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Research article

Checklist and molecular phylogenetics reveal three taxonomic novelties in *Habenaria* (Orchidaceae, Orchidoideae) from Chapada dos Veadeiros, Goiás, Brazil

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Abstract. A comprehensive checklist of *Habenaria* from Chapada dos Veadeiros, State of Goiás, was performed alongside morphologic and molecular phylogenetic studies, revealing three new taxa endemic to this region. A total of 61 taxa (59 species and two varieties) of *Habenaria* are recorded for Chapada dos Veadeiros, representing a two-fold increase compared to previous lists and comprising one of the greatest diversities of the genus in Brazil. Of this total, four taxa are locally endemic. *Habenaria cultellifolia*, until recently known only from the type collection, was rediscovered in the region after 127 years without records and represents this species' only known extant population. Three proposed new taxa of *Habenaria* (*H. minuticalcar* J.A.N. Bat. & Bianch. sp. nov., *H. proiteana* J.A.N. Bat., A.A. Vale & Bianch. sp. nov., and *H. lavrensis* var. *xanthodactyla* J.A.N. Bat. & Bianch. var. nov.) are corroborated by molecular phylogenetic analyses based on nuclear and plastid markers. They are described, illustrated, tentatively assessed as threatened, and compared to phylogenetically and morphologically related species. Since some areas of this mountain range have not yet been floristically sampled, additional taxonomic novelties and new records are still expected in the future.

Keywords. Biodiversity, Cerrado, grasslands, Habenariinae, taxonomy.

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Introduction

Habenaria Willd. (Orchidaceae, Habenariinae) is a genus of terrestrial orchids distributed throughout tropical and subtropical regions of the world (Cribb 2001). Currently, around 910 species of *Habenaria* are accepted (Govaerts *et al.* 2021), making the genus one of the largest in the family. Brazil, with 164 to 177 taxa (Flora e Funga do Brasil 2022; Govaerts *et al.* 2021) has the largest number of *Habenaria* species in the New World and possibly also worldwide and this genus is the largest of the family Orchidaceae in the country. Despite the recent progress made in the taxonomic knowledge of this genus in Brazil, there are still large gaps in almost all other aspects, such as the distribution, the regional and the local diversity, the ecology, the pollination and the reproductive biology of the species. Higher regions, with altitudes above 1,000 m a.s.l., are the ones with the greatest diversity (Brade 1951; Toscano de Brito 1995; Batista & Bianchetti 2003; Carvalho *et al.* 2013; Vieira & Barros 2017), but surveys are few and a systematic analysis of the distribution of the genus diversity in Brazil is lacking.

A particular area with great potential for terrestrial orchids is the Chapada dos Veadeiros region, State of Goiás, due to the large extension of savannic grasslands and the mild climate. The floristic knowledge of this region is still incomplete, with the first floristic inventory being the study of Munhoz & Proença (1998) for the municipality of Alto Paraíso de Goiás. This study listed 1,310 species and 120 families of vascular plants, including 47 species and 18 genera of Orchidaceae. In a more recent checklist for Chapada dos Veadeiros, Mendonça *et al.* (2007) listed 2,661 species distributed in 150 families. Orchidaceae, with 84 species, appears as the eighth family with the highest number of species. Both in the checklist by Munhoz & Proença (1998) and by Mendonça *et al.* (2007), *Habenaria* corresponds to the Orchidaceae genus with the largest number of species in this region, with 17 (35% of the family total) and 28 (33%) species, respectively. Besides checklists, some studies described new species of Orchidaceae for Chapada dos Veadeiros, including *Encyclia chapadensis* L.C.Menezes (Menezes 1992), *Cleistes pusilla* Pansarin (Pansarin 2004), *Sacoila cerradicola* Meneguzzo (Meneguzzo 2013), *Habenaria irwiniana* J.A.N.Bat. & Bianch., and *H. minuta* J.A.N.Bat. & Bianch. (Batista *et al.* 2017). Additionally, taxonomic treatments for *Bulbophyllum* Thouars (Santos & Silva 2019) and *Epidendrum* L. (Santos & Silva 2020) are also available for the Chapada dos Veadeiros National Park (PNCV).

Since 1991, two of the authors of the present study (JANB and LBB) have been exploring the diversity of Orchidaceae from the Chapada dos Veadeiros region within the scope of a larger Orchidaceae study from the Cerrado biome. Among the specimens found and collected throughout these years, several specimens of *Habenaria* did not correspond to any of the currently recognised Neotropical species and could not be assigned to any species rank. Subsequent morphological and molecular studies of these specimens revealed that some correspond to new species. In this study, we present the results of the *Habenaria* checklist from Chapada dos Veadeiros alongside morphological and molecular studies revealing three new taxa of *Habenaria* endemic to this region.

Material and methods

Study area

The Chapada dos Veadeiros covers an area of 21 475.60 km² comprising eight municipalities in the north of the State of Goiás: São João d'Aliança, Alto Paraíso de Goiás, Campos Belos, Cavalcante, Colinas do Sul, Monte Alegre de Goiás, Nova Roma, and Teresina de Goiás (Lima & Franco 2014). The region's climate is Aw type (according to the Köppen classification system), and the relief is mild to strongly undulate with altitudes ranging from 500–1 650 m a.s.l., including the highest point of the Brazilian Central Plateau. The region has a notable variety of phytophysionomies such as 'cerrado' sensu stricto (semideciduous xeromorphic medium tall forest), 'cerrado rupestre', 'campo rupestre' (rocky fields), 'campo úmido' (wet grasslands), 'campo limpo' (savannic grasslands), 'campo sujo' (grass-herb-sub-

shrub field), ‘veredas’ (damp grasslands with buriti palm trees *Mauritia flexuosa* L.f.), ‘mata seca’ (dry forests), and gallery forests (Munhoz & Proença 1998).

Checklist of *Habenaria* from Chapada dos Veadeiros

The checklist was carried out through 14 botanical expeditions from 1991 to 2018, complemented with the examination of specimens from the AMES, BHCB, CEN, ESA, HB, HBG, HEPH, HRBC, HTO, HUEFS, HUFU, IAN, IBGE, K, M, MBM, NY, P, RB, SP, SPF, UB, UEC and UFG herbaria (acronyms according to Thiers 2022). For herbarium specimens with a mixture of species in the same sheet, each species was counted as a separate record. Species identification was made by comparisons with protologues and type specimens. The main consulted taxonomic studies were those of Lindley (1830–1840), Reichenbach (1847, 1881), Barbosa Rodrigues (1877, 1882), Cogniaux (1893–1896, 1906), Kränzlin (1892, 1911), Renz (1992), and Batista *et al.* (2017). Type specimens and images of them were examined from the AMES, BHCB, BM, BR, C, CEN, G, HB, HBG, K, M, MBM, MPU, NY, P, R, RB, RENZ, S, SP, U, UPS, US and W herbaria. The taxonomic status and circumscription of the species follow a synopsis of the Neotropical species of *Habenaria* (Batista *et al.* 2011a, b).

Taxon sampling for the phylogenetic analyses

To investigate the phylogenetic relationships of the putative new taxa we performed an initial analysis using the same dataset as Batista *et al.* (2013), which comprises ca 51% of the Neotropical species of *Habenaria* and 1,389 bp of the complete nuclear internal transcribed spacer (ITS) region and part of the plastid *matK* gene. Based on this analysis, we selected some terminals for a second analysis using the dataset of Pedron *et al.* (2014), including some species addressed in later studies (Batista *et al.* 2014, 2016, 2017; Cruz-Lustre *et al.* 2022; Lau *et al.* 2021) and 37 new sequences from 11 species (12 terminals) that were generated for this study. For the outgroup, encompassing old world taxa, we used the same dataset as Batista *et al.* (2022a), which consists of a selected sampling based on the studies of Jin *et al.* (2017) and Ngugi *et al.* (2020), comprising representatives of the main clades of Asian and African *Habenaria* and related genera. The final data set consisted of the combined ITS, ETS, *matK-trnK* and *rps16-trnK* DNA sequences of 105 terminals of 103 taxa, including 85 terminals of 83 Neotropical *Habenaria* taxa containing most subclades of Neotropical *Habenaria* (Batista *et al.* 2013). We used *Stenoglottis longifolia* Hook.f., from subtribe Orchidinae, to root all trees based on the results of Ngugi *et al.* (2020). Voucher information, geographic origins, and GenBank accession numbers are provided in Supplementary file 1.

DNA extraction and molecular markers

The rDNA ITS, *matK-trnK*, *rps16-trnK* (Batista *et al.* 2013; Pedron *et al.* 2014) and rDNA ETS (Monteiro *et al.* 2010; Lau *et al.* 2021) markers were used. DNA was extracted from fresh or silica gel-dried samples, according to Doyle & Doyle (1987), and amplified the samples using the same primers and conditions described by Batista *et al.* (2013), Pedron *et al.* (2014) and Monteiro *et al.* (2010). PCR products were purified by precipitation with 20% PEG (2.5 M NaCl and 20% polyethene glycol 8000) and sequenced with the same primers used for amplification at Macrogen Inc. (Korea). All samples were bidirectionally sequenced, assembled with the Staden software package (Staden 1996), and aligned with MAFFT v.7 (Katoh & Standley 2013) as implemented in the EMBL-EBI bioinformatics web (Madeira *et al.* 2019).

Phylogenetic analyses

DNA sequence alignments were analysed using parsimony and Bayesian inference. To assess incongruences between the datasets we performed searches using both individual and combined matrices. In most cases, the concatenation was done with the same individuals sequenced for the different markers. We performed maximum parsimony (MP) phylogenetic analyses using PAUP* version 4 (Swofford 2002) with Fitch parsimony (equal weights, unordered characters; Fitch 1971) as the optimality criterion. Each search

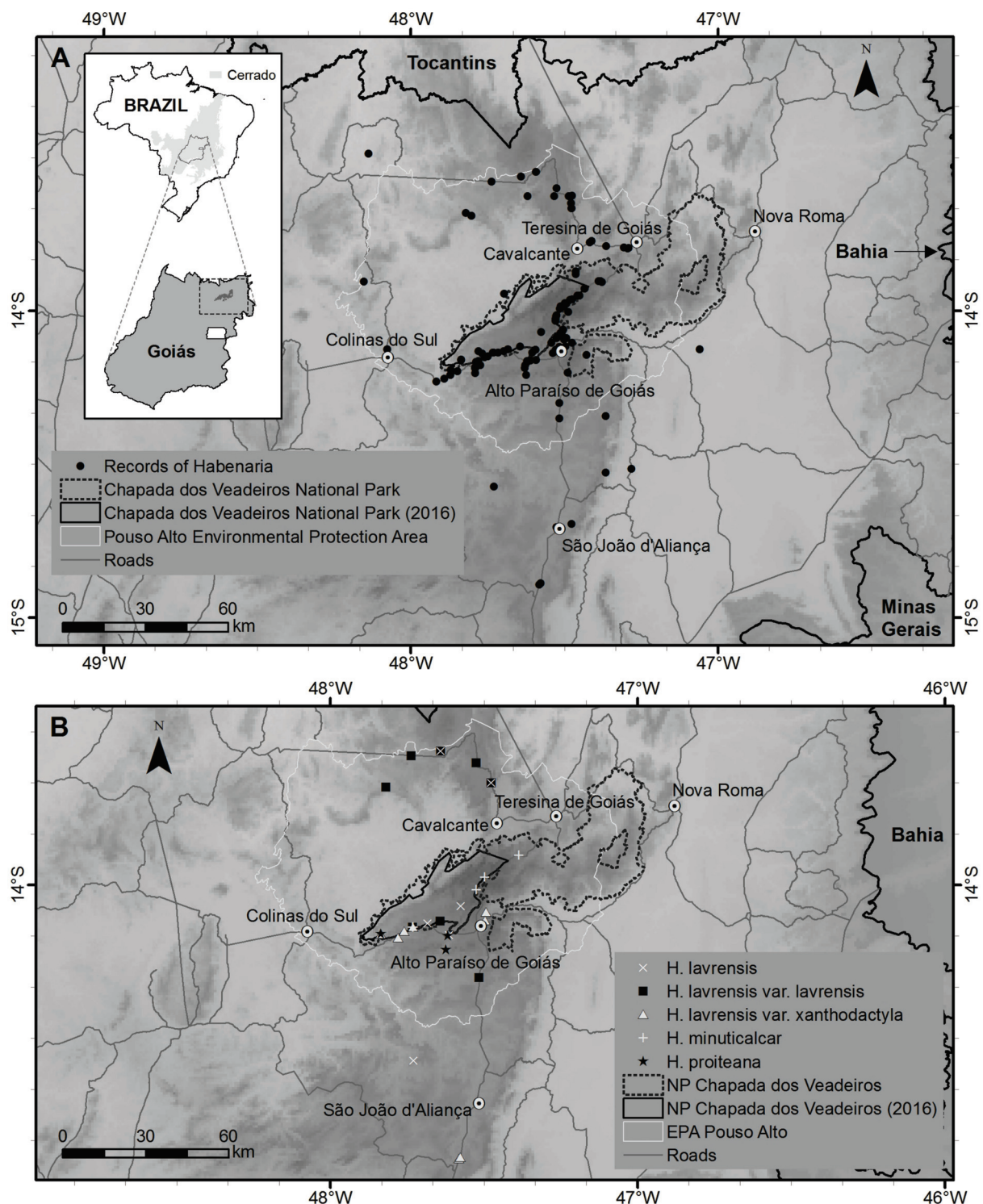


Fig. 1. Distribution map of *Habenaria* records from Chapada dos Veadeiros. **A.** Map with all records. **B.** Map with records of new taxa plus *H. lavrensis* Hoehne var. *lavrensis*. The area incorporated into the park in 2016 is indicated by a dashed line. *Habenaria lavrensis* denotes specimens with unknown intraspecific identification.

consisted of 2,000 replicates of random taxon additions, with branch swapping using the tree-bisection and reconnection (TBR) algorithm, saving ≤ 10 trees per replicate to avoid extensive swapping on suboptimal islands. We evaluated internal support by character bootstrapping (Felsenstein 1985) using 2,000 replicates, simple addition, and TBR branch swapping, saving ≤ 15 trees per replicate. For bootstrap support levels, we considered bootstrap percentages (BP) of 50–70% as weak, 71–85% as moderate, and $> 85\%$ as strong (Kress *et al.* 2002).

We performed Bayesian analyses using MrBayes 3.2.7a (Ronquist *et al.* 2012) as implemented in the Cyberinfrastructure for Phylogenetic Research (CIPRES) Portal 2.0 (Miller *et al.* 2010), treating each DNA region as a separate partition. We selected an evolutionary model for each DNA region in MrModeltest 2 (Nylander 2004) using the Hierarchical Likelihood Ratio Tests (hLRTs). The unlink command was used to estimate model parameters separately for each partition. Each analysis consisted of two independent runs with four chains for 5,000,000 generations, sampling one tree every 1,000 generations and a temperature parameter of 0.2. Convergence between the runs was evaluated using the average standard deviation of split frequencies (< 0.01) and the Potential Scale Reduction Factor – PRSF (= 1.0) and was achieved after 705,000 generations. After discarding the first 1,500 trees (30%) as the burn-in, the remaining trees were used to assess topology and posterior probabilities (PP) in a majority-rule consensus. Because PP in Bayesian analysis are not equivalent to BP but are generally much higher (Erixon *et al.* 2003), we used criteria similar to a standard statistical test, considering groups with $PP > 0.95$ as strongly supported, groups with PP ranging from 0.90–0.95 as moderately supported, and groups with $PP < 0.90$ as weakly supported.

Morphological analyses

Characters regarding habit, perianth, and gynostemium were examined from herbarium specimens, dissected flowers, and alcohol-fixed flowers. For the analyses of the gynostemium, we used a stereoscopic microscope and measured it using a digital calliper. Data concerning flowering times, habitats, and geographic distributions were based on herbarium labels and field observations. Morphological terminology is based on Stearn (1992) and Simpson (2006).

