Leaf anatomical features of three *Theobroma* species (Malvaceae s.l.) native to the Brazilian Amazon

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**ABSTRACT**

*Theobroma* species have economic importance due to their use in the cosmetic and food industries, mainly in the production of chocolate. However, the anatomy of their vegetative structures remains poorly studied. The goal of this study was to describe the anatomical features of *Theobroma grandiflorum*, *T. speciosum* and *T. subincanum* to contribute to the biological knowledge of these species, as well as provide support to the biotechnological studies of native fruit plants of the Amazon. Leaves at different developmental stages were collected and analyzed under light microscopy and scanning electron microscopy. Sessile and stalked stellate trichomes and digitiform glandular trichomes were observed in the expanded leaves of *T. grandiflorum* and *T. subincanum*. These species were also similar in the morphology of the midrib, the organization of the mesophyll and the presence of starch grains in the midrib pith cells. Claviform glandular trichomes and mucilage cells in the epidermis occurred only in the expanded leaves of *T. speciosum*. The presence of mucilage secretory trichomes in shoot apices (colleters) of all species is a new finding for the genus *Theobroma*.

**KEYWORDS:** colleters, cupuassu, glandular trichomes, mucilage, stellate trichomes.

Caracteres anatômicos foliares de três espécies de *Theobroma* (Malvaceae s.l.) nativas da Amazônia Brasileira

**RESUMO**

As espécies de *Theobroma* têm importância econômica devido a sua utilização nas indústrias cosmética e alimentícia, principalmente na produção de chocolate. Entretanto, a anatomia de suas estruturas vegetativas permanece pouco conhecida. O presente estudo teve por objetivo descrever as características anatômicas de *Theobroma grandiflorum*, *T. speciosum* e *T. subincanum*, como contribuição ao conhecimento biológico dessas espécies, bem como, fornecer subsídios aos estudos biotecnológicos de fruteiras nativas da Amazônia. Folhas em diferentes estágios de desenvolvimento foram coletadas e analisadas sob microscopia de luz e eletrônica de varredura. Nas folhas expandidas de *T. grandiflorum* e *T. subincanum* foram observados tricomas dos tipos estrelado sésil, pedunculado e glandular digitiforme. Estas espécies também foram similares quanto à morfologia da nervura central, à organização do mesofilo e à presença de grãos de amido no parênquima medular da nervura central. Tricomas glandulares claviformes e células mucilaginosas na epiderme do limbo foliar ocorreram somente em *T. speciosum*. A presença de tricomas secretores de mucilagem nos ápices vegetativos (coléteres) de todas as espécies estudadas é um novo registro para o gênero *Theobroma*.

**PALAVRAS-CHAVE:** coléteres, cupuaçu, mucilagem, tricomas estrelados, tricomas glandulares.
INTRODUCTION

The genus *Theobroma* L. (Malvaceae s.l.) includes 22 species distributed from the Amazon basin to southern Mexico (Cuatrecasas 1964, APG III 2009). In Brazil, it is represented by 13 species, all native to the Amazon (Esteves 2013). In this genus *Theobroma cacao* L. and *T. grandiflorum* (cupuassu) are the most economically important species, used, respectively, in the cosmetic and food industries, mainly in the production of chocolate and “cupulate” (a seed-derived product similar to chocolate). The seeds of *T. grandiflorum* have a juicy pulp with an acid-sweet flavor, which is used in the preparation of beverages, candies and ice creams (Venturieri 2011). Other species of the genus, such as *T. speciosum* (cacaui) and *T. subincanum* (cupuí) are not marketed, but their sweet pulp is consumed fresh and their seeds are used in the preparation of chocolate (Cuatrecasas 1964, Cavalcante 2010).

*Theobroma grandiflorum* is widely cultivated in northern Brazil, the largest production in the state of Pará, followed by Amazonas, Rondônia and Acre. However, the cultivation of this species is limited by insect pests and several fungal pathologies, mainly “witches’ broom” (*Crinipellis perniciosa* (Stahel) Singer), which affects the vegetative and reproductive organs (Hanada et al. 2010).

The maintenance of the genetic diversity through germplasm collections, either in situ or ex situ, allows for the development of disease and pest resistant cultivars in addition to genetic breeding. Two in vivo collections of *Theobroma* are located in the state of Pará “George O’Neill Addison” and “Basil George David Bartley”. These collections are essential for the in situ conservation of genetic resources since the seeds of *Theobroma* are recalcitrant, which limits the viability of conventional ex situ conservation (Silva et al. 2004).

