

# Home at last III: Transferring *Uechtritzia* and Asian *Gerbera* species into *Oreoseris* (Compositae, Mutisieae)

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## Abstract

Recently the Asian *Gerbera* species were shown to form a clade that was not the sister group of the African *Gerbera*. In this study, the position of the Asian *Gerbera* species was further assessed based on morphology and molecular phylogenetic analyses that included six Asian *Gerbera* and 26 other species from the *Gerbera*-complex. Morphological results showed that the six Asian *Gerbera* species, which were sampled, bear leaves with the adaxial epidermal surface lacking stomates, possess bracteate scapes and lack inner ray florets. These characters suggest that the Asian *Gerbera* species are most closely related to the species of *Uechtritzia*, which also share similar pollen grain size and shape with the Asian *Gerbera*, rather than to the African *Gerbera*. Furthermore, the phylogenetic results based on two nuclear (ITS and ETS) and three chloroplast (*trnL-trnF*, *trnL-rpl32* and *trnC-petN*) sequences strongly support the Asian *Gerbera* and *Uechtritzia* forming a clade, with the latter nested within the Asian *Gerbera* species. Both morphological and molecular phylogenetic data thus confirmed the taxonomic identity of the Asian *Gerbera* and *Uechtritzia*. The authors herein formally treat the nine species of the Asian *Gerbera* and the three species of *Uechtritzia* as members of the genus *Oreoseris*, which is the earliest generic name of this lineage and has the nomenclatural priority.

## Keywords

Compositae, *Gerbera*-complex, *Oreoseris*, *Uechtritzia*, SEM, stomata, pollen, South America, Africa, Asia

## Introduction

The *Gerbera*-complex (Compositae: Mutisieae) contains eight genera: *Gerbera* L., *Leibnitzia* Cass., *Uechtritzia* Freyn, *Amblyspurma* Benth., *Chaptalia* Vent., *Trichocline* Cass., *Perdicium* L. and *Lulia* Zardini. *Gerbera* currently contains about 31 species, which belong to six sections: the five African sections: sect. *Gerbera* (8 species), sect. *Parva* H.V.Hansen (1 species), sect. *Lasiopus* (Cass.) Sch.Bip. (6 species), sect. *Pseudoseris* (Baill.) C.Jeffrey (8 species, distributed in Madagascar) and sect. *Piloselloides* Less. (2 species, one of which is widespread in Asia and Africa) and the Asian sect. *Isanthus* (Less.) Jeffrey (6 species; Hansen 1985a, 1985b, 1988, Johnson et al. 2014, Funk et al. 2016). One South American species *G. hieracioides* (Kunth) Zardini was not included in any of the above-mentioned sections of *Gerbera* (Zardini 1974) and the authors have recently transferred it to *Chaptalia* based on both morphological and molecular data (Xu et al. 2018).

The Asian *Gerbera* section *Isanthus* is characterised mainly by campanulate involucres, naked receptacles and rostrate achenes (Hansen 1988). A recent molecular phylogenetic analysis showed that the Asian *Gerbera* species did not form a clade with the African species (Pasini et al. 2016): the Asian *Gerbera* + *Uechtritzia* formed a clade and the African *Gerbera* and *Amblyspurma* constituted another clade. Some earlier workers also suggested treating the Asian section as an entity separate from the African *Gerbera* (Candolle 1838, Jeffrey 1967). Hansen (1990), however, argued that, while the Asian *Gerbera* sect. *Isanthus* differed somewhat from the African *Gerbera*, it shared four apomorphies as well as 11 plesiomorphies with *Uechtritzia* and the three entities could not be discerned from one another.

Species of *Uechtritzia* have hemispherical involucres, fimbriate receptacles and slightly rostrate achenes (Hansen 1988). This genus contains three species, namely *U. armena* Freyn endemic to Turkey (Doganet al. 2016) and Armenia, *U. kokanica* (Regel et Schmalh.) Pobel. from Central Asia (Kazakhstan, Uzbekistan, Tajikistan, Kyrgyzstan, Turkmenistan to Afghanistan) and *U. lacei* (G.Watt) C.Jeffrey of the Himalayan region (Hansen 1988).

Pasini et al. (2016) included one *Uechtritzia* species, *U. kokanica* and showed that the species was nested within the two sampled Asian *Gerbera* species based on nuclear (ITS) and chloroplast (*trnL*–*trnF*) sequence data. This result indicated the possibility that the Asian *Gerbera* may belong to the genus *Uechtritzia*. However, the phylogenetic position and the taxonomic identity of the Asian *Gerbera* need to be tested with an expanded taxon sampling by adding more Asian and African species of *Gerbera* and *Uechtritzia* before any taxonomic decisions can be made.

In this study, the phylogenetic position of the Asian *Gerbera* was tested by expanding the taxon sampling of the Asian and African *Gerbera* and the *Uechtritzia* species and using both molecular (two nuclear markers: ITS and ETS and three chloroplast markers: *trnL*–*trnF*, *trnL*–*rpl32* and *trnC*–*petN*) and morphological data (leaf adaxial surface, pollen, scape and floral morphology).

## Materials and methods

A total of 32 species from eight genera of the *Gerbera*-complex and *Adenocaulon chilense* (outgroup) were sampled for this study (Tables 1, 2). The morphological data were taken from specimens at the United States National Herbarium (US) and included characters of the leaf epidermis, pollen, flowers and scapes.

**Adaxial leaf epidermal and pollen morphology.** A small area of the leaf lamina (about 0.5–1.0 cm<sup>2</sup>) was placed with the adaxial side exposed, on carbon tape over stubs for the scanning electron microscopy (SEM). For the pollen analysis, samples were dehydrated and were then placed on aluminium stubs using double-sided adhesive tape following Wen and Nowicke (1999). The stubs bearing the leaf sample and pollen were treated with gold-palladium to 16.6 µm thickness and were examined under a Philips XL-30 scanning electron microscope at the SEM Lab of the National Museum of Natural History (NMNH), Smithsonian Institution. The 22 samples were subsequently observed and photographed under the SEM using the proprietary software associated with the Philips SEM. Images of at least 15 different areas of the adaxial leaf surface were captured for each sample, as well as 20 pollen grains. The polar and equatorial axes of pollens were measured by ImageJ 1.8.0.

**DNA extraction, amplification and sequencing.** The DNA molecular work was undertaken in the Laboratory of Analytical Biology (LAB) of NMNH. DNA from 16 samples (15 species) was extracted through AutoGen (AutoGen Inc., Holliston, Massachusetts, USA) or the DNeasy Plant Mini Kit (Qiagen, Valencia, California, USA). Leaf tissue samples, along with 1.0 and 2.3 mm diameter beads, were dipped in liquid nitrogen then immediately shaken for 60 seconds at 1800 rpm by Tissuelyser. About 500 µl of the CTAB extraction buffer was added to the tubes, vortexed and incubated overnight (500 rpm at 55 °C). Then 300 µl of the supernatant was transferred to an AutoGen plate. The AutoGen was run according to the manufacturer's default settings.

Five markers including two nuclear ribosomal ITS and ETS and three chloroplast *trnL-trnF*, *trnL-rpl32* and *trnC-petN* intergenic spacers were amplified. The ITS primers were designed by Downie and Katz-Downie (1996) and White et al. (1990), ETS primers by Baldwin and Markos (1998); *trnL-trnF* primers by Taberlet et al. (1991), *trnL-rpl32* spacer primers by Timme et al. (2007) and *trnC-petN* spacer primers by Lee and Wen (2004) (Table 3). The PCR reaction mixture had a total of 25 µl volume, comprising 14.05 µl nuclease free water, 2.5 µl 10x buffer, 2 µl dNTPs, 1.25 µl MgCl<sub>2</sub>, 1 µl of both forward and reverse primers, 0.5 µl BSA, 0.2 µl Taq DNA polymerase and 2.5 µl of template DNA. The PCR reactions were performed in a Veriti PCR Thermal Cycler. The amplification protocols for all markers are summarised in Table 3. The amplified products were purified with ExoSapIT enzyme with activation at 37 °C and deactivation at 95 °C. 4 µl of the purified product and same primers (1 µl, 1 µM) were cycle-sequenced in a mixture containing 0.8 µl Big Dye (Applied Biosystems, Foster City, California, USA) and 2.0 µl 5x Big Dye buffer and 4.2 µl water.

