




Standard Paper

Marchantiana pyramus, *M. ramulicola* and *Austroplaca thisbe* (Teloschistaceae, lichenized Ascomycota) – three new twig lichens from southern Patagonia

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Abstract

Three twig-growing lichen species belonging to the family *Teloschistaceae* from southern Patagonia are described as new to science: *Marchantiana pyramus* sp. nov., with minute apothecia, orange due to dominant content of emodin, *M. ramulicola* sp. nov., with minute olive apothecia with dominant parietin and *Austroplaca thisbe* sp. nov., with clear yellow apothecia also with dominant parietin. *Marchantiana subpyracea*, *M. epibrya* and *M. queenslandica* are new combinations for species which, so far, are known only from New Zealand and Australia. *Blastenia circumpolaris* is shown to be very common in Patagonia and *Marchantiana asserigena* is documented for the first time from the Southern Hemisphere, viz. the Falkland Islands. The genus *Marchantiana* is analyzed here using three genes and is shown to be closely related to *Yoshimuria*; although appearing as paraphyletic, a monophyletic origin cannot be ruled out. Morphology, ecology and distribution support a monophyletic treatment and *Marchantiana* is therefore treated as such.

Keywords: biogeography; *Blastenia*; Chile; Falkland Islands; molecular taxonomy; Tierra del Fuego; *Yoshimuria*

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Introduction

Southern Patagonia including Tierra del Fuego is the southernmost region bearing trees and shrubs on Earth, and epiphytic lichens growing here are thus at the southern limit of their distribution. Only Antarctica, with no epiphytes, is further south, and the distance to similar climates and vegetation in New Zealand is more than 7000 km. The *Teloschistaceae* of southern Patagonia have been critically studied during the past decades (Lumbsch *et al.* 2011; Söchting & Sancho 2012; Söchting *et al.* 2014, 2016, 2021). Recently, an extensive catalogue of lichen species known from Navarino Island in Tierra del Fuego was published (Etayo *et al.* 2021), but *Teloschistaceae* on twigs were only superficially treated at the time. It might be expected that the biogeographical history of the southern continents could result in a high degree of endemism among twig lichens, therefore we decided to focus particularly on *Teloschistaceae* on twigs and thin branches of trees and shrubs. Results from the whole Southern Hemisphere show very little overlap between the twig species composition in Tasmania/New Zealand and Patagonia. Therefore this paper focuses on Patagonian species, and the New Zealand species will be dealt with in a forthcoming paper.

Twigs and thin branches often host a diverse and characteristic lichen community (Degelius 1964, 1978; Ott *et al.* 2000), but in the industrialized and agricultural world twig lichens often suffer from

being exposed to low air quality (Larsen *et al.* 2009). In southern Patagonia, including Tierra del Fuego, and in the Falkland Islands, the air is, however, generally clean and lichens thrive on both living and dead twigs. For this study, twigs were intensively collected by the first author during recent decades. The two very common species *Austroplaca thisbe* sp. nov. and *Marchantiana pyramus* sp. nov. are proposed as new to science, together with the less frequent *M. ramulicola* sp. nov. Molecular analyses were carried out to disclose the generic structure of the genus *Marchantiana* and its relationship with the neighbouring genus *Yoshimuria*, but a more comprehensive study of the genus will be published in a forthcoming paper based on species from Australia and New Zealand. The field studies also revealed new distribution data for *Blastenia circumpolaris* Söchting *et al.* and *Marchantiana asserigena* (J. Lahm) Söchting & Arup.

Material and Methods

Material

Twigs with lichens were collected in southern Patagonia, mainly in Tierra del Fuego but also on the Falkland Islands. Phorophytes were mostly *Berberis*, *Chilodictyon*, *Nothofagus*, *Ribes* and the parasitic *Misodendrum* on *Nothofagus*. Cited specimens collected by U. Söchting and M. Z. Søgaard were deposited in C.

Morphology and anatomy

Macroscopic descriptions are based on observations made with an Olympus SZH dissecting microscope with an Olympus OM-D

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camera. Sections were made by hand or with a Reichert-Jung Cryostat 2800 Frigocut E microtome and studied with a Leitz Orthoplan microscope. All measurements were made on material mounted in water. Spores were measured outside the asci and measurements are given as the mean (\bar{x}) and standard deviation (SD) of 'n' measurements, with minimum and maximum measurements in parentheses.

Secondary chemistry

Secondary metabolites were identified and quantified using HPLC according to Söchting (1997); thallus and apothecia were analyzed separately. The relative composition of the secondary compounds was calculated based on absorbance at 270 nm, according to Söchting (1997). All yellow, orange or reddish-pigmented parts are K+ purple.

Molecular analyses

PCR amplification was carried out on DNA extracts or using direct PCR following Arup *et al.* (2015). Amplifications were made of the internal transcribed spacer regions (nrITS) and the large subunit (nrLSU) of the nuclear ribosomal RNA genes, and the small subunit of the mitochondrial ribosomal RNA gene (mtSSU). Primers for amplification were ITS1F (Gardes & Bruns 1993), ITS4 (White *et al.* 1990), AL1R (Döring *et al.* 2000), LR5 or LR6 (Vilgalys & Hester 1990), mrSSU1 (Zoller *et al.* 1999) and mrSSU7 (Zhou & Stanosz 2001). The PCR parameters included an initial hold at 94 °C for 5 min, then denaturation at 94 °C for 1 min, annealing at 50 or 54 °C (mtSSU) or 53–56 °C (nrITS and nrLSU) for 1 min, decreasing 1 °C per cycle for the first six of the 39 cycles (touch-down), and an extension at 72 °C for 3 min.