The morphological and anatomical knowledge of *Theobroma* is important for the establishment and maintenance of germplasm collections, providing valuable information to breeding programs and agronomic studies regarding desirable attributes (Santos et al. 2012). Furthermore, the anatomical characters of *Theobroma* may provide useful information for phytopathological studies, contributing to the selection of disease resistant mother trees, mainly to combat witches’ broom (Rio et al. 2008).

Although anatomical studies of this genus have recently been performed, most are related to seeds (Santos 2003, Martini et al. 2008), whereas the vegetative organs remain poorly investigated (Metcalfe and Chalk 1950, Nakayama et al. 1996, Ferreira et al. 2008). Therefore, the goal of the present study was to investigate the leaf anatomy of *T. grandiflorum*, *T. speciosum* and *T. subincanum* in order to contribute to the biological knowledge of species of *Theobroma* and provide support for the biotechnological studies of native fruit plants of the Amazon.

MATERIALS AND METHODS

Plant material

Expanded leaves were collected from *Theobroma grandiflorum* (Willd. ex Spreng) Schum., *T. speciosum* Willd. ex Spreng, and *T. subincanum* Mart. plants growing in the city of Moju, Pará, Brazil (1’44’S; 48°30’W) and at the Estação de Recursos Genéticos do Cacau José Haroldo/Comissão Executiva do Plano da Lavoura Cacaueira (ERJOH/CEPLAC) (1’12’S; 49°13’W), located in the city of Marituba, Pará, Brazil. Voucher materials were deposited in the João Murça Pires Herbarium at Museu Paraense Emílio Goeldi: *T. grandiflorum* (MG 200926, MG 196837), *T. speciosum* (MG 202760, MG 200901) and *T. subincanum* (MG 198499, MG 200900). Samples were selected from the third median of the petiole and the leaf blade from the fifth visible node. Vegetative shoot apices and non-expanded leaves were also collected for the analysis of the glandular trichomes.

Light microscopy

Samples were fixed in FAA (formalin, acetic acid, 70% alcohol) for 24 h (Johansen 1940) or NBF (buffered neutral formalin) for 48 h (Lillie 1965), dehydrated in a tertiary butanol series and embedded in Paraplast (Johansen 1940). Transverse and paradermal sections (10-12 µm thick) were cut on a rotatory microtome (RM 2245, Leica Biosystem Nussloch GmbH, Nussloch, Germany) and stained with astra blue (0,5%) and safranin (1%) (Gerlach 1984).

In addition, samples were fixed in 2,5% glutaraldehyde in 0.1 M sodium phosphate buffer pH 7.2 for 24 h, dehydrated in an ethanol series and embedded in hydroxyethyl methacrylate (Leica-Historessin), following the manufacturer’s recommendations. Then, sections (8-10 µm) were stained with toluidine blue O, pH 6.8 (O’Brien et al. 1964). For epidermal analysis, leaf blade segments were dissociated using a hydrogen peroxide and acetic acid solution (Franklin 1945). Afterwards, epidermis were stained with astra blue (0,5%) and safranin (1%) (Gerlach 1984) and mounted in 50% aqueous glycerin (Purvis et al. 1964).

Samples of both fixed and fresh material were submitted to histochemical tests using the following stains: periodic acid-Schiff’s (PAS) reaction to identify total polysaccharides (McManus 1948), ruthenium red to detect mucilage (Gregory and Baas 1989), sudan black B to identify total lipids (Pearse 1950), and Lugol’s solution to identify starch (Johansen 1940). Phenolic substances were detected through ferric chloride and by fixing fresh samples in a ferrous sulphate solution in formalin for 48 h (Johansen 1940). Photomicrographs were

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taken using a digital camera coupled to a light microscope (Axiolab, Zeiss, Jena, Germany).

**Scanning electron microscopy**

For scanning electron microscopy (SEM), the samples were fixed in 2.5% glutaraldehyde in 0.1 M sodium phosphate buffer pH 7.2 for 24 h. They were then dehydrated in an ethanol series, critical-point dried using CO₂, sputtered with gold (Robards 1978), and examined with a scanning electron microscope (Leo Electron Microscopy Ltd, 1450 VP, Cambridge, England).