Conservation assessment

The distribution map was elaborated with ArcMap 10.5 (ESRI 2016). Occurrence records were obtained from herbaria and all records included had their identification revised according to the species protologues and the taxonomic circumscriptions of Batista *et al.* (2011 a, b). For specimens without coordinates, the latitude and longitude were inferred using Google Earth. We estimated the extent of occurrence (EOO) and area of occupancy (AOO) using a beta version of GeoCAT (Bachman *et al.* 2011) as implemented in the Royal Botanic Gardens, Kew Web site, with the default setting of 2 km cell width for AOO. Conservation status was inferred using the World Conservation Union Red List Categories and Criteria (IUCN 2012) and the guidelines for using the IUCN red list categories and criteria (IUCN 2016). To estimate the number of populations of each species, we considered each known record, if geographically separated, as a population.

Results

Survey of *Habenaria* in Chapada dos Veadeiros

A total of 61 taxa (59 species and two varieties) of *Habenaria* were recorded for Chapada dos Veadeiros (Table 1) in the municipalities of Alto Paraíso de Goiás, Cavalcante, Colinas do Sul, São João d'Aliança, and Teresina de Goiás (Fig. 1). The species with the highest number of records and the most common were: *H. obtusa* Lindl. (50 records), *H. magniscutata* Catling (32), *H. subviridis* Hoehne & Schltr. (29), *H. pabstii* J.A.N. Bat. & Bianch. (27), *H. hamata* Barb. Rodr. (25), *H. trifida* Kunth (25), *H. schwackei* Barb. Rodr. (24), *H. irwiniana* J.A.N. Bat. & Bianch. (20), *H. psammophila* J.A.N. Bat., Bianch. & B.M.

Table 1 (continued on the next page). *Habenaria* Willd. taxa recorded for Chapada dos Veadeiros, Goiás. Endemic taxa are indicated by an asterisk. Vouchers correspond to the first record of the taxon for the region. PNCV = Parque Nacional da Chapada dos Veadeiros (Chapada dos Veadeiros National Park).

| Taxa | N ° records | N ° records PNCV | Voucher (Herbaria) |
|---|-------------|------------------|---------------------------------------|
| <i>H. anisitsii</i> Kraenzl. | 7 | 0 | Batista et al. 719 (CEN, SP) |
| <i>H. armata</i> Rchb.f. | 5 | 0 | Anderson 6412 (UB) |
| <i>H. ayangannensis</i> Renz | 2 | 1 | Irwin et al. 32069 (UB) |
| <i>H. balansae</i> Cogn. | 3 | 3 | Batista & Pansarin 1144 (CEN, UEC) |
| <i>H. brevidens</i> Lindl. | 1 | 0 | Irwin et al. 24106 (NY) |
| <i>H. caldensis</i> Kraenzl. | 15 | 7 | Irwin et al. 12406 (HB) |
| <i>H. candolleana</i> Cogn. | 4 | 0 | Batista et al. 710 (BHCB, CEN, SP) |
| <i>H. ciliatisepala</i> J.A.N.Bat. & Bianch. | 5 | 2 | Gates & Estabrook 43 (HB, RB, UB) |
| <i>H. cryptophila</i> Barb.Rodr. | 1 | 0 | Batista et al. 1488 (CEN) |
| <i>H. cultellifolia</i> Barb.Rodr. | 5 | 0 | Batista et al. 1487 (CEN) |
| <i>H. curvilabria</i> Barb.Rodr. | 4 | 0 | Anderson 6542 (HB, SP, UB) |
| <i>H. depressifolia</i> Hoehne | 1 | 0 | Batista 837 (CEN, HUFU, SP) |
| <i>H. edwallii</i> Cogn. | 14 | 4 | Irwin et al. 12385 (UB UB0018542) |
| <i>H. eglariana</i> J.A.N.Bat. & Bianch. | 13 | 6 | Irwin et al. 12385 (UB UB0018542_1) |
| <i>H. gourlieana</i> Gill. ex Lindl. | 2 | 0 | Hatschbach et al. 54669 (MBM) |
| <i>H. guilleminii</i> Rchb.f. | 5 | 2 | Irwin et al. 12384a (HB, K) |
| <i>H. hamata</i> Barb. Rodr. | 25 | 7 | Irwin et al. 12725 (HB, K, UB) |
| <i>H. heptadactyla</i> Rchb.f. | 3 | 1 | Batista et al. 718 (CEN) |
| <i>H. heringeri</i> Pabst | 2 | 0 | Batista 839 (CEN, SP) |
| <i>H. hexaptera</i> Lindl. | 2 | 1 | Glaziou 22153a (K, P) |
| <i>H. imbricata</i> Lindl. | 8 | 2 | Batista et al. 705 (CEN) |
| <i>H. irwiniana</i> J.A.N.Bat. & Bianch. | 19 | 10 | Irwin et al. 12383 (NY, UB) |
| <i>H. jaguariahyaevae</i> Kraenzl. | 2 | 0 | Irwin et al. 12309 (NY, UB) |
| <i>H. johannensis</i> Barb.Rodr. | 1 | 0 | Batista 836 (CEN) |
| <i>H. juruenensis</i> Hoehne | 5 | 0 | Batista & Pansarin 1166 (CEN) |
| <i>H. lavrensis</i> Hoehne var. <i>lavrensis</i> | 15 | 4 | Glaziou 2215 (P) |
| <i>H. lavrensis</i> var. <i>xanthodactyla</i> J.A.N.Bat. & Bianch. var. nov. * | 10 | 4 | Gates & Estabrook 12 (HB, RB, SP, UB) |
| <i>H. leprieurii</i> Rchb.f. | 15 | 5 | Hatschbach et al. 36335 (MBM) |
| <i>H. leucosantha</i> Barb.Rodr. | 3 | 1 | De Haas et al. 356 (HB, M) |
| <i>H. ludibundiciliata</i> J.A.N.Bat. & Bianch. | 3 | 1 | Batista et al. 1539 (BHCB) |
| <i>H. macilenta</i> (Lindl.) Rchb.f. | 10 | 2 | Glaziou 22153 (K, P) |
| <i>H. magniscutata</i> Catling | 31 | 11 | Irwin et al. 12525 (HB, IAN, UB) |
| <i>H. minuta</i> J.A.N.Bat. & Bianch. * | 3 | 1 | Gates & Estabrook 216 (RB) |
| <i>H. minuticalcar</i> J.A.N.Bat. & Bianch. sp. nov. * | 5 | 4 | Batista 843 (CEN) |
| <i>H. cf. montis-wilhelminae</i> Renz | 2 | 0 | Pereira-Silva et al. 4600 (CEN) |
| <i>H. nasuta</i> Rchb.f. & Warm. | 1 | 0 | Anderson 7610 (UB) |
| <i>H. nuda</i> Lindl. var. <i>nuda</i> | 2 | 2 | Batista et al. 1217 (CEN) |

Table 1 (continued). *Habenaria* taxa recorded for Chapada dos Veadeiros, Goiás. Endemic taxa are indicated by an asterisk. Vouchers correspond to the first record of the taxon for the region. PNCV = Parque Nacional da Chapada dos Veadeiros (Chapada dos Veadeiros National Park).

| | | | |
|--|----|----|---|
| <i>H. nuda</i> var. <i>pygmaea</i> Hoehne | 2 | 2 | Walter 2142 (CEN) |
| <i>H. aff. nuda</i> Lindl. (1) | 1 | 0 | Pastore 1870 (HUEFS) |
| <i>H. aff. nuda</i> Lindl. (2) | 5 | 3 | Irwin <i>et al.</i> 12854 (UB) |
| <i>H. aff. nuda</i> Lindl. (3) | 3 | 0 | Batista <i>et al.</i> 1490 (BHCB) |
| <i>H. obtusa</i> Lindl. | 50 | 10 | Irwin <i>et al.</i> 24045 (HB, RB, UB) |
| <i>H. omissa</i> J.A.N.Bat. & Bianch. | 2 | 0 | Batista <i>et al.</i> 728 (CEN) |
| <i>H. orchioalcar</i> Hoehne | 14 | 5 | Irwin <i>et al.</i> 24996 (HB, K, UB) |
| <i>H. pabstii</i> J.A.N.Bat. & Bianch. | 28 | 13 | Irwin <i>et al.</i> 12382 (IAN, NY, UB) |
| <i>H. petalodes</i> Lindl. | 2 | 0 | Anderson 6579 (UB) |
| <i>H. pratensis</i> (Salzm. ex Lindl.) Rchb.f. | 1 | 1 | Martinelli 16578 (RB) |
| <i>H. proiteana</i> J.A.N.Bat., A.A.Vale & Bianch. sp. nov.* | 3 | 1 | Hatschbach <i>et al.</i> 58383 (MBM) |
| <i>H. psammophila</i> J.A.N.Bat., Bianch. & B.M.Carvalho | 21 | 7 | Irwin <i>et al.</i> 12360 (UB) |
| <i>H. cf. pseudoculicina</i> J.A.N.Bat. & B.M.Carvalho | 13 | 8 | Gates & Estabrook 156 (HB) |
| <i>H. aff. rodeiensis</i> Barb.Rodr. | 2 | 0 | Hatschbach <i>et al.</i> 58372 (HBG, MBM) |
| <i>H. schwackei</i> Barb.Rodr. | 24 | 12 | Wanderley <i>et al.</i> 1695 (SP) |
| <i>H. secundiflora</i> Barb.Rodr. | 4 | 1 | Pereira-Silva <i>et al.</i> 5887 (CEN) |
| <i>H. spanophytica</i> J.A.N.Bat. & Bianch. | 1 | 0 | Pastore & Suganuma 579 (CEN) |
| <i>H. spathulifera</i> Cogn. | 11 | 1 | Irwin <i>et al.</i> 24119 (HB, K, P) |
| <i>H. sprucei</i> Cogn. | 2 | 0 | Batista & Bianchetti 3086 (BHCB) |
| <i>H. subfiliformis</i> Cogn. | 7 | 4 | Irwin <i>et al.</i> 24201 (UB) |
| <i>H. subrepens</i> J.A.N.Bat., B.L.Lau & I.G.C.Machado-Costa | 5 | 2 | Hatschbach 36847 (MBM) |
| <i>H. subviridis</i> Hoehne & Schltr. | 29 | 13 | Irwin <i>et al.</i> 12384 (HB, UB) |
| <i>H. tamanduensis</i> Schltr. | 20 | 4 | Glaziou 22156 (K, P) |
| <i>H. trifida</i> Kunth | 24 | 7 | Irwin <i>et al.</i> 12957 (K, UB) |

Carvalho (20), and *H. tamanduensis* Schltr. (20) (Table 1). 27 taxa in this study are known from three or fewer records and can be considered rare or uncommon in the region. Among these stands out, *H. minuta*, endemic to Chapada dos Veadeiros and known so far from only three collections. Another important result was the rediscovery of *H. cultellifolia* (Fig. 5G–I). This species was described in 1877 from material collected in Uberaba, located 600 km to the south, in the State of Minas Gerais, and never collected again, and until its collection in Chapada dos Veadeiros was known only from the type collection.

Regarding distribution, four taxa (7% of the total) are known only from Chapada dos Veadeiros: *H. lavrensis* var. nov. *xanthodactyla* J.A.N. Bat. & Bianch., *H. minuta*, *H. minuticalcar* J.A.N. Bat. & Bianch. sp. nov., and *H. proiteana* J.A.N.Bat., A.A.Vale & Bianch. sp. nov. (Table 1). On a larger scale, four taxa are restricted to the Brazilian Central Plateau: *H. eglariana* J.A.N.Bat. & Bianch., *H. irwiniana*, *H. pabstii* J.A.N.Bat. & Bianch., and *H. spanophytica* J.A.N.Bat. & Bianch. (Flora e Funga

do Brasil 2022), while six taxa are shared with other montane regions of the States of Minas Gerais and Bahia: *H. ciliatisepala* J.A.N.Bat. & Bianch., *H. lavrensis* Hoehne, *H. magniscutata* Catling, *H. nuda* Lindl. var. *nuda*, *H. psammophila*, and *H. subrepens* J.A.N.Bat., B.L.Lau & I.G.C.Machado-Costa (Flora e Funga do Brasil 2022; Batista *et al.* 2022b). Regarding distribution of records and taxa within the PNCV, of the total records, 175 (33%) were made in the PNCV, and of the total taxa recorded for the region, 22 (36%) have no record for the PNCV, while another 17 (28%) taxa have only 1–2 records for this park (Table 1).

From a historical perspective, the first records for the region were made by A.F.M. Glaziou, who collected four species in ‘Chapadão dos Viadeiros’ between 1894 and 1895. The following records were only made in the 60s, when H.S. Irwin and colleagues intensively collected in the region between 1966 and 1971, resulting in 19 new records of *Habenaria* for Chapada dos Veadeiros (Table 1). In the 70s, the main collectors were W.R. Anderson, G. Hatschbach, and Gates & Estabrook, whose collections resulted in 13 new records for the region. From the 1990s onwards, the collections in the region were made mainly by researchers and postgraduate students from regional institutions such as CENARGEN, Universidade de Brasília and Jardim Botânico de Brasília, contributing with five new records. And from 1991 onwards, field collections within this study began and lasted until 2018, contributing to 17 new records of *Habenaria* for this region. According to our data, of a total of 63 collectors, the ones with the highest number of collections are J.A.N. Batista (165 collections), H.S. Irwin (46), A.A. Vale (32), J.F.B. Pastore (29), A.E.H. Salles (28), and G. Hatschbach (25). The number of collections of the genus in the region is high (542 herbarium collections; 9 collections/taxa), which reflects both the specific sampling effort for Orchidaceae carried out by the authors and the high frequency of the genus in the region.

Phylogenetic analyses

The concatenated ITS, ETS, *matK-trnK*, and *rps16-trnK* DNA matrix consisted of 4,372 aligned characters, of which 841 (19%) were parsimony-informative. Table 2 presents the general features of the datasets and parsimony statistics along with a summary of the models implemented for each partition. Missing data accounted for 8.4% of the combined matrix, most of which were from the complete *matK-trnK*, *rps16-trnK* and ETS sequences that were missing from Old World taxa. Missing data for the Neotropical sequences accounted for just 2.4% of the combined matrix. Comparable to previous molecular phylogenetic analyses with similar datasets (Pedron *et al.* 2014; Lau *et al.* 2021), no significant incongruences were detected between nuclear and plastid datasets, therefore only the results of the combined matrix are presented and discussed. The parsimony analyses retained a total of 17,551 most parsimonious trees, with a tree length of 3,013 steps, a consistency index (CI) of 0.63, and a retention index (RI) of 0.78. The strict consensus tree from the parsimony analysis was, for the most part, congruent with the Bayesian majority-rule consensus tree. Still, as the latter was more fully resolved and had stronger overall support, it was chosen for presentation and discussion (Fig. 2). For comparison, bootstrap percentages from the bootstrap analysis are shown in the Bayesian tree.

The relationships recovered were similar to those of earlier molecular phylogenetic studies (Batista *et al.* 2013, 2014, 2016, 2017, 2022a; Pedron *et al.* 2014; Lau *et al.* 2021), with the New World *Habenaria* species forming a well-supported monophyletic group (1.00 PP, 83% BP) and recovering several well-supported subclades within the New World clade that are numbered accordingly (subclades 1 to 21) (Fig. 2). The two new species addressed in this study (see Taxonomic treatment) were recovered within the Cerrado clades (1.00 PP, 96% BP), which comprises clades and species distributed mainly in the Cerrado Phytogeographic Domain of Central Brazil (Batista *et al.* 2013). *Habenaria mimuticalcar* sp. nov. (Figs. 33A–E, 4) was recovered with high support (1.00 PP, 83% BP) as sister to *H. pseudoculicina* J.A.N.Bat. & B.M.Carvalho, while *H. proiteana* sp. nov. (Figs. 5A–F, 6) was recovered with high support (1.00 PP, 100% BP) as sister to *H. psammophila*. *Habenaria lavrensis* var. *xanthodactyla* var. nov. was not sampled in this study, but *Habenaria lavrensis* var. *lavrensis* was also recovered in the Cerrado clades.

