Evolution of the Anther Gland in Early-Branching Papilionoids (ADA Clade, Papilionoideae, Leguminosae)

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Abstract: Papilionoideae is the most diverse subfamily of Leguminosae, especially in terms of floral morphology. The ADA clade shows some exciting floral features among papilionoids, such as anther glands. However, the evolution of the anther glands in such early-branching papilionoids remains unknown. Thus, we compared the occurrence, distribution, morphology, and evolutionary history of the anther glands in species of the ADA clade. Floral buds and/or flowers in 50 species were collected from herbarium specimens and investigated using scanning electron and light microscopy and reconstruction of ancestral character states. The anther apex has a secretory cavity, secretory duct, and phenolic idioblast. The lumen shape of the cavity and duct is closely related to the shape of the anther apex. The oval lumen is located between two thecae, the spherical lumen in the prominent anther apex and the elongated lumen in anthers with a long apex. The occurrence of cavities/ducts in the anther in only two phylogenetically closely related subclades is a unifying character-state. The floral architecture is not correlated with cavity/ducts in the anther but is possibly related to the type of pollinator. Future research needs to combine floral morphology and pollination systems to understand the evolution of floral designs and their diversification.

Keywords: anther; anatomy; Angylocalyceae; Amburaneae; Dipterygeae; secretory structure

1. Introduction

Leguminosae is a species-rich family with more than 19,000 species distributed into about 765 genera [1]. The currently accepted classification of the family includes six subfamilies: Cercidioideae LPWG (12 genera/ca. 335 species), Detarioideae Burmeist., Handb. Naturgesch (84 genera/ca. 760 species), Dialioideae LPWG (17 genera/85 species), Duparquetioideae LPWG (1 genera/1 species), Caesalpinioideae DC (12 genera/ca. 335 species) and Papilionoideae DC (503 genera/ca. 14,000 species).

Among the Leguminosae subfamilies, Papilionoideae is considered the most diverse and ecologically successful [2], with a recent history of diversification during the Cenozoic [3,4]. Its diversity is expressed in floral morphology because the presence of papilionaceous flowers characterizes legumes, which also exhibit flowers with other architectural types see [5–10].

The early-branching papilionoids comprise plants with exciting flower morphology. They were included in the ADA clade and comprised about 74 species [11,12] (see Figure 1) distributed into three subclades: Amburaneae (eight genera), Angylocalyceae (four genera), and Dipterygeae (four genera) (Figure 1) [1,10,12]. Some members such as *Dipteryx alata* and *Pterodon pubescens* exhibit an unusual condition in the family, which is the presence of glandular appendages in the anther containing a secretory cavity. The secretory cavity...
consists of an isodiametric to elliptical lumen delimited by a uniseriate secretory epithelium and a parenchyma sheath [7,13]. It secretes sticky substances (oleoresins and polysaccharides) with a key role in plant reproduction, aggregating pollen grains and attaching them to the floral visitor’s body [13].

**Figure 1.** Phylogenetic relationships between the three subclades of the ADA clade modified from [1,10,12]. The Angylocalyceae subclade is a sister group to the Amburaneae and Dipterygeae subclades.

Among the groups within the ADA clade, the Dipterygeae subclade stands out by sharing the presence of secretory cavities and ducts on some parts of the plant body other than anthers, such as the bracteole, sepal, petal [7], pulvine, petiole, rachis [14], leaf [15,16], stem [17], fruit [18], and even early during plant development [19].

For groups closely related to Dipterygeae, there are no records of secretory ducts/cavities in the anther or other floral organs (see [5,8,9]). Interestingly secretory cavities have been found in the leaflets of *Cordyla, Myrocarpus, Myroxylon,* and *Myropermum* [20,21], in genera of the Amburaneae clade, a sister group to the Dipterygeae subclade [1,10,12,22].

The absence of biological information for most species in the ADA clade is due to the sole use of surface analysis (scanning electron microscopy) in the study of flowers without including anatomical sections providing additional information concerning the internal anatomy and intra- and extracellular contents.

In the current study, we present a detailed morphological and evolutionary investigation into the anther glandular appendages of the ADA clade species. We intended to (i) compare the occurrence, distribution and morphology of the anther glandular appendages in the species of the ADA clade; (ii) to trace the evolutionary history of the secretory structures of the anther based on the recent phylogenetic hypothesis of the ADA clade (Figure 1) [1,10,12]; (iii) and to evaluate whether the presence of this condition is homologous in some groups.

The hypothesis we tested was that the presence of a secretory cavity/duct in the anther is widely distributed in the ADA clade, and thus, it was acquired by these taxa and can be considered as a synapomorphy for the group.

**2. Results**

Table 1 summarizes the results from our morphological analysis of the ADA clade species and other selected species. Additional information was obtained from the literature (Table 1). Because the presence of glands in the anther of this group is an uncommon condition, we analyzed and compared other characteristics related to the anther (for example, apex shape) and to other floral organs (for example, number per whorl and connation) in order to better understand their function in the flower and their evolutionary history.
Table 1. Matrix of morphological data obtained from information in the literature and analysis of herbarium material for the characters selected in the present study. Empty cells mean missing information.