Sequence alignment

Two different alignments were prepared, one for a combined analysis of the genes nrITS, nrLSU and mtSSU and one alignment of only nrITS sequences. The combined analysis included 72 sequences from most genera of the subfamily *Caloplacoideae* and the ITS alignment included 29 sequences of the genus *Marchantiana*. *Xanthoria parietina* (L.) Th. Fr. from subfamily *Xanthorioideae* was used as outgroup for the first analysis and *Xanthopeltis rupicola* R. Sant. for the ITS analysis. The sequencing was carried out by Macrogen Inc. (the Netherlands), using the same primers as for the PCR. The two resulting strands were assembled using CLC Main Workbench 4.1.2™ or Geneious v. 11.1.5. Subsequent alignments were performed in Geneious v. 11.1.15 using the MAFFT option (auto) and adjusted manually. Unalignable ends, introns in all the aligned genes and ambiguously aligned parts were excluded from the alignment. Sequences have been submitted to GenBank as indicated in Table 1. The alignments of the three different genes were first analyzed separately to check for incongruence between genes. A conflict between the datasets was assumed to be significant if two different relationships were both supported with posterior probabilities ≥ 0.95 .

Phylogenetic analysis

Phylogenetic relationships were inferred using maximum likelihood (ML) as implemented in IQ-TREE (Nguyen *et al.* 2015) and Bayesian tree inference was carried out using Markov chain

Monte Carlo (MCMC) as implemented in MrBayes v. 3.2 (Ronquist *et al.* 2012). In the combined analysis, the three included genes were treated as separate partitions. A suitable likelihood model for each of the genes was selected, using BIC as implemented in the software jModelTest v. 2.1.4 (Guindon & Gascuel 2003; Darriba *et al.* 2012), evaluating only the 24 models available in MrBayes (Ronquist *et al.* 2012). For the concatenated dataset, the SYM + I + G model was found to be optimal for the nrITS dataset and GTR + I + G for both the nrLSU and the mtSSU datasets. For the pure nrITS dataset, the evolutionary model SYM + G was found to be optimal. The parameters used in the analyses followed those of Arup *et al.* (2013), except for the branch length prior that was set to an exponential with mean 1/10. No molecular clock was assumed. Three parallel runs with 20 000 000 generations starting with a random tree and employing six simultaneous chains were executed, five of which were incrementally heated with a temperature of 0.10. Analyses were diagnosed every 1000 generations in the last 50% of the tree sample and automatically halted when convergence was reached. Convergence was defined as a standard deviation of splits (of frequency 0.1) between runs below 0.01. Every 2000th tree was sampled. A majority-rule consensus tree was constructed from the post-burn-in tree samples. The consensus trees were visualized using FigTree v. 1.4.4 and redrawn in Adobe Illustrator. The maximum likelihood analyses used the same evolutionary models as those used in the Bayesian analyses. Branch support values were computed via 1000 non-parametric bootstrap replicates.

Results

We generated 28 new nrITS sequences, three new nrLSU sequences and five new mtSSU sequences for this study. The alignment for the first analysis consisted of 73 terminals of 2082 aligned nucleotide sites, of which 543 were parsimony-informative. The nrITS partition consisted of 502 sites (260 informative), the nrLSU partition of 768 sites (137 informative) and the mtSSU partition of 812 sites (146 informative). The Bayesian analysis halted after 1 280 000 generations and a 50% majority-rule tree with posterior probabilities is shown in Fig. 1. The second analysis of only nrITS data consisted of 30 terminals of 533 sites, of which 121 were parsimony-informative. This analysis halted after 2 060 000 generations and the 50% majority-rule tree with posterior probabilities is shown above the branches in Fig. 2. The maximum likelihood and Bayesian analyses yielded very similar trees and so only the Bayesian trees are shown. Bootstrap values are shown under the branches in the same figure.

The analysis of the combined three-gene alignment (Fig. 1) recovers all genera recognized in the subfamily *Caloplacoideae* by Arup *et al.* (2013). The tree also includes many of the genera as supported monophyletic clades proposed in subsequent papers concerning the subfamily: *Lacrima*, *Obscuroplaca*, *Oceanoplaca* and *Sucioplaca* (Bungartz *et al.* 2020, 2021), *Marchantiana* and *Yoshimuria* (Kondratyuk *et al.* 2014), *Jasonhuria*, *Loekoesia* and *Olegblumia* (Kondratyuk *et al.* 2015), *Fauriea* (Kondratyuk *et al.* (2016), *Gintarasiella* (Kondratyuk *et al.* (2017), *Upretia* (Kondratyuk *et al.* 2018) and *Lendemeriella* (Kondratyuk *et al.* 2020). It also recovers *Pyrenodesmia* as monophyletic sister to the recently resurrected genus *Kuettlingeria* and the newly described *Sanguineodiscus* (Vondrák *et al.* 2020).

Yoshimuria appears as monophyletic and is supported in the tree as sister to another strongly supported clade that includes

Table 1. Sequences from species of *Teloschistaceae*, newly produced (in bold) or downloaded from GenBank used in the analyses.