**Table I.** Voucher information and morphological characters of *Gerbera* and related species.

Species	Section	Locality	Voucher information	Adaxial leaf stomata	Bracts on scape	Inner rays	Pollens	
							Polar axis (μm)	P/E ratio
<i>Gerbera viridifolia</i> (DC.) Sch.Bip.	<i>Lasiopus</i>	Kenya	<i>T.H. Trinder-Smith s.n.</i> (US)	+	-	+	44.12	1.21
<i>G. jamesonii</i> Adlam	<i>Lasiopus</i>	Cultivar	<i>T. Derby s.n.</i> (US)	+	-	+	45.77	1.29
<i>G. aurantiaca</i> Sch. Bip.	<i>Lasiopus</i>	South Africa	<i>Bayliss 2505</i> (US)	+	-	+	43.48	1.20
<i>G. ambigua</i> Sch. Bip.	<i>Lasiopus</i>	South Africa	<i>M. Koekemoer 2097</i> (US)	+	-	+	44.98	1.38
<i>G. piloselloides</i> Cass.	<i>Piloselloides</i>	Swaziland	<i>M. Koekemoer 2590</i> (US)	+	-	+	42.09	1.28
<i>G. cordata</i> Less.	<i>Piloselloides</i>	Madagascar	<i>T.B. Croat 29083</i> (MO)	+	-	+	43.19	1.27
<i>G. perrieri</i> Humbert	<i>Pseudoseris</i>	Madagascar	<i>L. Gautier 3110</i> (MO)	+	-	+	44.04	1.29
<i>G. diversifolia</i> Humbert	<i>Pseudoseris</i>	Madagascar	<i>B. Lewis 1201</i> (MO)	+	-	+	45.31	1.20
<i>G. crocea</i> Kuntze	<i>Gerbera</i>	South Africa	<i>M. Koekemoer 2029</i> (US)	+	+	-	53.83	1.39
<i>G. linnaei</i> Cass.	<i>Gerbera</i>	South Africa	<i>E. Werdermann 749</i> (US)	+	+	-	47.01	1.25
<i>G. tomentosa</i> DC.	<i>Gerbera</i>	South Africa	<i>P. Bond 745</i> (US)	+	+	-	50.43	1.26
<i>G. wrightii</i> Harv.	<i>Gerbera</i>	South Africa	<i>P. Goldblatt 5287</i> (US)	+	+	-	N	N
<i>G. serrata</i> Druce	<i>Gerbera</i>	South Africa	<i>M. Koekemoer 2001</i> (PRE)	+	+	-	N	N
<i>G. gossypina</i> Beauverd	<i>Isanthus</i>	India	<i>W.N. Koelz 4828</i> (US)	-	+	-	50.05	1.40
<i>G. maxima</i> Beauverd	<i>Isanthus</i>	India	<i>D.H. Nicolson 2755</i> (US)	-	+	-	50.41	1.26
<i>G. delavayi</i> Franch.	<i>Isanthus</i>	China	<i>X. Xu 1102</i> (KMUST)	-	+	-	51.90	1.27
<i>G. nivea</i> Sch.Bip.	<i>Isanthus</i>	China	<i>J.F. Rock 6430</i> (US)	-	+	-	50.30	1.39
<i>G. raphanifolia</i> Franch.	<i>Isanthus</i>	China	<i>J.F. Rock 10504</i> (US)	-	+	-	51.74	1.28
<i>G. henryi</i> Dunn	<i>Isanthus</i>	China	<i>W.B. Hemsley 1903</i> (US)	-	+	-	51.91	1.33
<i>Uechtritzia armena</i> Freyn	N	Turkey	<i>A. Kaya 1835</i> (EU)	N	+	-	N	N
<i>U. lacei</i> (G.Watt) C.Jeffrey	N	India	<i>W. Koelz 8710</i> (NA)	-	+	-	50.86	1.36
<i>U. kokanica</i> (Regel et Schmalh.) Pobed.	N	Tajikistan	<i>F.L. Zaprijagaev 4682</i> (US)	-	+	-	55.80	1.31
<i>Leibnitzia anandria</i> (L.) Nakai	N	China	<i>I. Thomas 8183</i> (US)	+	+	-	34.45	1.10
<i>L. nepalensis</i> (Kunze) Kitam.	N	China	<i>J. Wen 542</i> (US)	+	+	-	32.16	1.20
<i>L. occimadrensis</i> G.L.Nesom	N	Mexico	<i>H.S. Gentry 7189</i> (US)	+	+	-	37.33	1.16
<i>Amblyspurma scapigera</i> Benth.	N	Australia	<i>A. Morrison s.n.</i> (US)	+	+	-	51.60	1.17
<i>A. spathulata</i> (A.Cunn. ex DC.) D.J.N.Hind	N	Australia	<i>R.A. Davis 8267</i> (US)	+	+	-	55.10	1.23

Notes: + designates those mentioned present; - designates those mentioned absent; N represents data not available.

