit type?
102. What is the common name of the Ericaceae, and how is the family distinctive with regard to (a) associated soil chemistry; (b) leaf morphology; and (c) leaf duration?
103. What is the anther dehiscence of many Ericaceae and how do these anthers develop?
104. Name some economically important members of the Ericaceae.
105. What is the common name of the Fouquieriaceae, and what is the (a) perianth cycle; (b) corolla fusion; (c) carpel number?
106. Describe in detail the shoot and leaf morphology of the Fouquieriaceae. How are these adaptive?
107. What is the common name and floral formula of the Polemoniaceae?

EUASTERIDS I: BORAGINACEAE, HYDROPHYLLACEAE, GENTIANALES, LAMIALES, AND SOLANALES

108. For the Boraginaceae, s.s., what is the inflorescence unit, corolla fusion, and corolla symmetry?
109. Describe the gynoecial morphology (including carpel, locule, and ovule number), style position, and fruit type for the Boraginaceae, s.s.
110. How does the Hydrophyllaceae resemble the Boraginaceae in corolla symmetry, stamen merosity, ovary position, and carpel number?
111. How is the Hydrophyllaceae different from the Boraginaceae in ovary shape, style position, and fruit type?
112. What class of chemical compounds are found in the tissues of the Apocynaceae?
113. What is unusual about the gynoecial fusion in many members of the Apocynaceae?
114. Describe the distinctive androecium and pollen fusion type found in the Asclepiadoids (milkweeds).
115. Name two medicinally important members of the Apocynaceae, including the compounds used and diseases these are used to treat.
116. Review the diagnostic features of the gentian family (Gentianeae), noting the floral nectaries and glands.
117. Describe the leaf arrangement, stipular morphology, and ovary position of the Rubiaceae.
118. Name economically important members of the Rubiaceae with respect to uses as a medicine, beverage, and sexual stimulant.
119. For the Acanthaceae, describe the leaf arrangement, flower symmetry, and modified funiculus.
120. For the Bignoniaceae, describe the plant habit, leaf arrangement, flower symmetry, and seed morphology and nutrition.
121. How can the Lamiaceae be recognized with respect to (a) plant chemistry; (b) stem shape; (c) leaf type and arrangement; (d) corolla type; (e) style position (in most); and (f) fruit type (in most)?
122. What are the general characteristics of the traditionally defined Scrophulariaceae s.l.?
123. Name at least four families that the traditionally defined Scrophulariaceae s.l. has been subdivided into.
124. What is distinctive about the Orobanchaceae, as defined here? the Orobanchoids within that family?
125. What traditionally defined families are now included within the (expanded) Plantaginaceae?
126. What relatively large, mostly western North American genus is now classified in the Phymaceae s.l.?
127. How are the Convolvulaceae and Solanaceae similar and how different?
128. Name an important agricultural species in the Convolvulaceae.
129. For the Solanaceae, what is the (a) flower symmetry; (b) corolla aestivation (in bud); (c) stamen number; (d) ovary position; (e) carpel number; and (f) ovule number (per carpel)?
130. Name three members of the Solanaceae of great economic importance.

EUASTERIDS II: APIALES, ASTERALES, AND DIPSACALES

131. Give the common name and list three economically important members of the family Apiaceae.
132. For the Apiaceae, what is the (a) leaf base; (b) inflorescence type; (c) ovary position; (d) fruit type?
133. How does the Araliaceae resemble the Apiaceae? How does it differ?
134. What is the alternate traditional/classical name for the Asteraceae?
135. What is the inflorescence type of the Asteraceae? On what criteria can these be subdivided?

136. Define (a) involucre; (b) phyllary; (c) chaff; (d) pappus.
137. Name and describe the three corolla types of the Asteraceae.
138. For the Asteraceae, what is the (a) stamen fusion; (b) ovary position; (c) fruit type?
139. Name two economically important members of the Asteraceae (used for food).
140. Name the diagnostic features of the Goodeniaceae.
141. Name two alternative ways to classify the Dipsacales.
142. What is the common name of the Caprifoliaceae?
143. For the Caprifoliaceae, name the leaf arrangement, inflorescence type, and ovary position.
144. What is distinctive about the carpel number, locule number, and fruit type in the Dipsacaceae?
145. What is distinctive about the carpel number, locule number, and corolla type in the Valerianaceae?

EXERCISES

1. Select a family of eudicots and learn everything you can about it. Perform a literature search (e.g., family name + systematics) on journal articles published in the last five years. Consult family descriptions, recent data on phylogenetic relationships, and information on intrafamilial groupings.
2. From this same family, collect living material of an exemplar. Describe this species in detail, using the character list of Appendix 1 as a guide (see Chapter 9). Illustrate the vegetative and reproductive parts (see Appendix 2).
3. Assimilate all of your information in a written report and computerized slide show and present it.

REFERENCES FOR FURTHER STUDY

GENERAL REFERENCES ON ANGIOSPERM RELATIONSHIPS AND EVOLUTION

- Angiosperm Phylogeny Group. 1998. An ordinal classification for the families of flowering plants. *Annals of the Missouri Botanical Garden* 85: 531-553.
- Angiosperm Phylogeny Group. 2003. An update of the Angiosperm Phylogeny Group classification for the orders and families of flowering plants: APG II. *Botanical Journal of the Linnean Society* 141: 399-436.
- Barkman, T. J., G. Chenery, J. R. McNeal, J. Lyons-Weiler, W. J. Ellisens, G. Moore, A. D. Wolfe, and C. W. dePamphilis. 2000. Independent and combined analyses of sequences from all three genomic compartments converge on the root of flowering plant phylogeny. *Proceedings of the National Academy of Sciences of the United States of America* 97: 13166-13171.
- Behnke, H. D. 1972. Sieve-tube plastids in relation to Angiosperm systematics: an attempt towards a classification by ultrastructural analysis. *Botanical Review* 38: 155-197.
- Borsch, T., K. W. Hilu, D. Quandt, V. Wilde, C. Neinhuis, and W. Barthlott. 2003. Noncoding plastid trnT-trnF sequences reveal a well resolved phylogeny of basal angiosperms. *Journal of Evolutionary Biology* 16: 558-576.
- Chase, M. W., D. E. Soltis, R. G. Olmstead, D. Morgan, D. H. Les, B. D. Mishler, M. R. Duvall, R. A. Price, H. G. Hills, Y.-L. Qiu, K. A. Kron, J. H. Rettig, E. Conti, J. D. Palmer, J. R. Manhart, K. J. Sytsma, H. J. Michaels, W. J. Kress, K. G. Karol, W. D. Clark, M. Hedrøn, B. S. Gaut, R. K. Jansen, K.-J. Kim, C. F. Wimpee, J. F. Smith, G. R. Furnier, S. H. Strauss, Q.-Y. Xiang, G. M. Plunkett, P. S. Soltis, S. M. Swensen, S. E. Williams, P. A. Gadek, C. J. Quinn, L. E. Eguiarte, E. Golenberg, G. H. Learn Jr., S. W. Graham, S. C. H. Barrett, S. Dayanandan, and V. A. Albert. 1993. Phylogenetics of seed plants: an analysis of nucleotide sequences from the plastid gene rbcL. *Annals of the Missouri Botanical Garden* 80: 528-580.
- Chase, M. W., M. F. Fay, and V. Savolainen. 2000. Plant systematics: a half-century of progress (1950-2000) and future challenges: higher-level classification in the angiosperms: new insights from the perspective of DNA sequence data. *Taxon* 49: 685-704.
- Graham, S. W., and R. G. Olmstead. 2000. Utility of 17 chloroplast genes for inferring the phylogeny of the basal angiosperms. *American Journal of Botany* 87: 1712-1730.
- Hilu, K. W., Borsch, T., Muller, K., Soltis, D. E., Soltis, P. S., Savolainen, V., Chase, M. W., Powell, M. P., Alice, L. A., Evans, R., Sauquet, H., Neinhuis, C., Slotta, T. A. B., Rohwer, J. G., Campbell, C. S., and Chatrou, L. W. 2003. Angiosperm phylogeny based on matK sequence information. *American Journal of Botany* 90: 1758-1766.
- Kubitzki, K., Rohwer, J. G., and Bittrich, V. 1993. *The Families and Genera of Vascular Plants. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families.* Springer-Verlag, Berlin.
- Kubitzki, K., and C. Bayer. 2002. *The Families and Genera of Vascular Plants. V. Flowering Plants. Dicotyledons: Malvales, Capparales, and non-betain Caryophyllales.* Springer, Berlin.
- Kubitzki, K. 2004. *The Families and Genera of Vascular Plants. VI. Flowering Plants. Dicotyledons. Celastrales, Oxalidales, Rosales, Cornales, Ericales.* Springer, Berlin.

- Nandi, O. I., M. W. Chase, and P. K. Endress. 1998. A combined cladistic analysis of angiosperms using *rbcL* and non-molecular data sets. *Annals of the Missouri Botanical Garden* 85: 137-212.
- Savolainen, V., Chase, M. W., Hoot, S. B., Morton, C. M., Soltis, D. E., Bayer, C., Fay, M. F., de Bruijn, A. Y., Sullivan, S., and Qiu, Y.-L. 2000a. Phylogenetics of flowering plants based on combined analysis of plastid *atpB* and *rbcL* sequences. *Systematic Biology* 49: 306-362.
- Soltis, D. E., P. S. Soltis, D. L. Nickrent, L. A. Johnson, W. J. Hahn, S. B. Hoot, J. A. Sweere, R. K. Kuzoff, K. A. Kron, M. W. Chase, S. M. Swensen, E. A. Zimmer, S.-M. Chaw, L. J. Gillespie, W. J. Kress, and K. J. Sytsma. 1997. Angiosperm phylogeny inferred from 18S ribosomal DNA sequences. *Annals of the Missouri Botanical Garden* 84: 1-49.
- Soltis, D. E., P. S. Soltis, M. W. Chase, M. E. Mort, D. C. Albach, M. Zanis, V. Savolainen, W. H. Hahn, S. B. Hoot, M. F. Fay, M. Axtell, S. M. Swensen, L. M. Prince, W. J. Kress, K. C. Nixon, and J. S. Farris. 2000. Angiosperm phylogeny inferred from 18S rDNA, *rbcL*, and *atpB* sequences. *Botanical Journal of the Linnean Society* 133(4): 381-461.
- Takhtajan, A. L. 1991. *Evolutionary Trends in Flowering Plants*. Columbia University Press, New York.
- Thorne, Robert F. 2000. The classification and geography of the flowering plants: Dicotyledons of the class Angiospermae: (Subclasses Magnoliidae, Ranunculidae, Caryophyllidae, Dilleniidae, Rosidae, Asteridae, and Lamiidae). *Botanical Review* 66(4): 441-647.

WEB SITES

- Reveal, J. L. (2001 onwards). Indices Nominum Supragenericorum Plantarum Vascularium. Alphabetical Listing by Family of Validly Published Suprageneric Names. <http://www.inform.umd.edu/PBIO/fam/revfam.html>
An index of accepted family names.
- Royal Botanic Garden Vascular Plant Families and Genera Database <http://www.rbgekew.org.uk/data/vascplnt.html>
A searchable database taken from *Vascular Plant Families and Genera*, compiled by R. K. Brummitt and published by the Royal Botanic Gardens, Kew, in 1992.
- Stevens, P. F. (2001 onwards). Angiosperm Phylogeny Website. Version 5, May 2004. <http://www.mobot.org/MOBOT/research/APweb>
An excellent graphic representation of the Angiosperm Phylogeny Group classification, up-to-date cladograms, family characteristics, references, and apomorphies.

ANGIOSPERM FAMILY DESCRIPTIONS AND ILLUSTRATIONS

- Cronquist, A. 1981. *An Integrated System of Classification of Flowering Plants*. Columbia University Press, New York.
- Heywood, V. H. 1985. *Flowering Plants of the World*. Prentice-Hall, Englewood Cliffs, NJ.
- Mabberly, D. J. 1997. *The Plant-Book*. 2nd ed. Cambridge University Press, Cambridge, UK.
- Takhtajan A. 1997. *Diversity and Classification of Flowering Plants*. Columbia University Press, New York.
- Zomlefer, W. B. 1994. *Guide to Flowering Plant Families*. University of North Carolina Press, Chapel Hill.

REFERENCES ON SPECIFIC EUDICOT GROUPS

- Albach, D. C., P. S. Soltis, D. E. Soltis, and R. G. Olmstead. 2001. Phylogenetic analysis of Asterids based on sequences of four genes. *Annals of the Missouri Botanical Garden* 88: 163-212.
- Bayer, C. 1999. The bicolor unit-homology and transformation of an inflorescence structure unique to core Malvales. *Plant Systematics and Evolution* 214: 187-198.
- Bayer, C., M. F. Fay, A. Y. D. Bruijn, V. Savolainen, C. M. Morton, K. Kubitzki, W. S. Alverson, and M. W. Chase. 1999. Support for an expanded family concept of Malvaceae within a circumscribed order Malvales: a combined analysis of plastid *atpB* and *rbcL* DNA sequences. *Botanical Journal of the Linnean Society* 129:267-303.
- Beardsley, P. M., and R. G. Olmstead. 2002. Redefining Phrymaceae: the placement of *Mimulus*, tribe Mimuleae, and *Phryma*. *American Journal of Botany* 89: 1093-1102.
- Bremer, K. 1994. *Asteraceae: Cladistics and Classification*. Timber Press, Portland, OR.
- Bremer, K. 1996. Major clades and grades of the Asteraceae. In: Hind, D. J. N., and H. J. Beentje (eds.), *Compositae: Systematics*. Pp. 1-7. *Proceedings of the International Compositae Conference*, Kew. Royal Botanic Gardens, Kew.
- Bremer, K., A. Backlund, B. Sennblad, U. Swenson, K. Andreassen, M. Hjertson, J. Lundberg, M. Backlund, and B. Bremer. 2001. A phylogenetic analysis of 100+ genera and 50+ families of euasterids based on morphological and molecular data with notes on possible higher level morphological synapomorphies. *Plant Systematics and Evolution* 229: 137-169.
- Bremer, B., K. Bremer, N. Heidari, P. Erixon, R. G. Olmstead, A. A. Anderberg, M. K. Ilersj, and E. Barkhordarian. 2002. Phylogenetics of asterids based on 3 coding and 3 non-coding chloroplast DNA markers and the utility of non-coding DNA at higher taxonomic levels. *Molecular Phylogenetics and Evolution* 24: 274-301.
- Bruneau, A., F. Forest, P. S. Herendeen, B. B. Klitgaard, and G. P. Lewis. 2001. Phylogenetic relationships in the Caesalpinoideae (Leguminosae) as inferred from chloroplast *trnL* intron sequences. *Systematic Botany* 26: 487-514.
- Cantino, P. D. 1992. Evidence for a polyphyletic origin of the Labiales. *Annals of the Missouri Botanical Garden* 79: 361-379.
- Cantino, P. D. 2004. Phrymaceae. In: Kadereit, J. (ed.), *The Families and Genera of Vascular Plants. VII. Flowering Plants: Dicotyledons: Lamiales (except Acanthaceae including Avicenniaceae)*. Pp. 323-326. Springer, Berlin.
- Chandler, G. T., and G. M. Plunkett. 2004. Evolution in Apiales: nuclear and chloroplast markers together in (almost) perfect harmony. *Botanical Journal of the Linnean Society* 144: 123-147.

- Chase, M. W., C. M. Morton, and J. A. Kallunki. 1999. Phylogenetic relationships of Rutaceae: a cladistic analysis of the subfamilies using evidence from *rbcL* and *atpB* sequence variation. *American Journal of Botany* 86: 1191-1199.
- Chase, M. W., S. Zmarzty, M. D. Lled, K. J. Wurdack, S. M. Swensen, and M. F. Fay. 2002. When in doubt, put it in Flacourtiaceae: A molecular phylogenetic analysis based on plastid L DNA sequences. *Kew Bulletin* 57: 141-181.
- Clausing, G., and S. S. Renner. 2001. Molecular phylogenetics of Melastomataceae and Memecylaceae: implications for character evolution. *American Journal of Botany* 88: 486-498.
- Conti, E., A. Litt, and K. J. Sytsma. 1996. Circumscription of Myrtales and their relationships to other Rosids: evidence from *rbcL* sequence data. *American Journal of Botany* 83: 221-233.
- Conti, E., A. Litt, P. G. Wilson, S. A. Graham, B. G. Briggs, L. A. S. Johnson, and K. J. Sytsma. 1998. Interfamilial relationships in Myrtales: molecular phylogeny and patterns of morphological evolution. *Systematic Botany* 22: 629-647.
- Donoghue, M.J., T. Eriksson, P. A. Reeves, and R. G. Olmstead. 2001. Phylogeny and phylogenetic taxonomy of Dipsacales, with special reference to *Sinadoxa* and *Tetradoxa* (Adoxaceae). *Harvard Papers in Botany* 6: 459-479.
- Doyle, J., J. A. Chappill, C. D. Bailey, and T. Kajita. 2000. Towards a comprehensive phylogeny of legumes: Evidence from *rbcL* sequences and non-molecular data. In: Herendeen, P. S., and Bruneau, A. (eds.), *Advances in Legume Systematics, Part 9*, Pp. 1-20. Royal Botanic Gardens, Kew.
- Endress, M. E., and P. V. Bruyns. 2000. A revised classification of the Apocynaceae s. l. *Botanical Review* 66: 1-56.
- Fischer, E. 2004. Scrophulariaceae. In: Kadereit, J. (ed.), *The Families and Genera of Vascular Plants. VII. Flowering Plants. Dicotyledons. Lamiales (except Acanthaceae including Avicenniaceae)*. Pp. 333-432. Springer, Berlin.
- Fishbein, M., C. Hirsch-Jetter, D. E. Soltis, and L. Hufford. 2001. Phylogeny of Saxifragales (Angiosperms, Eudicots): analysis of a rapid, ancient radiation. *Systematic Biology* 50: 817-847.
- Forest, F., A. Bruneau, J. Hawkins, T. Kajita, J. J. Doyle, and P. R. Crane. 2002. The sister of the Leguminosae revealed: Phylogenetic relationships in the Fabales determined using *trnL* and *rbcL* sequences. In: *Botany 2002: Botany in the Curriculum, Abstracts*. Madison, WI (<http://www.botany2002.org>).
- Gadek, P. A., E. S. Fernando, C. J. Quinn, S. B. Hoot, T. Terrazas, M. C. Sheahan, and M. W. Chase. 1996. Sapindales: molecular delimitation and infraordinal groups. *American Journal of Botany* 83: 802-811.
- Geuten, K., E. Smets, P. Schols, Y.-M. Yuan, S. Janssens, P. K. pfer, and N. Pyck. 2004. Conflicting phylogenies of balsamoid families and the polytomy in Ericales: Combining data in a Bayesian framework. *Molecular Phylogenetics and Evolution* 31: 711-729.
- Gottschling, M., H. H. Hilger, M. Wolf, and N. Diane. 2001. Secondary structure of the ITS1 transcript and its application in a reconstruction of the phylogeny of Boraginales. *Plant Biology* 3: 629-636.
- Hall, J. C., K. J. Sytsma, and H. H. Iltis. 2002. Phylogeny of Capparaceae and Brassicaceae based on chloroplast sequence data. *American Journal of Botany* 89: 1826-1842.
- Ham, R. C., H. J. van, and H. t Hart. 1998. Phylogenetic relationships in the Crassulaceae inferred from chloroplast DNA restriction-site variation. *American Journal of Botany* 85: 123-134.
- Harley, R., S. Atkins, A. L. Budantsev, P. D. Cantino, B. J. Conn, M. Grayer, M. M. Harley, R. De Kok, T. Krestovskaya, R. Morales, A. J. Paton, O. Ryding, and T. Upson. 2004. Labiatae. In: Kadereit, J. (ed.), *The Families and Genera of Vascular Plants. VII. Flowering Plants. Dicotyledons. Lamiales (except Acanthaceae including Avicenniaceae)*. Pp. 167-275. Springer, Berlin.
- Hoot, S. B. 1991. Phylogeny of the Ranunculaceae based on epidermal characters and micromorphology. *Systematic Botany* 16: 741-755.
- Hoot, S. B. 1995. Phylogeny of the Ranunculaceae based on preliminary *atpB*, *rbcL* and 18S ribosomal DNA sequence data. In: Jensen, U., and J. W. Kadereit (eds.), *Systematics and Evolution of the Ranunculiflorae*. Pp. 241-251. Springer, Vienna. [Plant Syst. Evol. Suppl. 9.]
- Hoot, S. B., and P. R., Crane. 1995. Inter-familial relationships in the Ranunculidae based on molecular systematics. In: Jensen, U., and J. W. Kadereit (eds.), *Systematics and Evolution of the Ranunculiflorae*. Springer, Vienna. Pp. 119-131 [Plant Syst. Ev ol. Suppl. 9.]
- Hoot, S. B., and A. W. Douglas. 1998. Phylogeny of the Proteaceae based on *atpB* and *atpB-rbcL* intergenic spacer regions. *Australian Systematic Botany* 11: 301-320.
- Hoot, S. B., J. W. Kadereit, F. R. Blattner, K. B. Jork, A. E. Schwarzbach, and P. R. Crane. 1997. Data congruence and phylogeny of the Papaveraceae s.l. based on four data sets: *atpB* and *rbcL* sequences, *trnK* restriction sites, and morphological characters. *Systematic Botany* 22:575-590.
- Hoot, S. B., S. Magallon-Puebla, and P. R. Crane. 1999. Phylogeny of basal eudicots based on three molecular data sets: *atpB*, *rbcL* and 18S nuclear ribosomal DNA sequences. *Annals of the Missouri Botanical Garden* 86: 119-131.
- Kadereit, J. W., F. R. Blattner, K. B. Jork, and A. Schwarzbach. 1994. Phylogenetic analysis of the Papaveraceae s.l. (inc. Fumariaceae, Hypochoeraceae, and Pteridophyllum) based on morphological characters. *Botanische Jahrb cher f r Systematik* 116: 361-390.
- Kadereit, J. W., F. R. Blattner, K. B. Jork, and A. Schwarzbach. 1995. The phylogeny of the Papaveraceae sensu lato: morphological, geographical, and ecological implications. In: Jensen, U., and J. W. Kadereit (eds.), *Systematics and Evolution of the Ranunculiflorae*. Pp. 133-145. Springer, Vienna. [Plant Syst. Evol. Suppl. 9.]
- Kadereit, J. (ed.). 2004. *The Families and Genera of Vascular Plants. VII. Flowering Plants. Dicotyledons. Lamiales (except Acanthaceae including Avicenniaceae)*. Springer, Berlin.
- Klak, C., A. Khunou, G. Reeves, and T. Hedderson. 2003. A phylogenetic hypothesis for the Aizoaceae (Caryophyllales) based on four plastid DNA regions. *American Journal of Botany* 90: 1433-1445.
- Kron, K. A., W. S. Judd, P. F. Stevens, D. M. Crayn, A. A. Anderberg, P. A. Gadek, C. J. Quinn, and J. L. Luteyn. 2002. A phylogenetic classification of Ericaceae: molecular and morphological evidence. *Botanical Review* 68: 335-423.

- Kubitzki, K. 1993. Platanaceae. In: Kubitzki, K., J. G. Rohwer, and V. Bittrich (eds.). The Families and Genera of Vascular Plants. II. Flowering plants: dicotyledons, magnoliid, hamamelid, and caryophyllid families. Pp. 521-522. Springer-Verlag, Berlin, New York.
- Kubitzki, K., and M. W. Chase. 2002. Introduction to Malvales. In: The Families and Genera of Vascular Plants. V. Flowering Plants. Dicotyledons. Malvales, Capparales and Non-betain Caryophyllales. K. Kubitzki and C. Bayer (eds.). Pp. 12-16. Springer, Berlin.
- Lamb Frye, A. S., and K. A. Kron. 2003. *RbcL* phylogeny and character evolution in Polygonaceae. *Systematic Botany* 28:326-332.
- Levin, R. A., W. L. Wagner, P. C. Hoch, M. Nepokroeff, J. C. Pires, E. A. Zimmer, and K. J. Sytsma. 2003. Family-level relationships of Onagraceae based on chloroplast *rbcL* and *ndhF* data. *American Journal of Botany* 90: 107-115.
- Li, R.-Q., Z.-D. Chen, Y.-P. Hong, and A.-M. Lu. 2002. Phylogenetic relationships of the higher hamamelids based on chloroplast *trnL-F* sequences. *Acta Botanica Sinica* 44: 1462-1468.
- Lis-Balchin, M. (ed.). 2002. *Geranium* and *Pelargonium*: The Genera *Geranium* and *Pelargonium*. Taylor and Francis, London.
- Lundberg, J. and K. Bremer. 2003. A phylogenetic study of the order Asterales using one morphological and three molecular data sets. *International Journal of Plant Sciences* 164: 553-578.
- Manos, P. S., and K. P. Steele. 1997. Phylogenetic analysis of higher Hamamelididae based on plastid sequence data. *American Journal of Botany* 84: 1407-1419.
- Manos, P. S., Z.-K. Zhou, and C. H. Cannon. 2001. Systematics of Fagaceae: Phylogenetic tests of reproductive trait evolution. *International Journal of Plant Sciences* 162: 1361-1379.
- Matthews, M. L., and P. K. Endress. 2002. Comparative floral morphology and systematics in Oxalidales (Oxalidaceae, Connaraceae, Brunelliaceae, Cephalotaceae, Cunoniaceae, Elaeocarpaceae, Tremandraceae). *Botanical Journal of the Linnean Society* 140: 321-381.
- Matthews, M. L., and P. K. Endress. 2005. Comparative floral structure and systematics in Crossosomataceae, Stachyuraceae, Staphyleaceae, Aphloiaceae, Geissolomataceae, Ixerbaceae, Strasburgeriaceae). *Botanical Journal of the Linnean Society* 147: 1-46.
- McDade, L. A., S. E. Masta, M. L. Moody, and E. Waters. 2000. Phylogenetic relationships among Acanthaceae: Evidence from two genomes. *Systematic Botany* 25: 106-121.
- Meimberg, H., P. Dittrich, G. Bringmann, J. Schlauer, and G. Heubl. 2000. Molecular phylogeny of Caryophyllidae s.l. based on *matK* sequences with special emphasis on carnivorous taxa. *Plant Biology* 2: 218-228.
- Mort, M. E., D. E. Soltis, P. S. Soltis, J. Francisco-Ortega, and A. Santos-Guerra. 2001. Phylogenetic relationships and evolution of Crassulaceae inferred from *matK* sequence data. *American Journal of Botany* 88: 76-91.
- Nickrent, D. L., and V. MalØcot. 2001. A molecular phylogeny of Santalales. In: Fer, A., P. Thalouarn, D. M. Joel, L. J. Musselman, C. Parker, and J. A. C. Verkleij (eds.), *Proceedings of the 7th International Parasitic Weed Symposium*. FacultØ des Sciences, UniversitØ de Nantes, Nantes, France. See <http://www.science.siu.edu/parasitic-plants/Santalales.IPWC/Sants.IPWC.html> for an on-line version of the paper.
- Nyffeler, R. 2002. Phylogenetic relationships in the cactus family (Cactaceae) based on evidence from *trnK/matK* and *trnL-trnF* sequences. *American Journal of Botany* 89: 312-326.
- Olmstead, R. G., and P. A. Reeves. 1995. Evidence for the polyphyly of the Scrophulariaceae based on chloroplast *rbcL* and *ndhF* sequences. *Annals of the Missouri Botanical Garden* 82: 176-193.
- Olmstead, R. G., J. A. Sweere, R. E. Spangler, L. Bohs, and J. D. Palmer. 1999. Phylogeny and provisional classification of the Solanaceae based on chloroplast DNA. In: Nee, M., D. Symon, R. N. Lester, and J. P. Jessop (eds.), *Solanaceae IV: Advances in Biology and Utilization*. Pp. 111-137. Royal Botanic Gardens, Kew.
- Olmstead, Richard G., Claude W. DePamphilis, Andrea D. Wolfe, Nelson D. Young, Wayne J. Elisons, and Patrick A. Reeves. 2001. Disintegration of the Scrophulariaceae. *American Journal of Botany* 88: 348.
- Olmstead, R. G., M. L. Zihra, S. O. Grose, A. J. Eckert, and R. E. Spangler. 2002. Molecular phylogeny, evolution and classification of Bignoniaceae. In: *Botany 2002: Botany in the Curriculum*, Abstracts Madison, WI. (<http://www.botany2002.org>)
- Oxelman, B., P. Kornhall, R. G. Olmstead, and B. Bremer. 2005. Further disintegration of the Scrophulariaceae. *Taxon* (in press).
- Panero, J. L., and V. A. Funk. 2002. Toward a phylogenetic classification for the Compositae (Asteraceae). *Proceedings of the Biological Society of Washington* 115: 909-922.
- Pell, S. K., and L. Urbatsch. 2001. Tribal relationships and character evolution in the cashew family (Anacardiaceae): Inferences from three regions of the chloroplast genome. In: *Botany 2001: Plants and People*. Abstracts. Albuquerque, New Mexico. (<http://www.botany2001.org>)
- Persson, C. 2001. Phylogenetic relationships in Polygalaceae based on plastid DNA sequences from the *trnL-F* region. *Taxon* 50: 763-779.
- Plunkett, G. M., D. E. Soltis, and P. S. Soltis. 1997. Evolutionary patterns in Apiaceae: Inferences based on *matK* sequence data. *Systematic Botany* 21: 477-495.
- Porter, J. M. 1997. Phylogeny of Polemoniaceae based on nuclear ribosomal internal transcribed DNA sequences. *Aliso* 15: 57-77.
- Porter, J. M., and L. A. Johnson. 1998. Phylogenetic relationships of Polemoniaceae: inferences from mitochondrial *nad1b* intron sequences. *Aliso* 17: 157-188.
- Potter, D., F. Gao, P. E. Bortiri, S.-H. Oh, and S. Baggett. 2002. Phylogenetic relationships in Rosaceae inferred from chloroplast *matK* and *trnL-trnF* nucleotide sequence data. *Plant Systematics and Evolution* 231: 77-89.
- Prather, C. A., C. J. Ferguson, and R. K. Jansen. 2000. Polemoniaceae phylogeny and classification: Implications of sequence data from the chloroplast gene *ndhF*. *American Journal of Botany* 87: 1300-1308.
- Qui, Y.-L., M. W. Chase, S. B. Hoot, E. Conti, P. R. Crane, K. J. Sytsma, and C. R. Parks. 1998. Phylogenetics of Hamamelidae and their allies: parsimony analyses of nucleotide sequences of the plastid gene *rbcL*. *International Journal of Plant Sciences* 159: 891-905.
- Reeves, P. A., and R. G. Olmstead. 1998. Evolution of novel morphological, ecological, and reproductive traits in a clade containing *Antirrhinum*. *American Journal of Botany* 85: 1047-1056.

- Renner, S. S., A. Weerasooriya, and M. E. Olson. 2002. Phylogeny of Cucurbitaceae inferred from multiple chloroplast loci. In: Botany 2002: Botany in the Curriculum, Abstracts. Madison, WI. (<http://www.botany2002.org>)
- Richardson, J. E., M. E. Fay, Q. C. B. Cronk, and M. W. Chase. 2000. A revision of the tribal classification of Rhamnaceae. *Kew Bulletin* 55: 311-340.
- Rodman, J. E., K. G. Karol, R. A. Price, and K. J. Sytsma. 1997. Molecules, morphology, and Dahlgren's expanded order Capparales. *Systematic Biology* 21: 289-307.
- Rodman, J. E., P. S. Soltis, D. E. Soltis, K. J. Sytsma, and K. G. Karol. 1998. Parallel evolution of glucosinolate biosynthesis inferred from congruent nuclear and plastid gene phylogenies. *American Journal of Botany* 85: 997-1007.
- Savolainen, V., M. F. Fay, D. C. Albach, A. Backlund, M. v. d. Bank, K. M. Cameron, S. A. Johnson, M. D. Lledo, J. C. Pintaud, M. Powell, M. C. Sheahan, D. E. Soltis, P. S. Soltis, P. Weston, W. M. Whitten, K. J. Wurdack, and M. W. Chase. 2000b. Phylogeny of the eudicots: a nearly complete familial analysis based on *rbcL* gene sequences. *Kew Bulletin* 55: 257-310.
- Schultheis, L., and B. G. Baldwin. 1999. Molecular phylogenetics of Fouquieriaceae: evidence from nuclear rDNA ITS studies. *American Journal of Botany* 86: 578-589.
- Schwarzbach, A. E. 2004. Plantaginaceae. In: Kadereit, J. (ed.), *The Families and Genera of Vascular Plants. VII. Flowering Plants. Dicotyledons. Lamiales (except Acanthaceae including Avicenniaceae)*. Pp. 327-329. Springer, Berlin.
- Smissen, R. D., J. C. Clement, P. J. Garnock-Jones, and G. K. Chambers. 2002. Subfamilial relationships in Caryophyllaceae as inferred from 5' *ndhF* sequences. *American Journal of Botany* 89: 1336-1341.
- Soltis, D. E., and P. S. Soltis. 1997. Phylogenetic relationships in Saxifragaceae sensu lato: a comparison of topologies based on 18S rDNA and *rbcL* sequences. *American Journal of Botany* 84: 504-522.
- Soltis, D. E., R. K. Kuzoff, M. E. Mort, M. Zanis, M. Fishbein, L. Hufford, J. Koontz, and M. K. Arroyo. 2001. Elucidating deep-level phylogenetic relationships in Saxifragaceae using sequences for six chloroplastic and nuclear DNA regions. *Annals of the Missouri Botanical Garden* 88: 669-693.
- Spangler, R. E., and R. G. Olmstead. 1999. Phylogenetic analysis of the Bignoniaceae based on the cpDNA gene sequence *rbcL* and *ndhF*. *Annals of the Missouri Botanical Garden* 86: 33-46.
- Wallace, R. S., and A. C. Gibson. 2002. Evolution and systematics. In: Nobel, P. S. (ed.), *Cacti: Biology and Uses*. Pp. 1-21. University of California Press, Berkeley.
- Wen, J., G. M. Plunkett, A. D. Mitchell, and S. J. Wagstaff. 2000. The evolution of Araliaceae: a phylogenetic analysis based on ITS sequences of nuclear ribosomal DNA. *Systematic Botany* 26: 144-167.
- Wilken, D. H. 2004. Polemoniaceae. In: Kubitzki, K. (ed.), *The Families and Genera of Vascular Plants. VI. Flowering Plants. Dicotyledons. Celastrales, Oxalidales, Rosales, Cornales, Ericales*. Pp. 300-312. Springer, Berlin.
- Williamson, P. S., and E. L. Schneider. 1993. Nelumbonaceae. In: Kubitzki, K., Rohwer, J. G., and Bittrich, V. (eds.), *The Families and Genera of Vascular Plants. II. Flowering Plants: Dicotyledons, Magnoliid, Hamamelid and Caryophyllid Families*. Pp. 470-472. Springer, Berlin.
- Wilson, P. G., M. M. O'Brien, P. A. Gadek, and C. J. Quinn. 2001. Myrtaceae revisited: A reassessment of infrafamilial groups. *American Journal of Botany* 88: 2013-2025.
- Wurdack, K. J., and M. W. Chase. 2002. Phylogenetics of Euphorbiaceae s.s. using plastid (*rbcL* and *trnL-F*) DNA sequences. In: Botany 2002 Botany in the Curriculum: Integrating Research and Teaching. Abstract. Madison, WI. (<http://www.botany2002.org>)
- Xiang, Q.-Y., M. Moody, D. E. Soltis, C.-Z. Fan, and P. S. Soltis. 2002. Relationships within Cornales and circumscription of Cornaceae: *matK* and *rbcL* sequence data and effects of outgroups and long branches. *Molecular Phylogenetics and Evolution* 24: 35-57.
- Xiang, Q.-Y., S. J. Brunfeldt, and P. S. Soltis. 1998. Phylogenetic relationships of Cornaceae and close relatives inferred from *matK* and *rbcL* sequences. *American Journal of Botany* 85: 285-297.
- Xiang, Q.-Y., S. J. Brunfeldt, D. E. Soltis, and P. S. Soltis. 1997. Phylogenetic relationships in *Cornus* based on chloroplast DNA restriction sites: implications for biogeography and character evolution. *Systematic Botany* 21: 515-534.
- Zhang, L.-B., and S. S. Renner. 2003. Phylogeny of Cucurbitales inferred from seven chloroplast and mitochondrial loci. In: Botany 2003: Aquatic and Wetland Plants: Wet and Wild. Mobile, Alabama. (<http://www.botany2003.org>)
- Zhang, W.-H., H. Li, H.-B. Chen, and Y.-C. Tang. 2003. Phylogeny of the Dipsacales s.l. based on chloroplast *trnL-F* and *ndhF* sequences. *Molecular Phylogenetics and Evolution* 26:176-189.

This Page Intentionally Left Blank

III

SYSTEMATIC EVIDENCE AND DESCRIPTIVE TERMINOLOGY



This Page Intentionally Left Blank

9

PLANT MORPHOLOGY

PLANT STRUCTURE	348	In florescences	379
Plant Organs	348	In florescence Parts	379
Plant Habit	350	In florescence Position	379
Plant Habitat	350	In florescence Development	379
Roots	350	In florescence Type	380
Root Types	350	Secondary In florescences	382
Stems and Shoots	351	Specialized In florescences	382
Stem Types	352	Fruits	382
Stem Habit	354	Simple Fruit Types	384
Stem Branching Pattern	354	Aggregate Fruit Types	386
Twigs, Trunks, and Buds	355	Multiple Fruit Types	386
Leaves	356	Seeds	387
Leaf Parts	356	Seed Endosperm Type	388
Leaf Structural Types	357	Seed Germination Type	389
Leaf Type	359	GENERAL TERMINOLOGY	389
Leaf Attachment	361	Color	389
Leaf Venation	361	Size	389
Flowers	364	Number	389
Flower Parts	364	Texture	389
Flower Sex and Plant Sex	365	Fusion	389
Flower Attachment	365	Shape	390
Flower Cycle	366	Shape: Solid (Three-Dimensional)	390
Flower Symmetry	366	Shape: Plane (Two-Dimensional)	390
Flower Maturation	366	Base	392
Perianth	366	Apex	392
Perianth Parts	366	Shape Combinations	393
Perianth Arrangement/Cycle/Merosity	367	Margin	393
Perianth Fusion	369	Division	394
Perianth Type	369	Disposition	396
Perianth Aestivation	370	Position	396
Androecium	371	Arrangement	396
Stamen Type	371	Orientation	397
Stamen Arrangement, Cycle, Position	371	Posture	398
Stamen Attachment and Insertion	372	Surface	399
Stamen Fusion	373	Conspicuity	399
Anther Parts, Type, and Attachment	373	Epidermal Excrescence	399
Anther Dehiscence	373	Vestiture	400
Nectaries	374	Trichome and Bristle Type	401
Gynoecium, Carpel, and Pistil	374	Symmetry	402
Gynoecial Fusion	376	Temporal Phenomena	402
Carpel/Locule Number	376	REVIEW QUESTIONS	403
Ovary Attachment and Position	377	EXERCISES	406
Perianth/Androecial Position	377	REFERENCES FOR FURTHER STUDY	406
Placentation	377		
Style Position/Structural Type	378		
Stigma/Stigmatic Region Types	379		

Plant morphology is a field of study dealing with the external and gross internal structure of plant organs. Morphology intergrades somewhat with plant anatomy, which is the study of tissue and cell structure of plant organs (see Chapter 10). Morphology forms the basis of taxonomic descriptions and generally constitutes the most important data in delimiting and circumscribing taxa.

The terms cited here are largely descended from those used by herbalists and botanists of the past, beginning perhaps with Theophrastus (370 ca. 285 BC), one of the first to write detailed plant descriptions using technical terminology (as in *Historia Plantarum*). The terms have evolved tremendously over the years, especially since the invention of microscopes, having become more detailed and specific. Many of these terms were borrowed from classical Latin (or Greek converted to Latin) and used in new meanings; some were modified from preexisting terms, many invented, and some discarded along the way.

As with all fields of evidence, the terms used in plant morphological descriptions may vary from one source to the next. In using a particular flora, for example, its glossary (if present) should be checked to verify usage of terms. The terms cited below are from a number of sources but are largely derived and classified, with some exceptions, from Radford et al. (1974), which is a precise and logical system of organizing morphological terms (see also Bell 1991). By this classification scheme, the section on **PLANT STRUCTURE** includes characters and character states for specific plant organs and parts. This is followed by a section on **GENERAL TERMINOLOGY**, which lists characters and states that can be used to describe a variety of plant organs.

Note that some terms may need to be explained using other terms (especially those from the general terminology section). Please refer to the Glossary if an unknown term is encountered.

PLANT STRUCTURE

PLANT ORGANS

The basic structural components, or organs, of plants are delimited by and strongly correlated with their specific functions. Among the liverworts, hornworts, and mosses (see Chapter 3), these organs are components of the haploid gametophyte. The gametophyte of these taxa contain **rhizoids**, which are uniseriate, filamentous chains of cells functioning in anchorage and water/mineral absorption. The basic body of the gametophyte can either be a flat mass of cells, termed a **thallus** (found in some liverworts and all hornworts) or a **shoot**, consisting of a generally cylindrical stem bearing leaves (found in some liverworts and all mosses; see Chapter 3).

It should be noted that the shoot systems of liverworts and mosses are gametophytic tissue.

The major organs of vascular plants are sporophytic roots and shoots. **Roots** are present in almost all vascular plants and typically function in anchorage and absorption of water and minerals. Roots consist of an apical meristem that gives rise to a protective root cap, a central endodermis-bounded vascular system, absorptive epidermal root hairs, and endogenously developed lateral roots (Figure 9.1).

The sporophytic **shoots** of vascular plants consist of stem plus leaves (Figure 9.1). Shoots contain an apical meristem of actively dividing cells that, through continued differentiation, result in the elongation of the stem and formation of leaves and buds (see later discussion). The **stem** is a generally cylindrical organ that bears the photosynthetic leaves. Stems typically function in conduction of water and minerals from the roots and in support and elevation of both leaves and reproductive structures, although some stems are highly modified for other functions (see later discussion). The **leaf** is that organ of the shoot that is generally dorsiventrally flattened and that usually functions in photosynthesis and transpiration. Leaves are derived from **leaf primordia** within the shoot apex and are often variously modified. In vascular plants, leaves contain one to many vascular bundles, the **veins**; in some mosses, the gametophytic leaves may contain a veinlike **costa**, consisting of specialized (although not truly vascular) conductive tissue. **Buds** are immature shoot systems, typically located in the axils of leaves. Buds may grow to form lateral vegetative branches or reproductive structures (see later discussion).

Among reproductive plant organs, the **sporangium** is the basic spore-producing part of all land plants. In heterosporous plants (including all of the seed plants) sporangia are of two types: male (microsporangium) or female (megasporengium). The sporangium of liverworts, hornworts, and mosses is known as a **capsule** and typically makes up most of the sporophyte. A **cone**, also called a **strobilus**, is a modified, determinate, reproductive shoot system of many nonflowering vascular plants, consisting of a stem axis bearing either sporophylls (in simple cones) or modified shoot systems (in compound cones). An **ovule** is a megasporengium enveloped by one or more protective integuments. A **seed** is the mature ovule of the seed plants, consisting of an internal embryo surrounded by nutritive tissue (comprising female gametophyte or endosperm) and enveloped by a protective seed coat. The reproductive organ of angiosperms is the **flower**, a modified, determinate shoot bearing sporophylls called stamens and carpels, with or without outer modified leaves (the perianth). An **inflorescence** is an aggregate of one or more flowers, the boundaries of which generally occur with the presence of vegetative leaves. A **fruit** is the mature

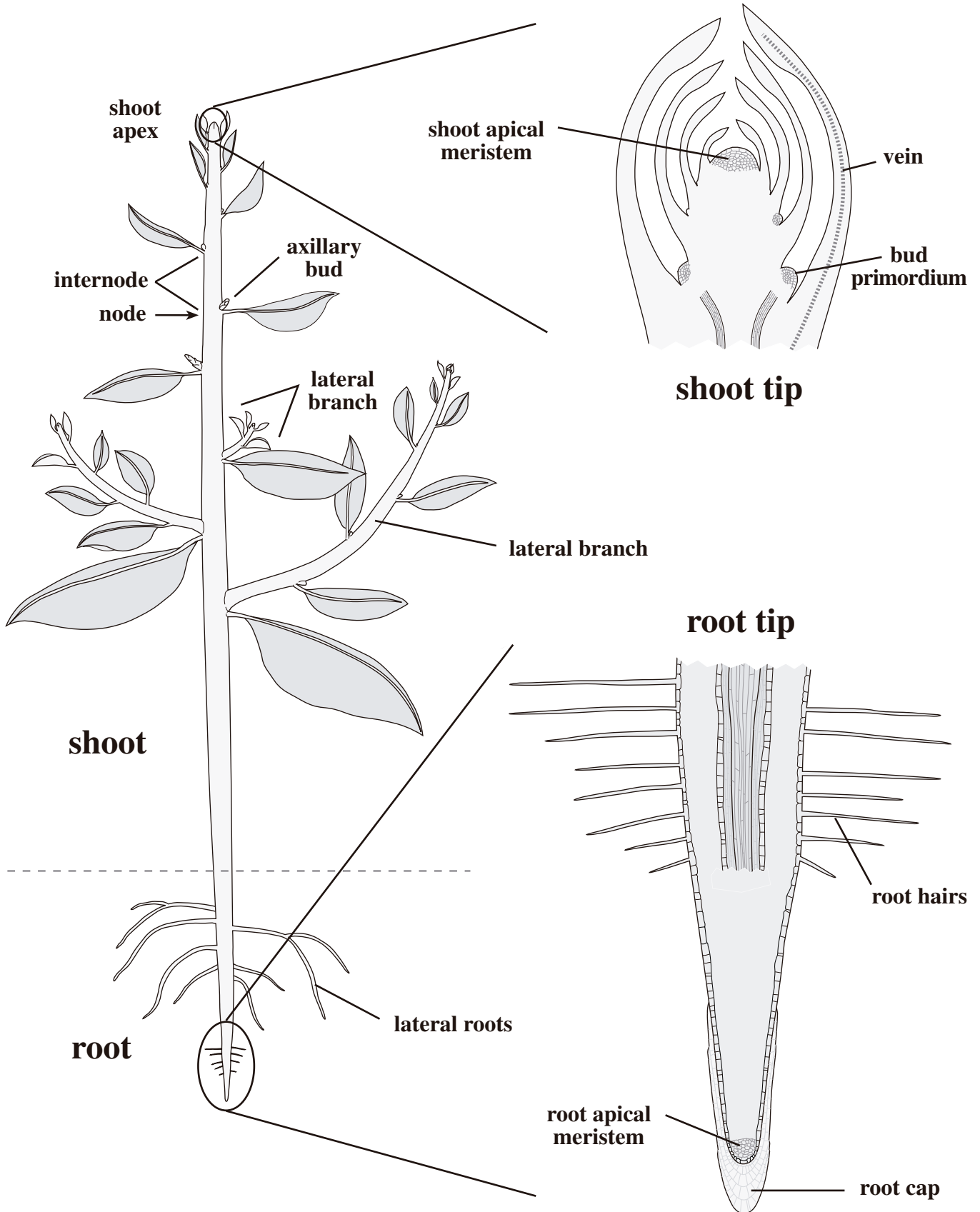


FIGURE 9.1 General plant structure, showing primary root and primary shoot. Note that all parts of both root and shoot are derived from cell divisions of the root or shoot apical meristem.

ovary of flowering plants, consisting of the pericarp (mature ovary wall), seeds, and (if present) accessory parts.

PLANT HABIT

Plant habit refers to the general form of a plant, encompassing a variety of components such as stem duration and branching pattern, development, or texture. Most plants can be clearly designated as an herb, vine, liana, shrub, or tree (with some subcategories; see later discussion); however, some species are difficult to accommodate into these categories. An **herb** is a plant in which any aboveground shoots, whether vegetative or reproductive, die back at the end of an annual growth season. Although the aboveground shoots are annual, the herb itself may be annual, biennial, or perennial, the last by means of long-lived underground rootstocks. Such perennial herbs, having a bulb, corm, rhizome, or tuber as the underground stem, are termed **geophytes**. A **vine** is a plant with elongate, weak stems, that are generally supported by means of scrambling, twining, tendrils, or roots; vines may be annual or perennial, herbaceous or woody. A **liana** (also spelled *liane*) is a vine that is perennial and woody; lianas are major components in the tree canopy layer of some tropical forests. A **shrub** is a perennial, woody plant with several main stems arising at ground level. A **subshrub** is a short shrub that is woody only at the base and that seasonally bears new, nonwoody, annual shoots above. Finally, a **tree** is defined as a generally tall, perennial, woody plant having one main stem (the trunk) arising at ground level. (Some plant ecologists will sometimes distinguish between shrubs and trees based primarily on an arbitrary height.)

PLANT HABITAT

Plant habitat refers to the general environment where the plant is growing. General habitat terms include whether the plant is **terrestrial**, growing on land; **aquatic**, growing in water; or **epiphytic**, growing on another plant. If aquatic, a plant can be **emersed**, occurring under water; **floating**, occurring at the water surface; or **emergent**, having roots or stems anchored to the substrate under water and aerial shoots growing above water. Other aspects of the habitat include the type of substrate that the plant is growing in (e.g., whether on sandy, loam, clay, gravelly, or rocky soil or **saxicolous**, directly on or in the cracks of rocks or boulders), the slope, aspect, elevation, moisture regime, and surrounding vegetation, community, or ecosystem. (See Chapter 17, Plant Collecting and Documentation .)

ROOTS

Roots are plant organs that function in anchorage and in absorption of water and minerals. Roots are found in all of

the vascular land plants except for the Psilophytes *Psilotum* and relatives. (As discussed earlier, nonvascular land plants generally have **rhizoids** that assume a similar function.)

Roots, like shoots, develop by the formation of new cells within the actively growing **apical meristem** of the root tip. The apical meristem is covered on the outside by a **rootcap**, functioning both to protect the root apical meristem and to provide lubrication as the root grows into the soil. The epidermal cells away from the root tip develop hairlike extensions called **root hairs**; these function in greatly increasing the surface area available for water and mineral absorption. Roots of many (if not most) species of plants have an interesting symbiotic interaction with a species of fungus, known as **mycorrhizae**. Although the exact function of mycorrhizae is often unclear, in some species at least the fungus host aids the plant both in increasing overall surface area for absorption and in increasing the efficiency of mineral uptake, particularly phosphorus. Roots have a central **vascular cylinder** of conductive cells, xylem and phloem. This vascular cylinder is surrounded by a special cylinder of cells known as the **endodermis**. Lateral roots develop by cell divisions within the **pericycle**, a cylindrical layer of parenchyma cells located just inside the endodermis itself. (See Chapter 10 for more details of root anatomy.)

The first root to develop in a vascular plant is the **radicle** of the embryo. If the radicle continues to develop after embryo growth, it is known as the **primary root**. Additional roots may arise from internal tissue of either another root, the stem/shoot (often near buds), or (rarely) a leaf. Roots that arise from other roots are called **lateral roots**. Roots that arise from a nonroot organ (stem or leaf) are **adventitious roots**.

ROOT TYPES (Figure 9.2)

Various modifications of roots have evolved. If the primary root becomes dominant, it is called a **taproot**, and the plant is described as having a **taproot system**. If the primary root soon withers and subsequent roots are adventitious, the plant has a **fibrous root system**. Several plant species, particularly those that are biennials, have **storage roots** in which the taproot has become greatly thickened, accumulating reservoirs of high-energy storage compounds (usually starch). Many plants that are epiphytic (grow on another plant), particularly tropical members of the monocot families Araceae and Orchidaceae, have **aerial roots**. These are adventitious roots that generally do not enter the soil and may absorb water and minerals from the air or from runoff from plants. Many plant species with bulbs or corms have **contractile roots**, roots that actually contract vertically, functioning to pull the rootstock further into the soil. Parasitic plants have specialized roots called **haustoria** that penetrate the tissues of a host plant.

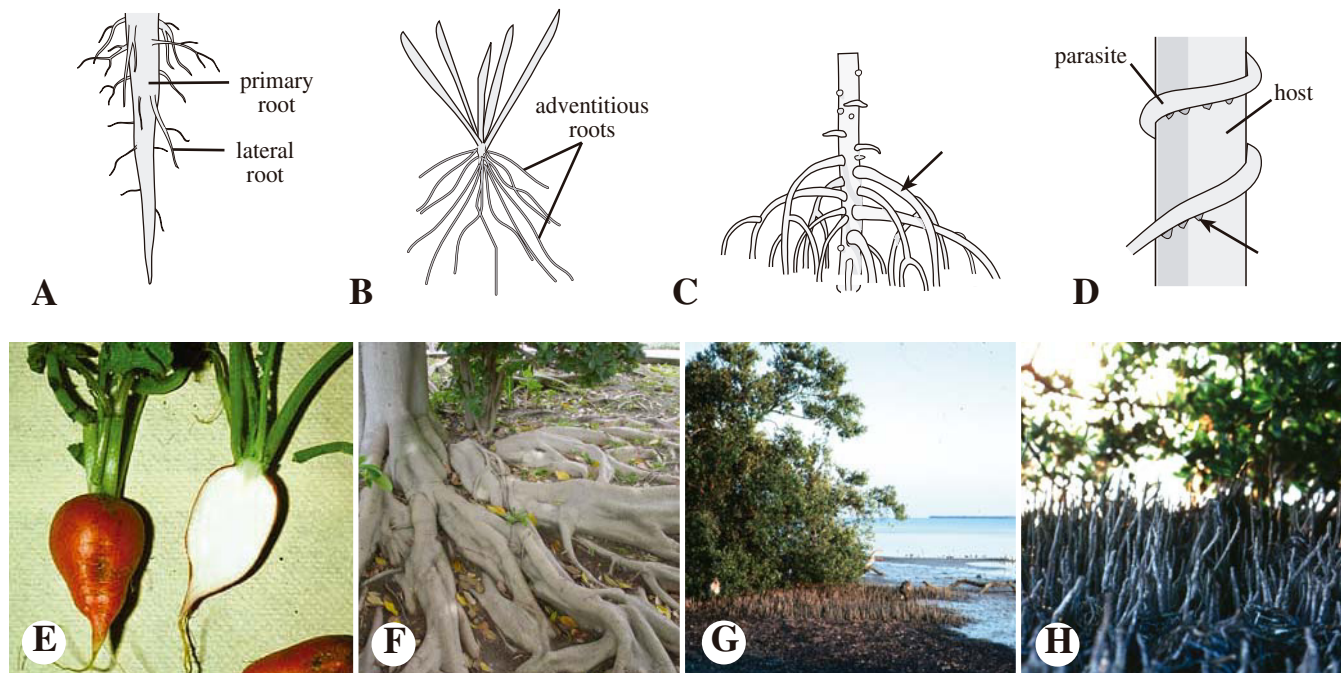


FIGURE 9.2 Root types. **A.** Tap root. **B.** Fibrous root system. **C.** Prop roots. **D.** Haustorial roots. **E.** Storage roots; *Raphanus sativus*, radish. **F.** Buttress roots; *Ficus rubiginosa*, g. **G,H.** Pneumatophores; *Avicennia germinans*, black mangrove.

Some adventitious roots called **prop roots** grow from the base of the stem and function to further support the plant. Some plant species that grow in swamps or marshes have **pneumatophores**, roots that grow upwardly from soil to air that function to obtain additional oxygen. **Buttress roots** are enlarged, horizontally spreading and often vertically thickened roots at the base of trees that aid in mechanical support; they are found in certain tropical or marsh/swamp tree species.

STEMS AND SHOOTS

Stems function both as supportive organs (supporting and usually elevating leaves and reproductive organs) and as conductive organs (conducting both water/minerals and sugars through the vascular tissue between leaves, roots, and reproductive organs). Structurally, stems can be distinguished from roots based on several anatomical features (see Chapter 10). As mentioned earlier, a **shoot** is a stem plus its associated leaves. Sporophytic shoots that are branched and bear leaves are an apomorphy for all extant vascular plants; the leafy shootlike structures of mosses and some liverworts are gametophytic and not directly homologous with shoots of vascular plants.

The first shoot of a seed plant develops from the **epicotyl** of the embryo (see Seeds). The epicotyl elongates after

embryo growth into an axis (the stem) that bears leaves from its tip, which contains the actively dividing cells of the shoot **apical meristem**. Further cell divisions and growth results in the formation of a mass of tissue that develops into the immature leaf, called a **leaf primordium** (Figure 9.1). The point of attachment of a leaf to a stem is called the **node**. The region between two adjacent nodes is the **internode** (Figure 9.1). A bit later in development, the tissue at the upper (adaxial) junction of leaf and stem (called the **axil**) begins to divide and differentiate into a **bud primordium**. As the shoot matures, the leaves fully differentiate into an amazing variety of forms. The bud primordium matures into a **bud**, defined as an immature shoot system, often surrounded by protective scale leaves (see Buds). Buds have an architecture identical to the original shoot. They may develop into a lateral branch or may terminate by developing into a flower or inflorescence. **Vascular strands** run between stem and leaf, providing a vascular connection, composed of xylem and phloem, for water, mineral, and sugar transport. The vascular strands of leaves are termed **veins**.

The mostly parenchymatous tissue external to the vascular (conductive) tissue of a stem is termed the **cortex**. The **pith** is the central, mostly parenchymatous tissue, internal to the stem vasculature (e.g., in siphonosteles and eusteles). In monocots, in which there are numerous, scattered vascular

bundles (an atactostele), the intervening parenchymatous tissue is termed **ground meristem** (see Chapter 10).

The stems of some vascular plants, notably the conifers and nonmonocot flowering plants, contain **wood**, which technically is secondary xylem tissue, derived from a vascular cambium (see Chapter 10). In these woody plants **bark** refers to all the tissues external to the vascular cambium, consisting of secondary phloem (inner bark), leftover cortex, and derivatives of the cork cambium (the last comprising the outer bark, or periderm; see Chapter 10).

STEM TYPES (Figure 9.3)

Various modifications of stems and shoots have evolved, many representing specific adaptations. For example, perennial and some biennial herbs have underground stems, which are generally known as **rootstocks**. Rootstocks function as storage and protective organs, remaining alive underground during harsh conditions of cold or drought. When environmental conditions improve, rootstocks serve as the site of new shoot growth, sending out new adventitious roots and new aerial shoots from the apical meristem or from previously dormant buds. Different types of rootstocks have evolved in various taxonomic groups. These include the following:

1. **Bulb**, in which the shoot consists of a small amount of vertical stem tissue (bearing roots below) and a massive quantity of thick, fleshy storage leaves (e.g., *Allium* spp., onions)
2. **Corm**, in which the shoot consists mostly of generally globose stem tissue surrounded by scanty, scale-like leaves (e.g., some *Iris* spp., irises)
3. **Caudex**, in which the rootstock consists of a relatively undifferentiated but vertically oriented stem
4. **Rhizome**, in which the stem is horizontal and underground, typically with short internodes (compare stolon, below) and bearing scalelike leaves (e.g., *Zingiber officinale*, ginger)
5. **Tuber**, which consists of a thick, underground storage stem, usually not upright, typically bearing outer buds and lacking surrounding storage leaves or protective scales (e.g., *Solanum tuberosum*, potato)

Rootstocks may function as reproductive structures in vegetative (clonal) propagation, either by splitting apart into separate plants or by forming proliferative structures that subsequently separate (and may even be dispersed by animals). For example, buds in the axils of the leaves of bulbs can develop into proliferative **bulbels** (e.g., garlic); some taxa (e.g., certain onions) can even form tiny, propagative bulbels within the aerial shoots or inflorescence of the plant, these

termed **bulbels**. Cormose plants can, similarly from axillary buds, form proliferative corms, termed **cormels**. Tuberous plants typically form numerous tubers at the tips of elongate stems; these tubers can become easily separated, growing into an individual plant. Tubers can even form on aerial shoots (e.g., *Dioscorea*, true yams), ultimately falling off and growing into a new individual. Rhizomes frequently become highly branched; when older parts die or become broken, the separated rhizomes function as separate individuals.

A **stolon** or **runner** is a stem with long internodes that runs on or just below the surface of the ground, typically terminating in a new plantlet, as in *Fragaria* (strawberry). Because stolons can be underground, they are sometimes termed rootstocks and resemble narrow, elongate rhizomes. Stolons function specifically as vegetative propagative structures, however, as the terminal plantlet often becomes separated from the parent plant.

Many modified types of stems that are aerial (above-ground) also have specific functions. For example, a **cladode** is a flattened, photosynthetic stem that may resemble and function as a leaf, found, e.g., in prickly-pear cacti, *Asparagus*, and *Ruscus*. Cladodes take over the primary photosynthetic function of leaves and may function to reduce water loss.

Some aerial stems may function for storage of food reserves or water. So-called succulent stems (the plants often referred to as stem succulents) contain a high percentage of parenchyma tissue that may store great quantities of water, allowing the plant to survive subsequent drought periods. The cacti of the New World and the stem succulent euphorbs of South Africa are classic examples of plants with succulent stems. Some of these, most notably the barrel cacti and the large columnar cacti such as saguaro's or cardon's, have fluted trunks that can expand rapidly following a rain, enabling the plant to store more water. Other aerial, storage stems include:

1. A **caudiciform stem**, which is a low, swollen, perennial storage stem (at or above-ground level), from which arise annual or nonpersistent photosynthetic shoots (e.g., *Calibanus*, some *Dioscorea* spp.)
2. A **pachycaul**, which is a woody, trunklike stem that is swollen basally, the swollen region functioning in storage (e.g., bottle trees, *Brachychiton* spp., and the boojum tree, *Fouquieria columnaris*)

Some stems or shoot types function as protective devices by deterring an herbivore from taking a bite of the plant. A **thorn** is a sharp-pointed stem or shoot. (A thorn is not to be confused with a spine, which is a sharp-pointed leaf or leaf part, or a prickle, which is a sharp-pointed epidermal structure found anywhere on the plant; see later discussion).

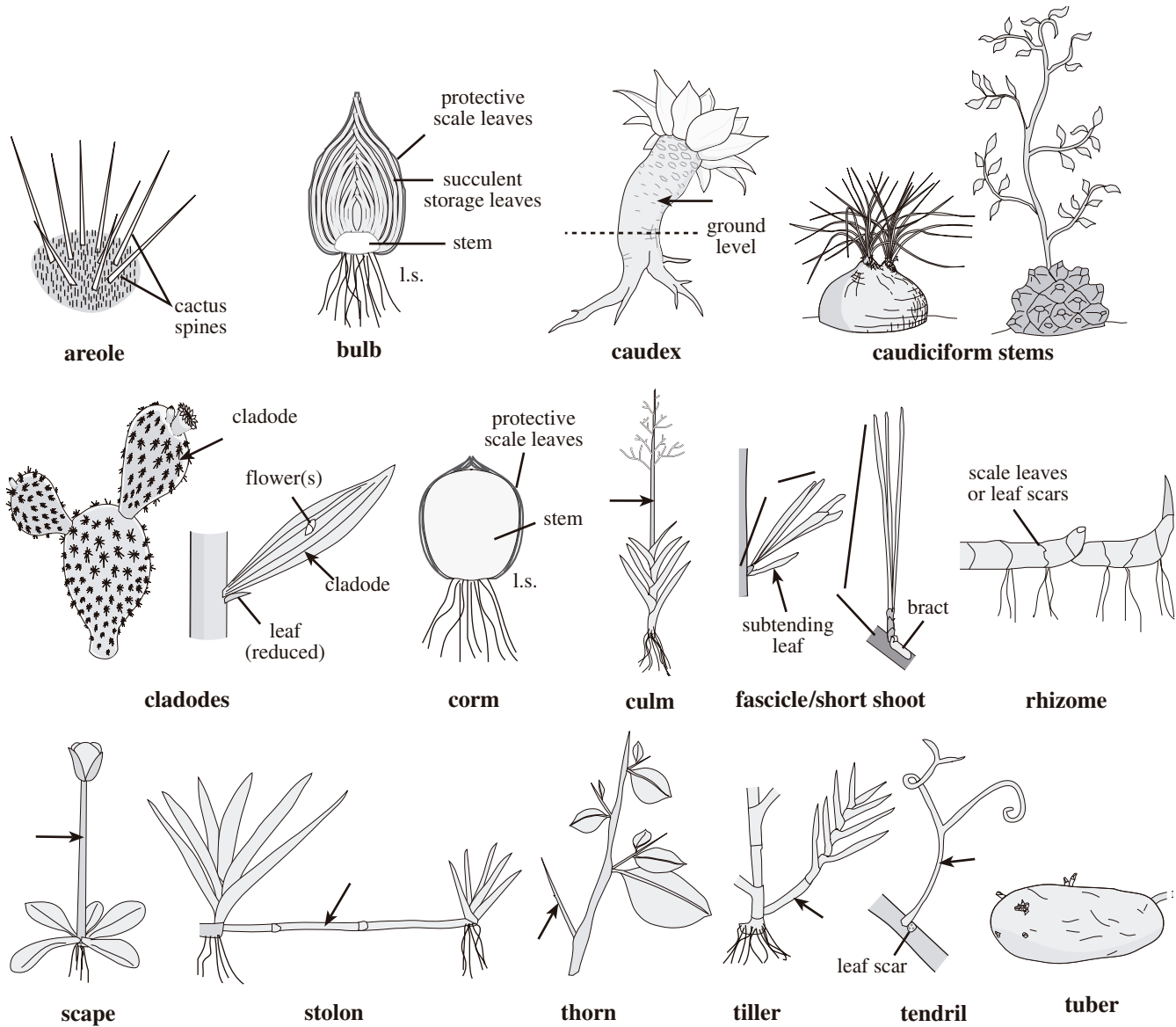


FIGURE 9.3 Stem types. (l.s. = longitudinal section)

A very specialized type of shoot is the **areole**, a modified, reduced, nonelongating shoot apical meristem bearing leaf spines. Areoles are characteristic of the cactus family, Cactaceae.

Some stems are specialized for reproduction. For example, a **scape** is a naked (lacking vegetative leaves) peduncle (inflorescence axis), generally arising from a basal rosette of vegetative leaves and functioning to elevate flowers well above the ground. A **culm** refers to the flowering and fruiting stem(s) of grasses and sedges. A **tiller** is the general term for a proliferative grass shoot, typically growing in masses from axillary buds at the base of the stem.

Stems may have multiple or varied functions. A **lignotuber** or **burl** is largely a protective and regenerative stem following fires. Lignotubers or burls are typically swollen, woody stems, at or slightly below ground level, from which arise persistent, woody, aerial branches (e.g., some *Manzanita* spp.). A **pseudobulb** is a short, erect, aerial storage or propagative stem of certain epiphytic orchids. A **short shoot** or **fascicle** (also called a spur shoot or dwarf shoot) is a modified shoot with very short internodes from which flowers or leaves are borne. Short shoots enable the production of leaves or reproductive organs relatively quickly, with minimal stem (branch) tissue being formed. Short shoots may be found on

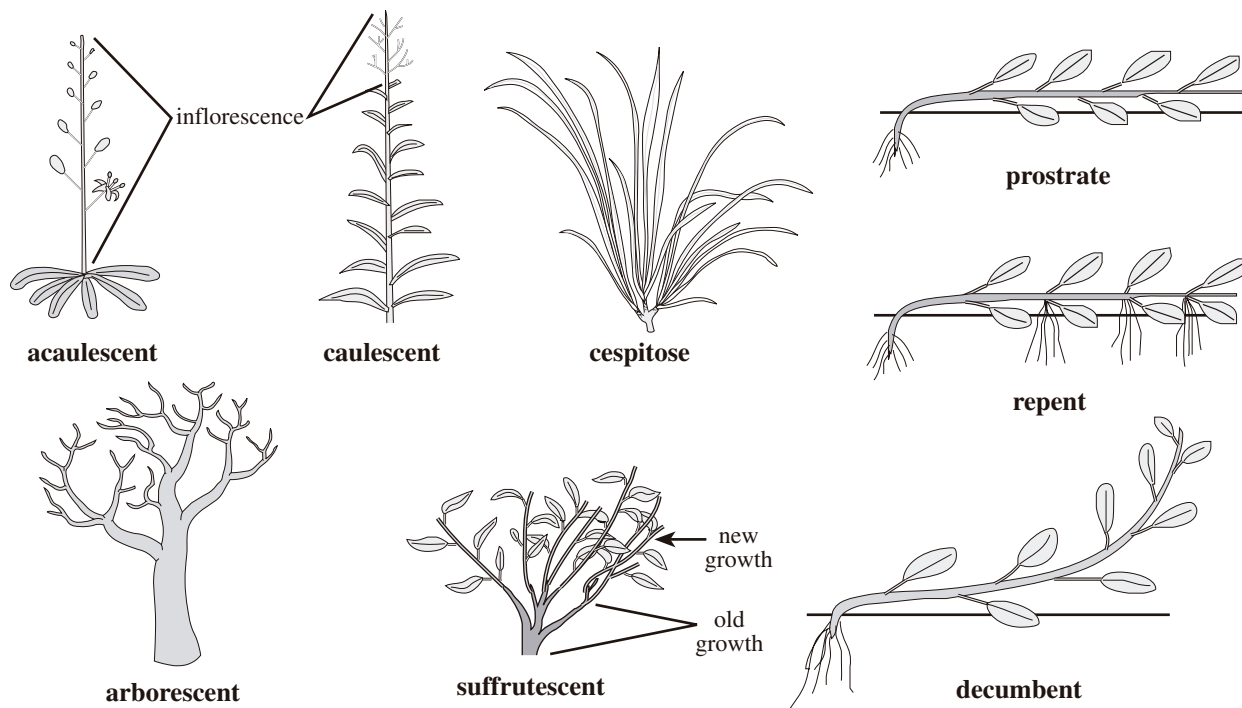


FIGURE 9.4 Stem habit.

so-called drought deciduous plants, which produce a quick flush of leaves from short shoots following a rain. Short shoots arise from the buds of more typical shoots (branches) with longer internodes, the latter termed, in contrast, *long shoots*. Finally, a **tendrils** is a long, slender, coiling branch, adapted for climbing. Tendrils are typically found on weak-stemmed vines and function in support. (Note that most tendrils are leaves or leaf parts; see Leaf Structural Type.)

STEM HABIT (Figure 9.4)

Stem habit is a character describing the relative position of the stem or shoot, but may also be based on stem structure, growth, and orientation. Stem habit features, like stem types, represent adaptations that enhance survival and reproduction. For example, a plant with an above-ground stem is **caulescent**; one that lacks an above-ground stem, other than the inflorescence axis, is termed **acaulescent**. Acaulescent plants bear major photosynthetic leaves only at ground level, often in a basal rosette, with the only shoot becoming aerial, being an inflorescence that eventually dies off. Acaulescent plants are often biennial herbs, in which a storage root develops in the first year and flowering (bolting) occurs in the second, or perennial herbs, in which the persistent stem remains underground and protected during extreme environmental conditions. Plants with caulescent stem habits include shrubs, trees, and herbs with aerial vegetative shoots and leaves.

Some corresponding stem habit terms are **arborescent**, treelike in appearance and size; **frutescent**, having the habit of a shrub, with numerous, woody, aerial trunks; and **suffrutescent**, being basally woody and herbaceous apically, the habit of a subshrub. Vines are also types of caulescent plants. The stem habit of vines can be either **clambering** (also called **scandent**), sprawling across objects without specialized climbing structures, or **climbing**, growing upward by means of tendrils, petioles, or adventitious roots. Some plants are adapted to lying on the ground, at least in part. These include those that are **prostrate**, trailing or lying flat, not rooting at the nodes; **repent**, creeping or lying flat but rooting at the nodes; or **decumbent**, being basally prostrate but apically ascending. Finally, some plants have a **cespitate** stem habit, in which multiple aerial but short-stemmed shoots arise from the base, forming a much-branched cushion. Many grasses are cespitose, these being the so-called bunch grasses.

STEM BRANCHING PATTERN (Figure 9.5)

The below- or above-ground stems or shoots of a plant often exhibit characteristic branching patterns. Branching pattern is determined by the relative activity of apical meristems, both the original shoot apical meristem derived from the seedling epicotyl and apical meristems subsequently derived from lateral buds. One major feature of branching pattern has to do with the duration of apical meristematic growth of a

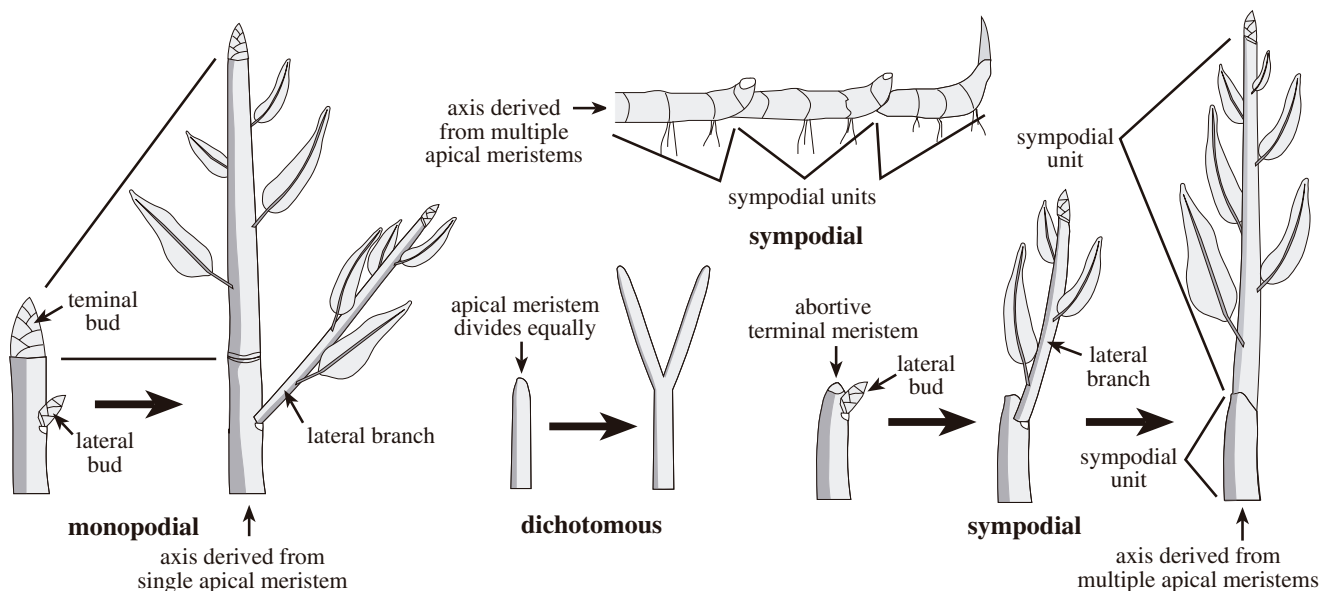


FIGURE 9.5 Stem branching patterns.

shoot. If a given shoot has the potential for unlimited growth, such that the apical meristem is continuously active, the growth is termed **indeterminate**. If instead a shoot terminates growth after a period of time, with either the abortion of the apical meristem or its conversion into a flower, inflorescence or specialized structure (such as a thorn or tendril), the growth is termed **determinate**. (Note that these same terms are used for inflorescence development; see later discussion.) Two other, related terms have to do with flowering. A determinate shoot that completely transforms into a flower or inflorescence is called **hapaxanthic**. An indeterminate shoot that bears lateral flowers but that continues vegetative growth is termed **pleoanthic**.

Many different models of stem branching pattern have been described (e.g., see Hall *et al.* 1978). These models may be of taxonomic value and are interesting from a biomechanical perspective, as they may represent evolutionary adaptations to a given environment or life strategy. Three general terms focus on the developmental origin of a given branch or axis. If a given stem axis is derived from growth of a *single* apical meristem, the pattern is termed **monopodial**. The monopodial axis may grow indefinitely and thus be indeterminate. In contrast, if a given axis (which may appear to be a single, continuous structure) is made up of numerous units that are derived from *separate* apical meristems, the branching pattern is **sympodial**. These sympodial units arise from lateral buds that are proximal to the apical meristem of the original shoot. Many rhizomes have sympodial growth. Finally, a rare type of branching is **dichotomous**, in which a

single apical meristem divides equally into branches, e.g., *Psilotum*.

TWIGS, TRUNKS, AND BUDS (Figure 9.6)

Twigs are the woody, recent-growth branches of trees or shrubs. **Buds** are immature shoot systems that develop from meristematic regions. In deciduous woody plants the leaves fall off at the end of the growing season and the outermost leaves of the buds may develop into protective **bracts** (modified leaves) known as **bud scales**. The bud of a twig that contains the original apical meristem of the shoot (which by later growth may result in further extension of the shoot) is called the **terminal** or **apical** bud. Buds formed in the axils of leaves are called **axillary** or **lateral** buds.

A given bud may be **vegetative**, if it develops into a vegetative shoot bearing leaves; **floral** or **inflorescence**, if it develops into a flower or inflorescence; or **mixed**, if it develops into both flower(s) and leaves. In some species more than one axillary bud forms per node. Two or more axillary buds that are oriented sideways are called **collateral buds**; two or more axillary buds oriented vertically are called **superposed buds**. If the original terminal apical meristem of a shoot aborts (e.g., by ceasing growth or maturing into a flower), then an axillary bud near the shoot apex may continue extension growth; because this axillary bud assumes the function of a terminal bud, it is called a **pseudoterminal bud**.

Several scars may be identified on a woody, deciduous twig. These include the **leaf scar**, **leaf vascular bundle scars**, **stipule scars** (if present), and **bud scale scars**. Bud scale

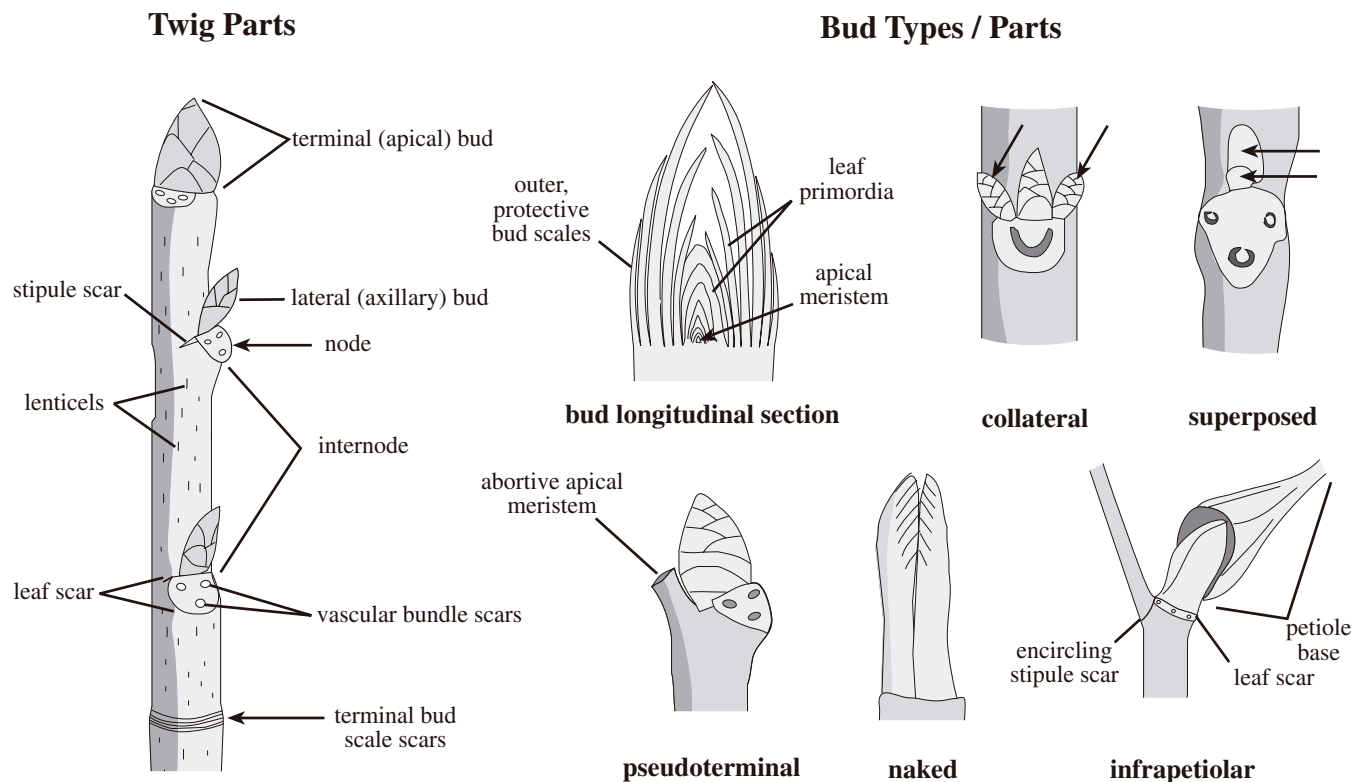


FIGURE 9.6 Twigs parts and bud types. (l.s. = longitudinal section)

scars represent the point of attachment of the bud scales of the original terminal bud after resumption of growth during the new season. Thus, bud scale scars represent the point where the branch ceased elongation the previous growing season; the region between adjacent bud scale scars represents a single year's growth.

Bark technically comprises all the tissue outside the vascular cambium of a plant with true wood (see Chapter 10). The outer bark, or periderm, are the tissues derived from the cork cambium itself. Morphologically, bark may refer to the outermost protective tissues of the stems or roots of a plant with some sort of secondary growth, whether derived from a true cork cambium or not. Bark types are often good identifying characteristics of plant taxa, particularly of deciduous trees during the time that the leaves have fallen. Various bark types include:

1. **Exfoliating**, a bark that cracks or splits into large sheets
2. **Fissured**, a bark split or cracked into vertical or horizontal grooves
3. **Plated**, a bark split or cracked, with flat plates between the fissures
4. **Shreddy**, bark coarsely fibrous

5. **Smooth**, a non-fibrous bark without fissures, fibers, plates, or exfoliating sheets

LEAVES

Leaves are the primary photosynthetic organs of plants, functioning also as the main site of transpiration. Leaves are derived from leaf primordia of the shoot apex and are, at least early in development, generally dorsiventrally flattened (i.e., with dorsal and ventral sides; see Position). A leaf can be gametophytic, in the leafy liverworts and mosses, or sporophytic, in the vascular plants. As mentioned earlier, sporophytic leaves characteristically are associated with **buds**, immature shoot systems, typically located in the axils of leaves. Buds may grow to form lateral vegetative branches or reproductive structures (see later discussion).

LEAF PARTS (Figures 9.7, 9.8, 9.9)

The expanded, flat portion of the leaf, which contains the bulk of the chloroplasts, is termed the **blade** or **lamina**. Many leaves also have a proximal stalk, the **petiole** (e.g., in ferns) the **stipe**. A leaf or leaf part (typically at the base) that partially or fully clasps the stem above the node is a leaf **sheath**, such as in the Poaceae (grasses) and many Apiaceae.

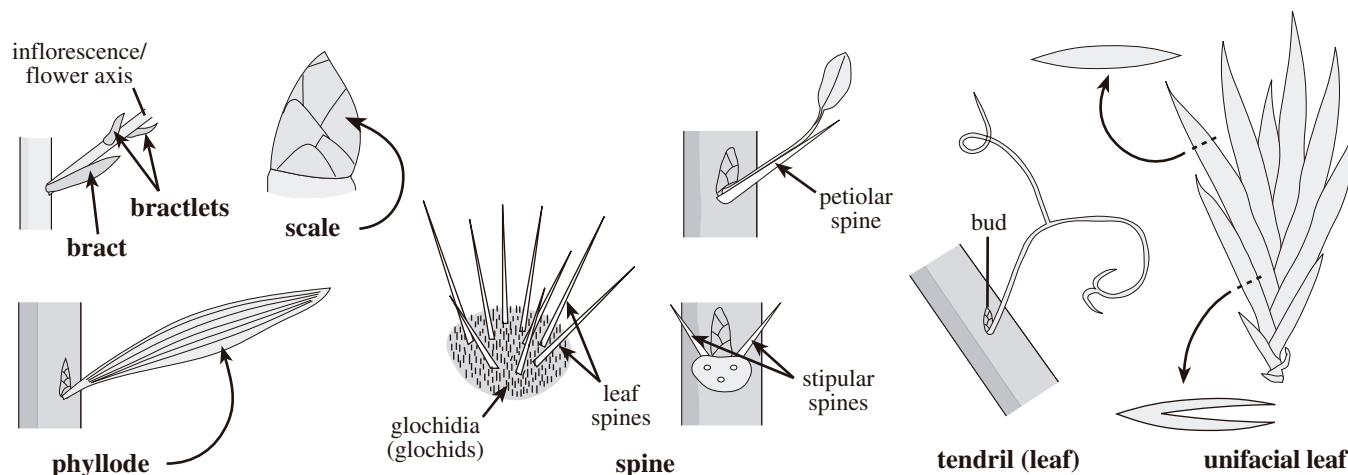


FIGURE 9.7 Leaf structural types.

A **pseudopetiole** is a petiole-like structure arising between a leaf sheath and blade, found in several monocots, such as bananas and bamboos. As mentioned earlier, leaves contain one to many vascular bundles, the **veins** (also sometimes called **nerves**); similar specialized (although not truly vascular) conductive tissue is present in mosses.

Many leaves have **stipules**, a pair of leaflike appendages, which may be modified as spines or glands, at either side of the base of a leaf. If stipules are present, the leaves are **stipulate**; if absent, they are **exstipulate**. A specialized, scarious, sheathlike structure arising above the node in some members of the family Polygonaceae, interpreted as modified stipules, is termed an **ocrea** (see Polygonaceae treatment in Chapter 8). **Stipels** are paired leaflike structures, which may also be modified as spines or glands, at either side of the base of the leaflet of a compound leaf, as in some Fabaceae. If stipels are present, the leaves are **stipellate**; if absent, they are **exstipellate**. Stipules and stipels may, in some cases, function to protect the young, developing leaf primordia. They often are small and fall off (are *caducous*) soon after leaf maturation. In some taxa, stipules or stipels may be highly modified into spines or glands. Extreme examples are some African acacias, in which the swollen stipular spines function as a home for protective populations of ants. In the Rubiaceae the inner surface of the connate stipules (from opposite leaves) bear **colleters**, structures that secrete mucilage (aiding to protect young, developing shoots).

Some leaves are compound (as discussed later), i.e., divided into discrete components called **leaflets**. The stalk of a leaflet is termed the **petiolule**. Some other specialized leaf parts, restricted to certain taxa, are:

1. **Hastula**, an appendage or projection at the junction of petiole and blade, as in some palms

2. **Ligule**, an outgrowth or projection from the inner, top of the sheath, at its junction with the blade, as in the Poaceae
3. **Pulvinus**, the swollen base of a petiole or petiolule, as in some Fabaceae

The pulvinus may, in some taxa, e.g., some Fabaceae (legumes), function in *seismonasty*, which is movement (closing) of the leaflets of a compound leaf as a response to touch or heat (e.g., as in *Mimosa pudica*, sensitive plant); a similar physiological response due to photoperiodism (darkness) is termed *nyctinasty*. These physiological responses may protect the leaf from mechanical damage or help to inhibit water loss through transpiration.

LEAF STRUCTURAL TYPES (Figures 9.7, 9.8)

Leaf structural type (in contrast to leaf type, discussed later) deals with specialized modifications of leaves. One basic leaf structural type in vascular plants is whether the leaves are lycophyllous or euphyllous. **Lycophylls** are small, simple leaves with intercalary growth and a single, central vein that joins to the stem without a leaf gap (below). Lycophylls are found only in lycophytes and are similar to the type of leaf found in the earliest ancestors of vascular plants. **Euphylls** are larger, simple or compound leaves with marginal or apical growth, a leaf gap (region of parenchymatous tissue above the junction of the leaf and stem vasculature), and generally multiple veins. Euphylls are found in ferns (in the broad sense), gymnosperms, and angiosperms (see Chapter 4).

A leaf that is modified in shape and usually smaller than the major photosynthetic leaves is called a **bract**. In angiosperms bracts are typically associated with flowers (flower bracts) or the axes of inflorescences (inflorescence bracts).

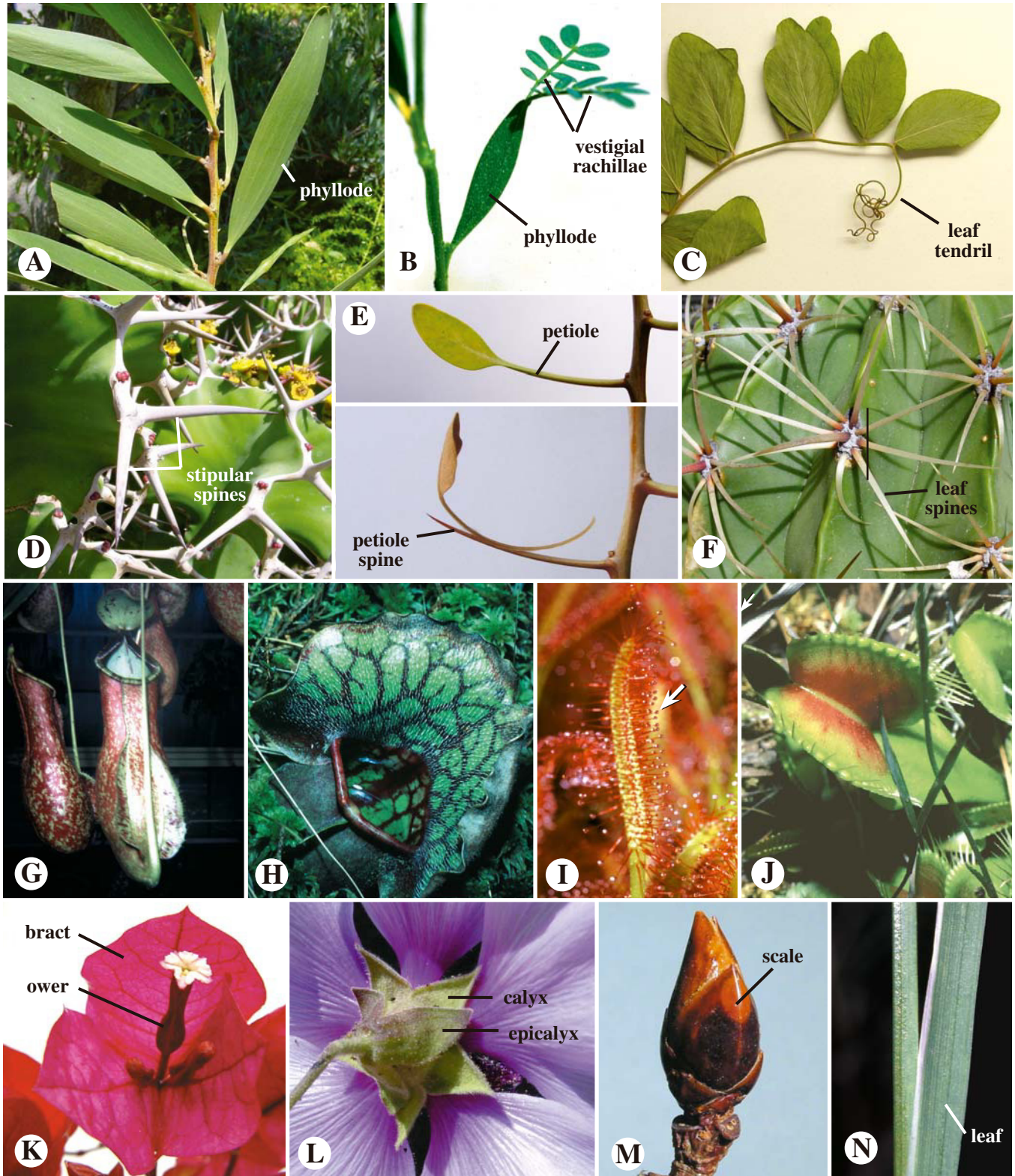


FIGURE 9.8 Leaf structural types. **A,B.** Phyllode, *Acacia longifolia*. **A.** Mature. **B.** Young, with vestigial, caducous rachillae, representative of ancestral condition. **C.** Tendril, *Lathyrus vestitus*. **D,F.** Spines. **D.** Stipular spines, *Euphorbia* sp. **E.** Petiolar spines, *Fouquieria splendens*. Note mature leaf (above), dehiscence of blade and upper tissue of petiole, leaving petiolar spine (below). **F.** Leaf spines, cactus areole. **G–K.** Leaf modifications of carnivorous plants. **G,H.** Pitcher leaves. **G.** *Nepenthes* sp. **H.** *Sarracenia purpurea*. **I.** Tentacular leaves of *Drosera capensis*. Note glandular trichomes (arrow). **J.** Trap leaf, *Dionaea muscipula*. **K.** Showy flower bracts, *Bougainvillea*. **L.** Epicalyx, *Lavatera bicolor*. **M.** Bud scale, *Liquidambar styraciflua*. **N.** Unifacial leaf, *Juncus phaeocephalus*.

A **bractlet** or **bracteole** (also called a **prophyll** or **prophyllum**) is a smaller or secondary bract often borne on the side of a pedicel in flowering plants. The term *bract* is also used for the largely nonphotosynthetic leaves that subtend the ovuliferous scales in conifer cones or that subtend the fascicles or short shoots of members of the pine family (Pinaceae). The term **scale** is used for a small, non-green leaf, either of a bud (**bud scales**), functioning to protect the delicate apical meristem and leaf primordia, or of an underground rootstock, e.g., along the internodes of a rhizome. Scales can also refer to the reduced bracts of sedge spikelets (Cyperaceae). A rudimentary scale leaf found in usually hypogeous (cryptocotylar) seedlings is termed a **cataphyll**.

Some bractlike leaves are found in specific taxonomic groups and are given specialized names. A group of bracts resembling sepals immediately below the true calyx is termed an **epicalyx**, found, e.g., in many members of the Malvaceae. Bracts subtending individual flowers of composites (Asteraceae) are collectively termed **chaff** or **paleae** (singular, **palea**), e.g., as found in the tribe Heliantheae of that family. The specialized bracts of the grass (Poaceae) spikelet are given different terms: **glumes**, the two bracts occurring at the base of a grass spikelet; **lemma**, the outer and lower bract at the base of the grass floret; and **palea**, the inner and upper bract at the base of the grass floret (See **Inflorescence Type**, later, and treatment of Poaceae in Chapter 7.)

A **phyllary** is one of the involucre bracts subtending a head (see later discussion), as in the Asteraceae. A **spathe** is an enlarged, sometimes colored bract subtending and usually enclosing an inflorescence, e.g., that subtending the spadix of the Araceae.

Phyllodes are leaves that consist of a flattened, bladelike petiole. Phyllodes are found in a group of mostly Australian *Acacia* species (the phyllodinous Acacias) and are derived from ancestrally compound leaves by loss of the rachis and leaflets. A **tendrill** is a coiled and twining leaf or leaf part, usually a modified rachis or leaflet. (*Tendrill* can also refer to a modified, coiling stem; see Stem Type).

A **spine** is a sharp-pointed leaf or leaf part. The typical spines of cacti (Cactaceae) are **leaf spines**, as they develop from the entire leaf primordia. A very small, deciduous leaf spine with numerous, retrorse barbs along its length is a **glochidium** (plural, **glochidia** or **glochids**), as found in the areoles of opuntoid cacti. Some taxa have spines that develop from a petiole, midrib, or secondary vein of a leaf, e.g., the **petiolar spines** of *Foquieria* spp. In some palms, e.g., *Phoenix*, the leaflets may be modified into sharp-pointed **leaflet spines**. Many plants, such as the stem-succulent *Euphorbias*, have **stipular spines**; these are typically paired, at the base of a leaf.

A **unifacial leaf** is *isobilateral*, i.e., flattened side-to-side and having a left and right side, except at the base, where they are often sheathing. Some monocots belonging to several different families have unifacial leaves, notably members of the Iridaceae, the Iris family. A **centric leaf** is one that is cylindrical in shape, e.g., *Fenestraria* of the Aizoaceae. Centric leaves are sometimes a subcategory of unifacial leaves.

Three types of leaves are very specialized adaptations of carnivorous plants. **Pitcher** leaves are those that are shaped like a container, which bears an internal fluid and functions in the capture and digestion of small animals. Several taxa have pitcher leaves, including *Darlingtonia*, *Nepenthes*, and *Sarracenia*, the pitcher plants. **Tentacular** leaves are those bearing numerous, sticky, glandular hairs or bristles that function in capturing and digesting small animals; these are characteristic of *Drosera* spp., the sundews. **Trap** leaves are those that mechanically move after being triggered, in the process capturing and digesting small animals; trap leaves are found in *Dionaea muscipula*, the Venus fly trap.

LEAF TYPE (Figure 9.9)

The pattern of division of a leaf into discrete components or segments is termed **leaf type**. A **simple** leaf is one bearing a single, continuous blade. A **compound** leaf is one divided into two or more, discrete **leaflets**. Leaf type should not be confused with leaf division; a simple leaf may be highly divided, but as long as the divisions are not discrete leaflets, it is still technically a simple leaf; see General Terminology. For either compound or divided leaves of ferns, the first (largest) division of a leaf is termed a **pinna**; the ultimate divisions are termed **pinnules**. If the leaves are compound or divided into more than two orders, the terms primary pinna, secondary pinna, etc. can be used, with the ultimate divisions or leaflets always being pinnules.

Simple leaves were the ancestral condition in the vascular plants, as in the lycophylls of the lycopods. Simple leaves are also the norm among the psilophytes, equisetophytes, *Ginkgo*, and conifers (including the Gnetales). Compound leaves are characteristic of many ferns, and all of the cycads. Angiosperms have the greatest diversity of leaves, ranging from simple to highly compound.

Various types of compound leaves have evolved, perhaps as a means of increasing total blade area without sacrificing structural integrity. For example, the blade tissue of a compound leaf generally may have better structural support (e.g., under windy conditions) than that of a comparably sized simple leaf. Compound leaves tend to be more common in mesic to wet environments and simple leaves in dry environments, but there are many exceptions to this and no clear trends.

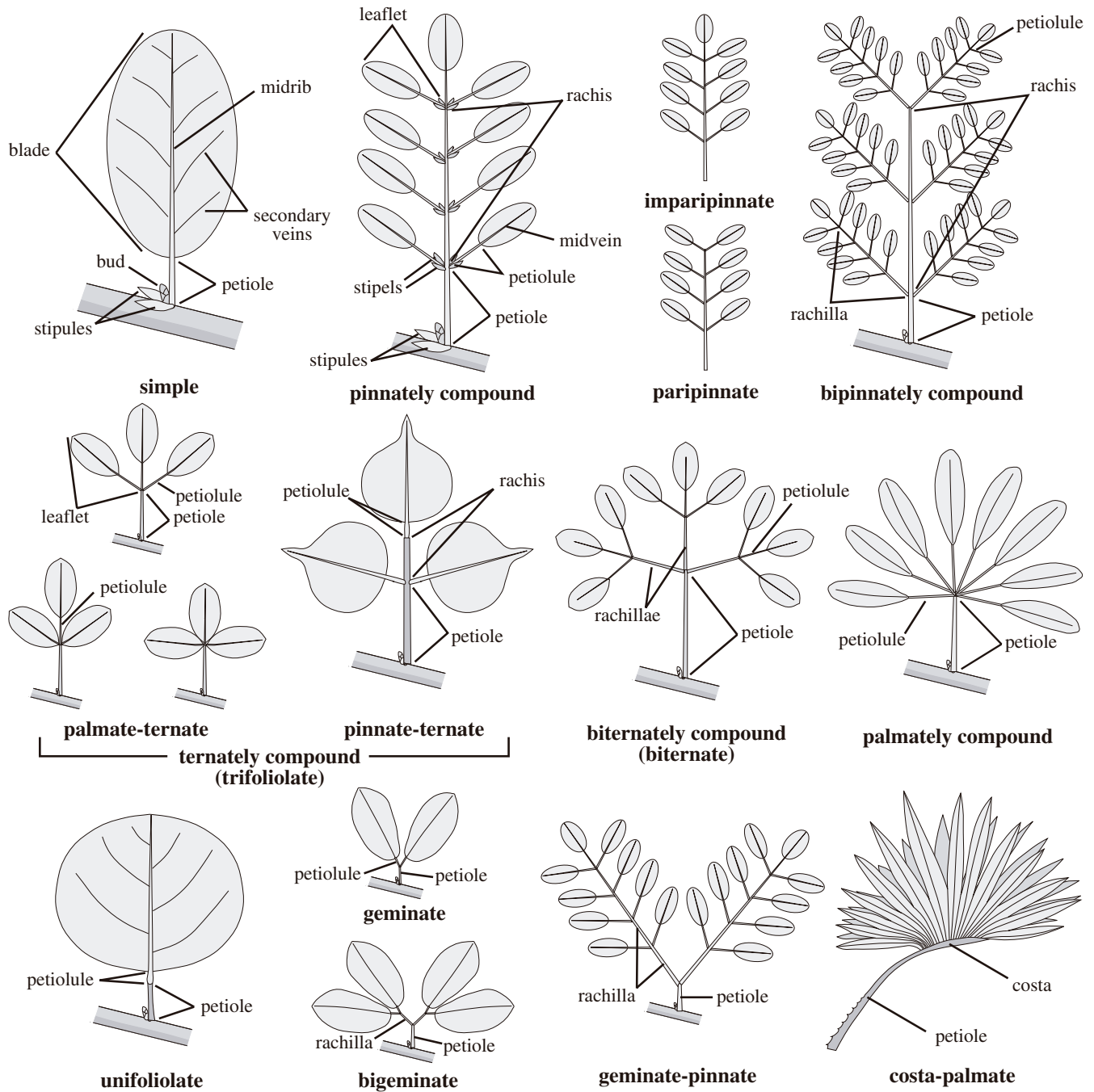


FIGURE 9.9 Leaf types/parts.

Compound leaves are defined based on the number and arrangement of leaflets. A **pinnately compound** or **pinnate** leaf is one with leaflets arranged (either oppositely or alternately) along a central axis, the **rachis**. If a pinnate leaf has a terminal leaflet (and typically an odd number of leaflets), it is **imparipinnate**; if it lacks a terminal leaflet (and has an even number of leaflets), it is **paripinnate**. A **bipinnately compound** or **bipinnate** leaf is with two orders of axes, each

of which is pinnate (equivalent to a compound leaf of compound leaves). The central axis of a bipinnate leaf is still termed the **rachis**; the lateral axes that bear leaflets are termed **rachillae** (singular **rachilla**). Similarly, a compound leaf with three orders of axes, each pinnate, is termed **tripinnately compound** or **tripinnate**; etc.

A compound leaf in which four or more leaflets arise from a common point, typically at the end of the petiole, is termed

palmately compound or **palmate**. A **costapalmate** leaf type is one that is essentially palmately compound to divided, but has an elongate, rachislike extension of the petiole (termed the **costa**), as occurs in some palms.

A compound leaf with only three leaflets is termed **trifoliolate** or **ternately compound**. (A leaf with two orders of axes, each ternately compound, is termed **biternately compound**. Further orders, e.g., **triternately compound**, can also occur.) Most ternately compound leaves are **palmate-ternate**, in which the three leaflets join at a common point (whether petiolulate or sessile). Rarely, ternately compound leaves can be **pinnate-ternate**, in which the terminal leaflet arises from the tip of a rachis. Pinnate-ternate leaves are actually derived (by reduction) from an ancestral pinnately compound leaf; they are found, e.g., in some members of the Fabaceae.

Decompound is a general term for a leaf that is more than once compound, i.e., with two or more orders, being bi-, tri-, etc. pinnately, palmately, or ternately, compound. However, *decompound* is also used for a highly divided leaf; see Division).

A compound leaf consisting of only two leaflets is termed **geminat**e (after Gemini, the twins, in Greek mythology). A compound leaf with two rachillae, each bearing two leaflets, is termed **bigeminat**e. A compound leaf with two rachillae, each of these bearing a pinnate arrangement of leaflets, is termed **geminat-pinnate**. Finally, a very specialized type of leaf is one that appears superficially to be simple, but actually consists of a single leaflet attached to the apex of a petiole, the junction between them clearly defined. This leaf type, known as **unifoliolate**, is interpreted as being derived by reduction of an ancestrally compound leaf.

In some taxa, e.g., many Araceae, the leaves exhibit **heteroblasty** (adjective, **heteroblastic**), in which the juvenile leaves are distinctly different in size or shape from the adult leaves (making species identification difficult).

LEAF ATTACHMENT (Figure 9.10)

The nature of the joining of the leaf to the stem is termed leaf attachment (sometimes treated under Base; see **General**

Morphology). In general, leaves may be **petiolate**, with a petiole, or **sessile**, without a petiole. Leaflets of a compound leaf are, correspondingly, either **petiolulate** or **sessile**. (The term **subsessile** is sometimes used for a leaf/leaflet with a small, rudimentary petiole/petiolute.) Sessile or petiolate leaves can also have a **sheathing** leaf attachment, in which a flattened leaf base (the *sheath*) partially or wholly clasps the stem, typical of the Poaceae (grasses) and many Apiaceae. If a leaf appears to extend down the stem from the point of attachment, as if fused to the stem, the leaf attachment is **decurent** (e.g., as in many Cupressaceae). A decurrent leaf base is not actually caused by later fusion of the leaf to the stem, but by extension growth of actively dividing cells of the leaf primordium at the leaf stem junction. Last, specializations of sessile leaves may occur. If a leaf is sessile and clasps the stem most, but not all, of its circumference, the attachment is termed **amplexicaul**. If the leaf is sessile with the base of the blade completely surrounding the stem, it is termed **perfoliate**. A special case of the latter (involving fusion of leaves) is **connate-perfoliate**, whereby typically two opposite leaves fuse basally, such that the blade bases of the fusion product completely surrounds the stem.

LEAF VENATION (Figures 9.11, 9.12)

The sporophytic leaves of vascular plants contain vascular bundles, known as **veins**, which conduct water, minerals, and sugars between the leaf and the stem. The leaves of some vascular plants have only a single vein, but in most the veins are branched (termed ramified or anastomosing), sometimes in a very intricate pattern. **Venation** refers to this pattern of veins and vein branching. Although venation is usually described for vegetative leaves, it can also be assessed in other leaf homologues, such as bracts, sepals, petals, stamens, or carpels.

The major vein (or veins) of a leaf, with respect to size, is termed the **primary vein**. From the primary vein(s), smaller, lateral veins may branch off, these known as **secondary veins**; from secondary veins, even smaller **tertiary veins** may

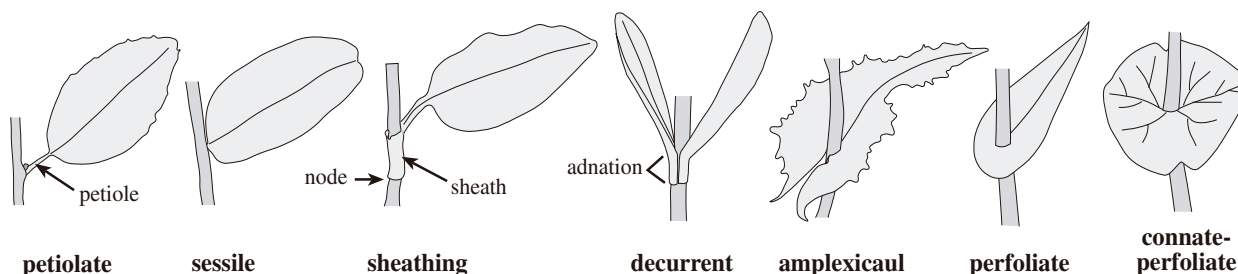


FIGURE 9.10 Leaf attachment.

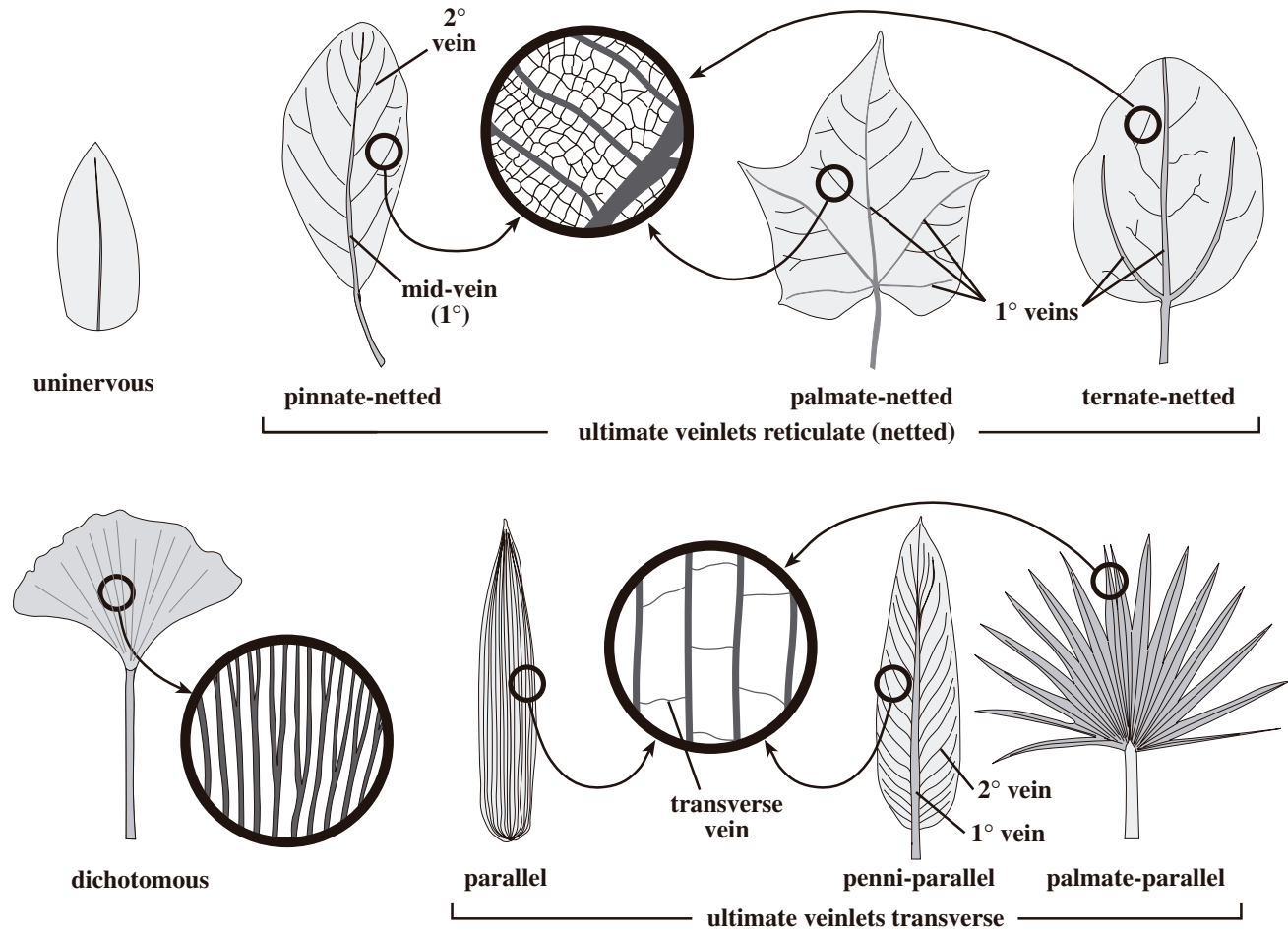


FIGURE 9.11 Leaf venation, generalized terminology.

arise, and so forth. [The distinctions between these vein classes can be difficult to determine in some taxa.] If a simple leaf has a single, primary vein, that vein is termed the **midrib** or **costa** (although *costa* may also be used for the nonvascularized conductive tissue found in the gametophytic leaves of mosses). The central, primary vein of the leaflet of a compound leaf is termed the **midvein**.

Venation patterns can be quite complex, and the terminology formidable (see later discussion). Four, very general venation classes are as follows (Figure 9.11):

1. **Uninervous**, in which there is a central midrib with no lateral veins, e.g., as in the lycophytes, psilophytes, and equisetophytes, as well as many conifers
2. **Dichotomous**, in which veins successively branch distally into a pair of veins of equal size and orientation, e.g., in *Ginkgo biloba*, in which there is no actual midrib
3. **Parallel**, in which the primary and secondary veins are essentially parallel to one another, the ultimate

veinlets being transverse (at right angles), e.g., in most monocots

4. **Netted** or **reticulate**, in which the ultimate veinlets form an interconnecting netlike pattern, e.g., most nonmonocot flowering plants

Reticulate leaves can be **pinnately veined (pinnate-netted)**, with secondary veins arising along length of a single primary vein (the midrib or, in a compound leaf, midvein); **palmately veined (palmate-netted)**, with four or more primary veins arising from a common basal point; or **ternately veined (ternate-netted)**, with three primary veins arising from a common basal point.

Similar to parallel venation in having transverse ultimate veinlets are **penni-parallel** (also called **pinnate-parallel**), with secondary veins arising from a single primary vein region, the former essentially parallel to one another (e.g., the Zingiberales); and **palmate-parallel**, with several primary veins (of leaflets or leaf lobes) arising from

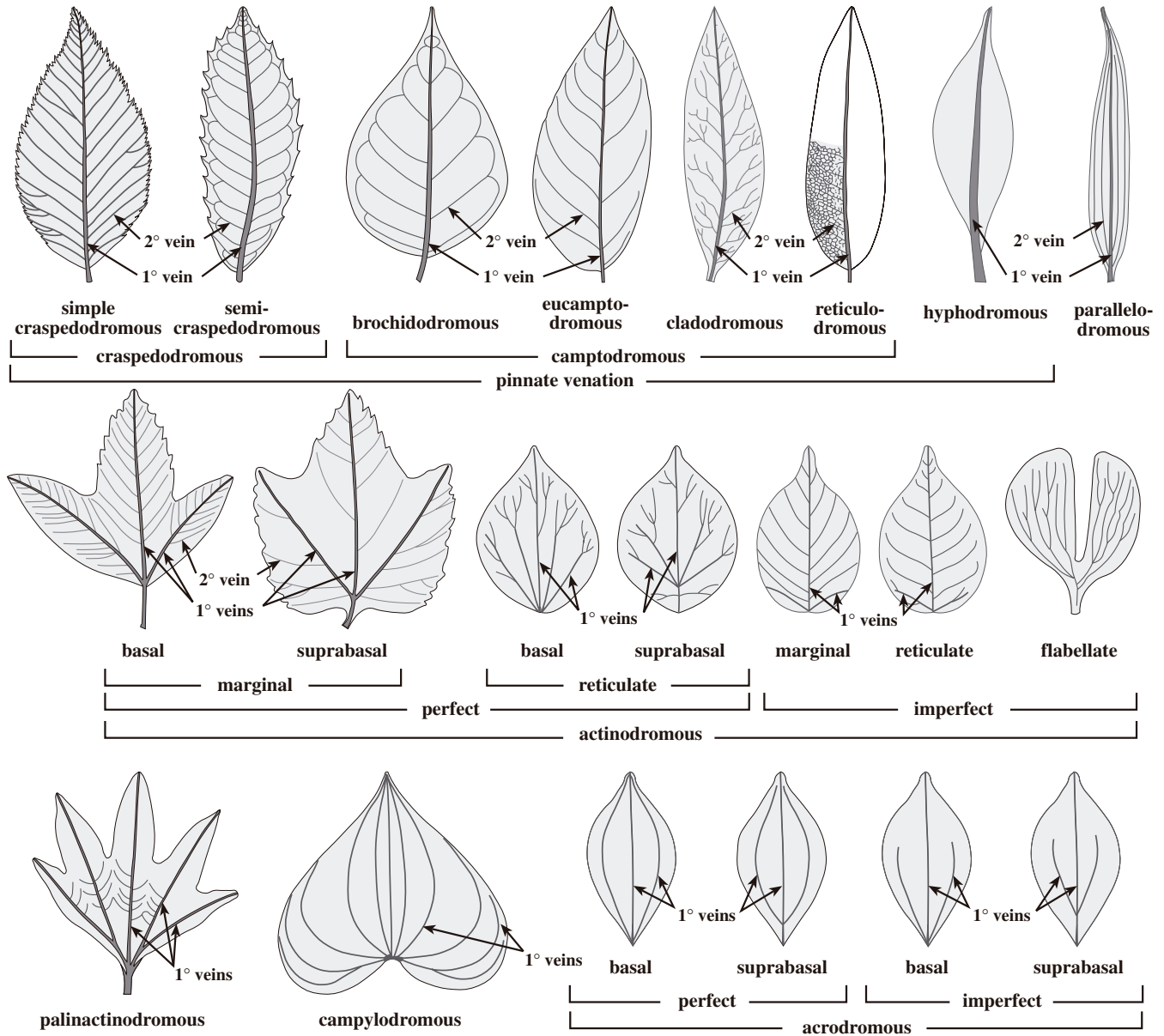


FIGURE 9.12 Leaf venation, specialized terminology, redrawn from Hickey (1973), by permission.

one point, the adjacent secondary veins parallel to these (e.g., fan palms).

A more detailed classification system of venation (and many other leaf features) is that of Hickey (1973) and Hickey and Wolf (1975). This system is based on the pattern of primary, secondary, and tertiary venation. The following is a summary of the terms used in this system, illustrated in Figure 9.12.

Three general venation categories are used for a basically pinnate venation: **craspedodromous**, in which secondary veins terminate at the leaf margin; **camptodromous**, in which secondary veins do not terminate at the margin; and **hyphodromous**, with only the primary midrib vein present

or evident and secondary veins either absent, very reduced, or hidden within the leaf mesophyll.

Subcategories of craspedodromous venation include **simple craspedodromous**, in which all secondary veins terminate at the margin; **semicraspedodromous**, in which the secondary veins branch near the margin, one terminating at the margin, the other looping upward to join the next secondary vein; and **mixed craspedodromous** (not illustrated), with some secondary veins terminating at the margin, but with many terminating away from the margin.

Subtypes of camptodromous venation include **brochidodromous**, in which secondary veins form prominent upward

loops near the margin, joining other, more distal, secondary veins; **eucamptodromous**, in which secondary veins curve upward near the margin but do not directly join adjacent secondaries; **cladodromous**, in which secondary veins branch toward the margin; and **reticulodromous**, in which secondary veins branch repeatedly, forming a very dense, netlike structure.

Parallelodromous venation is equivalent to parallel (defined earlier), in which two or more primary or secondary veins run parallel to one another, converging at the apex.

Venation is **actinodromous** if three or more primary veins diverge from one point (equivalent to ternate or palmate venation). **Palinactinodromous** is similar, but the primary veins have additional branching above the main point of divergence of the primaries.

For actinodromous and palinactinodromous types, the venation is **marginal** if the main, primary veins reach the blade margin, and **reticulate** (not to be confused with reticulate in the more general venation terminology) if they do not. **Flabellate** venation is that in which several equal, fine veins branch toward the apex of the leaf.

Campylodromous venation is that in which several primary veins run in prominent, recurved arches at the base, curving upward to converge at the leaf apex.

Finally, venation is **acrodromous**, if two or more primary veins (or strongly developed secondary veins) run in convergent arches toward the leaf apex (but are not recurved at the base, as in campylodromous).

For actinodromous, palinactinodromous, and acrodromous types, the venation is **basal** if the primaries are joined at the blade base, and **suprabasal** if the primaries diverge above the

blade base. The venation is **perfect** if branching of the lateral primary veins and their branches cover at least two thirds of the leaf blade area (or reach at least two thirds of the distance toward the leaf apex), and **imperfect** if these veins cover less than two thirds of the leaf blade area (or reach less than two thirds of the way toward the leaf apex).

These complex venation types, along with many other details of the leaf, can be specific to certain taxonomic groups of plants. Although they are not widely used in standard morphological descriptions, their recognition can be important in identification (e.g., of many tropical and fossil plants) and classification (see Hickey and Wolf 1975).

FLOWERS

A major diagnostic feature of angiosperms is the flower. As discussed in Chapter 6, a **flower** is a modified reproductive shoot, basically a stem with an apical meristem that gives rise to leaf primordia. Unlike a typical vegetative shoot, however, the flower shoot is determinate, such that the apical meristem stops growing after the floral parts have formed. At least some of the leaf primordia of a flower are modified as reproductive sporophylls (leaves bearing sporangia). Flowers are unique, differing, e.g., from the cones of gymnosperms, in that the sporophylls develop either as stamens or carpels (see Chapter 6, and later discussion).

FLOWER PARTS (Figure 9.13)

The basic parts of a flower, from the base to the apex, are as follows. The **pedicel** is the flower stalk. (If a pedicel is absent, the flower attachment is *sessile*.) Flowers may be subtended by a **bract**, a modified, generally reduced leaf; a smaller or

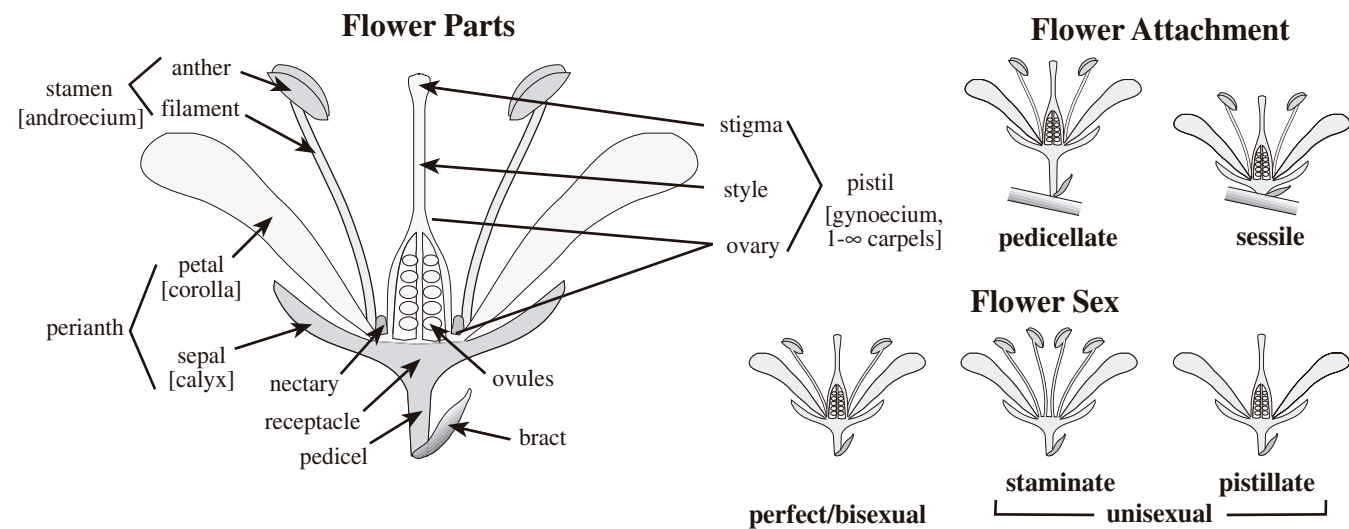


FIGURE 9.13 Flower parts, sex, and attachment.

secondary bract, often borne on the side of a pedicel, is termed a **bracteole** or **bractlet** (also called a **prophyll** or **prophyllum**). Bracteoles, where present, are typically paired. [In some taxa, a series of bracts, known as the **epicalyx**, immediately subtends the calyx (see later discussion), as in *Hibiscus* and other members of the Malvaceae.] The **receptacle** is the tissue or region of a flower to which the other floral parts are attached. The receptacle is typically a small, obscure region (derived from the original apical meristem). In some taxa the receptacle can grow significantly and assume an additional function. From the receptacle arises the basic floral parts. The **perianth** (also termed the **perigonium**) is the outermost, nonreproductive group of modified leaves of a flower. If the perianth is relatively undifferentiated, or if its components intergrade in form, the individual leaflike parts are termed **tepals**. In most flowers the perianth is differentiated into two groups. The **calyx** is the outermost series or whorl of modified leaves. Individual units of the calyx are **sepals**, which are typically green, leaflike, and function to protect the young flower. The **corolla** is the innermost series or whorl of modified leaves in the perianth. Individual units of the corolla are **petals**, which are typically colored (nongreen) and function as an attractant for pollination. Some flowers have a **hypanthium** (floral tube), a cup-like or tubular structure, around or atop the ovary, bearing along its margin the sepals, petals, and stamens.

Many flowers have a **nectary**, a specialized structure that secretes nectar. Nectaries may develop on the perianth parts, within the receptacle, on or within the androecium or gynoecium (below), or as a separate structure altogether. Some flowers have a **disk**, a discoid or doughnut-shaped structure arising from the receptacle. Disks can form at the outside and surrounding the stamens (termed an *extrastaminal disk*), at the base of the stamens (*staminal disk*), or at the inside of the stamens and/or base of the ovary (*intrastaminal disk*). Disks may be nectar-bearing, called a *nectariferous disk*.

The **androecium** refers to all of the male organs of a flower, collectively all the stamens. A **stamen** is a microsporophyll, which characteristically bears two thecae (each theca comprising a pair of microsporangia; see Chapter 6). Stamens can be leaflike (laminar), but typically develop as a stalk-like **filament**, bearing the pollen-bearing **anther**, the latter generally equivalent to two fused thecae.

The **gynoecium** refers to all of the female organs of a flower, collectively all the carpels. A **carpel** is the unit of the gynoecium, consisting of a modified megasporophyll that encloses one or more ovules. Carpels typically develop in a conduplicate manner. A **pistil** is that part of the gynoecium composed of an **ovary**, one or more **styles** (which may be absent), and one or more **stigmas** (see later discussion).

In some taxa, e.g. Aristolochiaceae and Orchidaceae, the androecium and gynoecium are fused into a common structure, known variously as a **column**, **gynandrium**, **gynostegium**, or **gynostemium**. A stalk that bears the androecium and gynoecium is an **androgynophore**, e.g., Passifloraceae.

FLOWER SEX AND PLANT SEX (Figure 9.13)

Flower sex refers to the presence or absence of male and female parts within a flower. Most flowers are **perfect** or **bisexual**, having both stamens and carpels. Bisexual flower sex is likely the ancestral condition in angiosperms.

Many angiosperm taxa, however, have **imperfect** or **unisexual** flower sex. In this case, flowers are either **pistillate/female**, in which only carpels develop, or **staminate/male**, in which only stamens develop.

Plant sex refers to the presence and distribution of perfect or imperfect flowers on individuals of a species. A **hermaphroditic** plant is one with only bisexual flowers. A **monoecious** (*mono*, one + *oikos*, house) plant is one with only unisexual flowers, both staminate and pistillate on the same individual plant; e.g., *Quercus* spp., oaks. A **dioecious** (*di*, two + *oikos*, house) plant is one with unisexual flowers, but with staminate and pistillate on separate individual plants (i.e., having separate male and female individuals; e.g., *Salix* spp., willows). Plant sex can vary within individuals of a species, and there may also be a combination of perfect and imperfect flowers in different individuals. **Polygamous** is a general term for a plant with both bisexual and unisexual flowers. **Andromonoecious** refers to a plant with both staminate and perfect flowers on the same individual, and **gynomonoecious** is a plant with both pistillate and perfect flowers on the same individual. **Trimonoecious** refers to a plant with pistillate, staminate, and perfect flowers on the same individual. **Androdioecious** refers to a plant with male flowers on some individuals and perfect flowers on other individuals. **Gynodioecious** refers to a plant with female flowers on some individuals and perfect flowers on other individuals. **Trioecious** refers to a plant with pistillate, staminate, and perfect flowers on different individuals. All of these types of nonhermaphroditic plant sex may function as a mechanism of promoting increased outcrossing between individuals of a species. (However, many hermaphroditic plants can outcross by other means; see Chapter 13).

FLOWER ATTACHMENT (Figure 9.13)

Flower attachment is **pedicellate**, having a pedicel; **sessile**, lacking a pedicel; or **subsessile**, having a short, rudimentary pedicel. The terms **bracteate**, with bracts, and **ebracteate**, lacking bracts, may also be used with respect to flower attachment. The adaptive significance of pedicels is likely

correlated with the spatial positioning of flowers relative to pollination or eventual fruit or seed dispersal.

FLOWER CYCLY

Flower cyclcy refers to the number of cycles (series or whorls) or floral parts. The two basic terms used are **complete**, for a flower having all four major series of parts (sepals, petals, stamens, and carpels) and **incomplete**, for a flowering lacking one or more of the four major whorls of parts (e.g., any unisexual flower, or a bisexual flower lacking a corolla).

FLOWER SYMMETRY (Figures 9.14, 9.15)

Flower symmetry is an assessment of the presence and number of mirror-image planes of symmetry. **Actinomorphic** or **radial** symmetry (also called **regular**) is that in which there are three or more planes of symmetry, such that there is a repeating structural morphology when rotated less than 360° about an axis. **Biradial symmetry** means having two (and only two) planes of symmetry. (The difference between biradial and radial symmetry is sometimes not recognized, both being termed *radial symmetry* or *actinomorphy*; however, the distinction can be useful and is recognized here.) **Zygomorphic** or **bilateral** symmetry (also called **irregular**) is that in which there is only one plane of symmetry. An **asymmetric** flower lacks any plane of symmetry, usually the result of twisting of parts. Flower symmetry can sometimes be subtle and can even vary within a flower; if so, it should be separately described for calyx, corolla, androecium, and gynoecium to avoid confusion.

Flower symmetry can be an important adaptation relative to pollination systems. Actinomorphic flower symmetry is likely the ancestral condition in angiosperms and is found in a large number of groups. Zygomorphy has evolved repeatedly in many groups, typically as a means of more efficiently transferring pollen to an animal (usually insect) pollinator. Zygomorphy is typically correlated with a more horizontal floral orientation, and there are many different ways that zygomorphy can come about developmentally and morphologically.

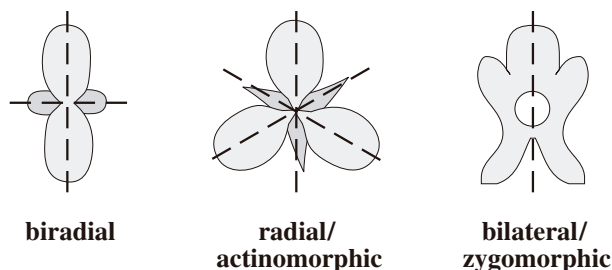


FIGURE 9.14 Flower symmetry types.

FLOWER MATURATION

Flower maturation refers to the time of development of flowers or flower parts (see also General Terminology). **Anthesis** is the general time of flowering, the opening of flowers with parts available for pollination. The relative timing of development of male versus female flowers or floral parts can be an important feature in reproductive biology. **Protandrous** refers to stamens developing, or pollen release occurring, prior to the maturation of carpels or stigmas being receptive. **Protogynous** is the reverse, with carpels or stigmas developing before stamens mature or pollen is released. Both protandry and protogyny may function to promote outcrossing (and thus inhibit selfing) within individuals of a species.

Two flower maturation terms dealing with the relative direction of development of parts can be important in describing taxonomic groups. **Centrifugal** refers to developing from the center toward the outside or periphery, whereas **centripetal** is development from the outside or periphery toward the center region. Both *centrifugal* and *centripetal* can be applied to parts of the perianth, calyx, corolla, androecium, or gynoecium; the terms are often used to describe the direction of development of stamens in a multiwhorled androecium.

PERIANTH

The **perianth** (or *perigonium*) is the outermost, nonreproductive group of modified leaves of a flower. (The term *perianth* has also been used for components of the reproductive structures of various Gnetales, but these are not homologous.) A perianth is absent in some flowering plants, typically those taxa that have very small, reduced flowers. The perianth, where present, functions both to protect the young flowering parts and to aid in pollination.

The units of the perianth arise like leaves as primordia from the apical meristem of the flower. Typically, they may retain leaflike characters. Sepals, in fact, are usually green with stomata and veins; even petals will have veins and may have vestigial stomata. However, the perianth can undergo significant developmental changes and be highly modified (and unleaflike) at maturity.

PERIANTH PARTS (Figure 9.19)

Various specialized terms are used for parts of the perianth. These include the following: **anterior** or **ventral**, referring to the lower, abaxial lobe(s) or side, toward a subtending bract; **beard**, a tuft, line, or zone of trichomes on a perianth or perianth part (see Vestiture); **claw**, an attenuate base of a sepal or petal; **corona**, a crownlike outgrowth between stamens and corolla, which may be petaline or staminal in

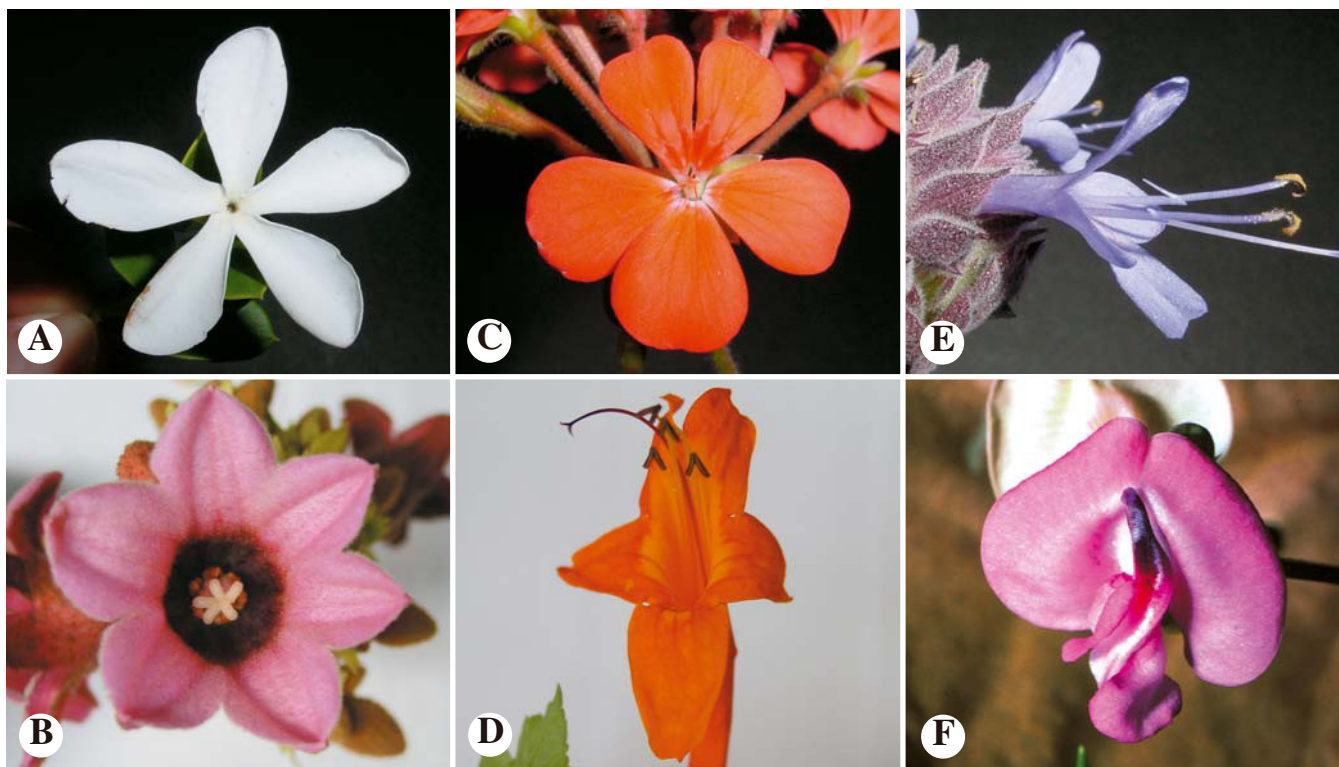


FIGURE 9.15 Flower symmetry examples. **A,B.** Actinomorphic/radial symmetry. **A.** Five planes of symmetry. **B.** Six planes of symmetry. **C.** Slight zygomorphy (bilateral symmetry), vertical plane of symmetry. **D,E.** Strong zygomorphy of all floral parts. **F.** Asymmetry, caused by twisting of floral parts.

origin; **hypanthium** or **floral cup**, a generally tubular or cup-shaped structure at the top rim of which are attached the calyx, corolla, and androecium; **labellum**, a modified, typically expanded, median petal, tepal, or perianth lobe, such as in the Orchidaceae; **limb**, the expanded portion of corolla or calyx above the tube, throat, or claw; **lip**, either of two variously shaped parts into which a calyx or corolla is divided, usually into upper (posterior) and/or lower (anterior) lips, such as most Lamiaceae, Orchidaceae (Note: each lip may be composed of one or more lobes); **lobe**, a segment of a synsepalous calyx or sympetalous corolla; **petal**, a corolla member or segment; a unit of the corolla; **posterior** or **dorsal**, referring to the upper, adaxial lobe(s) or side, nearest to the axis, away from the subtending bract; **sepal**, a calyx member or segment, a unit of the calyx; **spur**, a tubular, rounded or pointed projection from the calyx or corolla, functioning to contain nectar; **tepal**, a perianth member or segment not differentiated into distinct sepals or petals; **throat**, an open, expanded region of a perianth, usually of a sympetalous corolla; **tube**, a cylindrically shaped perianth or region of the perianth, usually of a sympetalous corolla.

PERIANTH ARRANGEMENT/CYCLY/MEROSITY

(Figures 9.16, 9.17)

A fundamental aspect of perianth structure is **perianth arrangement**, the position of perianth parts relative to one another. In some taxa, such as some magnolias and water lilies, the perianth parts have a **spiral** arrangement, i.e., spirally arranged with only one perianth part per node, not in distinct whorls. Typically, flowers with a spiral perianth arrangement have parts that are either undifferentiated (similar to one another) or that grade from an outer, sepal-like form to an inner petal-like form. In either case, the term **tepal** is used to describe undifferentiated or intergrading perianth parts. In most flowering plants the perianth parts have a **whorled** arrangement, in which the parts appear to arise from the same nodal region. (Note that, developmentally, the perianth parts may actually initiate as primordia at slightly different times and positions; however, at maturity, this is usually undetectable.)