Species	Country, collector, collector no., herbarium	nrITS	nrLSU	mtSSU
<i>Blastenia ammiopila</i>	Austria, <i>Søchting</i> 9345, C (ITS, SSU); Norway, <i>Søchting</i> 10092, C (LSU)	KC179413	KC179161	KC179491
<i>B. catalinae</i>	Sweden, <i>Arup</i> L06075, LD	FJ866792	KT291532	KT291477
<i>B. crenularia</i>	Iceland, <i>Søchting</i> 7523, C	KC179415	KC179162	KC179492
<i>Bryoplaca jungermanniae</i>	Greenland, <i>Søchting</i> 10451, C	KC179420	MT952895	MT95295
<i>B. sinapisperma</i>	Norway, <i>Arup</i> L08184, LD (ITS, SSU); Norway, <i>Arup</i> L08184, LD (LSU)	KC179421	MT952896	KC179495
<i>B. tetraspora</i>	Antarctica, <i>Søchting</i> 7979, C (ITS); Greenland, <i>Søchting</i> 10480, C (LSU, SSU)	KC179422	MT952897	KC179496
<i>Caloplaca cerina</i>	Svalbard, <i>Elvebakk</i> 03:109, TROM	KC179425	KC179168	KC179499
<i>C. chlorina</i>	Denmark, <i>Søchting</i> 7321, C	KC179426	KC179169	KC179500
<i>C. turkuensis</i>	Sweden, <i>Frödén</i> 1909, LD	KC179432	MT952899	KC179501
<i>Eilifdahlia dahlia</i>	Australia, <i>Kärnefelt</i> 20043101, LD	KJ021318	KJ021253	KJ021279
<i>E. wirthii</i>	Australia, <i>Wirth</i> et al. 05.10.2011, STU	KJ021320	KJH021255	KJ021281
<i>Fauria chuaensis</i>	South Korea, KoLRI 023698	KX793095	KX793098	KX793101
<i>F. orientochinensis</i>	China, 2011, <i>Wang</i> , KoLRI 013959	KX793096	KX793099	KX793102
<i>Franwilsia bastowii</i>	Australia, <i>Kärnefelt</i> 994301, LD	KJ021324	KJ021257	KJ021284
<i>F. kilcundaensis</i>	Australia, <i>Kärnefelt</i> 20047101, LD	KJ021327	KJ021260	KJ021287
<i>Gintarsiella aggregata</i>	Australia, <i>Kantvilas</i> 476/12, HO	KY614392	KY614450	
<i>Gyalolechia aurea</i>	<i>Arup</i> L97493, LD	KC179434		KC179530
<i>G. aurea</i>	Austria, 1993, <i>Poelt & Grube</i> , GZU		KC179196	
<i>G. flavorubescens</i>	Estonia, <i>Søchting</i> 10127, C	KC179439	KC179197	KC179531
<i>G. flavovirescens</i>	Russia, <i>Søchting</i> 8648, C	AF353966	KC179198	KC179532
<i>G. fulgens</i>	Spain, <i>Søchting</i> 7306, C (ITS); Sweden, <i>Arup</i> L06206, LD (LSU);	KC179440	KC179199	
<i>G. fulgens</i>	Sweden, <i>Søchting</i> 10586, C (SSU)			KC179533
<i>Huneckia pollinii</i>	USA, Kansas, <i>Morse</i> 14464, LD	KJ021336	KJ021265	KJ021296
<i>H. wrightii</i>	Galapagos, <i>Aptroot</i> 63246, CDS (ITS); <i>Miranda</i> 962, CDS (LSU, SSU)	MT967388	MT952903	MT952929
<i>Jasonhuria bogilana</i>	South Korea, KoLRI 120454	KT220196	KT220205	KT220214
<i>Kuettlingeria albolutescens</i>	Sweden, <i>Arup</i> L09030, LD	KC179423	MT952998	KC179502
<i>K. atroflava</i>	Sweden, <i>Arup</i> L06010, LD	KC179424	KC179171	KC179504
<i>K. erythrocarpa</i>	Italy, <i>Arup</i> L07109, LD	KC179427	KC179173	KC179506
<i>K. teicholyta</i>	Denmark, <i>Søchting</i> 11195, C (ITS, LSU); Denmark, <i>Søchting</i> 9772, C (SSU)	KC179431	KC179176	KC179510
<i>Lacrima epiphora</i>	Panama, <i>van den Boom</i> 43698, hb. van den Boom	MT967392	MT952904	MT952930
<i>L. galapagoensis</i>	Galapagos, <i>Bungartz</i> 4091, CDS (ITS); <i>Ertz</i> 22855, CDS (LSU, SSU)	MT967297	MT952905	MT952931
<i>Lendemeriella reptans</i>	USA, <i>Lendemer</i> 48186, NY	MH104934	MH100766	MH100796
<i>Leproplaca chrysodeta</i>	Sweden, <i>Arup</i> L07107, LD (ITS, LSU); Sweden, <i>Arup</i> L13261, LD (SSU)	KC179448	KC179206	MT952933
<i>L. xantholyta</i>	Austria, <i>Arup</i> L97278, LD (ITS); Spain, <i>Søchting</i> 9675, C (LSU, SSU)	KC179451	KC179208	KC179542
<i>Loekoesia austrocoreana</i>	South Korea, KoLRI 120523	KT220020	KT220210	KT220219
<i>L. yuchiorum</i>	USA, <i>Lendemer</i> 22026, LD	OQ678329	OQ678282	OQ678286
<i>Marchantiana asserigena</i>	Scotland, <i>Arup</i> L10184, LD	MT967307	MT952907	MT952934
<i>M. epibrya</i> 1	New Zealand, <i>Søchting</i> 11956, C	OQ678330		
<i>M. epibrya</i> 2	Tasmania, <i>Kantvilas</i> 95/12, C	OQ678331	OQ678283	OQ678287
<i>M. epibrya</i> 3	New Zealand, <i>Søchting</i> 11957, C	OQ678332		
<i>M. michelagoensis</i> 1	New Zealand, <i>Søchting</i> 11987, C	OQ678333		
<i>M. michelagoensis</i> 2	Australia, <i>Elix</i> 37402, LD	OQ678334		OQ678288
<i>M. occidentalis</i> 1	Australia, <i>Elix</i> 32479, CANB	OQ678335		

(Continued)

Table 1. (Continued)