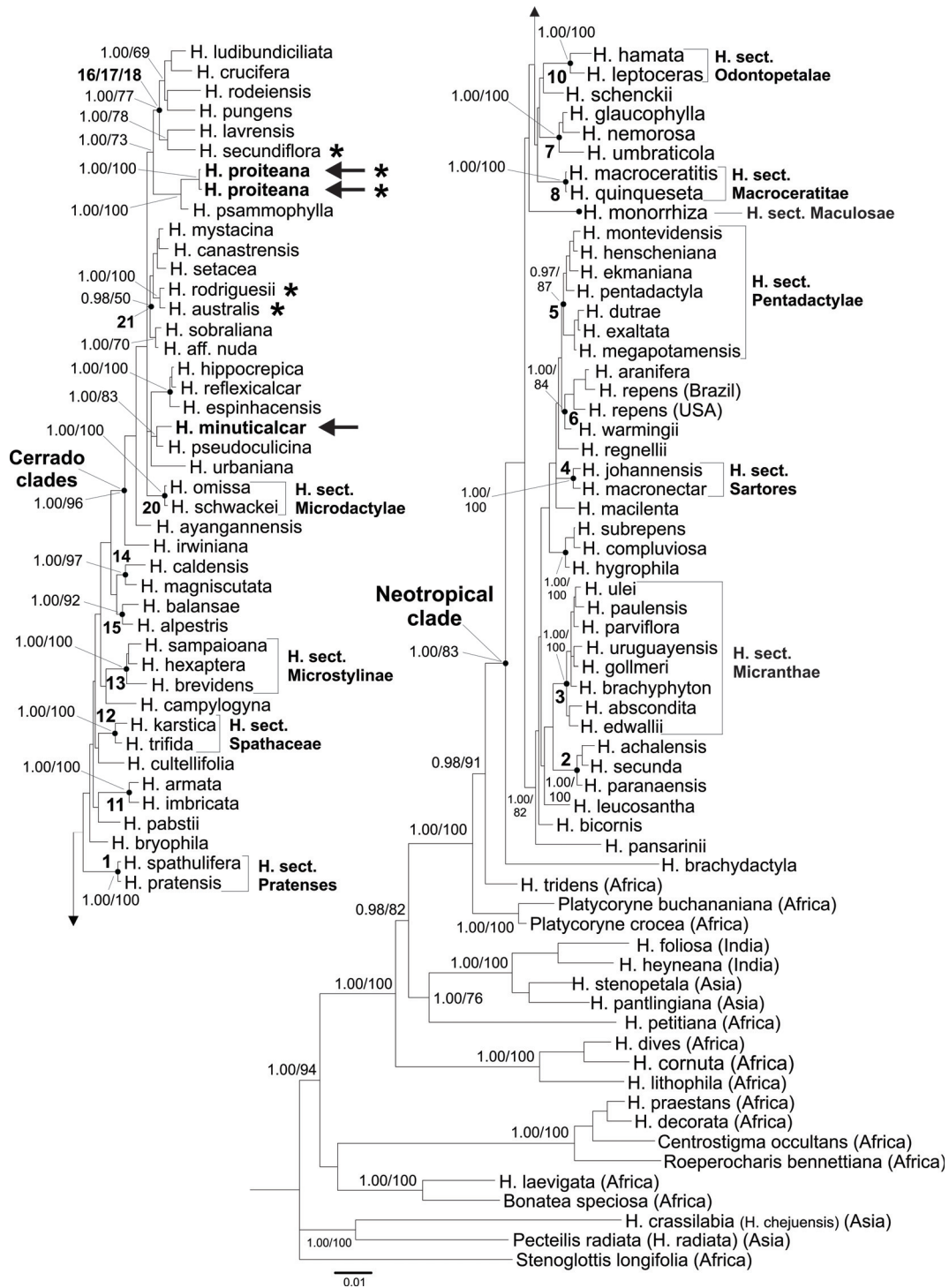


Fig. 2. Bayesian 50% majority-rule consensus tree of the combined ITS, ETS, *matK-trnK* and *rps16-trnK* datasets. Numbers next to the nodes represent the posterior probabilities (PP) from the Bayesian analyses and bootstrap percentages (BP) from the parsimony analyses. Only the values of the main clades are shown. Neotropical subclades are numbered according to Batista *et al.* (2013). The new species described here are highlighted in bold and indicated by an arrow. Species with the petal anterior segment inserted above the base of the posterior segment are highlighted by an asterisk. Subclades in which it was possible to associate with one of the sections of Kränzlin's sectional treatment are identified accordingly.

Taxonomy

Class Equisetopsida C.Agardh
Order Asparagales Link
Family Orchidaceae Juss.
Subfamily Orchidoideae Lindl.
Tribe Orchideae Verm.
Subtribe Habenariinae Benth.
Genus *Habenaria* Willd.

Habenaria minuticalcar J.A.N.Bat. & Bianch. sp. nov.

urn:lsid:ipni.org:names:77325126-1

Figs 3A–E, 4

Diagnosis

Similar to *H. guilleminii* Rchb. f. in the short spur, but distinguished by the leaves linear, narrow, 2–3 mm wide (vs lanceolate or lance-ovate, 5–23 mm wide), pedicellate ovary 7–12 mm long (vs 5–7 mm long), lateral sepals linear, 0.9–1.2 mm wide (vs obliquely ovate or obliquely lance-ovate, 2–3 mm wide), petal anterior segment filiform, 1.5–2.2 mm long. (vs reduced to a tooth-like projection, 0.2–0.7 mm long), and lip lateral segments shorter than the median segment (vs longer).

Etymology

From the Latin ‘minutus’ (very small) and ‘calcar’ (spur), referring to the very small spur, an unusual feature in the genus and one of the smallest among New World *Habenaria*.

Material examined

Holotype

BRAZIL – **Goiás** • Alto Paraíso de Goiás, 21 km N. de Alto Paraíso, campo a margem esquerda da GO-118, em direção a Teresina de Goiás; Parque Nacional da Chapada dos Veadeiros; 1 Jan. 1999; fl.; *J.A.N. Batista 843*; GenBank nrITS: HM777691; nrETS: OP589155; *matK* gene and *trnK* intron: OP589167; *rps16-trnK* intergenic spacer: OP589998; holotype: CEN[32585].

Paratypes

BRAZIL – **Goiás** • Teresina de Goiás, Chapada dos Veadeiros, GO-118, em direção a Alto Paraíso, ca de 37 km ao N e antes de Alto Paraíso; 9 Jan. 2001; fl.; *J.A.N. Batista & E.R. Pansarin 1149*; CEN[CEN00037136] • Alto Paraíso de Goiás, Chapada dos Veadeiros; 30 Dec. 2010; fl.; *J.A.N. Batista & L.B. Bianchetti 3099*; BHC[B151375] • Alto Paraíso de Goiás, Chapada dos Veadeiros; 30 Dec. 2010; fl.; *J.A.N. Batista & L.B. Bianchetti 3104*; BHC[B151377] • Alto Paraíso de Goiás, estrada para a Vila de São Jorge, ca 26,5 km da GO-118; 23 Jan. 2005; fl.; *J. Paula-Souza et al. 4528*; ESA[090123].

Description

Geophytic herb, caulescent, sympodial. Roots and tuberoid not examined. *Stem* 13–28 cm long including the inflorescence, 0.8–1.2 mm wide, slender, erect, straight to somewhat sinuose. *Leaves* 5–8, spirally-alternate, linear, narrow, the largest concentrated in the middle of the stem, 1.5–8 × 0.2–0.3 cm, reducing towards the apex of the stem, base sheathing, sheath closed. *Inflorescence* 2–6 cm long, spiral, few flowered, lax; floral bracts 5–17 × 3–4 mm, elliptical, caudate, apex acuminate, about the same size as the pedicellate ovary, decreasing in size towards the inflorescence apex, green. *Flowers* 2–7, resupinate, green; pedicellate ovary 7–12 mm long, mostly straight, more or less parallel to the rachis axis or ascending, apex curved; ovary 6–10 mm long, pedicel 1–2 mm long. *Sepals* green, margin smooth, apex

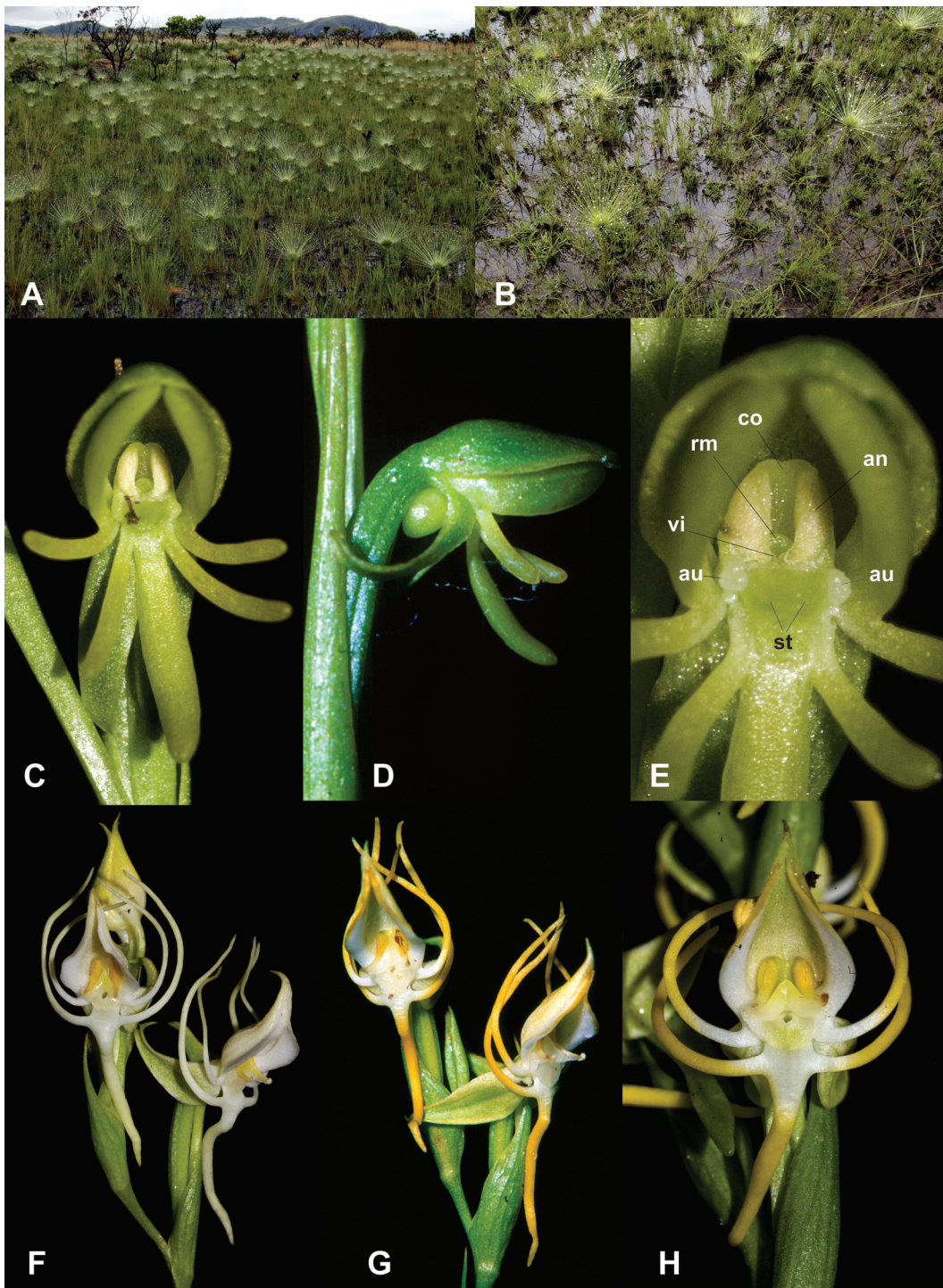


Fig. 3. *Habenaria minuticalcar* J.A.N. Bat. & Bianch. sp. nov. **A–B.** Habitat: seasonally humid grassland with *Paepalanthus* sp. during the rainy season in December 2010, in Chapada dos Veadeiros. **C.** Flower, front view. **D.** Flower, side view. **E.** Gynostemium, front view. *Habenaria lavrensis* var. *lavrensis*. **F.** Inflorescence. *Habenaria lavrensis* var. *xanthodactyla* var. nov. **G.** Inflorescence. **H.** Flower, front view. C, E from *Batista & Bianchetti 3099* (BHCB); D from *Batista 843* (CEN); F from *Batista 2940* (BHCB); G from *Batista & Proite 1494* (CEN); H from *Batista & Bianchetti 3095* (BHCB). Abbreviations: an = anther; au = auricles; co = connective; rm = rostellum mid lobe; st = stigmatophores; vi = viscidia. All photographs by J.A.N. Batista.

discreetly mucronate or obtuse; dorsal sepal 3–5 × 3–3.5 mm, concave, ovate or elliptical when flattened; lateral sepals 3–4.5 × 0.9–1.2 mm, obliquely linear to obliquely linear-lanceolate, deflexed or reflexed, apex obtuse. Corolla base whitish, towards the apex light green or greenish yellow. *Petals* bipartite; posterior segment 2.5–4.2 × 1–1.2 mm, oblong, slightly falcate, apex obtuse, connivent with the dorsal sepal; anterior segment 1.5–2.2 mm long, shorter than the posterior lobe, linear-filiform, inserted at the base of the posterior segment. *Lip* tripartite; undivided basal part 0.5 mm long; lateral segments 1.9–3 mm long, shorter than the median segment, linear-filiform, bent forward; median segment 3–4.5 × 0.7–0.9 mm, linear, bent forward; spur 1–1.6 × 1–1.3 mm, shorter than the pedicellate ovary, free from the bract, ovate, green. *Gynostemium* 1.5 mm high, reclinate; connective green, apex emarginate; lateral appendages (auricles) 0.9–1.2 × 0.4 mm, fleshy, verrucose, apex obtuse, whitish. *Anther* 0.7–1 mm high, bilocular, loculi parallel, translucent; canals very short, slightly convergent; pollinaria 2, 0.7–0.8 mm long each, joined by the viscidia; caudicles 0.2 mm long, filiform; pollinia 0.5–0.6 mm long, yellow; viscidia 0.15 × 0.15 mm, enclosed between the anthers. *Stigmatophores* (stigma lobes) 2, ca 1.6 mm long, closely parallel, green, receptive surface 0.9 mm long, convex, turned frontwards, 0.4 mm wide each, apex obtuse. *Rostellum* ca 1 mm long, green; mid lobe ca 0.6 mm long, 0.4–0.5 mm high, triangular, erect, held between the anther loculi, fleshy, apex obtuse; side-lobes 0.3–0.4 mm long, convergent towards the apex.

Distribution, Habitat and Phenology

Habenaria minuticalcar sp. nov. is restricted to Chapada dos Veadeiros, State of Goiás, central Brazil. The new species grows in seasonally humid grassland (Fig. 3A), associated with ‘murundus’ or ‘campos rupestres’ (rocky fields), with dark, shallow, rocky, sandy-clay soils, at 1,100 to 1,600 m a.s.l. The soil can be humid during some periods in the rainy season (Fig. 3B) but usually dries out completely during the dry season. The plants are difficult to find in their habitat as they are small and slender, grow among grasses and can only be found when they are in flower. Flowering occurs during the peak of the rainy season, from December to January. Like other *Habenaria* and terrestrial grassland orchids, the flowering of the new species is enhanced by seasonal brush fires. All our collections of the species were made in areas burned from the previous dry season.

Conservation Assessment

Habenaria minuticalcar sp. nov. is currently known from only four populations, all from Chapada dos Veadeiros, and shows an EOO estimated at 194 km² and a small AOO of 16 km². Based on the IUCN Red List Categories and Criteria and its guidelines (IUCN 2012, 2016), the species can be tentatively classified as Endangered EN B1ab(iii)+2ab(iii); C2a(i).

Taxonomic notes

Habenaria minuticalcar sp. nov. resembles *H. guillemirii* in the size and morphology of the spur. However, it is distinguished by the linear, narrow, 2–3 mm wide leaves (Fig. 4D–F) (vs lanceolate or elliptical, 5–23 mm wide in *H. guillemirii*), narrow, 0.9–1.2 mm wide lateral sepals (vs 2–3 mm wide), petal anterior segment 1.5–2.2 mm long, reaching half the size of the petal posterior segment (vs reduced to a tooth-like projection) and lip lateral segments at most half the length of the median segment (Fig. 4G–I) (vs lateral segments longer than the median segment). *Habenaria minuticalcar* sp. nov. is also similar to *H. pansarinii* var. *minuscula* J.A.N.Bat. & Bianch. in general morphology and size of the spur. However, it differs from this taxon by the floral bract measuring 5–17 mm long, approximately the same length as the pedicellate ovary (Fig. 4J–K) (vs 2.2–3.5 mm long, shorter than the pedicellate ovary), the slightly larger flowers (e.g., dorsal sepal 3–5 × 3–3.5 mm vs 2.4–3 × 1.8–2.4 mm in *H. pansarinii* var. *minuscula*), the petal posterior segment 2.5–4.2 mm long, oblong with an obtuse apex (vs 2.1–2.7 mm long, lanceolate, apex acute), and the geographical distribution restricted to Chapada dos Veadeiros (vs restricted to the southern and central regions of the Espinhaço Range in the State of Minas Gerais).

