



On growth habits and forms: the utility of stem anatomy to define growth habits of Melastomataceae

Sobre hábitos de crecimiento y formas: el uso de la anatomía del tallo para definir hábitos de crecimiento de Melastomataceae

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ABSTRACT:

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Background and Aims: The growth habit is a character commonly used in taxonomic descriptions of plants and one of the first to be taught to students of plant morphology. Defining growth habits has turned into a complex matter due to the proliferation of terms. By definition, some habits can be distinguished by plant stem anatomy, but there are actually few studies that evaluate this correspondence. The family Melastomataceae is an excellent group to investigate how stem anatomy varies among plants with contrasting growth habits because it shows considerable variation in growth habits and stem anatomy. In this study, we analyzed whether stem anatomy characteristics vary consistently across growth habits in species of Melastomataceae.

Methods: Eighteen species of Melastomataceae were collected in different Colombian ecosystems. Observation of 125 stem anatomy characters was performed on fixed internode slices, which were cut by freehand in transverse, radial longitudinal, and tangential planes. We used as a prior a classification of growth habits that we consolidated from a literature review, aiming to reconcile multiple schemes of classification of habits across vascular plants.

Key results: Few anatomical characteristics were shared exclusively by species of one particular growth habit. Therefore, the Melastomataceae species analyzed here could be grouped in three general growth habits: trees/shrubs, herbaceous, and climbers.

Conclusions: Many of the current definitions of growth habits are a mixture of different topics including ecology, specimen size, and the presence of specialized organs. For the concept of growth habits to be useful, it should be avoided to elevate ranges of size, habitats, or specific functional morphology adaptations to the generalized category of growth habits.

Key words: anatomy, climber, growth habit, herb, Melastomataceae, shrub, tree.

RESUMEN:

Antecedentes y Objetivos: El hábito de crecimiento es un carácter comúnmente usado en descripciones taxonómicas de plantas y uno de los primeros enseñados a estudiantes de morfología vegetal. La definición de hábitos de crecimiento se ha convertido en un aspecto complejo debido a la proliferación de términos. Por definición, algunos hábitos pueden distinguirse por medio de anatomía caulinar, pero hay pocos estudios que hayan evaluado esta correspondencia. La familia Melastomataceae es un excelente grupo para investigar cómo la anatomía del tallo varía entre plantas con hábitos de crecimiento contrastantes, ya que presenta una variación considerable en hábitos de crecimiento y anatomía caulinar. En este estudio analizamos si las características de la anatomía caulinar varían de manera consistente en especies de Melastomataceae de distintos hábitos de crecimiento.

Métodos: Recolectamos 18 especies de la familia Melastomataceae en diferentes ecosistemas colombianos. Se registraron observaciones de 125 caracteres de anatomía caulinar a partir de cortes histológicos del entrenudo, fijados, los cuales fueron cortados a mano alzada de los planos transversal, longitudinal y tangencial radial. Empleamos una clasificación a priori de hábitos de crecimiento, la cual se consolidó a partir de una revisión de literatura, tratando de reconciliar diferentes esquemas de clasificación de hábitos utilizados en plantas vasculares.

Resultados clave: Hubo muy pocas características compartidas exclusivamente por especies de un hábito de crecimiento particular. Por lo tanto, las especies de Melastomataceae analizadas podrían clasificarse en tres hábitos de crecimiento: árboles-arbustos, herbáceas y trepadoras.

Conclusiones: Muchas de las definiciones actuales sobre los hábitos de crecimiento son una mezcla de diferentes temas, incluyendo la ecología, el tamaño del espécimen y la presencia de órganos especializados. Para que este concepto sea útil, deben evitarse incluir rangos de tamaño, hábitats o adaptaciones morfológicas funcionales específicas en la categoría generalizada de hábitos de crecimiento.

Palabras clave: anatomía, árbol, arbusto, hábitos de crecimiento, hierba, Melastomataceae, trepadora.

INTRODUCTION

Growth habit is usually the first characteristic used to describe a plant. Whether a plant is a tree, a herb or a liana seems like a simple question, and yet, many times the answer is not straightforward because there are multiple, sometimes contradictory, or poorly delimited definitions of growth habits. The term habit comes from the Latin word *habitus*, which means external appearance (Font Quer, 2001); in botany, the habit refers to the external aspect of a plant (Vázquez-Sánchez et al., 2012). Defining growth habits can be quite challenging, even for a single species. For example, 80 randomly selected records of *Blakea quadrangularis* Triana (Melastomataceae) deposited in the Herbarium of Universidad de Antioquia (HUA, Colombia), were registered as trees (44%), shrubs (39%), vines (12%), lianas (3%), hemiepiphytes (1%), and herbs (1%). For growth habits to be a more useful concept in taxonomy and functional analysis, it is necessary to clarify their definition.

Defining growth habits has turned into a complex matter due to multiple factors. One source of confusion is the proliferation of terms. For example, a climbing plant is commonly described as a liana, vine, twiner, creeper, sprawler, or scrambler (Putz, 1984; Mendoza and Ramírez, 2006). However, these terms are not mutually exclusive and are related to different features: how they grow, climb, or hold to the substrate. Another issue is that growth habits have been confused with habitats where plants grow. For example, the following terms have been considered as growth habits: epiphytic, when plants grow on another plant without subtracting water or nutrient from it (Font-Quer, 2001); parasitic, when plants grow on another plant absorbing its resources; and aquatic, when plants live in the water. However, these terms may not be referring to growth habits because they do not allude directly to plant appearance, but instead to their habitat (Simpson, 2006).

Stem anatomy can be useful to delimit growth habits. Plants with different growth habits vary in their stem stiffness, which is related to the lignification of

their stem (Isnard et al., 2012). By definition, some habits can be distinguished by the plant stem anatomy, i.e. herbs are plants that lack woody tissue at any level of their aerial stems (Eiten, 1991). Plants with the lianous growth habit can show anomalous cambial growth, vessels dimorphism, and abundant unlignified parenchyma (Obaton, 1960; Carlquist, 1985; Angyalossy et al., 2011; 2014). Trees and shrubs self-supporting constraints lead to the development of xylem with abundant fibers and longer axial elements (Rowe and Speck, 2004). Despite the evident relationship between stem anatomy and growth habits, there are few studies on how stem anatomy can distinguish growth habits. The morphological observations that define growth habits can greatly benefit from a detailed study of anatomy that helps to better characterize these habits.

The family Melastomataceae is an excellent group to investigate how the stem anatomy varies among plants with contrasting growth habits. It includes nearly 5000 species in 170 genera, standing as one of the largest families of Angiosperms (Penneys and Judd, 2011). Melastomataceae species exhibit a wide variety of growth habits. However, these habits are poorly defined due to various issues, one of them the use of different terminology in growth habit classifications. For example, Vliet (1981) suggested only two anatomically recognizable growth habits (erect stem plants, and climbers). Renner (1993) recognized six habits (trees, treelets (small stature trees), shrubs, herbs, root climbers, and facultative epiphytes). Almeda (2001) reduced to five the growth habits recognized within the family (trees, shrubs, herbs, scandent climbers, and epiphytes). Later, Mendoza and Ramírez (2006) increased to eight the growth habits observed in the family (tree, shrub, suffrutex (=a tall herb), herb, liana, vine (=a liana with a non-woody stem), epiphyte (=a plant with roots that does not actively extract water or nutrients from the ground or from the live tissues of the host), and hemiepiphyte (=epiphyte with roots reaching the ground later in life)). A unified classifi-



cation of growth habits can clarify the discrepancies among these different groupings.

The stem anatomy variation of Melastomataceae is considerable. Most of the studies on the stem anatomy of members of the family have been realized on trees and shrubs, the most common growth habits in the family. In Melastomataceae, the length of xylem axial cells tends to be longer in trees than in shrubs (Ter Welle and Koek-Noorman, 1981). In some lianas of the family, the proportion of non-lignified xylem parenchyma and the diameter and number of vessels per mm^2 tend to be larger than in trees and shrubs (Koek-Noorman et al., 1979; Ter Welle and Koek-Noorman, 1981; Vliet, 1981). In epiphytes of the family, there is a cortical parenchyma in the stem that was interpreted as a hygroscopic tissue that helped to prevent desiccation, a common issue in epiphytes (Reginato et al., 2009). Other studies on the anatomy of the family have focused on the relationship between anatomical variations and environmental factors such as light radiation, precipitation, and average annual temperature (Ely et al., 2005a, b; Gluzek, 2005; Gardoni et al., 2007; Bosio, 2008). In general, the anatomical descriptions at the growth habit level in Melastomataceae are rare, and the comparisons of anatomical features among species with different growth habits are even more unusual and limited to brief annotations. Moreover, some publications do not even recognize the existence of special adaptations related with growth habits in Melastomataceae (Carlquist, 1985).

In this study, we analyzed whether stem anatomy characteristics vary consistently across growth habits of Melastomataceae. To do so we compared 18 species of this family, which were chosen to sample separate growth habits within the family, accounting for phylogeny by including representatives from different tribes. We used here a growth habit classification that we consolidated from a literature review, trying to reconcile multiple schemes of classification of habits across vascular plants. The stem anatomy of the 18 species was analyzed using 125 characters observed from histological preparations.

MATERIALS AND METHODS

Specimen collection

We collected one specimen from each of 18 species of Melastomataceae (Table 1) for descriptions and comparisons. We chose these species aiming to sample across different ecosystems of Colombian Andes (tropical wet forest, tropical rainforest, tropical moist forest, and páramo), different clades of Melastomataceae (five out of the 11 major clades recognized by Clausing and Renner (2001)), and especially all the different growth habits known to occur in the family. Voucher specimens were identified, described, and deposited at the Herbarium of University of Antioquia (HUA); duplicate specimens were sent to the Herbarium of the Universidad Católica de Oriente (HUCO), both in Colombia.

Anatomical sample preparation

Stem portions of the fifth most apical internode were collected for analysis. The portions were fixed with FAA (10% of 40% formaldehyde, 50% of 96% ethanol, 5% of 96% acetic acid, and 35% of distilled water). Transverse, radial and tangential sections of each stem were obtained by free-hand slicing. Sections were stained with toluidine blue 0.05% and observed and photographed using a setting formed by an Olympus BH2 microscope (Olympus, Tokio, Japan), an Olympus e620 camera (Olympus, Tokio, Japan), and a NDPL-1 (2X) camera adapter (Amscope, Los Angeles, USA). This setting was located in the Palynology Laboratory of the Universidad de Antioquia, Medellín, Colombia.

ANATOMICAL ANALYSIS

To describe the stem anatomy from the anatomical preparations we used 125 characters based on Metcalfe and Chalk (1950), Dickison (1975), Carlquist (1985), Wheeler et al. (1989), and Evert (2006). These characters describe the epidermis (12 characters), peridermis (15), cortex in general (3), cortical parenchyma (8), cortical collenchyma (6), cortical sclerenchyma (6), vascular system in general (8), phloem (9), xylem vessels (14), xylem

Table 1: List of species collected for this study and the ecosystem where they were sampled. All collections were performed in Colombia. Voucher specimens were deposited in the Herbarium of the Universidad de Antioquia (HUA).

Species	Voucher HUA	Locality	Ecosystem
<i>Adelobotrys adscendens</i> Triana	193782	Cerro La Judia, Florida blanca, Santander	tropical moist forest
<i>Arthrostemma ciliatum</i> Pav. ex D. Don	193781	Frontino, Antioquia	tropical rainforest
<i>Blakea fasciculata</i> Gleason	193791	Frontino, Antioquia	tropical rainforest
<i>Blakea granatensis</i> Naudin	193792	Cerro La Judia, Florida blanca, Santander	tropical moist forest
<i>Blakea quadrangularis</i> Triana	193788	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest
<i>Blakea rosea</i> (Ruiz & Pav.) D. Don	193793	San Luis, Antioquia	tropical wet forest
<i>Blakea watsonii</i> (Cogn.) Penneys & Almeda	193787	San Luis, Antioquia	tropical wet forest
<i>Chaetolepis microphylla</i> (Bonpl.) Miq.	193789	Belmira, Antioquia	tropical moist forest
<i>Clidemia epiphytica</i> (Triana) Cogn.	193790	San Luis, Antioquia	tropical wet forest
<i>Graffenreida emarginata</i> (Ruiz & Pav.) Triana	193785	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest
<i>Heterocentron subtriplinervium</i> (Link & Otto) A. Braun & C.D. Bouché	193794	Medellín, Antioquia	tropical moist forest
<i>Meriania nobilis</i> Triana	193779	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest
<i>Miconia jahnii</i> Pittier	193786	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest
<i>Miconia salicifolia</i> Naudin	193795	Urrao, Antioquia	paramo
<i>Monochaetum multiflorum</i> (Bonpl.) Naudin	193783	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest
<i>Monolena primuliflora</i> Hook. f.	192632	Amalfi, Antioquia	tropical wet forest
<i>Tibouchina kingii</i> Wurdack	193780	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest
<i>Tibouchina lepidota</i> (Bonpl.) Baill.	193784	Cerro Bonifacio, Carmen de Viboral, Antioquia	tropical moist forest

fibers (6), xylem axial parenchyma (6), xylem rays (9), miscellaneous xylem characters (7), pith in general (4), medullary parenchyma (8), and medullary sclerenchyma (4). The quantitative characters were the tangential diameter of solitary vessels, the number of vessels per mm^2 , and the vessel axial length. The number of measurements per specimen were 30 for the tangential diameter of solitary vessels, 10 for the number of vessels per mm^2 , and 20 for the vessel axial length. See the appendix for further details on these characters.

Description of growth habits

Growth habit of analyzed specimens was determined using field observations (Table 2), according to the six growth habits that we condensed from definitions in the literature (Eiten, 1991; Renner, 1993; Moffet, 2000; Al-

meda, 2001; Font-Quer, 2001; Mendoza and Ramírez, 2006; Simpson, 2006):

Trees/shrubs: characterized by germination and growth on the ground, the presence of stem and branches with secondary growth (Simpson, 2006), self-support and erect orientation (Font-Quer, 2001; Figs. 1A-B).

Herbs: characterized by the thin and brittle stem, mainly unbranched, with decumbent or prostrate orientation. If present, branches are basally inserted (Simpson, 2006), and, if they have an erect orientation, their height is less than two meters (Mendoza and Ramírez, 2006) (Fig. 1C). Herbs show the most variable features, making it a group which is hard to define (Eiten, 1991).

Lianas: exhibit a stem with initial erect orientation, when they are surrounded by abundant vegetation, they can lean or climb on a phorophyte or other

Table 2: Established growth habits for analyzed species. Note: “1st” primary, “2nd” secondary, “-” non-registered character.

Species	Rooting	Position of stem base	Stem orientation	Type of growth	Type of branchs	Height of 1st aerial branch (m)	Principal axis evident	Flower (inflorescence) position	Heighth of plant (m)	Support structures	Growth habit
<i>Adelobotrys adscendens</i> Triana	Soil	Soil	Prostrate	2nd	Woody aerial	4	Yes	Terminal	8	Adventitious roots	Vine
<i>Arthrostemma ciliatum</i> Pav. ex D. Don	Soil	Soil	Decumbent	1st	Non-woody basal	0	Yes	Terminal	1	-	Herb
<i>Blakea fasciculata</i> Gleason	Soil	Soil	Fickle stem supported	2nd	Woody aerial	0.4	Yes	Axillary	15	Adventitious roots	Liana
<i>Blakea granatensis</i> Naudin	Soil	Soil	Prostrate	2nd	Woody aerial	3	Yes	Axillary	6	Adventitious roots	Vine
<i>Blakea quadrangularis</i> Triana	Soil	Soil	Fickle stem supported	2nd	Woody aerial	2	Yes	Axillary	5	Adventitious roots	Liana
<i>Blakea rosea</i> (Ruiz & Pav.) D. Don	Soil	On Phorophyte	Decumbent	2nd	Woody basal	0	Yes	Axillary	4	Roots	Hemiepiphyte
<i>Blakea watsonii</i> (Cogn.) Penneys & Almeda	Soil	On Phorophyte	Decumbent	2nd	Woody basal	0	Yes	Axillary	7	Roots	Hemiepiphyte
<i>Chaetolepis microphylla</i> (Bonpl.) Miq.	Soil	Different substrates	Erect	2nd	Woody basal	0	No	Terminal	0.3	-	Herb
<i>Clidemia epiphytica</i> (Triana) Cogn.	Soil	Soil	Prostrate	2nd	Woody aerial	0.5	Yes	Axillary	10	Adventitious roots	Vine
<i>Graffenreida emarginata</i> (Ruiz & Pav.) Triana	Soil	Soil	Erect	2nd	Woody aerial	2	Yes	Terminal	7	-	Tree
<i>Heterocentron subtriplinervium</i> (Link & Otto) A. Braun & C.D. Bouché	Soil	Soil	Decumbent	2nd	Woody aerial	0.3	Yes	Terminal	1	-	Herb
<i>Meriania nobilis</i> Triana	Soil	Soil	Erect	2nd	Woody aerial	3	Yes	Terminal	5	-	Tree
<i>Miconia jahnii</i> Pittier	Soil	Soil	Erect	2nd	Woody aerial	1.5	Yes	Terminal	4	-	Tree
<i>Miconia salicifolia</i> Naudin	Soil	Soil	Erect	2nd	Woody aerial	0.2	Yes	Terminal	3	-	Shrub
<i>Monochaetum multiflorum</i> (Bonpl.) Naudin	Soil	Soil	Fickle stem supported	2nd	Woody aerial	2.5	Yes	Terminal	5	-	Liana
<i>Monolena primuliflora</i> Hook. f.	Superficial	Different substrates	Decumbent	1st	-	-	Yes	Axillary	0.15	Roots	Epiphyte
<i>Tibouchina kingii</i> Wurdack	Soil	Soil	Erect	2nd	-	-	Yes	Terminal	1	-	Herb
<i>Tibouchina lepidota</i> (Bonpl.) Baill.	Soil	Soil	Erect	2nd	Woody aerial	1.5	Yes	Terminal	6	-	Tree