Species	Country, collector, collector no., herbarium	nrITS	nrLSU	mtSSU
<i>M. occidentalis</i> 2	Australia, <i>Kärnefelt</i> 20042502, LD	MT964708		KJ021304
<i>M. pyramus</i> 1	Chile, <i>Søchting</i> 11312, C	OQ678336		
<i>M. pyramus</i> 2	Chile, <i>Søgaard</i> 58, C	OQ678337		
<i>M. pyramus</i> 3	Chile, <i>Søgaard</i> 28, C	OQ678338		
<i>M. pyramus</i> 4	Chile, <i>Søgaard</i> 27B, C	OQ678339		
<i>M. pyramus</i> 5	Chile, <i>Søchting</i> 12241, C	OQ678340		
<i>M. pyramus</i> 6	Chile, <i>Søchting</i> 11363, C	OQ678341		
<i>M. pyramus</i> 7	Falkland Islands, <i>Upson</i> HSG 091109-25, C	OQ678342		
<i>M. pyramus</i> 8	Falkland Islands, <i>Søchting</i> 12654, C	OQ678343		
<i>M. pyramus</i> 9	Falkland Islands, <i>Fryday</i> 11279, C	OQ678344		
<i>M. pyramus</i> 10	Chile, <i>Søchting</i> 12667, C	OQ678345		
<i>M. pyramus</i> 11	Chile, <i>Søchting</i> 10178, C	OQ678346		
<i>M. queenslandica</i>	Australia, <i>Kalb</i> 27764, CANB	MT967409	MT952908	
<i>M. ramulicola</i> 1	Chile, <i>Søchting</i> 11353, C	OQ678347		
<i>M. ramulicola</i> 2	Chile, <i>Søchting</i> 11377, C	OQ678348		
<i>M. ramulicola</i> 3	Chile, <i>Søchting</i> 11367, C	OQ678349		
<i>M. ramulicola</i> 4	Chile, <i>Søchting</i> 12308, C	OQ678350		OQ678289
<i>M. subpyracea</i> 1	New Zealand, <i>Frödén</i> 1317, LD	OQ678351	OQ678284	OQ678290
<i>M. subpyracea</i> 2	New Zealand, <i>Søchting</i> 12013, C	OQ678352		
<i>M. subpyracea</i> 3	New Zealand, <i>Søchting</i> 11958, C	OQ678353		
<i>M. subpyracea</i> 4	New Zealand, <i>Søchting</i> 11959, C	OQ678354		
<i>Marchantiana</i> sp. 1	New Zealand, <i>Søchting</i> 11961, C	OQ678355		
<i>Marchantiana</i> sp. 2	New Zealand, <i>De Lange</i> CH 2322, AK	OQ678356		
<i>Obscuroplaca camptidia</i>	USA, <i>Morse</i> 14420, LD	MT967431	MT952918	MT952942
<i>O. ochrolechioides</i>	Australia, <i>Kalb & Rogers</i> 18982, CANB	MT967432	MT952919	
<i>O. tortuca</i>	Galapagos, <i>Aptroot</i> 65189, CDS	MT967433		MT952943
<i>Oceanoplaca caesiosorediata</i>	Cape Verde, <i>van den Boom</i> 36346, LD	MT967412	MT952911	MT952936
<i>O. catillarioides</i>	Cape Verde, <i>van den Boom</i> 36365, LD	MT967410	MT952909	MT952935
<i>O. isidiosa</i>	Galapagos, <i>Adsersen</i> , LAM8-1, C	MT967416	MT952913	MT952938
<i>Olegblumia demissa</i>	Italy, <i>Arup</i> L97911, LD	AF353960	KC179172	KC179505
<i>Pyrenodesmia alociza</i>	Sweden, <i>Arup</i> L10185, LD	MT967436	MT952920	MT952944
<i>P. chalybaea</i>	Austria, <i>Søchting</i> 9351, C	KC179454	MT952921	KC179571
<i>P. variabilis</i>	Austria, <i>Arup</i> s. n., LD (ITS); Sweden, <i>Arup</i> L03134, LD (LSU, SSU)	AF353963	KC179234	KC179572
<i>Rufoplaca scotoplaca</i>	Sweden, <i>Arup</i> L10032, LD	KC179457	KC179235	KC179573
<i>R. tristiuscula</i>	Norway, <i>Arup</i> L08171, LD	KC179460	KC179237	KC179575
<i>Rufoplaca</i> sp.	California, <i>Arup</i> L09201, LD	KC179458	KC179236	KC179574
<i>Sanguinodiscus aractinus</i>	Czech Republic, <i>Vondrák</i> 6702, PRA	MH104919		MH100773
<i>S. haematites</i>	Ukraine, <i>Vondrák</i> 7278, PRA	MH104928	MH100789	MH100756
<i>Seiophora blumii</i>	Iran, <i>Haji Moniri</i> AL_4, KW	KT456219	KT456234	KT456249
<i>S. californica</i>	Mexico, <i>Gaya</i> 03.04.10-9 & <i>Lutzoni</i> , DUKE	KT291470	KT291564	KT291521
<i>S. lacunosa</i>	Kazakhstan, <i>Moberg & Nordin</i> K18:04, LD	KC179465	KC179243	KC179582
<i>S. scorigena</i>	Lanzarote, <i>Snogerup, S. & B.</i> 17201, LD	KC179466	KC179244	KC179583
<i>Upretia amarkantakana</i>	India, 2010 LWG: 10-013313/B	MG652764		MG652767

(Continued)

Table 1. (Continued)

Species	Country, collector, collector no., herbarium	nrITS	nrLSU	mtSSU
<i>U. squamulosa</i>	China, <i>Lisong Wang</i> et al. 17-56088, KUN (ITS); China, <i>Lisong Wang</i> et al. 16-50174, KUN (LSU)	MH497054	MH497052	
<i>Usnochroma carphinea</i>	France, <i>Roux</i> 1998, C	KC179468	KC179259	KC179598
<i>U. scoriofila</i>	Tenerife, 1995, <i>Gomez-Bloea</i> , C	KC179469	KC179260	KC179599
<i>Variospora aurantia</i>	Spain, 1998, <i>Llimona</i> , C (ITS, SSU); Italy, 2006, <i>Lange</i> , C (LSU)	KC179470	KC179261	KC179600
<i>V. dolomiticola</i>	Spain, <i>Thell</i> SP0514, LD	KC179471	KC179262	KC179601
<i>V. thallincola</i>	Sweden, <i>Søchting</i> 7481, C (ITS); Sweden, <i>Gaya</i> et al. s. n., BCN (LSU)	KC179475	JQ301563	
<i>V. thallincola</i>	UK, Wales, <i>Arup</i> L92148, LD (SSU)			KC179604
<i>V. velana</i>	Italy, <i>Arup</i> L07194, LD (ITS); Italy, <i>Arup</i> L07123, LD (LSU, SSU)	KC179476	KC179265	KC179605
<i>Xanthopeltis rupicola</i>	Chile, <i>Frödén</i> 1654, LD	KC179146	KC179286	KC179626
<i>Xanthoria parietina</i>	Denmark, 2002, <i>Søchting</i> , C (ITS, SSU); Denmark, <i>Søchting</i> 7157, C (LSU)	KC179411	KC179289	KC179629
<i>Yoshimuria galbina</i>	South Korea, <i>Arup</i> L15370, LD (ITS); <i>Arup</i> 15500, LD (LSU, SSU)	MT967482	MT952923	MT952946
<i>Y. spodioplaça</i>	South Korea, <i>Wang & Ryu</i> 110364, KoLRI (ITS, SSU); <i>Arup</i> L15575, LD (LSU)	KJ021249	MT952924	KJ023194

the type of the genus *Marchantiana*, but this position lacks support. A third, fully supported clade, included here in *Marchantiana*, holds a sister position to these two genera. These three clades seem closely related to one another with strong or rather strong support (PP = 0.984 and BS = 82).