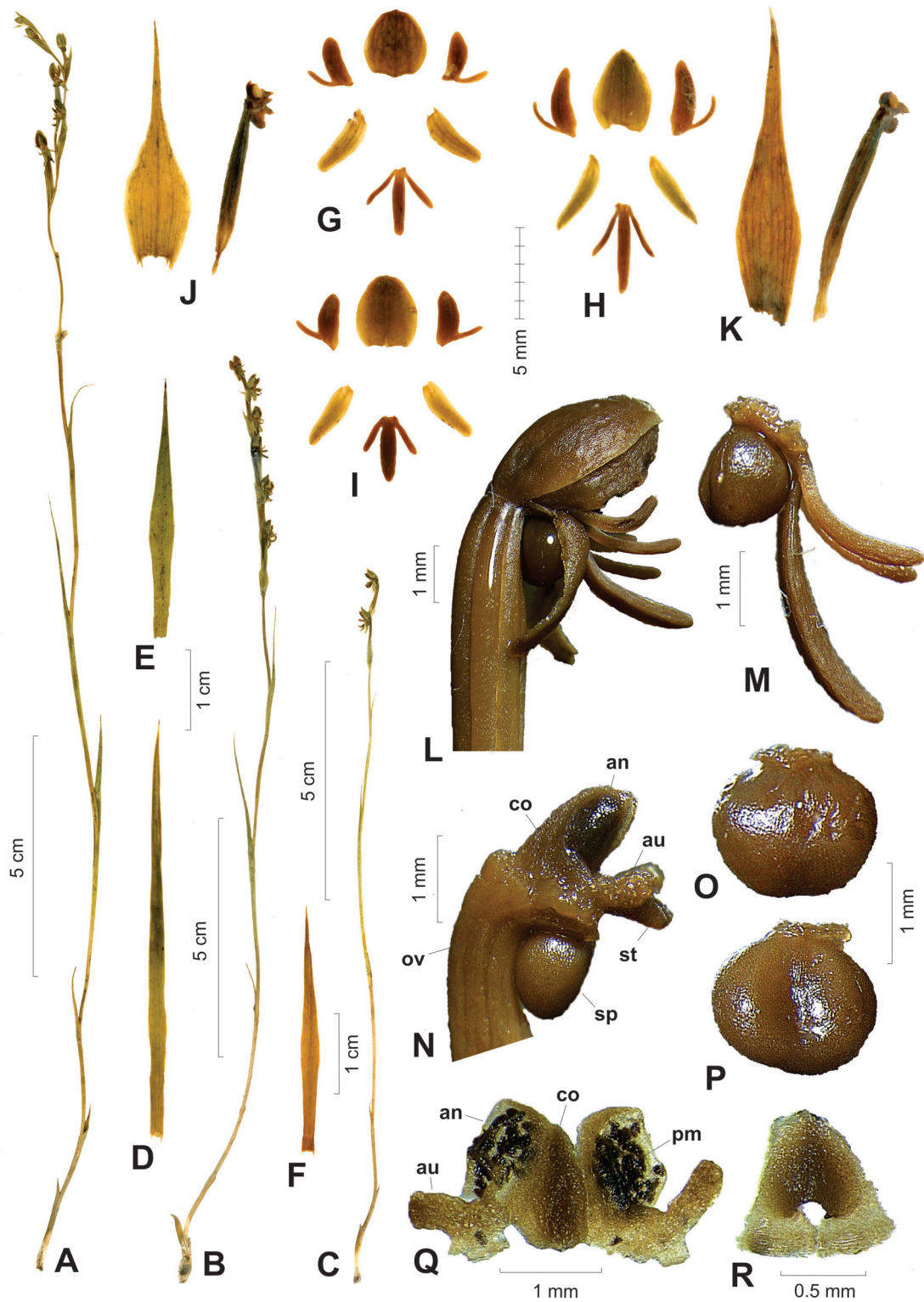


Fig. 4. *Habenaria minuticalcar* J.A.N. Bat. & Bianch. sp. nov. A–C. Habit. D–F. Leaves. G–I. Dissected perianth. J–K. Bract, pedicellate ovary, gynostemium and spur, lateral view. L. Flower, side view. M. Lip and spur, side view. N. Distal part of the ovary, gynostemium and spur, side view. O–P. Spur. Q. Flattened connective, anther and lateral appendages (auricles). R. Rostellum, upper view. A–D and F–R from *Batista* 843 (CEN); E from *Batista & Bianchetti* 3099 (BHCB). Abbreviations: an = anther; au = auricles; co = connective; ov = ovary; sp = spur; st = stigmatophores; pm = pollen massulae.

Key to the Brazilian species of *Habenaria* with a reduced spur

1. Pedicel shorter than the ovary; dorsal sepal ≤ 5 mm long 2
– Pedicel the same length or longer than the ovary; dorsal sepal ≥ 8 mm long 4
2. Leaves lanceolate or elliptical, 5–23 mm wide; lip lateral segments longer than the median segment; viscidia separated..... *H. guillemini* Rchb.f.
– Leaves linear, narrow, 1–4 mm wide; lip lateral segments shorter than the median segment; viscidia united 3
3. Floral bracts 5–17 \times 3–4 mm, elliptical, about the same size as the pedicellate ovary, decreasing in size towards the inflorescence apex; pedicellate ovary 7–12 mm long; dorsal sepal 3–5 \times 3–3.5 mm; petal posterior segment 2.5–4.2 mm long, oblong, apex obtuse; restricted to the Chapada dos Veadeiros, State of Goiás, Brazil *H. minuticalcar* sp. nov.
– Floral bracts 2.2–3.5 \times 1.2–1.8 mm, ovate, shorter than the pedicellate ovary; pedicellate ovary 6.7–8.1 mm long; dorsal sepal 2.4–3 \times 1.8–2.4 mm; petal posterior segment 2.1–2.7 mm long, lanceolate, apex acute; restricted to the southern and central regions of the Espinhaço Range in the State of Minas Gerais, Brazil *H. pansarinii* var. *minuscula* J.A.N.Bat. & Bianch
4. Dorsal sepal 11–20 mm long; petal anterior segment 8–11(–16) mm long, about the same size as the posterior segment; spur ovoid, scrotiform, ventrally compressed, 3–3.5 mm long.....
..... *H. orchioalcar* Hoehne
– Dorsal sepal 8–12 mm long; petal anterior segment 1–4 mm long, less than half the length of the posterior segment; spur elliptical, 5–6 mm long *H. heringeri* Pabst

Habenaria lavrensis Hoehne var. *xanthodactyla* J.A.N.Bat. & Bianch. var. nov.

urn:lsid:ipni.org:names:77326615-1

Figs 3G–H

Diagnosis

Similar to *H. lavrensis* var. *lavrensis* in general morphology, habitat and phenology, but distinguished by the yellow segments of the petals and lip (vs white) and by the geographical distribution restricted to the Chapada dos Veadeiros region (vs broadly distributed in the States of Goiás, Minas Gerais, and Distrito Federal).

Etymology

From the Greek ‘xanthos’ (yellow) and ‘dactylos’ (finger), referring to the yellow segments of the petals and lip.

Material examined

Type

BRAZIL – **Goiás** • São João da Aliança. GO-118, cerca de 22 km S de São João da Aliança; 9 Feb. 2004; fl.; J.A.N. Batista & K. Proite 1494; holotype: BHC[BHCB199457]; isotypes: CEN, RB, SP.

Paratypes

BRAZIL – **Goiás** • Alto Paraíso de Goiás, Chapada dos Veadeiros; 27 Jan. 1993; fl.; J.A.N. Batista & L.B. Bianchetti 386; CEN[CEN00018264]•Alto Paraíso de Goiás, Chapada dos Veadeiros, na estrada de terra em direção a cachoeira da Água Fria; 8 Jan. 2003; fl.; J.A.N. Batista et al. 1365; CEN[CEN00046058]•Alto Paraíso, Chapada dos Veadeiros, em direção ao Vale da Lua, cerca de 1,7 km do entroncamento com a GO-239; 30 Dec. 2010; fl.; J.A.N. Batista & L.B. Bianchetti 3095;

BHCB[151368]•Alto Paraíso de Goiás, Chapada dos Veadeiros; 30 Dec. 2010; fl.; *J.A.N. Batista & L.B. Bianchetti 3100*; BHCB[151373]•Chapada dos Veadeiros, 5 km E of Alto Paraíso; 1500 m; 24 Jan. 1979; fl.; *Gates & Estabrook 12*; HB[70548], NY[NY01031853], RB[RB00257176], SP[195098], UB[UB0018813]•Alto Paraíso de Goiás, Parque Nacional da Chapada dos Veadeiros, trilha para as corredeiras; 23 Jan. 2009; fl.; *P.L. Ribeiro et al. 396*; HUEFS[HUEFS0162888]•Alto Paraíso de Goiás, estrada para a Vila de São Jorge; 23 Jan. 2005; *J. Paula-Souza 4555*; ESA[090150]•Alto Paraíso de Goiás, Parque Nacional da Chapada dos Veadeiros, trilha para as Cachoeiras; 24 Jan. 2005; *J. Paula-Souza 4581*; ESA[090176]•São João da Aliança, 21 km ao Sul de São João da Aliança; 23 Feb. 1997; fl.; *J.A.N. Batista et al. 729*; CEN[CEN00026177]•ca 20,5 km S de São João da Aliança; 5 Feb. 2010; *J.A.N. Batista et al. 2965*; BHCB[137850].

Distribution, Habitat and Phenology

Habenaria lavrensis var. *xanthodactyla* var. nov. is so far only known from Chapada dos Veadeiros, Goiás, in the municipalities of Alto Paraíso de Goiás and São João da Aliança. In the northern part of Chapada dos Veadeiros, in the Distrito Federal and the state of Minas Gerais, where we have collected or examined specimens of this species, only *H. lavrensis* var. *lavrensis* is known. *Habenaria lavrensis* var. *xanthodactyla* var. nov. grows in dry, hillside or *campo rupestre*, in shallow, dry, gravelly soil. Flowering occurs during the peak of the rainy season, from January to February. In common with other grassland species of the genus, bushfires favoured flowering during the dry period. All populations we observed in flowering grew in areas that burned in the previous dry period. The habitat and the flowering period are about the same between the two varieties.

Conservation Assessment

Habenaria lavrensis var. *xanthodactyla* var. nov. is currently known from only seven populations from six localities and shows an EOO estimated at 1,647 km² and an AOO of 32 km². Based on the IUCN Red List Categories and Criteria and its guidelines (IUCN 2012, 2016), the species can be tentatively classified as Vulnerable VU B1ab(iii)+2ab(iii).

Taxonomic notes

Habenaria lavrensis var. *xanthodactyla* var. nov. is identical in vegetative and flower morphology to *H. lavrensis* var. *lavrensis* but distinct by the yellow lateral segments of petals and lip (Fig. 3G–H). In contrast, the entire flower is completely white or, at most segments, slightly creamy-white in *H. lavrensis* var. *lavrensis* (Fig. 3F). Both *H. lavrensis* var. *lavrensis* and *H. lavrensis* var. *xanthodactyla* var. nov. grow at the Chapada dos Veadeiros as homogeneous populations, spatially separated, and we have never found mixed populations or intermediate specimens. In defining the taxonomic rank of this taxon, we followed Stace (1989), who defined a variety as "a population of one or several biotypes, forming more or less distinct local races of a species."

Habenaria proiteana J.A.N. Bat., A.A.Vale & Bianch. sp. nov.

urn:lsid:ipni.org:names:77325125-1

Figs 5A–F, 6

Diagnosis

Similar to *H. rodriguesii* Cogn. in general morphology, but distinguished by the dorsal sepal 6–9 mm long (vs 10–14 mm long), petal anterior segment 11–16 mm long (vs 20–33 mm long), spur 13–17 mm long, genuflexus (vs 16–21 mm long, straight or only slightly curved), and anther canals curved upwards (vs straight, perpendicular to the anther loculi). Also similar to *H. sobraliana* J.A.N.Bat., Vale & Menini in general morphology, but distinct by the petal anterior segment 11–16 mm long, inserted a few millimetres above the posterior segment's base (vs 5–8 mm long, inserted at the base of the posterior segment).

Etymology

We named this species after Karina Proite for her support, companionship, help in the field and contribution to numerous collections, and for finding the population used to characterise, describe, and typify the newly proposed species.

Material examined

Type

BRAZIL – **Goiás** • Alto Paraíso de Goiás, Parque Nacional da Chapada dos Veadeiros, próximo as corredeiras do Rio Preto; 3 Jan. 2008; fl.; *J.A.N. Batista, K. Proite & P. Proite 2376*; GenBank nrITS: OP584473 and OP584474; nrETS: OP589157 and OP589158; *matK* gene and *trnK* intron: OP589169 and OP589170; *rps16-trnK* intergenic spacer: OP590000; holotype: BHC[B120057]; isotypes: CEN, RB, SP.

Paratypes

BRAZIL – **Goiás** • Alto Paraíso de Goiás, rodovia para Colinas, Rio das Cobras; 2 Dec. 1992; fl.; *G. Hatschbach et al. 58383*; MBM[158574]•Alto Paraíso de Goiás, Chapada dos Veadeiros, GO-239, entre Alto Paraíso e São Jorge, cerca de 3,1 km após a entrada para a fazenda São Bento; 30 Dec. 2010; fl.; *J.A.N. Batista & L.B. Bianchetii 3101*; BHC[B151374]•Alto Paraíso de Goiás, Parque Nacional Chapada dos Veadeiros, Corredeiras; 19 Dec. 2012; *A.A. Vale & J.A. Vale 180*; BHC[B174606].

Description

Geophytic herb, caulescent, sympodial. *Roots* 1–1.6 mm wide, few, sparsely hairy. *Tuberoïd* not examined. *Stem* 31–64 cm long including the inflorescence, 1.7–2.5 mm wide, erect, straight. *Leaves* 4–7, linear, narrow, the largest in the middle of the stem, 2.5–12 × 0.2–0.7 cm, reducing towards the apex of the stem, base sheathing, sheath closed, blade for the most part appressed to the stem, apex acuminate. *Inflorescence* 9–23 cm long, spiral, few flowered, lax; floral bracts 15–32 × 3.7–8 mm, ovate, lance-ovate, broadly elliptical or orbicular, apex acuminate, shorter than the pedicellate ovary, decreasing in size towards the inflorescence apex, green. *Flowers* 3–8, resupinate, green, glabrous; pedicellate ovary 19–21(–30) mm long, strongly arched; ovary 12–16 mm long, pedicel 5–15 mm long. Sepals green, apex apiculate; dorsal sepal 6–9 × 4.5–6.5 mm, concave, broadly elliptical when flattened; lateral sepals 8.5–12 × 3–4 mm, obliquely lance-ovate, deflexed. *Petals* bipartite; posterior segment 6–9 × 1.5–2.1 mm, falcate, apex acute-acuminate, connivent with the dorsal sepal; anterior segment 11–16 mm long, 1.6–2.2 times longer than the posterior segment, linear-filiform, inserted above the base of the posterior segment, directed backwards, bent upwards. *Lip* tripartite; undivided basal part 1–1.6 × 1.7–2.6 mm; lateral segments 13–18 mm long, 1.4–1.9 times longer than the median segment, linear-filiform, reflexed or bent upward; median segment 8–12 × 1–1.5 mm, linear, reflexed or slightly deflexed, more or less parallel to the distal part of the ovary; spur 13–17 mm long, shorter than the pedicellate ovary, clavate, base 1–1.3 mm wide, apex 1.6–2 mm wide, free or with the apex embraced by the bracts, more or less genuflexus, green. *Gynostemium* 4–6 mm high; connective 0.5 mm wide, green, apex retuse; lateral appendages (auricles) 0.5–1 × 1–1.6 mm, fleshy, verrucose, apex truncate, green. *Anther* 1.5–2 mm high, bilocular, loculi parallel, translucent; canals 1.8–2.3 mm long, parallels, curved upwards; pollinaria 2, 2.2–3.8 mm long each, separate; caudicles 3–4 mm long, filiform; pollinia 1.4–2 × 1–1.5 mm, yellow; viscidia ca 0.24 × 0.27 mm. *Stigmatophores* (stigma lobes) 2, 2–3 mm long, closely parallel, green, receptive surface 1.3–1.7 mm long, slightly convex, turned frontwards, inner margins around the spur entrance thickened, apex 1.6–2.3 mm wide, truncate or rounded. *Rostellum* 4–6 mm long, green; mid lobe 1–1.4 mm long, 1.8–2.3 mm high, triangular, fleshy, apex narrowly acute, held between the anther loculi; side-lobes 2.5–3.7 mm long, base ca 1 mm wide, apex ca 0.5 mm wide, curved, apex facing upwards, parallels. Fruit not seen.