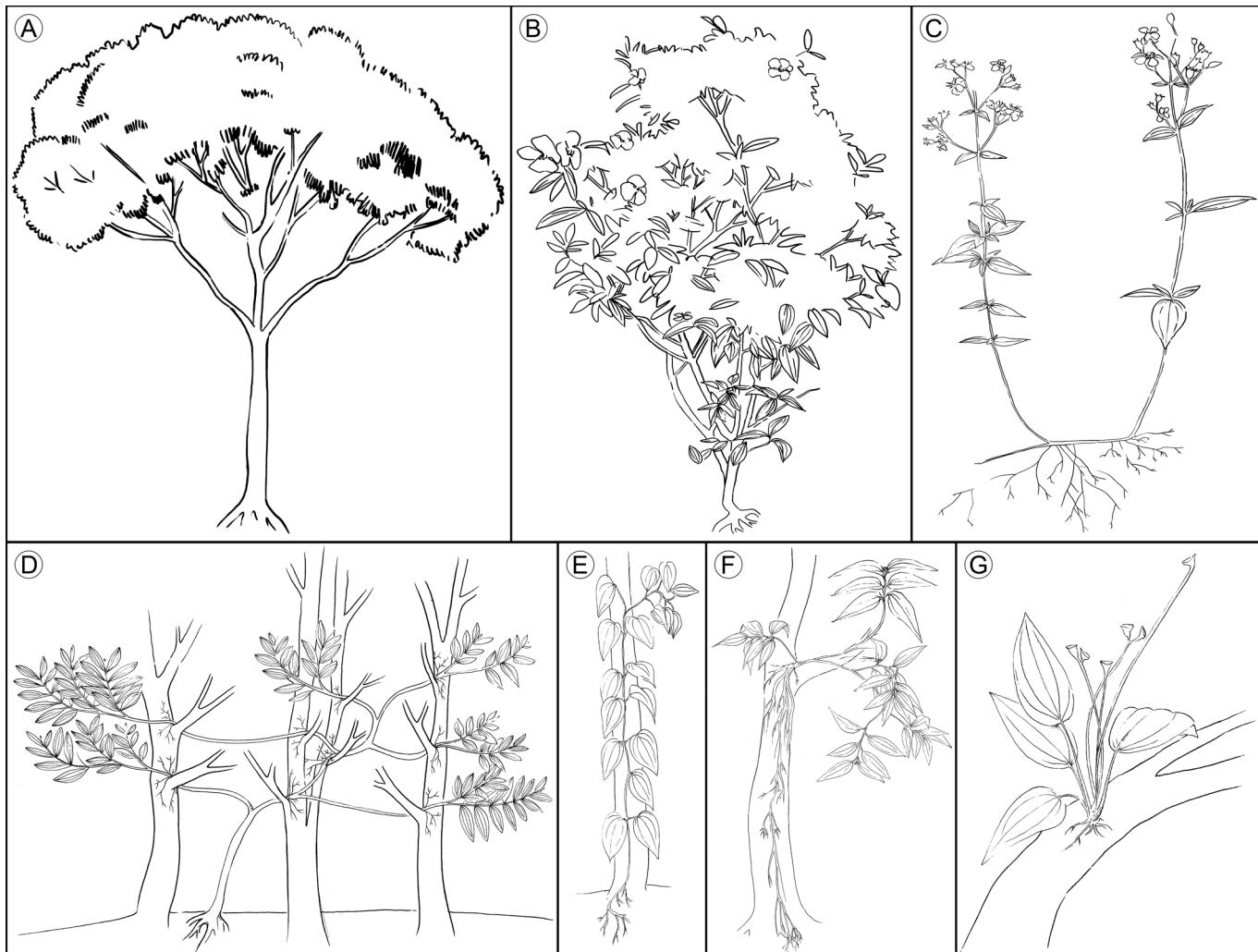


Figure 1: Growth habits of vascular plants compiled from the literature. A. trees; B. shrubs; C. herbs; D. lianas; E. vines; F. hemiepiphytes; G. epiphytes. Illustrations by Elizabeth Builes.

object to ascend; some can develop adventitious roots on supported branches (Rowe and Speck, 2004). The growth strategy of this group is similar to that of vines, differing in the vertical orientation of the liana's stem, their self-support in early stages of development, and their thicker stems (Rowe and Speck, 2004). The species of this group are also described as climbers, scandent shrubs, and creepers (Putz, 1984; Moffet, 2000) (Fig. 1D).

Vines: characterized by the prostrated orientation of stem, never self-supported. Early in life, the stem creeps until it finds vertical support to grow vertically, usually

developing adventitious roots (Moffet, 2000; Rowe and Speck, 2004). In these plants, the principal stem is thin and flexible (Fig. 1E).

Hemiepiphytes: stem base above the ground, supported on a host plant, with roots reaching the ground (Moffet, 2000; Mendoza and Ramírez, 2006). The stem base is thicker than the rest of the stem and branches, acting as the root insertion point, basally prostrated and usually erect at the branch level. The rest of the stem is short and ramified. Commonly, species identified as hemiepiphytes can grow on the ground, with stem and branches vertically oriented like a shrub (Fig. 1F).



Epiphytes: roots and adventitious roots attach to the substrate without reaching the ground (Moffet, 2000; Mendoza and Ramírez, 2006), although some species can also grow on the ground (facultative epiphytes (Renner, 1993)) (Fig. 1G).

Statistical analyses

Group comparisons of quantitative characters (tangential diameter of solitary vessels, the number of vessels per mm^2 , and the vessel axial length) were performed with nested ANOVAs, one for each variable, with growth habit as an independent factor, and species within growth habit as the nested factor. When group differences were significant, the Tukey test was used for multiple comparisons of groups. Due to the remarkable differences in diameter and number of vessels of *Chaetolepis microphylla* (Bonpl.) Miq., we repeated the analyses excluding that species. Our results were almost identical after excluding that species; therefore, we show here the analyses without it. Statistical analyses were conducted using R (R Core Team, 2017).

RESULTS

Anatomical descriptions

Trees/shrubs

Epidermis: Trichomes are present in *Graffenrieda emarginata* (Ruiz & Pav.) Triana, *Miconia jahnii* Pittier and *Miconia salicifolia* Naudin only. Dendritic hairs are showed by *Graffenrieda emarginata*, *Miconia jahnii*, and *Miconia salicifolia*; glands are present only in the two former species. Epidermis is absent in the fifth internode of *Tibouchina lepidota* (Bonpl.) Baill. and *Miconia salicifolia*, where it is replaced by periderm. Epidermis in *Meriania nobilis* Triana, *Graffenrieda emarginata*, and *Miconia jahnii* is single-layered and with epicuticular wax like granules (Figs. 2A-D).

Periderm: Periderm is absent in the fifth internode of *Graffenrieda emarginata* and *Meriania nobilis*. Scaly bark is only present in *Tibouchina lepidota*. Periderm in *Tibouchina lepidota*, *Miconia jahnii* and *Miconia sali-*

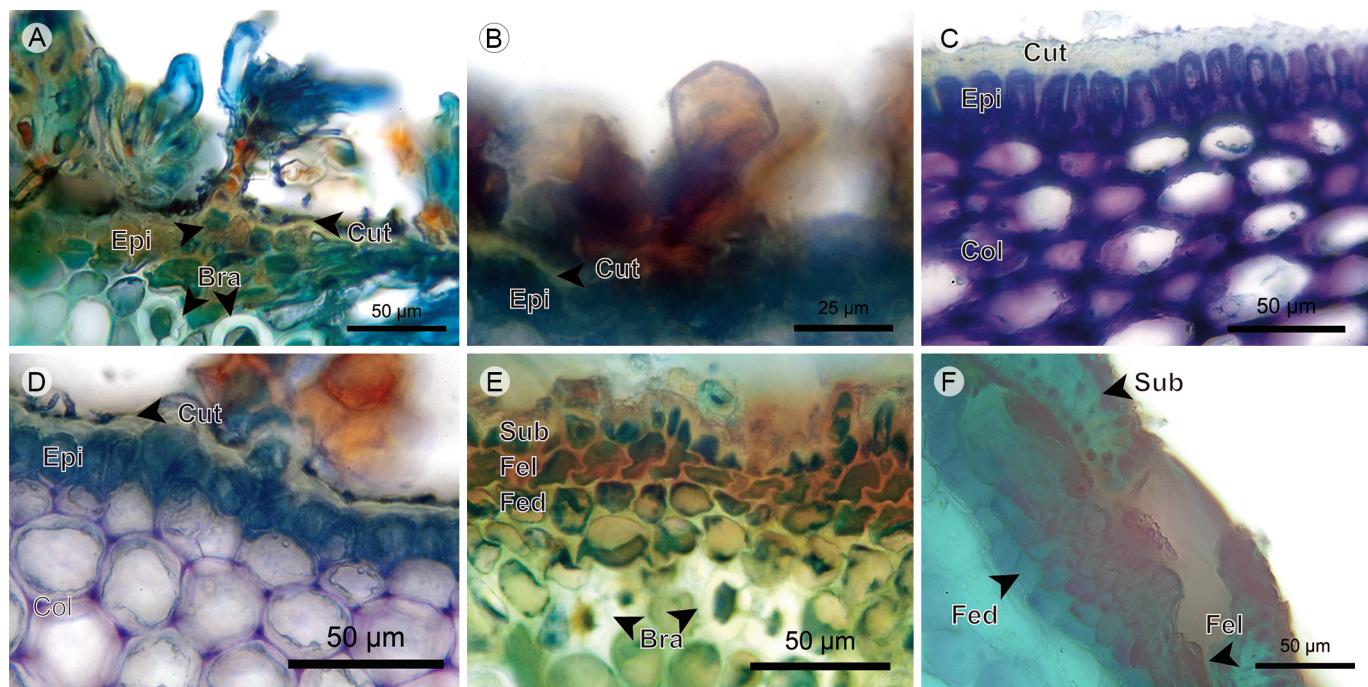


Figure 2: Trees/shrubs: Epidermis. A. dendritic hair in *Miconia jahnii* Pittier; B. glandular hair in *Graffenrieda emarginata* (Ruiz & Pav.) Triana; C. epidermis and cortex in *Meriania nobilis* Triana; D. epidermis and cortex in *Graffenrieda emarginata* (Ruiz & Pav.) Triana; E. periderm and cortex in *Miconia jahnii* Pittier; F. lenticel in *Tibouchina lepidota* (Bonpl.) Baill. (Bra, stone cell; Col, angular collenchyma; Cut, cuticle; Epi, epidermis; Fed, phellogen; Fel, phellogen; Sub, phellem).

cifolia is superficial, single-layered and the formation is complete around the stem. The phellem is one- to six-layered with cells in radial disposition without intercellular spaces; the cells have pits only in *Tibouchina lepidota*. The phellogen is simple and one-layered. The phellogen is one- to three-layered. Lenticels only occur in *Tibouchina lepidota* periderm (Figs. 2E, F).

Cortex: The cortex is mainly formed by parenchymatous cells with isodiametric shape, intercellular spaces, diverse materials, and druse accumulation (except in *Tibouchina lepidota* and *Miconia jahnii*). Photosynthetic parenchyma only occurs in *Meriania nobilis* cortex. The collenchyma is of the angular type, forming a continuous multiple-layered tissue around the stem inside to epidermis/periderm in *Meriania nobilis*, *Graffenrieda emarginata*, and *Miconia salicifolia*. Stone cells with simple and ramiform pits are found in *Meriania nobilis*, *Graffenrieda emarginata* and *Miconia jahnii*. In *Meriania nobilis* and *Graffenrieda emarginata*, the stone cells form a continuous three-layered tissue around the endodermis. In *Miconia jahnii* the stone cells form clusters ranging from endodermis to periderm. Vascular bundles compound by just xylem are only seen in *Tibouchina lepidota* (Figs. 2C-E, 3).

Endodermis and pericycle: The endodermis is only absent in *Miconia salicifolia*. In the rest of the species, the endodermis forms a continuous one- to three-lay-

ered tissue around the stem. Casparyan strip is only present in *Tibouchina lepidota* and *Miconia jahnii*. The pericycle forms a continuous two- to the three-layered tissue between the endodermis and the phloem with cells in radial disposition (Figs. 4A-C).

Phloem: Sieve-tube elements are arranged in radial disposition in *Meriania nobilis*, *Miconia jahnii* and *Miconia salicifolia*, scattered in *Tibouchina lepidota*, and *Graffenrieda emarginata*. Sieve plates are transverse, except for *Tibouchina lepidota*, where they are inclined. Fibers are only present in *Miconia jahnii* and *Miconia salicifolia*; in the former, grouped in tangential bands; in the latter, forming clusters of six to eight cells located on the distal part of phloem. Axial parenchyma is composed of strands of short cells containing druses, except for *Graffenrieda emarginata* (Figs. 4B-E).

Vascular cambium: The vascular cambium is a one- to five-layered tissue without discontinuities, variations or irregularities; xylem and phloem form continuous tissues with constant width (Figs. 4B-D).

Xylem vessels: The vessel arrangement has radial to diagonal patterns and the grouping is in multiples of two to four with a variable proportion of solitary vessels. The outline of solitary vessels is rounded in *Meriania nobilis* and *Miconia jahnii*, and angular in *Tibouchina lepidota*, *Graffenrieda emarginata*, and *Miconia salicifolia*. Perforation plates are simple and inclined. Inter-

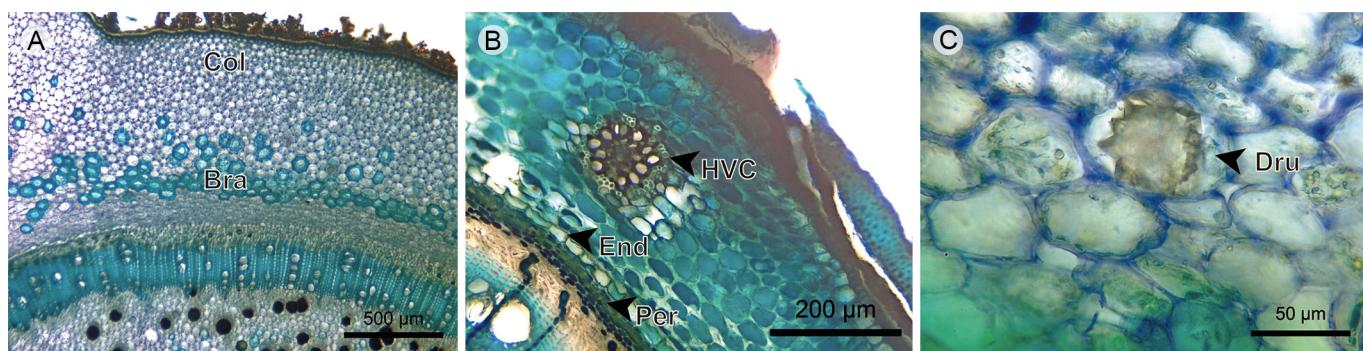


Figure 3: Trees/shrubs: Cortex. A. transverse section of *Graffenrieda emarginata* (Ruiz & Pav.) Triana; B. cortex of *Tibouchina lepidota* (Bonpl.) Baill.; C. cortex of *Meriania nobilis* Triana. (Bra, stone cell; Col, angular collenchyma; Dru, druse; End, endodermis; HVC, cortical vascular bundle; Per, pericycle).

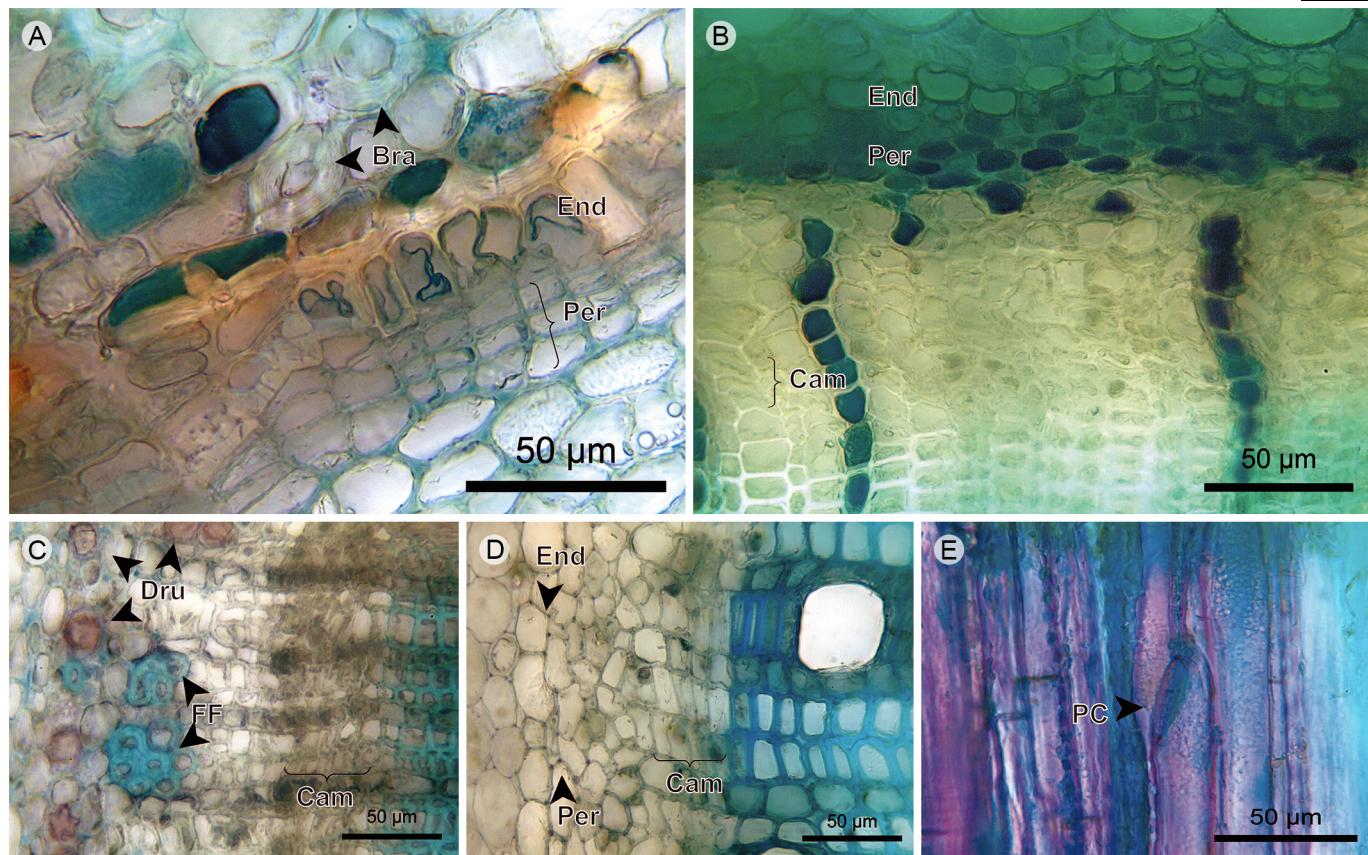


Figure 4: Trees/shrubs: Endodermis, pericycle, phloem and vascular cambium. A. endodermis and pericycle of *Miconia jahnnii* Pittier; B. endodermis, pericycle, and phloem of *Tibouchina lepidota* (Bonpl.) Baill.; C. phloem of *Miconia salicifolia* Naudin; D. phloem and cambium of *Graffenrieda emarginata* (Ruiz & Pav.) Triana; E. phloem in radial section of *Tibouchina lepidota* (Bonpl.) Baill. (Bra, stone cell; Cam, vascular cambium; Dru, druse; End, endodermis; FF, phloem fiber; PC, sieve plate; Per, pericycle).

vessel pits are rounded and bordered, have an opposite arrangement in *Meriania nobilis* and alternate in the other species (Fig. 5).

Xylem fibers: The fibers are of the libriform type and have transversal septa. The xylem has fiber dimorphism with two morphotypes of libriform fibers. The difference between morphotypes is in the lumen diameter and the thickness of the secondary wall. Dimorphic fibers are grouped in two to five layers formed by units that can range from short clusters to long tangential bands. The distance between dimorphic fibers is larger than the distance between rays (Figs. 5A, B, E, F).