**Table 2.** Voucher information and GenBank accessions of *Gerbera* and the related species.

Species	Locality	Voucher information	ITS	ETS	trnL-trnF	trnL-rpl32	trnC-petN
<i>Gerbera viridifolia</i> (DC.) Sch.Bip.	South Africa	T.H. Thinder-Smith s.n. (US)	MG6611696*	MG661588*	MG6611639*	MG6611670*	MG6611628*
<i>G. crocea</i> Kunze	South Africa	M. Koekemoer 2029 (US)	MG6611709*	MG6611606*	MG6611645*	MG6611683*	MG6611618*
<i>G. delavayi</i> Franch.	China	X. Xu 1102 (KMUST)	MG6611708*	MG6611605*	MG6611659*	MG6611682*	MG6611619*
<i>G. henryi</i> Dunn	China	X. Xu 1103 (KMUST)	MG6611706*	MG6611602*	MG6611655*	MG6611681*	MG6611621*
<i>G. nivea</i> Sch.Bip.	China	Y.S. Chen 2674 (PE)	MG6611703*	MG6611598*	MG6611648*	MG6611678*	N
<i>G. aurantiaca</i> Sch.Bip.	South Africa	Bailliss 2505 (US)	MG6611711*	MG6611610*	MG6611637*	MG6611687*	MG6611615*
<i>G. ambigua</i> Sch.Bip.	South Africa	M. Koekemoer 2097 (US)	MG6611712*	MG6611611*	MG6611636*	MG6611688*	N
<i>G. jamesonii</i> Adlam	Cultivar	T. Derby s.n. (US)	MG6611704*	MG6611599*	MG6611638*	MG6611679*	MG6611624*
<i>G. cordata</i> Less.	South Africa	J. Wen 10067 (US)	N	MG6611608*	MG6611661*	MG6611685*	MG6611617*
<i>G. piloselloides</i> Cass.	Swaziland	M. Koekemoer 1972 (US)	MG6611701*	MG6611592*	MG6611650*	MG6611675*	MG6611625*
<i>G. wrightii</i> Harv.	South Africa	P. Goldblatt 5287 (US)	MG6611695*	MG6611587*	MG6611642*	N	N
<i>G. serrata</i> Druce	South Africa	M. Koekemoer 2001 (PRE)	MG6611697*	MG6611590*	MG6611656*	MG6611671*	N
<i>G. diversifolia</i> Humbert	Madagascar	B. Lewis 1201 (MO)	N	MG6611604*	MG6611640*	N	N
<i>G. raphanifolia</i> Franch.	China	Rock [F] 10504 (US)	N	N	MG6611658*	N	MG6611626*
<i>G. gosypina</i> Beauverd	India	W.N. Koch 4824 (US)	MG6611707*	MG6611603*	MG6611646*	N	MG6611620*
<i>G. maxima</i> Beauverd	India	F. Kingdom-Ward 18199 (NY)	KX349402	N	KX349371	N	N
<i>Uechtritzia laevis</i> (G. Watt) C. Jeffrey	India	W. Koch 8710 (NA)	N	N	MG6611644*	N	N
<i>U. kohanicola</i> (Regel & Schmalz) Pobed.	Tajikistan	F.L. Zaprijaqev 4382 (US)	N	MG6611580*	MG6611643*	N	MG6611635*
<i>U. kohanicola</i> (Regel & Schmalz) Pobed.	Tajikistan	Zaprijaqev s.n. (NY)	KX349400	N	KX349401	N	N
<i>Amblyserpasma scapigera</i> Benth.	Australia	A. Morrison s.n. (US)	MG6611713*	MG6611612*	N	MG6611689*	N
<i>A. spathulata</i> (A. Cunn. ex DC.) D.J.N.Hind	Australia	Craven 16197 (CANB)	JX564167	N	KF3989620	N	N
<i>Adenocaulon chilense</i> Less.	Chile	G.L. Sobel 2558 (US)	MG6611714*	N	N	MG6611690*	N
<i>Chapmania pringlei</i> Greene	Mexico	G. Nesom 4405 (US)	GU126773	N	N	N	N
<i>C. brenesioides</i> (Kunth) X.-D.Xu & W.Zheng	Ecuador	P.M. Peterson 9287 (US)	MG6611705*	MG6611601*	MG6611657*	MG6611680*	N
<i>Trichocline repans</i> (Wedd.) Hieron	Argentina	E. Pasini & F. Tørshøien 1025 (ICN)	KX349398	N	KX349399	KX349410	N
<i>Leibnitzia anandria</i> (L.) Nakai	China	I. Thomas 8183 (US)	MG6611694*	MG6611585*	MG6611662*	MG6611668*	MG6611629*
<i>L. anandria</i> (L.) Nakai	Japan	Z.Y. Wu 8985 (KUN)	MG6611692*	MG6611584*	MG6611664*	MG6611667*	MG6611631*
<i>L. occimadensis</i> G.L.Nesom	Mexico	H.S. Gentry 7189 (US)	GU126784	MG6611583*	N	MG6611666*	MG6611632*
<i>L. nepalensis</i> (Kunze) Kitam.	China	J. Wen 542 (US)	KX349373	MG6611582*	KU126734	MG6611659	MG6611653*
<i>L. lytnia</i> (Sch.Bip.) G.L.Nesom	USA	G. Nesom 24778 (ARIZ)	GU126779	N	N	GU126757	N

Notes: \* designates the new sequences from this study; N represents data not available.

**Table 3.** Amplification protocols for all markers.

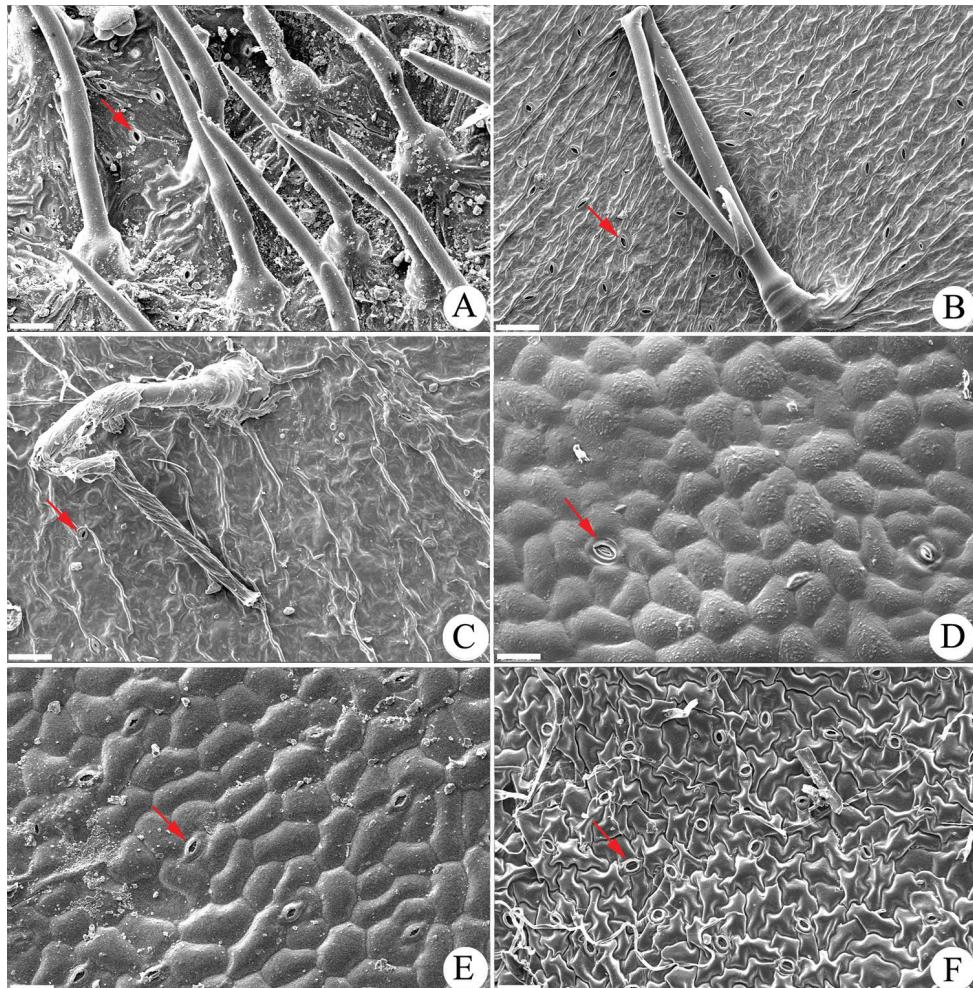
Marker	Primers and sequences 5'-3'	PCR protocol: initial pre-heating; DNA denaturation; primer annealing; DNA extension; final extension
ITS	ITS5A: GGAAGGAGAACGTCATAACAAGG ITS4: TCCTCCGCTTATTGATATGC	95 °C 1 min; 54 °C 1 min; 72 °C 1 min; 72 °C 10 min; 35 cycles
ETS	18s-ETS: ACTTACACATGCATGGCTTAATCT ETS-Hel-1: GCTCTTGCTTGCACAACT	94 °C 0:30 min; 60 °C 0:40 min; 72 °C 1:20 min; 72 °C 5 min; 30 cycles
<i>trnL</i> - <i>trnF</i>	<i>trnL</i> -Fc: CGAAATCGGTAGACGCTACG <i>trnL</i> -Ff: ATTTGAACTGGTACACGAG	94 °C 1 min; 53 °C 1 min; 72 °C 2 min; 72 °C 10 min; 35 cycles
<i>trnL</i> - <i>rpl32</i>	<i>trnL</i> : TACCGATTCACCATAGCGG <i>rpl32</i> : AGGAAAGGATATTGGCGGG	95 °C 3 min; 51 °C 40 s; 72 °C 1:20 min; 72 °C 5 min; 35 cycles
<i>trnC</i> - <i>petN</i>	<i>trnC</i> : CCAGTTCAAATCTGGGTGTC <i>petN</i> : GGATATAGTAAGTCTTGCTTGGG	95 °C 3 min; 54 °C 45 s; 72 °C 1:20 min; 72 °C 8 min; 35 cycles

The cycle sequencing programme was 30 cycles of 95 °C for 30 s, 50 °C for 30 s and 60 °C for 4 min. The resultant product was sephadex-filtered and sequenced through an ABI 3730 automated sequencer (Applied Biosystems, Foster City, USA). Sequences were aligned by using MAFFT (Katoh and Standley 2013) in Geneious 10.0.9. (Biomatters Ltd., Auckland, New Zealand) and checked manually. A total of 90 newly generated sequences from the 23 samples were deposited in GenBank (Table 2).