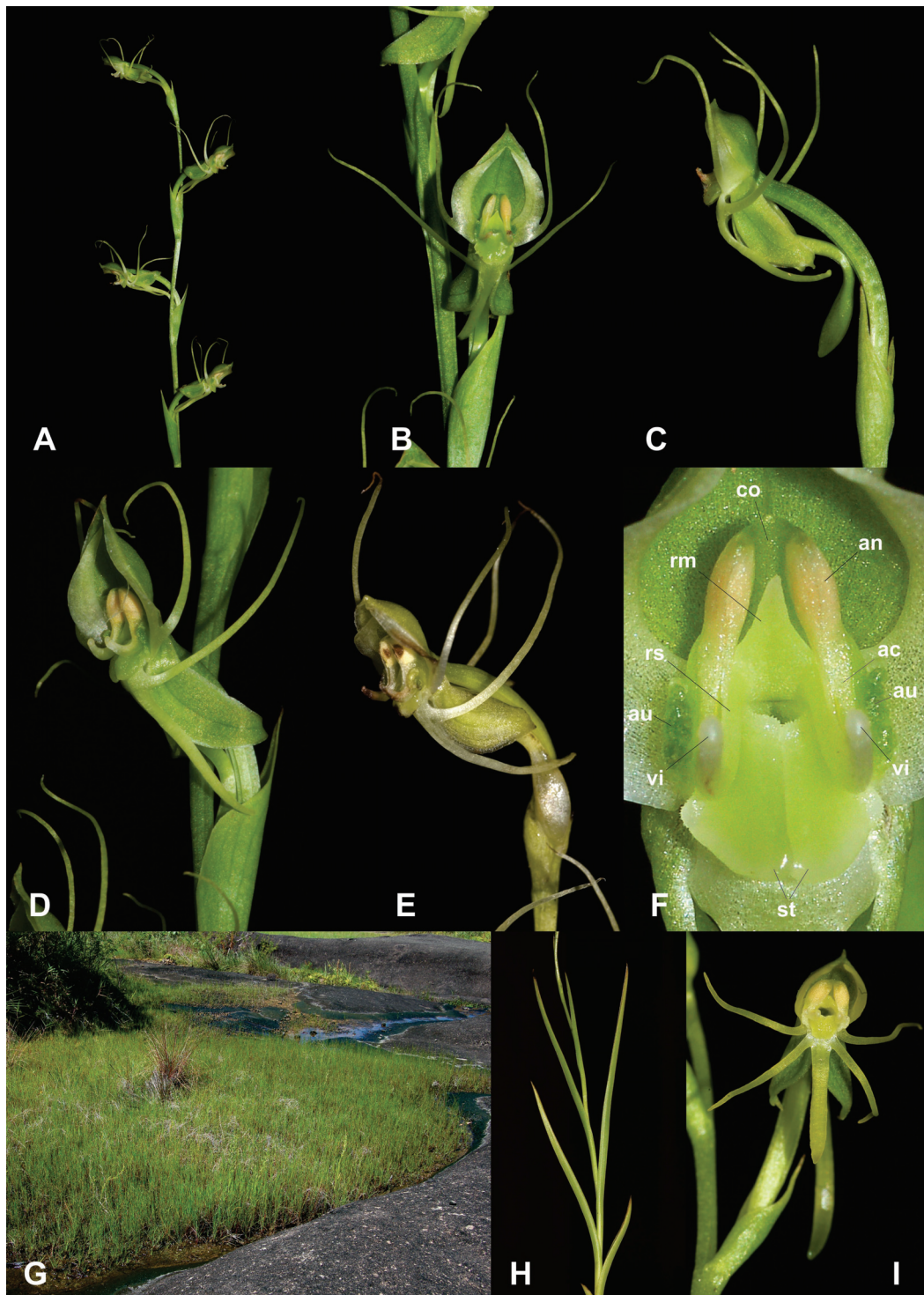


Fig. 5. *Habenaria proiteana* J.A.N. Bat., A.A.Vale & Bianch. sp. nov. **A.** Inflorescence. **B.** Flower, front view. **C.** Flower, side view. **D–E.** Flower, $\frac{3}{4}$ lateral view. **F.** Gynostemium, front view. *Habenaria cultellifolia*. **G.** Habitat (Chapadas dos Veadeiros, Goiás). **H.** Vegetative part. **I.** Flower, front view. A–D, F from Batista *et al.* 2376 (BHCB); E from Batista & Bianchetii 3101 (BHCB); G–I from Batista *et al.* 2959 (BHCB). Abbreviations: ac = anther canals; an = anther; au = auricles; co = connective; rm = rostellum mid lobe; rs = rostellum side-lobes; st = stigmatophores; vi = viscidium. All photographs by J.A.N. Batista.

Distribution, Habitat and Phenology

It is restricted to Chapada dos Veadeiros in the State of Goiás, occurring in *campos rupestres*, amid rocky outcrops in sandy, dry soil or seasonally humid campo limpo, in dark, sandy-clay soils. It flowers during the peak of the rainy season, in December and January.

Conservation Assessment

Habenaria proiteana sp. nov. is currently known from only three populations, all from Chapada dos Veadeiros, and shows a small EOO estimated at 37 km² and a small AOO of 12 km². Based on the IUCN Red List Categories and Criteria and its guidelines (IUCN 2012, 2016), the species can be tentatively classified as Critically Endangered CR B1ab(iii)+2ab(iii); C2a(i).

Taxonomic notes

Habenaria proiteana sp. nov., *H. australis* J.A.N.Bat., Vale & Menini, and *H. rodriguesii* present the anterior segment of the petal inserted a few millimetres above the posterior segment's base, not at the base as in the great majority of Neotropical *Habenaria*. However, *H. proiteana* sp. nov. differs from these species in the following characters: the petal anterior segment is proportionally longer (2.2–2.8 times longer than the posterior segment vs 1.2–1.8 times longer in *H. australis*), and the spur genuflexus (Figs. 5C, 6J–L) (vs straight or only slightly curved in *H. australis* and *H. rodriguesii*). In addition to these three species, only two other Neotropical species also have this distinctive character: *H. secundiflora* and *H. meeana* Toscano. Nonetheless, *H. secundiflora* has smaller flowers (dorsal sepal 4–6 × 3–4.5 mm), the lateral segments of the lip deflexed, and pollinaria joined by the viscidia, while *H. meeana* also has smaller flowers (dorsal sepal 5–6 × 3–4 mm), the anterior segment of the petal inserted only about 2 mm above the base of the posterior segment, and lateral sepals and lip segments strongly reflexed, embracing the ovary.

Habenaria proiteana sp. nov. is also similar to *H. sobraliana* in general morphology, but distinct by the petal anterior segment 11–16 mm long, inserted a few millimetres above the posterior segment's base (vs 5–8 mm long, inserted at the base of the posterior segment in *H. sobraliana*) and distribution restricted to the Chapada dos Veadeiros (vs restricted to the State of Rio Grande do Sul).

Key to the Brazilian species of *Habenaria* with the petal anterior segment inserted above the base of the posterior segment

1. Pedicellate ovary 19–28 mm long; dorsal sepal 10–14 mm long; petal anterior segment 20–33 mm long; spur 16–21 mm long..... *H. rodriguesii* Cogn.
– Pedicellate ovary 12–21(–30) mm long; dorsal sepal 5–10 mm long; petal anterior segment 8–16 mm long, occasionally absent; spur 8.5–17 mm long..... 2
2. Lip lateral segments erect (turned upwards)..... 3
– Lip lateral segments deflexed (turned downwards)..... 4
3. Pedicellate ovary 19–21(–30) mm long, strongly arched; dorsal sepal broadly elliptical when flattened; petal anterior segment 11–16 mm long; spur 13–17 mm long, more or less genuflexus; restricted to the Chapada dos Veadeiros, State of Goiás, Brazil..... *H. proiteana* sp. nov.
– Pedicellate ovary 12–18 mm long, slightly arched; dorsal sepal ovate or elliptical when flattened; petal anterior segment (0)8–12 mm; spur 8.7–10.3 mm long, more or less straight; restricted to the Serra Gaúcha in the State of Rio Grande do Sul, Brazil.....
..... *H. australis* J.A.N. Bat., A.A. Vale & Menini
4. Pedicellate ovary erect, more or less parallel with the rachis; dorsal sepal ovate or elliptical not reclinate, covering the gynostemium; lateral sepal 2–2.5 mm wide; lip lateral segments deflexed, longer

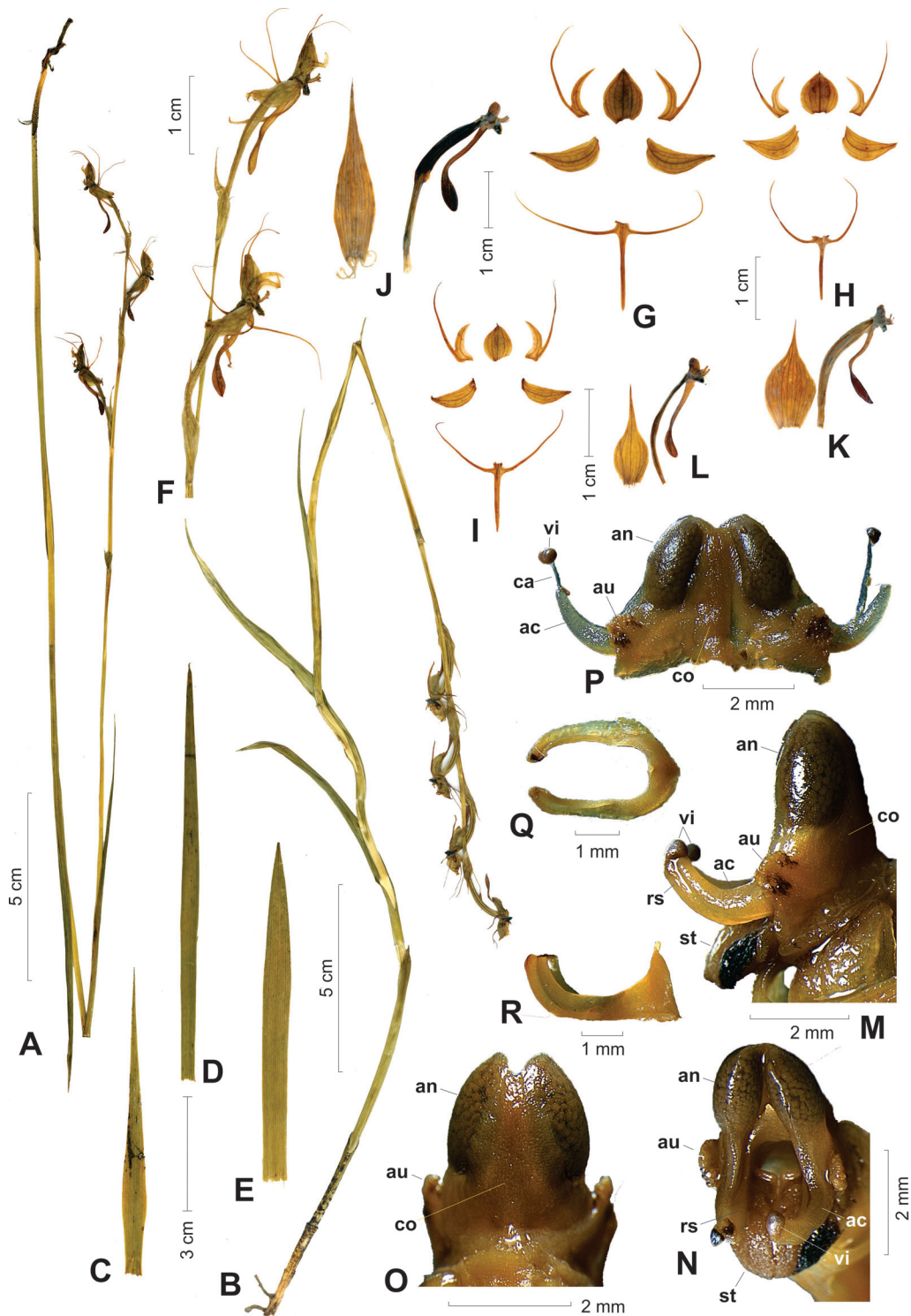


Fig. 6. *Habenaria proiteana* J.A.N. Bat., A.A.Vale & Bianch. sp. nov. **A–B.** Habit. **C–E.** Leaves. **F.** Flowers. **G–I.** Dissected perianth. **J–L.** Bract, pedicellate ovary, gynostemium and spur, lateral view. **M.** Gynostemium, side view. **N.** Gynostemium, front view. **O.** Gynostemium, back view. **P.** flattened connective, anther and lateral appendages (auricles). **Q.** Rostellum, upper view. **R.** Rostellum, lateral view. **A–H, J–K** and **M–R** from Batista *et al.* 2376 (BHCB); **I** and **L** from Hatschbach *et al.* 58383 (MBM). Abbreviations: ac = anther canal; an = anther; au = auricles; ca = caudicle; co = connective; rs = rostellum side-lobes; st = stigmatophores; vi = viscidium.

- than the median segment; anther canals and rostellum side-lobes held between the anther loculi; viscidia united; widely distributed from northern South America to southern Brazil.....
..... *H. secundiflora* Barb.Rodr.
- Pedicellate ovary perpendicular to the rachis; dorsal sepal obovate, partially reclinate, not covering the gynostemium; lateral sepals 3–4 mm wide; lip lateral segments reflexed, about the same size as the median segment; anther canals and rostellum side-lobes projected beyond the anther loculi; viscidia separated; restricted to the Chapada Diamantina in the State of Bahia, Brazil.....
..... *H. meeana* Toscano

Discussion

Survey and conservation of *Habenaria* in Chapada dos Veadeiros

The 61 taxa of *Habenaria* recorded for Chapada dos Veadeiros correspond to 74% of the 80 species of the genus recorded for the State of Goiás, according to Flora e Funga do Brasil (2022), while the region's area corresponds to only 6.3% of the state's total, reinforcing its importance for the preservation of the diversity of the genus in the state. The Chapada dos Veadeiros is one of the regions with the greatest diversity of the genus in Brazil, surpassed only by the Distrito Federal with 74 species (Batista & Bianchetti 2003), and followed by Serra do Cipó with 53 species (J.A.N. Batista, pers. comm.), and Serra da Canastra with 28 species (Carvalho *et al.* 2013) (Table 3).

Similar to other regions with surveys for *Habenaria* (Brade 1951; Batista & Bianchetti 2003; Batista *et al.* 2004, 2008; Carvalho *et al.* 2013), the number of endemic species is small, reinforcing that cases of restricted endemism are rare for the genus in Brazil (Table 3). This result contrasts with other groups that present high diversity in mountainous complexes of Southeastern and Central Western Brazil, such as Eriocaulaceae and the genus *Mimosa* (Simon & Proença 2000), which present a high number of endemic species. For the subfamily Paepalanthoideae (Eriocaulaceae), for example, more than 50% of the 42 species that occur in Chapada dos Veadeiros are endemic to the area (Trovó & Echternacht 2022).