In the combined analyses, *Gintarasiella aggregata* (Kantvilas & S. Y. Kondr.) S. Y. Kondr. & Hur, type of that genus, is found within the genus *Variospora* with full support. However, this position is based only on part of an nrITS sequence (5.8S and partial ITS2) and nrLSU data, and should therefore be considered preliminary with confirmation required using more data.

The analysis of the dataset containing only nrITS sequences (Fig. 2) shows the internal relationship of the two newly proposed species, *M. ramulicola* and *M. pyramus*, and related species of the clade. The new species and *M. epibrya* are monophyletic and well supported, whereas *M. subpyracea* appears as monophyletic except for one sequence at the base of the clade which is not supported as part of the clade.

Discussion

In accordance with *Wilk et al.* (2021), the genus *Marchantiana* is shown here to belong to the subfamily *Caloplacoideae*. In the combined analysis, the two paraphyletic clades of *Marchantiana* form a supported clade together with *Yoshimuria*. From a purely genetic perspective it would be possible to either unite all three genera into one or treat them as three independent genera. The two clades, treated here as *Marchantiana*, show great similarities in morphology, ecology and distribution. They are mainly small species with small apothecia and a thin or inconspicuous thallus, all but one (*M. epibrya*) grow on bark, often twigs, and they are mainly distributed in the Southern Hemisphere (Patagonia, Australia and New Zealand), with only *M. asserigena* seeming to have migrated to Europe in the Northern Hemisphere (*Søchting & Arup* 2018). The species of the genus *Yoshimuria* on the other hand occur in South-East Asia, are saxicolous, and have a well-developed thallus with rather large apothecia. Thus, it seems logical to recognize two genera rather than three or one. This hypothesis can be assessed using the AU test as implemented in IQ-TREE (*Shimodaira* 2002), where the null hypothesis is that a tree where

the two branches of *Marchantiana* form one monophyletic clade with *Yoshimuria* outside of it is significantly worse ($P < 0.05$) than the tree presented in Fig. 1. Such a test, forcing *Marchantiana* to be monophyletic, returned a P -value of 0.5, which means that a tree with *Marchantiana* as monophyletic cannot be rejected. This makes sense since the branch including *Marchantiana* in a strict sense and *Yoshimuria* is not supported. We therefore recognize *Marchantiana* in the wider sense and describe the new species within this genus.

Taxonomy

Austroplaca thisbe Søchting & Arup sp. nov.

Mycobank No.: MB 848021

Similar to *Austroplaca sibirica* (H. Magn.) Søchting & Arup but molecularly distinct and growing on twigs instead of detritus. Thallus crustose, inconspicuous; apothecia zeorine to biatorine, yellow; ascospores polardiblastic, $13.5 \times 7.5 \mu\text{m}$, septum $5 \mu\text{m}$; on twigs in Patagonia.

Type: Chile, XII Región de Magallanes y de la Antártica Chilena, Beagle Channel, Seno Pia IV, 54.710°S , 69.696°W , 5 m, on dead twigs of *Misodendrum* on *Nothofagus antarctica*, 15 December 2009, *U. Søchting* 11317 (C—holotype; LD—isotype).

(Fig. 3)

Thallus crustose, inconspicuous.

Apothecia regular, vivid yellow, zeorine to biatorine, dispersed to mostly aggregated, sessile, never immersed or erumpent, constricted at base, 0.2–0.5 mm diam.; disc flat, with fine epipsamma, vivid yellow; thalline margin where present excluded, discontinuous, thin; proper margin distinct, c. $50 \mu\text{m}$ thick, concolorous or slightly brighter than disc, initially prominent, when older \pm level with disc; thalline exciple cortex poorly differentiated; true exciple fan-shaped, laterally up to c. $50 \mu\text{m}$; hypothecium very thin; hymenium 50–70 μm ; paraphyses simple to apically branched, c. $1.5 \mu\text{m}$ thick, terminal cells moniliform, up to $3 \mu\text{m}$ thick. Asci clavate, with 8 spores. Ascospores ellipsoid,