A total of 16 sequences of eight species were retrieved from NCBI for the related taxa within the tribe Mutisieae (Table 2). Phylogenetic relationships were inferred based on the concatenated ITS+ETS+*trnL*-*rpl32*+*trnL*-*trnF*+*trnC*-*petN* data with MrBayes v. 3.2.2 (Ronquist et al. 2012) by using the substitution model of GTR based on the best-fitting model determined by jModelTest 2.1.6 (Posada 2008), the chain length of 10,000,000, rate variation of gamma, gamma categories of 4, heated chains of 4, heated chain temp of 0.2, subsampling freq. of 200 and burn-in length of 100,000. Tracer v. 1.5 (Rambaut and Drummond 2009) was used to confirm that the effective sample size (ESS) for all relevant parameters was > 200. After discarding the trees as burn-in, a 50 % majority-rule consensus tree and posterior probabilities (PP) for node support were calculated using the remaining trees.

## Results

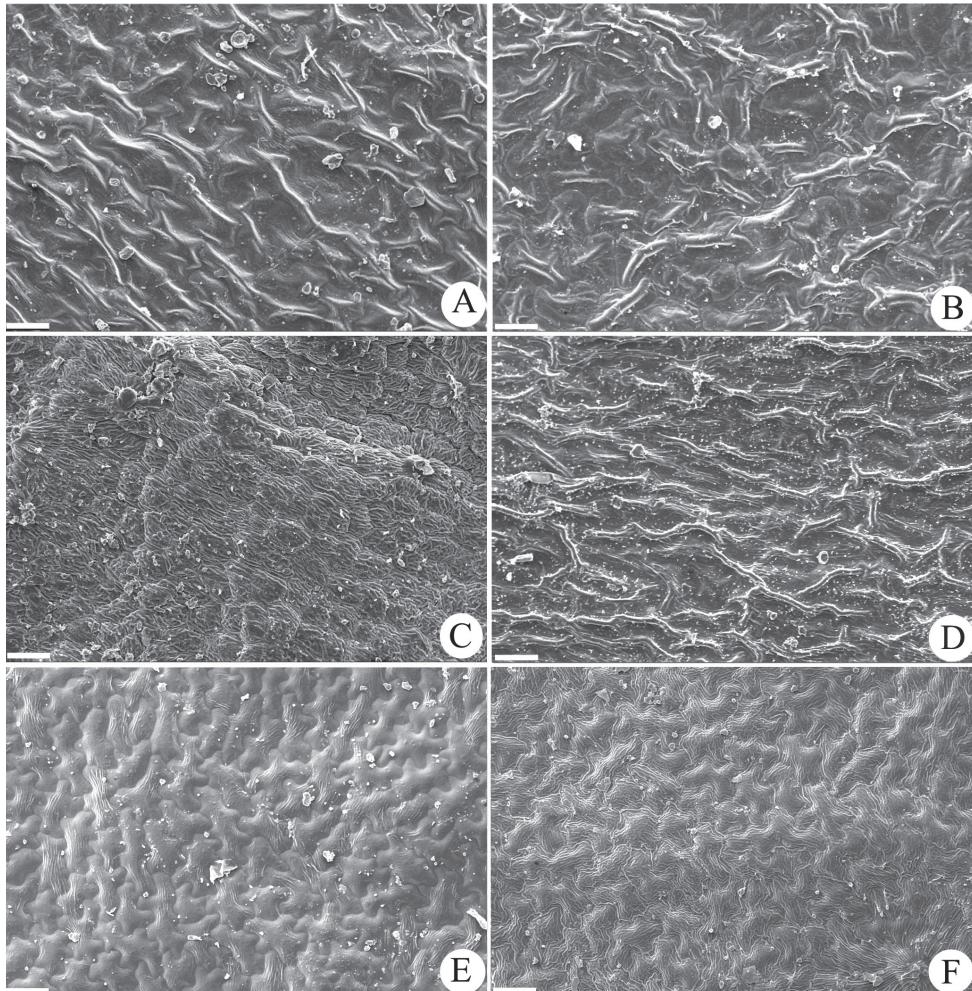
**Adaxial leaf epidermal morphology.** The results of the SEM work (Table 1) showed that the six tested Asian *Gerbera* species have no stomates on the adaxial leaf surface (Figure 2A, B, C, D). This adaxial leaf morphological trait differs from that of the African *Gerbera* species: (1) Three East African *Gerbera* sections sampled [sect. *Lasiopus* (4 species), sect. *Piloselloides* (2 species) and sect. *Pseudoseris* (2 species)] have stomates and stiff, straight, upright trichomes on the adaxial surface. Figure 1 has representative images for each of the above sections: *G. ambigua* (Fig. 1A), *G. piloselloides* (Fig. 1B) and *G. perrieri* (Fig. 1C), respectively. (2) Members of the South African sect. *Gerbera* have stomates.



**Figure 1.** Adaxial leaf epidermal surface morphology of African *Gerbera* and Asian *Leibnitzia*. **A** *G. ambigua* (sect. *Lasiopus*) **B** *G. piloselloides* (sect. *Piloselloides*) **C** *G. perrieri* (sect. *Pseudoseris*) **D** *G. serra*ta (sect. *Gerbera*) **E** *G. crocea* (sect. *Gerbera*) **F** *L. nepalensis*. Bar=50 µm.

Five species were examined and the epidermal characters are represented by *G. serra*ta (Fig. 1D) and *G. crocea* (Fig. 1E). Furthermore, the adaxial leaf morphological traits of the Asian *Gerbera* species also deviate from two Asian-American disjunct *Leibnitzia* species, which have stomates on the adaxial leaf epidermal, as represented by *L. nepalensis* (Fig. 1F). Nevertheless, the Asian *Gerbera* samples share similar adaxial leaf epidermal characters of lacking stomates with the two examined *Uechtritzia* species, *U. kokanica* (Fig. 2E) and *U. lacei* (Fig. 2F). Based on the adaxial leaf epidermal morphology, the Asian *Gerbera* is most closely related to *Uechtritzia* rather than to the African *Gerbera*.

**Pollen morphology.** The pollen grains of the examined species of the *Gerbera*-complex are very similar to one another, differing only in the size of the grains as well

