

Molecular phylogenetic and morphological support for the recognition of *Friesodielsia lalisae* (Annonaceae), a new species from S Thailand

Authors: Damthongdee, Anissara, Khunarak, Natthanon, Kaeokula, Suphaloek, Saengpho, Chanwut, Wiya, Chattida, et al.

Source: *Willdenowia*, 53(1-2) : 45-55

Published By: Botanic Garden and Botanical Museum Berlin (BGBM)

URL: <https://doi.org/10.3372/wi.53.53103>

BioOne Complete (complete.BioOne.org) is a full-text database of 200 subscribed and open-access titles in the biological, ecological, and environmental sciences published by nonprofit societies, associations, museums, institutions, and presses.

Your use of this PDF, the BioOne Complete website, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at www.bioone.org/terms-of-use.

Usage of BioOne Complete content is strictly limited to personal, educational, and non - commercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

BioOne sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.

Molecular phylogenetic and morphological support for the recognition of *Friesodielsia lalisae* (Annonaceae), a new species from S Thailand

Anissara Damthongdee¹, Natthanon Khunarak¹, Suphaloek Kaeokula^{1,2}, Chanwut Saengpho¹, Chattida Wiya¹, Phasit Ue-aree¹, Abdulromea Baka³, Kithisak Aongyong³ & Tanawat Chaowasku¹

Version of record first published online on 10 April 2023 ahead of inclusion in April 2023 issue.

Abstract: Thirteen species of *Friesodielsia* Steenis (*Annonaceae*) and 11 representatives of related genera were included in molecular phylogenetic analyses using up to six plastid DNA regions (*psbA-trnH*, *trnL-trnF* intergenic spacers; *trnL* intron; *matK*, *ndhF*, *rbcL* exons). The results support the recognition of a new species, *F. lalisae* Damth., Baka & Chaowasku from Narathiwat, S Thailand, as belonging to one of the two major clades of *Friesodielsia*. The members of this clade show outer petals that separate at anthesis and have a concave basal portion, while members in the other major clade exhibit outer petals that separate early during developmental stages and have a flat base. *Friesodielsia lalisae* is described and illustrated. It is most morphologically similar to *F. argentea* (J. Sinclair) Steenis and *F. glauca* (Hook. f. & Thomson) Steenis but differs from the two by having dissimilar sepal shape, higher proportion of inner petal to outer petal length and longer inner petals. The new species additionally differs from the former by having different leaf base and from the latter by having denser indumentum on young twigs and shorter flowering pedicels. Narathiwat, a province to which the new species is endemic, seems to be one of the most underexplored areas in Thailand as evidenced by a number of species described based on recent material. The conservation status of the new species is provisionally assessed as Critically Endangered.

Keywords: *Annonaceae*, *Friesodielsia*, molecular phylogeny, Narathiwat, new species, systematics, taxonomy, Thailand, *Uvarieae*

Article history: Received 9 December 2022; peer-review completed 7 February 2023; received in revised form 2 March 2023; accepted for publication 10 March 2023.

Citation: Damthongdee A., Khunarak N., Kaeokula S., Saengpho C., Wiya C., Ue-aree P., Baka A., Aongyong K. & Chaowasku T. 2023: Molecular phylogenetic and morphological support for the recognition of *Friesodielsia lalisae* (*Annonaceae*), a new species from S Thailand. – *Willdenowia* 53: 45–55. <https://doi.org/10.3372/wi.53.53103>

Introduction

The pantropical family *Annonaceae* consists of trees, shrubs and lianas classified in 109 genera (Jaikhamseub & al. 2022 plus subsequent publications: Cheek & al. 2022 [*Lukea* Cheek & Gosline, a recently established genus]; Couvreur & al. 2022 [acceptance of *Dennettia* Baker f.] and about 2550 species (Couvreur & al. 2022). According to Bangkomnate & al. (2021), the liana genera *Pyramidanthe* Miq. and *Mitrella* Miq. are considered congeneric, with the acceptance of the former name. *Friesodielsia* Steenis (*Annonoideae*, *Uvarieae*), comprising c. 48 species (Turner 2018; Saunders & al. 2020; Leeratiwong & al. 2021a, 2023), is also one of the genera with a liana habit (Turner 2012; Guo & al. 2017a). The genus underwent recent realignments based on molecular phylogenetic inferences, with the recombination of most African species names into *Monanthes* Baill. and a few into *Afroguatteria* Boutique and *Sphaerocoryne* (Boerl.) Scheff. ex Ridl.; therefore, its distribution is restricted to tropical Asia plus New Guinea (Guo & al.

2017a; Saunders & al. 2020). In addition to the liana habit, *Friesodielsia* can also be circumscribed by the presence of (1) a more or less glaucous lower leaf surface, (2) a usually obvious pair of glands at the base of each leaf blade, (3) initially terminal inflorescences developing to become internodal (or sometimes leaf-opposed), (4) inner petals that are smaller than the outer petals and remain connivent at maturity and (5) usually single-seeded monocarps (e.g. Leeratiwong & al. 2021a). In *Uvarieae*, *Friesodielsia* and three related genera: *Dasymaschalon* (Hook. f. & Thomson) Dalla Torre & Harms, *Desmos* Lour. and *Monanthes* are clustered in a strongly supported clade; *Monanthes* is then a sister group of a strongly supported clade composed of the other three genera (e.g. Guo & al. 2017a, 2017b). In Thailand, 18 species of *Friesodielsia* were reported (Leeratiwong & al. 2023), but based on preliminary observations on recently collected specimens, several undescribed species seem to exist. In this study we determine the taxonomic status of an unidentifiable gathering from Narathiwat, one of the southernmost and inadequately explored prov-

1 Herbarium, Department of Biology, Faculty of Science, Chiang Mai University, 239 Huay Kaew Rd., Chiang Mai 50200, Thailand.

2 Current address: NYU Grossman School of Medicine, New York, NY 10016, U.S.A.

3 Independent Research Group on Plant Diversity in Thailand, 13 Moo 1, Khaonoi, Sichon, Nakhon Si Thammarat 80120, Thailand.

Author for correspondence: Tanawat Chaowasku, tanawat.chaowasku@cmu.ac.th

inces of Thailand by morphological investigations and comparisons in combination with molecular phylogenetic analyses.