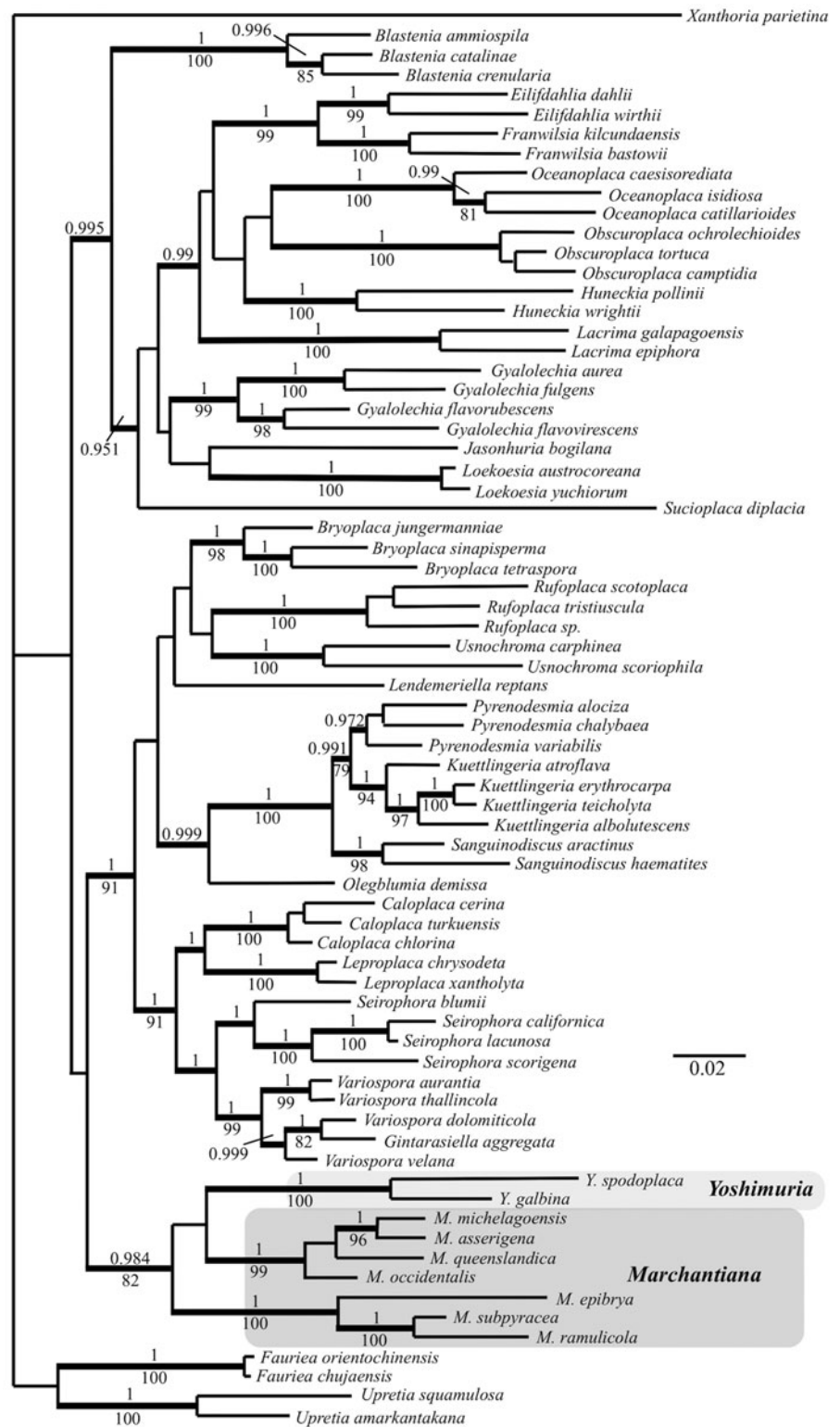


Figure 1. Majority-rule consensus tree based on a Bayesian MCMC analysis of a combined dataset of the nrITS, nrLSU and mtSSU genes showing the position of *Marchantiana* s. lat. Branches with posterior probabilities ≥ 0.95 are shown in bold. Posterior probabilities ≥ 0.95 are shown above the branches and bootstrap values ≥ 75 are shown below the branches.

polardiblastic, $(11.5)13.3 \pm 1.2(14.5) \times (6)7.6 \pm 0.7(8) \mu\text{m}$, length/width ratio 1.8 ± 0.1 , septum $(4)4.9 \pm 0.4(5.5) \mu\text{m}$, length/septum ratio 2.7 ± 0.3 ($n = 8$).

Chemistry. Chemosyndrome A of Söchting (1997): 6–10% teloschistin, 0–5% fallacinal, 0–4% parietinic acid and 85–90% parietin ($n = 3$).

Etymology. *Austroplaca thisbe* is named after Thisbe, the love of Pyramus in Ovid's classic play 'Metamorphoses'. It very often grows together with *Marchantiana pyramus* (Fig. 3).

Ecology and distribution. In *Nothofagus* forest, grassland and alpine heathland on living or dead twigs and occasionally rough bark. Phorophytes include *Berberis*, *Nothofagus* and its parasite

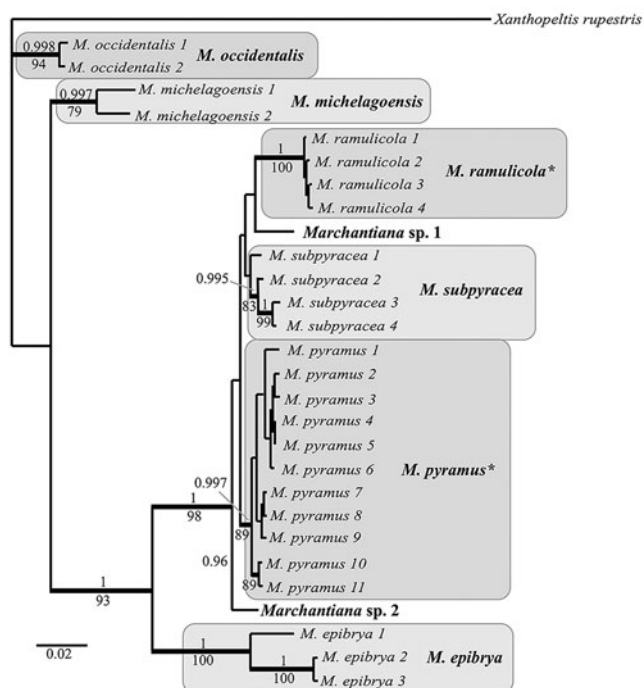


Figure 2. Majority-rule consensus tree based on a Bayesian MCMC analysis of ITS data of *Marchantiana*. Branches with posterior probabilities ≥ 0.95 are shown in bold. Posterior probabilities ≥ 0.95 are shown above the branches and bootstrap values ≥ 75 are shown below the branches. New species are marked with *.

Misodendrum. It is often associated with *Marchantiana pyramus* (see above). So far reported only from Chile but is expected to also occur in Argentina.

Notes. *Austroplaca thisbe* was included as sequence *Austroplaca* 9 in Arup *et al.* (2013). The species is molecularly well delimited and located at the base of the genus *Austroplaca*, where it is sister species to *Austroplaca sibirica*. They are morphologically very similar but are well separated by their ITS sequences and their different habitats and distribution; *A. thisbe* grows on bark and *A. sibirica* grows on dead plant material, bones, lignum and dead *Peltigera* sp. (Söchting & Arup 2021).



Figure 3. *Austroplaca thisbe* (U. Söchting 11272,2b), with one apothecium of *Marchantiana pyramus* (marked with arrow). Scale = 0.5 mm. In colour online.