Cycly refers to the number of whorls (cycles, series) of parts. (See General Terminology.) Thus, **perianth cycl**y is the number of whorls of perianth parts. The most common

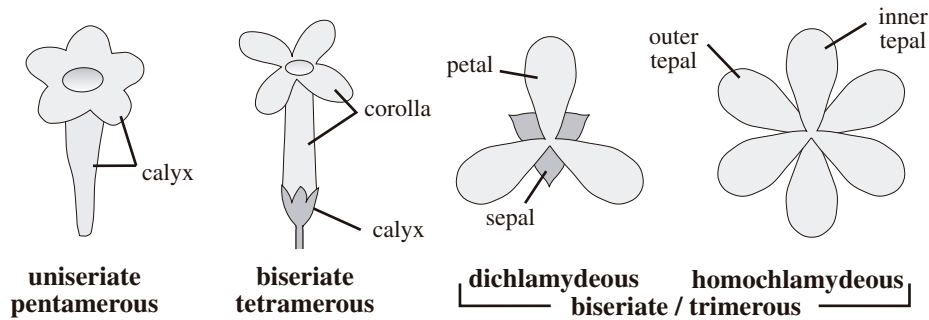
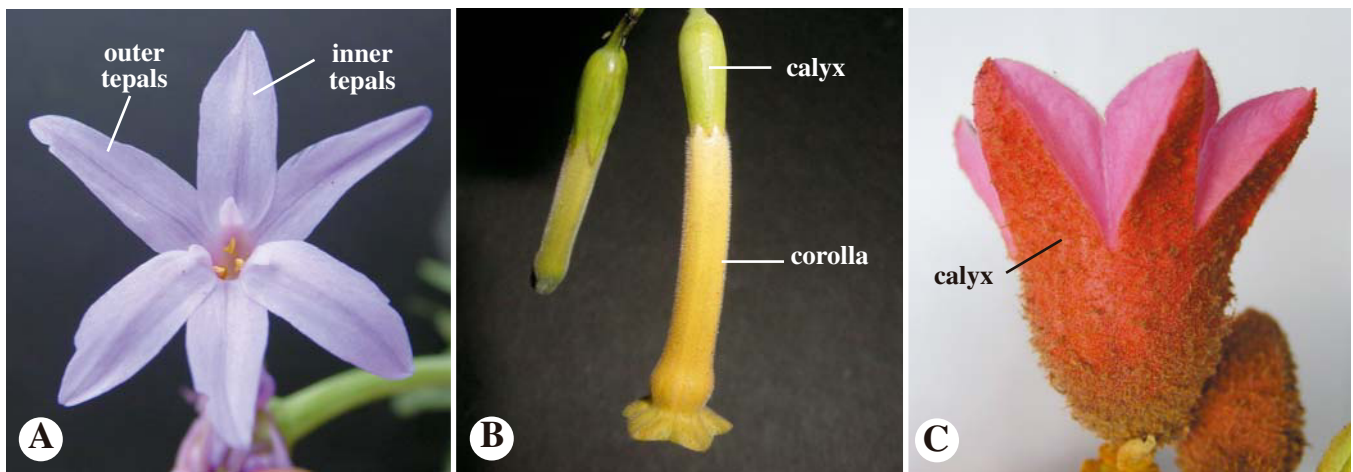


FIGURE 9.16 Perianth cyclcy and merosity.

type of perianth cyclcy by far is **biseriate** (also called *dicyclic*), in which there are two discrete whorls, an outer (= lower) and inner (= upper). A less common condition in flowering plants is a **uniseriate** perianth cyclcy, with perianth parts in a single whorl. Uniseriate perianths may arise by loss or reduction of one of the whorls of an ancestrally biseriata perianth. If it is known that the calyx was evolutionarily lost, what remains should be called a *corolla*; if the corolla was lost, what remains should be termed a *calyx*. If this directionality is not known, a uniseriate perianth is usually termed a calyx by tradition (although it may simply be called a *perianth*). Perianths may also rarely be **triseriate** (or *tricyclic*) = three-whorled, **tetraseriata** (or *tetracyclic*) = four-whorled, etc. The term **multiseriate** may be used to mean composed of three or more whorls. Other cyclcy terms evaluate the similarity of the whorls of parts to one another. **Dichlamydeous** describes a perianth composed of a distinct outer calyx and inner corolla; in most cases, a dichlamydeous perianth is also biseriata, but it may be multiseriate (i.e., the calyx or corolla

containing more than one whorl). **Homochlamydeous** refers to a perianth composed of similar parts, each part a tepal. Most monocots have a homochlamydeous perianth, whereas most eudicots have a dichlamydeous one. In some cases, the distinction between *dichlamydeous* and *homochlamydeous* can be difficult, as it may be difficult to assess whether outer and inner series are similar or different.

Merosity refers to the number of parts per whorl or cycle. (See General Terminology.) Thus, **perianth merosity** is the number of parts per whorl of the perianth. General terms for perianth merosity are **isomerous**, having the same number of members in different whorls (e.g., five sepals and five petals) and **anisomerous**, having a different number of members in different whorls (e.g., two sepals and five petals). Merosity may be described separately for each whorl of the perianth, e.g., **calyx merosity** and **corolla merosity**. It is assessed for numbers of discrete petals, sepals, and tepals, or, if perianth fusion occurs, for numbers of calyx, corolla, or perianth lobes (see later discussion). Perianth, calyx, or corolla merosity is

FIGURE 9.17 Perianth cyclcy. **A.** Biseriate, homochlamydeous, with outer and inner perianth parts (tepals) similar. **B.** Biseriate, dichlamydeous, with distinct calyx and corolla. **C.** Uniseriate, with a single whorl of perianth parts, by default termed a calyx.

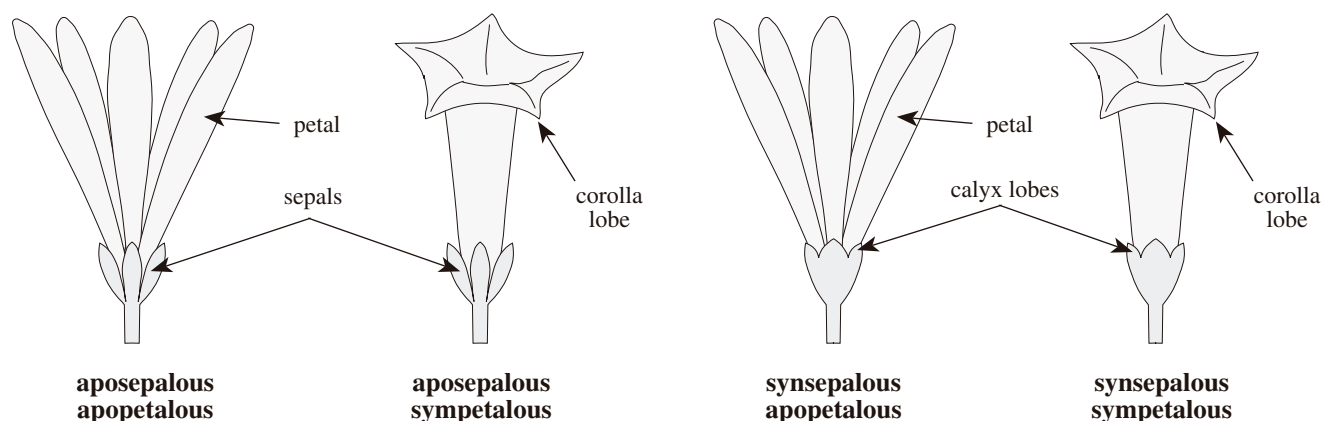


FIGURE 9.18 Perianth fusion.

usually designated as a simple number, although terms such as **bimerous** (a whorl with two members), **trimerous** (a whorl with three members), **tetramerous** (a whorl with four members), **pentamerous** (a whorl with five members), etc., can be used. Terms for absence of parts include **achlamydeous**, lacking a perianth altogether, **apetalous**, having no petals or corolla, and **asepalous**, having no sepals or calyx.

PERIANTH FUSION (Figure 9.18)

The term **perianth fusion** deals with the apparent fusion of perianth parts to one another. (This character may be treated separately as *calyx* or *corolla fusion*.) If sepals, petals, or tepals are discrete and unfused, the respective terms **aposepalous**, **apopetalous**, and **apotepalous** may be used. If sepals, petals, or tepals appear to be fused (even slightly at the base), the respective terms **synsepalous**, **sympetalous**, and **syntepalous** are used. The fusion of perianth parts does not usually occur as a separate event, e.g., petals fusing together after they are individually formed. The fusion is apparent, and typically results by the growth of a common floral primordium at the base of the calyx, corolla, or perianth.

Perianth fusion results in the development of a tubelike or cuplike structure (the region of fusion) in the calyx, corolla, or perianth. If little fusion occurs, the tubelike region occurs only at the base and gives rise to calyx, corolla, or perianth lobes.

PERIANTH TYPE (Figure 9.19)

Perianth type can include aspects of the entire perianth; however it could include aspects of only the calyx, corolla, or hypanthium (if present). Generally, perianth type is based on the structure of the corolla alone, in which case it could

logically be termed *corolla type*. The terminology for perianth type takes into account various aspects of shape, fusion, orientation, and merosity. Perianth type is often of systematic value and may be diagnostic for certain clades of angiosperms. The perianth type typically reflects adaptive features related to pollination biology, such as attracting a pollinator or better effecting the transfer of pollen. Some perianths are highly modified for other functions, such as the **lodicules** of grasses, which are reduced perianth parts that, upon swelling, open up the grass floret (see Inflorescence Type, later, and Poaceae of Chapter 7).

Specific perianth types include the following: **bilabiate**, two-lipped, with two, generally upper and lower segments, as in many Lamiaceae; **calyptrate/operculate**, having calyx and corolla fused into a cap that falls off as a unit, as in *Eucalyptus*; **campanulate**, bell-shaped, with a basally rounded flaring tube about as broad as long and flaring lobes, as in *Campanula* (may also be used for bell-shaped apopetalous corolla or apotepalous perianth); **carinate**, keeled, with a sharp median fold, usually on the abaxial side; **coronate**, with a tubular or flaring perianth or staminal outgrowth, as in *Narcissus*, *Asclepias* spp.; **cruciate**, with four distinct petals in cross form, as in many Brassicaceae; **cucullate/galeate**, hooded, with an abaxially concave posterior lip; **disk**, having an actinomorphic, tubular corolla with flaring lobes, as in some Asteraceae; **infundibular**, funnel-shaped, with a tubular base and continuously expanded apex, as in *Ipomoea*, morning glory; **lingulate/ray**, strap- or tongue-shaped, as in certain Asteraceae; **papilionaceous**, with one large posterior petal (banner or standard), two inner, lateral petals (wings), and two usually apically connate lower petals (keel), the floral structure of the Faboideae (Fabaceae); **personate**, two-lipped, with the upper arched and the lower protruding into the corolla throat, as in *Antirrhinum*, snapdragon; **ray**, having

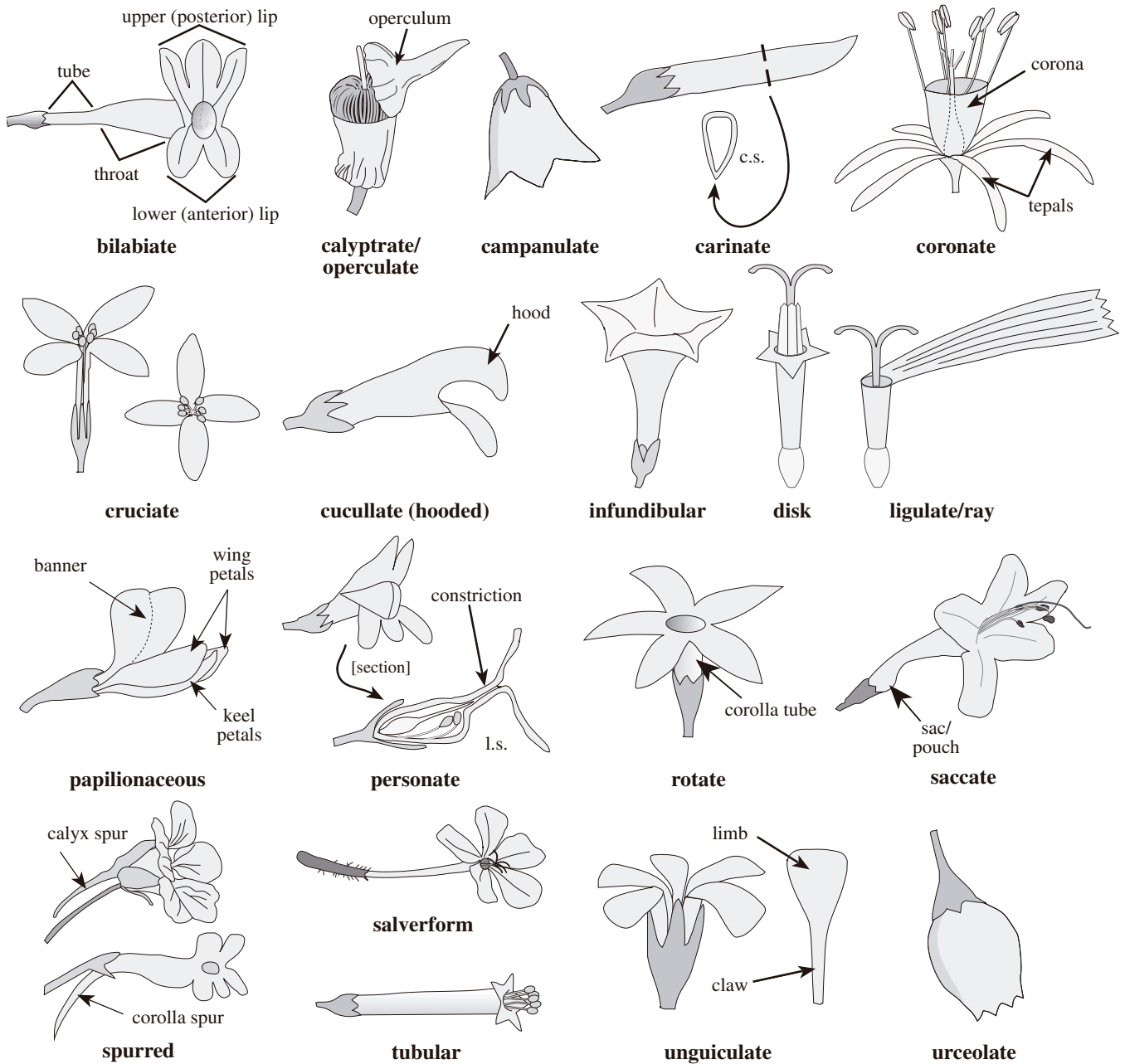


FIGURE 9.19 Perianth types/parts. (c.s. = cross-section; l.s. = longitudinal section)

a short, tubular corolla with a single, elongate, strap-like apical extension, as in some Asteraceae; **rotate**, with a short tube and wide limbs oriented at right angles to the tube, as in *Phlox*; **saccate**, having a pouchlike evagination; **salverform**, trumpet-shaped; with a long, slender tube and flaring limbs at right angles to tube; **tubular**, mostly cylindrical; **unguiculate**, clawed, as in many Brassicaceae, Caryophyllaceae;

unilabiate, one-lipped; and **urceolate**, urn-shaped, expanded at base and constricted at apex, as in many Ericaceae.

PERIANTH AESTIVATION (Figure 9.20)

Perianth aestivation is defined by the position, arrangement, and overlapping of floral perianth parts. Aestivation can be an important systematic character for delimiting or diagnosing

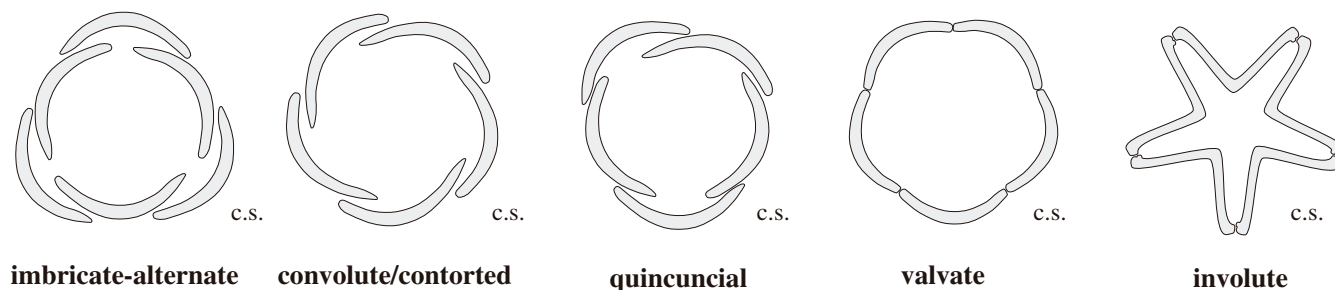


FIGURE 9.20 Perianth aestivation.

some flowering plant taxa. In practice, aestivation is best observed by making hand sections of mature flower buds, because after anthesis, the perianth aestivation may be obscured. For very small flowers, histological sectioning may be needed to clearly see the aestivation type.

Some standard perianth aestivation terms are as follows: **imbricate**, general term for overlapping perianth parts; **convolute** or **contorted**, imbricate with perianth parts of a single whorl overlapping at one margin, being overlapped at the other, as in the corolla of many Malvaceae; **imbricate-alternate**, imbricate with the outer whorl of perianth parts (sepals or outer tepals) alternating with (along different radii) the inner whorl of perianth parts (petals or inner tepals); **quincuncial**, imbricate with perianth parts of a single pentamerous whorl having two members overlapping at both margins, two being overlapped at both margins, and one overlapping only at one margin; **valvate**, with a whorl of perianth parts meeting at the margins, not overlapping; and **involute**, valvate with each perianth part induplicate (folded longitudinally inward along central axis).

ANDROECIUM

The androecium consists of all the floral male (pollen-producing) reproductive organs, the units of which are stamens. Stamens are interpreted as being modified, sporangia-bearing leaves or microsporophylls. Stamens initiate as primordia from the flower apical meristem, but at maturity are attached to the receptacle, corolla (having an epipetalous stamen fusion; see below), hypanthium rim, or **staminal disk**, a fleshy, elevated, often nectariferous cushion of tissue.

STAMEN TYPE (Figure 9.21)

There are two basic **stamen types**: laminar and filamentous (although intermediates can occur). **Laminar** stamens possess a leaflike, dorsiventrally flattened structure bearing two **thecae** (pairs of microsporangia), these typically on the adaxial surface. Laminar stamens may represent the ancestral type in flowering plants, although they have evolved secondarily

in some groups. **Filamentous** stamens are far more common, having a stalklike, generally terete **filament** with a discrete pollen-bearing part, the **anther**.

In some taxa one or more stamens will initially form but will be nonfertile. Such a sterile stamen is termed a **staminode** or **staminodium**. Staminodes may resemble the fertile stamens and can only be identified by determining if viable pollen is released. Other staminodes may be highly modified in structure, being petaloid, clavate (clublike), nectariferous, or very reduced and vestigial. Staminodes may or may not possess an **antherode**, a sterile antherlike structure.

STAMEN ARRANGEMENT, CYCLE, AND POSITION

Stamen arrangement (Figure 9.22) is the placement of stamens relative to one another (see General Terminology). Two basic stamen arrangements are **spiral**, with stamens arranged in a spiral, and **whorled**, with stamens in one or more discrete whorls or series. Additional stamen arrangement types consider the relative lengths of stamens to one another: **didymous**, with stamens in two equal pairs; **didynamous**, with stamens in two unequal pairs (as in many Bignoniaceae, Lamiaceae, Scrophulariaceae, etc.); and **tetradynamous**, with stamens in two groups of four long and two short (typical of the Brassicaceae).

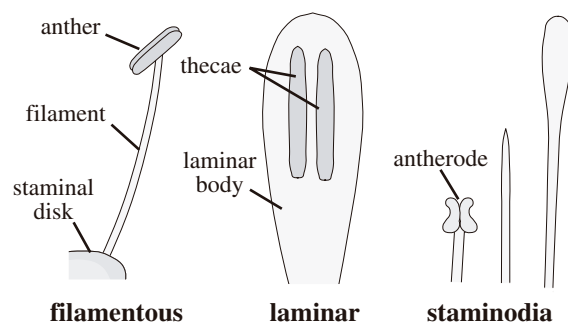


FIGURE 9.21 Stamen types and parts.

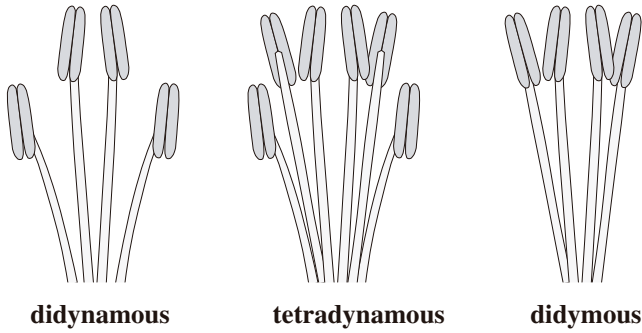


FIGURE 9.22 Stamen arrangement.

Stamen cycly (Figure 9.23) refers to the number of whorls or series of stamens present (applying only if the stamens are whorled to begin with). The two major types of stamen cycly are **uniserial**, having a single whorl of stamens, and **biseriate**, with two whorls of stamens. If additional whorls are present, the terms *triseriate*, *tetraseriate*, etc., can be used.

Stamen position (Figure 9.23) is the placement of stamens relative to other, unlike floral parts, in particular to the sepals and petals. An **antisepalous** (also called *antesepalous*) stamen position is one in which the point of stamen attachment is in line with (opposite) the sepals or calyx lobes; similarly, **alternipetalous** means having the stamens positioned between the petals or corolla lobes. Antisepalous and alternipetalous are usually synonymous because (in a biserial perianth) petals/corolla lobes are almost always inserted between sepals/calyx lobes; however, one should describe only what is evident, such that either or both terms may be used. Antisepalous or alternipetalous stamens are very common in taxa with uniserial stamens.

An **antipetalous** (also called *antepetalous*) stamen position is one in which the point of attachment is in line with (opposite) the petals or corolla lobes; **alternisepalous** means

that the stamens are positioned between the sepals or calyx lobes. Antipetalous and alternisepalous are usually synonymous (for the same reason cited earlier). An antipetalous/alternisepalous stamen position is relatively rare and may be diagnostic for specific groups, such as the Primulaceae and Rhamnaceae.

Two other stamen position terms apply to stamens that are biserial: **diplostemonous**, in which the outer whorl is opposite the sepals and the inner opposite the petals, and **obdiplostemonous**, in which the outer whorl is opposite petals, the inner opposite sepals. Among taxa with biserial stamen cycly, a diplostemonous position is much more common; obdiplostemonous stamens are relatively rare, being diagnostic, e.g., for some Crassulaceae.

STAMEN ATTACHMENT AND INSERTION

Stamen attachment refers to the presence or absence of a stalk, being either **filamentous**, with a filament present, **sessile**, with filament absent, or **subsessile**, with filament very short and rudimentary. Laminal stamens are, by default, sessile.

Stamen insertion (Figure 9.24) can refer to either of two things. First, it can indicate whether stamens extend past the perianth or not, the two terms being **exserted** (also termed *phaneranthous*), with stamens protruding beyond the perianth, and **inserted** (also termed *cryptanthous*), with stamens included within the perianth. Insertion may also correspond to the *point of insertion*, which is the point of adnation of an epipetalous stamen to the corolla (see later discussion). Examples of the latter usage are the stamens are inserted halfway up the corolla tube or stamens are inserted unequally (meaning they are inserted at different levels along the length of, say, a corolla tube). Stamen insertion, by either usage, is generally indicative of an adaptation for some particular pollination mechanism, functioning to present the anthers effectively to an animal pollinator.

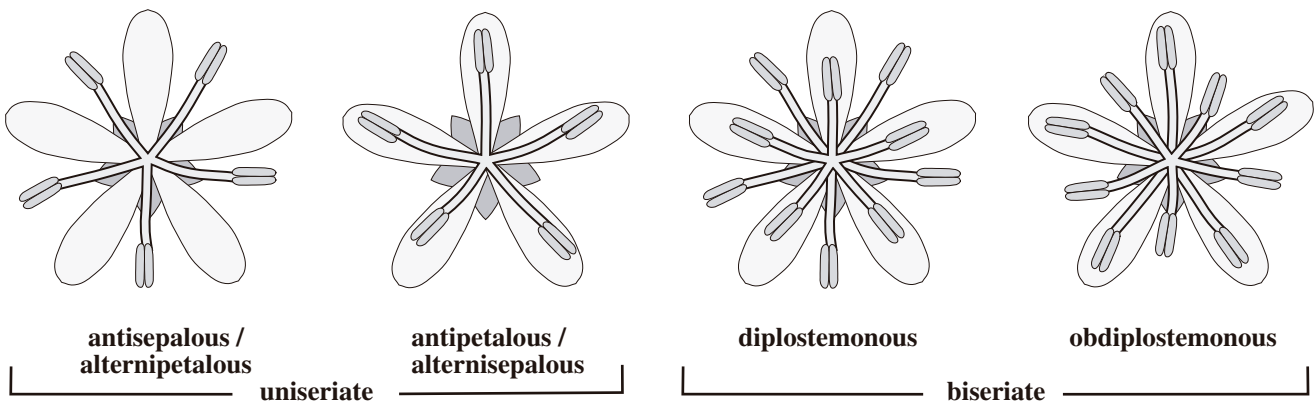


FIGURE 9.23 Stamen cycly (uniserial or biserial) and position.

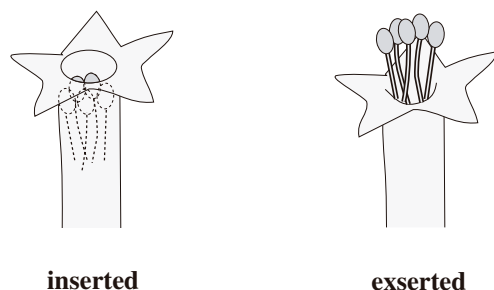


FIGURE 9.24 Stamen insertion.

STAMEN FUSION (Figure 9.25)

Stamen fusion refers to whether and how stamens are fused. The general terms *distinct* (unfused to one another), *connate* (fused to one another), *free* (unfused to a different structure), and *adnate* (fused to a different structure) may be used (see General Terminology). Common specialized terms are **apostemonous**, with stamens unfused (both distinct and free); **diadelphous**, with two groups of stamens, each connate by filaments only, as in many Faboideae (Fabaceae), which typically have nine stamens fused most of their length and one fused only at the base or not at all; **epipetalous** (also called *petalostemonous*), with stamens adnate to (inserted on) petals or the corolla (the terms *epitepalous* and *episepalous* can be used for adnation of stamens to tepals or sepals, respectively); **monadelphous**, with one group of stamens connate by their filaments, as in Malvaceae; and **syngenesious**, with anthers connate but filaments distinct, diagnostic of the Asteraceae. Stamen fusion, like stamen insertion, typically functions as a presentation mechanism for animal pollination.

ANTHER PARTS, TYPE, AND ATTACHMENT (Figure 9.26)

Anthers are discrete pollen containing units, found in the stamens of the great majority of angiosperms. Anthers typically consist of two compartments called **thecae** (singular *theca*),

with each theca containing two microsporangia (the fusion product of which is a *locule*). (Thus, anthers are typically tetrasporangiate.) The tissue between and interconnecting the two thecae is termed the **connective**, to which the filament (if present) is attached. **Microsporangia** are the sites of production of pollen grains, the immature male gametophytes of seed plants.

Various anther types occur, as determined by their internal structure. The typical anther is **dithecal**, having two thecae with typically four microsporangia. In a very few taxa, such as the Cannaceae and Malvaceae, anthers are **monotheical**, having one theca with typically two microsporangia. Finally, an extreme type of anther is the **pollinium**, a typically dithecal anther in which all the pollen grains of both thecae (Orchidaceae) or of adjacent thecae (*Asclepias*) are fused together as a single mass. The pollinia of the Orchidaceae and *Asclepias* have different developmental origins and structures.

Anther attachment refers to the position or morphology of attachment of the filament to the anther. Standard anther attachment types are **basifixed**, anther attached at its base to apex of the filament; **dorsifixed**, anther attached dorsally and medially to the apex of the filament; and **subbasifixed**, anther attached near its base to the apex of the filament. A **versatile** anther attachment is one in which the anther freely pivots (teeter-totter) at the point of attachment with the filament; versatile anthers may be dorsifixed, basifixed, or subbasifixed.

ANTHER DEHISCENCE

Anther dehiscence refers to the opening of the anther in releasing pollen grains. **Anther dehiscence type** (Figure 9.27) is the physical mechanism of anther dehiscence. The most common, and ancestral, anther dehiscence type is **longitudinal**, dehiscing along a suture parallel to the long axis of the thecae. Other types are rare and specific to given groups, including **poricidal**, dehiscing by a pore at one end of the thecae, such as the Ericaceae; **transverse**, dehiscing at right angles to the

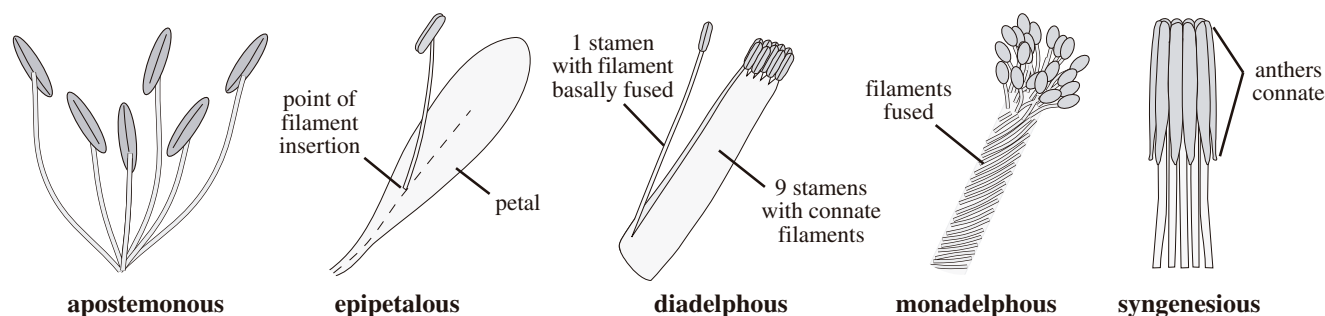


FIGURE 9.25 Stamen fusion.

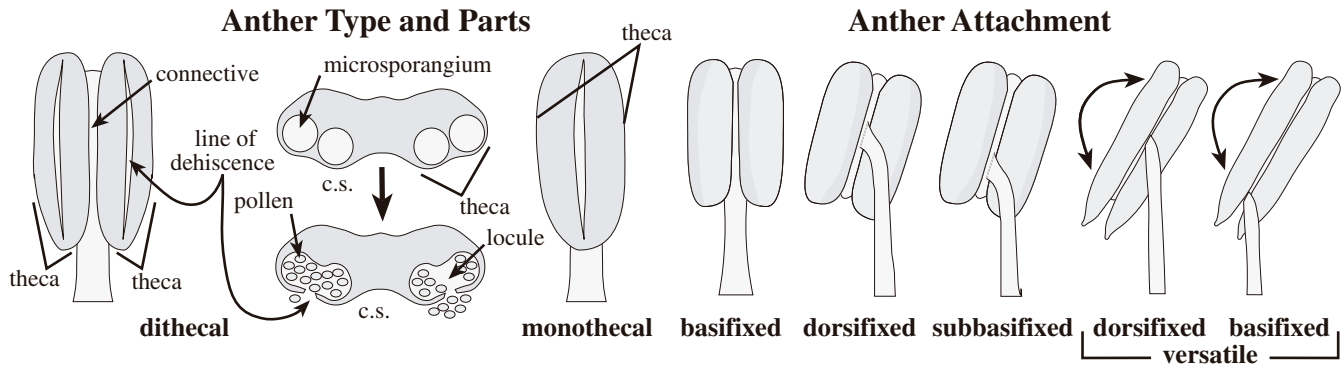


FIGURE 9.26 Anther types, parts, and attachment (c.s. = cross-section)

long axis of the theca; and **valvular**, dehiscing through a pore covered by a flap of tissue, as in the Lauraceae.

Anther dehiscence direction (Figure 9.28) indicates the position of the anther opening relative to the center of the flower or to the ground. Anther dehiscence direction is best detected when the anthers are immature (e.g., in bud) or just beginning to open. After dehiscence, the anthers usually shrivel and twist, obscuring the original direction in which they opened. Common types of dehiscence direction are: **extrorse**, dehiscing outward, away from the flower center; **introrse**, dehiscing inward, toward the flower center; and **latrorse**, dehiscing laterally, to the sides. In horizontally oriented flowers, anthers may face **upward** or **downward**, relative to the ground.

One fine point of anther dehiscence direction concerns some flowers, in which at least some of the stamens have one direction early in development but become reoriented to another direction at maturity. In such a case, the dehiscence direction can be described both in the early developmental stage and in the mature stage. For example, a common condition is one in which the anthers are introrse early in

development, but reorient to the top of the flower, with all the anthers facing downward. Such a dehiscence direction can be described as **introrse** early in development (based on observation of buds), and **downward** at maturity (see Figure 9.28). (In another example, the poricidal anthers of members of the Ericaceae are extrorse early in development, but introrse at maturity by inversion; see Chapter 8.)

NECTARIES

Nectaries are specialized nectar-producing structures of the flower (Figure 9.13). Nectar is a solution of one or more sugars and various other compounds and functions as an attractant (a reward) to promote animal pollination. Nectaries may be **padlike**, developing as a discrete pad of tissue extending only part-way around the base of the flower. Commonly, a floral **disk**, consisting of a disk-like or doughnut-shaped mass of tissue surrounding the ovary base or top, functions as a nectary. These **nectariferous disks** may be inner to (**intrastaminal**), beneath (**staminal**), or outer to (**extrastaminal**) the androecium. A **perigonal nectary** is one on the perianth, usually at the base of sepals, petals, or tepals. **Septal nectaries** are specialized tissues embedded *within* the septae of an ovary, secreting nectar via a pore at the ovary base or apex.

Note that other specialized glands may secrete non-sugar compounds that function as a pollination reward, such as waxes by members of the Krameriaceae. These are not termed nectaries, but are simply called glands, e.g., wax glands.

GYNOECIUM, CARPEL, AND PISTIL

The **gynoecium** refers to all female organs of a flower (Figure 9.29). The unit of the gynoecium is the **carpel**, defined as a modified, typically conduplicate megasporophyll that encloses one or more ovules (see Chapter 6). The carpel is one of the major features (apomorphies) that make angiosperms

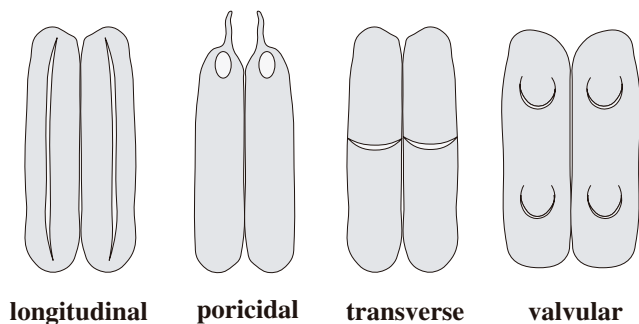


FIGURE 9.27 Anther dehiscence types.

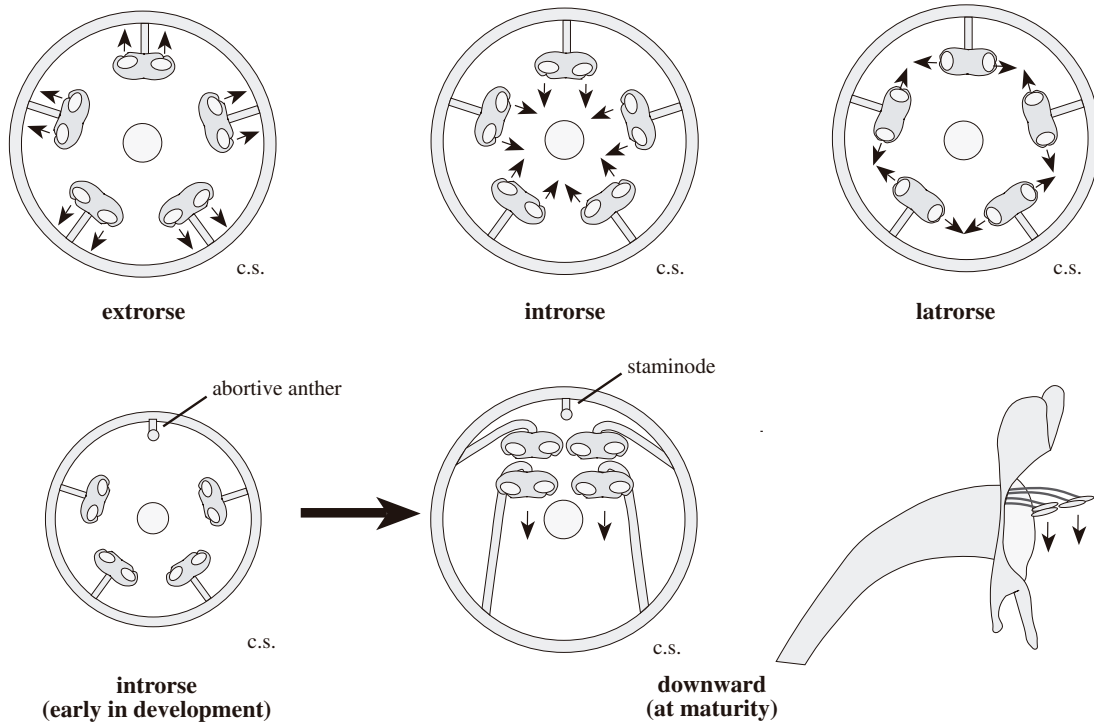


FIGURE 9.28 Anther dehiscence direction. (c.s. = cross-section)

unique within the seed plants. Like all flower parts, a carpel is interpreted as a modified leaf, in this case a megasporophyll, defined as a reproductive leaf bearing megasporangia (which in seed plants are components of the ovules). Carpels, in fact, may develop as dorsiventrally flattened leaves that fold conduplicately, ultimately enclosing the ovules.

A **pistil** is that part of the gynoecium composed of an **ovary**, one or more **styles**, and/or one or more **stigmas** (see later discussion). The **ovary** is the part of the pistil containing the ovules. A **style** is a generally stalklike, non-ovule-bearing portion of the pistil between the stigma and ovary. Styles may

be absent in some pistils. A **stigma** is the pollen-receptive portion of the pistil. Stigmas may be discrete structures or they may be a region (the *stigmatic region*) of a style or style branch, e.g., the stigmatic lines on the styles of Asteraceae pistils. Finally, the term **stipe** is used for a basal stalk of the pistil; stipes are usually absent. [Note that *stipe* is also used as a synonym for a leaf petiole, especially that of ferns.]

Pistils or ovaries may be **simple**, composed of one carpel, or **compound**, composed of two or more carpels (see Carpel Number). By convention, if there is more than one ovary, style, or stigma, but if any of these appear fused in any way

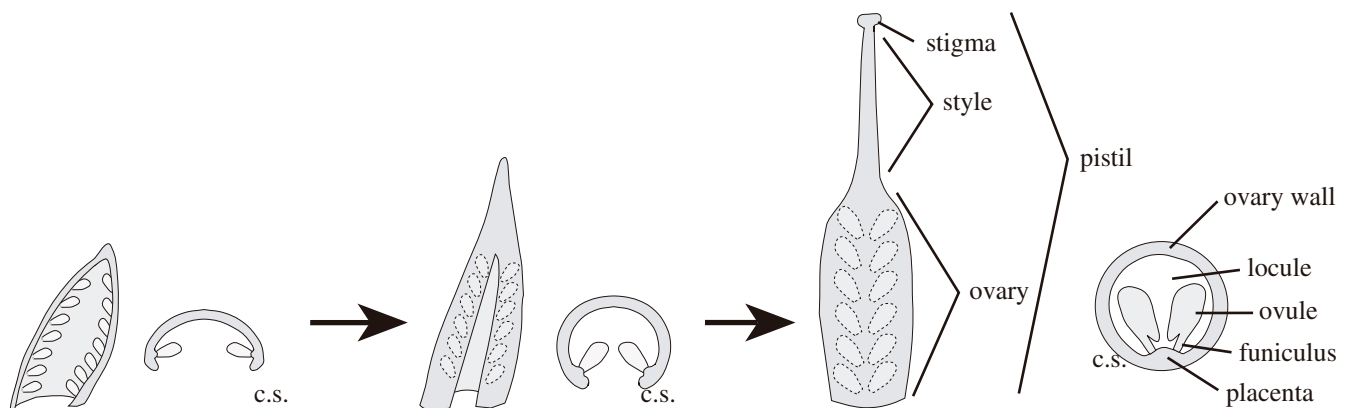


FIGURE 9.29 Gynoecium: carpel development. (c.s. = cross-section; l.s. = longitudinal section)

(e.g., three apparent ovaries fused at the base), they are all part of the same pistil. (One unique case are the Asclepiadoids, in which the gynoecium consists of two carpels made up of two distinct ovaries and styles but a single stigma joining the styles; because the stigmas of the two carpels are connate, the whole structure is termed a single pistil.)

Within the ovary, a **septum** (plural *septa*) is a partition or cross-wall. A **locule** is an ovary cavity, enclosed by the ovary walls and septa. Locule number may be an important systematic character; the term *unilocular* may be used for an ovary with one locule, *pleurilocular* for an ovary with two or more (typically many) locules. **Placenta** are the tissues of the ovary that bear the ovules, the immature seeds. A **funiculus** is a stalk that may lead from the placenta to the ovule. A **column** is the central axis to which septae and/or placentae are attached in axile or free-central placentation (see later discussion).

GYNOECIAL FUSION (Figures 9.30, 9.31)

Fusion of carpels is a very important systematic character, the features of which are characteristic of major taxonomic groups. An **apocarpous** gynoecial fusion is one in which the carpels are distinct. An apocarpous gynoecium is generally thought to be the ancestral condition in the angiosperms. In contrast, a **syncarpous** gynoecial fusion is one in which carpels are connate (the pistil *compound*) and is the most common type in flowering plants. In a syncarpous gynoecium, the degree of carpel fusion can vary considerably; from connation only at the extreme base (having a strongly lobed ovary), to fusion into one, unlobed ovary but distinct styles and/or stigmas, to complete fusion with one ovary, style, and stigma. Fusion of carpels can determine the placentation type (Figure 9.31; see later discussion). Last, if the gynoecium is composed of a single carpel (in which fusion is really inapplicable), the term **unicarpellous** is used.

CARPEL/LOCULE NUMBER (Figure 9.30)

Carpel and locule number are important characters in angiosperm systematics. **Locule number** is generally easy to determine from ovary cross- and/or longitudinal sections, being equivalent to the number of wall-enclosed chambers within the ovary. In a general sense, ovaries may be **unilocular**, with a single locule, or **plurilocular**, having two or more locules. In some angiosperms, septa may divide the ovary into chambers in one region, such as the ovary base, but not in another region, such as the ovary apex; in such a case, the chambers below are continuous with one chamber above, and the locule number is technically 1, or unilocular.

Carpel number is often critical in classification and identification of flowering plants. It is determined as follows. If the gynoecium is apocarpous, the number of carpels is equal to the number of pistils; this is because each pistil is equivalent to a single carpel in any apocarpous gynoecium. If there is a single pistil, that pistil can be equivalent to one carpel (i.e., unicarpellous) or be composed of any number of fused carpels. For one pistil the carpel number is determined (in sequence) as follows. First, carpel number is equal to the number of styles or stigmas, if either of these is greater than 1. This is true regardless of the structure of the ovary because each of the styles or stigmas is a part of a carpel or is interpreted as a vestige of an ancestral carpel. (For example, pistils of all members of the Asteraceae have two styles and stigmas, and thus carpel number is interpreted as 2. This is true even though there is but one locule, ovule, and placenta; the two styles are interpreted as ancestral vestiges of a two-carpellate pistil, which became evolutionarily reduced to a single ovuled and loculed structure.) Second, if a single pistil has only one style and stigma, the ovary must be dissected to reveal the carpel number. If the ovary is plurilocular, then locule number is generally equal to the number of carpels. Each locule, in such a case, represents the chamber of the

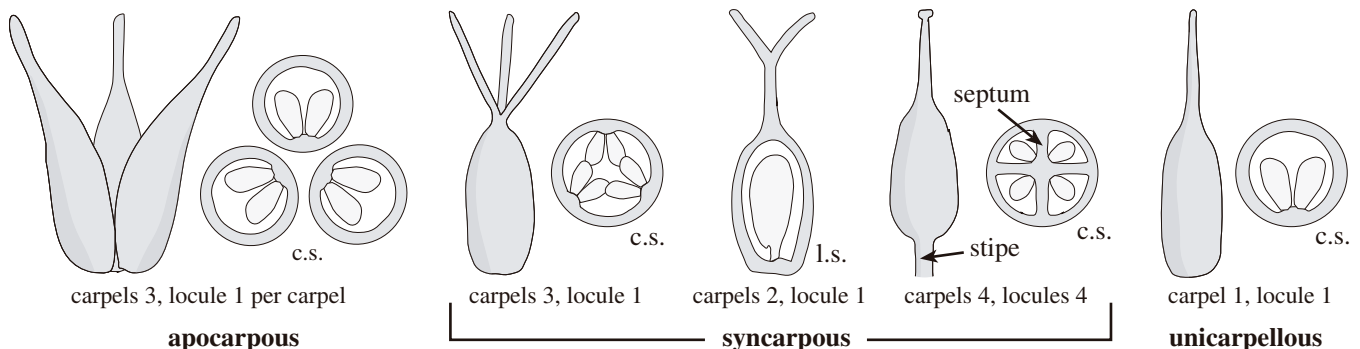


FIGURE 9.30 Gynoecial fusion, carpel number, and locule number. (c.s. = cross section, l.s. = longitudinal section)

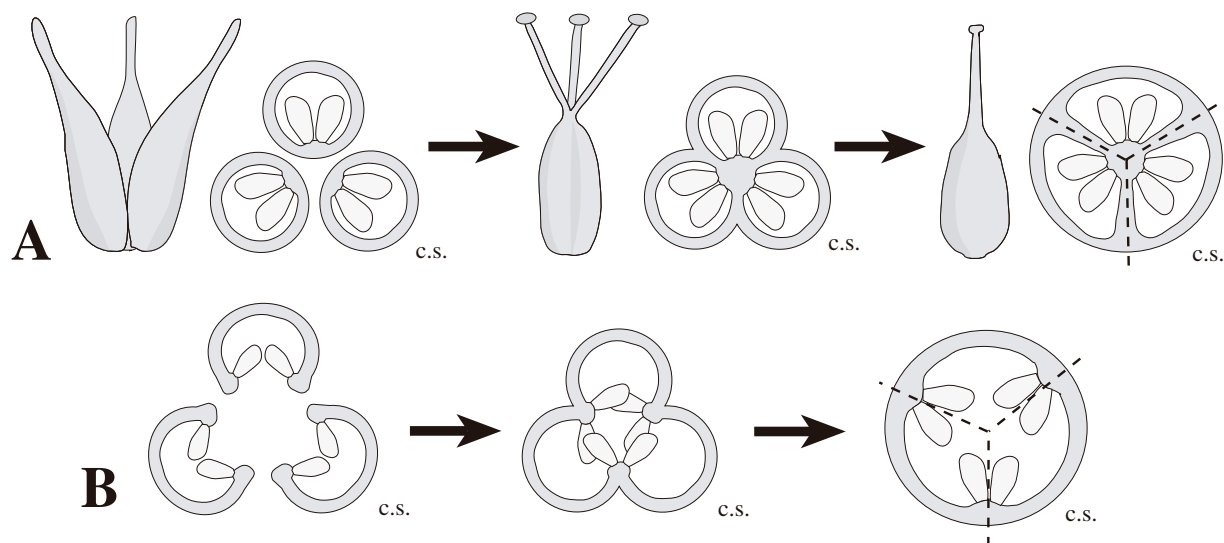


FIGURE 9.31 Evolutionary sequences of carpel fusion. **A.** Sequence leading to axile placentation. **B.** Sequence leading to parietal placentation. Carpel boundaries shown with dashed lines. (c.s. = cross-section)

original ancestral or developmental carpel (except in some gynobasic taxa; see later discussion). Finally, if the ovary is unilocular, the number of carpels is equal to the number of placentae. For example, a violet, with one pistil, one style/stigma, and one locule, has three carpels because of the three placentae (having parietal placentation). (Exceptions to the last two rules are the gynobasic taxa of the Lamiaceae and Boraginaceae, s.s. In both of these groups, each of the two carpels is bisected early in development by a so-called false septum, such that the mature ovary typically has four locules, each with a single placenta and ovule. Thus, in this case, the number of locules and placentae, which is four, is twice that of the number of carpels.)

OVARY ATTACHMENT AND POSITION

Ovary attachment deals with the presence or absence of a basal stalk or **stipe**. A **sessile** ovary is one lacking a stipe and is by far the most common situation. A **stipitate** ovary is one having a **stipe** and is relatively rare (Figure 9.30).

Ovary position (Figure 9.32) assesses the position or placement of the ovary relative to the other floral parts: hypanthium, calyx, corolla, and androecium. A **superior** ovary is one with sepals, petals, and stamens, and/or hypanthium attached at the base of the ovary. An **inferior** ovary position has sepals, petals, stamens, and/or hypanthium attached at the ovary apex. A range of intermediates between superior and inferior ovaries can occur; the term **half-inferior** is used for sepals, petals, stamens and/or hypanthium attached at the middle of the ovary.

PERIANTH/ANDROECIAL POSITION (Figure 9.32)

Perianth/androecial position describes placement of the perianth and androecium relative both to the ovary and to a hypanthium, if present. Although used widely, perianth/androecial position may be simply substituted with a description of both ovary position and hypanthium presence/absence.

Three perianth/androecial position terms describe a flower without a hypanthium (and are rather repetitious with ovary position). The term **hypogynous** is used for sepals, petals, and stamens attached at base of a superior ovary. **Epigynous** refers to the sepals, petals, and stamens attached at apex of an inferior ovary. **Epihypogynous** is used for sepals, petals, and stamens attached at middle of the ovary, the ovary being half-inferior.

Other perianth/androecial position terms denote the presence of a hypanthium, with the sepals, petals, and stamens attached to the hypanthium rim. **Perigynous** denotes a hypanthium attached at the base of a superior ovary. **Epiperigynous** denotes a hypanthium attached at the apex of an inferior ovary. (The awkward term **epihypoperigynous** may be used to describe a hypanthium attached at the middle of a half-inferior ovary.)

PLACENTATION (Figure 9.33)

Placentation refers to the positioning of the ovules and takes into account the number and position of placentae, septa, and locules. Determining placentation requires probing or making a cross and/or longitudinal section of the ovary.

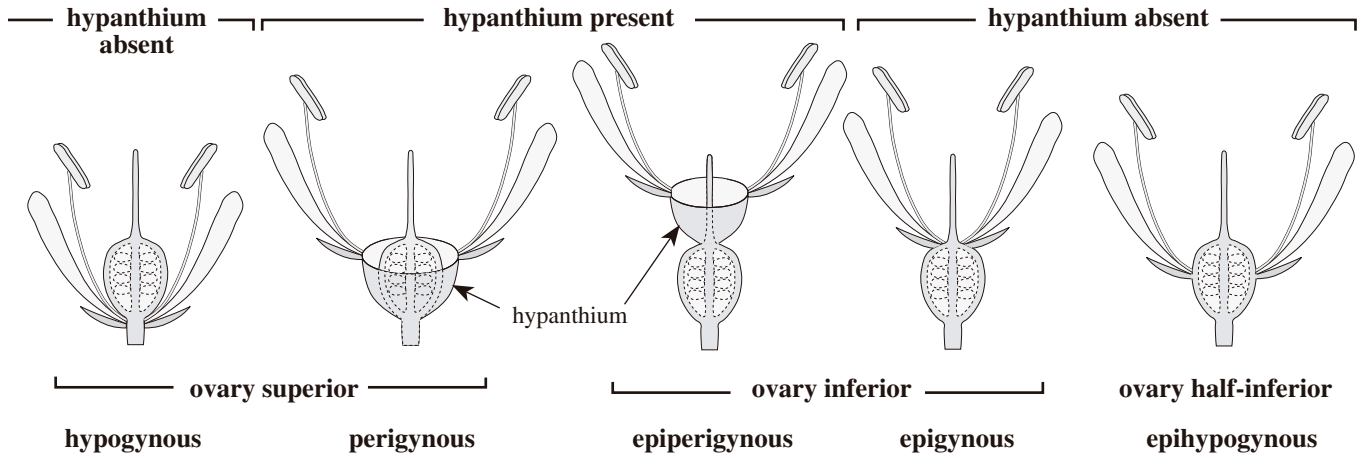


FIGURE 9.32 Ovary position and perianth/androecial position.

Standard placentation types are **axile**, with the placentae arising from the column in a compound ovary with septa, common in many flowering plants such as the Liliaceae; **apical** or **pendulous**, with a placenta at the top of the ovary; **apical-axile**, with two or more placentae at the top of a septate ovary, as occurs in the Apiaceae; **basal**, with a placenta at the base of the ovary, as occurs in the Asteraceae and Poaceae; **free-central**, with the placentae along the column in a compound ovary without septa, such as in the Caryophyllaceae; **laminar**, with ovules arising from the surface of the septae; **marginal**, with the placentae along the margin of a unilocular (simple) ovary, as in the Fabaceae; **parietal**, with the placentae on the ovary walls or upon intruding partitions of a unilocular, compound ovary, such as in the Violaceae; **parietal-axile**, with the placentae at the junction of the septum and ovary wall of a two or more loculate ovary, such as in the Brassicaceae; and **parietal-septate**, with placentae

on the inner ovary walls but within septate locules, as in some Aizoaceae.

STYLE POSITION/STRUCTURAL TYPE (Figure 9.34)

Style position is the placement of the style relative to the body of the ovary. A **terminal** or **apical** style position is one arising at the ovary apex; this is by far the most common type. A **subapical** style arises to one side, near and slightly below the ovary apex. A **lateral** style position is one arising at the side of an ovary, as in members of the Rosaceae, such as *Fragaria*. Finally, a **gynobasic** style arises from the base of the ovary. Gynobasic styles are characteristic of the Boraginaceae, s.s. and of most Lamiaceae, in which the style arises from the base and center of a strongly lobed ovary.

Styles may be structurally specialized in some taxa. One specialized **style structural type** is a stylar **beak**, a persistent, extended style or basal (to subbasal) stylar region.

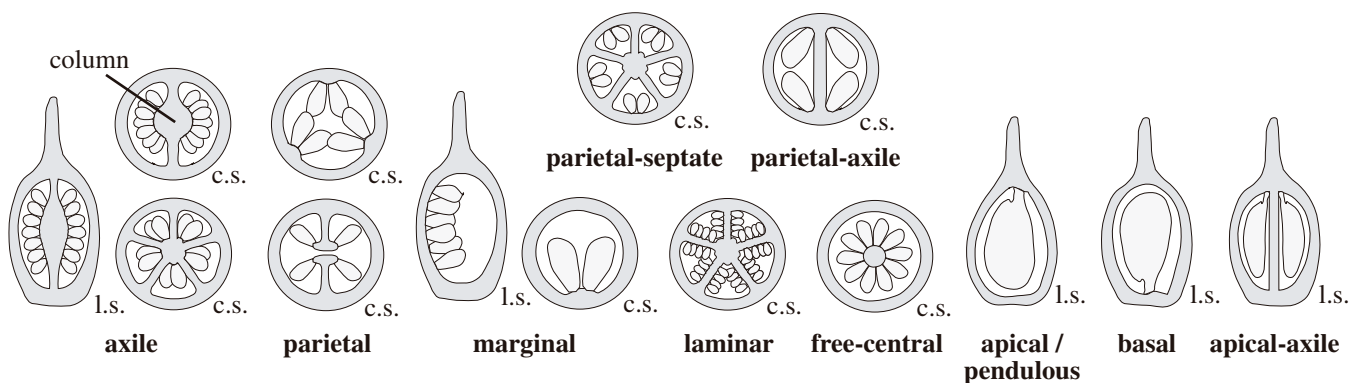


FIGURE 9.33 Gynoecium: placentation. (c.s. = cross-section; l.s. = longitudinal section).

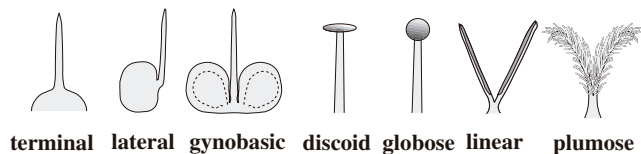


FIGURE 9.34 Gynoecium: style position (left) and stigma/stigmatic region type (right).

A beak is typically accrescent and elongates during fruit formation. Beaks function in fruit dispersal, as in members of the Asteraceae (e.g., *Taraxacum*, dandelion) or Geraniaceae (e.g., *Geranium*).

STIGMA/STIGMATIC REGION TYPES (Figure 9.34)

The term **stigma** is used for a discrete structure that is receptive to pollen on the entire surface, whereas *stigmatic region* may be used for that portion of a larger structure (generally a style or style branch) that is receptive to pollen. General shape terms may be used to describe **stigma** or **stigmatic region types**. A few common stigma or stigmatic region types are **discoic**, with stigma(s) disk-shaped; **globose**, with stigma(s) spherical in shape; **linear**, with stigmas or stigmatic tissue long and narrow in shape; and **plumose**, stigmas with feathery, trichome-like extensions, often found in wind-pollinated taxa (e.g., in Cyperaceae, Poaceae).

INFLORESCENCES

An inflorescence is a collection or aggregation of flowers on an individual plant. Inflorescences often function to enhance reproduction. For example, the aggregation of flowers in one location will make them visually more attractive to potential pollinators. Other inflorescences are related to very specialized reproductive mechanisms, examples being the spadices and associated spathes of some Araceae or the syconia of figs (see later discussion).

The structure of an inflorescence can be complicated, its elucidation requiring detailed developmental study.

INFLORESCENCE PARTS

Several terms deal with leaflike structures found in the inflorescence. An **inflorescence bract** is one that subtends not an individual flower but an inflorescence axis or a group of flowers. (Bracts that subtend an individual flower should be termed *floral bracts*; however, some sources do not make the distinction or will use *inflorescence bract* to refer to either.) A group or cluster of bracts subtending an entire inflorescence is termed an **involucre** (adjective *involute*);

a similar group of bracts subtending a unit of the inflorescence is an **involucel**. A **spathe** (adjective *spathaceous*) is an enlarged, sometimes colored bract subtending and usually enclosing an inflorescence; many Araceae are good examples of spathes, which subtend the spadix inflorescence (see later discussion). An **awn** is a bristlelike, apical appendage on the glumes or lemmas of grass (Poaceae) spikelets.

Other terms deal with various (stem) axes in an inflorescence. A **peduncle** (adjective *pedunculate*) is the stalk of an entire inflorescence. A **compound receptacle** (also called a *torus*) is a mass of tissue at the apex of a peduncle that bears more than one flower. A peduncle that lacks well-developed leaves, arising from a basal rosette of vegetative leaves is termed a **scape** (adjective *scapose*), the plant habit in such a case being acaulescent. A **rachis** is a major, central axis within an inflorescence. However, the central axis of a grass or sedge spikelet is a **rachilla**. Finally, a **ray** is a secondary axis of a compound umbel (see later discussion).

INFLORESCENCE POSITION

There are three major **inflorescence positions**, defined based on where the inflorescence develops: (1) **axillary**, in which the entire inflorescence is positioned in the axil of the nearest vegetative leaf; (2) **terminal**, in which the inflorescence develops as part of a terminal shoot that gave rise to the nearest vegetative leaves; and (3) **cauliflorous**, in which the inflorescence grows directly from a woody trunk. Three specialized inflorescence position terms for palms are **infrafoliar**, in which the inflorescence arises below the crownshaft, **interfoliar**, in which it arises within the crownshaft, and **suprafoliar**, in which it arises above the leaves of the crownshaft.

INFLORESCENCE DEVELOPMENT

Inflorescence development is a major aspect of defining inflorescence type. The two major inflorescence developmental types are determinate and indeterminate. A **determinate** inflorescence is one in which the apical meristem of the primary inflorescence axis terminates in a flower; typically, the terminal flower matures first, with subsequent maturation occurring from apex to base. Determinate inflorescences are characteristic of cymes. An **indeterminate** inflorescence is one in which the apical meristem of the primary inflorescence axis does not develop into a flower; typically, the basal flower matures first, with maturation occurring from base to apex. Indeterminate inflorescences include a number of types, such as spikes, racemes, and panicles (see later discussion).

INFLORESCENCE TYPE (Figures 9.35–9.37)

Inflorescences that have a common development and structure with respect to presence, number, arrangement, or orientation of bracts, axes, and certain specialized structures, define an **inflorescence type**. One difficulty with determining inflorescence type is simply delimiting its boundaries. Generally, an inflorescence is bounded by the lowest vegetative leaf. However, there may be a gradation between lower or basal vegetative leaves and small floral bracts, such that the delimitation of the inflorescence is somewhat arbitrary.

(Note that if an inflorescence consists of a single flower, it is termed **solitary**; a **scapose** inflorescence is one with one or more flowers on an essentially leafless peduncle or **scape**, usually arising from a basal rosette.)

Inflorescence types are valuable characters in systematics and are often characteristic of specific groups, such as the compound umbels of the Apiaceae, heads of the Asteraceae, and helicoid or scorpioid cymes of the Boraginaceae. Some inflorescence types are quite specialized adaptations for reproduction, such as the cyathia of Euphorbioids.

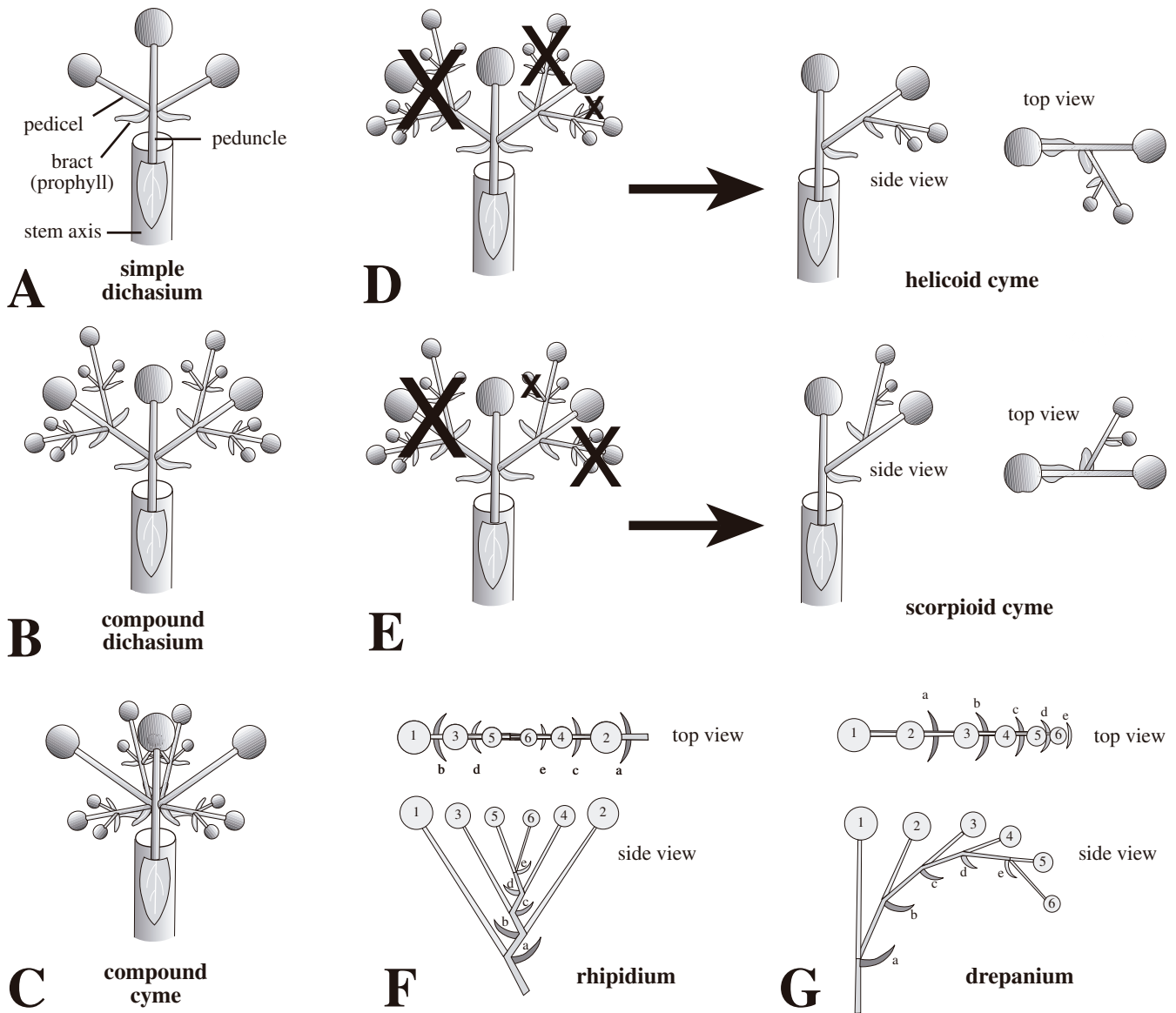


FIGURE 9.35 Determinate (cymose) inflorescences. **A–C.** Dichasial cymes. **A.** Simple dichasium. **B.** Compound dichasium. **C.** Compound cyme. **D–G.** Monochasial cymes. **D.** Helicoid cyme, showing derivation from compound dichasium by development of one axis on the side of the primary axis. **E.** Scorpioid cyme, showing derivation from compound dichasium by development of one axis on alternating sides of the primary axis. **F.** Rhipidium. **G.** Drepanium. (Terminology after and redrawn from Weberling 1989, by permission.)

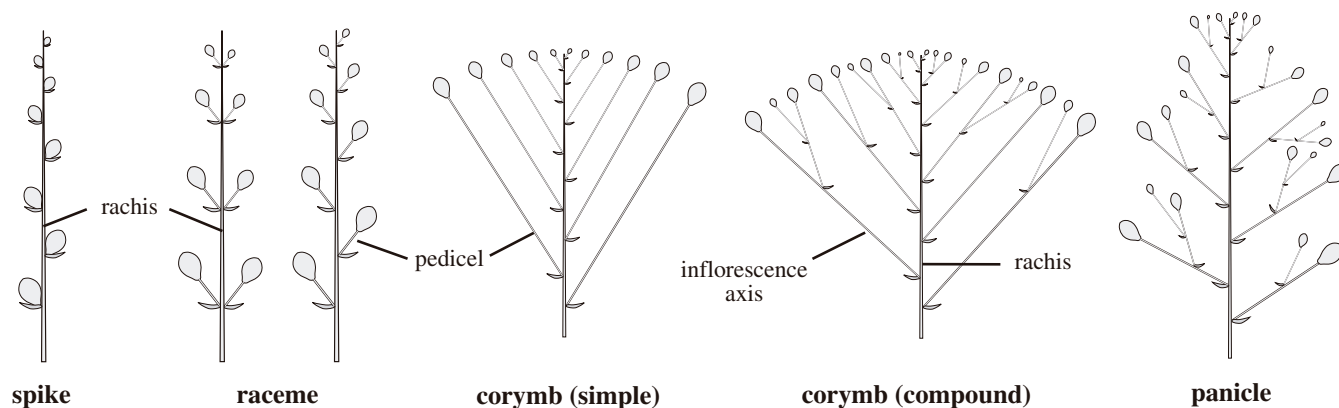


FIGURE 9.36 Indeterminate inflorescence types.

The term **cyme** (Figure 9.35) can be used as a general term to denote a determinate inflorescence. One type of cyme is the **dichasium**, one that develops along two axes, forming one or more pairs of opposite, lateral axes. A **simple dichasium** is a three-flowered cyme, having a single terminal flower and two, opposite lateral flowers, the pedicels of all of equal length; bracts typically subtend the two lateral flowers, although the bracts may be absent. (The term **cymule** may be used for a small, simple dichasium.) A **compound dichasium** is a many-flowered cyme of repeatedly branching simple dichasia units. In a compound dichasium, the branches are typically decussately arranged and are thus in multiple planes. Finally, a **compound cyme** is a branched cyme, similar to a compound dichasium but lacking a consistent dichasial branching pattern. Some compound cymes actually have the same branching pattern as a compound dichasium but with certain internodal axes being reduced or missing, yielding a more congested appearance.

A **monochasium** (Figure 9.35) is a cyme that develops along one axis only. (The terminology for monochasial cymes can vary from author to author, the following being just one.)

A **helicoid cyme** or **bostryx** is a monochasium in which the axes develop on only one side of each sequential axis, appearing coiled at least early in development. A **scorpioid cyme** or **cincinnus** is a monochasium in which the branches develop on alternating sides of each sequential axis, typically resulting in a geniculate (zig-zag) appearance. Both helicoid cymes and scorpioid cymes have branches or axes that are in more than one plane and can be viewed as being derived by reduction from the decussate branches of a compound dichasium. Two other monochasial cymes have, by definition, axes that are in one plane. A **drepanium** is a monochasium in which the axes develop on only one side of each sequential axis; like a helicoid cyme, drepania typically appear coiled at least early in development. (Drepania are treated as helicoid cymes in some terminology.) A **rhipidium** is a monochasium in which the branches develop on alternating sides of each sequential axis; like scorpioid cymes, rhipidia typically have a geniculate (zig-zag) appearance. (Rhipidia are treated as scorpioid cymes in some terminology.) In reality, these four monochasial structures may intergrade with one another. For example, a monochasium intermediate between a helicoid

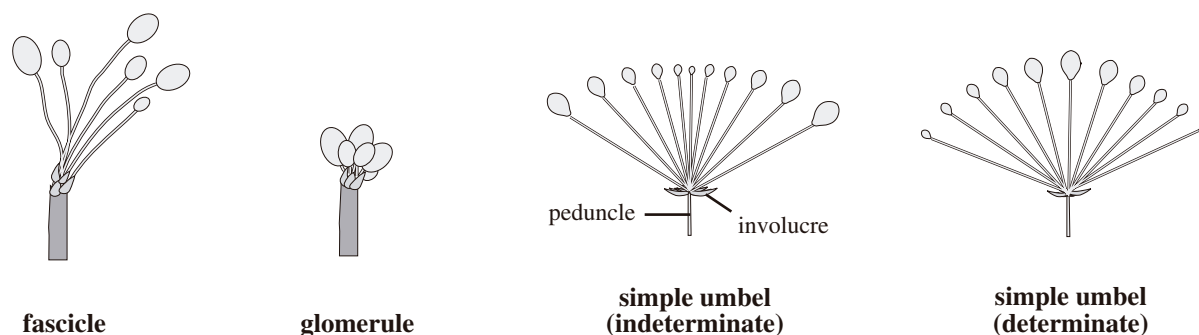


FIGURE 9.37 Indeterminate or determinate inflorescence types.

cyme and a drepanium may occur. Thus, simply using the term *monochasial cyme* may be best in lieu of more detailed observations and descriptions.

Several indeterminate inflorescence types are recognized (Figure 9.36). All of these generally lack a flower at the top of the main axis and develop from base to apex. A **spike** is an indeterminate inflorescence, consisting of a single axis bearing sessile flowers. Similarly, a **raceme** is an indeterminate inflorescence in which the single axis bears pedicellate flowers. A **panicle** is like a branched raceme, defined as an indeterminate inflorescence having several branched axes bearing pedicellate flowers. Finally, a **corymb** is an indeterminate inflorescence consisting of a single axis with lateral axes and/or pedicels bearing flat-topped or convex flowers. Corymbs can be either simple or compound. A **simple corymb** is unbranched, consisting of a central axis bearing pedicellate flowers, the collection of flowers being flat-topped or convex; simple corymbs are like racemes in which the lower pedicels are much more elongate than the upper. A **compound corymb** is branched, consisting of two or more orders of inflorescence axes bearing flat-topped or convex, pedicellate flowers; compound corymbs are like panicles in which the lower axes and pedicels are much more elongate than the upper.

Some inflorescences may be either determinate or indeterminate (Figure 9.37). A **simple umbel** is a determinate or indeterminate, flat-topped or convex inflorescence with pedicels attached at one point to a peduncle. Two inflorescences in which the flowers at the point of attachment appear congested are the fascicle and glomerule. A **fascicle** is a racemelike or paniclelike inflorescence with pedicellate flowers in which internodes between flowers are very short. A **glomerule** is an inflorescence of sessile or subsessile flowers in which the internodes between flowers are very short.

In some taxa an inflorescence will appear to be one type, but (upon detailed examination) is actually a modification of another type. For example, the term **pseudoumbel** is used for an inflorescence appearing like a simple umbel, but actually composed of condensed, monochasial cymes, as in the Alliaceae and Amaryllidaceae.

SECONDARY INFLORESCENCES (Figure 9.38)

Secondary inflorescences are defined as aggregates of unit inflorescences (also called primary or partial inflorescences); each **unit inflorescence** is a subunit of the secondary inflorescence that resembles an inflorescence type, per se. Examples of secondary inflorescences are a panicle of spikelets, a corymb of heads, or a raceme of spikes.

Two specific types of secondary inflorescences are the thyrse and verticillaster. A **thyrse** is essentially a raceme of cymes, in which the main axis is indeterminate but the opposite, lateral,

unit inflorescences are pedicellate cymes, typically either simple dichasia, compound dichasia, or compound cymes, occasionally monochasial cymes. A **verticillaster** is essentially a spike of opposite cymes, similar to a thyrse in having an indeterminate main axis but differing in that the lateral cymes have very reduced to absent internodal axes and pedicels, giving a congested appearance. Verticillasters are found in several members of the Lamiaceae, the mint family. A **compound umbel** is another secondary inflorescence in which the peduncle bears secondary axes called **rays** that are attached at one point and unit, simple umbels attached at the tip of the rays, as in many Apiaceae.

SPECIALIZED INFLORESCENCES (Figure 9.39)

Some inflorescences are quite specialized and often restricted to certain taxonomic groups. A **catkin** (also called an *ament*) is a unisexual, typically male spike or elongate axis that falls as a unit after flowering or fruiting, as in *Quercus*, *Salix*. A **cyathium** is an inflorescence bearing small, unisexual flowers and subtended by an involucre (frequently with petaloid glands), the entire inflorescence resembling a single flower, as in *Euphorbia* and relatives. A **head** or **capitulum** is a determinate or indeterminate, crowded group of sessile or subsessile flowers on a compound receptacle, often subtended by an involucre. Heads are typical of the Asteraceae and some other groups. (Note that some inflorescences resemble a head but lack a compound receptacle; these can be termed *head-like*.) A **hypanthodium** is an inflorescence bearing numerous flowers on the inside of a convex or involuted compound receptacle, as in *Ficus*. A **pseudanthium** is a unit that appears as and may function like a single flower, but that typically consists of two or more flowers fused or grouped together. A **spadix** is a spike with a thickened or fleshy central axis, typically with congested flowers and usually subtended by a spathe, as in the Araceae. A **spikelet** literally means a small spike and refers to the basic inflorescence unit in the Cyperaceae, the sedges, and Poaceae, the grasses. Sedge spikelets are like a small spike, with sessile (reduced) flowers on an axis (**rachilla**), each flower subtended by a bract (also called a *scale*). A grass spikelet consists of an axis (**rachilla**), typically bearing two basal bracts (*glumes*) and one or more short lateral branch units called **florets**, each of which bears two bracts (*lemma* and *palea*) that subtend a terminal, reduced flower. (See family treatments of Cyperaceae and Poaceae in Chapter 7.)

FRUITS

Fruits are the mature ovaries or pistils of flowering plants plus any associated accessory parts. **Accessory parts** are organs attached to a fruit but not derived directly from the ovary or ovaries, including the bracts, axes, receptacle,

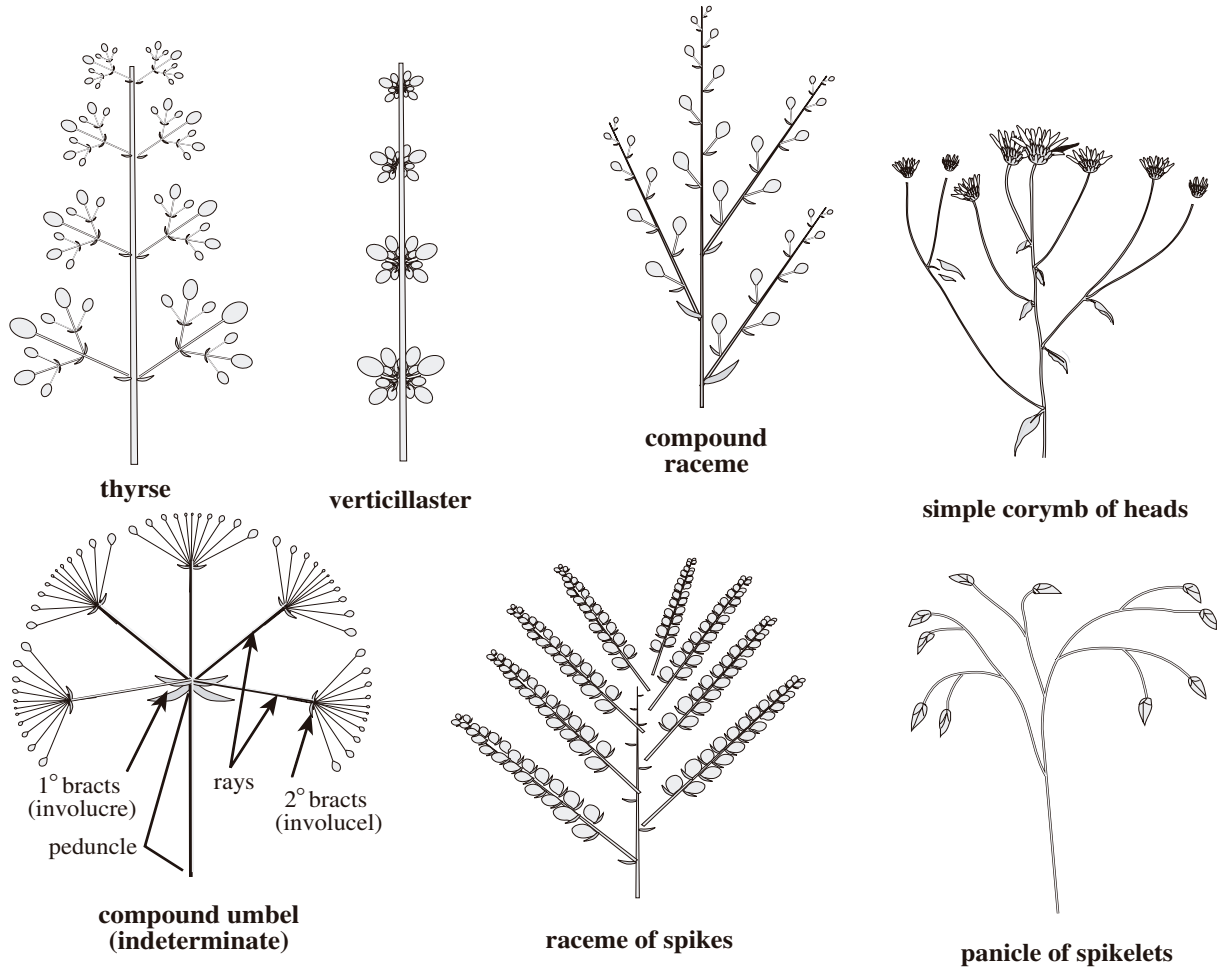


FIGURE 9.38 Secondary inflorescences.

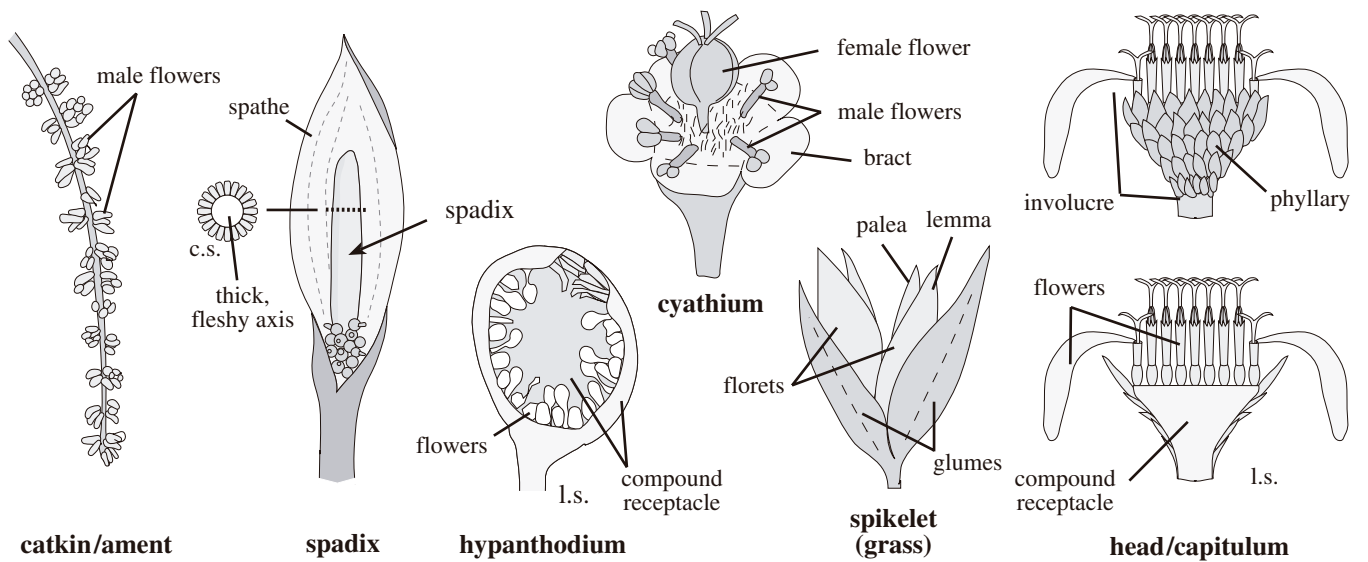


FIGURE 9.39 Specialized inflorescence types.

compound receptacle (in multiple fruits), hypanthium, or perianth. The term **pericarp** (rind, in the vernacular) is used for the fruit wall, derived from the mature ovary wall. The pericarp is sometimes divisible into layers: endocarp, mesocarp, and exocarp (see fleshy fruit types, discussed later).

Fruit types are based first on fruit development. The three major fruit developments are **simple** (derived from a single pistil of one flower), **aggregate** (derived from multiple pistils of a single flower, thus having an apocarpous gynoecium), or **multiple** (derived from many coalescent flowers; see later discussion). In aggregate or multiple fruits, the component derived from an individual pistil is called a **unit fruit**. The term **infructescence** may be used to denote a mature inflorescence in fruit.

As mentioned in Chapter 6, the evolution of fruits was correlated with the evolution of carpels and is a significant adaptation for seed dispersal in the angiosperms.

SIMPLE FRUIT TYPES (Figures 9.40, 9.41)

The simple fruit type, as well as unit fruit types of aggregate and multiple fruits, are classified based on a number of criteria, including (1) whether fleshy (succulent) or dry *at maturity*; (2) whether **indehiscent** (not splitting open at maturity) or **dehiscent** (splitting open along definite pores, slits, or sutures); (3) if dehiscent, the type (e.g., location, shape, and direction) of dehiscence; (4) carpel and locule number, including presence of septa; (5) seed/ovule number; (6) placentation; (7) structure of the pericarp wall; and (8) ovary position.

One class of simple fruits are those that are dry and indehiscent at maturity (Figure 9.40). An **achene** is a one-seeded, dry, indehiscent fruit with seed attached to the pericarp at one point only, such as the unit fruits of sunflowers. An **anthocarp** is a generally simple, dry, indehiscent fruit in which one or more flower parts function as accessory tissues, as in *Pontederia*, in which an accrescent perianth surrounds and fuses to the achene. A **grain** or **caryopsis** is a one-seeded, dry, indehiscent fruit with the seed coat adnate to pericarp

wall; grains are the fruit type of all Poaceae (grasses). (The embryo of grain crops is known as *germ*, as in wheat germ; the pericarp and seed coat together are the *bran*.) A **nut** is a one-seeded, dry indehiscent fruit with a hard pericarp, usually derived from a one-loculed ovary. (Nuts and achenes may intergrade; the terms are sometimes used interchangeably.) A **nutlet** is a small nutlike fruit; for example, the mericarps (see schizocarp) of the Boraginaceae and Lamiaceae are termed nutlets. A **samara** is a winged, dry, usually indehiscent fruit, as in *Acer* (maple) and *Ulmus* (elm). A **tryma** is a nut surrounded by an involucre that dehisces at maturity, such as in *Carya* (pecan). Finally, a **utricle** is a small, bladderly or inflated, one-seeded, dry fruit; utricles are essentially achenes in which the pericarp is significantly larger than the mature seed, as in *Atriplex* (salt bush).

Other simple fruits are dry and dehiscent at maturity (Figure 9.41). Most dry, dehiscent fruits open by means of a valve, pore, or mericarp (see later discussion). However, some, of various fruit types, are **explosively dehiscent**, i.e., will open with force (by various mechanisms), functioning to eject the seeds.

A general type of dry, dehiscent fruit is the capsule. **Capsules** are generally dry (rarely fleshy), dehiscent fruits derived from compound (multicarpelled) ovaries. Four types of capsules can be recognized based on the type or location of dehiscence. **Loculicidal capsules** have longitudinal lines of dehiscence radially aligned with the locules (or between the placentae, if septa are absent). **Septicidal capsules** have longitudinal lines of dehiscence radially aligned with the ovary septa (or with the placentae, if septa are absent). Both loculicidal and septicidal capsules split into **valves**, a portion of the pericarp wall that splits off, but does not enclose the seed(s); valves may remain attached to the fruit or may fall off, depending on the taxon. A **circumscissile capsule** (also called a **pyxis** or **pyxide**) has a transverse (as opposed to longitudinal) line of dehiscence, typically forming a terminal lid or operculum, as in *Plantago*. A **septifragal** or **valvular**

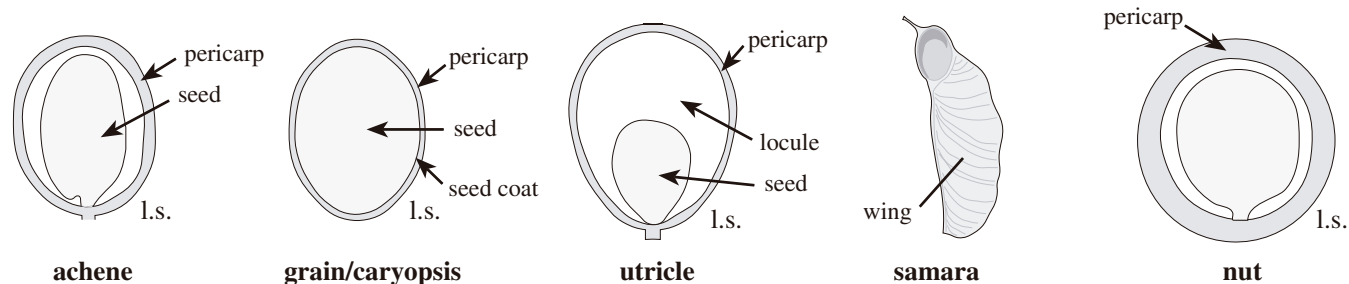


FIGURE 9.40 Fruits: simple, dry, and indehiscent fruit types. (l.s. = longitudinal section)

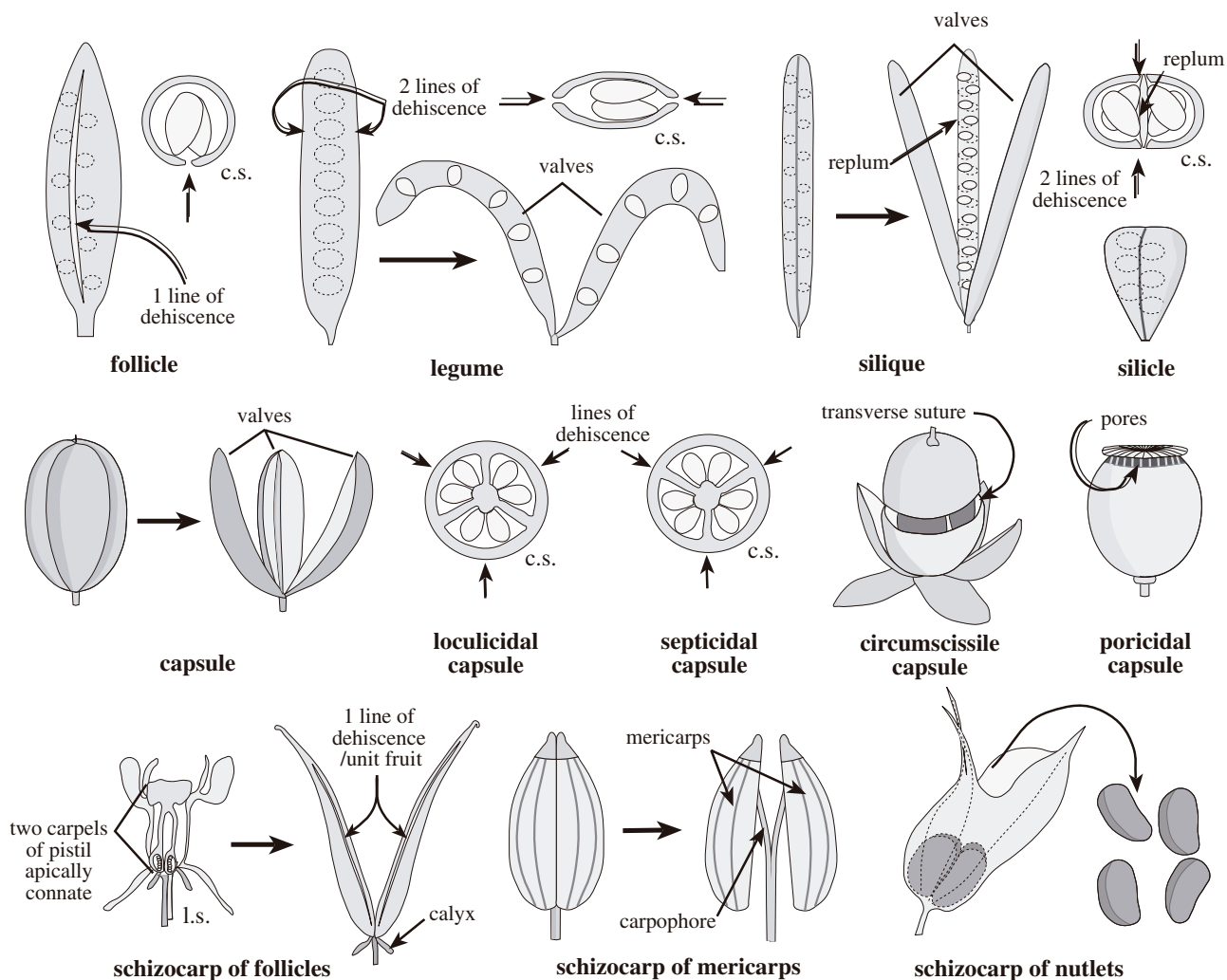


FIGURE 9.41 Fruits: simple, dry, and dehiscent fruit types. (c.s. = cross-section; l.s. = longitudinal section)

capsule is one in which the valves break off from the septa, as in *Ipomoea*, morning glory. **Poricidal capsules** have dehiscence occurring by means of pores, as in *Papaver*, poppy. Other capsules can be defined by the location of dehiscence, such as **acrocidal capsules**, dehiscing by means of apical slits, or **basicidal capsules**, dehiscing by means of basal slits, as in *Aristolochia* spp.

Some other dry, dehiscent fruit types are really just specialized capsulelike structures. A **follicle** is a dry, dehiscent fruit derived from one carpel that splits along one suture, such as in the unit fruits of *Magnolia*. A **legume** is a dry, dehiscent fruit derived from one carpel that splits along two longitudinal sutures; legumes are the diagnostic fruit type of the Fabaceae, the legume family. Some legumes retain the vestige of the two, longitudinal sutures, but have become secondarily modified,

such as **loment**s, which split transversely into one-seeded segments, and **indehiscent legumes**, which don't split open at all (e.g., peanut). **Silicles** and **siliques** are dry, dehiscent fruits derived from a two-carpeled ovary that dehisces along two sutures but that has a persistent partition, the **replum** (the mature septum, generally with attached seeds). The two fruit types differ is that a silicle is about as broad or broader than long, a silique is longer than broad; both are characteristic fruit types of the Brassicaceae, the mustard family.

Finally, a **schizocarp** is a dry, dehiscent fruit type derived from a two or more loculed compound ovary in which the locules separate at maturity. The individual unit fruits containing each locule can be defined based on other simple fruit types. For example, a **schizocarp of follicles** is a fruit in which the (generally two) carpels of a pistil split at maturity,

each carpel developing into a unit follicle, as in *Asclepias*, milkweed. A **schizocarp of mericarps** is one in which the carpels of a single ovary split during fruit maturation, each carpel developing into a unit mericarp, as in the Apiaceae. **Mericarps** are portions of the fruit that separate from the ovary as a distinct unit completely enclosing the seed(s); in the Apiaceae the two mericarps are typically attached to one another via a stalklike structure called the **carpopore**. Lastly, a **schizocarp of nutlets** is distinct in that a single ovary becomes lobed during development, the lobes developing at maturity into nutlets, which split off. (Note that nutlets here may be viewed as specialized types of mericarps.) Schizocarpic nutlets are typical of the Boraginaceae and most Lamiaceae, which have gynobasic styles attached between the ovary lobes.

Another class of simple fruits includes those that, at maturity, are fleshy or succulent (also termed baccate or carnosae; see Texture) (Figure 9.42). Fleshy fruits are general adaptations for seed dispersal by animals, the succulent pericarp being the reward (with at least some seeds either falling out or passing through the animal's gut unharmed). Fleshy fruits are generally indehiscent, but may rarely be dehiscent, as in some *Yucca* spp. The pericarp of some fleshy fruits may be divided into layers. These pericarp wall layers, if present, are termed the **endocarp** (the innermost wall layer), **mesocarp** (the middle wall layer), and **exocarp** (the outermost wall layer); if only two layers are evident, the terms *endocarp* and *exocarp* alone are used. A **berry** is the general, unspecialized term for a fruit with a succulent pericarp, as in *Vitis*, grape. A **drupe** is a fruit with a hard, stony endocarp and a fleshy mesocarp, as in *Prunus* (peach, plum, cherry, etc.). The term **pyrene** can be used either for a fleshy fruit in which each of two or more seeds is enclosed by a usually bony-textured endocarp, or **pyrene** can refer to the seed covered by a hard endocarp unit itself, regardless of the number. A **hesperidium** is a septate fleshy fruit with a thick-skinned, leathery outer pericarp wall and fleshy modified trichomes (juice sacs) arising from the inner walls, as in *Citrus*

(orange, lemon, grapefruit, etc.). A **pepo** is a nonseptate fleshy fruit with parietal placentation and a leathery exocarp derived from an inferior ovary, the fruit type of the Cucurbitaceae. A **pome** is a fleshy fruit with a cartilaginous endocarp derived from an inferior ovary, with the bulk of the fleshy tissue derived from the outer, adnate hypanthial tissue, as in *Malus* (apple) and *Pyrus* (pear). Finally, a **pseudodrupe** is a nut surrounded by a fleshy, indehiscent involucre, as in *Juglans* (walnut); thus, pseudodrupes have accessory tissue serving as the fleshy component.

AGGREGATE FRUIT TYPES (Figure 9.43)

An **aggregate fruit** is one derived from two or more pistils (ovaries) of one flower. In determining the aggregate fruit type, one first identifies the **unit fruit** that corresponds to a single pistil. The aggregate fruit type is then indicated either as aggregate fruit of the particular unit fruits or by adding the suffix *-acetum* to the unit fruit term.

An **achenecetum** is an aggregate fruit of achenes. A common example is *Fragaria*, strawberry, in which the achenes are on the surface of accessory tissue, an enlarged, fleshy receptacle. A **drupecetum** is an aggregate fruit of drupes, as in *Rubus*, raspberry or blackberry. A **follicetum** is an aggregate fruit of follicles, as occurs in *Magnolia*. A **syncarp** is an aggregate fruit, typically of berries, in which the unit fruits fuse together, as in *Annona*.

MULTIPLE FRUIT TYPES (Figure 9.44)

A **multiple fruit** is one derived from two or more flowers that coalesce. In determining the multiple fruit type, one may also identify the unit fruit corresponding to a single pistil of a single flower; the fruit type may be indicated as a multiple fruit of the particular unit fruit present.

Some specialized multiple fruit types are as follows. A **bur** is a multiple fruit of achenes or grains surrounded by a prickly involucre, such as in *Cenchrus*, sandbur (Poaceae), or

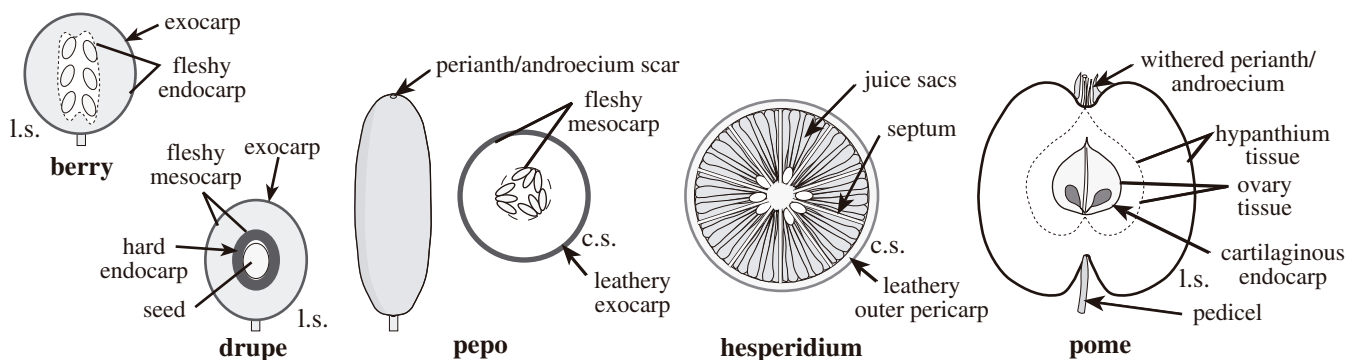


FIGURE 9.42 Fruits: simple, fleshy fruit types. (c.s. = cross-section; l.s. = longitudinal section)

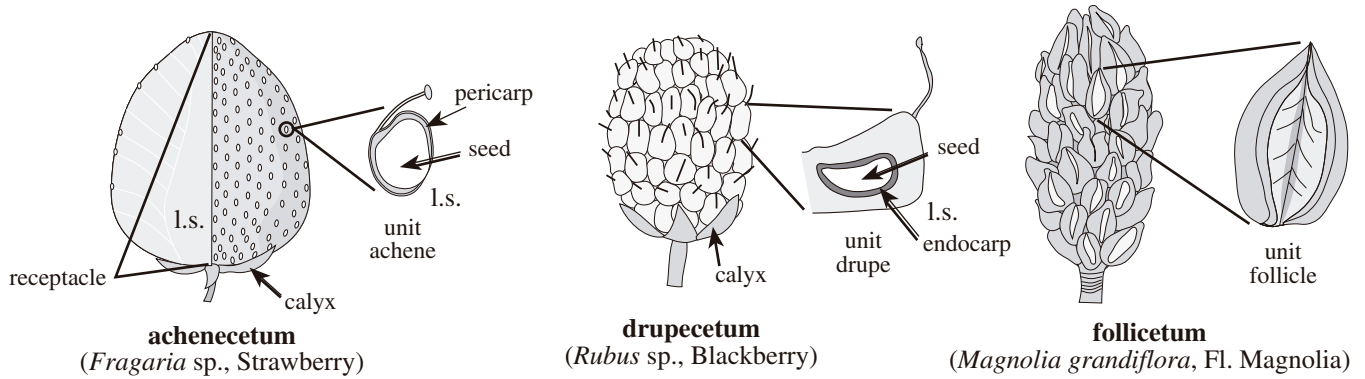


FIGURE 9.43 Fruits: aggregate fruit types. (l.s. = longitudinal section)

Xanthium, cocklebur (Asteraceae). A **soros** is a multiple fruit in which the unit fruits are fleshy berries and are laterally fused along a central axis, as in *Ananas*, pineapple. A **syconium** is a multiple fruit in which the unit fruits are small achenes covering the surface of a fleshy, inverted compound receptacle (derived from a hypanthodium), as in *Ficus*, fig.

SEEDS (Figure 9.45)

Aspects of seed morphology can be important systematic characters used in plant classification and identification. Some valuable aspects of seed morphology are size and shape, as well as the color and surface features of the **seed coat** or **testa**, the outer protective covering of seed derived

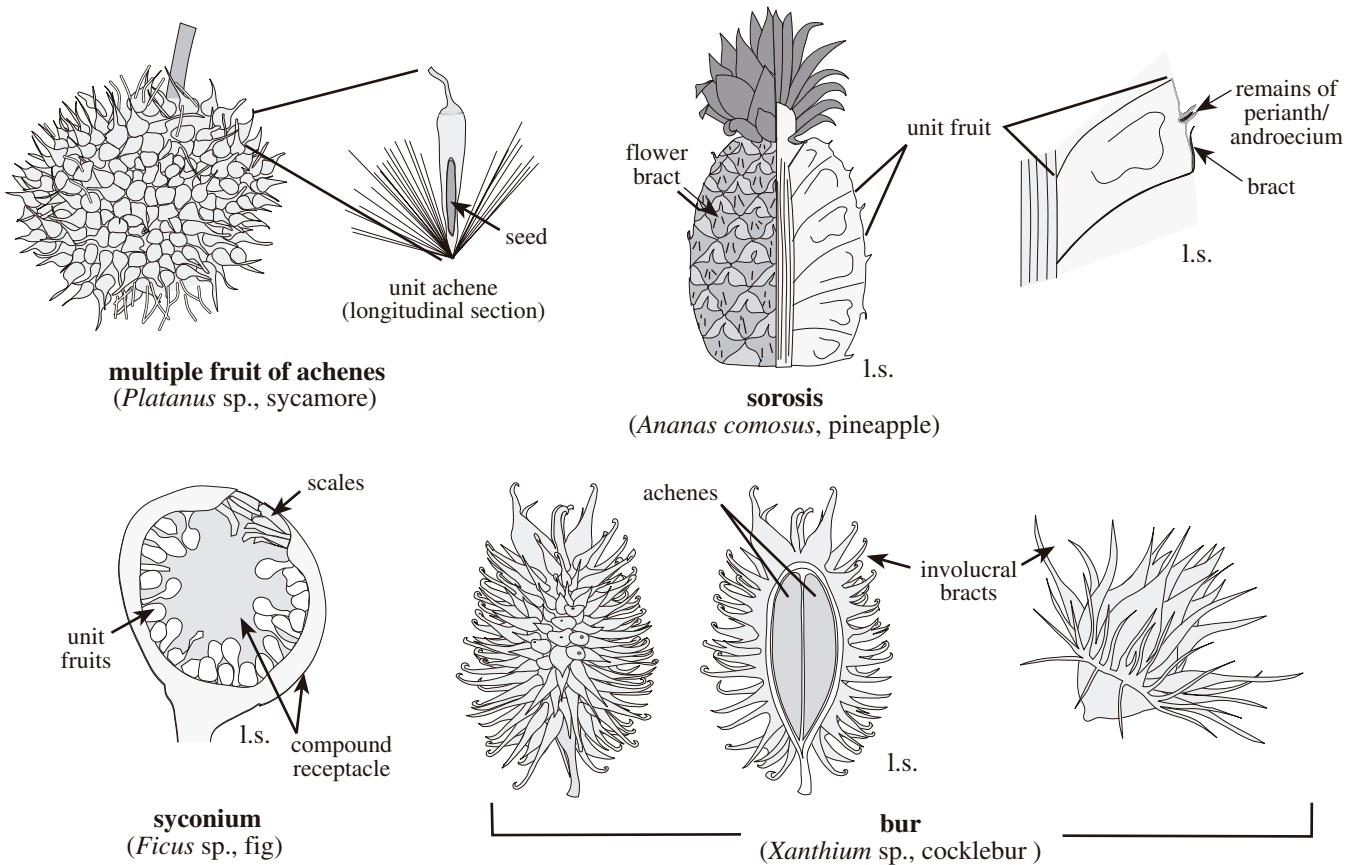


FIGURE 9.44 Fruits: multiple fruit types. (l.s. = longitudinal section)

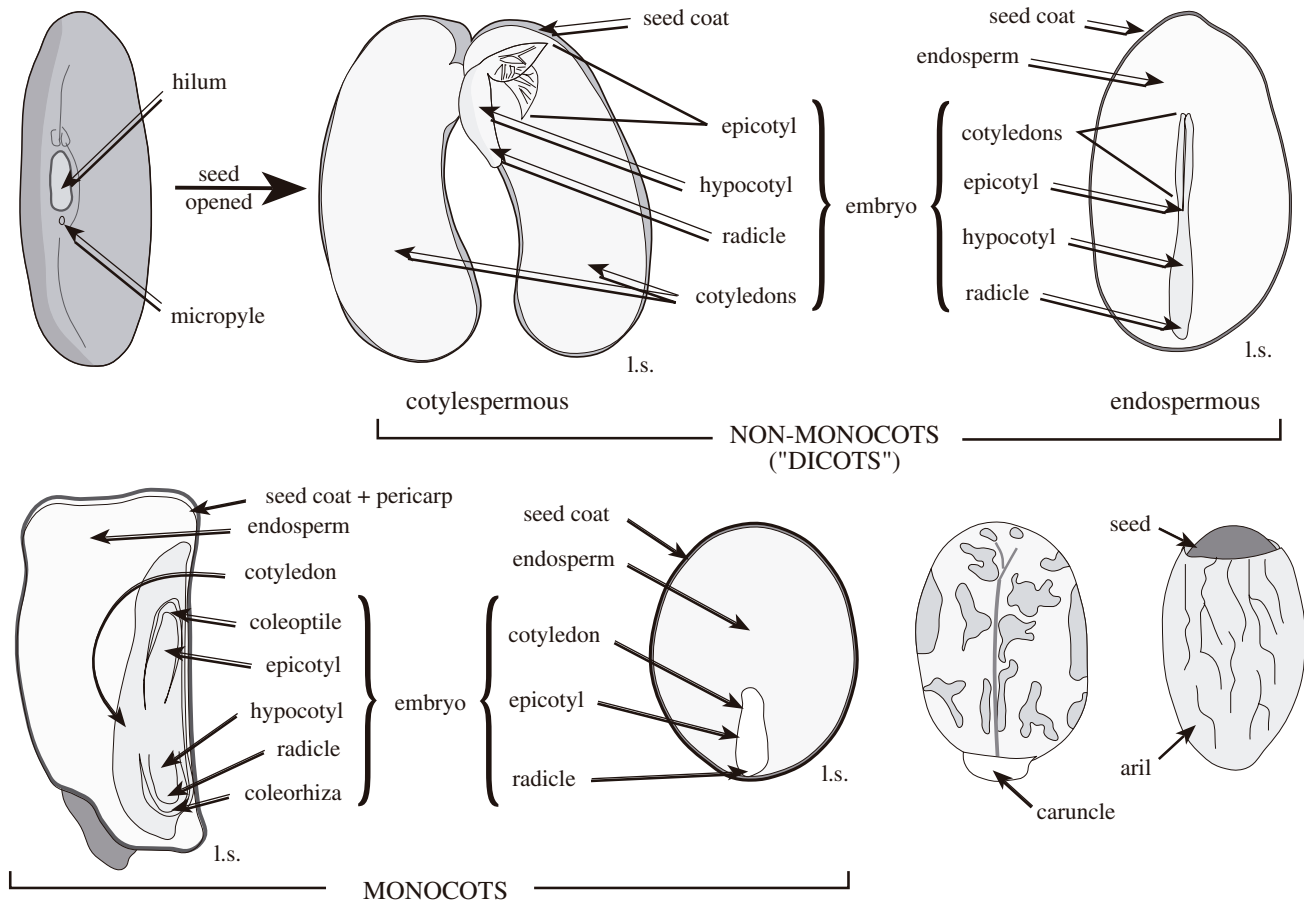


FIGURE 9.45 Seed parts, endosperm types. (l.s. = longitudinal section)

from the integument(s). A seed coat that is fleshy at maturity may be termed a **sarcotesta** (although this may be confused with an **aril**, which is separate from the integuments; see later discussion). Also important are the shape, size, and color of the **hilum**, the scar of attachment of the funiculus on the seed coat, and of the **raphe**, a ridge on the seed coat formed from an adnate funiculus. Some seeds have an **aril** (adj. *arillate*), a fleshy outgrowth of the funiculus, raphe, or integuments (but separate from the integuments) that generally functions in animal seed dispersal. Arils may be characteristic of certain groups, such as the Sapindaceae. Similar to the aril is a **caruncle**, a fleshy outgrowth at the base of the seed; caruncles also function in animal seed dispersal, such as the carunculate seeds of *Viola*, violets, with regard to seed dispersal by ants.

Specific details of the **embryo**, the immature sporophyte, can be studied. These include aspects of the **epicotyl** (the immature shoot), **radicle** (the immature root; not to be confused with a

radical position; see later discussion), **hypocotyl** (the transition region between the root and epicotyl), and **cotyledon(s)** (the first leaf/leaves of the embryo, often functioning in storage of food reserves). Some members of the Poaceae, the grass family, have the epicotyl surrounded by a protective sheath known as the **coleoptile**, and the radicle surrounded by a protective sheath known as the **coleorhiza**. Cotyledon aestivation (or *ptyxis*) can be a valuable systematic feature.

SEED ENDOSPERM TYPE

All angiosperms form endosperm, the food reserve tissue derived from fusion of sperm with the polar nuclei of the female gametophyte. The typical angiosperm seed is **albuminous** or **endospermous**, having endosperm as the food reserve in mature seeds. In some angiosperms endosperm develops, but very little to none is deposited in mature seeds, a feature termed **exalbuminous** or **nonendospermous**, as in orchid seeds. Finally, some flowering plants are **cotyledonous**,

in which the main food reserve is stored in the cotyledons. Cotyledonous seeds are typical of beans and peas.

SEED GERMINATION TYPE

Seed germination type requires observation of young seedlings during germination and describes positioning of the cotyledons. **Hypogeous** [**cryptocotylar**] refers to a type in which the cotyledon(s) remain in the ground during germination. **Epigeous** [**phanerocotylar**] has cotyledon(s) elevated above the ground during germination.

GENERAL TERMINOLOGY

Many plant morphological terms can apply to a number of different plant organs (or even to features of other types of organisms). These general terms are defined below.

COLOR

Color is a measure of the wavelengths of light reflected from or transmitted through an object. When describing color, that of each component organ or part should be precisely designated. For example, instead of just stating flowers yellow, describe as corolla and filaments yellow, anthers maroon, pollen white, ovary green. Color itself may be defined in a very precise way, utilizing components of hue, value, and chroma. For precise designation of color, a color chart is invaluable (see Tucker et al. 1991).

Color pattern is a measure of the distribution of colors on an object. Common color pattern terms are **maculate**, spotted, with small spots on a more or less uniform background; **pellucid**, having translucent spots or patches; and **variegated**, with two or more colors occurring in various irregular patterns, generally used for leaves.

SIZE

Of course, measuring the size of plant organs and parts is important in description and identification. Generally, size of parts refers to linear measurements, as in leaf length or corolla width. Metric units should be used throughout.

NUMBER

Number refers to a simple count of parts. Of course, number of parts can be very valuable information in systematic studies. With whorled structures, a distinction is made between cycly and merosity.

Cycly is the number of cycles or whorls of parts. It may simply be designated as a number, or terms may be used such as **monocyclic** or **uniseriate**, with a single whorl of parts; **dicyclic** or **biseriate**, with two whorls of parts; **tricyclic** or

triseriate, with three whorls of parts; etc. Cycly is most commonly used for parts of the perianth or androecium (see earlier discussion).

Merosity is the number of parts per whorl or cycle. Merosity may also be designated as a simple number, or the terms **bimerous**, a whorl with two members, **trimerous**, a whorl with three members, **tetramerous**, a whorl with four members, **pentamerous**, a whorl with five members, etc., may be used. Two general merosity terms are **isomerous**, having the same number of members in different whorls, and **anisomerous**, having a different number of members in different whorls. Merosity is most commonly designated for floral parts: the calyx, corolla, androecium, and gynoecium (equivalent to carpel number in a syncarpous gynoecium).

TEXTURE

Texture is the internal structural consistency of an object; some texture terms also take color into account. Texture is often described for leaves but can be used for any plant part, such as bracts or flower parts. Texture may be correlated with plant habitat and can be representative of the amount of water storage tissue (as in leaf or stem succulent plants), fibers, vascular bundles, lignin, suberin, or other internal anatomical features of a plant organ. Common texture terms include **cartilaginous**, with the texture of cartilage; hard and tough but flexible, usually whitish; **chartaceous**, opaque and of the texture of writing paper; **coriaceous**, thick and leathery, but somewhat flexible; **herbaceous**, having a soft or slightly succulent texture; **indurate**, hardened and inflexible; **membranous**, thin and somewhat translucent, membranous; **mesophytic**, having an intermediate texture, between coriaceous and membranous (typical of many, common leaves); **ruminant**, unevenly textured, coarsely wrinkled, looking as if chewed (e.g., the endosperm of the Annonaceae); **scariosus**, thin and appearing dry, usually whitish or brownish; **succulent** [*baccate* or *carnose*], fleshy and juicy; and **woody**, having a hard, woodlike texture.

FUSION

Fusion refers to the apparent joining (or lack of joining) of two or more discrete plant organs or parts. Objects that are fused may have developed separately and then come into contact and joined later. This process, known as *congenital fusion*, may happen, e.g., when anthers are fused. However, it is more likely that apparently fused objects actually developed from the same meristematic tissue. In this process, the primordia that would normally grow into separate objects are elevated by the more basal, actively growing tissue, a typical example being the fusion of corolla and stamens (epipetalous). (See also Perianth Fusion, Stamen Fusion, and Gynoecial Fusion.)

Fusion terms are distinguished as to whether fusion is between like or unlike parts. **Connate** is integral fusion of *like* parts, such that the parts are not easily separable. **Adnate** is a similar integral fusion of *unlike* parts. Thus, saying stamens are connate means that they are fused to one another (e.g., monadelphous, diadelphous, syngenesious, etc.), whereas stamens adnate means they are fused to something else (e.g., to the corolla). Two similar terms to represent partial or incomplete fusion are **coherent**, with *like* parts joined but only superficially and easily separable; and **adherent**, with *unlike* parts joined, but likewise only superficially and easily separable.

Some terms designate lack of fusion. **Distinct** means with *like* parts unfused and separate. **Free** is with *unlike* parts unfused and separate. Lastly, **contiguous** means with parts touching but not connate, adnate, coherent, or adherent. Contiguous plant parts may appear fused, but are only in close contact.

SHAPE

Shape terms may be used for stems, leaves, leaflets or other leaf parts, bracts, sepals, petals, stamens, pistils, trichomes, or other plant parts. Shape is an important feature in plant description and identification. Shape may be classified as solid (three-dimensional) versus plane (two-dimensional). The latter, plane shape, may be divided into overall plane shape, base, margin, apex shape, apical process, and division.

SHAPE: SOLID (THREE-DIMENSIONAL) (Figure 9.46)

Several specific three-dimensional shapes are widely used. **Capitate** is head-shaped, spherical with a short basal stalk. The term for spherical is **globose**; that for half-sphere-shaped is **hemispheric**. An ellipsoid shape with the long axis parallel to the point of attachment is termed **prolate**; one extended perpendicular to the point of attachment is **oblate**. **Clavate** means club-shaped, cylindrical with a gradually tapering, thickened and rounded end. **Discoïd** is disk-shaped, and **fusiform** is spindle-shaped, narrowly ellipsoid with two attenuate ends. **Filiform** means threadlike or filamentous, being

long, thin, and typically flexuous. **Fistulose** or **fistular** means cylindrical and hollow within. **Lenticular** means lens-shaped, disk-shaped with two convex sides. **Ligulate** is tongue-shaped; flattened and somewhat oblong in shape, as in the ligulate (ray) corollas of some Asteraceae. **Pilate** means with a long cylindrical stalk terminating in a globose or ellipsoid apical thickening, as in pilate-glandular trichomes (see later discussion). **Terete** is the general plant term meaning cylindrical. **Turbinate** means turban or top-shaped, as in turbinate heads or compound receptacles of the Asteraceae.

In addition to these specific terms, other three-dimensional shape terms can be derived from those for two-dimensional shapes (next) by adding the suffix -oid; as in ellipsoid, oblongoid, ovoid.

SHAPE: PLANE (TWO-DIMENSIONAL) (Figure 9.47)

Overall plane or two-dimensional shape has been standardized (see Systematics Association Committee for Descriptive Terminology, 1962). These shape terms are based, in part, on the ratio of the length to the width of the shape outline, the common length width ratios being >12:1, 12:1 6:1, 6:1 3:1, 2:1 3:2, approximately 6:5, and approximately 1:1. (Note that the bases, apices, and details of the margin can vary in these general planar shape terms; see later discussion.)

Shapes in which the margins (sides) of the object are straight and approximately parallel are **acicular**, needle-like with length : width ratio greater than 12:1; **ensiform**, sword-shaped, with length : width ratio greater than 12:1, e.g., leaves of *Iris* spp.; **strap-shaped**, flat, not needle-like but with length : width ratio greater than 12:1; **linear**, length : width ratio between 12:1 and 6:1; **narrowly oblong**, length : width ratio between 6:1 and 3:1; and **oblong**, length : width ratio between 2:1 and 3:2.

Shapes in which the margins are symmetrically curved, with the widest point near the midpoint of the object, are **narrowly elliptic**, length : width ratio between 6:1 and 3:1; **elliptic**, length : width ratio between 2:1 and 3:2; **widely elliptic**, length : width ratio approximately 6:5; and **orbicular** (*circular*), length : width ratio approximately 1:1.

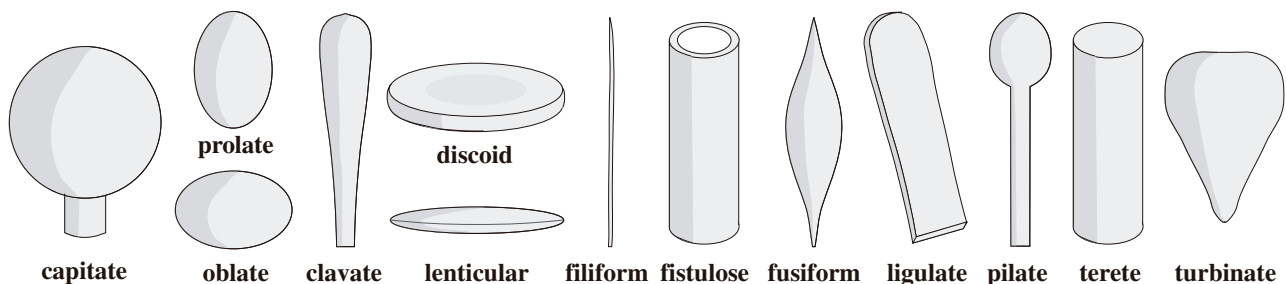


FIGURE 9.46 Solid (three-dimensional) shapes.

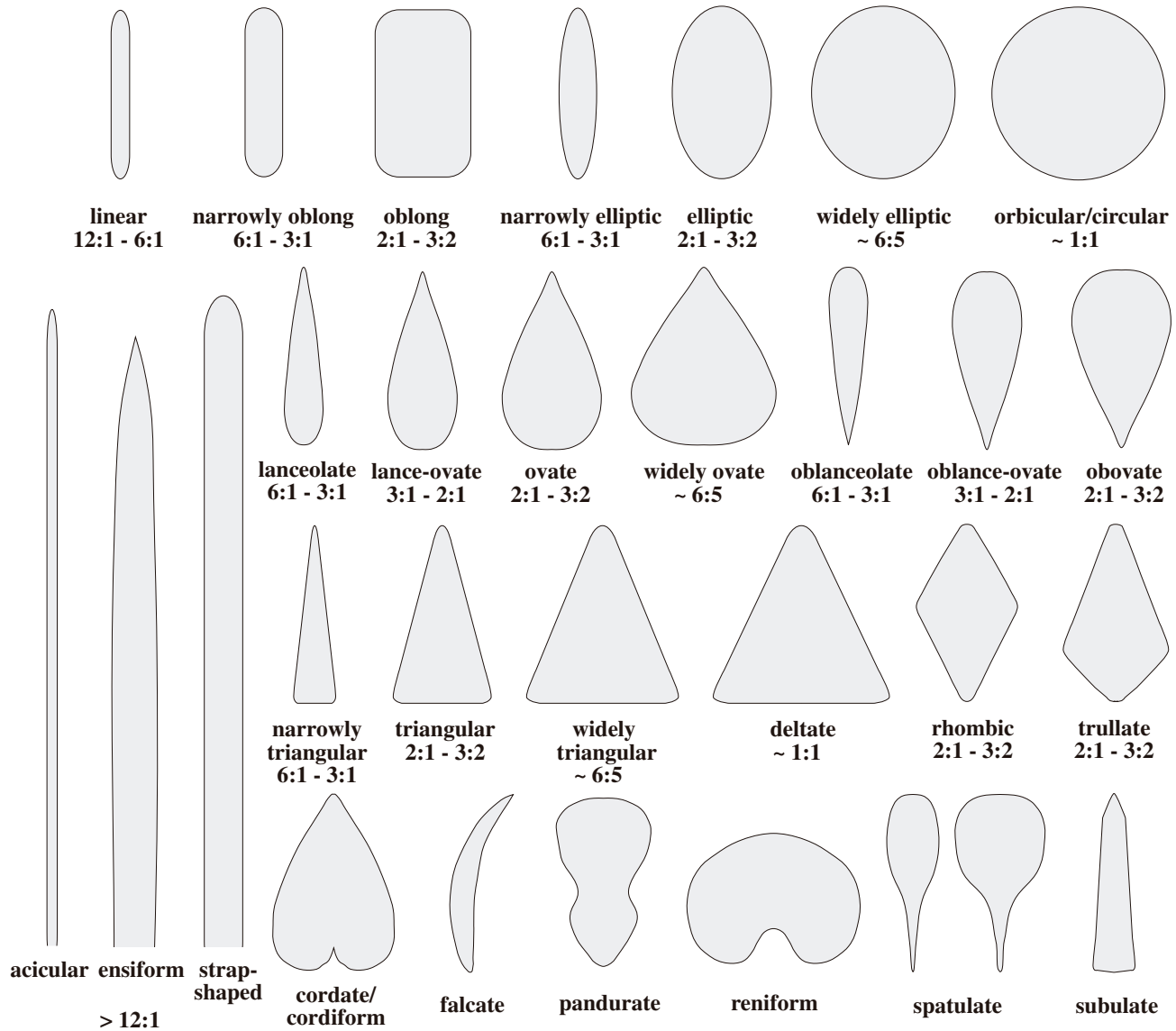


FIGURE 9.47 Planar (two-dimensional) shapes. Note that bottoms of drawings are points of attachment.

Shapes in which the margins are curved, with the widest point near the base, are **lanceolate**, length : width ratio between 6:1 and 3:1; **lance-ovate**, length : width ratio between 3:1 and 2:1; **ovate**, length : width ratio between 2:1 and 3:2; **widely ovate**, length : width ratio approximately 6:5; and **very widely ovate**, length : width ratio close to 1.

Shapes in which the margins are curved, with the widest point near the apex, are **oblanceolate**, length : width ratio between 6:1 and 3:1; **oblance-ovate**, length : width ratio between 3:1 and 2:1; **obovate**, length : width ratio between 2:1 and 3:2; **widely obovate**, length : width ratio approximately 6:5; and **very widely obovate**, length : width ratio close to 1.

Three-sided shapes, in which the sides are approximately straight, are **narrowly triangular**, length : width ratio between 6:1 and 3:1; **triangular**, length : width ratio between 3:2 and 2:1; **widely triangular**, length : width ratio approximately 6:5; and **deltate**, length : width ratio approximately 1.

Four-sided, parallelogram-like shapes are **rhombic**, widest near middle, length : width ratio between 2:1 and 3:2; and **trullate**, widest near base; length : width ratio between 2:1 and 3:2.

Finally, some specialized shapes are **cordate (cordiform)**, shaped like an inverted Valentine heart, approximately ovate with a cordate base (see Base); **falcate (falciform)**, scimitar-shaped, lanceolate to linear and curved to one side; **lyrate**,

pinnatifid, but with a large terminal lobe and smaller basal and lateral lobes; **pandurate**, violin-shaped, obovate with the side margins concave; **reniform**, kidney-shaped, wider than long with a rounded apex and reniform base (see Base); **spatulate**, oblong, obovate, or oblanceolate with a long attenuate base; and **subulate**, awl-shaped, approximately narrowly oblong to narrowly triangular.

BASE (Figure 9.48)

Base shapes in which the sides are incurved or are approximately straight are **attenuate**, basal margins abruptly incurved (concave), intersection angle less than 45° ; **narrowly cuneate**, basal margins approximately straight, intersection angle less than 45° ; **cuneate**, basal margins approximately straight, intersection angle $45^\circ - 90^\circ$; **obtuse**, basal margins approximately straight, intersection angle greater than 90° ; and **truncate**, basal margin cut straight across, intersection angle approximately 180° .

Base shapes in which the sides are curved are **rounded**, basal margins convex, forming a single, smooth arc; **cordate**, with two rounded, basal lobes intersecting at sharp angle, the margins above lobes smoothly rounded; and **reniform**, with two rounded, basal lobes, smoothly concave at intersection of lobes.

Bases in which there are two protruding lobes are **auriculate**, with two rounded, basal lobes, the margins above lobes concave; **hastate**, with two basal lobes, more or less pointed and oriented outwardly, approximately 90° relative to central axis; and **sagittate**, with two basal lobes, more or less pointed and oriented downward, away from the apex.

Finally, some other, specialized base shapes are **oblique**, having an asymmetrical base; **peltate**, with the petiole attached away from the margin, on the underside of the blade, as in *Tropaeolum*; and **sheathing**, having a basal, clasping leaf sheath. (Note: see also Leaf Attachment.)

APEX (Figures 9.49, 9.50)

Apex shape (Figure 9.49) refers to the shape of the apical region below the apical process or vein extension, if present (see later discussion). For a leaf or bract, this refers to the shape of the blade tissue at the apex.

An **acuminate** apex is one with the apical margins abruptly incurved (concave), the apical intersection angle $<45^\circ$. Two other apex shapes are specialized variants of acuminate: **caudate**, abruptly acuminate into a long, narrowly triangular (tail-like) apical region; and **cuspidate**, abruptly acuminate into a triangular, stiff or sharp apex.

Four apex shapes have straight, not curved, sides. A **narrowly acute** apex is one with the margins approximately straight, the intersection angle less than 45° . (Thus, narrowly acute differs from acuminate, caudate, and cuspidate in part by having straight margins.) An **acute** apex also has more or less straight margins, with the intersection angle between 45° and 90° . An **obtuse** apex shape has apical margins approximately straight, the intersection angle greater than 90° . A **truncate** apex has the apical margin cut straight across, the angle approximately 180° .

A **rounded** apex has convex apical margins, forming a single, smooth arc. An **oblique** apex has an asymmetrical

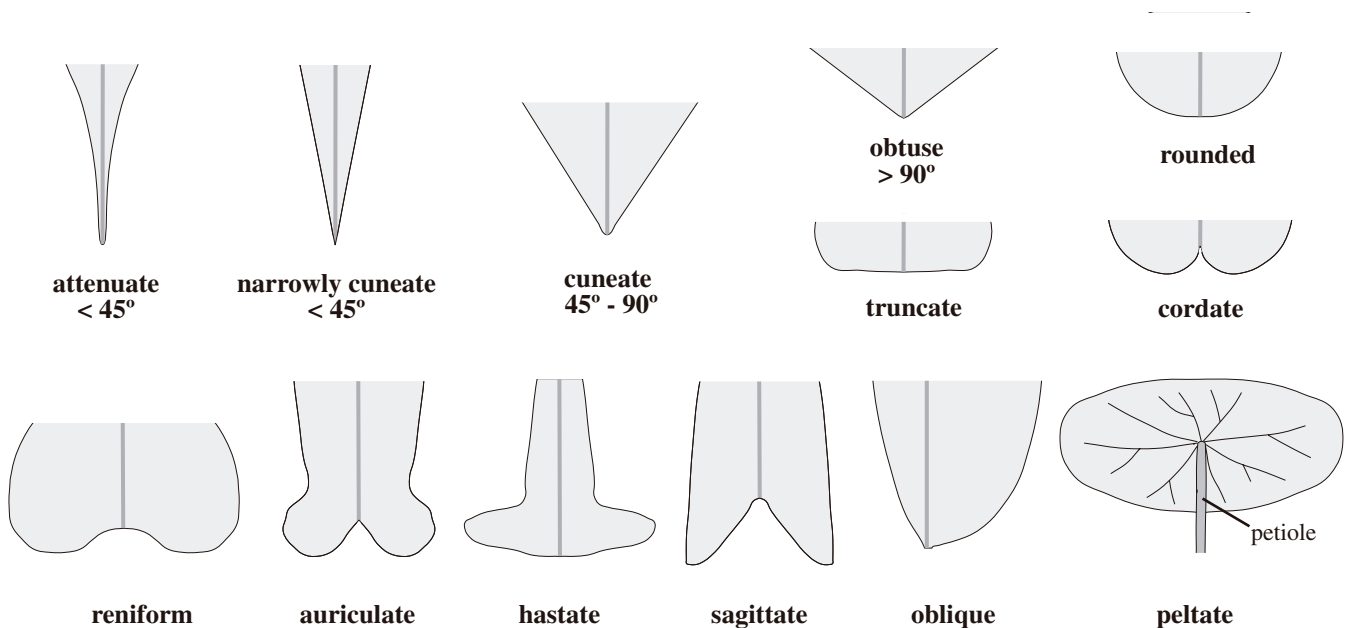


FIGURE 9.48 Bases. Note that bottoms of drawings are points of attachment.

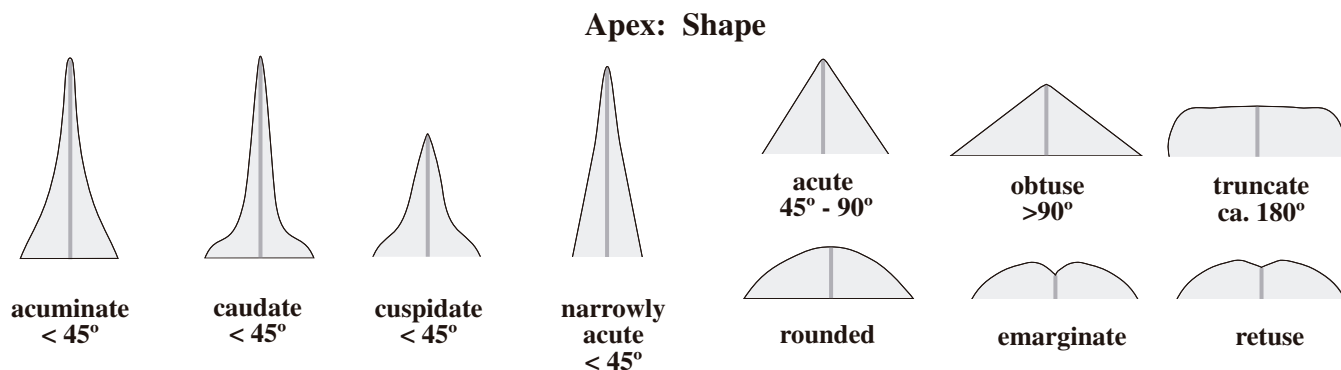


FIGURE 9.49 Apices. Note that bottoms of drawings are basal.

shape (see Base). Finally, two terms that describe an apical cleft (differing only in the depth of that cleft) are **emarginate**, having an apical incision cut $\frac{1}{16}$ - $\frac{1}{8}$ of the distance to midrib, midvein, or junction of primary veins; and **retuse**, having an apical incision cut up to $\frac{1}{16}$ of the distance to midrib, midvein, or junction of primary veins.

Apical process (Figure 9.50) generally denotes an extension of a vein (typically the midvein); thus, most of the apical process is vascular tissue. A given apical process can be associated with virtually any type of apical shape.

Common apical processes are **apiculate**, with a flexible apical process, length : width ratio $>3:1$, usually slightly curled; **aristate**, with a stiff apical process, length : width ratio $>3:1$, usually prolonged and straight; **cirrhose**, with a flexible, greatly curled apical process; **mucronate**, with a stiff, straight apical process, the length : width ratio 1:1 - 3:1; **mucronulate**, with a stiff, straight apical process, length : width ratio $\leq 1:1$; and **spinose** or **pungent**, with a sharp, stiff, spinelike apical process.

SHAPE COMBINATIONS (Figure 9.51)

The overall shape, base shape, and apex shape can be used in combination to describe a variety of two-dimensional forms. For example, Figure 9.51 shows five leaves, all with a more or less elliptic overall shape, but differing in the shape of the base and apex.

MARGIN (Figure 9.52)

Margin refers to the sides of an object, usually a leaf, bract, sepal, or petal. Many margin terms describe the presence and morphology of *teeth*, small sharp-pointed or rounded projections or lobes along the sides. Technically, teeth extend no more than $\frac{1}{8}$ of the distance to the midrib, midvein, or (in a palmately lobed leaf) junction of the primary veins; if further than $\frac{1}{8}$ of this distance, then the object is described as *lobed*, *cleft*, *parted*, or *divided* (see Division).

A margin without teeth is termed **entire**. (However, the plane may be divided; see later discussion.) A margin with teeth can be generally termed toothed, but more specific terms are preferable. Margin terms describing sawlike teeth, i.e., sharp-pointed and ascending (the lower side longer than the upper) are **serrate**, teeth cut $\frac{1}{16}$ - $\frac{1}{8}$ of the distance to midrib, midvein, or junction of primary veins; **serrulate**, diminutive of serrate, teeth cut to $\frac{1}{16}$ of the distance to midrib, midvein, or junction of primary veins; and **doubly serrate**, with large, serrate teeth having along the margin smaller, serrate teeth.

Margin terms describing sharklike teeth that point outward at right angles to the margin outline (the upper and lower sides about the same length) are **dentate**, with teeth cut $\frac{1}{16}$ - $\frac{1}{8}$ of the distance to midrib, midvein, or junction of primary veins; and **denticulate**, diminutive of dentate, cut to $\frac{1}{16}$ of the distance to midrib, midvein, or junction of primary veins.

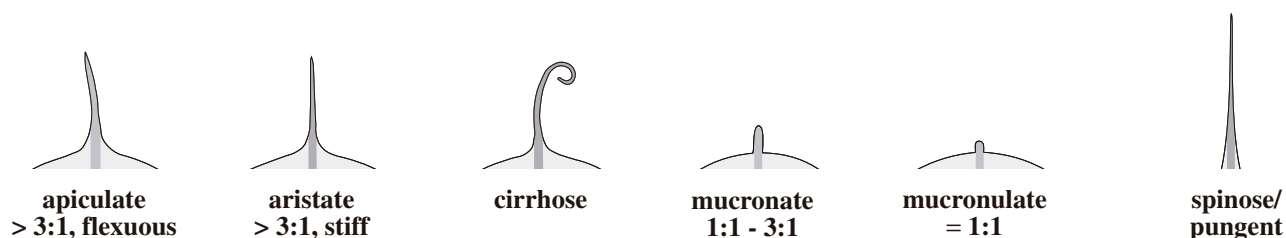


FIGURE 9.50 Apical processes.

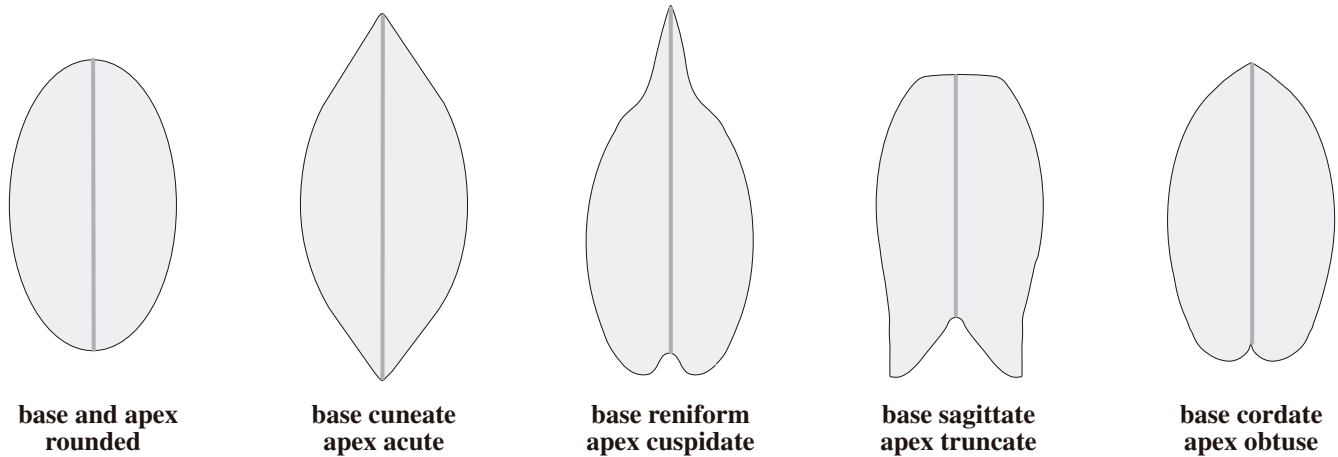


FIGURE 9.51 Shape combinations. Note that the five shapes are all elliptic, but differ in the base and apex shapes.

Margin terms describing rounded to obtuse teeth, that point outward at right angles or shallowly ascend, are **crenate**, with teeth cut $\frac{1}{16}$ – $\frac{1}{8}$ of the distance to midrib, midvein, or junction of primary veins; and **crenulate**, diminutive of crenate, teeth cut to $\frac{1}{16}$ of the distance to midrib, midvein, or junction of primary veins.