| Subclade       | Species                               | Number of Sepals | Number of Petals | Number of Stamens | Stamen Connation | Occurrence of Cavity/Canal at the Anther | Position of Cavity/Canal at the Anther | Shape of Lumen of Cavity/Canal | Shape of the Anther Apex | Occurrence of Phenolic Cells at the Anther Apex | References |
|----------------|----------------------------------------|------------------|------------------|-------------------|------------------|-----------------------------------------|----------------------------------------|-------------------------------|--------------------------|----------------------------------------------------|------------|
| Amburaneae      | *Amburana acreana* (Ducke) A.C. Sm.    | 5                | 1                | 10                | free             | absent                                   | absent                                  | -                            | absence                  | absence                             | [23], Present study |
|                | *Amburana cearensis* (Allemão) A.C. Sm. | 5                | 1                | 10                | free             | absent                                   | absent                                  | -                            | absence                  | absence                             | [7]        |
|                | *Amburana erythrosperma* E. P. Selem., C. H. Stirt. & V.F. Mansano | 5                | 1                | 10                | free             | absent                                   | absent                                  | -                            | absence                  | absence                             | Present study |
|                | *Cordyla africana* Lour.               | 3                | 0                | numerous          | cavity           | apical                                   | spherical                               | prominent                     | prominent                | Present study                             | Present study |
|                | *Cordyla harakaii* Capuron             | 3                | 0                | numerous          | free             | cavity                                   | apical                                  | spherical                     | prominent                | Present study                             | Present study |
|                | *Cordyla madagascariensis* R. Vig.     | 3                | 0                | numerous          | free             | cavity                                   | apical                                  | spherical                     | prominent                | Present study                             | Present study |
|                | *Dussia discolor* (Berth.) Amshoff      | 5                | 5                | 10                | fused at the base | absent                                   | not prominent                          | oval                         | not prominent            | not prominent                         | [6], Present study |
|                | *Dussia lehmannii* Harms               | 5                | 5                | 10                | fused at the base | absent                                   | not prominent                          | oval                         | not prominent            | not prominent                         | [24], Present study |
|                | *Dussia macroprophylata* (Dorr. Sm.) Harms | 5                | 5                | 10                | fused at the base | absent                                   | not prominent                          | oval                         | not prominent            | not prominent                         | [24], Present study |
|                | *Dussia martincensis* Krug & Urb. ex Taub | 5                | 5                | 10                | fused at the base | absent                                   | not prominent                          | oval                         | not prominent            | not prominent                         | [24], Present study |
|                | *Dussia tessmannii* Harms              | 5                | 5                | 10                | fused at the base | absent                                   | not prominent                          | oval                         | not prominent            | not prominent                         | Present study |
|                | *Mildbraediodendron excelsum* Harms     | 5                | 0                | numerous          | free             | absent                                   | not prominent                          | oval                         | not prominent            | absence                             | [25,26], Present study |
|                | *Myrocarpus emarginatus* A.L.B. Sartori & A.M.G. Azevedo | 5                | 5                | 10                | fused at the base | cavity                                   | apical                                  | oval                         | not prominent            | [27,28], Present study |
|                | *Myrocarpus fastigiatus* Allemão       | 5                | 5                | 10                | fused at the base | cavity                                   | apical                                  | oval                         | not prominent            | [27,28], Present study |
|                | *Myrocarpus feroxius* Allemão          | 5                | 5                | 10                | fused at the base | cavity                                   | apical                                  | oval                         | not prominent            | [20,27], Present study |
|                | *Myrspermum frutescens* Jacq           | 5                | 5                | 10                | free             | cavity                                   | apical                                  | oval                         | prominent                | [27,28], Present study |
| Subclade                      | Species                                | Number of Sepals | Number of Petals | Number of Stamens | Stamen Connation | Occurrence of Cavity/Canal at the Anther | Position of Cavity/Canal at the Anther | Shape of Lumen of Cavity/Canal | Shape of the Anther Apex | Occurrence of Phenolic Cells at the Anther Apex | References                             |
|-----------------------------|-----------------------------------------|------------------|------------------|-------------------|------------------|-----------------------------------------|--------------------------------------|----------------------------------|----------------------------|------------------------------------------|-------------------------------------------|
| Myroxylon                   | Myroxylon balsamum (L.) Harms            | 5                | 5                | 10                | free             | duct                                    | apical, distal                      | elongate                         | prominent                  | Present study                           | [5, 27], Present study                        |
| Myroxylon                   | Myroxylon peruaferum L. L.              | 5                | 5                | 10                | free             | duct                                    | apical, distal                      | elongate                         | prominent                  | Present study                           | [27], Present study                         |
| Petaladenium                | Petaladenium urceoliferum Ducke         | 5                | 5                | 10                | (fused at the base = nearly free) | absent                                |                                      | not prominent                   | absence                    | Present study                           | [9], Present study                         |
| Alexa                       | Alexa bouchitiiflora Ducke              | 5                | 5                | 8/10              | free             | absent                                  |                                      | not prominent                   | presence                   | Present study                           | [30], Present study                         |
| Alexa                       | Alexa canaracencenis Pittier            | 5                | 5                | 10                | free             | absent                                  |                                      | not prominent                   | presence                   | Present study                           | [30], Present study                         |
| Alexa                       | Alexa couanzii Yakovlev                 | 5                | 5                | 10                | free             | absent                                  |                                      | not prominent                   | presence                   | Present study                           | [30], Present study                         |
| Alexa                       | Alexa grandiflora Ducke                 | 5                | 5                | 10                | free             | absent                                  |                                      | not prominent                   | presence                   | Present study                           | [30], Present study                         |
| Alexa                       | Alexa imperatricis (R.H. Schomb.) Baill.| 5                | 5                | 8/10              | free             | absent                                  |                                      | not prominent                   | absence                    | Present study                           | [30], Present study                         |
| Alexa                       | Alexa leiopetala Sandwith               | 5                | 5                |                    |                  | absent                                  |                                      | not prominent                   | presence                   | Present study                           | [30], Present study                         |
| Alexa                       | Alexa superba R.S. Cowan                | 5                | 5                | 10/15             | free             | absent                                  |                                      | prominent                       | absence                    | Present study                           | [30], Present study                         |
| Alexa                       | Alexa wachenheimii Benoist              | 5                | 5                | 10                | free             | absent                                  |                                      | not prominent                   | presence                   | Present study                           | [30], Present study                         |
| Angylocalyx                  | Angylocalyx pynaertii De Wild.          | 5                | 5                | 10                | monadelphous     | absent                                  |                                      | not prominent                   | absence                    | Present study                           | [31], Present study                         |
| Angylocalyx                  | Angylocalyx talbotii Hutch. & Dalziel   | 5                | 5                | 10                | monadelphous     | absent                                  |                                      | not prominent                   | absence                    | Present study                           | Present study                             |
| Castanospermum              | Castanospermum australis A. Cunn. ex Mudie| 5                | 5                | 10                | free             | absent                                  |                                      | prominent                       | absence                    | Present study                           | [5], Present study                         |
| Xanthocercis                | Xanthocercis zambesica (Baker) Dumaz-le-Grand| 5                | 5                | 10                | fused at the base | absent                                  |                                      | not prominent                   | absence                    | Present study                           | [32], Present study                         |
## Table 1. Cont.

| Subclade       | Species                        | Number of Sepals | Number of Petals | Number of Stamens | Stamen Connation | Occurrence of Cavity/Canal at the Anther | Position of Cavity/Canal at the Anther | Shape of Lumen of Cavity/Canal | Shape of the Anther Apex | Occurrence of Phenolic Cells at the Anther Apex | References |
|----------------|--------------------------------|------------------|------------------|-------------------|------------------|------------------------------------------|----------------------------------------|-------------------------------|----------------------------------|-----------------------------------------------|-------------|
| Dipterygeae    | Dipteryx alata Vogel           | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Dipteryx lacunifera Ducke      | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Dipteryx magnifica (Ducke) Ducke | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Dipteryx micrantha Harms       | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Dipteryx odorata (Aubl.) Willd. | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Dipteryx pulchellla (Fibber) Ducke | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | oval                          | not prominent                    | Present study                   |             |
|                | Dipteryx punctata (Blake) Amshoff | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Dipteryx roosa Spruce ex Benth | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Pterodon abruptus (Moric.) Benth. | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Pterodon emarginatus Vogel     | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Pterodon pubescens (Benth.) Benth. | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | spherical                      | prominent                       | Present study                   |             |
|                | Taralea cordata Ducke          | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | oval                          | not prominent                    | Present study                   |             |
|                | Taralea crassitolia (Benth.) Ducke | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | oval                          | not prominent                    | Present study                   |             |
|                | Taralea nudipes (Tul.) Ducke   | 5                | 5                | 10                | monadelphous     | cavity                                    | apical                                 | oval                          | not prominent                    | Present study                   |             |
|                | Taralea oppositifolia Aubl.    | 5                | 5                | 10                | monadelphous     | absent                                    | not prominent                         |                               |                                 |                                 | Present study                   |             |
|                | Taralea reticulata (Benth.) Ducke | 5                | 5                | 10                | monadelphous     | absent                                    | not prominent                         |                               |                                 |                                 | Present study                   |             |
|                | Taralea rigida Schery          | 5                | 5                | 10                | monadelphous     | absent                                    | not prominent                         |                               |                                 |                                 | Present study                   |             |
|                | Monopteryx inpae W.A.Rodrigues  | 5                | 5                | 10                | free            | absent                                    | apical                                |                               | not prominent                    | presence                       | [10,33], Present study |
|                | Monopteryx urucu Spruce ex Benth. | 5                | 5                | 10                | free            | absent                                    | apical                                |                               | prominent                       | presence                       | [10,33], Present study |
### Table 1. Cont.