Material and methods

Phylogenetic reconstructions

The ingroup was composed of 24 accessions, divided into four genera: *Dasymaschalon* (four accessions representing four species), *Desmos* (five accessions representing five species), *Friesodielsia* (13 accessions representing 12 species plus an unidentifiable accession from Narathiwat, S Thailand [= *Friesodielsia* sp. TH]) and *Monanthotaxis* (two accessions representing two species). Outgroups consisted of two species in the tribe *Uvarieae*: *Pyramidanthella elegans* (Hook. f. & Thomson) Bangk. & Chaowasku and *Uvaria dasoclema* L. L. Zhou & al. The information of voucher specimens and GenBank accession numbers used in this study is shown in Appendix 1. Up to six plastome regions (*psbA-trnH*, *trnL-trnF* intergenic spacers; *trnL* intron; *matK*, *ndhF*, *rbcL* exons) were included. The methods for DNA extraction, amplification and sequencing used in the present study, including primer information, followed Chaowasku & al. (2018a, 2018b, 2020). Sequences were edited using the Staden package (Staden & al. 2000) and the data matrix was aligned by Multiple Sequence Comparison by Log-Expectation (MUSCLE; Edgar 2004) in MEGA7 (with default settings; Kumar & al. 2016). The aligned data matrix was subsequently manually checked and realigned where necessary using the similarity criterion (Simmons 2004). In total, there were 5496 nucleotide plus six binary-coded indel characters. The simple method for indel coding of Simmons & Ochoterena (2000) was used, with the emphasis on non-autapomorphic and less homoplastic indel structures.

Parsimony analysis was performed in TNT version 1.5 (Goloboff & Catalano 2016). All characters were equally weighted and unordered. The setting concerning collapsing rules was set to “max. length = 0”. Incongruence among plastid DNA regions was evaluated by analysing each region individually to detect if there was any significant topological conflict (e.g. Wiens 1998). Most parsimonious trees were generated by a heuristic search of the combined data, with 9000 replicates of random sequence addition, saving 10 trees per replicate and using the tree bisection and reconnection (TBR) branch-swapping algorithm. Clade support was measured by symmetric resampling (SR; Goloboff & al. 2003), with default change probability ($P=33$). Two hundred thousand replicates were run, each with four replicates of random sequence addition, saving four trees per replicate. A clade with SR $\geq 85\%$, 70–84% or 50–69% was considered strongly, moderately or weakly supported, respectively.

Maximum likelihood analysis was carried out in IQ-TREE version 2.1.2 (Minh & al. 2020) under partition

models (Chernomor & al. 2016) employed with the “-p” command, whereas Bayesian Markov chain Monte Carlo (MCMC; Yang & Rannala 1997) phylogenetic analysis was conducted in MrBayes version 3.2.7a (Ronquist & al. 2012). Both methods of phylogenetic reconstruction were analysed via the CIPRES Science Gateway version 3.3 (Miller & al. 2010). The aligned data matrix was divided into five partitions based on DNA-region identity (the *trnL* intron and adjacent *trnL-trnF* intergenic spacer were combined into a single partition). The most suitable model of sequence evolution for each DNA partition was selected using the Akaike Information Criterion (AIC; Akaike 1974) scores calculated in jModelTest version 2.1.10 (Darriba & al. 2012), with the following selections: +F, +G (nCat 4), ML optimized (base tree for likelihood calculations) and Best (base tree search). The General Time Reversible (GTR; Tavaré 1986) substitution model with a gamma distribution for among-site rate variation was selected for *matK* and *ndhF* partitions, whereas the Hasegawa-Kishino-Yano (HKY; Hasegawa & al. 1985) substitution model with a gamma distribution for among-site rate variation was chosen for the remaining partitions (*psbA-trnH*, *rbcL* and *trnL-trnF* [= *trnL* intron + *trnL-trnF* intergenic spacer]). In the maximum likelihood analysis, the model “JC2+FQ+ASC” was selected using the corrected AIC scores for the binary indel partition. Clade support was assessed by a non-parametric bootstrap resampling (BS; Felsenstein 1985) with 2000 replicates. A clade with BS $\geq 85\%$, 70–84% or 50–69% was considered strongly, moderately or weakly supported, respectively. In the Bayesian analysis, the “coding=variable” setting was assigned to the binary indel partition, which was implemented under a simple F81-like model without a gamma distribution for among-site rate variation. Four independent runs, each with four MCMC chains, were simultaneously performed; each run was set for 10 million generations. The default prior settings were used except for the prior parameter of rate multiplier (“ratepr” [=variable]). The temperature parameter was set to 0.08. Trees and all parameter values were sampled every 1000th generation. Convergence was assessed by checking the standard deviation of split frequencies of the runs with values < 0.01 interpreted as indicative of a good convergence and by checking for adequate effective sample sizes (ESS > 200) using Tracer version 1.7.1 (Rambaut & al. 2018). The first 25% of all trees sampled were removed as burn-in and the 50% majority-rule consensus tree was produced from the remaining trees. A clade with posterior probabilities (PP) ≥ 0.95 , 0.9–0.94 or 0.5–0.89 was considered strongly supported, weakly supported or unsupported, respectively.

Morphology

The morphological information of relevant *Friesodielsia* species was derived from literature (Sinclair 1955; Leeratiwong & al. 2021a; Johnson & al. 2022) as well as