Xylem rays: Rays are uniserial, but *Tibouchina lepidota* shows multiseriate portions. The cellular composition is upright and square cells. Vessel-ray pits are

rounded, oval or scalariform and half-bordered, with border only in the vessel lumen (Figs. 5A, B, E-G).

Pith: The pith has internal primary phloem towards the periphery; in the center, it has parenchymatous cells with isodiametric shape and diverse materials and accumulation of druses. Vascular bundles scattered throughout the pith are concentric amphicribal of primary growth where the phloem is predominant in *Miconia salicifolia*, being the xylem very reduced in *Graffenrieda emarginata* and *Miconia jahnnii*, and non-existent in *Tibouchina lepidota* and *Meriania nobilis*. The *Miconia jahnnii* pith is formed largely by stone cells with a wide lumen and simple and ramiform pits. The piths of *Graffenrieda emarginata* are common idioblasts that are axially elongated and related to the phloem, distributed around and within the internal phloem and vascular bundles (Fig. 6).

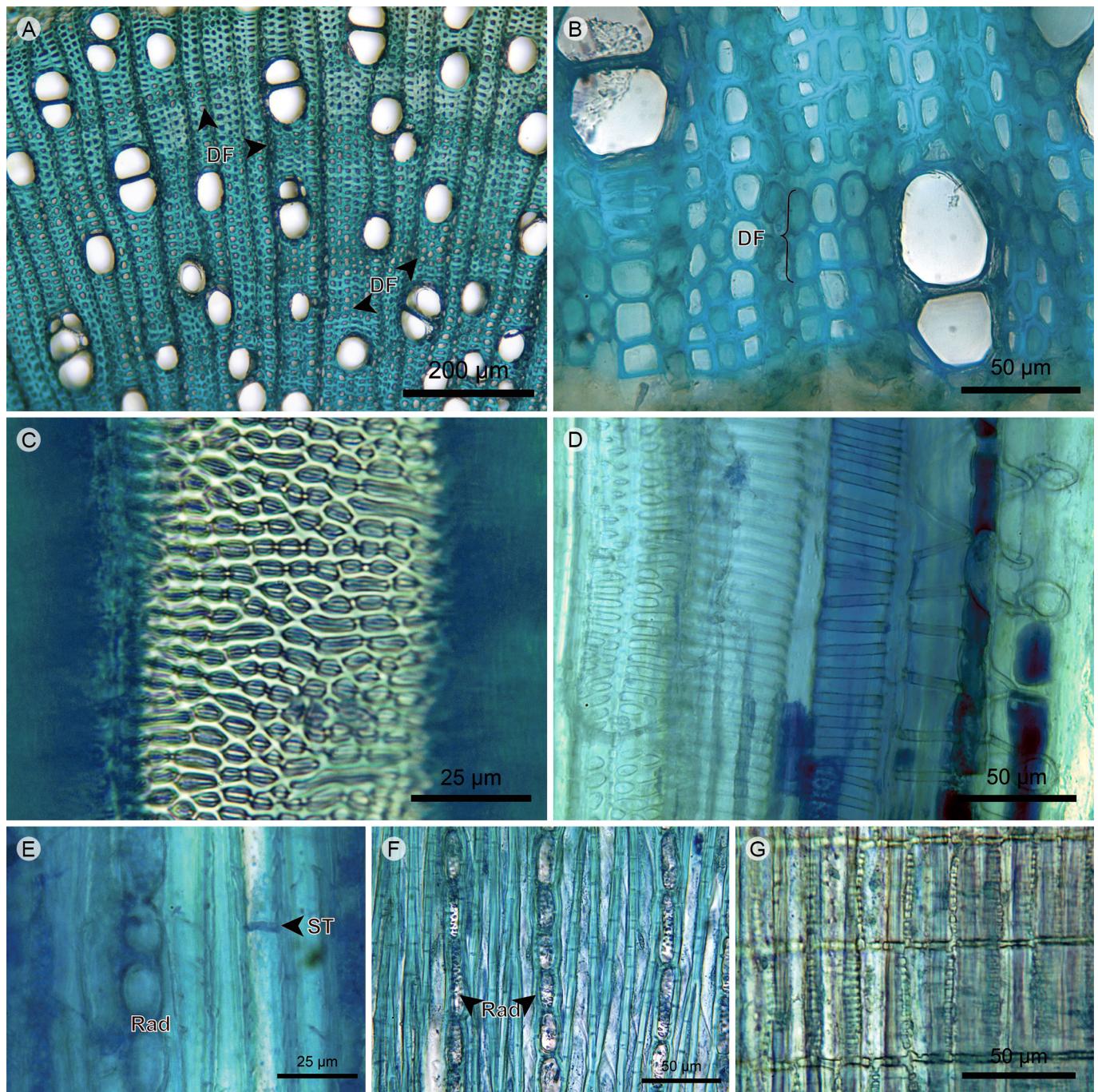


Figure 5: Trees/shrubs: Xylem. A. xylem of *Miconia jahnii* Pittier; B. xylem of *Graffenrieda emarginata* (Ruiz & Pav.) Triana; C. vessel of *Meriania nobilis* Triana in tangential section; D. protoxylem in radial section of *Tibouchina lepidota* (Bonpl.) Baill.; E. xylem in tangential section of *Meriania nobilis* Triana; F. xylem in tangential section of *Miconia salicifolia* Naudin; G. rays in radial section of *Miconia salicifolia* Naudin. (DF, fiber dimorphism; Rad, ray; ST, fiber transversal septum).

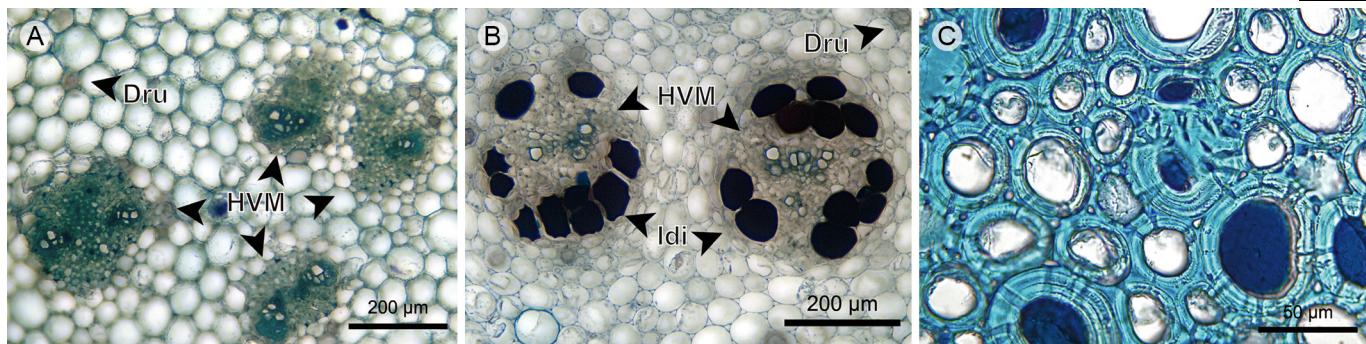


Figure 6: Trees/shrubs: Pith. A. pith of *Miconia salicifolia* Naudin; B. pith of *Graffenreida emarginata* (Ruiz & Pav.) Triana; C. stone cells in pith of *Miconia jahni* Pittier. (Dru, druse; HVM, medullary vascular bundle; Idi, idioblast).

Herbs

Epidermis: The trichomes are pluricellular; simple unbranched in *Heterocentron subtriplinervium* (Link & Otto) A. Braun & C.D. Bouché and *Arthrostemma ciliatum* Pav. ex D. Don, dendritic in *Tibouchina kingii* Wurdack and *Chaetolepis microphylla*. Trichomes of *Arthrostemma ciliatum* are scarce and longer than in the other

species. The epidermis is absent in the fifth internode of *Tibouchina kingii*. The epidermis is single-layered and has an epicuticular wax with the form of granules in *Heterocentron subtriplinervium* and *Arthrostemma ciliatum*, whereas in *Chaetolepis microphylla* the epicuticular wax consists of short and thin filaments in perpendicular direction to the epidermis (Figs. 7A-D).

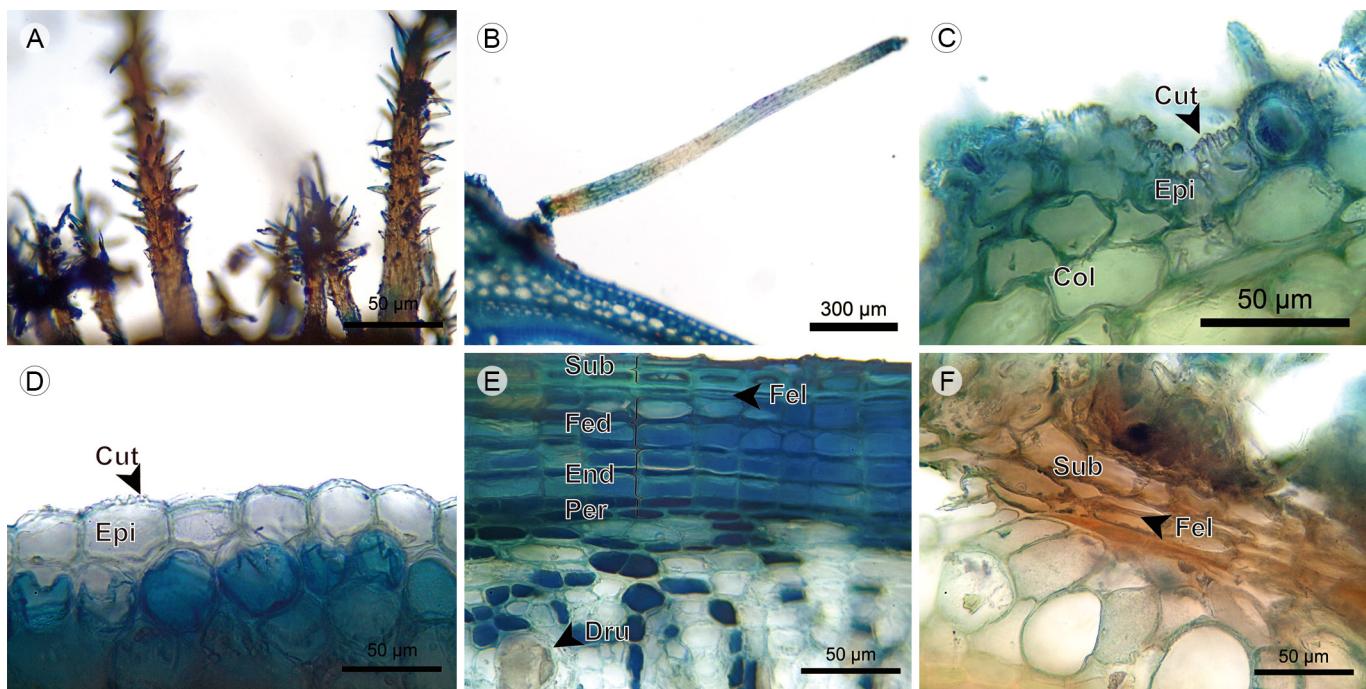


Figure 7: Herbs: Epidermis. A. dendritic hairs of *Tibouchina kingii* Wurdack; B. unbranched hair and epidermis of *Arthrostemma ciliatum* Pav. ex D. Don; C. epidermis and cortex of *Chaetolepis microphylla* (Bonpl.) Miq.; D. epidermis of *Arthrostemma ciliatum* Pav. ex D. Don; E. periderm, endodermis, pericycle, and phloem of *Tibouchina kingii* Wurdack; F. periderm and cortex of *Chaetolepis microphylla* (Bonpl.) Miq. (Col, angular collenchyma; Cut, cuticle; Dru, druse; End, endodermis; Epi, epidermis; Fed, phellogen; Fel, phellogen; Per, pericycle; Sub, phellem).

Periderm: Periderm is present only in *Tibouchina kingii* and *Chaetolepis microphylla*; in this latter species the transition where the periderm is forming below the epidermis is observed. Scaly bark is present in *Tibouchina kingii* only. The periderm is superficial, single-layered and the formation is complete around the stem. The phellem is two- to four-layered with cells in radial disposition. The phellogen is one-layered. The phellogerm is four- to six-layered with cells in radial disposition (Figs. 7E, F).

Cortex: The cortex is absent in the fifth internode of *Tibouchina kingii*, just below the phellogerm is the endodermis. In other species, the cortex is mainly formed by parenchymatous cells with isodiametric shape, with diverse materials and accumulation of druses in *Heterocentron subtriplinervium* and *Arthrostemma ciliatum*, and presence of chloroplasts in *Arthrostemma ciliatum*. The collenchyma is of the angular type, forming a multi-layered tissue below the epidermis in *Heterocentron subtriplinervium* and *Chaetolepis microphylla*, and in the middle of the cortex in *Arthrostemma ciliatum*. In *Heterocentron subtriplinervium* and *Arthrostemma ciliatum*, vascular bundles of primary growth are concentric amphicribral and those located in the stem angles are quadrangular (Figs. 7C-F, 8).

Endodermis and pericycle: The endodermis is below the cortex, it is one- to three-layered and the tangential walls have more contact area. The pericycle is a

five-layered continuous tissue between the endodermis and phloem, only present in *Heterocentron subtriplinervium* (Figs. 8A, 9A,B).

Phloem: Sieve-tube elements are scattered in *Tibouchina kingii*, *Heterocentron subtriplinervium*, and *Chaetolepis microphylla*, and in radial disposition in *Arthrostemma ciliatum*. Sieve plates are transverse, but inclined in *Tibouchina kingii*. Axial parenchyma is organized in fusiform cells in *Heterocentron subtriplinervium* and *Chaetolepis microphylla*, and in a strand of short cells containing druses in *Tibouchina kingii* (Figs. 7E, 8B, C, 9).

Vascular cambium (procambium in *Arthrostemma ciliatum*): The vascular cambium is not observed in the fifth internode of *Arthrostemma ciliatum*, the procambium is a one-layered and simple tissue, thus the phloem and xylem are primary. In the other species, the vascular cambium is a one- to four-layered tissue without discontinuities, variations or irregularities (Figs. 8B, C, 9A-C).

Xylem vessels: Vessel arrangement has a radial to diagonal pattern, mainly solitary vessels with a little proportion of radial multiples, more common in *Heterocentron subtriplinervium*. The outline of solitary vessels is rounded in *T. kingii* and *Chaetolepis microphylla*, the outline is angular in *Heterocentron subtriplinervium*. The perforation plates are simple and inclined. Intervessel pits are rounded and bordered and have an alternate arrangement (Figs. 10A-D).

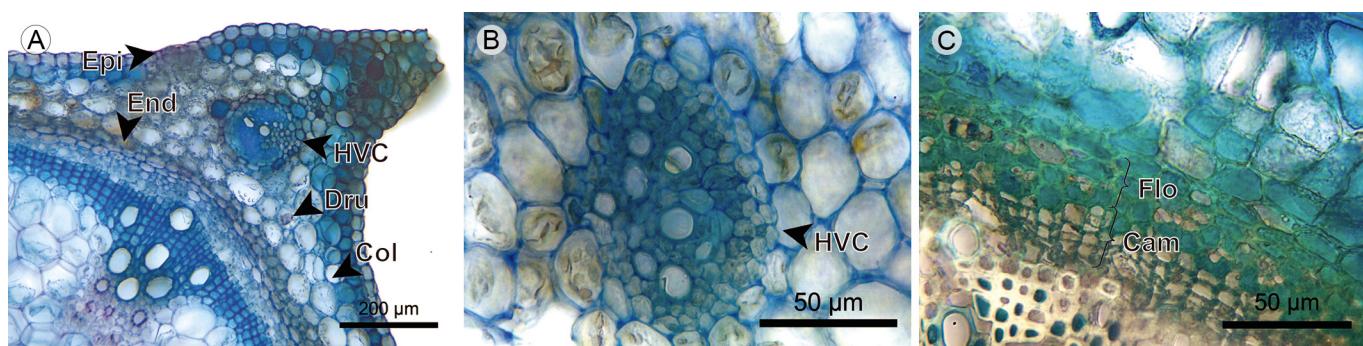


Figure 8: Herbs: Cortex. A. transverse section of *Arthrostemma ciliatum* Pav. ex D. Don; B. cortex and cortical vascular bundle of *Heterocentron subtriplinervium* (Link & Otto) A. Braun & C.D. Bouché; C. cortex, phloem, and vascular cambium of *Chaetolepis microphylla* (Bonpl.) Miq. (Cam, vascular cambium; Col, angular collenchyma; Dru, druse; End, endodermis; Epi, epidermis; Flo, phloem; HVC, cortical vascular bundle).

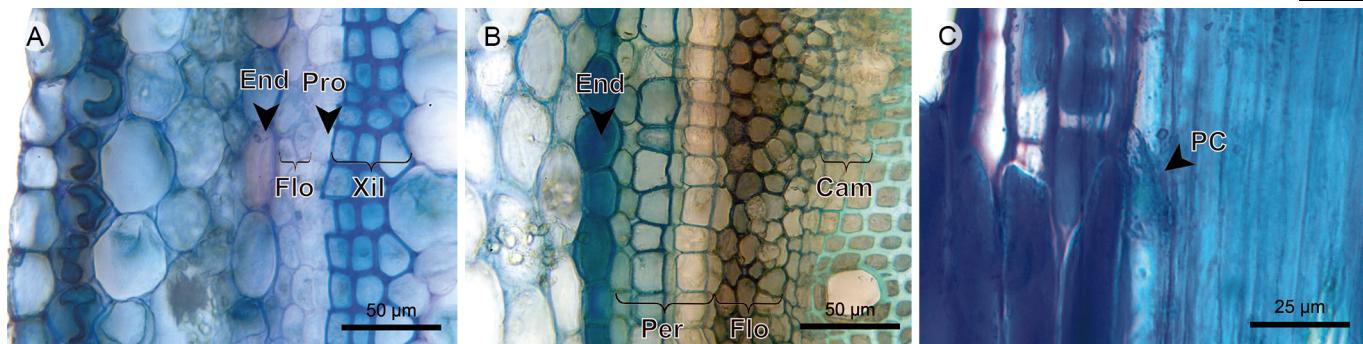


Figure 9: Herbs: Endodermis, pericycle, phloem, and vascular cambium. A. transverse section of *Arthrostemma ciliatum* Pav. ex D. Don; B. transverse section of *Heterocentron subtriplinervium* (Link & Otto) A. Braun & C.D. Bouché; C. phloem in radial section of *Tibouchina kingii* Wurdack. (Cam, vascular cambium; End, endodermis; Flo, phloem; Per, pericycle; PC, sieve plate; Pro, procambium; Xil, xylem).