| Subclade                   | Species                        | Number of Sepals | Number of Petals | Number of Stamens | Stamen Connation | Occurrence of Cavity/Canal at the Anther | Position of Cavity/Canal at the Anther | Shape of Lumen of Cavity/Canal | Shape of the Anther Apex | Occurrence of Phenolic Cells at the Anther Apex | References |
|----------------------------|--------------------------------|------------------|------------------|-------------------|------------------|------------------------------------------|---------------------------------------|-------------------------------|--------------------------|---------------------------------------------|------------|
|                           | **Outgroup**                   |                  |                  |                   |                  |                                          |                                       |                               |                          |                                            |            |
|                           | Ateleia glazioverana Baill.     | 5                | 1                | 10                | free             | absent                                    | not prominent                      | absence                       | not prominent             | absence                      | Present study |
|                           | Ateleia guaraya Herzog          | 5                | 1                | 10                | free             | absent                                    | not prominent                      | absence                       | not prominent             | absence                      | Present study |
|                           | Candolleodendron brachystachyum (DC.) R.S. Cowan | 5                | 1                | numerous         | free             | absent                                    | prominent                          | absence                       | not prominent             | absence                      | Present study |
|                           | Cyathostegia mathewsii (Benth.) Schery | 5                | 1                | numerous         | free             | absent                                    | not prominent                      | absence                       | not prominent             | absence                      | Present study |
|                           | Swartzia langsdorffii Raddi     | 4                | 0                | 2–3 (smaller stamens), numerous (larger stamens) | free             | absent                                    | prominent                          | presence                       | not prominent             | absence                      | Present study |
|                           | Ulleanthus erythrinoides Harms  | 5                | 4                | 10                | free             | absent                                    | not prominent                      | absence                       | not prominent             | absence                      | Present study |
2.1. Flower Morphology

The Amburaneae subclade species (Figures 2 and 3) have flowers with a one-petalled corolla and free stamens (*Amburana* and *Mildbraediodendron*), no corolla and free stamens (*Cordyla*), a papilionaceous corolla and basally united stamens (*Dussia* and *Petaladenium*), a non-papilionaceous corolla (five equal petals) and basally united stamens (*Myrocarpus*), with a papilionaceous corolla and free stamens (*Myrospermum*), and a non-papilionaceous corolla (widely oval standard) and 10 free homogeneous stamens (*Myroxylon*). A prominent anther apex occurs in species of *Cordyla*, *Myrospermum*, *Myroxylon* (Figure 2F,G,I, and Figure 3G,I,J), and a non-prominent anther apex occurs in species of *Amburana*, *Dussia*, *Mildbraediodendron*, *Myrocarpus*, and *Petaladenium* (Figures 2A,C,D,J–N and 3A,D,E,L).

![Figure 2](https://example.com/figure2)

**Figure 2.** Aspects of the anther apex morphology in the species of the Amburaneae subclade. (A,C,D,F,G,I–N) = SEM, (B,E,H,O) = light microscopy (LM). (A,B) *Amburana acreana*. (A) Frontal view of the anther. Note that the anther apex is non-prominent. (B) Longitudinal section showing the anther. Note the absence of a secretory cavity between the two thecae. (C) *A. cearensis*. Adaxial view of the anther. Note that the anther apex is non-prominent. (D) *A. erythrosperma*. Frontal view of the anther. Note that the anther apex is non-prominent. (E) Longitudinal section showing the anther. Note the absence of a secretory cavity between the two thecae. (F) *Cordyla africana*. Adaxial view of the anther. Note that the anther apex is non-prominent (arrow). (G,H) *C. madagascariensis*. (G) Frontal view of the anther. Note that the anther apex is prominent (arrow). (H) Longitudinal section showing the apex with a secretory cavity between the two thecae. (I) *C. haraka*. Frontal view of the anther. Note that the anther apex is prominent (arrow). (J) *Dussia discolor*. Abaxial view of the anther. Note that the anther apex is non-prominent. (K) *D. lehmannii*. Adaxial view of the anther. Note that the anther apex is non-prominent. (L) *D. macrophylla*. Adaxial view of the anther. Note that the anther apex is non-prominent. (M) *D. martinecensis*. Adaxial view of the anther. Note that the anther apex is non-prominent. (N,O) *D. tesmannii*. (N) Lateral view of the anther. (O) Longitudinal section showing the anther. Note the absence of a secretory cavity between the two thecae. Scale bars: (A,B,N), 200 µm; (C–E), 400 µm; (F–O), 100 µm; (H) 50 µm.
Figure 3. Aspects of the anther apex morphology in the species of the Amburaneae subclade. (A,D,E,G,I,J,L) = SEM, (B,C,F,H,K,M) = light microscopy, ML. (A,B) *Mildbraediodendron excelsum*. (A) Abaxial view of the anther. Note that the anther apex is non-prominent. (B) Longitudinal section showing the anther. Note the absence of a secretory cavity between the two thecae. (C) *Myrocarpus emarginatus*. Longitudinal section showing the anther apex with a secretory cavity. (D) *M. fastigiatus*. Frontal view of the anther. Note that the anther apex is non-prominent. (E,F) *M. frondosus*. (E) Adaxial view of anther. Note that the anther apex is non-prominent. (F) Longitudinal section showing the anther apex with a secretory cavity. (G,H) *Myrospermum frutescens*. (G) Frontal view of the anther. (H) Longitudinal section showing the anther apex with a secretory cavity. (I) *Myroxylon balsamum*. Lateral view of the anther. Note that the anther apex is prominent. (J,K) *M. peruferum*. (J) Adaxial view of the anther. Note that the anther apex is prominent. (K) Longitudinal section showing the anther apex with a secretory duct. Note that the lumen shape of the secretory duct follows the shape of the anther apex, rounded in its apical portion (arrowhead) and elongated in the lower portion. (L,M) *Petaladenium urceoliferum*. (L) Adaxial view of the anther. Note that the anther apex is non-prominent. (M) Longitudinal section of the anther showing the phenolic cells. Scale bars: (A,C–L), 100 µm; (B), 50 µm.
The Angylocalyceae subclade species (Figures 4 and 5) have flowers with a papilionaceous corolla and basally united stamens (Angylocalyx), a non-papilionaceous corolla that consists of a large vexillum and reduced abaxial petals (Alexa and Castanospermum), a non-papilionaceous corolla that consists of a slightly wider vexillum and four equal abaxial petals and united stamens (Xanthocercis). A prominent anther apex occurs in the two species of Alexa (Alexa grandiflora and A. superba) and Castanospermum australe (Figures 4E,I and 5C,E); a non-prominent anther apex occurs in species of Angylocalyx, in Xanthocercis zambesiaca and six species of Alexa (Figures 4A–C,G,H,L and 5A,G).