The relative size and density of teeth may also be described, with terms such as *coarsely*, to describe large and uneven teeth (e.g., coarsely crenate), *finely*, to describe relatively small, evenly spaced teeth (e.g., finely denticulate), or *sparsely*, to describe teeth that are few in number or spaced well apart (e.g., sparsely serrate).

Margins with teeth bearing sharp, stiff, spinelike processes are termed **spinose**. **Praemorse** describes a margin having a jagged, chewed appearance, as in some palms.

Terms describing margins with trichomes (plant hairs; see later discussion) are **ciliate**, with trichomes protruding from margins, and **ciliolate**, with minute trichomes protruding from margins, minutely ciliate. The term **eciliate** describes a

margin without trichomes, regardless of presence or absence of teeth. Finally, the term **filiferous** refers to margins bearing coarse, fiberlike structures (e.g., fibrovascular bundles, as in the leaf margins of some *Yucca* species).

(Terms that are often treated as features of margin, but treated here as longitudinal posture, are **involute**, with margins rolled upward, and **revolute**, with margins rolled under; see later discussion.)

DIVISION (Figure 9.53)

Division is a shape character referring to the presence and characteristics of **sinuses** (incisions), the sinuses defining **lobes** or **segments**. Division character states are sometimes treated as features of margin.

Four division terms that precisely denote the degree of division are: **lobed**, sinuses extending $\frac{1}{8}$ to $\frac{1}{4}$ of the distance to the midrib, midvein, or vein junction; **cleft**, sinuses extending $\frac{1}{4}$ to $\frac{1}{2}$ of the distance to the midrib, midvein, or vein junction; **parted**, sinuses extending $\frac{1}{2}$ to $\frac{3}{4}$ of the distance to the midrib,

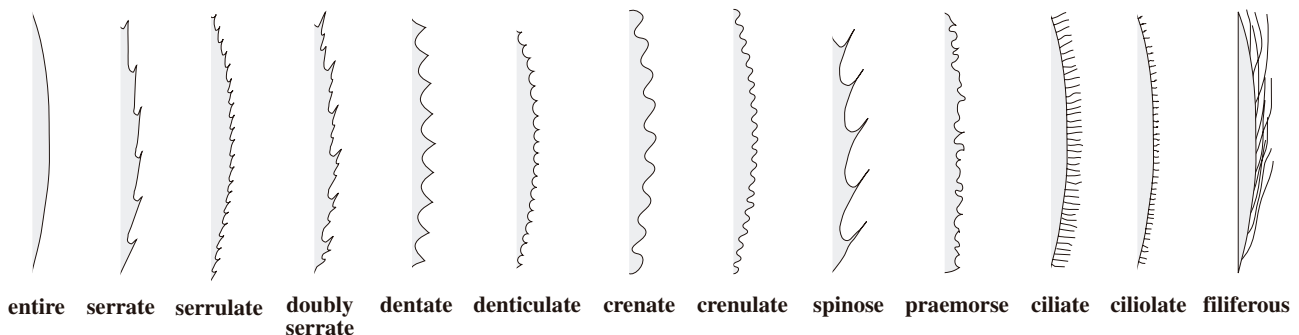


FIGURE 9.52 Margins.

midvein, or vein junction; and **divided**, sinuses extending $\frac{3}{4}$ to almost to the midrib, midvein, or vein junction.

Lobed, cleft, parted, and divided should be prefaced by terms that denote further the type of division: **pinnately** (e.g., pinnately lobed or pinnately cleft) to specify a division along a central axis (typically the midvein), and **palmately** (e.g., palmately divided) to specify a division relative to a point (typically the basal junction of major veins). (Note that the terms *lobed* and *divided* are sometimes used generally, for any extent of division; as used here, these terms refer to specific degrees of division.)

Some useful, general terms that indicate the general form, but not the extent, of division are **pinnatifid**, pinnately lobed to divided; **pinnatisect**, pinnately divided, almost into discrete

leaflets but confluent at the midrib; **bipinnatifid**, bipinnately lobed to divided; **palmatifid**, palmately lobed to divided; and **palmatisect**, palmately divided, almost into discrete leaflets but confluent at the lobe bases. **Decompound** denotes deeply divided into numerous segments such that leaflets are not clearly defined. (Note that *decompound* can also be used for a multiply compound leaf; see Leaf Type.)

Some division terms refer specifically to the shape of the sinuses and lobes. **Dissected** means divided into very fine, often indistinct segments. **Bifid** means 2-lobed to 2-divided, especially at the apex. **Incised** means the sinuses are sharp and deeply cut, usually jaggedly. **Sinuate**, in contrast, refers to sinuses being shallow and smooth, wavy in a horizontal plane (compare with *undulate*, under Longitudinal Posture).

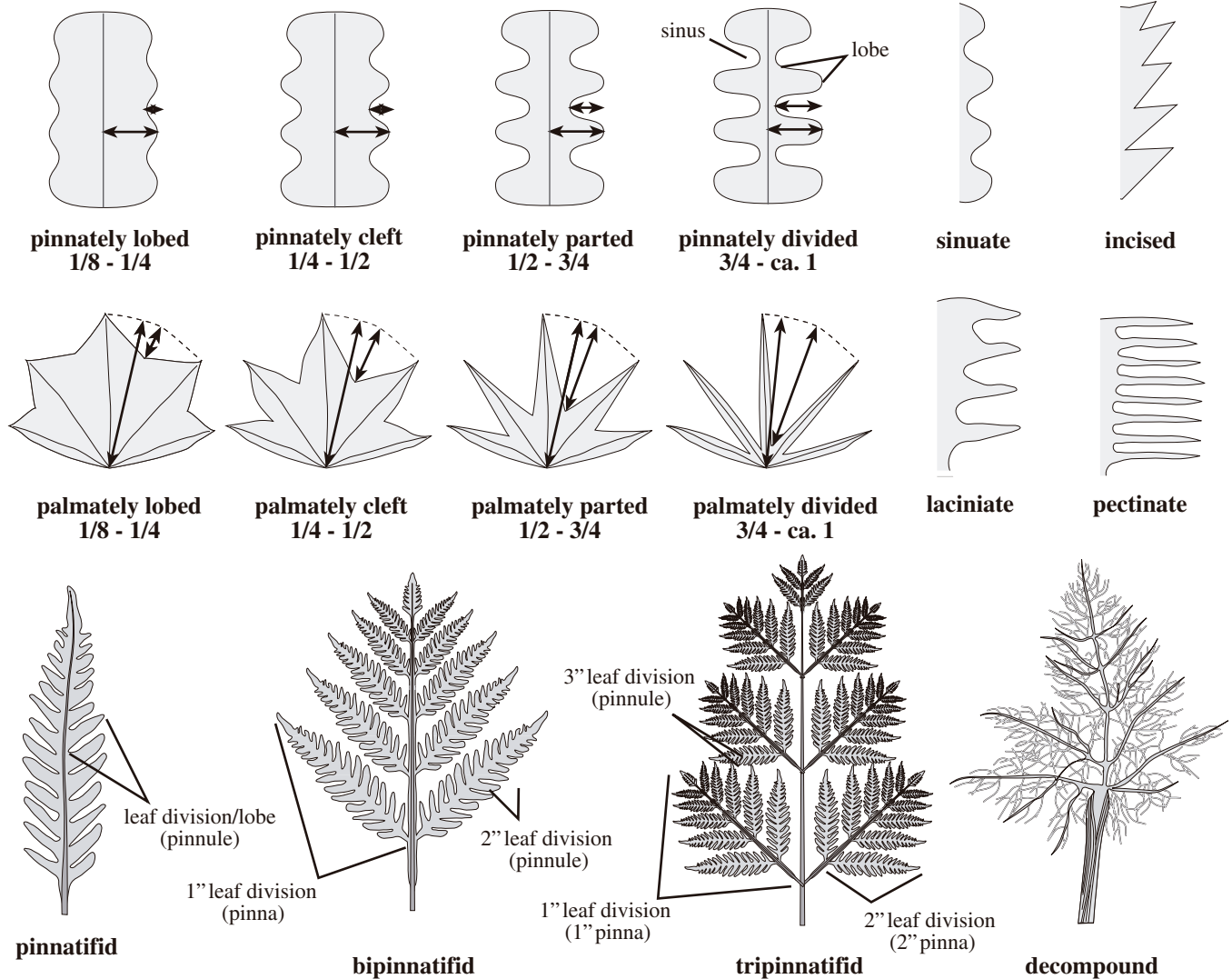


FIGURE 9.53 Division.

Lacerate refers to sinuses that are irregularly cut, the lobes appearing torn. **Laciniate** denotes lobes that are cut into narrow, ribbonlike segments. **Pectinate** means comblike, being pinnately divided with close, very narrow lobes.

DISPOSITION

Disposition refers to the relative placement of objects, e.g., of plant organs or parts of plant organs. Disposition is logically broken down into position, arrangement, orientation, and posture.

POSITION (Figure 9.54)

Position is the placement of parts relative to other, *unlike* parts. Some general position terms, which may apply to leaves, bracts, and flower parts, have to do with development. **Adaxial** (also known as **ventral**) corresponds to the *upper* or *inner* surface of an organ. Adaxial literally means toward the axis; in early development of the primordia of leaves or floral parts, the surface that is initially facing toward or nearest the axis will typically become the upper surface. Confusion arises when the organ in question bends downward or twists later in development; in such cases, it is clearer to state that a particular surface is *developmentally adaxial*. Correspondingly, **abaxial** (also known as **dorsal**) corresponds to the lower or outer surface of an organ, i.e., the surface most distant from the axis early in development. (Note that *ventral* and *dorsal* are used in an opposite sense to that for animals; for this reason, these terms are best avoided in plant descriptions, although they are still frequently used to refer to certain inflorescence, floral, or fruit features.)

With respect to a horizontally oriented structure, **posterior** refers to the upper lobe or part; **anterior** refers to the lower lobe or part. *Posterior* and *anterior* are widely used for

horizontally oriented floral parts and correspond to *adaxial* and *abaxial*, respectively. **Basal** or **radical** (not to be confused with the *radicle* of a seed embryo) indicates at or near the bottom or base of a structure. **Proximal** is similar to basal or radical and means near the point of origin or attachment, as in the point of attachment of a leaf. **Apical** or **terminal** means at or near the top, tip, or end of a structure. **Distal** is similar to *apical* or *terminal* and means away from the point of origin or attachment, e.g., the apex of a structure. Proximal and distal always refer to the point of attachment of an organ and are especially valuable for structures that loop around, in which the original base and apex are obscured. Some other general position terms are **lateral** or **axillary**, on the side of a structure or at the **axil** (the adaxial region of a node), as in a lateral or axillary bud; **central**, at or near the middle or middle plane of a structure; and **circumferential**, at or near the circumference of a rounded structure.

Some position terms are used primarily to describe the position of structures relative to the stem. In this sense, the general terms **radical** or **basal** mean positioned at the base of the stem; **cauline** means positioned along the length of the stem (as in cauline leaves or flowers); and **acrocaulis** means positioned at the apex of the stem.

ARRANGEMENT (Figure 9.55)

Arrangement is the placement of parts with respect to *similar, like* parts. Some arrangement terms, used primarily for leaves, bracts, or flower parts, describe the number of organs per node. **Alternate** refers to one leaf or other structure per node. Subcategories of alternate are **monostichous**, alternate with points of attachment in one, vertical row/rank, e.g., as in the Costaceae; **distichous**, alternate, with points of attachment in two vertical rows/ranks, e.g., as in the grasses

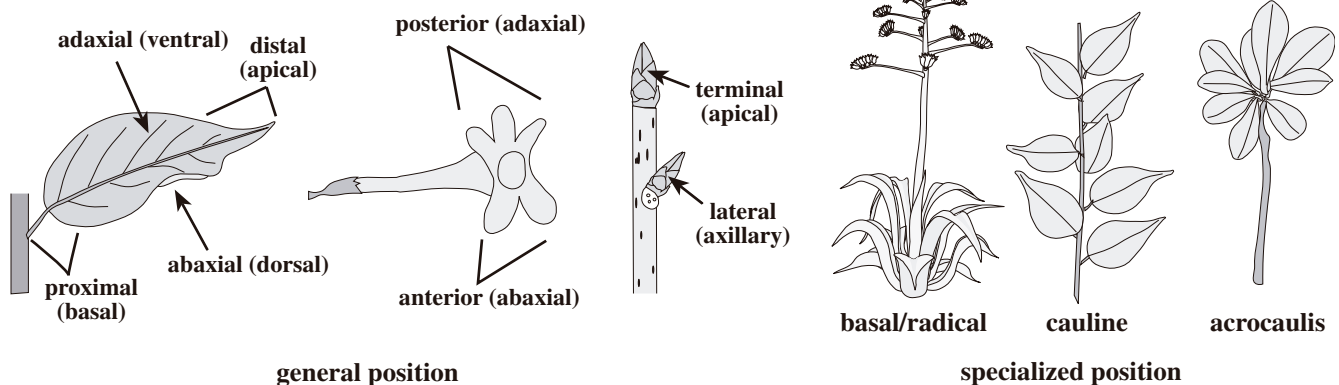


FIGURE 9.54 Position.

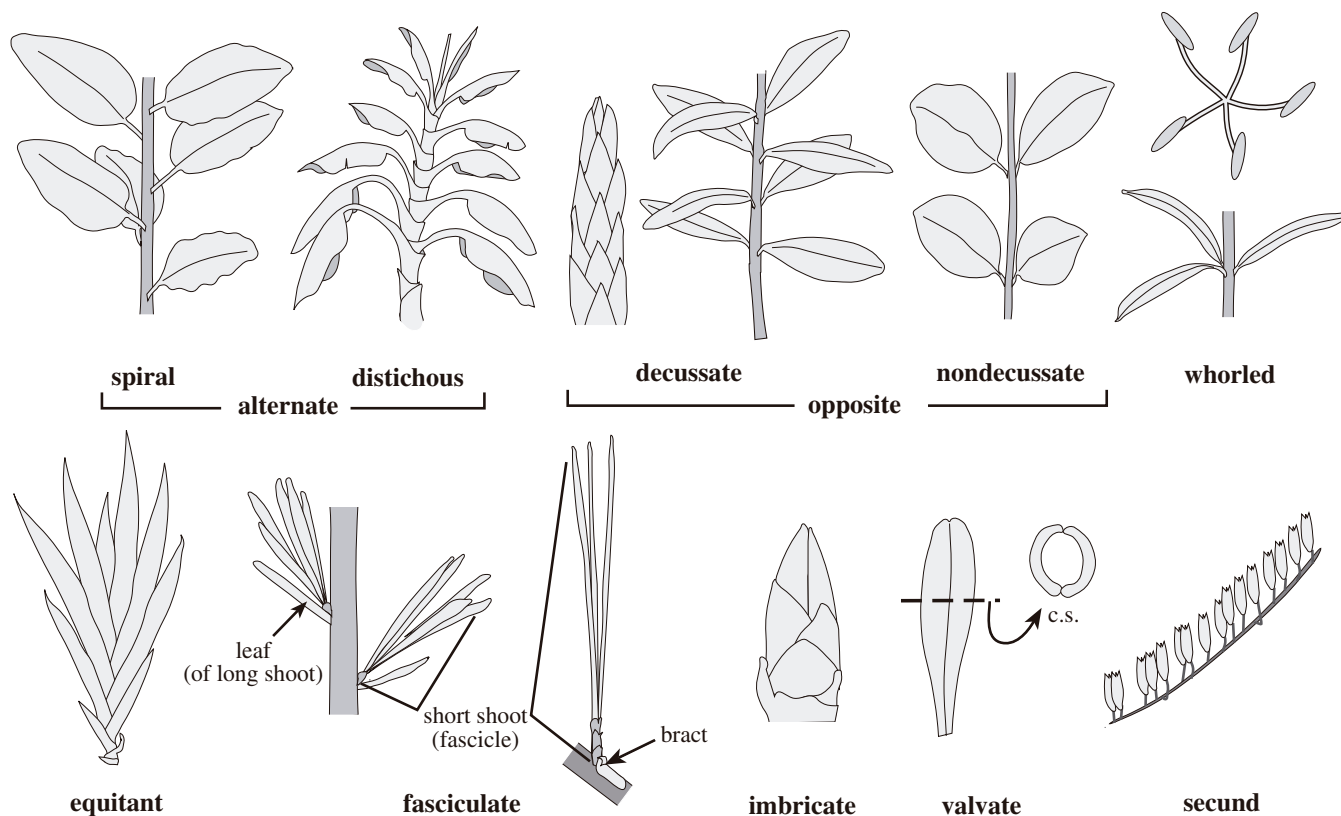


FIGURE 9.55 Arrangement. (c.s. = cross-section)

(Poaceae); **tristichous**, alternate, with points of attachment in three rows/ranks, as in the sedges (Cyperaceae); and **spiral** (also termed **polystichous**), alternate, with points of attachment in more than three rows/ranks.

Opposite describes two leaves or other structures per node, i.e., on opposite sides of a stem or central axis. Two subcategories of opposite are **decussate**, opposite leaves or other structures at right angles to preceding pair; and **nondecussate**, opposite leaves or other structures not at right angles to preceding pair. Most leaves, if opposite, are decussate; in fact, nondecussate leaves may be superficially the result of stem twisting. Leaflets of a compound leaf are typically nondecussate.

The term **subopposite** refers to two leaves or other structures on opposite sides of stem or central axis but at different nodes slightly displaced relative to one another. **Whorled** or **verticillate** means having three or more leaves or other structures per node.

More arrangement terms denote more specialized conditions. **Equitant** refers to leaves with overlapping bases, usually sharply folded along the midrib. **Fasciculate** refers to leaves or other structures in a fascicle or short shoot, a cluster with short internodes. **Imbricate** is a general term for leaves or other structures overlapping. **Valvate** means the sides are

enrolled, so that the margins touch. **Rosulate** means in a **rosette**, an arrangement in which parts (usually leaves) radiate from a central point at ground level (e.g., the leaves of *Taraxacum officinale*, dandelion). **Secund** or **unilateral** refers generally to flowers, inflorescences, or other structures on one side of the axis, often due to twisting of stalks.

ORIENTATION (Figure 9.56)

Orientation denotes the angle of a structure relative to a central (often vertical) axis. Precise orientation terms utilize ranges of angles in degrees, $0\ 15^\circ$ or $15^\circ\ 45^\circ$, from the upper axis, the horizontal axis, or the lower axis. These terms are (from top to bottom): **appressed**, pressed closely to axis upward, with divergence angle of $0\ 15^\circ$ from upper axis; **ascending**, directed upward, with divergence angle of $15^\circ\ 45^\circ$ from upper axis; **inclined**, directed upward, with divergence angle of $15^\circ\ 45^\circ$ from horizontal axis; **divergent** or **horizontal**, more or less horizontally spreading with divergence angle of $\leq 15^\circ$ up or down from the horizontal axis (also termed *divaricate* or *patent*); **reclined** or **reclinate**, directed downward, with divergence angle of $15^\circ\ 45^\circ$ from horizontal axis; **descending**, directed downward, with divergence angle of $15^\circ\ 45^\circ$ from lower axis; and **depressed**,

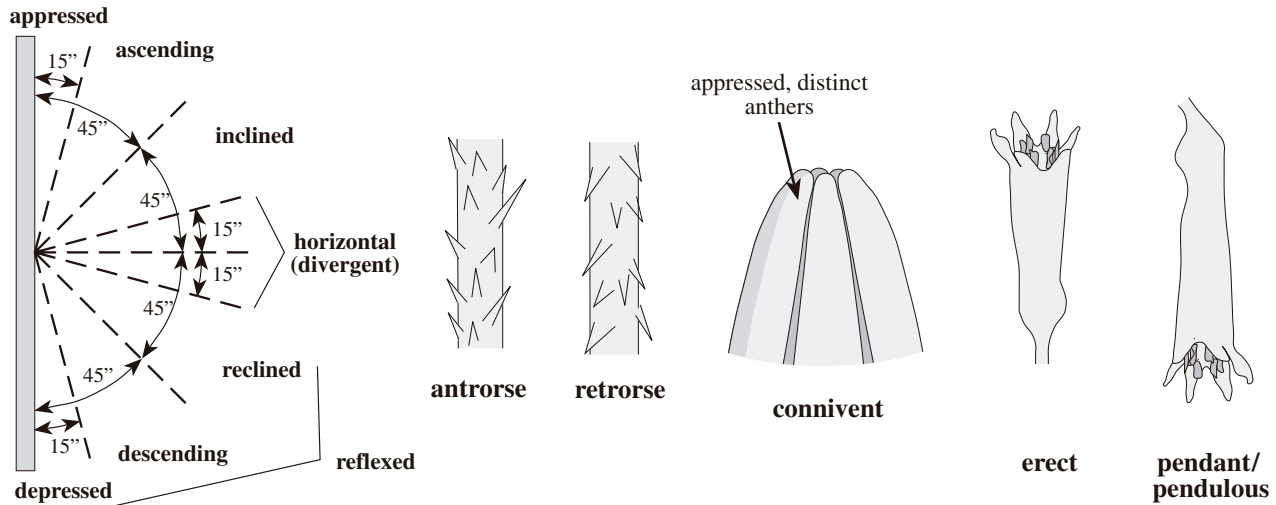


FIGURE 9.56 Orientation.

pressed closely to axis downward, with divergence angle of 0 – 15° from lower axis.

Other orientation terms are more general. **Antorse** means bent or directed upward, usually referring to small appendages; **retorse** means bent or directed downward. **Connivent** means convergent apically without fusion, as in anthers that come together in a flower (e.g., *Solanum*). **Erect** is pointing upward (usually without reference to an axis). **Pendant** or **pendulous** means hanging downward loosely or freely. **Deflexed** means bent abruptly downward, and **reflexed** means bent or turned downward.

POSTURE (Figure 9.57)

Posture refers to the placement relative to a flat plane. It may be further classified as transverse, longitudinal, twisting/bending posture, or ptyxis/vernation.

Transverse posture is the placement of the *tip* (distal end) of an object with respect to a starting plane. Transverse posture terms are **recurved**, tip gradually curved outward or downward (abaxially); **cernuous**, tip *drooping* downward (abaxially); **squarrose**, sharply curved downward or outward (abaxially) near the apex, as phyllaries of some Asteraceae; **incurved**, tip gradually curved inward or upward (adaxially); **plane** or **straight**, flat, without vertical curves or bends; and **flexuous**, the central axis and tip curved up and down.

Longitudinal posture is the placement of the *margins* of an object with respect to a starting plane. (Note: see also Aestivation.) Common longitudinal posture terms are **conduplicate**, longitudinally folded at central axis, with adjacent adaxial sides facing one another; **revolute**, with margins or outer portion of sides rolled outward or downward over

the abaxial surface; **involute**, with margins or outer portion of sides rolled inward or upward over adaxial surface; **cup-shaped**, concave-convex along entire surface (may be abaxially or adaxially concave); **plicate**, pleated, with a series of longitudinal folds (subcategories of plicate used, e.g., for palm leaves are **induplicate**, plicate with adjacent adaxial sides facing one another, V-shaped in cross-section; or **reduplicate**, plicate with adjacent abaxial sides facing one another, A-shaped in cross-section); and **undulate** or **repand**, the margins wavy in a vertical plane (compare *sinuate*, under Division).

Twisting/bending posture refers to the posture of a twisting or bending object relative to a starting plane. **Resupinate** means inverted or twisted 180° , as in leaves of Alstroemeriaceae or ovaries of most Orchidaceae flowers. **Geniculate** is having a zig-zag posture, as in the inflorescence rachis of some grasses. **Twining** is twisted around a central axis, as in many vines. The stems of twining vines may be **dextrorse**, twining helically like a typical, right-handed screw, as in some Convolvulaceae; or **sinistrorse**, twining helically like a left-handed screw, as in some Caprifoliaceae.

Ptyxis, also termed **vernation**, refers to the posture of embryonic structures, such as cotyledons within a seed or immature leaves or leaf parts. Many of the same terms used for posture of mature organs can be used to designate ptyxis. Some specialized ptyxis terms include **circinate**, with the blade (including rachis and rachillae, if present) coiled from apex to base, as in young fern and cycad leaves (see Chapters 4, 5); and **supervolute**, with one half of a simple leaf coiled tightly around the midrib, the other half coiled (in the opposite direction) around the first half, as in members of the Zingiberales (see Chapter 7).

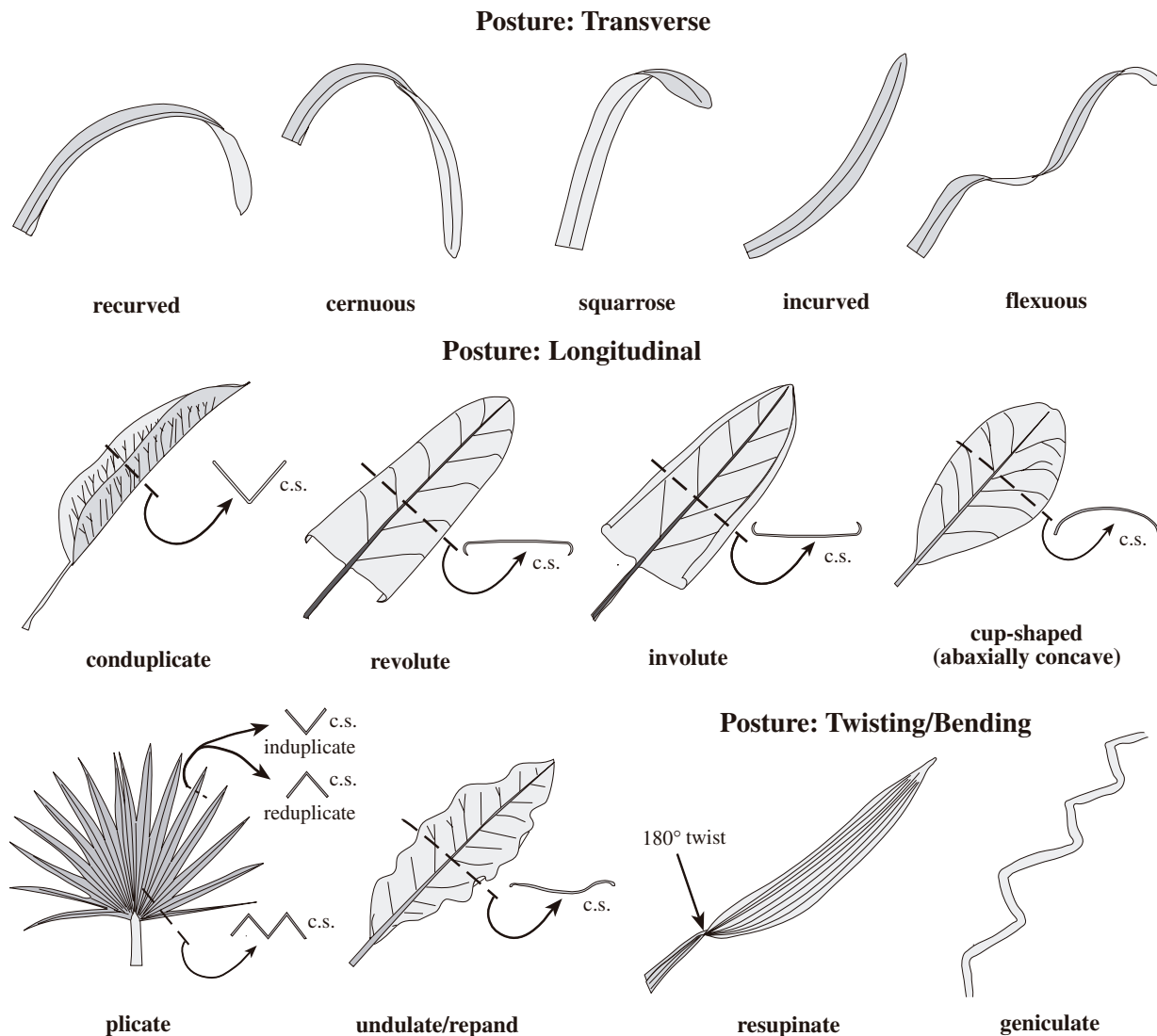


FIGURE 9.57 Posture. (c.s. = cross-section)

SURFACE

Numerous terms describe the surface of organs or plant parts. Surface features can be broken down into three characters: configuration, epidermal excrescence, and vestiture. Aspects of all three characters may be described as surface features. In addition, trichome type and bristle type may be described as surface features.

CONFIGURATION (Figure 9.58)

Configuration refers to the gross surface patterns of the epidermal cells other than that caused by venation (see Leaf Venation, earlier), or excrescences (next). Configuration terms include **canaliculate**, longitudinally grooved, usually in relation to

petioles or midribs; **fenestrate**, having windowlike holes in the surface (e.g., *Monstera deliciosa*, Araceae); **punctate**, covered with minute, pitlike depressions; **rugose** or **bullate**, covered with coarse reticulate lines, usually with raised blisterlike areas between; **ruminate**, unevenly textured, coarsely wrinkled, looking as if chewed (also used for texture); **smooth** or **plane**, with a smooth configuration; **striate**, with fine longitudinal lines; **wrinkled**, with irregular, fine lines or deformations.

EPIDERMAL EXCRESCENCE (Figure 9.58)

Epidermal excrescence refers to surface patterns caused by secretions or structural outgrowths of the epidermis (other than trichomes or bristles). Terms that denote epidermal

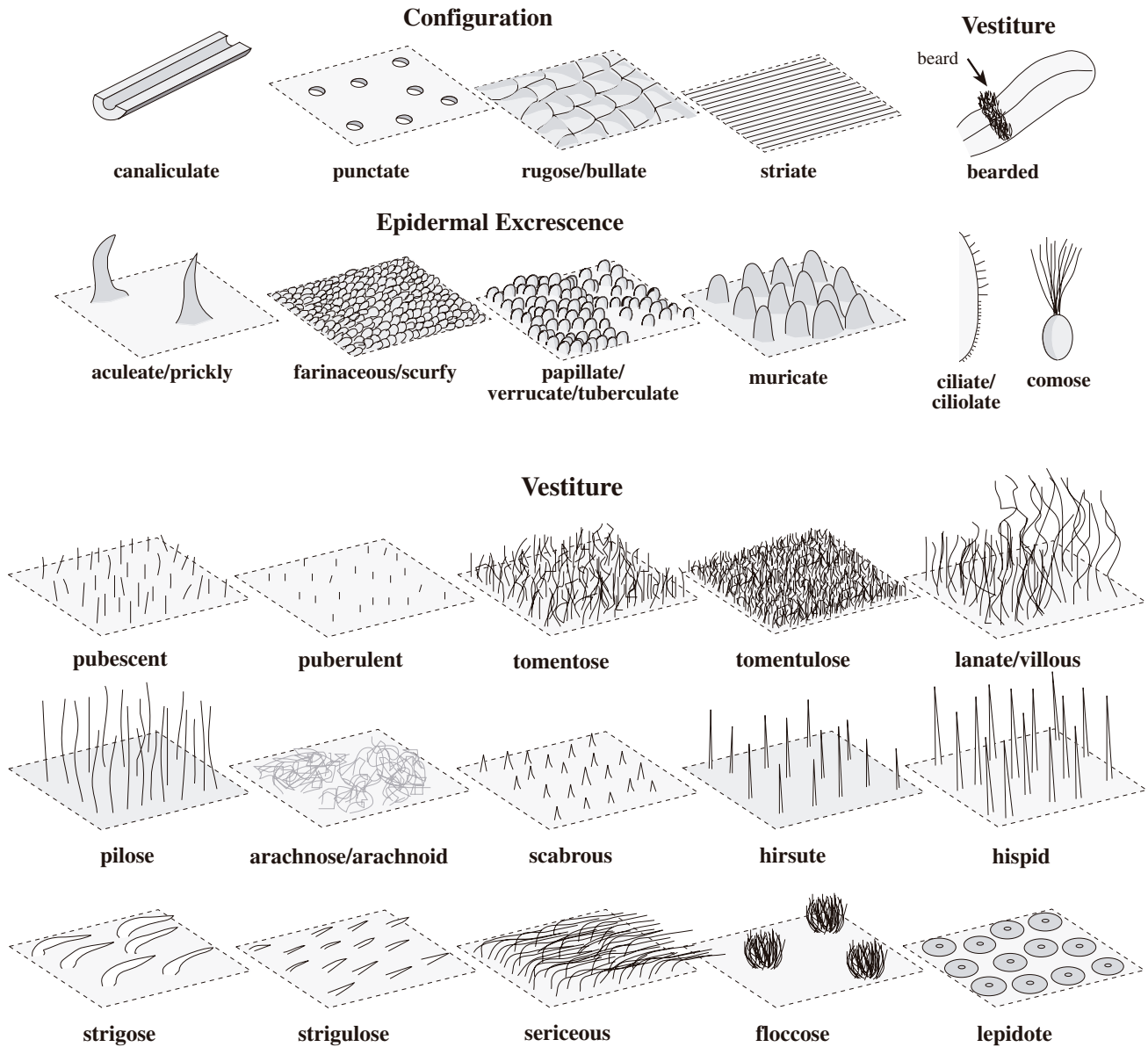


FIGURE 9.58 Surface features: con uration, epidermal excrecence, and vestiture.

secretions are **glandular**, covered with minute, blackish to translucent glands; **glaucous**, covered with a smooth, usually whitish, waxy coating (that can be rubbed off with touch); **shining (nitid or laevigate)**, appearing lustrous or polished; and **viscid or glutinous**, having a shiny, sticky surface. Terms that denote epidermal outgrowths are **aculeate** or **prickly**, with prickles, sharp nonspine, nonhorn appendages; **farinaceous (scurfy or granular)**, finely mealy, covered with small granules; **muriccate**, having coarse, radially elongate, rounded protuberances; **papillate, tuberculate, or verrucate**,

having minute, rounded protuberances; and **scabrous**, having a rough surface, like that of sandpaper (also treated under Vestiture).

VESTITURE (Figure 9.58)

Vestiture denotes trichome cover. Trichomes are surface hair-like structures that may function by protecting the plant from herbivory, reflecting visible and UV light, and inhibiting water transpiration. Vestiture terms encompass a combination of trichome type, length, strength, shape, density, and color.

Glabrous is the term meaning without trichomes at all. (A glabrous surface is often, but not necessarily, smooth or plane, which is used to denote a flat configuration.) **Subglabrous** means nearly glabrous (with just a few, scattered trichomes) and **glabrate** means the same or can mean becoming glabrous with age by loss of trichomes.

Bearded means with a single tuft or patch of trichomes arising from the surface of an object, e.g., from a petal. **Comose** is similar, but refers to an apical tuft of trichomes, e.g., from a seed. Two terms that are also treated under margin are **ciliate**, with conspicuous marginal trichomes, and **ciliolate**, with tiny or small marginal trichomes.

Pubescent is a common vestiture type meaning with more or less straight, short, soft, somewhat scattered, slender trichomes. (Note that *pubescent* can be used as a general term, meaning simply having trichomes.) **Puberulent** means minutely pubescent, i.e., pubescent but with very short or sparse (scattered) trichomes. **Canescent** or **incanous** means whitish-pubescent, covered with dense, fine grayish to white trichomes. **Tomentose** means covered with very dense, interwoven trichomes. **Tomentulose** is minutely tomentose, i.e., tomentose but with very short trichomes.

Villous or **lanate** means covered with long, soft, crooked trichomes; although *lanate* has shorter trichomes than *villous*, these terms intergrade and are probably best treated synonymously. **Pilose** means having soft, straight to slightly shaggy trichomes, generally at right angles to the surface. **Arachnose** or **arachnoid** means having trichomes forming a dense, cobwebby mass (but which resemble *villous* and can be confused with that type).

Scabrous means having rough trichomes, like that of sandpaper; **scaberulous** means minutely scabrous. (*Scabrous* is also treated under Epidermal Excrescence, because the scabrosity can be caused by either outgrowths or trichomes.)

Hirsute means having long, rather stiff trichomes (but not quite skin-penetrating), whereas **hispid** means having very long, stiff trichomes, often capable of penetrating skin. The term **urent** means having hispid trichomes that are stinging, as in *Urtica*, stinging nettle.

Strigose is covered with dense, coarse, bent and mostly flat trichomes often with a bulbous base. **Strigulose** is minutely strigose, i.e., with the same morphology but a much smaller size. **Sericeous** describes long, appressed trichomes that have a silky appearance or sheen.

Floccose means having dense trichomes that are in several patches or tufts. **Lepidote** means covered with scales or scalelike structures (intergrading with an epidermal excrescence character).

TRICHOME AND BRISTLE TYPE (Figure 9.59)

Trichome type refers to the specific, microscopic structure of individual trichomes and may come under the realm of plant anatomy (see Chapter 10) and constitute a systematically valuable character. Although trichome type and vestiture may be correlated, *vestiture* refers to the gross appearance of masses of trichome. For example, a tomentose vestiture could have any number of trichome types, e.g., dendritic, stellate, multiseriate tapering, or uniseriate tapering.

Trichome type may assess the number of cells per trichome. A **unicellular** trichome consists of a single cell and is

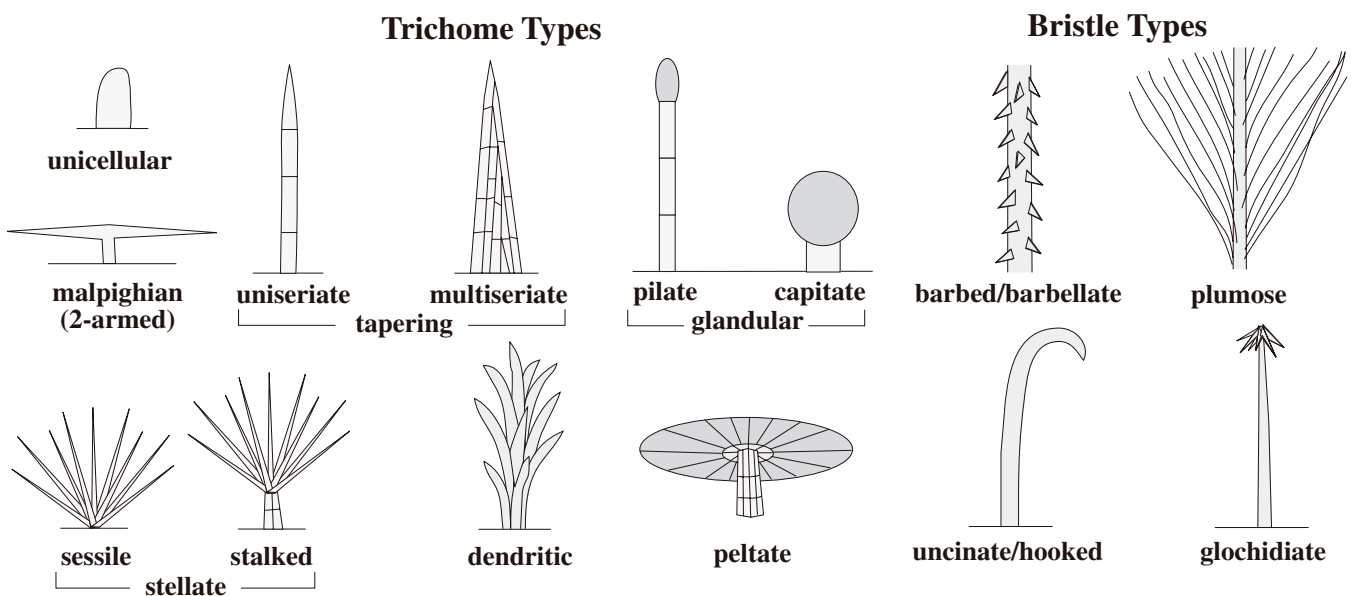


FIGURE 9.59 Trichome and bristle types.

usually quite small. A **multicellular** trichome contains two or more cells. Multicellular trichomes can be either **uniseriate**, having a single vertical row of cells, or **multiseriate**, having more than one vertical row of cells. The number of cell layers in a trichome can also be diagnostic.

Many trichomes are diagnosed based on their general shape and morphology. **Tapering** trichomes are those ending in a sharp apex. **Malpighian** or **two-armed** trichomes are those with two arms arising from a common base. (Malpighian is named after the family Malpighiaceae, where this trichome type is common.) **Glandular** trichomes are secretory or excretory trichomes, usually having an apical glandular cell. Glandular trichomes can be **pilate-glandular**, with a glandular cell atop an elongate basal stalk, or **capitate-glandular**, with a glandular cell having a very short or no basal stalk. Branched trichomes include two types: **stellate**, which are star-shaped trichomes having several arms arising from a common base (either stalked or sessile); and **dendritic**, which are treelike trichomes with multiple lateral branches. **Peltate** trichomes are those with a disk-shaped apical portion atop a peltately attached stalk.

Trichomes may also be delimited based on their position and function. For example, trichomes found in the axils of typically sheathing leaves, which may function in secreting protective mucilage, are termed **intravaginal** (or **axillary**) **squamules** (found, e.g., in many Alismatales).

Bristles are similar to trichomes but are generally much stouter (although bristles and trichomes may intergrade). Some so-called bristles are actually modified leaves, such as the glochidia of cacti. Major bristle types include **barbed** or **barbellate**, with minute, lateral, sharp appendages (*barbs*, which may be antrorse or retrorse in orientation) arising along the entire bristle surface; **plumose**, featherlike, covered with fine, elongate, ciliate appendages; **uncinate** or **hooked**, with an apical hooklike structure; and **glochidiate**, with apical, clustered barblike structures.

SYMMETRY

Although symmetry is usually used with reference to flowers (see Flower Symmetry), it can be a general feature to describe any plant organ or part. Symmetry is defined by the presence and number of mirror-image planes of symmetry. **Zygomorphic** or **bilateral symmetry** (also termed **irregular**) is that with one plane of symmetry. **Biradial symmetry** means having two (and only two) planes of symmetry. **Actinomorphic** or **radial symmetry** (also termed **regular**) is that with three or more planes of symmetry. **Asymmetric** describes a structure lacking any plane of symmetry. (Note that the distinction between biradial and radial symmetry is sometimes not recognized, both being termed *radial symmetry*

or *actinomorphy*; however, that distinction is often useful and is recognized here.)

TEMPORAL PHENOMENA

Temporal phenomena deal with any consideration specifically time-based. These are logically broken down into duration, maturation, and periodicity.

Duration refers to the length of life of a plant or part of a plant. **Plant duration** describes the length of life of an entire plant: annual, biennial, and perennial. An **annual** is a plant living 1 year or less, typically living for one growing season within the year. Annual plants are herbs (although herbs can be either annuals, biennials, or perennials). Annuals can usually be detected in that they lack an underground rootstock and show no evidence of growth from a previous season (e.g., there are no thickened, woody stems, dormant buds, or old fruits). **Biennials** are plants living 2 years (or two seasons), usually flowering in the second year. Biennial plants typically form a basal rosette of leaves during the first year and bolt (grow an elongate inflorescence stalk) in the second year. Biennials may be hard to detect without actually observing plants over two seasons. A **perennial** is a plant living more than 2 years. Perennials include herbs with rootstocks, shrubs, lianas, and trees.

Other duration terms describe plant parts, e.g., of leaves (in which the term **leaf duration** is used). **Evergreen** means persistent two or more growing seasons, as in the leaves of most conifers. **Deciduous** means parts persistent for one growing season, then falling off, as the leaves of *Acer*, maples. (Note that *evergreen* and *deciduous* can refer to the plants themselves, as in eastern deciduous forest.) **Caducous** means dropping off very early (compared with what is typical) and usually applies to floral parts. **Marcrescent** means ephemeral but with persistent remains, withering but persistent, such as corollas that remain attached during fruit formation. **Accrescent** refers to plant parts that persist and continue to grow beyond what is normal or typical, as with the calyx of *Physalis* (Solanaceae), which expands considerably and functions as an accessory part enclosing the fruit. Finally, **monocarpic** refers to a plant that flowers and fruits only once, then dies; the plant itself can be an annual or perennial, but the term is usually used only for perennials (because all annuals are, by definition, monocarpic).

Maturation refers to the relative time of development of plant parts. The term **anthesis** refers to the time of flowering, when flowers open with parts available for pollination. **Protandrous** (meaning male first) refers to stamens or anthers developing before the carpels or stigma. **Protogynous** (meaning female first) refers to the stigma or carpels maturing before the stamens or anthers. Both protandry and protogyny are general mechanisms to promote outcrossing within a species (see Chapter 13).

Periodicity refers to periodically repeating phenomena. Terms that refer to the time of day are **diurnal** (during the day), **nocturnal** (at night), **matutinal** (in the morning), and **vespertine** (in the evening). These terms are usually used

with respect to when flowers of a given taxon open. Other terms correspond to seasons, such as **vernal** (appearing in spring), **aestival** (appearing in summer), or **autumnal** (appearing in fall).

REVIEW QUESTIONS

PLANT STRUCTURE: GENERAL, ROOTS, AND STEMS/SHOOTS

1. Name the major plant organs.
2. What are the continuously actively dividing cell regions of a plant called and where are they located?
3. What is meant by plant habit and what are the types of plant habit?
4. Name various types of plant habitat.
5. What is the function of roots?
6. What are the root cap, root hair, adventitious root, and lateral root?
7. What is the difference between a taproot and a fibrous root system?
8. What is a shoot?
9. What is a bud, where do buds typically develop, and what do they develop into?
10. Define node, internode.
11. What is the difference between a bulb, corm, and tuber? between a rhizome, caudex, and stolon (runner)?
12. What is the difference between a caudiciform stem and a pachycaul?
13. What is thorn and how does it differ from a spine or prickly?
14. Define: tiller, burl, pseudobulb, short shoot, tendrils.
15. Name the difference between acaulescent and caulescent; between prostrate, repent, and decumbent. What is the corresponding character for all of these?
16. What is the difference between monopodial and sympodial?
17. Draw a typical twig and label terminal bud, axillary bud, leaf scar, vascular bundle scars, lenticels.
18. What is the difference between an axillary, terminal, and pseudoterminal bud? a collateral and superposed bud?

PLANT STRUCTURE: LEAVES

19. What is the difference between a bract and a scale?
20. Name some specialized modifications of leaves associated with flowers or inflorescences.
21. From what is a phyllode derived?
22. What is a spine and what are the three major types?
23. Name three modifications of leaves found in carnivorous plants.
24. Name five leaf types.
25. What are the basic components of a simple leaf?
26. Draw a bipinnately compound leaf and label: leaflet, petiole, petiolule, rachis, rachilla, stipule, stipel.
27. What is the difference between imparipinnate and paripinnate? trifoliolate and palmate? geminate-pinnate and bipinnately compound? unifoliolate and simple?
28. Name four different types of leaf attachment.
29. What is the difference between parallel and penni-parallel? between pinnate-netted, palmate-netted, and ternate-netted?
30. Name four major types of specialized venation types.

PLANT STRUCTURE: FLOWERS AND PERIANTH

31. Draw a typical flower and label all the parts, including collective terms.
32. Name the two basic types of flower sex.
33. Name the three basic types of plant sex. What is the corresponding type of flower sex for each?
34. Draw a zygomorphic corolla and label anterior lobe(s) and posterior lobe(s).
35. What is the difference between radial and biradial symmetry?

36. What is the difference between protandrous and protogynous? between centrifugal and centripetal?
37. What is a claw, corona, hypanthium, limb, lip, lobe, spur, throat, tube?
38. What are the two major types of perianth arrangement?
39. What is perianth cycly?
40. What is the difference between dichlamydeous and homochlamydeous?
41. Name two types of calyx fusion; of corolla fusion.
42. Define or draw the following perianth types: bilabiate, campanulate, rotate, salverform, urceolate.
43. Draw and label a petal with a claw and limb. What is the name of this perianth type?
44. Define convolute, imbricate, and valvate. What is the corresponding character?

PLANT STRUCTURE: ANDROECIUM

45. Name the two parts of a stamen; the two parts of an anther.
46. What is the difference between stamen arrangement and stamen position?
47. What is the difference between didymous, didynamous, and tetradynamous? What is the character?
48. What is the difference between antipetalous, antisepalous, and diplostemonous?
49. Do the above terms refer to stamen arrangement or to stamen position?
50. What is the difference between exerted and inserted? What is the character?
51. What is the term for fusion of stamens to the corolla?
52. What is the term for fusion of all the filaments together?
53. What is the term for fusion of the filaments into two groups?
54. What is a monotheical anther?
55. Name three types of anther attachment.
56. Name two types of anther dehiscence with regard to (a) the shape of the opening; (b) the direction of the opening.
57. What is a nectary and what are some types of nectaries?

PLANT STRUCTURE: GYNOECIUM

58. What is the difference between a gynoecium, carpel, and pistil?
59. What are the three parts of a pistil? What is a locule?
60. Name the two types of gynoecial fusion.
61. How is carpel number determined?
62. Name and draw the two basic types of ovary attachment and ovary position.
63. What does perianth/androecial position mean? Name and distinguish between four of these.
64. What is the difference between axile and parietal placentation? between basal and apical?
65. What is a gynobasic style?

PLANT STRUCTURE: INFLORESCENCES

66. What are two types of specialized bracts associated with inflorescences?
67. What is the difference between a pedicel and a peduncle?
68. Define compound receptacle.
69. What are three types of inflorescence position?
70. What is the difference between determinate and indeterminate inflorescence development?
71. What is a dichasium?
72. How does a monochasium differ and what are two major types?
73. What is a ray in an inflorescence?
74. What is the difference between a raceme and a spike? What is the inflorescence development of both?
75. What is the difference between a raceme and a panicle?
76. What is the difference between an umbel and a corymb? between an umbel and compound umbel?
77. What is the difference between a thyrse and verticillaster?
78. Name a taxonomic group characterized by a compound umbel; cyathium; head; hypanthodium; spadix.

PLANT STRUCTURE: FRUITS

79. What are the differences between simple, aggregate, and multiple fruits?
80. What features are used to define and classify fruit types?
81. What is a schizocarp? a mericarp? a valve?
82. What are the similarities and differences between an achene grain (caryopsis) and a nut?
83. What are the differences between loculidial, septicidal, and circumscissile capsules?
84. What are the similarities and differences between a follicle, legume, and silique?
85. What is the difference between a silique and a silicle? What family do they occur in?
86. What is the name given to a winged fruit?
87. How does a berry differ from a drupe or a hesperidium?
88. What is the placentation, ovary position, and texture of a pepo? In what family are they found?
89. A pome consists of much outer fleshy tissue derived from what? What is the ovary position? What is an example of a plant with pomes?
90. Name two types of aggregate fruits.
91. What types of fruits are burs, soroses, and syconia?
92. Name two types of seed based on endosperm type; seed germination type.

GENERAL TERMINOLOGY: COLOR, NUMBER, TEXTURE, AND FUSION

93. What is the difference between color and color pattern? Name and define a color pattern character state.
94. What is the difference between cycly and merosity? Give an example of each.
95. What is the difference between coriaceous and indurate? between scarios and succulent? What is the character for these?
96. What is the difference between connate and distinct? between adnate and free? between adherent and coherent? What is the character for these?

GENERAL TERMINOLOGY: SHAPE

97. Define the following terms for three-dimensional shapes: capitate, clavate, filiform, pilate, terete.
98. What is the difference between lanceolate, ovate, and lance-ovate? between lanceolate and oblanceolate? (Draw.)
99. What is the difference between ovate and obovate? between oblanceolate and spatulate? (Draw.)
100. What is the difference between elliptic and oblong? between oblong and linear? (Draw.)
101. What is the difference between cordate and reniform? (Draw.)
102. What is the difference between hastate and sagittate? (Draw.)
103. What does peltate mean?
104. What does an attenuate base mean? an oblique base? a sagittate base? a cuneate base?
105. What is the difference between entire, crenate, serrate, and dentate? What character do these refer to?
106. What is the difference between crenate and crenulate? serrate and serrulate? dentate and denticulate?
107. Define ciliate, ciliolate, filiferous.
108. What is the difference between acuminate, narrowly acute, acute, and obtuse (apex)? What are the corresponding base terms?
109. What is the difference between rounded and truncate (apex and base)?
110. What is the difference between mucronate, aristate, and apiculate?
111. What does emarginate or retuse mean (apex)?
112. What is the difference between lobed, cleft, parted, and divided?
113. What do pinnatifid and bipinnatifid mean?
114. Draw the following: (a) simple, sessile, ovate, acute, crenate leaf; (b) simple, petiolate, oblanceolate, serrulate leaf; (c) pinnately compound, petiolate, stipulate and stipellate leaf with sessile, entire, narrowly elliptic, cuneate, acuminate leaflets; (d) trifoliolate (ternately compound), petiolate leaf with petiolulate obovate, narrowly cuneate, apically obtuse, mucronate leaflets; (e) simple, lanceolate, mucronate, sagittate, dentate leaf.

GENERAL TERMINOLOGY: DISPOSITION

115. What is the difference between position and arrangement?
116. What is the difference between abaxial (dorsal) and adaxial (ventral)?
117. What is the difference between proximal and distal?
118. What does radical mean?
119. Name the three basic (general) types of arrangement (e.g., for leaves).
120. What is the difference between alternate and distichous? between opposite and decussate?
121. Describe the difference between equitant, imbricate, secund, valvate, and rosulate.
122. Define orientation and name three types (character states).
123. What is the difference between transverse posture and longitudinal posture? Give two examples of each.
124. Distinguish between conduplicate, revolute, sinuate, and undulate.

GENERAL TERMINOLOGY: SURFACE, SYMMETRY, TEMPORAL PHENOMENA

125. Surface refers to three features: configuration, epidermal excrescence, and vestiture. How do they differ?
126. Define rugose. For what character is this a character state?
127. What is the difference between glaucous, scabrous, and viscid? For what character are these character states?
128. What is the difference between hirsute, pubescent, and tomentose? For what character are these character states?
129. What do stellate, pilate, and uniseriate refer to? For what character are these character states?
130. What is the difference between actinomorphic (radial) and zygomorphic (bilateral)? For what character are these character states?
131. What is the difference between annual, biennial, and perennial? For what character are these character states?
132. What is the difference between caducous and accrescent? For what character are these character states?
133. What is the difference between protandrous and protogynous?

EXERCISES

1. Select a plant species and thoroughly describe its morphology using the comprehensive character list of Appendix 1. Fill in every applicable character with a character state, noting that several characters will not apply to your taxon. Try to examine several populations, individuals, or plant organs/parts and note the range of variation. For characters that are variable, either list the range of variation (e.g., *Leaves* oblanceolate to narrowly elliptic, crenate to dentate . . .) or list the most common morphology and in brackets list the exceptions (e.g., *Leaves* trifoliolate [rarely pinnate with 5 leaflets] or *Leaves* 4–7 [2.5–10] cm long . . .).
2. From the character listing of Appendix 1, write a detailed description, using the family descriptions of Chapters 7 and 8 as a model. Note to list only the plant organ or plant part, not the character. For example, the description format should be **Leaves** are opposite, simple, and evergreen and *not* **Leaf** arrangement is opposite, leaf type is simple, leaf duration is evergreen. (Note that a word processing merge file is often useful for this.) Edit this description such that it reads smoothly and avoids repetition.
3. Make detailed drawings, using a hard (2H or 3H) pencil, of various parts of your species, such as leaves, inflorescence, whole flower, flower in longitudinal section, anther close-up, ovary close-up, ovary cross- or longitudinal section, fruit, and seed. Be sure to include a scale bar, in metric measurements, beside each drawing. Make copies or tracings of these drawings and trace the outlines with a fine, black ink rapidograph. Attempt to do some fine stippling in various regions to show venation, shading, and depth. (See Appendix 2.)
4. Compare your description and drawings with that of standard references, including floras and monographic treatments. Note that yours is probably much more detailed and comprehensive than that of most floras, but perhaps comparable in detail to a monograph.

REFERENCES FOR FURTHER STUDY

- Bell, A. D. 1991. *Plant Form, an Illustrated Guide to Flowering Plant Morphology*. Oxford University Press, Oxford.
- Brouk, B. 1975. *Plants Consumed by Man*. Academic Press, London.

- Hallø, E. R. A. A. Oldeman, and P. B. Tomlinson. 1978. *Trophic Trees and Forests: An Architectural Analysis*. Springer, Berlin.
- Hickey, L. J. 1973. Classification of the architecture of dicotyledonous leaves. *American Journal of Botany* 60: 17-33.
- Hickey, L. J., and J. A. Wolf. 1975. The bases of angiosperm phylogeny: vegetative morphology. *Annals of the Missouri Botanical Garden* 62: 538-589.
- Lawrence, G. H. M. 1951. *Taxonomy of Vascular Plants*. Macmillan, New York.
- Radford, A. E., W. C. Dickison, J. R. Massey, and C. R. Bell. 1974. *Vascular Plant Systematics*. Harper & Row, New York.
- Rowley, Gordon D. 1987. *Caudiciform and Pachycaul Succulents Pachycauls, Bottle-, Barrel- and Elephant-Trees and Their Kin: A Collector's Miscellany*. Strawberry Press, Mill Valley, CA.
- Tucker, A. O., M. J. Maciarello, and S. S. Tucker. 1991. A survey of color charts for biological descriptions. *Taxon* 40: 201-214.
- Systematics Association Committee for Descriptive Terminology. 1962. II. Terminology of Simple Symmetrical Plane Shapes (Chart I). *Taxon* 11(5): 145-156.
- Weberling, F. 1989. *Morphology of Flowers and Inflorescences*. Cambridge University Press, Cambridge, United Kingdom.

This Page Intentionally Left Blank

10

PLANT ANATOMY AND PHYSIOLOGY

PLANT CELL STRUCTURE	409	PHOTOSYNTHESIS	429
PLANT GROWTH	411	ANATOMY AND SYSTEMATICS	430
PLANT TISSUES AND SPECIALIZED CELL TYPES	412	PLANT ANATOMY TECHNIQUE	432
ROOT STRUCTURE AND FUNCTION	417	REVIEW QUESTIONS	434
SHOOT AND STEM STRUCTURE AND FUNCTION	419	EXERCISES	436
SECONDARY GROWTH	421	REFERENCES FOR FURTHER STUDY	436
LEAF STRUCTURE AND FUNCTION	426		

Plant anatomy is the study of the tissue and cell structure of plant organs. The term *anatomy*, as applied to plants, generally deals with structures that are observed under a high-powered light microscope or electron microscope. (In zoology, the term *anatomy* refers to the study of internal organs; *histology* is the study of cells and tissues of animals.) **Plant physiology** is the study of metabolic processes in plants. A limited explanation of plant physiology is presented, dealing specifically with photosynthesis. Physiology and anatomy are tightly correlated, as cell and tissue structure has changed with respect to the evolution of novel functional mechanisms.

The following is a summary of basic plant anatomy and physiology, with a focus on the sporophytes of vascular plants. Plant anatomical and physiological features may provide valuable characters and character states in assessing homology and elucidating phylogenetic relationships among plants. Moreover, the evolution of anatomical and physiological characters is of great interest, as many of these features are of significant adaptive value and have been among the major selective pressures in plant evolution.

PLANT CELL STRUCTURE

In the 1600s the English biologist Robert Hook first coined the term **cell** after observing that plant tissues, such as

cork, are divided into little cavities separated by walls. Since then, the **cell theory** has been perfected, providing a unifying theme in biology, stating the following. First, all life is composed of one or more cells. Second, cells arise only from preexisting cells, occurring either through cell division (meiosis or mitosis) or cell fusion (e.g., fertilization of egg and sperm). Third, cells are the units of metabolic processes; thus, each cell contains the necessary chemical compounds and cellular components to carry on the biosynthetic pathways needed for basic physiological processes. Fourth, each cell contains a set of DNA, the hereditary material that is transferred from one cell to another that codes for the structural and functional features of the organism.

Plant cells are bounded by a **plasma membrane** that is composed of a phospholipid bilayer with embedded proteins. The membrane functions as the boundary of the cell, to contain the cellular components. It also functions in cell recognition and in transport of compounds. Everything inside the plasma membrane is called the **protoplasm**. The protoplasm contains **organelles**, which carry on some vital metabolic function, and **ergastic substances**, which do not function in metabolism and have a variety of functions such as storage, waste secretion, and protection.

Cellular organelles include the following (Figure 10.1): (1) a **nucleus**, which is double-membrane bound and contains DNA, the hereditary material of the cell (Note: Everything

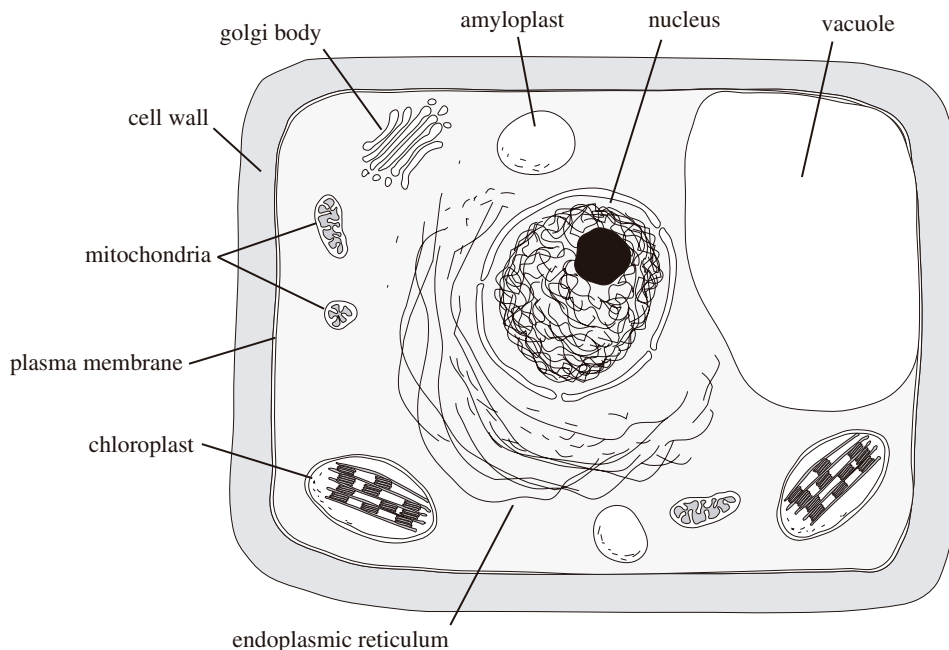


FIGURE 10.1 Cell structure of a typical, live plant cell.

inside the plasma membrane but not including the nucleus is called **cytoplasm**); (2) **mitochondria** (singular, *mitochondrion*), which are double membrane-bound, with invaginations called *cristae* that function in the electron transport reactions of respiration; (3) a **vacuole**, which is a large (often occupying *most* of the volume of plant cells), internal, membrane-bound sac that functions in storage of compounds such as pigments (e.g., anthocyanins or betalains), acids (e.g., malic acid involved in CAM photosynthesis), or ergastic substances (see later discussion); (4) **endoplasmic reticulum**, which is composed of interconnected phospholipid membranes and functions as the site of protein synthesis and material transport; (5) **golgi bodies**, which are composed of parallel stacks of flattened membranes and function in transport and modification of compounds; (6) **chloroplasts**, which are double membrane-bound with internal thylakoid membranes (composed of lamellae and grana in the green plants), functioning in the reactions of photosynthesis; and (7) **ribosomes**, which function as the site of protein synthesis.

Ergastic substances are cellular materials that are not actively metabolized, functioning, e.g., as storage reserves or wastes. Ergastic substances include (1) **chromoplasts** (Figure 10.2A), which are carotenoid-containing bodies that function to provide yellow, orange, or red pigmentation for a plant organ, as in petals or fruits; (2) **amyloplasts** or **starch grains** (Figure 10.2B,C), which are lamellate deposits of starch (alpha-1,4-glucopyranoside, a polysaccharide polymer

of glucose units with alpha-1,4 chemical bonding) which functions as the high-energy storage compound in green plants; (3) **aleurone grains** (=proteinoplasts), which are granular protein deposits, functioning as storage compounds; (4) **tannins**, which are phenol derivatives that may function to deter herbivory and parasite growth; (5) **fats, oils, waxes**, which are types of triglyceride compounds that may function as high energy storage compounds or secretion products; and (6) **crystals**, which may be composed of silica or calcium oxalate in various forms, such as **druses** (spherical crystals with protruding spikes; Figure 10.2D), **raphides** (bundles of needle-like crystals; Figure 10.2E), **styloids** (single, elongate, angular crystals; Figure 10.2F), or **prismatic** (shorter, prism-shaped crystals; Figure 10.2G). Crystals may function as waste products, as calcium ion sinks (a means of removing excess calcium for certain cellular functions), or as an irritant to deter herbivory.

In land plants, a pectic-rich **middle lamella** layer is formed between the plasma membrane of adjacent cells (Figure 10.3). The middle lamella functions to bind adjacent cells together. During plant cell development, a **cell wall** is secreted outside the plasma membrane. The cell wall that is secreted soon after cell division and that is maintained during cell growth is called the **primary** (1°) **cell wall** (Figure 10.3). As discussed earlier, an apomorphy for the green plants is a cell wall composed of **cellulose**, a polysaccharide polymer of glucose sugar units chemically bonded in the beta-1,4 position (=beta-1,4-glucopyranoside). Recall that cellulose is constructed of

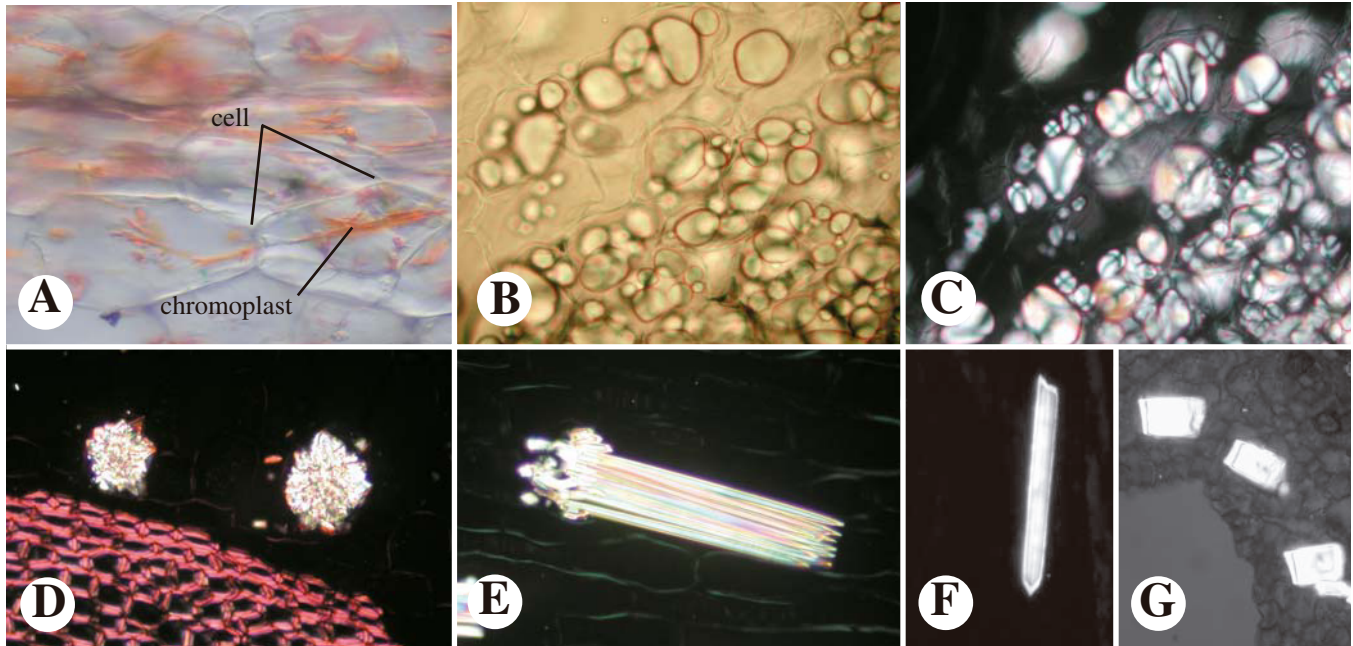


FIGURE 10.2 A. Chromoplasts, beta-carotene deposits in cultivated root of carrot (*Daucus carota*). B,C. Amyloplasts of potato (*Solanum tuberosum*). B. Bright field microscopy. C. Polarization microscopy, showing typical Maltese cross optical pattern of starch. D–G. Calcium oxalate rystals, all viewed with polarization microscopy. D. Druses. E. Raphides. F. Styloids. G. Prismatic crystals.

microscopic fiberlike units (called **microfibrils**) that are further intertwined into larger fibril units, forming a meshwork outside the plasma membrane. Its function is to impart rigidity to the cells, acting as a cellular skeleton. Within the primary cell wall, ultramicroscopic pores may form, termed **plasmodesmata**. These tiny openings function to allow for a continuity of membranes between cells, allowing for intercell exchange of compounds. A group of plasmodesmata is called a **primary pit field** (Figure 10.3).

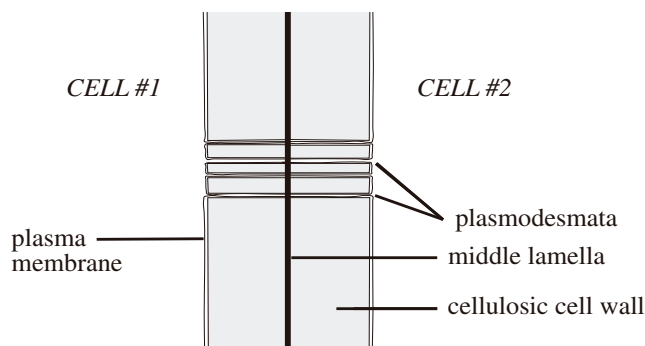


FIGURE 10.3 Cellulose cell wall. Note plasmodesmata, small pores or cavities in the cellulose structure of the primary cell wall.

In certain plant cells (e.g., sclerenchyma and tracheary elements) an additional wall layer, called a **secondary (2°) wall**, is secreted externally, between the primary cell wall and plasma membrane (Figure 10.4). A secondary wall is generally formed after the plant cell has ceased growth. In vascular land plants the secondary wall is composed partly of cellulose but also contains **lignin**, a complex polymer of phenolic compounds that binds the cellulose microfibrils together. Lignin imparts significant strength and rigidity to the cell wall.

In virtually all plant cells with lignified cell walls, there are holes in the secondary wall called **pits**. Pits of adjacent cells often occur opposite one another, as **pit-pairs** (Figure 10.4). The actual chamber and opening of a pit may assume different morphological forms. Pits function in allowing communication between cells during their development and differentiation. They may also have specialized functions in fluid conducting cells (see later discussion).

PLANT GROWTH

Plant cell growth is defined here in the broad sense as the initiation, expansion, and specialization of cells. The haploid spore or diploid zygote of land plants initially undergoes more or less continuous, sequential mitotic cell divisions.

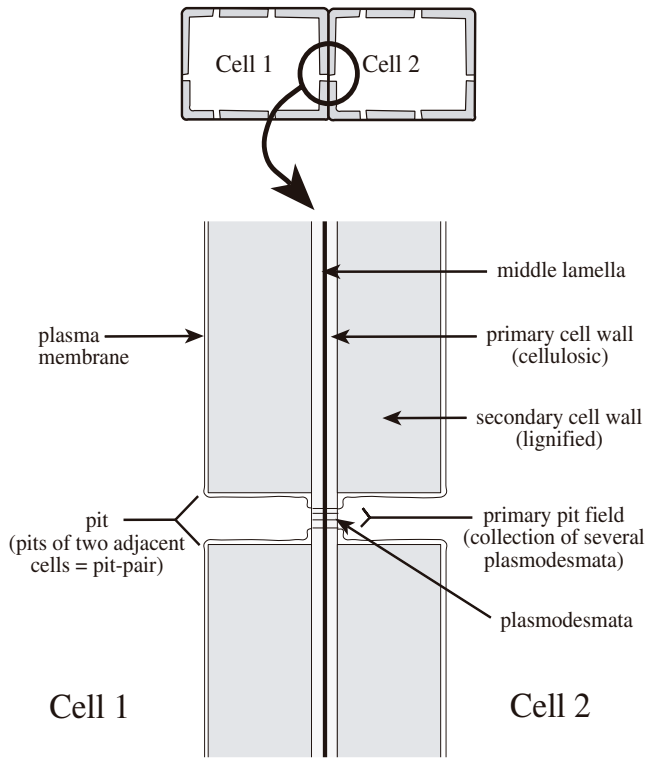


FIGURE 10.4 Lignified secondary cell wall of specialized cells of vascular plants. Note pit-pair.

Later, as gametophytes or sporophytes mature, active cell divisions become restricted to certain regions of the plant. This region of actively dividing cells is known as a **meristem**. In the vascular plants **apical meristems** are located at the apices of roots and shoots (Figure 10.5), resulting in growth in height or length. In woody plants both apical meristems

and lateral meristems occur. **Lateral meristems** are cylindrical sheaths of cells (Figures 10.18, 10.19), which function in growth that increases width or girth (see later discussion).

Apical meristematic cells are small and roughly isodiametric in shape, whereas those of the lateral meristems are elongate. A single meristematic cell undergoes a mitotic cell division, by which two cells are derived from one. Each of the two daughter cells undergoes some initial expansion. The derivatives themselves may continue to divide several more times, but only those cells that remain near the meristem will do so indefinitely. The others eventually cease mitosis and undergo further differentiation.

Cell differentiation refers to the series of changes that a cell undergoes from the point of inception to maturity, involving the transformation of a meristematic cell into one that assumes a particular structure and function. Differentiation involves two processes: **cell expansion**, in which the cell grows in size (often by **elongation**, in which growth in the axial direction is greatest); and **maturation** or **specialization**, in which the cell acquires the structural and functional features at maturity. Cell specialization means simply that cells may differ from one another, becoming specialized for a particular structure and function within the whole plant. Cell differentiation results in the development of various **cell types** (discussed next).

PLANT TISSUES AND SPECIALIZED CELL TYPES

A **tissue** is a group of cells having a common function or structure. Plant tissues of the vascular plants are often categorized

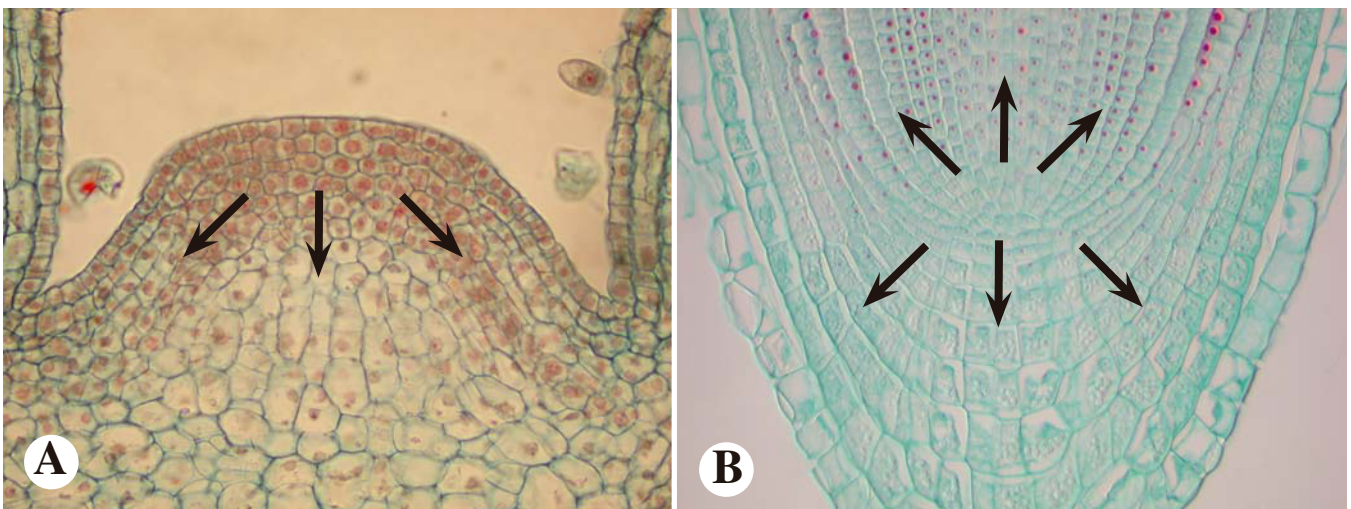


FIGURE 10.5 Meristematic cells. **A.** Shoot. **B.** Root. Arrows indicate general directions of cell growth.

into three broad classes: **ground**, **vascular**, and **dermal**; see later discussion. In addition, tissues may be classified as simple or complex. A **simple tissue** consists of only one type of cell; thus, a particular term may refer either to the simple tissue or the cell type. A **complex tissue** contains more than one cell type.

Ground tissue is that occurring inside the epidermis but not part of the vascular tissue. Three cell types (which are simple tissues) make up the ground tissue: parenchyma, collenchyma, and sclerenchyma. **Parenchyma** (Gr. *para*, beside + *en-chein*, to pour; in reference to the analogy that parenchyma is poured beside other tissues to fill up space) are cells that most resemble the unspecialized, undifferentiated cells of actively dividing meristematic tissue (Figure 10.6A). Structurally, parenchyma cells are (1) isodiametric to elongate; (2) have a primary (1°) cell wall only (rarely with secondary wall); and (3) are living at maturity and potentially capable of cell division. Parenchyma cells function in metabolic activities (e.g., respiration, photosynthesis, transport, storage) and in wound healing and regeneration, being capable of transforming into a meristem to form new roots or shoots.

Collenchyma (Gr. *colla*, glue + *enchyma*, infusion; in ref. to thick, glistening cell walls) are cells that structurally (1) are elongate; (2) have only a primary cell wall that is unevenly thickened and rich in pectins (glistening white in the light microscope); and (3) are living at maturity (Figure 10.6B). Collenchyma cells function in mechanical support and are often found at the periphery of stems or leaves. They can be stretched during elongation growth of the organ.

Sclerenchyma (Gr. *scleros*, hard + *enchyma*, infusion, in ref. to hard, lignified cell walls) are cells that structurally (1) have thick, lignified secondary (2°) cell walls, which may have pits; and (2) are (usually) dead at maturity. There are two general types of sclerenchyma, which sometimes intergrade: (a) **fibers**, which are long, very narrow cells with sharply tapering end wall (Figure 10.7A C); and (b) **scleireids**, which are isodiametric to irregular in shape and often branched (Figure 10.7D F). Fibers function in mechanical support in various organs and tissues, sometimes making up the bulk of the tissue. Fibers often occur in groups (or bundles) and may be components of the xylem and/or phloem or may occur independently of the vascular tissue. Scleireids may also function in structural support, but their role in some plant organs is unclear (e.g., possibly aiding in providing protection from herbivory).

Vascular tissues are made up of xylem and phloem, each of which are complex tissues (having more than one cell type). **Xylem** (Gr. *xylos*, wood) is a tissue composed of **tracheary elements** plus some parenchyma and sometimes sclerenchyma. Structurally, **tracheary elements** (1) are elongate to short; (2) have lignified, secondary (2°) cell walls, with **pits**; and (3) are dead at maturity, in which protoplasts degrade, leaving only cell walls (Figure 10.8). Tracheary elements are joined end-to-end, forming a tubelike continuum. They function to conduct water and dissolved essential mineral nutrients, generally from the roots to other parts of the plant.

There are two types of tracheary elements: **tracheids** and **vessel members**. These differ with regard to the junction between adjacent end-to-end cells, whether *imperforate* or *perforate*. **Tracheids** are imperforate, meaning that water

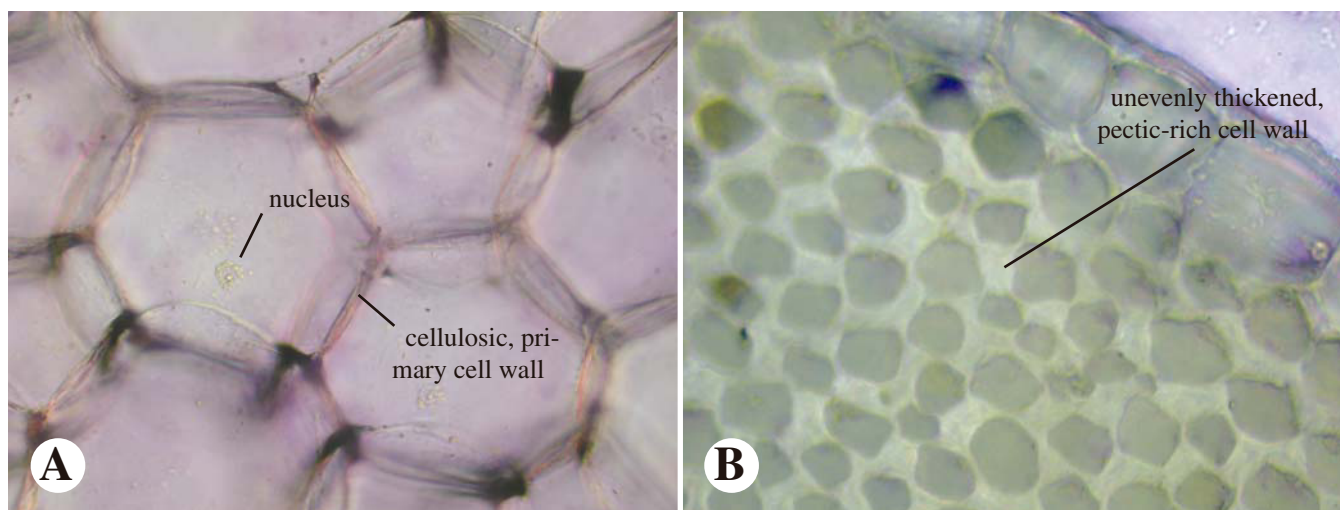


FIGURE 10.6 **A.** Parenchyma cells, live at maturity, with cellulose primary cell wall. Note nucleus. **B.** Collenchyma cells (cross-sectional view) live at maturity, with unevenly thickened, pectic-rich primary cell walls.

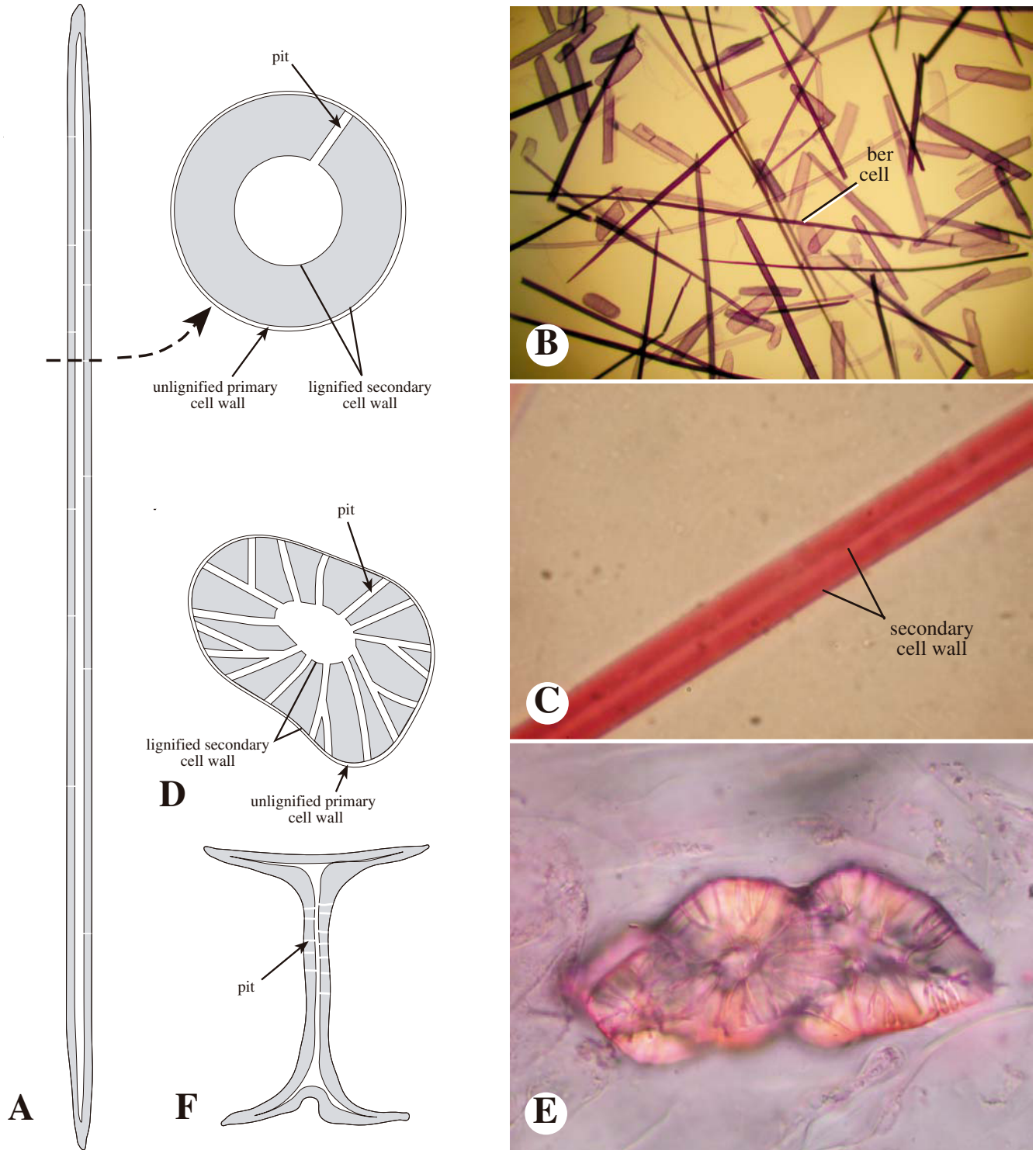


FIGURE 10.7 Sclerenchyma. **A–C.** Fiber cells. **A.** Diagram of fiber cell in longitudinal and cross-section. **B.** Whole mount of fiber cells, from macerated tissue of *Yucca*. **C.** Close-up of fiber cell, side view. Note thick, lignified secondary cell wall. **D–F.** Sclereid cells. **D,E.** Sclereids of pear (*Pyrus*) fruit tissue. Note thick, lignified secondary cell wall with numerous, canal-like pits. **F.** Sclereid of *Ficus* leaf.

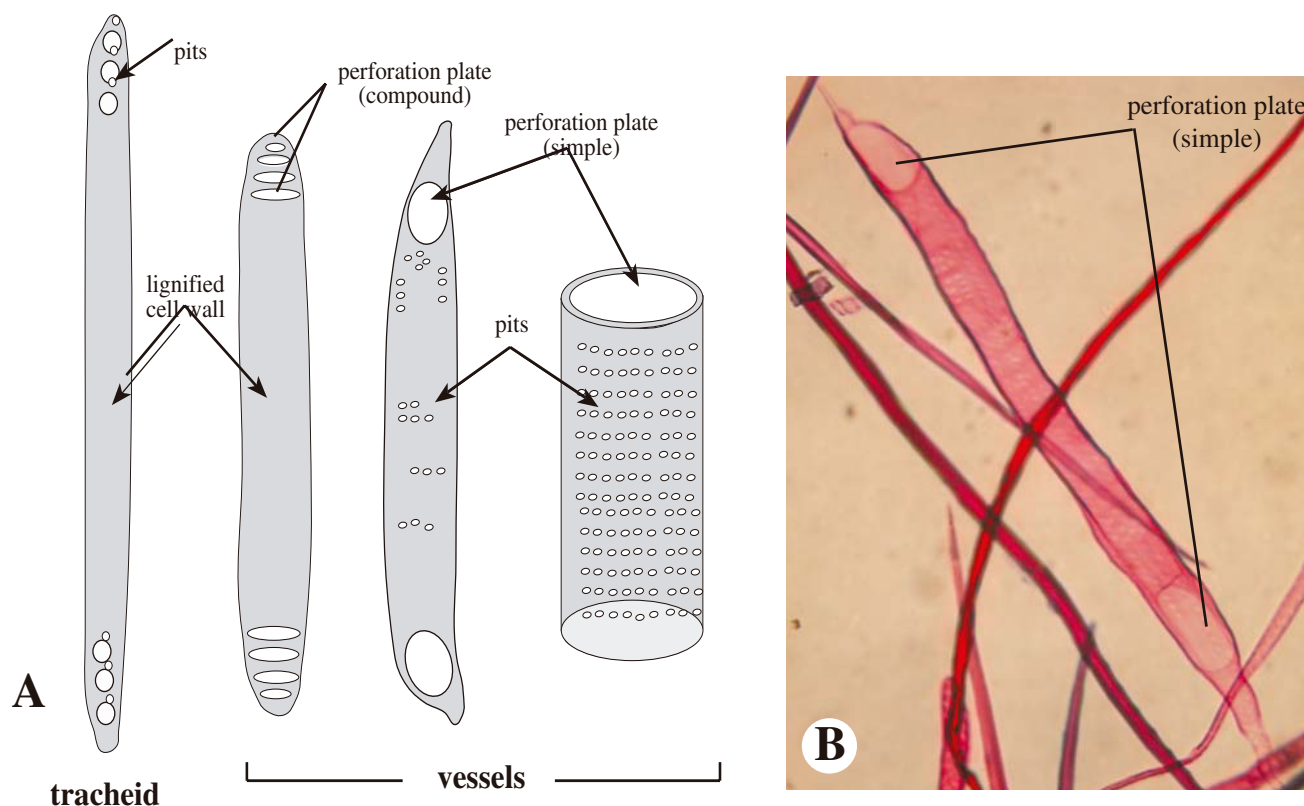


FIGURE 10.8 Tracheary elements. **A.** Diagrams of tracheids and vessel members. **B.** Photograph of vessel member, showing simple perforation plates.

and mineral nutrients flow between adjacent cells through pit-pairs (holes in the lignified 2° cell wall), in which there are intermediate primary cell walls (Figure 10.8A). **Vessel members** are perforate, meaning that there are one or more continuous holes between adjacent cells through which water and minerals may pass. (The term **vessel** refers to several vessel members attached end-to-end, forming a continuous, conductive tube.) The contact area of two adjacent vessel members is called the **perforation plate**. The perforation plate may be **compound**, if composed of several pores (pit-pairs with no primary cell walls) or **simple** if composed of a single opening (Figure 10.8A,B). Vessel members may differ considerably in length, width, angle of the end walls, and degree of perforation. As previously discussed, tracheids are the primitive type of tracheary element. Vessels are thought to have evolved from preexisting tracheids *independently* in several different groups, including a few species of *Equisetum*, a few leptosporangiate ferns, all Gnetales, and almost all angiosperms (although not always found in all plant organs).

Phloem (Gr. *phloem*, bark) is a tissue composed of specialized cells called **sieve elements** plus some parenchyma and

often some sclerenchyma. Structurally, **sieve elements** (1) are elongate cells; (2) have only a primary wall (no lignified 2° wall); (3) are semialive at maturity, losing their nucleus and other organelles but retaining the endoplasmic reticulum, mitochondria, and plastids; and (4) have specialized pores, aggregated together into **sieve areas** (Figure 10.9). Each pore of the sieve area is a continuous hole that is lined with a substance called **callose**, a polysaccharide composed of beta-1,3-glucose. (Note the different linkage from cellulose, which is a polymer of beta-1,4-glucose.) Like tracheary elements, sieve elements are oriented end-to-end, forming a tubelike continuum. Sieve elements function to conduct dissolved sugars from sugar-rich to sugar-poor regions of the plant. Sugar-rich regions include the leaves or other photosynthetic regions, where sugars are synthesized during photosynthesis, or storage roots or stems, where sugars may be produced by the hydrolysis of starch.

There are two types of sieve elements: **sieve cells** and **sieve tube members**. **Sieve cells** have only sieve areas on both end and side walls (Figure 10.9A). **Sieve tube members** have both sieve areas and **sieve plates** (Figure 10.9A C). Sieve plates consist of one or more sieve areas at the end wall

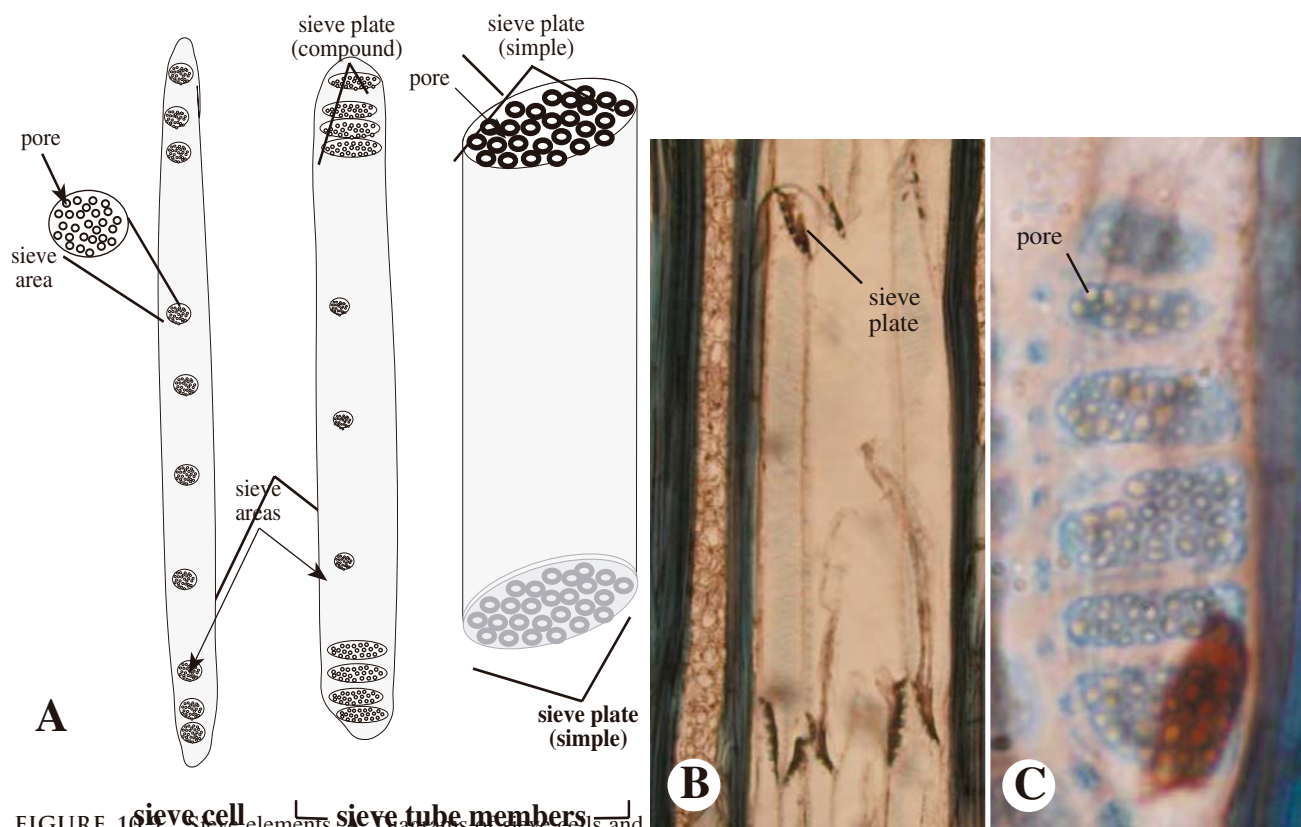


FIGURE 10.9 Sieve elements. A. Diagrams of sieve cells and sieve tube members. B. Sieve tube members, showing sieve plates at end walls. C. Close-up of compound sieve plate, showing callose-lined pores.

junction of two sieve tube members; the sieve pores of a sieve plate, however, are significantly larger than are those of sieve areas located on the side wall (Figure 10.9C). Both sieve cells and sieve tube members have parenchyma cells associated with them. Parenchyma cells associated with sieve cells are called **albuminous cells**; those associated with sieve tube members are called **companion cells**. The two differ in that companion cells are derived from the same parent cell as sieve tube members, whereas albuminous cells and sieve cells are usually derived from different parent cells. Both albuminous cells and companion cells function to load and unload sugars into the cavity of the sieve cells or sieve tube members. Sieve cells (and associated albuminous cells) are the primitive sugar-conducting cell and are found in all nonflowering vascular plants. Sieve tube members (and associated companion cells) are found only in angiosperms, the flowering plants.

Dermal tissue makes up the outer region of the plant and functions in mechanical protection of inner tissues and inhibition of water loss. Dermal tissue consists of the **epidermis** or, in woody plants, the **periderm** (see later discussion).

The **epidermis** (Figure 10.10) makes up the outermost layer of all primary plant organs. Structurally, epidermal cells (1) are usually tabular (flattened, tilelike) in shape; (2) have a cutinized (infiltrated with **cutin**, a polymer of fatty acids) or suberized (infiltrated with **suberin**) outer cell wall; (3) secrete a layer of cutin (called a **cuticle**) *outside* the cell wall; and (4) are usually living at maturity. As previously discussed, the **cuticle** (Figure 10.10) was a major innovation in the evolution of land plants, providing the primary protection from desiccation.

Specialized types of epidermal cells include stomates and trichomes. **Trichomes** (plant hairs) are cellular appendages that grow from the epidermal cells. They come in an amazing variety of shapes, sizes, and densities (see Chapter 9 for vestiture and trichome types). Trichomes may function in providing protection from UV light or herbivory; trichomes of carnivorous plants even function in digestion. **Stomates** are epidermal cells specialized for gas exchange. (See Leaf Structure and Function.)

Secretory structures are cells that secrete compounds, either internally (and stored within the cell) or externally

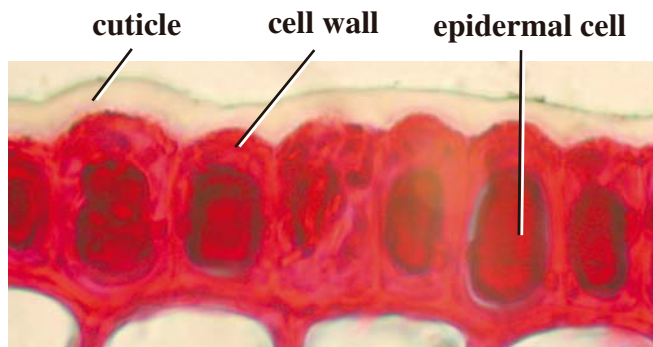


FIGURE 10.10 Epidermis, showing outer cuticle.

(outside epidermis or into a canal or duct). These include (1) **glandular** (Figure 10.11A) or **stinging** (urent) trichomes that secrete fluid to outside at tip of trichome; (2) **nectaries** (Figure 10.11B), specialized cells secreting sugar (or protein)-rich fluids to the outside that may be floral (associated with flowers as a reward for pollination) or extrafloral (often as a reward for protection); (3) **hydathodes**, which secrete excess transported water (usually due to root pressure) from leaf margins; (4) **resin/oil/mucilage** ducts or canals, which contain cells a lining of cells that secrete resin, oil, or mucilage; and (5) **laticifers** (Figure 10.11C), cells located in the periphery of some tissues that secrete and store latex. The last two may function both to deter herbivory and to seal and protect plant tissue upon wounding.

ROOT STRUCTURE AND FUNCTION

Roots are plant organs that function in anchorage and in absorption of water and minerals. Roots are found in the sporophytes of all land plants except for the nonvascular liverworts, hornworts, and mosses (in which the sporophytes are attached to the gametophytes) and the psilophytes (e.g., *Psilotum*). Land plants lacking roots generally have

uniseriate (one-cell-thick), filamentous **rhizoids** that assume a similar function.

The first root to develop, in the embryo, is termed the **radicle**. If the radical continues to develop after embryo growth, it is known as the **primary root**. Additional roots may arise from internal tissue of either another root, the stem/shoot (often near buds), or (rarely) a leaf. Roots that arise from other roots are called **lateral roots**. Roots that arise from a nonroot organ (stem or leaf) are called **adventitious roots**. Various modifications of roots have evolved, such as storage roots, aerial roots, fibrous roots, tap roots, contractile roots, haustoria, prop roots, and pneumatophores (see Chapter 9).

Roots, like shoots, develop by the formation of new cells within the actively growing **apical meristem** of the root tip (Figures 10.5B, 10.12A), a region of continuous mitotic divisions. At a later age (and further up the root) these cell derivatives elongate significantly. This cell growth, which occurs by considerable expansion both horizontally and vertically, literally pushes the apical meristem tissue downward. Even later in age and further up the root, the fully grown cells differentiate into specialized cells.

Roots can be characterized by several anatomical features. First, the apical meristem is covered on the outside by a **root-cap** (Figure 10.5B, 10.12A). The rootcap functions both to protect the root apical meristem from mechanical damage as the root grows into the soil and to provide lubrication as the outer cells slough off. Second, the epidermal cells proximal to the root tip develop hair-like extensions called **root hairs** (Figure 10.12A; see Figure 4.10A); root hairs function to *greatly* increase the surface area available for water and mineral absorption. Third, roots have no exogenous (externally developing) organs; all **lateral roots arise endogenously** from the internal tissues of the root. Lateral roots grow from cell divisions within the **pericycle**, a cylindrical layer of parenchyma cells located just inside the endodermis itself.

Two other features of roots may or may not distinguish them from stems, as the stems of some land plants are very

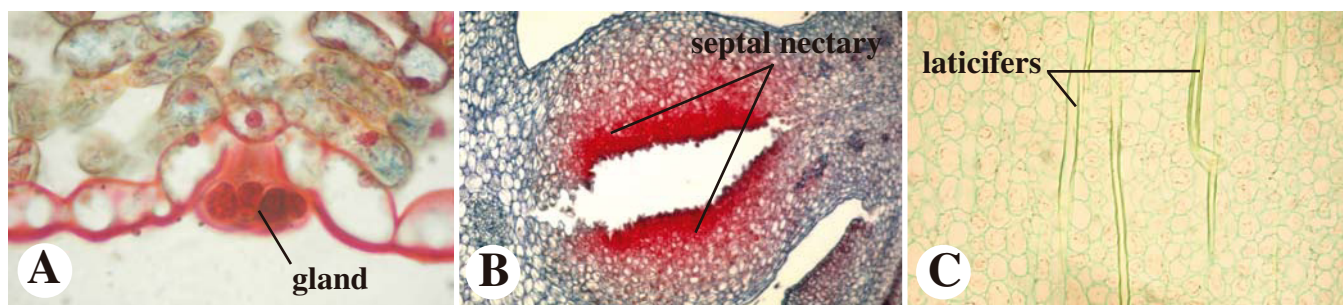


FIGURE 10.11 A. Secretory gland, on leaf surface. B. Septal nectary, the cells of which secrete sugar-rich nectar into cavity. C. Laticifers.

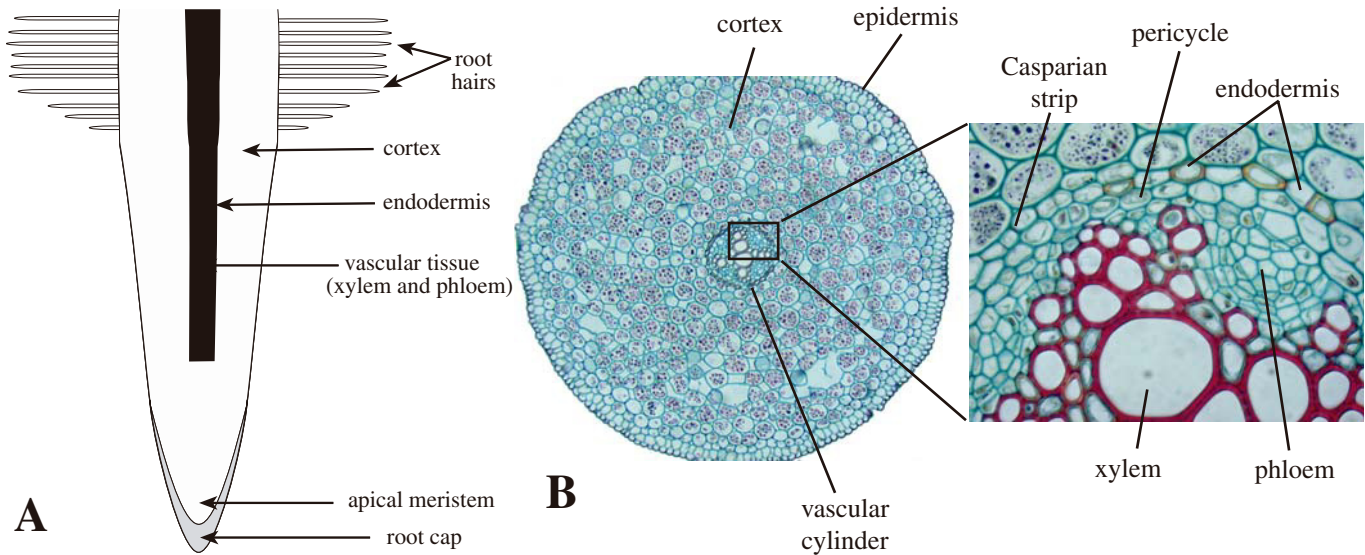


FIGURE 10.12 Root anatomy. **A.** Root longitudinal section. **B.** Root cross-section, close-up at right.

similar in these features to roots. All roots have a central **vascular cylinder** of xylem and phloem. Often, ridges of xylem alternate with cylinders of phloem (i.e., the **xylem and phloem are on alternate radii**). As with stems, the mostly parenchymatous region between the vasculature and epidermis is called the **cortex**; the center of the vascular cylinder, if vascular tissue is lacking there, is called a **pith**. In addition, the vascular tissue of all roots is surrounded by a special cylinder of cells known as the **endodermis**. In the general region of the root hairs, where most absorption takes place, each cell of the endodermis has a **Casparian strip**, which is a tangential band of suberin that infiltrates the cell wall (Figure 10.13). As discussed in Chapter 4, the Casparian strip functions as a water-impermeable binding material to the plasma membrane of the endodermal cells. This forces absorbed water and nutrients to flow through the endodermal plasma membrane, as opposed to within the intercellular spaces (between the cells or through the cell wall). The function of the endodermis and Casparian strips is to allow selectivity as to what mineral nutrients are and are not absorbed by the plant; e.g., toxic minerals may be selectively excluded.

Some anatomical specializations of roots are found in certain taxa. For example, the aerial roots of many Orchidaceae and Araceae lack root hairs and have a specialized, multilayered epidermis called a **velamen**. The velamen may function in protection, prevention of water loss, or water and mineral absorption.

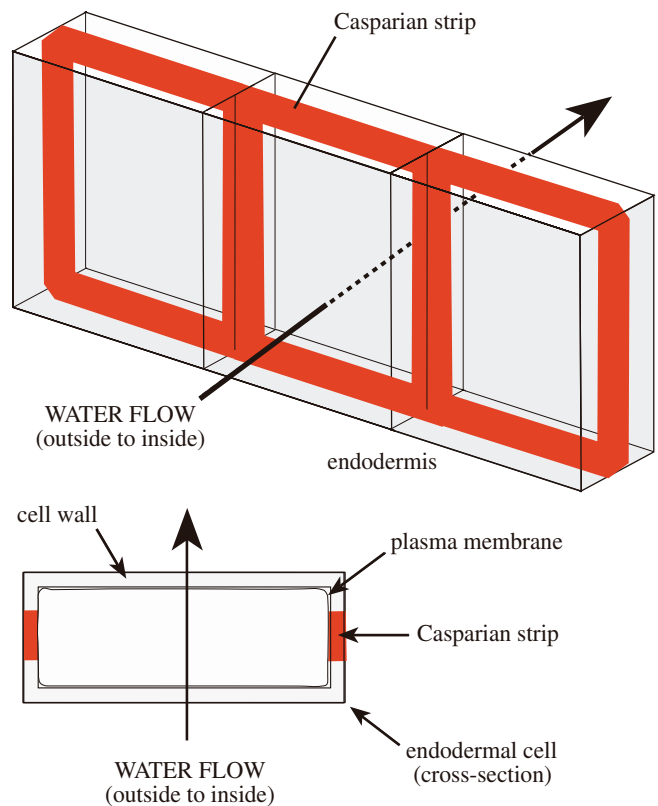


FIGURE 10.13 The Casparian strip, a specialized feature of cells of the endodermis.

SHOOT AND STEM STRUCTURE AND FUNCTION

A **shoot** is a stem plus its associated leaves. Sporophytic shoots are an apomorphy for all vascular plants. The leafy shootlike structures of mosses and some liverworts are gametophytic and not homologous with shoots of vascular plants.

The first shoot of a vascular plant develops from the **epicotyl** of the embryo. The epicotyl elongates after embryo growth into an axis (the stem), which bears leaves from its outer surface. The tip of a shoot contains the actively dividing cells of the **apical meristem** (Figure 10.14). As in the root, these cells undergo continuous mitotic divisions. A bit down from the apical meristem, the cells undergo considerable expansion, literally pushing the cells of the apical meristem upward (or forward). Proximal to the shoot tip, the fully expanded cells differentiate into specialized cell types.

Slightly down from the apical meristem region, the outermost cell layers of a shoot begin to repeatedly divide. Further cell divisions and growth result in the development of a mass of tissue that forms an immature leaf, the **leaf primordium**. **Vascular strands** run between stem and leaf, providing a connection for fluid transport (Figures 10.14, 10.15). As the shoot matures, the leaves fully differentiate into an amazing variety of forms. A bit later in development, the tissue at the upper junction of leaf and stem (called the **axil**) begins to divide and differentiate into a **bud primordium**. The bud primordium matures into a **bud**, defined as an immature shoot system. Buds have an architecture identical to that of

the original parent shoot. Buds may develop into a vegetative, lateral branch or may terminate by developing into a flower or inflorescence.

Stems function both as supportive organs (supporting and usually elevating leaves and reproductive organs) and as conductive organs (conducting both water/minerals and sugars through the vascular tissue between leaves, roots, and reproductive organs). Stems can be distinguished from roots in at least three ways. First, the apical meristem of stems is not covered by an outer protective layer (like the root cap; Figure 10.14). Second, the epidermal cells of the stem do not form structures resembling root hairs. However, the epidermal cells of stems and leaves may divide and differentiate into separate, one-to-many-celled **trichomes**, described earlier (see also Chapter 9). Third, **stems bear leaves exogenously**; no organs are born endogenously (except in cases of adventitious roots potentially arising from the internal parenchyma cells of stems).

Stems, particularly underground stems, may possess an endodermis similar to that of roots in structure and function. The aerial stems of many plants lack an endodermis. Numerous modifications of stems and shoots have evolved, such as bulbs, corms, caudices, rhizomes, stolons (=runners), cladodes, pachycauls, and thorns (see Chapter 9).

The primary vasculature of stems is organized into arrangements of xylem and phloem known as **steles** (Figure 10.16). In some groups of non-seed vascular plants, such as the lycophytes, the stem stelar type is a **protostele** (Figures 10.15A, 10.16A), in which there is a central cylinder of vascular tissue.

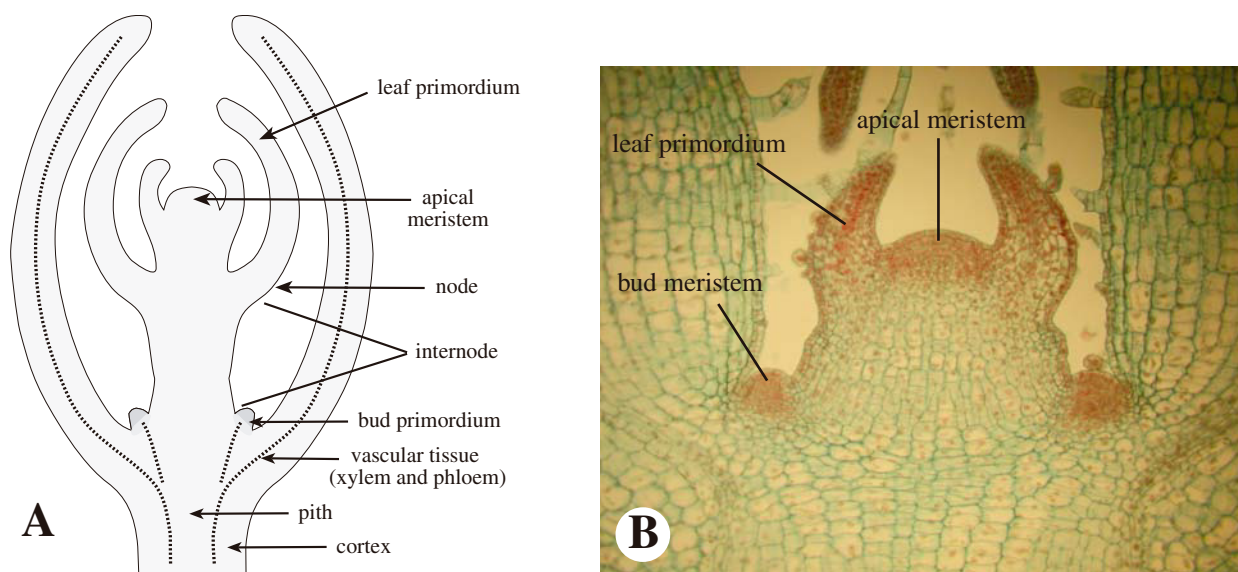


FIGURE 10.14 Shoot longitudinal section. **A.** Diagram. **B.** Photograph, *Plectranthus* [*Coleus*] shoot.

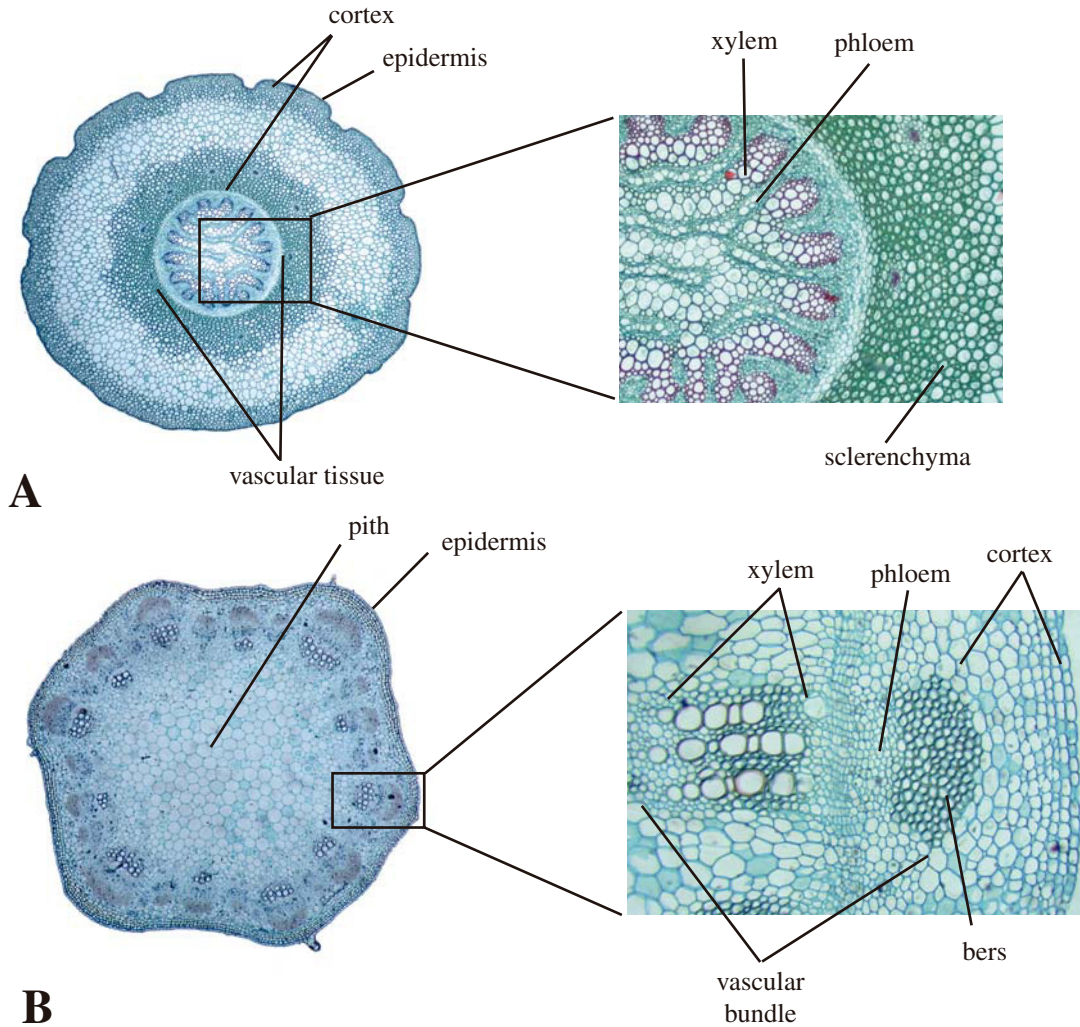


FIGURE 10.15 Stem anatomy. **A.** *Lycopodium* stem cross-section, showing close-up of vascular tissue, with xylem, phloem, and outer fibers. **B.** *Helianthus* stem cross-section, showing close-up of vascular bundle, with xylem, phloem, and associated fibers.

Protosteles are the most ancestral type of stem vasculature, one that most resembles the vasculature of a root. The vasculature of fern stems is typically a **siphonostele**, in which a ring of xylem is surrounded by a continuous layer of phloem (Figure 10.16B,C). The stems of seed plants contain discrete **vascular bundles** in which xylem and phloem are grouped together along a common radius, usually with xylem to the inside and phloem to the outside, a type known as a **collateral** vascular bundle. (In some angiosperms the stem vascular bundles have phloem to both the inside and outside of the xylem, a type known as **bicollateral**.) These collateral vascular bundles may be organized as a single ring, known as a **eustele** (Figures 10.15B, 10.16D). The eustele is an apomorphy for many seed plants, including all that are extant. For both siphonosteles and eusteles, the central region of tissue in

the stem is called **pith**; the region between the vasculature and the outer epidermis is called the **cortex**. Stems of monocots (of the angiosperms) have a modification of the eustele called an **atactostele** (Figure 10.16E). The atactostele, which represents an apomorphy for the monocots, consists of numerous, collateral vascular bundles positioned throughout the stem tissue (appearing scattered but actually having a precise and complex disposition). In an atactostele, there is no pith; the region of tissue between vascular bundles is called **ground meristem**. Vascular bundles typically are associated with sclerenchyma fibers, which may surround the entire bundle or occur in outer patches called **bundle caps** (e.g., Figure 10.15B). Parenchyma, collenchyma, or sclerenchyma cells make up the tissues of the pith, cortex, and ground meristem.

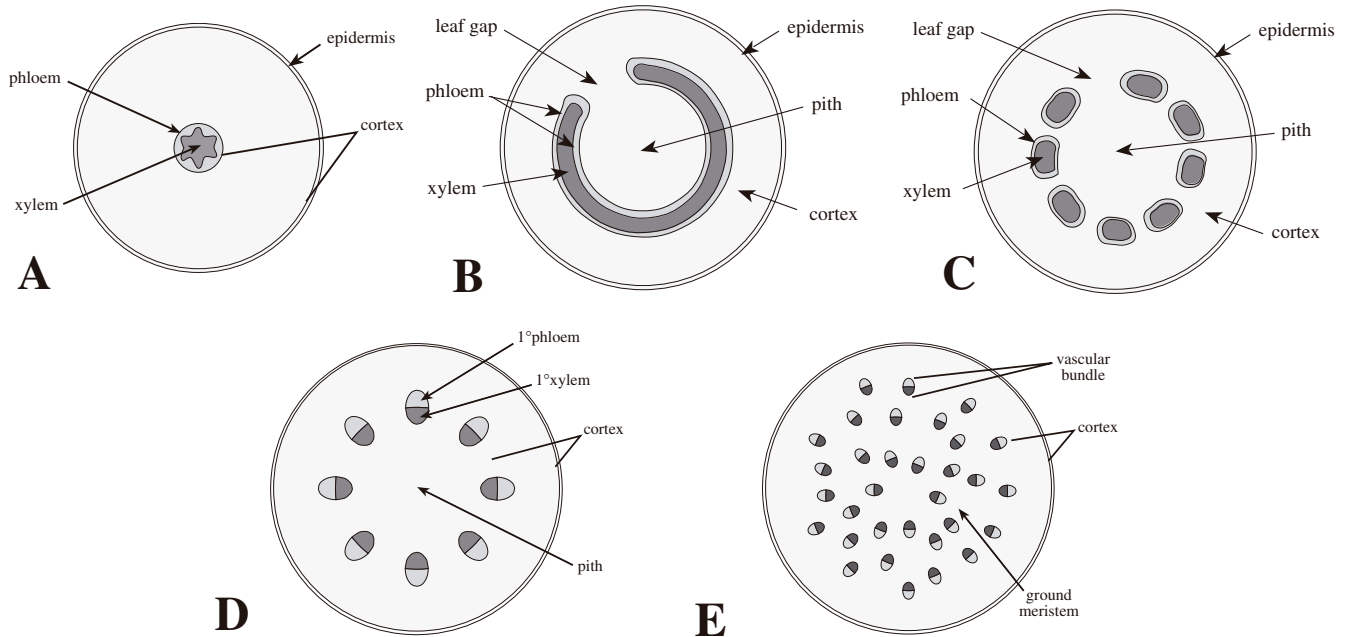


FIGURE 10.16 Stellar types in Vascular Plants. Shading: black=xylem; stippled=phloem. **A.** Protostele. **B,C.** Siphonostele, that at **C** termed a dictyostelic siphonostele. **D.** Eustele. **E.** Atactostele.

The vasculature of stems (and roots) can vary also with respect to the sequence of maturation of tracheary elements in the xylem. **Protoxylem** refers to the first xylem that matures in a group of vascular tissue; protoxylem cells are often smaller in diameter. **Metaxylem** is the xylem that develops later and usually consists of larger-diameter cells. Three general types of protoxylem orientation are recognized: (1) **exarch**, in which the protoxylem is oriented toward the outside relative to metaxylem, as occurs in some protosteles (Figure 10.17A); (2) **endarch**, in which the protoxylem is oriented toward the center of the stele, relative to the metaxylem,

as occurs in eusteles and atactosteles (Figure 10.17B); and (3) **mesarch**, in which the protoxylem is surrounded by metaxylem within the vascular tissue, as can occur in siphonosteles.

SECONDARY GROWTH

In vascular plants, the growth in height or length of a stem or root is brought about by the elongation and differentiation of cells derived from the apical meristem. This is termed

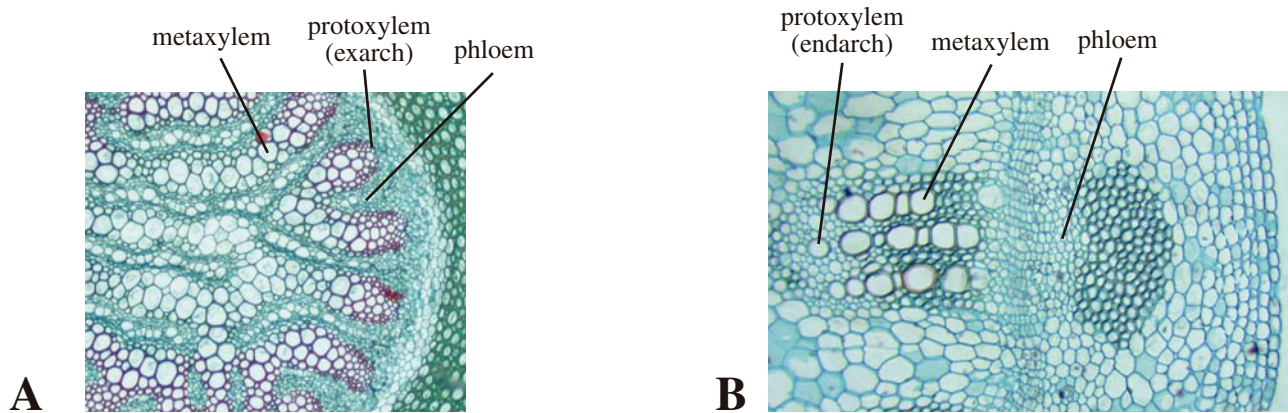


FIGURE 10.17 **A.** *Lycopodium* stem, showing exarch protoxylem development of protostele. **B.** *Helianthus* stem, showing endarch protoxylem development of eustele.

primary growth, and the tissues formed by primary growth are called **primary tissues** (e.g., as in primary xylem or primary phloem). However, in many seed plants, roots and stems may grow in girth or width by means of cells produced not from the apical meristems, but from **lateral meristems**. This process is termed **secondary growth**, and the tissues formed by secondary growth are called **secondary tissues**.

Two types of lateral meristems function in secondary growth: the **vascular cambium** and the **cork cambium**. These lateral meristems represent apomorphies for the woody plants, including all extant seed plants plus several fossil groups (although lateral meristems have been lost in some angiosperms, most notably the monocots). The **vascular cambium** is a cylindrical sheath of cells that typically forms by cell divisions of undifferentiated parenchyma cells. In eustelic stems the vascular cambium forms from parenchyma

cells both between the primary xylem and phloem of vascular bundles and in the adjacent region between the bundles (Figures 10.18,10.19). In woody roots the vascular cambium develops from parenchyma cells between xylem and phloem and from the adjacent pericycle. The cells of the vascular cambium divide more or less synchronously, and mostly in a tangential plane, the initial result being the formation of two layers of cells (Figure 10.18). One of these layers continues as the vascular cambium and divides indefinitely; the other layer eventually differentiates into either **secondary xylem** (=wood), if produced to the inside of the cambium, or **secondary phloem**, if produced to the outside of the cambium (Figures 10.18,10.20A). Generally, much more secondary xylem is produced than is secondary phloem. As the secondary tissue is formed, the inner cylinder of wood expands. As this growth in girth continues, some cells of the

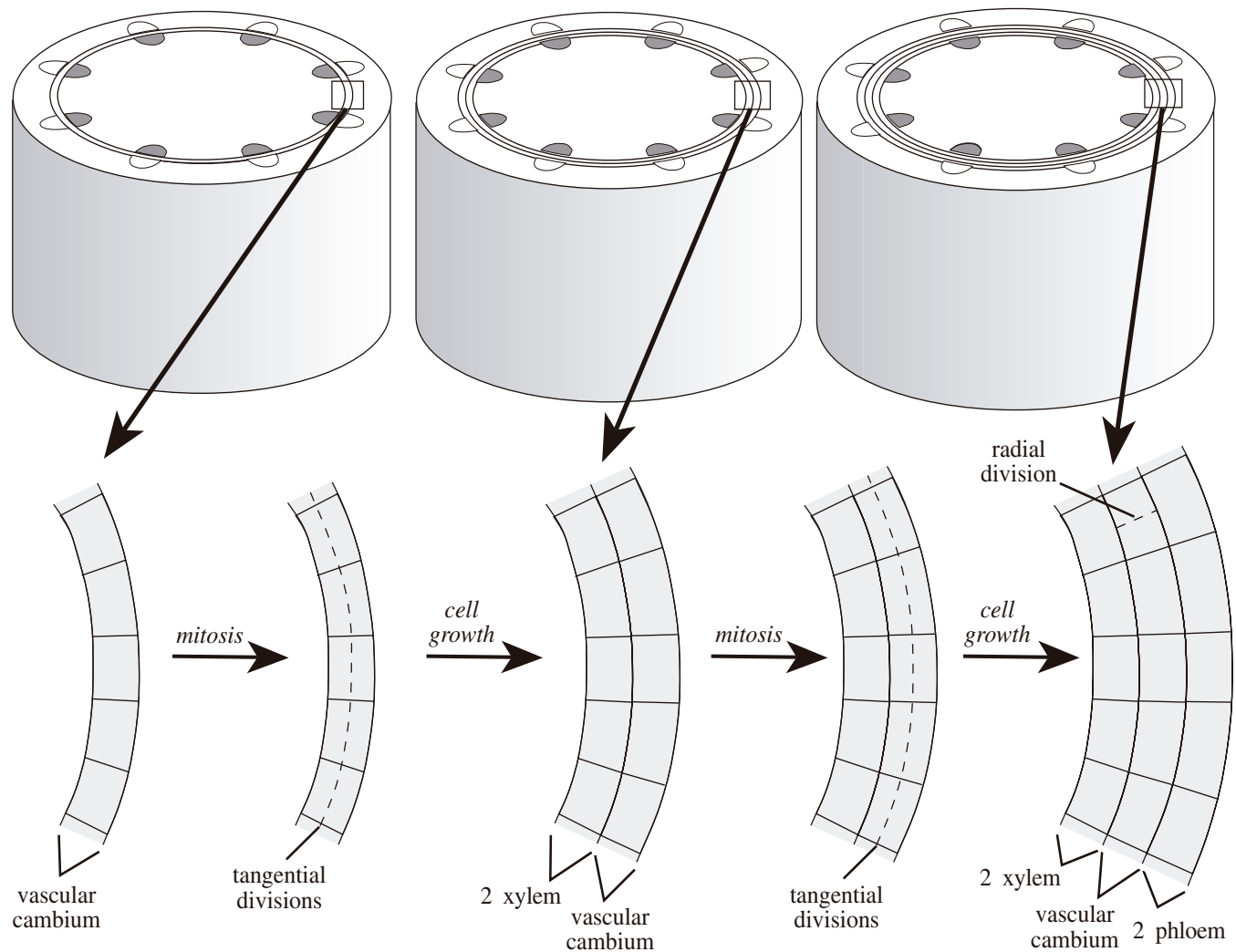


FIGURE 10.18 The vascular cambium.

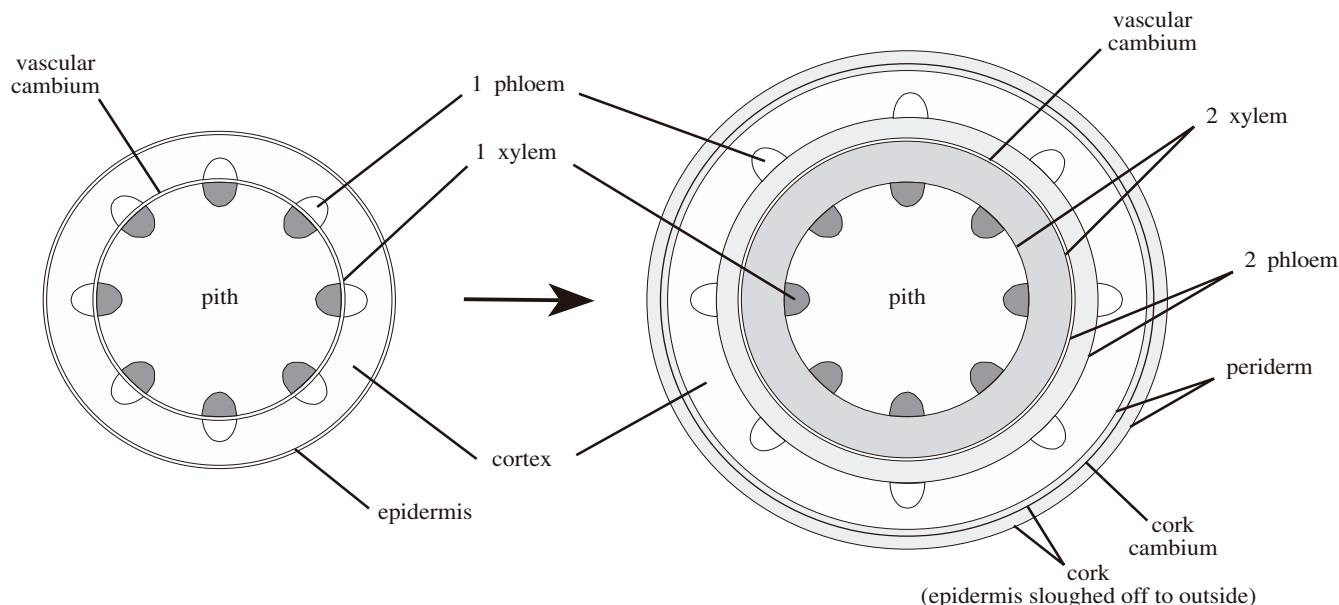


FIGURE 10.19 Secondary growth in stems.

vascular cambium undergo radial divisions (parallel to a radius), enabling the vascular cambium to grow larger in circumference.

The **cork cambium** is similar to the vascular cambium, only it differentiates near the periphery of the stem or root axis. The cork cambium forms **cork** to the outside and **phelloderm** to the inside, the latter usually much thinner (Figure 10.20G). The cork cambium and all of its derivatives constitute the **periderm**. The outer cork cells contain a waxy polymer called **suberin** (chemically related to cutin), which is quite resistant to water loss. In the wood industry, the term **inner bark** refers to all the tissue between the vascular cambium and the periderm (including all of the secondary phloem). **Outer bark** is equivalent to the periderm.

The vascular cambium and cork cambium are of significant adaptive value. Secondary xylem (wood) functions in structural support, enabling the plant to grow tall and acquire massive systems of lateral branches. Thus, the vascular cambium was a precursor to the formation of intricately branched shrubs or trees with tall overstory canopies, a significant ecological adaptation. Cork produced by the cork cambium functions as a thick layer of dermal tissue cells that protects the delicate vascular cambium and secondary phloem from mechanical damage, predation, and desiccation.

Secondary xylem, or **wood**, consists mostly of longitudinally oriented tracheary elements, either tracheids (in cycads, *Ginkgo*, and conifers, excluding Gnetales) or vessels (Gnetales and almost all angiosperms; Figures 10.18 10.21). Other longitudinally oriented cells may include **fibers** and

axial parenchyma. The vascular cambium also forms cells that are radially oriented (parallel with a stem or root radius). These radially oriented cells occur in bandlike strands called **rays** (Figure 10.19B E); their function is lateral translocation of water, minerals, and sugars.

In most woody plants with regular, annual growth seasons (in temperate regions caused by seasonal cold, in tropical regions by seasonal drought), the vascular cambium and cork cambium actively divide only near the start of the growing season; further secondary growth is delayed until the next growing season. As a result of this periodic growth, there are differences in the structure of the secondary xylem from the first part of the growing season (**spring wood**) versus the latter part of the growth season (**summer wood**). For example, the tracheary elements of spring wood tend to be larger in diameter with thinner walls; those of summer wood tend to be smaller in diameter with thicker walls (Figure 10.20C,D). The overall result of this discrepancy in structure between spring and summer wood results in the formation of **annual rings** (Figures 10.20A,B, 10.21A,B). Each annual ring represents the accumulation of secondary xylem (or phloem) over a single growing season. Annual rings are evident because of the structural difference between the last cells of the summer wood and the first cells of the subsequent spring wood.

Wood may be cut in three major planes: transverse (cross-sectional), radial (longitudinal and ca. parallel to a stem radius), or tangential (longitudinal and perpendicular to a stem radius); these planes of section are often abbreviated **X**, **R**, **T**, respectively (Figure 10.22). These different cuts

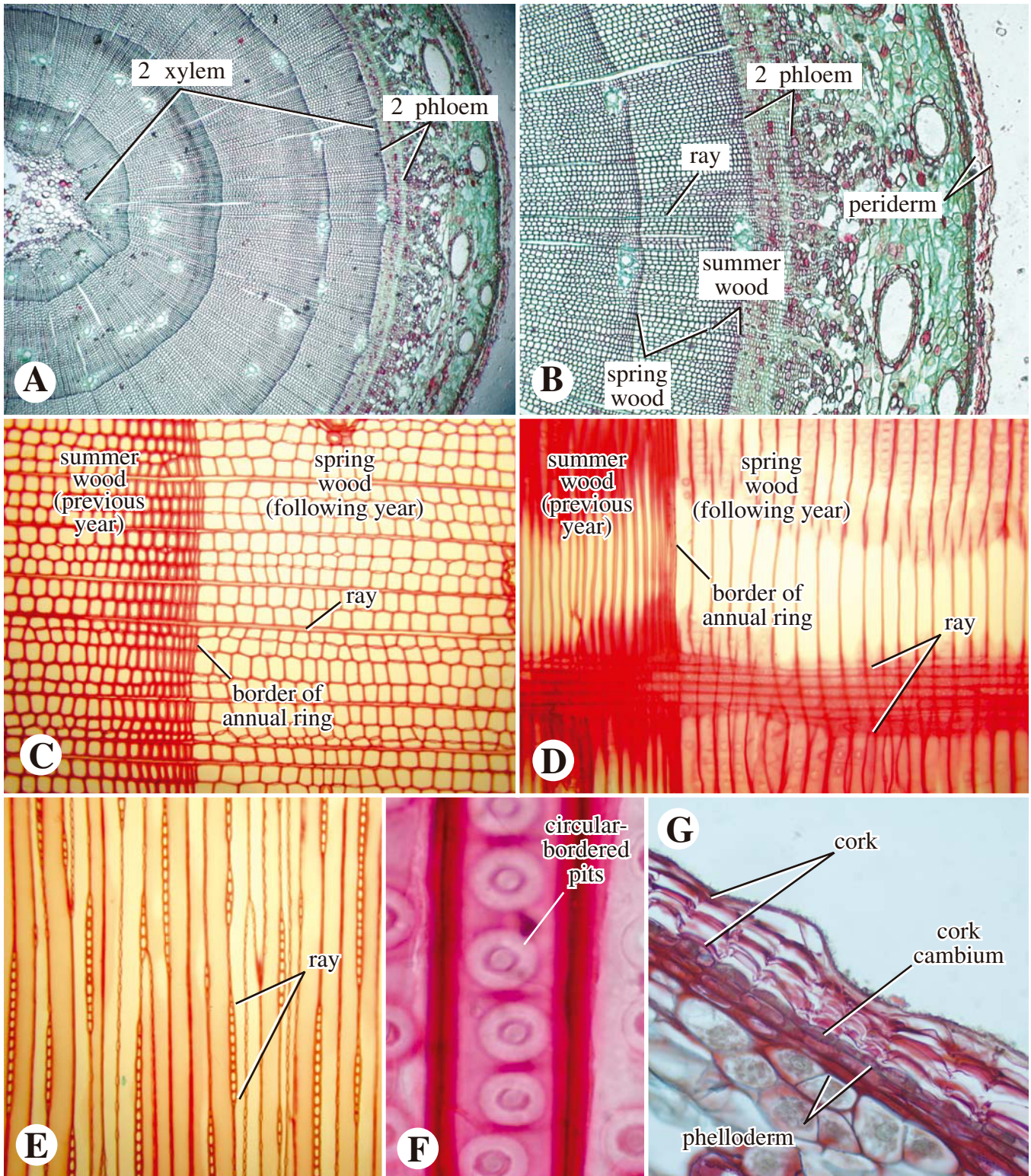


FIGURE 10.20 Conifer wood. **A,B.** *Pinus* sp. stem cross-section, showing 3 years' growth. **C–E.** *Libocedrus decurrens* wood sections. **C.** Transverse- or cross-section, showing junction of summer wood (of previous year) and spring wood (of following year). **D.** Radial-section, showing transverse ray and border of annual ring (summer wood to left; spring wood to right). **E.** Tangential section, showing vertical tracheids and rays. **F.** *Pinus*, circular bordered pits of tracheids (radial face). **G.** *Pinus* periderm, showing cork cambium, phelloderm, and cork.