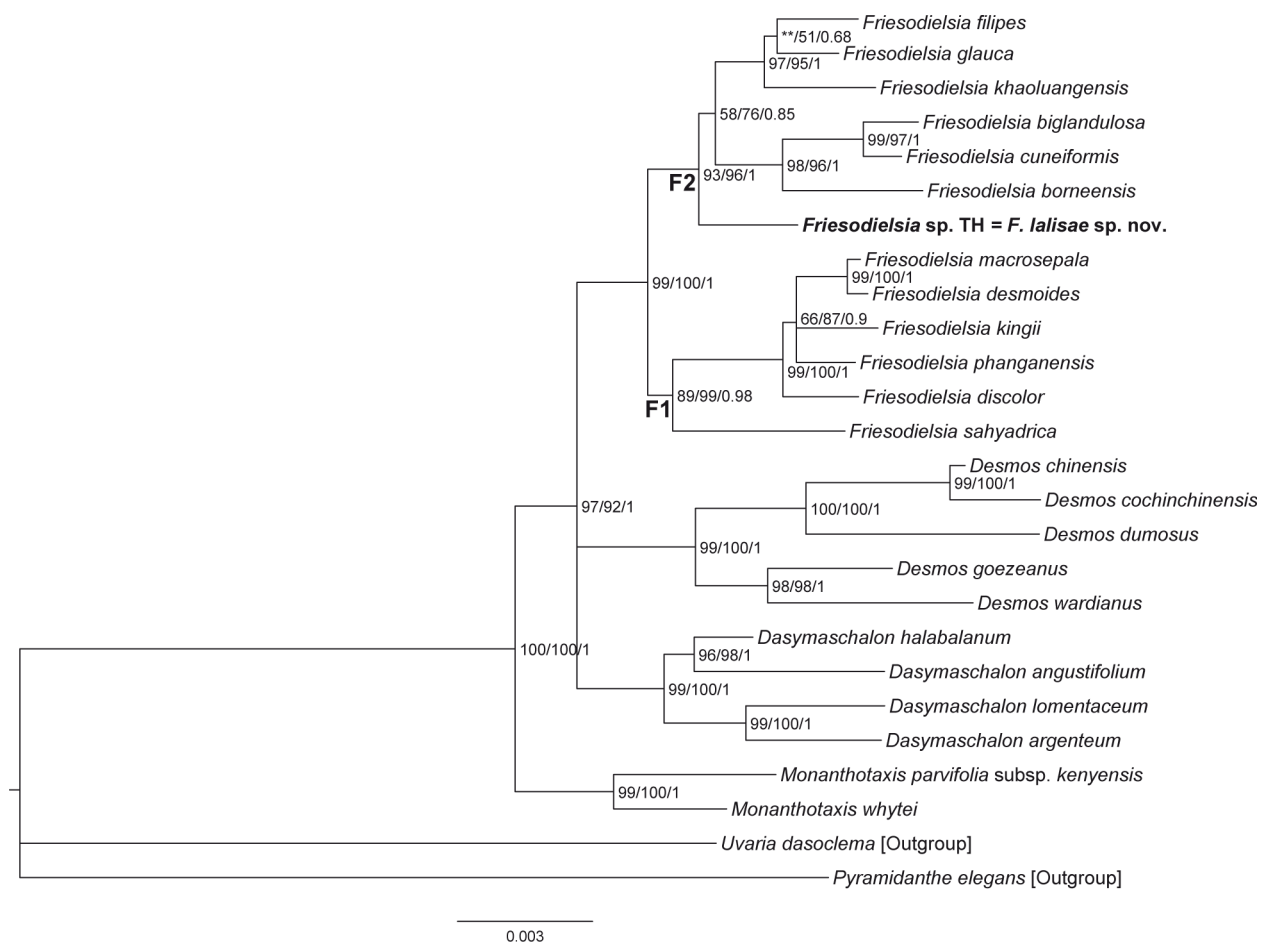


Fig. 1. Phylogram derived from Bayesian inference. Parsimony symmetric resampling values (SR), maximum likelihood bootstrap values (BS) and Bayesian posterior probabilities (PP) are indicated: SR/BS/PP. ** = SR < 50%. Scale bar unit = substitutions per site.

their type specimens at the herbaria K, L and SING (herbarium codes according to Thiers 2022+). Some floral organs of the unidentifiable gathering from Narathiwat, S Thailand (*Friesodielsia* sp. TH) were measured from spirit material (in square brackets in the description of the new species below). The indumentum terminology used followed Hewson (1988).

Results and Discussion

The parsimony analysis resulted in 18 most parsimonious trees with 689 steps. The consistency and retention indices (CI and RI) were 0.84 and 0.8, respectively. There was no strong incongruence (SR \geq 85%) among the analysis of each plastome region. The ingroup, comprising four genera: *Dasymaschalon*, *Desmos*, *Friesodielsia* and *Monanthataxis*, received maximal support (Fig. 1). Accessions of *Monanthataxis* formed a strongly supported (SR 99%, BS 100%, PP 1) clade sister to a larger strongly supported (SR 97%, BS 92%, PP 1) clade embracing the remaining accessions. In the latter clade, there was a polytomy consisting of three genera (*Dasymaschalon*, *Desmos* and *Friesodielsia*), each receiving strong support (SR 99%, BS 100%, PP

1). In *Friesodielsia*, two major clades were retrieved, each with strong support (Fig. 1): clades F1 (SR 89%, BS 99%, PP 0.98) and F2 (SR 93%, BS 96%, PP 1). In clade F1, *F. sahyadrica* N. V. Page & Survesw. was sister to a strongly supported (SR 99%, BS 100%, PP 1) clade composed of the remaining accessions in this clade. In clade F2, *Friesodielsia* sp. TH was recovered as a sister group of an unsupported to moderately supported (SR 58%, BS 76%, PP 0.85) clade consisting of the rest of this clade.

It is noteworthy that the two major clades of *Friesodielsia* correspond well with their morphology, i.e. the outer petals of members of clade F1, including the recently described *F. macrosepala* Leerat. & Aongyong and *F. phanganensis* Leerat., separate early during developmental stages and have a flat base, whereas those of members of clade F2, including *Friesodielsia* sp. TH and the recently described *F. khaoluangensis* Leerat. & Aongyong, separate at anthesis and have a concave basal portion, appearing as an excavation (Fig. 2B [left]; Guo & al. 2017a; Leeratiwong & al. 2021a). Upon morphological comparisons, *F. argentea* (J. Sinclair) Steenis native to Peninsular Malaysia and *F. glauca* (Hook. f. & Thomson) Steenis native to S Thailand and Peninsular Malaysia are most morphologically similar to *Friesodielsia* sp. TH; the three can