Figure 4. Aspects of the anther apex morphology in the species of the Angylocalyceae subclade. (A–C,E,G–I,K) = light microscopy. (D,F,J,L) = SEM. (A) Alexa bauhiniflora. Adaxial view of the anther. Note that the anther apex is non-prominent. (B) A. canaracunensis. Adaxial view of the anther. Note that the anther apex is non-prominent. (C,D) A. cowanii. (C) Lateral view of the anther. Note that the anther apex is non-prominent. (D) Longitudinal section of the anther showing the phenolic cells. (E,F) A. grandiflora. (E) Lateral view of the anther. Note that the anther apex is a small prominence. (F) Longitudinal section of the anther showing the phenolic cells. (G) A. imperatrizes. Adaxial view of the anther. Note that the anther apex is non-prominent. (H) A. leiopetala. Adaxial view of the anther. Note that the anther apex is non-prominent. (I,J) A. superba. (I) Lateral view of anther. Note that the anther apex is a small prominence. (J) Longitudinal section of the anther. Note the absence of a secretory cavity between the two thecae. (K,L) A. wachenheimii. (K) Lateral view of the anther. Note that the anther apex is non-prominent. (L) Longitudinal section of the anther showing the phenolic cells. Scale bars: (A,C,D), 200 µm; (B,G,L), 50 µm; (E,H), 200 µm; (F,J,K) 100 µm.
Figure 5. Aspects of the anther apex morphology in the species of the Angylocalyceae subclade. (A,C,E,F) = SEM, (B,D,F,H) = light microscopy, LM. (A,B) *Angylocalyx pynaertii*. (A) Adaxial view of the anther. Note that the anther apex is non-prominent. (B) Longitudinal section of the anther. Note the absence of a secretory cavity between the two thecae. (C,D) *A. talbotii*. (C) Lateral view of the anther. Note that the anther apex is non-prominent. (D) Longitudinal section of the anther. Note the absence of a secretory cavity between the two thecae. (E,F) *Castanospermum australe*. (E) Adaxial view of the anther. Note that the anther apex is a small prominence. (F) Longitudinal section of the anther. Note the absence of a secretory cavity between the two thecae. (G,H) *Xanthocercis zambesiaca*. (G) Adaxial view of the anther. Note that the anther apex is non-prominent. (H) Longitudinal section of the anther. Note the absence of a secretory cavity between the two thecae. Scale bars: (A), 200 μm; (B–H), 100 μm.
The Dipterygeae subclade species (Figures 6 and 7) have flowers with a papilionaceous corolla formed by a vexillum, two wings, and two keels, and a monadelphous androecium (Dipteryx, Pterodon, and Taralea); and a non-papilionaceous corolla that consists of a vexillum, two reduced wings, connate and open keels exposing the free stamens (Monopteryx). A prominent anther apex occurs in seven species of Dipteryx, all species of Pterodon and Monopteryx uaucu (Figures 6A,C,D–F,H,J–M and 7L); a non-prominent anther apex occurs in Dipteryx polyphylla, in all species of Taralea and Monopteryx iniae (Figures 6G and 7A,C,E,G,I–K).

![Figure 6](image_url)

**Figure 6.** Aspects of the anther apex morphology in the species of the Dipterygeae subclade. (A,C–H,J–M) = SEM, (B,I,N) = light microscopy, LM. (A,B) *Dipteryx alata.* (A) Adaxial view of the anther with longitudinal dehiscence and pollen grains adhering to the anther apical region. (B) Longitudinal section showing the anther apex with a secretory cavity. (C) *D. lacunifera.* Abaxial view of the anther. Note that the anther apex is prominent. (D) *D. magnifica.* Abaxial view. Note that the anther apex is prominent. (E) *D. micrantha.* Abaxial view. Note that the anther apex is prominent. (F) *D. odorata.* Abaxial view of the anther. Note that the anther apex is prominent. (G) *D. polyphylla.* Abaxial view of the anther. Note that the anther apex is non-prominent. (H,J) *D. punctata.* H. Lateral view of the anther. Note that the anther apex is prominent. (I) Longitudinal section showing the anther apex with a secretory cavity with a spherical lumen. (J) *D. rosea.* Lateral view of the anther. Note that the anther apex is prominent. (K) *Pterodon abruptus.* Adaxial view of the anther. Note that the anther apex is prominent. (L) *P. emarginatus.* Frontal view of the anther. Note that the anther apex is prominent. (M,N) *P. pubescens.* (M) Lateral view of the anther. Note that the anther apex is prominent. (N) Longitudinal section showing the anther apex with a secretory cavity. Scale bars (A,D,F,G,J,K,M), 100 μm; (B), 20 μm; (C,E,I,L,H), 50 μm; (N), 200 μm.
Figure 7. Aspects of the anther apex morphology in the species of the Dipterygeae subclade. (A,C,E,G,I–L) = SEM, (B,D,F,H,M) = light microscopy, LM. (A,B) Taralea cordata. (A) Abaxial view of the anther. Note that the anther apex is non-prominent. (B) Longitudinal section showing the anther apex with a secretory cavity with an oval lumen. (C,D) T. crassifolia. (C) Abaxial view of the anther. Note that the anther apex is non-prominent. (D) Longitudinal section showing the anther apex with a secretory cavity with an oval lumen. (E,F) T. nudipes. (E) Adaxial view of the anther. Note that the anther apex is non-prominent. (F) Longitudinal section showing the anther apex with a secretory cavity with an oval lumen. (G,H) T. oppositifolia. (G) Adaxial view of the anther. Note that the anther apex is non-prominent. (H) Longitudinal section showing parenchymal cells (circle) of the anther. (I) T. reticulata. Adaxial view of the anther. Note that the anther apex is non-prominent. (J) T. rigida. Adaxial view of the anther. Note that the anther apex is non-prominent. (K) Monopteryx impae. Lateral view of the anther. Note that the anther apex is non-prominent (arrowhead). (L,M) M. uaucu. (L) Lateral view of the anther. Note that the anther apex is prominent (arrowhead). (M) Longitudinal section showing the anther apex with phenolic cells. Scale bars: (A,E–I,K–M), 100 μm; (B,D,J), 200 μm; (C). 50 μm.
The additional species (Figure 8), *Ateleia glazioveana*, *A. guaraya*, *Cyathostegia mathewsii*, and *Uleanthus erythrinoides*, exhibit non-prominent anther apices (Figure 8A,C,G,M). In contrast, *Candolleodendron brachystachyum* and *Swartzia langsdorffii* have prominent anther apices (Figure 8E,K,I).