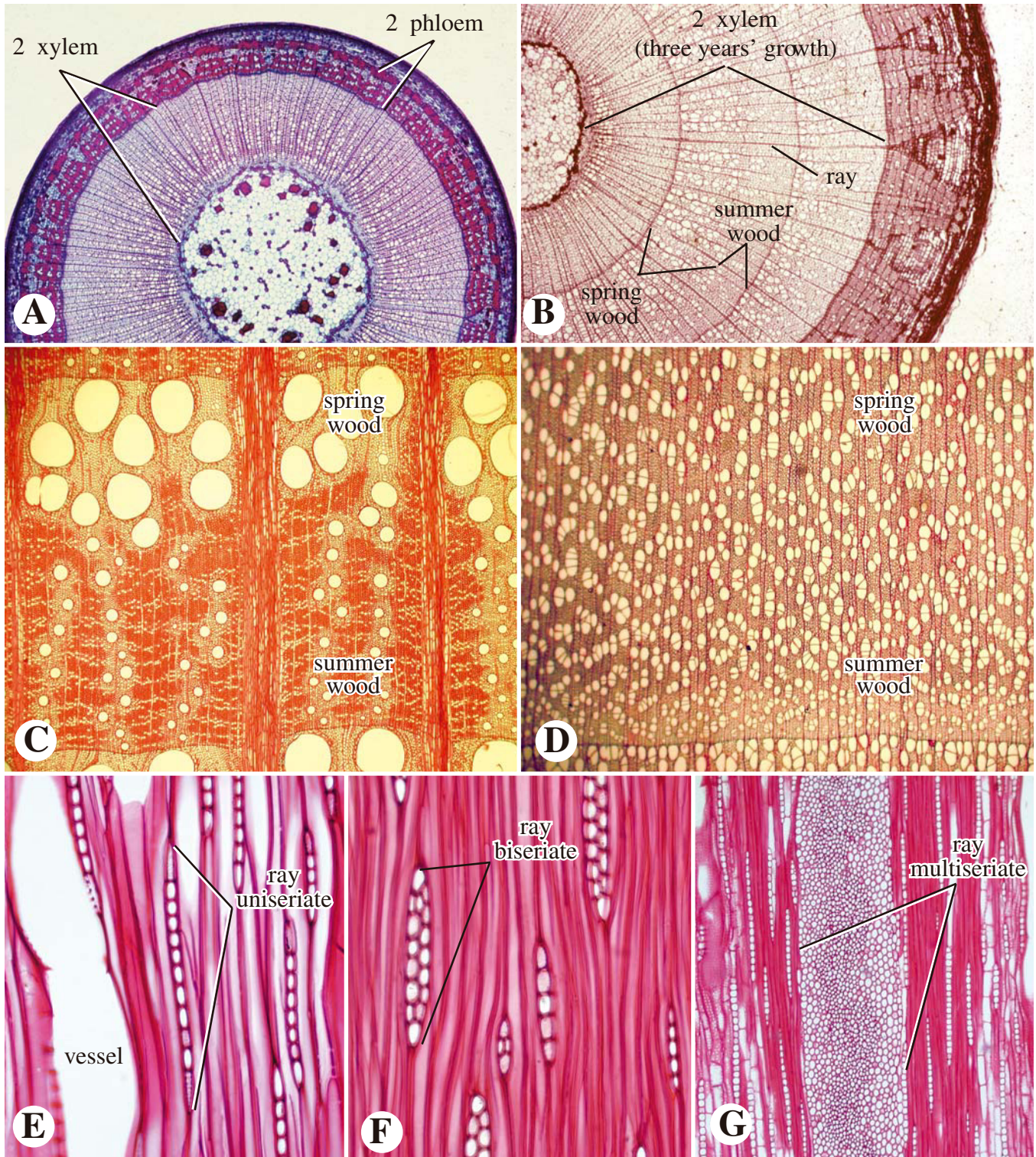


FIGURE 10.21 Eudicot wood. **A,B.** Woody stem cross-section, *Tilia*. sp. **A.** One year's growth. **B.** Three years' growth. Note rays and rings with spring and summer wood. **C.** Ring-porous wood, *Quercus*, with vessels much larger in spring wood (above), smaller to absent in summer wood (below). **D.** Diffuse-porous wood, *Salix*, having vessels evenly distributed in annual ring. **E,F.** Ray types. **E.** Uniseriate rays. **F.** Biseriate rays (some uniseriate also present). **G.** Multiseriate rays (some uniseriate also present).

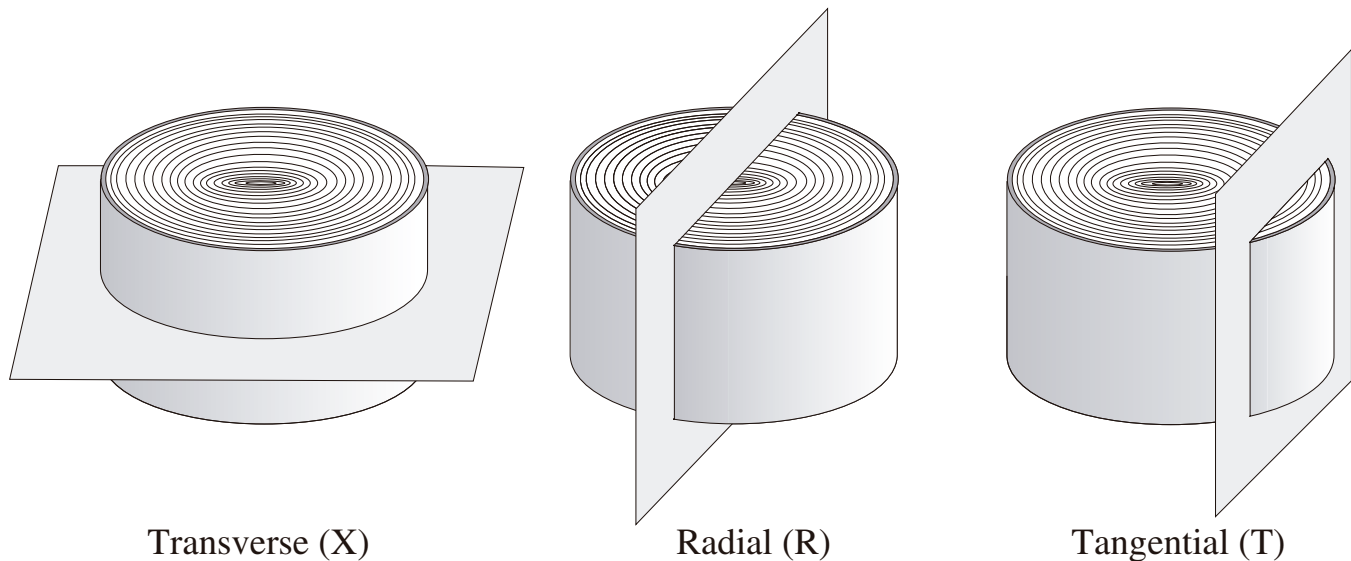


FIGURE 10.22 Wood sections: transverse (X), radial (R), and tangential (T).

are used for different purposes in the wood industry and influence the *figure*, or general appearance of the wood. The three cuts are also used by wood anatomists to view the cells from three different directions, often necessary for precise description or identification of wood samples (see Figures 10.20, 10.21).

In the wood industry, the term **softwood** is used for a wood product derived from a conifer and **hardwood** is used for one derived from a nonmonocotyledonous angiosperm. Softwoods from conifers (such as pine) are indeed usually softer and easier to work with than hardwoods (such as oak), as the latter typically contain numerous wood fiber cells. However, there are exceptions; some so-called hardwoods, such as balsa, are quite soft.

Wood anatomy may be very complex. Its structure may provide several characters that may be of systematic importance; these characters include tracheary element type (whether having only tracheids, termed **nonporous**, or having vessels, termed **porous**), tracheary element anatomy (size, shape, and pit or perforation plate structure), distribution of vessels (if present), ray anatomy, presence of resin ducts, distribution of axial parenchyma, and presence/distribution of fibers or fiberlike cells. For example, in some angiosperms there may be differences in the formation of vessel elements associated with the annual rings. The vessels may form only in the spring wood, with summer wood either lacking or having relatively small vessels and usually containing mostly fibers; this type of growth is called **ring-porous** (the term *porous* referring to the presence of **pores**, the vernacular term for vessels; Figure 10.21C). The alternative, in which

vessels develop more or less uniformly throughout the growth season, is called **diffuse-porous** (Figure 10.21D). Another feature of systematic importance is ray anatomy. Rays can be **uniseriate** (with a single, vertical row of cells, as in Figure 10.21E), **biseriate** (with two vertical rows of cells, as in Figure 10.21F), or **multiseriate** (with many vertical rows of cells, as in Figure 10.21G). Wood anatomical characters may be useful in phylogenetic inference and are valuable for microscopic identification of the species.

Some aspects of wood anatomy are ecologically significant. In fact, wood (both extant and fossil) may be used to trace the history of climatic conditions in a given region. This field of study is called **dendrochronology**. When growth conditions are good (e.g., high rainfall), annual rings will be wide; when conditions are poor, they will be narrow. By correlating the width of annual rings with time, assessment of past conditions may be made, e.g., cycles of cold or drought and even sunspot cycles.

LEAF STRUCTURE AND FUNCTION

Leaves are the plant organs that function primarily in photosynthesis. However, leaves or leaflike homologs have been co-opted for innumerable other functions in plants.

As discussed earlier, leaflike structures occur on the gametophytes of mosses and leafy liverworts. However, sporophytic leaves evolved first in the vascular plants; thus, *leaf* is here equated with sporophytic leaf. True leaves evolved with the development of a continuous strand of

vascular tissue running from the stem into the leaf. As discussed in Chapter 4, the first sporophytic leaves, having only a single, generally unbranched vein and lacking a leaf gap, are called **lycophylls** (essentially equivalent to microphylls); lycophylls also possess an intercalary meristem (at the proximal side of the leaf base; Figure 10.23). Among extant plants lycophylls are restricted to the lycophytes. A major innovation occurred in the monilophytes and lignophytes/seed plants, the evolution of **euphylls** (essentially synonymous with megaphyll, a more traditional term). **Euphylls** are leaves that (1) have multiple, branched vascular strands in the leaf blade; (2) have a **leaf gap**, in which parenchymatous tissue replaces vascular tissue in the region just distal to the point of departure of the vasculature from stem to leaf (Figure 10.23); and (3) grow by means of either marginal or apical meristems. The evolution of megaphylls allowed for a much bigger, broader, morphologically diverse leaf structure. This has undoubtedly been adaptive in several habitats, permitting, e.g., maximization of photosynthetic rate.

Leaves have a characteristic development and structure (Figure 10.24). As previously discussed, leaves arise as **leaf primordia** on shoots. During development, the cells along the margins generally divide more rapidly, resulting in a flattened, dorsiventral structure with an upper (adaxial) and lower (abaxial) surface. Thus, leaves have both an **upper epidermis** and **lower epidermis**. The cuticle, which is an

apomorphy of all land plants, is often quite thickened on leaf epidermal cells. As discussed in Chapter 3, the **stomate** was a major innovation in the evolution of land plants. Stomates consist of two chlorophyllous **guard cells** with an opening or **stoma** between (Figure 10.25A,B). The guard cells can alter turgor pressure by changes in ion gradients, which results in opening or closing the stoma. In vascular plants, stomates occur mostly on the leaves, and there they are predominate on the lower (abaxial) surface. Stomates function to regulate gas exchange. An open stomate permits carbon dioxide to enter the leaf, and oxygen and water to exit. Stomates are the only epidermal cell to have chloroplasts (which function in regulation of the stoma). Stomates are often associated with **subsidiary cells**, specialized epidermal cells that are contiguous with the stomate and that may function in ion exchange and therefore stomate opening and closing. The number, size, and placement of subsidiary cells varies between taxa and can be a useful anatomical systematic character (Figure 10.25C E).

The nonvascular cells located between upper and lower epidermal layers comprise the **mesophyll** (Figure 10.24). The mesophyll is composed primarily of chlorophyllous cells, the chloroplast-containing parenchyma cells that function as the site of photosynthesis. Typically (but not always), they are of two morphological types: (1) columnar **palisade mesophyll** cells, which occur in the upper (adaxial) region and have relatively small intercellular spaces; and (2) irregularly

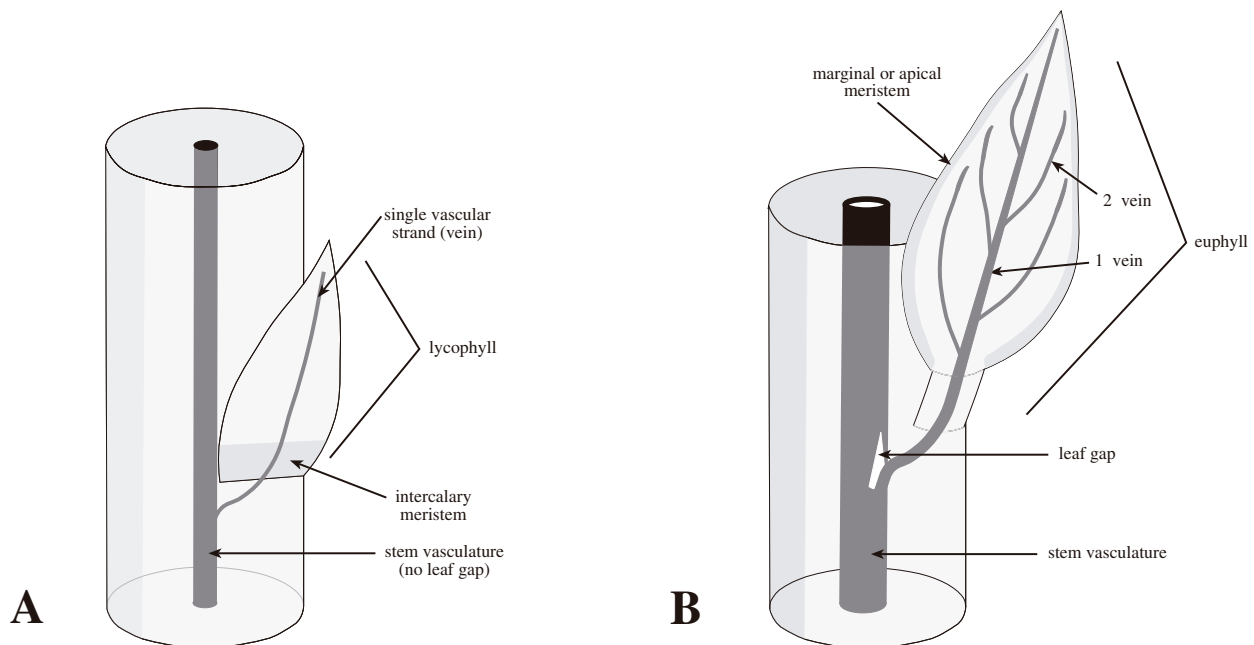


FIGURE 10.23 A. Lycophyll. B. Euphyll. Note leaf gap and branched venation.

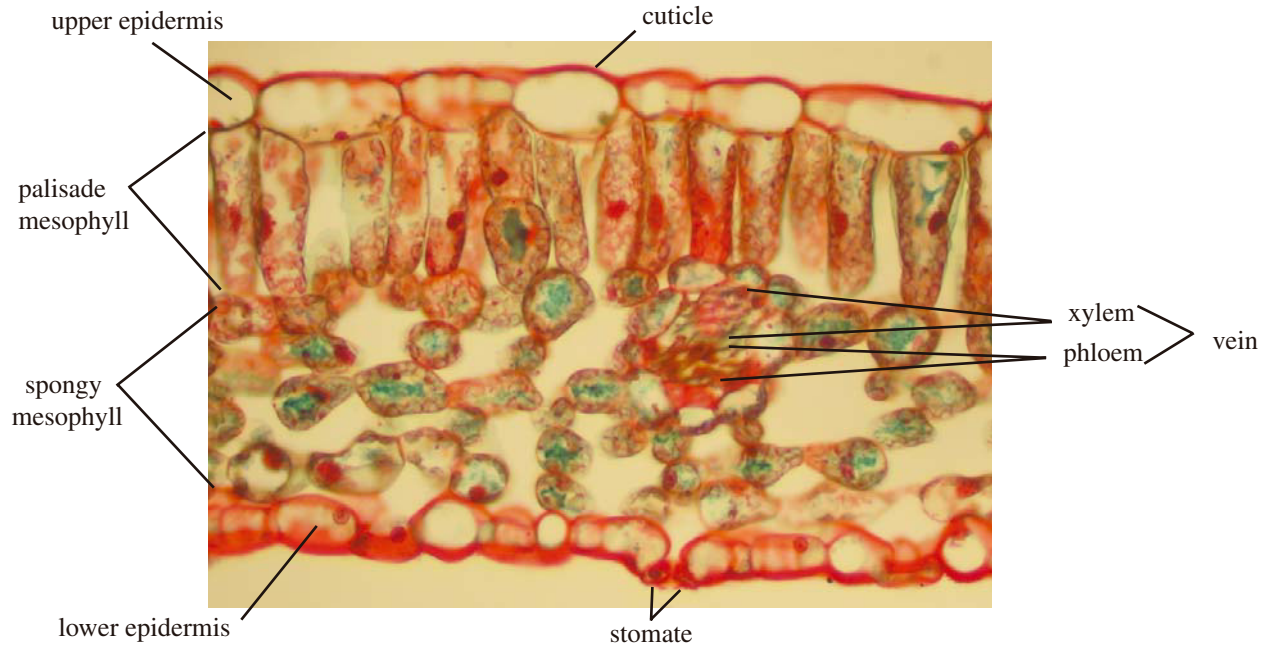


FIGURE 10.24 Cross-section of a typical vascular plant leaf.

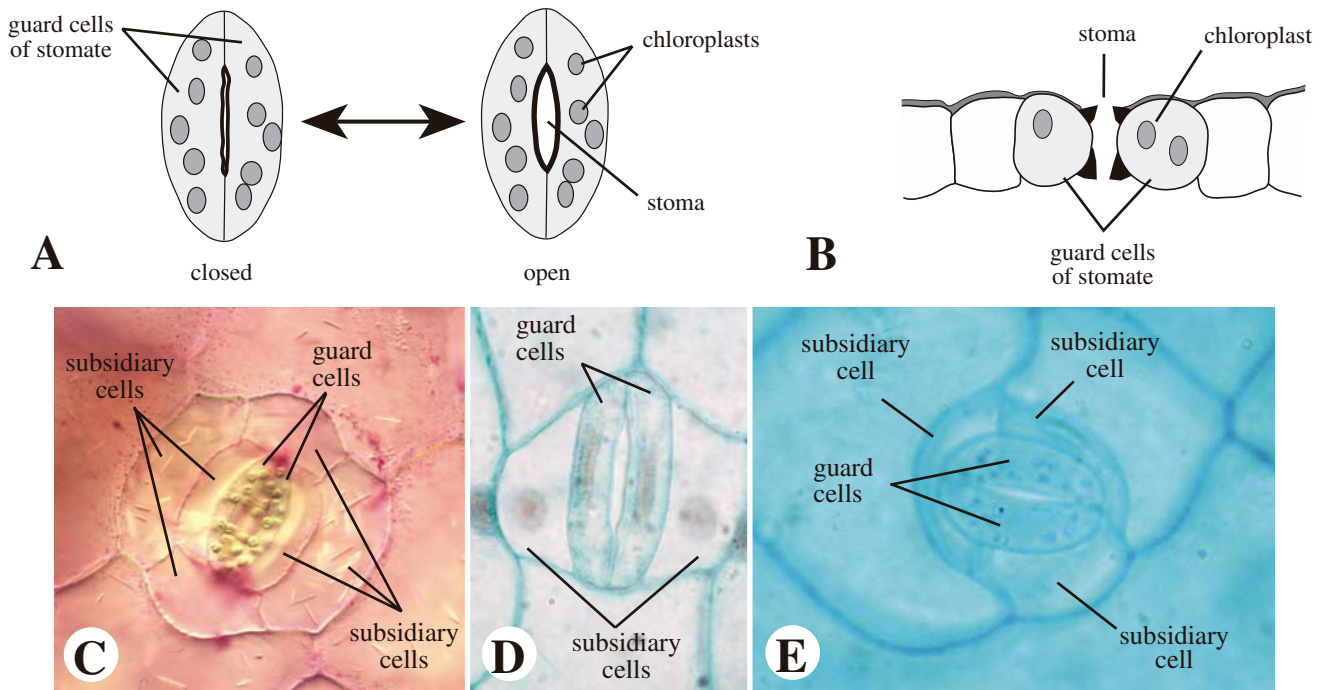


FIGURE 10.25 Stomata. **A**. Illustration of stomate in face view, closed and open. **B**. Stomate in cross-sectional view. **C–E**. Stomates of various taxa, showing the differences in subsidiary cells.

shaped **spongy mesophyll** cells, which occur in the lower (abaxial) region and have large intercellular spaces. The **veins** of a leaf have the anatomy of typical vascular bundles. In almost all veins the xylem is oriented to the adaxial side, phloem to the abaxial side, corresponding to their orientation in the stem. Veins may very often have a ring of cells surrounding the xylem and phloem called a **vascular bundle sheath**. This sheath may be composed of fiber cells, which function in structural support of the vein within the leaf tissue, or of parenchyma cells. The parenchymatous, chloroplast-containing bundle sheath cells of some plants function in C4 photosynthesis (discussed later).

PHOTOSYNTHESIS

The tremendous importance of plants is directly related to the photosynthetic process. Photosynthesis occurs by the fixation of carbon dioxide in the following net reaction: $n\text{CO}_2 + n\text{H}_2\text{O} \rightarrow (\text{CH}_2\text{O})_n + n\text{O}_2$. Interestingly, this net reaction actually occurs via two series of interdependent reactions: light reactions and dark reactions. In the **light reactions**, which occur within the thylakoid membranes and require photons of light, water (H_2O) is broken down into hydrogen ions (H^+), electrons ($-$), and molecular oxygen (also called dioxygen, O_2). This splitting of water molecules occurs via a complex series of enzymes and cofactors embedded within the thylakoid membranes of the chloroplast (Figure 10.26). The hydrogen ions resulting from the splitting of water become concentrated in the space within the thylakoids. These hydrogen ions are transported across the thylakoid membrane into the outer region called the stroma; that transport results in a net transfer of energy, used to synthesize a high-energy molecule of ATP (adenosine triphosphate; Figure 10.26). The electrons produced by the splitting of water are also transported across the thylakoid membrane to the stroma. There, the electrons react with hydrogen ions and a compound called NADP^+ (nicotinamide adenine dinucleotide phosphate) to produce a higher energy product, NADPH.

In the **dark reactions** (or Calvin cycle) atmospheric carbon dioxide (CO_2) makes its way into the stroma of the chloroplast, where it reacts with a five-carbon molecule to form two molecules, each containing three carbon atoms; hence, photosynthesis in these plants is called **C3 photosynthesis** and the plants are called **C3 plants** (Figure 10.26). This initial binding, or *fixation*, of CO_2 is catalyzed by a very important enzyme called **ribulose-bisphosphate carboxylase (RuBP-carboxylase)**, which is thought to be the enzyme with the greatest worldwide biomass). The two three-carbon molecules then undergo a series of further reactions, each catalyzed by

a separate enzyme, to ultimately produce a net molecule of glucose. The chemical reactions resulting in glucose production require the input of high-energy compounds, notably ATP and NADPH. As these compounds are converted into lower energy products in the dark reactions, they are regenerated in the light reactions. Thus, light and dark reactions are interdependent; each comes to a halt without the concerted action of the other (Figure 10.26).

In some species of vascular plants, the parenchymatous bundle sheath cells function in a different type of photosynthesis called **C4 photosynthesis** (Figure 10.27). In C4 plants carbon dioxide is initially fixed in the mesophyll cells by a different enzyme, **PEP carboxylase**. The initial molecule of carbon fixation is a four-carbon molecule, which, in the form of malic acid, is then transported to the bundle sheath cells. Chloroplasts of the bundle sheath cells are typically much larger than those of the mesophyll cells, this type of anatomy termed **Kranz anatomy** (Figure 10.28). In the bundle sheath cells, the carbon dioxide is released and fixed by the typical (and ancestral) enzyme, **ribulose-bisphosphate carboxylase (RuBP-carboxylase)**. C4 photosynthesis actually requires more energy (one more ATP per CO_2 molecule) than C3 photosynthesis. However, C4 photosynthesis has apparently been selected for in plants growing under conditions of high light intensity or drought. Under water-stressed conditions, the stomata of plants generally remain closed to inhibit excess water loss, but this also inhibits the flow of CO_2 into the leaf. The enzyme PEP carboxylase has a much greater affinity for CO_2 molecules than does the enzyme RuBP carboxylase. Thus, under conditions of low CO_2 (occurring under drought conditions), the initial fixation of CO_2 is much more efficient in C4 plants than in C3 plants. By fixing, transporting, and releasing CO_2 into the bundle sheath cells, it becomes more concentrated than in the mesophyll cells and can more readily be catalyzed by RuBP carboxylase in the Calvin Cycle. C4 photosynthesis has evolved in a number of angiosperms, one common example being corn (*Zea mays*, Figure 10.28).

Another different mechanism of photosynthesis is **CAM: crassulacean acid metabolism** (Figure 10.29). CAM plants are often succulents; as with C4 plants, CAM plants are generally adapted to xeric conditions. CAM photosynthesis is adaptive in minimizing water loss due to evapotranspiration. In CAM plants initial fixation of CO_2 occurs at night, when (unlike other plants) stomata are open. The CO_2 is initially fixed by the enzyme **PEP carboxylase** to form malic acid, which is temporarily stored within vacuoles of the mesophyll cells. (This is experimentally detected by a lowering of the pH.) During the day the stomata close and CO_2 is released from the vacuoles into the cytoplasm, where it is fixed in the chloroplasts by the Calvin cycle.

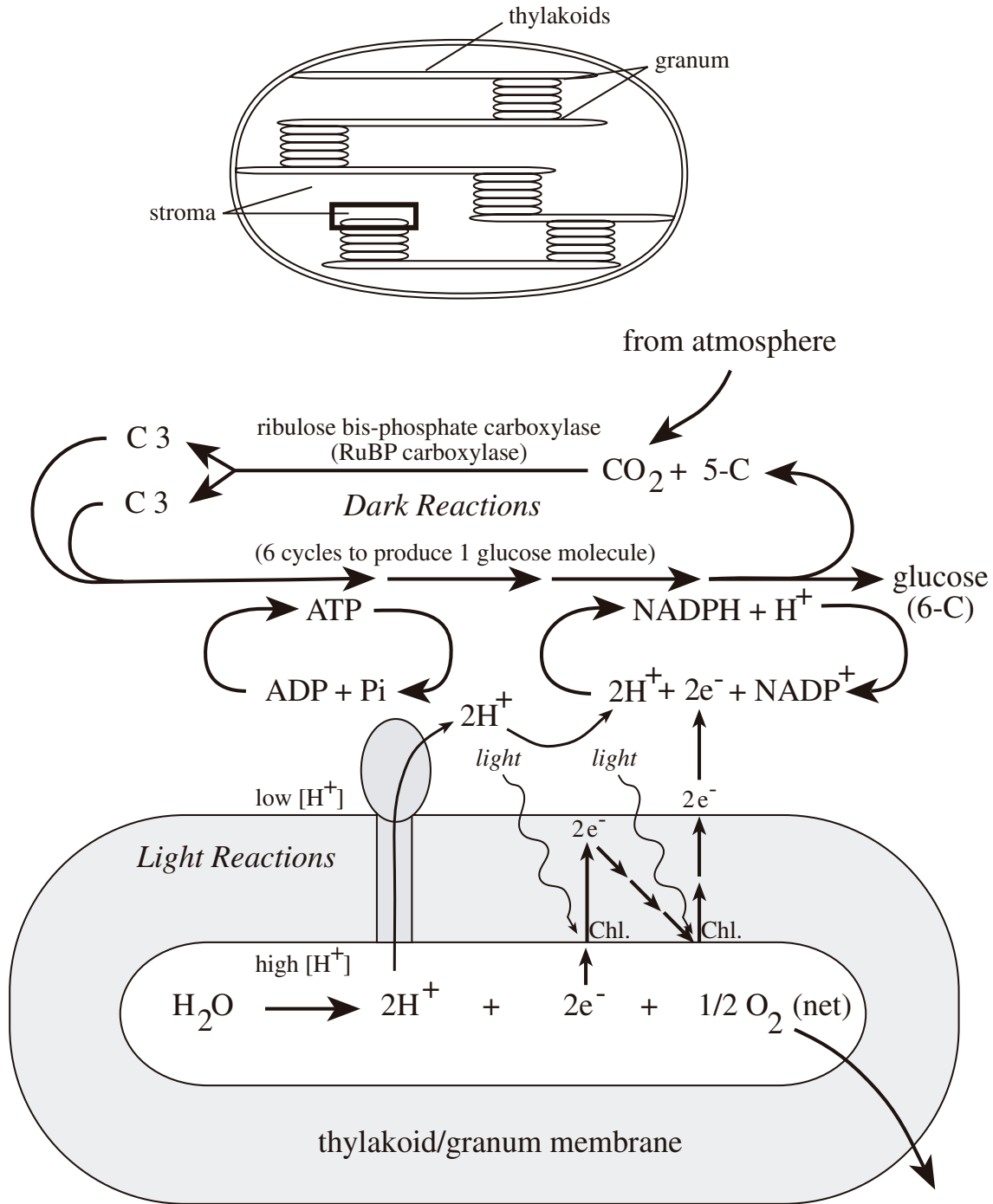


FIGURE 10.26 Photosynthesis (C3). ADP = adenosine diphosphate; Pi = phosphate. See text for other abbreviations.

CAM and C4 photosynthesis are very similar to one another. Both are adaptations to xeric conditions and involve initial fixation of CO_2 utilizing the enzyme PEP carboxylase and final fixation of CO_2 with RuBP carboxylase. The essential difference between the two is that initial and final CO_2 fixation differ spatially in C4 plants and temporally in CAM plants.

ANATOMY AND SYSTEMATICS

Plant anatomy can provide valuable characteristics in phylogenetic analyses, but these are less frequently acquired today than in the past. However, anatomical features, whether used directly to generate a cladogram or merely traced on an existing

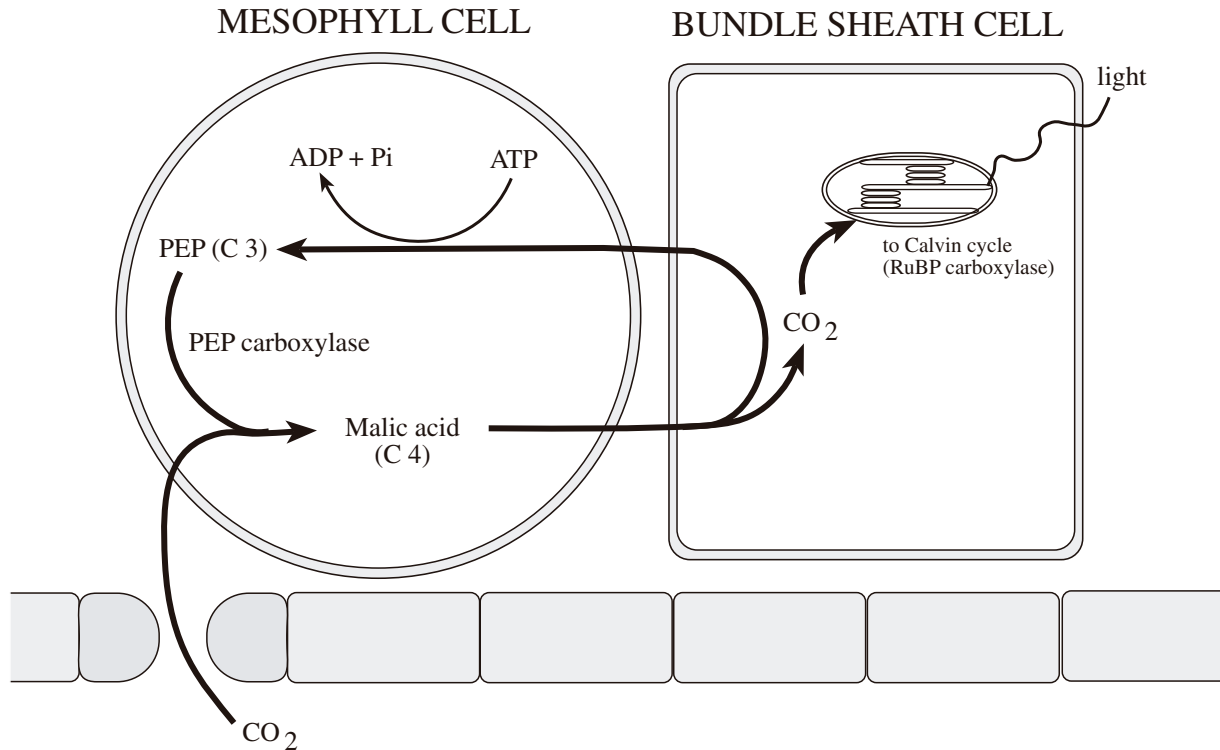


FIGURE 10.27 C₄ photosynthesis. ADP = adenosine diphosphate; PEP = phosphoenolpyruvate; Pi = phosphate. See text for other abbreviations.

cladogram, can give insight into major adaptational shifts. In that sense, they are quite important in understanding different selective pressures.

A summary of major anatomical apomorphies for the land plants is seen in Figure 10.30, taken from the cladograms

of Chapters 3.6. As can be seen, many of the apomorphies discussed and presented in these chapters are anatomical. Anatomical and physiological traits are worthy of study at a lower taxonomic level as well, and are often correlated with adaptational strategies and ecological shifts.

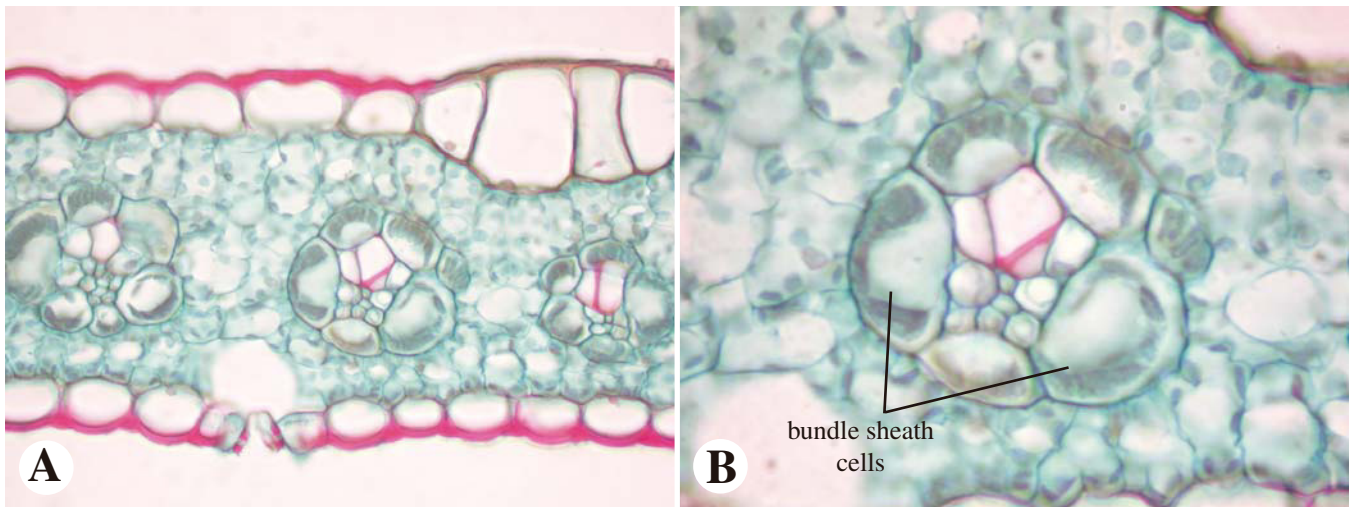


Figure 10.28 Kranz anatomy, illustrated by *Zea mays* leaf cross-section. **A.** Low magnification. **B.** Close-up of vascular bundle with enlarged bundle sheath cells, surrounded by mesophyll cells.

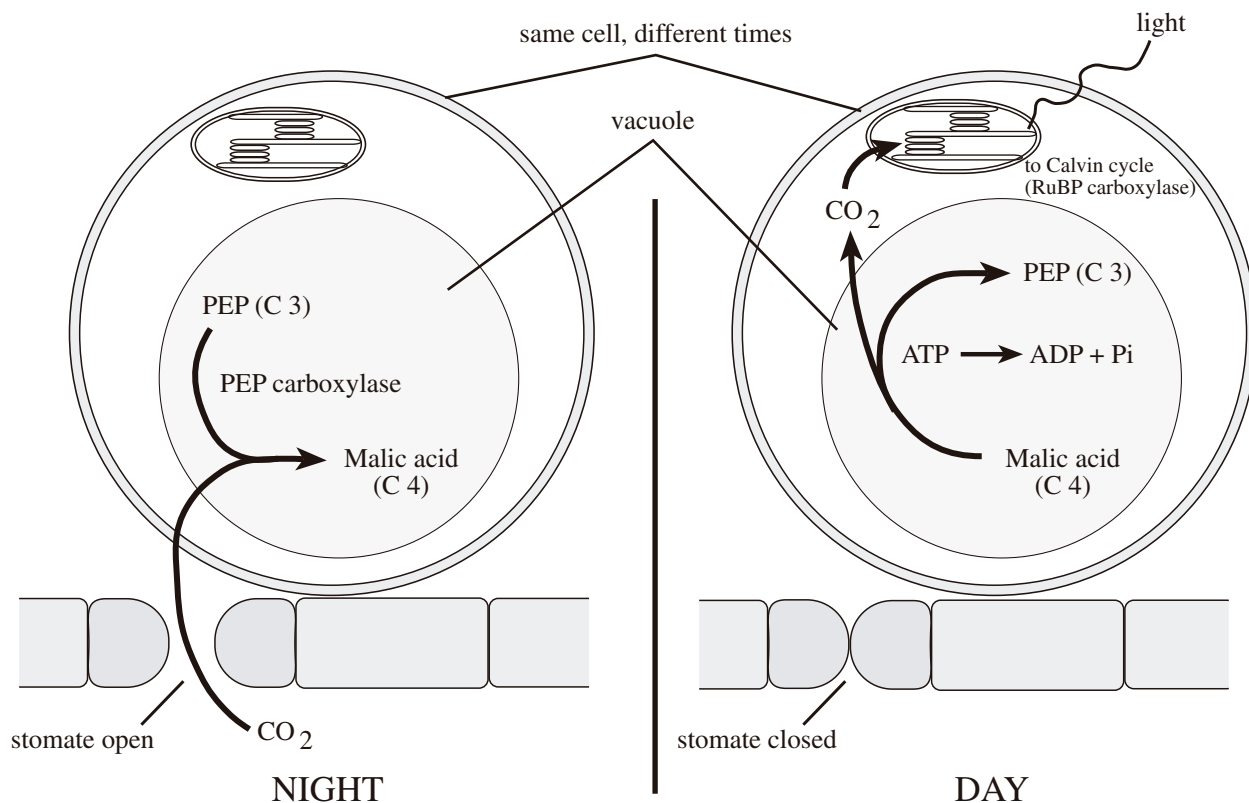


FIGURE 10.29 CAM photosynthesis. ADP = adenosine diphosphate; PEP = phosphoenolpyruvate; Pi = phosphate.

PLANT ANATOMY TECHNIQUE

Material dissection and preparation:

A wealth of information can be gained by careful dissection and observation of plants. Look first at the outer form of the plant, noting the basic plant organs (root, stem, leaves, buds, flowers, fruits) and specific aspects of these organs. Gently pull apart the plant organs to better see their morphology. For flowers and fruits, use both your hands and naked eye and dissecting needles under a dissecting scope to examine the components.

Careful anatomical studies usually involve time-consuming embedding and microtome sectioning. However, a simple technique of hand sectioning with a razor blade will allow you to see considerable detail of cell and tissue anatomy. Stout material, such as an herbaceous stem, can be held *upright* in the left hand between thumb and index finger (assuming you are right-handed). More flimsy material, such as a leaf, can be sandwiched between two small pieces (cut only slightly larger than the material) of Styrofoam; the end is moistened and both Styrofoam and plant material are sectioned together. In either case, rest the side of the razor blade on your index finger and position your thumb a bit lower (so that if you do slip, you won't cut yourself). There are tricks of the trade to successful sections:

1. As you cut, move the razor blade toward you, as well as across the material; thus, the cut is somewhat diagonal.
2. Make an initial cut to level off (discarding this piece) and then make several thin slices, keeping the sections on the razor blade until they get too crowded; then, transfer the sections to water in a Syracuse dish or Petri plate. Clean your razor blade and make a few more sections.
3. Select the thinnest sections, pull out with a brush, and place in a few drops of stain in another dish. After staining, rinse your sections very briefly in water and place in a drop of water or (for a semipermanent mount) 50% glycerol. Cover with a cover slip, avoiding air bubbles and adding more fluid to the side if necessary.

Most important is to make those sections **THIN!!** Although you will want at least one complete section, other sections may be partial, as long as they are thin. Clean your razor blade afterward and you may reuse it.

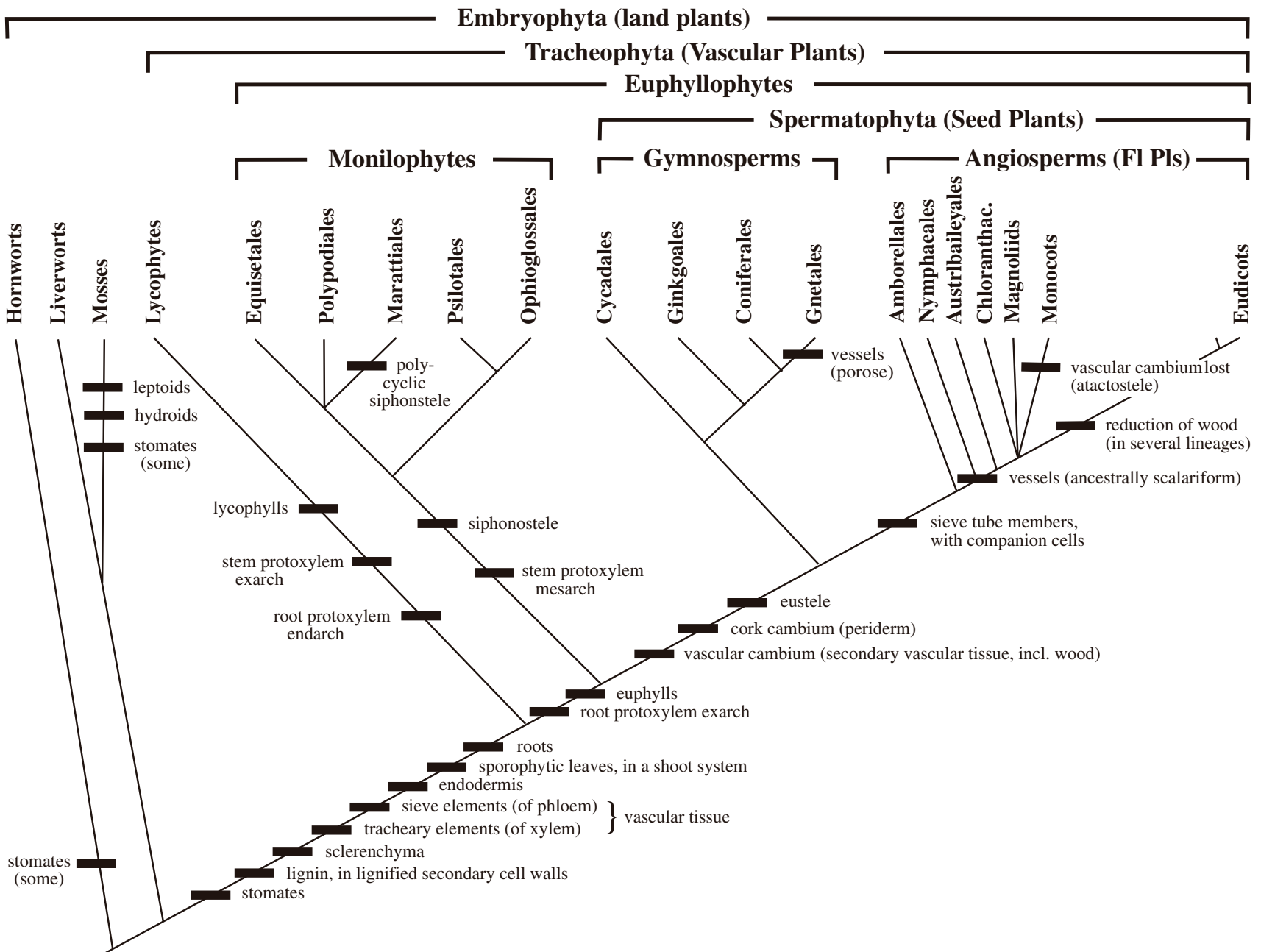


FIGURE 10.30 Summary cladogram of Land Plants, showing major anatomical apomorphies.

For tough, fibrous, or woody tissue place the material down on a plastic Petri plate and make downward slices with your razor blade. This same technique can be used with softer, small plant material if it is sandwiched between two layers of Parafilm and the material sectioned in a dicing motion.

The following are some vital stains (i.e., used with live material):

Stain	Compound for which stain is specific	Color
Alcian blue	Pectins	Blue
Aniline blue	Callose	Blue (UV-fluoresces Yellow)
IKI	Starch	Blue to black
Phloroglucinol/HCl	Lignin	Red (NOTE: Takes sev. mins. to react)
Sudan III or Sudan IV	Oil droplets	Reddish
Toluidine blue	Metachromatic (will stain a variety of cell walls different shades of blue/green):	
	Lignified tracheary elements	Dark blue
	Sclerenchyma	Blue to blue-green
	Parenchyma	Light blue
	Collenchyma	Reddish-purple
	Sieve tubes and companion cells	Greenish
	Callose/starch	Unstained

Drawings:

Making careful drawings not only gives you a record of what you observe, it also helps you become a careful observer. When forced to draw it, you often see more than you otherwise would. Make drawings with a #2 or #3 hard lead pencil. Draw the outlines of organs or tissues (e.g., of a root cross-section) at low magnification to record the overall structure. Then draw a portion of the whole (e.g., a pie slice of the root section, showing some of the individual cells of a vascular bundle) to show details.

REVIEW QUESTIONS

PLANT CELL STRUCTURE AND PLANT GROWTH

1. What is plant anatomy and how does it differ from animal anatomy?
2. What is the cell theory and its four tenets?
3. Name and give the function of the major components and organelles of a typical plant cell.
4. Name the various types of ergastic substances.
5. What are chromoplasts and what is their function?
6. What is an amyloplast and what is its chemical composition?
7. Of what is an aleurone grain composed?
8. Name four types of crystals based on their shape. What two different substances make up plant crystals?
9. Characterize a primary cell wall in terms of development and structure.
10. How does cellulose differ from starch?
11. What are plasmodesmata and what is their function?
12. Where is the secondary cell wall formed in relation to the plasma membrane and the primary cell wall?
13. What are the name, properties, and function of the compound (other than cellulose) making up a secondary cell wall?
14. What is a pit and what is the function of pit-pairs?
15. What are meristems and what are the two major types?
16. Explain the processes of cell differentiation.

PLANT TISSUES AND SPECIALIZED CELL TYPES

17. What is a tissue and what are the three general tissue types?
18. How are parenchyma and collenchyma similar and how different with respect to structure and function?
19. What are the characteristics and two general cell types of sclerenchyma?

20. What is the function of xylem and why is it a complex tissue?
21. What are the names and characteristics of the two types of water-conductive cells of xylem?
22. What is the function of phloem and why is it a complex tissue?
23. What are the names and characteristics of the two types of sugar-conductive cells of xylem?
24. What tissue occurs as the outermost cell layer of plant organs?
25. Describe the characteristics and function of the epidermis, stomata, trichomes, and secretory structures.

ROOT STRUCTURE AND FUNCTION

26. From what in an embryo does the first root arise?
27. Distinguish between a primary, lateral, and adventitious root.
28. Name three ways that roots can be distinguished from shoots/stems.
29. What is a Casparian strip and endodermis and what are their function?
30. What is the function of the pericycle?
31. What is a velamen?

SHOOT/STEM STRUCTURE AND FUNCTION

32. What is the definition of a shoot?
33. From what in an embryo does the first shoot arise?
34. What is a bud primordium and where are buds typically located?
35. What are three ways that stems differ from roots?
36. What is a stele? Name five stele types and distinguish between them.
37. Distinguish between protoxylem and metaxylem; between exarch, endarch, and mesarch.

SECONDARY GROWTH

38. What is secondary growth and from what general type of meristem does it arise?
39. Where does the vascular cambium arise?
40. What two products does the vascular cambium give rise to and in what direction?
41. What is the technical name for wood?
42. Where does the cork cambium form, and what two tissues does it give rise to?
43. Describe the adaptive significance of the lateral meristems.
44. What is a ray and what is its function?
45. What is an annual ring and what is the structural difference between spring wood and summer wood?
46. Define and draw the three major sections of wood.
47. What is the difference between a softwood and a hardwood?
48. Distinguish between nonporous, ring-porous, and diffuse-porous wood.
49. Distinguish between uniseriate, biseriate, and multiseriate rays.
50. What is dendrochronology and for what can it be used?

LEAF STRUCTURE AND FUNCTION

51. What is the difference between a lycoophyll and a euphyll?
52. What are the structure and function of stomates and of subsidiary cells?
53. What is the name of the leaf cells located between upper and lower epidermal layers? What are the two types called?

PHOTOSYNTHESIS

54. Describe the basic pathway of C₃ photosynthesis.
55. What enzyme functions to fix carbon dioxide in C₃ photosynthesis?
56. How does C₄ photosynthesis differ from C₃?
57. What is Kranz anatomy?

58. What enzyme functions to fix carbon dioxide in C₄ photosynthesis?
59. How does CAM photosynthesis differ from C₃ and C₄ and how does this function for plants living in dry conditions?

ANATOMY AND SYSTEMATICS

60. Draw a general cladogram of land plants, illustrating several anatomical apomorphies.

EXERCISES

1. Obtain live material of a plant species and prepare hand sections of the root, stem, and leaf, if feasible. Stain these with the appropriate stain (see PLANT ANATOMY TECHNIQUE), and describe all the cell and tissue types. Note the differences between the three organs.
2. Obtain live material of the leaves of a few monocot and eudicot species. Prepare epidermal peels of the leaves and note the differences between the stomata and subsidiary cells. Can you determine a correlation with taxonomic group?
3. Observe the trichomes of various plant organs (e.g., leaves, stem axes, or flower parts) by peeling the epidermal tissue bearing the trichomes or scraping them from the surface. Place this material on a microscope slide in a drop of water or (to preserve for some time) 50% glycerol. The material may be stained with, e.g., toluidine blue. Carefully draw the various trichome types. Is the trichome anatomy the same from organ to organ or does it vary? What might be the adaptive significance of trichomes?
4. Peruse journal articles in plant systematics, e.g., *American Journal of Botany*, *Systematic Botany*, or *International Journal of Plant Sciences*, and note those that describe plant anatomical features in relation to systematic studies. Identify all anatomical characters and character states used.

REFERENCES FOR FURTHER STUDY

- Dickison, W. C. 2000. *Integrative Plant Anatomy*. Harcourt/Academic Press, New York.
- Esau, K. 1967. *Plant Anatomy*, 2nd ed. John Wiley and Sons, New York.
- Esau, K. 1977. *Anatomy of Seed Plants*, 2nd ed. John Wiley and Sons, New York.
- Fahn, A. 1967. *Plant Anatomy*. Pergamon Press, Oxford.
- Mauseth, J. D. 1988. *Plant Anatomy*. Benjamin Cummings.
- Metcalfe, C. R., and L. Chalk. 1950. *Anatomy of Dicotyledons*, Vols. 1 and 2. Clarendon Press, Oxford.

11

PLANT EMBRYOLOGY

ANTHER AND POLLEN DEVELOPMENT	437	SEED DEVELOPMENT	446
Anther Type	437	Embryogeny	446
Anther Wall Development	437	Embryo Type	447
Pollen Development	439	Endosperm Development	447
OVULE DEVELOPMENT	439	Seed Storage Tissue Origin	448
Ovule Parts	439	Seed Storage Tissue Composition	448
Nucellus Type	440	Seed Coat Anatomy	448
Megasporogenesis	441	Seed Morphology	449
Megagametogenesis	441	EMBRYOLOGY AND SYSTEMATICS	449
Integument Type	441	EMBRYOLOGICAL TECHNIQUE	449
Micropyle Type	441	REVIEW QUESTIONS	450
Ovule Type	442	EXERCISES	450
Ovule Position	445	REFERENCES FOR FURTHER STUDY	451
Obturator Presence/Absence	446		

Plant embryology is the study of the development of sporangia, gametophytes, and embryos in the land plants, the embryophytes. Among the seed plants, the spermatophytes, embryological studies encompass the development of microsporangia (within anthers in the angiosperms), microspores, pollen grains, ovules, megaspores, female gametophytes, and seeds. Because most plant embryological data have been acquired and utilized for the flowering plants, this chapter focuses on processes and terminology for the angiosperms.

As characters used in phylogenetic studies, plant embryological data are generally most useful at higher taxonomic levels, as in the characterization of traditional plant families. However, the data may be useful at any taxonomic level.

ANTHER AND POLLEN DEVELOPMENT

ANTHER TYPE

In the angiosperms an important embryological character, one often treated as a standard morphological character, is the number of microsporangia per anther. Microsporangia are typically tubular in shape and occur in pairs, which coalesce

during development by the breakdown of the cell layers between them. Each pair of microsporangia is termed a **theca**. The great majority of angiosperm species have anthers composed of two thecae, termed **bithecal** or **tetrasporangiate** (Figure 11.1A), which is the ancestral condition. However, some angiosperm taxa, such as the Malvaceae, Cannaceae, Marantaceae, and species of *Salvia* (Lamiaceae), have a derived anther type with only one theca, termed **monothecal** or **bisporangiate** (Figure 11.1B).

ANTHER WALL DEVELOPMENT

A cross-section of an anther reveals a division between the internal microsporangium, the cells of which undergo meiosis, and an outer anther wall. The development of the anther wall has provided some useful embryological features. A mature anther wall consists of few to several layers of cells. The outermost cell layer (just inside the epidermis) is termed the **endothecium**, which typically consists of enlarged cells with secondary wall thickenings functioning in anther dehiscence. The secondary wall thickenings function by providing tensile force that pulls back the anther walls from the line or region of dehiscence. The innermost cell layer is termed the **tapetum**, which consists of metabolically active cells that

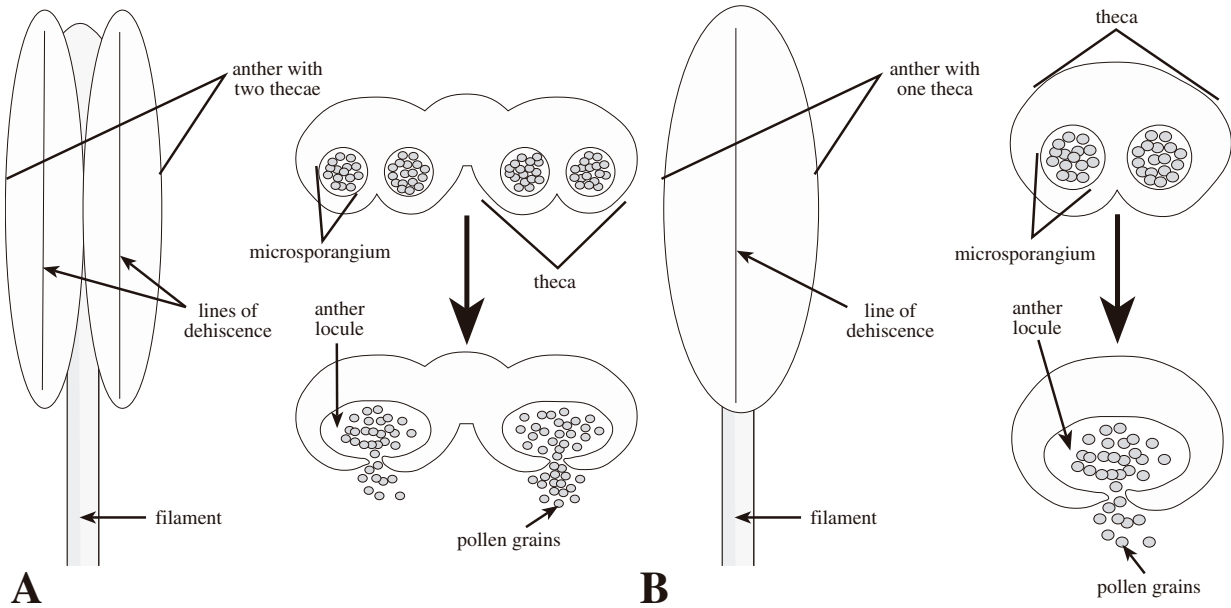


FIGURE 11.1 Anther types in the angiosperms. **A.** Dithecal. **B.** Monotheical.

function in the development of pollen grains. Additional wall layers, termed middle layers, may occur between the endothecium and tapetum. Both the total number of wall layers and their developmental origin define various anther wall types. Early in development an anther contains *two* layers of cells, an outer epidermis and an inner layer of **primary parietal cells**. Cells of the primary parietal layer

divide tangentially (parallel to the outer surface) to give rise to two layers of cells, **secondary parietal cells**. Based on the derivation of cell lineages, four general types of anther wall development have been defined (Figure 11.2): (1) **basic**, in which both secondary parietal cell layers divide to yield two middle layers; (2) **dicotyledonous**, in which only the *outer* secondary parietal cell layer divides to yield the endothecium

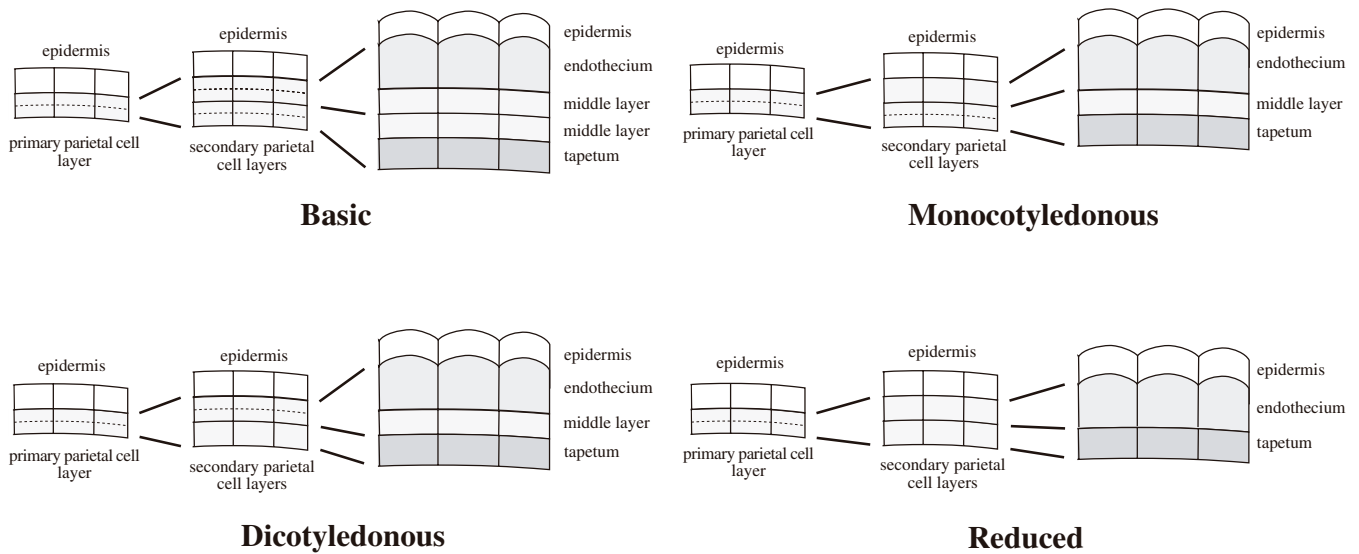


FIGURE 11.2 Anther wall development, outer epidermis at top.

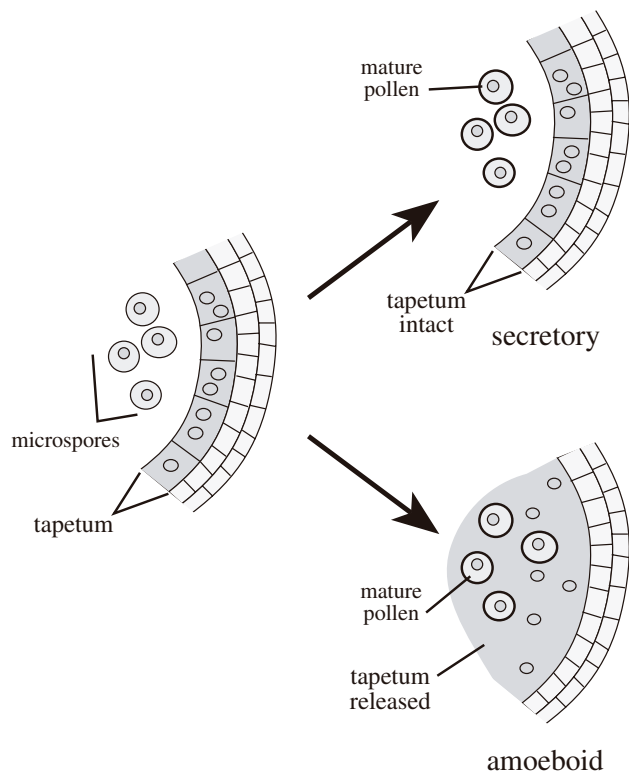


FIGURE 11.3 Tapetum types.

and a single middle layer; (3) **monocotyledonous**, in which only the *inner* secondary parietal cell layer divides to yield the tapetum and a single middle layer; and (4) **reduced**, in which the secondary parietal cells do not divide further and develop directly into the endothecium and tapetum, respectively.

Another embryological character concerns the development of the tapetum, with two basic types defined (Figure 11.3). In some angiosperms the tapetum remains intact with no breakdown of cell walls. This tapetal type is called **secretory** (or glandular; Figure 11.4A,B) because of the implication that compounds are secreted into the locule of the anther that function in pollen development. In other angiosperm taxa the tapetal cell walls break down, with release of the cytoplasm of the tapetal cells into the locule. This latter tapetal type is called **amoeboid** (plasmodial or periplasmodial; Figure 11.4C,D) because the cytoplasmic contents surround developing pollen grains like an amoeba surrounds food. Subtypes of the secretory and amoeboid tapetal types have been proposed by some, based on fine developmental differences.

A final embryological character dealing with the anther wall is endothelial anatomy. Two basic types of endothelial cells have been defined based on the structure of the secondary wall thickenings. A **girdling** endothecium is one in which the secondary wall thickenings form rings with cross bridges

between them (Figure 11.5). A **spiral** endothecium is one in which the secondary wall thickenings are spiral or helical in shape.

POLLEN DEVELOPMENT

Development of microspores from microsporocytes is termed **microsporogenesis**. There are two basic types of microsporogenesis as determined by the timing of cytokinesis, which is the formation of a plasma membrane and cell wall that divides one cell into two (Figure 11.6A). If cytokinesis occurs after meiosis I, then microsporogenesis is **successive** (Figure 11.6B). Successive microsporogenesis results in two cells after meiosis I and four cells after meiosis II. If cytokinesis doesn't occur until after meiosis II, then microsporogenesis is **simultaneous** (Figure 11.6C). Simultaneous microsporogenesis results in cell formation only after meiosis II.

Development of pollen grains (male gametophytes) from microspores is called **microgametogenesis**, technically beginning with the first mitotic division of the single microspore nucleus. One embryological character concerning microgametogenesis is the number of nuclei present in the pollen grain at the time of anthesis, or flower maturation (Figure 11.7). Most angiosperms have pollen grains that are **binucleate** (Figure 11.7), containing one tube cell/nucleus and one generative cell/nucleus. The generative cell divides to form two sperm cells only after pollen tube formation. In many angiosperm taxa, however, the pollen at anthesis is **trinnucleate** (Figure 11.7), caused by division of the generative cell prior to pollen release.

OVULE DEVELOPMENT

The development of the ovule provides a number of significant embryological characters used in plant systematics studies.

OVULE PARTS

Ovules are immature seeds, consisting of a stalk, the **funiculus**, a **megasporangium** (also called the **nucellus**), from which develops the megasporocyte and female gametophyte, plus one or two surrounding **integuments**. The **micropyle** is the pore or canal within one or more integuments through which (in angiosperms) a pollen tube traverses prior to fertilization. (In nonangiospermous seed plants, the micropyle receives pollen grains directly.) The structure of the outer versus inner integument can be used to define various micropylar types (below).

The region of the nucellus where the micropyle is located is called the **micropylar** region; that opposite the micropyle

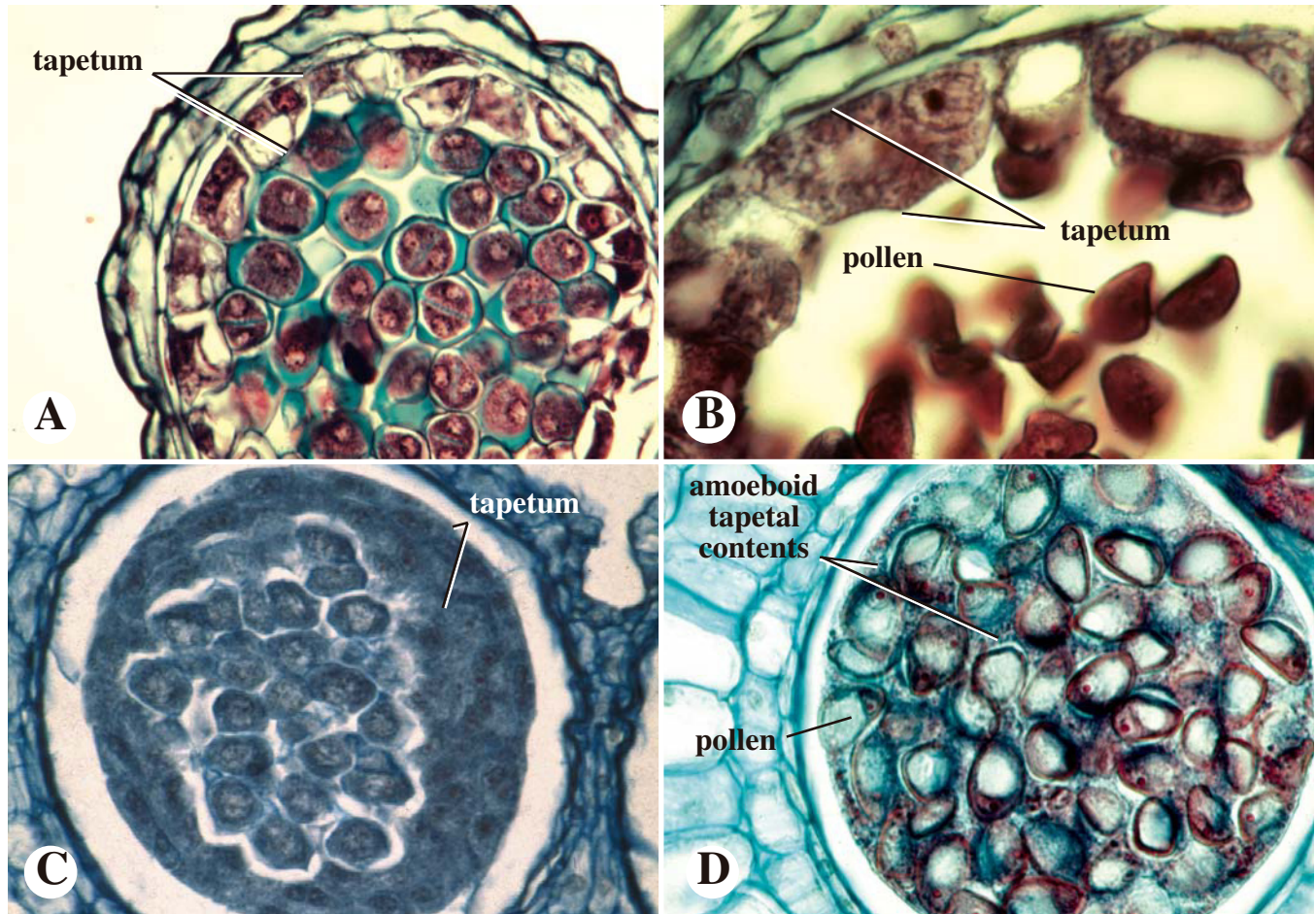


FIGURE 11.4 Anther cross-sections, showing different tapetum types. **A,B.** Secretory tapetum, the cells of which remain intact during pollen development. **C,D.** Amoeboid tapetum, in which the cells break down, releasing their cytoplasmic contents into the anther locule. Early stage at left, later stage at right for both types.

is called the **chalazal** region. A vascular strand typically traverses from the base of the funiculus to the nucellus. In most angiosperm ovules, the ovule curves during development (see Ovule Type), displacing the micropyle to a location near the funiculus base. In this type, the body of the funiculus appears fused to the body of the nucellus. This region where the funiculus is adnate (or decurrent) to the nucellus is called the **raphe**, which is sometimes visible in the mature seed as a ridge (see later discussion).

NUCELLUS TYPE

The type of nucellus, or megasporangium, is defined based on the number of cell layers comprising it (especially at the micropylar end) and the derivation of those cells. An immature ovule contains a single large cell known as an **archesporial cell**. In some taxa the archesporial cell undergoes a single

periclinal division, resulting in the formation of an outer **parietal cell** and inner **megasporocyte** (Figure 11.8). The parietal cell undergoes additional mitotic divisions, the products of which form an inner layer of nucellus cells; this type of nucellus is called **crassinucellate**, composed of two (sometimes more by additional divisions) layers of cells (Figure 11.8). On the contrary, if the archesporial cell does not divide and develops directly into the megasporocyte, the nucellus will generally be composed of a single layer of cells, the original outer layer; this type of nucellus is called **tenuinucellate** (Figure 11.8). However, in a few taxa, no parietal cell is formed, yet periclinal divisions occur in the single outer layer, forming an additional inner layer of nucellar cells; this type of nucellus is called **pseudocrassinucellate** because it appears at maturity to resemble the crassinucellate type in having two nucellar layers, but the inner layer has a

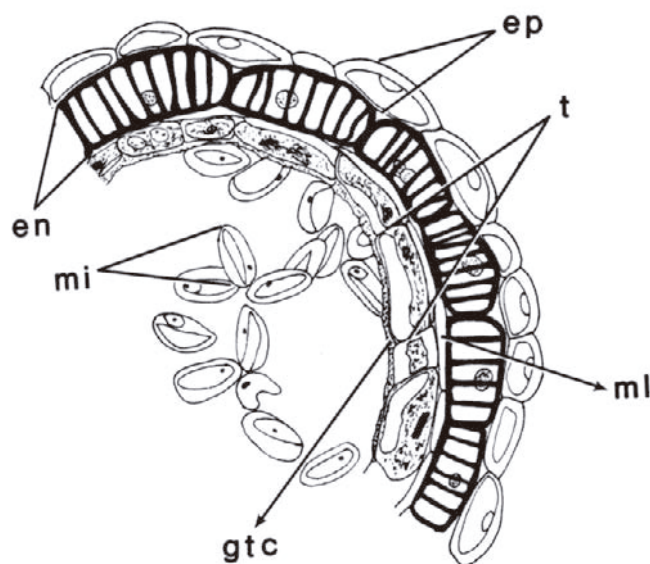


FIGURE 11.5 A girdling anther endothecium type. Symbols: **en**=endothecium; **ep**=epidermis; **gtc**=glandular tapetal cell; **mi**=microspore; **ml**=middle layer; **t**=tapetum.

different derivation. The fact that crassinucellate and pseudo-crassinucellate ovules resemble one another at maturity emphasizes the need for early developmental studies to distinguish between them.

Some taxa may have a proliferation of cell divisions of the nucellus at the micropylar region of the ovule; this mass of cells is typically termed a **nucellar beak**.

MEGASPOROGENESIS

Megasporogenesis refers to the development of megaspores from the megasporocyte, the cell that undergoes meiosis. Meiosis of the megasporocyte nucleus results in the formation of four haploid megaspore nuclei. In most taxa, meiosis is followed by cytokinesis, resulting in four megaspore cells. This pattern is termed **monosporic** megasporogenesis; because of the four megaspores produced, only one of them contributes to the female gametophyte (Figures 11.9, 11.10A D). In some angiosperm taxa, however, cytokinesis occurs after the first meiotic division, but not the second, resulting in two cells, each of which contain two haploid nuclei. This developmental pattern is termed **bisporic** megasporogenesis because one of the binucleate cells, containing two megaspore nuclei, contributes to the female gametophyte (Figure 11.9). Finally, in other taxa cytokinesis does not occur at all after meiosis, resulting in a single cell with four haploid nuclei. Because all four haploid megaspore nuclei contribute to the female gametophyte, this pattern is termed **tetrasporic** megasporogenesis (Figure 11.9).

MEGAGAMETOGENESIS

Megagametogenesis is development of the female gametophyte from the haploid product(s) of meiosis. The particular type of megagametogenesis is a function of mitotic divisions, the formation of new cells, and the fusion of existing nuclei or cells. This sequence of events defines what are termed **female gametophyte** (or **embryo sac**) **types**. The type of female gametophyte is dependent in part on the pattern of megasporogenesis, whether tetrasporic, bisporic, or monosporic. The most common and presumably ancestral type of female gametophyte in the angiosperms is one that develops from the chalazal haploid megaspore, the result of monosporic megasporogenesis. This haploid megaspore nucleus then divides mitotically to yield two nuclei, each of those two nuclei divide to yield four, and each of those four divide to yield eight. The eight nuclei arrange themselves into seven cells: three **antipodals** at the chalazal end, a large **central cell** having two **polar nuclei**, and one **egg cell** flanked by two **synergids** at the micropylar end. (The egg and two synergid cells are together termed the *egg apparatus*.) This sequence of nuclear and cell divisions gives rise to the **Polygonum type** of female gametophyte (named after the genus *Polygonum* where it was first described), the most common and the ancestral type among the angiosperms (Figures 11.9, 11.10E). However, numerous other types of female gametophytes occur in various taxa of angiosperms (Figure 11.9). For example, the *Fritillaria* type develops from a tetrasporic megasporogenesis in which three of the four megaspores fuse to form a triploid nucleus (Figures 11.9, 11.10F,G). Two sequential mitotic divisions of the haploid and triploid nuclei ultimately result in an 8-nucleate female gametophyte in which the three antipodals and one of the polar nuclei are triploid (the other polar nucleus and the cells of the egg apparatus remaining haploid).

INTEGUMENT TYPE (Figure 11.11)

The ovules of angiosperms have either one or two integuments. If two, the ancestral condition for the angiosperms, the ovule is called **bitegmic**. If one, the ovule is **unitegmic**. Unitegmic ovules have evolved in several different angiosperm groups, including the bulk of the Asteridae. Very rarely, ovules may lack any integument; this condition is termed **ategmic**.

MICROPYLE TYPE (Figure 11.12)

In a typical, bitegmic angiosperm ovule, the micropyle is typically formed or delimited by both integuments; this is termed an **amphistomal** micropyle type. If the micropyle is delimited by only the inner integument (the outer one being foreshortened), it is termed **endostomal**; if by only the

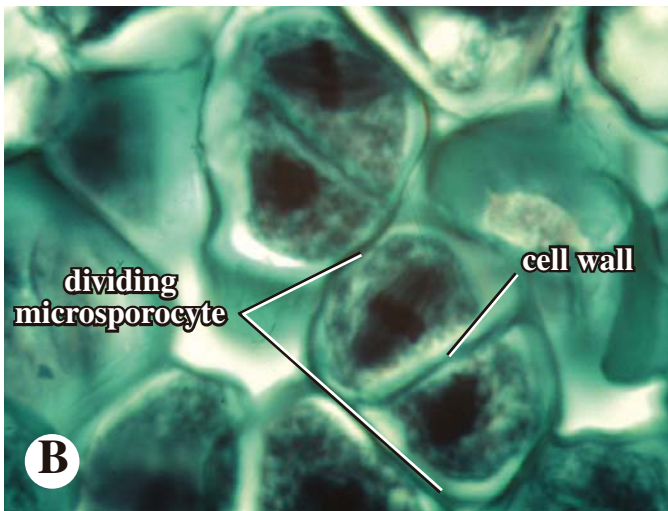
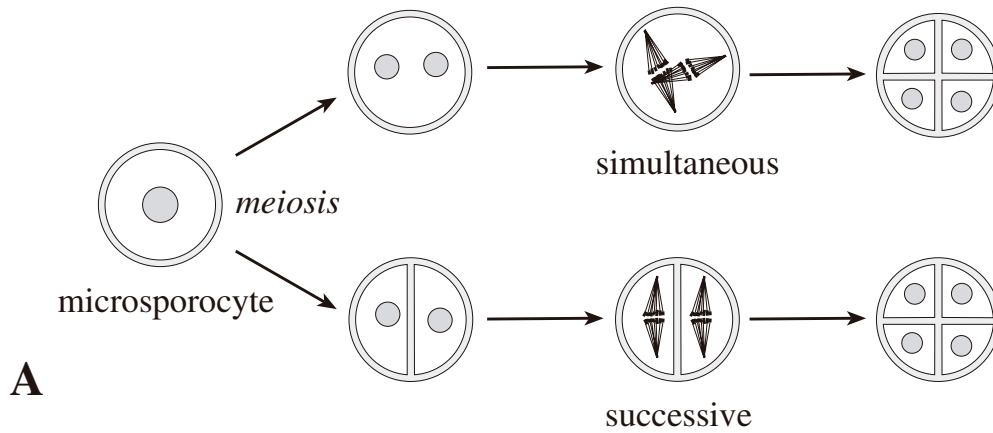


FIGURE 11.6 Microsporogenesis. **A.** Diagram showing two major types, simultaneous and successive. **B.** Successive microsporogenesis. Microsporocyte at anaphase II of meiosis. Note that cytokinesis, resulting in cell wall formation, has occurred after meiosis I. **C.** Simultaneous microsporogenesis. Note lack of cell wall after anaphase II of meiosis.

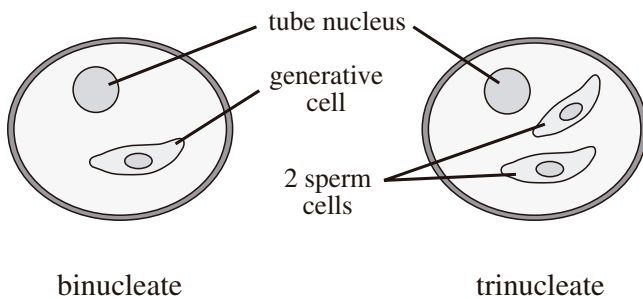


FIGURE 11.7 Microgametogenesis. Pollen nucleus number at anthesis.

outer integument (the inner one foreshortened), it is termed **exostomal**. In some angiosperms the micropyle is **zig-zag**, meaning that the micropylar pore of the outer integument is spatially displaced relative to the inner integument. If the ovule is unitegmic, the micropyle type may be called **unistomal** by default.

OVULE TYPE (Figure 11.13)

Ovule types are defined primarily on the curvature of the funiculus and nucellus/female gametophyte. The following terms are useful, yet different ovule types can be difficult to define and may require quantitative analyses. An **anatropous** ovule is one in which curvature during development results in displacement of the micropyle to a position adjacent to the funiculus base; a vasculature strand traverses from the base of

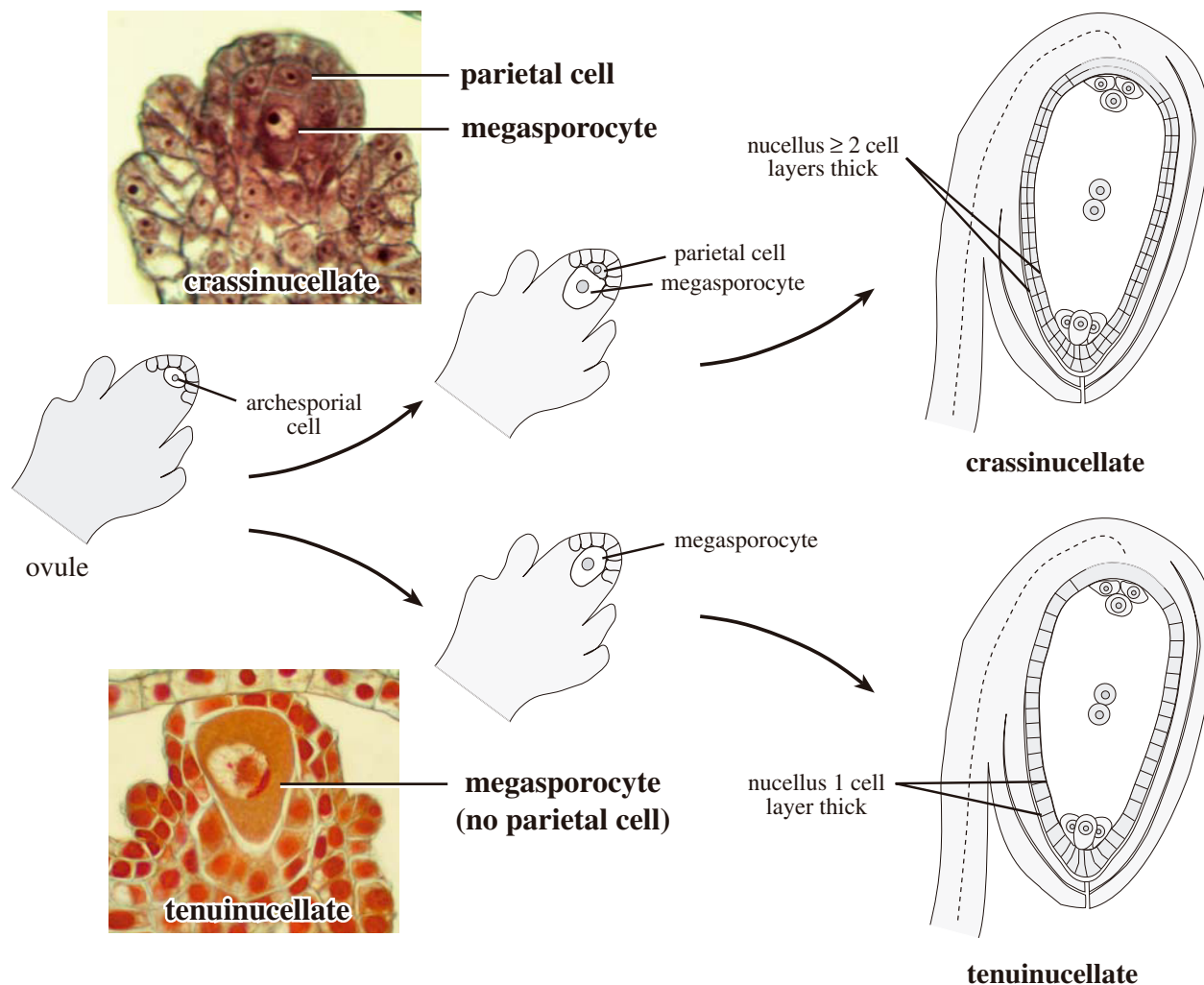


FIGURE 11.8 Nucellar type.

the funiculus to the nucellar region opposite the micropyle. The anatropous ovule type is the most common in the angiosperms and is presumed to be ancestral for the group. An **orthotropous** ovule is one in which no curvature takes place during development; the micropyle is positioned opposite the funiculus base, and the vasculature traverses from the base of the funiculus to the chalazal nucellar region. Orthotropous ovules have evolved independently in various groups of angiosperms. Both anatropous and orthotropous ovules have a straight (unbent) nucellus. (An ovule somewhat intermediate in curvature between anatropous and orthotropous is sometimes termed **hemitropous** or **hemianatropous**.)

Four other ovule types that have been defined exhibit a curvature of the ovule during development such that the micropyle is displaced adjacent to the funiculus base, similar to an anatropous ovule. These four additional ovule types

differ from an anatropous ovule in having a bent or curved nucellus, as viewed in mid-sagittal section (i.e., a section along the plane of symmetry). Traditionally, these four types were divided into only two: the **amphitropous** type, in which the nucellus is bent along both upper and lower sides, and the **campylotropous** type, in which the nucellus is bent only along the lower side. The amphitropous and campylotropous ovule types may often be cited in plant systematic literature. However, these may be subdivided into additional types (ana- and ortho-) based on the orientation of the vasculature. An **ana-amphitropous** ovule is one in which a vascular strand curves, traversing from the base of funiculus to the chalazal region of the nucellus; the nucellus is bent sharply in the middle along both the lower and upper sides, often with differentiated cells (called a basal body) at the angle of the bend. An **ana-campylotropous** ovule is similar


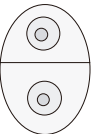
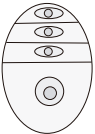
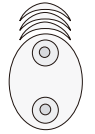
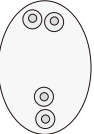



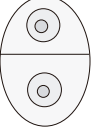
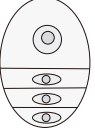
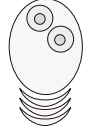


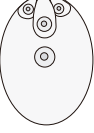

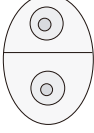
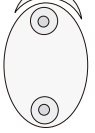
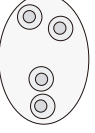
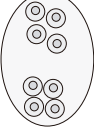

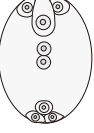

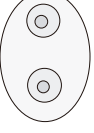
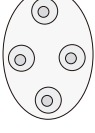
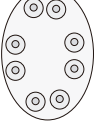
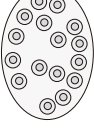



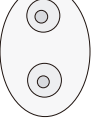
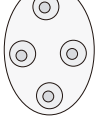
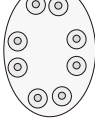
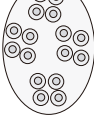

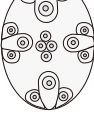

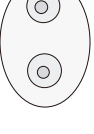

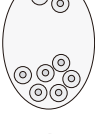


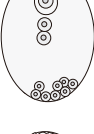
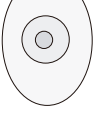



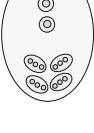

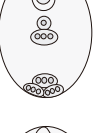
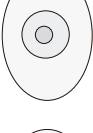






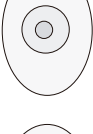

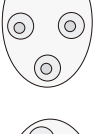
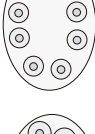


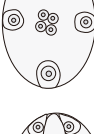


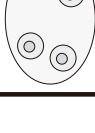
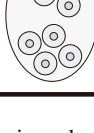



Female Gametophyte Type	Megaspороgenesis			Megagametogenesis			
	Mega-sporocyte	Meiosis I	Meiosis II	Mitosis I	Mitosis II	Mitosis III	Mature female gametophyte
Monosporic 8-nucleate <i>Polygonum</i> type							
Monosporic 4-nucleate <i>Oenothera</i> type							
Bisporic 8-nucleate <i>Allium</i> type							
Tetrasporic 16-nucleate <i>Peperomia</i> type							
Tetrasporic 16-nucleate <i>Penaea</i> type							
Tetrasporic 16-nucleate <i>Drusa</i> type							
Tetrasporic 8-nucleate <i>Fritillaria</i> type							
Tetrasporic 8-nucleate <i>Plumbagella</i> type							
Tetrasporic 8-nucleate <i>Plumbago</i> type							
Tetrasporic 8-nucleate <i>Adoxa</i> type							

FIGURE 11.9 Female gametophyte types, based on type of megasporogenesis and sequence of divisions and cell fusions during megagametogenesis. Note: micropyle above in all illustrations. Terminology after Maheshwari, 1950.

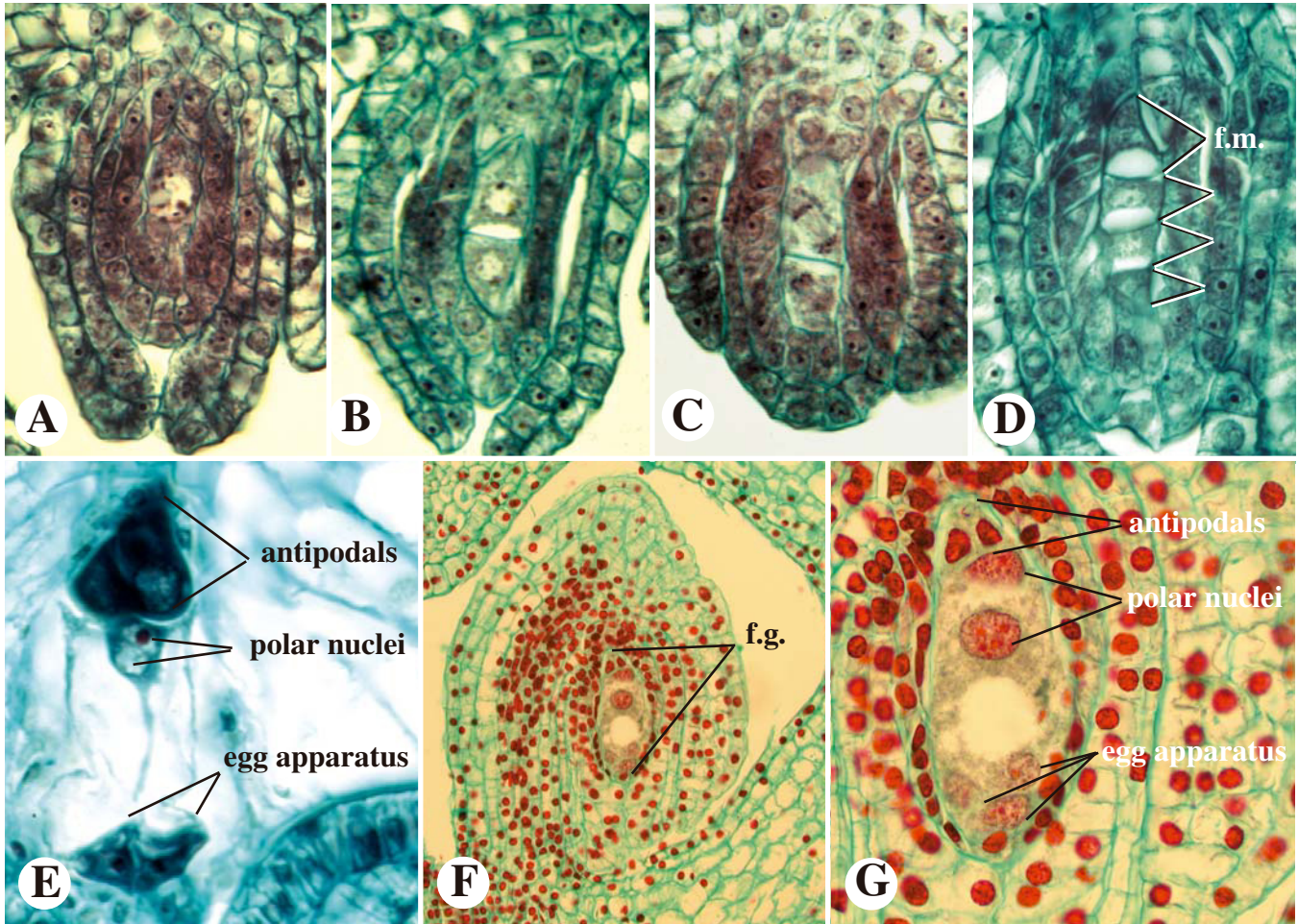


FIGURE 11.10 Examples of embryological development. (Note: micropyle below in all.) **A–D.** Monosporic megasporogenesis. **A.** Ovule with single megasporocyte. **B.** After first meiotic division. **C.** Second meiotic division. **D.** After second meiotic division, with four megaspores; f.m. = functional (proximal) megaspore. **E.** *Polygonum*-type female gametophyte (embryo sac), showing 3 antipodals, 2 polar nuclei, and 2 of 3 cells of egg apparatus. **F,G.** Megagametogenesis of *Lilium* sp., having tetrasporic, *Fritillaria*-type female gametophyte development. **F.** Whole (anatropous) ovule, with mature female gametophyte (f.g.). **G.** Mature female gametophyte close-up, showing 2 of 3 triploid antipodals, 2 polar nuclei (one haploid, the other triploid), and haploid cells of egg apparatus.

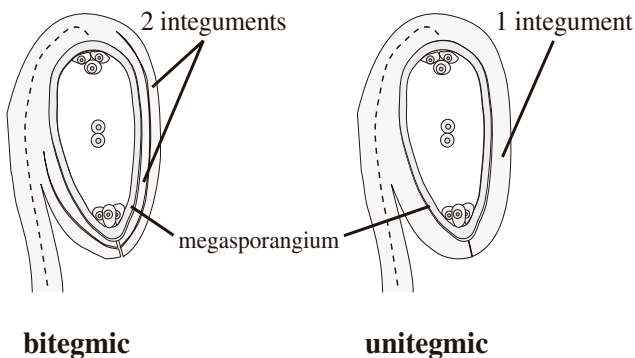


FIGURE 11.11 Ovule integument types.

to the ana-amphitropous type in vasculature, differing in that the nucellus is bent only along the lower side, with no basal body. An **ortho-amphitropous** ovule is one in which the vasculature is straight, leading from the funiculus base to the middle of the nucellus; the nucellus is bent sharply in the middle along both the lower and upper sides, often with a basal body present. An **ortho-campylotropous** ovule is similar to that of the ortho-amphitropous type, except that the nucellar body is bent only along the lower side, with no basal body.

OVULE POSITION (Figure 11.14)

Ovule position refers to the placement of the micropyle and raphe relative to the distal end (apex), proximal end (base), or sides of the floral axis.

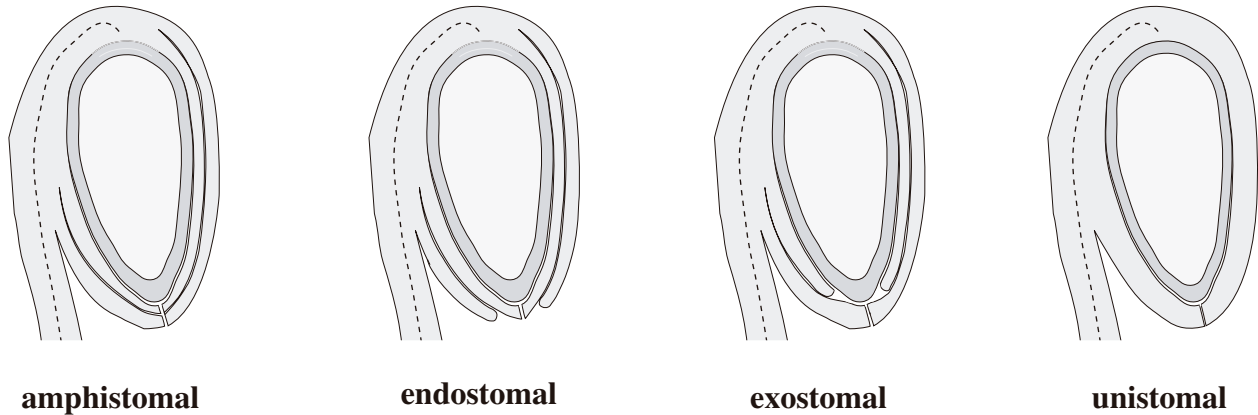


FIGURE 11.12 Ovule micropyle types.

An **epitropous** ovule is one in which the micropyle points distally. This type can be further divided into **epitropous-dorsal**, in which the raphe is dorsal (abaxial, pointing away from the central floral or ovary axis) or **epitropous-ventral**, in which the raphe is ventral (adaxial, pointing toward the central floral or ovary axis).

A **hypotropous** ovule is one in which the micropyle points proximally. This type can be further divided into **hypotropous-dorsal**, in which the raphe is dorsal (abaxial, pointing away from the central floral or ovary axis) or **hypotropous-ventral**, in which the raphe is ventral (adaxial, pointing toward the central floral or ovary axis).

A **pleurotropous** ovule is one in which the micropyle points to the side. This type can be further divided into **pleurotropous-dorsal**, in which the raphe is above or **pleurotropous-ventral**, in which the raphe is below.

A **heterotropous** ovule is one that varies in orientation.

OBTURATOR PRESENCE/ABSENCE

Rarely, a protuberance of tissue, typically arising from the funiculus or placenta, may develop at the base of the ovule.

This mound of tissue, termed an **obturator**, may be typical of certain groups, e.g., the Euphorbiaceae.

SEED DEVELOPMENT

EMBRYOGENY

Embryogeny refers to the development of the embryo within the seed. The sequence of divisions of the zygote (the product of fertilization of egg and sperm) can define various embryogeny types, which have been named after the major taxonomic groups where they occur.

Typically, the first division of the zygote is transverse (perpendicular to the long axis of the female gametophyte and nucellus), initiating the formation of a very young embryo, often termed the **proembryo**. This transverse division delimits two cells, a basal cell at the chalazal end and an apical (terminal) cell at the micropylar end. The terminal cell will divide prolifically, generally forming all or most of the **embryo proper**, which will eventually grow into the new sporophyte. Mitotic divisions of the original basal cell may

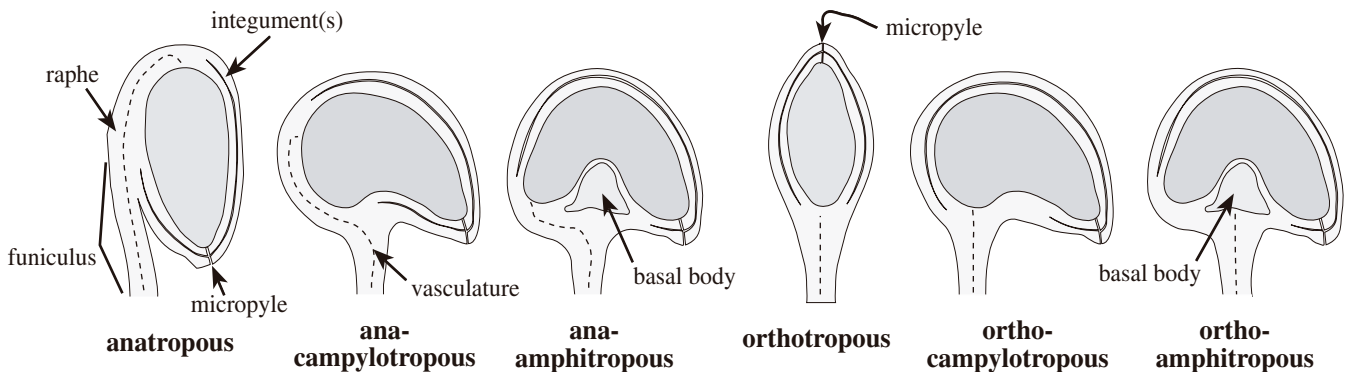


FIGURE 11.13 Ovule types.

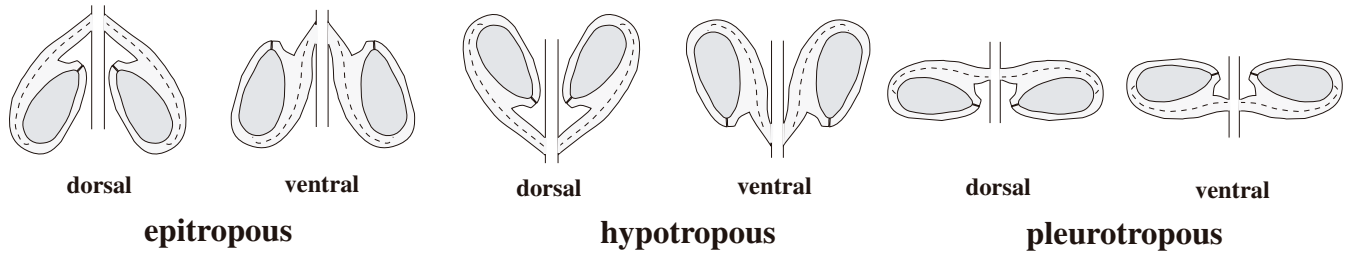


FIGURE 11.14 Ovule position, illustrated with anatropous ovules.

also contribute to the mature embryo and/or may develop into column of cells termed the **suspensor** (Figure 11.15A,B), a nonpersistent structure that functions in transport of nutrients to the mature embryo during its development.

Five embryogeny types have been defined based on the sequence of divisions of the basal and terminal cells and which cell derivatives contribute to the mature embryo: (1) **asterad type**, in which the terminal cell divides longitudinally, with both basal and terminal cell derivatives contributing to the mature embryo; (2) **caryophyllid type**, in which the terminal cell divides transversely, with only terminal cell derivatives contributing to the mature embryo; (3) **chenopodiad type**, in which the terminal cell divides transversely, with both basal and terminal cell derivatives contributing to the mature embryo; (4) **crucifer or onagrad type**, in which the terminal cell divides longitudinally, with only terminal cell derivatives contributing to the mature embryo; (5) **solanad type**, in which the terminal cell divides transversely, the basal

cells derivatives forming a suspensor but otherwise not contributing to mature embryo development.

Finally, a sixth embryogeny type, the **piperad type**, is defined if the zygote divides longitudinally (i.e., parallel to the axis of the female gametophyte and nucellus), thus not forming a basal and terminal cell.

EMBRYO TYPE

The mature embryo type is based on its form and size. The shape and size of the radicle and cotyledons is most important. Various embryo types have been defined primarily on **ptyxis**, the aestivation of the cotyledons (Chapter 9). An embryo may be either **achlorophyllous** (lacking chloroplasts) or **chlorophyllous** (green, having chloroplasts) at maturity.

ENDOSPERM DEVELOPMENT

Development of the endosperm (Figure 11.15A C) is described based on early mitosis and cytokinesis of the usually triploid,

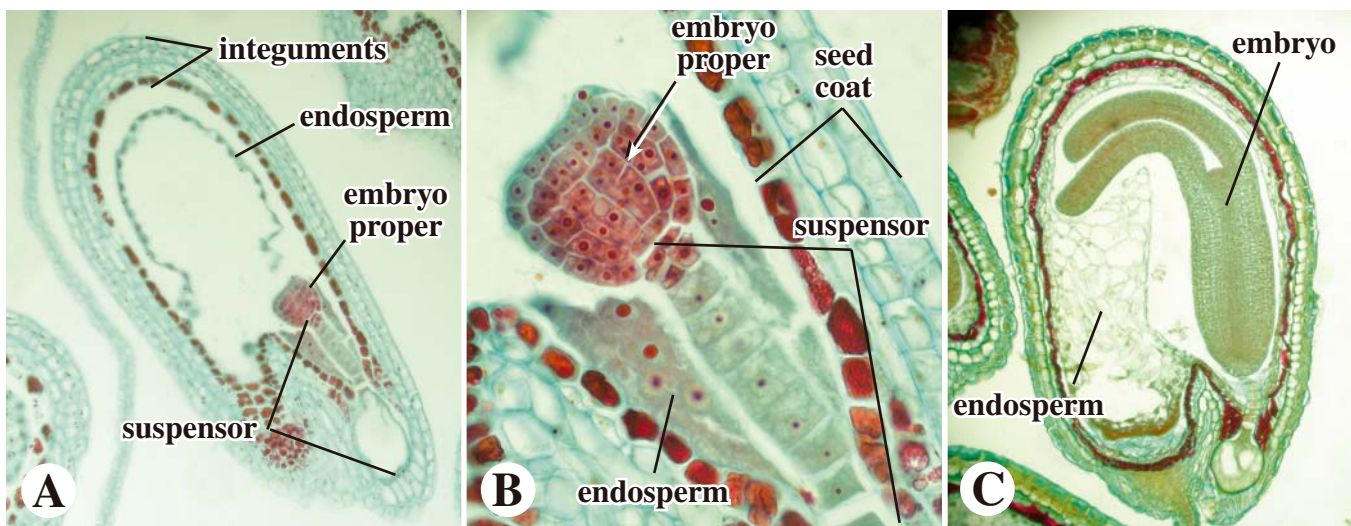


FIGURE 11.15 Endosperm and embryo development, *Capsella bursa-pastoris*. **A.** Very young seed, showing immature embryo (composed of suspensor and embryo proper), early endosperm, and seed coat. **B.** Close-up of young embryo, showing basal suspensor cells and terminal, actively dividing cells of globular embryo proper. **C.** More mature seed, showing embryo with two cotyledons.

endosperm cell (the second product of double fertilization). A **cellular** endosperm is one in which the endosperm cell divides mitotically, regularly followed by cytokinesis. Thus, each endosperm nucleus is contained within a cell wall from the beginning. A **nuclear** endosperm is one in which the early mitotic divisions are not followed by cytokinesis. Thus, numerous nuclei are contained within a single cell, at least early in development; later, cell walls typically surround the nuclei. A **helobial** endosperm is one in which the first mitotic division is followed by cytokinesis, delimiting two cells. However, the nucleus of one cell continues a nuclear type of development; that of the other cell divides in a cellular fashion.

SEED STORAGE TISSUE ORIGIN

The most common, and ancestral, type of storage tissue in angiospermous seeds is endosperm. This typical seed type is called **endospermous** or **albuminous** (Figures 11.15, 11.16). In some taxa, however, double fertilization and endosperm development occur, but the endosperm soon stops growing; the mature seed is termed **exalbuminous**. This is typical,

for example, of all orchids, which have very reduced seeds in general. In other exalbuminous taxa, the early endosperm tissue may be absorbed, with other tissues taking its place as a storage tissue. A **cotyledonous** seed storage tissue type is one in which the cotyledons enlarge and assume the function of storage tissue (Figure 11.16). Cotyledonous seeds are found, e.g., in many legumes, such as peas and beans. A **perispermous** type of seed storage tissue is one in which the chalazal nucellar cells enlarge and store energy-rich compounds.

SEED STORAGE TISSUE COMPOSITION

The storage tissue of a seed (usually endosperm) can be defined by the chemical composition of the energy-rich compounds within its cells. Storage tissue can contain primarily **starch** (in the form of starch grains or amyloplasts), **oil** (in the form of oil bodies), or **protein** (in the form of protein bodies).

SEED COAT ANATOMY

The integument(s) of the ovule matures into the **seed coat** (also called the **testa**) of the seed. Features of the anatomy of

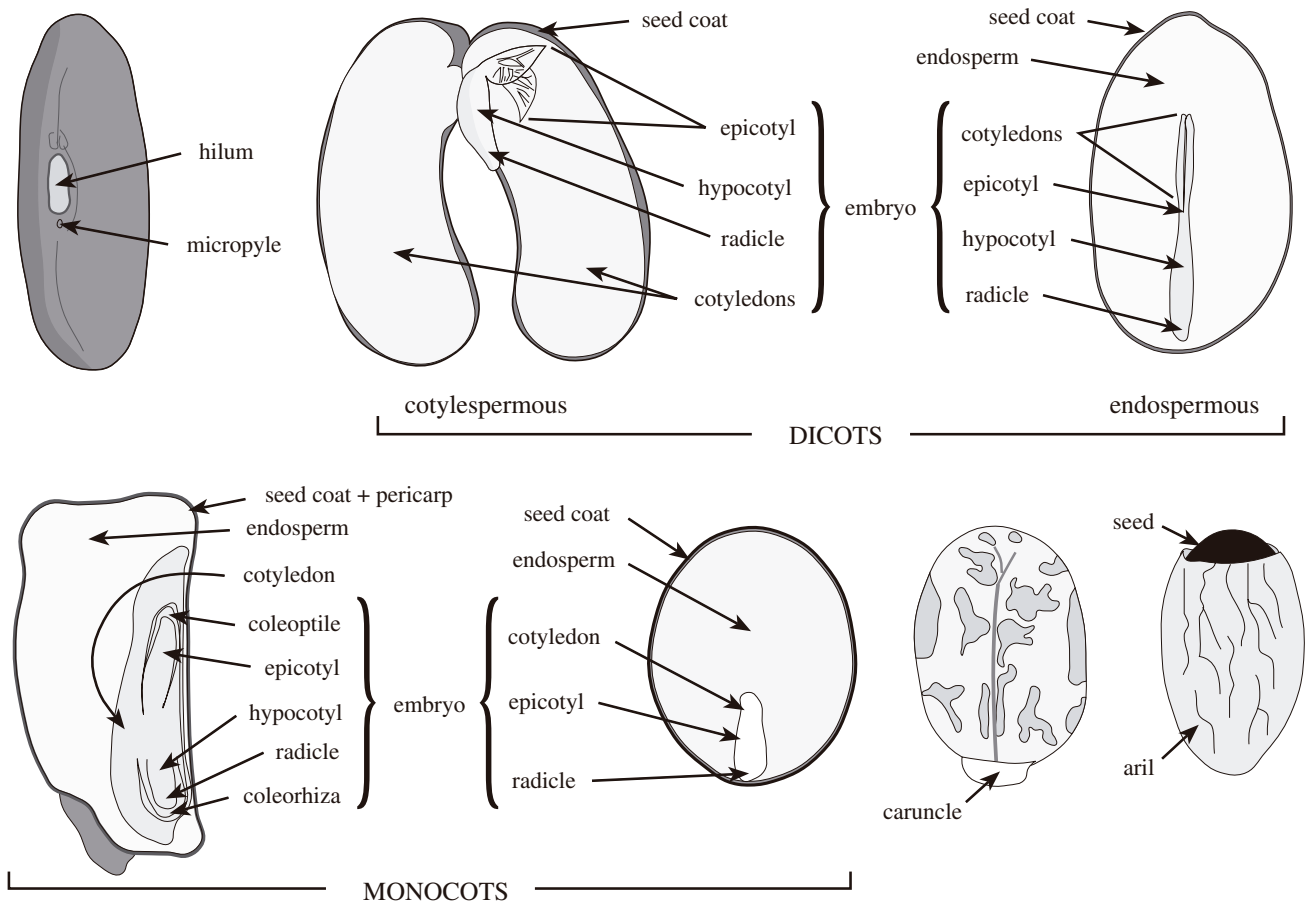


FIGURE 11.16 Seed morphology.

the mature seed coat can be significant embryological characters. These include the number of cell layers in each seed coat layer (versus the number in the integuments) and specialized cell anatomy of the cells (including cell shape, cell wall thickness, and cell wall composition) of each seed coat layer. In addition, in some taxa an extra, fleshy layer may form outside the seed coat. If the fleshy layer more or less envelops the seed coat, it is known as an **aril** (Figure 11.16). The aril generally functions as an attractant in animal dispersal. A **caruncle** is a fleshy layer that doesn't surround the seed coat, but forms as a basal appendage, typically near the **hilum** (the scar of the funiculus). The caruncle functions like an aril, as a food reward in animal seed dispersal (Figure 11.16).

SEED MORPHOLOGY

Aspects of mature seed morphology include shape, size, color, and sculpturing. In addition, some taxa have a prominent raphe, funicular scar, or hilum (micropylar scar; Figure 11.15).

EMBRYOLOGY AND SYSTEMATICS

As noted earlier, the collection of embryological features can be very valuable in delimiting or aiding in phylogenetic inference. An example of embryological features in a group of angiosperms is portrayed in Table 11.1.

TABLE 11.1 Example of embryological features of the Caryophyllids, after Rodman, 1990. Centrospermae revisited, Part I. Taxon 39: 353–393. Character names are bolded in brackets.

Ovule anatropous, (ana-)campylotropous, or orthotropous [Ovule Type]
Micropyle formed by both integuments or inner only [Micropyle Type]
Nucellar beak (or cap) absent or present [Nucellus type]
Embryogeny: onagrad, asterad, solanad, caryophyllad, or chenopodiad type [Embryogeny]
Embryo sac development Polygonum-type or Plumbago-type [Female Gametophyte (Embryo Sac) type]
Funicular obturator absent or present [Obturator Presence/Absence]
Endosperm present or absent in seed [Seed Storage Tissue Origin]
Perisperm not persisting or persisting in seed [Seed Storage Tissue Origin]
Embryo without chlorophyll or with chlorophyll [Embryo type]
Seeds arillate (funicular aril) or not [Seed coat anatomy]
Exotestal layer of seed not thickened or thickened [Seed coat anatomy]
Endotestal layer of seed thickened or not [Seed coat anatomy]
Exotegmic layer of seed tracheidal, fibrous or palisade, or unspecialized [Seed coat anatomy]
Endotegmic layer of seed not thickened or thickened [Seed coat anatomy]

EMBRYOLOGICAL TECHNIQUE

Material Dissection and Preparation:

As with anatomical studies, the study of plant embryology can involve very time-consuming embedding, microtome sectioning, staining, and slide preparation. Often, it is critical to obtain a range of developmental stages, in order to trace the changes that occur from inception to maturity. Female gametophyte development is particularly difficult to study, as some developmental changes occur rapidly and are hard to catch.

For all embryological studies, plant material must first be *fixed* in a chemical solution. The *fixative* preserves the material close to its original state and often clears it somewhat such that it can be better resolved under the microscope. Fix a number of flowers, from very young buds to developing fruits, by placing the material in a jar or vial of 70%–95% ethanol. The buds and flowers (or large ovaries or anthers) should generally be cut open to allow for better penetration of the fixative. Fix for a minimum of 10–15 minutes, although 1–2 days is better; store in 70% ethanol. [Note that, for more detailed studies, the material should be fixed in *FAA*, which is a mixture of formalin, acetic acid, alcohol. One recipe for *FAA* is: 66 ml of 95% ethanol, 21 ml water, 8 ml commercial (37%) formalin, and 5 ml glacial acetic acid. Formalin is dangerous to inhale, and glacial acetic acid is very caustic, so the solution should be mixed very carefully in a laboratory hood. Once mixed, it can be stored indefinitely, in a properly sealed container.]

Ovule Morphology:

Many features of the ovule can be observed with some relatively simple techniques. Place mature flowers that have been chemically fixed in a small dish (Petri or Syracuse dish) filled with 70% ethanol. Dissect the material with needles and forceps to remove and open up the ovary. Use fine needles to detach the ovules. During the dissection, observe the general ovule type (e.g., is the micropyle pointing toward the point of attachment of the funiculus [anatropous] or away from it [orthotropous]) and ovule position (the placement of the micropyle and funicular raphe relative to the flower axis). Place some ovules (using forceps or a pipette) on a microscope slide in a drop of water or 50% glycerol and cover with a cover slip.

For more detailed studies, the ovules can be cleared and observed using phase contrast or (preferably) differential interference contrast (DIC or Nomarski) optics. One useful clearing fluid is Herr's solution. [A recipe is 1 part 85% lactic acid : 1 part chloral hydrate : 1 part phenol crystals : 1 part clove oil : 100 part xylene (all parts by weight; after Herr, J. M., Jr. 1971. *American Journal of Botany* 58: 785–790.)]

Microspore and Pollen Development:

One simple technique to observe the development of pollen grains is to dissect the internal contents of fixed anthers onto a microscope slide, stain, and cover with a cover slip. The material may be stained with toluidine blue or (to observe

meiotic stages) acetocarmine. After staining, the material may be squashed by placing a cork on top of the cover slip and applying gentle, firm pressure. Squashing spreads the cells out into a thin layer, allowing for better observation of cell divisions and morphology.

REVIEW QUESTIONS

ANTHER AND POLLEN DEVELOPMENT

1. What are the two major anther types and how do they differ?
2. What criteria are used to define the four types of anther wall development?
3. What is the tapetum? What are the two types of tapetum development and how do they differ?
4. What are two types of anther endothecial anatomy?
5. What is microsporogenesis and what are the two major types?
6. What is microgametogenesis and what are the two major types?

OVULE DEVELOPMENT

7. Name the parts of a typical ovule.
8. What is meant by the chalazal region? a raphe?
9. Name and distinguish between the three types of nucellus. Which two resemble one another at maturity?
10. Name and distinguish between the three types of megasporogenesis.
11. What criteria are used to distinguish between the numerous female gametophyte development types?
12. Which female gametophyte type is most common and probably ancestral in the angiosperms?
13. What are the two major integument types?
14. What are the differences between endostomal, exostomal, amphistomal, unistomal, and zig-zag micropylar types?
15. What criteria are used to distinguish between ovule types? Which type is most common and ancestral in the angiosperms?
16. Define and give three examples of ovule position.

SEED DEVELOPMENT

17. What is embryogeny and on what criteria are different embryogeny types based?
18. What is ptyxis and what does it define?
19. Name the three basic types of endosperm development and describe how they differ.
20. Other than endosperm, what two other seed storage tissue origins occur in angiosperms?
21. Name four seed storage tissue origin types. Of what three major chemicals are seed storage tissue composed?
22. What are arils and caruncles, and what is their function?

EMBRYOLOGY AND SYSTEMATICS

23. Name some features of embryology that may be valuable in plant systematics.

EXERCISES

1. Obtain flowering material of a species and fix the material according to the procedures noted above (Embryological Technique). Dissect ovules from the ovaries and prepare a slide for light microscope observations. If possible, clear the ovules for phase contrast or differential interference contrast (DIC) microscopy. Note, from the dissected or cleared ovules, the (a) ovule position (epitropous, hypotropous, pleurotropous (and whether dorsal or ventral), or heterotropous; (b) integument type (bitegmic or unitegmic); (c) female gametophyte shape; (d) vasculature; and (e) specific ovule type (ana-amphitropous, ana-campylotropous, ortho-amphitropous, or ortho-campylotropous). Draw and record this information.

2. Obtain anthers of various stages and dissect them open to make slide preparations. Stain with acetocarmine. Observe stages of meiosis and pollen development.
3. Obtain mature seeds of various flowering plants. Observe outer components of the seed, including seed coat morphology, funicular scar, raphe, caruncle, or aril (if present). Dissect the seeds by cutting with a razor blade. Observe the embryo and the seed nutritive type (endospermous or albuminous, exalbuminous, cotyledonous, or perispermous). Stain the seed sections with IKI, which stains starch a dark purple or brown, to determine if the nutritive tissue is starchy at maturity.
4. Peruse journal articles in plant systematics, e.g., *American Journal of Botany*, *Systematic Botany*, or *International Journal of Plant Sciences*, and note those that describe plant embryological features in relation to systematic studies. Identify all embryological characters and character states described.

REFERENCES FOR FURTHER STUDY

- Bjrnstad, I. N. 1970. Comparative embryology of Asparagoideae-Polygonateae, Liliaceae. *Nytt Magasin for Botanikk* 17(3 4): 169–207.
- Davis, G. L. 1966. *Systematic Embryology of the Angiosperms*. John Wiley and Sons, New York.
- Johri, B. M. (ed.). 1984. *Embryology of angiosperms*. Springer-Verlag, Berlin, New York.
- Johri, B. M., K. B. Ambegaokar, and P. S. Srivastava. 1992. *Comparative embryology of angiosperms*. Springer-Verlag, Berlin, New York.
- Maheshwari, P. 1950. *An Introduction to the Embryology of Angiosperms*. McGraw-Hill, New York.
- Maheshwari, P. (ed.). 1963. *Recent Advances in the Embryology of Angiosperms*. International Society of Plant Morphologists, Delhi.

This Page Intentionally Left Blank

12

PALYNOLOGY

INTRODUCTION	453	Pollen Sculpturing	458
PALYNOLOGICAL TERMINOLOGY	453	Pollen Wall Structure	460
Pollen Nucleus Number	453	Pollenkit and Viscin Threads	462
Pollen Storage Product	453	PALYNOLOGY AND SYSTEMATICS	462
Pollen Unit	453	PALYNOLOGICAL TECHNIQUE	462
Pollen Polarity	455	REVIEW QUESTIONS	462
Pollen Aperture	455	EXERCISES	463
Pollen Symmetry	458	REFERENCES FOR FURTHER STUDY	463
Pollen Size	458		
Pollen Shape	458		

INTRODUCTION

Palynology (Gr. *palynos*, dust) is the study of spores and pollen grains. Spores and pollen grains have a number of morphological and ultrastructural features. These palynological features have provided a wealth of characters that have been important in inferring phylogenetic relationships of plants. In addition, the features of spores and pollen grains can often be used to identify a particular plant taxon. For this reason, palynological studies are used extensively to examine the fossil record, a field called paleo-palynology. The identity, density, and frequency of pollen grains at a particular stratigraphic level can give information as to the plant species present at that time and place. Paleo-palynological studies are thus used to determine plant community structure and to gauge, by extrapolation over time, shifts in climate.

PALYNOLOGICAL TERMINOLOGY

The terminology applied to pollen morphology and ultrastructure varies from author to author. The following terminology follows the suggestions of Reitsma (1970), Walker and Doyle (1975), and Punt et al. (1994).

POLLEN NUCLEUS NUMBER

The number of nuclei at the time of pollen release can be phylogenetically informative. Two types occur in angiosperms. **Binucleate** grains (Figure 12.1) contain one tube cell and nucleus and one generative cell and nucleus; this is the most common and ancestral type in the angiosperms. **Trinucleate** grains contain one tube cell and nucleus and two sperm cells, the latter resulting from precocious division of the generative cell. Trinucleate grains are relatively rare in the angiosperms, being a diagnostic feature and possible apomorphy for some Caryophyllales. Pollen nuclear number is also listed as an embryological character (see Chapter 11).

POLLEN STORAGE PRODUCT

Pollen grains contain high-energy storage reserves. These are composed of either **starch** or **oil**. The distribution of storage product type can be phylogenetically informative in the angiosperms.

POLLEN UNIT

Pollen unit refers to the number of pollen grains united together at the time of release. Most commonly, the four microspores formed after microsporogenesis separate prior to pollen (or spore) release. Such single, unfused pollen grains are called **monads**, found in the great majority of angiosperms.

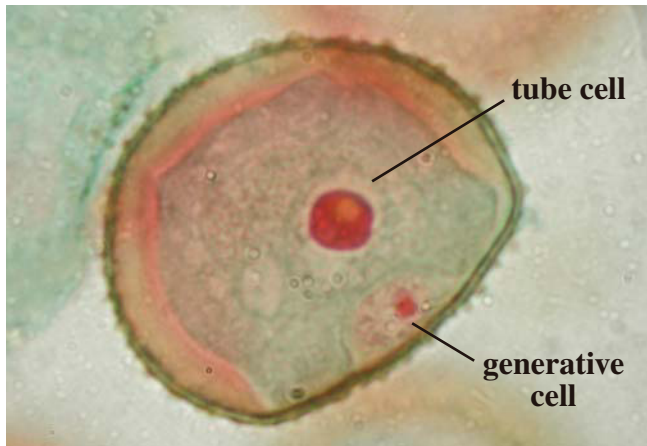


FIGURE 12.1 Binucleate pollen nuclear number. Note generative cell and tube cell nuclei.

Rarely, pollen grains will fuse in pairs, each pair known as a **dyad**. More commonly, the four haploid products of meiosis remain fused together, comprising a **tetrad**. Five types of tetrads are recognized, based on the arrangement of pollen grains: (1) **tetrahedral tetrads** (Figure 12.2A), in which the four grains form the points of a tetrahedron, e.g., as in members of the family Ericaceae; (2) **linear tetrads**, in which the four pollen grains are arranged in a straight line, e.g., as in *Typha* spp.; (3) **rhomboidal tetrads**, in which the four grains are in one plane, with two of the grains separated from one another by the close contact of the other two; (4) **tetragonal tetrads** (Figure 12.2B), in which the four grains are in one plane and are equally spaced apart; and (5) **decussate tetrads** (Figure 12.2C), in which the four grains are in two pairs arranged at right angles to one another.

Pollen grains that are connate in precise units of more than four are called **polyads** (Figure 12.2D). Polyads are common in the Mimosoideae of the Fabaceae and generally consist of a multiple of eight fused grains. Fusion of pollen grains in large, often irregular numbers, but less than an entire theca,

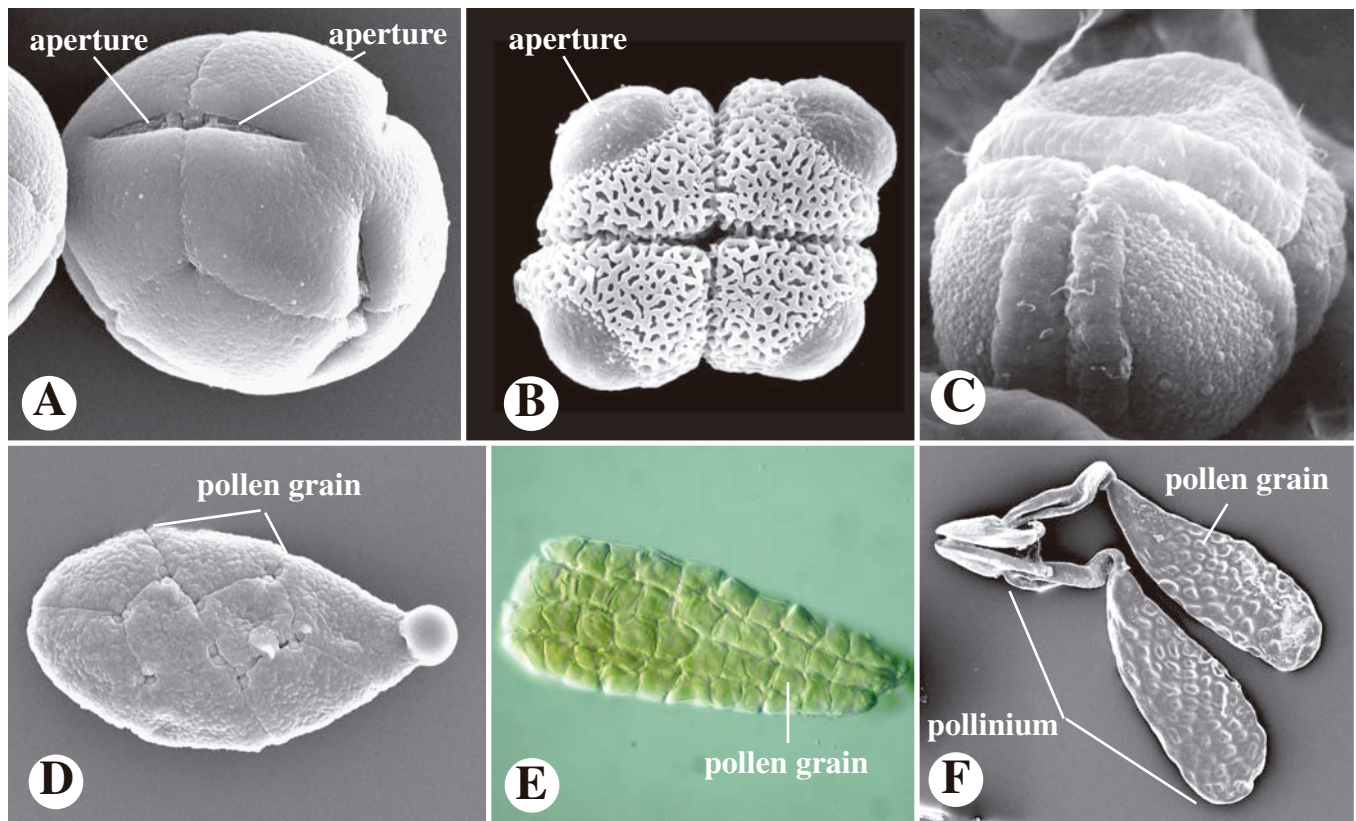


FIGURE 12.2 Pollen unit. **A.** Tetrahedral tetrad, *Arbutus*, Ericaceae. **B.** Tetragonal tetrad, *Philydrum*, Philydraceae. **C.** Decussate tetrad, early developmental stage, *Lachnanthes*, Haemodoraceae. **D.** Polyad of eight fused pollen grains, *Calliandra*, Mimosoideae, Fabaceae. **E.** Massula of *Piperia*, Orchidaceae. **F.** Pollinium of *Asclepias*, Apocynaceae.

are called **massulae** (singular **massula**) (Figure 12.2E). Finally, the fusion of all pollen grains of an entire theca is called a **pollinium** (plural **pollinia**), found in the families Apocynaceae (Figure 12.2F) and Orchidaceae (Chapter 7).

POLLEN POLARITY

Pollen polarity refers to the position of one or more apertures (see later discussion) relative to a spatial reference. This spatial reference defines a **polar axis** as the extended pollen grain diameter that passes through the center of the original pollen tetrad (Figure 12.3). The intersection of the polar axis with the grain surface near the center of the tetrad is the **proximal pole**, the surrounding area being the proximal face or proximal hemisphere; that away from the tetrad center is the **distal pole**, the surrounding area being the distal face or distal hemisphere. Just as with a globe, the intersection with the pollen surface of a plane at a right angle to the pole and passing through the center of the grain defines the pollen **equator**, the surrounding area being the equatorial region. Observing a pollen grain from the direction of either pole is

known as a **polar view**; observing from the equatorial direction is an **equatorial view** (Figure 12.3).

The three general types of pollen polarity are (1) **isopolar**, in which the two polar hemispheres are the same but can be distinguished from the equatorial region; (2) **heteropolar**, in which the two polar hemispheres are different, because of differential displacement of one or more apertures; and (3) **apolar**, in which polar and equatorial regions cannot be distinguished after pollen grain separation from the tetrad. Note that pollen polarity is with reference to the microspore or pollen tetrad. Unless the mature pollen unit is a tetrad (above), pollen grain polarity can be directly determined only by observing the position of apertures during the early tetrad stage. Because this is rarely observed, polarity is generally inferred by comparison with taxa for which polarity has been directly observed.

POLLEN APERTURE

A pollen **aperture** (Figures 12.4, 12.5) is a specially delimited region of the pollen grain wall. (See Pollen Wall Structure.)

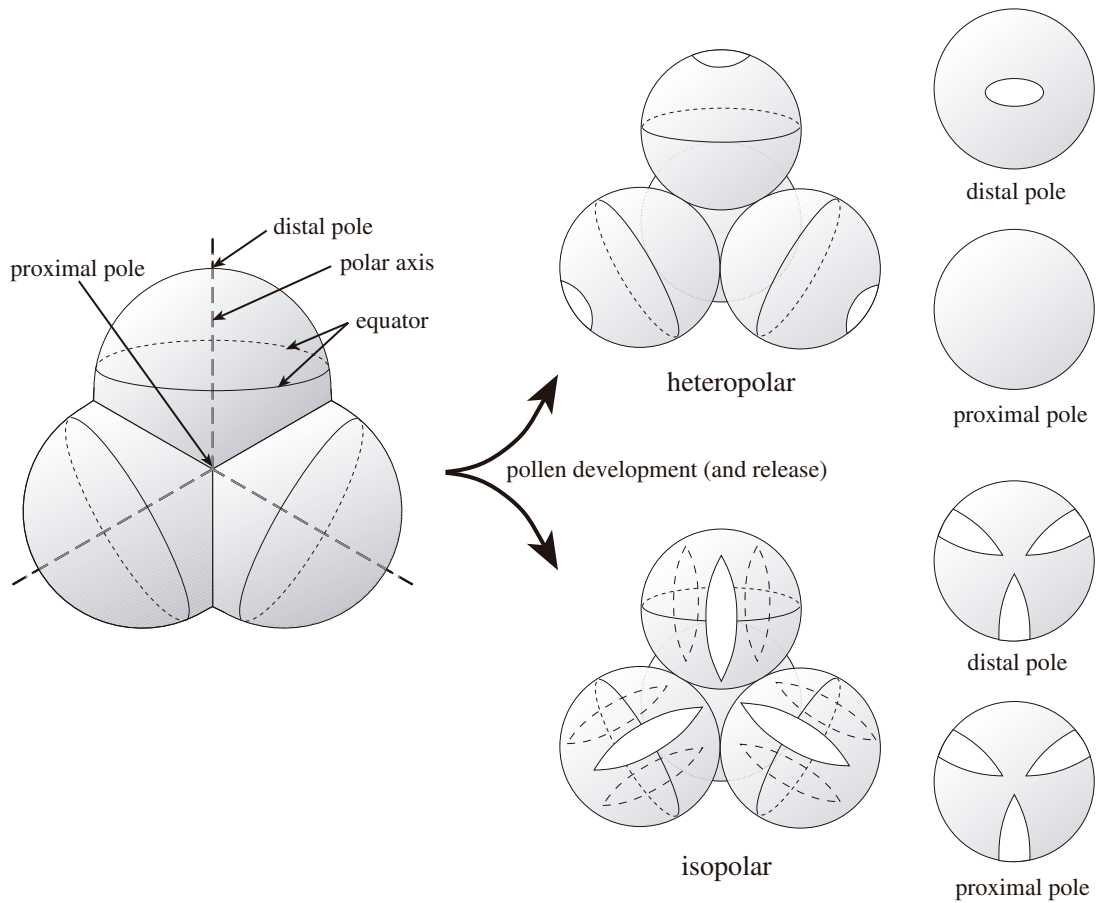


FIGURE 12.3 Pollen polarity.

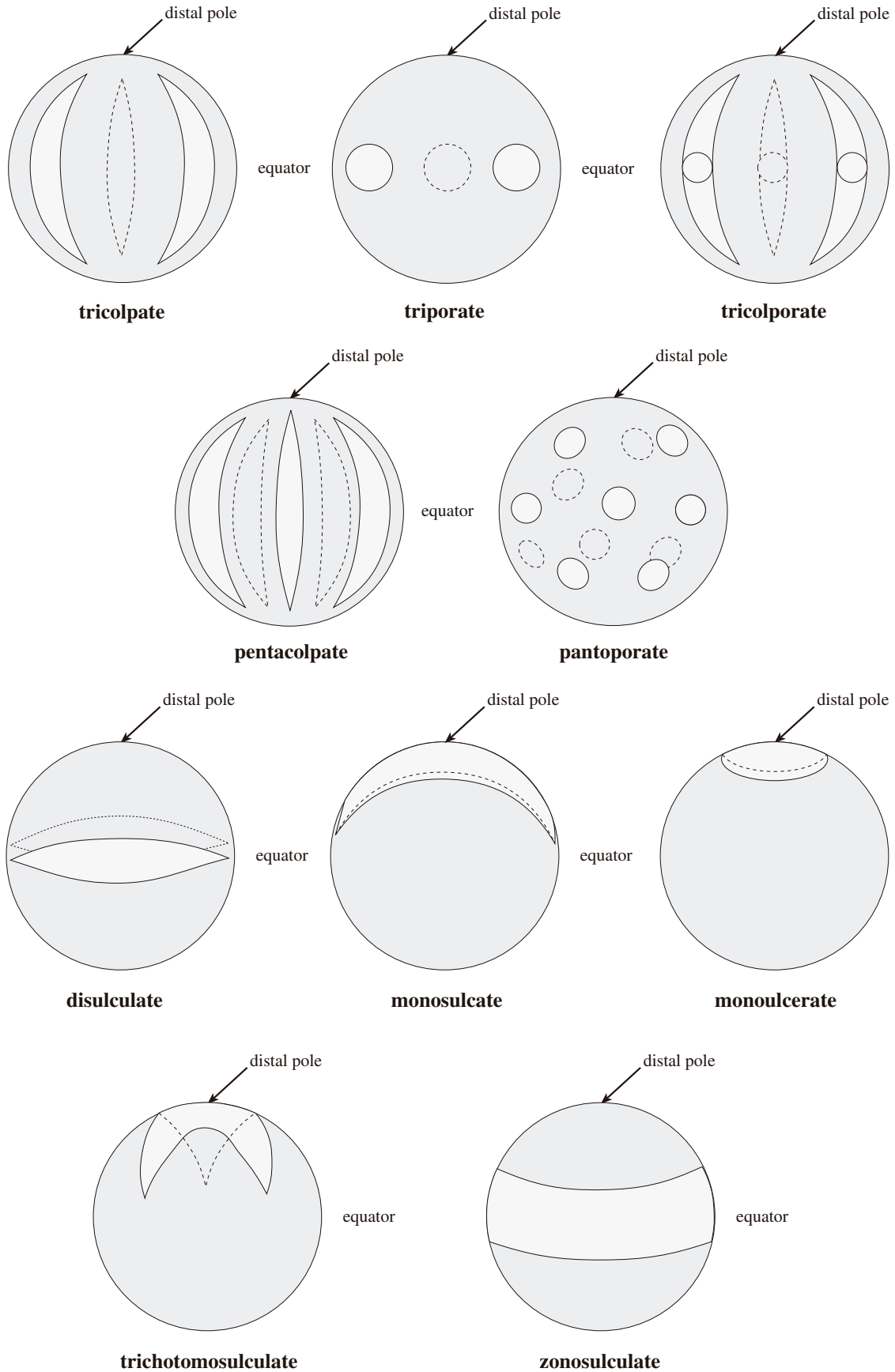


FIGURE 12.4 Pollen aperture.