**RESULTS**

**Leaf blade**

In surface view, the anticlinal walls of epidermal cells presented a slightly sinuous contour in *T. grandiflorum* and *T. speciosum* (Figures 1A, 1B) and were straight in *T. subincanum* (Figure 1C; Table 1). In all studied species, the epidermis was covered by a smooth cuticle layer with plate-like epicuticular wax deposits on the adaxial surface (Figure 1D). Abaxial surface had a dense indumentum (Figure 1E), mainly in *T. subincanum* (Figure 1F). All studied species had a leaf blade with a single-layered epidermis in cross section, with cells

![Figure 1](image-url)
containing phenolic substances in both surfaces (Figure 1G). Histochemical tests showed that in addition to phenolic cells, the adaxial epidermis of *T. speciosum* contained mucilage idioblasts (Figures 1H, 1I; Table 1).

Glandular and non-glandular trichomes were distributed on both surfaces of the expanded leaves of all species but they were more abundant on the abaxial surface (Figure 2A). Two morphological types of glandular trichomes were observed in this region: digitiform (Figures 2A, 2B), occurring on the leaf surface of *T. grandiflorum* and *T. subincanum*, and clavate (Figure 2C), occurring only on the leaf surface of *T. speciosum*. The two types of trichomes consisted of a basal cell, a bicellular stalk, and a multicellular secretory head (Table 1).

A number of clavate glandular trichomes (colleters) were also observed on both surfaces of leaf primordia (Figure 2D) and non-expanded leaves of all species (Figures 2E-H), but they differed from those described above by their size and cell number on the stalk and on the secretory head (Table 1). On the expanded leaves of the fifth visible node, only the basal region of these trichomes could be observed, indicating that they were deciduous. The secretion produced by these trichomes during the initial phases of leaf expansion was highly hygroscopic and viscous. In all species, PAS and ruthenium red reactions for polysaccharides and mucilage, respectively, were positive for the substances secreted by the head cells. The non-glandular trichomes were multicellular, stellate, four- to eight-armed, sessile (Figure 2I) or stalked (Figure 2J). The sessile stellate trichomes were abundant on the abaxial surface in all studied species, whereas the stalked stellate trichomes were exclusive to the veins on both leaf surfaces of the *T. grandiflorum* and *T. subincanum* (Table 1).

In cross section, the species had hypostomatic leaves with anomocytic stomata located above the other epidermal cells. The mesophyll in all species was dorsiventral with collateral vascular bundles surrounded by a sheath of sclerenchymatous and parenchymatous cells with phenolic content, which extended towards one or both surfaces (Figures 3A, 3B). The palisade parenchyma consisted of a single layer in *T. speciosum*, one to three layers in *T. subincanum* and *T. grandiflorum*. The spongy parenchyma was composed of three layers of transversally elongated cells in *T. speciosum* and three to five layers of isodiametric cells in *T. subincanum* and *T. grandiflorum* (Table 1). Idioblasts containing calcium oxalate druses (Figure 3C) and prismatic crystals (Figure 3D) were observed among palisade parenchyma cells, particularly in *T. subincanum* and *T. grandiflorum*.

### Table 1. Anatomical characters of leaf blade of *T. grandiflorum*, *T. speciosum* and *T. subincanum*. (+): present, (-): absent. (S1: leaf primordia and non-expanded leaves; S2: fully expanded leaves of fifth visible node).

<table>
<thead>
<tr>
<th>Character/Species</th>
<th><em>T. grandiflorum</em></th>
<th><em>T. speciosum</em></th>
<th><em>T. subincanum</em></th>
</tr>
</thead>
<tbody>
<tr>
<td>Epidermal anticlinal cell walls</td>
<td>Slightly sinuous</td>
<td>Slightly sinuous</td>
<td>Straight</td>
</tr>
<tr>
<td>Mucilage idioblasts in the epidermis</td>
<td>-</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>Glandular trichome S1</td>
<td>Clavate</td>
<td>Clavate</td>
<td>Clavate</td>
</tr>
<tr>
<td>Size (µm)</td>
<td>50 – 90</td>
<td>40 – 90</td>
<td>40 – 90</td>
</tr>
<tr>
<td>Secretory head (number of cells)</td>
<td>7 – 12</td>
<td>5 – 10</td>
<td>6 – 7</td>
</tr>
<tr>
<td>Stalk (number of cells)</td>
<td>3 – 4</td>
<td>2 – 3</td>
<td>3 – 4</td>
</tr>
<tr>
<td>Glandular trichome S2</td>
<td>Digitiform</td>
<td>Clavate</td>
<td>Digitiform</td>
</tr>
<tr>
<td>Size (µm)</td>
<td>30 – 35</td>
<td>30 – 35</td>
<td>25 – 30</td>
</tr>
<tr>
<td>Secretory head (number of cells)</td>
<td>3 – 4</td>
<td>4 – 6</td>
<td>2 – 3</td>
</tr>
<tr>
<td>Stalk (number of cells)</td>
<td>2</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>Stellate trichome</td>
<td>Sessile and stalked</td>
<td>Sessile</td>
<td>Sessile and stalked</td>
</tr>
<tr>
<td>Spongy parenchyma cells</td>
<td>Isodiametric</td>
<td>Elongated</td>
<td>Isodiametric</td>
</tr>
<tr>
<td>Midrib</td>
<td>Concave-convex</td>
<td>Biconvex</td>
<td>Concave-convex</td>
</tr>
<tr>
<td>Cortical region (predominantly)</td>
<td>Sclerenchymatous +</td>
<td>Parenchymatous</td>
<td>Parenchymatous</td>
</tr>
<tr>
<td>Starch grains in the pith</td>
<td></td>
<td>-</td>
<td>+</td>
</tr>
</tbody>
</table>