Our results suggest that a significant portion of the diversity of the genus in the region does not occur or is underrepresented within the PNCV (Table 1). However, this data must be considered with reservations since many records reported in this study do not show coordinates or precise geographical locations, making it impossible to know exactly where they were collected. Collections in the region are centred along access roads, mainly on the GO-118 highway between Alto Paraíso de Goiás and Teresina de Goiás and GO-239 between Alto Paraíso and the village of São Jorge (Fig. 1). Large extensions of the region have not yet been sufficiently botanically collected or have no collections at all, including the central and north-western parts of the park as well as the extensive area added to the park in 2016 (Fig. 1), which includes some localities with the highest altitudes in the region and for which there are currently no herbarium collections available. More comprehensive and systematic explorations are expected to reveal new records and eventual taxonomic novelties within the *Habenaria* from this region, considering the exceptional richness of the genus in the Chapada dos Veadeiros. In this work, for example, we describe three new taxa and record the only known extant population of *H. cultellifolia*, a species known only from type material collected at least 127 years ago. These discoveries indicate that although some parts of the region have been extensively explored, new novelties are still to be expected, particularly among the terrestrial, small and ephemeral species.

Efforts to resolve the taxonomy of Brazilian species of *Habenaria* have intensified in the last years with the study of species complexes and the description of several new species (Batista *et al.* 2017, 2022a, b; Cruz-Lustre *et al.* 2020, 2022; Lau *et al.* 2021). Nevertheless, for the Chapada dos Veadeiros at least three morphotypes of the *Habenaria nuda* complex and some taxa associated with *H. montis-*

Table 2. Characterization of the markers used in the phylogenetic analyses. Ter = number of terminals; Char = number of aligned characters in base pairs; MD = percentage of missing data; VU = number of variable uninformative characters; VI = number of variable informative characters; CI = consistency index; RI = retention index. hLRTs = hierarchical likelihood ratio tests.

| Marker | Ter | Char | Total MD | Neotropical MD | VU | VI (%) | CI | RI | Model (hLRTs) |
|-------------------|-----|------|----------|----------------|-----|----------------|------|------|---------------|
| ITS | 105 | 760 | 0.2% | 0% | 111 | 291 (38.3%) | 0.60 | 0.77 | GTR+I+G |
| ETS | 82 | 554 | 24.5% | 9% | 157 | 157 (28.4%) | 0.72 | 0.79 | HKY+G |
| Nuclear | 105 | 1314 | 10.4% | 3.8% | 268 | 448 (34%) | 0.64 | 0.77 | – |
| <i>trnK</i> | 100 | 419 | 9% | 2.8% | 49 | 52 (12.4%) | 0.70 | 0.85 | F81+G |
| <i>matK</i> | 105 | 1594 | 3.8% | 1.6% | 156 | 207 (13%) | 0.66 | 0.83 | GTR+I+G |
| <i>rps16-trnK</i> | 86 | 1045 | 12.6% | 1.9% | 114 | 134 (12.8%) | 0.70 | 0.84 | GTR+I+G |
| Plastid | 105 | 3058 | 7.5% | 1.8% | 319 | 393 (12.8%) | 0.66 | 0.82 | – |
| Total | 105 | 4372 | 8.4% | 2.4% | 587 | 841 (19.2%) | 0.63 | 0.78 | – |

wilhelminae Renz, *H. rodeiensis* Barb. Rodr., and *H.pseudoculicina* (Table 1) are still unresolved and additional studies are needed to uncover the identity and taxonomy of these taxa.

Phylogenetic analyses

In terms of overall results of the phylogenetic analyses, the main differences between our results and some of the earlier molecular phylogenetic studies (Batista *et al.* 2013, 2014, 2016, 2018; Cruz-Lustre *et al.* 2022; Lau *et al.* 2021) were the position of *H. monorrhiza* (Sw.) Rchb.f. and subclade 1 (*H. sect. Pratenses* Kraenzl.), which were recovered in a polytomy at the base of the Neotropical clade, while in our results, they were internally recovered in the Neotropical clade (Fig. 2). Nevertheless, in both cases, the results had low support, showing that the phylogenetic relationships of these taxa within the Neotropical clade are not yet resolved. In another difference, species from subclades 16, 17 and 18 were intermingled into a single subclade, while in the topology of Batista *et al.* (2013) these subclades were recovered separately with high support. Although our dataset included a greater number of markers and characters (five markers and 4,372 characters vs two markers and 1,389 characters in the study of Batista *et al.* 2013), our sampling of taxa from these subclades was smaller (6 taxa vs 15 taxa), which may have reduced the resolution of the relationships between them.

As for the new species described here, both were recovered with high support as sister to species with different floral morphology, but with similar vegetative morphology, habitat and ecology. Both *H. minuticalcar* sp. nov./ *H. pseudoculicina* and *H. proiteana* sp. nov./ *H. psammophila*, occur in ‘campos rupestres’, in seasonally humid, sandy soils in isolated montane areas of central Brazil. In the pair *H. minuticalcar* sp. nov./ *H. pseudoculicina* both plants are slender with linear and reduced leaves, while in *H. pseudoculicina* the anterior segment of the petals (5.0–6.5 mm long) is about 1.7 times longer than the posterior segment (2.8–3.8 mm long), the lateral segments of the lip (5.0–7.5 mm long) are 1.3 to 1.4 times longer than the median segment (3.5–5.0 mm long) and the spur

Table 3. *Habenaria* Willd. data for localities or regions in Brazil with checklists of the genus with records above 10 species. Abbreviations: APA = Área de Proteção Ambiental; P.E. = Parque Estadual; P.N. = Parque Nacional; R.P.P.N. = Reserva Particular do Patrimônio Natural. Abbreviations for Brazilian states: BA = Bahia; GO = Goiás; MG = Minas Gerais; RJ = Rio de Janeiro.

| Locality or region (State) | N ° spp. (infra-specific taxa) | N ° endemic taxa (% total) | Area (km ²) | Reference |
|--|--------------------------------------|----------------------------------|----------------------------|-----------------------------|
| Distrito Federal | 74 (3) | 8 (10.4%) | 5,783 | Batista & Bianchetti 2003 |
| Chapada dos Veadeiros (GO) | 59 (2) | 4 (6.6%) | 21,475 | This work |
| Serra do Cipó/APA Morro da Pedreira (MG) | 52 (3) | 2 (3.8%) | 1,317 | Batista, pers. comm. |
| R.P.P.N. Santuário do Caraça (MG) | 33 (1) | 1 (2.9%) | 112 | Mota 2006 |
| P.N. da Serra da Canastra (MG) | 28 (0) | 2 (7.1%) | 2,000 | Carvalho <i>et al.</i> 2013 |
| Lagoa Santa (MG) | 17 (0) | 0 | 108 | Warming 1892 |
| P.N. do Itatiaia (RJ) | 17 (0) | 0 | 119 | Brade 1951 |
| P.E. da Serra do Ouro Branco (MG) | 17 (1) | 0 | 27 | Vieira & Barros 2017 |
| P.E. do Itacolomi (MG) | 15 (0) | 1 (6.7%) | 75 | Batista <i>et al.</i> 2004 |
| Pico das Almas (BA) | 12 (0) | 0 | 170 | Toscano-de-Brito 1995 |

(8.0–11.7 mm long) is about the same length as the pedicellate ovary (12–15 mm long). On the other hand, the morphology of the column is similar between the two species. In particular, the rostellum is very similar, is completely held between the anther loculi, and the pollinaria are joined by the viscidia in both species. However, these features are not unique and are shared with other non-directly related species groups, such as the species in subclade 20 (*H. sect. Microdactylae* Kraenzl.) and subclade 15, among others.

For the pair *H. proiteana* sp. nov./ *H. psammophila* the differences are greater than the similarities. Vegetatively, *H. psammophila* has sub-patent, papery, somewhat rigid leaves with the limb mostly detached from the stem. In floral morphology, the flowers of *H. psammophila* are smaller, the spur (3.0–4.5 mm long) is shorter than the pedicellate ovary (7–11 mm long), and the morphology of the column is quite different between the two species. The only obvious similarities between *H. proiteana* sp. nov./ *H. psammophila* are the anterior segment of the petals longer than the posterior segment and the lateral segments of the lip longer than the median segment. However, these features are also found in all species of the *H. nuda* complex and in several other Neotropical groups and species of the genus. An additional and perhaps more striking, but less obvious similarity, is the spur of *H. psammophila*. Although much smaller, it is also clavate and occasionally also more or less genuflexus, which are distinctive features of *H. proiteana* sp. nov.. Our results show that *H. proiteana* sp. nov. is not directly related to the species of the *H. nuda* complex (subclade 21), with which it shares a similar overall morphology in both vegetative and floral terms. This result suggests that both this general morphology as also the anterior petal segment inserted above the posterior segment's base, a characteristic shared with *H. rodriguesii*, and *H. australis*, are probably a result of convergence. On the other hand, *H. rodriguesii* and *H. australis* form a highly supported species pair and this morphological feature probably constitutes a synapomorphy for this pair within subclade 21.

Most subclades of Neotropical *Habenaria*, such as subclades 1, 2, 3 (Cruz-Lustre *et al.* 2022), 4, 8, 15, 20 (Batista *et al.* 2018) and the subclade formed by *H. espinhacensis* J.A.N.Bat. & A.A.Vale and

related species (Batista *et al.* 2016) have a high morphological uniformity. On the other hand, some subclades are otherwise homogeneous, but present one morphologically distinct species, as is the case of *H. montevidensis* Spreng. in subclade 5 (*H. sect. Pentadactylae* Kraenzl.) (Pedron *et al.* 2014) and *H. karstica* J.A.N.Bat. in subclade 12 (*H. sect. Spathaceae* Kraenzl.) (Batista *et al.* 2022a). In addition to these differences within the subclades, some species pairs from the Cerrado Phytogeographic Domain recovered as sisters with high support, such as *H. caldensis* Kraenzl. / *H. magniscutata* and now *H. minuticalcar* sp. nov. / *H. pseudoculicina* and *H. proiteana* sp. nov. / *H. psammophila* (Fig. 2) differ greatly in floral morphology. Since habitat, ecology and vegetative morphology are similar between these species pairs, and the pairs are often sympatrical, these results suggest speciation driven by different pollinators (Harder & Johnson 2009). In line with this interpretation, Pedron *et al.* (2012) showed that *H. montevidensis* is pollinated by butterflies, while other species of *H. sect. Pentadactylae*, are pollinated by moths or present floral morphology associated with moth pollination, suggesting that the morphological differences of *H. montevidensis* are a result of a shift to diurnal pollinators.

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References

- Bachman S., Moat J., Hill A.W., de la Torre J. & Scott B. 2011. Supporting red list threat assessments with GeoCAT: Geospatial conservation assessment tool. *ZooKeys* 150: 117–126. <https://doi.org/10.3897/zookeys.150.2109>
- Barbosa Rodrigues J. 1877. *Genera et Species Orchidearum Novarum I*. C. & H. Fleiuss, Rio de Janeiro.
- Barbosa Rodrigues J. 1882. *Genera et Species Orchidearum Novarum II*. Typographia Nacional, Rio de Janeiro.
- Batista J.A.N. & Bianchetti L.B. 2003. Lista atualizada das Orchidaceae do Distrito Federal, Brasil. *Acta Botanica Brasilica* 17 (2): 183–201. <https://doi.org/10.1590/S0102-33062003000200003>
- Batista J.A.N., Bianchetti L.B., Nogueira R.E., Pellizzaro K.F. & Ferreira F.E. 2004. The genus *Habenaria* (Orchidaceae) in the Itacolomi state park, Minas Gerais, Brazil. *Sitientibus Série Ciências Biológicas* 4 (1–2): 25–36.
- Batista J.A.N., da Silva J.B.F. & Bianchetti L.B. 2008. The genus *Habenaria* (Orchidaceae) in the Brazilian Amazon. *Brazilian Journal of Botany* 31 (1): 105–134. <https://doi.org/10.1590/S0100-84042008000100011>
- Batista J.A.N., Bianchetti L.B., R. González-Tamayo R., Figueroa X.M.C. & Cribb P.J. 2011a. A synopsis of new world *Habenaria* (Orchidaceae) I. *Harvard Papers in Botany* 16: 1–47. <https://doi.org/10.3100/025.016.0101>
- Batista J.A.N., Bianchetti L.B., R. González-Tamayo R., Figueroa X.M.C. & Cribb P.J. 2011b. A synopsis of new world *Habenaria* (Orchidaceae) II. *Harvard Papers in Botany* 16: 233–273. <https://doi.org/10.3100/0.25.016.0202>
- Batista J.A.N., Borges K.S., Faria M.W., Proite K., Ramalho A.J., Salazar G.A. & van den Berg C. 2013. Molecular phylogenetics of the species-rich genus *Habenaria* (Orchidaceae) in the New World based on nuclear and plastid DNA sequences. *Molecular Phylogenetics and Evolution* 67: 95–109.

<https://doi.org/10.1016/j.ympcv.2013.01.008>

Batista J.A.N., Proite K., Carvalho B.M., Vale A.A. & Felix L.P. 2014. From Cuba to most of the Neotropic: *Habenaria Bicornis* (Orchidaceae) is widespread from Mexico to Southeastern Brazil. *Lankesteriana* 13: 165–184. <https://doi.org/10.15517/lank.v13i3.14351>

Batista J.A.N., Vale A.A., Carvalho B.M., Proite K., Ramalho A.J., Munhoz A.C.D., van den Berg C. & Bianchetti L.B. 2016. Four new species in *Habenaria* (Orchidaceae) from the Espinhaço Range, Brazil. *Systematic Botany* 41: 275–292. <https://doi.org/10.1600/036364416X691858>

Batista J.A.N., Proite K. & Bianchetti L.B. 2017. Descriptions and phylogenetic relationships of four new species and a new name of *Habenaria* (Orchidaceae) from the cerrado and campos rupestres of Brazil. *Plant Systematics and Evolution* 303: 873–899. <https://doi.org/10.1007/s00606-017-1415-x>

Batista J.A.N., Reis A.F.S., Leite J.L.Jr. & Bianchetti L.B. 2018. Phylogeny and taxonomy of *Habenaria leprieurii*, *H. alpestris* and *H. sect. Microdactylae* (Orchidaceae), with descriptions of two new taxa. *Phytotaxa* 373: 241–271. <https://doi.org/10.11646/phytotaxa.373.4.1>

Batista J.A.N., Ferreira V.L., Alves C.I.G. & Stehmann J.R. 2022a. A new species of *Habenaria* (Orchidaceae, Asparagales) and a checklist of Orchidaceae from limestone outcrops of Brazil. *European Journal of Taxonomy* 828: 16–44. <https://doi.org/10.5852/ejt.2022.828.1855>

Batista J.A.N., Lau B.L., Costa I.G.C.M., Massensini Junior A., Mota R.C. & Borba E.L. 2022b. Description of four new species of the *Habenaria repens* complex (Orchidaceae) from Brazil based on a biosystematic and molecular phylogenetic approach. *Systematic Botany* 47: 215–231. <https://doi.org/10.1600/036364422X16442668423536>

Brade A.C. 1951. O Gênero *Habenaria* (Orchidaceae) no Itatiaia. *Rodriguésia* 14 (26): 7–21.

Carvalho B.M., Ramalho A.J. & Batista J.A.N. 2013. O gênero *Habenaria* (Orchidaceae) na Serra da Canastra, Minas Gerais, Brasil. *Rodriguésia* 64 (2): 223–245. <https://doi.org/10.1590/S2175-78602013000200003>

Cogniaux A. 1893–1896. Orchidaceae. In: Martius C.F.P., Eichler A.G. & Urban I. (eds) *Flora brasiliensis*, vol. 3: 1–672. F. Fleischer, München.

Cogniaux A. 1906. Notes sur les orchidées du Brésil et des régions voisines. *Bulletin de la Société Royale de Botanique de Belgique* 43: 266–350.

Cribb P.J. 2001. *Habenaria*. In: Pridgeon A.M., Cribb P.J., Chase M.W. & Rasmussen F.N. (eds) *Genera orchidacearum*, vol. 2, *Orchidoideae* (part1) : 298–303. Oxford University Press Inc., Oxford.