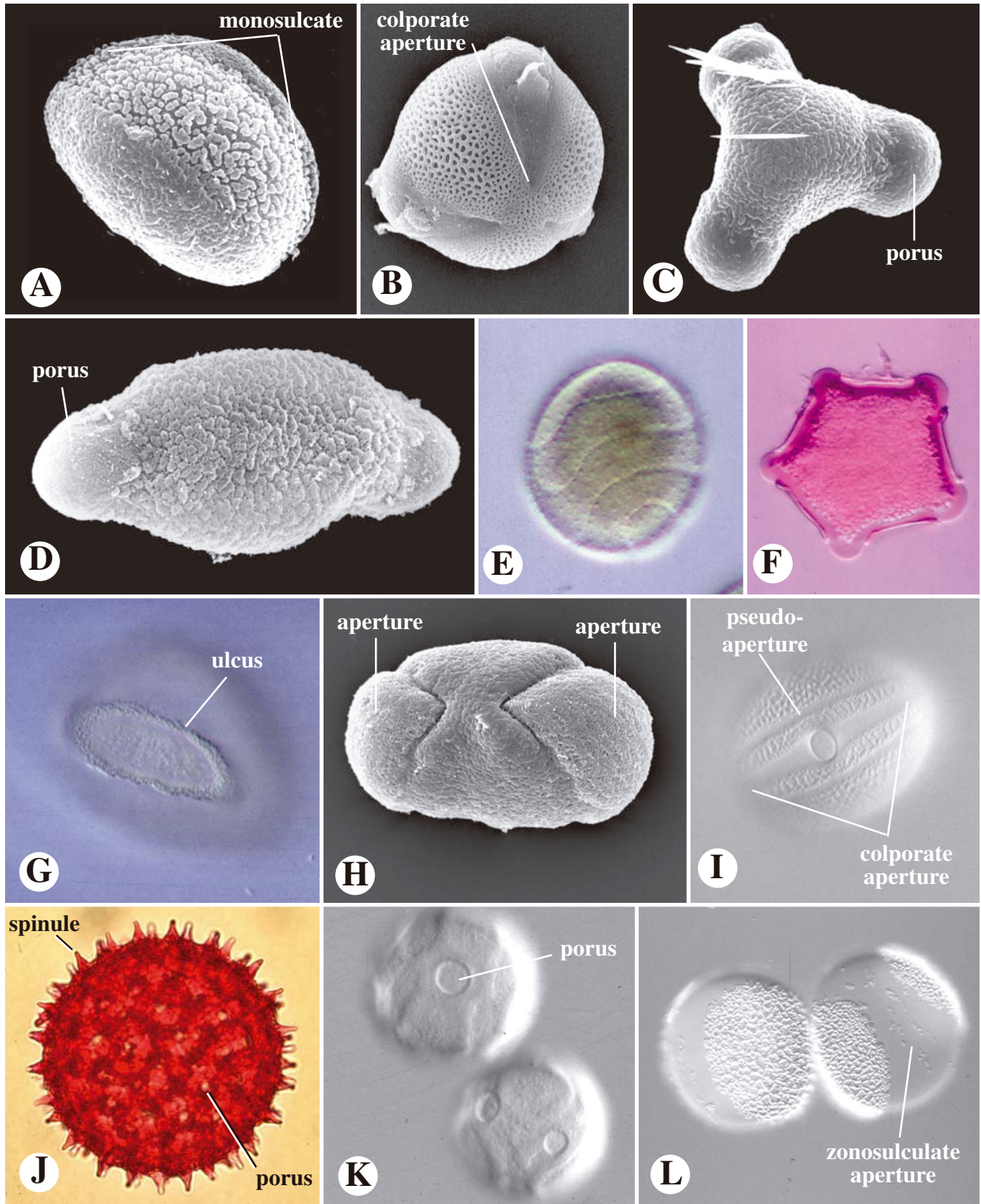


FIGURE 12.5 Pollen aperture examples. A. Monosulcate. B. Tricolporate. C. Triporate. D. Diporate. E. Spiraperturate. F. Pentaporate. G. Ulcerate. H. Disulcate. I. Tricolporate (one aperture visible, with flanking pseudo-apertures). J. Pantoporate, with echinate sculpturing. K. Pantoporate. L. Zonosulcate.

The function of the aperture is primarily to serve as the site of formation of a pollen tube exiting from the pollen grain body. Apertures may also function to allow volume changes of the pollen grain with changes in water content, e.g., humidity. This feature is known as **harmomegathy**. Harmomegathy allows the pollen grain apertures to contract with water loss, effectively sealing the apertures via the surrounding desiccation resistant exine wall (see later discussion).

Pollen aperture type refers to the shape, number, position, and arrangement of the aperture(s) of a pollen grain, often with an implied reference to the polar axis. Rarely, pollen grains lack any recognizable aperture; these are termed **inaperturate**.

Two general types of apertures correspond to shape. A **colpus** (plural, colpi) is an elongate aperture with a length/width ratio of greater than 2:1 (Figure 12.4). Colpi can be elliptic, oblong, or fusiform in outline shape. A **porus** (plural, pori) is a circular to slightly elliptic aperture with a length/width ratio of less than 2:1 (Figures 12.4, 12.5C,D,F); if pori occur globally on the pollen grain surface, the aperture type is called **pantoporate** (Figures 12.4, 12.5J,K). An aperture that is shaped like a colpus but has a circular region in the center is termed **colporate** (Figures 12.4, 12.5B,I).

Pollen grains with apertures occurring in the equatorial region may be termed **zonoaperturate** (or stephanoaperturate), e.g., as in **zonocolpate** or **zonoporate**. The terms **colpus** and **porus** are often restricted to apertures occurring in a region of the pollen grain other than the poles, often in the equatorial region. By this terminology an elongate aperture similar in shape to a colpus (length/width ratio >2:1) occurring at the (usually distal) pole is called a **sulcus** (Figure 12.5A). Comparably, a circular to slightly elliptic aperture similar in shape to a porus (length/width ratio <2:1) occurring at the (usually distal) pole is called an **ulcus** (Figure 12.5G). **Disulcate** refers to a pollen grain with two elongate apertures on opposite sides of the grain, e.g., parallel to the equatorial plane (Figure 12.5H); **trisulcate** is similar but with three apertures.

The number of apertures of any shape can be designated by appending the prefix mono-, di-, tri-, tetra-, penta-, hexa-, or poly- (more than six) to the terms colpate or porate. Thus, a tricolpate pollen grain is one with three, elongate apertures occurring in the equatorial region. A pentaporate pollen grain is one with five, approximately circular apertures occurring in the equatorial region.

Some aperture types are rather rare and specialized. **Syncolpate** refers to a pollen grain in which the colpi are joined, e.g., at the poles. **Trichotomosulcate** refers to an aperture type that is three-branched. Sulcate and ulcerate pollen grains typically have only a single aperture; these

terms are usually equivalent to monosulcate and monoulcerate, respectively. **Spiraperturate** refers to one or more apertures that are spirally shaped (Figure 12.5E).

POLLEN SYMMETRY

Pollen symmetry is generally either radially symmetric, i.e., with two or more planes of symmetry, or bilaterally symmetric, with a single plane of symmetry. Symmetry is often incorporated or assumed as part of a shape term (see Pollen Shape).

POLLEN SIZE

Pollen size can vary tremendously across taxa. Size is typically measured in terms of *both* the polar diameter and the equatorial diameter (see Pollen Shape).

Typical pollen grains are ca. 25–50 μm in diameter, but pollen diameter can range from < 5 μm (approaching the size of some bacteria!) to > 200 μm .

POLLEN SHAPE

Pollen shape (Figure 12.6) may refer to the three-dimensional shape of a pollen grain; e.g., **boat-shaped**, **ellipsoid**, **fusiform**, or **globose/spheroidal**. Shape may also be assessed by the two-dimensional outline shape either in polar view or equatorial view, e.g., as viewed by focusing under a light microscope. The outline shape in polar view is known as **amb**. Amb can be nonangular, e.g., circular, elliptic, or angular, e.g., triangular, rhombic, rectangular, five-angled. For angular amb, the shape of the sides may be described as straight, concave, or convex (Figure 12.6).

Another measure of pollen shape is the ratio of the polar diameter to the equatorial diameter, termed the **P/E ratio**. If the P/E ratio is approximately equal to 1, the grains are termed **spheroidal**. If P/E is > ca. 1.2, the grains are termed **prolate** (i.e., elongate along the polar axis, like a cucumber); if < ca. 0.8, the grains are **oblate** (compressed along the polar axis, like a tangerine). (The subcategories **prolate-spheroidal** and **oblate-spheroidal** are sometimes used for grains that are slightly prolate or oblate, respectively.)

POLLEN SCULPTURING

Pollen sculpturing (Figure 12.7) refers to the *external* features of the pollen grain wall. Sculpturing features may be viewed with light microscopy, but much greater detail can be detected with scanning electron microscopy.

Specialized pollen sculpturing terms include:

baculate, having rod-shaped elements, each element termed a baculum, plural baculi;

clavate, having club-shaped elements, each element called a clava, plural clavae;

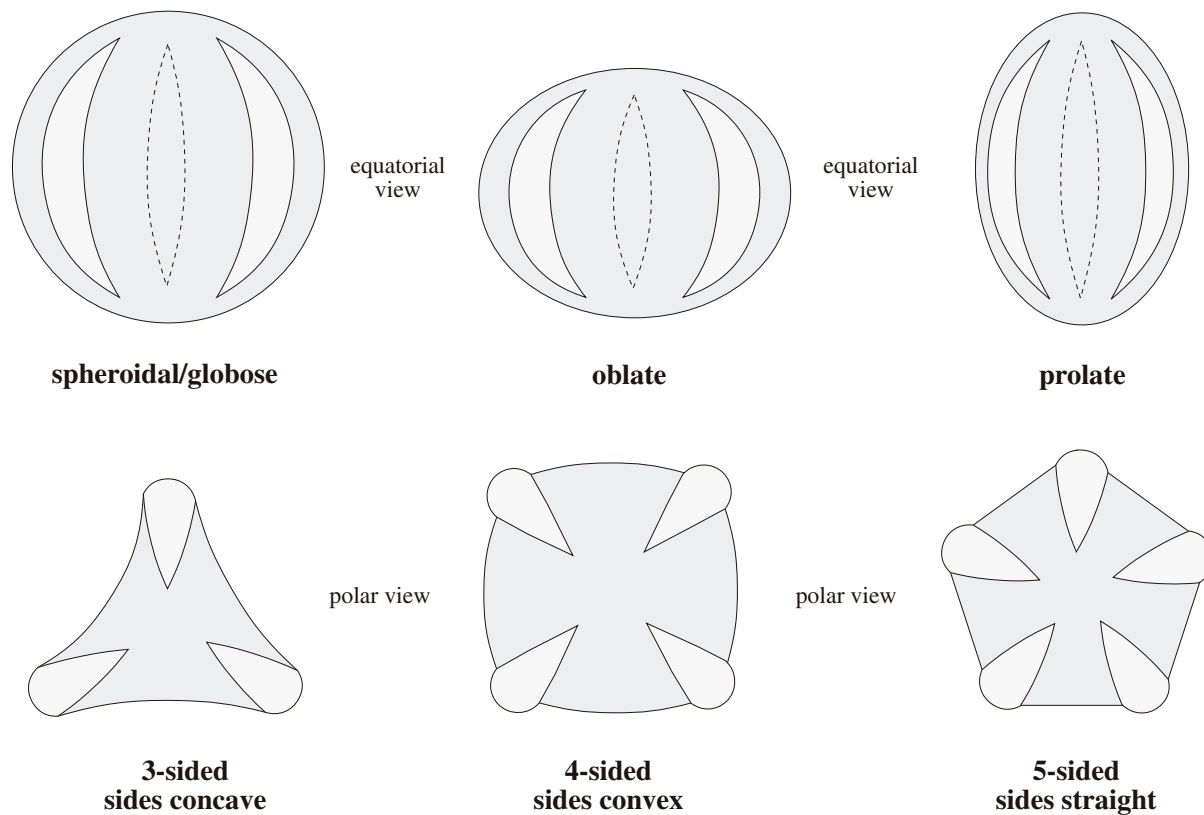


FIGURE 12.6 Pollen shape.

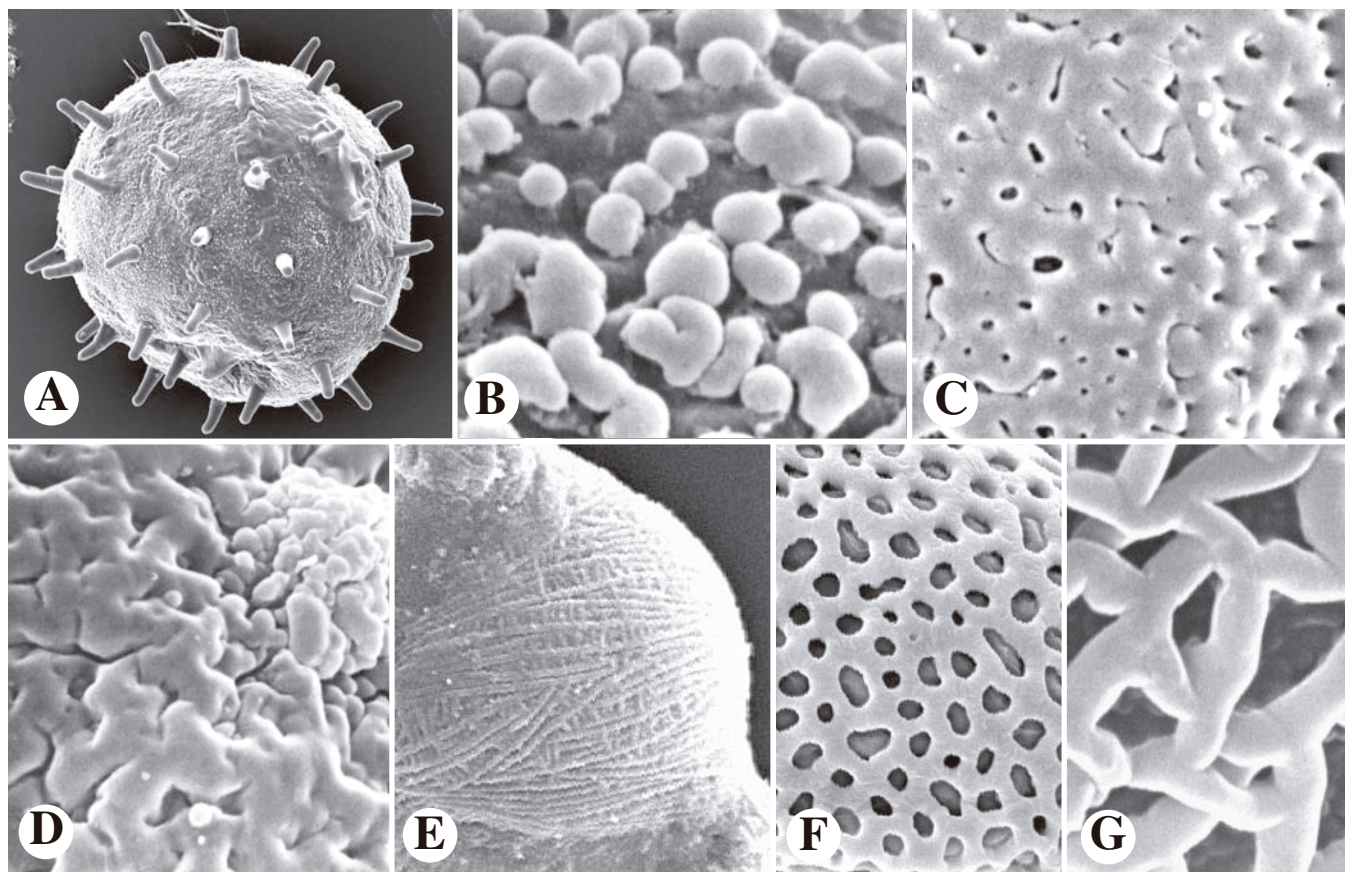


FIGURE 12.7 Pollen sculpturing. A. Echinate. B. Verrucate. C. Foveolate. D. Rugulose. E. Striate. F,G. Reticulate.

- echinate**, having spinelike elements $> 1 \mu\text{m}$ long, each element termed an echina, plural echinae;
fossulate, having longitudinal grooves.
foveolate, having a pitted surface caused by pores in the surface;
gemmate, having globose or ellipsoid elements, each element termed a gemma, plural gemmae;
psilate, having a smooth sculpturing;
reticulate, having a netlike sculpturing, each element termed a murus (plural muri) and the space between muri termed a lumen (plural lumina);
rugulate, having irregular to sinuous, tangentially oriented elements, often appearing brainlike;
spinulose (also termed **scabrate**), having spinelike elements $< 1 \mu\text{m}$ long, each element termed a spinulum, plural spinuli;
striate, having thin, cylindrical, tangentially oriented elements;
verrucate, having short, wart-like elements, each element termed a verruca, plural verrucae.

POLLEN WALL STRUCTURE

The pollen grain wall functions primarily to provide structural support and protection of the cytoplasm from mechanical damage and dessication. The wall may also function to facilitate pollination. For example, entomophilous (insect-pollinated) flowers tend to have elaborately sculptured pollen; these sculpturing elements may function to attach pollen grains to one another in masses and to appendages on the insect. Anemophilous (wind-pollinated) flowers tend to be smooth (psilate), functioning as a more efficient aerodynamic mechanism for wind transport.

The pollen grain wall may also function to store proteins involved in incompatibility reactions. Sporophytic incompatible taxa tend to have incompatibility proteins stored in cavities of the exine, derived from the sporophyte tapetum.

Gametophyte incompatible taxa tend to have incompatibility proteins stored in the intine, derived from the microspore/pollen cytoplasm.

Pollen wall structure (Figure 12.8) refers to the *internal* form of the pollen grain wall. Early in development, microspores typically have a thick cell wall composed of callose, the same substance that lines the pores of sieve elements. During pollen development, however, the callose wall breaks down completely. Mature pollen walls almost always consist of two major layers: intine and exine. The **intine** is the innermost layer, which is composed primarily of cellulose and pectines, resembling the primary cell wall of a typical parenchyma cell. The **exine** is the hard, outermost, desiccation-resistant wall layer that provides the major structural support for the cytoplasm. Exine is impregnated with a substance called **sporopollenin**, a polymer of oxidative carotenoids and/or carotenoid esters. Sporopollenin is very tough and resistant to mechanical damage and decay. The presence of sporopollenin accounts for the fact that pollen grains may often be preserved in the fossil record. The sporopollenin-impregnated exine is also resistant to **acetolysis**, which is a standard acid treatment used to dissolve all but the exine in order to better observe pollen wall structure with the light microscope.

The exine of many taxa may be divided into two layers, an inner **endexine** and an outer **ektexine**. These two layers differ chemically and have different staining properties as viewed with transmission electron microscopy. The endexine typically forms a more or less homogeneous, inner layer. The ektexine may exhibit a variety of structural forms. The most common type of ektexine in angiosperms is termed **tectate-columellate** (Figures 12.8, 12.9) and consists of an inner **foot layer**, a middle layer of radially elongate **columellae**, and an outer, rooflike layer called the **tectum**. In some taxa, the middle layer (given the generalized term *interstitium*) may not be

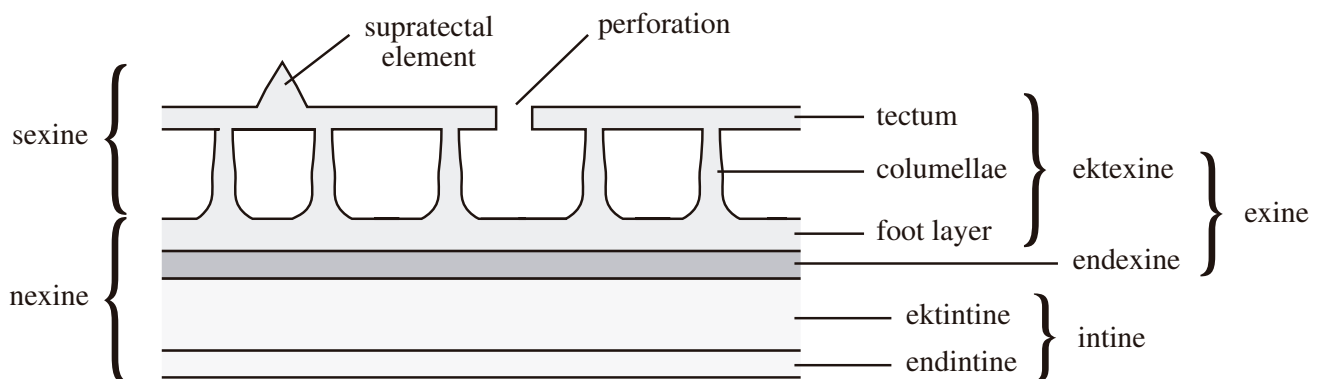


FIGURE 12.8 Pollen wall structure.

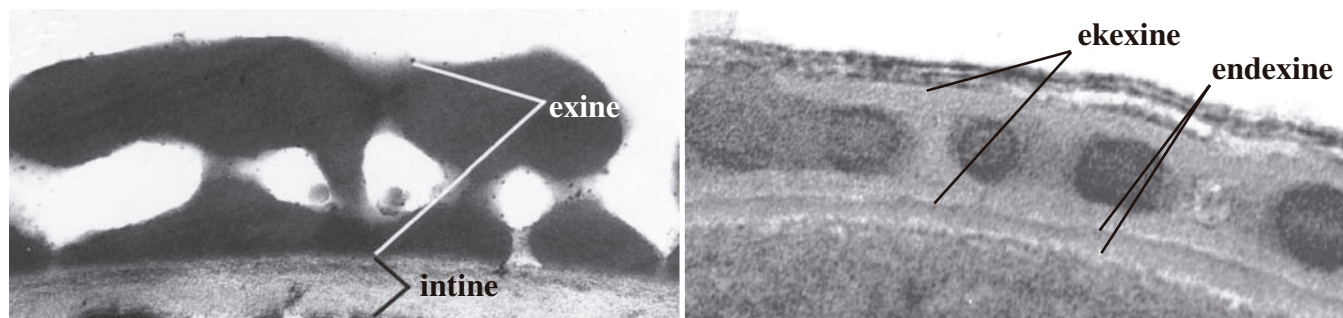


FIGURE 12.9 Example of tectate-columellate wall structure.

composed of columellae, but may instead consist of irregular or granular elements.

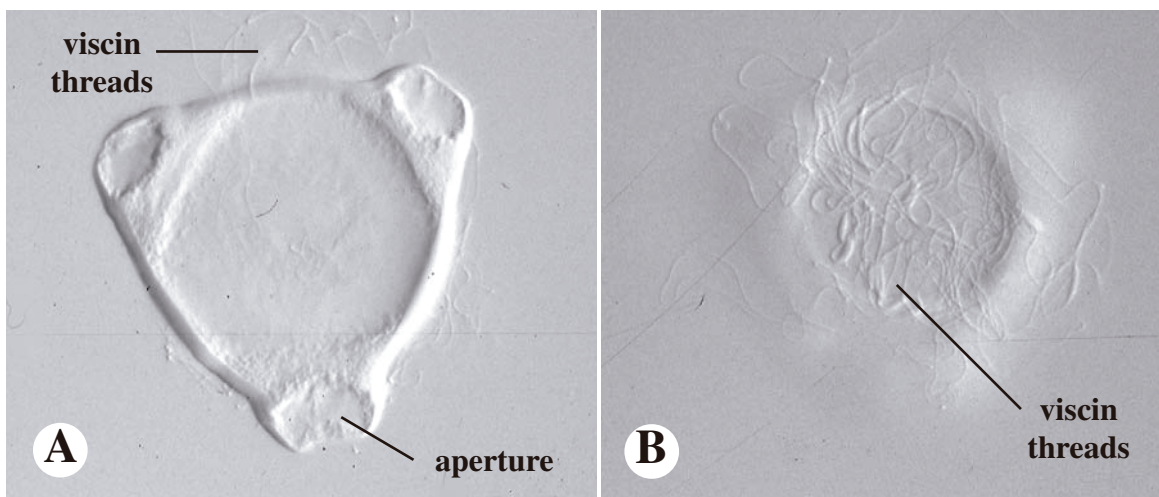
A tectate-columellate wall structure that lacks pores or perforations in the tectum is termed tectate-imperforate. In some taxa, the tectum contains tiny pores, a structure known as semi-*tectate*. A semi-*tectate* structure typically corresponds with a foveolate sculpturing type. A wall structure in which the tectum has large openings is called tectate-perforate. This may correspond, e.g., with a reticulate sculpturing.

Exinous elements on top of the tectum (described as supratectal) may account for sculpturing types such as baculate or echinate. However, in some taxa, a tectum may be absent; in these taxa protruding sculpturing elements such as baculae or echinae may be homologous to modified columellae. Only by viewing the wall structure internally may these differences be noted.

In addition to the tectate-columellate wall structure, various angiosperm taxa may have a wall that ancestrally lacks

a tectum, termed an **atectate** wall structure. The exine wall of atectate taxa may be structurally solid, termed **homogeneous**, or **granular**, containing small, granular elements with intervening air spaces. Additional types of exine wall structure include **lamellar**, having stacked, tangentially oriented, planar structures, often constituting the inner wall layer; and **alveolar**, having numerous, spherical air pockets within the exine.

The layers of the exine can be precisely observed only using transmission electron microscopy. However, because many aspects of pollen wall structure may be observed using light microscopy, the terms **nexine** and **sexine** are sometimes applied to describe exine wall layers (Figure 12.8). **Nexine** refers to the inner layers, which may include both endexine and the foot-layer of the *ektexine*. **Sexine** refers to the outer, protruding layers, which may include columellae, tectum, and supratectal sculpturing elements (if present).

FIGURE 12.10 *Fuchsia* sp. (Onagraceae), having triangular, triporate pollen grains with viscin threads (arrow). **A.** Equatorial region in focus. **B.** Polar region in focus.

POLLENKIT AND VISCIN THREADS

Pollenkit is a yellowish or orange, carotenoid-like material adhering to the exine. It functions to stick pollen grains in masses, better effecting transfer of pollen by animal (esp. insect) pollinators.

Viscin threads (Figure 12.10) are long strands of carbohydrate material that, like pollenkit, function in sticking pollen grains together.

PALYNOLOGY AND SYSTEMATICS

Palynological features have been very valuable in delimiting taxa or aiding in phylogenetic inference. An example of palynological features in a group of angiosperms is portrayed in Table 12.1.

TABLE 12.1 Example of palynological features of the Caryophyllales (Caryophyllidae), after Rodman, 1990. Centrospermae revisited, Part I. Taxon 39: 353-393. Character names are in brackets.

Pollen shed in binucleate or trinucleate condition [Pollen nuclear number]
Plastid-DNA not transmitted through pollen or transmitted
Aperturate pollen with furrows or not [Pollen aperture]
Aperturate pollen without pores or with pores [Pollen aperture]
Apertures simple or compound [Pollen aperture]
Number of apertures basically three, 4, 7, or >7 [Pollen aperture]
Zonocolpate apertures (>3) absent or present [Pollen aperture]
Pollen surface not spinulose or spinulose [Pollen sculpturing]
Pollen surface punctate/perforate or not punctate/perforate [Pollen sculpturing]
Pollen surface not reticulate or reticulate [Pollen sculpturing]

PALYNOLOGICAL TECHNIQUE

Material preparation:

For studies of pollen morphology, it is best to obtain living material of anthers, just at the time they are dehiscing, and fix these in a chemical solution, such as alcohol or FAA (see Chapter 10). [For transmission electron microscope studies, other fixatives, such as gluteraldehyde, formalin, or osmium tetroxide are used.] Collect plenty of material, and store in vials.

Light microscopic observations:

Pollen grains can be observed simply by making a wet mount on a microscope slide. A single anther can be removed from the fixative material, placed in a drop of water or 50% glycerol (the latter to prevent the material from drying out), and dissected with needles to extrude the pollen grains; the anther wall material should then be removed and a cover slip applied.

In addition, the pollen can be stained with either toluidine blue or acetocarmine, in order to better visualize details of the apertures and wall sculpturing. Simply dissect the anthers in a drop of stain, remove the anther wall, and add a cover slip.

Another technique is to clear the pollen grains in a clearing solution and visualize them using phase contrast or differential interference contrast (DIC, also called Nomarski) optics. A useful clearing solution is called Hoyer's clearing fluid. (Recipe: soak 30 g arabic gum lump in 50 g of water for 24 hours; add 200 g chloral hydrate (note: a controlled substance) until all the material dissolves; then add 20 g glycerine.) Dissect the anthers in a drop of Hoyer's as before, add a cover slip, and observe under phase contrast or DIC optics. The pollen grains may need time to clear, but once they do, you can visualize many details of the wall and apertures.

The presence or absence of starch in pollen grains can be examined by staining the pollen with IKI stain; starch changes to a dark blue or black in the presence of this stain. In addition, pollen grains can be mounted in 50% glycerol and viewed with polarization optics; starch grains are birefringent and show a Maltese cross type pattern under polarized light.

REVIEW QUESTIONS

1. What is the study of spores and pollen called?
2. What are the two types of pollen nuclear number? of pollen storage product?
3. What does pollen unit refer to?
4. What is the difference between a monad, tetrad, polyad, and pollinium?
5. What is pollen polarity?
6. What is the difference between an isopolar and heteropolar pollen grain?

7. What is the definition and function of a pollen aperture?
8. What is the difference between a colpus, porus, sulcus, and ulcerus?
9. What is a tricolporate pollen grain? a pentaporate pollen grain?
10. What is the size range of angiospermous pollen grains?
11. Name and define six terms that specify pollen sculpturing.
12. Name three functions of the pollen grain wall.
13. What are the two major layers of a pollen grain wall and how do they differ in chemical composition?
14. Name the two layers of exine.
15. Name and describe the most common type of exine wall structure.
16. What do nexine and sexine refer to?
17. What is the function of pollenkit or viscin threads?

EXERCISES

1. Using the simple procedures described earlier (Palynological Technique), examine pollen grains of various groups of angiosperms, including Magnoliids, Monocots, and several Eudicots, including a member of the Ericaceae (with permanent tetrads). Tabulate the differences in pollen unit, aperture type, aperture number, sculpturing type. Also, note the presence or absence of starch in the pollen grains.
2. Peruse journal articles in plant systematics, e.g., *American Journal of Botany*, *Systematic Botany*, or *International Journal of Plant Sciences*, or in specific palynological journals such as *Grana* or *Pollen et Spores* (see Appendix 3: Scientific Journals in Plant Systematics). Note those that describe palynological features in relation to systematic studies. Identify all pollen characters and character states described.

REFERENCES FOR FURTHER STUDY

- Erdtman, G. 1966. *Pollen Morphology and Plant Taxonomy. Angiosperms*. Corrected reprint and new addendum. Hafner, New York.
- Faegri, K., and J. Iversen. 1964. *Textbook of Pollen Analysis*. 1964. Blackwell Scientific, Oxford.
- Hoen, P. *Glossary of Pollen and Spore Terminology*. Second and revised edition. <http://www.bio.uu.nl/~palaeo/glossary/glos-txt.htm>
(An excellent Web page for pollen terminology, based on Punt et al. 1994).
- Lewis, W. H., P. Vinay, and V. E. Zenger. 1983. *Airborne and Allergic Pollen of North America*. The Johns Hopkins University Press, Baltimore. [QK 658 L48 1983]
- Moore, P. D., J. A. Webb, and M. E. Collinson. 1991. *Pollen Analysis*, 2nd ed. Blackwell Scientific, Oxford.
- Nilsson, S., and J. Praglowski (eds). 1992. *Erdtman's Handbook of Palynology*. Munksgaard International, Copenhagen.
- Punt, W., S. Blackmore, S. Nilsson, and A. le Thomas. 1994. *Glossary of Pollen and Spore Terminology*. LPP Foundation, Utrecht.
- Radford, A. E., W. C. Dickison, J. R. Massey, and C. R. Bell. 1974. *Vascular Plant Systematics*. Harper & Row, New York.
- Reitsma, T. 1970. Suggestions towards unification of descriptive terminology of angiosperm pollen grains. *Reviews in Palaeobotany and Palynology* 10: 39-60.
- Walker, J. W., and J. A. Doyle. 1975. The Bases of Angiosperm Phylogeny: Palynology. *Annals of the Missouri Botanical Garden* 62: 664-723.

This Page Intentionally Left Blank

13

PLANT REPRODUCTIVE BIOLOGY

SEXUAL REPRODUCTION	465	HYBRIDIZATION, INTROGRESSION, AND POLYPLOIDY	472
Flowering Plants	466	TESTING FOR BREEDING MECHANISM	474
Pollination Mechanisms	466	REVIEW QUESTIONS	475
Breeding Systems	468	EXERCISES	476
Seed and Fruit Dispersal	472	REFERENCES FOR FURTHER STUDY	476
ASEXUAL REPRODUCTION	472		

Plant reproductive biology is the study of the mechanisms and processes of sexual and asexual reproduction in plants. It may encompass study of pollination mechanisms, gene flow, genetic variation, and propagule dispersal between and within populations. A knowledge of the reproductive mechanisms of plants can help assess the adaptive significance and homology of descriptive characters used in plant systematics. Studying reproductive biology can also give insight into the delimitation and classification of species and infraspecies.

The following is a very abbreviated summary of the concepts and terms used in reproductive biology as they may be significant in studies of plant systematics.

SEXUAL REPRODUCTION

In nonseed plants, sexual reproduction entails the release of motile sperm from a free-living gametophyte into the outside environment. The sperm swims in a film of water into the neck of an archegonium, fertilizing the egg to form a zygote and then embryo. Completion of this phase of the life cycle is dependent on survivorship of the gametophytes, on the effective development and operation of antheridia and archegonia, and on the proper external conditions. The sporophytes of nonseed plants generally release massive numbers of spores into the environment, which are transported by wind or, more rarely, by water. These spores may, upon encountering

the proper environmental conditions, germinate and grow into a gametophyte, completing the cycle (see Chapters 3, 4).

In seed plants, separate male and female gametophytes are produced within male and female spores (microspores and megaspores). Sex involves the transfer of endosporic male gametophytes, the pollen grains, either to the micropyle of an ovule (in gymnosperms) or to the stigma of a pistil (in angiosperms). Sperm cells are ultimately released (into or just outside of the female gametophyte of the ovule), where one sperm cell fuses with the egg, initiating development of an embryo within the seed. Seeds are then transported by a variety of mechanisms to a new environment (see Chapters 5, 6).

Two major processes in sexual reproduction of seed plants, then, are **pollination**, the transfer of pollen grains from microsporangia to the ovule or stigma, and **fertilization**, union of sperm and egg. Many of the structural modifications of seed plants function in this transfer of pollen and the subsequent development and propagation of seeds.

In gymnosperms *cycads*, *Ginkgo*, conifers, and Gnetales pollen grains are almost entirely transported by wind. Because transport by wind is indirect, it necessitates the production of relatively large numbers of pollen grains to overcome the very low probability that any given pollen grain will make it to the ovule. In contrast, the great majority of angiosperms are animal (mostly insect) pollinated, which appears to be the ancestral condition for the family (Chapter 6), although wind

pollination has arisen secondarily in several groups of flowering plants (see later discussion).

FLOWERING PLANTS

Angiosperms have largely evolved very specialized floral structures that are adaptive in promoting animal pollination. Animal pollination is much more directed and precise, necessitating the synthesis of many fewer pollen grains to effect fertilization of the eggs within ovules.

The basic adaptive strategy of animal-pollinated flowering plants has been the evolution of an **attractant** and a **reward**. The attractant works to entice the animal to the flower, either by vision or by odor. A visual attractant is usually a showy perianth (corolla and/or calyx) that may be brightly colored or otherwise contrasting with the external environment, e.g., a white perianth at night. Other floral parts, such as stamens (e.g., *Hibiscus*), staminodes (e.g., members of the Aizoaceae, Cannaceae, or Zingiberaceae), corona (e.g., *Crinum*, *Narcissus*, *Passiflora*), or even the gynoecium, may replace or augment the perianth as a visual attractant. Individual flowers may actually be small, but the accumulation of flowers in an inflorescence may provide a significant visual attractant. Olfactory attractants include the volatile compounds emitted by flowers, usually from the surface of the perianth. Most odiferous flowers have a sweetish smell (e.g., *Jasminum*), but others emit compounds that mimic the smell of rotting flesh (e.g., *Aristolochia*, *Arum*, *Stapelia*).

Many species of flowering plants have evolved structures or exudates that act as a reward, ensuring that the animal pollinator will consistently return to transport pollen. The most common floral reward is nectar, a fluid primarily rich in sugars, secreted from specialized regions or organs of the flower called **nectaries** (Chapter 9). Nectaries are specialized tissues or organs that may be located within the gynoecium (e.g., the septal nectaries of many monocots), on the perianth, or at the base of and often surrounding the gynoecium or androecium. (Although nectar usually functions as a food source and reward for the prospective pollinator, some nectaries are extra-floral and may function as a reward for insects, such as ants, that protect the plant from herbivory by other animals.) Another pollination reward is pollen itself, which is a relatively rich source of protein. Some flowering plants produce waxes (e.g., *Krameria*) or oils as a reward. Finally, in some rare cases, insects may obtain specific chemical compounds that are used to attract a mate.

Although the general strategy of pollination in most flowering plants is to provide a reward (thus, presumably, increasing the fitness of both plant and animal), not all animal-pollinated flowers do this. Some flowers have evolved structures or mechanisms to trick the animal to transport pollen, possibly

with an adverse effect on the reproductive success of the animal. For example, in certain water lilies and orchids, the nectar may actually function to trap or even drown the insect to promote pollination. Other species of orchids actually mimic (visually and olfactorily) the female of an insect (usually a wasp), fooling the male to attempt to copulate with the flower, which, in the process, transports pollen.

POLLINATION MECHANISMS

Many, if not most, species of angiosperms have evolved specialized pollination mechanisms in which structural modifications are correlated with a specific agent of transferring pollen. Knowledge of the pollination agent can give insight into the function, homology, and evolution of associated floral features. The following are a summary of these general correlations or syndromes.

Insect pollination (or entomophily) is undoubtedly the most common type in angiosperms. **Bee pollination (melittophily or hymenopterophily)** is correlated with flowers that tend to be showy, colorful, and fragrant. The flowers often have specialized color patterns called nectar guides (Figure 13.1A), which function to attract and orient the bee to maximally effect pollination. In many bee-pollinated flowers, nectar guides may be correlated with the anterior perianth part(s) (usually petals or corolla lobes) modified as landing platforms (Figure 13.1A), on which the bee lands to more efficiently gather nectar or pollen and more effectively cause pollination.

Butterfly pollination (psychophily) is associated with showy, colorful, and fragrant flowers, usually with no nectar guides. The flowers tend to have long, nectar-filled tubes or spurs (Figure 13.1B), preventing all but an insect with a long proboscis from acquiring the nectar.

Moth pollination (phalaenophily) is correlated with large, white, and fragrant flowers, with no nectar guides; as with butterfly pollination, the flowers often have long, nectar-filled tubes (Figure 13.1C) or spurs. One interesting example of moth pollination occurs in species of the monocots *Hesperoyucca* and *Yucca* (Agavaceae), which are exclusively pollinated by yucca moths (*Parategeticula* and *Tegeticula* spp.). Yucca moths, in addition to pollinating *Yucca* flowers, deposit their eggs only within the ovary of *Yucca* plant species. Thus, the *Yucca* plant and yucca moths are obligately dependent upon each other for procreation.

Fly pollination (sapromyiophily) is correlated with flowers that are often maroon or brown in color and emit a fetid odor that simulates the smell of rotting flesh. Examples of these are *Arum* and *Stapelia* spp. (Figure 13.1D). In some of these flowers, flies may lay their eggs, which will fail to develop because of the absence of a suitable food source.



FIGURE 13.1 Flower modifications associated with specific pollination mechanisms. **A.** *Penstemon eximus*, bee-pollinated. Note nectar guide and landing platform. **B.** *Linaria canadensis*, toad-ax, butterfly pollinated, with corolla-spur. **C.** *Calonyction aculeatum*, moonflower, moth pollinated, white with long corolla tube. **D.** *Stapelia gigantea*, starflower, fly-pollinated. **E.** *Selenicereus* sp., night-blooming cereus, bat-pollinated. **F.** *Couroupida guianensis*, cannonball tree, bat-pollinated. **G.** *Erythrina caffra*, coral bean tree, bird-pollinated. **H.** *Anigozanthos manglesii*, kangaroo-paw, bird-pollinated. **I.** *Bromus* sp., brome grass, wind-pollinated.

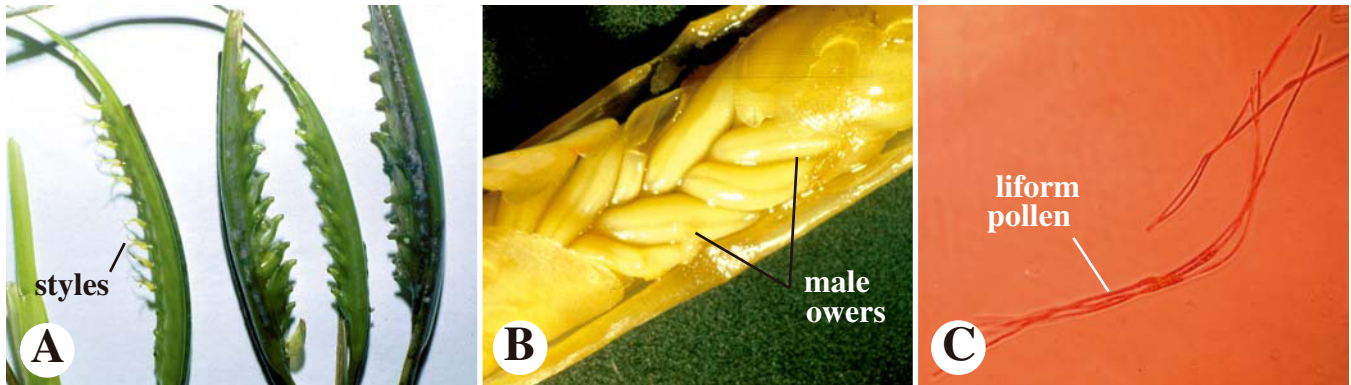


FIGURE 13.2 *Phyllospadix torreyi*, surf-grass, a water-pollinated angiosperm. **A.** Female plant. Note styles at left. **B.** Close-up of male flowers. **C.** Elongate, filiform pollen grains.

Bat pollination (cheiropterophily) is correlated with flowers that open at night (have a nocturnal anthesis), and are large, white or colorful, with copious production of pollen or nectar (often secreted into a hypanthium or perianth tube), either or both of which may serve as a reward. When pollen is the reward, stamens tend to be numerous (Figure 13.1E,F).

Bird pollination (ornithophily) tends to occur in red, relatively large, and often tubular flowers that secrete copious nectar (Figure 13.1G,H). Sometimes the tube results from tightly wrapped but distinct perianth parts (e.g., the cactus *Cleistocactus*, pollinated by hummingbirds).

Wind pollination (or anemophily) is correlated with small, numerous, often unisexual flowers that tend to have a reduced, nonshowy, or absent perianth (Figure 13.1I). Pollen is produced in large quantities and pollen grains tend to have a smooth (*psilate*) wall sculpturing. Styles tend to be highly branched as a more efficient means of catching pollen grains in air currents. Anthers and styles may be erect or pendant. Wind pollination is found in several flowering plant groups, such as the Fagaceae (e.g., oaks), Betulaceae (e.g., birches), Salicaceae (poplars and willows), and many Poales (grasses and their close relatives). Some wind-pollinated taxa are quite specialized, such as *Alexgeorgea* (Restionaceae, a grass relative), in which the flowers are underground but in which the emergent styles and stamens undergo wind pollination.

Water pollination (hydrophily) may occur in aquatic plants with flowers either at or under the water surface. For example, *Vallisneria* (Hydrocharitaceae) releases tiny male flowers that float to the surface, where they may float to the enlarged stigmas of a relatively large female flower. Some sea grasses, such as *Phyllospadix* (Figure 13.2) have very elongate, filiform pollen grains (Figure 13.2C), making them considerably more efficient in being captured

by the styles and stigmas of female flowers in ocean currents (Figure 13.2A).

Other, less common pollination syndromes include beetle pollination (*cantharophily*), carrion beetle pollination (*necrocoleopterophily*), and ant pollination (*myrmecophily*).

BREEDING SYSTEMS

Plants can be predominately outbreeding, inbreeding, or some mixture of the two. In many flowering plants specific mechanisms have evolved that promote one of these systems.

Outbreeding, also called **outcrossing**, **allogamy**, or **xenogamy**, is the transfer of gametes from one individual to another, genetically different individual. The general advantage of outbreeding is to promote an increase in phenotypic variability within a population. This generally enables plants to adapt to a wider range of environmental conditions and increases the likelihood for survival and evolutionary change. One disadvantage of outbreeding is that it requires a transfer of gametes between individuals. If individuals are far apart, or if pollinators are scarce, sexual reproduction may not occur at all in obligately outbreeding species.

The probability of outbreeding can be increased by a variety of mechanisms. **Dioecy**, in which individual plants have either male (staminate) or female (pistillate) flowers, ensures that outbreeding will always occur. Many flowering plants exhibit a modified type of dioecy in which some individuals have flowers of one sex but others have bisexual (perfect) flowers. These include **gynodioecy** (some individuals with pistillate flowers only, others with perfect flowers), **androdioecy** (some individuals with staminate flowers only, others with perfect flowers), and **trioecy** (some individuals with staminate flowers only, some with pistillate flowers only, and some with perfect flowers). These alternative mechanisms may

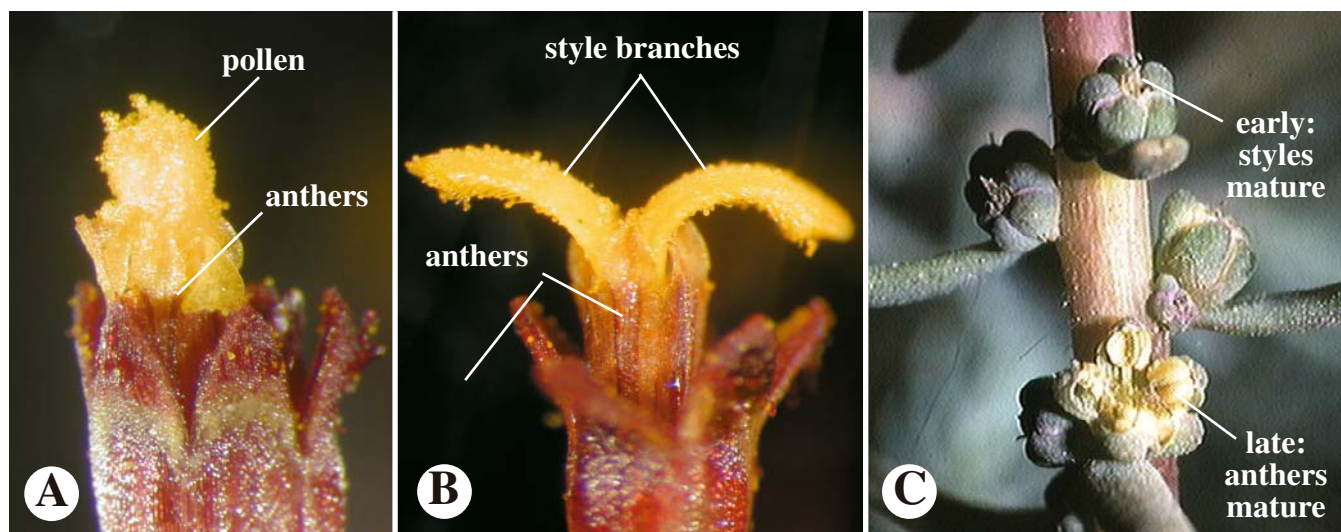


FIGURE 13.3 A,B. Protandry in *Encelia californica* (Asteraceae). A. Style is elongating through anther tube, pushing pollen outward. B. Style has finished elongating, and style branches have unfolded. C. Protogyny in *Suaeda* (Chenopodiaceae). Styles have already matured and are drying up at time of pollen release.

promote outcrossing but also allow for some inbreeding (see later discussion), ensuring that at least some seed will be set.

Another outcrossing mechanism is the result of differences in timing of maturation of male and female floral parts, a feature known as **dichogamy**, of which there are two general types. **Protandry** (Figure 13.3A,B) is the precocious development of the androecium, as occurs, e.g., in many members of the Apiaceae, Asteraceae, and Campanulaceae. In protandrous species the pollen matures and is released prior to the maturation and receptivity of the gynoecium. **Protogyny** (Figure 13.3C) is the precocious development of the gynoecium, as occurs, e.g., in some Chenopodiaceae. Both protandry and protogyny promote outcrossing when flowers of different individuals mature at slightly different rates. Thus, the pollen from one flower will not normally pollinate that same flower, but can pollinate a different flower in which the gynoecium is receptive. In protandrous and protogynous species, outcrossing is ensured only if the flowers from a given individual mature at the same time. In reality, most of these species have flowers aggregated together into inflorescences, in which a range of developmental stages may be present. In any case, at least some outcrossing may occur, and pollination within a single flower is normally prevented. However, if the pollen is not removed from a flower, it may in some cases pollinate the gynoecium of that same flower. This provides a fail-safe mechanism for producing seeds even in times or environments where pollinators are lacking.

Outcrossing has also been enhanced by evolutionary changes in floral structure, particularly the spatial separation

of anthers and stigmas, a phenomenon known as **hercogamy**, also spelled herkogamy. (However, hercogamy may also function to prevent interference of male and female functions in the flower, by physically separating them.) One type of hercogamy is **heterostyly**, in which the relative lengths or heights of stigmas versus anthers vary among different flowers. (Most flowers are monomorphic or homostylous, whereby the height of stigmas and anthers is relatively constant.) In so-called **distylous** species, two floral morphologies occur: **pin** flowers, with a long style and short stamens, and **thrum** flowers, with a short style and long stamens (Figure 13.4A). In this syndrome, an insect visiting a pin flower is likely to have pollen deposited on its body in a location that would effect pollination of a thrum flower rather than another pin flower, and vice versa. This increases the probability of pollination between flowers rather than within flowers. If individuals tend to have one floral type or the other, outcrossing would be ensured. A rarer, more complex situation occurs in species that are **tristylous**, with three heights of styles and stamens; the principle for cross-pollination is the same (Figure 13.4B).

Another type of hercogamy is **enantiostyly** (or *enantiomorphy*), the curvature of the style to either the left or the right (as viewed from the front), defining left-handed and right-handed flowers (Figure 13.5). This style curvature usually corresponds with a curvature of at least one stamen to the side opposite the style. As with heterostyly, enantiostyly results in the preferential deposition of pollen on one side of, say, an insect pollinator's body. For example, an insect visiting

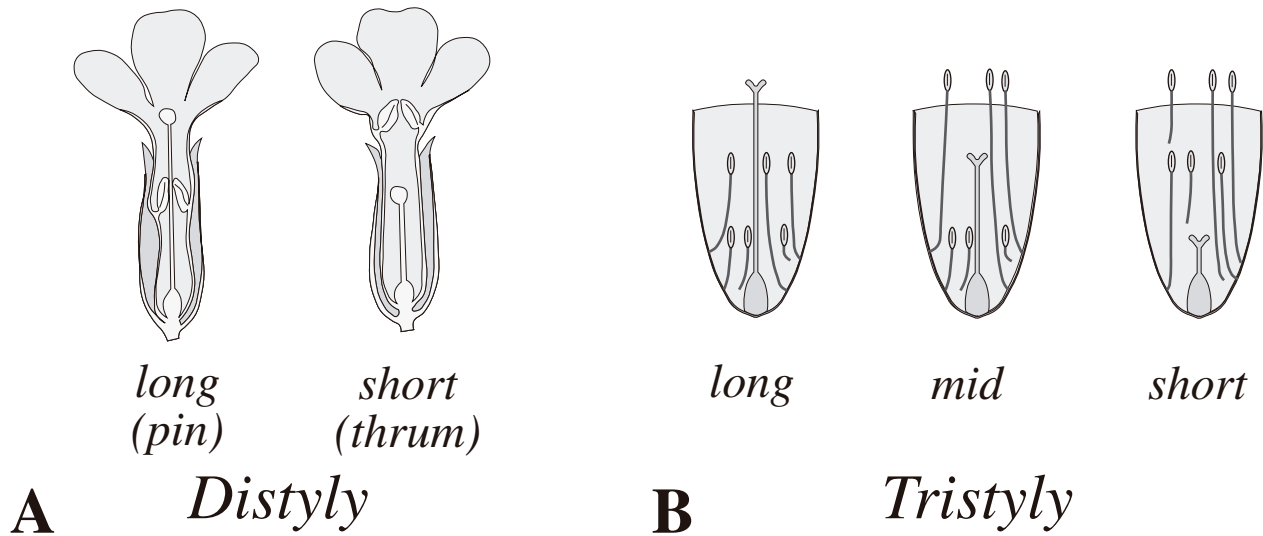


FIGURE 13.4 Flower heteromorphism. **A.** Distyly. (Redrawn from Weberling, 1989. *Morphology of flowers and inflorescences*. Cambridge University press, Cambridge, New York.) **B.** Tristyly (Redrawn from Kohn et al., 1996. *Evolution* 50: 1454–1469.)

a left-handed flower would tend to get pollen deposited on the right side of its body. Thus, the insect would more likely pollinate a right-handed flower as opposed to another left-handed flower. If an individual plant is relatively constant as to floral handedness, enantiostyly will greatly promote the probability of outcrossing.

Yet another type of herkogamy involves movement of floral parts. One type of **movement herkogamy** is the rapid closure of the stigmas upon their being touched by a potential animal pollinator (e.g., *Mimulus*, Phymaceae). If the stigmas

first receive pollen from a pollinator, their rapid closure can physically prevent pollen from the same flower being transferred to the stigmatic region, effectively preventing intrafloral self-pollination. Another type of movement herkogamy involves **trigger mechanisms** (e.g., *Kalmia*, *Stylidium*), in which an insect pollinator triggers the sudden movement of one or more stamens, dusting the insect with pollen at the point of contact (Figure 13.6). This pollen is then at a position to be more effectively transmitted to the stigma of another flower.

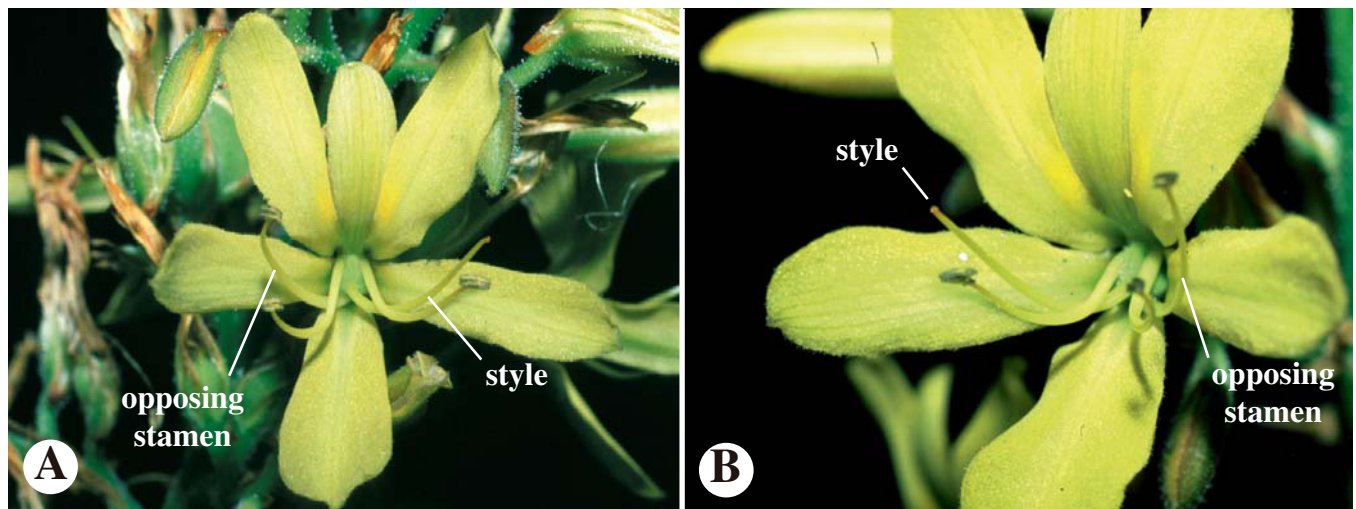


FIGURE 13.5 Example of enantiostyly in *Wachendorfia thyrsiflora* (Haemodoraceae). Note curvature of style with corresponding opposite positioning of stamen. **A.** Right-handed flower. **B.** Left-handed flower.



FIGURE 13.6 Example of a trigger mechanism in *Stylidium* (Stylidiaceae). **A.** Prior to being triggered. **B.** Pencil has triggered single stamen to release pollen grains.

Finally, outcrossing can be promoted by genetically determined self-incompatibility mechanisms. **Self-incompatibility** refers to the inability for fertilization to occur between gametes derived from an individual genotype. Because this is genetically determined, the incompatibility operates both within a single flower and between flowers of one individual. There are two basic types of self-incompatibility. Gametophytic self-incompatibility is controlled by the genetic composition of the male gametophyte. Sporophytic self-incompatibility is controlled by the genetic composition of the sporophyte, specifically the stigma and style of the pistil.

Inbreeding, also called **selfing**, is the union of gametes derived from a single individual. In flowering plants, inbreeding may occur either within a single flower, known as **autogamy (intrafloral selfing)** or between flowers derived from one individual, known as **geitonogamy**. (The genetic product of autogamy is identical with that of geitonogamy.) A major evolutionary advantage of inbreeding is enabling reproduction to occur when there are relatively few (or even one) individuals present in a population or at times when pollinators are rare, e.g., in ephemeral habitats. The disadvantage of inbreeding is that it reduces variation in a population and can even result in the accumulation of deleterious alleles, a phenomenon known as inbreeding depression.

Some plant species have both outcrossing and selfing flowers, a breeding system known as **allautogamy**. For example, species of *Viola* (Violets) and *Clarkia* have two types of flowers. **Chasmogamous** flowers are typical ones in which the perianth opens and exposes the sexual organs, with subsequent cross-pollination common. Other flowers, however, are **cleistogamous**, in which the perianth remains

closed, such that pollen produced from within the flower pollinates only the stigma(s) of that flower. In still other species, both self- and cross-pollination may occur within the same flower. For example, *Myosurus*, mousetail, has numerous, small pistils born on a receptacle that elongates during flower maturation. When the flower first blooms, the receptacle is short, and the pistils that mature tend to be pollinated by the low, surrounding anthers. As the receptacle elongates, however, the pistils are positioned high above the anthers and are more likely to be pollinated by an insect visitor carrying pollen from another flower. All autogamous breeding systems are adaptive in promoting some outcrossing, which may increase overall genetic variation, but also ensuring seed set regardless of availability of pollinators via inbreeding.

Breeding systems may be correlated with overall timing of sexual reproduction. **Iteroparous** plants are those that reproduce more than one time in the life of the plant, typically in regular cycles. Iteroparity is very common in angiosperms, ensuring regular seed set. **Semelparous** plants are those in which plant resources are utilized entirely for one episode of reproduction, followed by degeneration and death of the entire plant (i.e., the plants are **monocarpic**; Chapter 9). Semelparity occurs in all annual and biennial plants, but in very few perennials (occurring, e.g., in *Agave* and *Bambusa* spp.). Semelparity in perennial plants may be a strategy for deceiving or overwhelming potential seed predators, the former by not reproducing seasonally, the latter by producing seeds in such numbers that predators cannot consume them all.

Other temporal phenomena may be correlated with the breeding system of a plant species. For example, in annual or deciduous plants, the relative timing of leaf versus flower

development may influence pollination and/or seed dispersal. Two general temporal patterns are **synanthous**, in which leaves and flowers develop at the same time, and **hysteranthous**, in which leaf and flower development do not coincide. Wind-pollinated plants are sometimes hysteranthous (e.g., in the willows, *Salix* spp.), with flowers maturing and releasing pollen before leaves form, thus, more effectively transmitting pollen in the canopy region of a community of trees.

SEED AND FRUIT DISPERSAL

The evolution of the numerous types of fruits and seeds (Chapter 9), which are used to delimit many taxa, is strongly correlated with their function as dispersal devices. Many mechanisms for dispersal of seeds and fruits have evolved in the angiosperms, including (1) **wind dispersal** (or **anemochory**), as in the samaras of *Ulmus*, elms, and *Acer*, maples, and the winged seeds of *Liquidambar*, sweetgum, including those wind dispersed by tumbling (e.g., the tumble weeds, such as *Salsola*); (2) **water dispersal** (or **hydrochory**), as occurs in the ocean-dispersed fruits of *Cocos nucifera*, coconut; (3) dispersal by **explosive dehiscence** of fruits, as in the explosively dehiscent capsules of *Ceanothus* or *Impatiens*; (4) **self-dispersal** (or **autochory**), as in *Arachis hypogaea*, peanut, which buries its own fruits; or (5) **animal dispersal** (or **zoochory**). Zoochory is divided into **ecto-zoochory**, in which propagules are carried on the outside of an animal (as in the burs of *Xanthium*, cocklebur, or the loment of *Desmodium*, sticktight), and **endo-zoochory**, in which seeds are eaten (the fruit pericarp or fleshy seed coat or aril being an award or attractant) but are passed through the gut of the animal unharmed.

ASEXUAL REPRODUCTION

Many species of land plants will regularly produce offspring without sex. The advantage of asexual reproduction is that numerous propagules can be generated relatively quickly and efficiently, without reliance on the transfer of gametes. However, the major evolutionary disadvantage is that no genetic variability results. One type of asexual reproduction is vegetative reproduction, the production of genetic clones from vegetative tissue. (Clones of an individual are known as **ramets**, whereas genetically different individuals are called **genets**.) Vegetative plant clones are produced by the formation of aerial plantlets (e.g., develop along the leaf margins of *Kalanchoe daigremontiana*, maternity plant). Cloning may also result from stolons, rhizomes, bulbels, cormels, etc., that may become dispersed or physically detached from the genetically identical parent plant. Some clones of creosote

bush (*Larrea* sp.) are calculated to have persisted in nature for several thousands of years and may represent the oldest known clonal organisms on earth.

Agamospermy is the production of seeds without fertilization. In some species, agamospermy requires pollination to form seeds, though fertilization never occurs. Embryos of agamospermic seeds are genetically identical to the parent plant. The embryo may develop from a cell of an abnormal, diploid female gametophyte, such as a diploid egg, this being **parthenogenesis**. Alternatively, the embryo may arise from a cell of the surrounding tissue, such as megasporangial or integument tissue, which is called **adventive embryony**. Agamospermy may be facultative, occurring in addition to normal sexual reproduction if flowers are unpollinated. Alternatively, some species are obligate agamosperms, an example being *Taraxacum officinale*, the common dandelion. Evidence for agamospermy includes (1) the occurrence of viable seed in absence of males or after experimental bagging or emasculation of flowers (see later discussion); (2) precocious embryo formation, i.e., prior to anthesis or pollen maturation; (3) adventive embryos, budding from nucellar or integument tissues; (4) multiple embryo and seedling formation from a single seed; (5) rarity or absence of males in nature, e.g., in dioecious species.

HYBRIDIZATION, INTROGRESSION, AND POLYPLOIDY

Hybridization is usually defined as sexual reproduction between different species, specifically termed *interspecific hybridization* (although the term can be used for sexual reproduction between different populations or infraspecific taxa within a species). Hybridization is thought to be relatively common in plants, more so than in most groups of animals.

Two different species of plants will not interbreed if they are geographically isolated or if they exhibit one or more of a variety of genetically determined traits that prevent or inhibit gene exchange. These genetic, reproductive isolating features include (1) differences in habitat; (2) differences in timing of reproduction; (3) differences in floral morphology; or (4) differences in pollinators. (For example, one species that is adapted to a wet environment may not be capable of interbreeding with one that is adapted to a dry environment, simply because the two species are rarely in close enough proximity to allow pollination.)

Hybridization between two plant taxa may only occur if these taxa are genetically similar enough. Any hybrid progeny that are produced may be fertile (capable of sexual reproduction) or sterile, the latter often the result of

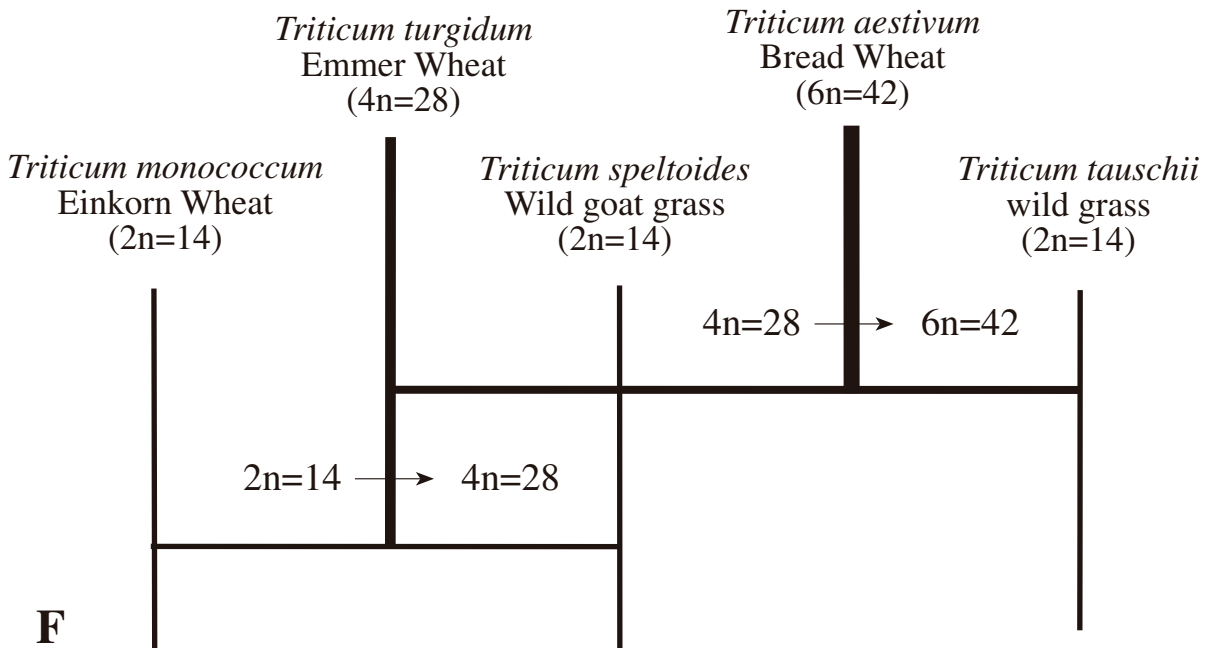
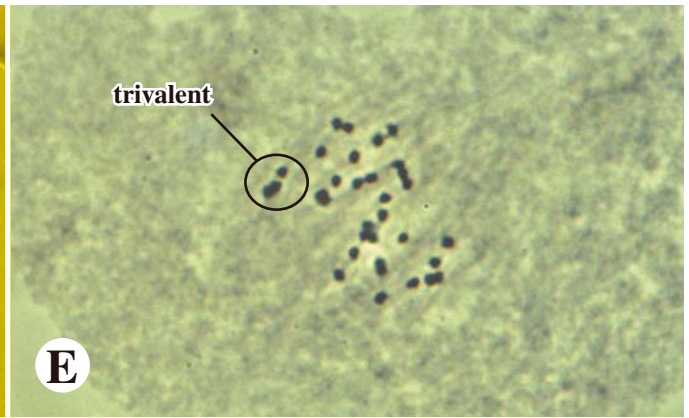
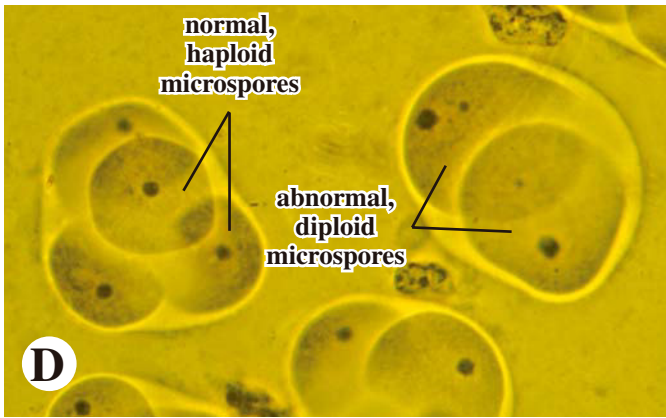
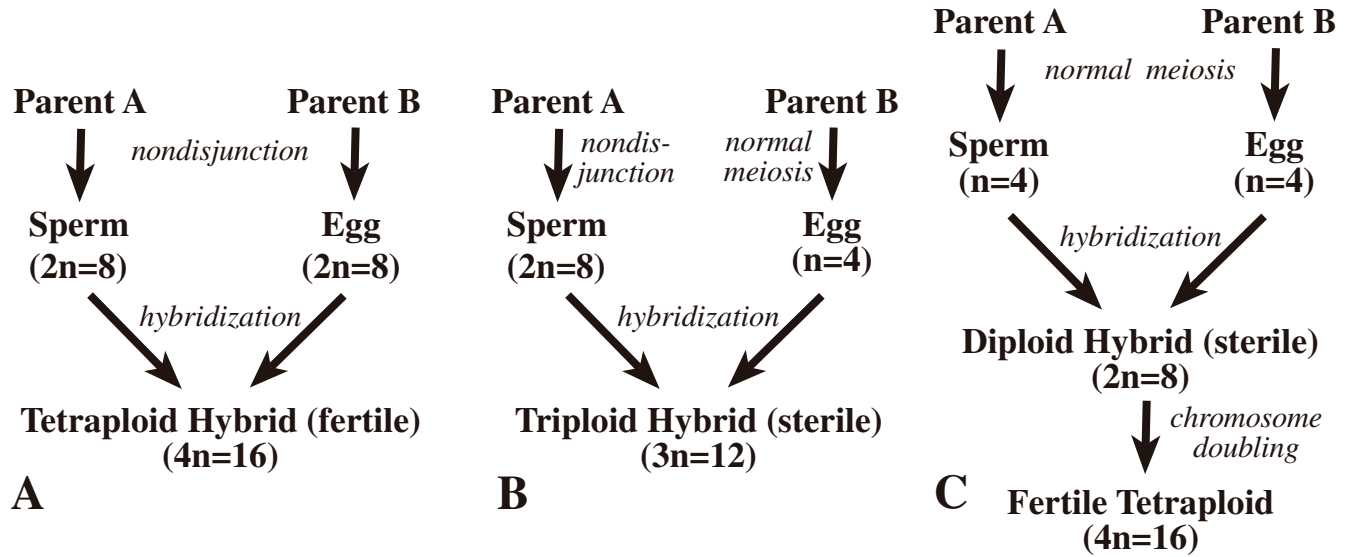


FIGURE 13.7 Polyploidy. **A,B.** Mechanisms by which tetraploidy and triploidy can arise in nature by meiotic nondisjunction, resulting in diploid gametes. **C.** Mechanism by which a tetraploid individual can arise by somatic chromosome doubling in a sterile hybrid. **D.** Pollen development in *Cylindropuntia* sp., showing normal tetrad of haploid microspores (left) and abnormal dyad of diploid microspores (right), the latter precursors to diploid pollen grains. **E.** Chromosome squashes (*Cylindropuntia prolifera*, a sterile triploid), showing groupings of three homologous chromosomes (trivalents) during meiosis, indicative of triploidy. **F.** Evolution of wheat, *Triticum* spp., via polyploid events. (Photos at D and E courtesy of Jon Rebman.)

irregularities in meiosis, resulting in sterile or noncompatible gametes.

Introgression is hybridization between two species followed by backcrossing to one or both parents. The importance of introgression is that it can be a mechanism of promoting some gene flow between two different species, ultimately increasing the genetic variability or fitness of one or the other species.

Polyploidy is a mutation in which offspring have a multiple of some ancestral set of chromosomes. Polyploidy can occur either within a species (**autopolyploidy**) or between different species (**allopolyploidy**).

Polyploidy can occur in two general ways. One way that polyploidy can occur is by the production of gametes that have more than one set of chromosomes (Figure 13.7A,B). Diploid gametes can result from an irregularity during meiosis termed **nondisjunction**, in which homologous chromosomes do not segregate; if this occurs with all homologous chromosome pairs, then the daughter cells may be unreduced (i.e., diploid, not haploid). [An unreduced (diploid) pollen grain can sometimes be detected microscopically, whereby only two, larger microspores (see Chapter 12) are detected in the tetrad phase (Figure 13.7D).] If both parents (either of the same or different species) produce diploid gametes, then the offspring possesses four sets of chromosomes, which is a **tetraploid** (Figure 13.7A). Tetraploids are normally fertile, as they can produce viable, diploid gametes. If, however, one parent contributes a haploid gamete and the other a diploid gamete, the offspring will be **triploid** (Figure 13.7B). Triploids are generally sterile, as any gametes they produce will generally lack a full complement of chromosomes because of meiotic irregularities, forming groupings of three homologs (trivalents; e.g., Figure 13.7E), instead of the normal two (bivalents). Triploids might persist as a population, however, if they can continue to reproduce asexually.

A second way that polyploidy can occur is by the spontaneous doubling of chromosome number in an individual plant *after* normal sexual reproduction. For example, hybridization between two different species might produce living diploid offspring, but the offspring often cannot produce viable gametes because of the genetic dissimilarities between the two

parents (Figure 13.7C). However, if this sterile offspring can persist, e.g., if it can also reproduce vegetatively, it might (rarely) undergo a rare **somatic** (i.e. in a nonreproductive cell) chromosome doubling during mitosis in a critical region, e.g., the apical meristem of a shoot, such that this entire shoot becomes tetraploid. The tetraploid is now potentially capable of producing viable, diploid gametes and, therefore, fertile offspring (Figure 13.7C). This type of polyploid event may be rare, but has been documented in species of *Spartina* (cordgrass), in which a new tetraploid species evolved from two, separate diploid parents.

Polyploidy is thought to be a major mechanism in plant evolution, as chromosome studies have demonstrated that most plants have undergone a polyploid event during some time of their history. The evolution of both emmer wheat and bread wheat occurred via ancestral, allopolyploidy events, resulting in tetraploid (4n) and hexaploid (6n) individuals (Figure 13.7F).

TESTING FOR BREEDING MECHANISM

Experimental methods may be used to assess the type of breeding mechanism. Flower buds may be bagged or caged, i.e., covered with a fine netting that excludes potential pollinators. Also, flower buds may be emasculated, in which the anthers are removed prior to pollen release. An example of an experimental regime to test the breeding system is seen in Table 13.1. The determination of seed set for each experiment allows inference as to the breeding system.

In addition to these manipulative experiments, embryological observations may be made to determine, e.g., if pollen tubes are growing through the style of the flower. (Fluorescence microscopy may be used to detect pollen tubes; Figure 13.8.) Absence or inhibition of pollen tube growth indicates some type of genetic incompatibility. In addition, observations of female gametophyte or embryo development and/or chromosome counts of these tissues may detect the occurrence of agamospermy.

TABLE 13.1 Five experimental manipulations 1–5, with explanation of results. Normal seed set = + and greatly reduced to zero seed set = –.

SEED PRODUCTION		
1. Flowers left to develop normally, a control.	+ Fertile	– Infertile
2. Flowers caged, then self-pollinated by hand.	+ Self-fertile	– Not Self-fertile
3. Flowers caged, then left alone.	+ Self-pollinating	– Not Self-pollinating
4. Flowers emasculated and caged.	+ Autogamous	– Not Autogamous
5. Flowers caged, emasculated, and outcrossed.	+ Outcrossing	– Not Outcrossing

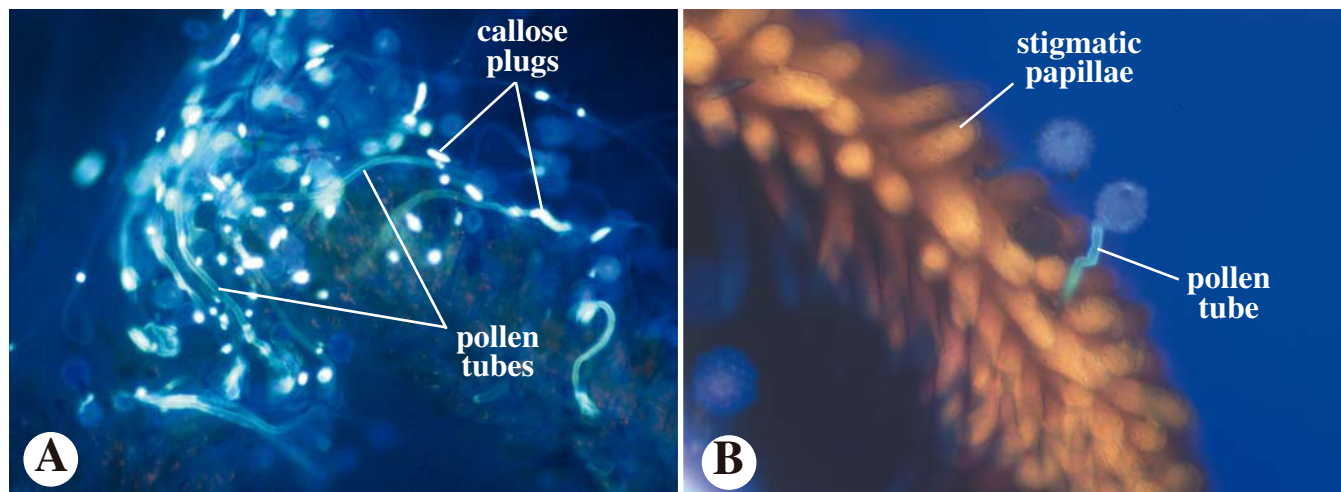


FIGURE 13.8 A,B. Pollen tube growth. Fluorescent stain, positive for callose, allows visualization of pollen tubes down length of style.

REVIEW QUESTIONS

1. What is plant reproductive biology?
2. What is pollination?
3. What two general features have evolved in flowers that function to effect animal pollination?
4. What products serve as a reward for animal pollinators?
5. What is a nectar guide? A landing platform?
6. For the following pollination mechanisms, name some floral syndromes (correlated structural modifications): (a) bee; (b) butterfly; (c) moth; (d) wind; (e) bird; (f) bat.
7. What are the two major or extreme types of breeding systems?
8. What are the advantages and disadvantages of outbreeding?
9. What is dichogamy? Name two specific types of dichogamy that can promote outcrossing.
10. What is herkogamy?
11. Define and explain: heterostyly, distyly (pin and thrum), tristyly, enantiostyly, movement herkogamy, trigger mechanisms. What is the overall function of these floral mechanisms?
12. What is self-incompatibility and what is its significance in plant reproductive biology?
13. What are the advantages and disadvantages of inbreeding?
14. What is the difference between allogamy, autogamy, geitonogamy, and allautogamy?
15. Name some types of inbreeding mechanisms.
16. What is agamospermy? How can it be detected?
17. What is hybridization? introgression?
18. Define polyploidy.
19. Cite the ways that polyploidy can occur.
20. How can one test the breeding mechanism in plants? (cite specific ways to test)
21. If the following experiments are performed for plant species A D, what can you say about the breeding mechanism based on the pattern of seed set?

Seed Production

	A	B	C	D
1. Flowers left to develop normally, a control.	+	+	+	+
2. Flowers caged, then self-pollinated by hand.	-	+	+	+
3. Flowers caged, then left alone.	-	-	+	+
4. Flowers emasculated and caged.	-	-	-	+
5. Flowers caged, emasculated, and outcrossed.	+	+	+	+

EXERCISES

1. Examine specimens of two species of plants plus any putative hybrids between them. (a) Study both vegetative and floral characters, from original observations or using a manual of the area, and note which diagnostic features distinguish the two species. (b) Decide upon which characters to measure in the specimens available. (c) Record 10–25 measurements of each of the parameters chosen. Compare these by preparing graphs in order to recognize discontinuities (or lack thereof) of the three taxa.
2. Locate a population of a Composite (Asteraceae) species that has both disk and ray flowers. Observe insect visitors (potential pollinators) in each of two subsets of plants (or inflorescences): one undisturbed and another with all ray flowers removed. Count the number and type of visitors over a time period (e.g., 10–30 minutes) and record.
3. If material is available, observe ultraviolet light sensitive regions in the perianth by placing a flower into a jar saturated with ammonium vapors. Bees can detect these UV-reflective regions of the flower, enabling them to find flowers and orient to pollen or nectar more efficiently.
4. Fix the styles of a species of flowering plant in 70% alcohol. Remove the style and place in drops of aniline blue on a microscope slide, covered by a cover slip. If this style is small enough, it may be squashed by applying firm pressure on the cover slip (using, e.g., a cork). Observe under fluorescence microscopy. Pollen tubes regularly deposit callose, which differentially picks up the aniline blue stain. This method allows for detection of pollen tube growth and can be used to test whether self-incompatibility is occurring.
5. If time permits, select a plant species and perform the crossing and caging experiments described in the text. These techniques are used to test the potential and degree of self-pollination versus cross-pollination.
6. Peruse journal articles on plant systematics, e.g., *American Journal of Botany*, *Systematic Botany*, or *International Journal of Plant Sciences*, and note those that describe aspects of reproductive biology in relation to systematic studies. Identify the techniques used and the problems addressed.

REFERENCES FOR FURTHER STUDY

- Bernhardt, P. 1989. *Wily Violets and Underground Orchids*. W. Morro & Co., New York.
- Chittka, L., A. Shmida, N. Troje, and R. Menzel. 1994. Ultraviolet as a component of flower reflections, and the colour perception of Hymenoptera. *Vision Research* 34: 1489–1508.
- Faegri, K., and L. van der Pijl. 1979. *The Principles of Pollination Ecology*, 3rd ed. Pergamon Press, Oxford.
- Stace, C. A. 1989. *Plant Taxonomy and Biosystematics*, 2nd ed. Edward Arnold. Distributed in the USA by Routledge, Chapman, and Hall, New York.
- van der Pijl, L. 1982. *Principles of Dispersal in Higher Plants*. Springer-Verlag, Berlin.

14

PLANT MOLECULAR SYSTEMATICS

ACQUISITION OF MOLECULAR DATA	477	RANDOM AMPLIFIED POLYMORPHIC DNA (RAPDs)	487
DNA SEQUENCE DATA	477	AMPLIFIED FRAGMENT LENGTH POLYMORPHISM (AFLPs)	487
Polymerase Chain Reaction	477	REVIEW QUESTIONS	490
DNA Sequencing Reaction	479	EXERCISES	491
Types of DNA Sequence Data	479	REFERENCES FOR FURTHER STUDY	491
Analysis of DNA Sequence Data	482		
RESTRICTION SITE ANALYSIS (RFLPs)	483		
ALLOZYMES	484		
MICROSATELLITE DNA	487		

Molecular systematics encompasses a series of approaches in which phylogenetic relationships are inferred using information from macromolecules of the organisms under study. Specifically, the types of molecular data acquired include that from DNA sequences, DNA restriction sites, allozymes, microsatellites, RAPDs, and AFLPs. (The use of data from other, generally smaller molecules, such as secondary compounds in plants, is usually relegated to the field of chemosystematics and will not be reviewed here.)

A revolution in inferring the phylogenetic relationships of life is occurring with the use of molecular data. The following is a review of the types of data, methods of acquisition, and methods of analysis of molecular systematics.

ACQUISITION OF MOLECULAR DATA

Plant samples from which DNA is to be isolated may be acquired by various means. It is vital to always collect a proper voucher specimen, properly mounted and accessioned in an accredited herbarium, to serve as documentation for any molecular systematic study (see Chapter 17). Live samples may be collected and immediately subjected to chemical processing, e.g., for allozyme analysis (see later discussion). For many DNA methods, pieces of leaves (from which chloroplast, mitochondrial, and nuclear DNA can be isolated) are removed from the live plant and immediately dried, typically

in a container of silica gel. Alternatively, plant samples may be frozen or placed in concentrated extraction buffer. With any of these procedures, DNA is usually preserved intact. Useable DNA is often successfully isolated from dried herbarium sheets, attesting to the toughness of the molecule.

DNA SEQUENCE DATA

Perhaps the most important method for inferring phylogenetic relationships of life is that of acquiring DNA sequences. DNA sequence data basically refers to the sequence of nucleotides (adenine = A, cytosine = C, guanine = G, or thymine = T; Figure 14.1) in a particular region of the DNA of a given taxon. Comparisons of homologous regions of DNA among the taxa under study yield the characters and character states that are used to infer relationships in phylogenetic analyses.

The first step of acquiring DNA sequence data is to identify a particular region of DNA to be compared between species. Much prior research goes into identifying these regions and determining their efficacy in phylogenetic analysis.

POLYMERASE CHAIN REACTION

After a gene sequence of interest is identified, the DNA from a given plant sample is first isolated and purified by various

These short, conserved, flanking regions are used as a template for the synthesis of multiple, *complementary* copies, known as **primers**. Primers ideally are constructed such that they do not bind with one another.

In the polymerase chain reaction, a solution is prepared, made up of (1) the isolated and purified DNA of a sample; (2) multiple copies of primers; (3) free nucleotides; (4) DNA polymerase molecules (typically *Taq* polymerase, which can tolerate heat); and (5) buffer and salts. This solution is heated to a point at which the sample DNA denatures, whereby the two strands of DNA separate from one another. Once the sample DNA denatures, the primers in solution may bind with the corresponding, complementary DNA of the sample (Figure 14.2). Following binding of the primer to the sample DNA, individual nucleotides in solution attach to the 3' end of the primer, with the sample DNA acting as a template; DNA polymerase catalyzes this reaction. A second primer, at the opposite end of the DNA sequence of importance, is used for the complementary, denatured DNA strand. Thus, the two denatured strands of DNA are replicated. After replication, the solution is cooled to allow for annealing of the replicated DNA with the complementary DNA single strands. This is followed by heating to the point of DNA denaturation, and repeating the process. A typical PCR reaction can produce more than a million copies of DNA in a matter of hours.

DNA SEQUENCING REACTION

After DNA is replicated, it is sequenced. The most common sequencing technology involves a machine that reads fluorescent dyes with a laser detector. The production of dye-labeled DNA is very similar to DNA replication using the PCR. The replicated DNA is placed into solution with DNA polymerase, primers, free nucleotides, and a small concentration of synthesized compounds called dideoxynucleotides (discussed later) that are each attached to a different type of fluorescent dye. As in the polymerase chain reaction, the sample DNA is heated until the double helix unwinds and the two complementary DNA chains separate (Figure 14.3). At this point, a primer attaches to a conserved region of one of the strands of DNA, and free nucleotides in solution join to the 3' end of the primer, using the sample DNA as a template and catalyzed by DNA polymerase (Figure 14.3). Thus, a replicated copy of the DNA strand begins to form. However, at some point a dideoxynucleotide joins to the new strand instead of a nucleotide doing so. The dideoxynucleotides (dideoxyadenine, dideoxycytosine, dideoxyguanine, and dideoxythymine) resemble the four nucleotides, except that they lack a hydroxyl group. Once a dideoxynucleotide is joined to the chain, absence of the hydroxyl group prevents the DNA polymerase from joining it to anything else. Thus, with the addition of

a dideoxynucleotide, synthesis of the new DNA strand terminates (Figure 14.3).

The ratio of dideoxynucleotides to nucleotides in the reaction mixture is carefully set and is such that the concentration of dideoxynucleotides is always much smaller than that of normal nucleotides. Thus, the dideoxynucleotides may terminate the new DNA strand at any point along the gene being replicated. For example, some of the new DNA strands will be the length of the primer plus one additional base (in this case the dideoxynucleotide); some will be the primer length plus two bases (a nucleotide plus the terminal dideoxynucleotide); some will be the primer length plus three bases (two nucleotides plus the terminal dideoxynucleotide); etc. There are many thousands, if not millions, of copies of the sample DNA. Thus, there will be an equivalent number of newly replicated DNA strands, of all different lengths.

The final step of DNA sequencing entails subjecting the DNA strands to electrophoresis, in which the DNA is loaded onto a flat gel plate or in a thin capillary subjected to an electric current. Because the phosphate components of nucleic acids give DNA a net negative charge, the molecules are attracted to the positive pole. The DNA strands migrate through the medium over time, the amount of migration inversely proportional to the molecular weight of the strand (i.e., lighter strands migrate further). Each strand is terminated with a dideoxynucleotide to which a fluorescent dye is attached; each of the four dideoxynucleotides has a different type of fluorescent dye, which (upon excitation) emits light of a different wavelength. Thus, as the multiple copies of DNA of one particular length migrate along the gel or capillary, the wavelength of emitted light is detected and recorded as a peak, which measures the light intensity. Because a given emitted wavelength (color) is determined by one of the four dideoxynucleotides, the corresponding nucleotide can be inferred and its position identified by the timing of migration of the DNA strands. In this way, the sequence of nucleotides of the DNA strand can be inferred (Figure 14.3).

TYPES OF DNA SEQUENCE DATA

For plants, the three basic types of DNA sequence data stem from the three major sources of DNA: nuclear (nDNA), chloroplast (cpDNA), and mitochondrial (mtDNA). Nuclear DNA is, of course, transmitted from parent(s) to offspring by nuclear division (meiosis or mitosis) via sexual or asexual (somatic) reproduction. Chloroplasts and mitochondria, however, replicate and divide independently of the nucleus and may be transmitted to offspring in a different fashion. For example, in angiosperms these organelles are usually (with some exceptions) sexually transmitted only maternally, being

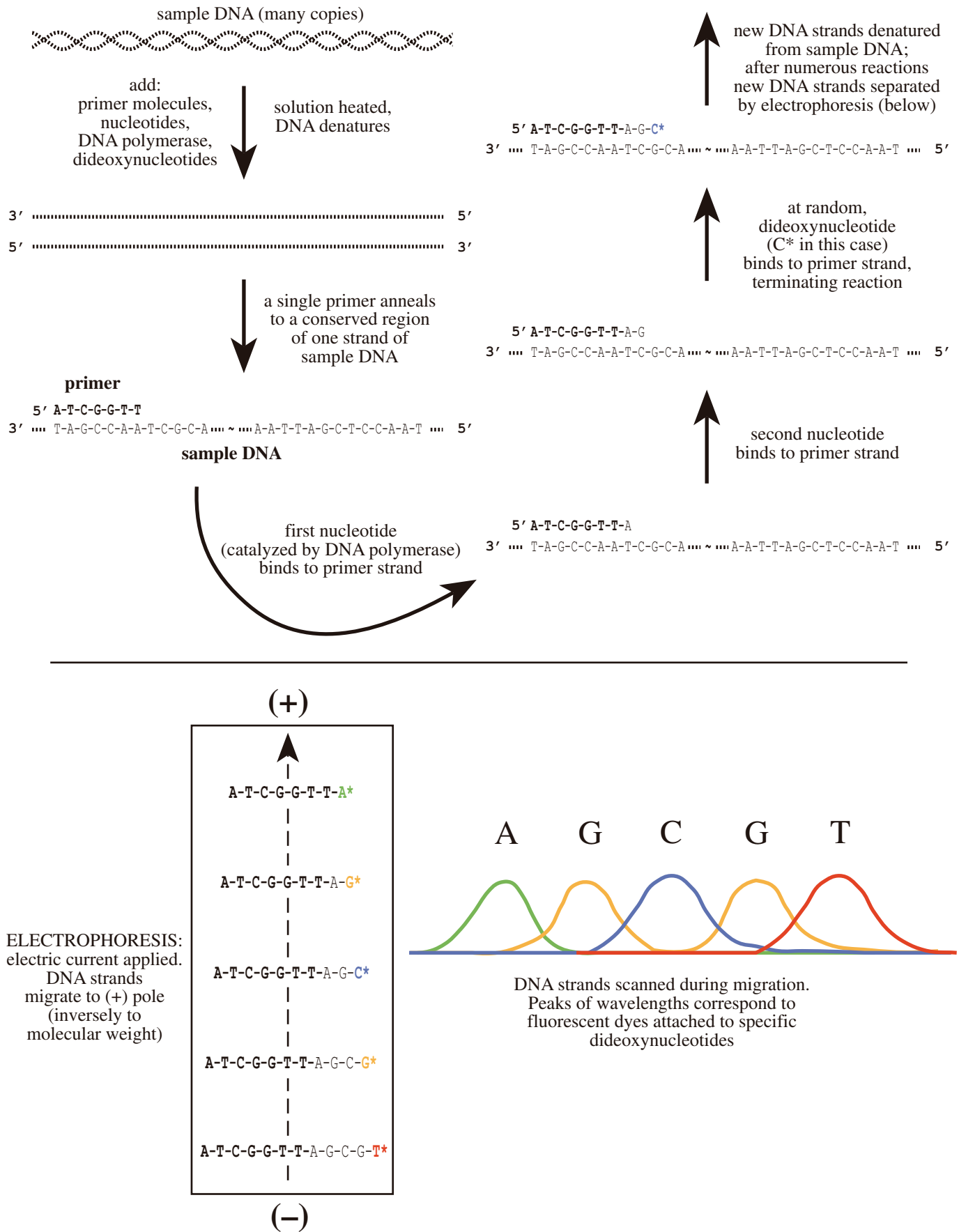


FIGURE 14.3 DNA sequencing reactions. A* = dideoxyadenine; C* = dideoxycytosine; G* = dideoxyguanine; T* = dideoxythymine.

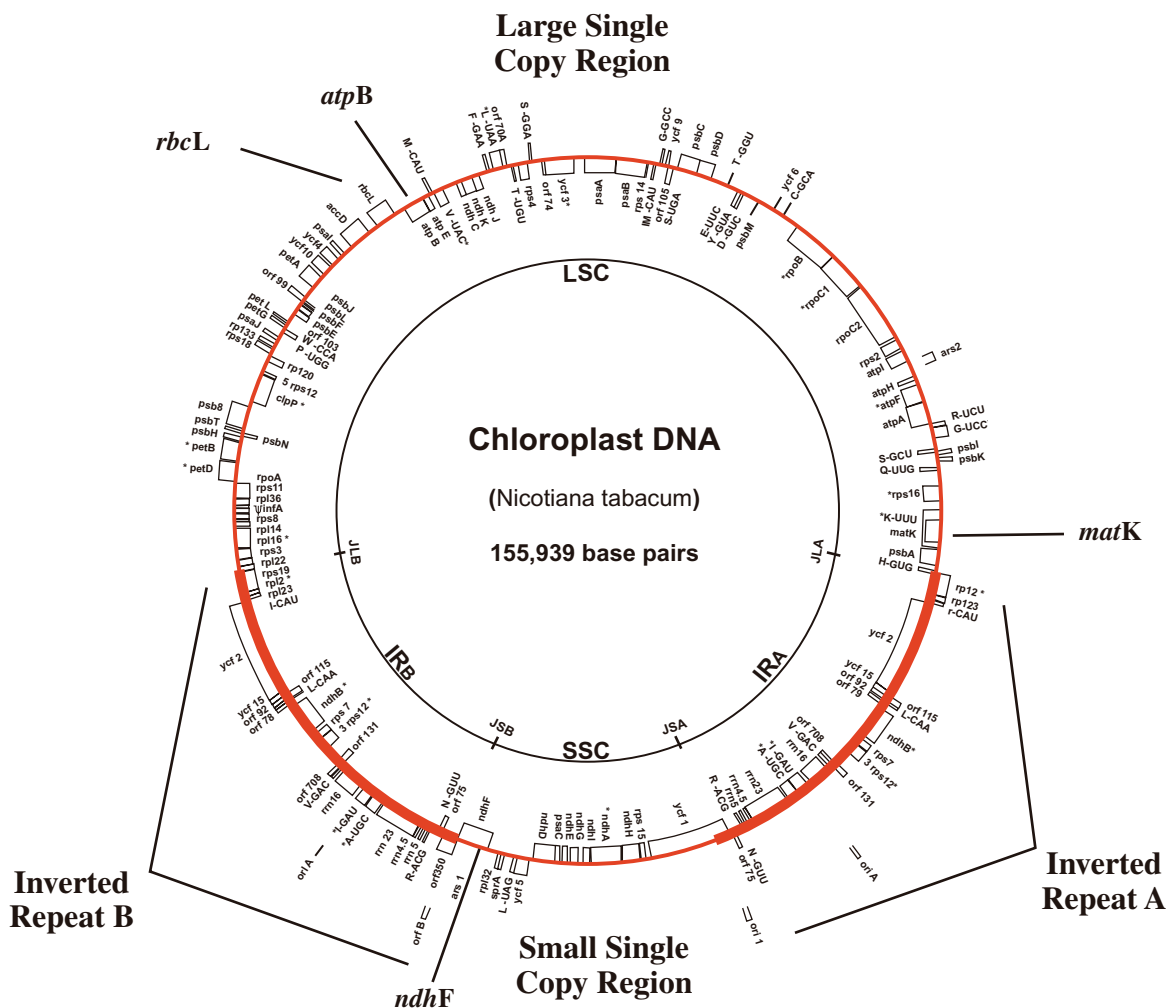


FIGURE 14.4 Molecular structure of the chloroplast DNA of tobacco (*Nicotiana tabacum*). Note large single-copy region (LSC), small single-copy region (SSC), and the two inverted repeats (IRA and IRB). Also note location of *atpB*, *rbcL*, *matK*, and *ndhF* genes (see Table 14.1). (Redrawn from Wakasugi, T., M. Sugita, T. Tsudzuki, and M. Sugiura. 1998. Updated gene map of tobacco chloroplast DNA. *Plant Molecular Biology Reporter* 16: 231–241, by permission.)

retained in the egg but excluded in sperm cells. (In conifers, interestingly, chloroplast DNA is transmitted paternally, not maternally.)

The use of sequence data from the DNA of chloroplasts has proven to be highly useful in elucidating both lower and higher level relationships. The basic structure of chloroplast DNA is shown in Figure 14.4. Like all organelle and prokaryotic DNA, it is circular. Curiously, most angiosperms have a region of chloroplast DNA known as the inverted repeat, which is the mirror image of the corresponding region (Figure 14.4). Some of the more commonly sequenced chloroplast DNA genes are listed in Table 14.1, although new ones are being utilized frequently.

Nuclear DNA sequencing has been used to a lesser degree in plant systematics. Some nuclear genes such as alcohol dehydrogenase (*Adh*), which has traditionally been used in allozyme studies, are becoming more frequently used. One of the more useful types of nuclear DNA sequences has been the internal transcribed spacer (ITS) region, which contains multiple DNA copies (as opposed to single copies found in most protein-coding genes). The ITS region lies between the 18S and 26S nuclear ribosomal DNA (nrDNA) (Figure 14.5). ITS sequence data has been most valuable for inferring phylogenetic relationships at a lower level, e.g., between closely related species. However, it has also been used in elucidating higher level relationships.

TABLE 14.1 Some chloroplast genes that have been used in plant molecular systematics, after Soltis et al. 1998.

<i>CHLOROPLAST GENES</i>		
GENE	LOCATION	FUNCTION
<i>atpB</i>	Large single-copy region of chloroplast	Beta subunit of ATP synthetase, which functions in the synthesis of ATP via proton translocation
<i>rbcL</i>	Large single-copy region of chloroplast	Large subunit of ribulose-1,5-bisphosphate carboxylase/oxygenase (RUBISCO), which functions in the initial fixation of carbon dioxide in the dark reactions
<i>matK</i>	Large single-copy region of chloroplast	Maturase, which functions in splicing type II introns from RNA transcripts
<i>ndhF</i>	Small single-copy region of chloroplast	Subunit of chloroplast NADH dehydrogenase, which functions in converting NADH to NAD + H ⁺ , driving various reactions of respiration

ANALYSIS OF DNA SEQUENCE DATA

DNA sequence data is converted to characters and character states to be used in phylogenetic analyses. First, the sequences of a given length of DNA are **aligned**, in which homologous nucleotide positions (e.g., corresponding to the same codon position of a given gene) are arranged in corresponding columns (Figure 14.6). For some genes that are relatively conserved, alignment is straightforward, as all taxa have the same number of nucleotides per gene. For other genes or DNA segments, some taxa may have one or more additions, deletions, inversions, or translocations relative to other taxa. The occurrence of these mutations, and/or the occurrence of considerable homoplasy among taxa, can make alignment of DNA sequences difficult. In addition, multiple copies of a gene can make homology assessment difficult. Various computer algorithms can be used to automatically align sequences of the taxa being studied, but these have assumptions that must be carefully assessed.

Generally, in using DNA sequence data in a phylogenetic analysis, a character is equivalent to the nucleotide position, and a character state of that character is the specific nucleotide at that position (there being four possible character states, corresponding to the four nucleotides; see Figure 14.6). A large number (often the great majority) of nucleotide positions are generally invariable among taxa, and some of the variable ones are often uninformative by being autapomorphic for a given taxon; thus, relatively few sites are informative and therefore useful in phylogenetic reconstruction (Figure 14.6).

However, a major addition, deletion, inversion, or translocation can in itself be identified as an evolutionary novelty (apomorphy), used in grouping lineages together. For example, members of the Faboideae (of the Fabaceae) lack, by deletion, one of the inverted repeats found in the chloroplasts of most angiosperms (see Figure 14.4). Chromosomal mutations such as these may be coded separately from single base

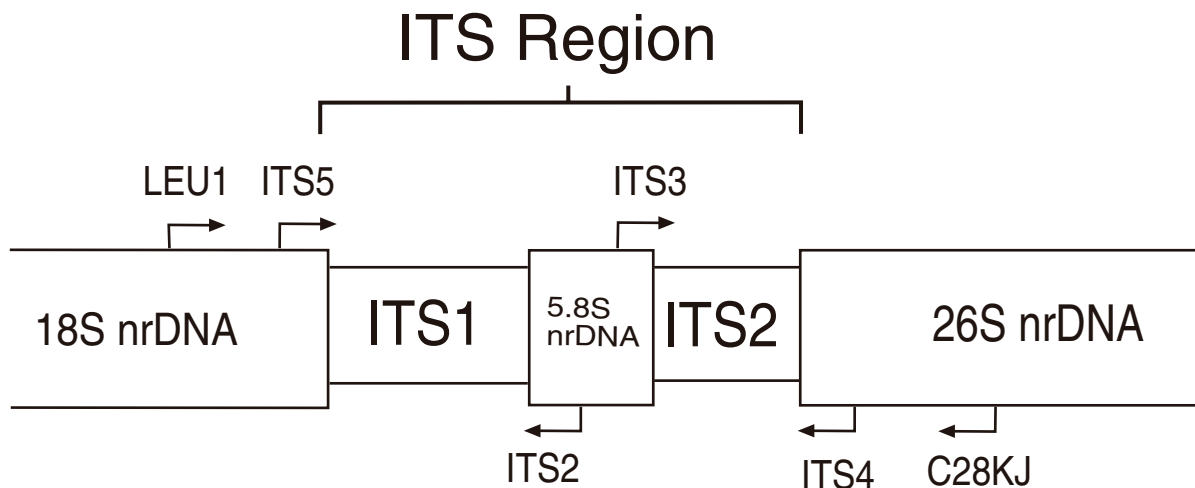


FIGURE 14.5 Internal transcribed spacers (ITSs) of nuclear ribosomal DNA, illustrating the ITS region and flanking subunits, and showing the orientations and locations of primer sites. After Baldwin et al. (1995).

DNA Alignment

Character Coding

	0000000000000000000011111111111111111111					
	888888888899999999990000000000111111111122					
	12345678901234567890123456789012345678901					
Taxon 1	GCCTAGCC AAAGCTCT TTCCAAG GTGACTCTCAGTTCAAGCT	2	0	3	2	0
Taxon 2	GCCTAGCC AAAGCTCT TTCCAAG CTGACTCTCA -----GCT	2	0	3	1	0
Taxon 3	GCCTAGCC TAAGCTCA ACCAAG GTGTCTCTCAGTTCAAGCT	2	3	0	2	3
Taxon 4	GCCTAGCC TAAGCTCT TTCCAAG GTGTCTCTCAGTTCAAGCT	2	3	3	2	3
Taxon 5	GCCTAGCC AAAGCTCT TTCCAAG CTGACTCTCA -----GCT	2	0	3	1	0
Taxon 6	CCCTAGCC AAAGCTCT TTCCAAG CTGACTCTCAGTTCAAGCT	1	0	3	1	0
Taxon 7	CCCTAGCC AAAGCTCT TTCCAAG CTGACTCTCAGTTCAAGCT	1	0	3	1	0
Taxon 8	GCCTAGCC TAAGCTCT TTCCAAG CTGACTCTCAGTTCAAGCT	2	3	3	1	0

FIGURE 14.6 Example of alignment of DNA sequences of 41 nucleotide sites (positions 81 121) from eight taxa. Variable nucleotide sites are in **bold**. Note deletion of six bases in taxon 2 and taxon 5. Possible character coding of variable sites is seen at right. Coding of nucleotides is as follows: A = state 0; C = state 1; G = state 2; T = state 3. In this example, the deletion is coded as a single binary character (character 6), coded differently from nucleotides, as state 4 = deletion absent and state 5 = deletion present.

differences (e.g., as in the example of Figure 14.6) and may be given relatively greater weight in inferring relationships.

Several types of weighting schemes may be done with molecular data. For protein encoding genes, the codon position may be differentially weighted. For example, because of redundancy of the genetic code, the third codon position is generally more labile (a change more likely to have occurred randomly) than the second, and the second may be more labile than the first. Thus, the first and second codon positions may be given relatively greater weight, respectively (such as a weight of 10 for the first codon position, 5 for the second position, and 1 for the third position). The logic here is that a change in codon position 1 or 2 is less likely to have occurred at random within a taxon and more likely represents evolutionary novelties that are shared among taxa. Weighting by codon position may be based on empirical data. For a given data set, the number of changes occurring for codon positions 1, 2, and 3 may be used (inversely) to establish the relative weights.

Another weighting parameter that may be used with DNA sequence data concerns transitions versus transversions. Transitions are evolutionary changes from one purine to another purine (A → G or G → A) or from one pyrimidine to another pyrimidine (C → T or T → C); see Figure 14.1. Transversions are evolutionary changes from a purine to a pyrimidine (A → C, A → T, G → C, or G → T) or from a pyrimidine to a purine (C → A, C → G, T → A, or T → G). Weighting using transitions versus transversions may be based on empirical data. For a given data set, the relative frequency of transitions versus transversions may be used (inversely) to establish the relative weights. For example, for a given group under study, if transitions occur 5× more frequently than

transversions, the latter may be given a weight of 5 and the former a weight of 1, as illustrated in the step matrix of Figure 14.7.

DNA sequence data can also be used to evaluate the secondary structure of a molecule. Thus, nucleotide differences that result in major changes in the conformation of the product (whether ribosomal RNA or protein) may have a much greater physiological effect than those that do not and might receive a higher weight. Computer algorithms can evaluate this to some degree.

Parsimony, maximum likelihood, and Bayesian methods are commonly used to infer phylogenetic relationships using DNA sequence data (Chapter 2). The most robust hypotheses of relationship have arisen from studies using sequence data from multiple (e.g., anywhere from 3 to 18), ideally unlinked, genes.

RESTRICTION SITE ANALYSIS (RFLPs)

A restriction site is a sequence of approximately 6 8 base pairs of DNA that binds to a given restriction enzyme.

	A	G	C	T
A	0	1	5	5
G	1	0	5	5
C	5	5	0	1
T	5	5	1	0

FIGURE 14.7 Step matrix of nucleotide changes, showing weighting scheme in which transversions are given a weight 5 times greater than that of transitions.

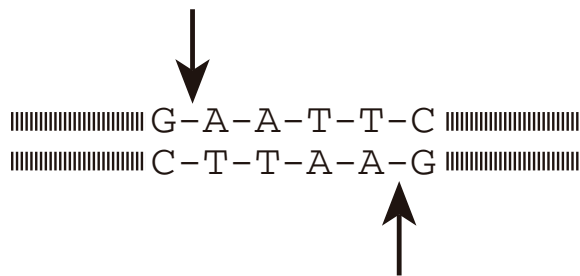


FIGURE 14.8 A DNA restriction site, cleaved (at arrows) by the restriction site enzyme *EcoRI*.

These restriction enzymes, of which there are many, have been isolated from bacteria. Their natural function is to inactivate invading viruses by cleaving the viral DNA. Restriction enzymes known as type II recognize restriction sites and cleave the DNA at particular locations within or near the restriction site. An example is the restriction enzyme *EcoRI* (named after *E. coli*, from which it was first isolated), which recognizes the DNA sequence seen in Figure 14.8 and cleaves the DNA at the sites indicated by the arrows in this figure.

Restriction fragment length polymorphism, or RFLP, refers to differences between taxa in restriction sites, and therefore the lengths of fragments of DNA following cleavage with restriction enzymes. For example, Figure 14.9 shows, for two hypothetical species, amplified DNA lengths of 10,000 base pairs that are subjected to (digested with) the restriction enzyme *EcoRI*. Note, after a reaction with the *EcoRI* enzyme, that the DNA of species *A* is cleaved into three fragments, corresponding to two *EcoRI* restriction sites, whereas that of species *B* is cleaved into four fragments, corresponding to three *EcoRI* restriction sites. The relative locations of these restriction sites on the DNA can be mapped; one possibility is seen at the bottom of Figure 14.9. (Note that there are other possibilities for this map; precise mapping requires additional work.) Additional restriction enzymes can be used. Figure 14.10 illustrates how each of the DNA fragments from the *EcoRI* digests can be digested with the *Bam* HI restriction enzyme, yielding different fragments for the two species. These data can be added to the original in preparing a map (one possible map is shown in lower part of Figure 14.10).

Restriction site fragment data can be coded as characters and character states in a phylogenetic analysis. For example, given that the restriction site maps of Figure 14.10 are correct, the presence or absence of these sites can be coded as characters, as seen in Figure 14.11. Restriction site analysis contains far less data than complete DNA sequencing, accounting only for the presence or absence of sites 6–8 base

pairs long. It has the advantage, however, of surveying considerably larger segments of DNA. However, with improved and less expensive sequencing techniques, it is less valuable and less often used than in the past.

ALLOZYMES

Allozymes are different molecular forms of an enzyme that correspond to different alleles of a common gene (*locus*). (This is not to be confused with *isozymes*, which are forms of an enzyme that are derived from separate genes or loci.) Allozymes are traditionally detected using electrophoresis, in which the enzymes are extracted and placed on a medium (e.g., starch) through which an electric current runs (similar to gel electrophoresis in DNA sequencing). A given enzyme will migrate toward one pole or the other depending on its charge. Similarly, different allozymes of an enzyme will migrate differentially because they differ slightly in amino acid composition and therefore have somewhat different electrical charges. Allozymes subjected to electrophoresis are identified with a stain specific to that enzyme and the bands marked by their relative position on the electrophoresis medium.

Allozymes have traditionally been used to assess genetic variation within a population or species, but they can also be used as data in phylogenetic analyses of closely related species, e.g., species within a monophyletic genus. Figure 14.12A illustrates an example of electrophoretic allozyme banding data for five species and an outgroup.

There are several ways to code polymorphic allozyme data. One way is to code each allele as a character and the presence or absence of that allele as a character state. A second way to code allozyme data is to treat the locus (corresponding to the gene coding for the enzyme) as the character and all unique combinations of alleles as character states (as in Figure 14.12B). The number of state changes between these unique allelic combinations can be a default of one. However, another method of coding is to treat the loss of each allele as one state change and the gain of an allele as a separate state change. Thus, the number of state changes between different allelic combinations can vary, as seen in Figure 14.12C. Step matrices (see Chapter 2) are used to code these in a cladistic analysis.

Yet another way to code allozyme data is to take into account the frequency of alleles present in a given taxon. For example, by this method, species *A*, which has allele X present with a frequency of 95% and allele Y with a frequency of 5%, would receive a different coding from species *B*, which has the same alleles, but in frequencies of 55% and 45%, respectively.

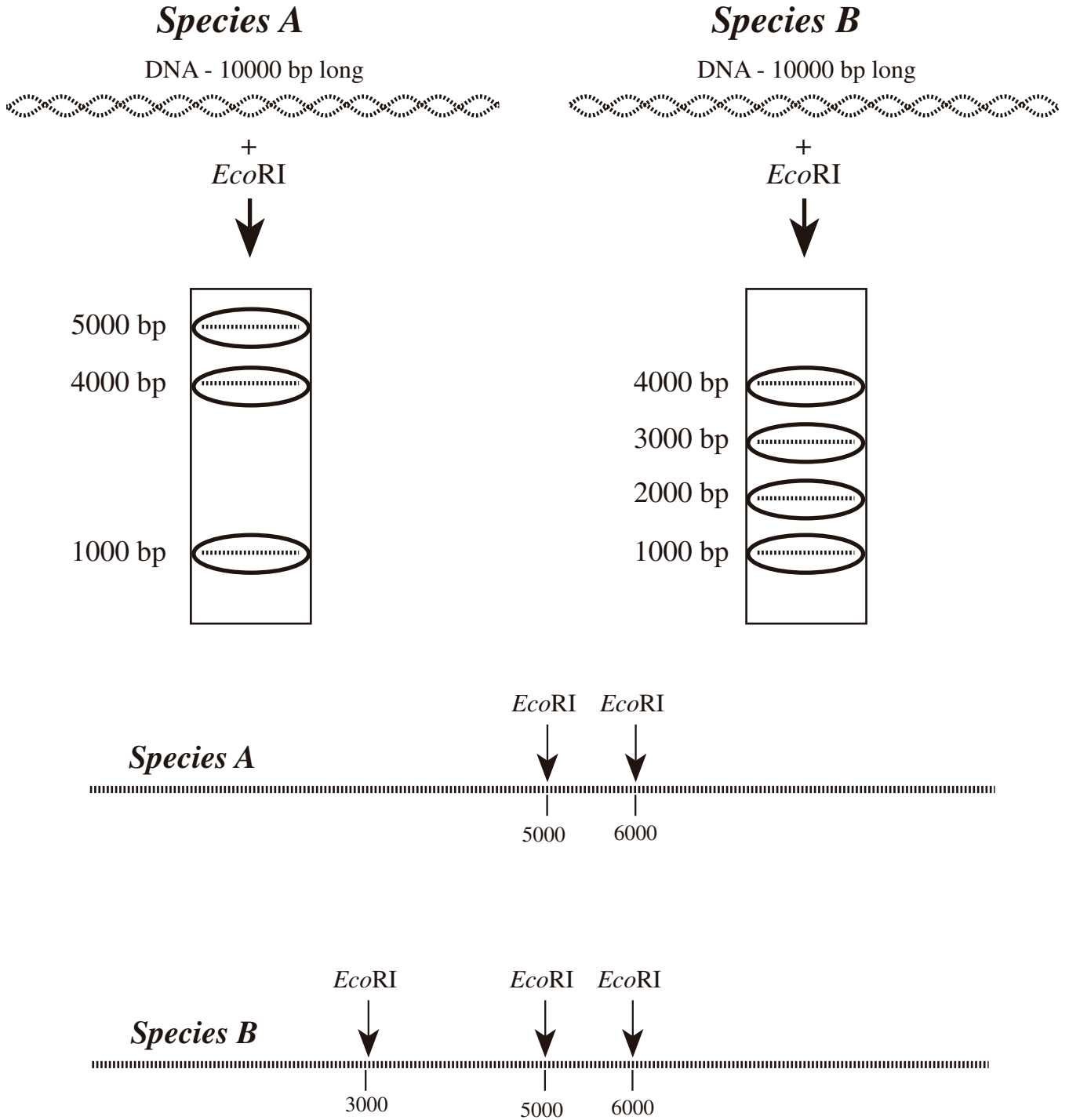


FIGURE 14.9 Example of restriction site analysis of species A and B, using restriction site enzyme *EcoRI*. Note differences in fragment lengths. Possible restriction site maps of species A and B are shown in the lower portion of the figure.

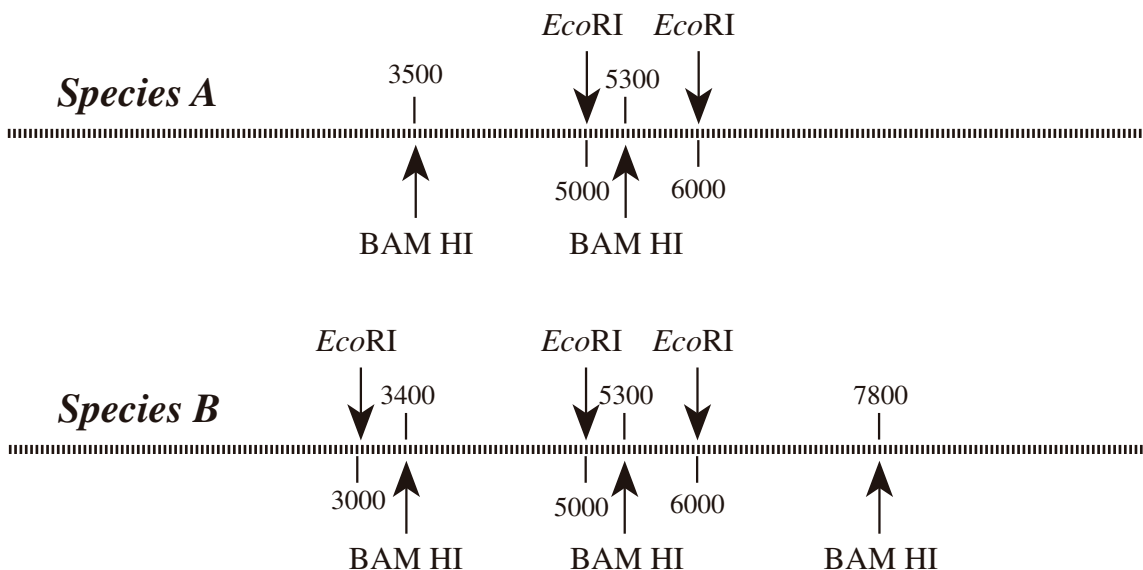
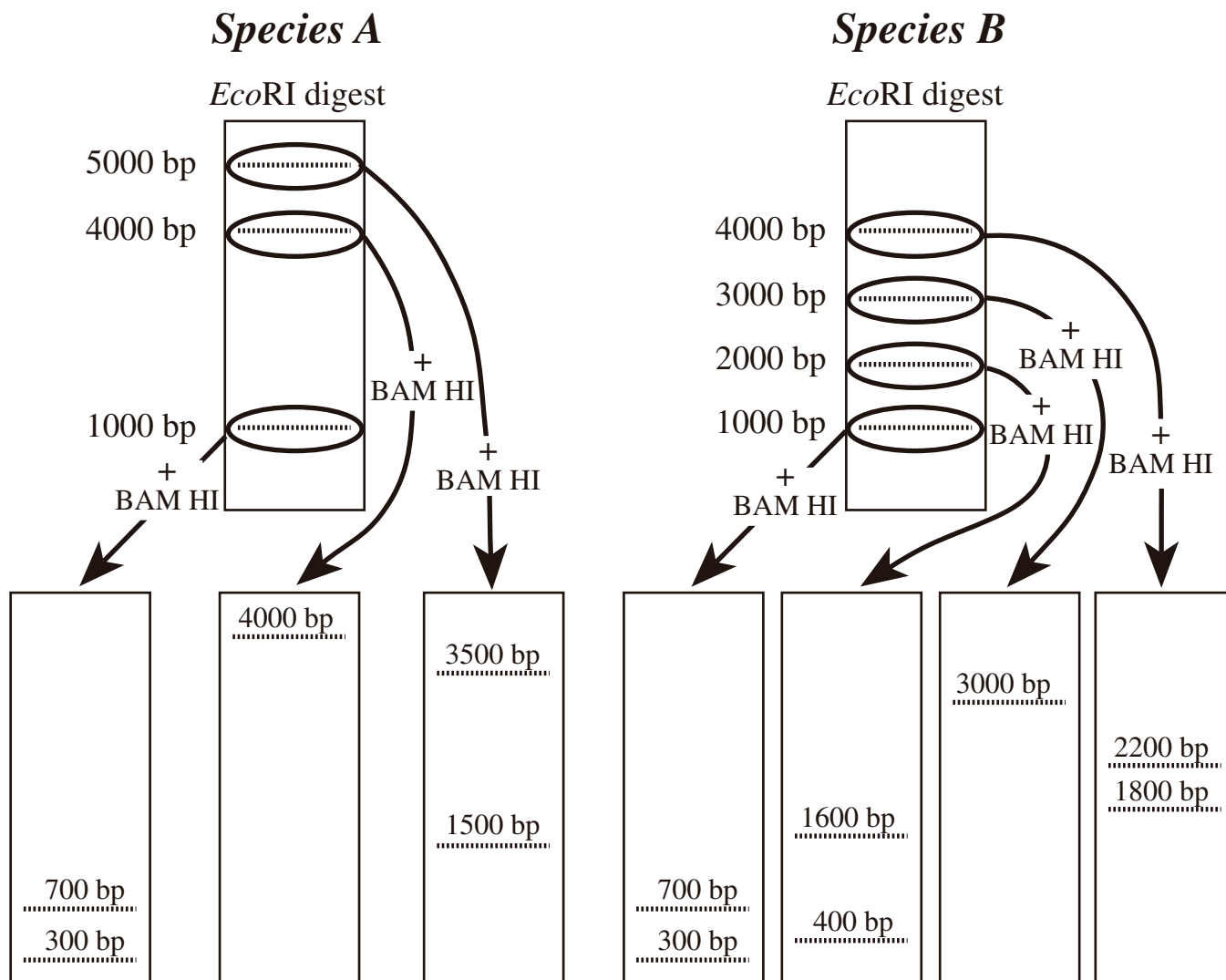


FIGURE 14.10 Example of restriction site analysis of species A and B, using restriction site enzyme *EcoRI*, followed by restriction site enzyme *BAM HI*. Possible restriction site maps of species A and B are shown in the lower portion of the figure.

TAXA	CHARACTERS							
	<i>Eco</i> RI	BAM	BAM	<i>Eco</i> RI	BAM	<i>Eco</i> RI	BAM	
	3000	3400	3500	5000	5300	6000	7800	
Species A	–	–	+	+	+	+	–	
Species B	+	+	–	+	+	+	+	

FIGURE 14.11 Character coding of restriction site map data of Figure 14.10, derived by presence or absence of *Eco*RI or BAM sites at specific locations along DNA.

MICROSATELLITE DNA

Microsatellites are regions of DNA that contain short (usually 2–5) repeats of nucleotides, an example being TGTGTG, in which two base pairs repeat. The regions are termed **tandem repeats**; if they vary within a population or species, they are called **variable-number tandem repeats (VNTR)**. (Other designations and acronyms are used, depending on the particular field of study.) These tandem repeats can be located all across the genome; at a given location (locus), the repeat will tend to be of a certain length. However, individuals within or between populations may vary in the number of tandem repeats at a given locus (or even show allelic variation) because of irregularities in crossing-over and replication. Thus, variable-number tandem repeats can be used as a genetic marker.

Microsatellites are identified by constructing primers that flank the tandem repeats and then using PCR technology. (The primers are initially identified for a species by the time-consuming process of synthesizing genetic probes of a tandem repeat, screening DNA for binding to these probes, and sequencing these regions to design primers that flank the tandem repeats.) Once the primers are identified, PCR can be used to quickly generate multiple copies of the tandem repeat DNA, the length of which (for a given individual at a given locus or allele) can be determined by gel electrophoresis. (See example in Figure 14.13.)

Microsatellite analysis can generate data quickly and efficiently (once the primers are identified for a given group) for a large number of individuals. It is most often used for population studies, e.g., to assess genetic variation or homozygosity. Its use in systematics is largely in examining relationships within a species (such as to assess infraspecific classifications) or between very closely related species.

RANDOM AMPLIFIED POLYMORPHIC DNA (RAPDs)

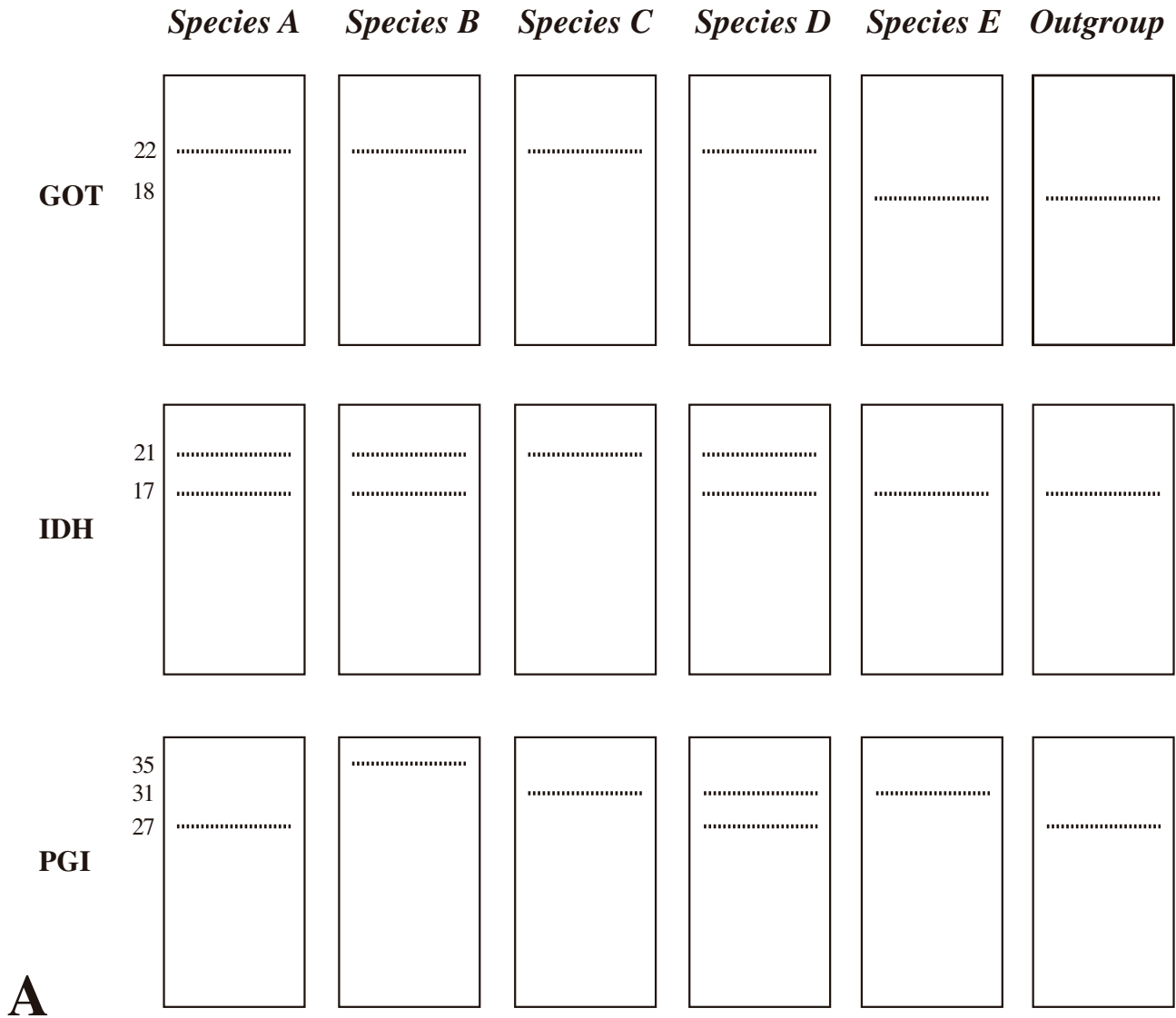
Another method of identifying genetic markers is by using a *randomly* synthesized primer to amplify DNA in a PCR reaction.

In this method, the primer will anneal to complementary regions located in various locations of isolated DNA. If another complementary site is present on the opposing DNA strand at a distance that is not too great (i.e., within the limits of PCR), then the reaction will amplify this region of DNA (Figure 14.14). Because many sections of DNA complementary to the primer may occur, the PCR reaction will result in DNA strands of many different lengths, which can be size-separated by electrophoresis. Because even closely related individuals may show some sequence variation that may determine potential primer sites, these different individuals will show different amplification products. Thus, **RAPD** refers to using **randomly** generated primers for the **amplification** of DNA to identify **polymorphic DNA** regions of different individuals or taxa. (See example in Figure 14.14.)

RAPDs, like microsatellites, may often be used for within-species genetic studies, but may also be successfully employed in phylogenetic studies to address relationships within a species or between closely related species. However, RAPD analysis has the major disadvantages in that results are difficult to replicate (being very sensitive to PCR conditions) and in that the homology of similar bands in different taxa may be unclear.

AMPLIFIED FRAGMENT LENGTH POLYMORPHISM (AFLPs)

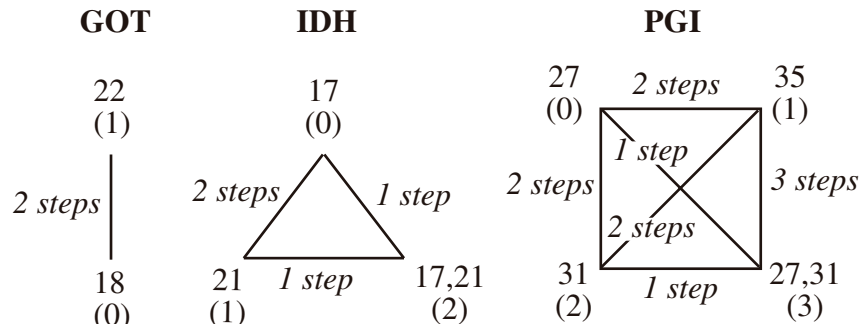
This method is similar to that of identifying RFLPs in that a restriction enzyme is used (Figure 14.15A) to cut DNA into numerous, smaller pieces, each of which (because of the action of the restriction enzymes) terminates in a characteristic nucleotide sequence (Figure 14.15B). However, the numerous, cut DNA fragments are then modified by binding to each end (using DNA ligase) a synthesized, double-stranded piece of DNA, known as a **primer adapter** (Figure 14.15C). The primer adapters are designed to insert at the cut ends (corresponding to the complementary sequences of the restriction enzymes). Primers are then constructed that bind to the primer adapters and amplify the DNA fragments using



A

	GOT	IDH	PGI
A	1	2	0
B	1	2	1
C	1	1	2
D	1	2	3
E	0	0	2
OUT	0	0	0

B



C

FIGURE 14.12 Allozyme data. **A.** Hypothetical allozyme banding data for taxa A–E and *Outgroup* and enzymes GOT, IDH, and PGI. **B.** Coding of data using the locus as the character and unique allelic combinations as character states. **C.** One possible coding of data (after Mabee and Humphries, 1993). Diagrams illustrating number of state changes between character states (state number in parentheses); each loss or gain of an allele counts for one step change.

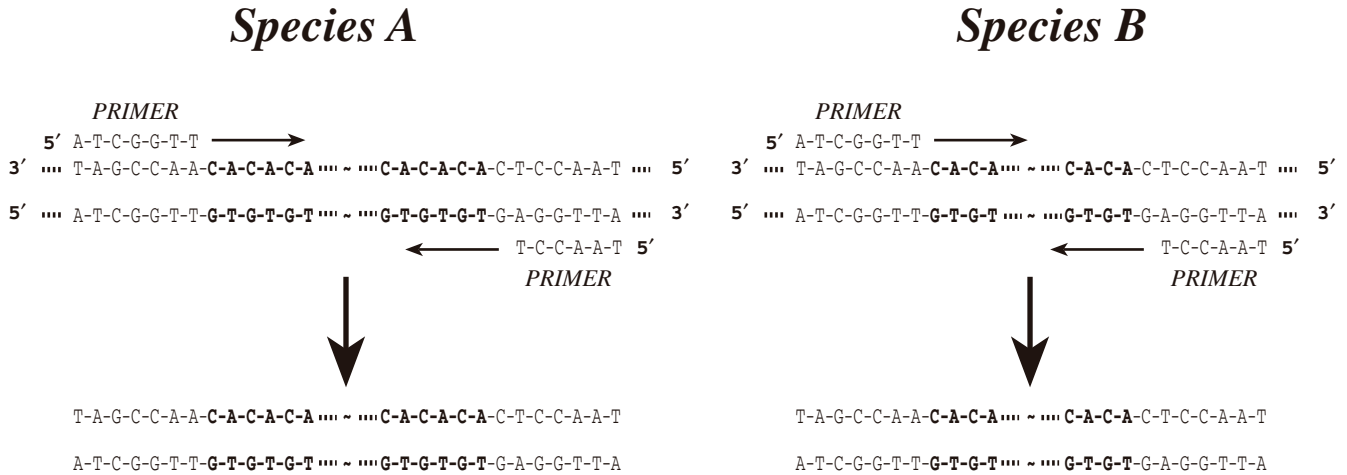


FIGURE 14.13 Microsatellite data. Primers were constructed to flank regions of tandem repeats. Note that tandem repeat region of species A is longer than that of species B and is thus a genetic difference between the two.

a polymerase chain reaction (Figure 14.15D). Electrophoresis separates the **amplified DNA fragments** that exhibit **length polymorphism** (hence, **AFLP**), enabling the recognition of numerous genetic markers.

AFLP data are more experimentally replicable than are RAPD data and can be used to identify genetic differences among individuals using large pieces of DNA. AFLP has one disadvantage

in that so many fragments may be generated that it is hard to distinguish them on an electrophoretic gel. However, a slight modification of the primers used may limit the number of fragments that are amplified, enabling them to be more easily identified. AFLP is largely used for population genetics studies, but has been used in studies of closely related species and even, in some cases, for higher-level, cladistic analyses.

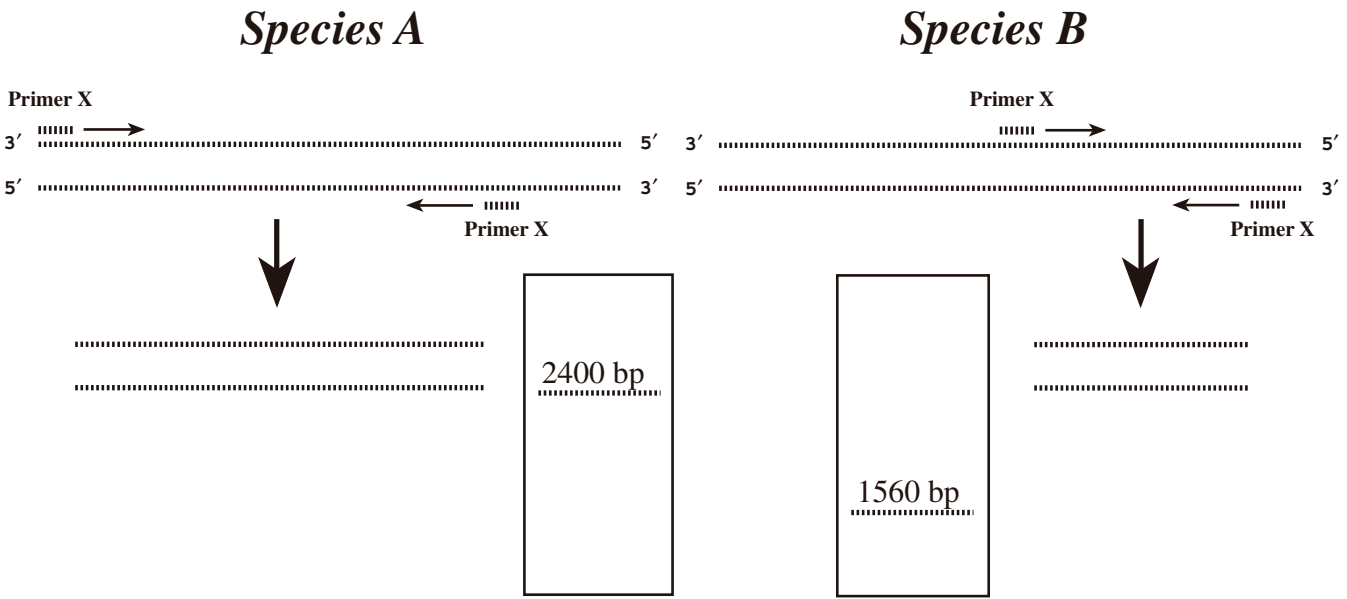


FIGURE 14.14 RAPDs data. In this example the same DNA regions for species A and B anneal to different randomly generated primers, resulting in amplified DNA of different lengths, a genetic difference between the two taxa.

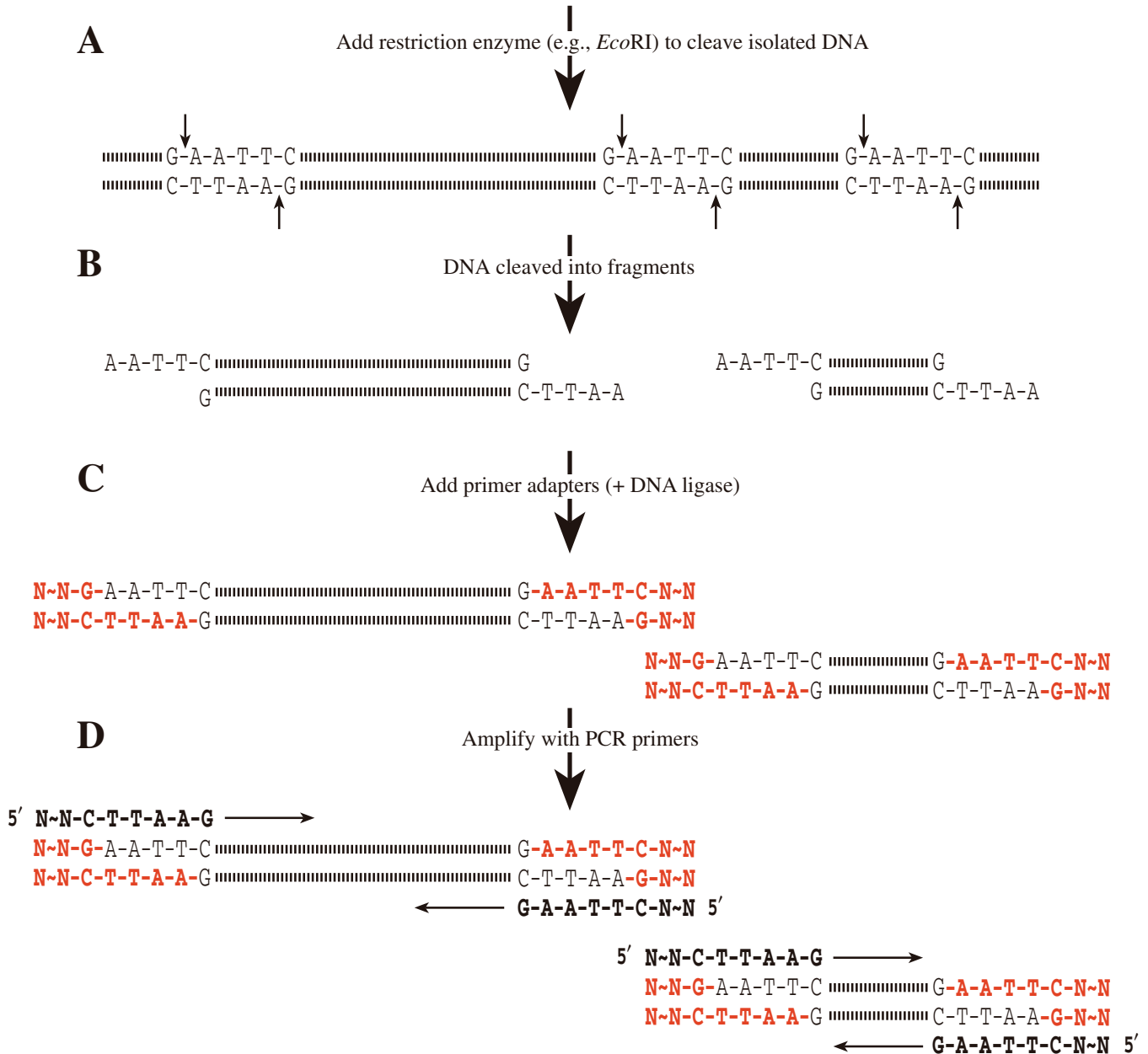


FIGURE 14.15 AFLP technique. The letters N~N- represent a length of nucleotides.