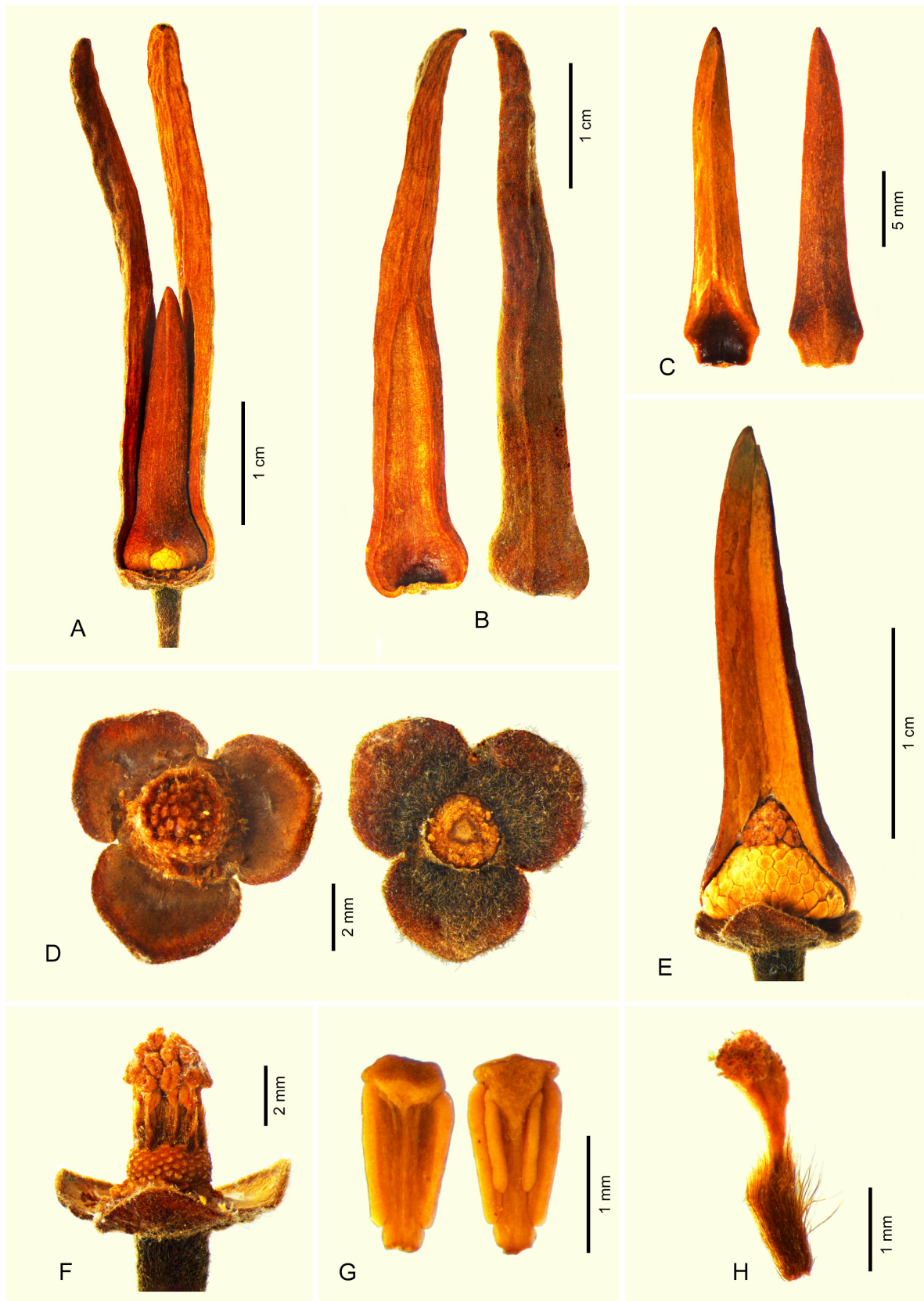


Fig. 2. Flower and floral organs of *Friesodielsia lalisae* – A: flower with one outer petal removed showing connivent inner petals; B: adaxial (left) and abaxial (right) sides of outer petal; C: adaxial (left) and abaxial (right) sides of inner petal; D: flower with petals, stamens and carpels removed showing adaxial (left) and abaxial (right) sides of sepals; E: flower with outer petals and one inner petal removed showing stamens surrounding carpels; F: flower with petals and stamens removed showing carpels and torus (side view); G: stamen, adaxial (left) and abaxial (right) sides; H: carpel. – All from Aongyong & Baka 57 (CMUB – spirit material).



FLORA OF THAILAND

Herbarium of Department of Biology, Faculty of Science, Chiang Mai University (CMUB)

Family: Annonaceae

Botanical name: *Friesodielsia lalisae* Damth., Baka & Chaowasku

Location: Chanae District, Narathiwat Province

Habitat: Occurring near a stream in secondary forests adjacent to rubber tree plantations

Elevation: c. 90 m

Notes: A woody climber; flowers fragrant, petals + yellow, inner petals remaining connivent at maturity

Date: May 2022

Collected by: Kithisak Aongyong & Abdulromea Baka

Number: 57

ibom

Fig. 3. Holotype of *Friesodielsia lalisae*: Aongyong & Baka 57 (CMUB).

Table 1. Morphological comparisons between *Friesodielsia lalisae*, *F. argentea* and *F. glauca*. Square brackets indicate measurements from spirit material.

Feature	<i>F. lalisae</i>	<i>F. argentea</i>	<i>F. glauca</i>
Young twigs	densely hairy	densely hairy	glabrous to sparsely hairy
Leaf base	obtuse to rounded	usually cuneate	obtuse to rounded
Pediceal length (mm, in flower)	9–10	c. 10	(14–)20–28
Sepal shape	transversely ovate	triangular	triangular
Proportion of inner petal to outer petal length	c. ½	c. ⅓	c. ⅓
Inner petal length (mm)	19 [23–25]	6–10	5–10

be distinguished by several features as shown in Table 1. It should be noted that, following Leeratiwong & al. (2021a) and Johnson & al. (2022), we consider the morphological features of *F. glauca* as based on material that only corresponds to the type of *Oxymitra glauca* Hook. f. & Thomson (the basionym of *F. glauca*), not the type of its several heterotypic synonyms, including *O. argentea* J. Sinclair (the basionym of *F. argentea*) (see Turner 2018). Based on morphological differences between *F. argentea* and *F. glauca* as indicated in Table 1, we believe the former should be regarded as distinct from *F. glauca*. Because *F. argentea*, *F. glauca* and two other species closely related to *F. glauca*: *F. filipes* (Hook. f. & Thomson) Steenis and *F. khaoluangensis* (Fig. 1) all have rather similar petal traits, i.e. proportion of inner petal to outer petal length (c. 1/3 or lower) and inner petal length (not exceeding 10 mm) (Table 1; Leeratiwong & al. 2021a; Johnson & al. 2022), it is hypothesized that *F. argentea* is also phylogenetically allied to *F. glauca*, which is distantly related to *Friesodielsia* sp. TH (Fig. 1). On the basis of these findings, *Friesodielsia* sp. TH deserves recognition as a new species, which is described below. It is worthwhile to note that Narathiwat constitutes one of the most insufficiently explored areas in Thailand as deduced from several species of *Annonaceae* described based on recently collected material (e.g. Jongsook & al. 2020; Bunchalee & al. 2021; Leeratiwong & al. 2021b; Wiya & al. 2021).