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In cross section, the leaf margins were revolute and had a rounded shape, thick cuticle, chlorenchyma consisting of isodiametric cells, and phenolic and crystal idioblasts in all of the species (Figure 3E). The midrib in cross section was biconvex in \textit{T. speciosum} (Figure 3F) and concave-convex in the remaining species (Figures 3G, 3H; Table 1). In all of the species, two or three layers of angular collenchyma cells were located below the abaxial epidermis. Cortical region was composed predominantly of sclerenchymatous cells in \textit{T. grandiflorum} (Figure 4A) and parenchymatous cells in \textit{T. subincanum} and \textit{T. speciosum} (Figure 4B). The vascular system of the midrib of all species was collateral, surrounded by a sclerenchymatous sheath (Figures 3H, 4A).

Cells containing phenolic compounds (Figure 4C) and crystals (prismatic and druses) were common in the dermal, vascular and ground tissue systems. Numerous starch grains were observed in the parenchyma cells of the midrib pith in \textit{T. grandiflorum} and \textit{T. subincanum} (Figure 4D; Table 1). Glands occurred both in the midrib cortex and pith (Figures 3F-H, 4B, 4D). The histochemical tests performed on these glands to detect total polysaccharides and mucilage showed positive results. However, lipophilic and phenolic substances were not identified. The anatomical characters of the leaf...
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Figure 3. Mesophyll, margin and midrib of *T. speciosum* (A, F), *T. grandiflorum* (B, C-E, H) and *T. subincanum* (G) in cross section. A-B. Dorsiventral mesophyll, note the of sheath extensions bundles (arrows). C-D. Idioblasts containing calcium oxalate crystals in the mesophyll, druses (C) and prismatic (D). E. Margin. F-H. Midrib (abe: abaxial epidermis, ade: adaxial epidermis, g: gland, s: stomata, ph: phloem, pp: palisade parenchyma, sp: spongy parenchyma, ss: sclerenchymatous sheath, x: xylem). Scale bars = 40µm (A-B, E), 20µm (C, D), 300µm (F, G), 200µm (H). This figure is in color in the electronic version.

blades of *T. grandiflorum*, *T. speciosum* and *T. subincanum* were summarized in Table 1, where it was possible to verify the main similarities and differences among these species.

**Petiole**

The petiole of all studied species presented, in cross section, one layered epidermis with stomata located above the other epidermal cells (Figure 4E). The glandular and non-glandular trichomes were similar to those described for the leaf blade. The glandular trichomes were clavate and composed of a basal cell, two- or three-celled uniseriate stalk and a five- to nine-celled secretory head. The non-glandular trichomes could be sessile in *T. speciosum* and stalked in *T. grandiflorum* and *T. subincanum* (Figure 2L).

In addition, numerous idioblasts containing phenolic compounds and calcium oxalate crystals (prismatic and druses) occurred in the petiole cortex (Figures 4E, 4F). Druses were more numerous than prismatic crystals, mainly in *T. subincanum*. Mucilage glands, similar to those of the leaf blade were found in the petiole cortical (Figure 4F) and pith regions (Figure 4G). The vascular bundles of petiole were collateral in all studied species (Figure 4G).
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**DISCUSSION**

The leaf anatomical features examined in the present study are in accordance with those previously described for Sterculiaceae (currently circumscribed in Malvaceae s.l.) regarding the vascular structure of petiole, stomatal type and distribution, presence of crystals usually in form of druses, mucilage-secreting structures, and stellate trichomes (Metcalfe and Chalk 1950). The last two characteristics are synapomorphies of Malvaceae s.l. (APG III 2009).