REVIEW QUESTIONS

1. Name the specific types of data used in studies of molecular systematics.
2. How are samples used to acquire molecular data typically processed?
3. Why is collection of a voucher specimen in molecular studies essential?
4. What does *DNA sequence data* refer to?
5. Explain the polymerase chain reaction and its importance in molecular systematics.
6. What is a primer?
7. Explain the basic process of automated DNA sequencing. What is the significance of dideoxynucleotides?
8. What are the three major types of DNA used in DNA sequence (and other molecular) studies?

9. In chloroplast DNA, what are the large single-copy region, small single-copy region, and inverted repeats?
10. Name some useful chloroplast genes used in plant molecular systematics.
11. What is the internal transcribed spacer region (ITS) and what is its efficacy in plant molecular systematics?
12. What is DNA alignment, and what are potential problems with this?
13. In general, what are the characters and character states for DNA sequence data?
14. Name the ways that DNA sequence data may be weighted in a cladistic analysis.
15. What is a restriction site?
16. What does restriction fragment length polymorphism (RFLP) refer to?
17. How is RFLP data acquired and how is it used in a cladistic analysis?
18. What is an allozyme?
19. How are allozyme data acquired?
20. Explain the different ways to code allozyme data in a cladistic analysis.
21. What are microsatellites and how are these data obtained?
22. What are random amplified polymorphic DNAs (RAPDs) and how are these data obtained?
23. Describe the technique for generating amplified fragment length polymorphisms (AFLPs), citing how this differs from that of generating RFLPs.

EXERCISES

1. If possible, get a demonstration of the various techniques of molecular systematics, e.g., DNA extraction and sequencing. Consider a special topics project in which you define a problem and use these techniques to acquire the data to answer the problem.
2. Access GenBank (<http://www.ncbi.nih.gov/Genbank>) and acquire molecular data on a particular group of choice. Consider analyzing these data using phylogenetic inference software (see Chapter 2).
3. Peruse journal articles in plant systematics, e.g., *American Journal of Botany*, *Annals of the Missouri Botanical Garden*, *Systematic Botany*, or *Taxon*, and note those that describe the use of molecular data in relation to systematic studies. Identify the techniques used, data acquired, and problems addressed.

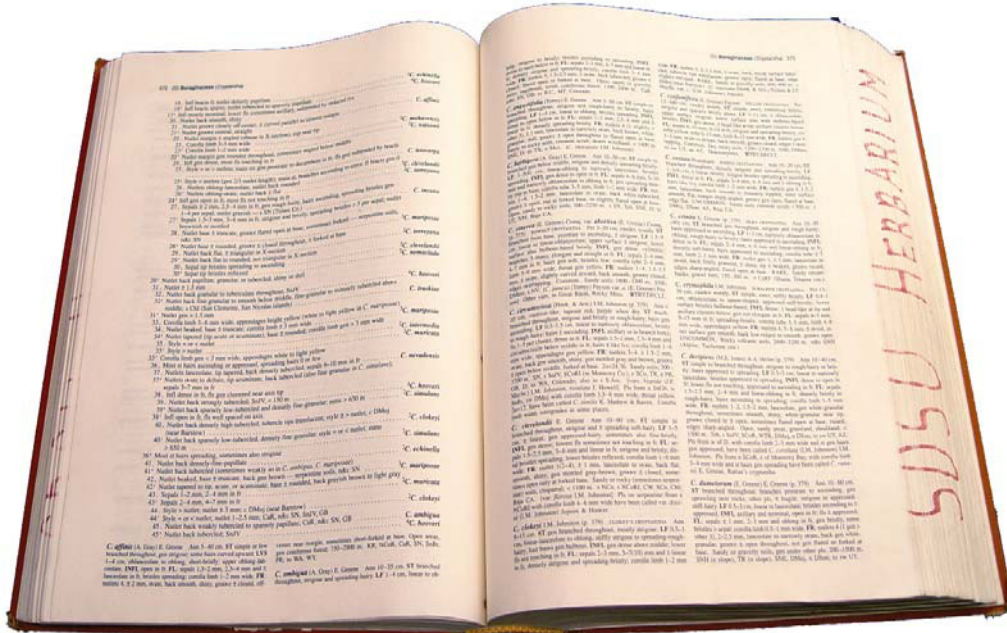
REFERENCES FOR FURTHER STUDY

- Baldwin, B. G., M. J. Sanderson, J. M. Porter, M. F. Wojciechowski, C. S. Campbell, and M. J. Donoghue. 1995. The ITS region of nuclear ribosomal DNA: a valuable source of evidence on angiosperm phylogeny. *Annals of the Missouri Botanical Garden* 82: 247-277.
- Hillis, D. M., C. Moritz, and B. K. Mable. 1996. *Molecular Systematics*. 2nd ed. Sinauer, Sunderland, Massachusetts.
- Mabee, P. M., and J. Humphries. 1993. Coding polymorphic data: examples from allozymes and ontogeny. *Systematic Biology* 42: 166-181.
- Soltis, D. E., P. S. Soltis, and J. J. Doyle (eds.). 1998. *Molecular Systematics of Plants II: DNA Sequencing*. Kluwer Academic, Boston.
- Soltis, P. S., D. E. Soltis, and J. J. Doyle (eds.). 1992. *Molecular Systematics of Plants*. Chapman and Hall, New York.

This Page Intentionally Left Blank

IV

RESOURCES IN PLANT SYSTEMATICS



This Page Intentionally Left Blank

15

PLANT IDENTIFICATION

METHODS OF PLANT IDENTIFICATION	495	PRACTICAL IDENTIFICATION	498
Taxonomic Keys	495	REVIEW QUESTIONS	498
Written Description	497	EXERCISES	499
Specimen Comparison	497	REFERENCES FOR FURTHER STUDY	499
Image Comparison	498		
Expert Determination	498		

Identification is the process of associating an unknown entity with a known entity (or recognizing that the unknown entity does not have a known counterpart). In other words, identification is a judgement that some perceived object is similar enough to a known entity that it falls within the criteria of belonging to the same class as the known entity. Identification is a basic activity of humans and other animals (and perhaps of all life at some level of organization). The ability to recognize, e.g., edible from toxic or friend from foe has undoubtedly evolved via strong selective pressure.

Because two entities are never exactly the same, a critical consideration in identification is determining the characteristics or boundaries of the known. For example, plant identification entails studying a plant or plant specimen and making a decision as to whether the plant belongs to a particular taxon, e.g., to a species. This identification rests on the prior description, both of the unknown plant and of the taxa that are possibilities. In considering the possible taxa, it is critical to evaluate the **diagnostic characterization** (= **diagnosis**) of each, which is a listing of all the features of a taxon that distinguish it from all other taxa. If the characteristics of the unknown entity fall within the range of the diagnosis of a known one, then an identification is made.

METHODS OF PLANT IDENTIFICATION

As mentioned earlier, identification first necessitates describing the plant in question. This can involve anything from a quick glance of a flower to a complete morphological

characterization of vegetative and reproductive features. The second criterion of plant identification is having a list of the possibilities. Thus, it is important that whatever references are used, they incorporate all of the possibilities. For native or naturalized plants, regional floras are typically used in the process of identification. One should check the geographic range of the flora used to be sure that it encompasses that of the unknown plant.

Cultivated plants can be particularly difficult to identify. This is true in part because the number of plants taken into cultivation is quite large and continues to expand every year. Because a cultivated plant can be native to any region of the world, one can rarely be certain of having a reference that will include the correct taxon. Cultivated plant species may also be difficult to identify because they may include a great number of cultivars, hybrids, or other breeds that are continually being introduced and may be quite different in appearance from an original native species.

There are several methods of identifying plants. These are described below.

TAXONOMIC KEYS

Perhaps the most useful method of identification is a taxonomic key. A **key** is an identification device that consists of sequentially choosing among a list of possibilities until the possibilities are narrowed down to one. Most keys are practical, narrowing down the identity of a taxon in the most efficient and effective means. The key may or may not split a larger group into smaller, natural (monophyletic) subgroups.

As reviewed in Chapter 1, the most common type of key, particularly in floras and monographic treatments, is a

dichotomous key. This consists of a sequence of two contrasting statements, each statement known as a **lead**; the two leads together comprise a **couplet**. The leads of a couplet may be indented and/or numbered. Identification proceeds by choosing between the contrasting leads of a couplet. That lead which best fits the organism to be identified is selected; then all couplets hierarchically beneath that lead (either by indentation or numbering) are sequentially checked until an identification is obtained (see Figure 15.1). A well-written dichotomous key may have several types of evidence presented, with every character of the first lead matched, respectively, in the second lead (Figure 15.1).

Most keys are *artificial* or *practical*, meaning that the sequential groupings of the key do not intentionally reflect natural groups; their goal is to most easily and efficiently identify a given taxon, with no concern about classification

into other groups. Rarely, a key may be *natural* or *phylogenetic*, in which diagnostic (or even apomorphic) features are used to delimit natural groups, which are usually formal taxa. An example of a natural key might be one to the tribes of the Asteraceae. More technical, but less obvious, characters are used in natural keys, so they are generally less useful in practical identification, but may denote the features used to separate taxonomic groups.

Some precautions should be taken in using a dichotomous key. Most important is to *read all parts of both leads* before making a decision as to which fits the plant best. Never read just the first lead; although it might seem to fit, the second lead may fit even better. If, after reading both leads of a couplet, you are not certain which is correct, both should be considered. The two (or more) possibilities attained can then be checked against descriptions, illustrations, or specimen comparisons.

Haemodoraceae BLOODWORT FAMILY

1. Fertile stamens 3 or 1 (Subfamily Haemodoroideae)
 2. Ovary superior
 3. Fertile stamen 1 *Pyrrorhiza*
 - 3 Fertile stamens 3
 4. Corolla actinomorphic
 5. In orescence a simple raceme; functional carpel 1; ovule 1; style subterminal *Barberetta*
 - 5 In orescence an elongate thyse with lateral monochasial cymes; functional carpels 3; ovules 20-30 per carpel; style terminal *Xiphidium*
 - 4 Corolla zygomorphic
 6. Stamens unequal, the 2 latero-posterior anthers reduced; ovules 3-4 per carpel *Schiekia*
 - 6 Stamens equal; ovule 1 per carpel *Wachendor a*
 - 2 Ovary inferior
 7. Ovule 1 per carpel *Dilatris*
 - 7 Ovules 2 or more per carpel
 8. Ovules 2 per carpel; perianth glabrous *Haemodorum*
 - 8 Ovules 5-7 per carpel; perianth abaxially tomentose *Lachnanthes*
- 1 Fertile stamens 6 (Subfamily Conostylidoideae)
 9. Flowers actinomorphic; perianth not splitting along mid-anterior line
 10. Perianth glabrous to glabrate *Phlebocarya*
 - 10 Perianth lanate to tomentose
 11. Perianth lanate; trichomes simple to sparsely branched, white-whitish; anthers with broad, apical connective appendage *Tribonanthes*
 - 11 Perianth tomentose, trichomes dendritic, yellow, whitish, reddish, pink, orange, or purplish; anthers without broad, apical connective appendage
 12. Flowers pendulous; perianth reddish to pink-orange *Blancoa*
 - 12 Flowers generally ascending; perianth usually yellow or whitish, rarely orange to purplish *Conostylis*
 - 9 Flowers zygomorphic; perianth tube splitting along mid-anterior line
 13. Ovule 1 per carpel; perianth trichomes black *Macropidia*
 - 13 Ovules >1 per carpel; perianth trichomes red, yellow, orange or green *Anigozanthos*

FIGURE 15.1 Example of an indented and numbered dichotomous key: to the genera of the family Haemodoraceae. Note that character states of the first lead of a couplet are matched by corresponding character states in the second lead. From the author.

- | | |
|--|---|
| <ul style="list-style-type: none"> 1 Woody plants (excl. suffrutices) 2 Herbaceous plants (incl. suffrutices) 3 Aquatic plants, leaves floating or submerged 4 Chlorophyll absent (parasites or saprophytes) 5 Bulb present (monocots only) 6 Milky juice present 7 Spiny stems or leaves 8 Tendrils present 9 Cladodes or phyllodes (modified branches or petioles) 10 Hairs glandular 11 Hairs stellate (also 2-armed, branched and tufted) 12 Hairs stellate (not 2-armed, branched and tufted) 13 Hairs 2-armed or t-shaped, nonglandular 14 Hairs branched 15 Hairs tufted, nonglandular 16 Hairs peltate or scalelike 17 Hairs vesicular or bladderlike | <ul style="list-style-type: none"> 18 Hairs stinging 19 Cystoliths present (dicots only) 20 Leaves opposite or verticillate 21 Leaves alternate (excl. distichous monocots) 22 Leaves distichous (monocots only) 23 Leaves equitant (e.g. Iris) 24 Leaves not compound 25 Leaves compound 26 Leaves pinnately compound (4 or more leaflets) 27 Leaves ternately compound (3 leaflets) 28 Leaves palmately compound (4 or more leaflets) 29 Venation pinnate or hardly visible in leaves or leaflets (incl. no. 30) 30 Venation invisible or leaves 1-nerved (monocots only) 31 Venation longitudinal in leaves or leaflets (incl. 3-nerved leaves) 32 Venation palmate in leaves or leaflets |
|--|---|

FIGURE 15.2 Example of a polyclave key. Note that the key consists of a list of numerous character states. Identification proceeds by indicating which subset of character states describe the unknown plant. Not all characters need be considered.

Another type of identification device is the polyclave key. A **polyclave key** consists of a list of numerous character states, whereby the user selects all of states that match the specimen (e.g., Figure 15.2). Based on which of the many character states are a match, the correct taxon (or closest match) can be determined or narrowed down to a smaller subset of the possibilities. All polyclave keys in use today are implemented by a computer algorithm.

The great advantage of polyclave keys over dichotomous ones is that they permit the use of a limited subset of information to at least narrow down the possibilities. For example, if a dichotomous key lists only floral characters, its usefulness may be limited if your plant specimen lacks flowers. A polyclave key, however, will have a listing of not only floral characters, but also features of the roots, stems, leaves, fruits, and seeds. Thus, the polyclave key will often enable the user to identify the plant, even if one or more types of data are missing from the specimen. A second advantage of polyclave keys is that if the specimen cannot be absolutely identified, its identity may at least be narrowed down to a few alternatives, which can then be checked by other means. The only major disadvantage of polyclave keys is their availability; they have generally been written only for a limited number of taxonomic groups.

Although keys are probably the most practical and utilized method of identification, they should be regarded as guides, not foolproof methods. Any identification attained should be checked by other means, such as specimen comparison or expert determination (discussed later).

WRITTEN DESCRIPTION

A second means of identification is to compare features of the unknown plant with written descriptions of the possible known taxa. This is a good method of determining with certainty whether the range of variation of the unknown plant corresponds to that listed in the description of a known plant. However, because reading all of the written descriptions of a flora is impractical, this method relies on narrowing down the possibilities first. In addition, gleaning the diagnostic characteristics from a long list of features may be difficult. Thus, written descriptions are best used to verify an identity after one or a few possibilities are presented.

SPECIMEN COMPARISON

A third method of identification is to compare the plant in question to a live or preserved plant collection, usually an identified herbarium specimen. This is an excellent method of identification, as many features of a plant (e.g., coloration and surface features) are often not adequately denoted in written descriptions or visible from photographs or illustrations. As with the foregoing methods, comparison to an herbarium specimen is practically limited to verifying an identity after a subset of possibilities is narrowed down. **Synoptic collections**, which house generally one specimen of each taxon for a given region (e.g., a county), are very useful in this regard. If a taxon can be narrowed to a smaller group, such as a family or genus, a quick search through a synoptic collection for that region may often allow for site identification of the unknown. One precaution about this method,

however, is that it is dependent on the fact that the herbarium specimens are themselves correctly identified. Thus, a possible match should always be verified with a written description.

IMAGE COMPARISON

A fourth method by which an unknown plant may be identified is by visually comparing it to photographs or illustrations of known taxa. These are usually obtained from books, although Webpage images have now become a very useful resource. A practical problem with this method is that photographs and illustrations are usually available only for a small subset of possible taxa. In addition, it may be cumbersome to locate the matching photograph or illustration, necessitating an examination of all those available. However, visual comparison to an image can still be an excellent way to identify a plant, particularly if the possibilities can be narrowed down beforehand. The major precaution about this method is that two or more taxa may look very similar to one another as based on a photograph or illustration; the differences between them may reside on obscure morphological features that are not easily visible. Thus, any match of the unknown to a visual image should be confirmed with a technical description of the plant.

EXPERT DETERMINATION

A fifth and final means of identification is simply to ask someone else, preferably an expert in the group in question. This method may be time-consuming, as it usually requires sending a specimen away for identification (as well as knowing *who* the experts of a given group are). However, expert identification is perhaps the best way to identify a specimen, as the expert will usually know the taxa of that group over a wide geographic range. If the expert is familiar with all recent literature on the group, his or her determination is often more accurate and current than any flora.

Expert determination is often essential for certain groups in which species or infraspecific identification is very difficult.

PRACTICAL IDENTIFICATION

The practical steps taken in identification of plant taxa often depend on the experience of the person making the determination. Obviously, the more you know, the easier it is to identify a plant. For example, most floras begin with a key to the plant families, which may be cumbersome because they must take into account the variation within the total flora. Thus, knowing the general characteristics of several families ahead of time helps, as you may proceed directly to the key of genera within that family. Similarly, if you have an idea as to the general group within a family to which the taxon belongs (e.g., a suspected genus), you may wish to check the keys, illustrations, descriptions, or specimens within that group first. However, when in doubt, it is best to start from the beginning to be certain of eliminating the close but incorrect choices.

The importance of correctly identifying a specimen cannot be overstated. Once a determination is made, it should be viewed as only tentative. Never assume you have reached the correct answer in using any one method; it is important to check your determination by all available means. Be your own devil's advocate; check and recheck yourself. Verify every identification against a written description and comparison to an herbarium specimen. Some groups may be particularly difficult to identify, being composed of a great number of taxa that differ from one another by obscure features. Proceed very carefully, and don't hesitate to send off specimens for expert determination if needed.

Finally, one should always be conscious of the possibility that the identification process points to a new taxon. If a thorough evaluation of available references indicates that the unknown plant in question does not match any known taxa listed in a flora, then the plant may be a new record (either native or naturalized) for the geographic range of that flora. In some cases, the unidentifiable taxon may be new to science, warranting the valid publication of a new taxon.

REVIEW QUESTIONS

1. What is identification?
2. Describe how identification is used in your everyday life.
3. What two procedures are necessary before an identification can be made?
4. Define diagnostic characterization (= diagnosis).
5. What are the difficulties with identifying cultivated plant taxa?
6. Name the five major ways to identify plant taxa, citing the advantages and disadvantages of each.
7. What is a dichotomous key?
8. What is a couplet? a lead?
9. What precautions should be made in using a dichotomous key?

10. What is the difference between a *natural* and an *artificial* key?
11. What is a polyclave key?
12. What are the advantages and disadvantages of a polyclave key?
13. Name ways, other than a taxonomic key, to identify a plant specimen.
14. What is a synoptic collection and what is its advantage in plant identification?
15. State the practical steps made in identifying a plant specimen.

EXERCISES

1. Select an unknown cultivated plant and attempt to identify it using all of the five methods discussed in the chapter. What difficulties did you encounter with any of these?
2. Select an unknown native plant and attempt to identify it using local floras or manuals or by using an herbarium collection, such as a synoptic collection.
3. Do a Web search for a polyclave key (see Reference below), either one on-line or one that may be downloaded. Test this on a given unknown. If possible, create a key for a set of 5-10 plants.

REFERENCES FOR FURTHER STUDY

Radford, A. E., W. C. Dickison, J. R. Massey, and C. R. Bell. 1974. *Vascular Plant Systematics*. Harper and Row, New York. 891 pp.

WEB PAGES:

Dallwitz, M. J. (1996 onwards). Programs for interactive identification and information retrieval. <http://delta-intkey.com/www/idprogs.htm>

This Page Intentionally Left Blank

16

PLANT NOMENCLATURE

PRINCIPLES OF NOMENCLATURE	502	BOTANICAL NAMES	510
Scientific Names	502	Gender	510
Ranks	503	Number	510
Authorship	504	Commemoratives	510
Learning Scientific Names	504	PRONUNCIATION OF NAMES	511
Nomenclatural Types	504	Diphthongs	511
Priority of Publication	505	Syllables	511
Conservation of Names	505	Accenting	512
Name Changes	505	Commemoratives	513
Valid Publication	507	REVIEW QUESTIONS	513
Synonyms	507	EXERCISES	514
Abbreviations	509	REFERENCES FOR FURTHER STUDY	515
Independence of Botanical Nomenclature	510		
Retroactivity of the ICBN	510		

Nomenclature is the assignment of names utilizing a formal system. The criteria for formally naming land plants, algae, and fungi are based on the rules and recommendations of the **International Code of Botanical Nomenclature** or **ICBN**. (Note that a separate code is utilized for cultivated plants, the International Code of Nomenclature for Cultivated Plants.) Botanical names serve as symbols of a group of natural entities for the purpose of communication and data reference.

Of interest is the fact that the ICBN deals with the names of extant or extinct (fossil) organisms traditionally treated as plants, i.e., encompassed by the field of botany (see Chapter 1 for a definition of *botany*). These include not only the land plants, but also the blue-green algae (Cyanobacteria); fungi, including chytrids, oomycetes, and slime moulds; photosynthetic protists and taxonomically related non-photosynthetic groups. As discussed in Chapter 1, it is now known that many of these groups are not closely related phylogenetically. Yet, the ICBN still deals with these taxa.

Separate nomenclatural codes exist for traditional zoology (International Code of Zoological Nomenclature) and for prokaryotes (International Code of Nomenclature of Bacteria). One difficulty with this is that photosynthetic bacteria are named both under the ICBN and under the Bacteria Code.

Similarly, some of the so-called protists (itself a paraphyletic assemblage) are named both under the ICBN and the Zoological Code. Thus, some organisms have two names, from two different nomenclatural codes. A future universal code, covering all forms of life, has been discussed recently and may be of advantage.

The International Code of Botanical Nomenclature governs the rules both for the specific names assigned to taxa and for the name endings that denote taxon rank (see below). The ICBN is utilized in two basic activities: (1) naming new taxa, which were previously unnamed and often not described; and (2) determining the correct name for previously named taxa, which may have been divided, united, transferred, or changed in rank (see later discussion).

Legitimate names are those that are in accordance with the rules of the International Code of Botanical Nomenclature. Any name that violates one or more rules of the ICBN is known as an **illegitimate name**. A **valid name** is one that is **validly published** (see later discussion). The rules of the ICBN can be somewhat complex, often necessitating careful scrutiny (and a lawyerlike mentality).

Points of controversy and periodic changes to the International Code of Botanical Nomenclature are voted upon

during meetings of the International Botanical Congress, which assembles about every 6 years in some city around the world. As of this writing, the last International Botanical Congress was held in St. Louis, Missouri, in 1999. The following summary is based on the ICBN resulting from that Congress (Greuter et al. 2000).

PRINCIPLES OF NOMENCLATURE

The **Principles** of the International Code of Botanical Nomenclature are stated verbatim below from the 2000 St. Louis Code. Each of these will be covered in detail.

- I. Botanical nomenclature is independent of zoological and bacteriological nomenclature. The Code applies equally to names of taxonomic groups treated as plants whether or not these groups were originally so treated.
- II. The application of names of taxonomic groups is determined by means of nomenclatural types.
- III. The nomenclature of a taxonomic group is based upon priority of publication.
- IV. Each taxonomic group with a particular circumscription, position, and rank can bear only one correct name, the earliest that is in accordance with the Rules, except in specified cases.
- V. Scientific names of taxonomic groups are treated as Latin regardless of their derivation.
- VI. The Rules of nomenclature are retroactive unless expressly limited.

The details of the International Code of Botanical Nomenclature are organized into a number of **Rules** (which are mandatory and written out as Articles), **Notes** (which are binding and clarifying), **Recommendations** (which are not binding but suggested), and explanatory **Examples** and **Footnotes**.

Currently, the entire International Code of Botanical Nomenclature is available on a Web site (Greuter et al. 2000).

SCIENTIFIC NAMES

The *fundamental* principle of nomenclature is the fourth principle of the ICBN, stating that every taxon, whether species, genus, family, etc., can bear only one correct name (see below for precise definition of *correct* name). This is only common sense. Confusion would reign if taxonomic entities could bear more than one name or if one name could refer to more than one entity. The names assigned by the rules of the ICBN are known as **scientific names**. Scientific names are, by convention, in the Latin language (see later discussion).

As reviewed in Chapter 1, the scientific names of species are **binomials**, i.e., composed of two names. The binomial convention was first consistently used by Carolus Linnaeus (also known as Carl Linn \ddot{O} or Carl von Linn \ddot{O}), a Swedish botanist, who is often referred to as the father of taxonomy. Prior to the use of binomials, the designation of species was inconsistent and may have utilized several words.

As an example of a binomial, the species commonly known as sweetgum has the scientific name *Liquidambar styraciflua*. The first name of the binomial, *Liquidambar* in this case, is the **genus name** and is always capitalized. The genus name may be abbreviated by its first letter, but only after it is spelled out in its entirety (and would not be confused with another genus name); thus, the above may be abbreviated as *L. styraciflua*. The second name of the binomial, *styraciflua* in this example, is the **specific epithet**. The specific epithet may be capitalized if it is a commemorative (named after a person or place), but this is optional; the trend today is to never capitalize the specific epithet. Binomial species names are always either italicized or underlined. Recall from Chapter 1 that a species name is always the entire binomial. It is incorrect to say that the species name for sweetgum is *styraciflua*, as this is the specific epithet; the species name is *Liquidambar styraciflua*.

In contrast to scientific names, many taxa also bear **common names** (also called vernacular names), which are generally used by people within a limited geographic region. Common names are not formally published and are governed by no rules. Scientific names are much preferable to common names for several reasons. First, only scientific names are universal, used the same world-wide; common names may vary from region to region, even within a country or within regions of a country. For example, species of the genus *Ipomoea* are known commonly as morning glory in the United States, but as woadbine in England. Differences in language will, of course, further increase the number of different common names. Second, common names are not consistent. One taxon may bear more than one common name, these often varying in different regions. For example, *Adenostoma fasciculatum* of the Rosaceae is known by at least two common names, chamise and grease wood. Alternatively, a single common name may refer to more than one taxon. Hemlock may refer to two quite different plants, either a species of *Tsuga*, a coniferous tree of the Pinaceae, or *Conium maculatum*, an herb of the Apiaceae (the extract of which Socrates drank in execution). Third, common names tell nothing about rank and often nothing about classification, whereas scientific names automatically indicate rank and yield at least some information about their classification. For example, sea-blite tells nothing about rank; it could be

variety, species, genus, or family. However, one immediately knows that *Suaeda californica* is at the rank of species and is a close relative to other species of *Suaeda*. Fourth, many, if not most, organisms have no common name in any language; thus, scientific names alone must be used to refer to them. This is especially true for plants that are nonshowy, occur in remote areas, or belong to groups whose members are difficult to distinguish from one another.

There is a tendency in some works to arbitrarily convert all scientific species names into common names by translating from the Latin, even when these common names are *not* used by the native people. For example, *Carex aurea* might be designated golden carex or golden sedge, even if these names are not in common usage. It is the author's opinion that this is less than ideal policy and that it is preferable simply to utilize scientific names and refer to common names only if they are, in fact, commonly used.

RANKS

Recall from Chapter 1 that taxa are classified hierarchically by **rank**, in which a higher rank is inclusive of all lower ranks (Figure 16.1). [Note that there are principal ranks, secondary ranks, and additional ranks (if needed) that may

be used by adding the prefix *sub*; see Figure 16.1.] Each scientific name of a particular rank must end in a certain suffix according to the rules and recommendations of the ICBN (Figure 16.1). For example, Asteridae is a taxon at the rank of subclass, Asterales is at the rank of order, and Asteraceae is at the rank of family, etc. Note that taxa above the rank of genus are *not* underlined or italicized.

One exception to these rank endings of taxa is the acceptance of eight *alternative* family names, none of which end in *-aceae*. These are Compositae (= Asteraceae), Cruciferae (= Brassicaceae), Gramineae (= Poaceae), Guttiferae (= Clusiaceae/Hypericaceae), Labiatae (= Lamiaceae), Leguminosae (= Fabaceae), Palmae (= Arecaceae), and Umbelliferae (= Apiaceae). In addition, within the Fabaceae (= Leguminosae), the subfamily name Papilionoideae is an acceptable alternative to the Faboideae. The trend today is to consistently apply the type principle (see later discussion) by using the standardized family names that end in *-aceae* and to use subfamily names that are based on these (e.g., to use Faboideae over Papilionoideae). However, plant taxonomists should know these alternative names, as they are often used in older, as well as some current, floras and other taxonomic works.

TAXONOMIC RANKS OF LAND PLANTS

	ENDING	EXAMPLE TAXON
Kingdom	(various)	Plantae
Phylum [Division]	-phyta	Magnoliophyta
Subphylum [Subdivision]	-phytina	Magnoliophytina
Class	-opsida	Asteropsida
Subclass	-idae	Asteridae
Order	-ales	Asterales
Suborder	-ineae	Asterineae
Family	-aceae	Asteraceae
Subfamily	-oideae	Asteroideae
<u>Tribe</u>	-eae	Heliantheae
Subtribe	-inae	Helianthinae
Genus	(various)	<i>Helianthus</i>
Subgenus	(various)	<i>Helianthus</i>
<u>Section</u>	(various)	<i>Helianthus</i>
<u>Series</u>	(various)	<i>Helianthus</i>
Species [abbr. sp. (sing.), spp. (pl.)]	(various)	<i>Helianthus annuus</i>
Subspecies [abbr. subsp. or ssp. (sing.), subsp. or sspp. (pl.)]	(various)	<i>Helianthus annuus</i> ssp. <i>annuus</i>
<u>Variety</u> [abbr. var. (sing.), vars. (pl.)]	(various)	<i>Helianthus annuus</i> var. <i>annuus</i>
<u>Form</u> [abbr. f.]	(various)	<i>Helianthus annuus</i> f. <i>annuus</i>

FIGURE 16.1 Taxonomic ranks recognized by the International Code of Botanical Nomenclature. Principal ranks are in bold. Secondary ranks are underlined. Sub ranks may be used as needed, some of the possibilities indicated. Phylum, subphylum, class, and subclass may utilize different endings for Fungi or Algae. Ranks at and below the level of order have endings specified in the ICBN rules; those above the rank of order have endings recommended by the ICBN. Division may be used interchangeably with Phylum, but this is not recommended in order to promote consistency of use with Animals and Prokaryotes.

Position is the placement of a taxon as a member of another taxon of the next higher rank. For example, the position of the genus *Aster* is as a member of the family Asteraceae. Taxa may be the same in rank but differ in position. *Rosa* and *Aster* are both at the rank of genus but differ in position, the former in the Rosaceae, the latter in the Asteraceae.

As mentioned earlier, the prefix *sub-* can be used formally in a rank name in more categories are needed, such as *subgenus* or *subspecies*. In a less formal sense, the prefix *sub-* or *infra-* can be used to denote taxa below one of the major ranks. For example, *subfamilial* or *infrafamilial* taxa are those *below* family, including subfamily, tribe, subtribe, genus, subgenus, section, species, subspecies, variety, etc.

A subspecies or variety name is a **trinomial** (three names), e.g., *Toxicodendron radicans* ssp. *diversilobum* or *Brickellia arguta* var. *odontolepis*. In these examples, the **subspecific epithet** is *diversilobum*; the **varietal epithet** is *odontolepis*. Note that, technically, the rank of subspecies is above that of variety (Figure 16.1). In practice, subspecies and variety are sometimes used interchangeably. However, it is possible, but very rare, to have a subspecies that is divided into varieties itself (these constituting quadriminomial!).

AUTHORSHIP

All scientific names at and below the rank of family have an **author**, the name of the person who first validly published the name (see later discussion). For example, the full name (including authorship) of the family Rosaceae is Rosaceae Jussieu because de Jussieu first formally named the family. In other examples, the full name of the tribe Conostylideae is Conostylideae Lindley; that of the genus *Mohavea* is *Mohavea* A. Gray; that of the species *Mohavea confertiflora* is *Mohavea confertiflora* (Bentham) Heller; and that of the subspecies *Monardella linoides* ssp. *viminea* is *Monardella linoides* A. Gray ssp. *viminea* (Greene) Abrams. Author names are often abbreviated, such as Haemodoraceae R. Br. (for Robert Brown) or *Liquidambar styraciflua* L. (L. being the standardized abbreviation for Linnaeus). (See Name Changes for an explanation of author names appearing in parentheses.)

Although authorship is part of a scientific name and should be cited in all scientific publications, in practice the author is not typically memorized or recited as part of a scientific name. The authors of higher taxa are sometimes omitted in print even in scientific publications, except in detailed monographic treatments in which the nomenclatural history of the taxa under study is described. In many floras and journal publications, only species and infraspecific taxa (subspecies or varieties) are listed with full authorship, and these generally only once, when the name is first cited.

LEARNING SCIENTIFIC NAMES

As argued earlier, it is important to learn the scientific names of plants, correctly spelled. The serious plant taxonomist will learn many hundreds of scientific names in his/her lifetime, still just a tiny fraction of the more than 250,000 described plant species. Beginners may at first have difficulty learning scientific names. Some suggestions for mastering them are as follows.

First, learn to divide into syllables and accent scientific names (see **BOTANICAL NAMES**). It is often easier to recite and spell a scientific name if it is consciously broken down into syllables, each of which is separately pronounced.

Second, use mnemonic devices. Select one distinctive feature about the plant. Then find a common word that *sounds* somewhat similar to the scientific name. Link the distinctive plant feature with the similar sounding word in an active, vivid mental image, the weirder and more active the better. Thus, when you see the plant, you associate it with the mental image, which sounds like (and reminds you of) the scientific name. For example, visualizing liquid amber flowing from the distinctive, ball-shaped fruits of sweetgum may help you remember the genus name, *Liquidambar*.

Third, learn the etymology (meaning) of scientific names. Scientific names often are descriptive about the morphology of the plant. Once you know, for example, that the Latin word *alba* means white or that *leptophylla* means narrow-leaved, you can better associate the name with the organism. Other scientific names may be named after a person or place of significance; learning the history of these commemorative names may be helpful in memorizing them.

Finally, there is no substitute for continual practice and review. Use a combination of both oral and (for correct spelling) written recitation, with the plant, plant specimen, photograph, or mental image in view.

NOMENCLATORIAL TYPES

The second principle of the ICBN states that scientific names must be associated with some physical entity, known as a **nomenclatural type** or simply **type**. A nomenclatural type is almost always a specimen, e.g., a standard herbarium sheet specimen, but it may also be an illustration. The type serves the purpose of acting as a reference for the name, upon which the name is based. If there is ever any doubt as to whether a name is correct or not, the type may be studied.

There are different types of types. **Aholotype** is the one specimen or illustration upon which a name is based, originally used or designated at the time of publication. It serves as the definitive reference source for any questions of identity or nomenclature. It is recommended that a holotype be deposited in an internationally recognized herbarium

(see Chapter 18) and cited as one of the criteria for the valid publication of a name (see later discussion). Holotypes constitute the most valuable of specimens and are kept under safe keeping in one (usually a major) herbarium. An **isotype** is a duplicate specimen of the holotype, collected at the same time by the same person from the same population. The ICBN recommends that isotypes be designated in the valid publication of a new name. Isotypes are valuable in that they are reliable duplicates of the same taxon and may be distributed to numerous other herbaria to make it easier for taxonomists of various regions to obtain a specimen of the new taxon. A **lectotype** is a specimen that is selected from the original material to serve as the type when no holotype was designated at the time of publication, if the holotype is missing, or if the original type consisted of more than one specimen or taxon. A **neotype** is a specimen derived from a nonoriginal collection that is selected to serve as the type as long as all of the material on which the name was originally based is missing. [Other types of types include (1) **syntype**, which is any specimen that was cited in the original work when a holotype was not designated; alternatively, a syntype can be one of two or more specimens that were all designated as types; (2) **paratype**, a specimen cited but that is not a holotype, isotype, or syntype; and (3) **epitype**, a specimen (or illustration) that is selected to serve as the type if the holotype, lectotype, or neotype is ambiguous with respect to the identification and diagnosis of the taxon.]

Normally, we think of types as referring to a species or infraspecific taxon. However, type specimens may serve as references for higher taxonomic ranks as well. For example, the type specimen for a genus name is the same as the one for the species within the genus that was published *first* (see **Priority of Publication**). The type specimen for a family name is the same as the one for the genus within the family that was published *first*.

PRIORITY OF PUBLICATION

The third principle of the ICBN is priority of publication, which generally states that of two or more competing possibilities for a name, the one published *first* is the correct one, with some exceptions. Priority of publication only applies to taxa at the rank of family and below and does not apply outside a particular rank (with a transfer in rank; see later discussion). For example, of two competing names (both legitimate and validly published) *Mimulus* (published in 1753) and *Diplacus* (published in 1838) the genus *Mimulus* has priority and is the correct name when the two genera are combined into one. The principle of priority for vascular plants starts 1 May 1753 with the publication of *Species Plantarum* by Linnaeus; names published prior to that are not considered

for priority. (Different groups covered by the ICBN have various starting dates.)

CONSERVATION OF NAMES

One adverse effect of the principle of priority is that scientific names that are well known and frequently used may be replaced by some other name if the latter was discovered to have been published earlier. This lends a degree of instability to nomenclature. However, in such a case, a petition may be presented (in the botanical journal *Taxon*) and voted upon at the International Botanical Congress to conserve one name over another that actually has priority. Such a procedure is outlined as three Amendments to the ICBN: *Nomina familiarum conservanda*, *Nomina generica conservanda et rejicienda*, and *Nomina species conservanda*. The rationale for the conservation of names is to provide greater stability in nomenclature by permitting names that are well known and widely used to persist, even upon the discovery of an earlier, but more obscure, name.

NAME CHANGES

Occasionally, the name of a taxon will change. Name changes can occur for only two reasons: (1) because of the recognition that one name is contrary to the rules (i.e., is illegitimate), and, thus, another name must take its place; or (2) because additional taxonomic study or research (for example, a cladistic analysis) has resulted in a change of the definition and delimitation of a taxon; this process is known as a **taxonomic revision**.

There are four basic types of nomenclatural activities that can result in a name change. First, a single taxon may be **divided** into two or more taxa, often called *segregate taxa* because they are segregated from one another relative to the original classification. This is done generally via the recognition of features that clearly distinguish two or more groups from one another. For example, the genus *Langloisia* has been split into two genera, *Langloisia* and *Loeseliastrum*, based on a number of morphological, anatomical, and palynological (pollen) features that distinguish them. Ideally, the segregate groups should be monophyletic, as based upon a rigorous cladistic analysis (see Chapter 2). Other examples of taxa being divided are:

1. The genus *Carduus* of the family Asteraceae is often split into two genera: *Carduus*, having barbellate pappus bristles, and *Cirsium*, having plumose pappus bristles
2. The genus *Rhus* of the Anacardiaceae has been split into several segregate genera, such as *Malosma*, *Rhus*, and *Toxicodendron*, the last including poison-oak and poison-ivy
3. The classical family Liliaceae has been split into numerous families, such as the Alliaceae, Hyacinthaceae, Liliaceae s.s., and Melanthiaceae

4. The large genus *Haplopappus* of the Asteraceae has been split into several genera, including *Anisocoma*, *Ericameria*, *Hazardia*, *Haplopappus*, and *Isocoma*

Note that when a larger taxon is divided into two or more smaller taxa of the same rank, the terms **sensu lato** (abbreviated **s.l.**) and **sensu stricto** (abbreviated **s.str.** or **s.s.**) may be used to distinguish the more inclusive and less inclusive treatments, respectively. For example, *Haplopappus* s.l. contains many more species than *Haplopappus* s.s., the latter of which is what remains after *Haplopappus* s.l. is split into many segregate genera.

A second, major name change occurs when two or more separate taxa are **united** into one. One reason for uniting taxa is the recognition that features previously used to distinguish them are, upon more detailed study, unresponsive of their being different; i.e., there is no clear character state discontinuity. Another reason to unite taxa may be based on cladistic studies, in which of two or more separate taxa, one (or more) is demonstrated to be paraphyletic; thus, one way to eliminate a paraphyletic taxon is to unite it with other taxa such that the new inclusive group is now monophyletic (see Chapter 1). In cases of taxa being united, the final name used is that which was *published earliest*, according to the principle of priority. Examples of taxa being united are:

1. The species *Bebbia juncea* and *Bebbia aspera*, which were considered indistinct and were united into one species, *B. juncea*
2. The genera *Diplacus* and *Mimulus*, which were united into one genus, *Mimulus*
3. The families Apocynaceae and Asclepiadaceae, which have been united into one family, the Apocynaceae (which could be designated Apocynaceae s.l. to distinguish it from the earlier circumscribed less inclusive family)

Third, a taxon may be **transferred in position**, i.e., from one taxon to another of the *same rank*. Examples of this are:

1. The species *Rhus laurina* was transferred in position as a member of the genus *Malosma*, the new species name being *Malosma laurina*
2. The species *Sedum variegata* was transferred to the genus *Dudleya*, the new species name being *Dudleya variegata*

Note that a transfer in position may be an automatic result of uniting or dividing taxa of higher rank. For example, if the genera *Diplacus* and *Mimulus* are united into the genus *Mimulus*, then the *species* of *Diplacus* must be transferred in position.

Fourth, a taxon may be **changed in rank**. Examples include:

1. The species *Eruca sativa* was changed to the rank of subspecies (of the species *E. vesicaria*), the new combination being *Eruca vesicaria* ssp. *sativa*
2. The variety *Viguiera deltoidea* var. *parishii* was changed to the rank of species, the new name being *Viguiera parishii* (with *V. deltoidea* persisting as a separate species equivalent in circumscription to the earlier *V. deltoidea* var. *deltoidea*)

Note in the two rank change examples just given that the original names for the epithets are retained. The retention of a name that is changed in rank is recommended (but not required) by the ICBN, but only if an earlier name for the same taxon had not already been published at that rank (and also, only if the *same* name had not already been used for another taxon; see **homonym**). The principle of priority does not apply outside the rank of a taxon, however; this means that if a name is changed in rank, the date of publication of the original name (before being changed in rank) cannot be considered in evaluating priority of publication with respect to the change.

In some cases a taxonomic study results in the **remodeling** of a taxon, i.e., a change in diagnostic characteristics, those that distinguish the taxon from other taxa. In these cases, a name change is not warranted and the rules of the ICBN need not apply.

A **basionym** is the name-bringing or epithet-bringing synonym, i.e., the original (but now rejected) name, part of which has been used in a new combination. As seen earlier, if a species or infraspecific name is transferred in position or rank, the specific or infraspecific epithet of the (now-rejected) basionym may be retained (unless violating another rule of the code, such as priority of publication, e.g., if the taxon had already been named, or if the name had already been used for another taxon at that rank). The name of the author(s) who originally named the basionym is also retained and placed in parentheses ahead of the author who made the change. Thus, botanical names may have two sets of authors: the author(s) set in parentheses who originally named the basionym, and the author(s) who made the name change. From some of the examples cited previously:

1. When *Sedum variegata* Wats. was transferred to the genus *Dudleya* by Moran, the new species name became *Dudleya variegata* (Wats.) Moran. The original epithet, *variegata*, is retained, and the author associated with that epithet, Watson in this case, is also retained, but is

placed in parentheses preceding the new author. The basionym in this case is *Sedum variegata* Wats., the original name.

2. When *Dilatris caroliniana* Lam. was transferred to the genus *Lachnanthes* by Dandy, the new species name became *Lachnanthes caroliniana* (Lam.) Dandy. The basionym in this case is *Dilatris caroliniana* Lam.
3. When *Fumaria bulbosa* L. var. *solida* L. was elevated to the rank of species by Miller, the new name became *Fumaria solida* (L.) Miller. The basionym in this case is *Fumaria bulbosa* L. var. *solida* L. Subsequent to this change, *Fumaria solida* (L.) Miller was transferred in position by Clairv to the genus *Corydalis*, the new name becoming *C. solida* (L.) Clairv [not *C. solida* (Mill.) Clairv]. Note that it is the author of the varietal name of the basionym, *Fumaria bulbosa* L. var. *solida* L., that is retained in parentheses.

An **autonym** is an automatically created name for infrafamilial, infrageneric, and infraspecific taxa. Autonyms are used whenever a family is divided into subfamilies, tribes, or subtribes; a genus is divided into subgenera or sections; or a species is divided into subspecies or varieties. Of the two or more subtaxa formed, the autonym is assigned based on priority, i.e., to the group containing the taxon that was published first. Autonyms have no authors; only the higher taxa upon which they are based and the other subtaxa have formal authorship. For example, Isely split *Lotus stipularis* (Benth.) E. Greene into two varieties: *L. stipularis* (Benth.) E. Greene var. *ottleyi* Isely and *L. stipularis* (Benth.) E. Greene var. *stipularis*; note that the latter variety, containing the autonym, lacks authorship because its type is the same as that for the originally described species.

For infrafamilial taxa, the autonym has the same root name as the family but a different ending that corresponds to the infrafamilial rank. For example, the family Euphorbiaceae is usually divided into subfamilies, one of which, the Euphorbioideae, is the autonym; this subfamily, of course, contains the genus *Euphorbia*, the type for the family. For infrageneric taxa, the autonym is identical to the genus name and should be preceded by the name of the rank to avoid confusion. For example, *Ceanothus* (a genus) consists of two subgenera, subgenus *Ceanothus* and subgenus *Cerastes*; subgenus *Ceanothus* is the one that includes the type for the genus itself. For infraspecific taxa, autonyms are identical to the specific epithet. For example, *Eriogonum fasciculatum* is divided into several varieties, one of which, *Eriogonum fasciculatum* var. *fasciculatum*, includes the autonym (and is based on the original type specimen for the species).

VALID PUBLICATION

According to the ICBN, in order for a scientific name to be formally recognized, it must be **validly published**. There are four general criteria for valid publication of a name. First, the name must be effectively published, which means that it must be published in a journal commonly available to botanists (not, say, in the local newspaper or *National Enquirer* magazine). Second, the name must be published in the correct form, i.e., properly Latinized (see later discussion), with the rank indicated (e.g., as sp. no v. or gen. no v.; see **Abbreviations**). Such a legitimate name in correct form is known as an *admissible name*. Third, the name must be published with a Latin description or diagnosis or with a reference to such. The Latin description may be brief, e.g., listing how the new taxon is different from a similar, related taxon. (In addition, a more detailed description in some vernacular language, or a reference to a previous description, is usually included but not required.) Fourth, for taxa of the rank of genus and below, a nomenclatural type must be indicated; the location of this type is also indicated (using the acronyms of Index Herbariorum; see Holmgren et al. 1990; Chapter 18). An example of a valid publication, illustrating these criteria, is seen in Figure 16.2.

The full citation of a scientific name includes not only the authorship, but also the place and date of publication. For example, the full citation for the species cited in Figure 16.2 is *Perityle vigilans* Spellb. & A. Powell, Syst. Bot. 15: 252. 1990. Full citations are listed in the citations of the International Plant Names Index (see References for Further Study).

SYNONYMS

A **synonym** is a rejected name, *by a particular author or authors*. Synonyms are rejected for either of two reasons: (1) because they are illegitimate, i.e., contrary to the rules of the ICBN; or (2) because of taxonomic judgment, i.e., a particular author rejects the classification represented by the synonym. Synonyms may be based on the same or on a different type specimen from the correct name.

A **correct name** is a legitimate (and therefore validly published) name that is accepted *by a particular author or authors*. Recall that the fundamental principle of the ICBN states that each taxon can have only *one* correct name. Thus, if there are two or more competing names for the same taxon, e.g., *Malosma laurina* (Nutt.) Abrams and *Rhus laurina* Nutt., only one of them can be correct. However, *which* name is correct may depend on the author(s) of a given reference book or journal. For example, according to one author, *Rhus laurina* Nutt. is the correct name and *Malosma laurina* (Nutt.) Abrams

Effective publication, in a journal commonly available to botanists

Systematic Botany 1990, 15(2): pp. 252-255
 Copyright 1990 by the American Society of Plant Taxonomists

A New Species of *Perityle* (Asteraceae) from Southwestern Chihuahua, Mexico

RICHARD SPELLENBERG

Department of Biology, New Mexico State University,
 Las Cruces, New Mexico 88003

A. MICHAEL POWELL

Department of Biology, Sul Ross State University,
 Alpine Texas 79832

ABSTRACT. *Perityle vigilans* is described from the Sierra Madre Occidental of southwestern Chihuahua. It differs from other *Perityle* by the combination of its white ligules, sparsely setose-hispid achenes, finely grayish villosulous tomentum, and absence of pappus bristles. The new species is believed to be most closely related to *P. rosei* and *P. trichodonta*, which occur about 800 km to the south. A key is presented that distinguishes the 12 known taxa of *Perityle* occurring in the northern Sierra Madre Occidental, an area of high diversity in the genus.

In a series of three papers Powell (1969, 1973, 1974) revised the genus *Perityle*, recognizing 53 species. Except for one amphitropical disjunct, *P. emoryi* Torrey, the genus is restricted to southwestern North America. The treatment by Niles (1970) is in concurrence with regard to the circumscription of species. In his series of papers Powell proposed that speciation in *Perityle* occurs primarily by geographic isolation, that the populations of species, many of which inhabit nearly barren rock cliffs, were derived from more widespread ancestral species whose ranges were divided by geologic uplift, igneous intrusion, and subsequent erosion. Powell explained that this isolation on island-like habitats of exposed rock has resulted in a high degree of endemism, and he noted (1974) that the diversity in the genus is highest in the Sierra Madre Occidental of northwestern Mexico. He postulated that this may be the center of origin for the genus. Since that revision, seven narrowly endemic species have been added to the genus as remote areas have been explored: *P. ajoensis* Todsén (1974), *P. batopilensis* Powell (1983), *P. carmenensis* Powell (1976), *P. fosteri* Powell (1983), *P. huecoensis* Powell (1983), *P. specuicola* Welsh and Neese (1983), and *P. vandevenderi* B. Turner (1989). A few varieties also have been described. This article adds an eighth species, this from the isolated mid-elevations of the west slope of the Sierra Madre Occidental.

to 3 mi. NE along road, just below oak zone and in lower edge of zone, elev. 1700 m, 27° 52' 30" N, 107° 59' 30" W, 25 Apr 1985, Spellenberg, Soreng, Corral, Todsén 8104 (holotype: NMC; isotypes: ENCB, NY, MEXU, SRSC, TEX, UC, Escuela Superior Agricultura Hermanos Escobar [Cd. Juárez]).

Type specimen indicated

Plantae perennes suffruticosa subpulvinatae, caulis 2.5-8.5 cm longis. Indumentum densum, griseum, minute villosulum. Folia opposita, periolata, laminae ovatae rhombeae-ovatae, 2.5-5.5 mm longae, 1.7-4 mm latae, dentibus 0-3 brevibus obtusis in uterque marginibus. Capitula terminalia solitaria radiata; corollae radii 6-8, ligulis 2.5-3.0 mm longis, 1.5-2.5 mm latis, in tubis extus et lamina subtus glandulis aureis sparsis obsitis; corollae disci ca. 35-40, ca. 2.0 mm longae. Achaenia anguste obdeltata nigra 2.2-2.5 mm longa modice pilosa-hispida. Pappus obsolete constans ex corona dentibus tenuibus triangularibus minoribus quam 0.1 mm longis. Fig. 1.

Latin description

Plants suffruticose, more or less cushion-like, the stem tips ascending. Stems 2.5-8.5 cm long, densely and finely grayish-villosulous, the fine kinked hairs over-topping yellowish spherical glands. Leaves opposite, pubescent as the stem, the upper leaves slightly more densely so than the lower; petioles slender, expanding into the cuneate leaf base, 2-3 mm long; blades ovate to rhombic-ovate, broadly cuneate at the base, 2.5-5.5 mm long, 1.7-4.0 mm wide, with 0-3 low blunt teeth on each margin, when only 1 tooth the blade then subhastate. Capitulescence of a

Vernacular description

Name in Latin — ***Perityle vigilans*** Spellenberg & A. Powell, *sp. nov.* (Fig. 1). — TYPE: Mexico, Chihuahua, Municipio Maguarichi, on igneous rocks at Maguarichi and

Rank indicated

FIGURE 16.2 Example of a new species description, illustrating the components of validly publication. Note the abbreviation *sp. nov.* after the scientific name, meaning *species nova*, Latin for new species. Rewritten from *Systematic Botany*, by permission.

is the synonym. According to other authors, *Malosma laurina* (Nutt.) Abrams is the correct name and *Rhus laurina* Nutt. is the synonym.

Synonyms are typically indicated in brackets following the correct name, such as *Malosma laurina* (Nutt.) Abrams [*Rhus laurina* Nutt.] or *Machaeranthera juncea* (Greene) Hartman [*Haplopappus juncea* Greene]. Alternatively, if a synonym (according to one author) is cited or referenced, the correct name is often indicated preceded by an = sign. For example, in *Cyrtanthera* Nees = *Justicia*, the correct name is *Justicia* and the synonym is *Cyrtanthera*.

A **homonym** is one of two (or more) *identical* names (not including authorship) that are based on different type specimens. The later homonym, based on publication date, is illegitimate (unless it is conserved; see earlier discussion). For example, *Tapeinanthus* Herb. (1837), of the Amaryllidaceae, and *Tapeinanthus* Boiss. ex Benth. (1848), of the Lamiaceae, are homonyms. The later homonym in the Lamiaceae is illegitimate [and was renamed *Thuspeinanta* T. Durand (1888)].

A **tautonym** is a binomial in which the genus name and specific epithet are identical in spelling. Tautonyms are not permitted in botanical nomenclature. For example, the name *Helianthus helianthus* is a tautonym and illegitimate, whereas *Helianthus helianthoides* is not a tautonym and would be permitted. (Note that zoological nomenclature does permit tautonyms, as in *Gorilla gorilla*.)

ABBREVIATIONS

Certain abbreviations are used in scientific names. For example, the word **ex** means validly published by. For example, *Microseris elegans* Greene ex A. Gray means that Asa Gray validly published the name *Microseris elegans* that was originally proposed (but not validly published by) Greene. The **ex** plus the author(s) preceding it may be omitted, as in *Microseris elegans* A. Gray.

The word **in** means in the publication of, referring to a name published within a larger work authored by the person(s) following the **in**. For example, *Arabis sparsiflora* Nutt. in T. & G. means that Nuttall validly published the name *Arabis sparsiflora* in another work authored by Torrey & Gray. The **in** plus the author(s) following it may be omitted for brevity, as in *Arabis sparsiflora* Nutt. (The use of **in** is not recommended by the ICBN.)

An **×** indicates a hybrid. For example, *Salvia* × *palmeri* (A. Gray) E. Greene is a named (validly published) taxon representing a hybrid between two species: *S. apiana* Jepson and *S. clevelandii* (A. Gray) E. Greene. Alternatively, this hybrid could be represented as *S. apiana* Jepson × *S. clevelandii* (A. Gray) E. Greene. Hybrids may also be indicated by placing the prefix *notho-* prior to the rank name, as in *Polypodium*

vulgare nothosubsp. *mantoniae* (Rothm.) Schidlay (indicating that the named subspecies is of hybrid origin).

The abbreviation **sp. nov.** following a binomial (e.g., *Eryngium pendletonensis*, sp. nov.) refers to the Latin *species nova* and means that the species is new to science. Similarly, **gen. nov.** (*genus novum*) cites a new genus name. The abbreviation **comb. nov.** following a binomial refers to the Latin *combinatio nova* and means that the taxon has recently been transferred to a new position or rank. An example is *Porella acutifolia* (Lehm. & Lindenb.) Trevis. var. *ligulifolia* (Steph.) M. L. So, comb. nov.

Two abbreviations **af** and **cf.** are used to describe plant specimens whose identity is uncertain. The distinction between the two abbreviations is unclear, as different taxonomists have used them with slightly different meanings. The abbreviation **aff.** preceding a taxon name literally means related to (Latin *affinis*, related, connected), as in *Calyptridium* aff. *monandrum* or **af** *Calyptridium monandrum*. This abbreviation implies some type of close relationship, presumably an evolutionary relationship, but also that the specimen differs from the cited taxon in some way, e.g., beyond the described range of variation for one or more characters; the cited specimen might, in fact, be indicative of a new taxon. The abbreviation **cf.** (Latin *confer*, compare) preceding a taxon name, as in *Calyptridium* cf. *monandrum* or **cf.** *Calyptridium monandrum*, indicates that the identity of a specimen is more questionable or uncertain (perhaps because references or comparative specimens are not available), and should be compared with specimens of the taxon indicated (i.e., the name following **cf.**) for more detailed study.

As mentioned earlier, **s.l.** (*sensu lato*) means in the broad sense, referring to a broad, inclusive taxon circumscription, and **s.str.** or **s.s.** (*sensu stricto*) means in the strict sense, referring to a narrow, exclusive taxon circumscription.

Other, minor abbreviations or specialized words include:

1. **auct. non** (*auctorum non*) refers to a misapplication of a name, such that the type specimen of the name does not fall within the circumscription of the taxon being referred to by that name
2. **emend.** (*emendatio*) means a correction or amendment
3. **et** is Latin for and
4. **nom. cons.** (*nomen conservandum*) means a conserved name
5. **nom. nov.** (*nomen novum*) means a new name
6. **nom. nud.** (*nomen nudum*) means published without a description or diagnosis, making the name invalid
7. **non** is Latin for not
8. **orth. cons.** (*orthographia conservanda*) means a conserved spelling

- 9 **stat. nov.** (*status novus*) means a change in rank, e.g., elevating a varietal name to specific status
- 10 **typ. cons.** (*typus conservandus*) means a conserved type specimen
- 11 **typ. des.** (*typus designatus*) means the designation of a type specimen
- 12 **vide** (*video*) means to cite a reference
- 13 **!** (symbol for *vidi*, I have seen it) means a confirmation of a name

INDEPENDENCE OF BOTANICAL NOMENCLATURE

The International Code of Botanical Nomenclature is independent of the International Code of Zoological Nomenclature. Thus, there may be some names of plants, algae, or fungi that are identical to those of some animals (and Protista). For example, the genus *Morus* refers both to a flowering plant, the mulberry, and to a bird, the gannet; *Ficus* is the genus name of the figs and of a group of gastropods. A separate code is also used for the Prokaryotes (including the bacteria) and viruses.

RETROACTIVITY OF THE ICBN

The Rules of the International Code of Botanical Nomenclature are retroactive, except in specified cases.

BOTANICAL NAMES

Botanical Latin is best described as a modern Romance language of special technical application (Stern 1992). The fourth principle of the ICBN is that botanical names are treated as Latin, a language chosen because of its classical history (in the past being the language of scholars) and because it is no longer in active use. No matter what the language of the person who published a name, the name itself must consist of direct Latin words or be Latinized, i.e., converted from the vernacular to the Latin. Thus, the Latin alphabet (which is almost identical to the English alphabet) and grammatical rules must be used.

GENDER

All Latin words have a gender: masculine, feminine, or neuter. Gender determination is usually only of concern for names at the rank of genus or below. The standardized gender endings are:

Masculine	Feminine	Neuter
-us	-a	-um
-er	-ra	-rum
-is	-is	-e
-r	-ris	-re

The first row of endings (-us, -a, and -um) are those most commonly used. For example, the gender of the genus *Amaranthus* is masculine, *Crassula* is feminine, and *Polygonum* is neuter. Specific or infraspecific epithets are usually adjectives, the endings of which must agree in gender with that of the genus name, as in *Amaranthus albus*, *Crassula connata*, and *Eriogonum fasciculatum* ssp. *polifolium*. However, in rare cases the specific or infraspecific epithet is a noun (in apposition), in which case it retains its original gender. For example, in *Cypripedium calceolus*, *calceolus* is a noun and retains the masculine gender despite the neuter gender of the genus name.

One exception to the standardized gender endings is that many tree genera are typically treated as *feminine*, regardless of the ending. For example, the genera *Quercus*, *Pinus*, and *Liquidambar* are feminine in gender, even though they have masculine endings. Thus, specific or infraspecific adjective epithets of these genera names must be feminine (to agree in gender), as in *Quercus alba*, *Pinus ponderosa*, and *Liquidambar styraciflua*.

Note that a **name change** (divided, united, transferred in position, or changed in rank) can necessitate a *change in the gender ending of a specific epithet*. For example, for species *Haplopappus squarrosus*, the ending (-us) is masculine. When this species is transferred to the genus *Hazardia*, the new name becomes *Hazardia squarrosa*. Although the root of the specific epithet does not change, its ending may, in order to agree in gender with the new genus name.

NUMBER

Names of genera, infrageneric names (such as subgenera or sections), and species or infraspecific combinations are all treated as the singular case in Latin. However, all taxon names above the rank of genus are treated as Latin plural nouns. Thus, it is correct to say, e.g., The Orchidaceae *are* a large family of monocotyledons and The Rosales *consist* of many species.

COMMEMORATIVES

Commemorative names are those named after a person or place. Specific or infraspecific commemorative names are usually treated as the genitive case (denoting possession) and must have genitive endings. For male commemoratives, the ending is (1) *-ii*, if the name ends in a consonant, as in *Isoetes orcuttii* (unless the terminal consonant is *-r* or *-y*, in which case a single *-i* is used, as in *Erigeron breweri*); (2) *-i*, if the name ends in a vowel other than *a*, as in *Arctostaphylos pringlei*. For male commemorative names that end in *-a* and for all female commemorative names (regardless of ending) an *-e* is added, as in *Baccharis vanessae* or *Carex barbarae*.

In some cases a commemorative name is treated as an adjective, in which case the endings *-ianus*, *-iana*, or *-ianum* may be used. These endings agree in gender with the

genus name, as in *Lotus nuttallianus* (named after Thomas Nuttall), *Prunus caroliniana* (named after the Carolinas), or *Antirrhinum coulterianum* (named after John M. Coulter). Another suffix ending used for commemoratives is *-ensis*, etc., as in *virginiensis* (of Virginia) or *capensis* (of the Cape).

PRONUNCIATION OF NAMES

Although scientific names are universal, their pronunciation may vary from region to region, especially between different countries. For example, European pronunciations are often different from those of most American botanists. There are no firm rules as to how scientific names should be pronounced. Very often, pronunciations are influenced by one's native language. One should be flexible and adaptive with regard to pronunciations, as the overriding goal is communication.

The rules discussed below are recommended. These generally use traditional English for pronunciation of diphthongs, vowels, and consonants and reformed academic pronunciation (based on classical Latin) for converting to syllables and for accenting. (See Stern, 1992; however, see also Weber, 1986.)

DIPHTHONGS

Diphthongs are two vowels that are combined together and treated as the equivalent of a single vowel. The Latin diphthongs and their traditional English pronunciations are:

Diphthong	English Pronunciation	Example
ae	long e	<i>Tropaeolum</i>
oe	long e	<i>Kallstroemia</i>
au	a w	<i>Daucus</i>

ei	long i	<i>Eichhornia</i>
eu	long u	<i>Teucrium</i>
ui	as in quick	<i>Equisetum</i>

Note that *ie* is not a Latin diphthong, but two separate vowels, each of which would be pronounced separately, as in the genus *Parietaria* (Pa-ri-e-ta-ri-a). Also note that *oi* is not a Latin diphthong. Technically, each vowel should be pronounced separately, as in *Langloisia* (Lan-glo-i-si-a). However, by convention *oi* is often pronounced like the English language diphthong, as in *oil*. Thus, the genus *Langloisia* is often heard as Lan-gloi-si-a.

Occasionally, adjacent vowels will resemble a diphthong, but are actually separate vowels. In ligatured typesetting, the two letters of a diphthong are connected together, such as *oe*, to distinguish the diphthong from two adjacent vowels. However, in cases where the diphthong is not specially indicated (most print these days), a diaeresis (¨) is permitted to indicate that the vowel combination is not a diphthong. For example, in the genera *Aloë*, *Kalanchoë*, and *Monanthochloë*, there is no diphthong; the diaeresis shows that the *o* and *e* are separate vowels and are pronounced separately. (Sometimes these are ignored in practice; for example *Aloë* is usually pronounced as if the ¨ were absent, as in Ah-loh).

SYLLABLES

Latin words have as many syllables as there are vowels and diphthongs. *Every* syllable of a Latin word is pronounced. Thus, it is often valuable to convert scientific names to syllables in order to pronounce them properly and better memorize them. Some of the rules for this are enumerated in Figure 16.3.

Rule	Example	Syllables
A single consonant between two vowels or diphthongs goes with the second one Exception: an "x" between two vowels or diphthongs goes with the preceding one	<i>Tridens</i> <i>exaltataus</i>	Tri-dens ex-al-ta-tus
Two adjacent consonants between vowels or diphthongs are split evenly Exceptions: the combinations <i>bl</i> , <i>cl</i> , <i>dl</i> , <i>gl</i> , <i>kl</i> , <i>pl</i> , <i>tl</i> and <i>br</i> , <i>cr</i> , <i>dr</i> , <i>gr</i> , <i>kr</i> , <i>pr</i> , and <i>tr</i> go together with the following vowel Exceptions: the consonants <i>ch</i> , <i>ph</i> , and <i>th</i> always stay together	<i>guttatus</i> <i>scabra</i> <i>leptocladus</i> <i>Ephedra</i> <i>agrifolia</i> <i>brachypoda</i> <i>eremophila</i> <i>Notholaena</i>	gut-ta-tus sca-bra lep-to-cla-dus E-phe-dra a-gri-fo-li-a bra-chy-po-da e-re-mo-phi-la No-tho-lae-na
Of three or more consonants between two vowels or diphthongs, all but the first goes with the second vowel or diphthong	<i>absconditus</i>	ab-scon-di-tus

FIGURE 16.3 Rules for converting Latinized scientific names into syllables.

Pronunciation Rule	Example	Syllabizing and Accenting	Sounds like
C or g is hard (pronounced like k or a hard g, respectively)	<i>Cakile</i> <i>Garcinia</i>	Ca-k -le Gar-c -ni-a	Kah-k -lee Gar-c h-nee-ah
Exceptions: c or g is soft (pronounced like s or j, respectively) when followed by the letters/diphthongs e, i, y, ae, or oe	<i>Cedrus</i> <i>cinerea</i> <i>coccinea</i> <i>cyaneus</i> <i>caerulea</i> <i>Geranium</i> <i>Gibasis</i> <i>Gypsophila</i>	CŁ-drus ci-nŁ-re-a coc-c -ne-a cy- -ne-us cae-rø-le-a Ge-r -ni-um Gi-b -sis Gyp-s -phi-la	SeŁ-druhs sigh-neŁ-ree-ah Kahk-s h-nee-ah sigh- -nee-us see-rø-lee-ah Jeh-r -nee-uhm Jih-b -sis Jip-s - -lah
When a word or root begins with cn, ct, gn, mn, pn, ps, pt, or tm, the first letter is silent; only the second letter is pronounced.	<i>Cneoridium</i> <i>Ctenium</i> <i>Gnetales</i> <i>Mniodes</i> <i>Pneumatopteris</i> <i>Psilotum</i> <i>Pteridium</i> <i>Tmesipteris</i>	Cne-o-r -di-um CtŁ-ni-um Gne-t -les Mni- -des Pneu-ma-to-ptŁ-ris Psi-l -tum Pte-r -di-um Tme-s -pte-ris	Nee-oh-r -di-um TeŁ-nih-um Nee-t y-lees Ni- -des Noo-ma-to-tŁ-ris Sigh-l -tum TŁh-r -di-um Meh-s -te-ris
Ch is hard, pronounced like k	<i>Chilopsis</i>	Chi-l p-sis	Ki-l p-sis
X at the beginning of word or root is pronounced like a z	<i>Xylococcus</i> <i>Xanthium</i>	Xy-lo-c c-cus XŁEn-thi-um	Zy-lo-c c-cus ZŁEn-thi-um
An x within a word is pronounced like ks	<i>Zanthoxylum</i>	Zan-th x-y-lum	Zan-th ks-i-lum
Anal e or es is long	<i>Anemone</i> <i>Rosales</i>	A-nŁ-mo-ne Ro-s -les	A-nŁ-mo-nee Ro-s y-lees
Anal a is short	<i>Nicotiana</i>	Ni-co-ti- -na	Ni-co-ti- -nah
A y is pronounced like a short i	<i>argophyllus</i>	ar-go-phy-l-lus	ar-go- -l-lus
For uu, both u s are pronounced, the first long, the second short	<i>Carduus</i>	CŁÆr-du-us	cŁÆr-doo-us
An i at the end of a syllable is short	<i>crassifolius</i>	cras-si-f -li-us	cras-si-f h-li-us
An e is long if it is derived from the Greek diphthong ei	<i>Achillea</i>	A-chil-ŁŁ-a	a-kil-leŁ-a

FIGURE 16.4 Rules for pronunciation of Latinized scientific names, ‘ representing a grave accent denoting a long vowel, ‘ representing an acute accent denoting a short vowel.

Special rules for the pronunciation of consonants and vowels are cited in Figure 16.4.

ACCENTING

A standard format for denoting accent is ‘ for a (grave) accent denoting a long vowel, ‘ for an (acute) accent denoting

a short vowel. Determining the accent of a scientific name may be difficult without actually looking up the word in a flora or other reference. However, if these are not available, the following general rules may be used to determine which syllable is accented and whether the vowel of that syllable is long or short.

Determining whether a vowel is long or short generally requires consulting a Latin dictionary.

The last syllable of a word is *never accented* unless the word has only one syllable; e.g., *max* of *Glycine máx*.

If a word has two syllables, the accent always goes with the next to the last (called the penult); e.g., *Ácer*.

If a word has three or more syllables, the accent always goes either with the next to the last (penult) or the third from the last (called the antepenult). The next to the last (penult) is accented if (1) it ends in a consonant (in which case the vowel is short) as in *perennis* pe-rŃn-nis; (2) it ends in a diphthong (which is treated as long), as in *amoenus* a-moŁ-nus; or (3) it ends in a long vowel, e.g., *alsine* al-s -ne. If none of these conditions is met, then the accent goes with the third from the last syllable (antepenult); e.g., *dracontium* dra-c n-ti-um.

COMMEMORATIVES

Although commemoratives may be divided into syllables and accented according to the rules of Latin, they also may be pronounced as the person or place would be pronounced in the native language. For example, the specific epithet of *Hesperoyucca whipplei* may be pronounced w p-pull-i (as the person's name is pronounced plus the letter i) as opposed to the Latinized pronunciation w p-plee-i. The general pronunciation rule is to simply pronounce the commemorative as it would be pronounced in the language of that person, then add the ending. However, in practice the commemorative pronunciation is usually converted to the language of the speaker, as pronunciation in the original language of that person may be unknown or unwieldy. (Remember, the overriding goal is communication!)

REVIEW QUESTIONS

NOMENCLATURE

1. What is nomenclature?
2. What is the name (and abbreviation) of the work that provides the rules and recommendations for plant nomenclature?
3. What groups of organisms are covered by this reference? What organisms are not?
4. What are the two basic activities governed by nomenclature (and the ICBN)?
5. What are legitimate and illegitimate names?
6. How are changes to the ICBN made?
7. Name the six principles of botanical nomenclature. Which of these is considered the fundamental principle?
8. What is the difference between the rules and the recommendations of the ICBN?
9. What is a scientific name? Give an example of such.
10. Which scientific names (i.e., at which rank) are always binomials? Give an example of a binomial.
11. Who first consistently used the binomial and is called the father of taxonomy?
12. What is the correct form of a binomial?
13. For *Quercus dumosa* Nuttall, what is (a) *Quercus*; (b) *dumosa*; (c) *Quercus dumosa*; (d) Nuttall?
14. What are common names?
15. Name the reasons that common names are disadvantageous.
16. What is the difference between rank and position?
17. Name the standardized endings for scientific names at the ranks of phylum, class, subclass, superorder, order, family, subfamily, tribe.
18. What is the rank of the following: (a) Conostyloideae; (b) Flacourtiaceae; (c) Haemodoreae; (d) Hamamelidae; (e) *Linnaea borealis* var. *longiflora*.; (f) Liliopsida; (g) Magnoliophyta; (h) Rosales; (i) *Tribonanthes*; (j) *Tribonanthes variegata*; (k) *Phlebocarya ciliata* ssp. *pilosissima*?
19. What is an alternate, acceptable name for the Apiaceae; Arecaceae; Asteraceae; Brassicaceae; Fabaceae; Faboideae; Clusiaceae/Hypericaceae; Lamiaceae; Poaceae?
20. What is the difference between subgenus and subgeneric?
21. What is a trinomial? What are the two infraspecific ranks and which is higher?
22. What is the author of a scientific name? Which scientific names do not have official authorship?
23. Name four suggestions for memorizing scientific names.
24. What is meant by a nomenclatural type?
25. What is the difference between a holotype, isotype, lectotype, and neotype?
26. Taxa of which ranks have nomenclatural types?

27. What is meant by priority of publication ?
28. When and with what publication does priority of publication officially begin?
29. What is an adverse consequence of priority of publication?
30. What is conservation of names and how is this accomplished?
31. What are the two basic reasons for changing a scientific name?
32. Give the four major ways that names are changed and give an example of each.
33. What is remodeling? Does it require a name change?
34. What is a basionym?
35. What does it mean if an author's name is in parentheses, e.g., *Machaeranthera juncea* (Greene) Hartman?
36. You decide that the taxon *Xiphidium coeruleum* Aublet should be transferred in position to the genus *Schiekia*. What is the required new name (including authorship)? What if the name *Schiekia coeruleum* had already been validly published?
37. You decide that the taxon *Quercus albiniana* (C. Jones) G. Smith ssp. *tomentosa* H. Carlisle should be elevated to the rank of species. What is the new name called (including authorship)?
38. What is an autonym? Give an example of an autonym at the rank of subfamily, subgenus, or subspecies.
39. What are the main criteria of a validly published name?
40. What is a synonym?
41. What are the two major reasons that a name may be rejected?
42. How can a name be legitimate yet not be correct?
43. What can you infer from: *Malacothrix incana* (Nutt.) T. & G. [*Malacomeris i.* Nutt.] ?
44. What can you infer from: *Porophyllum gracile* Benth. [*P. caesium* Greene; *P. vaseyi* Greene] ?
45. What can you infer from: *Gilia diegensis* (Munz) A. & V. Grant [*G. inconspicua* (Sm.) Sweet var. *diegensis* Munz] ?
46. What is a homonym?
47. What is a tautonym? Are tautonyms illegitimate in (a) botanical nomenclature; (b) zoological nomenclature?
48. What is meant by in in authorship designations? How may such a designation be simplified?
49. What is meant by ex in authorship designations? How may such a designation be simplified?
50. What is the meaning of an × in a scientific name, as in *Quercus ×morehus*?
51. How does the fact that a plant and a bird have the same scientific name *not* violate the principles of the ICBN?

BOTANICAL NAMES

52. In what language are scientific names treated?
53. Name the three Latin genders and give the standardized genus endings.
54. What is one prominent exception to these gender endings?
55. Names at which taxonomic ranks are always Latin plurals?
56. What is a commemorative name?
57. What endings may commemorative names have?
58. Are there universal rules for the pronunciation of scientific names?
59. What are the Latin diphthongs and how are they pronounced?
60. How is the combination oi properly pronounced in Latin?
61. What is the rule determining the number of syllables in a scientific name?
62. Name some of the specific rules for converting scientific names to syllables (refer to Figure 16.3).
63. Name some of the specific rules for pronouncing scientific names (refer to Figure 16.4).
64. Name the basic rules for accenting scientific names.
65. Convert to syllables and pronounce the following names: *Cleistis*, *Eucalyptus*, *microcarpa*, *Oenothera*, *Pyrus*.
66. A commemorative (named after a person or place) may be pronounced in what two basic ways?

EXERCISES

1. Using a manual or flora of local, native plants, record 12 scientific names plus the listed synonymy for these names. Trace the nomenclatural history of these taxa names as best you can from the data given, especially noting author names in parentheses.

2. Look up these 12 scientific names using the International Plant Names Index (<http://www.ipni.org>). Record the date and journal/book of publication of these names. Also record all synonymy. Does this added information elucidate the nomenclatural history of the taxa?
3. Divide into syllables, accent, and pronounce these 12 scientific names, using any available references.

REFERENCES FOR FURTHER STUDY

- Bailey, L. H. 1963. *How Plants Get Their Names*. Dover Publications, Inc., New York.
- Borror, Donald J. 1960. *Dictionary of Word Roots and Combining Forms*. Mayfield, Palo Alto, California.
- Brummitt, R. K., and C. E. Powell (eds.). 1992. *Authors of Plant Names: A List of Authors of Scientific Names of Plants, with Recommended Standard Forms of Their Names, Including Abbreviations*. Royal Botanic Gardens, Kew, London.
- Gledhill, D. 1989. *The Names of Plants*. 2nd ed. Cambridge University Press, Cambridge, UK.
- Jaeger, E. C. 1959. *A Source Book of Biological Names and Terms*. 3rd ed. Charles C. Thomas, Springfield, Illinois.
- McVaugh, R., R. Ross, and F. A. Stafleu. 1964. An annotated glossary of botanical nomenclature. *Regnum Vegetabile* 56: 1-31.
- Radford, A. E., W. C. Dickison, J. R. Massey, and C. R. Bell. 1974. *Vascular Plant Systematics*. Harper and Row, New York.
- Stern, William T. 1992. *Botanical Latin: History, Grammar, Syntax, Terminology and Vocabulary*. David & Charles, Brunel House, Newton Abbot, Devon, UK.
- Weber, William. 1986. Pronunciation of scientific names. *Madroæo* 33(3): 234-235.

WEB PAGES:

- Greuter, W., J. McNeill, F. R. Barrie, H.-M. Burdet, V. Demoulin, T. S. Filgueiras, D. H. Nicolson, P. C. Silva, J. E. Skog, P. Trehane, N. J. Turland, and D. L. Hawksworth (eds). 2000. *International Code of Botanical Nomenclature (St. Louis Code)*. *Regnum Vegetabile* 131. Koeltz Scientific Books, K nigstein. <http://www.bgbm.fu-berlin.de/iapt/nomenclature/code/SaintLouis/0000St.Luistitle.htm>
- Griffith, C. 1996 onwards. *Dictionary of Botanical Epithets*. <http://www.winternet.com/~chuckg/dictionary.html>
- International Plant Names Index. A list of current plant names from three sources: the Index Kewensis (IK), the Gray Card Index (GCI) and the Australian Plant Names Index (APNI). <http://www.ipni.org/index.html>
- Taxacom Discussion List. 1992 onwards. A listserv of discussions and queries of taxonomic issues, for plants and animals. Useful for querying specific information, listing images for identification, and searching for topics discussed. <http://listserv.nhm.ku.edu/archives/taxacom.html>

This Page Intentionally Left Blank

17

PLANT COLLECTING AND DOCUMENTATION

INTRODUCTION	517	Living Collections	523
METHODS OF COLLECTING PLANTS	517	Collections for Molecular Studies	524
Field Collecting	518	REVIEW QUESTIONS	524
Preparation of Plant Specimens	518	EXERCISES	524
Documentation of Plant Collections	519	REFERENCES FOR FURTHER STUDY	524
Liquid-Preserved Collections	523		

INTRODUCTION

Plant collections are essential components of systematic research. Collections generally consist of samples of plants that are preserved by drying or by means of liquid preservation. They may also include live plants or propagules taken from the wild and grown in an artificial environment.

Collections of plants serve several purposes. One is to provide resource material in systematic research. Although systematists should attempt to study plants in the wild, in practice almost all research in plant systematics is done using preserved or living plant collections.

Another function of plant collections is to serve as reference material for named taxa. Such a reference plant collection is known as a **voucher specimen**. (Voucher specimens are almost always dried herbarium specimens; see later discussion.) Voucher specimens are required by the International Code of Botanical Nomenclature to serve as types in the valid publication of new taxa names (Chapter 16). Thus, every botanical name at and below the rank of family is associated with the **type specimen** (generally the holotype; see Chapter 16), which is almost always a voucher specimen selected from the original plant collection.

In addition, voucher specimens may serve as a reference in verifying the identity of a plant taxon. If there is ever any doubt as to a taxon's identity, the voucher can be studied to check a prior identification. Reference voucher specimens are essential to obtain and cite in any systematic study. This is

true whether the actual data are acquired from study of morphology, chemistry, anatomy, ultrastructure, reproduction, or molecular biology. Reference vouchers are also essential in field surveys involving the species composition of a given region. Thus, studies of floristic diversity, ecological mechanisms, or environmental assessment (e.g., environmental impact reports) must include plant collections and voucher specimens as a component of the study. Otherwise, the scientific validity of the conclusions may be in doubt.

Finally, the information recorded in the field as part of a plant collection is very important and can be utilized for a number of purposes. Many larger herbaria and some smaller ones have now initiated projects to input data from the labels of herbarium collections into a computerized database system. The database system allows information on plant morphology, ecology, phenology, and geography to be summarized and categorized in order to gain insight on a number of biological questions (see Documentation of Plant Collections). Thus, as these database systems are implemented, plant collections and their associated data are becoming increasingly valuable for fully documenting biodiversity in studies of systematics, ecology, and conservation biology.

METHODS OF COLLECTING PLANTS

Documentation of plant taxa necessitates not only thoroughly recording data in the field about the plant and its habitat but

also procuring a physical specimen. This specimen is obtained by (1) collecting the plant; (2) pressing and drying the plant; and (3) preparing a mounted herbarium specimen by gluing the plants and a label (listing the field data) to a sheet of standard herbarium paper. The specimen is deposited and maintained in an herbarium in order to be accessible for future study, e.g., to verify its identity or prepare a taxonomic revision.

FIELD COLLECTING

Locating specific plants may be by chance or can involve prior checking of specific collection records (e.g., herbarium sheet label information) or pertinent maps to locate the likely location of a plant in a specialized habitat. The collector should obtain prior permission or the proper permit for collecting on a tract of land.

Once a plant of interest is located in the field, the conscientious botanist must evaluate whether or not the species *should* be collected. The first guideline is to become aware of and be able to recognize any possible sensitive species, i.e., those that are rare, threatened, or endangered. These are typically protected by law and may not be collected legally without special permits. Second, regardless of the legal status of a plant species, any collecting should not endanger the local population. A good rule of thumb is the so-called 1 to 20 rule: for every one plant sample you collect, there should be at least 20 more present in the surrounding population. (For herbs, the 1 to 20 rule applies to individual plants; for shrubs and trees, it applies to shoots removed.)

In collecting an herb, at least one whole plant must be completely dug up to show roots and/or rootstocks. (The exception might be a plant that is extremely rare or endangered.) This is often necessary to determine whether the plant is an annual, biennial, or perennial and to determine the type of root (e.g., fibrous or tap) or underground stem (e.g., corm, bulb, or rhizome). With shrubs, trees, or vines, only one or more branches need be clipped off, using hand clippers to minimize damage to the plant. An attempt should be made to collect plants at flowering and/or fruiting stage and to collect enough individual specimens (population size permitting) to represent the range of individual variation.

It is strongly urged that plants be pressed immediately upon collecting in the field. Portable **field plant presses** can be obtained from herbarium supply companies. A simple, inexpensive field plant press can be made by placing several single, folded sheets of used newspaper (preferably ca. 11.5" × 16.5" when closed), between two adjacent 12" × 18" cardboards, all secured with two small elastic cords or straps. Plants are pressed by placing the specimen inside one of the single sheets of folded newspapers, all of which are

temporarily stacked atop one another and sandwiched between the two cardboards. Each newspaper should be labeled with the collection number, referencing that recorded in the field notebook. Plants initially pressed in the field are then later transferred to a standard plant press prior to drying. (See later discussion for details of preparing pressed plant specimens.)

If collected plants are not immediately pressed, they should be stored to prevent wilting. Identifying string tags may be attached to the plant with the collector's name and collection number. Plants then may be stored in a plastic bag. Alternatively, plants may be wrapped in newspaper (open at the top end), wetted, and stored in a large plastic bag; evaporation from the newspaper keeps the plants cool and moist. Ideally, unpressed plants should be kept in an ice chest or refrigerator.

PREPARATION OF PLANT SPECIMENS

The standard method of preserving plants for future study and reference is by the preparation of a specimen that is deposited in an herbarium. An **herbarium specimen** (see Chapter 18) consists of a pressed and dried plant sample that is permanently glued and strapped to a sheet of paper (of standard weight and type, measuring 11.5" × 16.5" in most U.S. herbaria) along with a documentation label (see later discussion). Herbarium specimens or sheets will last for hundreds of years if properly maintained. They are still the most efficient and economical means of preserving a sample of plant diversity.

To prepare an herbarium specimen, material from the field plant press or bag is transferred to a standard plant press to be pressed flat and air dried. A **plant press** consists of several 12" × 18" pieces of standard cardboard that are placed between two outer 12" × 18" frames or 1/4" plywood pieces all secured by two straps (Figure 17.1). Optionally, two 12" × 18" felts may be placed between adjacent cardboards to help absorb moisture, but good results can be obtained without felt. Plants are pressed by placing the specimen inside a single page of folded paper (again, used newspaper, preferably close to the size of herbarium paper, ca. 11.5" × 16.5"), which is then placed between two adjacent cardboards (or felts and cardboards) in the plant press.

The plants to be pressed should be positioned on the newspaper in a way that best represents the plant in the wild and maximizes information content, according to the following guidelines. Open the single sheet of newspaper and carefully place plant organs in a position that allows full view of morphology. Press herbs to show roots and underground stems, which should first be rinsed to remove dirt. Place whole, small herbs on the newspaper with several plants on a single sheet, enough to fill up the space (Figure 17.2B). Taller herbs

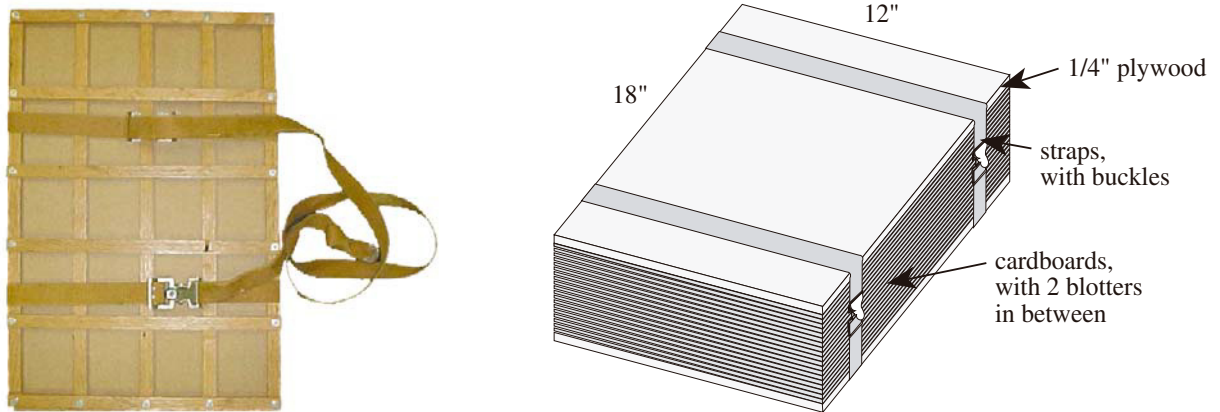


FIGURE 17.1 A standard herbarium plant press.

may be bent into a V, N, or M shape (Figure 17.2A) in order to fit the entire plant on one sheet. If necessary, cut a tall herb into two or more pieces, preparing a separate newspaper for each. Slice large rhizomes, corms, or bulbs longitudinally and place one cut side face down and the other face *up* to show internal structure. For larger or highly branched specimens (Figure 17.2C E), clip back the shoots or leaf ves (leaving the shoot or leaf base) in order to minimize overlapping of parts. Orient at least one leaf up and one leaf down, so that both leaf surfaces will be in full view upon drying. To dry succulent plants properly, cut their leaves or stems longitudinally and, if large, scoop out the fleshy tissue, placing the cut side face down. Cacti and other succulents may be soaked in 95% alcohol for 1–2 days before drying. Arrange flowers or flower parts carefully; section larger flowers to allow viewing of internal organs. Place extra flowers or inflorescences to one side in order to provide extra material for morphological study. Fruits may be sectioned to illustrate internal wall layers or placentation and to facilitate drying. Use wax paper on both sides of fleshy, aquatic, or delicate plant samples in order to prevent adhering to the newspaper. Place folded sections of newspaper on top of leaves or flowers in order to press them flat when the adjacent stems are thick. For all pressed plants, keep the space at the lower right corner (ca. 3" × 5" area) free, as this is where the herbarium label will be glued on the herbarium specimen. After final positioning of the plant sample, carefully fold the newspaper over the plant and place between two cardboards in the press.

After all plants have been placed in the plant press, the straps are tightened and the press is positioned on its long edge (with buckles on the opposite side) inside a plant drier. The plant drier consists of a ventilated box or cabinet having at its base either heating elements or lightbulbs plus a fan to provide air circulation. Because modern techniques permit

removal and amplification of DNA from herbarium material, it is important that plant specimens be dried at not too high a temperature, to prevent DNA degradation. Heated and circulated air rises through the cardboards and newspapers, drying most plants in 2 or 3 days. After this time, the plant specimens should be removed and checked individually; if any specimen feels cool to the touch, water is still evaporating from its tissues, necessitating a longer drying time.

DOCUMENTATION OF PLANT COLLECTIONS

It is critical that certain data be recorded at the time of collecting a plant. Such data will be typed onto an herbarium label and may be entered into computerized database systems. The following is an explanation of the data categories to be recorded at the time of collecting. Figure 17.3 illustrates an example data page for this documentation.

Field Site Data

[List a locality number to cross-reference to other collections.]

Date of collection: List day month (*spell out* to avoid confusion) year

Time (optional): Sometimes important for noting the time of flowering.

Country/state/province/county/city: List as needed.

Specific locality information: List complete locality data for possible relocation of habitat in the future, including measured or estimated distance on roads or trails.

Latitude and longitude: Important to list for biogeographic data systems. Use GPS device or put dot on topographic map to reference plant collection numbers.

Source/accuracy of lat./long.: List how lat./long. is determined, e.g., by USGS 7.5 quad or GPS device. List (in seconds) accuracy of determination.



FIGURE 17.2 Examples of plants collected and pressed. **A.** Herb, stem bent twice to fit on newspaper. **B.** Herb, in which whole plant is collected, including rootstock. **C.** Small shrub, whole plant collected, including roots. **D.** Tree, in which a branch (in fruit, in this example) is collected. **E.** Vine, in flower; rootstock not collected.

LOCALITY #: *2-A*

FIELD SITE DATA	Date of collection: <i>24 April 1994</i>	Time (optional): <i>10 AM</i>
	Country/State/Province/County/City: <i>CA San Diego Co.</i>	
	Specific Locality information: <i>Near hiking trail, just east of Oak Canyon.</i>	
	<i>ca. 1.5 miles north of trailhead at Hwy 83 and Ventura Rd.,</i>	
	<i>Pickwood State Reserve</i>	
	Latitude: <i>32° 50' 28" (N) S</i>	Longitude: <i>117° 02' 59" (W) E</i>
	Source/Accuracy of Lat./Long.: <i>USGS 7.5' topo. quad (La Mesa); +/- 1"</i>	
	Township & Range:	
	Elevation (ft or m): <i>1,100 feet</i>	
Landmark information: <i>Ca. 4.7 miles northwest of Wilson Peak</i>		

PLANT DATA	Coll. No.: <i>702</i>	Photo. doc.: <i>Roll #2, slides 13-14</i>
	Collector (primary): <i>Cynthia D. Jones</i>	
	Associated collector(s): <i>John J. Smith</i>	
	Taxon: <i>Porophyllum gracile Benth.</i>	
	Ann./Bien. <input checked="" type="checkbox"/> Per. <input type="checkbox"/> Habit, Height, Branching: <i>Subshrub, 30-40 cm tall,</i>	
	<i>with several branches from base, densely branched above</i>	
	<input checked="" type="checkbox"/> Fl. <input checked="" type="checkbox"/> Fr. colors, other notes: <i>Involucre purple. Corolla white to greenish-yellow. Pappus bristles white to purplish. Leaves strongly pungent.</i>	
	<i>NOTE: Fls. visited by Checkerspot butterflies. Material preserved in Carnoy's fixative for chromosomal studies</i>	
	Physical Habitat, Substrate: <i>Mountain slope. Rocky, sandy loam soil.</i>	
	Slope, Aspect, Exposure: <i>Slope ca. 30 degrees, south-facing, exposed.</i>	
	Community/Vegetation type: <i>Open Eriogonum fasciculatum - mixed (Artemisia californica, Malosma laurina) scrub.</i>	
	I.d. by: <i>Cynthia D. Jones</i>	Date: <i>4/1994</i>
Accession Number: <i>SDSU 12837</i>		

FIGURE 17.3 Plant collecting documentation sheet.

Township and range: May be listed instead of lat./long., but less preferable.

Elevation (ft or m): List in units appropriate for source of determination.

Landmark information: Describe nearest major landmark (preferably one listed on standard topographic map) and list distance and direction from landmark.

Plant Data

Collection number: A unique number associated with the primary collector. Standard format is for a given person to begin with 1 for the first plant collected, 2 for the second, etc. Another format is to transform the date into a collection number, e.g., 10VI94A, in which the month is in Roman numerals, A represents the first plant collected that day, B the second plant, etc. Note: Duplicate specimens of a taxon collected at the same site and time receive the same collection number. If one plant specimen is divided into two (or more) parts, the labels for the pressed sheets are listed as 1 of 2, 2 of 2, etc.

Photograph documentation: For keeping track of photos or other images.

Collector (primary): The *one* person associated with a plant collection.

Associated collector(s): Other people present or aiding in collecting. These names are not directly associated with the collection number.

Taxon: Scientific name of species (a binomial of genus + specific epithet), including authorship. If applicable, also list the subspecies (ssp.) or variety (var.) name, including authorship. In final form, the scientific name is always underlined or italicized.

Ann./bien./per., habit, height, branching: Circle or list duration (annual, biennial, or perennial), habit (herb, shrub, subshrub, vine, or tree), height from ground level (in metric, not essential if entire plant is collected), and any distinguishing features of the branching pattern that are not apparent from the specimen itself.

Fl./Fr., colors, other notes: Circle or list phenology, whether plants are in flower and/or fruit. Precisely describe the colors of unusual vegetative parts and of all flower parts (e.g., of calyx, corolla, anthers). If precision needed for colors, use a chart (e.g., Royal Horticultural Society Color Charts; see Tucker, et al. 1991. *Taxon* 40: 201-214.) Describe features that are obscure or might be lost from specimen upon drying.

Other field notes may include references to additional research studies or additional field observations, such as observed visitors/pollinators.

Population size/distribution: A few notes about the size and distribution of the population are useful, such as very rare, population very large (≥ 1000 individuals per hectare), or plants locally common.

Physical habitat/substrate: Physical habitat refers to abiotic features, such as dry creek bed, granite outcrop, or flood plain. For substrate, list color and basic soil type (e.g., clay, clay-loam, loam, sandy-loam, sand, gravel, boulder, or rock). More detailed information can include soil series and/or rock type.

Slope/aspect/exposure: List angle of slope, from none (flat) to 90° (cliff face). Aspect is general compass direction toward which slope is facing. Exposure is either exposed, partly shaded, or shaded.

Community/vegetation type: Both immediate and surrounding plant communities/vegetation types may be listed for a single plant collection. Community/vegetation type may be general (e.g., chaparral or woodland) or precise. A precise designation of community type (modified from Radford, A. E., et al. 1981. *Natural Heritage: Classification, Inventory, and Information*. University of North Carolina Press, Chapel Hill) is as follows:

1. Determine the **boundaries** of the community, based on overall similarity of species composition. This may not be clear cut, as one community may intergrade with another or show much variation.
2. Identify **layers** present in the community: *canopy* (tall tree and lianas, if present) / *subcanopy* (smaller tree layer under canopy) / *shrub* or *subshrub* / *herb*. A vine, epiphyte, moss, or lichen layer also may be defined if a major component of the community.
3. For each layer of vegetation, assess the **total cover**, measured as the degree to which the total area of the community is covered by members of a given layer. Designations of cover are (1) *closed* (50-100% cover); (2) *open* (25-50% cover); or (3) *sparse* (<25% cover).
4. For each of the common species of a given layer (e.g., the shrub layer), assess **relative cover**, measured as the degree to which each species of the layer contributes to the total cover *of that layer alone*. (Other ecological measures, such as importance value, may be used instead, but relative cover is perhaps easiest to eyeball in the field.) Assess relative cover as (a) *dominant* = > 50% relative cover; (b) *codominant* = 25-50% relative cover.
5. Summarize the **community type** by listing layers separated by a / in sequence from tall to short layers, e.g., tree / shrub / herb / moss. Note: Dominant or co-dominant vines and epiphytes are listed at the end separated by, respectively, a double slash (//) or triple slash (///).

List as follows: (a) Total cover; (b) Dominant species, if *one* species is dominant (50-100% cover); or (c) codominant species, if two or more species are codominant (25-50% cover), each species present separated by a hyphen.

Note: You may use mixed trees, mixed shrubs, or mixed herbs as a layer designation where collectively the group of mixed species is dominant or codominant, but each individual species is <25% relative cover. This designation may be followed by a listing of the more common species (those with at least 10% relative cover) in parentheses.

6. Follow the community type with a designation of **vegetation type**. This is based on habit, habitat, and cover of species present. Examples include *forest* (closed trees), *woodland* (open trees), *savanna* (sparse trees with intervening grassland), *chaparral* (closed, evergreen, sclerophyllous shrubs), *scrub* (open to sparse shrubs), *grassland* or *meadow*, *strand* (sparse, low shrubs/herbs), *marsh* (aquatic shrubs and/or herbs in slow-moving water), *swamp* (closed to sparse aquatic trees), *pond*, *vernal pool*.

Example: Open *Malosma laurina* - *Artemisia californica* / closed *Erodium botrys* scrub community

Meaning: The total cover of the shrub layer is open (25-50% total site area). *Malosma laurina* and *Artemisia californica* are a codominant in the shrub layer (25-50% relative cover). *Erodium botrys* is a dominant in the herb layer, which is closed (>50% relative cover). The vegetation type is a scrub (open shrubs).

I.D. by / date / source: List the person who identified the taxon, even if it is the same as the primary collector. Also list the date, usually just the month and year, and the source or reference of determining the taxon identity. The source will generally be a flora of the region, but could include monographic treatments or expert determination.

Accession number: After the plant collection is processed into an herbarium sheet and deposited in an herbarium, list the herbarium acronym and accession number for a complete record of the collection. Accession numbers are usually cited in publications to document a collection (see later discussion).

LIQUID-PRESERVED COLLECTIONS

It is often valuable to preserve samples of a plant collection in a liquid preservative. Liquid preservation maintains the shape, size, and internal structure of plant tissues. This is particularly valuable to do for delicate floral parts, whose form is easily distorted or even destroyed from standard herbarium specimen drying techniques. Liquid preservation is also essential for anatomical, developmental, or ultrastructural studies, in which the internal structure of cells and tissues must be maintained.

The most commonly used, general liquid preservative (known as a fixative) is FAA, one recipe being 10 parts 70% ethanol : 1 part commercial (37%) formalin : 1 part glacial acetic acid (all by volume). (Note: FAA is toxic; avoid getting on skin or breathing the fumes!) Plant samples are simply placed into a glass or plastic vial or jar filled with FAA. Although FAA penetrates most plant tissues rapidly, some plant samples should be cut open to allow the fixative to fully infiltrate into the tissues. At least some closed flower buds or ovaries, leaves, and stems should generally be sectioned with a razor blade prior to fixation.

For cytological studies, e.g., chromosome counts, flower buds or root tips may be fixed in Carnoy's fixative (3 parts 100% ethanol : 1 part glacial acetic acid). For detailed ultrastructural studies, e.g., using electron microscopy, other fixatives may be needed, such as glutaraldehyde or osmium tetroxide. These compounds are dangerously toxic and should only be handled in a laboratory hood. Because they penetrate less rapidly than FAA, the material must be cut into much smaller pieces, generally 1 mm or less.

Plant material may be fixed in 70-100% ethanol and used for general morphological studies and sometimes DNA analysis. This is not commonly done for the latter, as material dried in silica gel is better preserved (see later discussion).

Any liquid preserved material should have a corresponding herbarium voucher specimen to serve as a reference for identification. The vial or jar should be labeled both on the outside and on a strip of paper (using a pencil) placed into the fixative. Label information should include the species name and collector and collection number; other data are optional and can be obtained from the field collection notebook or voucher.

LIVING COLLECTIONS

A very valuable type of plant collection is a live specimen removed from the wild. This may be either a whole plant, a vegetative propagule, or a seed. Living plant collections are typically grown in a greenhouse or botanic garden, where they can be accessible to a researcher. Growing them and keeping them alive requires some horticultural experience and may involve trial and error under different regimes of potting or soil mixture, moisture, and photoperiod. As with liquid-preserved collections, they should be properly labeled with permanent metal or plastic tags, with collection information corresponding to a voucher specimen deposited in an herbarium.

A living plant collection has the great advantage of permitting long-term observations, e.g., through an entire reproductive stage, or experimental manipulations, such as breeding studies. It also permits removing fresh samples of material for study over an extended period of time (rather than from a single field expedition). However, one precaution about

studying live plant collections is that their morphology may be altered in cultivation from that in the wild. In addition, pollinators normally present in the wild will not normally be present in an artificial environment, perhaps preventing normal seed set.

COLLECTIONS FOR MOLECULAR STUDIES

A standard method for collecting material for studies of DNA is to cut pieces of leaves or other plant tissue and immerse these in a container (vial or plastic bag) of silica gel. A paper label, indicating the taxon and the name and number of the collector (corresponding to an herbarium voucher collection),

is placed in the container. The silica gel rapidly dehydrates the material, preserving the DNA for future extraction, purification, and amplification. Extracted plant material is usually frozen at -80°C to prevent degradation of the DNA. Plant material to be used for DNA analysis may also be fixed in 70–100% ethanol, but this may not preserve the DNA as well.

For allozyme analysis, fresh material must be used, as enzymes degrade very rapidly. Extra plant material is placed in a plastic bag (again with a slip of paper or label indicating the voucher information) and kept in a cooler until it is transported to the lab.

REVIEW QUESTIONS

1. What are the different types of plant collections?
2. List the several uses of plant collections.
3. What is a voucher specimen?
4. What is the purpose of a voucher specimen?
5. Review the preparations needed for collecting plants in the field.
6. What are the general rules for assessing whether a plant *should* be collected?
7. List the guidelines for properly collecting plants in the field.
8. How should plant collections be stored prior to processing?
9. What is a standard herbarium specimen?
10. What are the components of a plant press?
11. Review the guidelines for properly pressing plants, including special requirements for processing (a) herbs; (b) tall herbs; (c) shrubs or trees; (d) highly branched specimens; (e) rootstocks (such as rhizomes, bulbs, or corms); (f) succulent plants; (g) flowers and fruits.
12. How are herbarium specimens dried?
13. List all the data that should be recorded in the field at the time of collecting.
14. Review in detail how specific plant community types can be assessed.
15. Why is it important to list the person who determined the identity of the plant and the determination source?
16. Review the guidelines for preparing liquid-preserved collections.
17. What is the most common type of liquid preservative?
18. What liquid preservatives must be used in ultrastructural studies?
19. Review the guidelines for obtaining living plant collections.
20. What are the advantages and disadvantages of living plant collections?
21. How is material for molecular studies normally collected?

EXERCISES

1. Collect six plants, including at least two herbs, one shrub, and one tree. Record all pertinent information in the field, using Figure 17.3 as a guide.
2. For at least one of the above, collect liquid-preserved material for both anatomical and cytological studies.
3. For at least one of the above, collect material for DNA sequence studies.

REFERENCES FOR FURTHER STUDY

- Radford, A. E., W. C. Dickison, J. R. Massey, and C. R. Bell. 1974. *Vascular Plant Systematics*. Harper & Row, New York.
- Radford, A. E., et al. 1981. *Natural Heritage: Classification, Inventory, and Information*. University of North Carolina Press, Chapel Hill.
- Tucker, A. O., M. J. Maciarello, and S. S. Tucker. 1991. A survey of color charts for biological descriptions. *Taxon* 40: 201–214.

18

HERBARIA AND DATA INFORMATION SYSTEMS

HERBARIUM SPECIMENS	525	Removing Plant Material	531
Herbarium Labels	525	Re-lining Herbarium Specimens	532
Mounting Herbarium Specimens	526	Herbarium Loans	532
		Insect Control	532
HERBARIUM OPERATIONS	529	DATA INFORMATION SYSTEMS	532
Curators	529	FLORAS AND MONOGRAPHS	533
Accessioning	529	REVIEW QUESTIONS	533
Storage and Classification of Specimens	529	EXERCISES	534
Using Herbaria	529	REFERENCES FOR FURTHER STUDY	534
Removing and Handling Specimens	529		
Synoptic Collections	530		
Annotation Labels	530		

Herbaria are repositories of preserved plant collections, these usually in the form of pressed and dried plant specimens mounted on a sheet of paper. The purpose of herbaria is both to physically contain the plant collections and to act as centers for research. The plant collections themselves function as vouchers for identification and as sources of material for systematic work. Herbaria also may house numerous geographic and taxonomic references, particularly floras or manuals that may aid in plant identification. In addition to housing plant collections, many herbaria today have initiated computerized data information systems to record and access the collection information of the plant specimens, as well as to access information from other collections worldwide (see **Data Information Systems**).

Information about herbaria is contained in *Index Herbariorum* (Holmgren et al. 1990; see also listing of on-line computer access in References for Further Study), which lists the names, addresses, curators, and number and types of specimens. Each herbarium listed in *Index Herbariorum* is assigned an acronym. It is this acronym that is cited in publications in order to specify where voucher specimens are deposited. Herbaria are typically associated with universities or colleges, botanic gardens, museums, or other research institutions.

The 15 largest herbaria, their acronyms, and the number of specimens they contain are listed in Figure 18.1.

HERBARIUM SPECIMENS

An **herbarium specimen** consists of a pressed and dried plant sample that is permanently glued and/or strapped to a sheet of paper along with a documentation label. The herbarium paper is high quality, heavyweight, and acid-free to inhibit yellowing. In most American herbaria, standard herbarium paper measures 11.5" wide × 16.5" tall; in other countries the dimensions may be slightly different. An herbarium label (see below) is glued to the lower right corner of the herbarium specimen. An example of an herbarium specimen is seen in Figure 18.2. Herbarium specimens (also called *herbarium sheets*) will last for hundreds of years if properly maintained. They are still the most efficient and economical means of preserving a record of plant diversity.

HERBARIUM LABELS

An **herbarium label** is affixed to each specimen, usually at the lower right hand corner. Herbarium labels are typically

Herbarium	Year Founded	Acronym	Number of Specimens
Muséum National d'Histoire Naturelle, Paris France	1635	P,PC	9,500,000
New York Botanical Garden, Bronx, New York, USA	1891	NY	7,000,000
Royal Botanic Gardens, Kew, England, UK	1841	K	7,000,000
Conservatoire et Jardin Botaniques, Genève, Switzerland	1824	G	6,000,000
Komarov Botanical Institute, St. Petersburg (Leningrad), Russia	1823	LE	5,770,000
Missouri Botanical Garden, Saint Louis, Missouri, USA	1859	MO	5,400,000
The Natural History Museum, London, England, UK	1753	BM	5,200,000
Harvard University, Cambridge, Massachusetts, USA	1864	GH	5,005,000
		(incl. A,AMES,ECON,FH,NEBC)	
Naturhistorisches Museum, Wien, Austria	1807	W	5,000,000
Swedish Museum of Natural History, Stockholm, Sweden	1739	S	4,400,000
Smithsonian Institution, Washington, DC, USA	1848	US, USNC	4,368,000
Institute de Botanique, Montpellier, France	1845	MPU	4,000,000
Université de Lyon, Villeurbanne, France	1924	LY	4,000,000
Universitatis Florentinae, Museo Botanico, Firenze, Italy	1842	FI	3,650,000
Friedrich-Schiller-Universität, Jena, Germany	1895	JE	3,000,000

FIGURE 18.1 Names, acronyms, and number of specimens of the 15 largest herbaria in the world. After Holmgren et al. (1990), updated from Web page (see **References for Further Study**).

computer generated using a laser or ink jet printer. Label sizes vary, but are generally about 4.5" (10.2 cm) wide and 2.3" (5.7 cm) tall, using high-quality, thick-weight (20- or 24-lb), acid-free bond paper. Virtually all of the information recorded at the time of collecting should be placed on the herbarium label. An example of a typical label format, containing all information from the collecting event, is seen in Figure 18.3. A convenient formatting is to list (following the taxon name) all characteristics about the plant itself in the first paragraph, including duration/habit/height/branching pattern and phenology, colors, and other features. The second paragraph contains information about the habitat and locality of the plant, including physical habitat/substrate, slope/aspect/exposure, community/vegetation type, specific locality information, landmark information, latitude and longitude, source/accuracy of lat./long., and elevation. A third paragraph may include other field notes and photograph/image documentation. At the bottom of the label, the collector, collection number, and date of collection is listed. (The abbreviation **s.n.**, Latin for *sine numero*, without a number, sometimes follows a collector's name to indicate that the collector did not designate a personal collecting number.) The last item on the herbarium label may list by whom and when the identity was determined (even if by the same person who collected the material) and what the source of that identification was. Information on taxon determination is important to include on the label, as it cannot be assumed that the person who collected a plant identified it. In addition, the source or means of identification (whether a flora, monograph, or expert

determination) may constitute valuable information in verification of identities.

If the plant specimen is so large that it must be divided between two or more herbarium sheets, a separate label must be prepared for each of these parts. All labels referring to the same plant have the same collection number (but different accession numbers; see later discussion). The two herbarium sheets may be differentiated by the designation, e.g., 1 of 2, 2 of 2.

MOUNTING HERBARIUM SPECIMENS

Plant specimens are affixed to herbarium paper with glue and/or straps. The glue used may be standard white glue or a solution of methyl cellulose, available from chemical supply and some herbarium supply companies. White glue is best diluted slightly, ca. 9 parts glue to 1 part tap water, stirred well. Methyl cellulose is prepared by adding about 70 grams of methyl cellulose powder to a liter of warm tap water and stirring briskly until well mixed; more water or powder may be added to achieve a thick, viscous solution. The advantage of methyl cellulose is that, with minimal moistening, it will soften or dissolve, allowing for relatively easy removal of dried plant material from the herbarium specimen (see later discussion). Glues containing organic solvents are not recommended, as they are toxic and require special ventilation.

The following is one useful method to glue a dried plant specimen and label to a sheet of herbarium paper. Have the following supplies on hand: herbarium paper, cardboard (12" × 18"), a flat sheet (ca. 12" × 18"), paintbrush (2.4" wide),

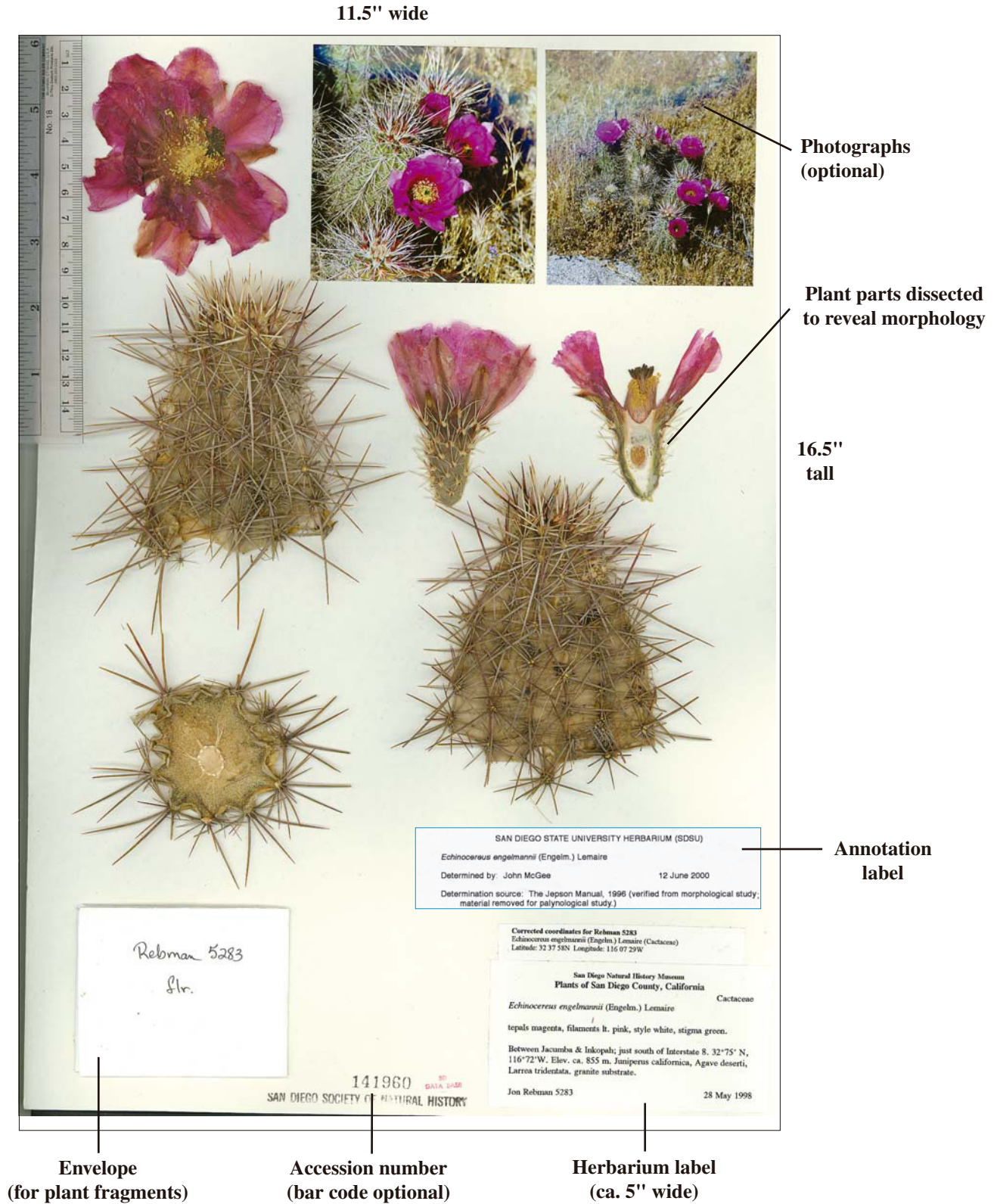


FIGURE 18.2 Example of a typical herbarium sheet. (Photo courtesy of Jon Rebman.)

ca. 5" wide

SAN DIEGO STATE UNIVERSITY HERBARIUM
 USA
 CALIFORNIA
 San Diego Co.

Porophyllum gracile Benth.

Perennial subshrub, 30–40 cm tall, with several branches from base, densely branched above. In flower and fruit. Involucre purple. Corolla white to greenish yellow. Pappus bristles white to purplish. Leaves strongly pungent. Note: Flowers visited by checkerspot butterflies. Material preserved in Carnoy's xative for chromosomal studies.

Near hiking trail, just east of Oak Canyon, ca. 1.5 miles north of trailhead at Hwy 83 and Ventura Rd., Pickwood State Reserve. Mountain slope. Rocky, sandy loam soil. Slope ca. 30 degrees, south facing, exposed. Open *Eriogonum fasciculatum* mixed (*Artemisia californica*, *Malosma laurina*) scrub. 32°50'28" N 117°02'59" W (USGS 7.5' La Mesa quad, ±1" accuracy.) Elevation 1100 ft. Ca 4.7 miles northwest of Wilson Peak.

Cynthia D. Jones 702 24 April 1994
 with John J. Smith

FIGURE 18.3 Example of a typical herbarium label (this one hypothetical).

glue, two pairs of forceps, spatula, and weights (standard bathroom tiles, measuring 4", 6", and 8" square work well). First, place a sheet of herbarium paper on top of a cardboard. Position the herbarium label (without gluing yet) on the lower right corner of the herbarium paper, leaving ca. 1/8–1/4" space between the label and the margins of the paper. Place the pressed plant specimen (also without gluing yet) on the paper in order to test the final positioning. Make sure the specimen does not overlap the label or go beyond the edges of the herbarium paper; if overlap occurs, the plant must be cut. Also, try to leave some room above and to the left of the label for placing an accession number or barcode and possible annotation labels (see later discussion). Extra pieces of the plant specimen (e.g., individual flowers, fruits, or inflorescence) may be placed on the sheet as well. Smaller pieces are best placed in a separate small envelope that may be glued to the final specimen, such that it may be opened to remove the material for study. (Envelopes may be constructed by cutting heavyweight, 100% bond typing/printing paper (e.g., 8.5" × 11") into two pieces; each 8.5" × 5.5" piece is then folded to make a 4.25" × 5.5" rectangle, which is then folded to overlap ca. 1/4" along the three cut margins.)

Next, using a paintbrush, coat a large (at least 12" × 18") sheet (e.g., of glass, Plexiglas, or a cookie sheet) thoroughly with a layer of glue. Transfer the plant specimen from the paper to the glue-covered sheet, gently press down, carefully remove (using forceps for delicate material to prevent damage),

and place back onto the herbarium paper, positioning the plant as originally placed. You may use a scalpel or squirt jar to transfer glue directly to plant surfaces that require greater adhesion. Continue this until all plant components are glued to the sheet. Finally, in a smaller region of the sheet, paint a very thin layer of glue (preferably white glue, diluted as specified above) on the sheet, place and press down the herbarium label onto this region, and transfer back to the herbarium specimen with forceps, being careful to correctly position it about 1/4" from the edges. Flatten and smooth the herbarium label by placing a used paper towel or sheet of paper (to absorb excess glue) over the label and pressing down firmly. Finally, place weights (e.g. different-sized ceramic tiles or lead weights) over both herbarium label and various locations of the plant material. Leave the specimen overnight to dry thoroughly. Specimens, with underlying cardboard, may be stacked if needed to conserve space.

After the glue has dried, remove the weights and check the specimen. Reapply glue to individual spots as needed. Place narrow (ca. 1/8" wide) strips of strapping tape (available from herbarium supply companies) over stout stems to better secure them to the sheet. Some herbaria use little to no glue, relying on heavy use of strips of strapping tape to secure the specimen. Although this may not secure some plant specimens as well, it has that advantage of making removal of plant material from a mounted specimen (e.g., for detailed study) much easier.

HERBARIUM OPERATIONS

CURATORS

The person in charge of the day-to-day running of an herbarium is known as a **curator**. The duties of the curator (and assistant curators or collections managers, if any) are to (1) manage the existing collection, including the mechanics of proper storage and regular treatment to control insect pests; (2) mount, label, and accession new additions to the herbarium collection; (3) distribute requested loans from scientific institutions and receive loans from other herbaria; and (4) act as a resource person for the identification of regional plants or plants of special collections. In addition, curators today are often involved in transferring herbarium collection data to a computerized data information system for interactive access to that information.

ACCESSIONING

Accessioning refers to the designation of a number to all specimens placed into the permanent herbarium collection. This **accession number** is assigned to each specimen of the collection. The accession number is written or imprinted onto the herbarium sheet, often along with the international acronym of the herbarium (e.g., **UC 218485**, where **UC** is the herbarium acronym, in this case referring to University of California at Berkeley). With the advent of computerized data information systems, accession numbers are now sometimes imprinted with a bar code label that may be scanned (see later discussion).

The purpose of the accession number is to provide a permanent reference for each specimen of the plant collection. Accession numbers (plus the collector and collection number) are often cited in journal publications and may be valuable in tracking down the exact specimen for purposes of identity verification.

STORAGE AND CLASSIFICATION OF SPECIMENS

Standard procedure is to store herbarium specimens by genus in a **genus folder**. There are different constructions of genus folders, but all consist of stout, heavyweight paper folded along at least one crease. The genus folder is labeled, typically on the lower right corner of the outer cover. The specimens within a genus folder are usually arranged alphabetically by species. If a particular genus has numerous collections, two to many genus folders may be used to house them.

Genus folders are often **color-coded** to represent different geographic regions for the plant collections. For example, different colors may represent various counties, states, regional areas (e.g., Southwestern states), countries, blocks of countries, or continents. Thus, specimens of the same

species may occur in two or more genus folders if these were collected from different geographic regions. Color-coded genus folders are typically stacked one on top of the other according to a standard order.

Plant specimens are usually stored in **herbarium cabinets** (Figure 18.4). Herbarium cabinets are usually made of metal and have sealed doors to inhibit insect migration or to prevent possible diffusion of pesticides. A standard, full-sized herbarium cabinet is typically 7 feet tall and 2.5 feet wide, with approximately 26 shelved compartments arranged in two columns, having a capacity of approximately 500 herbarium specimens (depending on the bulkiness of the plants). In many of the larger herbaria, standard herbarium cabinets have been replaced by **compactors**, which allow for a greater number of specimens to be stored. Compactor systems consist of rows of attached cabinets (or shelves), each row mounted on floor tracks. Entire rows of cabinets can be moved as a unit to abut against an adjacent row. Thus, compactors generally allow for only one (temporary) aisle space, maximizing the storage space available.

Genus folders are usually classified alphabetically within a given plant family. In many herbaria, families are arranged according to someone's formal classification system, a common one still in use being the *Dalle Torre and Harms*, which is based on the antiquated Englerian system. In other herbaria, families are simply classified alphabetically, with the exception that the major plant groups (e.g., Lycophytes, Equisetophytes, Leptosporangiate Ferns, Conifers, Gnetales, Monocots, or Eudicots) are usually stored separately.

USING HERBARIA

In general, use of an herbarium requires prior approval and/or an appointment made through the herbarium curator. When using the herbarium, please be considerate of dissecting microscopes, tools, and references in the collection. Clean up after yourself; brush the table clean of debris (into a trash can) as needed.

REMOVING AND HANDLING SPECIMENS

You will, of course, need to remove herbarium specimens from the collection for observation, for example, to check the identity of your own plant or to study a given taxonomic group. Herbarium specimens may be rather fragile and should be handled very carefully, as follows.

Note taxa lists or maps to locate the family and genus of interest. Remove the entire genus folder from the cabinet. You may wish to *slightly* pull out the genus folders above or below the desired folder to mark the location. (However, always recheck the labels when filing!) *Close the herbarium cabinet door* immediately in order to inhibit insect infestation.



FIGURE 18.4 Example of a standard herbarium cabinet. Note genus folder (color-coded) on cart.

Carefully transfer the genus folder to a table (with plenty of space) for observation and open it.

Always hold an individual specimen with both hands to prevent it from inadvertently bending. Never place anything (e.g., books) on top of a specimen. *Never turn a sheet upside down*, as this may result in the plant material breaking or falling off the sheet. Remove each specimen, one at a time from the top, and stack (in reverse order) to the side. Avoid sliding stacked specimens against one another, as this can result in damage to the plants. To find a specific collection, you may very gently shuffle through the labels at the lower right-hand corner of the specimens. Then, move aside the group of sheets on top of the desired specimen to expose it. When finished, replace the removed specimens in the genus folder, generally classified in alphabetical order by species or infraspecific name.

SYNOPTIC COLLECTIONS

Synoptic collections are those that contain generally one specimen (of all available specimens) of each taxon for a given region. Synoptic collections are very useful for quickly perusing the possible taxa in a region, such as a state, county, park, reserve, or some other political boundary. The disadvantage of synoptic collections is that they are generally limited to one specimen per taxon. Thus, it is imperative to always

check an identity with the main collection, to note the entire range of variation of the taxon.

ANNOTATION LABELS

An **annotation label** is a label that verifies or changes the identity of a specimen or that documents the removal of plant material from the specimen (Figure 18.5). Annotation labels are permanently glued to the plant specimen, typically just above the standard herbarium label (see Figure 18.2). Annotation labels are typically placed on herbarium specimens by experts in a particular group, often as part of a research project. The labels vary in format, but generally measure about 4" wide and 1" high, using paper like that of herbarium labels.

Annotation labels that verify identity (Figure 18.5A,B) generally include (1) the name of the herbarium; (2) the species, subspecies, or variety name, including full authorship; an exclamation point ! (symbol for the Latin *vidi*, I have seen it) indicates a confirmation and is sometimes written instead of the full taxon name, if the annotated name is the *same* as that on the herbarium label; (3) the name of the person who made the correct name determination (often listed after Det.); (4) the date of the determination; and (5) the determination source. As with herbarium labels, the determination source is often omitted from annotation labels, but is

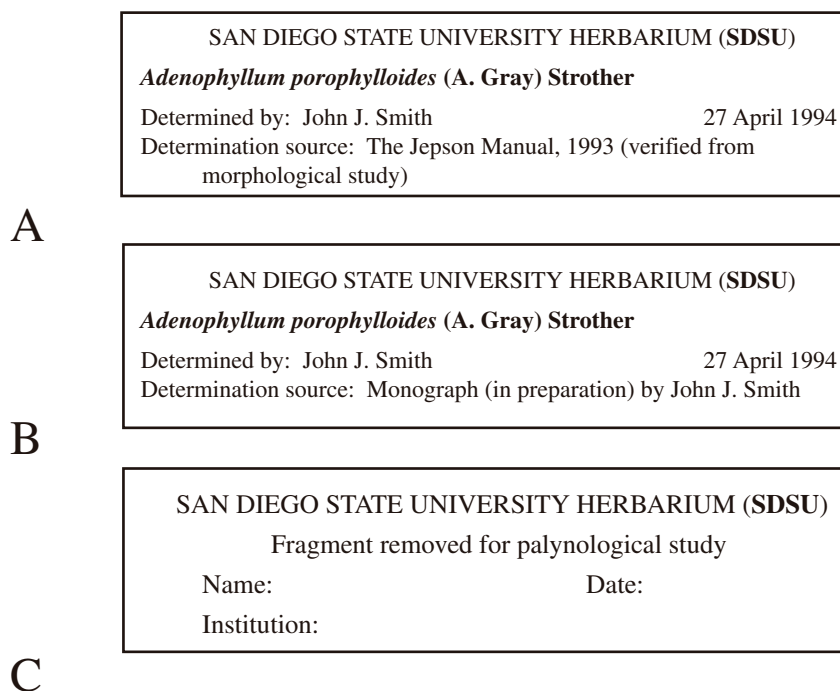


FIGURE 18.5 Examples of annotation labels citing identity (A,B) or documenting removal of plant material (C).

nonetheless valuable to include. The determination source refers to which, if any, references were used in the determination (usually a specific flora) and how the correct identity was ascertained.

One type of annotation is an update of the nomenclature of a species, without verifying the identity by morphological examination. Such annotations are not ideal, but may be necessary in large collections in order to cite the specimens in terms of the nomenclature of a recent flora or monograph. In such a case, the original identity is assumed to have been correct and now a synonym of the new name. Another type of annotation would list the reference and indicate that the person making the determination did examine the specimen critically, examining its morphology. An example of this type of annotation label is seen in Figure 18.5A. A third type of determination source might cite an original monographic treatment, published or unpublished. An example of this type of annotation label is seen in Figure 18.5B.

REMOVING PLANT MATERIAL

Annotation labels may also document the removal of dried plant material from an herbarium specimen. Removal of material may be needed to verify the identity of the specimen or to study some detail of the plant, e.g., anatomy or palynology.

Even DNA may be successfully extracted from dried plant fragments.

You must *always get permission* from the herbarium curator before removing any material. Once permission is obtained, first see if the material you need is contained in an envelope attached to the sheet. If not, you will need to remove a piece from the plant specimen that is affixed to the sheet. Be very careful and conscientious doing this, trying to minimize the damage done to the specimen. Generally, material may simply be clipped, cut, or pulled off with forceps. In some cases the material that you need will be directly glued to the sheet. This may be difficult to remove, requiring the use of a razor blade to gently cut under the specimen (but above the paper). Plant material that is glued with methyl cellulose is easily removed by adding a few drops of water to soften the adhesive. Dried material can be reconstituted in boiling water and/or a detergent solution (such as Aerosol OT). It may then be observed and dissected in water or fixed in a liquid preservative for long-term storage.

Annotation labels should be used to document the removal of plant material. The person, institute, date, and reason for removing material should be indicated. Additional information indicates the type or purpose of the study, e.g., for anatomical, morphological, palynological (pollen), embryological, ultrastructural, or molecular analysis (Figure 18.5C).

REFILING HERBARIUM SPECIMENS

At some herbaria the staff do all specimen refileing; others allow (and expect) the user to refile anything removed. Generally, if a genus folder was removed for a short period of time and *not* placed in proximity to plant debris, it may be refilled immediately into the collection. However, if possible insect contamination is suspected, the genus folder and its contents should be treated for insect control (see later discussion).

If you are refileing a folder yourself, you should be extremely careful to file it in the correct location, both by taxonomic category and geographic region (color-coding).

HERBARIUM LOANS

Those doing research on a plant group do not generally need to visit herbaria to examine the specimens. Typically, herbarium specimens may be sent out (via standard mail) on loan. Loans are typically granted only to members of universities, museums, or other research institutions. A request for a loan requires a letter to the curator, justifying the research needs for examining the specimens. The period of a loan is often 6 months, but this varies at different herbaria and may be extended upon request and approval.

INSECT CONTROL

An essential component of maintenance of herbarium collections is insect control. If herbarium specimens are kept dry and free of insects, they may be preserved in good condition for hundreds, if not thousands, of years. However, if insects infect a specimen, it may quickly be reduced to rubble.

There are two general ways to control insects: by chemicals and by freezing. Chemical control generally involves placing a volatile chemical insecticide within each sealed cabinet. Moth balls or moth chips have been used in the past,

but these have been shown to be extremely dangerous to people. Other types of insecticides include insecticide strips, which can be placed directly into a cabinet and which may last for 3–6 months.

Another method of insect control is freezing. Genus folders are periodically placed in a freezer for 3 to 7 days at at least -20°C before being refilled back into the herbarium cabinet. The advantage of freezing is that it eliminates toxic fumes, which could cause health problems to those working in the herbarium. The disadvantage of freezing is that it is more labor-intensive and potentially may result in greater damage to the specimens because of the regular removal and refileing required.

DATA INFORMATION SYSTEMS

A **data information system** (or **database system**) refers to the organization, inputting, and accessing of information. The accumulation of separate pieces of data (known as the database) may be manipulated such that general or specific questions in plant taxonomy may be addressed. It is important that students today be trained in the basics of accessing and manipulating information available from these systems.

All data information systems utilize computer hardware and software to record information. The data are organized as discrete units, generally known as fields (Figure 18.6). For example, for information in an herbarium collection, typical fields might be species name; authorship; flower color; phenology; soil type; topography; community type; specific locality; latitude; longitude; collector; collection number; determinor; accession number; annotation information; etc. Basically all of the discrete items recorded at the time of

Taxonomic Number	District
Group	Locality
Family	Latitude
Genus	Longitude
Species Epithet	Elevation
Species Author	Habitat Description
Infraspecific Rank	Date of Collection
Infraspecific Name	Collector
Infraspecific Author	Collector Number
Determination	Associate Collectors
Determination Date	Morphology
Country	Phenology
State	Accession Number
County	Supplemental Information

FIGURE 18.6 Example of data fields of a computerized plant inventory data information system.

collection or as part of accessioning or identifying the plant may be entered into the database. In addition, a digitized image of the plant specimen may be recorded for future access.

The great advantage of computerized data management systems lies in the ability to retrieve or summarize information about the plant collection. For example, one may call up a listing of all plant specimens collected at a specific locality or within a certain geographic range, defined by latitude and longitude coordinates. Or, one may request a list of all species collected on serpentine soil, or of all species that flower in September, etc. Sophisticated systems may be able to generate a dot distribution map of all the collections of a given taxon.

One critical problem with data information systems resides not with the system itself but with the collections. Many, if not most, herbarium specimens lack much of the critical information needed. For example, information about plant characteristics, phenology, ecology, or latitude/longitude is often not recorded on labels. In fact, on many older herbarium specimens, locality information may be very scanty, corresponding to a very broad or ill-defined region. Thus, depending on the quality of the collection, the amount of useful information obtained from herbarium specimens may be quite limited.

Data management systems may also help in the day-to-day organization of herbarium operations. For example, accession numbers (which may be scanned with bar codes) may automatically keep track of both outgoing loans and incoming loan returns.

In addition to in-house data information systems, it is now commonplace to electronically connect directly to the

databases of other herbaria. Many of the major herbaria have an on-line connection that allows others to access this information over the internet, including searchable Web pages.

Finally, data management of natural collections has become invaluable in biodiversity studies. The data information system allows for the tabulation of presence, range, and distribution of taxa, especially important for studying rare or endangered species or sensitive habitats.

FLORAS AND MONOGRAPHS

Herbaria are particularly essential in two important activities in plant systematics: floristics and monographic treatments. **Floristics** is the documentation of all plant species in a given geographic region. Floristics may also entail documentation of plant communities and abiotic factors as well. Floristic studies may be published in taxonomic journals or may result in the publication of a **flora** or plant **manual** of a given region, such as *Flora of North America*. Floristic studies are vital in the documentation of plant biodiversity.

A **monograph** is a detailed taxonomic study of all species and infraspecific taxa of a given taxonomic group, generally a genus or family. Unlike floristic studies, the goal of which is to document taxa for a given area, monographic treatments focus on a particular taxonomic group, over its entire geographic range. For example, see the Systematic Botany Monographic series (American Society of Plant Taxonomy, <http://www.sysbot.org>).

REVIEW QUESTIONS

1. What is an herbarium?
2. What is the function or purpose of herbaria?
3. What is the name and most recent version of the reference book that lists the names, acronyms, and details of herbaria worldwide?
4. What is an herbarium specimen?
5. What are the characteristics of an herbarium specimen, including the standard size (in the United States)?
6. Describe a standard format and list the information that is contained in an herbarium label.
7. If a plant specimen is divided among two or more herbarium sheets, how is the herbarium label written?
8. Describe the procedure for mounting plants on herbarium paper.
9. What are two type of glues used for the above? How do they differ?
10. What is an herbarium curator and what are his/her duties?
11. What is an accession number and what is its function?
12. What is a genus folder?
13. Why are many genus folders color-coded?
14. How are herbarium specimens typically stored?
15. Review the procedures for handling herbarium specimens.

16. What is an annotation label?
17. What are the different types or designations on annotation labels?
18. Review the procedures for removing material from an herbarium specimen.
19. How may removed material be reconstituted for observations?
20. What is an herbarium loan and what is its purpose?
21. How may insects be controlled in herbaria?
22. What is a data information (database) system?
23. What types of data manipulation may be done with data information systems?
24. How might data information systems be valuable in conservation biology?
25. What is the difference between a flora (or floristic study) and a monograph, and what are examples of each?

EXERCISES

1. Obtain a list of plant species from your instructor, including a lycopod, fern, conifer, monocot, and eudicot. Become familiar with the system of classification of your herbarium (or a herbarium that you visit). Locate and remove the genus folder for this species. Carefully transfer the genus folder to an open-space table, remove one specimen, and study it. (Be sure to handle the herbarium sheets correctly.) Note how the specimen is attached to the herbarium paper. Also note the label, accession number (possibly on a bar code), and annotation labels (if present). Write down the collector, collection number, general region/locality where the specimen was collected, herbarium acronym (see Index Herbariorum), and accession number (e.g., Smith 762; NY 1120387). Do this for each of the species on the list.
2. When finished with each specimen, replace it in correct order (usually alphabetical by species) within the genus folder. If permitted, refile the genus folder in its correct location in the herbarium cabinet, both by taxonomy and (if used) color coding by region. [Caution: Be sure to check yourself carefully! If misfiled, specimens could be lost for some time.]
3. Access an herbarium database (e.g., U.C. Berkeley's Jepson Herbarium or the New York Botanical Garden herbarium) by doing a search on the Web. Look up several species. If available, get information for type specimens and/or images of herbarium specimens.
4. Do a search on a particular herbarium database for a given species or genus. Download all information to a spreadsheet. Optional: Determine the latitude and longitude for these specimens, either directly from what was recorded on the label or by tracking down localities using available hard-copy or computerized topographic maps. From your list of species with latitude/longitude information, generate a map, showing the place of collection of each specimen as a dot. Do you notice any correlation with region (e.g., elevation) or habitat?
5. Check Index Herbariorum on-line (see below) for several large herbaria (e.g., K, MO, NY, UC, US) and determine (a) the curator's name, address, e-mail; (b) the number of specimens currently accessioned; (c) the general types of collections in the herbarium; (d) any type specimens in the herbarium.
6. Obtain and study an example of a flora and one of a monograph. Copy an example entry from each, indicating how they differ.