***Friesodielsia lalisae* Damth., Baka & Chaowasku, sp. nov.** – Fig. 2–4.

Holotype: Thailand, Narathiwat Province, Chanae District, May 2022 [in flower], *Aongyong & Baka* 57 (CMUB [barcode CMUB003997901]; isotypes: B, CMUB, QBG).

Diagnosis — *Friesodielsia lalisae* is most morphologically similar to *F. argentea* and *F. glauca*. The new species differs from these two species by having a different sepal shape, longer inner petals and a higher proportion of inner petal to outer petal length. Furthermore, *F. lalisae* differs from *F. argentea* by having an obtuse to rounded (vs usually cuneate) leaf base and from *F. glauca* by having denser indumentum on young twigs and shorter flowering pedicels.

Description (square brackets indicate measurements from spirit material) — Woody climbers; *young twigs* tomentose with erect and appressed hairs. *Petiole* 3–5 mm long, tomentose with erect and appressed hairs, slightly grooved above; *leaf blade* chartaceous, 7.7–16.2 × 2.8–5.8 cm, elliptic to elliptic-obovate, seldom obovate, puberulous-tomentose with erect and appressed hairs above, puberulous-tomentose with erect hairs below, base obtuse to rounded, apex ± cuspidate, acute to acute-acuminate, rarely obtuse or rounded; *midrib* slightly sunken above, tomentose with mostly erect hairs, raised below, puberulous-tomentose with erect and appressed hairs; *secondary veins* prominent below, 12–15 per side, angle with midrib 37°–46° (at middle part of leaf blade). *Flowers* solitary, terminal developing to internodal, fragrant *in vivo*; *pedicel* 9–10 mm long, curly-tomentose, bearing 1 bract near pedicel midpoint (but a bit lower), ovate-triangular. *Sepals* free, [3–3.1 × 4.5–4.6] mm, transversely ovate, without visible veins on both sides, outside curly-tomentose on basal half, more sparsely so on apical half, margin curly-tomentose, inside tomentose with appressed hairs only near margin, remaining area glabrous. *Petals* ± yellow *in vivo*; *outer petals* 34 [46–47] × 6 [8] mm, narrowly ovate-triangular, outside puberulous-tomentose with mostly appressed hairs, margin tomentose with appressed hairs, inside glabrous, each outer petal with an excavation on ± basal half, apex of outer petals ± acute; *inner petals* 19 [23–25] × [5–5.5] mm, narrowly ovate, c. ½ as long as outer petals, outside puberulous with appressed hairs only along bilateral midline, remaining area glabrous, margin and inside glabrous, apex acute. *Torus* depressed subglobose, villous intermixed with tomentose (both with erect hairs) on area surrounding each carpel. *Stamens* c. 132 per flower, [1.5–2.1] mm long, connective apex ± truncate or with a slanted prolongation, covering thecae. *Carpels* c. 22 per flower, [2.6–3.8] mm long; *stigmas* ± elongated and irregular-shaped; *ovaries* villous with mostly appressed hairs; *ovule* 1 per ovary, basal. *Fruit* unknown.

Phenology and ecology — Flowering material was collected in May. The species appears to grow near streams in secondary forests adjacent to rubber-tree plantations at an elevation of c. 90 m.



Fig. 4. *Friesodielsia lalisae* – A: flower bud; B: flower at anthesis; both from *Aongyong & Baka 57* (CMUB). – Photographs taken at type locality: Thailand, Narathiwat Province, Chanai District, May 2022, by A. Baka.

Distribution — Endemic to Narathiwat, S Thailand.

Preliminary conservation assessment — So far, *Friesodielsia lalisae* is only known to occur in secondary forests adjacent to rubber-tree plantations. Its habitat is highly threatened by agricultural activities. Only two individuals in a single location were observed, one of which has been cut recently. The AOO (area of occupancy) based on this single location is estimated to be less than 10 km². Although more exploratory data seem crucial, we believe the category Critically Endangered: CR B2ab(iii) based on IUCN Standards and Petitions Committee (2022) is appropriate for now and any conservation effort should be immediately initiated.

Etymology — The new species is named in honour of Lalisa Manobal, a famous Thai rapper, singer and dancer, whose motivation has greatly inspired the first author to overcome any obstacles during her Ph.D. study.

Author contributions

T.C. conceived and coordinated the study, obtained the research grant and performed molecular phylogenetic analyses; A.D. performed morphological investigations and comparisons, as well as DNA amplification; N.K. performed DNA amplification; S.K., C.S., C.W. and P.U. assisted with morphological investigations and comparisons, as well as with manuscript preparations; A.B. and K.A. provided crucial plant specimens; all authors drafted every version of the manuscript.

Acknowledgements

We thank the herbaria B, CMUB, K, L, QBG and SING for the material studied. Aroon Sinbumroong, Den Roopkom, Isma-ael Sama-ae and Suhbukree Samae provided useful material for molecular phylogenetic analyses. Two reviewers, Thomas L. P. Couvreur (Pontificia Universidad Católica del Ecuador) and Daniel C. Thomas (Singapore Botanic Gardens), provided constructive comments for the improvement of this article. The first author is grateful to the Science Achievement Scholarship of Thailand (SAST) for granting the scholarship to study a doctoral degree at Chiang Mai University. This research was supported by Fundamental Fund 2022, Chiang Mai University, as well as Office of the Permanent Secretary, Ministry of Higher Education, Science, Research and Innovation (OPS MHESI), Thailand Science Research and Innovation (TSRI) and Chiang Mai University (grant no. RGNS 63-082).