Austroplaca thisbe may be mistaken for an *Athallia* but so far no molecular analysis has confirmed the presence of epiphytic *Athallia* species in Patagonia, even though it was suggested by Etayo *et al.* (2021).

Additional specimens examined. **Chile:** XI Región Aisén: c. 5 km along the road from Coyhaique to Coyhaique Alto, on dead twigs of *Berberis* sp. in open, exposed pastureland, 2001, P. Frödén 1550 (LD). XII Región de Magallanes y de la Antártica Chilena: Cueva de Miledon, 51.5611°S, 72.6160°W, 145 m, dead *Berberis* twigs, 2018, U. Söchting 12684; Morro Chico, 100 km N of Punta Arenas, basaltic rock of old volcano in steppe, 52.0575°S, 71.4212°W, 195 m, dead twigs of *Berberis*, 2015, U. Söchting 12366, 12674; 45 km SW of Punta Arenas, Reserva Nacional Laguna Parrillar, 53.4070°S, 71.2616°W, 300 m, base of *Nothofagus* sp. by lake shore, rough bark, 2005, U. Söchting 10144; Province Última Esperanza, Parque Nacional Torres del Paine, 51.0494°S, 72.9367°W, 169 m, bush (*Berberis* cf.), 2008, M. Z. Søgaard 35, 39; Parque Nacional Torres del Paine, tracks from Lago Grey, 51.1222°S, 073.1242°W, twigs, 2008, M. Z. Søgaard 46; Parque Nacional Torres del Paine, Mirador Lago Nordenskjold, halfway between Laguna Larga and Laguna Mellizas, 51.05°S, 72.92°W, 200 m, dry, sun-exposed, dead twigs of *Discaria chacaye*, 1997, A. Elvebakk 97:448 (TROM); Torres del Paine, 51.1278° S, 73.1426°W, dead twigs, 2018, U. Söchting 12690; Rio Serrano, 51.18°S, 72.95°W, dead bark of *Salix* sp., 2005, U. Söchting 10442,1; 20 km S of Puerto Natales, 51.9417°S, 72.3881°W, 100–200 m, twigs of bush, 2008, M. Z. Søgaard 27a; Hosteria Cabo San Isidro, 53.7822°S, 70.9737°W, 2 m, dead *Nothofagus betuloides* trunk on beach, 2009, U. Söchting 11351; Seno Almirantazgo, 54.703°S, 69.332°W, 25 m, old dead twigs of *Misodendrum*, 2009, U. Söchting 11287; Beagle Channel, Seno Pia, west arm, shore of fjord, 54.7897°S, 69.6946°W, 1 m, twigs of *Chilotrimum* hanging over the sea, 2 m from the sea, 2015, U. Söchting 11308; Cabo Hyades, Bahía Yendegaia, 54.9158°S, 68.7137°W, 1 m, dead twigs of *Berberis*, 2015, U. Söchting 12316; Isla Navarino, 30 km WNW of Puerto Williams, Wulaia, 55.0467°S, 68.1467°W, 2 m, twigs of *Berberis* sp., 2005, U. Söchting 10336; Isla Navarino, Puerto Williams around the airport, pasture on beach, 54.9291°S, 67.6372°W, 7 m, dead *Berberis* twigs, 2015, U. Söchting 12226; 37 km W of Puerto Williams, Caleta Honde, 54.93°S, 68.28°W, 5 m, dead *Chilotrimum*, 2005, U. Söchting 10179; 20 km W of Puerto Williams, Caleta Mejillones, 54.9067°S, 68.1180°W, 5 m, *Berberis* twigs, 2005, U. Söchting 10189; Laguna Amarga, 50.9783°S, 72.78°W, 990 m, twigs, 2005, U. Söchting 10443; 2 km E of Puerto Navarino, 54.9270°S, 68.2756°W, 0.5 m, dead twigs of *Berberis*, 2015, U. Söchting 12333.

Blastenia circumpolaris Söchting, Frödén & Arup

(Fig. 4)

Notes. *Blastenia circumpolaris* was originally described as *Caloplaca wilsonii* S. Y. Kondr. *et al.* from Australia (Kondratyuk *et al.* 2009) but was transferred to *Blastenia* by Arup *et al.* (2013). It is an extremely common epiphyte in Tierra del Fuego, where it is found as dispersed thalli with characteristic orange-brown soralia on the bark of *Nothofagus* in particular. It also colonizes twigs, where it forms thallus with just a small number of soralia; fertile specimens are found only rarely. In the Falkland Islands, Tasmania and New Zealand, it is much

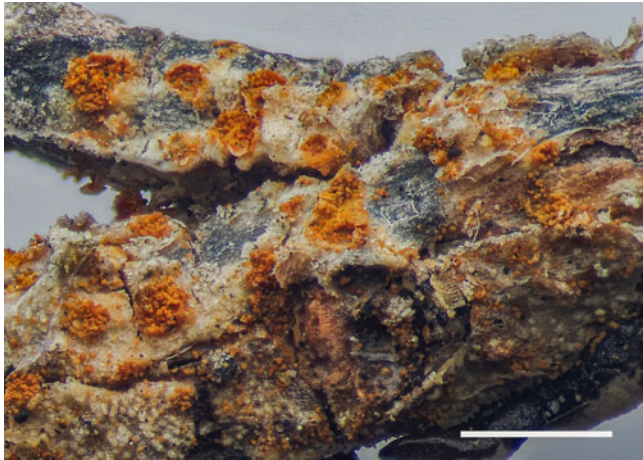


Figure 4. *Blastenia circumpolaris* (U. Søchting 11296). Scale = 1 mm. In colour online.

less frequent. Its currently known distribution outside mainland Australia is shown in Fig. 5.