Cruz-Lustre G., Batista J.A.N., Radins J.A., González A. & Borba E.L. 2020. Morphometric analysis of the *Habenaria parviflora* complex (Orchidaceae). *Plant Systematics and Evolution* 306: 37. <https://doi.org/10.1007/s00606-020-01634-2>

Cruz-Lustre G., Castro C., Borba E.L. & Batista J.A.N. 2022. Phylogenetics and taxonomy of *Habenaria* sect. *Micranthae* (Orchidaceae), with the description of an overlooked new species from the Espinhaço mountain range, Eastern Brazil. *Systematics and Biodiversity* 20 (1): 2030424. <https://doi.org/10.1080/14772000.2022.2030424>

Doyle J.F. & Doyle J.L. 1987. A rapid DNA isolation procedure for small quantities of fresh leaf tissue. *Phytochemical Bulletin* 19: 11–15.

Erixon P., Sennblad B., Britton T. & Oxelman B. 2003. Reliability of Bayesian posterior probabilities and bootstrap frequencies in phylogenetics. *Systematic Biology* 52: 665–673. <https://doi.org/10.1080/10635150390235485>

- ESRI(2016)ArcGISDesktop:Release10.EnvironmentalSystemsResearchInstitute,Redlands.[Computer software]. <https://www.redlands.edu/study/schools-and-centers/css/resources/arcgis-desktop/>
- 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>
- Fitch W.M. 1971. Toward defining the course of evolution: minimum change for a specific tree topology. *Systematic Zoology* 20: 406–416. <https://doi.org/10.1093/sysbio/20.4.406>
- Flora e Funga do Brasil 2022. Orchidaceae. Jardim Botânico do Rio de Janeiro. Available from <https://floradobrasil.jbrj.gov.br/FB11639> [accessed 12 August 2022].
- Govaerts R., Nic Lughadha E., Black N., Turner R. & Paton A. 2021. The World Checklist of Vascular Plants, a continuously updated resource for exploring global plant diversity. *Scientific Data* 8: 215. <https://doi.org/10.1038/s41597-021-00997-6>
- Harder L.D. & Johnson S.D. 2009. Darwin’s beautiful contrivances: evolutionary and functional evidence for floral adaptation. *New Phytologist* 183(3): 530–545. <https://doi.org/10.1111/j.1469-8137.2009.02914.x>
- IUCN2012.*IUCNredlistcategoriesandcriteriaversion3.1*.Ed.2.IUCN,Gland,SwitzerlandandCambridge, UK. Available from <https://www.iucnredlist.org/technical-documents/categories-and-criteria> [accessed 9 February 2017].
- IUCN 2016. *Guidelines for using the IUCN red list categories and criteria version 12*. Prepared by the standards and petitions subcommittee. IUCN Species Survival Commission. IUCN, Gland, Switzerland and Cambridge, UK. Available from <https://www.iucnredlist.org/documents/RedListGuidelines.pdf> [accessed 9 Feb. 2017].
- Jin W.-T., Schuiteman A., Chase M.W., Li J.-W., Chung S.-W., Hsu T.-C. & Jin X.-H. 2017. Phylogenetics of subtribe Orchidinae s.l. (Orchidaceae; Orchidoideae) based on seven markers (plastid *matK*, *psaB*, *rbcL*, *trnL-F*, *trnH-psba*, and nuclear nrITS, *Xdh*): implications for generic delimitation. *BMC Plant Biology* 17: 222. <https://doi.org/10.1186/s12870-017-1160-x>
- Katoh K. & Standley D.M. 2013. MAFFT multiple sequence alignment software version 7: improvements in performance and usability. *Molecular Biology and Evolution* 30: 772–780. <https://doi.org/10.1093/molbev/mst010>
- Kränzlin F. 1892. Beitrage zu einer monographie der gattung *Habenaria* Willd. *Botanisch Jahrbücher für Systematik* 16: 52–223.
- Kränzlin F. 1911. Beitrage zur Orchideenflora Südamerikas. *Kongliga Svenska Vetenskapsakademiens Handlingar* 46: 1–105.
- Kress W.J., Prince L.M. & Williams K.J. 2002. The phylogeny and a new classification of the gingers (Zingiberaceae): evidence from molecular data. *American Journal of Botany* 89: 1682–1696. <https://doi.org/10.3732/ajb.89.10.1682>
- Lau B.L., Batista J.A.N., Massensini Junior A., Whitten W.M. & Borba E.L. 2021. Unravelling the *Habenariarepens* (Orchidaceae) complex in Brazil: a biosystematic and molecular phylogenetic approach. *Botanical Journal of the Linnean Society* 197: 229–248. <https://doi.org/10.1093/botlinnean/boab>
- Lima P.C.A. & Franco J.L.A. 2014. As RPPNs como estratégia para a conservação da biodiversidade: O caso da chapada dos veadeiros. *Sociedade & Natureza (Uberlândia)* 26 (1): 113–125. <https://doi.org/10.1590/1982-451320140108>
- Lindley J. 1830–1840. *The genera and species of Orchidaceous plants*. Ridgways, London.

- Madeira F., Park Y.M., Lee J., Buso N., Gur T., Madhusoodanan N., Basutkar P., Tivey A.R.N., Potter S.C., Finn R.D. & Lopez R. 2019. The EMBL-EBI search and sequence analysis tools APIs in 2019. *Nucleic Acids Research* 47: W636–W641. <https://doi.org/10.1093/nar/gkz268>
- Mendonça R.C., Filgueiras T.S. & Fagg C.W. 2007. Análise florística da Chapada dos Veadeiros. In: Felfili J.M., Rezende A.V. & Silva-Junior M.C. (eds) *Biogeografia do bioma Cerrado: vegetação e solos da Chapada dos Veadeiros*: 119–237. Editora da Universidade de Brasília/Finattec, Brasília.
- Meneguzzo T.E.C. 2013. 752. *Sacoila cerradicola*: Orchidaceae. *Curtis's Botanical Magazine* 30 (1): 18–27. <https://doi.org/10.1111/curt.12013>
- Menezes L.C. 1992. *Encyclia chapadensis*. *Orchid Digest* 56 (3): 133.
- Miller M.A., Pfeiffer W. & Schwartz T. 2010. Creating the CIPRES Science Gateway for inference of large phylogenetic trees. In: *Proceedings of the Gateway Computing Environments Workshop (GCE)*: 1–8. Gateway Computing, New Orleans. <https://doi.org/10.1145/2016741.2016785>
- Monteiro S.H.N., Selbach-Schnadelbach A., Oliveira R.P. & van den Berg C. 2010. Molecular phylogenetics of *Galeandra* (Orchidaceae : Catasetinae) based on plastid and nuclear DNA sequences. *Systematic Botany* 35: 476–486. <https://doi.org/10.1600/036364410792495944>
- Mota R.C. 2006. *Orchidaceae na Serra do Caraça, Minas Gerais: Levantamento florístico com ênfase no estudo taxonômico da subfamília Epidendroideae*. Masters thesis, Universidade Federal de Minas Gerais, Brazil.
- Munhoz C.B.R. & Proença C.E.B. 1998. Composição florística do Município de Alto Paraíso de Goiás na Chapada dos Veadeiros. *Boletim do Herbário Ezechias Paulo Heringer* 3: 102–150.
- Ngugi G., Péchon T., Martos F., Paillere T., Bellstedt D.U. & Bytebier B. 2020. Phylogenetic relationships amongst the African genera of subtribe Orchidinae s.l. (Orchidaceae; Orchideae): Implications for subtribal and generic delimitations. *Molecular Phylogenetics and Evolution* 153: 106946. <https://doi.org/10.1016/j.ympev.2020.106946>
- Nylander, J.A.A. 2004. MrModeltest v2. Program distributed by the author. Uppsala: Evolutionary Biology Centre, Uppsala University.
- Pansarin E.R. 2004. *Cleistes pusilla* (Orchidaceae): A new species from central Brazil. *Kew Bulletin* 59 (4): 555–558. <https://doi.org/10.2307/4110910>
- Pedron M., Buzatto C.R., Singer R.B., Batista J.A.N. & Moser A. 2012. Pollination biology of four sympatric species of *Habenaria* (Orchidaceae: Orchidinae) from southern Brazil. *Botanical Journal of the Linnean Society* 170: 141–156. <https://doi.org/10.1111/j.1095-8339.2012.01285.x>
- Pedron M., Buzatto C.R., Ramalho A.J., Carvalho B.M., Radins J.A., Singer R.B. & Batista J.A.N. 2014. Molecular phylogenetics and taxonomic revision of *Habenaria* section *Pentadactylae* (Orchidaceae, Orchidinae). *Botanical Journal of the Linnean Society* 175: 47–73. <https://doi.org/10.1111/boj.12161>
- Reichenbach f. H.G. 1847. Orchidiographische Beiträge. *Linnaea* 19: 369–379.
- Reichenbach f. H.G. 1881. VII–Novitiae Orchidaceae Warmingianae. *Otia Botanica Hamburgensia* 2: 48–65.
- Renz J. 1992. The genus *Habenaria* (Orchidaceae) in the Guianas. *Candollea* 47: 483–512.
- 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. *Systematic Biology* 61: 539–542. <https://doi.org/10.1093/sysbio/sys029>

- Santos I.S. & Silva M.J. 2019. O gênero *Bulbophyllum* Thouars (Orchidaceae, Epidendroideae) no Parque Nacional da Chapada dos Veadeiros, Goiás, Brasil. *Hoehnea* 46: e872018. <https://doi.org/10.1590/2236-8906-87/2018>
- Santos I.S. & Silva M.J. 2020. *Epidendrum* L. (Orchidaceae, Epidendroideae) no Parque Nacional da Chapada dos Veadeiros, Estado de Goiás, Brasil. *Hoehnea* 47: e202020. <https://doi.org/10.1590/2236-8906-20/2020>
- Simon M.F. & Proença C. 2000. Phytogeographic patterns of *Mimosa* (Mimosoideae, Leguminosae) in the Cerrado biome of Brazil: an indicator genus of high-altitude centers of endemism? *Biological Conservation* 96: 279–296. [https://doi.org/10.1016/S0006-3207\(00\)00085-9](https://doi.org/10.1016/S0006-3207(00)00085-9)
- Simpson M.G. 2006. *Plant Systematics*. Elsevier Academic Press, Burlington, San Diego, and London.
- Stace C.A. 1989. *Plant taxonomy and biosystematics*. 2nd Ed. Edward Arnold, Hodder and Stoughton Limited, London.
- Staden R. 1996. The Staden sequence analysis package. *Molecular Biotechnology* 5: 233–241. <https://doi.org/10.1007/BF02900361>
- Stearn W.T. 1992. *Botanical Latin*. 4th Ed. Timber Press, Portland, Oregon.
- Swofford D.L. 2002. PAUP* Phylogenetic analysis using parsimony (*and other methods). Version 4. Sinauer Associates, Sunderland.
- Thiers B. 2022. [continuously updated] Index Herbariorum: A global directory of public herbaria and associated staff. New York Botanical Garden's Virtual Herbarium. Available from <https://sweetgum.nybg.org/science/ih/> [accessed 12 August 2022].
- Toscano de Brito A.L.V. 1995. Orchidaceae. In: Stannard B.L. (ed) *Flora of the Pico das Almas, Chapada Diamantina – Bahia, Brazil*: 725–767. Royal Botanic Gardens, Kew.
- Trovó M. & Echternacht L. 2022. The Paepalanthoideae (Eriocaulaceae) of the Chapada dos Veadeiros National Park, Brazil: taxonomic novelties, identification key, and illustrated list of species. *European Journal of Taxonomy* 834: 58–93. <https://doi.org/10.5852/ejt.2022.834.1899>
- Vieira T.L & Barros F. 2017. Orchidaceae na Serra do Ouro Branco, Minas Gerais, Brasil. *Rodriguésia* 68(2): 691–747. <https://doi.org/10.1590/2175-7860201768224>
- Warming E. 1892. Lagoa Santa: Et Bidrag til den biologiske Plantegeografi. Kongelige Danske videnskabernes Selskabs Skrifter, *Naturvidenskabeli Mathematisk Afdeling* 6: 159–455.