References

- Akaike H. 1974: A new look at the statistical model identification. – *IEEE Trans. Automat. Contr.* **19**: 716–723. <https://doi.org/10.1109/TAC.1974.1100705>
- Bangkomnate R., Damthongdee A., Baka A., Aongyong K. & Chaowasku T. 2021: *Pyramidanthe* and *Mitrella* (*Annonaceae*, *Uvarieae*) unified: molecular phylogenetic and morphological congruence, with new combinations in *Pyramidanthe*. – *Willdenowia* **51**: 383–394. <https://doi.org/10.3372/wi.51.51306>

- Bunchalee P., Leeratiwong C. & Johnson D. M. 2021: Two new species and a new record of the genus *Polyalthia* (*Annonaceae*) from Peninsular Thailand. – *Phytotaxa* **510**: 239–250. <https://doi.org/10.11646/phytotaxa.510.3.4>
- Chaowasku T., Aongyong K., Damthongdee A., Jongsook H. & Johnson D. M. 2020: Generic status of *Winitia* (*Annonaceae*, *Miliuseae*) reaffirmed by molecular phylogenetic analysis, including a new species and a new combination from Thailand. – *Eur. J. Taxon.* **659**: 1–23. <https://doi.org/10.5852/ejt.2020.659>
- Chaowasku T., Damthongdee A., Jongsook H., Ngo D. T., Le H. T., Tran D. M. & Suddee S. 2018a: Enlarging the monotypic *Monocarpieae* (*Annonaceae*, *Malmeloideae*): recognition of a second genus from Vietnam informed by morphology and molecular phylogenetics. – *Candollea* **73**: 261–275. <https://doi.org/10.15553/c2018v732a11>
- Chaowasku T., Damthongdee A., Jongsook H., Nuraliev M. S., Ngo D. T., Le H. T., Lithanatudom P., Osathanukul M., Deroin T., Xue B. & Wipasa J. 2018b: Genus *Huberantha* (*Annonaceae*) revisited: erection of *Polyalthiopsis*, a new genus for *H. floribunda*, with a new combination *H. luensis*. – *Ann. Bot. Fenn.* **55**: 121–136. <https://doi.org/10.5735/085.055.0114>
- Cheek M., Luke W. R. Q. & Gosline G. 2022: *Lukea* gen. nov. (*Monodoreae*–*Annonaceae*) with two new threatened species of shrub from the forests of the Udzungwas, Tanzania and Kaya Ribe, Kenya. – *Kew Bull.* **77**: 647–664. <https://doi.org/10.1007/s12225-022-10039-y>
- Chernomor O., von Haeseler A. & Minh B. Q. 2016: Terrace aware data structure for phylogenomic inference from supermatrices. – *Syst. Biol.* **65**: 997–1008. <https://doi.org/10.1093/sysbio/syw037>
- Couvreur T. L. P., Dagallier L. P. M. J., Crozier F., Ghogue J. P., Hoekstra P. H., Kamdem N. G., Johnson D. M., Murray N. A. & Sonké B. 2022: Flora of Cameroon – *Annonaceae* Vol 45. – *PhytoKeys* **207**: 1–532. <https://doi.org/10.3897/phytokeys.207.61432.figure45>
- Darriba D., Taboada G. L., Doallo R. & Posada D. 2012: jModelTest 2: more models, new heuristics and parallel computing. – *Nature Methods* **9**: 772–772. <https://doi.org/10.1038/nmeth.2109>
- Edgar R. C. 2004: MUSCLE: multiple sequence alignment with high accuracy and high throughput. – *Nucleic Acids Res.* **32**: 1792–1797. <https://doi.org/10.1093/nar/gkh340>
- Felsenstein J. 1985: Confidence limits on phylogenies: an approach using the bootstrap. – *Evolution* **39**: 783–791. <https://doi.org/10.1111/j.1558-5646.1985.tb00420.x>
- Goloboff P. A. & Catalano S. A. 2016: TNT version 1.5, including a full implementation of phylogenetic morphometrics. – *Cladistics* **32**: 221–238. <https://doi.org/10.1111/cla.12160>
- Goloboff P. A., Farris J. S., Källersjö M., Oxelman B., Ramírez M. J. & Szumik C. A. 2003: Improvements to resampling measures of group support. – *Cladistics* **19**: 324–332. <https://doi.org/10.1111/j.1096-0031.2003.tb00376.x>
- Guo X., Hoekstra P. H., Tang C. C., Thomas D. C., Wieringa J. J., Chatrou L. W. & Saunders R. M. K. 2017a: Cutting up the climbers: evidence for extensive polyphyly in *Friesodielsia* (*Annonaceae*) necessitates generic realignment across the tribe *Uvarieae*. – *Taxon* **66**: 3–19. <https://doi.org/10.12705/661.1>
- Guo X., Tang C. C., Thomas, D. C., Couvreur T. L. P. & Saunders R. M. K. 2017b: A mega-phylogeny of the *Annonaceae*: taxonomic placement of five enigmatic genera and support for a new tribe, *Phoenicantheae*. – *Sci. Rep.* **7**(7323). <https://doi.org/10.1038/s41598-017-07252-2>
- Hasegawa M., Kishino H. & Yano T. 1985: Dating of the human-ape splitting by a molecular clock of mitochondrial DNA. – *J. Molec. Evol.* **22**: 160–174. <https://doi.org/10.1007/BF02101694>
- Hewson H. J. 1988: Plant indumentum. A handbook of terminology. – Canberra: Australian Government Publishing Service. [Austral. Fl. Fauna Ser. 9].
- IUCN Standards and Petitions Committee 2022: Guidelines for using the IUCN Red List categories and criteria. Version 15.1. Prepared by the Standards and Petitions Committee. – Published at https://nc.iucnredlist.org/redlist/content/attachment_files/RedListGuidelines.pdf [accessed 24 Feb 2023].
- Jaikhamseub T., Le T. A., Damthongdee A., Huong T. T. T., Kuznetsov A. N., Kuznetsova S. P., Nuraliev M. S. & Chaowasku T. 2022: Two new species of *Meiogyne* (*Annonaceae*) from Vietnam, based on molecular phylogeny and morphology. – *Ann. Bot. Fenn.* **59**: 219–231. <https://doi.org/10.5735/085.059.0133>
- Johnson D. M., Murray N. A. & contributors 2022: *Annonaceae*. – In: Newman M. F., Barfod A. S., Esser H. J., Simpson D. A. & Parnell J. A. N. (ed.), *Flora of Thailand* **16** (part 1). – Bangkok: Prachachon Printing.
- Jongsook H., Samerpitak K., Damthongdee A. & Chaowasku T. 2020: The non-monophyly of *Dasymaschalon dasymaschalum* (*Annonaceae*) revealed by a plastid DNA phylogeny, with *D. halabalanum* sp. nov. from Thailand and *D. argenteum* comb. nov. – *Phytotaxa* **449**: 265–278. <https://doi.org/10.11646/phytotaxa.449.3.5>
- Kumar S., Stecher G. & Tamura K. 2016: MEGA7: molecular evolutionary genetics analysis version 7.0 for bigger datasets. – *Molec. Biol. Evol.* **33**: 1870–1874. <https://doi.org/10.1093/molbev/msw054>
- Leeratiwong C., Chalermglin P., Sattaphorn J., Aongyong K. & Johnson D. M. 2021a: New species and new records for the climber genus *Friesodielsia* (*Annonaceae*) in the flora of Thailand. – *Thai Forest Bull., Bot.* **49**: 212–230. <https://doi.org/10.20531/tfb.2021.49.2.09>
- Leeratiwong C., Chalermglin P. & Saunders R. M. K. 2021b: *Goniothalamus roseipetalus* and *G. sukhirii*