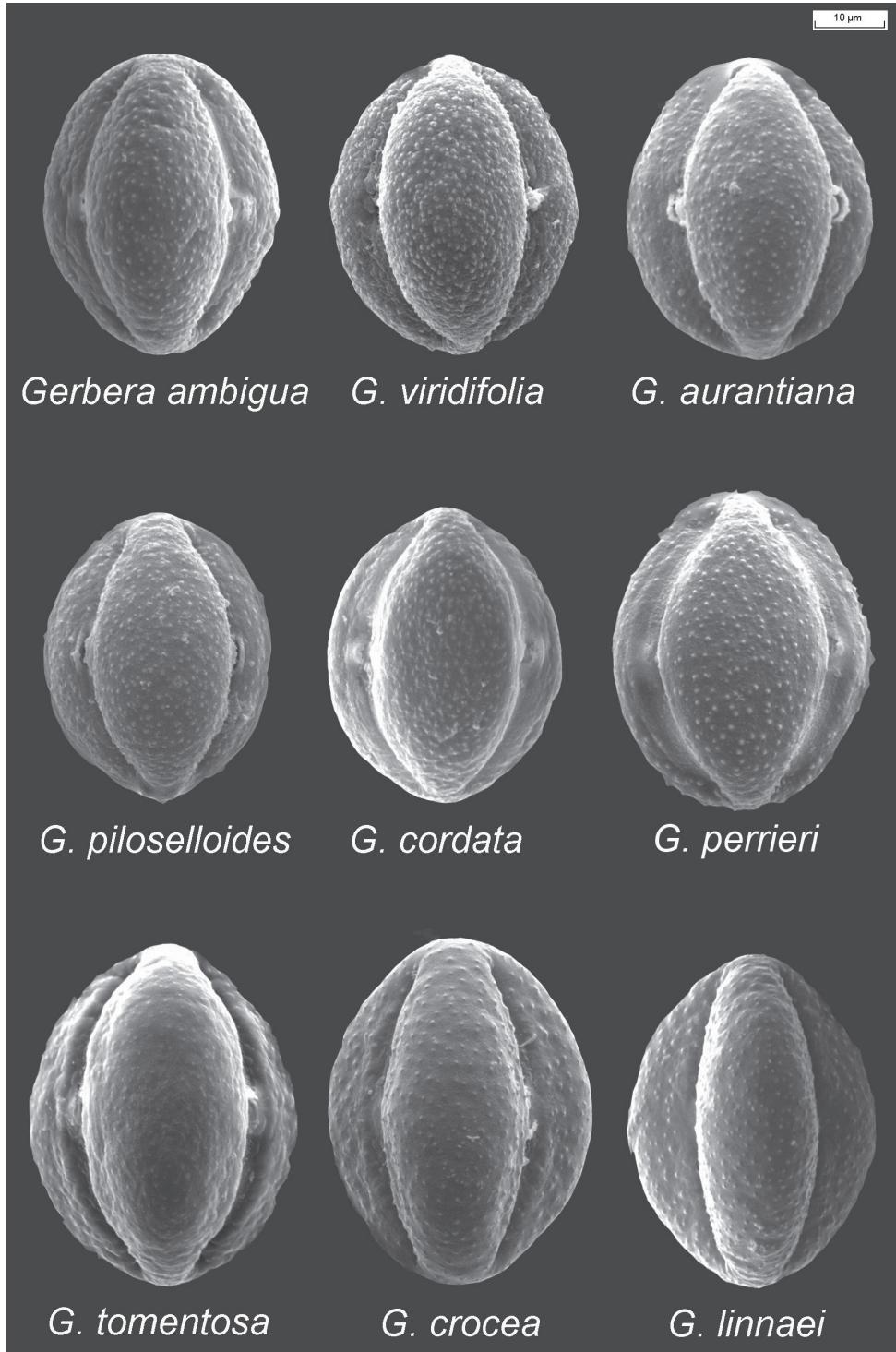


**Figure 2.** Adaxial leaf epidermal surface morphology of Asian *Gerbera* and *Uechtritzia*. **A** *G. maxima* **B** *G. delavayi* **C** *G. gossypina* **D** *G. nivea* **E** *U. kokanica* **F** *U. lacei*. Bar=50 µm.

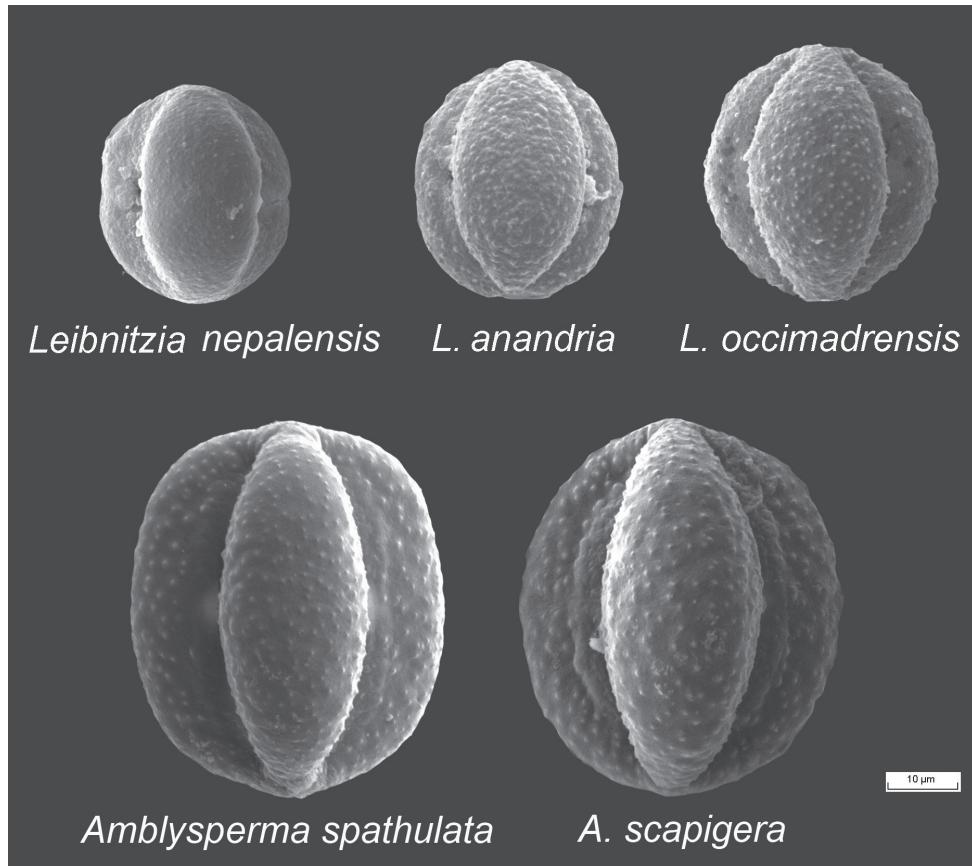
as the granules on the surfaces (Figs 3 and 4). They are tricolporate, have a granule exine and are prolate and subprolate in shape. The ratios of the polar axis and equatorial axis (P/E) are given in Table 1. For *Gerbera* and *Uechtritzia*, the P/E ratios are between 1.2–1.4. The average polar axis of the Asian *Gerbera* and *Uechtritzia* pollen grains is 50.05–55.80 µm. For the African *Gerbera*, however, the average polar axis of pollen grains is 42.09–45.77 µm in sects. *Lasiopus*, *Piloselloides* and *Pseudoseris* and 47.01–53.83 µm in sect. *Gerbera*. The P/E ratio of the pollen grains of the Asian *Gerbera* and *Uechtritzia* (Table 1) differs from that of the East Asian-North American *Leibnitzia* and the Australian *Amblyesperma*, which fall between 1.10–1.20 (Fig. 5). Furthermore, the average polar axis of the Asian *Gerbera* and *Uechtritzia* pollen grains is higher than that of *Leibnitzia* species, which has the range of 32.16–37.33 µm.