![Figure 8](image_url)

**Figure 8.** Aspects of the anther apex morphology in the species of the *Ateleia, Candolleodendron, Cyathostegia, Swartzia, Uleanthus*. (A,C,E,G,K,I,M) = SEM, (B,D,H,F,J,L,N) = light microscopy, LM. (A,B) *Ateleia glazioveana*. (A) Adaxial view of the anther. Note that the anther apex is non-prominent. (B) Longitudinal section showing parenchymal cells at the anther apex. Note that there is no secretory cavity between the anther’s thecae. (C,D) *Ateleia guaraya*. (C) Adaxial view of the anther. Note that the anther apex is non-prominent. (D) Longitudinal section showing parenchymal cells at the anther apex. (E,F) *Candolleodendron brachystachyum*. (E) Adaxial view of the anther. Note that the anther apex is prominent. (F) Longitudinal section showing parenchymal cells at the anther apex. (G,H) *Cyathostegia mathewsii*. (G) Abaxial view of the anther. Note that the anther apex is non-prominent. (H) Longitudinal section showing parenchymal cells of the anther. (I,J) *Swartzia langsdorffii* (I) Adaxial view of the anther of the small stamens with a small prominence apex. (J) Longitudinal section showing the anther apex with phenolic cells. (K) Adaxial view of the anther of the large stamens with a small prominent apex. (L) Longitudinal section showing the anther apex with phenolic cells. (M,N) *Uleanthus erythrinoides*. (M) Adaxial view of the anther. Note that the anther apex is non-prominent. (N) Longitudinal section showing parenchymal cells of the anther. Scale bars: (A–G,I,J,N), 100 µm; (H), 50 µm; (L,M), 200 µm.
2.2. Anther Glands

The anthers of the ADA clade species have three types of glands: secretory cavity, secretory duct, and phenolic idioblast (Table 1).

A secretory cavity occurs embedded in the anther apex of Cordyla, Myrocarpus, Myroxylon species (Amburaneae subclade), of all species of Dipteryx, Pterodon, and of three species of Taralea (T. cordata, T. crassifolia and T. nudipes) (Dipterygeae subclade) (Figures 2H, 3C,E,H, 6B, LN and 7B, D, F). The lumen shape varies from spherical to oval between Dipteryx and Taralea (compare Figures 6B and 7B).

A secretory duct occurs embedded in the anther apex and extends distally between thecae of the Myroxylon species (Amburaneae subclade) (Figure 3K). The lumen shape is elongated (Figure 3K).

In the remaining species, the anther apex does not show a secretory cavity or a secretory duct (Figures 2B, E, O, 3B, M, 4D, F, J, L, 5B, D, F, H, 7H, M and 8B, D, H, F, J, L, N).

Phenolic idioblasts are found in the anther apex of Petaladenium, six Alexa species, two Monopteryx species, and in Swartzia langsdorffii (Figures 3M, 4D, F, L, 7M and 8J, L). These species do not have cavities or ducts.

2.3. Distribution of Secretory Cavities in the Anther of the Amburaneae and Dipterygeae Subclades

By tracing the evolutionary history of the character “occurrence of secretory cavity/duct in the anther”, it was inferred that the presence of cavity/duct in the anther was acquired in most representatives of the Dipterygeae subclade and some of the Amburaneae subclade (Figure 9).

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Figure 9. Representation of the modified LPGW cladogram (2017) for the ADA clade, showing the reconstruction of the character history in the occurrence of glands in anthers of the group. The red color represents the presence of glands in the lineage, the blue color represents the absence of glands, and the gray color represents doubt about the occurrence of the glands. The reconstruction indicates that the presence of glands has two plausible hypotheses: the appearance of the glands in the subclades (Amburaneae and Dipterygeae) or their loss in some representatives of Amburaneae and Dipterygeae and in all representatives of Angylocalyceae.
2.4. Correlations between Character-States

The pairings of the character-state reconstructions related to the occurrence of cavity/duct at the anther apex vs. shape of the anther apex indicate a positive correlation between these two characters (Figure 10). However, it is noteworthy that in 5 of the 20 species (Myrocarpus emarginatus, M. fastigiatus, M. frondosus, Diploxyx polyphylla, and Taralea cordata), this correlation was negative that is, even without a prominent anther apex, there was a secretory cavity in the anther.

![Figure 10. Modified LPGW cladogram (2017) for the ADA clade showing the mirroring of the character state reconstruction for “secretory cavity/duct occurrence” vs. “shape of the anther apex.”](image-url)
Figure 10. Modified LPGW cladogram (2017) for the ADA clade showing the mirroring of the character state reconstruction for “secretory cavity/duct occurrence” vs. “shape of the anther apex”.

Figure 11. Modified LPGW cladogram (2017) for the ADA clade showing the mirroring of the character state reconstruction for “lumen shape” vs. “position of the secretory cavity/duct in the anther”.

3. Discussion

Our study highlights how glandular appendages occur in the anther of the ADA clade and provides an opportunity to clarify their enigmatic evolutionary history within early-branching papilionoids. The glandular appendage in the anther has been previously reported in the *Dipteryx alata* and *Pterodon pubescens* species of the Dipterygeae subclade [7,13], and also occurring in the species of the Amburaneae subclade.

3.1. Distribution and Location of Secretory Cavities/Ducts

The large ADA clade comprises morphologically eclectic genera with a diverse occurrence and structure of a glandular appendage in the anther, and the shape and location of this gland in the anther. Among 50 species analyzed within the ADA clade, 21 exhibit secretory cavities, two secretory ducts, and nine phenolic idioblasts, for a total of 64% species with a secretory structure in the anther apex.

Anatomical analyses of the anther in a longitudinal section showed that in most species analyzed, the gland is a secretory cavity with lumen shapes ranging from spherical to oval (see Table 1). *Myroxylon balsamum* and *M. peruiferum*, two species of the Amburaneae subclade, are the exceptions. The gland in the anther exhibits an elongated lumen so that, the term, secretory duct, becomes more appropriate. Variations in the lumen shape and, consequently, the difficulties generated in the typification of the gland, have been extensively explored in the literature, especially in studies with the leaf [15,20] and the stem [14,15,18,19].

The glandular appendage in the anther is most evident in the species of *Pterodon*, *Dipteryx* [7,13], present study, and *Cordyla*, which have secretory cavities with a spherical lumen, except for *D. polyphylla* in which the appendix is not very prominent, and the cavities have oval lumens. Similarly, *Taralea cordata*, *T. crassifolia*, and *T. nudipes* exhibit non-prominent anther appendages and contain a secretory cavity with an oval lumen, which is found in the region between the two thecae. In these species, the lumen of the cavity has the same shape as described for the vegetative organs of *Taralea oppositifolia* [15].

It is interesting to note that the secretory cavity located at the apex of the anther is subepidermal, and the epidermis cells have phenolic compounds see [13]. In *Monopteryx*
Plants 2022, 11, 835 and some species of Alexa, the cells of the anther appendix also exhibit phenolic content, although they do not have a secretory cavity. The appendix composed of phenolic cells that the anthers exhibit must be related to the floral structure, which is non-papilionaceous in the species of Alexa and Monopteryx. In these cases, the wing petals are reduced, the keel petals are united and opened, and the free stamens are exposed. The presence of phenolic compounds in the anther apex may be associated with the defense against herbivory or UV radiation since anthers are not protected by the petals as in a papilionaceous flower [34–37].