- nensis* (*Annonaceae*): two new species from Peninsular Thailand. – *PhytoKeys* **184**: 1–17. <https://doi.org/10.3897/phytokeys.184.73210>
- Leeratiwong C., Karapan S., Sattaphorn J. & Johnson D. M. 2023: Two new species of *Friesodielsia* (*Annonaceae*) from Peninsular Thailand. – *Phytotaxa* **589**: 73–82. <https://doi.org/10.11646/phytotaxa.589.1.7>
- Miller M. A., Pfeiffer W. & Schwartz T. 2010: Creating the CIPRES Science Gateway for inference of large phylogenetic trees. – Pp. 1–8 in: Gateway Computing Environments Workshop (GCE). – Piscataway: IEEE. <https://doi.org/10.1109/GCE.2010.5676129>
- Minh B. Q., Schmidt H. A., Chernomor O., Schrempf D., Woodhams M. D., von Haeseler A. & Lanfear R. 2020: IQ-TREE 2: new models and efficient methods for phylogenetic inference in the genomic era. – *Molec. Biol. Evol.* **37**: 1530–1534. <https://doi.org/10.1093/molbev/msaa015>
- Rambaut A., Drummond A. J., Xie D., Baele G. & Suchard M. A. 2018: Posterior summarization in Bayesian phylogenetics using Tracer 1.7. – *Syst. Biol.* **67**: 901–904. <https://doi.org/10.1093/sysbio/syy032>
- Ronquist F., Teslenko M., van der Mark P., Ayres D. L., Darling A., Höhna S., Larget B., Liu L., Suchard M. A. & Huelsenbeck J. P. 2012: MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across a large model space. – *Syst. Biol.* **61**: 539–542. <https://doi.org/10.1093/sysbio/sys029>
- Saunders R. M. K., Guo X. & Tang C. C. 2020: *Friesodielsia subaequalis* (*Annonaceae*): a new nomenclatural combination following conservation of the generic name against *Schefferomitra*. – *Phytotaxa* **464**: 183–184. <https://doi.org/10.11646/phytotaxa.464.2.5>
- Simmons M. P. 2004: Independence of alignment and tree search. – *Molec. Phylog. Evol.* **31**: 874–879. <https://doi.org/10.1016/j.ympev.2003.10.008>
- Simmons M. P. & Ochoterena H. 2000: Gaps as characters in sequence-based phylogenetic analyses. – *Syst. Biol.* **49**: 369–381. <https://doi.org/10.1093/sysbio/49.2.369>
- Sinclair J. 1955: A revision of the Malayan *Annonaceae*. – *Gard. Bull. Singapore* **14**: 149–516.
- Staden R., Beal K. F. & Bonfield J. K. 2000: The Staden Package, 1998. – Pp. 115–130 in: Misener S. & Krawetz S. A. (ed.), *Bioinformatics methods and protocols*. – Totowa: Humana Press. [*Meth. Molec. Biol.* **132**]. <https://doi.org/10.1385/1-59259-192-2:115>
- Tavaré S. 1986: Some probabilistic and statistical problems in the analysis of DNA sequences. – *Lectures Math. Life Sci.* **17**: 57–86.
- Thiers B. M. 2022+ [continuously updated]: Index herbariorum. – Published at <https://sweetgum.nybg.org/science/ih/> [accessed Dec 2022].
- Turner I. M. 2012: *Annonaceae* of Borneo: a review of the climbing species. – *Gard. Bull. Singapore* **64**: 371–479.
- Turner I. M. 2018: *Annonaceae* of the Asia-Pacific region: names, types and distributions. – *Gard. Bull. Singapore* **70**: 409–744. [https://doi.org/10.26492/gbs70\(2\).2018-11](https://doi.org/10.26492/gbs70(2).2018-11)
- Wiens J. J. 1998: Combining data sets with different phylogenetic histories. – *Syst. Biol.* **47**: 568–581. <https://doi.org/10.1080/106351598260581>
- Wiya C., Aongyong K., Damthongdee A., Baka A. & Chaowasku T. 2021: The genus *Phaeanthus* (*Annonaceae*, *Miliuseae*) in Thailand: *P. piyae* sp. nov. and resurrection of *P. lucidus*, with molecular phylogenetic analyses. – *Taiwania* **66**: 509–516. <https://doi.org/10.6165/tai.2021.66.509>
- Yang Z. & Rannala B. 1997: Bayesian phylogenetic inference using DNA sequences: a Markov Chain Monte Carlo method. – *Molec. Biol. Evol.* **14**: 717–724. <https://doi.org/10.1093/oxfordjournals.molbev.a025811>

Supplemental content online

See <https://doi.org/10.3372/wi.53.53103>

Original alignments and alignments for phylogenetic analyses in FASTA format.