Xylem fibers: The fibers are of the libriform type and have transversal septa only in *Heterocentron subtriplinervium* and *Arthrostemma ciliatum*. Fiber dimorphism only in *Tibouchina kingii* and *Chaetolepis microphylla*. Dimorphic fibers in *Tibouchina kingii* are grouped in four- to six-layered tangential bands and the distance between the bands is bigger than the distance between rays. In *Chaetolepis microphylla* the dimorphic fibers form one tangential band of five to seven layers (Figs. 10B, E).

Xylem rays: Rays uniserial, formed by upright and square cells. Vessel-ray pits are rounded to oval and half-bordered, with a border in the vessel lumen only (Figs. 10B, F, G).

Pith: The pith has internal primary phloem towards the periphery, which is continuous in *Chaetolepis microphylla* and discontinuous in other species. The parenchyma is formed by cells with isodiametric shape and diverse materials and accumulation of druses. In *Chaetolepis microphylla* the pith is reduced and without intercellular spaces between parenchymatous cells. Vascular bundles scattered throughout the pith are concentric amphiocribral of primary growth where the phloem is predominant, being the xylem very reduced in *Arthrostemma ciliatum* and non-existent in *Tibouchina kingii*. In the pith of *Chaetolepis microphylla* the vascular bundles are absent (Fig. 11).

Lianas

Epidermis: Trichomes have a dendritic form in *Monochaetum multiflorum* (Bonpl.) Naudin and *Blakea fasciculata* Gleason, and are absent in *Blakea quadrangularis*. The epidermis is single-layered, the transition with the periderm is formed below the epidermis in *Blakea quadrangularis* and *Blakea fasciculata*. The epidermis is absent in the fifth internode of *Monochaetum multiflorum*. The epicuticular wax has the shape of granules (Figs. 12A-D).

Periderm: Scaly bark is present in *Monochaetum multiflorum* only. The periderm is superficial and single-layered. The phellem is two- to ten-layered with cells in radial disposition, without intercellular spaces and with pits (except for *Blakea fasciculata*). The phellogen is one- to two-layered. The phellogen is one- to three-layered with cells in radial disposition, and is absent in *Blakea fasciculata* (Figs. 12D-F, 13A, 14B).

Cortex: The cortex in the fifth internode of *Monochaetum multiflorum* is absent, the endodermis is just below the phellogen. In other species, the cortex is mainly formed by parenchymatous cells with isodiametric shape, diverse materials, and accumulation of druses, with chloroplasts present only in *Blakea fasciculata*. Collenchyma is only present in *Blakea fasciculata* and of the angular type, forming a five- to six-layered continuous tissue below the phellogen, the latter containing chloroplasts and druses. The stone cells form a four-layered discontinuous

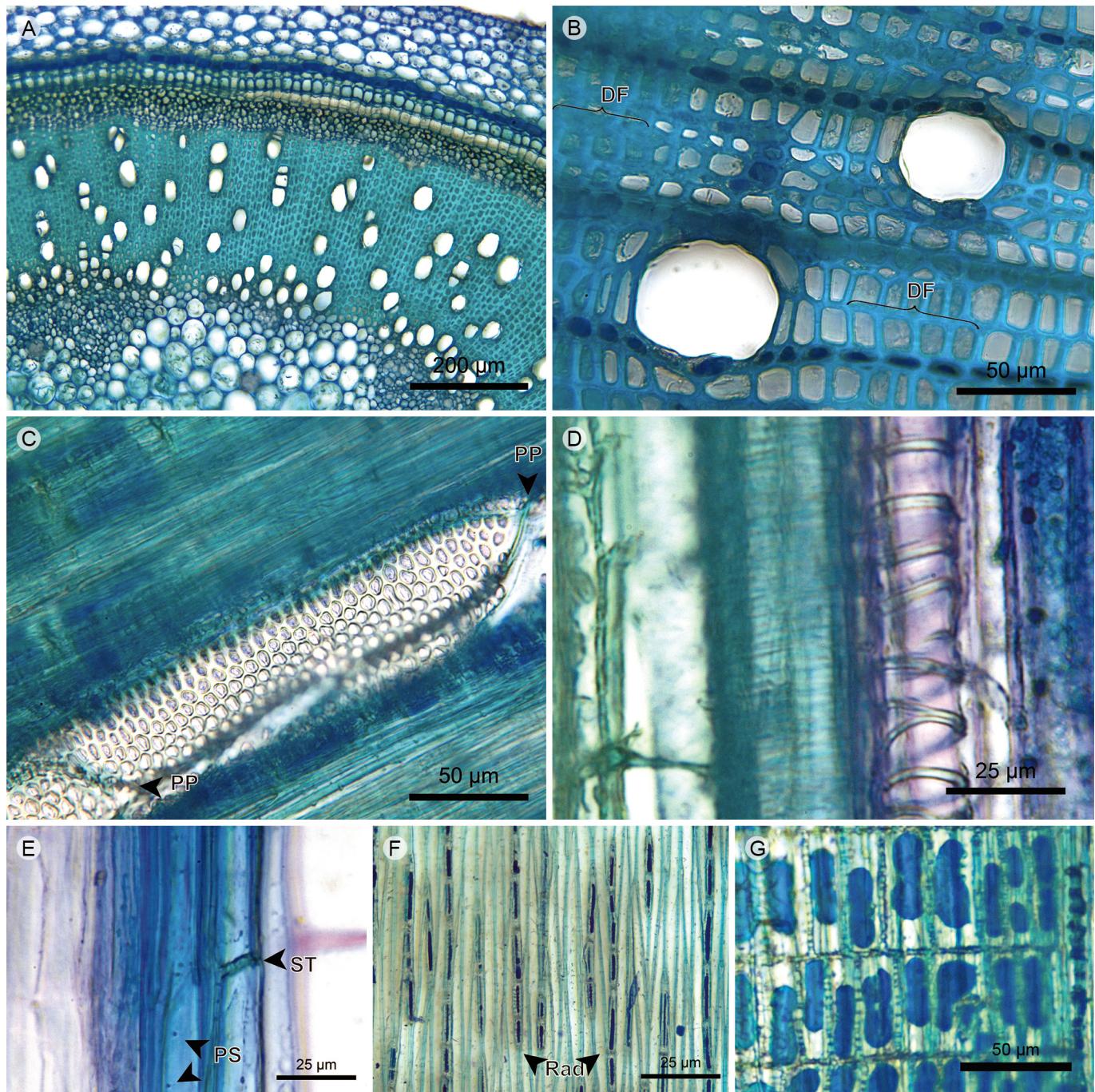


Figure 10: Herbs: Xylem. A. xylem of *Heterocentron subtriplinervium* (Link & Otto) A. Braun & C.D. Bouché; B. xylem of *Tibouchina kingii* Wurdack; C. vessel in tangential section of *Tibouchina kingii* Wurdack; D. protoxylem in radial section of *Heterocentron subtriplinervium* (Link & Otto) A. Braun & C.D. Bouché; E. xylem fibers in radial section of *Arthrostemma ciliatum* Pav. ex D. Don; F. xylem in tangential section of *Tibouchina kingii* Wurdack; G. rays in radial section of *Tibouchina kingii* Wurdack. (DF, fiber dimorphism; PP, perforation plate; PS, fiber simple pit; Rad, ray; ST, fiber transversal septum).

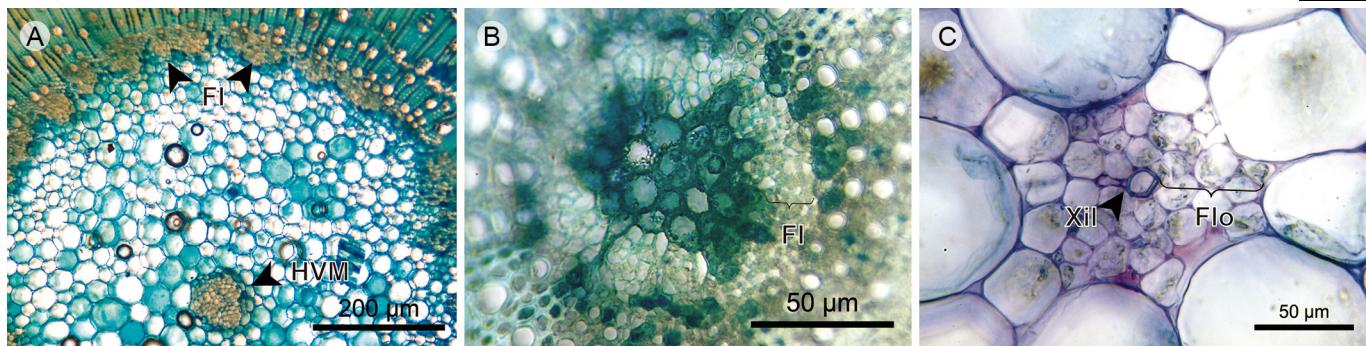


Figure 11: Herbs: Pith. A. pith of *Tibouchina kingii* Wurdack; B. pith of *Chaetolepis microphylla* (Bonpl.) Miq.; C. medullary vascular bundle of *Arthrostemma ciliatum* Pav. ex D. Don (FI, internal phloem; Flo, Phloem; HVM, medullary vascular bundle; Xil, xylem).

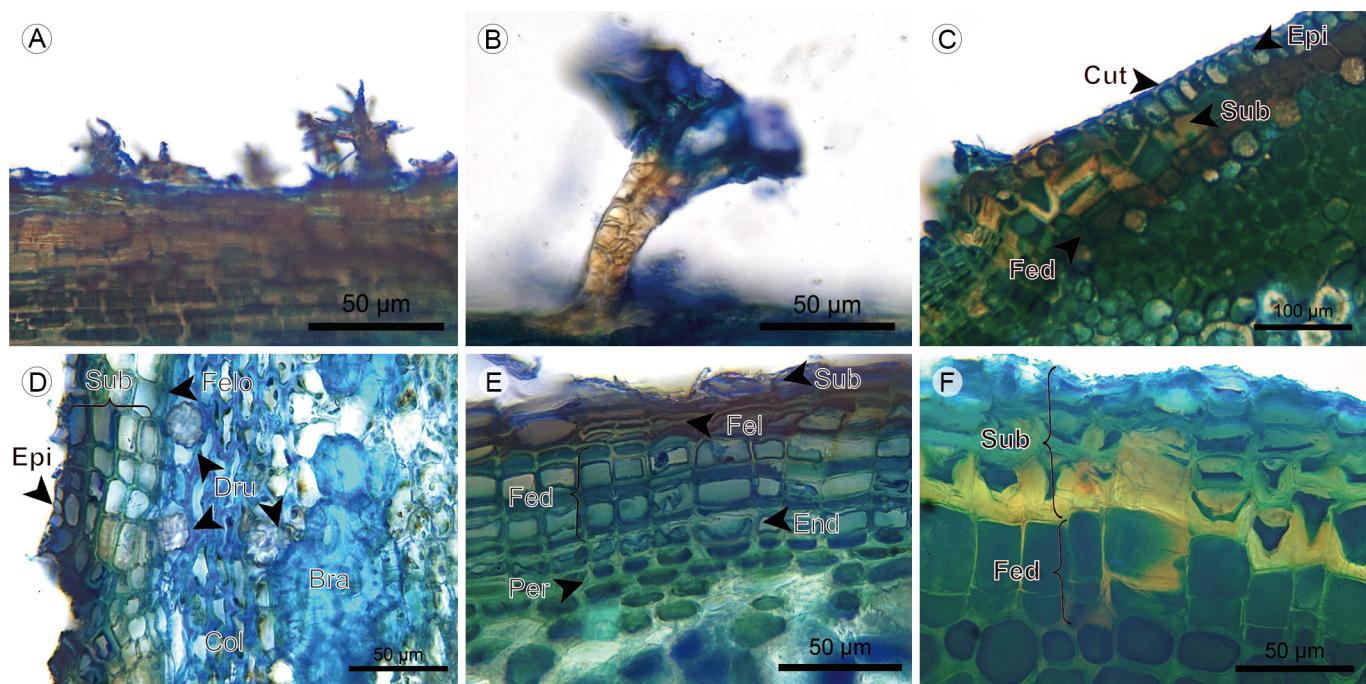


Figure 12: Lianas: Epidermis. A. dendritic hair of *Monochaetum multiflorum* (Bonpl.) Naudin; B. dendritic hair of *Blakea fasciculata* Gleason; C. epidermis and periderm beginning of *Blakea quadrangularis* Triana; D. epidermis, periderm and cortex of *Blakea fasciculata* Gleason; E. periderm, endodermis, and pericycle of *Monochaetum multiflorum* (Bonpl.) Naudin; F. periderm of *Blakea quadrangularis* Triana. (Bra, stone cell; Col, angular collenchyma; Cut, cuticle; Dru, druse; End, endodermis; Epi, epidermis; Fed, phellogen; Fel, phellogen; Per, pericycle; Sub, phellem).

tissue in the center of cortex, with simple and ramiform pits (Figs. 12D, E; 13; 14A, B).

Endodermis and pericycle: The endodermis is a continuous tissue between the cortex and the phloem. In *Monochaetum multiflorum* the endodermis cells show a Caspary strip with unequal thickening of cell walls and reduced lumen. In *Blakea quadrangularis* and *Blakea*

fasciculata the endodermis is subtle, formed by one layer of cells larger than the cortex parenchyma cells, with thin cellular walls and without cytoplasmic accumulations. The pericycle is one- to three-layered, only present in *Monochaetum multiflorum* and *Blakea quadrangularis*; in the latter the presence of solitary fibers with thick walls, reduced lumen and simple pits is common (Figs. 12E, 13A, 14A, B; 15A).

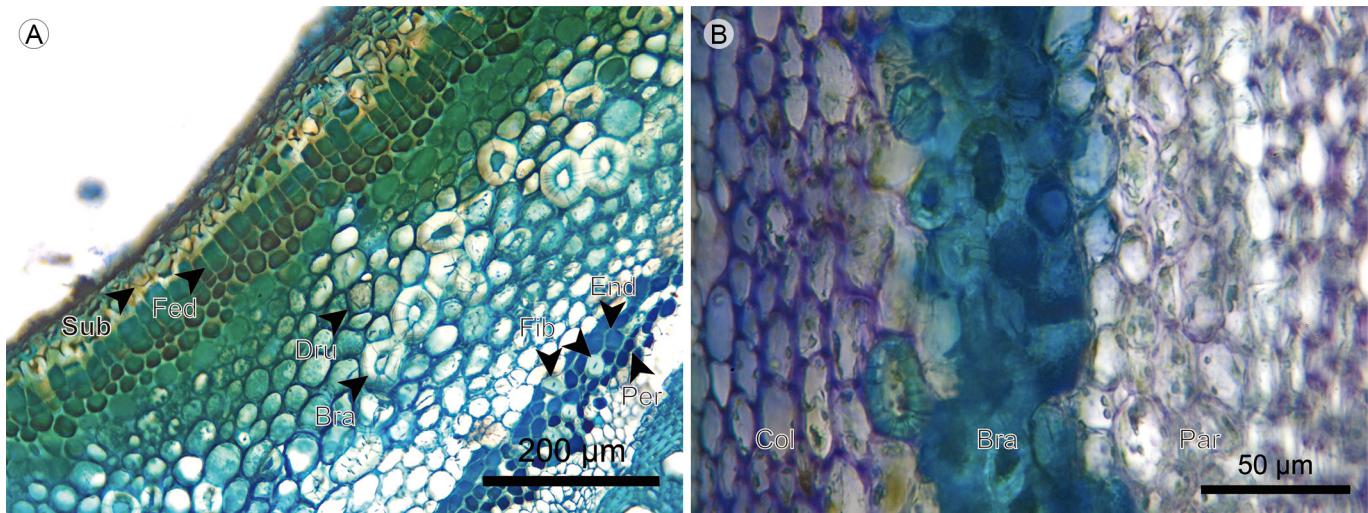


Figure 13: Lianas: Cortex. A. transverse section of *Blakea quadrangularis* Triana; B. cortex of *Blakea fasciculata* Gleason. (Bra, stone cell; Col, angular collenchyma; Dru, druse; End, endodermis; Fed, Pheloderm; Fib, fiber; Par, parenchyma; Per, pericycle; Sub, phellem).

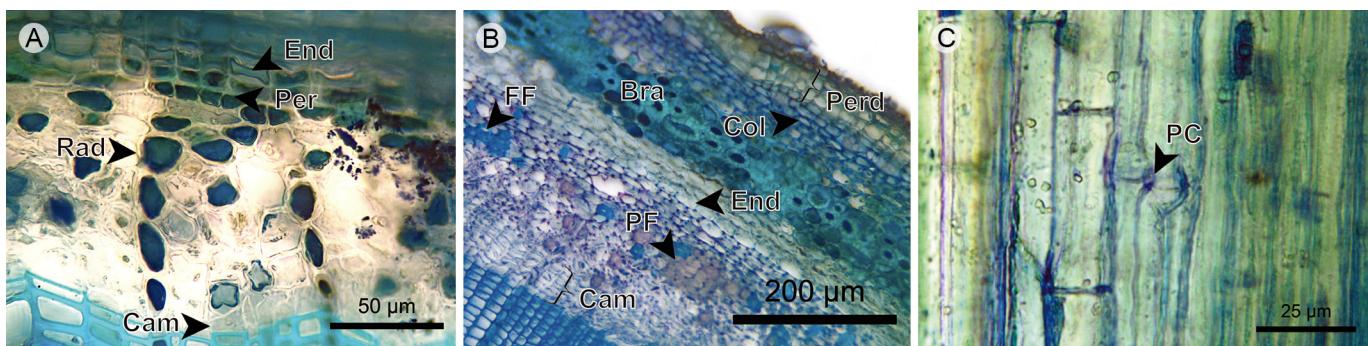


Figure 14: Lianas: Endodermis, pericycle, phloem, and vascular cambium. A. endodermis, pericycle, and phloem of *Monochaetum multiflorum* (Bonpl.) Naudin; B. transverse section of *Blakea fasciculata* Gleason; C. phloem in radial section of *Blakea quadrangularis* Triana. (Bra, stone cell; Cam, vascular cambium; Col, angular collenchyma; End, endodermis; FF, phloem fiber; PC, sieve plate; Per, pericycle; Perd, Periderm; PF, phloem parenchyma; Rad, ray).

Phloem: Sieve-tube elements scattered, with simple and transverse sieve plates. Phloem fibers present only in *Blakea fasciculata*, where they are scattered, mainly solitary, or rarely in clusters. Axial parenchyma in a strand of short cells with accumulation of druses, scattered in *Monochaetum multiflorum* and *Blakea quadrangularis*, and grouped in clusters in *Blakea fasciculata* (Figs. 13B, 14, 15A).

Vascular cambium: In the same way as trees, shrubs, and herbs, liana species show a vascular cambium lacking discontinuities, variations or irregularities that

may change the vascular system conformation. The vascular cambium of *Blakea fasciculata*, with eight layers of cells, is the thickest among the species studied here (Figs. 14, B, 15A).