**Figure 3.** Pollen morphology of Asian *Gerbera* and *Uechtritzia*.



**Figure 4.** Pollen morphology of African *Gerbera* species.

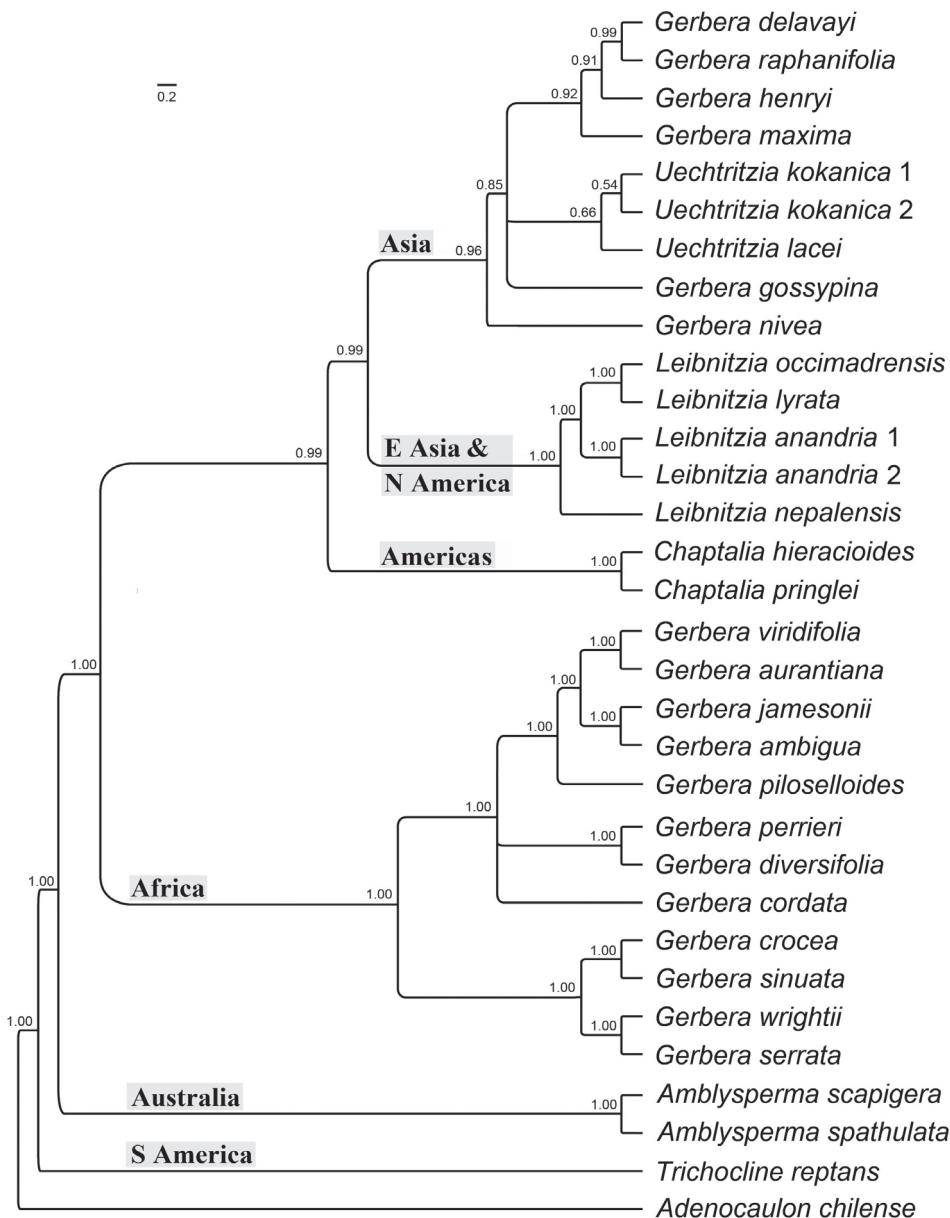


**Figure 5.** Pollen morphology of *Leibnitzia* and *Amblyserma* species.

**Phylogenetic analysis.** The Bayesian analysis of the combined nuclear markers and three plastid genes showed six clades of the sampled species of the *Gerbera*-complex, all showing a strong geographic signal (Fig. 6): (1) the Asian *Gerbera* and the *Uechtritzia* species, (2) the East Asian and North American *Leibnitzia* species, (3) the New World genus *Chaptalia*, (4) the African *Gerbera* species, (5) the Australian genus *Amblyserma* and (6) the South American genus *Trichocline*. The three samples of *Uechtritzia* (two species of *U. kokanica* and *U. lacei*) were clearly nested within the Asian *Gerbera* clade (Fig. 6).

## Discussion

Based on this study, the Asian *Gerbera* and the *Uechtritzia* species share several morphological characters, including bracteate scapes, absence of inner ray florets, no stomates on the adaxial leaf surface and similar pollen size and shape (Table 1). Hansen (1990) also commented that the Asian *Gerbera* (i.e. sect. *Isanthus*) is morphologically



**Figure 6.** Phylogeny of the *Gerbera*-complex. The phylogeny is based on the Bayesian inference of the combined ITS and ETS, *trnL-trnF*, *trnL-rp32* and *trnC-petN* markers. The posterior probabilities are shown next to branches.

similar to *Uechtritzia* and presented a key to distinguish *Gerbera* sect. *Isanthus*, *Uechtritzia* and *Leibnitzia*. The differences between the Asian *Gerbera* sect. *Isanthus* and *Uechtritzia* were minor. *Uechtritzia* species generally have hemispherical heads, alveo-

lar receptacles that are fimbriate-ciliate; margins of involucral bracts (at least the upper part) often with reddish hairs; achenes that are slightly or indistinctly tapering with hairs that are long-villose, ca. 1 mm long (Katinas 2004) and sericeous (Hansen 1988). *Gerbera* sect. *Isanthus* has heads campanulate; a receptacle that is alveolate and naked; the margins of involucral bracts are without reddish hairs; the achenes are tapering and pilose glabrous, with hairs that are shorter, tapered and not sericeous (Hansen 1988). The heads of *Uechtritzia* were reported as hemispherical, in contrast to the heads of the Asian *Gerbera* sect. *Isanthus* which are campanulate. However, the species of *U. armena* (the type species of the genus) from Turkey showed the heads as campanulate in the fresh plants (Dogan et al. 2016), which is the same as the Asian *Gerbera* sect. *Isanthus* (e.g. *G. delavayi*; Zheng et al. 2017).

Some previous workers argued that the species of Asian *Gerbera* (sect. *Isanthus*) should be treated as an entity, separate from African *Gerbera* (Candolle 1838, Jeffrey 1967, Pasini et al. 2016). The results presented here show that the Asian *Gerbera* sect. *Isanthus* differs from the African *Gerbera* sect. *Lasiopus*, sect. *Piloselloides* and sect. *Pseudoseris* in the ebracteate scapes, presence of inner ray florets, stomates on the adaxial leaf surface and smaller pollen size of the African *Gerbera* compared with the Asian *Gerbera*. Although the Asian *Gerbera* sect. *Isanthus* shares the traits of bracteates scapes, absence of inner ray florets and similar pollen size with *Gerbera* sect. *Gerbera*, the Asian species have no stomates on the adaxial leaf surface. Hansen (1990) stated that the Asian *Gerbera* sect. *Isanthus* shows style-arms laterally dilated and truncate achenes; in contrast, the African *Gerbera* sect. *Gerbera* has the style-arm slender and achenes tapering or beaked. Additionally, most species of the African *Gerbera* sect. *Gerbera* grow in open areas, have leathery leaves and flower only in the spring and summer (Manning et al. 2016), whereas the Asian species of *Gerbera* sect. *Isanthus* often grow in forest habitats, have herbaceous leaves and flower in the winter (Gao et al. 2011).

The two *Uechtritzia* species sampled in the molecular phylogeny (Fig. 6) were nested within the Asian *Gerbera* species based on two nuclear markers (ITS and ETS) and three chloroplast markers (*trnL-trnF*, *trnL-rpl32* and *trnC-perN*). This result, based on the authors' expanded taxon and character sampling, is consistent with the findings of Pasini et al. (2016). This study included two of the three species of *Uechtritzia* (Hansen, 1988) and six of the nine Asian *Gerbera* sect. *Isanthus* taxa (Gao et al. 2011). The phylogenetic analysis clearly supports the species of *Uechtritzia* as nested within the Asian *Gerbera* and this clade is the sister group of *Leibnitzia* with strong support (PP=0.99) (Fig. 6).

*Leibnitzia* is a genus containing about six species with a disjunct distribution: four species in Asia (Gao et al. 2011) and two species in Mexico (Baird et al. 2010). It shows the same characters of bracteate scapes and no inner ray florets as the Asian *Gerbera* + *Uechtritzia*. It differs from the latter by the presence of stomates on the adaxial leaf surface and smaller pollen size (polar axis of 32.16–37.33 µm) compared with Asian *Gerbera* + *Uechtritzia* (polar axis of 50.05–55.80 µm). Furthermore, *Leibnitzia* has two generations of heads (a vernal generation with chasmogamous capitula and an aestival generation with cleistogamous capitula), subseriate involucral bracts, slender style-arm, anthers of the ray flowers reduced to threads or wanting

and achenes that are tapering or beaked. The Asian *Gerbera* + *Uechtritzia*, on the other hand, have one generation of heads, imbricate involucral bracts, laterally dilated style-arms, a fully developed apex (and base) on the anthers in the ray flowers and truncate achenes (Hansen, 1990).