REFERENCES FOR FURTHER STUDY

- Allkin, R. and F. A., Bisby (eds.). 1984. Databases in Systematics. Academic Press, London. Systematics Association special volume, no. 26. Proceedings of an International Symposium held in Southampton, UK.
- Forman, L., and D. M., Bridson (eds.). 1992. The Herbarium Handbook, rev. ed. Royal Botanic Gardens, Kew, UK.
- Holmgren, P. K., N. H. Holmgren, and L. C. Barnett, eds. 1990. Index Herbariorum, Part I: The Herbaria of the World, 8th ed. New York Botanical Garden, New York. [For online searches, see <http://www.nybg.org/bsci/ih/ih.html> (continuously updated).]
- Radford, A. E., W. C. Dickison, J. R. Massey, and C. R. Bell. 1974. Vascular Plant Systematics. Harper & Row, New York.

APPENDIX 1

PLANT DESCRIPTION

WRITING A PLANT DESCRIPTION

The following list of characters can serve as the basis for a detailed plant description. The basic form of the description is to list the plant organ (noted in **bold** in the character list below), followed by a listing of all character states that apply for that plant organ, with each character state separated by commas. Note that, for any particular species, not all characters will apply; these are simply omitted. Also note that some characters are listed with multiple character names, e.g.,

Sepal/Calyx lobes/Outer tepals. This is designed as a guide, with the intention that only one of these three will be used, depending on whether the outer whorl of the perianth consists of distinct sepals (**Sepal** used), of fused sepals (**Calyx lobes** used), or of tepals (**Outer tepals** used).

There are different styles in writing a detailed plant description. Some use a telegraphic style, e.g., *Leaves simple, sessile, whorled, ovate, entire, glabrous*. This style is common in floras, where space for text may be at a premium. Other descriptions use complete sentences, e.g., *Leaves are simple, sessile, whorled, ovate, entire, and glabrous*. The use of the *at the beginning* of a sentence is optional, as in *The leaves are simple, sessile, whorled, ovate, entire, and glabrous*.

Some general suggestions are as follows:

1. Be sure to *only* list the plant organs (and list only once), followed by the character states that apply to that plant organ. The major plant organs are sometimes placed in **bold** text to highlight them. *Do not* list the specific character names, unless a clarification is needed. Examples:
Do write: Flowers are bisexual, actinomorphic, pedicellate, 1.5–2.2 cm long (including pedicel) ...
[Flowers refers to the plant organ; all other terms are character states.]

Do not write: Flower sex is bisexual, symmetry is actinomorphic, attachment is pedicellate, length is 1.5–2.2 cm [Sex, symmetry, attachment, and length are characters and should not be listed.]

However, *do write: Leaf blades are elliptic, serrate, rounded at base, obtuse at apex*. [Rounded and obtuse could refer to either of the characters base or apex, so these characters should be listed for clarification.]

2. Description of the major organs may be written in the singular or plural form, but the latter should be used only if more than one such organ occurs in an individual. If only one organ occurs per individual, the singular should be used.
Do write: Leaves are trifoliolate, alternate, ... or *The leaf is trifoliolate, alternate, ...* if there are multiple leaves.
Do write: The inflorescence is a solitary raceme, ... if there is a single raceme per individual.
3. Always use metric for plant or plant organ heights, lengths, and widths. Always abbreviate these: mm for millimeters, cm for centimeters, dm for decimeters, m for meters. Use mm and cm for smaller structures, dm or m for larger. Use the appropriate unit of measure to avoid values less than 1, if possible. (E.g., write 2.5 mm instead of 0.2–0.5 cm.) Always place a 0 before a decimal point, as in 0.5 mm. Be clear about what you are describing. Examples:
Do write: Flowers are 0.5–1.3 mm long (excluding pedicel), 2–3 mm wide when fully opened.
Do not write: Flowers are .5–1.3 mm.
4. For characters that are variable, either list the range of variation (e.g., **Leaves** oblanceolate to narrowly elliptic, crenate to dentate ...) or list the most common morphology and in brackets list the exceptions (e.g., **Leaves** trifoliolate [rarely pinnate with 5 leaflets] or **Leaves** 4–7 [2.5–10] cm long ...).

COMPLETE MORPHOLOGICAL CHARACTER LIST

[Available as download from Website; Note: Not all characters apply to a given taxon; add characters for specialized structures.]

Species/Infraspecies Name (with authorship) [Common Name]: _____

Family: _____ **Native locality:** _____

Plant Habitat: _____

Plant Duration: _____

Plant Sex if not hermaphroditic: _____

Plant Habit: _____

Plant Height: _____

Root Type: _____

Root Origin (e.g., primary, adventitious): _____

Underground Stem Type if specialized: _____

Underground Stem Branching Pattern: _____

Underground Stem Size: _____

Aerial Stem Habit: _____

Aerial Stem Branching Pattern: _____

Bark Type: _____

Bark Lenticels presence/shape: _____

Twig Surface/Shape: _____

Twig Lenticel presence/shape: _____

Twig Shape/Cross-Sectional Outline: _____

Pith Type: _____

Pith Cross-Sectional Outline: _____

Fruit Scar presence/shape: _____

Leaf Scar Size/Shape: _____

Vascular Bundle Scar Number/Pattern: _____

Stipule Scar presence: _____

Stipule Scar Position/Shape if present: _____

Terminal Bud Scale Scars presence/absence: _____

Bud Type: _____

Bud Orientation: _____

Bud Shape/Size: _____

Bud Position: _____

Bud Scale Arrangement: _____

Bud Scale Surface/Texture: _____

Thorns if present: _____

Spines if present: _____

Prickles if present: _____

Spur Shoot if present: _____

Leaves/Leaf Number if unusual: _____

Leaf Type: _____

Leaf Length/Width: _____

Leaf Attachment: _____

Leaf stipule presence: _____

Leaf Duration: _____

Leaf Position if not cauline: _____

Leaf Arrangement: _____

Leaf Orientation if discrete: _____

Leaf Posture if discrete: _____

Petiole Shape: _____

Petiole Color: _____

Petiole Length: _____

Petiole Orientation: _____

Stipule Shape: _____

Stipule Surface adaxial: _____

Stipule Surface abaxial: _____

Stipule Length: _____

IF LEAVES SIMPLE:

Leaf Blade Color if unusual: _____

Leaf Blade Shape: _____

Leaf Blade Length: _____

Leaf Blade Width: _____

Leaf Blade Base: _____

Leaf Blade Margin: _____

Leaf Blade Apex: _____

Leaf Blade Apical Process: _____

Leaf Blade Division: _____

Leaf Blade Venation: _____

Leaf Blade Surface adaxial: _____

Leaf Blade Surface abaxial: _____

Leaf Blade Texture: _____

IF LEAVES COMPOUND:

Leaf Outline Shape: _____

Rachillae Number if decomound: _____

Leaflets Number if not very large: _____

Leaflet Arrangement: _____

Leaflet Blade Shape: _____

Leaflet Blade Attachment: _____

Leaflet Blade Color if unusual: _____

Leaflet Blade Length: _____

Leaflet Blade Width: _____

Leaflet Blade Base: _____

Leaflet Blade Margin: _____

Leaflet Blade Apex: _____

Leaflet Blade Apical Process: _____

Leaflet Blade Division: _____

Leaflet Blade Venation: _____

Leaflet Blade Surface adaxial: _____

Leaflet Blade Surface abaxial: _____

Leaflet Blade Texture: _____

Petiolule Shape: _____

Petiolule Color: _____
 Petiolule Length: _____
Stipel presence: _____
 Stipel Shape: _____
 Stipel Surface adaxial: _____
 Stipel Surface abaxial: _____
 Stipel Length: _____

Inflorescence Position: _____
 Inflorescence Bract presence: _____
 Inflorescence Type: _____
 Inflorescence Length: _____
 Inflorescence Width: _____
 Inflorescence Branch Orientation: _____
 Inflorescence Sex: _____
 Inflorescence Axis Surface: _____

Flower Sex: _____
 Flower Bract presence: _____
 Flower Length minus pedicel: _____
 Flower Width minus pedicel: _____
 Flower Arrangement: _____
 Flower Orientation: _____
 Flower Posture: _____
 Flower Symmetry overall: _____
 Flower Attachment: _____

Pedicel if present Length: _____
 Pedicel if present Shape if unusual: _____

Bracts/Bractlets No (note inflorescence vs. flower): _____
 Bracts Position: _____
 Bracts Length: _____
 Bracts Color if unusual: _____
 Bracts Attachment: _____
 Bracts Shape: _____
 Bracts Base: _____
 Bracts Margin: _____
 Bracts Apex: _____
 Bracts Apical Process: _____
 Bracts Division: _____
 Bracts Venation if unusual: _____
 Bracts Texture if unusual: _____
 Bracts Surface adaxial: _____
 Bracts Surface abaxial: _____

Receptacle Size if unusual: _____
 Receptacle Shape if evident: _____

Hypanthium presence: _____
Hypanthium Shape: _____
 Hypanthium Length: _____
 Hypanthium Width: _____

Perianth Cycly: _____
 Perianth Arrangement if not whorled: _____

Perianth Type (if homochlamydeous): _____

Calyx/Outer Tepals Aestivation: _____

Calyx/Outer Tepals Fusion: _____

Calyx/Outer Tepals Symmetry: _____

Calyx/Outer Tepals Length: _____

Calyx/Outer Tepals Color if not green: _____

Calyx/Outer Tepals Surface adaxial: _____

Calyx/Outer Tepals Surface abaxial: _____

Calyx/Outer Tepals Venation if unusual: _____

Calyx/Outer Tepals Texture if unusual: _____

Sepal/Calyx Lobes/Outer Tepals Merosity: _____

Sepal/Calyx Lobes/O.T. Length: _____

Sepal/Calyx Lobes/O.T. Shape: _____

Sepal/Calyx Lobes/O.T. Base: _____

Sepal/Calyx Lobes/O.T. Margin: _____

Sepal/Calyx Lobes/O.T. Apex: _____

Sepal/Calyx Lobes/O.T. Apical Process: _____

Corolla Type (if dichlamydeous): _____

Corolla/Inner Tepals Aestivation: _____

Corolla/Inner Tepals Fusion: _____

Corolla/Inner Tepals Cycly if not uniseriate: _____

Corolla/Inner Tepals Color: _____

Corolla/Inner Tepals Symmetry: _____

Corolla/Inner Tepals Length: _____

Corolla/Inner Tepals Surface: _____

Corolla/Inner Tepals Venation if unusual: _____

Corolla/Inner Tepals Texture if unusual: _____

Petal/Corolla Lobes/Inner Tepals Merosity: _____

Petal/Corolla Lobe/I.T. Shape: _____

Petal/Corolla Lobe/I.T. Base: _____

Petal/Corolla Lobe/I.T. Margin: _____

Petal/Corolla Lobe/I.T. Apex: _____

Petal/Corolla Lobe/I.T. Length: _____

Petal/Corolla Lobe/I.T. Orientation: _____

Petal/Corolla Lobe/I.T. Posture: _____

Stamens (Androecium) Cycly: _____

Stamens (Androecium) Merosity: _____

Stamen Type: _____

Stamen Attachment: _____

Stamen Arrangement: _____

Stamen Position: _____

Stamen Insertion if applicable: _____

Stamen Fusion: _____

Staminodes if present No: _____

Staminodes if present Pos: _____

Staminodes if present Size: _____

Staminodes if present Shape: _____

Filament Shape/Color: _____

Filament Length: _____

Anthers Attachment: _____

Anther Type: _____
 Anther Dehiscence Type: _____
 Anther Dehiscence Direction: _____
 Anther Color: _____
 Anther Length: _____
 Anther Shape: _____
 Anther Thecae Arrangement: _____
Connective Morphology if unusual: _____
Pollen color: _____
Gynoecium Fusion: _____
Perianth Androecial Position: _____
Ovary Position: _____
 Ovary Attachment if not sessile: _____
 Ovary Color: _____
 Ovary Length: _____
 Ovary Shape: _____
 Ovary Surface: _____
Styles Number per pistil: _____
 Style Position: _____
 Style Shape/Color: _____
 Style Disposition/Length: _____
Stigmas Number: _____
 Stigmas Position: _____
 Stigmas Shape: _____
 Stigmas Surface: _____
Nectaries presence/absence: _____
 Nectary Type/Position: _____
Carpels Number: _____
 Median Carpel Position relative to stem axis: _____
Locules Number: _____
Placentation: _____
Placenta Shape/Position if unusual: _____
Ovules Number per carpel: _____
 Ovule Type: _____
 Ovule Position: _____
Fruit Type: _____
 Fruit Color: _____
 Fruit Shape: _____
 Fruit Length/Width: _____
 Fruit Surface: _____

Seed Color: _____
 Seed Shape: _____
 Seed Length: _____
 Seed Surface: _____
Funiculus Length: _____
 Funiculus Shape: _____
Aril presence: _____
 Aril Size: _____
 Aril Shape: _____
 Aril Position: _____
Seeds Nutritive Tissue: _____
Embryo Type Size/Shape/Position: _____
Cotyledon Position: _____
Radicle Position: _____
Seedling Type: _____

FLORAL FORMULA:

P _____ A _____ G
 or K _____ C _____ A _____ G

Note: List number of parts after each symbol:

P = # perianth parts or tepals (outer + inner whorls)
 or **K** = # sepals or calyx lobes **C** = # petals or corolla lobes
A = # stamens of androecium (outer + inner whorls)
G = # carpels of gynoecium (add ovary position)
 () = fusion of parts [] = rare numbers of parts
 Optional:
K_z = zygomorphic calyx; **C_z** = zygomorphic corolla; etc.

E. g., **K** (5) **C_z** (5) **A** 5 [4] **G** (2), inferior
 = calyx synsepalous with 5 lobes
 corolla zygomorphic, sympetalous with 5 lobes
 stamens 5, rarely 4, distinct, in one whorl
 gynoecium syncarpous, carpels 2, ovary inferior
 E.g., **P** 3+3 **A** 3+3 **G** 3, superior
 = perianth apotepalous with 3 outer and 3 inner tepals
 stamens 6, distinct, in two whorls: 3 outer + 3 inner
 gynoecium apocarpous, carpels (pistils) 3, ovaries
 superior

PLANT DESCRIPTION EXAMPLE

Tecomaria capensis (Thunb.) Spach, Cape-Honeysuckle (native to S. Africa). Bignoniaceae.

Plant a shrub, up to ca. 5 m tall. **Root** a woody taproot with numerous lateral roots. **Stems** (aerial) highly and sympodially branched by abortion of terminal inflorescence meristems, branches basally inclined. **Bark** brown, smooth to minutely furrowed, lenticels orbicular to vertically elliptic with raised borders, ca. 1-2 mm wide. **Twigs** terete, minutely puberulent. **Pith** solid, circular in outline. **Fruit scars** (of infructescence) raised, circular, typically at junction of two, lateral branches. **Leaf scars** slightly raised below, orbicular with truncate apex. **Vascular bundle scar** U-shaped. **Buds** in leaf axils small (ca. 2 mm long), with outer two scales in a plane tangential to stem axis, scales valvate, lance-ovate and strongly cup-shaped, densely pubescent; terminal buds naked, elongate, to 5 mm long. **Leaves** 10-12 cm long, imparipinnate, petiolate, exstipulate, evergreen, cauline, opposite-decussate, divergent to inclined, and planar to recurved. **Petiole** green, terete to canaliculate, 1-3 cm long. **Leaf outline** elliptic. **Leaflets** 9 [11], opposite. Lateral leaflets elliptic to widely elliptic, subsessile, 15-17 mm long, 10-14 mm wide, base attenuate to obtuse, sometimes oblique, margin usually proximally entire and serrate to crenate distally, apex acuminate (caudate), tip minutely mucronulate, Apical leaflet widely elliptic, usually petiolulate (**petiolule** green, narrowly winged, 3-13 mm long) 24-30 mm long, 15-20 mm wide, cuneate, entire at base and distally serrate to crenate, acute to acuminate, mucronulate. All leaflets pinnate-netted, midvein and secondary veins sunken above and raised below, mostly glabrous but with arachnose trichomes near abaxial vein junctions, mesophyllous. **Inflorescence** a terminal thyrse with several bracteate units of simple dichasia or of solitary flowers, the latter often with two abortive, lateral flower buds or with two sub-basal bracts (indicative of a vestigial dichasium). **Flowers** perfect, ca 50 mm long, ca 25 mm wide, opposite, appressed, recurved, zygomorphic, pedicellate. **Pedicel** ca 7 mm long, terete. **Bract** 1 subtending each unit of inflorescence, 1-5 mm long, lanceolate, mucronulate. **Bractlets** 2, sub-basal, subtending lateral flowers if simple dichasium present. **Perianth** biseriata, dichlamydeous. **Calyx** synsepalous, actinomorphic, ca 5 mm long. **Calyx lobes** acute, mucronulate, ca 1 mm long. **Corolla** sympetalous, orange, zygomorphic, salverform-bilabiate with enlarged throat, ca 45 mm long, recurved, inner surface pubescent. **Corolla lobes** 5 (2 posterior, 2 lateral, and 1 anterior), oblong to elliptic, apices rounded to emarginate, 7-12 mm long, 5-7 mm wide, inclined to divergent and recurved relative to floral axis. **Stamens** 4 fertile, uniseriate, filamentous, epipetalous, didynamous, alternipetalous, exserted, apostemonous. **Staminodium** 1, medio-posterior, reduced, up to ca 10 mm long. Filaments (of fertile stamens) terete, yellow-orange, 35-40 mm long. **Anthers** versatile, basifixed, longitudinally and introrsely dehiscent (downwardly dehiscent at maturity), ca. 3 mm long, thecae divergent. **Pollen** orange. **Gynoecium** syncarpous. **Perianth/Androecial position** hypogynous. **Ovary** superior, green, 4-5 mm long, narrowly obloid, glabrous. **Carpels** 2. **Locules** 2. **Placentation** parietal-axile. **Ovules** many. **Styles** 1, terminal, apically recurved, purple-brown. **Stigmas** 2, ovate, divergent to appressed. **Nectary** dark maroon, doughnut-shaped, surrounding ovary base. **Fruit** a brown loculicidal capsule (with persistent replum), narrowly oblong, up to ca. 1 cm wide and 6 cm long. **Seeds** flat, with surrounding, yellowish, translucent wing, ca. 15 mm long and 8 mm wide (including wing), seed body roughly orbicular, ca. 6 mm in diameter.

FLORAL FORMULA: K (5) C (5) A 2+2+1_{staminode} G (2), superior.

This Page Intentionally Left Blank

APPENDIX 2

BOTANICAL ILLUSTRATIONS

Botanical illustration is the preparation and presentation of line drawings or paintings of plants and plant parts. Although photography is perhaps the primary medium of image documentation (and should be encouraged in field and laboratory classes), illustration is an important component of plant systematics studies and is generally required in publications to document features that are described. Illustrations are also important didactic tools in that they promote careful and complete observation of features; going through the process of drawing often helps an investigator to see more and in greater detail.

Illustrations in plant systematic research are almost always line drawings. Line drawings typically begin with a pencil drawing, which alone may be sufficient for personal observations and data collection. For publication-quality illustrations, pencil drawings must be retraced in black ink and are typically stippled for a shaded, three-dimensional appearance (see Figure A2.1). Alternatively, line drawings with shading may be computer generated using graphics software (below).

Basic supplies needed for pencil line drawings include a 2H or 3H pencil, high-quality eraser, and drawing paper (ranging from generic white to artist's drawing paper). A clipboard or artist's drawing board is handy to secure the paper. Drawings may be made free-hand. More precise drawings can be made using a camera-lucida/drawing tube device. Such an optical device allows one to see a double image of the object to be drawn plus the hand and pencil, allowing quick and easy tracing of object features. Pencil drawings should consist of outlines of whole organs/parts and of individual components. Lines should be crisp and precise. Minimize shading; shade only when objects are darker and then only lightly with pencil.

Drawings should be labeled clearly, including (a) name of taxon and documentation of material (e.g., reference to

a voucher specimen), and (b) names of structures, indicating all pertinent terms, with lines or arrows leading from the structure to the term label. A metric magnification scale bar should accompany each drawing. A scale bar (e.g., 5 mm) is much preferred over a simple magnification listing (e.g., $25\times$) because the scale bar remains to scale with any subsequent enlargement or reduction of the drawing. The size of drawings should be planned for a final size reduction (after inking and stippling) of approximately 50–75%, which yields a better final product.

WHAT TO DRAW

A complete illustration of plant morphology may include drawings of the following (Figure A2.1): (a) the whole plant at low magnification, showing the plant habit, branching pattern and overall form; (b) one or more leaves, showing leaf attachment to the stem; (c) a flower in front, oblique, and/or side view; (d) a flower in median longitudinal (sagittal) section; (e) androecium, especially stamen/anther close-up; (f) gynoecium; (g) ovary longitudinal and/or cross-section; and (h) close-ups of other floral parts of significance.

In addition to drawing real views of plant parts, diagrams may be drawn to illustrate the relative position of parts. **Floral diagrams** (Figures A2.1H, A2.2) show the relative position, aestivation, and fusion of perianth parts, stamens, and pistil(s)/carpels. For showing the relative position of floral parts, a diagram of the floral axis is typically indicated at the top of the drawing, corresponding to the posterior side of the flower. The floral diagram begins at the center of the flower. On a sheet of paper, draw the pistil(s) as appearing in cross-sectional view, carefully denoting ovary wall, septa, ovules, and placentation. Next, stamens are drawn surrounding the gynoecium. Stamens are drawn as anther cross-sections (internal contents such as microsporangia usually not denoted),

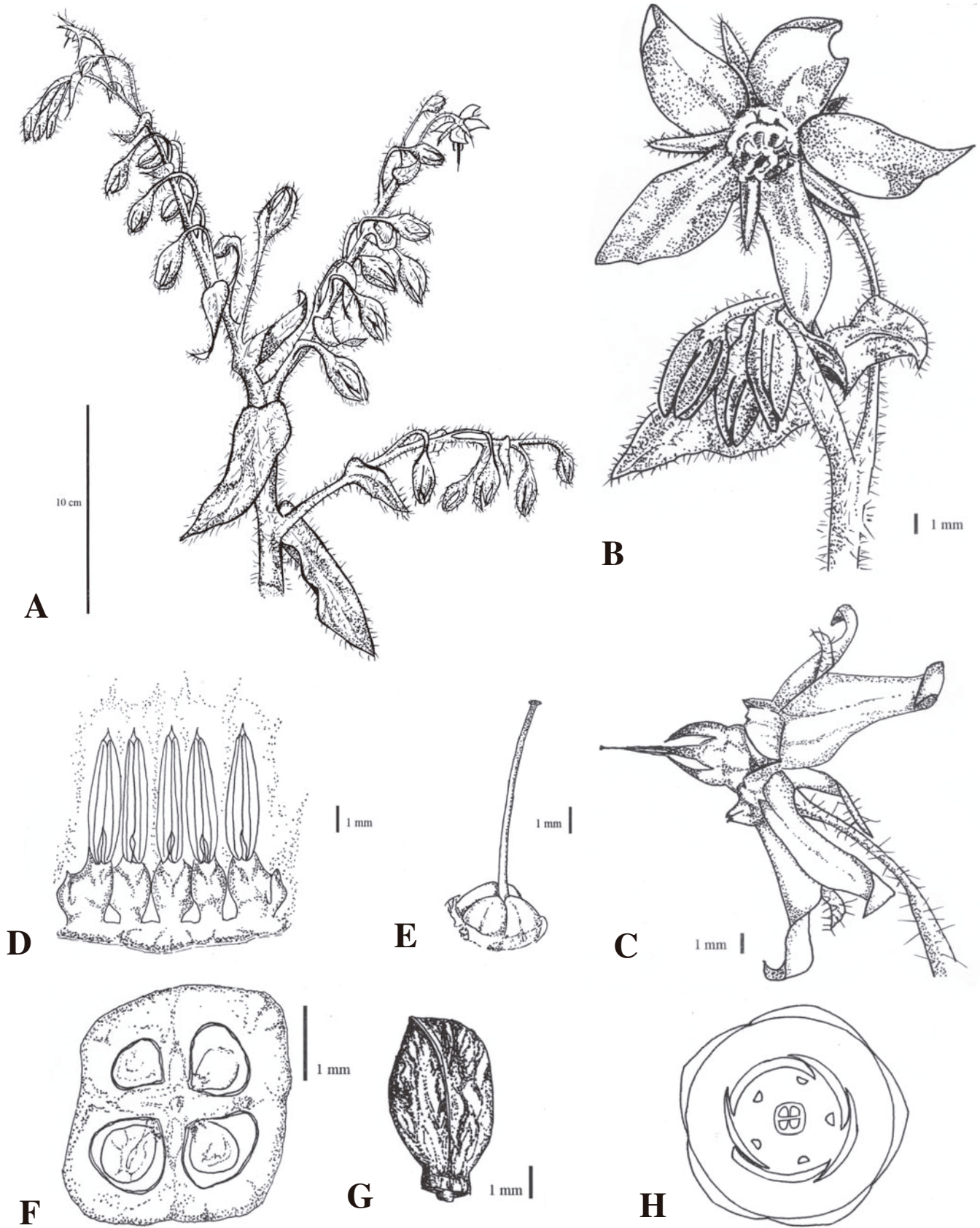


FIGURE A2.1 Example of illustrations, showing stippling and scale bars. *Borego of cinalis*. **A**. Whole in inflorescence and leaves. **B**. Flower, face view. **C**. Flower, side view. **D**. Androecium, spread position, adaxial view. **E**. Gynoecium. **F**. Ovary cross-section, base of ovary. **G**. Fruit. **H**. Floral diagram. (Contributed by Dinna Estrella, student in Plant Systematics class.)

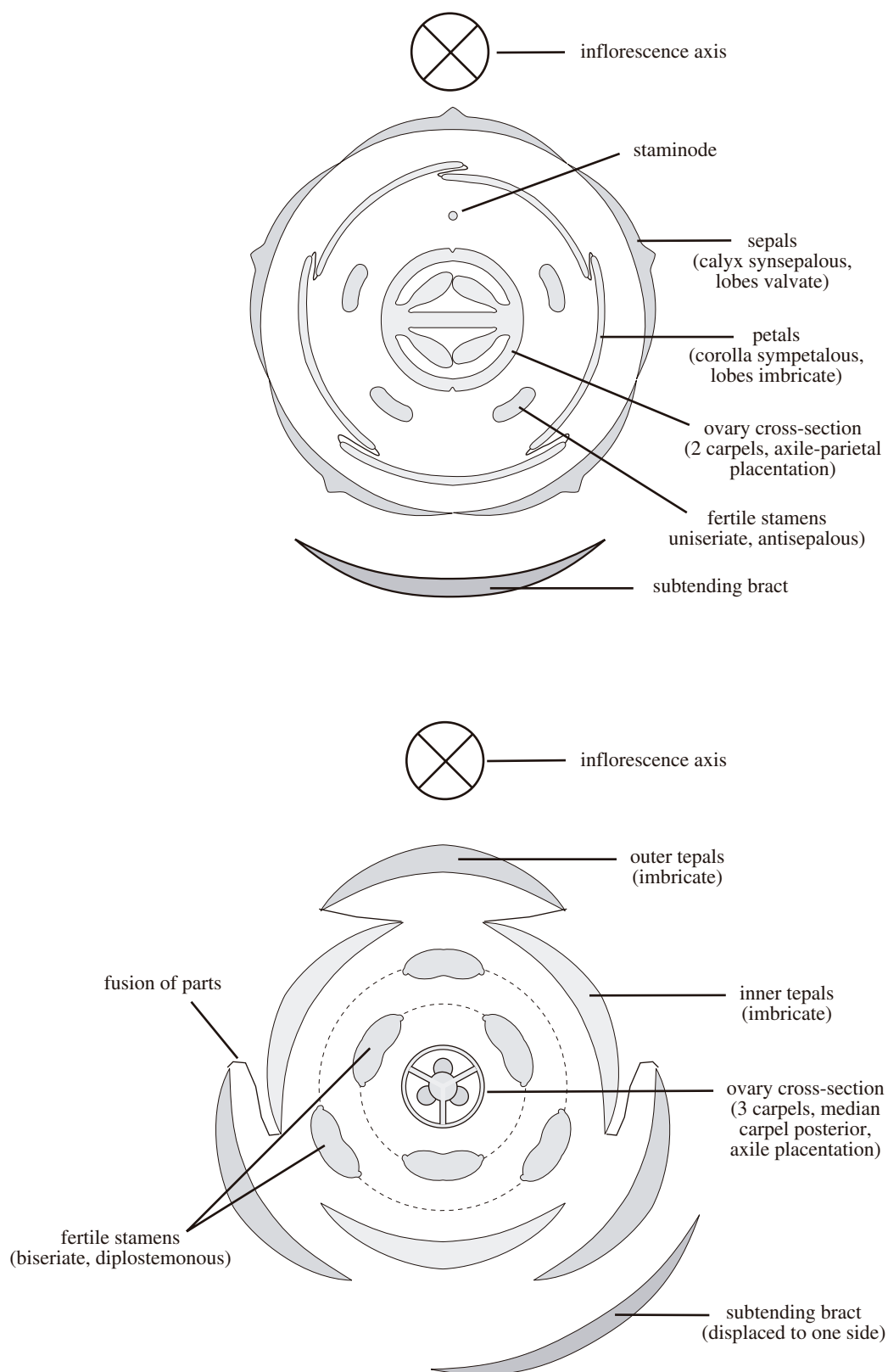


FIGURE A2.2 Examples of floral diagrams.

with the direction of dehiscence indicated. Petals (or corolla lobes) and sepals (or calyx lobes) are drawn surrounding the gynoecium and androecium. These perianth parts are drawn as in cross-section, with careful attention to relative position and aestivation. Bracts are drawn similar to perianth parts, at their position of attachment. Any connation or adnation of parts is drawn either as organs contacting one another or as

lines drawn between fused structures. If the flower is more or less erect, a circle is drawn that indicates the relative position of the axis to which the flower is attached. If the flower is horizontal in orientation, parts may be drawn as if the opening of the flower is facing the observer. If flowers are unisexual, male and female flowers should be drawn separately, of course.

REFERENCES FOR FURTHER STUDY

- Cook, C. D. K. 1998. A quick method for making accurate botanical illustrations. *Taxon* 47: 317-380.
- Holmgren, N. H., and B. Angell. 1986. *Botanical Illustration: Preparation for Publication*. The New York Botanical Garden, Bronx, New York.

APPENDIX 3

SCIENTIFIC JOURNALS IN PLANT SYSTEMATICS

JOURNALS

CARD CATALOG	JOURNAL	WEB SITE (at time of publication)
QK 149 A4	Aliso	http://www.rsabg.org/aliso.html
QK 1 B345	American Journal of Botany	http://www.amjbot.org
QK 1 M6	Annals of the Missouri Botanic Garden	http://www.mbgpress.org/
QK 1 A8	Australian Journal of Botany	http://www.publish.csiro.au/nid/65.htm
QK 1 L52	Botanical Journal of the Linnean Society	http://www.blackwellpublishing.com/jnl_default.asp
QK 1 B6	Botanical Review	http://sciweb.nybg.org/science2/BotanicalReview.asp
QK 1 B875	Brittonia	http://www.nybg.org/bsci/brit/
QK 1 C3	Canadian Journal of Botany	http://pubs.nrc-cnrc.gc.ca/cgi-bin/rp/rp2_desc_e?cjb
QK 1 F4	Fieldiana (Botany)	http://www.fortsasbooks.com/publish.htm
QK 149 F73	Fremontia	http://www.cnps.org/publications
QK 658 G72	Grana	http://www.tandf.co.uk/journals/titles/00173134.asp
QK 1 B3	International Journal of Plant Sciences	http://www.journals.uchicago.edu/IJPS/home.html
Q 11 E4	Journal of the Elisha Mitchell Scientific Society	http://www2.ncsu.edu/ncas/jemsstab.html
QK 1 M3	Madroño	http://www.calbotsoc.org/madrono.html
QH 506 M664	Molecular Biology & Evolution	http://mbe.oupjournals.org
QH 367.5 M56	Molecular Phylogenetics & Evolution	http://www.elsevier.com
QK 1 N75	Nordic Journal of Botany	http://www.nathimus.ku.dk/bot/b_nordic.htm
QK96 .N68	Novon	http://www.mbgpress.org/
QK 1 P3A26	Pollen et Spores	
Q 11 N26	Proceedings of the National Academy of Sciences of the United States of America (PNAS)	http://www.pnas.org
QK 1 S57	Smithsonian Contributions to Botany	http://www.sipress.si.edu/the_press/press_main.html
QL352 .S92	Systematic Biology	http://systbiol.org
QK 95 597	Systematic Botany	http://www.sysbot.org/
QK 93 T3	Taxon	http://www.botanik.univie.ac.at/iapt/s_taxon.php

EXERCISES

1. Examine several of the botanical journals listed above. For each of the ones selected by your instructor: (a) record the general types of papers for which it specializes by reading the journal's mission and by perusing the index or Web site (above); (b) cite one specific article (of interest to you) *that deals with plant systematics* (not other fields such as physiology or ecology). For the latter, copy the citation format used in that particular journal.
2. Of the plant systematics articles that you recorded above, select the one of greatest interest to you and make a photocopy of it. Read the abstract of this article thoroughly. Then look over all the figures and tables. Next, skim through the introduction, materials and methods, and discussion/conclusion. Be prepared to give a 5-minute oral presentation (with projected tables, figures, or illustrations) on this article.
3. Write the author(s) in charge of reprints for the above and request a reprint to be mailed to you. (Note: It's best to choose a recent article for this, as reprints may not be available for older ones.) Some reprints are now available electronically, usually as .pdf files. Hand this in to your instructor when it arrives.
4. Search the International Plant Names Index (<http://www.ipni.org/index.html>). Find and record the full citation (including author, journal, date, etc.) of *any one* of the species that were studied in the article you chose for discussion.
5. Optional: Do a literature search on a particular item in plant systematics. It may be of taxon or topic. (Note: Consult your library as to how to access literature databases; also try <http://www.scholar.google.com> for general scientific literature.)

GLOSSARY OF TERMS

The following is a glossary of terms used in all chapters of the book. For the terms from Chapter 9 (Plant Morphology), the character to which a character state belongs is noted in parentheses following the definition. Symbols used are: Abbr = abbreviation; Adj = adjective; Cf = compare; Pl = plural; Syn = synonym.

- ! Abbreviation for the confirmation of a name in an annotation label.
- abaxial** Surface most distant or away from the axis, the lower or outer surface of organ. Syn: *dorsal*. (position)
- ABC model** A model of floral development, in which gene products of the so-called A, B, and C classes combine to produce the four major floral organs: sepals, petals, stamens, and carpels.
- acaulescent** Lacking an above-ground stem other than the inflorescence axis. (stem habit)
- accession number** A number assigned to each specimen placed into a permanent herbarium collection.
- accessioning** The assignment of a number to all new specimens placed into a permanent herbarium collection.
- accessory bud** Bud(s) lateral to or above axillary buds. (bud type)
- accessory part** A portion of the mature fruit that is not directly derived from the ovary or ovaries, may include bract(s), stem axes, receptacle, hypanthium, or perianth. (fruit part)
- accrescent** Parts persistent and continuing to grow beyond what is normal or typical, e.g. calyx of *Physalis*, Solanaceae. (duration)
- acetolysis** A standard acid treatment used to dissolve all but the exine of pollen grains in order to better observe pollen wall structure with the light microscope.
- achene** A one-seeded, dry, indehiscent fruit with seed attached to pericarp at one point only, e.g. unit fruits of sunflowers and other Asteraceae. (fruit type)
- achenechetum** An aggregate fruit of achenes, e.g. *Fragaria* (strawberry), in which the achenes are on the surface of accessory tissue, an enlarged, fleshy receptacle. (fruit type)
- achlamydeous** Lacking a perianth. (perianth merosity)
- achlorophyllous** Lacking chlorophyll/chloroplasts.
- acicular** Needlelike, often round in cross-section, with margins straight and parallel, length:width ratio >12 : 1. (shape)
- acrocaulis** Positioned at the apex of the stem. (position)
- acrocidal capsule** A capsule dehiscing by means of apical slits. (fruit type)
- acrodromous** With two or more primary veins or strongly developed secondary veins running in convergent arches toward the leaf apex but not recurved at the base. (leaf venation)
- actinodromous** With three or more primary veins diverging from one point, inclusive of ternate or palmate venation. (leaf venation)
- actinomorphic** Radially symmetrical, with 3 or more planes of symmetry. (symmetry)
- aculeate** With prickles, sharp nonspine, nonthorn appendages. Syn: *prickly*. (epidermal excrescence)
- acuminate** Apical margins abruptly incurved (concave), the apical intersection angle <45°. (apex)
- acute** Apical margins approximately straight, the intersection angle 45° 90°. (apex)
- adaptation** a structure or feature that performs a particular function and which results in increased survival or reproduction.
- adaxial** Surface toward or nearest the axis, the upper or inner surface of organ. Syn: *ventral*. (position)
- adherent** With unlike parts joined, but only superficially and easily separable. (fusion)
- adnate** With unlike parts integrally fused, not easily separable. (fusion)
- advanced** *Derived*.
- adventitious roots** A root arising from an organ other than a root, usually from a stem. (root type)
- adventive embryony** Development of an embryo from a cell of the surrounding tissue, such as megasporangial or integument tissue.
- aerial roots** Adventitious roots that absorb moisture and minerals from the air or runoff, common in some epiphytic plants, e.g. of Araceae and Orchidaceae. (root type)
- aestival** Appearing in summer. (periodicity)
- aestivation** Referring to position, arrangement, and overlapping of floral perianth parts. (perianth aestivation)
- affinis** Implying, within a taxon name, some type of close relationship, presumably an evolutionary relationship, but also that the specimen differs from the cited taxon in some way, e.g. beyond the described range of variation for one or more characters. Abbr: *aff*.
- agamosperry** The production of seeds without fertilization.
- aggregate fruit** A fruit derived from two or more pistils (ovaries) of one flower. (fruit type)

- albuminous** *Endospermous*. (seed endosperm type)
- albuminous cell** A parenchyma cell associated with a sieve cell, derived from a different parent cell than is the sieve cell.
- aleurone grain** Granular protein deposits in plant cells, functioning as storage compounds; a type of ergastic substance. Syn: *proteinoplast*.
- alignment** The process of matching homologous nucleotide positions of two or more sequences of DNA in order to code the data for phylogenetic or other types of analysis.
- allautogamy** Having both outcrossing and selfing flowers. Adj: *allautogamous*.
- alogamy** *Outbreeding*. Adj: *alogamous*.
- allopolyploidy** Polyploidy occurring between different species.
- allozyme** One of two or more different molecular forms of an enzyme, corresponding to different alleles of a common gene.
- alternate** One leaf or other structure per node. (arrangement)
- alternation of generations** *haplodiplontic life cycle*.
- alternipetalous** Stamens with point of attachment between the petals or corolla lobes. (stamen position)
- alternisepalous** Stamens with point of attachment between the sepals or calyx lobes. (stamen position)
- alveolar** An exine wall structure having numerous, spherical air pockets within the exine.
- amb** The outline shape of a pollen grain in polar view.
- ament** *Catkin*. (inflorescence type)
- amoeboid** An anther tapetum type in which the tapetal cell walls break down, with release of the cytoplasm of the tapetal cells into the locule. Syn: *plasmodial*; *periplasmodial*.
- amphiphloic siphonostele** A siphonostele in which a ring of xylem is surrounded by an outer and inner layer of phloem.
- amphistomal** Referring to the micropyle of a bitegmic ovule formed or delimited by both integuments.
- amphitropous** A general ovule type in which curvature of the ovule during development displaces the micropyle adjacent to the funiculus base, with the nucellus bent along both upper and lower sides.
- amplexicaul** Sessile and clasping most of stem circumference. (leaf attachment)
- amplification** The replication of numerous copies of DNA.
- amplification fragment length polymorphism** The use of amplified DNA fragments that exhibit length polymorphism, enabling the recognition of numerous genetic markers. Abbr: *AFLP*.
- amyloplast** *Starch grain*.
- ana-amphitropous** An ovule type in which a vascular strand curves, traversing from the base of funiculus to the chalazal region of the nucellus, with the nucellus bent sharply in the middle along both the lower and upper sides, often with differentiated cells (basal body) at the angle of the bend.
- ana-campylotropous** An ovule type in which a vascular strand curves, traversing from the base of funiculus to the chalazal region of the nucellus, with the nucellus bent only along the lower side, with no basal body.
- anatropous** A type of ovule in which curvature during development results in displacement of micropyle to a position adjacent to the funiculus base; most common and apomorphic for the angiosperms.
- ancestral** Referring to a preexisting condition or character state. Syn: *plesiomorphic*; *primitive*.
- androdioecious/androdioecy** Having male flowers on some individuals and perfect flowers on other individuals. (plant sex)
- androecium** The male organ(s) of a flower; collectively all stamens of a flower. (flower part)
- androgynophore** A stalklike structure that bears the gynoecium and androecium; e.g. *Passiflora*. (flower part)
- andromonoecious** Having both staminate and perfect flowers on the same individual. (plant sex)
- anemochory** Dispersal of propagules by wind. Adj: *anemochorous*.
- anemophily** Pollination by wind. Adj: *anemophilous*.
- anisomeric** Having a different number of members in different whorls. (merosity, perianth merosity)
- annotated phylogenetic classification** A classification in which monophyletic groups are ordered by a sequential listing of derivative lineages from the base to the apex of the cladogram.
- annotation label** A label affixed to an herbarium specimen that verifies or changes the identity of a specimen or that documents the removal of plant material from the specimen.
- annual** Plant living 1 year or less. (duration)
- annual ring** The accumulation of secondary xylem (or phloem) over a single growing season, being evident because of the structural difference between the last cells of the summer wood and the first cells of the subsequent spring wood.
- annulus** A single row of specialized cells, having differentially thickened cell walls, on the outer rim of a *leptosporangium*, functioning in its dehiscence.
- antepetalous** *Antipetalous*. (stamen position)
- anterior** Referring to the lower lobe or part, especially in a horizontally oriented structure. (position)
- antesepalous** *Antisepalous*. (stamen position)
- anther** The pollen-bearing part of a filamentous stamen. (stamen part)
- anther sac** *Microsporangium*. (anther part)
- antheridial wall** The outer, sterile layer of cells of the antheridium. Syn: *jacket layer*; *sterile jacket layer*.
- antheridiophore** A specialized, stalked, generally peltate structure that grows from the gametophyte of some liverworts and bears antheridia.
- antheridium** The male gametangium of the gametophyte of land plants, producing sperm cells and surrounded by a outer layer of sterile cells, the antheridial wall or jacket layer.
- antherode** The sterile anther of some staminodes. (stamen type)
- anthesis** Time of flowering; the opening of flower with parts available for pollination. (maturation, flower maturation)
- anthocarp** A fruit in which one or more flower parts functions as accessory tissues, e.g. *Pontederia* (Pontederiaceae), in which an accrescent perianth surrounds and fuses to the achene. (fruit type)
- antipetalous** Stamens with point of attachment in line with (opposite) the petals or corolla lobes, e.g. Primulaceae, Rhamnaceae. Syn: *antepetalous*. (stamen position)
- antipodal cells** In a typical angiosperm female gametophyte, the three haploid cells that are positioned opposite the micropyle, i.e. in the chalazal region.
- antisepalous** Stamens with point of attachment in line with (opposite) the sepals or calyx lobes. Syn: *antesepalous*. (stamen position)
- antrorse** Bent or directed upward, usually referring to small appendages. (orientation)
- aperture** A specially delimited region of the pollen grain wall.
- apetalous** Having no petals or corolla. (perianth merosity)

- apical** (a) At or near the top, tip, or end of a structure. (position)
(b) Style arising at the apex of the ovary. (style position) (c) With the placenta at the top of the ovary. Syn: *pendulous*. (placentation)
- apical bud** *Terminal bud*. (bud type)
- apical meristem** A region of actively dividing cells in the land plants, located at the apex of the thallus, shoot, or root.
- apical-axile** With two or more placentae at the top of a septate ovary, e.g. Apiaceae. (placentation)
- apiculate** With a flexible, apical process, usually slightly curled, length : width ratio >3:1. (apical process)
- apocarpous** With carpels distinct, the pistil or ovary simple. (gynoecial fusion)
- apolar** Pollen polarity in which polar and equatorial regions cannot be distinguished after pollen grain separation from the microspore tetrad.
- apomorphic** *Derived*.
- apomorphy** A derived condition or character state, representing an evolutionary novelty. Adj: *apomorphic*.
- apomorphy-based** A type of phylogenetic classification in which an apomorphy is the basis for grouping, such that all members of a monophyletic group that share a given, unique evolutionary event are grouped together.
- apopetalous** With distinct petals. Syn: *polypetalous*. (perianth fusion)
- aposepalous** With distinct sepals. Syn: *polysepalous*. (perianth fusion)
- apostemonous** With distinct stamens. (stamen fusion)
- apotepalous** With distinct tepals. (perianth fusion)
- appressed** Pressed closely to an axis oriented upward, with a divergence angle of 0°–15° from upper axis. (orientation)
- aquatic** Growing in water. (plant habitat)
- arachnoid/arachnose** With trichomes forming a cobwebby mass. (vestiture)
- arborescent** Treelike in appearance and size. (stem habit)
- archegonial chamber** A cavity between the megasporangium and female gametophyte of gymnosperm seeds, into which sperm cells are released by the male gametophyte.
- archegonial wall** The outer, sterile layer of cells of the archegonium. Syn: *jacket layer*; *sterile jacket layer*.
- archegoniophore** A specialized, stalked, generally peltate structure that grows from the gametophyte of some liverworts and bears archegonia.
- archegonium** The female gametangium of the gametophyte, containing a basal egg cell and surrounded by an outer layer of sterile cells, the archegonial wall, which differentiates into a basal venter and proximal neck.
- archesporial cell** A single, large cell of an immature ovule that either directly becomes the megasporocyte or that divides once to form a parietal cell and a megasporocyte.
- areole** A modified, reduced, nonelongating shoot apical meristem bearing spines, e.g. Cactaceae. (stem/shoot type)
- aril** A fleshy outgrowth of the funiculus, raphe, or integuments (but separate from the integuments), generally functioning in animal seed dispersal, e.g. Sapindaceae. Adj: *arillate*. (seed part)
- aristate** With a stiff apical process, usually prolonged and straight, length : width ratio >3:1. (apical process)
- arrangement** Placement with respect to similar parts. (disposition)
- arylphenalenones** A class of chemical compounds common in the Haemodoraceae and also found
- ascending** Directed upward, with a divergence angle of 15°–45° from upper axis. (orientation)
- ascidiate** Referring to a carpel that is not leaflike, but develops from a ring of tissue that grows upward, sometimes assuming a somewhat peltate form.
- asepalous** Having no sepals or calyx. (perianth merosity)
- asterad type** A type of embryo development in which the terminal cell divides longitudinally, with both basal and terminal cell derivatives contributing to the mature embryo.
- asymmetric** Lacking a plane of symmetry. (symmetry)
- atactostele** A stele consisting of numerous, collateral vascular bundles positioned throughout the stem tissue.
- atectate** A pollen grain that ancestrally lacks a tectum.
- ategmic** An ovule lacking an integument.
- attenuate** Basal margins abruptly incurved (concave), the basal intersection angle <45°. (base)
- attractant** An aspect of floral morphology that functions to entice an animal pollinator to the flower, either by vision or by odor.
- auctorium non** The misapplication of a name, such that the type specimen of the name does not fall within the circumscription of the taxon being referred to by that name. Abbr: *auct. non*.
- auriculate** With two rounded, basal lobes, margins above lobes concave. (base)
- autapomorphy** An apomorphy that occurs for a single lineage or taxon.
- author** The name of the person who first validly published a scientific name.
- autochory** *Self-dispersal*. Adj: *autochorous*.
- autogamy** Inbreeding occurring within a single flower. Adj: *autogamous*. Syn: *infrafloral selfing*.
- autonym** An automatically created name for infrafamilial, infrageneric, and infraspecific taxa.
- autopolyploidy** Polyploidy occurring within a species.
- autumnal** Appearing in fall. (periodicity)
- awn** (a) bristlelike, apical appendage, e.g. on the glumes or lemmas of grass spikelets. (b) A unit of a pappus type in the Asteraceae that is narrow, elongate, straight, and stiff. (inflorescence part)
- axial parenchyma** Longitudinally oriented parenchyma cells that occur in some secondary xylem (wood) tissues.
- axil** The region at the upper (adaxial) junction of leaf and stem. (position)
- axile** With the placentae positioned along the column in a septate, compound ovary. (placentation)
- axillary** (a) On the side of a structure or at the nodes of an axis. (position). (b) With the inflorescence positioned in the axil of the nearest vegetative leaf. Syn: *lateral*. (inflorescence position)
- axillary bud** Bud in axils of leaves or leaf scars. Syn: *lateral bud*. (bud type)
- baccate** Fleshy or juicy, often with reference to a fruit. Syn: *succulent*, *carnose*. (texture)
- baculate** A pollen sculpturing with rod-shaped elements, each element termed a baculum.
- baculum** A rod-shaped element, as in the wall sculpturing of some pollen grains. Pl: *baculi*. Adj: *baculate*.
- barbed/barbellate** With minute, lateral, sharp appendages (barbs) arising along the surface or margin of a bristle, the barbs typically antrorse or retrorse in orientation. (bristle type)
- bark** Tissues external to the vascular cambium in stem (and roots) of woody plants, consisting of secondary phloem (inner bark) and

- derivatives of the cork cambium (outer bark or periderm). (stem/shoot parts)
- basal** (a) At or near the bottom or base of a structure. Syn: *radical*. (position) (b) With the placenta at the base of the ovary, e.g. Asteraceae, Poaceae. (placentation) (c) With three or more primary veins diverging from one point at the base of the blade, a subcategory of actinodromous and palinactinodromous. (leaf venation)
- basic** A type of anther wall development in which both secondary parietal cell layers divide to yield two middle layers.
- basical capsule** A capsule dehiscing by means of basal slits, as in *Aristolochia* spp. (fruit type)
- basifixed** Anther attached at its base to the apex of the filament. (anther attachment)
- basionym** The original, but now rejected, name, part of which has been used in a new combination.
- Bayesian analysis/inference** A method of phylogenetic inference based upon the posterior probability of a phylogenetic tree.
- beak** An extended, usually accrescent, basal styler region, typically functioning in fruit dispersal, e.g. *Taraxacum*, dandelion. (style structural type)
- beard** A tuft, line, or zone of trichomes on a perianth or perianth part. Adj: *bearded*. (perianth part, vestiture)
- berry** A fleshy fruit with a succulent pericarp, e.g. *Vitis*, grape. (fruit type)
- bicollateral bundle** A vascular bundle with phloem to both the inside and outside of the xylem.
- bicolor unit** An inflorescence unit, possibly an apomorphy of the Malvaceae s.l., consisting of a modified, three-bracted cyme, the bracts modified into an epicalyx in members of the group.
- biennial** A plant living 2 years, typically forming a basal rosette of leaves during the first year and flowering with an elongate inflorescence stalk in the second year. (duration)
- bifid** Two-lobed to two-divided, especially at the apex. (division)
- bigeminate** A compound leaf with two rachillae, each bearing two leaflets. (leaf type)
- bilabiate** Two-lipped, with two, generally upper and lower segments, e.g. many Lamiaceae. (perianth type)
- bilateral** *Zygomorphic, irregular*. (symmetry)
- bimerous** Referring to a whorl with two members. (merosity, perianth merosity)
- binary character** A character with only two character states.
- binomial** Format for the scientific name of species, composed of two names, the genus name and the specific epithet, italicized or underlined.
- binucleate** Having two nuclei, referring to some angiosperm pollen grains at the time of release.
- bipinnate/bipinnately compound** A compound leaf with two orders of axes, the second axes (rachillae) bearing leaflets. (leaf type)
- bipinnatifid** Bipinnately lobed to divided. (division)
- biradial** Having two planes of symmetry. (symmetry)
- biseriate** (a) With two whorls of parts. (cycly) (b) Perianth parts in two distinct whorls. (perianth cycly) (c) Having two whorls or cycles of stamens (stamen cycly) Syn (a c): *dicyclic*. (d) Rays in wood that are made up of two vertical rows of cells.
- bisexual** Flowers having both carpel(s) and stamen(s). Syn: *perfect*. (flower sex)
- bisporangiate** Anther with two microsporangia and typically one theca. Cf: *monothechal*. (anther type)
- bisporic** Megasporogenesis in which cytokinesis occurs after the first meiotic division, but not the second, resulting in two cells, each of which contain two haploid nuclei, with one of the binucleate cells contributing to the female gametophyte.
- bitegmic** An ovule with two integuments, apomorphic for the angiosperms.
- bitermate/bitermately compound** A compound leaf with three axes, each of which is ternately compound. (leaf type)
- blade** The flat, expanded portion of leaf. Syn: *lamina*. (leaf part)
- bootstrap/bootstrapping** A method of evaluating cladogram robustness that reanalyzes the data of the original character × taxon matrix by selecting (resampling) characters at random, such that a given character can be selected more than once.
- bostryx** *Helicoid cyme*. (inflorescence type)
- botany** The traditional study of photosynthetic organisms (including the green plants, red algae, brown plants, dinoflagellates, and euglenoids, but excluding the photosynthetic bacteria), the true fungi, and groups that used to be treated as fungi, such as the Oomycota and slime molds; inclusive of the plant sciences.
- bract** A modified, generally reduced leaf, generally found associated with reproductive organs, e.g. subtending the ovuliferous scale of conifers or the flowers or inflorescence axes of flowering plants. Adj: *bracteate*. (leaf structural type)
- bractlet/bracteole** A smaller or secondary bract, often borne on the side of a pedicel. Syn: *prophyllum*. (leaf structural type, flower part)
- Bremer support** *Decay index*.
- bristle** An external hairlike plant structure, but stouter than a trichome.
- brochidodromous** Pinnate venation in which secondary veins do not terminate at the margin, forming prominent upward loops near the margin, joining other, more distal, secondary veins. (leaf venation)
- bud** An immature shoot system, often surrounded by protective scale leaves, developing into a lateral branch, a flower, or an inflorescence; may be gametophytic or sporophytic. (plant part, stem/shoot parts, twig part)
- bud primordium** An immature bud of the shoot, typically located in the leaf axil. (stem/shoot parts)
- bulb** A short, erect, underground stem surrounded by fleshy leaves, e.g. *Allium* spp., onions. (stem/shoot type)
- bulbel** A proliferative bulb arising from existing bulbs at or below ground level. (stem/shoot type)
- bulbil** A proliferative bulb arising from shoots above ground, typically within an inflorescence. (stem/shoot type)
- bullate** *rugose*. (configuration)
- bundle cap** An outer patch of sclerenchyma fibers associated with a vascular bundle.
- bur** A multiple fruit of achenes or grains surrounded by a prickly involucre, e.g. *Cenchrus* (Poaceae), *Xanthium* (Asteraceae). (fruit type)
- burl** A swollen, woody underground stem from which arises persistent, woody, aerial branches, e.g. fire regenerative stems in some *Arctostaphylos* spp., Manzanita. Syn: *lignotuber*. (stem/shoot type)
- buttress roots** Enlarged, horizontally spreading and often vertically thickened roots at the base of trees that aid in mechanical support. (root type)
- C4 photosynthesis** An alternate photosynthetic pathway of some land plants in which carbon dioxide is initially fixed in the mesophyll cells by the enzyme PEP carboxylase, producing a

- four-carbon molecule, which is transported to bundle sheath cells, where the carbon dioxide is released and fixed by ribulose-bisphosphate carboxylase in the typical dark reactions.
- caducous** Dropping off very early compared with what is typical, usually applied to floral parts. (duration)
- callose** A polysaccharide, composed of beta-1,3-glucose units, which lines the pores of sieve areas and sieve plates of sieve elements and is commonly deposited within pollen tubes.
- calyptra** An apical region of archegonial tissue that is torn from and lifted up by the elongating sporophyte during the latter's development and that may function to protect the young sporophyte apex.
- calyptrate** Having calyx and corolla fused into a cap that falls off as a unit, e.g. *Eucalyptus*. (perianth type)
- calyx** The outermost series or whorl of modified leaves in the perianth, the units of which are sepals. (flower part)
- calyx lobes** The segments of a calyx that is synsepalous (with connate sepals).
- campanulate** Bell-shaped, i.e. with a basally rounded, flaring tube about as broad as long and flaring lobes, e.g. *Campanula*; may also be used for bell-shaped apopetalous corolla or apotepalous perianth. (perianth type)
- campitodromous** Pinnate venation in which secondary veins do not terminate at the margin. (leaf venation)
- campylodromous** With several primary veins running in prominent, recurved arches at the base, curving upward to converge at the leaf apex. (leaf venation)
- campylotropous** A general ovule type in which curvature of the ovule during development displaces the micropyle adjacent to the funiculus base, with the nucellus bent only along the lower side.
- canaliculate** Longitudinally grooved, usually in relation to petioles or midribs. (configuration)
- canescent** Covered with dense, fine grayish-white trichomes; whitish-pubescent. (vestiture)
- capillary bristle** A unit of a pappus type in the Asteraceae that is generally straight, very thin, and threadlike, often barbellate.
- capitate** Head-shaped; spherical with a short basal stalk. (shape)
- capitulum** *Head*. (inflorescence type)
- capsule** (a) The spore-producing component of the sporophytes of liverworts, hornworts, and mosses. (plant part). (b) A dry, dehiscent fruit derived from a compound ovary. (fruit type)
- carinate** Keeled, having a sharp, median fold projected on the abaxial side; sharply conduplicate. (perianth type)
- carnose** *Baccate*; *succulent*. (texture)
- carpel** The unit of the gynoecium of angiosperms; ancestrally, a modified, conduplicate female (mega-) sporophyll of a flower, at maturity enclosing one or more ovules. (flower part, gynoecium part)
- carpopophore** The stalklike structure that is attached to the mericarps in a schizocarp of mericarps, e.g. Apiaceae. (fruit part)
- cartilaginous** With the texture of cartilage; hard and tough but flexible, usually whitish. (texture)
- caruncle** A fleshy outgrowth at the base of the seed, generally functioning in animal seed dispersal, e.g. *Viola*. (seed part)
- caryophyllid type** A type of embryo development in which the terminal cell divides transversely, with only terminal cell derivatives contributing to the mature embryo.
- caryopsis** *Grain*. (fruit type)
- Casparian strip** A band or ring of mostly suberin that infiltrates the cell wall of endodermal cells, functioning to force water and mineral solutes to pass through the plasma membrane of these cells.
- cataphyll** A rudimentary scale leaf found in usually hypogeous (cryptocotylar) seedlings. (leaf structural type)
- catkin** A unisexual, typically male spike or elongate axis that falls as a unit after flowering or fruiting, e.g. *Quercus*. Syn: *ament*. (inflorescence type)
- caudate** Abruptly acuminate into a long, narrowly triangular (tail-like) apical region. (apex)
- caudex** A short, thick, vertical or branched perennial stem, underground or at/near ground level. (stem/shoot type)
- caudiciform stem** A low (at or above ground level), swollen, perennial storage stem from which arise annual or otherwise nonpersistent photosynthetic shoots, e.g. some *Dioscorea* spp., *Calibanus*. (stem/shoot type)
- caulescent** Having an above ground, vegetative stem. (stem habit)
- cauliflorous** Inflorescence growing directly from a woody trunk. (inflorescence position)
- cauline** Positioned along the length of the stem. (position)
- cell** The structural unit of all life.
- cell differentiation** The series of changes that a cell undergoes from the point of inception to maturity, involving the transformation of a meristematic cell into one that assumes a particular structure and function.
- cell expansion** Growth of a cell in size, often involving axial elongation.
- cell theory** The postulate that all life is composed of one or more cells, that cells arise only from preexisting cells, and that cells are the units of metabolic processes.
- cell wall** A layer of the plant cell that is secreted outside the plasma membrane.
- cellular endosperm** An endosperm in which the endosperm cell divides mitotically, regularly followed by cytokinesis, each endosperm nucleus contained within a cell wall.
- cellulose** A polymer of glucose sugar units (= polysaccharide) in which the glucose molecules are chemically bonded in the beta-1,4 position (= β -1,4-glucopyranoside); a major component of the cell wall of green plants.
- cellulose** A polysaccharide of glucose units (β -1,4-glucopyranoside), a primary component of plant cell walls.
- central** At or near the middle or middle plane of a structure. (position)
- central cell** In a typical angiosperm female gametophyte, the single, relatively large cell in the central region that contains the two polar nuclei.
- centric leaf** A leaf that is cylindrical in shape, e.g. *Fenestraria* of the Aizoaceae. (leaf structural type)
- centrifugal** Developing from the center region toward the outside or periphery; can be applied to parts of perianth, calyx, corolla, androecium, or gynoecium. (flower maturation)
- centripetal** Developing from the outside or periphery toward the center region; can be applied to parts of perianth, calyx, corolla, androecium, or gynoecium. (flower maturation)
- cernuous** With tip drooping downward, abaxially. (transverse posture)
- cespitose/caespitose** Referring to a generally short, bunched, much-branched plant forming a cushion. (stem habit)
- chaff** One of the bracts subtending flowers in some Asteraceae, e.g. tribe Heliantheae. Syn: *palea*. (leaf structural type)
- chalazal** Describing the region of an ovule that is opposite the micropyle.

- chalazal** Referring to the proximal region of the ovule, opposite the micropyle.
- character** a feature or attribute of a taxon
- character correlation** The condition in which one character is interdependent upon and influenced by another character.
- character evolution** The sequence of evolutionary changes occurring for a given character.
- character optimization** The representation of characters in a cladogram in a most parsimonious way, such that the minimal number of character state changes occur.
- character state** one of two or more forms of a character
- character step matrix** A numerical tabulation of the number of changes occurring between all pairwise combinations of character states for a given character.
- character × taxon matrix** A numerical tabulation of the characters and corresponding character states for each taxon in a phylogenetic analysis.
- chartaceous** Opaque and of the texture of writing paper. (texture)
- chasmogamy** Referring to typical flowers in which the perianth opens and exposes the sexual organs, with subsequent cross-pollination common. Adj: *chasmogamous*.
- cheiropterophily** Pollination by bats. Adj: *cheiropterophilous*.
- chenopodiad type** A type of embryo development in which the terminal cell divides transversely, with both basal and terminal cell derivatives contributing to the mature embryo.
- Chlorobionta** A monophyletic group of eukaryotes, consisting of the green algae and the land plants, united having chloroplasts with chlorophyll a and b, starch, and thylakoids stacked as grana [green plants].
- chlorophyll a** The primary pigment in the light reactions of photosynthesis, found in the chloroplasts of all photosynthetic eukaryotes and some bacteria.
- chlorophyll b** An accessory pigment in the light reactions of photosynthesis, an apomorphy of the green plant chloroplast.
- chlorophyllous** Having chloroplasts at maturity; green.
- chlorophyllous cell** 1. Any general, chloroplast-containing cell.
2. One of the chloroplast-containing cells in the specialized leaves of *Sphagnum* moss the border and surround a hyaline cell.
- chloroplast** A double membrane-bound organelle with internal thylakoid membranes (lamellae and grana in the green plants), functioning in the reactions of photosynthesis.
- chloroplast** An organelle of some eukaryotes that functions in photosynthesis.
- chromoplast** Carotenoid-containing bodies that function to provide yellow, orange, or red pigmentation for a plant organ, as in petals or fruits; a type of ergastic substance.
- ciliate** With conspicuous marginal trichomes. (margin, vestiture)
- ciliolate** With minute trichomes protruding from margins; minutely ciliate. (margin, vestiture)
- cincinnus** *Scorpioid cyme*. (inflorescence type)
- circinate** With the leaf (blade plus rachis and rachillae, if present) coiled from apex to base, as in young fern and cycad leaves. (posture: *ptyxis/aestivation*)
- circinate vernation** Descriptive of a leaf (including the blade and rachis/rachillae, if present) that is coiled from apex to base when immature, as in young fern and cycad leaves (posture: *ptyxis/aestivation*).
- circumferential** At or near the circumference; surrounding a rounded structure. (position)
- circumscissile capsule** A capsule having a transverse line of dehiscence, e.g. *Plantago*. Syn: *pyxis/pyxide*. (fruit type)
- cirrrose** With a flexible, greatly curled apical process. (apical process)
- clade** A sequence of ancestral-descendent populations through time; a set of organisms interconnected through time and space by the transfer of genetic material from parents to offspring, represented as a line on a cladogram. Syn: *lineage*.
- cladistics** A methodology for inferring the pattern of evolutionary history of a group of organisms, utilizing grouping of taxa by apomorphies [phylogenetic systematics].
- cladistics** *Phylogenetic systematics*.
- cladode/cladodophyll** A flattened photosynthetic stem, functioning as and resembling a leaf. Adj: *phylloclad*. (stem/shoot type)
- cladodromous** Pinnate venation in which secondary veins do not terminate at the margin and branch near the margin. (leaf venation)
- cladogram** A branching diagram that conceptually represents the best estimate of phylogeny. Syn: *phylogenetic tree*.
- cladogram robustness** A measure of the confidence for which a cladogram actually denotes phylogenetic relationships, e.g. by bootstrapping.
- clambering** Sprawling across objects, without specialized climbing structures. Syn: *scandent*. (stem habit)
- classification** The arrangement of taxa (or other entities) into some type of order or grouping.
- clava** A club-shaped element, as in the sculpturing of some pollen grains. Pl: *clavae*.
- clavate** (a) Club-shaped; terete with a gradually tapering thickened and rounded end. (shape) (b) A pollen sculpturing with club-shaped elements, each element termed a clava.
- claw** An attenuate base of a sepal or petal. (perianth part)
- cleft** Sinuses extending (pinnately or palmately) one quarter to one half the distance to the midrib, midvein, or vein junction. (division)
- cleistogamy** Referring to flowers in which the perianth remains closed, such that pollen produced from within the flower pollinates only the stigma(s) of that flower. Adj: *cleistogamous*.
- climbing** Growing upward by means of tendrils, petioles, or adventitious roots. (stem habit)
- coherent** With like parts joined, but only superficially and easily separable. (fusion)
- coleoptile** A protective sheath surrounding the epicotyl, e.g. in some members of the Poaceae. (seed/embryo part)
- coleorhiza** A protective sheath surrounding the radicle, e.g. in some members of the Poaceae. (seed/embryo part)
- collateral** A vascular bundle with an internal strand of xylem and an external strand of phloem.
- collateral bud** Bud(s) lateral to the axillary bud. (bud type)
- collateral bundle** A vascular bundle with xylem to the inside and phloem to the outside.
- collenchyma** A cell type that is live at maturity, has unevenly thickened, pectic-rich, primary cell walls, and functions in structural support, often found at the periphery of stems or leaves.
- colleter** A structure on the inner surface of connate stipules that secretes mucilage, aiding in protection of young, developing shoots, e.g. Rubiaceae. (leaf parts)
- color pattern** The distribution of colors on an object. (color)
- colporate** A pollen grain aperture type that is shaped like a colpus but has a circular region in the center.

- colpus** A pollen grain aperture that is elongate with a length:width ratio of greater than 2:1. Pl: *colpi*. Adj: *colpate*.
- columella** (a) A central column of sterile (non-spore-producing) tissue within the sporophyte capsule of hornworts. (b) One of the middle, radially elongate elements of a tectate-columellate pollen exine wall. Pl: *columellae*.
- column** (a) The central axis to which septae and/or placenta are attached in axile or free-central placentation. (gynoecium part) (b) *gynandrium*, *gynostegium*, *gynostemium*. (flower part)
- combinatio nova** Indication that a taxon has recently been transferred to a new position or rank. Abbr: *comb. nov.*
- commemorative** A name that is after a person or place.
- common name** A vernacular name, used by people within a limited geographic region, not formally published and governed by no rules.
- comose** With an apical tuft of trichomes. (vestiture)
- companion cell** A parenchyma cell associated with a sieve tube member, derived from the same parent cell as is the sieve tube member and functioning to load and unload sugars into the cavity of the sieve tube member.
- complete** Having all four major whorls or floral parts: sepals, petals, stamens, carpels. (flower cycle)
- complex tissue** A tissue that contains more than one cell type.
- compound cone** The cone of a conifer, consisting of an axis bearing bracts, each of which subtends a modified branch systems, the ovuliferous scale.
- compound corymb** A branched corymb, consisting of two or more orders of inflorescence axes bearing flat-topped or convex, pedicellate flowers. (inflorescence type)
- compound cyme** A branched determinate inflorescence, similar to a compound dichasium but lacking a consistent dichasial branching pattern, often by reduction of internodal axes. (inflorescence type)
- compound dichasium** A many-flowered, determinate inflorescence of repeatedly branching simple dichasia units. (inflorescence type)
- compound leaf** A leaf divided into two or more discrete leaflets. (leaf type)
- compound ovary/pistil** An ovary/pistil composed of a two or more carpels, the gynoecium syncarpous. (ovary/pistil type)
- compound perforation plate** A perforation plate composed of several openings.
- compound receptacle** A mass of tissue at the apex of a peduncle that bears more than one flower. Syn: *torus*. (inflorescence part)
- compound sieve plate** A sieve plate that is made up of two or more aggregations of pores.
- compound umbel** An umbel with the peduncle bearing rays attached at one point and unit simple umbels attached at the tip of the rays, e.g. many Apiaceae. (inflorescence type)
- conduplicate** Longitudinally folded at the central axis, with adjacent adaxial sides facing one another. (longitudinal posture)
- cone** A modified, determinate, reproductive shoot system of many nonflowering vascular plants, consisting of a stem axis bearing either sporophylls (in simple cones) or ovuliferous scales subtended by bracts (in compound cones of conifers). Syn: *strobilus*. (plant part)
- confer** Indication that the identity of a specimen is questionable or uncertain and should be compared with specimens of the taxon indicated for more detailed study. Abbr: *cf.*
- configuration** Referring to gross surface patterns other than venation or epidermal excrescences. (surface)
- connate** With like parts integrally fused, not easily separable. (fusion)
- connate-perfoliate** Two opposite leaves fused basally, such that the blade base of each leaf completely surrounds the stem. (leaf attachment)
- connective** The tissue or filament extension between the thecae of an anther. (anther part)
- connivent** Convergent apically without fusion. (orientation)
- consensus tree** A cladogram derived by combining features in common between two or more cladograms.
- conservation of names** A principle of the International Code of Botanical Nomenclature stating that scientific names that are well known and frequently used may be retained over other, earlier, but less well-known, names.
- consistency index (CI)** A measure of the relative amount of homoplasy in a cladogram, equal to the ratio of the minimum possible number of character state changes to the actual number of changes that occur.
- contiguous** With parts touching but not fused. (fusion)
- contorted** *Convolute*. (perianth aestivation)
- convergence** Homoplasy occurring by the independent evolution of a similar feature in two or more lineages. Syn: *parallelism*.
- convolute** Perianth parts of a single whorl overlapping at one margin, being overlapped at the other, e.g. corolla of Malvaceae. Syn: *contorted*. (perianth aestivation)
- cordate** With two rounded, basal lobes intersecting at sharp angle, the margins above lobes smoothly rounded. (base)
- cordate/cordiform** Shaped like an upside-down Valentine heart; approximately ovate with a cordate base. (shape)
- coriaceous** Thick and leathery, but somewhat flexible. (texture)
- cork** The outermost layer of the periderm, generated by the cork cambium.
- cork cambium** A sheath or hollow cylinder of cells that develops near the periphery of the stem or root, undergoing tangential divisions to form phelloderm to the inside and cork to the outside.
- corm** An enlarged, solid underground storage stem or stem base, with outer, protective scales. (stem/shoot type)
- cormel** A proliferative corm arising from existing corms. (stem/shoot type)
- corolla** The innermost series or whorl of modified leaves in the perianth, the units of which are petals. (flower part)
- corolla lobe** A segment of a sympetalous corolla (with connate petals).
- corona** A crownlike outgrowth between stamens and corolla; may originate from petals or stamens. (perianth part, perianth type)
- coronate** With a tubular or flaring perianth or staminal outgrowth, e.g. *Narscissus*, *Asclepias*. (perianth type)
- correct name** A legitimate, validly published name that is accepted by a particular author or authors.
- cortex** The outer, mostly parenchymatous tissue inner to the epidermis and external to the vasculature. (root part, stem/shoot parts)
- corymb** An indeterminate inflorescence consisting of a single axis with lateral axes and/or pedicels bearing flat-topped or convex flowers. (inflorescence type)
- costa** (a) *Midrib*. (b) The nonvascularized conductive tissue found in the gametophytic leaves of some mosses. (leaf part)
- costapalmate** Essentially palmately lobed to compound but with an elongate, rachislike extension of the petiole, as in some palms. (leaf type)

- cotyledon** A first (seed) leaf of the embryo, often functioning in storage of food reserves. (seed part)
- cotylespermous** With the food reserve in the cotyledon, part of the embryo. (seed endosperm type)
- couplet** The pair of contrasting leads in a dichotomous key.
- craspedodromous** Pinnate venation in which secondary veins terminate at the leaf margin. (leaf venation)
- crassinucellate** An ovule in which the nucellus develops two or more layers of cells, the inner ones from divisions of a parietal cell.
- crassulacean acid metabolism** An alternate photosynthetic pathway in some xeric, generally succulent plants and functioning to conserve water, in which initial fixation of carbon dioxide occurs at night (when stomata are open) by the enzyme PEP carboxylase to form malic acid, which is stored within vacuoles of the mesophyll cells; during the day the stomata close and CO₂ is released from the vacuoles into the cytoplasm, where it is fixed in the chloroplasts. Abbr: *CAM*.
- crenate** With rounded to obtuse, shallowly ascending teeth, cut $\frac{1}{16}$ to $\frac{1}{8}$ the distance to the midrib, midvein, or junction of primary veins. (margin)
- crenulate** Diminutive of crenate, teeth cut $\leq \frac{1}{16}$ the distance to the midrib, midvein, or junction of primary veins. (margin)
- crownsheath** The collection of overlapping, sheathing leaf bases at the apex of a palm trunk.
- crozier** A leaf that is coiled during its development, characteristic of the leptosporangiate ferns (Polypodiales) and Marattiales. Syn: *fiddlehead*.
- cruciate** With four, distinct petals in a cross form, e.g. many Brassicaceae. (perianth type)
- crucifer type** A type of embryo development in which the terminal cell divides longitudinally, with only terminal cell derivatives contributing to the mature embryo. Syn: *onagrad type*.
- cryptantherous** *Inserted*. (stamen insertion)
- cryptocotylar** *Hypogeous*. (seed germination type)
- crystal** A deposit of silica or calcium oxalate in plant cells that may function as waste products, as calcium ion sinks, or as an irritant to deter herbivory; a type of ergastic substance.
- cupulate** Hooded; with an abaxially concave posterior lip. (perianth type)
- culm** The flowering and fruiting stem(s) of grasses and sedges. (stem/shoot type)
- cuneate** With basal margins approximately straight, intersection angle 45°–90°. (base)
- cup-shaped** Concave-convex along entire surface; may be abaxially or adaxially concave. (longitudinal posture)
- cupule** A structure that encloses a cluster of unitegmic ovules/seeds, with a small opening near the proximal end, through which pollen grains entered; characteristic of some Pteridosperms, e.g. *Caytonia*.
- curator** The person in charge of the day-to-day running of an herbarium.
- cuspidate** Abruptly acuminate into a triangular, stiff or sharp apex. (apex)
- cuticle** A protective layer, containing cutin, that is secreted to the outside of epidermal cells and functions to inhibit water loss; found in all land plants.
- cutin** A polymer of fatty acids, functioning as a sealant in the cuticle layer of land plants, inhibiting water loss.
- cyathium** An inflorescence bearing small, unisexual flowers and subtended by an involucre (frequently with petaloid glands), the entire inflorescence resembling a single flower, e.g. *Euphorbia*. (inflorescence type)
- cycl** Number of cycles or whorls of parts. (number)
- cyme** General term for a determinate inflorescence. (inflorescence type)
- cymule** A small, simple dichasium. (inflorescence type)
- cypsela** An achene derived from an inferior ovary, e.g. Asteraceae. Syn: *achene* (in general sense).
- cytolith** A mass of calcium carbonate attached to a stalk from the cell wall, occurring within specialized cells termed *lithocysts*.
- cytoplasm** Everything inside the plasma membrane but not including the nucleus.
- dark reactions** A series of biochemical reactions of photosynthesis in plants, occurring in the stroma of the chloroplast, during which atmospheric carbon dioxide reacts to produce a molecule of glucose, requiring the input of the high-energy compounds ATP and NADPH₂.
- data information system/database system** Referring to the (computerized) organization, inputting, and accessing of information.
- decay index** A method of evaluating cladogram robustness by calculating how many extra steps are needed (beyond the number in the most parsimonious cladograms) before the original clade is no longer supported; the greater this value, the greater the confidence in a given clade. Syn: *Bremer support*.
- deciduous** Parts persistent for one growing season, then falling off, e.g. leaves of deciduous plants. (duration)
- decompound** A general term for a leaf having leaflets in two or more orders: bi-, tri-, and so on pinnately, palmately, or ternately, compound; also used for a highly divided leaf. (leaf type, division)
- decumbent** Basally prostrate but ascending apically. (stem habit)
- decurrent** Appearing to extend down the stem from the point of attachment, as if fused to the stem, e.g. many Cupressaceae. (leaf attachment)
- decussate** Opposite leaves or other structures at right angles to the preceding pair. (arrangement)
- decussate tetrad** A tetrad in which the four grains are in two pairs arranged at right angles to one another.
- deflexed** Bent abruptly downward. (orientation)
- deltate** Three-sided, length : width ratio 1:1. (shape)
- dendritic** Trichomes treelike, with multiple lateral branches. (trichome type)
- dendrochronology** The scientific study of wood anatomy to infer details about past events.
- dentate** With sharp, coarse teeth that point outward at right angles to margin outline, cut $\frac{1}{16}$ to $\frac{1}{8}$ distance to midrib, midvein, or junction of primary veins. (margin)
- denticulate** Diminutive of dentate, teeth cut to $\frac{1}{16}$ or less the distance to the midrib, midvein, or junction of primary veins. (margin)
- depressed** Pressed closely to axis downward, with divergence angle of 0°–15° from lower axis. (orientation)
- derived** Referring to a new condition or character state. Syn: *apomorphic*, *advanced*.
- dermal tissue** The outer region of plant organs, composed of the epidermis.
- descending** Directed downward, with divergence angle of 15°–45° from lower axis. (orientation)
- descent** The sequence of ancestral-descendant populations through time.

- description** The assignment of features or attributes to a taxon or other entity.
- determinate** (a) A shoot that terminates growth after a certain period, the apical meristem aborting or converting into a flower, inflorescence, or other specialized structure (stem branching pattern, stem type). (b) An inflorescence in which the terminal flower matures first, maturing from apex to base. (inflorescence development)
- dextrorse** Twining helically like a typical, right-handed screw, e.g. some Convolvaceae. (twisting/bending posture)
- diadelphous** With two groups of stamens, each connate by filaments, e.g. many Faboideae (Fabaceae). (stamen fusion)
- dichasium** A determinate inflorescence that develops along two axes, forming one or more pairs of opposite, lateral axes. (inflorescence type)
- dichlamydeous** Perianth composed of a distinct outer calyx and inner corolla, regardless of total number of whorls. (perianth cycle)
- dichogamy** A type of outcrossing mechanism that is the result of differences in timing of maturation of male and female floral parts. Adj: *dichogamous*.
- dichotomous** With veins successively branching distally into two veins of equal size and orientation, e.g. *Ginkgo biloba*. (leaf venation)
- dichotomous key** A key utilizing series of two contrasting statements, each statement a lead, the pair of leads a couplet.
- dicotyledonous** A type of anther wall development in which only the outer secondary parietal cell layer divides to yield the endothecium and a single middle layer.
- dictyostele** A dissected amphiphloic siphonostele.
- dicyclic** *Biseriate*. (cyclic, perianth cyclic, stamen cyclic)
- didymous** With stamens in two equal pairs. (stamen arrangement)
- didynamous** With stamens in two unequal pairs, e.g. many Bignoniaceae, Lamiaceae, Scrophulariaceae. (stamen arrangement)
- diffuse-porous** Wood in which vessels develop more or less uniformly throughout the growth season.
- dioecious/dioecy** Having unisexual flowers, staminate and pistillate on separate individual plants. (plant sex)
- diphthong** A two-vowel combination in Latin that is treated as the equivalent of a single vowel.
- diplohaplontic life cycle** *Haplodiplontic life cycle*.
- diplostemonous** Stamens in two whorls, the outer opposite the sepals, the inner opposite petals. (stamen position)
- discoid** (a) Discus-shaped. (shape). (b) Stigma(s) disk-shaped. (stigma/stigmatic region type)
- disk** A discoid or doughnut-shaped structure arising from the receptacle at the outside and surrounding the stamens. (*extrastaminal disk*), at the base of the stamens (*staminal disk*), or at the inside of the stamens and/or base of the ovary (*intrastaminal disk*); may be nectar-bearing (*nectariferous disk*). (flower part)
- disk flower** Having an actinomorphic, tubular corolla with flaring lobes, e.g. some Asteraceae. (perianth type)
- dispersal** The movement of an organism or propagule from one region to another, such as the transport of a seed or fruit (by wind, water, or bird) from a continent to an island.
- disposition** Relative placement of objects or parts of objects, inclusive of position, arrangement, and orientation.
- dissected** Divided into very fine, often indistinct segments. (division)
- distal** Away from the point of origin or attachment. (position)
- distal pole** The intersection of the pollen grain polar axis with the grain surface that is away from the center of the microspore tetrad.
- distichous** Alternate, with points of attachment in two vertical rows/ranks, e.g. the grasses, Poaceae. (arrangement)
- distinct** With like parts unfused and separate. (fusion)
- distyly** Hecogamy in which two floral morphologies occur: pin flowers, with a long style and short stamens, and thrum flowers, with a short style and long stamens. Adj: *distylous*.
- disulculate** A pollen grain with two elongate apertures on opposite sides of the grain, parallel to the equatorial plane.
- dithecal** Anther with two thecae and typically four microsporangia. Cf: *tetrasporangiate*. (anther type)
- diurnal** During the day, typically with respect to when flowers open. (periodicity)
- divaricate** *Divergent; horizontal; patent*. (orientation)
- divergence/diversification** The formation of two (or more), separate lineages from one common ancestor.
- divided** Sinuses extending (pinnately or palmately) $\frac{3}{4}$ to almost to midrib, midvein, or vein junction. (division)
- dorsal** *Abaxial*. (position)
- dorsal vein** The central vein of a carpel, corresponding to the midvein of a leaf. Syn: *median vein*.
- dorsifixed** Anther attached dorsally and medially to apex of the filament. (anther attachment)
- dorsiventral** Having a flattened shape, with an upper (adaxial) and lower (abaxial) surface, characteristic, e.g. of leaves.
- doubly serrate** With large, serrate teeth that have along the margin smaller, serrate teeth. (margin)
- downward** Anther dehiscing toward the ground in a horizontally oriented flower. (anther dehiscence direction)
- drepanium** A monochasium in which the axes develop on only one side of each sequential axis, typically appearing coiled at least early in development; sometimes equated with helicoid cymes. (inflorescence type)
- drupe** A fleshy fruit with a hard, stony endocarp, e.g. *Prunus*: peach, plum, cherry. (fruit type)
- drupectum** An aggregate fruit of drupes, e.g. *Rubus*: raspberry, blackberry. (fruit type)
- druse** A spherical crystal with protruding spikes, composed of calcium oxalate; a type of ergastic substance.
- duration** The length of life of a plant or plant part. (temporal phenomena)
- dyad** A fusion product of two pollen grains.
- ebracteate** Lacking bracts. (flower attachment)
- echina** A spinelike sculpuring element >1 μm long, as in some pollen grain walls. Pl: *echinae*. Adj: *echinate*.
- eciliate** Without trichomes on the margins, regardless of presence or absence of teeth. (margin)
- ectophloic siphonostele** A siphonostele in which a ring of xylem is surrounded by an outer layer of phloem only.
- ecto-zochoy** Dispersal by animals in which propagules are carried on the outside of an animal. Adj: *ecto-zochoyous*.
- egg** A nonmotile, evolutionarily enlarged gamete, the end product of oögamy.
- ektexine** An outer layer of the pollen grain exine wall.

- elater** (a) One of the hygroscopic appendages arising from the spores of *Equisetum*, functioning in spore dispersal. (b) A non-sporogenous, elongate, hygroscopic cell with spiral wall thickenings that develops within the sporangia of some liverworts and that functions in spore dispersal.
- elliptic** Margins curved, widest at the midpoint, the length : width ratio 2 : 1 to 3 : 2. (shape)
- emarginate** Having an apical incision, cut $\frac{1}{16}$ to $\frac{1}{8}$ the distance to midrib, midvein, or junction of primary veins. (apex)
- embryo** An immature, diploid sporophyte developing from the zygote of land plants. (seed part; plant part)
- embryo proper** That portion of the proembryo that will eventually grow into the new sporophyte.
- embryo sac** Term for the female gametophyte of angiosperms.
- embryogeny** The development of the embryo within the seed.
- Embryophyta/embryophytes** A monophyletic group of eukaryotes united by an outer cuticle, specialized gametangia antheridia and archegonia and an intercalated diploid phase in the life cycle, including the embryo [land plants].
- emend** Abbreviation of a correction or amendment of a name.
- emergent** With roots or stems anchored to substrate under water, the aerial shoots above water. (plant habitat)
- emersed** Occurring under water. (plant habitat)
- enantiostyly** A type of herkogamy in which the style of different flowers curves either the left or the right. Adj: *enantiostylous*.
- enation** A small appendage arising from the stem, resembling a rudimentary leaf but lacking vascular tissue.
- endarch** An orientation of xylem maturation in which the protoxylem is oriented toward the center of the stele relative to the metaxylem, as occurs in eusteles and atactosteles.
- endexine** An inner layer of the pollen grain exine wall.
- endocarp** The innermost wall layer of the pericarp, if the latter is divided into layers. (fruit part)
- endodermis** A hollow cylinder of cells in roots and some stems that surrounds the vasculature and functions to selectively control passage of solutes from the outside, via Casparian strips. Adj: *endodermal*. (root part, stem part)
- endogenous** Arising from the internal tissues, as in the growth of secondary roots from within a primary root.
- endoplasmic reticulum** A cellular organelle consisting of interconnected phospholipid membranes that may function in material transport and as the site of protein synthesis.
- endosperm** The triploid tissue that develops from mitotic divisions of the endosperm cell (the product of double fertilization), ultimately enveloping or abutting the embryo and functioning as the nutritive tissue of angiosperm seeds.
- endospermous** With endosperm as the food reserve in mature seeds. Syn: *albuminous*. (seed endosperm type)
- endospory** The development of a gametophyte within the original spore wall. Adj: *endosporic*.
- endostomal** Referring to the micropyle of a bitegmic ovule delimited by only the inner integument, the outer one being foreshortened.
- endosymbiosis** The intracellular cohabitation of one cell within another cell; the process that gave rise to mitochondria and chloroplasts by engulfment of a prokaryote by a eukaryotic cell.
- endothecium** The outermost cell layer of an anther, typically of enlarged cells with secondary wall thickenings functioning in anther dehiscence.
- endozoochory** Dispersal by animals in which propagules are eaten but are passed through the gut of the animal unharmed. Adj: *endozoochorous*.
- ensiform** Sword-shaped, with length : width ratio greater than 12 : 1, e.g. leaves of *Iris* spp. (shape)
- entire** Without teeth on margins; locally smooth. (Note, however, that surface may be divided.) (margin)
- entomophily** Pollination by insects. Adj: *entomophilous*.
- epicalyx** A series of bracts immediately subtending the calyx, e.g. *Hibiscus*, other Malvaceae. (leaf structural type)
- epicotyl** (a) The first shoot of a vascular plant that develops from the embryo. (b) The first shoot of a seed plant, derived from the embryo of the seed. (stem/shoot parts, seed part)
- epidermal excrescence** Referring to surface patterns from structural outgrowths or secretions of the epidermis. (surface)
- epidermis** The outermost cell layer of all land plant organs, functioning to provide mechanical protection of inner tissue and to inhibit water loss.
- epigeous** With cotyledon(s) elevated above the ground during germination. Syn: *phanerocotylar*. (seed germination type)
- epigynous** With sepals, petals, and stamens attached at the apex of the ovary, the ovary inferior. (perianth/androecial position)
- epihypogynous** With sepals, petals, and stamens attached at middle of the ovary, the ovary half-inferior. (perianth/androecial position)
- epihypoperigynous** Hypanthium present, attached at middle of ovary, with sepals, petals, and stamens attached to hypanthium rim, the ovary half-inferior. (perianth/androecial position)
- epiperigynous** Hypanthium present, attached at apex of ovary, with sepals, petals, and stamens attached to hypanthium rim, the ovary inferior. (perianth/androecial position)
- epipetalous** With stamens adnate to (inserted upon) petals or corolla. Syn: *petalostemonous*. (stamen fusion)
- epiphytic** Growing upon another plant. (plant habitat)
- episepalous** With stamens adnate to (inserted upon) sepals or calyx. (stamen fusion)
- epitpalous** With stamens adnate to tepals or the perianth as a whole. (stamen fusion)
- epitropous** An ovule position in which the micropyle points distally.
- epitropous-dorsal** An epitropous ovule position in which the raphe is dorsal (abaxial), pointing away from the central floral or ovary axis.
- epitropous-ventral** An epitropous ovule position in which the raphe is ventral (adaxial), pointing toward the central floral or ovary axis.
- epitype** A specimen (or illustration) that is selected to serve as the type if the holotype, lectotype, or neotype is ambiguous with respect to the identification and diagnosis of the taxon.
- equator** The intersection with the pollen surface of a plane at a right angle to the polar axis and passing through the center of the grain.
- equatorial view** Observing a pollen grain from the equatorial region.
- equitant** Leaves with overlapping bases, usually sharply folded along midrib. (arrangement)
- erect** Pointing upward. (orientation)
- ergastic substance** A cellular substance that does not function in metabolism, generally functioning in storage, waste secretion, and protection.

- et** Latin for and, used in scientific name combinations.
- eucauptodromous** Pinnate venation in which secondary veins do not terminate at the margin, curving upward near the margin but not directly joining adjacent secondaries. (leaf venation)
- euphyll** The sporophytic leaf of the euphyll group, growing by means of either marginal or apical meristems, having multiple, branched veins, and having an associated leaf gap. Adj: *euphyllous*. Syn: *megaphyll*. (leaf structural type)
- eusporangium/eusporangiate sporangium** A relatively large sporangium that is derived from several epidermal cells and having a sporangial wall composed of more than one cell layer.
- eustele** A primary stem vasculature that consists of a single ring of discrete collateral or bicollateral vascular bundles.
- evergreen** Persistent two or more growing seasons, e.g. leaves of most conifers. (duration)
- evolution** Descent with modification; the transfer of genetic material from parent(s) to offspring over time, with a corresponding change in that genetic material.
- exalbuminous** Lacking endosperm as the food reserve in mature seeds. Syn: *nonendospermous*. (seed endosperm type)
- exarch** An orientation of xylem maturation in which the protoxylem is oriented toward the organ periphery relative to metaxylem, as occurs in some protosteles.
- exfoliating** Bark cracking and splitting off in large sheets. (bark type)
- exindusiate** Referring to a sorus that lacks an indusium.
- exine** The hard, outermost, desiccation-resistant layer of a pollen grain wall, providing structural support and inhibiting desiccation.
- exocarp** The outermost wall layer of the pericarp, if the latter is divided into layers. (fruit part)
- exospory** The formation of a gametophyte external to the original spore wall. Adj: *exosporic*.
- exostomal** Referring to the micropyle of a bitegmic ovule delimited by only the outer integument, the inner one being foreshortened.
- explosively dehiscent** Referring to a dry, dehiscent fruit that opens with force, in the process ejecting the seeds some distance away. (fruit type)
- exserted** With stamens protruding beyond the perianth. Syn: *phaneranthrous*. (stamen insertion)
- exstipellate** Without stipels. (leaf part)
- exstipulate** Without stipules. (leaf part)
- extrastaminal disk** A discoid or doughnut-shaped structure arising from the receptacle at the outside and surrounding the stamens; may be nectar-bearing (nectariferous disk). (flower part)
- extorse** Dehiscing outward, away from the flower center (anther dehiscence direction)
- falcate/falciform** Lanceolate to linear and curved to one side; scimitar-shaped. (shape)
- false indusium** An extension of the blade margin that overlaps the sorus of a leptosporangiate fern.
- farinaceous** Finely mealy, covered with small granules. Adj: *granular*, *scurfy*. (epidermal excrescence)
- fascicle** (a) A shoot with very short internodes on which flowers or leaves are borne. Syn: *short shoot*; *spur*; *spur shoot*. (stem/shoot type). (b) A raceme-like or panicle-like inflorescence with pedicellate flowers in which internodes between flowers are very short, with pedicel bases appearing congested. (inflorescence type) Adj: *fasciculate*.
- fat** A type of triglyceride compound that may function as high-energy storage compounds or secretion products in plants; a type of ergastic substance.
- female** (a) Individual with female reproductive organs only. (plant sex) (b) *Pistillate*. (flower sex)
- female gametophyte** A gametophyte that bears only archegonia, housing the egg cell. Syn: *megagametophyte*, *embryo sac*.
- female sporophyll** A sporophyll that bears one or more megasporangia or seeds. Syn: *megasporophyll*.
- fenestrate** Having windowlike holes in the surface, e.g. *Monstera deliciosa*, Araceae (configuration)
- fertile segment** The sporangia containing component of the leaf of an ophioglossoid fern.
- fiber** A sclerenchyma cell that is long and very narrow, with sharply tapering end walls, functioning in mechanical support and often occurring in bundles.
- fibrous roots** Roots that are adventitious and typically fine and numerous. (root type)
- fiddlehead** *Crozier*.
- filament** A stamen stalk, generally terete in shape. (stamen part)
- filamentous** (a) With a more or less terete stamen stalk, as opposed to a laminar body. (stamen type) (b) Filament present, as opposed to absent and anther sessile. (stamen attachment)
- filiferous** Bearing coarse, fiberlike structures. (margin)
- filiform** Long, thin, and typically flexuous, threadlike, filamentous. (shape)
- fissured** Bark split or cracked into vertical or horizontal, usually coarse grooves. (bark type)
- fistulose/fistular** Cylindrical and hollow within. (shape)
- flabellate** With three or more primary veins diverging from one point and several, equal, fine veins branching toward the leaf apex, a subcategory of actinodromous. (leaf venation)
- flexuous** Central axis and tip alternately curved up and down. (transverse posture)
- floating** Occurring at the water surface. (plant habitat)
- floccose** With dense trichomes in several patches or tufts. (vestiture)
- flora** A listing of the plant taxa of a given region, usually accompanied by keys and descriptions. Syn: *manual*.
- floral diagram** A diagrammatic, cross-sectional view of a flower bud, showing the relative relationship of perianth, androecial, and gynoecial components, and illustrating things such as stamen position, placentation, and perianth, calyx, or corolla aestivation.
- floral formula** A symbolic representation of floral morphology, including cyclcy (number or whorls or series), merosity (number of parts per whorl), fusion of parts, and ovary position.
- floral tube** *Hypanthium*. (flower part)
- floret** A unit of a grass (Poaceae) spikelet, consisting of a short, lateral axis bearing two bracts (lemma and palea) that subtend a terminal, reduced flower. (inflorescence type)
- floristics** The documentation of all plant species in a given geographic region.
- flower** The reproductive organ of flowering plants; a modified, determinate shoot bearing sporophylls (stamens and/or carpels) with or without outer modified leaves, the perianth. (plant part, inflorescence part)
- flower bract** A modified, generally reduced leaf subtending a flower. (flower part)
- flower bud** A bud that develops into a flower. (bud type)

- follicetum** An aggregate fruit of follicles, e.g. *Magnolia*. (fruit type)
- follicle** A dry, dehiscent fruit derived from one carpel that splits along one suture, e.g. *Asclepias*, milkweed. (fruit type)
- foot-layer** The inner layer of a tectate-columellate pollen exine wall.
- form genus** A genus that corresponds to a particular organ of a fossil plant.
- fossulate** A pollen sculpturing with longitudinal grooves.
- foveolate** A pollen sculpturing with a pitted surface caused by pores in the surface.
- free** With unlike parts unfused, separate. (fusion)
- free-central** With the placenta along the column in a compound ovary lacking septa, e.g. Caryophyllaceae. (placentation)
- Frutillaria type** A type of tetrasporic female gametophyte in which three of the four megaspores fuse to form a triploid nucleus, followed by two sequential mitotic divisions of the haploid and triploid nuclei, resulting in an 8-nucleate female gametophyte in which the three antipodals and one of the polar nuclei are triploid, the other polar nucleus and the cells of the egg apparatus remaining haploid.
- frond** Specialized term for a fern leaf.
- fruit** The mature ovary of flowering plants, consisting of the pericarp (mature ovary wall), seeds, and (if present) accessory parts. (plant part)
- frutescent** Having the habit of a shrub, with numerous, woody, aerial trunks. (stem habit)
- fumatory** A substance that is smoked by humans, usually for its pleasing or euphoric effects, e.g. tobacco, *Nicotiana tabacum*.
- funiculus** A stalk that attaches the ovule to the placenta. (gynoecium part)
- fusiform** Spindle-shaped; narrowly ellipsoid with two attenuate ends. (shape)
- galeate** Hooded; with an abaxially concave posterior lip. (perianth type)
- gamete** A specialized, haploid cell that fuses with another gamete (in sexual reproduction) to form a diploid zygote.
- gametophyte** The haploid phase in the life cycle of all land plants.
- gamopetalous** *Sympetalous*. (perianth fusion)
- gamosepalous** *Synsepalous*. (perianth fusion)
- geitonogamy** Inbreeding occurring between different flowers derived from one individual. Adj: *geitonogamous*.
- geminat** A compound leaf with two leaflets arising from a petiole and no rachillae. (leaf type)
- geminat-pinnat** A compound leaf with two rachillae, each bearing a pinnate arrangement of leaflets. (leaf type)
- gemma** (a) An asexual propagule produced within the gemmae cups of some thalloid liverworts. (b) One of the globose or ellipsoid elements of a gemmate pollen grain. Pl: *gemmae*.
- gemma cup** A cup-shaped organ on the upper surface of the gametophytes of some thalloid liverworts, containing gemmae propagules.
- gemmate** A pollen sculpturing with globose or ellipsoid elements, each element termed a gemma.
- gender** The designation of masculine, feminine, or neuter in Latin names.
- generative cell** One of the two initial, haploid cells in the male gametophyte of angiosperms that mitotically divides to form two sperm cells.
- genetic drift** Random genetic modification of a population or species, not the result of natural selection.
- genet** A genetically different individual of a population. Cf: *ramet*.
- geniculate** Having a zig-zag posture, e.g. the inflorescence rachis of some grasses. (twisting/bending posture)
- genus name** The first component of a binomial, always capitalized.
- genus novum** Meaning that a taxon name, at the rank of genus, is new to science. Abbr: *gen. nov.*
- geophyte** A perennial herb, typically with a bulb, corm, rhizome, or tuber underground stem. (plant habit)
- girdling** A type of anther endothecium in which the secondary wall thickenings form rings with cross bridges between them.
- glabrate** Nearly glabrous or becoming glabrous with age. (vestiture)
- glabrous** Without trichomes. (vestiture)
- glandular** (a) Covered with minute, blackish to translucent glands (epidermal excrescence). (b) Trichomes secretory or excretory, usually having an apical glandular cell. (trichome type)
- glandular (tapetum type)** *Secretory*.
- glaucous** Covered with a smooth, usually whitish, waxy coating, which can be rubbed off with touch. (epidermal excrescence)
- globose** (a) Spherical in shape (shape). (b) Stigma(s) spherical in shape. (stigma/stigmatic region type)
- glochidiate** With apical, clustered barblike structures. (bristle type)
- glochidium** A very small leaf spine with numerous, retrorse barbs along its length, produced in the areoles of opuntoid cacti. Pl: *glochidia*; *glochids*. (leaf structural type)
- glomerule** An inflorescence of sessile or subsessile flowers in which internodes between flowers are very short, with flowers appearing congested. (inflorescence type)
- glucosinolate** A secondary chemical compound found in many Brassicales that functions to deter herbivory and parasitism and also serves as a flavoring agent in the commercially important members of the Brassicaceae.
- glume** One of usually two bracts occurring at the base of a grass spikelet. (leaf structural type)
- glutinous** *Viscid*. (epidermal excrescence)
- golgi body** A cellular organelle comprised of parallel stacks of flattened membranes, functioning in transport and modification of compounds.
- grain** A one-seeded, dry, indehiscent fruit with the seed coat adnate to pericarp wall, e.g. Poaceae, grasses. Syn: *caryopsis*. (fruit type)
- granular** *Farinaceous*.
- granum** A pancakelike aggregation of thylakoid membranes within the chloroplast of green plants. Pl: *grana*.
- grass spikelet** The inflorescence unit of the Poaceae, grass family, consisting of an axis (rachilla) bearing distichous parts: two basal bracts (glumes, sometimes modified or absent) and one or more florets, each floret consisting of a minute lateral axis with two additional bracts (lemma and palea) plus the flower.
- green plants** A monophyletic group of eukaryotes, consisting of the green algae and the land plants, united having chloroplasts with chlorophyll a and b, starch, and thylakoids stacked as grana [Chlorobionta].
- ground meristem** The nonvascular, usually parenchymatous tissue between and among the vascular bundles of an atactostele. (stem/shoot parts)
- ground tissue** Tissue that is inside the epidermis and not part of the vascular tissue, composed of parenchyma, sclerenchyma, and collenchyma cells.
- guard cell** One of the two cells that together make up a *stomate*.

- gynandrium** A fusion product of androecium and gynoecium, e.g. Aristolochiaceae, Orchidaceae. Syn: *column*, *gynostegium*, *gynostemium*. (flower part)
- gynobasic** With style arising at the base and center of a lobed ovary, e.g. Boraginaceae, Lamiaceae. (style position)
- gynodioecious/gynodioecy** Having female flowers on some individuals and perfect flowers on other individuals. (plant sex)
- gynoecium** The female organ(s) of a flower, collectively all carpels of a flower. (flower part)
- gynomonoecious/gynomonoecy** Having both pistillate and perfect flowers on the same individual. (plant sex)
- gynophore** A stalk of the pistil, usually absent. Syn: *stipe*. (gynoecium part)
- gynostegium** *column*, *gynandrium*, *gynostemium*. (flower part)
- gynostemium** *column*, *gynandrium*, *gynostegium*. (flower part)
- half-inferior** With sepals, petals, stamens, and/or hypanthium attached at the middle of the ovary. (ovary position)
- hapaxanthic** A determinate shoot that completely transforms into a flower or inflorescence. (stem/shoot type, stem branching pattern, inflorescence development)
- haplodiplontic life cycle** A life cycle having both haploid and diploid phases, occurring in all land plants. Syn: *alternation of generations*; *diplobiontic life cycle*.
- haplontic life cycle** A type of sexual life cycle in which the mature, adult phase is haploid, which produces gametes (egg and sperm) that fuse to form a diploid zygote, the latter undergoing meiosis to produce haploid spores, which develop into new haploid adults. Syn: *haplobiontic life cycle*.
- hardwood** Wood derived from a nonmonocotyledonous angiosperm, generally (but not always) harder than a softwood because of a greater concentration of fiber cells.
- harmomegathy** Volume changes of the pollen grain with changes in water content, e.g. humidity, functioning to inhibit desiccation.
- hastate** With two basal lobes, more or less pointed and oriented outwardly approximately 90° relative to central axis. (base)
- hastula** An appendage or projection at the junction of petiole and blade, as in some palms. (leaf part)
- haustoria** Parasitic roots that penetrate the tissues of a host plant. (root type)
- head** A determinate or indeterminate, crowded group of sessile or subsessile flowers on a compound receptacle, often subtended by an involucre, e.g. Asteraceae. Syn: *capitulum*. (inflorescence type)
- helicoid cyme** A monochasium in which the branches develop on only one side of each sequential axis, appearing coiled at least early in development; may intergrade with scorpioid cyme. Syn: *bostryx*. (inflorescence type)
- helobial** An endosperm in which the first mitotic division is followed by cytokinesis, delimiting two cells, with the nucleus of one of the cells dividing without cytokinesis, that of the other cell dividing with cytokinesis.
- hemiparasite** A chlorophyllous, parasitic plant.
- hemispheric** Half-sphere-shaped. (shape)
- hemitropous/hemianatropous** An ovule somewhat intermediate in curvature between anatropous and orthotropous types.
- herb** A plant with annual above-ground shoots, including a flower or inflorescence, the plant itself being annual, biennial, or perennial. (plant habit)
- herbaceous** Having a soft or slightly succulent texture. (texture)
- herbarium specimen** A pressed and dried plant sample that is permanently glued and/or strapped to a sheet of paper, along with a documentation label.
- hercogamy/herkogamy** The spatial separation of anthers and stigmas, generally enhancing outbreeding. Adj: *hercogamous*/*herkogamous*.
- hermaphroditic** A plant with bisexual flowers. (plant sex)
- hesperidium** A septate berry with a thick-skinned, leathery pericarp wall and fleshy modified trichomes (juice sacs) arising from the inner walls, e.g. *Citris* (orange, lemon, grapefruit, etc.). (fruit type)
- heteroblasty** The condition in which the juvenile leaves are distinctly different in size or shape from the adult leaves, e.g. many Araceae. Adj: *heteroblastic*. (leaf type)
- heterochrony** An evolutionary change in the rate or timing of development.
- heteropolar** Pollen polarity in which the two polar hemispheres are different because of displacement of one or more apertures.
- heterospory** The formation of two types of haploid spores, microspores and megaspores, within two types of sporangia. Adj: *heterosporic*.
- heterostyly** Hercogamy in which the relative lengths or heights of stigmas versus anthers vary among different flowers. Adj: *heterostylous*.
- heterotropous** An ovule that varies in orientation.
- hilum** Funicular scar on the seed coat. (seed part)
- hirsute** With long, rather stiff trichomes. (vestiture)
- hispid** With very long, stiff trichomes, often capable of penetrating skin. (vestiture)
- holotype** The one specimen or illustration upon which a name is based, originally used or designated at the time of publication.
- homochlamydeous** Perianth composed of similar parts, each part termed a tepal. (perianth cycle)
- homology** Similarity that is the result of common ancestry. Adj: *homologous*.
- homolog/homologue** A specific feature that is homologous to another, cited feature. Cf: *homology*.
- homonym** One of two (or more) identical names that are based on different type specimens.
- homoplasy** Similarity that is not due to homology or common ancestry, but the result of independent evolutionary change.
- homospory** The formation of one type of haploid spore, by one type of sporangium. Adj: *homosporic*.
- hood** A hoodlike appendage arising from the gynostegium of some Asclepiadoids of the Apocynaceae.
- hooked** With apical hooklike structure. Syn: *uncinate*. (bristle type)
- horizontal** More or less horizontally spreading with divergence angle of $\leq 15^\circ$ up or down from horizontal axis. Syn: *divaricate*; *divergent*; *patent*. (orientation)
- horn** A hornlike appendage, often associated with a hood, arising from the gynostegium of some Asclepiadoids of the Apocynaceae.
- hyaline cell** One of the nonchlorophyllous cells in the specialized leaves of *Sphagnum* moss, having characteristic pores and helical thickenings and functioning in water absorption and retention.
- hybridization** Sexual reproduction between different species (interspecific hybridization) or between different populations or infraspecific taxa within a species.
- hydathode** A group of specialized cells that secrete excess, transported water (usually due to root pressure) from leaf margins.
- hydrochory** Dispersal of propagules by water. Adj: *hydrochorous*.

- hydroid** A specialized cell that functions in water conduction in some mosses.
- hydrophily** Pollination by water. Adj: *hydrophilous*.
- hygroscopic** Absorbing moisture from the air, often resulting in movement.
- hymenopterophily** *Melittophily*.
- hypanthium** A cuplike or tubular structure around or atop the ovary, bearing along its margin the sepals, petals, and stamens. Syn: *floral tube*. (flower part)
- hypanthodium** An inflorescence bearing numerous flowers on the inside of a convex or involuted compound receptacle, e.g. *Ficus*. (inflorescence type)
- hyphodromous** Pinnate venation with only the primary midrib vein present or evident, the secondary veins absent, very reduced, or hidden within the leaf mesophyll. (leaf venation)
- hypocotyl** A region of the embryo between the root and epicotyl; may function in seedling development and as an anatomical transition between root and shoot. (seed part)
- hypogeous** With cotyledon(s) remaining in the ground during germination. Syn: *cryptocotylar*. (seed germination type)
- hypogynous** With sepals, petals, and stamens attached at the base of a superior ovary. (perianth/androecial position)
- hypotropous** An ovule in which the micropyle points proximally.
- hypotropous-dorsal** A hypotropous ovule in which the raphe is dorsal (abaxial), pointing away from the central floral or ovary axis.
- hypotropous-ventral** A hypotropous ovule in which the raphe is ventral (adaxial), pointing toward the central floral or ovary axis.
- hysteranthy** Timing in which leaf and flower development do not coincide. Adj: *hysteranthous*.
- identification** The process of associating an unknown taxon or other entity with a known one.
- illegitimate name** A name that violates one or more rules of the International Code of Botanical Nomenclature.
- imbricate** (a) Leaves or other structures overlapping (arrangement). (b) With overlapping perianth parts. (perianth aestivation)
- imbricate-alternate** Outer whorl of perianth parts (sepals or outer tepals) alternating with, along different radii, the inner whorl of perianth parts (petals or inner tepals), a typical perianth aestivation. (perianth aestivation)
- imparipinnate/imparipinnately compound** A pinnately compound leaf with a terminal leaflet, typically odd-pinnate. (leaf type)
- imperfect** (a) *Unisexual*. (flower sex) (b) With lateral primary veins covering less than two thirds of the leaf blade area, a subcategory of actinodromous and of acrodromous. (leaf venation)
- in** Abbreviation for in the publication of, referring to a name published within a larger work authored by the person(s) following the in.
- inaperturate** A pollen grain that lacks any recognizable aperture.
- inbreeding** The union of gametes derived from a single individual. Syn: *selfing*.
- incanous** Covered with dense, fine, grayish-white trichomes; whitish-pubescent. (vestiture)
- incised** With margins sharply and deeply cut, usually jaggedly. (margin, division)
- inclined** Directed upward, with a divergence angle of 15°–45° from horizontal axis. (orientation)
- incompatibility reaction** The inhibition of pollen germination or pollen tube growth between genetically similar individuals, mediated by incompatibility genes and functioning to promote outcrossing.
- incomplete** Lacking one or more of the four major whorls or floral parts: sepals, petals, stamens, carpels. (flower cycle)
- incurved** Tip gradually curved inward or upward (adaxially). (transverse posture)
- indehiscent legume** A secondarily modified legume does not split open, e.g. *Arachis hypogaea*, peanut. (fruit type)
- indented phylogenetic classification** A classification in which monophyletic groups are ordered in a sequential, hierarchical method.
- indeterminate** (a) A shoot that has the potential for unlimited growth, the apical meristem continuing to grow. (stem/shoot type, stem branching pattern). (b) An inflorescence in which the basal flower matures first; maturation from base to apex. (inflorescence development)
- induplicate** Plicate, with adjacent adaxial sides facing one another, being V-shaped in cross-section. (longitudinal posture)
- indurate** Hardened and inflexible. (texture)
- indusium** A flap of tissue that covers a *sorus*, found in some leptosporangiate ferns. Adj: *indusiate*.
- inferior** With sepals, petals, stamens, and/or hypanthium attached at the apex of the ovary. (ovary position)
- inflorescence** An aggregate of one or more flowers, the boundaries of which generally occur with the presence of vegetative leaves below; may be composed of unit inflorescences. (plant part)
- inflorescence bract** A modified, generally reduced leaf subtending an inflorescence axis. (inflorescence part)
- inflorescence bud** A bud that develops into an inflorescence. (bud type)
- intrafloral selfing** *Autogamy*.
- intrafoliar** Descriptive of a palm inflorescence that is positioned below the leaves of the crownshaft. (inflorescence position)
- infrapetiolar bud** An axillary bud surrounded by a petiole base, e.g. *Platanus*, sycamore. (bud type)
- infructescence** The complete inflorescence at the stage of fruiting. (fruit part)
- infundibular** Funnel-shaped; with a narrow base and greatly expanded apex, e.g. *Ipomoea*. (perianth type)
- ingroup** The study group as a whole in a phylogenetic analysis.
- inner bark** *Secondary phloem*.
- inserted** With stamens included within the perianth. Syn: *cryptantherous*. (stamen insertion)
- integument** A sheath or flap of tissue that surrounds the megasporangium (nucellus) of an ovule and develops into the seed coat of the seed.
- intercalary meristem** An indeterminate (having potentially continuous growth), basal or sub-basal region of actively dividing cells.
- interfoliar** Descriptive of a palm inflorescence that is positioned among the leaves of the crownshaft. (inflorescence position)
- International Code of Botanical Nomenclature** The standardized system of rules for naming plants, algae, fungi, and or animals traditionally treated as fungi, governing specific names assigned to taxa and the endings that denote taxon rank, and utilized for naming new taxa and determining the correct name for previously named taxa. Abbr: *ICBN*.
- internode** (a) The region between two adjacent nodes of a shoot. (stem/shoot parts, twig part) (b) A cladogram lineage that spaces between two nodes (points of divergence). Syn: *stem*.

- intine** The innermost layer of a pollen grain wall, composed primarily of cellulose and pectines.
- intrastaminal disk** A discoid or doughnut-shaped structure arising from the receptacle at the inside of the stamens and/or base of the ovary; may be nectar-bearing. (nectariferous disk) (flower part)
- intravaginal (axillary) squamules** Trichomes found in the axils of sheathing leaves, possibly functioning in secreting a protective mucilage, e.g. many Alismatales. (trichome type)
- introgression** Hybridization between two species followed by backcrossing to one or both parents.
- introrse** Dehiscing inward, toward the flower center. (anther dehiscence direction)
- involucl** A group or cluster of bracts subtending a unit of an inflorescence. (inflorescence part)
- involucral bract** *Phyllary*. (leaf structural type)
- involucre** A group or cluster of bracts subtending an inflorescence. Adj: *involucrate*. (inflorescence part)
- involute** (a) Margins or outer portion of sides rolled inward or upward over adaxial surface (longitudinal posture, margin). (b) Valvate with each perianth part induplicate, folded longitudinally inward along central axis. (perianth aestivation)
- iridoid** A secondary chemical compound characteristic of many Asterids.
- irregular** *Zygomorphic, bilateral*. (symmetry)
- isomerous** Having the same number of parts in different whorls. (merosity, perianth merosity)
- isomorphic** Appearing identical, e.g. the gametes of some green plants.
- isopolar** Pollen polarity in which the two polar hemispheres are the same but can be distinguished from the equatorial region.
- isotype** A duplicate specimen of the holotype, collected at the same time by the same person from the same population.
- iteropary** Referring to plants that reproduce more than one time in the life of the plant, typically in regular cycles. Adj: *iteroparous*.
- jacket layer** *Antheridial wall; archegonial wall*.
- jackknife/jackknifing** A method of evaluating cladogram robustness that reanalyzes the data of the original character \times taxon matrix by selecting (resampling) characters at random, such that a given character can be selected only once, the resultant resampled data matrix being smaller than the original.
- jaculator** Funiculi of the seeds that are modified into rigid, often hook-shaped structures that function to disperse the seeds by a catapulting mechanism, characteristic of the Acanthaceae. Syn: *retinaculum*.
- key/taxonomic key** An identification device, consisting of contrasting statements used to narrow down the identity of a taxon.
- Kranz anatomy** A leaf anatomy in which chloroplasts of the bundle sheath cells are typically much larger than those of the mesophyll cells, correlated with C4 photosynthesis.
- labellum** A modified, typically expanded, median petal, tepal, or perianth lobe, such as in the Orchidaceae. (perianth part)
- lacerate** With sinuses irregularly cut, lobes appearing torn. (division)
- lacinate** Cut into narrow, ribbonlike segments. (division)
- laevigate** Lustrous, polished. (epidermal excrecence)
- lamellar** An exine wall structure having stacked, tangentially oriented, planar structures, often constituting the inner wall layer.
- lamina** *Blade*. (leaf part)
- laminar** (a) With a dorsiventrally flattened, leaflike structure bearing the thecae. (stamen type) (b) With ovules arising from the surface of the septae. (placentation)
- lanate** *Villous*. (vestiture)
- lanceolate** Margins curved, widest near base, length:width ratio between 6:1 and 3:1. (shape)
- lance-ovate** Margins curved, widest near base, length:width ratio between 3:1 and 2:1. (shape)
- land plants** A monophyletic group of eukaryotes united by an outer cuticle, specialized gametangia antheridia and archegonia and an intercalated diploid phase in the life cycle, including the embryo [embryophytes/Embryophyta].
- lateral** (a) *Axillary* (position, inflorescence position). (b) Style arising at the side of an ovary. (style position)
- lateral bud** *Axillary bud* (bud type)
- lateral meristem** A cylindrical sheath of cells, functioning in secondary growth, that increases width or girth of stems or roots in woody plants; includes the vascular cambium and cork cambium.
- lateral root** A root that arises from another root, derived endogenously from the pericycle. (root type)
- lateral vein** *Ventral vein*.
- laticifer** Cells located in the periphery of some tissues that secrete and store latex, functioning to deter herbivory and to seal and protect plant tissue upon wounding.
- latrorse** Dehiscing laterally relative to the flower center. (anther dehiscence direction)
- layer** One of the ecological criteria of plant communities based on height and plant habit, including the canopy, subcanopy, shrub or subshrub layer, and herb layer.
- lead** One of the two contrasting statements in a dichotomous key.
- leaf** A generally dorsi-ventrally flattened organ, usually functioning in photosynthesis and transpiration, either gametophytic (in mosses and some liverworts) or sporophytic (in vascular plants), often variously modified. (plant part)
- leaf gap** A region of nonvascular parenchyma tissue interrupting the vasculature of the stem at a node, associated with euphylls.
- leaf primordium** An immature leaf of the shoot. (stem/shoot parts)
- leaf scar** A mark indicating the former place of attachment of a leaf. (twig part)
- leaf spine** A sharp-pointed leaf, e.g. cactus spines or glochidia. Cf: *prickle; thorn*. (leaf structural type)
- leaflet** A distinct and separate segment of a leaf. (leaf part)
- leaflet spine** A sharp-pointed leaflet, e.g. some palms, such as *Phoenix*. (leaf structural type)
- lectotype** A specimen that is selected from the original material to serve as the type when no holotype was designated at the time of publication, if the holotype is missing, or if the original type consisted of more than one specimen or taxon.
- legitimate name** A name that is in accordance with the rules of the International Code of Botanical Nomenclature.
- legume** A dry, dehiscent fruit derived from one carpel that splits along two sutures, e.g. Fabaceae. (fruit type)
- lemma** The outer and lower bract at the base of the grass floret. (leaf structural type)
- lenticel** A pore in the bark, generally functioning in gas exchange. (twig part)
- lenticular** Lens-shaped; disk-shaped with two convex sides. (shape)

- lepidote** Covered with scales or scalelike structures. (vestiture)
- leptoid** A specialized cell that functions in sugar conduction in some mosses.
- leptosporangium** The sporangia of the leptosporangiate ferns (Polypodiales), characterized by developing from a single cell and having a single layer of cells making up the sporangium wall. Pl: *leptosporangia*.
- liana/liane** A woody, perennial vine, in tropical forests often a component of the canopy layer. (plant habit)
- light reactions** A series of biochemical reactions of photosynthesis in plants, occurring in the thylakoid membranes of the chloroplast and requiring light as an energy source, during which water is broken down into hydrogen ions, electrons, and molecular oxygen, and producing high-energy ATP and NADPH₂, which are utilized in the dark reactions.
- lignin** A complex polymer of phenolic compounds that impregnates the *secondary cell wall* of some cells (including tracheary elements and sclerenchyma), functioning to impart strength and rigidity to the wall.
- lignotuber** *Burl*. (stem/shoot type)
- ligulate** Strap- or tongue-shaped; flattened and somewhat oblong in shape, e.g. ray flowers of some Asteraceae. (perianth type, shape)
- ligule** (a) A small appendage on the upper (adaxial) side of the leaf, near the leaf base, found in the Selaginellaceae and Isoetaceae of the lycophytes. (b) An outgrowth or projection from the top of a leaf sheath at its junction with the blade, as in the Poaceae. (leaf part)
- limb** The expanded portion of the corolla or calyx above the tube, throat, or claw. (perianth part)
- lineage** *Clade*.
- linear** (a) With margins straight, parallel, length : width ratio between 12:1 and 6:1. (shape) (b) Stigmas or stigmatic tissue long and narrow in shape. (stigma/stigmatic region type)
- linear tetrad** A tetrad in which the four pollen grains are arranged in a straight line, e.g. *Typha*.
- lip** Either of two variously shaped parts into which a calyx or corolla is divided, usually into upper (posterior) and/or lower (anterior) lips, each lip often composed of one or more lobes, e.g. Lamiaceae, Orchidaceae. Cf: *labellum*. (perianth part)
- lithocyst** A specialized cell that contains a cystolith. Cf: *cystolith*.
- lobe** (a) A segment of a synsepalous calyx or sympetalous corolla. (perianth part) (b) A segment of a divided leaf. (leaf part)
- lobed** (a) Sinuses extending (pinnately or palmately) one eighth to one fourth the distance to midrib, midvein, or vein junction. (b) A general term meaning having lobes. (division)
- locule** An ovary cavity, bounded by ovary walls and septa. (gynoecium part) (b) A compartment of the anther, usually the result of two microsporangia fusing within a theca. (anther part)
- loculicidal capsule** A capsule in which longitudinal lines of dehiscence are radially aligned with the locules. (fruit type)
- lodicule** One of the (2-3) modified perianth parts of a grass (Poaceae) flower, which collectively upon swelling function to open the floret by separating the lemma from palea. (perianth type)
- loment** A secondarily modified legume that splits transversely into 1-seeded segments. (fruit type)
- long-branch attraction** A condition in which taxa with relatively long branches (having numerous character state changes) tend to come out as close relatives of one another in a phylogenetic analysis because of random effects.
- longitudinal dehiscence** Dehiscing along a suture parallel to the long axis of the thecae. (anther dehiscence type)
- longitudinal posture** Placement of margins with respect to a horizontal plane. (disposition)
- lower epidermis** The abaxial epidermis of a leaf.
- lumen** The space between muri in a reticulate pollen grain. Pl: *lumina*.
- lycophyll** The sporophytic leaf of the lycophytes, characterized by an intercalary meristem, having a single vein, and lacking a gap in the vasculature of the stem. Adj: *lycophyllous*. Syn: *microphyll*. (leaf structural type)
- lyrate** Pinnatifid, but with a large terminal lobe and smaller basal and lateral lobes. (shape)
- maculate** Spotted; with small spots on a more or less uniform background. (color pattern)
- majority consensus tree** A consensus tree in which only those clades that are retained 50% or more of the time are retained (i.e. not collapsed to a polytomy).
- male** (a) Individual with male reproductive organs only. (plant sex) (b) *Staminate*. (flower sex)
- male gametophyte** A gametophyte that bears only antheridia. Syn: *microgametophyte*.
- male sporophyll** A sporophyll that bears one or more microsporangia. Syn: *microsporophyll*.
- malpighian** Trichomes with two arms arising from a common base, e.g. Malpighiaceae. (trichome type)
- manual** *Flora*.
- marcescent** Ephemeral, but with persistent remains; withering persistent. (duration)
- marginal** (a) With the placenta along the margin of a unilocular (simple) ovary, e.g. Fabaceae. (placentation) (b) With three or more primary veins diverging from one point and reaching the blade margin, a subcategory of actinodromous. (leaf venation)
- massula** A group of fused pollen grains in large, often irregular numbers, but less than an entire theca. Pl: *massulae*.
- masticatory** A substance that is chewed by humans, usually for its pleasing or euphoric effects, e.g. peyote, *Lophophora williamsii*.
- maturation** (a) Acquisition of the mature structural and functional features of a cell following cell expansion. Syn: *specialization*. (b) Relative time of development of parts. (temporal phenomena)
- matutinal** In the morning, typically with respect to when flowers open. (periodicity)
- maximum likelihood** A method of phylogenetic inference that considers the probability, based on some selected model of evolution, that each tree explains the data.
- mealy** Covered with small, fine granules. (epidermal excrescence)
- median vein** *Dorsal vein*.
- megagametogenesis** The process of development of the female gametophyte from the haploid product(s) of meiosis.
- megagametophyte** *Female gametophyte*.
- megaphyll** *Euphyll*.
- megasporangium** A female sporangium, within which megasporocytes undergo meiosis to produce haploid megaspores. Pl: *megasporangia*. Syn (in seed plants only): *nucellus*. (plant part)
- megaspore** A female spore, produced via meiosis in the megasporangium and giving rise to a female gametophyte.
- megaspore mother cell** *Megasporocyte*.
- megasporocyte** A cell within the megasporangium that undergoes meiosis, forming four megaspores. Syn: *megaspore mother cell*.
- megasporogenesis** The process of development of megaspores from the megasporocyte.

- megasporophyll** *Female sporophyll.*
- melittophily** Pollination by bees. Adj: *melittophilous*. Syn: *hymenopterophily*.
- membranous** Thin and somewhat translucent; membranelike. (texture)
- mericarp** A portion of a fruit that separates from the ovary as a distinct unit that completely encloses the seed(s). (fruit part)
- meristem** A region of actively dividing cells.
- merosity** Number of parts per whorl or cycle. (number)
- mesarch** An orientation of xylem maturation in which the protoxylem is surrounded by metaxylem within the vascular tissue, as can occur in siphonosteles.
- mesocarp** A middle wall layer of the pericarp, if the latter is divided into layers. (fruit part)
- mesophyll** The region of a sporophytic leaf between the outer epidermal layers and exclusive of the vasculature, containing the chlorophyllous cells.
- mesophytic** Having an intermediate texture, between coriaceous and membranous (texture)
- metaxylem** The xylem of a group of tracheary elements that matures later, consisting of larger-diameter cells.
- microfibril** Microscopic fiberlike units of intertwined cellulose molecules, forming a meshwork within the cell wall.
- microgametogenesis** The process of development of pollen grains from microspores.
- microgametophyte** *Male gametophyte.*
- microhair** A very small trichome, as in the three-celled, glandular microhairs of the Commelinaceae.
- microphyll** Essentially equivalent to a lycophyll.
- micropylar** Describing the region of an ovule that is near the micropyle.
- micropylar** Referring to the distal region of the ovule, near the micropyle.
- micropyle** A small pore at the distal end of the integument of a seed, functioning as the site of entry of pollen grains, or in angiosperms of pollen tubes.
- microsatellite** A region of DNA that contains **tandem repeats**, short (usually 2-5) repeats of nucleotides.
- microsporangium** A male sporangium, within which microspores undergo meiosis to produce haploid microspores. Pl: *microsporangia*. (plant part)
- microspore** A male spore, produced via meiosis in the microsporangium and giving rise to a male gametophyte.
- microsporogenesis** The process of development of microspores from microsporocytes.
- microsporophyll** A sporophyll bearing one or more microsporangia; a male sporophyll.
- middle lamella** A pectic-rich layer formed between the plasma membrane of adjacent cells in land plants, functioning to bind adjacent cells together.
- middle layers** Anther wall layers that may occur between the endothecium and tapetum.
- midrib** The central, main vein of the blade of a simple leaf or of a compound leaf in some palms. Syn: *costa*. (leaf part)
- midvein** The central, main vein of the blade of a leaflet. (leaf part)
- mitochondrion** A double membrane-bound, cellular organelle with invaginations called cristae that function in the electron transport reactions of respiration. Pl: *mitochondria*.
- mixed bud** A bud that produces both flowers and leaves. (bud type)
- mixed craspedodromous** Pinnate venation in which some secondary veins terminate at the margin, but with many terminating away from the margin. (leaf venation)
- monad** A single, unfused pollen grain.
- monadelphous** With one group of stamens connate by their filaments, e.g. Malvaceae. (stamen fusion)
- monistichous** Alternate, with points of attachment in one vertical row/rank, e.g. Costaceae. (arrangement)
- monocarpic** A perennial or annual plant, flowering and fruiting once, then dying. (duration)
- monochasium** A determinate inflorescence that develops along one axis only. (inflorescence type)
- monocotyledonous** A type of anther wall development in which only the inner secondary parietal cell layer divides to yield the tapetum and a single middle layer.
- monocyclic** *Uniseriate*. (cyclic, perianth cyclic)
- monoecious** Having unisexual flowers, both staminate and pistillate on the same individual plant. (plant sex)
- monograph** A detailed taxonomic study of all species and infraspecific taxa of a given taxonomic group.
- monomorphic character** A character that is invariable in character state values within an OTU.
- monophyletic/monophyly** Referring to a group that consists of a common ancestor plus all (and only all) descendants of that ancestor.
- monopodial** A branching pattern in which a given axis is derived from a single apical meristem. (stem branching pattern)
- monosporic** Megasporeogenesis in which meiosis of the megasporocyte nucleus results in the formation of four haploid megaspore nuclei, followed by cytokinesis, resulting in four megaspore cells, only one of which contributes to the female gametophyte.
- monosulcate** A pollen grain with a single, sulcate aperture.
- monothecal** Anther with one theca and typically two microsporangia. Cf: *bisporangiate*. (anther type)
- monoulcerate** A pollen grain with a single, ulcerate aperture.
- morphocline** *Transformation series.*
- movement hercogamy** A type of hercogamy involving movement of floral parts, e.g. the rapid closure of the stigmas upon their being touched by a potential animal pollinator.
- mucilage ducts/canals** Specialized cells that secrete mucilage.
- mucronate** With a stiff, straight apical process, length : width ratio 1:1 to 3:1. (apical process)
- mucronulate** With a stiff, straight apical process, length : width ratio \leq 1:1. (apical process)
- multicellular** Trichomes having two or more cells. (trichome type)
- multiple fruit** A fruit derived from two or more flowers. (fruit type)
- multiseriate** (a) Perianth composed of three or more distinct whorls. (perianth cyclic) (b) Trichomes having more than one vertical row of cells. (trichome type) (c) Rays in wood that are made up of many vertical rows of cells.
- multistate character** A character with three or more character states.
- muricate** Having coarse, radially elongate, rounded protuberances. (epidermal excrescence)
- murus** The structural element of a reticulate pollen grain. Pl: *muri*.

- mycorrhiza** A symbiotic association between a fungus and roots, functioning to increase absorptive surface area and mineral uptake. Pl: *mycorrhizae*. (root part)
- mycorrhizae** A symbiotic association between the root of a vascular plant and a fungus.
- mycotrophic** Obtaining nutrition from mycorrhizal fungi in the soil, as in some Ericaceae.
- naked bud** A bud lacking surrounding protective scales, e.g. *Viburnum*, Caprifoliaceae. (bud type)
- narrowly acute** Apical margins approximately straight, the intersection angle $<45^\circ$. (apex)
- narrowly cuneate** Basal margins approximately straight, the intersection angle $<45^\circ$. (base)
- narrowly elliptic** Margins curved, widest near midpoint, length : width ratio between 6:1 and 3:1. (shape)
- narrowly oblong** Margins straight, parallel, length : width ratio between 6:1 and 3:1. (shape)
- narrowly triangular** Three-sided, length : width ratio between 6:1 and 3:1. (shape)
- natural selection** The directed and nonrandom genetic modification of a population or species, in which genetic changes that result in an increase in survivorship and/or reproduction are contributed to the next generation more.
- neck** The distal, narrow extension of the sterile jacket cells of the archegonium, through which a sperm cell must travel to fertilize the egg.
- neck canal cells** Cells located within the neck of the archegonium that break down and are secreted from the pore of the neck at maturity.
- nectariferous disk** A nectary consisting of a disklike or doughnut-shaped mass of tissue surrounding the ovary base or top; may be inner to (intrastaminal), beneath (staminal), or outer to (extrastaminal) the androecium. (flower part, nectary type)
- nectary** A group of specialized cells that secrete sugar- (or protein-) rich fluids to the outside, as a reward for pollination or protection. (flower part)
- neotony** A type of paedomorphosis caused by a decrease in the rate of development of a feature.
- neotype** A specimen derived from a nonoriginal collection that is selected to serve as the type as long as all of the material on which the name was originally based is missing.
- nerve** *Vein*. (leaf part)
- netted** With ultimate veinlets forming a reticulum or netlike pattern. Syn: *reticulate*. (leaf venation)
- network** *Unrooted tree*.
- nexine** The inner layers of the exine, including both endexine and the foot-layer of the ectexine.
- nitid** Appearing lustrous, polished. Syn: *shining*. (epidermal excrescence)
- nocturnal** Occurring during the night, typically with respect to when flowers open. (periodicity)
- node** (a) The point of attachment of a leaf to a stem. (stem/shoot parts) (b) The region of stem at which leaf, leaves, or branches arise. (twig part) (c) The point of divergence of one clade into two; the point in time and space of the most common ancestor of the two divergent clades.
- node-based** A type of phylogenetic classification in which a node (common ancestor) of the cladogram (and all descendants of that common ancestor) serves as the basis for grouping.
- nomen conservandum** Meaning a conserved name. Abbr: *nom. cons.*
- nomen novum** Meaning a new name. Abbr: *nom. nov.*
- nomen nudum** Meaning a name published without a description or diagnosis, making the name invalid. Abbr: *nom. nud.*
- nomenclatural type** A specimen that acts as a reference for a scientific name, upon which the name is based. Syn: *type*; *type specimen*.
- nomenclature** The formal naming of taxa according to some standardized system; for plants, algae, fungi, and or ganisms traditionally treated as fungi, governed by the International Code of Botanical Nomenclature.
- non** Latin for not.
- nondecussate** Opposite leaves or other structures (e.g. leaflets) not at right angles to the preceding pair; may be superficially the result of stem twisting. (arrangement)
- nondisjunction** An irregularity during meiosis in which homologous chromosomes do not segregate, which may result in the production of gametes that are unreduced, i.e. have two sets of chromosomes.
- nonendospermous** *Exalbuminous*. (seed endosperm type)
- nonporous** Referring to wood having only tracheids.
- nucellar beak** A proliferation of cell divisions of the nucellus at the micropylar region of the ovule.
- nucellus** Term for the megasporangium of a seed. Adj: *nucellar*.
- nuclear endosperm** An endosperm in which the early mitotic divisions are not followed by cytokinesis.
- nucleus** A double membrane-bound, cellular organelle that contains DNA.
- number** Whether a Latin name is singular or plural.
- nut** A one-seeded, dry indehiscent fruit with a hard pericarp, usually derived from a 1-loculed ovary. (fruit type)
- nutlet** A small nut, e.g. mericarps of the Lamiaceae. (fruit type)
- nutritive tissue** Tissue that surrounds or abuts the embryo of a seed and that consists of female gametophyte (in nonangiosperms) or endosperm (in angiosperms).
- obdiplostemonous** Stamens in two whorls, the outer opposite petals, inner opposite the sepals, e.g. Simaroubaceae. (stamen position)
- oblanceolate** Margins curved, widest near the apex, length : width ratio 6:1 to 3:1. (shape)
- oblance-ovate** Margins curved, widest near the apex, length : width ratio 3:1 to 2:1. (shape)
- oblate** A pollen grain in which the P/E ratio is less than about 0.8.
- oblique** With an asymmetrical apex or base. (apex, base)
- oblong** Margins straight, parallel, length : width ratio 2:1 to 3:2. (shape)
- obovate** Margins curved, widest near apex, length : width ratio 2:1 to 3:2. (shape)
- oburator** A protuberance of tissue, typically arising from the funiculus or placenta at the base of the ovule, e.g. Euphorbiaceae.
- obtuse** Apical or basal margins approximately straight, intersection angle $>90^\circ$. (apex, base).
- ocrea** A specialized, scarious, sheathlike structure arising above the node in some members of the family Polygonaceae, interpreted as modified stipules. (leaf part)
- oil** A type of triglyceride compound that may function as high-energy storage compounds or secretion products; a type of ergastic substance.