Xylem vessels: The vessel arrangement has a radial to diagonal pattern, mainly solitary in *Monochaetum multiflorum* and *Blakea fasciculata*, and more common in multiples of two to three vessels in *Blakea quadrangularis*. The outline of solitary vessels is rounded in *Monochaetum multiflorum* and angular in *Blakea quadrangularis* and *Blakea fasciculata*. Perforation plates are simple

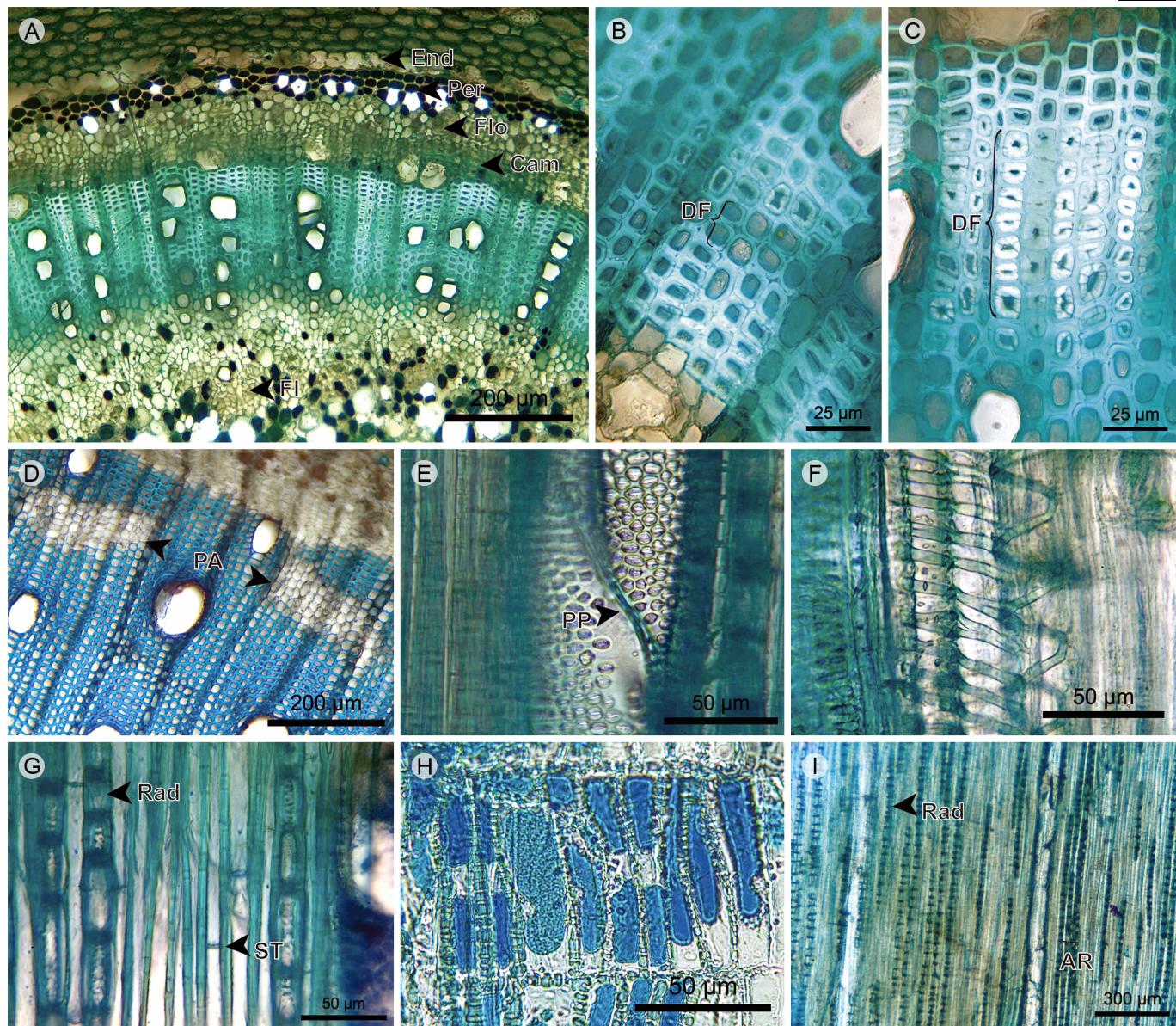


Figure 15: Liana: Xylem. A. transverse section of *Blakea quadrangularis* Triana; B. and C. two types of fiber dimorphism of *Blakea quadrangularis* Triana; D. xylem of *Blakea fasciculata* Gleason; E. vessels in radial section of *Blakea fasciculata* Gleason; F. protoxylem in radial section of *Blakea fasciculata* Gleason; G. xylem in tangential section of *Blakea fasciculata* Gleason; H. rays in radial section of *Monochaetum multiflorum* (Bonpl.) Naudin; I. xylem in tangential section of *Blakea fasciculata* Gleason. (AR, rays aggregates; Cam, vascular cambium; DF, fiber dimorphism; End, endodermis; FI, internal phloem; Flo, phloem; PA, axial parenchyma; Per, pericycle; PP, perforation plate; Rad, ray; ST, fiber transversal septum).

and inclined. Intervessel pits are bordered, from scalariform to opposite rounded in *Monochaetum multiflorum*, and alternate in *Blakea quadrangularis* and *Blakea fasciculata* (Figs. 15A, D-F).

Xylem fibers: The fibers are libriform and have transversal septa. Dimorphic fibers grouped in two- to

eight-layered tangential bands, with a wider lumen and thinner walls allowing intercellular spaces. Additionally, *Blakea quadrangularis* xylem shows a second type of dimorphic fiber, with a narrower lumen and thicker walls, grouped in two- to six-layered tangential bands (Figs. 15A-D, G).

Xylem axial parenchyma: The axial parenchyma is of the apotracheal type, grouped in three- to six-layered clusters close to vascular cambium. This tissue is absent in *Monochaetum multiflorum* (Fig. 15D).

Xylem rays: Rays are uniserial, but xylem of *Blakea fasciculata* also shows aggregates of two to four rays, which are formed by upright and square cells. Vessel-ray pits are rounded to oval and half-bordered, with border only in vessel lumen (Figs. 15G-I).

Pith: The periphery of the pith has internal primary phloem, forming a continuous band in *Blakea quadrangularis* and a discontinuous one in *Monochaetum multiflorum* and *Blakea fasciculata*. In the center, the pith is formed by parenchymatous cells with isodiametric shape, diverse materials, and accumulation of druses. Vascular bundles, which are scattered throughout the pith of prima-

ry growth, are concentric amphicribral in *Monochaetum multiflorum* and *Blakea quadrangularis*, and collateral in *Blakea fasciculata*. The pith of *Blakea quadrangularis* and *Blakea fasciculata* also shows fibers with a narrow lumen, thick walls, and simple pits; these fibers are located around the internal phloem and the vascular bundles (Fig. 16).

Vines

Epidermis: The trichomes in *Clidemia epiphytica* (Triana) Cogn. are simple unbranched, whereas they are simple unbranched and dendritic in *Adelobotrys adscendens* Triana. The epidermis is absent in the fifth internode of *Blakea granatensis* Naudin. The epidermis in other species is single-layered and has an epicuticular wax in form of granules (Figs. 17A, B).

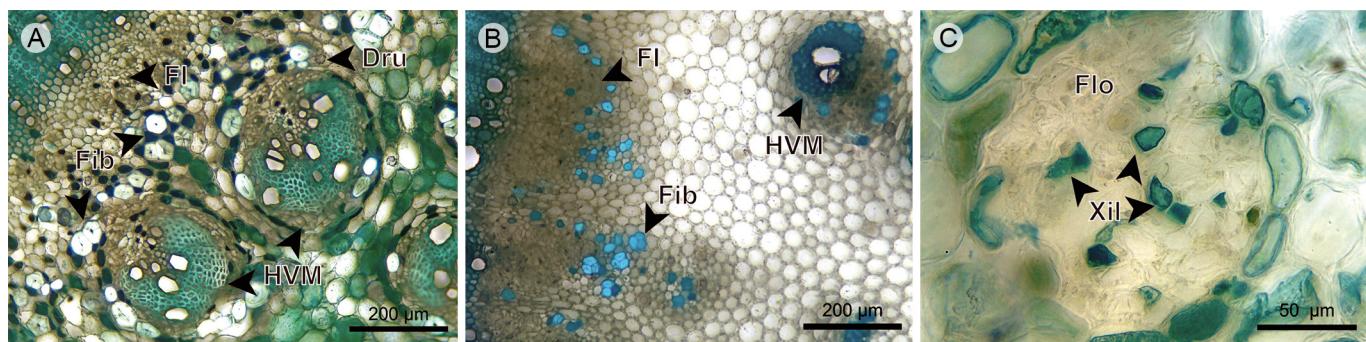


Figure 16: Liana: Pith. A. pith of *Blakea quadrangularis* Triana; B. pith of *Blakea fasciculata* Gleason; C. medullary vascular bundle of *Monochaetum multiflorum* (Bonpl.) Naudin. (Dru, druse; FI, internal phloem; Fib, fiber; Flo, phloem; HVM, medullary vascular bundle; Xil, xylem).

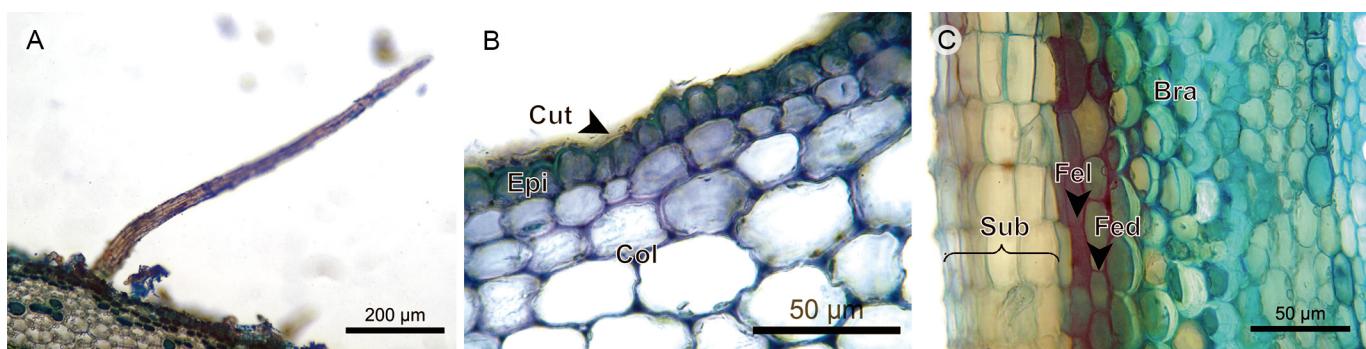


Figure 17: Vines: Epidermis and periderm. A. unbranched hair of *Clidemia epiphytica* (Triana) Cogn.; B. epidermis and cortex of *Adelobotrys adscendens* Triana; C. periderm and cortex of *Blakea granatensis* Naudin. (Bra, stone cell; Col, angular collenchyma; Cut, cuticle; Epi, epidermis; Fed, phellogen; Fel, phellogen; Sub, phellem).



Periderm: The periderm is absent in the fifth internode of *Adelobotrys adscendens*, whereas in *Clidemia epiphytica* the transition where the periderm is forming below the epidermis is observed. The periderm is single-layered, in *Blakea granatensis* it is a superficial and a continuous tissue, whereas in *Clidemia epiphytica* it is discontinuous. The phellem is one- to eight-layered with cells in radial disposition. The phellogen is one- to two-layered. The phellogen is one- to five-layered, with cells in radial disposition only in *Clidemia epiphytica* (Figs. 17C, 18A, B).

Cortex: The cortex is formed by parenchymatous cells with isodiametric shape, diverse materials, and accumulations of druses, with chloroplasts present in *Adelobotrys adscendens*. The collenchyma is of the angular type, forming a three- to four-layered continuous tissue below the epidermis in *Adelobotrys adscendens*. The stone cells form a four- to six-layered continuous tissue below the periderm in *Blakea granatensis*, and these have simple pits and tangential inner walls (in the direction of the pith) that are thicker than other walls (Figs. 18, 19A-C).

Endodermis and pericycle: The endodermis is a one- to two-layered continuous tissue; in *Adelobotrys adscendens* it is only formed by stone cells with simple and ramiform pits. The pericycle is a two- to three-layered continuous tissue; in *Blakea granatensis* it also consists of solitary fibers with simple pits and a narrow lumen.

In *Clidemia epiphytica* the cells between the cortex and phloem have no variation or specialization, except that they are smaller than cortical parenchyma cells and have no cytoplasmic accumulations (Figs. 17C, 18, 19A-C).

Phloem: Sieve-tube elements are scattered in *Blakea granatensis* and *Adelobotrys adscendens*, and in clusters in *Clidemia epiphytica*. Sieve plates are simple and inclined, transverse in *Clidemia epiphytica*. Fibers are scattered in the distal part of phloem in *Blakea granatensis*, and form a four- to six-layered continuous tissue on the distal part of phloem in *Clidemia epiphytica*. Axial parenchyma is formed by a strand of short cells containing druses (Figs. 18A, 19).

Vascular cambium: The vascular cambium is a two- to four-layered tissue. In this group of species, the vascular cambium has a sinuous form that is more prominent than in other species (Figs. 18A, B, 19).

Xylem vessels: The vessel arrangement is radial to diagonal and the grouping is in multiples of two to four, with a variable proportion of solitary vessels. Vessel diameters in *Blakea granatensis* show a marked increase on the newly formed xylem. The outline of solitary vessels is angular. Perforation plates are simple and inclined. Intervessel pits are rounded, bordered and have an alternate arrangement (Fig. 20).

Xylem fibers: The fibers are of the libriform type and have transversal septa. The dimorphic fibers are

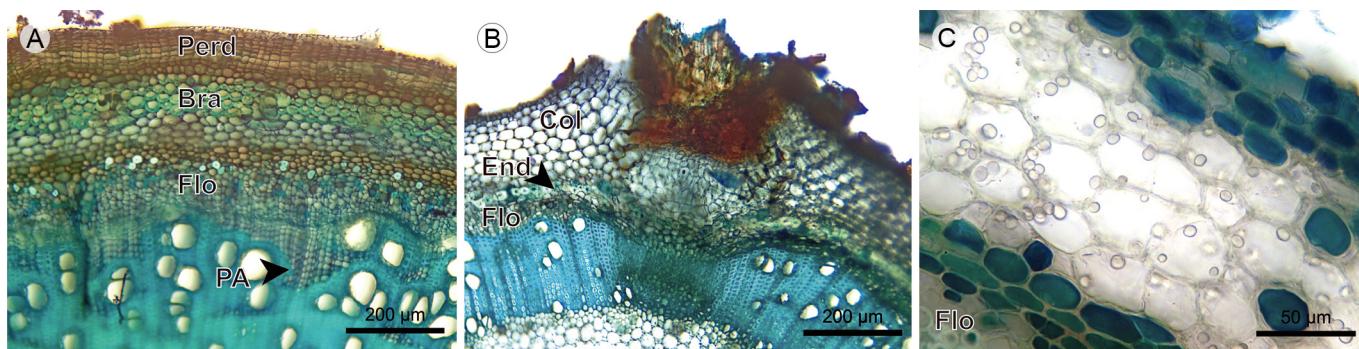


Figure 18: Vines: Cortex. A. transverse section of *Blakea granatensis* Naudin; B. adventitious root of *Adelobotrys adscendens* Triana; C. cortical parenchyma of *Clidemia epiphytica* (Triana) Cogn. (Bra, stone cell; Col, angular collenchyma; End, endodermis; Flo, phloem; Perd, periderm; PA, axial parenchyma).

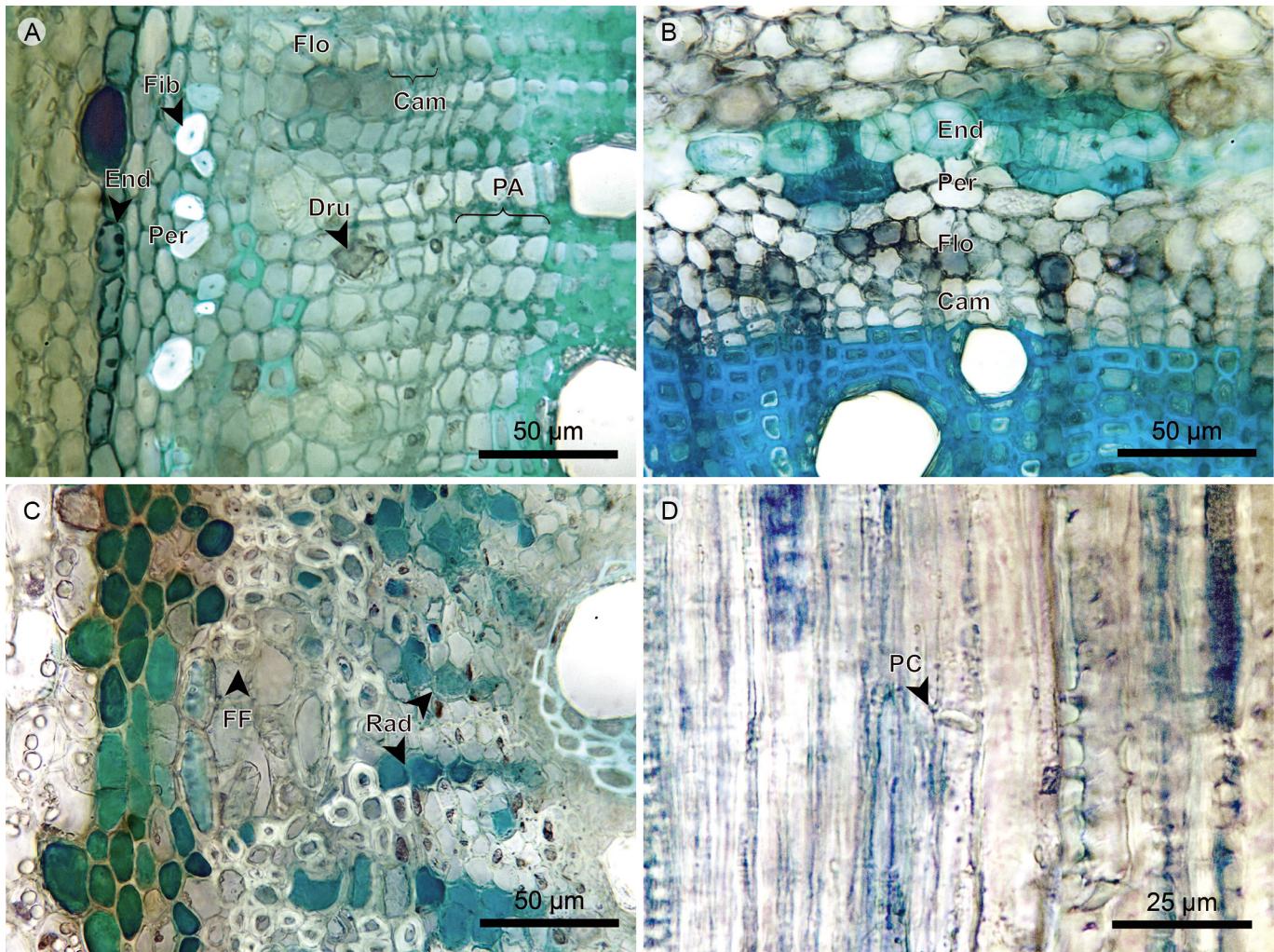


Figure 19: Vines: Endodermis, pericycle, phloem, and vascular cambium. A. transverse section of *Blakea granatensis* Naudin; B. transverse section of *Adelobotrys adscendens* Triana; C. phloem of *Clidemia epiphytica* (Triana) Cogn.; D. phloem in radial section of *Clidemia epiphytica* (Triana) Cogn. (Cam, vascular cambium; Dru, druse; End, endodermis; FF, phloem fiber; Fib, fiber; Flo, phloem; PA, axial parenchyma; Per, pericycle; PC, sieve plate; Rad, ray).

grouped in two- to five-layered tangential bands and the distance between the bands is larger than the distance between the rays. Like *Blakea quadrangularis*, *Adelobotrys adscendens* shows two different types of dimorphic fibers (Figs. 20A-C).