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Supplementary file 1 (continued on the next 7 pages). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|---|--|---------------|----------|------------------|-------------------|-----------|
| Neotropical | | | | | | |
| <i>Habenaria abscondita</i> J.A.N.Bat. & G.Cruz-Lustre | <i>Cruz-Lustre</i> 206 (BHCB) | Brazil, BA | MW567131 | MW675883 | MW675857 | MW675870 |
| <i>Habenaria achalensis</i> Kraenzl. | <i>Batista</i> 2506 (BHCB) | Brazil, RS | HM777526 | KJ021403 | KJ021457 | MT623244 |
| <i>Habenaria alpe stris</i> Cogn. | <i>Batista</i> 2539 (BHCB) | Brazil, MG | HM777688 | KJ021361 | KJ021416 | MT623218 |
| <i>Habenaria aranifera</i> Lindl. | <i>Batista</i> 2472 (BHCB) | Brazil, RS | HM777626 | KJ021394 | KJ021448 | MT623173 |
| <i>Habenaria armata</i> Rehb.f. | <i>Batista</i> 1297 (CEN) | Brazil, DF | HM777677 | KJ021367 | KJ021422 | – |
| <i>Habenaria australis</i> J.A.N. Bat., A.A. Vale & Menini | <i>Batista</i> 2496 (BHCB) | Brazil, RS | HM777724 | OP589163* | OP589994* | OP589151* |
| <i>Habenaria ayangannensis</i> Renz | <i>Batista</i> 1919 (BHCB) | Brazil, MG | HM777706 | KJ021360 | KJ021415 | – |
| <i>Habenaria balansae</i> Cogn. | <i>Batista</i> 2336 (BHCB) | Brazil, MG | HM777683 | KJ021362 | KJ021417 | MT623217 |
| <i>Habenaria bicornis</i> Lindl. | <i>Felix 10803</i> (EAN) | Brazil, PB | KF998087 | KX784170 | KX784174 | MT623247 |
| <i>Habenaria brachydactyla</i> J.A.N.Bat. & Bianch. | <i>Batista & Proite 3109</i> (BHCB) | Brazil, DF | KX784176 | KX784171 | KX784175 | MT623252 |
| <i>Habenaria brachyphyton</i> Schltr. | <i>Batista</i> 2515 (BHCB) | Brazil, RS | HM777557 | MW675872 | MW675846 | MW675859 |
| <i>Habenaria brevidens</i> Lindl. | <i>Batista</i> 2616 (BHCB) | Brazil, MG | HM777535 | MW675873 | MW675847 | MW675860 |
| <i>Habenaria bryophila</i> J.A.N.Bat., B.L.Lau & I.G.C.Machado-Costa | <i>Batista et al. 1962</i> (BHCB) | Brazil, MG | HM777629 | MT623320 | MT623273 | MT623199 |
| <i>Habenaria caldensis</i> Kraenzl. | <i>Batista 250</i> (CEN) | Brazil, GO | HM777645 | KJ021363 | KJ021418 | MT623219 |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|--|---------------------------|---------------|----------|------------------|-------------------|-----------|
| <i>Habenaria campylogyna</i> J.A.N.Bat. & Bianch. | Batista 1901 (BHCB) | Brazil, MG | HM777581 | KJ021364 | KJ021419 | MT623220 |
| <i>Habenaria canastrensis</i> J.A.N. Bat. & B.M.Carvalho | Batista 1806 (BHCB) | Brazil, MG | HM777726 | OP589164* | OP589995* | OP589152* |
| <i>Habenaria compluviosa</i> J.A.N.Bat., B.L.Lau & I.G.C.Machado-Costa | Lau 23 (BHCB) | Brazil, MG | MT582228 | MT623313 | MT623267 | MT623194 |
| <i>Habenaria crucifera</i> Rchb.f. & Warm. | Batista 1826 (BHCB) | Brazil, MG | HM777574 | KJ021356 | KJ021411 | MT623212 |
| <i>Habenaria cultellifolia</i> Barb.Rodr. | Batista 1487 (CEN) | Brazil, GO | HM777675 | MW675874 | MW675848 | MW675861 |
| <i>Habenaria dutrae</i> Schltr. | Pedron 3 (ICN) | Brazil, RS | KJ021339 | KJ021387 | KJ021441 | – |
| <i>Habenaria edwallii</i> Cogn. | Batista 1717 (BHCB) | Brazil, MG | HM777564 | KJ021402 | KJ021456 | MT623243 |
| <i>Habenaria ekmaniana</i> Kraenzl. | Radins s.n. (BHCB) | Argentina | KJ021340 | KJ021383 | KJ021438 | MT623233 |
| <i>Habenaria espinhacensis</i> J.A.N.Bat. & A.A.Vale | Batista 2814 (BHCB) | Brazil, MG | HM777701 | MT623325 | MT623277 | MT623211 |
| <i>Habenaria exaltata</i> Barb.Rodr. | Batista 2771 (BHCB) | Brazil, MG | HM777621 | KJ021388 | KJ021442 | MT623234 |
| <i>Habenaria glaucophylla</i> Barb. Rodr. | Batista 761 (CEN) | Brazil, DF | HM777631 | KJ021373 | KJ021428 | MT623231 |
| <i>Habenaria gollmeri</i> Schltr. | Castro 2000 (COL) | Colombia | MW567127 | MW675875 | MW675849 | MW675862 |
| <i>Habenaria hamata</i> Hoehne | Batista 1519 (CEN) | Brazil, DF | HM777586 | KJ021371 | KJ021426 | MT623228 |
| <i>Habenaria henscheniana</i> Barb. Rodr. | Batista 2802 (BHCB) | Brazil, MG | HM777623 | KJ021376 | KJ021431 | – |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|---|--|---------------|------------|-------------------------|--------------------------|------------|
| <i>Habenaria hexaptera</i> Lindl. | <i>Batista</i> 2399 (BHCB) | Brazil, MG | HM777538 | KJ021366 | KJ021421 | MT623221 |
| <i>Habenaria hippocrepica</i> J.A.N.Bat. & Bianch. | <i>Batista</i> 1948 (BHCB) | Brazil, MG | HM777698 | MT623326 | MT623278 | MT623209 |
| <i>Habenaria hygrophila</i> J.A.N.Bat., B.L.Lau & I.G.C.Machado-Costa | <i>Batista</i> 2538 (BHCB) | Brazil, MG | MT582221 | MT623321 | MT623274 | MT623200 |
| <i>Habenaria imbricata</i> Lindl. | <i>Batista</i> 1123 (CEN) | Brazil, DF | HM777648 | KJ021368 | KJ021423 | MT623226 |
| <i>Habenaria irwiniana</i> J.A.N.Bat. & Bianch. | <i>Batista</i> 249 (CEN) | Brazil, GO | HM777642 | KX784167 | KX784172 | MT623251 |
| <i>Habenaria johannensis</i> Barb.Rodr. | <i>Mota</i> 2777 (BHCB) | Brazil, MG | HM777609 | KJ021398 | KJ021452 | MT623240 |
| <i>Habenaria karstica</i> J.A.N.Bat. | <i>Batista et</i> <i>al.</i> 3649 (BHCB) | Brazil, MG | ON197322 | ON168967 | ON168969 | – |
| <i>Habenaria lavrensis</i> Hoehne | <i>Batista</i> 673 (CEN) & <i>Batista</i> 1497 (CEN) | Brazil, DF | HM777711 | OP589165* | OP589996* | OP589153* |
| <i>Habenaria leptoceras</i> Hook. | <i>Batista</i> 2658 (BHCB) | Brazil, RJ | HM777597 | KJ021370 | KJ021425 | MT623227 |
| <i>Habenaria leucosantha</i> Barb.Rodr. | <i>Batista</i> 1604 (BHCB) | Brazil, DF | HM777568 | KJ021406 | KJ021460 | MT623248 |
| <i>Habenaria ludibundiciliata</i> J.A.N.Bat. & Bianch. | <i>Batista</i> 1372 (CEN) | Brazil, MA | HM777638 | MT623328 | MT623280 | MT623215 |
| <i>Habenaria macilentata</i> (Lindl.) Rchb.f. | <i>Batista</i> 2393 (BHCB) | Brazil, MG | HM777606 | KJ021400 | KJ021454 | MT623241 |
| <i>Habenaria macroceratitis</i> Willd. | <i>Whitten</i> 2827 (FLAS) | USA, FL | MT582231 | MT623329 | MT623281 | MT623230 |
| <i>Habenaria macronectar</i> (Vell.) Hoehne | <i>Batista</i> 2519 (BHCB) | Brazil, PR | HM777614 | KJ021399 | KJ021453 | MT623239 |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|---|----------------------------|------------|----------|------------------|-------------------|-----------|
| <i>Habenaria magniscutata</i> Catling | Batista 1227 (CEN) | Brazil, GO | HM777641 | OP589166* | OP589997* | OP589154* |
| <i>Habenaria megapotamensis</i> Hoehne | Pedron 10 (ICN) | Brazil, RS | KJ021345 | KJ021384 | KJ021439 | MT623235 |
| <i>Habenaria minuticalcar</i> J.A.N.Bat. & Bianch. sp. nov. | Batista 843 (CEN) | Brazil, GO | HM777691 | OP589167* | OP589998* | OP589155* |
| <i>Habenaria monorrhiza</i> (Sw.)Rchb.f. | Whitten 2721 (FLAS) | Ecuador | MT582232 | MT623330 | MT623282 | MT623249 |
| <i>Habenaria montevidensis</i> Spreng. | Batista 2479 (BHCB) | Brazil, RS | HM777619 | KJ021378 | KJ021433 | – |
| <i>Habenaria mystacina</i> Lindl. | Batista 1812 (BHCB) | Brazil, MG | HM777728 | KJ021353 | KJ021408 | MT623205 |
| <i>Habenaria nemorosa</i> Barb.Rodr. | Batista 2567 (BHCB) | Brazil, MG | HM777634 | MW675876 | MW675850 | MW675863 |
| <i>Habenaria</i> aff. <i>nuda</i> Lindl. | Batista 2869 (BHCB) | Brazil, MG | HM777718 | KJ021354 | KJ021409 | MT623206 |
| <i>Habenaria omissa</i> J.A.N.Bat. & Bianch. | Batista 1576 (BHCB) | Brazil, DF | HM777655 | KJ021357 | KJ021412 | MT623207 |
| <i>Habenaria pabstii</i> J.A.N.Bat. & Bianch. | Batista 2360 (BHCB) | Brazil, DF | HM777666 | OP589168* | OP589999* | OP589156* |
| <i>Habenaria pansarinii</i> J.A.N.Bat. & Bianch. | Batista et al. 1926 (BHCB) | Brazil, MG | HM777520 | KX784168 | KX784173 | MT623250 |
| <i>Habenaria paranaensis</i> Barb.Rodr. | Batista 2436 (BHCB) | Brazil, MG | HM777528 | KJ021405 | KJ021459 | MT623246 |
| <i>Habenaria parviflora</i> Lindl. | Batista 1813 (BHCB) | Brazil, MG | HM777560 | KJ021401 | KJ021455 | MT623242 |
| <i>Habenaria paulensis</i> Porsch | Batista 2481 (BHCB) | Brazil, RS | HM777556 | MW675878 | MW675852 | MW675865 |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|--|---|---------------|-----------|------------------|-------------------|-----------|
| <i>Habenaria pentadactyla</i> Lindl. | <i>Pedron 11</i> (ICN) | Brazil, RS | KJ021348 | KJ021380 | KJ021435 | MT623236 |
| <i>Habenaria pratensis</i> (Salzm. ex Lindl.) Rechb.f. | <i>Batista</i> <i>2686</i> (BHCB) | Brazil, BA | HM777546 | KJ021374 | KJ021429 | MT623225 |
| <i>Habenaria proiteana</i> J.A.N.Bat., A.A.Vale & Bianch. sp. nov. | <i>Vale 180</i> (BHCB) | Brazil, GO | OP584473* | OP589169* | OP590000* | OP589157* |
| <i>Habenaria proiteana</i> J.A.N.Bat., A.A.Vale & Bianch. sp. nov. | <i>Batista</i> <i>2376</i> (BHCB) | Brazil, GO | OP584474* | OP589170* | – | OP589158* |
| <i>Habenaria psammophila</i> J.A.N. Bat., Bianch. & B.M. Carvalho | <i>Batista</i> <i>1794</i> (BHCB) | Brazil, MG | HM777550 | OP589171* | OP590001* | OP589159* |
| <i>Habenaria pseudoculicina</i> J.A.N.Bat. & B.M.Carvalho | <i>Batista</i> <i>1808</i> (BHCB) | Brazil, MG | HM777707 | OP589172* | OP590002* | OP589160* |
| <i>Habenaria pungens</i> Cogn. | <i>Batista</i> <i>2095</i> (BHCB) | Brazil, GO | HM777570 | KJ021355 | KJ021410 | MT623213 |
| <i>Habenaria quinqueseta</i> (Michx.) Sw. | <i>Whitten</i> <i>3324</i> (FLAS) | USA, FL | MT582230 | MT623331 | MT623283 | MT623229 |
| <i>Habenaria reflexicalcar</i> J.A.N.Bat. & B.M.Carvalho | <i>Batista</i> <i>1744</i> (BHCB) | Brazil, MG | HM777700 | MT623332 | MT623284 | MT623210 |
| <i>Habenaria regnellii</i> Cogn. | <i>Batista</i> <i>2801</i> (BHCB) | Brazil, MG | HM777603 | KJ021391 | KJ021445 | – |
| <i>Habenaria repens</i> Nutt. | <i>Braun 01</i> (FLAS) | USA, FL | MT582195 | MT623296 | – | MT623180 |
| <i>Habenaria repens</i> Nutt. | <i>Batista</i> <i>2522</i> (BHCB) | Brazil, PR | HM777627 | KJ021395 | KJ021449 | MT623178 |
| <i>Habenaria rodeiensis</i> Barb.Rodr. | <i>Mota 2824</i> (BHCB) | Brazil, MG | HM777577 | KJ021359 | KJ021414 | MT623216 |
| <i>Habenaria rodriguesii</i> Cogn. | <i>Vale 122</i> (BHCB) | Brazil, MG | OP584475* | OP589173* | OP590004* | OP589162* |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|--|-------------------------------|---------------|----------|------------------|-------------------|-----------|
| <i>Habenaria schenckii</i> Cogn. | Batista 2882 (BHCB) | Brazil, BA | HM777580 | KJ021372 | KJ021427 | MT623232 |
| <i>Habenaria schwackei</i> Barb.Rodr. | Batista 2524 (BHCB) | Brazil, PR | HM777657 | MT623327 | MT623279 | MT623208 |
| <i>Habenaria secunda</i> Lindl. | Batista 2640 (BHCB) | Brazil, RJ | HM777525 | KJ021404 | KJ021458 | MT623245 |
| <i>Habenaria secundiflora</i> Barb.Rodr. | Batista 2392 (BHCB) | Brazil, MG | HM777637 | KJ021358 | KJ021413 | MT623214 |
| <i>Habenaria setacea</i> Lindl. | Mota 3019 (BHCB) | Brazil, MG | HM777731 | KJ021352 | KJ021407 | MT623204 |
| <i>Habenaria sobraliana</i> J.A.N.Bat., Vale & Menini | Batista 2499 (BHCB) | Brazil, RS | HM777704 | HM777990 | OP590003* | OP589161* |
| <i>Habenaria spathulifera</i> Cogn. | Bringel 411 (CEN) | Brazil, GO | MT582233 | MT623333 | MT623285 | MT623224 |
| <i>Habenaria subrepens</i> J.A.N.Bat., B.L.Lau & I.G.C.Machado-Costa | Mota 2285 (BHCB) | Brazil, MG | MT582217 | MT623307 | MT623264 | MT623191 |
| <i>Habenaria trifida</i> Kunth | Batista 1783 (BHCB) | Brazil, DF | HM777672 | KJ021369 | KJ021424 | MT623223 |
| <i>Habenaria ulei</i> Cogn. | Cruz- Lustre 281 (BHCB) | Brazil, SC | MW567130 | MW675880 | MW675854 | MW675867 |
| <i>Habenaria umbraticola</i> Barb.Rodr. | Mota 3569 (BHCB) | Brazil, MG | HM777605 | MW675881 | MW675855 | MW675868 |
| <i>Habenaria urbaniana</i> Cogn. | Batista 911 (CEN) | Brazil, MG | HM777658 | MT623334 | MT623286 | – |
| <i>Habenaria uruguayensis</i> Garay | Batista 2477 (BHCB) | Brazil, RS | KC257475 | MW675882 | MW675856 | MW675869 |
| <i>Habenaria warmingii</i> Rchb.f. & Warm. | Batista 2409 (BHCB) | Brazil, MG | HM777616 | KJ021396 | KJ021450 | MT623238 |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|---|---------|--------|----------|------------------|-------------------|----------|
| Paleotropical | | | | | | |
| <i>Bonatea speciosa</i> (L.f.) Willd. | GenBank | Africa | MT500586 | MT533490 | – | – |
| <i>Centrostigma occultans</i> (Welw. ex Rchb.f.) Schltr. | GenBank | Africa | MT500664 | MT533569 | – | – |
| <i>Habenaria crassilabia</i> Kraenzl. (<i>H. chejuensis</i>) | GenBank | Korea | KT338765 | NC_046821 | NC_046821 | MN221405 |
| <i>Habenaria tridens</i> Lindl. | GenBank | Africa | DQ522080 | DQ522101 | – | – |
| <i>Habenaria cornuta</i> Lindl. | GenBank | Africa | MT500657 | MT533559 | – | – |
| <i>Habenaria decorata</i> Hochst. ex A.Rich. | GenBank | Africa | MT500609 | MT533514 | – | – |
| <i>Habenaria dives</i> Rchb.f. | GenBank | Africa | MT500592 | MT533497 | – | – |
| <i>Habenaria foliosa</i> A.Rich. | GenBank | India | JN114529 | JN004457 | – | – |
| <i>Habenaria heyneana</i> Lindl. | GenBank | India | JN114547 | JN004469 | – | – |
| <i>Habenaria laevigata</i> Lindl. | GenBank | Africa | MT500591 | MT533496 | – | – |
| <i>Habenaria lithophila</i> Schltr. | GenBank | Africa | MT500589 | MT533493 | – | – |
| <i>Habenaria petitiana</i> (A.Rich.) T.Durand & Schinz | GenBank | Africa | MT500613 | MT533518 | – | – |
| <i>Habenaria praestans</i> Rendle | GenBank | Africa | MT500605 | MT533510 | – | – |
| <i>Habenaria stenopetala</i> Lindl. | GenBank | China | MF944324 | MF945467 | – | – |
| <i>Habenaria stenopetala</i> Lindl. | GenBank | China | MF944309 | MF945513 | – | – |
| <i>Pecteilis radiata</i> (Thunb.) Raf. (<i>Habenaria radiata</i>) | GenBank | Korea | KT338720 | NC_035834 | NC_035834 | MN221407 |
| <i>Platycoryne buchaniana</i> (Kraenzl.) Rolfe | GenBank | Africa | MT500587 | MT533491 | – | – |

Supplementary file 1 (continued). GenBank accessions and voucher data of the terminals used in the phylogenetic analyses. Acronyms for the Brazilian states: BA=Bahia; GO=Goiás; MG=Minas Gerais; MT=Mato Grosso; PB=Paraíba; PR=Paraná; RJ=Rio de Janeiro; RS=Rio Grande do Sul; SC=Santa Catarina. DF=Distrito Federal. USA: United States of America; FL = Florida. New sequences generated for this work are indicated by an asterisk.

| Taxon | Voucher | Origin | ITS | <i>matK-trnK</i> | <i>rps16-trnK</i> | ETS |
|---|---------------------------|---------------|------------|-------------------------|--------------------------|------------|
| <i>Platycoryne crocea</i> Rolfe | GenBank | Africa | MT500614 | MT533519 | – | – |
| <i>Roeperocharis</i> <i>bennettiana</i> Rechb.f. | GenBank | Africa | MT500659 | MT533562 | – | – |
| <i>Stenoglottis longifolia</i> Hook.f. | Batista 3279 (BHCB) | Africa | MT582234 | MT623335 | MT623287 | MT623253 |

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Artikel/Article: [Checklist and molecular phylogenetics reveal three taxonomic novelties in *Habenaria* \(Orchidaceae, Orchidoideae\) from Chapada dos Veadeiros, Goiás, Brazil 51-86](#)