An interesting fact is the association between a secretory cavity in the anther and the leaf. Secretory cavities are present at the anther apex of closely related species such as Cordyla africana, C. haraka, C. madagascariensis, Myrocarpus emarginatus, M. fastigiatus, M. fndonosis, and Myroperum frutescens. In contrast, Myroxyylon balsamum, and M. peruiferum exhibit a secretory duct at the anther apex. The secretory cavity or duct is absent in the species of Amburana, Dussia, Mildbraediodendron, and Petaladenium (Amburaneae subclade), and in the species of Alexa, Angylocalyx, Castanospermum, and Xanthocercis (Angylocalyceae subclade). The presence of a secretory cavity and ducts in the leaflet is shared by species of Myrocarpus (M. emarginatus, M. fastigiatus, M. fndonosus, M. leprosus, M. venezuelensis), Myroperum (M. frutescens, M. sousanum), Myroxyylon (M. peruiferum and M. balsamum) and Cordyla (C. haraka, C. africana and C. madagascariensis [20,21]) which demonstrates that these structures also occur in the floral organs of these species.

3.2. Evolutionary History of the Presence of a Secretory Cavity in the Anther of the ADA Clade

Our hypotheses were postulated to explain the occurrence of glands in anthers of the ADA clade species and have two robust explanations: (1) the appearance of the anther glands in the Amburaneae and Dipterygeae subclades or (2) their loss in some species of Amburaneae and Dipterygeae and all species of Angylocalyceae (see Figure 9).

Considering that the secretory cavities are present in other genera of the Amburaneae subclade, it is concluded that they are not a synapomorphy of the Dipterygeae subclade, as suggested by Leite et al. [7].

In the Dipterygeae subclade, our data suggest that secretory cavities may have been acquired in Dipteryx + Pterodon and some species of Tarae. A phenolic glandular appendix may have been acquired in Monopteryx, a sister group of Dipteryx, Pterodon, and Tarae (see distribution in Figure 9). The glandular appendix with a phenolic epidermis in Dipteryx, Pterodon, and Tarae could be a remnant of the Monopteryx phenolic appendage.

The Amburaneae subclade is remarkable because of its high level of floral diversity, production of coumarins (Amburana), red resin from bark and twigs (Dussia), balsams (Myrocarpus, Myroperum, Myroxyylon), and punctate glandular leaves of several genera (Cordyla, Mildbraediodendron, Myrocarpus, Myroperum, and Myroxyylon) [27,38,39]. The presence of glands at the anther apex is also noteworthy. Our data suggest that secretory cavities/ducts may have been acquired in the well-supported clades Myroxyylon + Myrocarpus (non-papilionaceous flowers) and Myroperum (papilionaceous flower). Another interesting fact is the presence of a secretory duct only in Myroxyylon, and therefore, an autapomorphy of Myroxyylon. Although Cordyla and Mildbraediodendron have a swartzioid-like floral morphology [12,38], the flowers have an entire calyx, no petals, and numerous free stamens, only Cordyla exhibits a secretory cavity in the anther apex. In contrast, the genus Amburana, sister to Cordyla + Mildbraediodendron, does not have a secretory cavity in the anther and has a one-petalled corolla and 10 free stamens. The absence of a secretory cavity in the anther apex of the genera Petaladenium and Dussia reflects their positioning in the phylogenetic tree as sister genera. They also present a papilionaceous corolla and basally united stamens [10]. A peculiar characteristic of Petaladenium urceoliferum is its wing petals with glands, while in Dussia (its sister genus), the glands are found on the bract and bracteoles [9], although there are no anatomical studies on the composition of these structures.

In the Angylocalyceae subclade, our data suggest that secretory cavities were not acquired in the genera Alexa, Angylocalyx, Castanospermum, and Xanthocercis, defining them
as a sister group of the Amburaneae and Dipterygeae subclades. An appendix producing phenolic compounds may have been acquired in some *Alexa* species, a sister group of *Castañospermum*, both with similar floral morphology, non-papilionaceous corolla, and 10 free stamens [12]. *Angylocalyx* and *Xanthocercis*, sister genera, exhibit a distinct floral morphology [12].

Other relevant data are the occurrence of secretory cavity/duct vs. anther appendage shape (see Figure 10) and lumen shape vs. position of cavity/duct in the anther (see Figure 11). It is likely that the anthers with a prominent apex also exhibit secretory cavities with a spherical lumen, different from those with a non-prominent apex, which exhibit secretory cavities more internalized in the anther, between the thecae, with the lumen being oval. Thus, we suppose that the shape of the anther is related to the lumen shape in species with a secretory cavity in the anther (see Figures 10 and 11). Another interesting result is the lumen shape in the clades *Myrocarpus* + *Myrospermum*, and *Myroxylon*. *Myrocarpus* + *Myrospermum* exhibit secretory cavities more internalized in the anther between the thecae, with an oval lumen and a prominent/non-prominent apex, respectively, different from *Myroxylon*, which exhibits a secretory duct with a rounded lumen in its apical portion and is elongated in the lower portion between the thecae following the shape of the anther apex.

The secretory cavities of *Dipteryx* (except *D. polyphylla*) and *Pterodon* are anatomically more similar to each other than to those of *Taralea cordata*, *T. crassifolia* and *T. nudipes*, confirming previous data obtained about the flower [7], leaf, stem [15] and leaflets [16] and corroborating the phylogeny data (see LPGW [1]). This fact reflects the topology of a phylogenetic tree in the subclades, which places them as sister groups see [1,12,22].

The inclusion of other species with a non-papilionaceous corolla from the Swartzieae clade (*Ateleia glazioveana*, *A. guaraya*, *Candolleodendron brachystachyum*, *Cyathostegia mathewsii*, *Swartzia langsdorffii*) and genistoid clade (*Uleanthus erythrinoides*) [1] suggests that the presence of secretory cavities/ducts in the anther apex may be restricted to the members of the subclades Amburaneae and Dipterygeae. There are no reports on cavities secretory/duct in the anthers apex of other papilionoid legumes. *Hymenaea verrucosa* Gaertn. (subfamily Detarioideae) exhibits a secretory cavity in the connective region (ventral region) of the anthers [40]; and in *Stryphnodendron adstringens*, *Tetrapleura tetraptera*, *Adenanthera pavonine*, and *Pentaclethra macroloba* (Mimosoid clade, subfamily Caesalpinioideae) such anther glandular appendages can comprise secretory emergences [41].

3.3. Evolutionary Significance of the Corolla, Type of Androecium vs. Presence of a Secretory Cavity/Duct in the Anther

Our data plotted in phylogeny have not yet made it possible to correlate the type of corolla and androecium with a secretory cavity/duct in the anther. The presence of a secretory cavity/duct has been reported in species with apetalous, non-papilionaceous flowers with numerous free stamens, including *Cordyla africana*, *C. haraka* and *C. madagascariensis*. *Myrocarpus emarginatus*, *M. fastigiatus*, and *M. frondosus* exhibit five undifferentiated petals with 10 basally united stamens, *Myrospermum frutescens* exhibits a papilionaceous corolla with 10 free stamens; *Myroxylon balsamum* and *M. peruiferum* exhibit a non-papilionaceous corolla with the widely oval banner and 10 free stamens. Therefore, a correlation with pollination seems plausible.

In *Dipteryx alata* and *Pterodon pubescens*, species of the Dipterygeae subclade, the anther glands consist of a cavity secreting sticky substances (oleoresins and polysaccharides) that play a key role during the flower’s lifespan by aggregating pollen grains and attaching them to the floral visitor’s body, besides maximizing the pollen release mechanism that is intermediate between the valvular and the explosive [13]. The same mechanism probably occurs in *Myrospermum frutescens* because these species are pollinated by insects [42]; *Myrocarpus frondosus* has a non-papilionaceous corolla but is pollinated by insects [43], probably aggregating pollen grains.