Xylem axial parenchyma: Axial parenchyma is present in *Blakea granatensis* only, where it is the apotracheal type, grouped in four- to seven-layered tangential bands, more frequently close to the vascular cambium. The cells can be fusiform and short in their strand, showing an accumulation of druses (Figs. 18A, 19A, 20F).

Xylem rays: Rays uniserial, formed by upright and square cells. The vessel-ray pits are rounded to oval and half-bordered, with a border in the vessel lumen only (Figs. 20E-G).

Pith: The internal phloem is reduced in *Adelobotrys adscendens* and discontinuous in *Blakea granatensis*. The parenchyma cells have isodiametric shape and diverse materials and druses accumulation. Vascular bundles, which are scattered throughout the pith, are concentric amphicribal in *Adelobotrys adscendens* and *Clidemia epiphytica* (this one with a very reduced xylem), and just

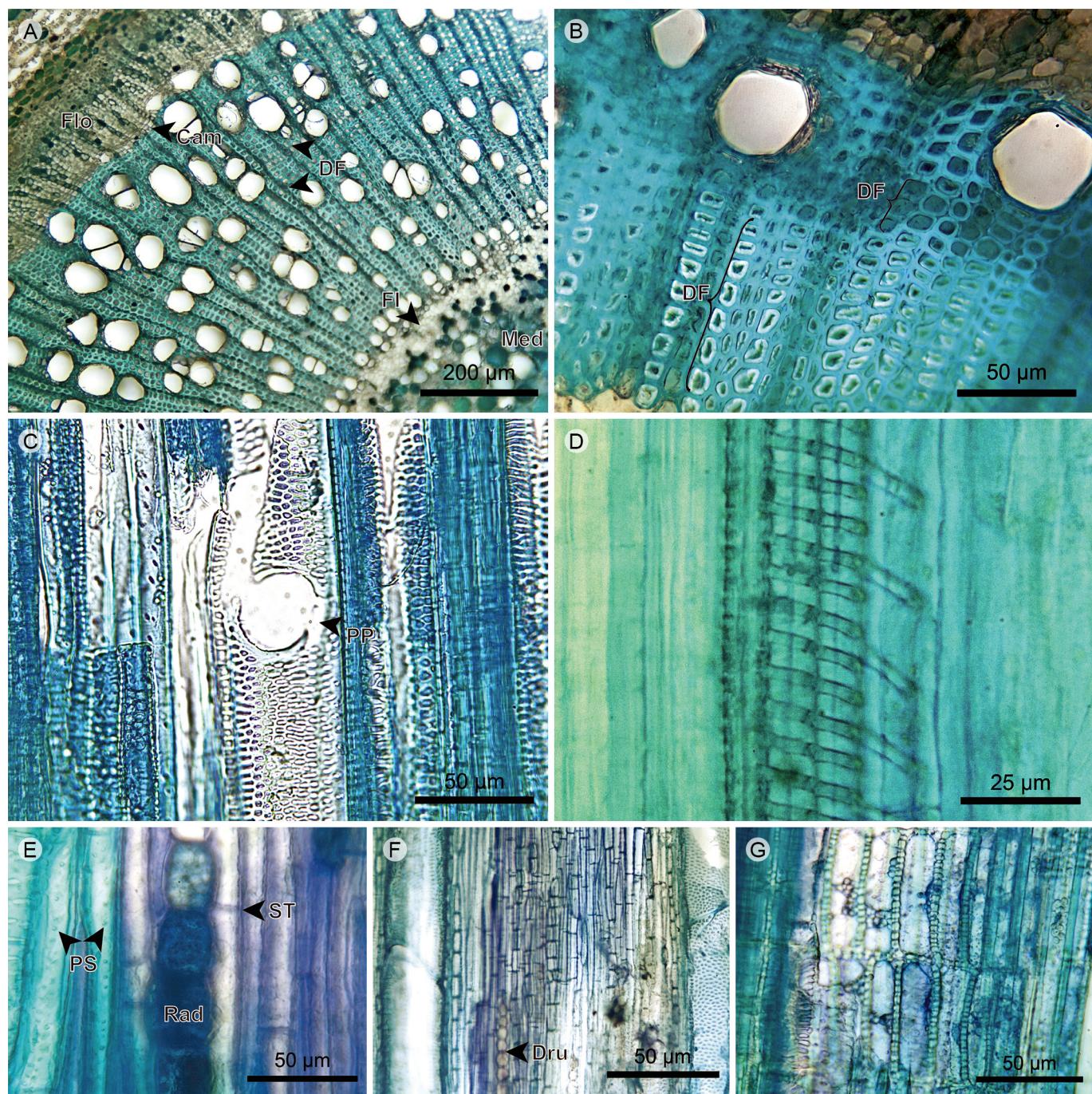


Figure 20: Vines: Xylem. A. Transverse section of *Clidemia epiphytica* (Triana) Cogn.; B. xylem of *Adelobotrys adscendens* Triana; C. xylem in radial section of *Clidemia epiphytica* (Triana) Cogn.; D. protoxylem in radial section of *Blakea granatensis* Naudin; E. xylem in tangential section of *Adelobotrys adscendens* Triana; F. axial parenchyma in tangential section of *Blakea granatensis* Naudin; G. rays in radial section of *Clidemia epiphytica* (Triana) Cogn. (Cam, vascular cambium; DF, fiber dimorphism; Dru, druse; FI, internal phloem; Flo, phloem; Med, pith; PP, perforation plate; PS, fiber simple pit; Rad, ray; ST, fiber transversal septum).

amphicribral in *Blakea granatensis*. *Blakea granatensis* pith has fibers with a narrow lumen, thick walls, and simple pits; these fibers are found around the internal phloem and vascular bundles (Fig. 21).

Hemiepiphytes

Epidermis: The epidermis is absent in the fifth internode of *Blakea rosea* (Ruiz & Pav.) D. Don. In *Blakea watsonii* (Cogn.) Penneys & Almeda, the epidermis is single-layered and shows small cells. The developing periderm is located just below the epidermis, and trichomes are absent (Fig. 22A).

Periderm: The periderm is superficial, a single layer of continuous tissue. The phellem is one-layered in *Blakea rosea*, and two- to five-layered with scattered stone cells in *Blakea watsonii*. The phellogen is a one- to three-layered tissue of thin cells. The phellogen cells have radial disposition; in *Blakea watsonii* it has one to two layers that accumulate druses; in *Blakea rosea* the number of layers varies from three to 15 layers, across the different regions of the transverse section (Figs. 22, 23).

Cortex: The cortex is mainly formed by parenchymatous cells with isodiametric shape, druses are accumu-

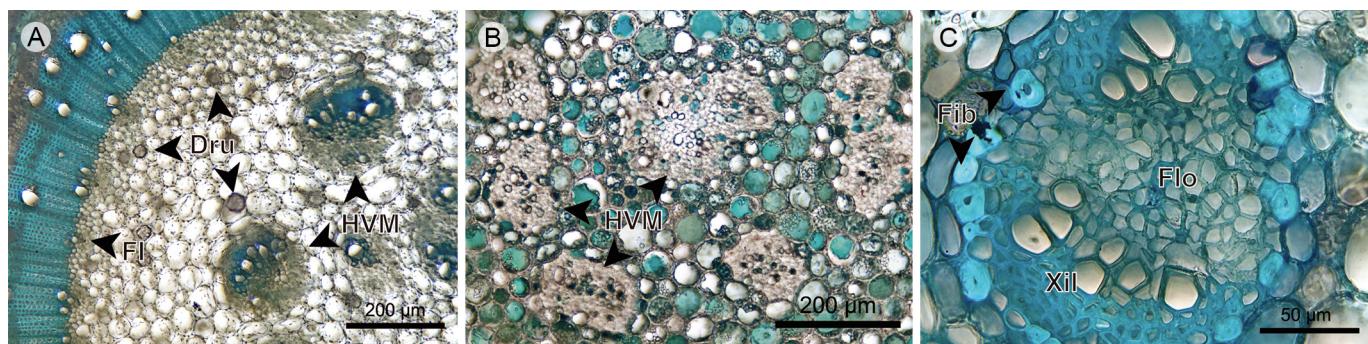


Figure 21: Vines: Pith. A. pith of *Adelobotrys adscendens* Triana; B. pith of *Clidemia epiphytica* (Triana) Cogn.; C. medullary vascular bundle of *Blakea granatensis* Naudin. (Dru, druse; FI, internal phloem; Fib, fiber; Flo, phloem; HVM, medullary vascular bundle; Xil, xylem).

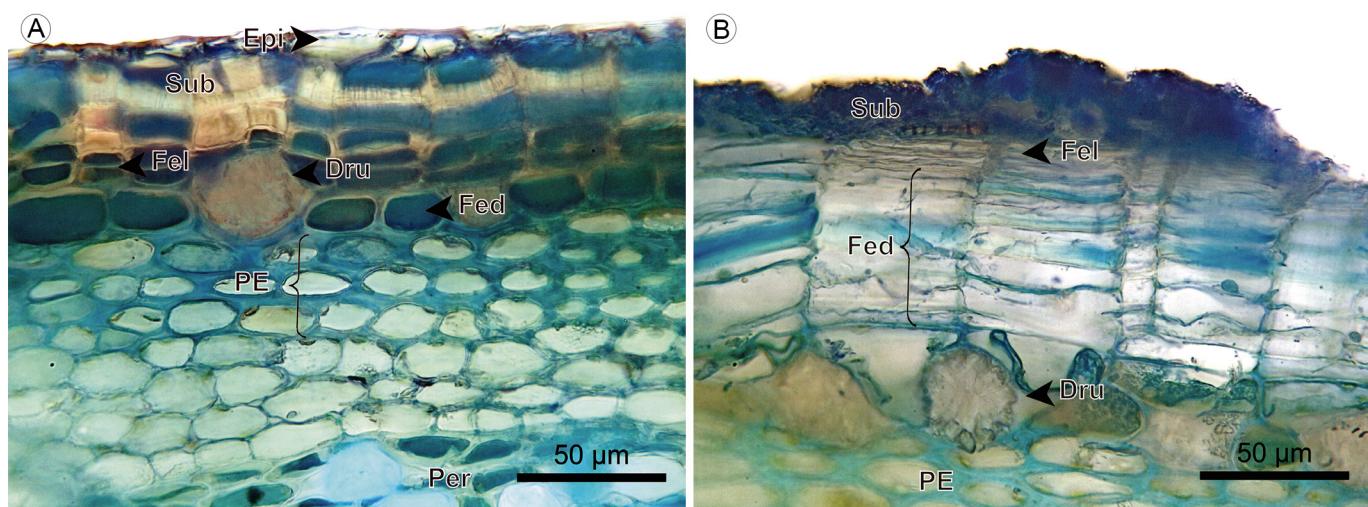


Figure 22: Hemiepiphytes: Epidermis and periderm. A. transverse section of *Blakea watsonii* (Cogn.) Penneys & Almeda; B. periderm of *Blakea rosea* (Ruiz & Pav.) D. Don. (Dru, druse; Epi, epidermis; Fed, phellogen; Fel, phellem; PE, sclerotic parenchyma; Sub, suberized tissue).

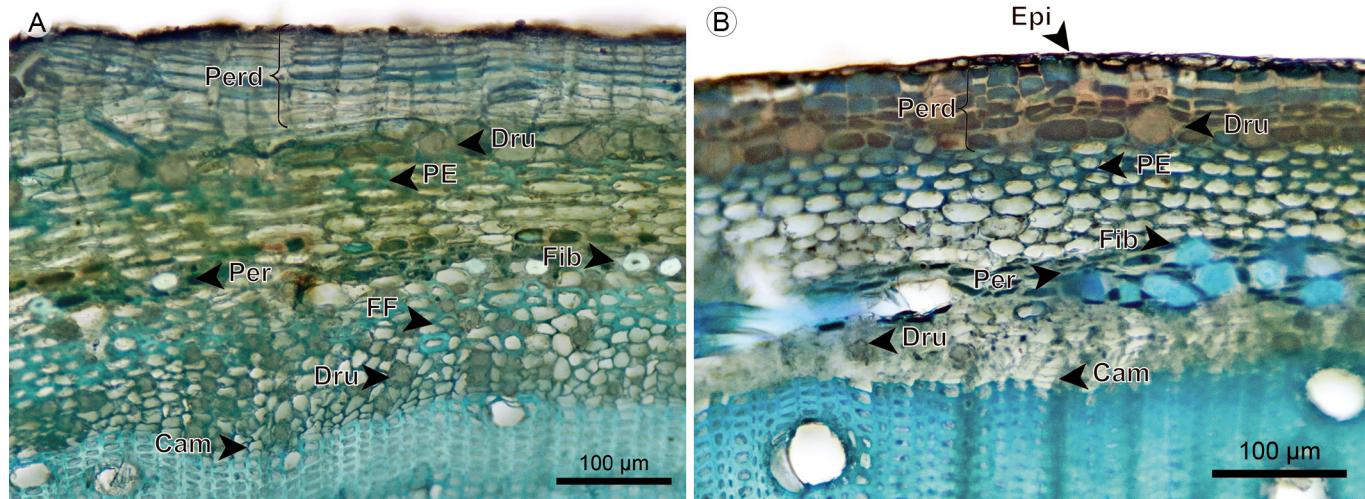


Figure 23: Hemiepiphytes: Cortex. A. transverse section of *Blakea rosea* (Ruiz & Pav.) D. Don.; B. transverse section of *B. watsonii* (Cogn.) Penneys & Almeda (Cam, vascular cambium; Dru, druse; Epi, epidermis; FF, phloem fiber; Fib, fiber; PE, sclerotic parenchyma; Perd, periderm).

lated only in *Blakea rosea* in a layer below the phellogen. Only in these two species, a sclerotic parenchyma forms a two- to four-layered continuous tissue below the phellogen (Figs. 22, 23).

Endodermis: In these two species, the pericycle was not observed. The limit between the cortex and the phloem is only taken up by the endodermis, having two- to six-layered tissues with cells smaller than cortical parenchyma and without cytoplasmic accumulations. Scattered fibers with thick walls, narrow lumen and simple pits are present (Figs. 22A, 23).

Phloem: Sieve-tube elements have a radial disposition in *Blakea rosea*, and are scattered in *Blakea watsonii*. Sieve plates are simple and transverse. Fibers in *Blakea rosea* form a one- to four-layered continuous tissue close to the endodermis, in *Blakea watsonii* the presence is rare. Axial parenchyma is present in a strand of short cells, some containing druses, more commonly close to the endodermis (Figs. 23A, 24).

Vascular cambium: The vascular cambium is a one- to three-layered continuous and uniform tissue of small cells that have walls with irregular shape (Fig. 24A, B).

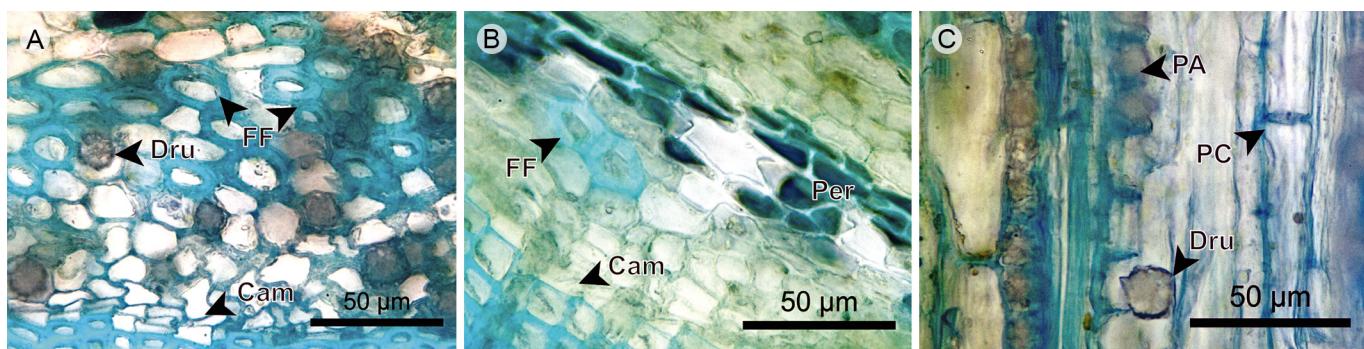


Figure 24: Hemiepiphyte: Endodermis, phloem, and vascular cambium. A. phloem of *Blakea rosea* (Ruiz & Pav.) D. Don.; B. phloem of *Blakea watsonii* (Cogn.) Penneys & Almeda; C. phloem in radial section of *Blakea rosea* (Ruiz & Pav.) D. Don. (Cam, vascular cambium; Dru, druse; FF, phloem fiber; PA, axial parenchyma; PC, sieve plate).

Xylem vessels: The arrangement of vessels has a radial to diagonal pattern and the grouping is in multiples of two to three, or more commonly solitary. The outline of solitary vessels is angular. Perforation plates are simple and inclined. Intervessel pits are rounded, bordered and have an alternate arrangement (Figs. 25A-D).

Xylem fibers: The fibers are of the libriform type and have transversal septa. *Blakea watsonii* xylem shows just one type of dimorphic fibers, and *Blakea rosea* show two, in the same way that occurs in *Blakea quadrangularis* and *Adelobotrys ascendens*, respectively. The tangential bands of dimorphic fibers have two to three layers and the distance between bands is larger than the distance between rays (Figs. 25A-C, E).

Xylem axial parenchyma: The axial parenchyma is of the apotracheal type and formed by strands of short cells. In the xylem of *Blakea watsonii*, the axial parenchyma forms a tangential five-layered band of cells that contain druses and have irregular and wide lumens. In *Blakea rosea* the axial parenchyma cells are scarce, isolated, and containing prismatic crystals (Figs. 25A, E).

Xylem rays: The rays are uniserial, formed by upright and square cells. Vessel-ray pits are rounded to oval and half-bordered, with border only in the vessel lumen (Figs. 25E, F).

Pith: The pith periphery incorporates a continuous tissue of internal phloem. In the center, it is formed by parenchymatous cells that show isodiametric shape and accumulation of druses. Vascular bundles are concentric and amphicribral, with primary growth, and are scattered throughout the pith. Fibers show a narrow lumen, thick walls, and simple pits, and are located around the internal phloem and the vascular bundles (Fig. 26).