Based on the molecular phylogenetic results, the Asian *Gerbera* species are closest to *Uechtritzia*, with the latter nested within the Asian *Gerbera* species. *Leibnitzia* shows significant morphological differences to the Asian *Gerbera* + *Uechtritzia*. The taxonomic identity of *Uechtritzia* and the Asian *Gerbera* is strongly supported by the morphology of inflorescences, scapes, capitula, pollen and the lack of stomates on the adaxial leaf surface. Therefore, the authors herein include the nine Asian *Gerbera* species and the three *Uechtritzia* species in *Oreoseris* DC. which is the earliest available name for the expanded Eurasian genus.

## Taxonomic synopsis with nomenclatural changes

In trying to determine the correct genus name for the Eurasian clade, it is necessary to investigate three relevant generic names. *Gerbera* L. was described in 1758; *Arnica gerbera* L. is the basionym of the African species *G. linnaei* Cass., the conserved type of *Gerbera* L. (lectotype designated by Hansen 1985a). *Gerbera* was named after Traugott Gerber, a German naturalist who died in 1743. *Oreoseris* DC. was described in 1838 and its type species is *O. nivea* DC. which was designated by Hansen in 1988. While de Candolle (1838) did not say why he named the genus, *Oreo* is from the Greek *oreos* for mountain and, in his description, de Candolle says that the genus is a “... perennial herb from the mountains of eastern India (translated).” *Uechtritzia* Freyn was described in 1892; the type species is *U. armena* Freyn (lectotype designated by Pobedimova, 1963). The genus was named in honour of Rudolf Karl Friedrich von Uechtritz (1838–1886), a botanist from Wroclaw, Poland (ex-Breslau) (Freyn 1892).

When *Oreoseris nivea* DC. was absorbed into *Gerbera*, the priority was given to *Gerbera* because the latter was the older generic name and, as long as this species stayed in *Gerbera*, the name *Oreoseris* was not available. *Uechtritzia* was described later in 1892; and, as long as *O. nivea* remained in *Gerbera*, then *Oreoseris* continued to be unavailable.

However, as soon as *Gerbera nivea* from Asia was removed from *Gerbera* and a separate genus was formed from the Asian species of *Gerbera* + *Uechtritzia*, then the name *Oreoseris* became available and it is the oldest available name. Hence, these species have been transferred into *Oreoseris*.

### *Oreoseris* DC., Prodr. 7(1): 17. 1838.

*Onoseris* Willd. sect. *Isanthus* Less., Linnaea 5: 338. 1830. *Onoseris* Willd. subgen. *Isanthus* (Less.) Less., Syn. Comp.: 119. 1832. *Gerbera* L. sect. *Isanthus* (Less.) C.Jeffrey, Kew Bull. 21: 213. 1967.

*Gerbera* L. sect. *Oreoseris* (DC.) Sch.Bip., Flora 27: 780. 1844.

*Uechtritzia* Freyn, Oesterr. Bot. Z. 42(7): 240. 1892. *Gerbera* sect. *Uechtritzia* (Freyn) Beauverd, Bull. Soc. Bot. Genève Ser. 2, 2: 43. 1910.

**Type species.** *Oreoseris nivea* DC., designated by Hansen (1988).

*Oreoseris* has the following 12 species from Eurasia.

**1. *Oreoseris armena* (Freyn et Sint.) V.A.Funk & J.Wen, comb. nov.**

urn:lsid:ipni.org:names:77176439-1

*Uechtritzia armena* Freyn et Sint., Oesterr. Bot. Z. 42(7): 241. 1892. *Gerbera armena*

Beauverd, Bull. Soc. Bot. Genève, ser. 2, 2: 43. 1910.

**Distribution.** Armenia and Turkey.

**2. *Oreoseris delavayi* (Franch.) X.D.Xu & W.Zheng, comb. nov.**

urn:lsid:ipni.org:names:77176440-1

*Gerbera delavayi* Franch., J. Bot. (Morot). 2: 68. 1888.

**Distribution.** China (Guizhou, Sichuan, Yunnan) and N Vietnam.

**3. *Oreoseris gossypina* (Royle) X.D.Xu & V.A.Funk, comb. nov.**

urn:lsid:ipni.org:names:77176441-1

*Chaptalia gossypina* Royle, Ill Bot. Himal. 251. T. 59. F. 2. 1835. *Gerbera gossypina*

(Royle) Beauverd, Bull. Soc. Bot. Genève Ser. 2, 2: 40. 1910.

*Oreoseris lanuginosa* DC., Prodr. 7(1): 17. 1838. *Gerbera lanuginosa* (DC.) Sch.Bip., Flora 27: 780. 1844.

**Distribution.** Karakoram, N and C Himalaya.

**4. *Oreoseris henryi* (Dunn) W.Zheng & J.Wen, comb. nov.**

urn:lsid:ipni.org:names:77176442-1

*Gerbera henryi* Dunn, J. Linn. Soc., Bot. 35: 511. 1903. *Gerbera delavayi* var. *henryi*

(Dunn) C.Y.Wu et H.Peng, Acta Bot. Yunnan. 24: 143. 2002.

**Distribution.** China (Yunnan).

**5. *Oreoseris kakanica* (Regel et Schmalh.) J.Wen & W.Zheng, comb. nov.**  
urn:lsid:ipni.org:names:77176443-1

*Gerbera kakanica* Regel et Schmalh., Descr. Pl. Nov. Rar. Fedtsch. 53. 1882 (published as Izv. Imp. Obsc. Ljubit. Estesv. Moskovsk. Univ. 34(2): 53. 1882). *Uechtritzia kakanica* (Regel et Schmalh.) Pobed., Fl. URSS 28: 597. 1963.

**Distribution.** Pamir-Altai and Tian-Shan regions of C Asia, south to Afghanistan and Kashmir.

**6. *Oreoseris lacei* (G.Watt) V.A.Funk & W.Zheng, comb. nov.**  
urn:lsid:ipni.org:names:77176444-1

*Gerbera lacei* G.Watt Bull. Misc. Inform. Kew 1911(6): 272. 1911. *Uechtritzia lacei* (G.Watt) C.Jeffrey, Kew Bull. 21(2): 213. 1967.

**Distribution.** N India (Himachal Pradesh), S Jammu and Kashmir (Nachar, Baspa, E and NE of Simla, Chamba and Kisthwar).

**7. *Oreoseris latiligulata* (Y.C.Tseng) W.Zheng & J.Wen, comb. nov.**  
urn:lsid:ipni.org:names:77176445-1

*Gerbera latiligulata* Y.C.Tseng, Acta Bot. Austro-Sin. 3: 11. 1986.

**Distribution.** China (in Qiaojia county of Yunnan).

**8. *Oreoseris maxima* (D.Don) X.D.Xu & W.Zheng, comb. nov.**  
urn:lsid:ipni.org:names:77176446-1

*Chaptalia maxima* D.Don, Prodr. Fl. Nepal. 166. 1825. *Gerbera maxima* (D.Don) Beauverd, Bull. Soc. Bot. Genève Ser. 2, 2: 44. 1910.

**Distribution.** China (Xizang), Bhutan, India, Nepal, Pakistan and Thailand.

**9. *Oreoseris nivea* DC., Prodr. 7: 18. 1838.**

*Gerbera nivea* (DC.) Sch.Bip., Flora 27: 780. 1844.

**Distribution.** China (W Sichuan, S Xizang, NW Yunnan), Bhutan, India and Nepal.

**10. *Oreoseris raphanifolia* (Franch.) V.A.Funk & J.Wen, comb. nov.**  
urn:lsid:ipni.org:names:77176447-1

*Gerbera raphanifolia* Franch., J. Bot. (Morot). 2: 67. 1888.

**Distribution.** China (NW Yunnan).

**11. *Oreoseris rupicola* (T.G.Gao & D.J.N.Hind) X.D.Xu & V.A.Funk, comb. nov.**  
urn:lsid:ipni.org:names:77176448-1

*Gerbera rupicola* T.G.Gao et D.J.N.Hind, Fl. China 20–21: 14. 2011.