In *Myroxylon peruiferum*, the ducts secrete a substance that aggregates pollen grains; however, when in contact with air, the resinous content secreted together with the pollen...
grains hardens, probably acting on pollination by sunbirds (personal observation). The same must occur in *Cordyla africana*, which has petaliferous flowers rich in nectar that are pollinated by sunbirds [44].

Species with phenolic idioblasts at the anther apex, such as those of *Alexa* and *Monopterix*, are ornithophilous [27] and entomophilous [45], respectively. There are also reports of sphingophily [46,47] and chiropterophily [48] for *Alexa*. The lack of information on the pollination biology for these species makes it difficult to understand the function of these glands in early-diverging papilionoids.

However, analyzing the corolla shape and the presence of a secretory structure in the anther concerning the pollinator (insect or bird), probably the corolla type associated with the material exuded by the secretory structure at the anther apex at different proportions found in each species acts by favoring the different pollinators.

3.4. Outlook

We studied the evolution of the anther’s glandular appendage in early-diverging papilionoid genera, reporting this condition in a large number of species.

The diversity of the subfamily Papilionoideae expressed in terms of floral morphology, such as loss of petals, undifferentiated petals, numerous free stamens, entire calyx in bud, and radially symmetric flowers [5,8,49–55] is particularly common among the early-diverging papilionoid genera [5–10]. Additionally, secretory cavities/ducts in the anther in some species of the Dipterygeae and Amburaneae subclades [7,13], present study, is a unifying character-state for these groups. This condition has not been previously reported for representatives of the most recent divergent papilionoids.

4. Materials and Methods

4.1. Sampling Plant Material

We examined at least one species in each genus of the Amburaneae (*Amburana*, *Cordyla*, *Dussia*, *Mildbraediadendron*, *Myrcaropus*, *Myrospermum*, *Myroxylon*, and *Petaladenium*), Angylocalyceae (*Alexa*, *Angylocalyx*, *Castanospermum*, and *Xanthocercis*), and Dipterygeae subclades (*Dipteryx*, *Pterodon*, and *Monopterix*). In addition, we included six species with non-papilionoid flowers outside the ADA clade to map the character of other distant taxa (Table S1).

Samples of flowers and floral buds (1 or/and 2) were obtained from herbarium specimens (Table S1) and treated with 2% KOH solution for 2 h, washed several times in distilled water [56], and stored in 70% ethanol. The anthers were removed and prepared for observations by scanning electron microscopy (SEM) and light microscopy (LM).

4.2. Scanning Electron Microscopy

For SEM analysis, anthers were critical point dried in a Balzers CPD 030 dryer (Balzers, Liechtenstein), mounted on aluminum stubs with colloidal carbon, coated with gold in a Bal Tec SCD 050 sputter coater, and observed with a Jeol JSM 6610LV scanning electron microscope (Tokyo, Japan).

4.3. Light Microscopy

For LM analysis, anthers were embedded in historesin [57] and longitudinally sectioned (2–3 µm thick) with a rotary microtome (Leica RM 2245, Wetzlar, Germany). Sections were stained with 0.05% toluidine blue in phosphate buffer, pH = 6.8 [58] and photographed with a light microscope (Leica DM5000 B) coupled to a digital camera (Leica DFC295).

4.4. Phylogenetic Analysis

The evolution of the glands in anthers from the ADA clade was investigated based on a recent phylogenetic hypothesis [1]. The characters selected to compose the data matrix were the absence or presence of secretory cavity/ducts, anther appendage shape (prominent, not prominent), anther appendix position (apical, distal, apical/distal), lumen
shape (spherical, oval, elongated), and absence or presence of phenolic compound at the apex of the anther (Table 2). We used the Mesquite program [59] to map the selected characters (Table 2) on the RAXML tree obtained by the Legume Phylogeny Working Group (LPWG) [1]. For this purpose, we chose “trace character history”, selecting the option “parsimony ancestral states”.

**Table 2.** Morphological characteristics evaluated in taxa of the Amburaneae, Angylocalyceae, Dipterogoeae subclades, *Ateleia*, *Candolleodendron*, *Cyathostegia*, *Swartizia* and *Uleanthus*.

| Variables | 1. Number of sepals on the mature flower | 0) five, (1) four, (2) three |
|-----------|-----------------------------------------|-----------------------------|
|           | 2. Number of petals on the mature flower | (0) zero, (1) one, (2) four, (3) five |
|           | 3. Number of stamens | (0) up to 10, (1) 10, (3) numerous (= over 10) |
|           | 4. Type of stamen connation | (0) free, (1) fused at the base, (2) monadelphous |
|           | 5. Occurrence of cavity/canals at the apex of the anther | (0) absence, (1) presence |
|           | 6. Position of cavity/canals in the anther | (0) apical, (1) distal, (2) apical and distal, (3) absence of position |
|           | 7. Anther appendage shape | (0) prominent, (1) not prominent |
|           | 8. Phenolic compound tissue at the apex of the anther | (0) absence, (1) presence |
|           | 9. Lumen shape (apex of anther in longitudinal section) | (0) spherical, (1) oval, (2) elongate, (3) not applicable |

**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/plants11070835/s1, Table S1: Species of Amburaneae, Angylocalyceae, Dipterogoeae subclades and outgroup. IAN = Herbário do Embrapa Amazônia Oriental; INPA = Herbário do Instituto Nacional de Pesquisas da Amazônia; KEW = Herbarium Royal Botanic Gardens; MNHN = Muséum national d’Histoire naturelle; MG = Herbário do Museu Paraense Emílio Goeldi; NYBG = The New York Botanical Garden; RB = Herbário do Jardim Botânico do Rio de Janeiro; RBSpirit = Herbário do Jardim Botânico do Rio de Janeiro; SPFR = Herbário do Departamento de Biologia da Faculdade de Filosofia Ciências e Letras de Ribeirão Preto da Universidade de São Paulo; UEC = Herbário da Universidade Estadual de Campinas; US (Smithsonian Institution).

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References

1. LPWG (The Legume Phylogeny Working Group). A new subfamily classification of the Leguminosae based on a taxonomically comprehensive phylogeny. *Taxon* 2017, 66, 44–77. [CrossRef]

2. LPWG (The Legume Phylogeny Working Group). Legume phylogeny and classification in the 21st century: Progress, prospects and lessons for other species-rich clades. *Taxon* 2013, 62, 217–248. [CrossRef]

3. Lavin, M.; Pennington, R.T.; Kiltigaard, B.B.; Sprent, J.I.; Lima, H.C.; Gasson, P.E. The dalbergioid legumes (Fabaceae): Delimitation of a pantropical monophyletic clade. *Am. J. Bot.* 2001, 88, 503–533. [CrossRef]

4. Schrire, B.D.; Lavin, M.; Lewis, G.P. Global distribution patterns of the Leguminosae: Insights from recent phylogenies. In *Plant Diversity and Complexity Patterns: Local, Regional and Global Dimensions*; Biologiske Skrifter 55; Friis, I., Balslev, H., Eds.; Kgl. Danske Videnskabernes Selskab: Copenhagen, Denmark, 2005; pp. 375–422.