Epiphytes

This group only contains *Monolena primuliflora* Hook. f. The stem is composed by a wide periderm that gives it external support. Below the periderm, the stem is almost exclusively formed by parenchyma and a few narrow vessels from the adventitious roots. Therefore, for this

species, it was not possible to delimit a cortex, vascular system, or pith in the inner area (Fig. 27A).

Periderm: The periderm is superficial and single-layered with lenticels. The phellem is five- to eight-layered with cells in radial disposition. Phellogen is one-layered with thin cells. The pheloderm is eight to 12-layered with cells containing druses, without radial disposition (Figs. 27B, C).

Parenchyma: Composed by large isodiametric cells that show little accumulation of material and lack intercellular spaces. In this inner tissue, there is neither growth direction, nor determined cellular arrangement (e.g. an endodermis). In the parenchyma, except for the xylem, no other cellular type or tissue is present (Figs. 27B-F).

Xylem: The xylem has primary growth and is formed by few narrow and short vessels. These vessels show helical thickenings that sometimes are reticulated, and reach from the adventitious roots base to the center of the stem and along the periphery below the periderm. The vessels split along their trajectory (Figs. 27C-F).

Summary of anatomical differences among growth habits

The stem anatomy characters that distinguished Melastomataceae growth habits are summarized here:

A higher proportion of vessels in radial multiples than single ones among trees/shrub (Figs. 5A, B) and vine species (Fig. 20A).

A higher proportion of single vessels than grouped ones among herbs (Figs. 10A, B), lianas (Fig. 15A-D), and hemiepiphytes (Figs. 25A, B).

Xylem with shorter vessels and without rays (except for *Tibouchina kingii*) among herb species (Fig. 10).

A higher density of xylem vessels and vascular cambium in bands with sinuous form among vine species (Figs. 19A-C; 20A-C, F).

More rows in bands of dimorphic xylem fibers and adventitious roots among liana (Figs. 15A-D) and vine species (Figs. 18B, 20A, B).

Discontinuous intraxylary phloem among herb (Figs. 8A, 10A, 11A) and vine species (Figs. 18B, 20A, 21A).

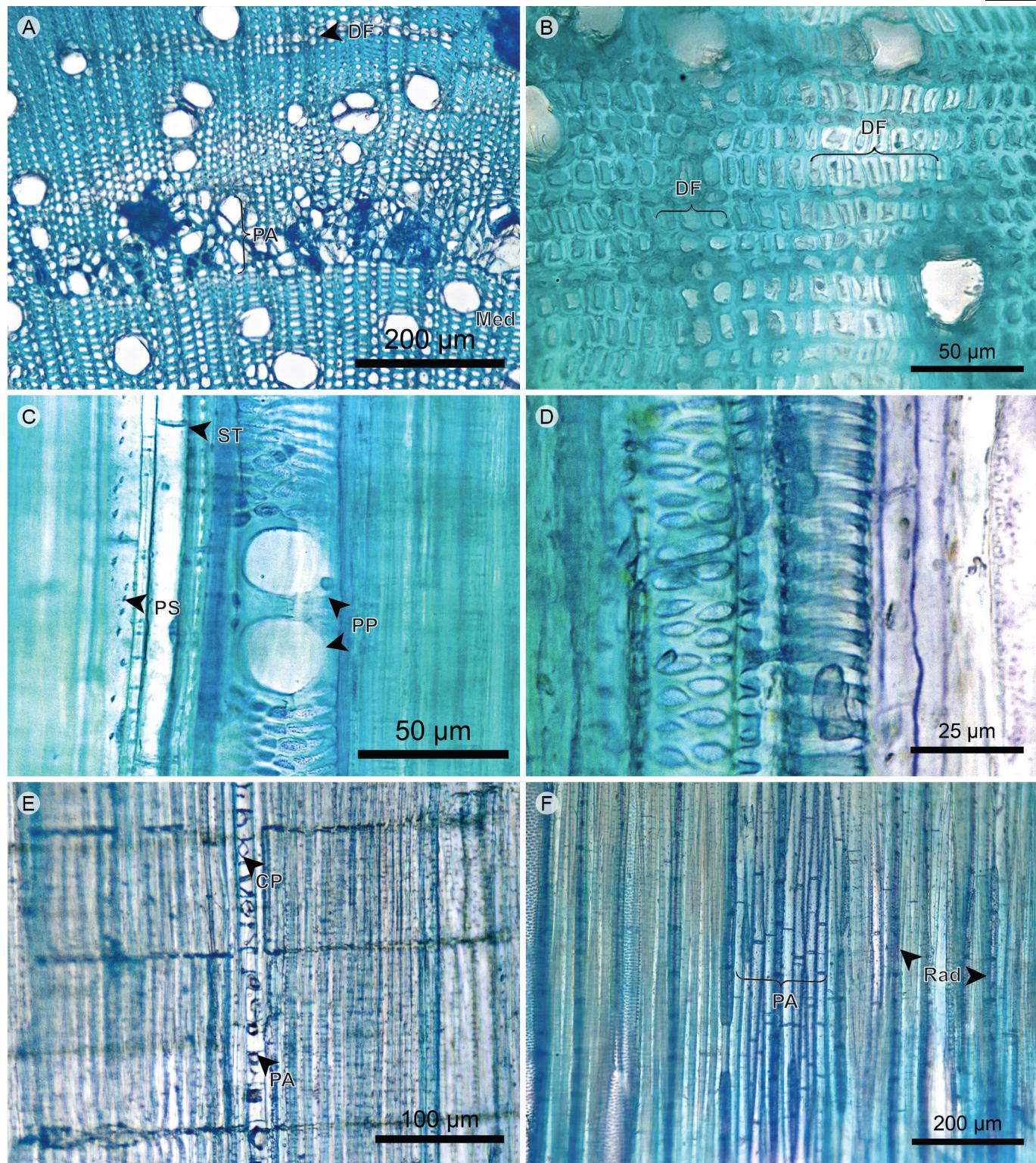


Figure 25: Hemiepiphyte: Xylem. A. xylem of *Blakea watsonii* (Cogn.) Penneys & Almeda; B. xylem of *Blakea rosea* (Ruiz & Pav.) D. Don.; C. xylem in tangential section of *Blakea watsonii* (Cogn.) Penneys & Almeda; D. protoxylem in radial section of *Blakea watsonii* Cogn.; E. xylem in radial section of *Blakea rosea* (Ruiz & Pav.) D. Don.; F. xylem in tangential section of *Blakea watsonii* (Cogn.) Penneys & Almeda (CP, prismatic crystal; DF, fiber dimorphism; PA, axial parenchyma; PP, perforation plate; PS, fiber simple pit; Rad, ray; ST, fiber transversal septum).

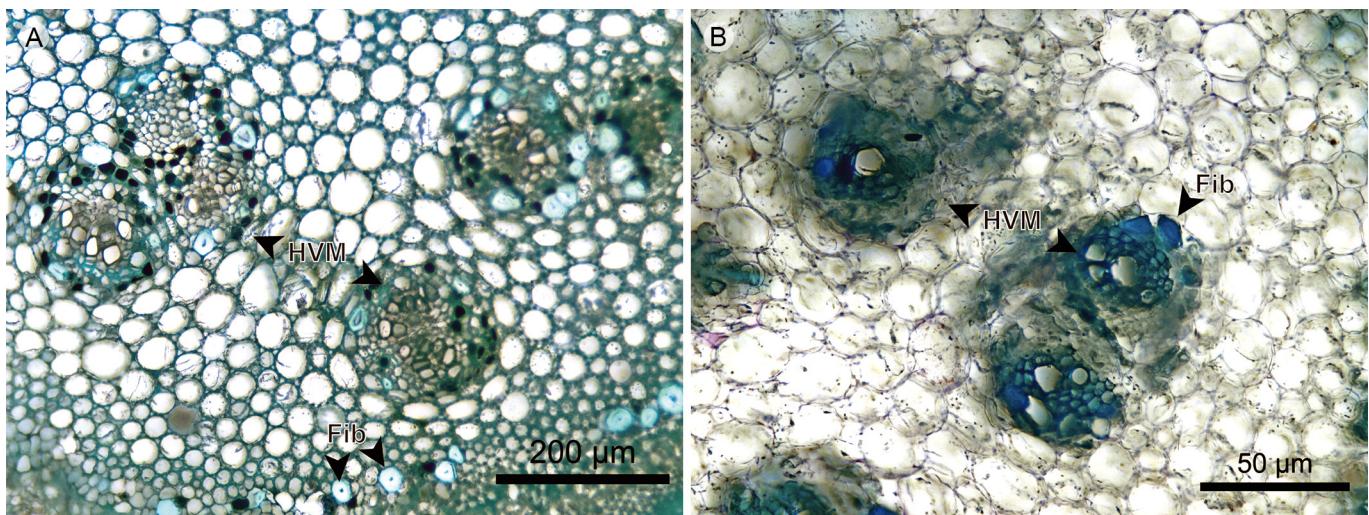


Figure 26: Hemiepiphytes: Pith. A. pith of *Blakea rosea* (Ruiz & Pav.) D. Don.; B. pith of *Blakea watsonii* (Cogn.) Penneys & Almeda (Fib, fiber; HVM, medullary vascular bundles).

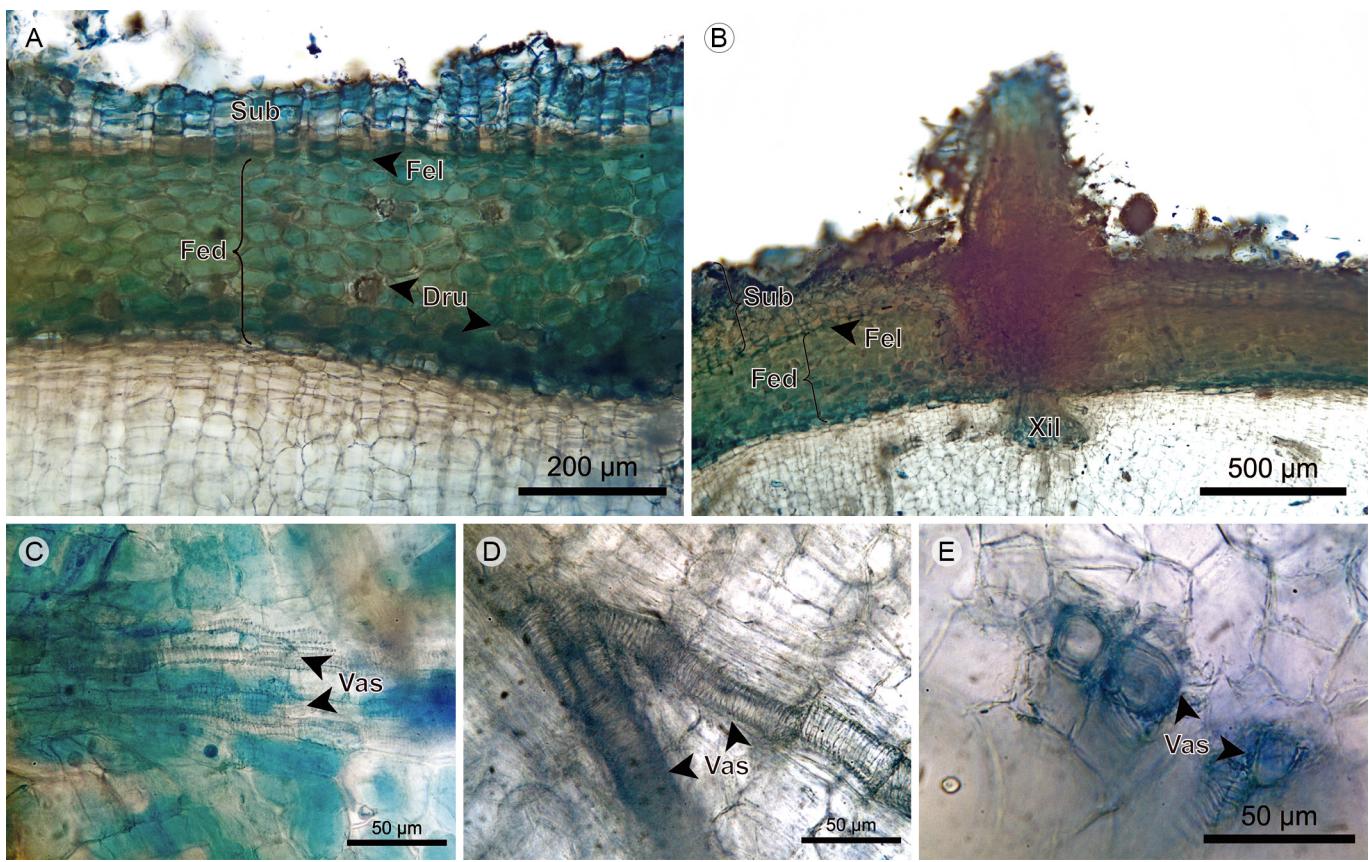


Figure 27: Epiphytes: Transverse section of *Monolena primuliflora* Hook. f. A. periderm; B. adventitious root; C. and D. xylem in longitudinal section; E. xylem in transverse section. (Dru, druse; Fed, phellogen; Fel, phellogen; Sub, phellem; Vas, vessel; Xil, xylem).



Sclerotic parenchyma acting like support elements in the cortex of hemiepiphyte species (Figs. 22, 23).

Angular collenchyma acting like support elements in the cortex of species without periderm (trees: *Meriania nobilis*, *Graffenrieda emarginata*, Figs. 2C, D, 3A-C; herbs: *Arthrostemma ciliatum*, *Heterocentron subtriplinervium*, Figs. 8A, 9B; vines: *Adelobotrys adscendens*, Figs. 17B, 18B, 19B).

Quantitative differences among growth habits

Growth habits differed in the tangential diameter of solitary vessels ($p<0.001$; tree/shrub>herbs=vines; Fig. 28A), the number of vessels per mm^2 varied among growth habits ($p<0.001$; vines>hemiepiphyte=tree/shrub=herb>liana; Fig. 28B) and the vessel axial length ($p<0.001$; hemiepiphyte>three/shrub=liana=vine>herb; Fig. 28C). Species was a significant factor in the three ANOVAs ($p<0.001$).

DISCUSSION

Botany practitioners continuously describe specimens using a range of growth habit terms, and yet the application of these can sometimes obscure rather than clarify the portrayal of a specimen. If growth habits are a natural depiction of a morphological pattern in plants, it can be safely assumed that anatomy is one of the character types that can be used to support classification of any group of plants into different growth habits. We described here 18 species of Melastomataceae using 125 anatomical characters of their stems. These plants were classified using external morphology into an initial classification of growth habits that we compiled from the literature. This preliminary classification was not very different from others previously proposed and we are not suggesting that it is better than any other; it was rather a tool for the further analysis. Our main finding was that stem anatomy did not support all the growth habits that we started with. Some stem anatomy characters were observed in species of some growth habits, but there were very few characteristics exclusively shared by species of

one particular growth habit. However, the species analyzed here could be grouped in fewer growth habits, which we hereafter describe and discuss.

Based on the anatomical study of 18 species of Melastomataceae, we recognize only three growth habits: trees/shrubs, herbaceous species, and climbers. Trees and shrubs grow their entire life cycle rooted in the soil, with erect orientation, self-supporting their weight with stems that have secondary growth (Font-Quer, 2001; Simpson, 2006). Most species of Melastomataceae belong to this group. One source of confusion about this growth habit in general is its inherent gradient of size, from suffrutex (a tall herb with woody stem base; Mendoza and Ramírez, 2006), to shrubs (in range from 2 to 5 m of height), to treelets (“small stature trees”; Opler et al., 1980), to trees growing up to 40 m high, like some *Tessmannianthus* Markgr. species. We observed in Melastomataceae that the stems of plants with sizes across this gradient share a structural design and cannot be distinguished using stem anatomy. We suggest that species that in the initial classification were labeled as herbs (*Tibouchina kingii*, *Arthrostemma ciliatum*, *Chaetolepis microphylla*, and *Heterocentron subtriplinervium*) and hemiepiphytes (*Blakea rosea* and *Blakea watsonii*) should be considered members of the trees/shrubs growth habit. These species not only showed an anatomy that was not distinguishable from trees/shrubs, they also develop secondary xylem, a characteristic absent from aerial stems of herbs (Eiten 1991; Font-Quer 2001). Therefore, among the 18 species studied here, only *Monolena primuliflora* should be catalogued as a herb because its stem does not develop organized inner tissues such as cortex, vascular system, pith, or secondary xylem.

We found that hemiepiphytes cannot be distinguished from trees/shrubs using stem anatomy characters. Hemiepiphytic species can be seen as shrubs accidentally sprouting and growing on a phorophyte. The definition of epiphytes and hemiepiphytes depends mainly on habitat (position above ground), and not on their habit (Simpson, 2006). It is confusing that the aspect of the same plant can be described with one term if it reached the ground and

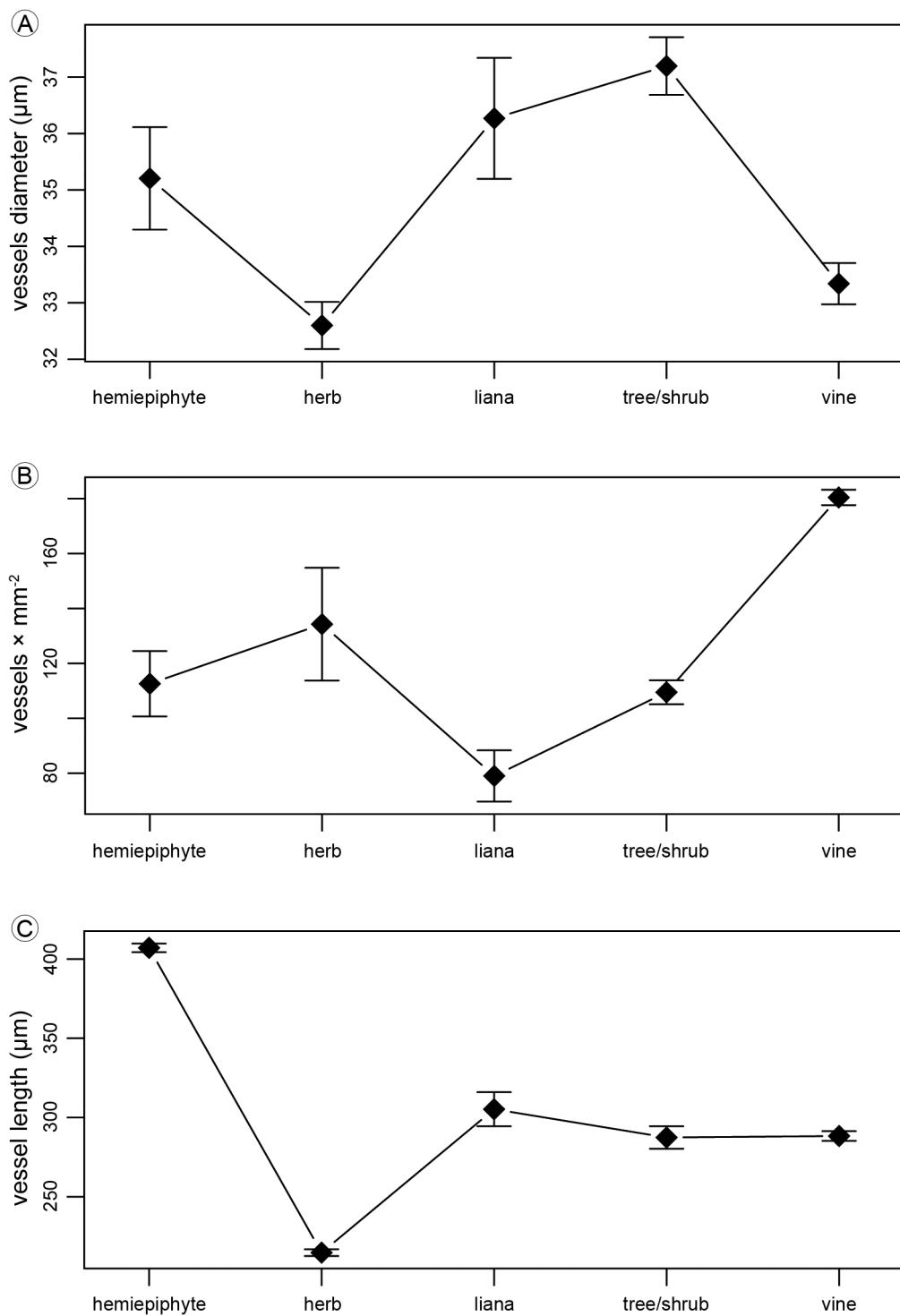


Figure 28: Vessel measurements. A. tangential diameter of vessel lumen in transverse section (values in μm); B. number of vessels per mm^2 in transverse section; C. axial length of vessels (values in μm).



with another if it did not. The inconsistency of this classification is clear in cases when an “epiphyte” is found growing on the ground (“facultative epiphytism”; Renner, 1993) or when a shrub is found thriving on the axial concavity of a tree branch (“accidental epiphytism” (Zotz, 2005)).