- oil bodies** Oil-containing structures found within certain cells of most liverworts.
- oil ducts/canals** Specialized cells that secrete oil.
- onagrad type** *Crucifer type*.
- ontogenetic sequence** The discrete stages of the developmental sequence of a given feature.
- ontogenetic trajectory** A plot of developmental change as a function of time.
- ontogeny** The developmental sequence of a given feature.
- oogamy** A type of sexual reproduction in which one gamete, the egg, becomes larger and nonflagellate and the other gamete, the sperm cell, remains relatively small and flagellate; found in all land plants and independently evolved in many other eukaryotes.
- operational taxonomic unit (OTU)** One of the individual, unit taxa of a phylogenetic analysis.
- operculate** Having calyx and corolla fused into a cap that falls off as a unit, e.g. *Eucalyptus*. (perianth type)
- operculum** An apical lid, as in the capsule of most mosses that falls off during spore release.
- opposite** With two leaves or other structures per node, on opposite sides of stem or central axis. (arrangement)
- orbicular** Circular in outline; margins curved, length : width ratio approximately 1:1. Adj: *circular* (shape)
- ordered** Referring to a transformation series in which the character states occur in a predetermined sequence.
- organelles** Structural, membrane-bound units of the cell that provide some vital metabolic function.
- orientation** Referring to the angle relative to a central, usually vertical, axis. (disposition)
- ornithophily** Pollination by birds. Adj: *ornithophilous*.
- ortho-amphitropous** An ovule type in which the vasculature is straight, leading from the funiculus base to the middle of the nucellus, with the nucellus bent sharply in the middle along both the lower and upper sides, often with a basal body present.
- ortho-campylotropous** An ovule type in which the vasculature is straight, leading from the funiculus base to the middle of the nucellus, with the nucellus bent only along the lower side, with no basal body.
- orthographia conservanda** Meaning a conserved spelling in a scientific name. Abbr: *orth. cons.*
- orthotropous** A type of ovule in which no curvature takes place during development, the micropyle being positioned opposite the funiculus base.
- outbreeding** The transfer of gametes from one individual to another, genetically different individual. Syn: *outcrossing*; *allogamy*; *xenogamy*.
- outcrossing** *Outbreeding*.
- outer bark** *Periderm*.
- outgroup** A taxon or group of taxa that is not a member of the ingroup.
- ovary** The part of the pistil containing the ovules. (gynoecium part)
- ovate** Margins curved, widest near base, length : width ratio 2:1 to 3:2. (shape)
- ovule** An immature seed, prior to fertilization; a megasporangium enveloped by one or more integuments. (plant part, gynoecium part)
- ovuliferous scale** The modified lateral branch system of a conifer cone that bears one or more seeds.
- P/E ratio** The ratio of the polar diameter to the equatorial diameter.
- pachycaul** An erect, woody, trunklike stem that is swollen basally, the swollen region functioning in storage of food reserves or water. (stem/shoot type)
- padlike nectary** Developing as a discrete pad of tissue extending partway around the base of the flower. (nectary type)
- paedomorphosis** A type of heterochrony in which the mature or adult stage of the derived ontogenetic sequence resembles a juvenile ontogenetic stage of the ancestral condition.
- palea** The inner and upper bract at the base of the grass floret, or the bracts subtending flowers in some Asteraceae, e.g. tribe Heliantheae. Pl: *paleae*. (leaf structural type)
- palinactinodromous** With three or more primary veins diverging from one point, the primary veins having additional branching above their main point of divergence. (leaf venation)
- palisade mesophyll** The usually upper (adaxial), columnar cells of the mesophyll of some leaves.
- palmate/palmately compound** A compound leaf with four or more leaflets arising from a common point, usually at the end of the petiole. (leaf type)
- palmately netted** Netted, with four or more primary veins arising from a common basal point. (leaf venation)
- palmately veined** With four or more primary veins arising from a common basal point. (leaf venation)
- palmate-netted** Palmately veined, with four or more primary veins arising from a common basal point, the ultimate veins forming a fine reticulum. (leaf venation)
- palmate-parallel** With several primary veins arising from one point, the adjacent secondary veins that are parallel to these having transverse, interconnecting veins, e.g. f an palms. (leaf venation).
- palmate-ternate** Ternate, with the three leaflets joined at a common point. (leaf type)
- palmatifid** Palmately lobed to divided. (division)
- palmatisect** Palmately divided, almost into discrete leaflets but confluent at the lobe bases. (division)
- palynology** The study of spores and pollen grains.
- pandurate** Violin-shaped, obovate with the side margins concave. (shape)
- panicle** An indeterminate inflorescence, consisting of several branched axes bearing pedicellate flowers. (inflorescence type)
- pantoporate** A pollen grain aperture type in which pori occur globally on the pollen grain surface.
- papilionaceous** A flower with one large posterior petal (banner or standard), two inner, lateral petals (wings), and two usually apically connate lower petals (keel); floral structure of the Faboideae-Fabaceae. (perianth type)
- papillate** Having minute, rounded protuberances. Syn: *tuberculate*; *verrucate*. (epidermal excrescence)
- pappus** The calyx of the Asteraceae, modified as awns, scales, or capillary bristles.
- parallel** With primary or secondary veins essentially parallel to one another, generally converging at the apex, the ultimate veinlets transverse. (leaf venation)
- parallelism** *Convergence*.
- parallelodromous** *Parallel*. (leaf venation)
- paraphyletic/paraphyly** Referring to a group that includes a common ancestor plus some, but not all, descendants of that common ancestor.

- paratype** A specimen cited in a publication but that is neither a holotype, isotype, or syntype.
- parenchyma** Cells that, at maturity, are alive and potentially capable of cell division, are isodiametric to elongate in shape, and have a primary cell wall only (rarely with secondary wall), forming a solid mass of tissue and functioning in metabolic activities and in wound healing and regeneration.
- parietal** With the placentae on the inner wall or on intruding partitions of a unilocular, compound ovary, e.g. *Violaceae*. (placentation)
- parietal cell** The outer cell formed if the archesporial cell of an ovule divides to form an inner megasporocyte.
- parietal-axile** With the placentae at the junction of the septum and pericarp of a 2- or more loculate ovary, e.g. *Brassicaceae*. (placentation)
- parietal-septate** With placentae on the inner ovary walls but within septate locules, as in some *Aizoaceae*. (placentation)
- paripinnate/paripinnately compound** A pinnately compound leaf without a terminal leaflet, typically even-pinnate. (leaf type)
- parsimony analysis** *Principle of parsimony*.
- parted** With sinuses extending (pinnately or palmately) one half to three quarters of the distance to the midrib, midvein, or vein junction. (division)
- parthenogenesis** Development of an embryo from a cell of an abnormal, diploid female gametophyte, such as a diploid egg.
- partial inflorescence** *Unit inflorescence*. (inflorescence part)
- patent** *Horizontal; divaricate; divergent*. (orientation)
- peat** Fossilized and partially decomposed *Sphagnum* (peat moss).
- pectinate** Comblike; pinnately divided with close, narrow lobes. (division)
- pedicel** A flower stalk. Adj: *pedicellate*. (flower part; inflorescence part)
- peduncle** The stalk of an entire inflorescence. Adj: *pedunculate*. (inflorescence part)
- pellucid** Having translucent spots or patches. (color pattern)
- peltate** (a) Having a stalk attached away from the margin of a flattened structure, e.g. a petiole attached on the underside of blade. (base) (b) Trichomes with stalk attached on the underside of a disk-shaped apical portion. (trichome type)
- pendant/pendulous** Hanging downward loosely or freely (orientation)
- pendulous** *Apical*. (placentation)
- penni-parallel** With secondary veins arising along the length of a single primary vein region, the former essentially parallel to one another and interconnected by transverse veins. Syn: *pinnate-parallel*. (leaf venation)
- pentamerous** Having a whorl with five members. (merosity, perianth merosity)
- pepo** A nonseptate, fleshy fruit with parietal placentation and a leathery exocarp derived from an inferior ovary, e.g. *Cucurbitaceae*. (fruit type)
- peramorphosis** A type of heterochrony in which ontogeny passes through and goes beyond the stages or trajectory of the ancestral condition.
- perennial** A plant living more than 2 years. (duration)
- perfect** (a) *Bisexual*. (flower sex) (b) With lateral primary veins covering at least two thirds of the leaf blade area, a subcategory of actinodromous and of acrodromous. (leaf venation)
- perfoliate** A leaf blade that is sessile with the base completely surrounding the stem. (leaf attachment)
- perforation plate** The contact area of two adjacent vessel members, may be compound, if composed of several pores or simple if composed of a single opening.
- perforation plate** The region of one or more perforations at the end wall of a vessel member, where one cell makes contact with another; may be simple or compound.
- perianth** The outermost, nonreproductive group of modified leaves of a flower, composed of the combined calyx and corolla or of tepals. Syn: *perigonium*. (flower part)
- pericarp** The fruit wall, derived from the mature ovary wall, sometimes divisible into layers: endocarp, mesocarp, and exocarp. Syn: *rind*. (fruit part)
- pericycle** A cylindrical sheath of parenchyma cells just inside the endodermis, functioning as the site of resumed meristematic growth, forming a secondary root or (in woody plants) contributing to the vascular cambium.
- periderm** The cork cambium and its derivatives: phelloderm and cork tissue. Syn: *outer bark*.
- perigonal nectary** Nectaries on the perianth, usually at the base of sepals, petals, or tepals. (nectary type)
- perigonium** *perianth*. (flower part)
- perigynous** Hypanthium present, attached at base of ovary, with sepals, petals, and stamens attached to hypanthium rim, the ovary superior. (perianth/androecial position)
- perine layer** A thick, outer layer in the spores of mosses.
- periodicity** Referring to periodically repeating phenomena. (temporal phenomena)
- periplasmodial** *Amoeboid*.
- perispermous** Having a seed storage tissue in which the chalazal nucellar cells enlarge and store energy-rich compounds.
- peristome teeth** Hygroscopic, toothlike structures that occur in a whorl along the margin of the opening of a moss capsule and that function in spore release.
- personate** Two-lipped, with the upper arched and the lower protruding into corolla throat, e.g. *Antirrhinum*. (perianth type)
- petal** An individual member or segment of the corolla, typically (nongreen) colored and functioning as an attractant for pollination. (flower part, perianth part)
- petalostemonous** *Epipetalous*. (stamen fusion)
- petiolar spine** A sharp-pointed leaf petiole, e.g. *Fouquieria* spp. Cf: *thorn; prickle*. (leaf structural type)
- petiole** A leaf stalk. Adj: *petiolate*. Syn: *stipe*. (leaf part)
- petiolule** A leaflet stalk. Adj: *petiolulate*. (leaf part)
- phalaenophily** Pollination by moths. Adj: *phalaenophilous*.
- phaneranthorous** *Exserted*. (stamen insertion)
- phanerocotylar** *Epigeous*. (seed germination type)
- phelloderm** The inner layers of cells produced by the cork cambium.
- phenetic** A classification system in which taxa are grouped by some measure of overall similarity.
- phenogram** A branching diagram representing a phenetic classification.
- ploem** A tissue composed of specialized sieve elements plus some parenchyma and often some sclerenchyma, functioning in conduction of sugars.
- photosynthesis** The series of biochemical reactions in which light energy is used to produce high-energy compounds, in land plants involving reactions of carbon dioxide and water to produce polysaccharides.

- phyllary** One of the involucre bracts subtending a head, as in the Asteraceae. Syn: *involucre bract*. (leaf structural type)
- phylloclad** *Cladode*; *cladophyll*. (stem/shoot type)
- phylloide** A leaf consisting of a flattened bladelike petiole. (leaf structural type)
- phylogenetic** Referring to a classification that is based on evolutionary history, or pattern of descent.
- phylogenetic systematics** A methodology for inferring the pattern of evolutionary history of a group of organisms by grouping taxa based on apomorphies. Syn: *cladistics*.
- phylogenetic tree** *Cladogram*.
- phylogeny** The evolutionary history or pattern of descent of a group of organisms.
- pilate** With a long terete stalk terminating in a globose or ellipsoid apical thickening. (shape)
- pilate-glandular** Having a glandular cell atop an elongate basal stalk. (trichome type)
- pilose** With soft, straight to slightly shaggy trichomes at right angles to the surface. (vestiture)
- pin** Flowers with a long style and short stamens, found in distylous flowers.
- pinna** The first discrete leaflets or blade divisions of a fern leaf. Pl: *pinnae*. (leaf part)
- pinnate/pinnately compound** A compound leaf with leaflets arranged oppositely or alternately along a central axis, the rachis. (leaf type)
- pinnately veined** With secondary veins arising along length of a single primary vein, the latter a midrib or leaflet midvein. (leaf venation)
- pinnate-netted** Pinnately veined, with secondary veins arising along length of a single primary vein, the latter a midrib or midvein, the ultimate veins forming a fine reticulum. (leaf venation)
- pinnate-parallel** *Penni-parallel*. (leaf venation)
- pinnate-ternate** Ternately compound, with the terminal leaflet arising from the tip of a rachis, evolutionarily derived from a pinnately compound leaf, e.g. some Fabaceae. (leaf type)
- pinnatifid** Pinnately lobed to divided (division)
- pinnatisect** Pinnately divided, almost into discrete leaflets but confluent at the midrib. (division)
- pinnule** The ultimate divisions or leaflets of a leaf, usually applied to ferns. (leaf part)
- piperad type** A type of embryo development in which the zygote divides longitudinally (i.e. parallel to the axis of the female gametophyte and nucellus), thus not forming a basal and terminal cell.
- pistil** That part of the gynoecium composed of an ovary, one or more styles (which may be absent), and one or more stigmas. (flower part, gynoecium part)
- pistillate** Flowers unisexual, with carpel(s) only, lacking fertile stamen(s). Syn: *female*. (flower sex)
- pit** A hole in a secondary cell wall that functions in cell-to-cell communication during development and that may function in water conduction in some tracheary elements.
- pit pairs** Adjacent holes in the lignified secondary cell walls of two adjacent cells.
- pitcher leaf** A leaf shaped like a container that bears an internal fluid and that functions in the capture and digestion of small animals, e.g. leaves of *Darlingtonia*, *Nepenthes*, *Sarracenia*, pitcher plants. (leaf structural type)
- pith** The central, mostly parenchymatous tissue, internal to vasculature of siphonosteles and eusteles and within the vascular cylinder of some roots. (root parts, stem/shoot parts)
- pit-pair** Pits of adjacent cells occurring opposite one another, functioning in allowing communication between cells during their development and differentiation.
- placenta** The ovule-bearing tissue of the ovary. (gynoecium part)
- placental vein** *Ventral vein*.
- placentation** Referring to the position of the ovules within the ovary. (gynoecium, carpel, and pistil)
- plane** (a) Flat, without vertical curves or bends. (transverse posture). (b) With a smooth configuration. Syn: *smooth*. (configuration)
- plant** A group of organisms, defined either by characteristics (possessing photosynthesis, cell walls, spores, and a more or less sedentary behavior), or by phylogenetic relationships, equivalent in this text to the land plants, Embryophyta.
- plant anatomy** The study of tissue and cell structure of plant organs.
- plant habit** General form of plant, including aspects of stem duration, branching pattern, development, and texture. (plant organs)
- plant physiology** The study of metabolic processes in plants.
- plant press** A device used to press and then dry plant specimens, such that they can be effectively used in an herbarium specimen.
- plant sciences** The study of plants, which are here equivalent to the land plants.
- plasma membrane** A phospholipid bilayer with embedded proteins that envelops all cells, functioning as the cell boundary, in cell cell recognition, and in transport of compounds.
- plasmodesmata** Minute pores in the primary cell wall through which membranes traverse between cells, allowing for interchange of compounds between cells; an apomorphy for the Charophytes of the green plants.
- plasmodial** *Amoeboid*.
- plated** Bark split or cracked, with flat plates between fissures. (bark type)
- pleonanthic** An indeterminate shoot that bears lateral flowers but that continues vegetative growth. (stem/shoot type, stem branching pattern, inflorescence type)
- plesiomorphic** *Ancestral*.
- pleurotropous** An ovule position in which the micropyle points to the side.
- pleurotropous-dorsal** A pleurotropous ovule in which the raphe is above.
- pleurotropous-ventral** A pleurotropous ovule in which the raphe is below.
- pligate** With a series of longitudinal folds; pleated. (longitudinal posture)
- plumose** (a) Covered with fine, elongate, ciliate appendages; featherlike, e.g. pappus of some Asteraceae. (bristle type) (b) Stigmas with feathery, trichomelike extensions, often found in wind-pollinated taxa such as Cyperaceae, Poaceae. (stigma/stigmatic region type)
- plurilocular** Referring to an ovary with two or more locules. (locule number)
- pneumatophores** Roots that grow upwardly from soil to air, functioning in obtaining additional oxygen for the plant. (root type)
- polar axis** An extended pollen grain diameter that passes through the center of the original pollen tetrad.

- polar nuclei** In a typical angiosperm female gametophyte, the two haploid nuclei of the central cell that ultimately fuse with a sperm cell (via double fertilization) to form a triploid endosperm cell.
- polar view** Observing a pollen grain from the direction of either pole.
- polarity** The designation of relative ancestry to the character states of a transformation series/morphocline.
- pollen grain** An immature, endosporic male gametophyte of seed plants. (anther part)
- pollen sac** A microsporangium, usually one-half of a theca in an angiosperm anther. (anther part)
- pollen tube** An exosporic process that grows from a pollen grain, functioning as a haustorial organ or to deliver sperm cells to the egg.
- pollenkit** A yellowish or orange, carotenoid-like material adhering to the exine, functioning to stick pollen grains in masses.
- pollinarium** In an orchid flower, the pollinia plus a sticky stalk (derived from either the anther or stigma), the unit of transport during pollination.
- pollination** The transfer of pollen grains from microsporangia either directly to the ovule (in gymnosperms) or to the stigma (in angiosperms).
- pollination chamber** A cavity formed by the breakdown of cells at the distal end of the megasporangium (nucellus) in gymnosperm seeds.
- pollination droplet** A droplet of liquid secreted by the young ovule through the micropyle, functioning to transport pollen grains by resorption.
- pollinium** Anther in which all pollen grains of both thecae (Orchidaceae) or of adjacent thecae (Asclepiadaceae) are fused together as a single mass. Pl: *pollinia*. (anther type)
- polyad** A group of pollen grains that are fused in precise units of more than four, e.g. Mimosoideae.
- polyclave key** A key in which all of the known character states that match a specimen are selected in order to narrow down the identity to a smaller subset of the possibilities.
- polygamous** A plant with both bisexual and unisexual flowers. (plant sex)
- Polygonum type** A type of monosporic female gametophyte in which the megaspore nucleus undergoes three, sequential mitotic divisions, producing eight nuclei and seven cells, the most common and ancestral type in the angiosperms.
- polymerase chain reaction** A procedure for the rapid amplification of DNA using primers, free nucleotides, and DNA polymerase in solution and heating the solution to effect denaturation and replication of the DNA. Abbr: *PCR*.
- polymorphic character** A character that has variable character state values within an OTU.
- polypetalous** *Apopetalous*. (perianth fusion)
- polyphyletic group** A group that consists of two or more, separate monophyletic or paraphyletic groups, each with a separate common ancestor; a group in which the common ancestor of all members is not itself a member of the group.
- polyploidy** A mutation in which offspring have a multiple of some ancestral set of chromosomes.
- polysepalous** *Aposepalous*. (perianth fusion)
- polystichous** *Spiral*. (arrangement)
- polytomy** Three or more lineages arising from a single common ancestor in a cladogram, representing conflicting data or the lack of resolution.
- pome** A fleshy fruit with a cartilaginous endocarp derived from an inferior ovary, with the bulk of the fleshy tissue derived from the outer, adnate hypanthial tissue, e.g. *Malus*, apple. (fruit type)
- population** A group of individuals of the same species that is usually geographically delimited and that typically have a significant amount of gene exchange.
- pore** (a) A specialized, permanent opening in the upper epidermis of the thallus of some liverworts, functioning in gas exchange. (b) A single hole in the primary cell wall of sieve elements that is lined with callose and through which solutes flow in sugar conduction. (c) Vernacular term for a wood vessel, used in the wood industry.
- poricidal** Dehiscing by a pore at one end of the thecae, e.g. Ericaceae. (anther dehiscence type)
- poricidal capsule** A capsule in which the dehiscence occurs by means of pores, e.g. *Papaver*, poppy. (fruit type)
- porose** Referring to vessel cells with more or less circular, porelike perforations.
- porous** Referring to wood that contains vessel cells.
- porus** A pollen grain aperture that is circular to slightly elliptic with a length:width ratio of less than 2:1. Pl: *pori*. Adj: *porate*.
- position** (a) Placement relative to other, unlike parts. (disposition) (b) The placement of a taxon as a member of another taxon of the next higher rank.
- posterior** Referring to the upper lobe or part, especially in a horizontally oriented structure. (position)
- posture** Placement relative to a flat plane. (disposition)
- praemorse** Having a jagged, chewed appearance, e.g. some palms. (margin)
- prickle** A nonspine, nonhorn, sharp-pointed outgrowth from the surface of any organ. Adj: *prickly*. (plant, twig part)
- primary cell wall** The first, mostly cellulosic cell wall layer that is secreted external to the plasma membrane during cell growth.
- primary endosymbiosis** Endosymbiosis involving the engulfment of an ancestral bacterium by a eukaryotic cell.
- primary growth** Growth in height or length of a stem or root, brought about by the elongation and differentiation of cells derived from the apical meristem.
- primary inflorescence** *Unit inflorescence*. (inflorescence part)
- primary parietal cells** The inner layer of cells in an early stage of anther wall development.
- primary pit field** A group of numerous plasmodesmata in the primary cell wall, spatially associated with secondary cell wall pit pairs.
- primary root** The root of the sporophyte that develops from the radical of the embryo. (root type)
- primary tissue** A tissue formed by primary growth, e.g. as in primary xylem or primary phloem.
- primary vein** The major vein or veins of a leaf with respect to size. (leaf venation)
- primer** A complementary copy of a short, conserved, flanking region of a region of DNA of interest, used to amplify and sequence the DNA.
- primitive** *Ancestral*.
- principle of parsimony** The principle stating that the cladogram exhibiting the fewest number of evolutionary steps is accepted as the best estimate of phylogeny; a corollary of the general principle of Ockham's Razor. Syn: *parsimony analysis*.
- priority of publication** A principle of the International Code of Botanical Nomenclature stating that of two or more competing

- possibilities for a name, the one published first is the correct one, with some exceptions.
- prismatic** A short, prism-shaped crystal; a type of ergastic substance.
- procumbent** *Prostrate*. (stem habit)
- proembryo** A very young embryo.
- prolate** A pollen grain in which the P/E ratio is greater than about 1.2.
- prop root** Above-ground, adventitious roots that function in supporting the stem (root type).
- prophyll/prophyllum** *Bracteole/bractlet*. (leaf structural type, flower part)
- prostrate** Trailing or lying flat, not rooting at the nodes. Syn: *procumbent*; *reclining*. (stem habit)
- protandry** With stamens or anthers developing before carpels or stigma. Adj: *protandrous*. (maturation, flower maturation)
- proteinoplast** *Aleurone grain*.
- protogyny** With carpels or stigma maturing before stamens or anthers. Adj: *protogynous*. (maturation, flower maturation)
- protonema** An initial, filamentous form of a gametophyte (e.g. in mosses), prior to its differentiation into parenchymatous tissue.
- protoplasm** Everything inside the plasma membrane of a cell.
- protostele** A stele with a central, solid cylinder of vascular tissue.
- protoxylem** Referring to the first tracheary elements that develop within a patch of xylem, being typically smaller and with thinner cell walls than the later formed metaxylem.
- proximal** Near the point of origin or attachment. (position)
- proximal pole** The intersection of the pollen grain polar axis with the grain surface that is near the center of the microspore tetrad.
- pseudanthium** A unit that appears as and may function like a single flower, but that typically consists of two or more flowers fused or grouped together. (inflorescence type)
- pseudobulb** A short, erect, aerial storage stem of certain epiphytic orchids. (stem/shoot type)
- pseudocrassinucellate** An ovule type in which an second inner layer of nucellar cells by periclinal divisions in the single outer layer, a parietal cell not forming.
- pseudodrupe** A nut surrounded by a fleshy, indehiscent involucre, resembling a true drupe, e.g. *Juglans*, walnut. (fruit type)
- pseudolaters** Groups of elongate, cohering, nonsporogenous, generally hygroscopic cells that develop within the sporangia of hornworts and function in spore dispersal.
- pseudopetiole** Term sometimes used for the petiole-like structure arising between a leaf sheath and blade, found in several monocots, such as bananas, bamboos. (leaf part).
- pseudoterminal bud** A bud appearing to be apical but that is actually lateral near the apex, assuming a terminal position with the death or nondevelopment of the true terminal bud. (bud type)
- pseudoumbel** An inflorescence appearing like a simple umbel, but actually composed of condensed, monochasial cymes, as in the Alliaceae and Amaryllidaceae. (inflorescence)
- psilate** A pollen grain having a smooth sculpturing.
- psychophily** Pollination by butterflies. Adj: *psychophilous*.
- puberulent** Minutely pubescent. (vestiture)
- pubescent** (a) With straight, short, soft, somewhat scattered, slender trichomes. (b) A general term, meaning having trichomes. (vestiture)
- pulvinus** The swollen base of a petiole or petiolule, e.g. in some Fabaceae. (leaf part)
- punctate** Covered with minute, pitlike depressions. (configuration)
- pungent** *Spinose*. (apical process)
- pyrene** A fleshy fruit in which each of two or more seeds is enclosed by a usually bony-textured endocarp, or the seed covered by a hard endocarp unit itself, regardless of the number. (fruit type)
- pyxide/pyxis** *Circumscissile capsule*. (fruit type)
- quincuncial** Perianth parts of a single pentamerous whorl having two members overlapping, two being overlapped, and one overlapping only at one margin. (perianth aestivation)
- raceme** An indeterminate inflorescence consisting of a single axis bearing pedicellate flowers. (inflorescence type)
- rachilla** A lateral or secondary axis of a bipinnate leaf. (leaf part) (b) The central axis of a grass or sedge spikelet. (inflorescence part) Pl: *rachillae*.
- rachis** (a) The main axis of a pinnately compound leaf. (leaf part) (b) A major, central axis within an inflorescence (inflorescence part)
- radial** (a) *Actinomorphic*. (b) Referring to a longitudinal section of wood, parallel to a stem radius.
- radical** *Basal*. (position)
- radicle** The first root of a seed embryo. (root type, seed part)
- ramet** A clonal unit of a genet that is, at least potentially, independent from other ramets. Cf. *genet*.
- random amplified polymorphic DNA** The use of using randomly generated primers for the amplification of DNA to identify polymorphic DNA regions of different individuals or taxa. Abbr: *RAPD*.
- rank** One of the hierarchical taxonomic categories, in which a higher rank is inclusive of all lower ranks.
- raphe** Ridge on seed coat formed from adnate funiculus. (seed part)
- raphide** A needlelike crystal of calcium oxalate, typically occurring in bundles; a type of ergastic substance.
- ray** (a) A secondary axis in a compound umbel. (inflorescence part) (b) A corolla with a short tube and a single, elongate, straplike apical extension, e.g. some Asteraceae. Syn: *ligulate*. (perianth type) (c) Radially oriented cells that occur in bandlike strands in the secondary xylem (wood), functioning in lateral translocation of water, minerals, and sugars.
- recency of common ancestry** A measure of phylogenetic relationship, stating that two taxa are more closely related to one another if they share a common ancestor that is more recent in time than the common ancestor they share with any other taxon.
- receptacle** The tissue or region of a flower to which the other floral parts are attached. (flower part)
- reclinate** *Reclined*. (orientation)
- reclined** Directed downward, with divergence angle of 15°–45° from horizontal axis. Syn: *reclinate*. (orientation)
- reclining** *Prostrate*. (stem habit)
- recurved** With tip gradually curved outward or downward, abaxially. (transverse posture)
- reduced** A type of anther wall development in which the secondary parietal cells do not divide further and develop directly into the endothecium and tapetum, respectively.
- reduplicate** Plicate, with adjacent abaxial sides facing one another, Λ -shaped in cross-section. (longitudinal posture)
- reflexed** Bent or turned downward. (orientation)
- regular** *Actinomorphic*.
- relative cover** A measure of the degree to which each species of a community layer contributes to the total cover of that layer alone.

- remodel/remodeling** To change the diagnostic characteristics of a taxon.
- reniform** (a) Kidney-shaped; wider than long with a rounded apex and reniform base. (shape) (b) With two rounded, basal lobes, smoothly concave at intersection of lobes. (base)
- repand** Margins wavy in a vertical plane. (longitudinal posture)
- repent** Creeping or lying flat and rooting at the nodes. (stem habit)
- replum** A septum, generally with attached seeds, that persists after fruit dehiscence, e.g. silicles and siliques of Brassicaceae. (fruit part)
- rescaled consistency index (RC)** A measure of the relative amount of homoplasy in a cladogram, equal to the product of the consistency index and retention index.
- resin ducts/canals** Specialized cells that secrete resin.
- restriction fragment length polymorphism** Differences between taxa in restriction sites, and therefore the lengths of fragments of DNA after cleavage with restriction enzymes. Abbr: *RFLP*.
- restriction site** A sequence of approximately six to eight base pairs of DNA that binds to a given restriction enzyme.
- resupinate** Inverted or twisted 180°, as leaves of Alstroemeriaceae or ovaries of most Orchidaceae flowers. (twisting/bending posture)
- retention index (RI)** A measure of the relative amount of homoplasy in a cladogram, equal to the ratio $(g - s)/(g - m)$, where g is the maximum possible number of state changes that could occur on any conceivable tree, m is the minimum possible number of character state changes, and s is the actual number of state changes that occur.
- reticulate** (a) *Netted*. (leaf venation) (b) A pollen sculpturing with a netlike sculpturing., each element termed a murus and the space between termed a lumen.
- reticulation** Hybridization of two previously divergent lineages, forming a new lineage.
- reticulodromous** (a) Pinnate venation in which secondary veins do not terminate at the margin and branch repeatedly, forming a very dense, netlike structure. (leaf venation) (b) With three or more primary veins diverging from one point but not reaching the blade margin, a subcategory of actinodromous. (leaf venation)
- retinaculum** *Jaculator*. Pl: *retinacula*.
- retorse** Bent or directed downward, usually referring to small appendages. (orientation)
- retuse** Having an apical incision, cut up to $\frac{1}{6}$ the distance to the midrib, midvein, or junction of primary veins. (apex)
- reversal** Homoplasy by the loss of a derived feature and the reestablishment of an ancestral feature.
- revolute** Margins or outer portion of sides rolled outward or downward over the abaxial surface. (longitudinal posture, margin)
- reward** A floral structure, such as pollen, or exudate, such as nectar, that functions to ensure that an animal pollinator will consistently return to transport pollen.
- rhizidium** A monochasium in which the branches develop on alternating sides of each sequential axis, typically having a geniculate (zig-zag) appearance; sometimes equated with scorpioid cymes. (inflorescence type)
- rhizoid** One of several uniseriate (one-cell-thick), filamentous processes that function in anchorage and water/mineral absorption, arising from the gametophytes of free-living gametophytes and from the underground, sporophytic stems of some ferns (e.g. the psilophytes).
- rhizome** A horizontal, underground stem, generally with short internodes and scalelike leaves, e.g. *Zingiber officinale*. (b) A horizontal stem of a fern that grows at ground level. (stem/shoot type)
- rhombic** Parallelogram-shaped, widest near the middle, the length : width ratio 2:1 to 3:2. (shape)
- rhomboidal tetrad** A tetrad in which the four grains are in one plane, with two of the grains separated from one another by the close contact of the other two.
- ribosome** A cellular organelle that functions in protein synthesis.
- ribulose-bisphosphate carboxylase** The enzyme that catalyzes the initial binding (fixation) of carbon dioxide in the dark reactions of photosynthesis. Abbr: *RuBP-carboxylase*.
- rind** *Pericarp*.
- ring-porous** Wood in which the vessels form only in the spring wood, with summer wood either lacking or having relatively small vessels and usually containing mostly fibers.
- root** A cylindrical organ of virtually all vascular plants, consisting of an apical meristem that gives rise to a protective root cap, a central endodermis-bounded vascular system, absorptive epidermal root hairs, and endogenously developed lateral roots; usually functioning in anchorage and absorption of water and minerals; initially derived from the radicle of the embryo and typically growing downward. (plant part)
- root apical meristem** Region of continuously dividing cells from which all cells of the root are derived. (root part)
- root cap** The outer cell layer at the root tip, functioning in protection and lubrication. (root part)
- root hair** One of several hairlike extensions from an epidermal cell of a root, functioning to greatly increase the surface area available for water and mineral absorption. (root part)
- root tuber** A swollen taproot containing concentrations of high-energy compounds such as starch. (root type)
- rootcap** An outer layer of cells at the tip of a root, functioning to protect the root apical meristem and to provide lubrication for the growing root.
- rootstock** A general term for an underground stem or shoot, these generally giving rise to aerial shoots either by direct conversion of the terminal apical meristem or via lateral buds. (stem/shoot type)
- rosette** An arrangement in which parts, usually leaves, radiate from a central point at ground level, e.g. the leaves of *Taraxacum officinale*, dandelion. Adj: *rosulate*. (arrangement)
- rotate** With a short tube and wide limbs oriented at right angles to the tube. (perianth type)
- rounded** With apical or basal margins convex, forming a single, smooth arc. (apex, base)
- rugose** Covered with coarse reticulate lines, usually with raised, blisterlike areas between. Syn: *bullate*. (configuration)
- rugulate** A pollen sculpturing having irregular to sinuous, tangentially oriented elements, often appearing brainlike.
- ruminant** Unevenly textured, coarsely wrinkled, looking as if chewed, e.g. the endosperm of the Annonaceae. (surface, texture)
- runner** *Stolon*. (stem/shoot type)
- saccate** Having a pouchlike evagination. (perianth type)
- sagittate** With two basal lobes, more or less pointed and oriented downward, away from apex. (base)
- salverform** Trumpet-shaped; with a long, slender tube and flaring limbs at right angles to tube. (perianth type)
- samara** A winged, dry fruit, e.g. *Acer*, maple; *Ulmus*, elm. (fruit type)

- sapromyophily** Pollination by flies. Adj: *sapromyophilous*.
- sarcotesta** A seed coat that is fleshy at maturity. (seed part)
- sarmentose** *Stoloniferous*. (stem habit)
- saxicolous** Occurring on rock. (plant habitat)
- scaberulous** Minutely scabrous. (epidermal excrescence, vestiture)
- scabrate** *Spinulose*.
- scabrous** Having a rough surface or trichomes, resembling sandpaper. (epidermal excrescence, vestiture)
- scale** (a) A small, nongreen leaf of a bud or underground rootstock. (leaf structural type) (b) A bract of a sedge spikelet. (inflorescence part)
- scaling** Assigning a weight in a phylogenetic analysis to correlated characters that is the inverse of the total number of the correlated characters. Adj: *scaled*.
- scandent** *Clambering*.
- scape** A naked (leafless) peduncle, generally arising from a basal rosette of vegetative leaves. Adj: *scapose*. (stem/shoot type, inflorescence part)
- scarious** Thin and appearing dry, usually whitish or brownish. (texture)
- schizocarp** Derived from a two or more loculed compound ovary in which the locules separate at maturity. (fruit type)
- schizocarp of follicles** A schizocarp in which the carpels of a pistil split at fruit maturity, each carpel developing into a unit follicle, e.g. *Asclepiadaceae*. (fruit type)
- schizocarp of mericarps** A schizocarp in which (generally two) carpels of a single, unlobed ovary split during fruit maturation, the carpels developing into unit mericarps and attached to one another via a stalklike carpophore, e.g. *Apiaceae*. (fruit type)
- schizocarp of nutlets** A schizocarp in which a single ovary becomes lobed during development, each lobe developing at maturity into a nutlet, e.g. *Boraginaceae*, *Lamiaceae*. (fruit type)
- scientific name** A formal, universally accepted name, the rules and regulations of which (for plants, algae, fungi, and organisms traditionally treated as such) are provided by the International Code of Botanical Nomenclature.
- sclereid** A sclerenchyma cell that is isodiametric to irregular in shape and often branched, functioning in structural support or possibly aiding in providing protection from herbivory.
- sclerenchyma** A tissue composed of nonconductive cells that are dead at maturity, that have a thick, lignified, generally pitted secondary cell wall, and that function in structural support and/or to deter herbivory; composed of fibers and sclereids.
- scorpioid cyme** A monochasium in which the branches develop on alternating sides of each sequential axis, typically appearing geniculate (zig-zag); may intergrade with helicoid cyme. Syn: *cincinnus*. (inflorescence type)
- scurfy** *Farinaceous*. (epidermal excrescence)
- secondary cell wall** An additional wall layer, composed of cellulose and lignin, secreted between the plasma membrane and the primary cell wall following cell elongation, found in some cell types (including tracheary elements and fibers).
- secondary endosymbiosis** Endosymbiosis involving the engulfment of an ancestral eukaryotic cell by another eukaryotic cell, a possible mechanism of chloroplast exchange between eukaryotes.
- secondary growth** Growth in girth or width by means of cells produced from lateral meristems.
- secondary parietal cells** Two layers of cells arising by tangential divisions of the primary parietal cells of the anther wall.
- secondary phloem** Sugar-conducting tissue produced by the vascular cambium to the outside of a woody stem or root. Syn: *inner bark*.
- secondary root** A root derived endogenously from the pericycle of an existing root. (root type)
- secondary tissue** A tissue formed by secondary growth, via lateral meristems.
- secondary vein** A lateral vein that branches from and is smaller than a primary vein. (leaf venation)
- secondary xylem** Water- and mineral-conducting tissue produced by the vascular cambium to the inside of a woody stem or root. Syn: *wood*.
- secretory** An anther tapetum type in which the tapetum remains intact with no breakdown of cell walls. Syn: *glandular*.
- secretory structure** A collection of cells that secrete compounds, either internally or externally.
- secund** Flowers, inflorescences, or other structures on one side of axis, often due to twisting of stalks. Syn: *unilateral*. (arrangement)
- sedge spikelet** The inflorescence unit of the *Cyperaceae*, sedge family, consisting of a central axis (the rachilla), bearing spiral or distichous bracts (also called scales or glumes), each subtending a single flower.
- seed** An embryo surrounded by nutritive tissue and enveloped by a seed coat; the propagule of the seed plants.
- seed coat** The outer protective covering of seed, developing from one or two integuments. Syn: *testa*. (seed part)
- self-dispersal** Process by which a flowering plant buries its own fruits, e.g. *Arachis hypogaea*, peanut. Syn: *autochory*.
- self-incompatibility** Outcrossing occurring by the genetic inability for fertilization to occur between gametes derived from a single genotype.
- selfing** *Inbreeding*.
- semelpary** Referring to plants that have one episode of reproduction, followed by degeneration and death of the entire plant. Adj: *semelparous*.
- semicraspedodromous** Pinnate venation in which the secondary veins branch near the margin, one terminating at the margin, the other looping upward to join the next secondary vein. (leaf venation)
- sensu lato** Meaning in the broad sense; designated for a taxon name that is used inclusively, for a broad, inclusive taxon circumscription to include other, previously recognized taxa. Abbr: *s.l.*
- sensu stricto** Meaning in the strict sense; designated for a taxon name used exclusively, to exclude other, previously recognized taxa. Abbr: *s.s.*; *s.st.*
- sepal** An individual member or segment of the calyx, typically green, leaflike, and functioning to protect the young flower. (flower part, perianth part)
- septal nectary** A nectary embedded within the ovary septae, secreting nectar via a pore at the ovary base or top. (nectary type)
- septicidal capsule** A capsule in which longitudinal lines of dehiscence are radially aligned with the septa. (fruit type)
- septifragal capsule** A capsule in which the valves break off from the septa, as in *Ipomoea*, morning glory. Syn: *valvular capsule*. (fruit type)
- septum** A partition or cross-wall of the ovary. (gynoecium part, fruit part)

- sericeous** With long, appressed, silky trichomes. (vestiture)
- series** *Whorl*. (flower part)
- serrate** Saw-toothed; teeth sharp and ascending, the lower side longer, cut $\frac{1}{6}$ to $\frac{1}{8}$ the distance to midrib, midvein, or junction of primary veins. (margin)
- serrulate** Diminutive of serrate, teeth cut to $\frac{1}{6}$ the distance to midrib, midvein, or junction of primary veins. (margin)
- sessile** (a) Without a petiole or, for leaflets, without a petiolule. (leaf attachment) (b) Lacking a pedicel. (flower attachment) (c) With filament absent, the anther attached directly. (stamen attachment) (d) Ovary lacking a basal stalk. (ovary attachment)
- sessile-glandular** Glandular cell with a very short or no basal stalk. (trichome type)
- sexine** The outer, protruding layers of the exine, including columellae, tectum, and suprategular sculpturing elements.
- sheath** A flattened leaf base or petiole that partially or fully clasps the stem, e.g. Poaceae and many Apiaceae. (leaf part)
- sheathing** Attached by a curved or tubular structure that partially or totally encloses the stem. (leaf attachment, base)
- shining** *Nitid*. (epidermal excrescence)
- shoot** A stem plus associated, derivative leaves, initially formed by an apical meristem that gives rise to the stem and external (exogenous) leaf primordia; may be gametophytic or sporophytic. (plant part)
- short shoot** *Fascicle; spur; spur shoot*. (stem/shoot type)
- shreddy** Bark coarsely fibrous, often fissured. (bark type)
- shrub** A perennial, woody plant having several main stems arising at ground level. (plant habit)
- sieve area** An aggregation of callose-lined pores in *sieve elements*.
- sieve cell** A type of sieve element that has only sieve areas on both end and side walls, found in nonangiospermous vascular plants.
- sieve element** A generally elongate cell that is semialive at maturity, has a nonlignified, primary cell wall with specialized, callose-lined pores aggregated into sieve areas and/or sieve plates, and that functions in conduction of sugars.
- sieve plate** One or more sieve areas at the end wall junction of two sieve tube members, the pores of which are significantly larger than are those of sieve areas located on the side wall; characteristic of angiosperms.
- sieve tube member** A type of sieve element that has both sieve areas and sieve plates, the latter at the end wall junction of two sieve tube members and having larger pores; an apomorphy of angiosperms.
- sieve tube plastid** A membrane-bound plastid found in sieve tube members, the contents of which (whether starch or protein) can vary between taxa and be systematically informative.
- silicle** A dry, dehiscent, 2-carpeled fruit that dehisces along two sutures, has a persistent partition (replum), and is as broad or broader than it is long, e.g. Brassicaceae. Cf: *siliqua*. (fruit type)
- siliqua** A dry, dehiscent, 2-carpeled fruit that dehisces along two sutures, has a persistent partition (replum), and is longer than it is broad, e.g. Brassicaceae. Cf: *silicle*. (fruit type)
- simple leaf** A leaf not divided into leaflets, bearing a single blade (leaf type)
- simple cone** A cone consisting of an axis bearing sporophylls.
- simple corymb** An unbranched corymb, consisting of a central axis bearing pedicellate flowers, the collection of flowers being flat-topped or convex. (inflorescence type)
- simple craspedodromous** Pinnate venation in which all secondary veins terminate at the margin. (leaf venation)
- simple dichasium** A three-flowered determinate inflorescence with a single terminal flower and two, opposite lateral flowers, the pedicels of all of equal length. (inflorescence type)
- simple fruit** A fruit derived from one pistil (ovary) of one flower. (fruit type)
- simple ovary/pistil** An ovary/pistil composed of a single carpel, the gynoeceum apocarpous. (ovary/pistil type)
- simple perforation plate** A perforation plate composed of a single opening.
- simple sieve plate** A sieve plate that is composed of one pore region.
- simple tissue** A tissue that consists of only one type of cell.
- simple umbel** An umbel with the peduncle bearing pedicellate flowers attached at one point, e.g. Alliaceae. (inflorescence type)
- simultaneous** A type of microsporogenesis in which cytokinesis does not occur until after meiosis II.
- sinistrorse** Twining helically like a left-handed screw, e.g. some Caprifoliaceae. (twisting/bending posture)
- sinuate** Sinuses shallow and smooth, wavy in a horizontal plane. (division)
- sinus** The indentation of a divided planar structure. (division)
- siphonogamy** The formation or evolution of pollen tubes.
- siphonostele** A type of stem vasculature in which a ring of xylem is surrounded by an outer layer of phloem (ectophloic siphonostele) or by an outer and inner layer of phloem (amphiphloic siphonostele; if dissected, called a dictyostele).
- sister group/sister taxa** Two taxa that are each other's closest relative, representing two descendant lineages from a common ancestor.
- smooth** (a) *Plane*. (configuration) (b) A nonfibrous bark without fissures, plates, or exfoliating sheets. (bark type)
- softwood** Wood derived from a conifer, generally (but not always) softer in texture than a hardwood.
- solanad type** A type of embryo development in which the terminal cell divides transversely, the basal-cell derivatives forming a suspensor but otherwise not contributing to mature embryo development.
- solitary** A one-flowered inflorescence. (inflorescence type)
- somatic** Vegetative; referring to a cell not directly involved in sexual reproduction.
- soros** A multiple fruit in which the unit fruits are fleshy berries and are laterally fused along a central axis, e.g. *Ananas*, pineapple. (fruit type)
- sorus** A discrete cluster or aggregation of leptosporangia. Pl: *sori*.
- spadix** A spike with a thickened or fleshy central axis, typically with congested flowers, e.g. Araceae. (inflorescence type)
- spathaceous** Having a spathe. (leaf structural type, inflorescence part)
- spathe** An enlarged, sometimes colored bract subtending and usually enclosing an inflorescence, e.g. in the Araceae. (leaf structural type, inflorescence part)
- spatulate** A shape that is oblong, obovate, or oblanceolate with a long attenuate base. (shape)
- speciation** The formation of new species from preexisting species.
- species** By one definition, a distinct lineage that, in sexually reproducing organisms, consists of a group of generally intergrading, interbreeding populations that are essentially reproductively isolated from other such groups.

- species nova** Meaning that a taxon name, at the rank of species, is new to science. Abbr: *sp. nov.*
- specific epithet** The second name of a binomial; may be capitalized or not.
- sperm** A haploid, motile or nonmotile, male gamete that functions to fuse with an egg cell, producing a diploid zygote.
- spheroidal** Approximately spherical in shape, e.g. referring to pollen grains.
- spike** An indeterminate inflorescence consisting of a single axis bearing sessile flowers. (inflorescence type)
- spikelet** A small spike; the basic inflorescence unit in the Cyperaceae (sedges) and Poaceae (grasses). (inflorescence type)
- spine** A sharp-pointed leaf (e.g. cactus spines or glochidia) or leaf part, including petiole (e.g. *Fouquieria* spp.), midrib, secondary vein, leaflet (e.g. *Phoenix* spp.) or stipule (e.g. *Euphorbia* spp.) Cf: *thorn*; *prickle*. (leaf structural type)
- spinose** (a) Margins with teeth bearing sharp, stiff, spinelike processes. (margin) (b) With a sharp, stiff, spinelike apical process. Syn: *pungent*. (apical process)
- spinulose** A pollen sculpturing having spinelike elements <1 m long, each element termed a spinulum. Syn: *scabrate*.
- spinulum** One of the spinelike elements of a spinulose pollen grain. Pl: *spinuli*.
- spiral** (a) Alternate, with points of attachment in more than three rows/ranks. Syn: *polystichous*. (arrangement) (b) Perianth parts arranged in a spiral, one per node, not in distinct whorls. (perianth arrangement) (c) Stamens arranged in a spiral, one per node; not whorled. (stamen arrangement) (c) A type of endothecium in which the secondary wall thickenings are spiral or helical in shape.
- spiraperturate** A pollen grain having one or more apertures that are spirally shaped.
- spongy mesophyll** The usually lower (abaxial), irregular cells of the mesophyll that generally have large intercellular spaces, found in some leaves.
- sporangial wall** One or more outer layers of sterile, non-spore-producing cells of the sporangium.
- sporangiophore** A unit of the strobilus of the Equisetales, consisting of a peltate axis bearing pendant (ancestrally recurved), longitudinally dehiscent sporangia.
- sporangium** The spore-producing organ of the sporophyte. Pl: *sporangia*. (plant part)
- spore** A haploid cell that, in the land plants, originates from meiotic divisions of sporocytes within a sporangium, ultimately growing into a gametophyte.
- sporocarp** The generally spherical reproductive structure of aquatic ferns, functioning in allowing the sporangia inside to remain dormant and resist desiccation for a long time.
- sporocyte** A cell of the sporangium that undergoes meiosis, producing (generally four) haploid spores.
- sporogenous tissue** Internal cells of the sporangium that mature into sporocytes, the latter forming spores by meiotic divisions.
- sporophyll** A specialized leaf that bears one or more sporangia.
- sporophyte** The diploid (2n) phase in the haplodiplontic life cycle of all land plants.
- sporophytic leaf** The leaf of a vascular plant sporophyte.
- sporopollenin** A polymer of oxidative carotenoids or carotenoid esters that impregnates the exine of pollen grain walls.
- spring wood** The secondary xylem from the first part of the growing season, the cells of which tend to be larger in diameter with thinner walls.
- spur** A tubular, rounded or pointed projection from the calyx or corolla, functioning to contain nectar. (perianth part)
- spur/spur shoot** *Fascicle*. (stem/shoot type)
- squarrose** Sharply curved downward or outward (abaxially) near the apex, as the phyllaries of some Asteraceae. (transverse posture)
- stamen** The unit of the androecium; a microsporophyll (typically modified as a filament) bearing generally two pollen-bearing thecae, fused into an anther. (flower part)
- stamen insertion** (a) Referring to whether or not stamens protrude beyond the perianth. (b) Denoting the point of insertion, the point of adnation of an epipetalous stamen to the corolla. (androecium)
- staminal disk** A discoid or doughnut-shaped structure arising from the receptacle at the base of the stamens; may be nectar-bearing. Syn: *nectariferous disk*. (flower part)
- staminate** Flowers unisexual, with stamen(s) only, lacking fertile carpel(s). Syn: *male*. (flower sex)
- staminode/staminodium** A sterile stamen, producing no functional pollen; may be modified as a nectary or petaloid structure; may or may not contain an antherode, a sterile anther. (stamen type)
- starch** A polysaccharide of glucose units (alpha-1,4-glucopyranoside) in green plants, functioning as the high-energy storage compound.
- starch grain** Lamellate deposits of starch in green plant cells, functioning as the high-energy storage compound; a type of ergastic substance. Syn: *amyloplast*.
- status novus** Indication of a change in rank of a scientific name. Abbr: *stat. nov.*
- stele** The spatial distribution of the primary vasculature of a stem, organized into arrangements of xylem and phloem.
- stellate** Star-shaped trichomes having several arms arising from a common base (stalked or sessile), e.g. many Malvaceae. (trichome type)
- stem** A generally cylindrical organ that bears leaves, typically functioning in support and elevation of leaves and reproductive structures and in conduction of water, minerals, and sugars; in vascular plants initially derived from the epicotyl of the embryo and generally growing upward. (plant part)
- stem (cladogram)** *Internode*.
- stem habit** Stem feature based on structure, position, growth, and orientation of above-ground stems. (stems and shoots)
- stem-based** A type of phylogenetic classification in which the stem (internode) region just above a common ancestor plus all descendants of that stem are the basis for grouping.
- sterile jacket layer** *Antheridial wal*; *archegonial wall*.
- sterile segment** The photosynthetic blade or lamina component of the leaf of an ophioglossoid fern.
- stigma** The pollen-receptive portion of a pistil; may be a discrete structure or a region of a style or style branch. (gynoecium part)
- stinging trichome** A sharp-pointed trichome that secretes an irritating fluid upon penetrating the tissue of an animal.
- stipe** (a) A leaf stalk, often used for ferns. Syn: *petiole*. (leaf part) (b) *Gynophore*. Adj: *stipitate*. (gynoecium part) (c) The stalk of the sporophyte of mosses.
- stipel** One of a pair of leaflike structures, which may be modified as spines or glands, at either side of the base of a petiolule, e.g. some Fabaceae. Adj: *stipellate*. (leaf part)

- stipular spine** A sharp-pointed stipule, e.g. *Euphorbia* spp. Cf. *thorn*; *prickle*. (leaf structural type)
- stipule** One of a pair of leaflike structures, which may be modified as spines or glands, at either side of the base of a petiole. Adj: *stipulate*. (leaf part)
- stipule scar** A mark indicating the former place of attachment of a stipule. (twig part)
- stolon** An indeterminate, elongate, slightly underground or above-ground propagative stem, with long internodes, rooting at the tip forming new plants. Syn: *runner*. (stem/shoot type)
- stoloniferous** With elongate propagative shoots (stolons) rooting at the tip producing new plants. Syn: *sarmentose*. (stem habit)
- stoma** The opening between two guard cells of a stomate.
- stomate** Specialized epidermal cells, consisting of two guard cells, which, by changes in turgor pressure, can increase or decrease the size of the opening (stoma) between them. Pl: *stomata*; *stomates*.
- storage root** A swollen taproot containing concentrations of high-energy compounds such as starch. (root type)
- straight** Flat, without vertical curves or bends. (transverse posture)
- strap-shaped** Flat, not needlelike but with length : width ratio greater than 12:1. (shape)
- striate** (a) With fine longitudinal lines. (configuration) (b) A pollen sculpturing having thin, cylindrical, tangentially oriented elements.
- strict consensus tree** A consensus tree that collapses any differences in branching pattern between two or more cladograms to a polytomy.
- strigose** Covered with dense, coarse, bent, and mostly flat trichomes, often with a bulbous base. (vestiture)
- strigulose** Minutely strigose. (vestiture)
- strobilus** *Cone*. (plant part)
- style** A stalklike portion of the pistil between the stigma and ovary; may be absent. (gynoecium part)
- styloid** A single, elongate, angular crystal of calcium oxalate; a type of ergastic substance.
- stylopodium** A swollen region at apex of the ovary, below the styles/stigmas, e.g. *Apiaceae*.
- subapical** With style arising at one side near the apex of the ovary. (style position)
- subbasifixed** Anther attached just above its base to the filament. (anther attachment)
- suberin** A waxy, water-resistant substance, found in stomata, Casparian strips of the endodermis, and the cell walls of cork cells.
- subglabrous** Nearly glabrous, with just a few, scattered trichomes. (vestiture)
- subopposite** With two leaves or other structures on opposite sides of stem or central axis but at different nodes, slightly displaced relative to one another. (arrangement)
- subsessile** Having a very short, rudimentary petiole (leaf attachment), petiolule (leaflet attachment), pedicel (flower attachment), or filament (stamen attachment).
- subshrub** A short shrub that is woody only at the base and that seasonally bears new, nonwoody, annual shoots above. (plant habit)
- subsidiary cell** An epidermal cell adjacent to a stomate and somewhat different from other epidermal cells in shape and size, functioning in regulating stomatal opening.
- subspecific epithet** The third component of a trinomial at the rank of subspecies.
- subulate** Awl-shaped; approximately narrowly oblong to narrowly triangular. (shape)
- successive** A type of microsporogenesis in which cytokinesis occurs after meiosis I.
- succulent** Fleshy or juicy. Syn: *baccate*, *carlose*. (texture)
- suffrutescent** Woody basally, herbaceous apically; the habit of a subshrub. (stem habit)
- sulcus** An elongate aperture similar in shape to a colpus (length : width ratio >2:1) occurring at the (usually distal) pole. Adj: *sulcate*.
- summer wood** The secondary xylem from the latter part of the growing season, the cells of which tend to be smaller in diameter with thicker walls.
- superior** With sepals, petals, stamens, and/or hypanthium attached at the base of the ovary. (ovary position)
- superposed bud** Bud(s) occurring above the axillary bud. (bud type)
- supervolute** With one half of a simple leaf coiled tightly around the midrib, the other half coiled (in the opposite direction) around the first half, as in members of the *Zingiberales*. (posture: *ptyxis*/aestivation)
- suprabasal** With three or more primary veins diverging from one point above the base of the blade, a subcategory of actinodromous and palinactinodromous. (leaf venation)
- suprafoliar** Descriptive of a palm inflorescence that is positioned above the leaves of the crownshaft.
- suspensor** A nonpersistent column of cells of the embryo, functioning in transport of nutrients to the mature, persistent embryo during its development.
- syconium** A multiple fruit in which the unit fruits are small achenes covering the surface of a fleshy, inverted compound receptacle, e.g. *Ficus*, fig. (fruit type)
- symmetry** Referring to the presence and number of mirror-image planes of symmetry.
- sympetalous** With connate petals. Syn: *gamopetalous*. (perianth fusion)
- symplesiomorphy** A shared, ancestral features among taxa, which may be used as a basis for grouping in a phenetic classification.
- sympodial** A branching pattern in which a given axis is made up of several units, each of which is derived from a separate apical meristem, the units themselves determinate or indeterminate. (stem branching pattern)
- syngangium** A fusion product of two or more sporangia, e.g. in the psilophytes and some marattioid ferns.
- synanthy** Timing in which leaves and flowers develop at the same time. Adj: *synanthous*.
- synapomorphy** An apomorphy that unites two or more taxa or lineages.
- syncarp** An aggregate fruit, typically of berries, in which the unit fruits fuse together, e.g. *Annona*. (fruit type)
- syncarpous** With carpels (at least at the base) connate, the pistil or ovary being compound. (gynoecial fusion)
- syncolpate** A pollen grain in which the colpi are joined, e.g. at the poles.
- synergid cells** In a typical angiosperm female gametophyte, the two haploid cells that flank the egg and that may function in pollen tube entry.
- syngenesious** With anthers connate and filaments distinct, e.g. *Asteraceae*. (stamen fusion)
- synonym** A rejected name, by a particular author or authors, either because the name is illegitimate or because of taxonomic judgment.

- synoptic collection** A collection of plant specimens that contain one specimen (of all available specimens) of each taxon for a given region.
- synsepalous** With connate sepals. Syn: *gamosepalous*. (perianth fusion)
- syntepalous** With connate tepals. (perianth fusion)
- syntype** A specimen that was cited in the original work when a holotype was not designated, or one of two or more specimens that were all designated as types.
- systematics** A science that includes and encompasses traditional taxonomy and that has as its primary goal the reconstruction of phylogeny.
- tandem repeat** A sequence of DNA that repeats multiple times.
- tangential** Referring to a longitudinal section of wood, perpendicular to a stem radius.
- tannin** Phenol derivatives in plant cells that may function to deter herbivory and parasite growth; a type of ergastic substance.
- tapering** Trichomes ending in a sharp apex. (trichome type)
- tapetum** The innermost cell layer of the anther wall, consisting of of metabolically active cells that function in the development of pollen grains.
- taproot** A persistent, well-developed primary root. (root type)
- tautonym** A binomial in which the genus name and specific epithet are identical in spelling, illegitimate in the International Code of Botanical Nomenclature.
- taxon** (*plural, taxa*) A group of organisms, ideally monophyletic and traditionally treated at a particular rank.
- taxonomic key** *Key*.
- taxonomic revision** A change of the definition and delimitation of a taxon.
- taxonomy** A field of science (and major component of systematics) that encompasses description, identification, nomenclature, and classification.
- tectate-columellate** A common exine structure that consists of an inner foot-layer, a middle layer of radially elongate columellae, and an outer, rooflike layer called the tectum.
- tectum** The outer, rooflike layer of a tectate-columellate pollen exine wall.
- tendrill** (a) A coiled and twining leaf part, usually a modified rachis or leaflet. (leaf structural type) (b) A long, slender, coiling branch, adapted for climbing. (stem/shoot type)
- tentacular leaf** A leaf bearing numerous, sticky, glandular hairs or bristles that function in capturing and digesting small animals, e.g. *Drosera*, sundew. (leaf structural type)
- tenuinucellate** An ovule in which the nucellus is composed of a single layer of cells, with no formation of a parietal cell.
- tepal** A component of the perianth in which the parts intergrade or in which the perianth is undifferentiated into distinctive sepals and petals. (flower part, perianth part)
- terete** Cylindrical. (shape)
- terminal** (a) At or near the top, tip, or end of a structure. (position) (b) Entire inflorescence positioned as a terminal shoot relative to the nearest vegetative leaves. (inflorescence position) (c) Style arising at the apex of the ovary. (style position)
- terminal bud** Bud at the apex or end of a stem. Syn: *apical bud*. (bud type)
- terminal bud scale scar** Ringlike marks indicating the former place of attachment of the terminal bud scales. (twig part)
- ternate/ternately compound** A compound leaf with three leaflets. Syn: *trifoliolate*. (leaf type)
- ternately veined** With three primary veins arising from a common basal point. (leaf venation)
- ternate-netted/ternately netted** Ternately veined, with three primary veins arising from a common basal point, the ultimate veins forming a fine reticulum. (leaf venation)
- terrestrial** Growing on land. (plant habitat)
- tertiary vein** A lateral vein that branches from and is smaller than a secondary vein. (leaf venation)
- testa** *Seed coat*. (seed part)
- tetrad** A fusion product of four pollen grains, developing from the four products of microsporogenesis.
- tetradynamous** With stamens in two groups of four long and two short, e.g. Brassicaceae. (stamen arrangement)
- tetragonal tetrad** A tetrad in which the four grains are in one plane and are equally spaced apart.
- tetrahedral tetrad** A tetrad in which the four grains form the points of a tetrahedron, e.g. members of the Ericaceae.
- tetramerous** A whorl with four members. (merosity, perianth merosity)
- tetraploid** A polyploid having four sets of chromosomes.
- tetrasporangiate** Anther with four microsporangia and typically two thecae. Cf: *dithecal*. (anther type)
- tetrasporic** Megasporogenesis in which cytokinesis does not occur after meiosis, resulting in a single cell with four haploid nuclei, all of which contribute to the female gametophyte.
- texture** Internal structural consistency, sometimes incorporating color.
- thalloid** Referring to or having a thallus.
- thallus** The flattened (dorsi-ventral) gametophyte of some liverworts and all hornworts. (plant part)
- theca** One half of typical anther containing two microsporangia. (anther part)
- thorn** A sharp-pointed stem or shoot (cf. prickle, spine). (stem/shoot type)
- throat** An open, expanded region of a perianth, usually of a sympetalous corolla. (perianth part)
- thrum** Flowers with a short style and long stamens, found in distylous flowers.
- thylakoid** A membrane of chloroplasts and photosynthetic bacteria that contains compounds involved in the light reactions of photosynthesis.
- thyrs** A secondary inflorescence with an indeterminate central axis bearing opposite, lateral, pedicellate cymes, e.g. *Echium* (Boraginaceae), *Penstemon* (Plantaginaceae). (inflorescence type)
- tiller** A grass shoot produced from the base of the stem. (stem/shoot type)
- tissue** A group of cells having a common function or structure; may be simple or complex.
- tomentose** Covered with dense, interwoven trichomes. (vestiture)
- tomentulose** Minutely tomentose. (vestiture)
- torus** *Compound receptacle*. (inflorescence part)
- total cover** A measure of the degree to which the total area of the community is covered by members of a given layer.
- tracheary element** A generally elongate cell that is dead at maturity, has a lignified, secondary cell wall, and is positioned end-to-end with other tracheary elements, forming a continuous

- tube that functions in water and mineral conduction; the major component of xylem tissue.
- tracheid** The ancestral type of tracheary element that is imperforate, in which water and mineral nutrients flow between adjacent cells through the primary cell walls at pit-pairs.
- transfer in position** To reclassify a taxon without a change in rank; to place within a different taxon of the same rank.
- transformation series** The hypothesized sequence of evolutionary change, from one character state to another, in terms of direction and probability. Syn: *morphocline*.
- transverse** (a) Dehiscing at right angles to the long axis of the theca. (anther dehiscence type) (b) Referring to a cross-section of wood.
- transverse posture** Placement of tip with respect to the horizontal plane of object. (disposition)
- trap leaf** A leaf that mechanically moves after being triggered, capturing and digesting small animals, e.g. *Aldrovanda*; *Dionaea* venus fly trap. (leaf structural type)
- tree** A generally tall, perennial, woody plant having one main stem (the trunk) arising at ground level. (plant habit)
- triangular** Three-sided, length : width ratio 2:1 to 3:2. (shape)
- trichome** An external, hairlike plant structure.
- trichotomosulcate** A pollen grain aperture type that is three-branched.
- tricyclic** *Triseriate*. (cycly)
- trifoliolate** *Tenat/ternately compound*. (leaf type)
- trigger mechanism** A type of movement hercogamy in which an insect pollinator triggers the sudden movement of one or more stamens, dusting the insect with pollen at the point of contact.
- trilete mark** A three-lined imprint on the spore wall of some land plants, being the remnant (scar) of attachment of the adjacent three spores of a tetrad.
- trimerous** A whorl with three members. (merosity, perianth merosity)
- trionoecious** A plant with pistillate, staminate, and perfect flowers on the same individual. (plant sex)
- trinomial** A scientific name composed of three names, such as a subspecies or variety name.
- trinucleate** Having three nuclei, referring to some angiosperm pollen grains at the time of release.
- trioecious/trioecy** A plant with pistillate, staminate, and perfect flowers on different individuals. (plant sex)
- tripinnate/tripinnately compound** A compound leaf with three orders of axes, the third (rachillae) bearing leaflets. (leaf type)
- triploid** A polyploid having three sets of chromosomes.
- triseriate** With three whorls of parts. Syn: *tricyclic*. (cycly)
- tristichous** Alternate, with points of attachment in three vertical rows/ranks, e.g. the sedges, Cyperaceae. (arrangement)
- tristyly** Hercogamy in which there are three heights of styles and stamens. Adj: *tristylous*.
- trisulcate** A pollen grain with three elongate apertures on opposite sides of the grain, parallel to the equatorial plane.
- trullate** Parallelogram-shaped, widest near base and the length: width ratio 2:1 to 3:2. (shape)
- truncate** Apical or basal margin cut straight across, the angle approximately 180°. (apex, base) .
- tryma** A nut surrounded by an involucre that dehisces at maturity, e.g. *Carya*, pecan. (fruit type)
- tube** A cylindrically shaped perianth or region of the perianth, usually of a sympetalous corolla. (perianth part)
- tube cell** One of the two initial, haploid cells in the male gametophyte of angiosperms that remains near the tip of the growing pollen tube and may function in its development.
- tuber** A thick, underground storage stem, usually not upright, typically bearing outer buds and lacking surrounding storage leaves or protective scales, e.g. *Solanum tuberosum*, potato. (stem/shoot type)
- tuberculate** *Papillate*. (epidermal excrescence)
- tubular** Cylindrical. (perianth type)
- turbinate** Top- or turban-shaped. (shape)
- twining** Twisted around a central axis. (twisting/bending posture)
- two-armed** Trichomes with two arms arising from a common base, e.g. Malpighiaceae. (trichome type)
- type/type specimen** *Nomenclatural type*.
- typus conservandus** Meaning a conserved type specimen. Abbr: *typ. cons.*
- typus designatus** Referring to the designation of a type specimen. Abbr: *typ. des.*
- ulcerus** A circular to slightly elliptic aperture similar in shape to a porus (length : width ratio <2:1) occurring at the (usually distal) pole. Adj: *ulcerate*.
- umbel** A determinate or indeterminate, flat-topped or convex inflorescence with pedicels attached at one point. (inflorescence type)
- uncinate** *Hooked*. (bristle type)
- undulate** Margins wavy in a vertical plane. (longitudinal posture)
- unguiculate** Clawed, e.g. many Brassicaceae, Caryophyllaceae. (perianth type)
- unicarpellous** With gynoecium composed of one carpel, the pistil or ovary being simple. (gynoecial fusion)
- unicellular** Referring to a trichome (or other structure) consisting of a single cell. (trichome type)
- unifacial leaf** A leaf that is isobilateral and flattened side-to-side, having left and right sides, except at the base, where it is often sheathing, e.g. members of the Iridaceae. (leaf structural type)
- unifoliolate** A leaf bearing a single leaflet with petiolule distinct from the petiole, interpreted as the derived reduction of an ancestral compound leaf, e.g. *Cercis*, redbud. (leaf type)
- unilateral** *Secund*. (arrangement)
- unilabiate** One-lipped; with one, generally lower, segment, e.g. many Goodeniaceae. (perianth type)
- unilocular** An ovary with a single locule. (locule number)
- uninervous** Having a central midrib with no lateral veins, e.g. lycopods, psilophytes, and equisetophytes, many conifers. (leaf venation)
- uniseriate** (a) With a single whorl of parts. (cycly, stamen cycly, perianth cycly) Syn: *monocyclic*. (b) Trichomes having a single vertical row of cells. (trichome type) (c) Rays in wood that are composed of a single, vertical row of cells.
- unisexual** Flowers having only carpel(s) or only stamen(s). Syn: *imperfect*. (flower sex)
- unistomal** Referring to the micropyle of a unitegmic ovule.
- unit inflorescence** A subunit of the entire secondary inflorescence. Syn: *partial inflorescence*; *primary inflorescence*. (inflorescence part)
- unitegmic** An ovule with a single integument, found in all gymnosperms and derived in some angiosperms.

- unordered** Referring to a character transformation series in which each character state is allowed to evolve into every other character state with equal probability, generally in a single evolutionary step.
- unrooted tree** A branching diagram in which polarity is not indicated, representing the relative character state changes between taxa. Syn: *network*.
- upper epidermis** The adaxial epidermis of a leaf.
- upward** Dehiscing toward the sky in a horizontally oriented flower. (anther dehiscence direction)
- urceolate** Urn-shaped; expanded at the base and constricted at the apex, e.g. many Ericaceae. (perianth type)
- urent** With hispid and stinging trichomes. (vestiture)
- utricle** A small, bladdery or inflated, one-seeded, dry fruit; an achene in which the pericarp is significantly larger than the mature seed, e.g. *Atriplex*, salt bush. (fruit type)
- vacuole** A large, internal, membrane-bound sac of plant cells that functions in storage of compounds such as pigments, acids, or ergastic substances.
- valid name** A name that is validly published.
- validly published** Referring to the criteria needed for a scientific name to be formally recognized, including effective publication, publication in correct form, publication with a Latin description or diagnosis or with a reference to such, and indication of a nomenclatural type.
- valvate** (a) Sides enrolled, so that margins touch. (arrangement) (b) Having a whorl of perianth parts meeting at the margins and not overlapping. (perianth aestivation)
- valve** A portion of the pericarp of a dehiscent fruit that splits off but does not enclose the seed(s). (fruit part)
- valvular** Anther dehiscing through a pore covered by a flap of tissue, e.g. Lauraceae. (anther dehiscence type)
- valvular capsule** *Septifragal capsule*. (fruit type)
- variegated** With two or more colors occurring in various irregular patterns, generally used for leaves. (color pattern)
- varietal epithet** The third component of a trinomial at the rank of variety.
- vascular bundle** A strand of associated xylem and phloem tissue. Syn: *vascular strand*.
- vascular bundle scar** A mark within the leaf scar indicating the former position of a vascular bundle that extended from stem to leaf. (twig part)
- vascular bundle sheath** A ring of cells, composed of fiber or parenchyma cells, surrounding the xylem and phloem of a vascular bundle, functioning in C4 photosynthesis.
- vascular cambium** A cylindrical sheath of cells that undergoes primarily tangential divisions, producing secondary xylem (wood) to the inside and secondary phloem (inner bark) to the outside.
- vascular cylinder** A central region of vascular tissue (xylem and phloem) in a root (root part) or in some stems (stem part).
- vascular cylinder** A central region of xylem and phloem in the root.
- vascular strand** *Vascular bundle*. (stem/shoot parts)
- vascular tissue** A tissue made up of xylem and phloem, functioning mainly in conduction of water, minerals, and sugars.
- vegetation type** An assessment of the habit, habitat, and cover of plant species present, e.g. forest, woodland, savanna, chaparral, scrub, grassland, meadow, strand marsh, swamp, pond, and vernal pool.
- vegetative bud** A bud that develops into a vegetative shoot, bearing leaves. (bud type)
- vein** The vascular bundle of a leaf or leaf homologue such as a sepal, petal, stamen, or carpel, containing the conductive tissues. Syn: *nerve*. (plant part, leaf part)
- velamen** A specialized, multilayered epidermis of some roots, functioning in protection, prevention of water loss, or water and mineral absorption, e.g. Araceae, Orchidaceae.
- venation** The pattern of veins and vein branching of a leaf or leaf homologue. (leaf venation)
- venter** The swollen, basal portion of an archegonium, containing the egg.
- ventral** *Adaxial*. (position)
- ventral canal cell** A second cell within the swollen base of the archegonium, located just distal to the egg,
- ventral vein** One of the two veins of a carpel near the carpel margins. Syn: *lateral vein*, *placental vein*.
- vernal** Appearing in spring. (periodicity)
- verruca** One of the short, wartlike elements of a verrucate pollen grain. Pl: *verrucae*.
- verrucate** (a) *Papillate*. (epidermal excrescence) (b) A pollen sculpturing having short, wartlike elements, each element termed a verruca.
- versatile** With anther freely pivoting at the point of attachment with the filament. (anther attachment)
- verticillaster** A secondary inflorescence with an indeterminate central axis bearing opposite, lateral, sessile cymes, the flowers appearing congested, e.g. some Lamiaceae. (inflorescence type)
- verticillate** Having three or more leaves or other structures per node. (arrangement)
- very widely obovate** Margins curved, widest near the apex, length : width ratio ca. 1:1. (shape)
- very widely ovate** Margins curved, widest near the base, length : width ratio ca. 1:1. (shape)
- vespertine** In the evening, typically with respect to when flowers open. (periodicity)
- vessel** Term for several vessel members attached end-to-end, forming a continuous, conductive tube. Syn: *pore*.
- vessel member** A type of tracheary element that is perforate, in which continuous holes or perforations in the cell walls occur through which water and mineral nutrients flow between cells.
- vestiture** Trichome cover, a combination of trichome type, length, strength, shape, density, and color. (surface)
- vicariance** The splitting of one ancestral population into two (or more) populations, such as by continental drift or the formation of a geographic barrier.
- vide** Meaning to cite a reference.
- villous** Covered with very long, soft, crooked trichomes. Syn: *lanate*. (vestiture)
- vine** A plant with elongate, weak stems, supported by means of scrambling, twining, tendrils, or roots; may be annual or perennial, herbaceous or woody. (plant habit)
- viscid** Having a shiny, sticky surface. Syn: *glutinous*. (epidermal excrescence)
- viscin thread** One of the long strands of carbohydrate material that function in sticking pollen grains together.
- voucher specimen** An herbarium specimen in a plant collection serving as reference material for a named taxon or as part of a research project.

wax A type of triglyceride compound that may function as high-energy storage compounds or secretion products; a type of ergastic substance.

weight The specific assignment of taxonomic importance when weighting a character in a phylogenetic analysis.

weighting/character weighting The assignment of greater or lesser taxonomic importance to certain characters over other characters in determining phylogenetic relationships.

whorl A cyclic group of floral parts, e.g. of sepals, petals, stamens, or carpels. Syn: *series*. (flower part)

whorled (a) With three or more leaves or other structures per node. (leaf arrangement) (b) Perianth parts in distinct whorls or series, with parts arising from the same nodal region. (perianth arrangement) (c) With stamens in one or more whorls or series. (stamen arrangement)

widely elliptic Margins curved, widest near the midpoint, length : width ratio ca. 6:5. (shape)

widely obovate Margins curved, widest near the apex, length : width ratio ca. 6:5. (shape)

widely ovate Margins curved, widest near the base, length : width ratio ca. 6:5. (shape)

widely triangular Three-sided, length : width ratio ca. 6:5. (shape)

wood *Secondary xylem*.

woody Having a hard, woodlike texture. (texture)

wrinkled With irregular, fine lines or deformations. (configuration)

× Abbreviation, used within a scientific name, that indicates a hybrid.

xenogamy *Outbreeding*. Adj: *xenogamous*.

xylem A tissue composed of tracheary elements plus some parenchyma and sometimes sclerenchyma, functioning in conduction of water and mineral nutrients.

zig-zag Referring to the micropyle of a bitegmic ovule in which the micropylar pore of the outer integument is spacially displaced relative to the inner integument.

zonoaperturate Descriptive of pollen grains with apertures occurring in the equatorial region. Syn: *stephanoaperturate*.

zonocolpate A pollen grain with colpi occurring in the equatorial region.

zonoporate A pollen grain with pori occurring in the equatorial region.

zoochory Dispersal of propagules by animals.

zygomorphic Bilaterally symmetrical, with one plane of symmetry. Syn: *bilateral, irregular*. (symmetry)

zygote A diploid cell that results from the fusion of two haploid cells (egg and sperm) and that ultimately matures into a new sporophyte in the land plants.

INDEX

Note: Page numbers in bold indicate presence of photographic images for specific taxa.

- Abbreviations (nomenclatural), 509-510
ABC model, 122
Abelia, **333**
Abelmoschus, 286
Abies, **116**
Abolboda, 212
Acacia, 262, **263**
Acanthaceae, 308-**309**
Acanthogilia, **299**
Acanthomintha, **312**
Acanthus, 308, **309**
Accenting, 512-513
Accessioning, 529
Acer, 289, **291**
Acetolysis, 460
Achariaceae, 255
Achatocarpaceae, 230
Achlyphila, 212
Acoraceae/*Acorus*, 157-**159**
Acorales, 157
Acourtia, **329**
Actaea, **234**
Actinidiaceae, 292-293
Actinodium, **258**
Adansonia, 286
Adaptation, 10
Adiantum, **83**, 88, **89-90**
Adoxaceae, 292, 331, **333**
Advanced, *see* Derived
Adventive embryony, 472
Aechmea, **204**
Aeonium, 252
Aextoxicaceae, 230
AFLP (Amplified Fragment Length Polymorphism), 487, 489-490
Agamospermy, 472
Agapanthaceae, 159, 165-**166**
Agapanthus, **166**
Agavaceae, 165, **167-168**
Agave, 165, **167**
Aglaonema, 160, **162**
Agrostis, **217**
Aizoaceae, 238, 240-241, **242**
Akaniaceae, 255
Albuminous cell, 73
Alectryon, **290**
Alethopteris, 107
Aleurites, **270**
Alifaria, 254
Alignment, 482-483
Alismataceae, 160, **164**
Alismatales, 160-161
Allenrolfea, **243**
Alliaceae, 165, **167-169**
Allium, 167, **169**
Allozyme, 484, 488
Alnus, 265, **267**
Alocasia, 160
Aloe, 171-**172**
Alpinia, 198, **200**
Alseuosmiaceae, 292
Alstroemeria, **181**
Alstromeriaceae, 159, 180-**181**
Alternation of generations, *see*
Haplodiplontic life cycle
Altingiaceae, 230, 250
Alyogyne, **284, 285**
Alzateaceae, 255
Amaranthaceae/*Amaranthus*, 241,
243, 244
Amaryllidaceae, 169-**170**
Amaryllis, 169
Amborellaceae/*Amborella*, **142-143**
Amborellales, 141-142
Ambrosia, **329**
Amoeboid (tapetum), 439
Amorpha, **263**
Amorphophallus, 160, **163**
Amphianthus, **318**
Amsinckia, **300**
Amyloplast, *see* Starch, grain
Anacardiaceae, 286-**287**
Anacardium, 286
Ananas, **204**, 205
Anarthriaceae, 185
Anatomy, 409-436
Anatropous, 445-446
Ancestral, 13, 18, 34-35
Ancistrocladaceae, 230
Androecium, 121, 371-374
Anemone, **234**
Anemopsis, **154**
Anethum, 322
Angiosperms, *see* Flowering plants
Angraecum, 177
Anigozanthos, 190, **191**
Anisophylleaceae, 255
Annonaceae/*Annona*, 146, **148-149**
Annotation labels, 530-531
Annotation, *see* Classification, annotated
Annual ring, 423-425
Annulus, 88
Anther, 124
attachment, 373
dehiscence, 373-374
parts, 373, 374
type, 373, 374, 437
wall development, 437
Antheridial wall, 58
Antheridiophore, 59
Antheridium, 58, 88
Anthoceros, **62**
Anthocerotae, *see* Hornworts
Anthurium, 160, **162**
Antigonon, **248**, 249
Antipodal cells, 130, 441, 445
Antirrhinum, 315, **318**
Apex, 392-393
Aphanopetalaceae, 230
Aphelandra, 308
Aphloiaceae, 255
Aphyllanthaceae, 159
Apiaceae (Umbelliferae), 322, **323-324**
Apiales, 322
Apical meristem, 57, 75, 77, 412
Apium, 322, **323**
Apocynaceae, 301, **303-304**, 305
Apodanthaceae, 230
Apomorphic, *see* Derived
Apomorphy, 13, 18, 24
Apomorphy-based, *see* Classification,
apomorphy-based
Aponogetonaceae, 159-160
Aptenia, **242**
Aquifoliaceae, 292
Aquifoliales, 292
Aquilegia, **234**
Araceae, 160, **162-163**
Arachis, 262
Araliaceae, 322-**325**
Aralidiaceae, 292
Aratitiopea, 212
Araucaria, **116**
Araucariaceae, 114, **116**
Arbutus, 294, **295**
Arceuthobium, 249, **250**
Archaea, 3
Archaeofructus, 134-**135**
Archegonial
chamber, 105
wall, 58
Archegoniophore, 59
Archegonium, 58, 88

- Archeopteris*, 107, **108**
Archontophoenix, **186**
Arctostaphylos, 294
Areca, 188
 Arecaeae (Palmae), 185, **186-187**, 188
 Arecales, 185
Arenaria, **247**
 Argophyllaceae, 292
Argyroxiphium, **326**
Arisaema, **162**
 Aristolochiaceae/*Aristolochia*, 149, **151**
Armeria, 238
 Arrangement, 396-397
Artocarpus, 275
Asarum, 149, **151**
 Asclepiadaceae, *see* Apocynaceae
Asclepias, **304**
 Asexual reproduction, 472
 Asparagaceae, 159, 165
 Asparagales, 163, 165-166
 Asphodelaceae, 171-**172**
Asphodelus, 171-**172**
Asplenium, **89**
 Asteliaceae, 159
Aster, **328**
 Asteraceae (Compositae), **326-330**, 331
 Asterales, 322
Asterella, **61**
 Asterids, 289, 291-293
 Asteropeiaceae, 230
 Atactostele, 155, 420, 421
 Atherospermataceae, 140
Atrichum, **65**
Atriplex, **243**
Atropa, 319
 Attractant, 466
 Austrobaileyaceae, 140
 Austrobaileyales, 143
 Autapomorphy, 18, 25
 Authorship, 504
 Autonym, 507
Avena, 216, **218**
Averrhoa, 272, **274**
 Axil, 76
Azalea, *see* *Rhododendron*
Azolla, 88, **92**
- Bacteria, 3
 Balanopaceae, 255
 Balanophoraceae, 230
 Balsaminaceae, 292
Banksia, **237**
 Barbeuiaceae, 230
 Barbeyaceae, 255
 Base, 392
 Basellaceae, 230
 Bataceae, 255
Bauhinia, **263**
 Bayesian analysis/inference, 36
Bazania, **61**
 Begoniaceae, 255
 Berberidaceae/*Berberis*, 228, 230-**231**
 Berberidopsidaceae, 230
 Berberidopsidales, 230
- Beta*, 244
 Betalains, 238-239, 241
 Betulaceae/*Betula*, 265, **267**
 Biebersteiniaceae, 255
 Bignoniaceae, 308, **310-311**
 Binomial, 12, 502
 Biogeography, 41
 Bisporic, 441, 444
 Bixaceae, 255, 280
Blandfordia, **166**
 Blandfordiaceae, 159, 165-**166**
Blighia, 289
Bloomeria, **178**
Bomarea, **181**
 Bonnetiaceae, 255
 Bootstrap, 37-38
 Boraginaceae s.s., 298, **300-301**
Borago, **300**, 301
 Boryaceae, 159
Boschniakia, **316**
 Botanical names, *see* Names, botanical
 Botany, 9
Botrychium, 83-**84**
Bougainvillea, **239**
Bowenia, 108, **109**
Brachychiton, **283**, 286
Brachypodium, **218**
 Bract, 114
Brasenia, 143, **145**
Brassaia, **325**
 Brassicaceae (Cruciferae)/*Brassica*,
280-281
 Brassicales, 278, 280
 Breeding
 mechanism, 474
 systems, 468-472
 Bremmer support, *see* Decay index
 Bristle type, 401-402
Brodiaea, **178**
 Bromeliaceae, 203, **204**, 205
 Brown plants, 4-5
Brugmansia, **321**
 Brunelliaceae, 255
 Bruniaceae, 292
Bryophyllum, **251**
 Bud, 76, 355-356
Bulbine, **172**
Burchardia, **181**
 Burmanniaceae, 159
 Burseraceae, 255, 286
 Butomaceae, 159
 Buxaceae, 230
 Buxales, 230
 Byblidaceae, 292
- Cabomba*, 143
 Cabombaceae, 143, **145**
 Cactaceae, 244, **245-246**
Caesalpinia, **263**
Cakile, **281**
Caladenia, **174**
Caladium, 160
 Calamites, **85**, 97
Calamus, **187-188**
- Calathea*, **202**
 Calceolariaceae, 292, 310
Calindrinia, **239**
Calliandra, **263**
Callistemon, **258**, 259
 Callitricheaceae, *see* Plantaginaceae
Callitriche, 315, **318**
 Callose, 415
Calochortus, **182**
Calodendrum, **288**
Calonyction, **320**
Calopogon, **174**
Calothamnus, **258**
 Calycanthaceae, 140
 Calyceraceae, 292
 Calyptra (=Operculum)
 flower, 256, 258
 moss, 63
Calystegia, **320**
 Calyx, 121; *see also* Perianth
Camassia, 165, **168**
Camissonia, **260**
 Campanulaceae, 292
Campsis, **311**
 Campylotropous, 446
 Campynemaceae, 159
 Canellales/Canellaceae, 140
 Cannabaceae, 255, 275
 Cannaceae/*Canna*, 198, **201**, 203
 Capparaceae, 255, 280, **282** *see also*
 Brassicaceae
 Caprifoliaceae, 331, **333**
 Capsule, 59, 348; *see* Sporangium
Cardamine, **281**
 Cardiopteridaceae, 292
Cardiospermum, **290**
Carduus, **327, 328, 329, 330**
Carex, 207, **210**
 Caricaceae, 255, 280
Carissa, **303**
 Carlemanniaceae, 292
Carnegiea, **246**
 Carpel, 121, 127-129, 374-379
 ascidiate, 128
 number, 376-377
Carpinus, 265
Carpobrotus, **242**
Carum, 322
 Caryocaraceae, 255
 Caryophyllaceae, 244, **247**, 249
 Caryophyllales, 238, **239**
 Casparian strip, 73-74, 77, 417-418
Cassia, **263**
Castanea, **268**, 269
Castilleja, **316**
 Casuarinaceae, 255, 265
Catharanthus, **303**, 305
Cattleya, **175**
Caulophyllum, **231**
 Caytonia/Caytoniales, 134-135
Ceanothus, **277**
Cedrus, **116**
Ceiba, 286
 Celastraceae, 255

- Celastrales, 255
- Cell
 differentiation, 412
 organelles, 410
 structure, 409-411
 theory, 409
 wall, 51
 primary, 410-411
 secondary, 69, 411, 412
- Cellulose/cellulosic, 51
- Celosia*, 244
- Centaurium*, **306**
- Centranthus*, **334**
- Centrolepidaceae, 185
- Centroplacaceae, 255
- Cephalotaceae, 255
- Cerastium*, **247**
- Ceratonia*, 262, **263**
- Ceratophyllaceae/*Ceratophyllum*,
 153, **155**
- Ceratophyllales, 153
- Ceratozamia*, **111**
- Cercidiphyllaceae, 230, 250
- Cercis*, **263**
- Chaenactis*, **329**
- Chamaesyce*, **270**
- Chamelaucium*, 259
- Chamerops*, **187**
- Chara*, 53, **55**
- Character analysis, 19-24
- Character state
 definition, 12, 19
 discreteness, 20-21
- Character
 binary, 22
 correlation, 21
 definition, 12, 19
 evolution, 40
 multistate, 22
 optimization, 30
 ordered, 22
 polymorphic, 31
 scaled, 23
 step matrix, 24
 unordered, 22
 weighting, *see* Weighting
 taxon matrix, 24
- Charales, 53
- Chasmanthe*, **173**
- Chelone*, 315
- Chenopodium*, **243**, 244
- Chilopsis*, **311**
- Chiranthodendron*, **284**
- Chlamydomonas*, **52**, **53**
- Chlorobionta, *see* Green Plants
- Chlorogalum*, 165, **168**
- Chlorophyceae/chlorophytes, 53
- Chlorophyll a, b, 51
- Chlorophyllous cells, 66
- Chloroplast
 definition, 4
 DNA, 82
 green plant, 51-53
 origin, 4-5
- Chondropetalum*, **214**
- Chorisia*, **284**, 286
- Chorizanthe*, **248**
- Chromoplast, 410-411
- Chrysanthemum*, **327**, **330**
- Chrysobalanaceae, 255
- Cibotium*, **90**
- Cichorium*, **328**
- Cinchona*, 305
- Cinnamomum*, 146-**147**
- Circaeasteraceae, 230
- Circinate vernation, 88, 108
- Cirsium*, **329**, **330**
- Cistaceae, 255, 280
- Citrullus*, 252
- Citrus*, **288**, 289
- Clade, *see* Lineage
- Cladistics, *see* Phylogenetic systematics
- Cladogram
 analysis, 38-43
 construction, 24-38
 definition, 13, 17
 of life, 4
 robustness, 37
- Clarkia*, 259, **260**
- Classification
 annotated, 39
 apomorphy-based, 39-40
 definition of, 12
 indented, 38
 node-based, 39
 phenetic, 13, 40
 phylogenetic, 13, 38-40
 stem-based, 39
- Clematis*, **234**
- Cleomaceae/*Cleome*, 255, 280, **282**
see also Brassicaceae
- Clethraceae, 292
- Clitoria*, **264**
- Clivia*, 169-**170**
- Clusiaceae, 255
- Cneoridium*, **288**
- Cobaea*, 298
- Cocoloba*, **248**
- Cocos*, 188
- Coffea*, 305, **307**
- Cola*, 286
- Colchicaceae, 159, 180-**181**
- Coleochaete*, **55**
- Collecting, 517-524
 for molecular studies, 524
 liquid preserved, 523
 living, 523-524
- Collection, synoptic, 497, 530
- Collenchyma, 71
- Colletia*, **277**
- Collinsia*, 315, **318**
- Colocasia*, 160
- Color, 389
- Columella, 63
- Columelliaceae, 292
- Comandra*, **250**
- Combretaceae, 254-255
- Comesperma*, **266**
- Commelinaceae/*Commelina*, 188, **189**, 190
- Commelinales, 184, 188
- Commelinid Monocots, 184-185
- Commemoratives, 510-511, 513
- Common ancestor, 13, 17
- Community type, 522-523
- Companion cell, 73
- Compositae, *see* Asteraceae
- Cone, 348
 conifer (compound), 114
 cycad, 108
 equisetophyte, 87
 lycophyte, 79-80
- Configuration, 399-400
- Conifers/Coniferophyta, 109-116
- Conium*, 322, **323**
- Connaraceae, 255
- Conocephalum*, **61**
- Conospermum*, **237**
- Conostylis*, **191**, **192**
- Consensus tree, 35
- Consistency index (CI), 36
- Convallariaceae, *see* Ruscaceae
- Convergence, 21
- Convolvulaceae, 319-**320**
- Convolvulus*, 319
- Copernicia*, 188
- Corchorus*, 286
- Cordiaceae, 292, 301 *see also* Boraginaceae
- Cordyline*, **166**
- Core eudicots, 238
- Corema*, **296**
- Coriandrum*, 322
- Coriariaceae, 255
- Cork/Cork cambium, 97, 422, 423
- Cornaceae/*Cornus*, 291, 293-**294**
- Cornales, 291
- Corolla, 121; *see also* Perianth
- Corsiaceae, 159
- Cortex, 74, 77, 420
- Corylus*, 265, **267**
- Corynocarpaceae, 255
- Costa, 63
- Costaceae, 185, **202**
- Costus*, **202**
- Cotoneaster*, 278
- Cotula*, **329**
- Cotyledon, 98, 105
 single (monocots), 155
- Couplet, 12
- Cover, 522
- Crassinucellate, 440, 443
- Crassulaceae/*Crassula*, 250, **251**, 252
- Cressa*, **320**
- Crinum*, 169-**170**
- Crocus*, 171, **173**
- Crossosomataceae, 255
- Crossosomatales, 255
- Crozier, *see* Fiddlehead
- Cruciferae, *see* Brassicaceae
- Cryptantha*, **300**
- Crytal, 410-411
- Cryteroniaceae, 255
- Ctenolophonaceae, 255

- Cucumis*, **261**, 262
 Cucurbitaceae/*Cucurbita*, 259, **261**, 262
 Cucurbitales, 259
Cuminum, 322
 Cunoniaceae, 255
 Cupressaceae/*Cupressus*, 114, **116**
 Cupule, 134
 Curator, 529
Curcuma, 198
 Curtissiaceae, 292
Cuscuta, **320**
 Cuticle/Cutin, 5, 55, 416
 Cyads/Cyadophyta, 108-109
Cyanotis, **189**
Cyathea, 88, **89**
Cycas, 108, **110**
 Cyclanthaceae, 159
Cylindropuntia, **246**
Cymbidium, **175**
 Cymodoceaceae, 159-160
Cynara, 331
Cynodon, **218**
 Cynomoriaceae, 230
 Cyperaceae, 206-207, **209-210**, 211
Cyperus, **209**, 211
Cypripedium, **175**
 Cyrillaceae, 292
 Cytinaceae, 230
- Dampiera*, **332**
Danaea, **87**
 Daphniphyllaceae, 230
Darwinia, **258**
 Data information systems, 532-533
 Datisceae, 255
Datura, 319, **321**
Daucus, 322, **323**
Daviesia, **264**
 Decay index, 38
Dedekera, **248**
 Degeneriaceae, 140
Delphinium, **234**
Dendrobium, **175**
Dendrocalamus, **216**
 Dendrochronology, 98, 426
Dendromecon, **232**
 Derived, 13, 18
 Descent, 10
 Description, 12, 535-539
Dianella, **166**
Dianthus, 246
 Diapensiaceae, 292
Dicentra, **232**
Dichelostemma, **178**
Dichondra, **320**
Dichorisandra, **189**
 Dichotomous key, *see* Key
 Didiereaceae, 230
 Didymelaceae, 230
Dieffenbachia, 160
 Diervillaceae, 292, 331
Dietes, **173**
Digitalis, 315, **318**
Dilatris, **191**
- Dilleniaceae, 230
 Dillineales, 230
Dimocarpus, 289
 Dinoflagellates, 4-5
 Dioecious/dioecy, 108, 109, 114, 365, 468
Dionaea, **239**
 Dioncophyllaceae, 230
Dioscorea, **179**, 180
 Dioscoreaceae/*Dioscorea*, 178, **179**, 180
 Dioscoreales, 178
 Dipentodontaceae, 255
Diphasiastrum, 79
Diphylleia, **231**
 Diplohaplontic life cycle, *see*
 Haplodiplontic life cycle
 Dipsacaceae/*Dipsacus*, 331, **334-335**
 Dipsacales, 331
 Dipterocarpaceae, 255, 280
 Dirachmaceae, 255
 Dispersal, 41
 Disposition, 396-399
Distichia, 206
Distichlis, **218**
Distictis, **311**
 Divergence/diversification, 10, 17
 Division, 394-396
 DNA
 amplification, 478
 analysis, 482-483
 chloroplast, 481
 evolution of, 3
 nuclear, 481
 sequence data, 477-478, 479-482
 sequencing reaction, 479-480
 Documentation, 519-523
Dodonaea, **290**
Dombeya, **283**, 286
 Donatiaceae, *see* Stylidiaceae
Dorstenia, 275, **276**
 Doryanthaceae, 159, 165
Doryanthes, **166**
 Double fertilization, 114
Drosera, **239**
 Droseraceae, 230, 238, **239**
 Drosophyllaceae, 230
Dryopteris, **90**
Dudleya, **251**
 Duration, 402
Durio, **283**, 286
Dyckia, **204**
- Ebenaceae, 292-293
 Ecdeiocolleaceae, 185
Echeveria, 252
Echinocereus, **245**, **246**
Echinodorus, **164**
Echium, **300**
 Ecology, 41
 Egg, 53
 angiosperm, 130, 441, 445
 Ehretiaceae, 292, 301 *see also* Boraginaceae
Eichhornia, 190, **193**, **194**
 Elaeagnaceae, 255
Elaeis, 188
- Elaeocarpaceae, 255
 Elaters, 59
 Elatinaceae, 255
Elegia, **214**
Eleocharis, **210**
Elettaria, 198
Elodea, **52**
Elymus, **217**
 Emblingiaceae, 255
 Embryo, 54, 98
 development, *see* Embryogeny type, 447
 Embryogeny, 446-447
 Embryology, 437-451
 Embryophyta/embryophytes, *see* Land Plants
Emmenanthe, **302**
 Empetraceae, *see* Ericaceae
Empodisma, **214**
 Enantiostyly, 469-470
 Enation, 78, 84
Encelia, **327**, **328**, **330**
Encephalartos, **109**, **111**
Encyclia, **175**
 Endarch, 78, 107, 421
 Endodermis, 73-74, 77, 417-418
 Endosperm, 130-132
 development, 447-448
 starchy, 188
 type, *see* Seed, endosperm type
 Endospore/endosporic/endospory,
 91, 97, 101
 Endosymbiosis, 4-5, 53
 Endothecium, 437
Ensete, 192, 195
 Epacridaceae, *see* Ericaceae
Epacris, **296**
 Ephedraceae/*Ephedra*, 114, **117**
Epidendrum, **175**
 Epidermal excrescence, 399-400
 Epidermis, 55, 416-417
Epifagus, **316**
Epilobium, **260**
Epipremnum, 160
 Equisetales/equisetophytes, 85, **86**, 87
Equisetum, 85, **86**, 87
 Eremosynaceae, 292
 Ergastic substances, 409-411
Erica, 294, 295, **296**
 Ericaceae, 293-294, **295**, **296**
 Ericales, 293
 Eriocaulaceae/*Eriocaulon*, **211-212**
Eriodictyon, **302**
Eriogonum, **248**
Eriostemon, **288**
Erodium, 254, **256**
Erythrina, **264**
Erythronium, **182**
 Erythropalaceae, 230
 Erythroxylaceae, 255, 269
 Escalloniaceae, 292
Eschscholzia, **232**
 Euasterids I, 298
 Euasterids II, 319
Eucalyptus, 256, **258**, 259
Eucharis, 169-170

- Eucommiaceae, 292
Eucrypta, **302**
 Eudicots, 228-230
 Euglenoids, 4-5
 Eukarya/eukaryotes/eucaryotes, 3
 Euphorbiaceae/*Euphorbia*, 269-**270**
 Euphyllphytes/euphyll, 82, 427
 Eupomatiaceae, 140
 Eupteleaceae, 230
 Eurosids I, 259
 Eurosids II, 278
 Eusporangium, 84, 87
 Eustele, 105-107, 420
Eustoma, 305
 Evolution
 biological, 10
 of chloroplasts, *see* Endosymbiosis
Exacum, 305, **306**
 Exarch, 78, 421
 Exine, 460
 Exospory, 99
- Fabaceae (Leguminosae), 262,
 263-264, 265
 Fabales, **262**
 Fagaceae, 265, **268-269**
 Fagales, 265
Fagopyrum, **248**, 249
Fagus, 269
 False indusium, 88
 Family names, alternative, 503
 Fascicle, *see* Short shoot
Fatsia, 322, **325**
Faucaria, **242**
 Female gametophyte, 99, 130
 Female sporophyll, *see* Megasporophyll
Fenestera, **242**
 Ferns, *see* Monilophytes
Ferocactus, **246**
 Fertile segment, 83
 Fertilization, 53-54, 465
 Fiber, 70-71, 413, 414
Ficus, 275, **276**
 Fiddlehead, 88
 Filament, 124
 Flagellariaceae, 185
 Flora, 533
 Floral diagram, 141, 541-543
 Floral formula, 140-141, 538
 Flower, 121-124, 364-366
 attachment, 364, 365-366
 cyclic, 366, 367
 development, 122, 124
 maturation, 366
 parts, 364-365
 sex, 364, 365
 symmetry, 366
 Flowering plants
 apomorphies and evolution, 121-133
 clades/classification, 138-140
 origin, 133-135
Foeniculum, 322, **323**, **324**
 Fouquieriaceae/*Fouquieria*, 295, **297**, 298
Fragaria, 278
- Frankenia*, **239**
 Frankeniaceae, 230, 238, **239**
Freesia, 171, **173**
Fremontodendron, **284**, **285**, 286
Fritillaria, 180, **182**
 Frond, 88
 Fruit, 129, 348, 382-387
 aggregate, 386, 387
 dispersal, 472
 multiple, 386-387
 simple, 384-386
 Fumariaceae, *see* Papaveraceae
Fuschia, 259, **260**
 Fusion, 389-390
- Galanthus*, 169
Galium, **307**
 Gametangia, 5; *see also* Antheridium,
 Archegonium
 Gamete, 53, 54
 Gametophyte, 54
 Garryaceae, 292
 Garryales, 292
Gasteria, 171-**172**
Gazania, **327**
 Geissolomataceae, 255
Geitonoplesium, **181**
 Gelsmiaceae, 292
 Gemmae/gemma cups, 59
 Gender, 510
 General terminology (morphology),
 389-403
 Generative cell, 126
 Genetic drift, 10
Gentiana, 305
 Gentianaceae, 305-**306**
 Gentianales, 301
Geosiris, 171, **173**
 Geraniaceae/*Geranium*, 252, 254, **256**
 Geraniales, 252
 Gesneriaceae, 292, 308
Gilia, 298, **299**
Ginkgo/Ginkgophyta, 109, **112**
 Gisekiaceae, 230
Gladiolus, 171
Glycine, 262
 Gnetaceae/*Gnetum*, 114, **117**
 Gnetales, 114, **117-118**
 Gomortegaceae, 140
Gomphocarpus, **304**
 Goodeniaceae, 331-**332**
Gossypium, 286
 Goupiaceae, 255
 Gramineae, *see* Poaceae
 Granum/grana, 51
Graptopetalum, **251**
 Gratiolaceae, 292, 310
 Green plants, 4-6, 51-54
Grevillea, **237**
Grewia, **283**
 Griselinaceae, 292
 Grossulariaceae, 230, 250
 Growth, 411-412
 secondary, *see* Secondary growth
- Grubbiaceae, 292
 Gryostemonaceae, 255
 Guard cells, 59
Guichenotia, **283**
 Gunneraceae, 230
 Gunnerales, 230
 Guttiferae, *see* Clusiaceae, 255
 Gymnosperms, 107-118
Gymnostachys, **162**
 Gynoecium, 121
 apocarpous, 129
 fusion, 376
 syncarpous, 129
- Haemodoraceae, 190, **191**, **192**
Haemodorum, **191**
 Halophytaceae, 230
 Haloragaceae, 230, 250
 Hamamelidaceae, 230, 250
 Hanguanaceae, 185
 Haplodiplontic life cycle, 54
 Haplontic life cycle, 53
 Harmomegathy, 458
Haworthia, 171-**172**
 Hectorellaceae, 230
Hedera, 322, **325**
Hedychium, 198, **200**
Helianthus, 331
Heliconia, **202**
 Heliconiaceae, 185, **202**
 Heliotropaceae, 292, 301 *see also*
 Boraginaceae
 Helwingiaceae, 292
 Hemerocallidaceae, 159, 165-**166**
Hemerocallis, **166**
Hemiandra, **312**
Hepatica, **234**
 Hepaticae, *see* Liverworts
Heracleum, **324**
 Herbarium, 525-534
 cabinet, 529
 insect control, 532
 labels, 525-526
 loans, 532
 use, 529-530
 Hercogamy, 469-470
Heritiera, **283**
 Hernandiaceae, 140
Hesperocallis, 165, **168**
Hesperoyucca, **168**
Heteranthera, 192, **193**, **194**
 Heterochrony, 43
 Heteropyxidaceae, 255
 Heterospory, 79, 91, 99
 Heterostyly, 469-470
Heuchera, **253**
Hevea, 269
Hexastylis, **151**
Hibiscus, **284**, **285**, 286
 Himantandraceae, 140
Hippeastrum, 169
Hippochaete, 87
 Homology, 21-22
 Homoplasy, 21

- Homospory, 79
 Hoplestigmataceae, 230
Hordeum, 216
Horneophyton, 66
 Hornworts, 59, 62-63
 Horsetails, *see* Equisetales
Hosta, 165
Houttuynia, **154**
 Huaceae, 255
Huernia, **304**
 Huerteales, 255
 Humiriaceae, 255
Huperzia, 79, **80**
 Hyacinthaceae, 159, 165
 Hyaline cells, 66
Hybanthus, **273**
 Hybridization, 472-473
 Hydatellaceae, 185
 Hydathode, 417
 Hydnoraceae, 140
 Hydrangeaceae, 291-292
 Hydrocharitaceae, 159, 160, **161**
Hydrocotyle, **324**
 Hydroid, 63
 Hydroleaceae, 292
 Hydrophyllaceae, 301-302
 Hydrostachyaceae, 292
Hydrothrix, 190
 Hypericaceae, 255
Hyphaene, 185
Hypoestes, **309**
 Hypoxidaceae, 159, 165-166
Hypoxis, **166**
Hyptis, **312**
- Icacinaceae, 292
 Identification, 12, 495-499
 Illiciaceae/*Illicium*, **145**
 Illustration, 541-544
 Inbreeding, 471
 Incompatibility reaction, 129
 Indusium, 88
 Inflorescence, 379-382
 development, 379
 parts, 379
 position, 379
 secondary, 382, 383
 specialized 382, 383
 type, 380-382
 Ingroup, 18
 Integument, 99, 439
 angiosperm, 130
 type, 441, 445
 Intercalary meristem, 63
 International Code of Botanical
 Nomenclature, 501-502, 510
 Internode
 cladogram, 18
 leaf, 75-76
 Intine, 460
 Introgression, 474
Iochroma, **321**
Ipheion, 169
Ipomoea, 319, **320**
- Ipomopsis*, 298
 Iridaceae, 171, **173**
Iris, 171, **173**
 Irvingiaceae, 255
Isatis, 280
Isocoma, **330**
 Isoetaceae/*Isoetes*, 79, **81-82**
Isomeris, **282**
 Isomorphic, 53
Isophysis, 171
 Iteaceae, 230
 ITS (Internal Transcribed
 Spacer), 481-482
 Ixerbaceae, 255
 Ixioliriaceae, 159
 Ixonanthaceae, 255
- Jacaranda*, 308, **310, 311**
 Jacket layer, *see* Antheridial wall,
 Archegonial wall
 Jackknife, 38
Jacquemontia, 319
Jagera, 289
Jaumea, **328**
Jepsonia, **253**
Johnsonia, **166**
 Joinvilleaceae, 185
 Journals, 545-546
Jubaea, **186**
 Juglandaceae, 255, 265
 Juncaceae/*Juncus*, 206, **208**
 Juncaginaceae, 159, 160, **161**
Juniperus, **116**
Justicia, 308, **309**
- Kalanchöe*, **251, 252**
Kalmia, **296**
 Key
 dichotomous, 12, 496
 polyclave, 497
Kigelia, **311**
 Kirkiaceae, 255
Kniphofia, 171-172
 Koeberliniaceae, 255
Koelreuteria, 289, **290**
 Krameriaceae, 255
- Labiatae, *see* Lamiaceae
Lachnanthes, **191**
 Lacistemataceae, 255
 Lactoridaceae, 140
Lactuca, **330**
Lambertia, **237**
 Lamiaceae (Labiatae), 308, 310, **312-313**
 Lamiales, 305, 308
 Lanariaceae, 159
 Land Plants, 5-6, 54-58
Langloisia, **299**
 Lardizabalaceae, 230
 Lateral meristem, 42, 422
 Laticifer, 417
 Lauraceae/*Laurus*, 146-**147**
 Laurales, 146
- Lavandula*, 310
 Laxmanniaceae, 159, 165-**166**
Layia, **328**
 Lead, 12
 Leaf, 348, 356-364
 anatomy, 426-429
 attachment, 361
 gametophytic, 59, 63
 gap, 82, 427
 parts, 356-357
 sporophytic, 75
 structural types, 357-359
 type, 359-361
 venation, 155, 361-364
 Lecythidaceae, 292
 Ledocarpaceae, 255
 Leguminosae, *see* Fabaceae
Leiothrix, 212
Lemna, **163**
 Lemnaceae, *see* Araceae
 Lennoaceae, 292
Lens, 262
 Lentibulariaceae, 292, 308
 Lepidobotryaceae, 255
Lepidodendron, **78, 82, 97**
Lepidozamia, **109**
Leptocarpus, **214**
 Leptoid, 63
Leptosiphon, **299**
Leptospermum, **258, 259**
 Leptosporangiate ferns, *see* Polypodiales
 Leptosporangium, 88
Leucocoryne, 169
Leucojum, 169
Leucopogon, **296**
Leucospermum, **237**
Licuala, **186**
 Lignin, 69
 Lignophytes, *see* Woody plants
 Ligule
 grass, 216
 leaf, 357
 lycophyte, 79
Lilaea, **161**
 Liliaceae, 180, **182**
 Liliales, 180-181
Lilium, 180, **182**
 Limeaceae, 230
 Limnanthaceae, 255
 Limnocharitaceae, 159
Limonium, 238
 Linaceae, 255, 269
Linanthus, **299**
Linaria, **318**
 Linderniaceae, 292, 310
 Lineage, 10, 17
 Linnaeaceae, 292, 331
Liriodendron, 149
Litchi, 289, **290**
Lithophragma, **253**
Lithops, **242**
 Liverworts, 59
Livistona, **186**
 Loasaceae, 292

- Lobeliaceae, *see* Campanulaceae
 Locule, 376
 Loganiaceae, 292
 Loganiaceae, *see* Scrophulariaceae s.s.
Lolium, **218**
Lomatium, **324**
 Long branch attraction, 35-36
Lonicera, **333**
Lophophora, 244
 Lophopyxidaceae, 255
 Loranthaceae, 230
Loseliastrum, **299**
 Lowiaceae, 185
Ludisia, **175**
Ludwigia, **260**
Luffa, 262
Lupinus, **264**
 Luzuriagaceae, 159
Lychnis, **247**
Lycopersicon, 319
Lycophyll, 78, 427
Lycopodiaceae, 79
Lycopodiella, 79
Lycopodium, **79, 80**
 Lycopodophyta/lycophytes, 78-82
Lycoris, 169
Lygodium, **89**
 Lythraceae, 254-255
- Macadamia*, 238
 Mackinlayaceae, 292
Maclura, 275, **276**
Macropidia, **192**
Macropiper, **152**
Macrozamia, **109**
 Maesaceae, 292
 Magnoliaceae/*Magnolia*, 149, **150**
 Magnoliales, 146
 Magnoliids, 146
Mahonia, **231**
 Majority/50% majority consensus tree,
see Consensus tree
Malacothrix, **328**
 Male gametophyte, 101
 Male sporophyll, *see* Microsporophyll
Malosma, **287**
 Malpighiaceae, 255, 269
 Malpighiales, 269
Malus, 278, **279**
 Malvaceae, 280, 282, **283-284**, 286
 Malvales, 280
Mammillaria, **245**
Mangifera, 286, **287**
Manihot, 269, **270**
Marah, **261**
 Marantaceae, 185, **202**
Marattia, **87**
 Marattiales/marattioid ferns, 87
 Marcgraviaceae, 292
Marchantia, **61**
 Margin, 393-394
Markhamia, **311**
Marrubium, **312**
Marsilea, 88, **92**
- Martyniaceae, 292
Matteuccia, 88
 Maturation, 402
 Maximum likelihood, 36
 Mayacaceae, 185
Medeola, **182**
Medicago, 262
Medinilla, **257**
Medullosa, 107, **108**
 Medusandraceae, 230
 Megagametogenesis, 441
 Megasporangium/megaspore, 79, 99,
 114, 439
 Megaspore mother cell, *see* Megasporocyte
 Megaspore, 79, 81-82, 99
 Megasporocyte, 97, 104, 440
 Megasporogenesis, 441
 Megasporophyll, 82, 108
 Melanophyllaceae, 292
 Melanthiaceae, 159, 180-181
Melaspheerula, **173**
 Melastomataceae, 254, **257**
 Meliaceae, 255
 Melianthaceae, 255
 Menispermaceae, 230
Mentha, 310
 Menyanthaceae, 292
 Meristem, *see* Apical m., Lateral m.
Mertensia, **300**
 Mesarch, 83, 421
 Mesophyll, 75, 427-429
 Metaxylem, 78, 421
Metroxylon, 188
 Metteniusaceae, 230
Michelia, 149, **150**
Microcyas, 108
 Microfibrils, 51
 Microgametogenesis, 439, 442
 Microgametophyte, *see* Male gametophyte
 Micropyle, 101, 130, 439
 type, 441, 446
 Microsatellite DNA, 487, 489
 Microsporangium, 79, 114
 Microspore, 79, 81-82, 99, 126
 Microsporogenesis, 439, 442
 Microsporophyll, 82, 108
 Middle lamella, 410, 411
Mimulus, 315, **317**
Mirabilis, **239**
 Misodendraceae, 230
Mitella, **253**
 Mitrastemonaceae, 292
Mnium, **65**
 Molecular systematics, 477-491
 Molluginaceae, 230
Monarda, **312**
Monardella, **313**
 Monilophytes, 82
 Monimiaceae, 140
Monochoria, 192
 Monocotyledons, 153-219
 apomorphies, 155-158
 classification, 156, 159
 Monoecy, 114, 365
- Monograph, 533
 Monophyly/monophyletic/monophyletic
 group, 13, 26
 Monosporic, 441, 444
 Monosulcate, 228
Monotropa, **296**
Monstera, 160, **162**
 Montiniaceae, 292
 Montropaceae, *see* Ericaceae
 Moraceae, 275-276
Moraea, **173**
 Morinaceae, 292, 331
 Moringaceae, 255
 Morphocline, *see* Transformation series
 Morphology, 347-407
Morus, 275, **276**
 Mosses, 63-66
Muelenbeckia, 249
Musa, 192, **195, 197**
 Musaceae, 192, **195, 197**
 Musci, *see* Mosses
Musella, 192
 Mutingiaceae, 255
 Myodocarpaceae, 292
 Myoporaceae, *see* Scrophulariaceae s.s.
Myoporum, **314**
Myosotis, 301
Myosurus, **234**
 Myricaceae, 255, 265
 Myristicaceae, 140
 Myrothamnaceae, 230
 Myrsinaceae, 292
 Myrtaceae/*Myrtus*, 256, **258**, 259
 Myrtales, 254
- Najas*, **161**
 Name
 botanical, 510-513
 changes, 505-507
 common, 502
 conservation of, 505
 correct, 507
 illegitimate, 501
 legitimate, 501
 pronunciation, 511
 remodeling, 506
 scientific, 502-503
 validly published, 501, 507, 508
Nandina, **231**
Narcissus, 169-170
 Nartheciaceae, 159
Nasella, **218**
 Natural selection, 10
 Neck/neck canal cells, 58
 Nectary, 374, 417, 466
 Nelumbonaceae/*Nelumbo*, 233, **235**
 Neotony, 43
 Nepenthaceae, 230, 238, **239**
Nepenthes, **239**
 Nephrolepis, 88, **89-90**
Nerium, **303**
 Network, *see* Unrooted tree
 Neuradaceae, 255
Neuropteris, 107, **108**

- Nexine, 461
Nicotiana, 319, **321**
Nitella, **53**
 Nitrariaceae, 255
 Node
 cladogram, 18
 leaf, 75
 Nomenclatural type, *see* Type (nomenclatural)
 Nomenclature, 12, 501-515
 independence of, 510
 principles, 502
 Nothofagaceae, 255, 265
 Nucellus, *see also* Megasporangium type, 440
 Nucleotides, 477-478
 Number
 in botanical names, 510
 in morphology, 389
Nuphar, 143
 Nyctaginaceae, 230, 238, **239**
 Nyctinasty, 357
 Nymphaeaceae/*Nymphaea*, 143-**144**
 Nymphaeales, 143
Nypa, 185, 188
 Nyssaceae, *see* Cornaceae
- Obolaria*, **306**
 Ochnaceae, 255
Ochroma, 286
Ocimum, 310
Oenothera, 259, **260**
 Oil bodies, 59
 Oil cells, 142
 Olacaceae, 230
 Oleaceae, 292, 308
 Oliniaceae, 255
 Onagraceae, 259, **260**
Oncidium, **174**
 Oncothecaceae, 292
 Ontogeny/ontogenetic, 41-43
 Oogamy, 53
 Oomycota, 4-5
 Operculum, *see* Calyptra
 Ophioglossales/ophioglossoid ferns, 83
Ophioglossum, 83, **84**
 Opiliaceae, 230
 Optimization, *see* Character optimization
Opuntia, 244, **245**, **246**
 Orchidaceae, 171, **174-175**, 176-177
Orchis, **175**
 Ordered, *see* Character, ordered
 Orectanthe, 212
 Organic acids, *see* UV-fluorescent acids
 Orientation, 397-398
 Orobanchaceae, 310, 315-**316**
Orobanche, **316**
Orthosiphon, **312**
 Orthotropous, 446
Oryza, **215**, 216, **218**
Osmunda, **89**
 Ostrya, 265
 OTU (Operational Taxonomic Unit), 18-19
- Outbreeding/outcrossing, 468
 Outgroup
 comparison, 31-34
 definition, 19
 Ovary
 attachment, 376, 377
 position, 377, 378
 Ovule, 98
 development, 439
 parts, 439
 position, 445-446, 447
 type, 442-443, 445-446
 Ovuliferous scale, 114
 Oxalidaceae/*Oxalis*, 272, **274**
 Oxalidales, **272**
- Pachira*, 286
Pachycereus, **246**
Pachyphytum, **251**
Pachypodium, **303**
Pachytesta, 107, **108**
 Paedomorphosis, 43
 Paeoniaceae, 230, 250
Paepalanthus, 212
Palafoxia, **329**, **330**
 Palmae, *see* Arecaceae
 Palynology, 453-463
Panax, 322
 Pandaceae, 255
 Pandanaceae/*Pandanus*, 180, **183-184**
 Pandanales, 180
Papaver, **232**
 Papaveraceae, 230, **232-233**
Paphiopedilum, **175**
 Paracryphiaceae, 292
 Parallelism, 21
 Paraphyletic group/paraphyly, 13, 26
 Parenchyma, 53-54, 57
 axial, 423
 Parnassiaceae, 255
 Parsimony analysis, 27-29
 Parthenogenesis, 472
 Passifloraceae/*Passiflora*, **271-272**
Paullinia, 289
 Paulowniaceae, 292
Pausinystalia, 305
 Peat moss, *see* *Sphagnum*
 Peat, 66
 Pedaliaceae, 292, 308
Pedicularis, **316**
Pelargonium, 254, **256**
 Penaeaceae, 255
 Pennantiaceae, 292
 Pentadiplandraceae, 255
 Pentaphragmataceae, 292
 Pentaphylacaceae, 292
Pentas, 305, **307**
 Penthoraceae, 230
 PEP carboxylase, 429-432
Peperomia, **152-153**
 Peramorphosis, 43
Pereskia, **245**, **246**
 Perforation plate, 72
- Perianth, 366-371
 aestivation, 370-371
 arrangement, 367
 cyclic, 367-368
 definition, 121
 fusion, 369
 merosity, 368-369
 parts, 366-367
 type, 121, 123, 369-370
 Perianth/Androecial position, 377, 378
 Pericarp, 129
 Pericycle, 77
 Periderm, *see* Cork
 Peridiscaceae, 230
 Perine layer, 63
 Periodicity, 403
 Peristome teeth, 63
 Persea, 146-**147**
 Petals, *see* Corolla
 Petermanniaceae, 159
 Petrosaviales/Petrosaviaceae, 159
Petroselinum, 322
Phacelia, **302**
Phalaris, **218**
Phaseolus, 262
 Phellinaceae, 292
Phenakospermum, 195
 Phenetic, *see* Classification
 Phenogram, 40
 Philesiaceae, 159, 180-**181**
Philodendron, 160, **163**
Philothea, **288**
 Philydraceae, 185
Phlebocarya, **191**
 Phloem, 72, 415
 secondary, 97, 422-425
Phlox, 298
Phoenix, **186**, **187**, 188
Pholistoma, **302**
Phorodendron, 249, **250**
Photinia, 278
 Photosynthesis, 4, 5-6, 429-432
 C4, 429, 431
 CAM, 429, 432
Phryma, 315, **317**
 Phrymaceae, 310, 315, **317**
 Phyllanthaceae, 255
Phylloglossum, 79, **80**
 Phyllonmaceae, 292
Phyllospadix, **161**
 Phylogenetic systematics
 definition, 13, 17
 perspective on, 43
 Phylogenetic tree, *see* Cladogram
 Phylogeny, 9, 13, 17
Physalis, 319, **321**
 Phytolaccaceae, 230
 Physiology, 409, 429-432
Phytolacca, **239**
 Phytolaccaceae, 230, 238, **239**
 Picramniaceae, 255
 Picrodendraceae, 255
Pillansia, **173**
Pilularia, 88, **92**

- Pimenta*, 259
Pinaceae, 114, **115-116**
Pinna/pinnae/pinnules, 88
Pinus, **113, 115**
Piper, **152-153**
Piperaceae, 149, **152-153**
Piperales, 149
Pistacia, 286
Pistia, **163**
Pistil, 121
 compound, 129
 simple, 129
Pistil, 374-379
Pisum, 262
Pit, 69, 411, 412
Pith, 77, 420
Pithecellobium, **263**
Pittosporaceae, 292
Placentation, 377-378
Plagiobothrys, **300**
Plant press, 518-519
Plant
 definition, 3
 habit, 350
 habitat, 350
 importance, 5-6
 organs, 348-350
 sciences, 9
 sex, 365
 structure, 348
 uses, 6-9
Plantaginaceae, 310, 315, 317, **318, 319**
Plantago, 317, **318**
Plasmodesmata, 53, 411
Plastids, sieve tube, *see* Sieve tube plastids
Platanaceae/*Platanus*, 235-**236**
Platyserium, **89**
Platystemon, **232**
Plectranthus [*Coleus*], **76**
Plectritis, **334**
Plesiomorphic, *see* Ancestral
Plocospermataceae, 292
Plumbaginaceae, 230, 238, **239**
Plumbago, **239**
Plumeria, 301, **303**
Poaceae (Gramineae), 213, **215-218, 219**
Poales, 203
Podocarpaceae, 114, **116**
Podocarpus, **116**
Podophyllum, 230, **231**
Podostemaceae, 255
Pogogyne, **313**
Pogostemon, 310
Polar nuclei, 130, 441, 445
Polarity, 23
Polemoniaceae, 298, **299**
Polemonium, 298
Pollen tube, 101, 126
Pollen, 101
 aperture, 455-458
 development, 439
 nucelus number, 453
 polarity, 455
 reduced, 126
Pollen Cont d
 sculpturing, 458-460
 shape, 458-459
 size, 458
 storage product, 453
 symmetry, 458
 unit, 453-455
 wall structure, 460-461
Pollenkit, 462
Pollination, 465
 chamber, 103
 droplet, 101, 103
 mechanisms, 466-468
Polyalthea, 149
Polyclave key, *see* Key
Polygalaceae/*Polygala*, 265-**266**
Polygonaceae/*Polygonum*, **248-249**
Polymerase chain reaction, 477-479
Polyosmaceae, 292
Polyphyletic group, 13, 26
Polyploidy, 473-474
Polypodiales, 87-92
Polypodium, **83, 89-90**
Polysporangiophytes, 66, 74
Polytomy, 25, 30-31
Polytrichum, **65**
Pontederia, 192, **193, 194**
Pontederiaceae, 190, **193-194**
Population, 10
Pore, liverwort, 59
Pore, wood, *see* Vessel
Porella, **61**
Portulacaceae, 230, 238, **239**
Posidoniaceae, 159-160
Position
 morphological, 396
 nomenclatural, 504
Posture, 398-399
Potamogeton, **161**
Potamogetonaceae, 159-**161**
Potentilla, **279**
Pothos, **163**
Primary cell wall, *see* Cell wall, primary
Primary pit field, 69, 411
Primer, 479
Primitive, *see* Ancestral
Primulaceae, 292-293
Principle of parsimony, *see* Parsimony
 analysis
Prioniaceae, 185
Priority of Publication, 505
Pronunciation, *see* Names, pronunciation
Protandry, 402, 469
Protea, 238
Proteaceae, 235, **237-238**
Proteales, 233
Protogyny, 402, 469
Protonema, 63
Protostele, 74, 419-421
Protoxylem, 78, 83, 421
Prunella, **313**
Prunus, 278, **279**
Psathyrotes, **329**
Pseudobombax, **283**
Pseudocrassinucellate, 440
Pseudo-elaters, 63
Pseudotsuga, 114, **115**
Psidium, 259
Psilotales, 84-85
Psilotum, 84-**85**
Psiloxylaceae, 255
Psychotria, **307**
Pteridosperm, *see* Seed fern
Pteris, 88
Pterostemonaceae, 230
Putranjivaceae, 255
Puya, **204**
Pyracantha, 278
Pyrola, **296**
Pyrolaceae, *see* Ericaceae
Pyrostegia, **311**
Pyrularia, **250**
Pyrus, 278, **279**

Quercus, **268, 269**
Quiabentia, **245**
Quillaja, **279**
Quillajaceae, 255

Rafflesiaceae, 255
Rafinesquia, **328**
Rank, 12, 503
Ranunculaceae/*Ranunculus*, 233-**234**
Ranunculales, 228
Rapateaceae, 185
RAPD (Random Amplified Polymorphic
 DNA), 487, 489
Raphanus, **281**
Rauwolfia, 305
Ravenala, 195
Recency of common ancestry, 26
Red algae, 4-5
Regnellidium, 88
Reproductive biology, 465-476
Rescaled consistency index (RC), 37
Resedaceae, 255
Restionaceae, 212-**214**
Restriction site analysis, 483-487
Retention index (RI), 37
Reticulation, 31
Reussia, 192
Reversal, 21
Reward, 466
RFLP, *see* Restriction site analysis
Rhabdodendraceae, 230
Rhamnaceae, 275, **277**
Rhamnus, **277**
Rheum, 249
Rhexia, **257**
Rhipsalis, 244
Rhizoid, 59, 76, 88, 348, 417
Rhizome, 88
Rhizophoraceae, 255, 269
Rhododendron, 294, 295, **296**
Rhoeo, 190
Rhopalostylis, **187**
Rhus, 286, **287**

- Rhynchoalycaceae, 255
Rhynia/Rhyniophyta/rhyniophytes, **78**
Ricinus, 269
 Ripogonaceae, 159
Romneya, **232**
 Root hairs, 77
 Root, 76, 348, 350, 417-418
 cap, 77, 417-418
 hairs, 417-418
 lateral, 417
 type, 350-351
 Roridulaceae, 292
Rorippa, **281**
Rosa, **278, 279**
 Rosaceae, 275, **278-279**
 Rosales, 275
 Rosids, **252, 254, 255**
Rosmarinus, 310, **312**
 Rousseaceae, 292
Rubia, 305
 Rubiaceae, 305, **307**
Rubus, 278
Ruellia, **309**
Rumex, **248**
 Ruppiaceae, 159-160
 Ruscaceae, 159
Ruta, **288, 289**
 Rutaceae, 286, **288-289**
- Sabatia*, 305, **306**
 Sabiaceae, 230
Sagittaria, **164**
 Salicaceae, 255, 269
Salicornia, **243**
Salsola, **243**
 Salvadoraceae, 255
Salvia, 310, **312, 313**
Salvinia, 88, **92**
Sambucus, **333**
Sanguinaria, **232**
Sanicula, **324**
 Santalaceae, 249-**250**
 Santalales, 249
Santalum, 249
 Sapindaceae/*Sapindus*, 289, **290, 291**
 Sapindales, 286
 Sapotaceae, 292-293
 Sarcobataceae, 230
 Sarcolaenaceae, 255
Sarcostemma, **304**
 Sarraceniaceae, 292-293
 Sassafras, **147**
 Saururaceae/*Saururus*, 153-**154**
 Saxifragaceae/*Saxifraga*, 252, **253**
 Saxifragales, 250, 253
Scaevola, **332**
 Scaled (character), *see* Character, scaled
Sceletium, 241
Schefflera, 322, **325**
 Scheuchzeriaceae, 159
Schiekia, **192**
Schinus, 286, **287**
 Schisandraceae, 140
 Schlegeliaceae, 292
- Schoepfiaceae, 230
 Scientific name, 12
Scirpus, **209, 210**
 Sclereid, 70-71, 414, 415
 Sclerenchyma, 70, 413, 414
Scrophularia, **314**
 Scrophulariaceae s.l., 310
 Scrophulariaceae s.s., 310, **314-315**
Scutellaria, **313**
Secale, 216
 Secondary cell wall, *see* Cell wall,
 secondary
 Secondary growth, 421-426
 Secondary phloem, *see* Phloem, secondary
 Secondary xylem, *see* Wood
 Secretary (tapetum), 439
 Secretary structures, 416-417
Sedum, **251, 252**
 Seed fern, 107, **108**
 Seed plants, 98-105
 Seed, 98, 348, 387-389
 adaptations, 105
 coat, 98, 448-449
 development, 104-105, 446-449
 dispersal, 472
 endosperm type, 388-389
 evolution, 98-101
 germination type, 389
 morphology, 449
 storage tissue composition, 448
 storage tissue origin, 448
 Seismonasty, 357
 Selaginellaceae/*Selaginella*, 79, **81-82**
Selenicereus, **245**
 Self-incompatibility, 471
 Selfing, *see* Inbreeding
Senecio, **328**
Senna, **263**
 Sepals, *see* Calyx
Sequoia, **116**
Sequoiadendron, **116**
 Setchellanthaceae, 255
 Sexine, 461
 Sexual reproduction, 465-472
 timing, 471
 Shape, 390-396
 combinations, 393, 394
 plane (two-dimensional), 390-392
 solid (three-dimensional), 390
 Shoot, 75, 348, 351, 419
 Short shoot, 110, 353
 Sieve tube plastids, 155
 of Caryophyllales, 238
 of monocots, 155, 157
 Sieve
 area, 72-73, 415-416
 cell, 73, 415-416
 element, 72, 415-416
 plate, 73, 132, 415-416
 tube member, 73, 132, 415-416
Sigillaria, **78**
Silene, **247**
 Simaroubaceae, 255
Simmondsia, **239**
- Simmondsiaceae, 230, 238, **239**
 Siparunaceae, 140
 Siphonogamy, 101, 111, 126
 Siphonostele, 83, 420-421
 Sister group/taxa, 26
Sisyrinchium, **173**
 Size, 389
 Sladeniaceae, 292
 Smilacaceae, 159, 180-**181**
Smilax, **181**
 Solanaceae, 319, **321**
 Solanales, 319
Solandra, **321**
Solanum, 319, **321**
Solidago, **328**
Sonchus, **327, 330**
Sorghum, **218**
 Sorus/sori, 88
 Sparganiaceae/*Sparganium*, 206-**207**
Spathiphyllum, 160
Spathodea, 308, **310, 311**
 Speciation, 10
 Species, 10
 Specimen, 518, 525
 mounting, 526-528
 storage, 529
Spergula, **247**
Spergularia, **247**
 Sperm, 53
 angiosperm, 126
 motile, 108-109
 Spermatophytes, *see* Seed plants
 Sphaerosepalaceae, 255
Sphagnum, 63, **65, 66**
 Sphenocleaceae, 292
 Sphenophytes/sphenopsids, *see* Equisetales
 Sphenostemonaceae, 292
Spinacea, 244
Spiraea, 278, **279**
Spirogyra, **53**
 Sporangiphore, 85
 Sporangium
 definition, 54
 female, *see* Megasporangium
 liverwort, 59
 male, *see* Microsporangium
 wall, 55
 Spore, 53
 Sporocarp, 91
 Sporocyte, 55
 Sporogenous tissue, 55
 Sporophyll, 79, 108
 Sporophyte, 54, 74
 Sporopollenin, 460
Stachys, **313**
 Stachyuraceae, 255
 Stamen, 124-126
 arrangement, 371-372
 attachment, 372
 cyclic, 372
 fusion, 373
 insertion, 372-373
 position, 372
 type, 371

- Stangeria*, **109**
Stanhopea, **174**
Stapelia, **301, 304**
 Staphyleaceae, 255
 Starch, 51, 53
 grain, 410-411
 Stegnospermataceae, 230
 Stele, 74, 419-421
Stellaria, **247**
 Stem, 74, 348, 351-356, 419-421
 branching pattern, 354-355
 habit, 354
 types, 352-354
 Stem-based (classification), *see*
 Classification, stem-based
 Stemonaceae, 159
 Stemonuraceae, 292
Stenocereus, **246**
Stenomeris, 180
 Sterile jacket layer, *see* Antheridial wall,
 Archegonial wall
 Sterile segment, 83
 Stigma/Stigmatic region, 129, 379
 Stilbaceae, 292, 310
 Stipe
 fern, 88
 moss, 63
 Stomate/stoma, 59, 62, 427, 428
 Stramenopiles, 4-5
 Strasburgeriaceae, 255
 Strelitziaceae/*Strelitzia*, **195-196,**
 198, 199
 Streptophyceae/streptophytes, 53
 Strict consensus tree, *see* Consensus tree,
 strict
 Strobilus, *see* Cone
Strophanthus, **303**
Strophostyles, **264**
 Style, 129
 position, 378-379
 structural type, 378-379
 Stylidiaceae, 292
 Styraceae, 292
Suaeda, **243**
 Suberin
 in bark, 97
 in endodermis, 73-74
 in stomates, 62
 Surface, 399-402
 Surianaceae, 255
Swertia, **306**
Syagrus, **186, 187**
 Syllables, 511
 Symmetry, 402
Symphoricarpos, **333**
 Sympleiomorphy, 40
 Symplocaceae, 292
Symplocarpus, **162**
 Synangium, 84, 87
 Synapomorphy, 18, 24
 Synergid cells, 130
Syngonanthus, 212
Syngonium, 160
 Synonym, 507
 Systematics
 definition, 9
 molecular, *see* Molecular systematics
 rationale and uses, 13-15
Syzygium, 259

Tacca, **179, 180**
 Tamaricaceae, 230, 238, **239**
Tamarix, **239**
 Tapetum, 437-438
 Tapisciaceae, 255
 Taxaceae, 114, **116**
Taxodium, **116**
 Taxon
 definition, 12-13
 selection, 18, 31
 Taxonomic key, *see* Key
 Taxonomy, 10-13
Taxus, **116**
Tecomaria, **310**
 Tecophilaeaceae, 159
 Tectate-columellate, 460
 Temporal phenomena, 402
 Tenuinucellate, 440, 443
 Tepal, 122
 Tetracarpaeaceae, 230
 Tetrachondraceae, 292
Tetragonia, 241
 Tetramelaceae, 255
 Tetrameristaceae, 292
Tetrapanax, 322
 Tetrasporic, 441, 444
Tetrazygia, **257**
 Texture, 389
Thalia, **202**
Thalictrum, **234**
 Thallus/thalloid, 59, 348
 Theaceae, 292-293
 Theca, 124, 437, 438
Thecium, **250**
Thelymitra, **175**
 Themidaceae, 177-**178**
Theobroma, **283**
 Theophrastaceae, 292
Thottea, 149
Thunbergia, **309**
 Thurniaceae, 185
 Thylakoids, 4, 51
 Thymelaeaceae, 255, 280
Thymus, 310
Thysanocarpus, **281**
Thysanotus, **166**
Tiarella, **253**
Tibouchina, 254, **257**
 Ticodendraceae, 255
Tilia, **283, 285, 286**
Tillandsia, **204**
 Tissue, 71, 412-417
 ground, 413
 vascular, 413, 415-416
Tithonia, **328**
Tmesipteris, 84
 Tofieldiaceae, 159-160

 Torricelliaceae, 292
 Tovariaceae, 255
Toxicodendron, 286, **287**
 Tracheary element, 72, 413, 415
 Tracheid, 72, 413, 414
 Tracheophyta/tracheophyte, *see*
 Vascular plant
Tradescantia, **189, 190**
 Transformation series, 22
 Tribelaceae, 292
Tribonanthes, 190, **191, 192**
 Trichome, 416
 type, 401-402
Trichopus, 180
 Tricolpate, 228
Trifolium, 262
 Trigger mechanism, 470
Triglochin, **161**
 Trilete mark, 63
Trillium, **181**
 Trimeniaceae, 140
 Trinomial, 504
Tristania, **258**
Triticum, **215, 216**
Tritoniopsis, **173**
 Triuridaceae, 159
Trixis, **327**
 Trochodendraceae, 230
 Trochodendrales, 230
 Tropaeolaceae, 255, 280
 Trunk, 355-356
 Tube cell, 126
Tulbaghia, **169**
Tulipa, 180, **182**
 Twig, 355-356
 Type (nomenclatural), 504-505, 517
 Typhaceae/*Typha*, **205-206**

 Ulmaceae, 255, 275
Ulva, **53**
 Umbelliferae, *see* Apiaceae
 Unordered, *see* Character, unordered
 Unrooted tree, 29-30
Uropappus, **330**
Ursinia, **328**
 Urticaceae, 255, 275
 UV-fluorescent acids, 184, 185

Vaccinium, 294, 295, **296**
 Vahliaceae, 292
 Valerianaceae, 331, **334-335**
 Valid publication, *see* Name, validly
 published
Vanilla, **174, 177**
 Vascular bundle, 75, 420
 Vascular cambium, 97, 422
 Vascular plants, 69-95
 Vegetation type, 523
 Vein, 75
 Velloziaceae, 159
 Venation, *see* Leaf, venation
 Venter/ventral canal cell, 58
Verbascum, **314**

- Verbenaceae, 292, 308
Veronica, **318**
Verticordia, **258**
 Vessel/vessel member, 72
 angiosperm, 133
 Gnetales, 114, 117
 Vestiture, 400-401
 Vicariance, 41
Victoria, 143-**144**
Viguiera, **327, 330**
Vinca, 301
 Violaceae/*Viola*, 272-**273**
 Viscin threads, 461-462
 Vitaceae, 255
 Vitales, 255
 Vivianiaceae, 255
 Vochysiaceae, 255
Volvox, **53**
 Voucher, 517

Wachendorfia, 190, **191**
Washingtonia, **186**
 Weighting (character), 22-23

 Welwitschiaceae/*Welwitschia*,
 114, **118**
Wigandia, **302**
 Winteraceae, 140
Wisteria, **264**
Wolffia, 160, **163**
 Wood, 97, 422-426
 cuts, 423, 426
 ray, 423-426
 spring, 423-425
 summer, 423-425
 Woody plants, 97-98

Xanthium, **330**
Xanthorhiza, **234**
Xanthorrhoea, **166**
 Xanthorrhoeaceae, 159, 165-**166**
Xanthosia, **324**
Xanthosoma, 160, **162**
 Xeronemataceae, 159
Xiphidium, **191**
 Xylem, 72, 413
 secondary, *see* Wood

Xylococcus, **295, 296**
 Xyridaceae, 212-**213**
Xyris, 212-**213**

Yucca, 165, **168**

Zamia, **111**
Zantedeschia, 160, **162**
Zea, **215**, 216
Zebrina, 190
Zigadenus, **181**
Zingiber, 197-198
 Zingiberaceae, 197-198, **200**
 Zingiberales, 192, 194-203
Zizyphus, 275, **277**
Zombia, **187**
 Zosteraceae, 159-**161**
Zygocactus, **246**
Zygopetalum, **174**
 Zygophyllaceae, 255
 Zygophyllales, 255
 Zygote, 53, 131