5. Tucker, S.C. Floral ontogeny in Sophoreae (Leguminosae: Papilionoideae). I. *Myrocarpus (Myrocarpus)* Group and *Castanospermum* (Angolycahy Group). *Am. J. Bot.* 1993, 80, 65–75. [CrossRef]

6. Prenner, G. New Aspects In Floral Development of Papilionoideae: Initiated but Suppressed Bracteoles and Variable Initiation of Sepals. *Ann. Bot.* 2004, 93, 537–545. [CrossRef] [PubMed]

7. Leite, V.G.; Mansano, V.F.; Teixeira, S.P. Floral ontogeny in Dipterygeae (Fabaceae) reveals new insights into one of the earliest branching tribes in papilionoid legumes. *Bot. J. Linn. Soc.* 2014, 174, 529–550. [CrossRef]

8. Leite, V.G.; Teixeira, S.P.; Mansano, V.F.; Prenner, G. Floral development of the early branching papilionoid legume *Amburana cearensis* (Leguminosae) reveals rare and novel characters. *Int. J. Plant Sci.* 2015, 176, 94–106. [CrossRef]

9. Prenner, G.; Cardoso, D.; Zartman, C.E.; Queiroz, L.P. Flowers of the early-branching papilionoid legume *Petaladenium urceoliferum* display morphological and ontogenetic features. *Am. J. Bot.* 2015, 102, 1780–1793. [CrossRef]

10. Cardoso, D.; São-Mateus, W.M.B.; Cruz, D.T.; Zartman, C.E.; Komura, D.L.; Kite, G.; Prenner, G.; Wieringa, J.J.; Clark, A.; Lewis, G.; et al. Filling in the gaps of the papilionoid legume phylogeny: The enigmatic Amazonian genus *Petaladenium* is a new branch of the early-diverging Amburanae clade. *Mol. Phylogenetics Evol.* 2015, 84, 112–124. [CrossRef]

11. Lewis, G.; Schrire, B.; Mackinder, B.; Lock, M. *Legumes of the World*; Royal Botanic Gardens, Kew: London, UK, 2005.

12. Cardoso, D.; Pennington, R.T.; Queiroz, L.P.; Boatwright, J.S.; Van Wyk, B.-E.; Wojciechowski, M.F.; Lavin, M. Reconstructing the deep-branching relationships of the papilionoid legumes. *S. Afr. J. Bot.* 2013, 89, 58–75. [CrossRef]

13. Leite, V.G.; Mansano, V.F.; Pansarin, E.; Teixeira, S.P. Presence of the anther gland is a key feature in pollination of the early-branching papilionoid legumes *Pterodon alata* and *Pterodon pubescens* (Leguminosae). *Plant Biol.* 2019, 21, 1016–1023. [CrossRef]

14. Rodrigues, T.M.; Machado, S.R. Anatomy comparada do pulvino, pecíolo e raque de *Pterodon pubescens* Benth (Fabaceae-Faboideae). *Acta Bot. Bras.* 2004, 18, 381–390. [CrossRef]

15. Palermo, F.; Teixeira, S.P.; Mansano, V.F.; Leite, V.G.; Rodrigues, T.M. Secretory spaces in species of clade Dipterygeae (Leguminosae, Papilionoideae). *Acta Bot. Bras.* 2017, 31, 374–381. [CrossRef]

16. Silva, N.F.; Arruda, R.C.O.; Alves, F.M.; Sartori, A.L.B. Leaflet anatomy of the Dipterygeae clade (Faboideae: Fabaceae): Evolutionary implications and systematics. *Bot. J. Linn. Soc.* 2018, 187, 99–117. [CrossRef]

17. Rodrigues, T.M.; Santos, D.C.; Machado, S.R. The role of the parenchyma sheath and PCD during the development of oil cavities in *Pterodon pubescens* (Leguminosae-Papilionoideae). *Comptes Rendus Bot.* 2011, 334, 535–543. [CrossRef] [PubMed]

18. Paiva, E.A.S.; Oliveira, D.M.T.; Machado, S.R. Anatomy and ontogeny of the pericarp of *Pterodon emarginatus* Vogel (Fabaceae, Faboideae), with emphasis on sepalory ducts. *An. Acad. Bras. Ciênc.* 2008, 80, 455–465. [CrossRef] [PubMed]

19. Rodrigues, T.M.; Machado, S.R. Oil glands in *Pterodon pubescens* Benth. (Leguminosae-Papilionoideae): Distribution, structure and secretion mechanisms. *Int. J. Plant Sci.* 2012, 173, 984–992. [CrossRef]

20. Sartori, A.L.B.; Tozzi, A.M.G.A. Comparative leaflet anatomy in *Myrcarpus Allemão, Myroxyilon L.* f and *Myxsypermum Jacq.* (Leguminosae-Papilionoideae-Sophoreae) species. *Bot. J. Linn. Soc.* 2002, 140, 249–259. [CrossRef]

21. Bento, J.P.S.P.; Scresmin-Dias, E.; Alves, F.M.; Mansano, V.F.; Sartori, A.L.B. Phylogenetic implications of the anatomical study of the Amburanae clade (Fabaceae: Faboideae). *Bot. J. Linn. Soc.* 2020, 192, 69–83. [CrossRef]

22. Cardoso, D.; Queiroz, L.P.; Pennington, R.T.; Lima, H.C.; Fonty, E.; Wojciechowski, M.F.; Lavin, M. Revisiting the phylogeny of Papilionoid Legumes: New insights from comprehensively sampled early-branching lineages. *Am. J. Bot.* 2012, 99, 1991–2013. [CrossRef] [PubMed]

23. Seleme, E.P.; Lewis, G.P.; Stirton, C.H.; Sartori, A.L.B.; Mansano, V.F. A Taxonomic review and a new species of the south American woody genus *Amburana* (Leguminosae, Papilionoideae). *Phytotaxa* 2015, 212, 249–263. [CrossRef]

24. Rudd, V.E. *The Genus Dussia* (Leguminosae): United States National Museum Contributions from the United States National Herbarium, Smithsonian Institution: Washington, DC, USA, 1963; pp. 247–277.

25. Cowan, R.S. *Swartzieae*. In *Advances in Legume Systematics, Part 1*; Polhill, R.M., Raven, P.H., Eds.; Royal Botanic Gardens, Kew: London, UK, 1981; pp. 209–212.

26. WFO. *Mildbraediodendron excelsum* Harms. Available online: http://www.worldfloraonline.org/taxon/wfo-0000171559 (accessed on 7 February 2022).

27. Polhill, R.M. *Sophoreae*. In *Advances in Legume Systematics, Part 1*; Polhill, R.M., Raven, P.H., Eds.; Royal Botanic Gardens, Kew: London, UK, 1981; pp. 213–230.

28. Sartori, A.L.B.; Tozzi, A.M.G.A. Revisão taxonómica do *Myxycarpus Allemão* (Leguminosae, Papilionoideae, Sophoreae). *Acta Bot. Bras.* 2004, 18, 521–535. [CrossRef]