Climbers are plants that require holding on a phorophyte to grow vertically. Lianas are climbers that can grow early in life in a self-supported manner, whereas vines always need the support of phorophytes to ascend (Rowe and Speck, 2004; Isnard et al., 2012). Vines are thin and flexible climbers, showing a xylem with more area occupied by vessels and fewer fibers (Ter Welle and Koek-Noorman, 1981; Araque et al., 2000; Fig. 28B). On the other hand, lianas reach altitude through surrounding vegetation, but they can grow some distance self-supported as if they were shrubs. In comparison to vines, lianas show thicker stems, cortical support tissues with more layers, and a xylem occupied by more fibers and fewer vessels. However, these differences in their stem anatomy are not so pronounced as to allow a clear separation of these two habits in Melastomataceae. Instead, climber plants show a gradient of robustness, from thin vines as *Clidemia epiphytica* to the woody lianas of the genus *Blakea* P. Browne; a thicker stem diameter increases stem stiffness in climbers (Isnard et al., 2012).

There is more than one way to be a climber. We found here that the characteristics shared only by the climber species of Melastomataceae were the presence of adventitious roots and more rows in bands of dimorphic xylem fibers. Adventitious roots are a typical character of climbing plants (Putz, 1984). The dimorphic xylem fibers bands have a variable composition of ordinary libriform fibers, dimorphic fibers (with a thinner wall, wider lumen, and different cell length), fusiform parenchyma cells (axially extended) and parenchyma strands (two-eight cells high; Ter Welle and Koek-Noorman 1978). This band occurs in about half of the genera of Neotropical Melastomataceae (where it is called pseudoparenchyma; Ter Welle and Koek-Noorman, 1981) and in a large proportion of Paleotropical Melastomataceae (where it is called

deviating fibers; Vliet, 1981). The presence of more rows in the bands of dimorphic fibers, an increment in axial parenchyma, is related to an increased flexibility characteristic of climber stems (Koek-Noorman et al., 1979; Ter Welle and Koek-Noorman, 1981; Vliet, 1981; Carlquist, 1985).

Other stem characteristics that have been reported as typical for climbers are the presence of auxiliary elements that prevent embolism (vessel dimorphism, narrow vessels, tracheids and vasicentric tracheids) and elements that allow mechanical flexibility (anomalous growth, cambial variants, paratracheal parenchyma, and wide rays; Carlquist, 1985; Angyalossy et al., 2011, 2014; Isnard et al., 2012; Quijano-Abril et al., 2013). However, climbers studied here did not share such traits. The diversity of functional strategies to climb has been recognized before by multiple authors (Putz 1984; Moffet, 2000; Rowe and Speck, 2004; Simpson, 2006). Climbers can have tendrils, which can be adhesive, they can twine with their main stem or with some branches, or they can sprawl leaning on rocks or trees without attaching to them (Putz, 1984). All these different morphological adaptations to climb could be translated into growth habit names, but that would be an unnecessary proliferation of terms. For the concept of growth habits to be useful it should be avoided to elevate specific functional morphology adaptations to the generalized category of growth habit.

Our study has two caveats that should be addressed in future analyses. Our definition of growth habits in Melastomataceae could change if more characters are used to compare species from apparently different growth habits. That is, a more accurate distinction of growth habits could be obtained with the use of additional characters that describe plant morphology, architecture, or biomechanics (Rowe and Speck, 2004; Isnard et al., 2012). The second issue is that we did not take into account the underlying evolutionary hierarchy when comparing the species of this study. Some characteristics have a phylogenetic component that we did not address here. The species of *Blakea* have characteristic longer

vessels (Figs. 15E, F, I, 20F, 25C, F), tangential bands exclusively composed by apotracheal parenchyma (Figs. 15A, D, 18A, 25A, E, F), prismatic crystals and druses in xylem parenchyma (Figs. 20F, 25E), and presence of fibers with very lignified walls and reduced lumen being part of pericycle, surrounding the intraxylary phloem and medullary vascular bundles (Figs. 13A, 14B, 15A, 18A, 19A, 21C, 23, 26). The species of the tribe Melastomeae (*Tibouchina lepidota*, *Tibouchina kingii*, *Arthrostemma ciliatum*, *Heterocentron subtriplinervium*, *Chaetolepis microphylla* and *Monochaetum multiflorum*) have cortical vascular bundles (Figs. 3B, 8A, B) and thin vascular cambium with few rows of little cells (Figs. 3B, 4B, 8A, C, 9A, B, 10A, 14A). The species of *Tibouchina* (and *Monochaetum multiflorum*) have scaly bark and have a very simplified cortex or even lack it (Figs. 3B, 7E, 12E). The species of the Merianieae tribe (*Meriania nobilis*, *Graffenrieda emarginata*, and *Adelobotrys adscendens*) show angular collenchyma (Figs. 2C, D, 3A, C, 17B, 18B, 19B) and sclereids (Figs. 3A, 18B, 19B) that act like support elements in their cortex. The species of *Blakea* and *Miconia* Ruiz & Pav. (and *Clidemia epiphytica*) have fibers in their phloem (Figs. 4C, 13A, 14B, 15A, 18A, 19A, C, 20A, 23, 24A, B). These characters may have taxonomic value (Ter Welle and Koek-Noorman, 1978; Koek-Noorman et al., 1979; Ter Welle and Koek-Noorman, 1981; Vliet, 1981; Vliet et al., 1981), to identify some lineages of the family, independently of their growth habit.

AUTHOR CONTRIBUTIONS

JP and MQA contributed to the conception of this work. JP performed fieldwork and anatomical descriptions. JP, MT, and MQA made analyses and wrote the manuscript. All authors contributed to the discussion, review, and approval of the final manuscript.

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APPENDIX: Anatomical characters used in this study, including characters states for each of them.

TRANSVERSE SECTION

Epidermis

1. Epidermis stratification: (0) one cell row, (1) two cell rows, (2) more than two cell rows.
2. Intercellular spaces among epidermal cells: (0) absent, (1) present.
3. Trichomes number of cells: (0) unicellular, (1) pluricellular.
4. Trichome form: (0) glandular, (1) filiform, (2) lepidote.
5. Trichome branching: (0) absent, (1) branched.
6. Type of trichome branch: (0) dendritic, (1) stellate.
7. Trichome inclination: (0) erect, (1) sub-erect
8. Epidermal idioblasts: (0) absent, (1) present.
9. Cuticle thickness: (0) cuticle width smaller than epidermal cell, (1) cuticle width same as epidermal cell, (2) cuticle width higher than epidermal cell.
10. Epicuticular wax form: (0) granules, (1) tubules, (2) solid rodlets, (3) filaments, (4) plates, (5) ribbons.
11. Substances accumulation in epidermal cells: (0) absent, (1) present.
12. Epidermal cell lignification: (0) absent, (1) present.

Periderm

13. Bark type: (0) smooth bark (superficial periderm without rhytidome), (1) scaly bark.
14. Periderm stratification: (0) absent, (1) present.
15. Periderm formation: (0) uniform around the stem (annular removable bark), (1) in discontinuous areas (scaly bark).
16. Superficial periderm formation: (0) subsequent periderm width smaller than initial periderm, (1) subsequent periderm width same to initial periderm, (2) subsequent periderm width higher than initial periderm.
17. Phellem stratification: (0) absent, (1) present.
18. Phellem width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.
19. Intercellular spaces among phellem cells: (0) absent, (1) present.
20. Phellem cells in radial disposition: (0) absent, (1) present.
21. Pits in phellem cell: (0) absent, (1) present.
22. Phelloid cells in phellem: (0) absent, (1) present.
23. Sclereid cells in phellem: (0) absent, (1) present.
24. Phellogen width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.
25. Phellogen width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.

26. Phellogen cells in radial disposition: (0) absent, (1) present.
27. Lenticels: (0) absent, (1) present.

Cortex

28. Cortical vascular bundles: (0) absent, (1) present.
29. Vascular bundle type: (0) collateral closed, (1) collateral open, (2) bicollateral, (3) concentric amphicribal, (4) concentric amphivasal.
30. Adventitious roots: (0) absent, (1) present.

Cortical parenchyma

31. Substances accumulation in parenchyma cells: (0) absent, (1) present.
32. Intercellular spaces among parenchyma cells: (0) absent, (1) present.
33. Photosynthetic parenchyma: (0) absent, (1) present.
34. Crystal accumulations in parenchyma cells: (0) absent, (1) present.
35. Crystal accumulation type: (0) prismatic crystals, (1) druses, (2) raphides.
36. Aerenchyma: (0) absent, (1) present.
37. Plastids in parenchyma cells: (0) absent, (1) present.
38. Mucilage conduits: (0) absent, (1) present.

Cortical collenchyma

39. Collenchyma distribution: (0) continuous band around the stem, (1) in discontinuous areas.
40. Collenchyma type: (0) angular, (1) lamellar, (2) lacunar.
41. Chloroplasts in collenchyma: (0) absent, (1) present.
42. Pits in collenchyma cells: (0) absent, (1) present.
43. Crystal accumulations in collenchyma cells: (0) absent, (1) present.
44. Crystal accumulation type: (0) prismatic crystals, (1) druses, (2) raphides.

Cortical sclerenchyma

45. Sclerenchyma cell type: (0) sclerenchyma fibers, (1) stone cells, (2) both cell types.
46. Sclerenchyma distribution: (0) adjacent to phellogen, (1) center of the cortex, (2) adjacent to phloem.
47. Pits in sclerenchyma cells: (0) absent, (1) present.
48. Ramiform pit in sclerenchyma cells: (0) simple, (1) ramiform.

Vascular system

49. Vascular cambium width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.

APPENDIX: Continuation.

50. Internal phloem: (0) absent, (1) present.
 51. Included phloem: (0) absent, (1) concentric, (2) diffuse.
 52. Endodermis: (0) absent, (1) present.
 53. Endodermis width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.
 54. Caspian strip in endodermis: (0) absent, (1) present.
 55. Pericycle: (0) absent, (1) present.
 56. Pericycle width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.

Phloem

57. Sieve-tube elements distribution: (0) radial disposition, (1) tangential bands, (2) scattered
 58. Phloem fibers: (0) absent, (1) radial and/or tangential disposition, (2) scattered, (3) sieve-tube elements immersed in a fiber matrix.
 59. Phloem parenchyma: (0) absent, (1) present.
 60. Crystal accumulations in phloem: (0) absent, (1) present.
 61. Crystal accumulation type: (0) prismatic crystals, (1) druses, (2) raphides.

Xylem

Vessels

62. Xylem porosity: (0) vessels in the earlywood larger than those in the latewood in previous and same growth ring, (1) gradual change in vessel width among earlywood and latewood, (2) vessels with more or less the same diameter throughout the growth ring.
 63. Vessel dimorphism independent of the growth rings: (0) absent, (1) present.
 64. Vessel arrangement: (0) tangential bands, (1) diagonal and/or radial pattern, (2) dendritic pattern.
 65. Vessel groupings: (0) solitary, (1) in radial multiples of two to four vessels with a variable proportion of solitary vessels, (2) in radial multiples of four or more common, (3) clusters common.
 66. Solitary vessel outline: (0) rounded, (1) angular.
 67. Vessel tangential diameter.
 68. Vessels number per mm².
 69. Tyloses and deposits in vessels: (0) absent, (1) present.

Fibers

70. Fiber dimorphism: (0) absent, (1) present.
 71. Fiber dimorphism bands composition: (0) only dimorphic fibers, (1) dimorphic fibers and axial parenchyma.
 72. Fiber dimorphism bands width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.

73. Fiber dimorphism bands distribution: (0) reticulate (tangential bands of approximately the same width as the rays), (1) radial scalariform (tangential bands width smaller than rays), (2) tangential scalariform (tangential bands width higher than rays).

Axial parenchyma

74. Axial parenchyma in xylem: (0) absent, (1) present.
 75. Axial parenchyma type: (0) extremely rare, (1) apotracheal, (2) paratracheal, (3) banded parenchyma, (4) bands in growth ring limits.
 76. Paratracheal axial parenchyma type: (0) scanty, (1) vascentric, (2) aliform, (3) confluent, (4) unilateral.
 77. Axial parenchyma bands width: (0) one cell row, (1) two cell rows, (2) more than two cell rows.
 78. Axial parenchyma bands distribution: reticulate (tangential bands of approximately the same width as the rays), (1) radial scalariform (tangential bands width smaller than rays), (2) tangential scalariform (tangential bands width higher than rays).

Rays

79. Rays in xylem: (0) absent, (1) present.
 80. Aggregate rays: (0) absent, (1) present.

Miscellaneous xylem characters

81. Growth rings: (0) absent, (1) present.

Pith

82. Medullary vascular bundles: (0) absent, (1) present.
 83. Vascular bundle type: (0) collateral closed, (1) collateral open, (2) bicollateral, (3) concentric amphicribal, (4) concentric amphivasal.

Medullary parenchyma

84. Substances accumulation in parenchyma cells: (0) absent, (1) present.
 85. Intercellular spaces among parenchyma cells: (0) absent, (1) present.
 86. Photosynthetic parenchyma: (0) absent, (1) present.
 87. Crystal accumulations in parenchyma cells: (0) absent, (1) present.
 88. Crystal accumulation type: (0) prismatic crystals, (1) druses, (2) raphides.
 89. Aerenchyma: (0) absent, (1) present.
 90. Plastids in parenchyma cells: (0) absent, (1) present.
 91. Mucilage conduits: (0) absent, (1) present.



APPENDIX: Continuation.

Medullary sclerenchyma

92. Sclerenchyma cell type: (0) sclerenchyma fibers, (1) stone cells, (2) both cell types.

93. Sclerenchyma distribution: (0) pith center, (1) concentric rings, (2) adjacent to vascular system, (3) scattered.

94. Pits in sclerenchyma cells: (0) absent, (1) present.

95. Ramiform pit in sclerenchyma cells: (0) simple, (1) ramiform.

RADIAL SECTION

Cortex

Cortical sclerenchyma

96. Sclerenchyma pit border: (0) simple, (1) bordered pits.

97. Septate fibers in cortex: (0) absent, (1) present.

Vascular system

Phloem

98. Sieve plate type: (0) simple, (1) scalariform, (2) reticulate.

99. Sieve plate orientation: (0) inclined, (1) transverse.

100. Phloem parenchyma cellular composition: (0) fusiform, (1) strands of short cells, (2) both of them.

101. Septate fibers in phloem: (0) absent, (1) present.

Xylem

Vessels

102. Vessels length.

103. Perforation plate type: (0) simple, (1) scalariform, (2) reticulate, (3) foraminate.

104. Perforation plate orientation: (0) inclined, (1) transverse.

Fibers

105. Fiber type in xylem: (0) fiber-tracheids, (1) libriform fiber.

106. Septate fibers in xylem: (0) absent, (1) present.

Axial parenchyma

107. Axial parenchyma cellular composition: (0) fusiform, (1) strands of short cells, (2) both of them.

Rays

108. Ray cellular composition: (0) procumbent cells, (1) upright and/or square cells, (2) body ray cells procumbent with a row of upright and/or square marginal cells, (3) body ray cells upright and/or square with a row of procumbent marginal cells, (4) rays

with procumbent, square and upright cells mixed throughout the ray.

109. Vessel-ray pit border: (0) absent, (1) half-bordered (border inside the vessel) (2) bordered pit in both cells.

110. Vessel-ray pit form: (0) rounded to angular, (1) scalariform, (2) two different sizes and forms in the same ray cell.

111. Vessel-ray pit arrangement: (0) vessel-ray pits in all ray cells, (1) vessel-ray pits restricted to marginal cells.

Miscellaneous xylem characters

112. Idioblasts: (0) absent, (1) associated to rays, (2) associated to axial parenchyma, (3) immersed in the fibers.

113. Intercellular canals: (0) absent, (1) associated to rays, (2) associated to axial parenchyma, (3) immersed in the fibers.

114. Crystal accumulations in xylem: (0) absent, (1) present.

115. Crystal accumulation type: (0) prismatic crystals, (1) druses, (2) raphides.

Pith

Medullary sclerenchyma

116. Sclerenchyma pit border: (0) simple, (1) bordered.

117. Septate fibers in pith: (0) absent, (1) present.

TANGENTIAL SECTION

Vascular system

Xylem

Vessels

118. Intervessel pit border: (0) simple, (1) bordered.

119. Intervessel pit form: (0) scalariform, (1) rounded to angular.

120. Intervessel pit arrangement: (0) opposite, (1) alternate.

Rays

121. Ray cells series: (0) uniseriate, (1) multiseriate.

122. Rays with uniseriate and multiseriate portions: (0) absent, (1) present.

123. Rays arranged in tiers: (0) absent, (1) only in uniseriate rays, (2) present in all rays.

Miscellaneous xylem characters

124. Axial elements are arranged in tiers (horizontal series): (0) absent, (1) present.

125. Intercellular horizontal canals: (0) absent, (1) independent to rays, (2) inside the rays.