*Gerbera macrocephala* Y.C.Tseng, Acta Bot. Austro Sin. 3: 12. 1986, not *Gerbera macrocephala* Less., Linnaea 5: 295. 1830.

**Distribution.** China (NW Yunnan).

**12. *Oreoseris tanantii* (Franch.) W.Zheng & X.D.Xu, comb. nov.**  
urn:lsid:ipni.org:names:77176449-1

*Gerbera tanantii* Franch., J. Bot. (Morot). 7: 155. 1893.

**Distribution.** China (Yunnan).

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## References

- Baird KE, Funk VA, Wen J, Weeks A (2010) Molecular phylogenetic analysis of *Leibnitzia* Cass. (Asteraceae: Mutisieae: *Gerbera*-complex): An Asian-North American disjunct genus. Journal of Systematics and Evolution 48: 161–174. <https://doi.org/10.1111/j.1759-6831.2010.00077.x>
- Baldwin BG, Markos S (1998) Phylogenetic utility of the external transcribed spacer (ETS) of 18S–26S rDNA: Congruence of ETS and ITS trees of *Calycadenia* (Compositae). Molecular Phylogenetics and Evolution 10: 449–463. <https://doi.org/10.1006/mpev.1998.0545>
- Candolle AP de (1838) *Prodromus systematis naturalis regni vegetabilis*, vol. 7(1). Treuttel & Würtz, Paris. <https://doi.org/10.5962/bhl.title.286>
- Dogan YN, Kandemir A, Osma E (2016) Genetic diversity and variability among populations and ecological characteristics of the *Uechtritzia armena* Freyn (Asteraceae) endemic to Turkey. Research & Reviews: Research Journal of Biology 4: 20–27.
- Downie SR, Katz-Downie DS (1996) A molecular phylogeny of Apiaceae subfamily Apioideae: Evidence from nuclear ribosomal DNA internal transcribed spacer sequences. American Journal of Botany 83: 234–251. <https://doi.org/10.1002/j.1537-2197.1996.tb12701.x>
- Freyn J (1892) *Plantae novae Orientales*. Österreichische Botanische Zeitschrift 42: 235–242. <https://doi.org/10.1007/BF01791047>
- Funk VA, Pasini E, Bonifacino JM, Katinas L (2016) Home at last: The enigmatic genera *Eriachaenium* and *Adenocaulon* (Compositae, Mutisioideae, Mutisieae, Adenocaulinae). PhytoKeys 60: 1–19. <https://doi.org/10.3897/phytokeys.60.6795>
- Gao TG, Chen YS, Hind DJN, Freire SE (2011) Mutisieae. In: Wu ZY, Raven PH, Hong DY (Eds) Flora of China, vol 20–21 (Asteraceae). Science Press, Beijing & Missouri Botanical Garden Press, St. Louis.
- Hansen HV (1985a) A taxonomic revision of the genus *Gerbera* (Compositae, Mutisieae) sections *Gerbera*, *Parva*, *Piloselloides* (in Africa) and *Lasiopus*. Opera Botanica 78: 5–36.
- Hansen HV (1985b) Notes on *Gerbera* sect. *Pseudoseris* (Compositae, Mutisieae). Nordic Journal of Botany 5: 451–453. <https://doi.org/10.1111/j.1756-1051.1985.tb01675.x>
- Hansen HV (1988) A taxonomic revision of the genera *Gerbera* sect. *Isanthus*, *Leibnitzia* (in Asia), and *Uechtritzia* (Compositae, Mutisieae). Nordic Journal of Botany 8: 61–76. <https://doi.org/10.1111/j.1756-1051.1988.tb01707.x>
- Hansen HV (1990) Phylogenetic studies in the *Gerbera*-complex (Compositae, tribe Mutisieae, subtribe Mutisiinae). Nordic Journal of Botany 9: 469–485. <https://doi.org/10.1111/j.1756-1051.1990.tb00537.x>
- Jeffrey C (1967) Notes on Compositae, III. The Cynareae in east tropical Africa. Kew Bulletin 22: 107–140. [Jstor.org/stable/4107829](https://doi.org/10.1080/002229967086829)
- Johnson IM, Crouch NR, Edwards TJ (2014) *Gerbera sylvicola* (Asteraceae: Mutisieae), a new forest species from KwaZulu-Natal, South Africa. Phytotaxa 186: 229–235. <https://doi.org/10.11646/phytotaxa.186.4.7>
- Katinas L (2004) The *Gerbera*-complex (Asteraceae, Mutisieae): To split or not to split. SIDA, Contributions to Botany, 935–940. [Jstor.org/stable/41968349](https://doi.org/10.11646/sida.22.935)

- Katoh K, Standley DM (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>
- Lee C, Wen J (2004) Phylogeny of *Panax* using chloroplast *trnC–trnD* intergenic region and the utility of *trnC–trnD* in interspecific studies of plants. Molecular Phylogenetics and Evolution 31: 894–903. <https://doi.org/10.1016/j.ympev.2003.10.009>
- Manning JC, Simka B, Boatwright JS, Magee AR (2016) A revised taxonomy of *Gerbera* sect. *Gerbera* (Asteraceae: Mutisieae). South African Journal of Botany 104: 142–57. <https://doi.org/10.1016/j.sajb.2015.10.002>
- Pasini E, Funk VA, de Souza-Chies TT, Miotto STS (2016) New insights into the phylogeny and biogeography of the *Gerbera*-complex (Asteraceae: Mutisieae). Taxon 65: 547–562. <https://doi.org/10.12705/653.7>
- Pobedimova YG (1963) Mutisieae. In: Brobov EG, Cherepanov SK, Komarov VL (Eds) Flora USSR 28: 588–598.
- Posada D (2008) jModelTest, phylogenetic model averaging. Molecular Biology and Evolution 25: 1253–1256. <https://doi.org/10.1093/molbev/msn083>
- Rambaut A, Drummond AJ (2009) Tracer v1.5. Available at: <http://beast.bio.ed.ac.uk/Tracer>.
- Ronquist F, Teslenko M, van der Mark P, Ayres DL, Darling A, Höhna S, Larget B, Liu L, Suchard MA, Huelsenbeck JP (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>
- Taberlet P, Gielly L, Pautou G, Bouvet J (1991) Universal primers for amplification of three non-coding regions of chloroplast DNA. Plant Molecular Biology 17: 1105–1109. <https://doi.org/10.1007/BF00037152>
- Timme R, Kuehl EJ, Boore JL, Jansen RK (2007) A comparative analysis of the *Lactuca* and *Helianthus* (Asteraceae) plastid genomes: Identification of divergent regions and categorization of shared repeats. American Journal of Botany 94: 302–313. <https://doi.org/10.3732/ajb.94.3.302>
- Wen J, Nowicke JW (1999) Pollen ultrastructure of *Panax* (the ginseng genus, Araliaceae), an eastern Asian and eastern North American disjunct genus. American Journal of Botany 86: 1624–1636. <https://doi.org/10.2307/2656799>
- White TJ, Bruns T, Lee S, Taylor J (1990) Amplification and direct sequencing of fungal ribosomal RNA genes for phylogenetics. In: Innis M, Gelfand D, Sninsky J, White T (Eds) PCR protocols: A guide to methods and applications. Academic Press, San Diego, 315–322. <https://doi.org/10.1016/B978-0-12-372180-8.50042-1>
- Xu X, Zheng W, Funk VA, Wen J (2018) Home at Last II: *Gerbera hieracioides* (Kunth) Zar-dini (Mutisieae, Asteraceae) is really a *Chaptalia*. PhytoKeys 95: 93–106. <https://doi.org/10.3897/phytokeys.95.22916>
- Zardini EM (1974) About the presence of the genus *Gerbera* in America. Boletin de la Sociedad Argentina de Botanica 16: 103–108.
- Zheng W, Xu X, Wen J (2017) The ethnic textile use of natural fibers from fireweed (*Gerbera delavayi*) in Southwest China. Economic Botany. <https://doi.org/10.1007/s12231-017-9394-y>