Willdenowia

Open-access online edition bioone.org/journals/willdenowia



Online ISSN 1868-6397 · Print ISSN 0511-9618 · 2021 Journal Impact Factor 1.460

Published by the Botanic Garden and Botanical Museum Berlin, Freie Universität Berlin

© 2023 The Authors · This open-access article is distributed under the CC BY 4.0 licence

Appendix 1. Specimens for molecular phylogenetic analyses and their GenBank accession numbers. Unavailable sequences are denoted with —, whereas newly generated sequences are denoted with ***.

Taxon	Location	Collector and number (herbarium)	matK	ndhF	psbA-trnH	rbcL	trnL-trnF
<i>Dasymaschalon angustifolium</i> Jing Wang & R. M. K. Saunders	Thailand	Chaowasku 145 (CMUB)	MT264029	MT263997	MT264005	MT264013	MT264021
<i>Dasymaschalon argenteum</i> (Griff.) Jongsook & Chaowasku	Thailand	Aongyong 14 (CMUB)	MT264030	MT263998	MT264006	MT264014	MT264022
<i>Dasymaschalon halabalanum</i> Jongsook & Chaowasku	Thailand	Chaowasku 180 (CMUB)	MT264033	MT264001	MT264009	MT264017	MT264025
<i>Dasymaschalon lomentaceum</i> Finet & Gagnep.	Thailand	Chaowasku 182 (CMUB)	MT264034	MT264002	MT264010	MT264018	MT264026
<i>Desmos chinensis</i> Lour.	Hong Kong	Pang N2 (HKU)	JQ768567	JQ768603	JQ768646	JQ768687	JQ768727
<i>Desmos cochinchinensis</i> Lour.	China	Wang 0612 (HKU)	JQ768568	JQ768604	JQ768647	JQ768688	JQ768728
<i>Desmos dumosus</i> (Roxb.) Safi.	China	Wang 068 (HKU)	JQ768570	JQ768606	JQ768649	JQ768689	JQ768730
<i>Desmos goezeanus</i> (F. Muell.) Jessup	Australia	Ford & Cinelli 04780 (BRI)	JQ768572	JQ768607	JQ768651	JQ768691	JQ768732
<i>Desmos wardianus</i> (F. M. Bailey) Jessup	Australia	Sankowsky 2664 (BRI)	JQ768574	JQ768608	JQ768653	JQ768693	JQ768734
<i>Friesodielsia biglandulosa</i> (Blume) Steenis	Indonesia	Sitik 3809 (L)	KX786592	JQ768610	JQ768655	—	JQ768736
<i>Friesodielsia borneensis</i> (Miq.) Steenis	Indonesia	Kessler 2018 (A)	KX786593	KX786608	KX786616	—	KX786633
<i>Friesodielsia cuneiformis</i> (Blume) Steenis	cultivated (Bogor Bot. Gard.)	Ardi 54 (HKU)	JQ768576	JQ768611	—	JQ768695	JQ768737
<i>Friesodielsia desmoides</i> (Craib) Steenis	cultivated (Queen Sirikit Bot. Gard.)	Khunarak 1 (CMUB)	OQ505965**	OQ505974**	OQ505983**	OQ505992**	OQ506001**
<i>Friesodielsia discolor</i> (Craib) D. Das	Thailand	Aongyong & Roopkom 48 (CMUB)	OQ505964**	OQ505973**	OQ505982**	OQ505991**	OQ506000**
<i>Friesodielsia filipes</i> (Hook. f. & Thomson) Steenis	Thailand	Aongyong & Baka 46 (CMUB)	OQ505966**	OQ505975**	OQ505984**	OQ505993**	OQ506002**
<i>Friesodielsia glauca</i> (Hook. f. & Thomson) Steenis	Thailand	Aongyong & Samae 49 (CMUB)	OQ505968**	OQ505977**	OQ505986**	OQ505995**	OQ506004**
<i>Friesodielsia khaoluangensis</i> Leerat. & Aongyong	Thailand	Aongyong 20 (CMUB)	OQ505961**	OQ505970**	OQ505979**	OQ505988**	OQ505997**
<i>Friesodielsia kingii</i> (J. Sinclair) Steenis	Thailand	Aongyong & Samae 43 (CMUB)	OQ505963**	OQ505972**	OQ505981**	OQ505990**	OQ505999**

Taxon	Location	Collector and number (herbarium)	matK	ndhF	psbA-trnH	rbcL	trnL-trnF
<i>Friesodielsia talisae</i> Damth., Baka & Chaowasku	Thailand	Aongyong & Baka 57 (CMUB)	OQ505969**	OQ505978**	OQ505987**	OQ505996**	OQ506005**
<i>Friesodielsia macrosepala</i> Leerat. & Aongyong	Thailand	Aongyong 23 (CMUB)	OQ505962**	OQ505971**	OQ505980**	OQ505989**	OQ505998**
<i>Friesodielsia phanganensis</i> Leerat.	Thailand	Sinbumroong 09082021 (CMUB)	OQ505967**	OQ505976**	OQ505985**	OQ505994**	OQ506003**
<i>Friesodielsia sahyadrica</i> N. V. Page & Survesw.	India	Page 110949 (CAL)	KC933936	KC933942	KC933940	KC933934	KC933938
<i>Monanithotaxis parvifolia</i> subsp. <i>kenyensis</i> Verdc.	Kenya	Luke 7299 (EA)	KX761312	KX787032	KX786971	KX761343	KX787001
<i>Monanithotaxis whytei</i> (Stapf) Verdc.	cultivated (Utrecht Univ. Bot. Gard.)	Chatrou 475 (U)	EF179278	EF179304	EF179315	AY841635	AY841713
<i>Pyramidanthe elegans</i> (Hook. f. & Thomson) Bangk. & Chaowasku	Thailand	Aongyong & Baka 37 (CMUB)	OL546479	OL546490	OL546506	OL546522	OL546538
<i>Uvaria dasoclema</i> L. L. Zhou & al.	Thailand	Damthongdee AD 1 (BKF)	OL546465	OL546481	OL546492	OL546508	OL546524