The studied species are characterized by a dense indumentum consisting of glandular and non-glandular trichomes. The morphological variation and distribution of these trichomes on the leaf surface are useful characters for the subgeneric classification in Malvaceae s.l., as reported for Mortoniodendron Standl. & Steyerm. (Rédon-Carmona et al. 2006), *Abutilon* Mill. (Shaheen et al. 2009a), *Hibiscus* L. (Shaheen et al. 2009b), *Sida* L. (Shaheen et al. 2009c), and in the present study for *Theobroma*.
To date, glandular trichomes in *Theobroma* are reported only for *T. cacao* (Nakayama et al. 1996). We are describing for the first time the presence of glandular trichomes in *T. grandiflorum, T. speciosum* and *T. subincanum*. In these species, although the trichomes found in the vegetative shoot apices are homologous to those of expanded leaves, they may play a different role. In the vegetative shoot apices, these trichomes are abundant, with a well developed secretory head, whose mucilage secretion may provide lubrication and protection against desiccation during early stages of leaf development. According to Paiva and Martins 2011, structures with this morphology are similar to colleters. In the family Malvaceae s.l. colleters are reported only in Hibiscus pernambucensis Arruda and H. silicaceus L. (Rocha et al. 2002).

Leaf trichomes may be an effective defense mechanism in plants. They may be involved in both chemical and mechanical defense, negatively affecting the oviposition rate and feeding of herbivorous insects and nutrition of larvae (Lokesh and Singh 2005). It is possible that the leaf trichome density in the studied species can play a role in protection against herbivores insects, which are considered pests in the cultivation of some species of this genus, such as beetle (Barbosa and Fonseca 2002), caterpillar (Lourido et al. 2007), and leaf-cutting ants (Delabie 1989). However, additional studies are needed to better understand the ecological role of these trichomes in *Theobroma*.

The occurrence of mucilaginous cells in the leaf epidermis is described in 69 families of eudicotyledons including Malvaceae s.l. (Metcalfe and Chalk 1950). In the present study, *T. speciosum* is the only species to present mucilaginous cells in the leaf epidermis. However, these mucilaginous cells are observed only on the adaxial surface of the leaf blade, a feature also reported in other representatives of Malvaceae s.l., such as *T. cacao* L. (Nakayama et al. 1996), Dombeya Desf. and Nesogordonia Baill. (Lersten and Curtis 1997), Sterculia Vent. (Hussin and Sani 1998), Hibiscus (Rocha et al. 2002) and Pavonia alnifolia A.St.-Hil. (Pimentel et al. 2011).

Another important feature found in the species examined for this study is the presence of numerous calcium oxalate crystals, mainly in the form of druses. Calcium oxalate crystals in plants are involved in several physiological functions and may play an ecological role as static or active defense structures (Franceschi and Nakata 2005). Their synthesis not only depends on calcium levels but can also be influenced by external pressures such as herbivory. For example, in Sida rhombifolia L. (Malvaceae), herbivory increases the synthesis of crystals in seedling leaves even if calcium availability is limited (Molano-Flores 2001).

Cells with phenolic content also have a wide distribution on leaf tissues of the species studied here. A previous study has shown that phenolic compounds in *T. cacao* have an inhibitory effect on *Crinipellis perniciosa* basidiospores germination, representing a mechanism of resistance to witches’ broom disease (Nojosa et al. 2003). The presence of crystals and phenolic idioblasts in the leaves of *Theobroma* species may thus have a protective function against herbivory and may increase the resistance to witches’ broom disease.

**CONCLUSION**

We have verified that *T. grandiflorum* and *T. subincanum* share the presence of digitiform glandular trichomes, stalked and sessile stellate trichomes, isodiametric cells in the spongy parenchyma, a concave-convex midrib, and starch grains in the midrib pith cells, while *T. speciosum* presents clavate glandular trichomes, only sessile stellate trichomes, elongated cells in the spongy parenchyma, a biconvex midrib, and mucilaginous cells in the epidermis. Such characteristics provide a useful tool for taxonomic studies of the species of *Theobroma*. We have evidenced that the presence of mucilage secretory trichomes in shoot apices (colleters) of all species is a new finding for this genus.

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