Specimens studied. **Argentina:** Provincia de Santa Cruz: Rio Turbida at border post NE of Puerto Natales, 51.5635°S, 72.3483°E, 620 m, 2005, U. Søchting 10425.—**Australia:** **Tasmania:** North-East Tasmania, 15 km S of St Helens, Shelley Point, *Eucalyptus* regrowth forest by coast, 41.435°S, 148.274°E, 5 m, live bark, 2011, U. Søchting 11627.—**Chile:** **Región del Biobío:** Alto Bio Bio, 38.635°S, 70.96°W, 2012, R. Vargas 3708 (C). **IX Región del Araucanía:** Malleco Province, Parque

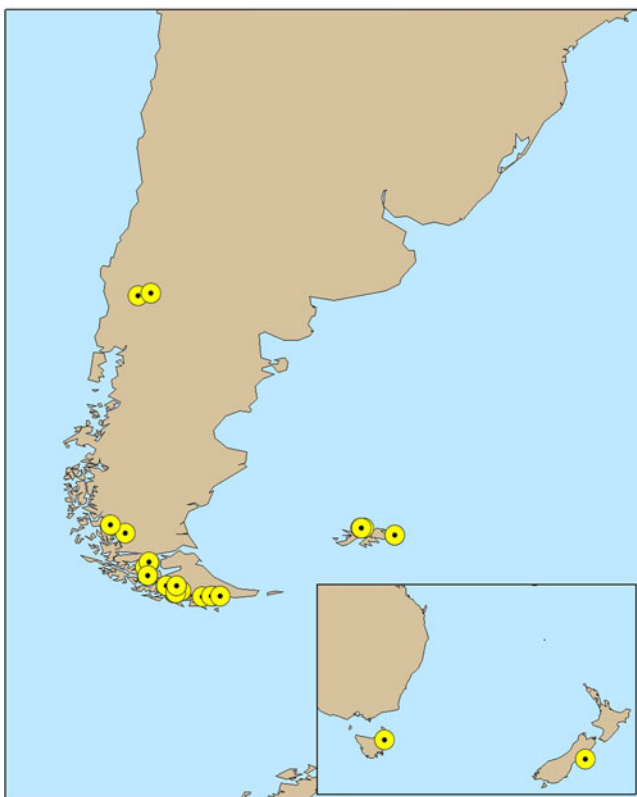


Figure 5. Distribution map of *Blastenia circumpolaris* occurring outside the Australian mainland. In colour online.

Nacional Conguillio, Sector Melipeuco, Sendero las Vertientes, 38.7669°S, 71.6344°W, bark of *Nothofagus*, 2011, R. Vargas (C). **XII Región de Magallanes y de la Antártica Chilena:** Parque Nacional Torres del Paine, walk from Lago Grey, 51.1181°S, 73.1419°W, trunk of dead *Nothofagus*, 2008, M. Z. Søgaard 55; 10 km north of Punta Arenas, 53.1403°S, 74.0544°W, living *Nothofagus*, 2008, M. Z. Søgaard 63, 65; 45 km SW of Punta Arenas, Reserva Nacional Laguna Parillar, 53.4161°S, 71.2614°W, base of *Nothofagus* by lake shore, 2005, U. Søchting 10142, 10143; Seno Almirantazgo, Ainsworth Bay, 54.420°S, 69.562°W, dead *Chillotrichum*, 2009, U. Søchting 11272; 1 m, dead *Berberis* twigs, U. Søchting 11273; *ibid.*, 54.703°S, 69.332°W, 25 m, bark of dead twigs of *Nothofagus antarctica*, 2009, U. Søchting 11288; *ibid.*, 54.432°S, 70.118°W, 1 m, dead twigs of *Pernettya mucronulata*, 2009, U. Søchting 11296; *ibid.*, 54.432°S, 70.118°W, 1 m, dead twigs of *Berberis*, 2009, U. Søchting 11297; Hosteria Cabo San Isidro, 53.7822°S, 70.9737°W, bark of *Nothofagus betuloides* on beach, 2009, U. Søchting 11350, 11357; Beagle Channel, Seno Pia, 54.780°S, 69.590°W, 3 m, dead *Misodendrum* on *Nothofagus antarctica*, 2009, U. Søchting 11312b; *ibid.*, 54.772°S, 69.606°W, 20 m, old trunk of *Nothofagus betuloides*, 2009, U. Søchting 11314; *ibid.*, 54.719°S, 69.705°W, 2 m, old *Nothofagus betuloides*, 2009, U. Søchting 11319; *ibid.*, 54.772°S, 69.606°W, 20 m, old trunk of *Nothofagus betuloides*, 2009, U. Søchting 11314; *ibid.*, 54.7634°S, 69.6028°W, 98 m, old trunk of *Nothofagus betuloides*, 2015, U. Søchting 12282; *ibid.*, 54.7942°S, 69.6987°W, 1 m, dead twigs of *Nothofagus betuloides* overhanging the sea, 2015, U. Søchting 12306; Isla Navarino, 30 km W of Puerto Williams, Caleta José, 55.008°S, 68.174°W, 2 m, dry bark of *Nothofagus betuloides*, 2005, U. Søchting 10372; 7 km SW of Puerto Williams, Valle Róbalo, 54.9528°S, 67.6386°W, 240 m, trunk of *Nothofagus betuloides*, 2005, U. Søchting 10152; *ibid.*, 54.9614°S, 67.1933°W, 2 m, dead *Nothofagus*, 2008, M. Z. Søgaard 74.—**New Zealand:** **South Island:** Canterbury, 1 km N of Akaroa, volcanic outcrops in pasture, 43.7973°S, 173.0274°E, 742 m, rotten wood, 2012, U. Søchting 11989.—**Falkland Islands:** **East Island:** Gipsy Bay, outcrops in heathland, 51.7042°S, 57.9165°W, dead *Bolax* cushions, 2018, U. Søchting 12606; *ibid.*, dead *Empetrum* twigs, U. Søchting 12605. **Pebble Island:** pebbles on protected beach, 51.3039°S, 59.6032°W, rotten wood, 2018, U. Søchting 12663; *ibid.*, 51.3095°S, 59.6123°W, dead *Empetrum* twigs, 2018, U. Søchting 12653; *ibid.*, 51.3090°S, 59.4721°W, driftwood on beach, 2018, U. Søchting 12662.

Marchantia asserigena (J. Lahm) Søchting & Arup

Notes. The very insignificant species *Marchantia asserigena* has recently been shown to be fairly common in Denmark and Norway (Søchting & Frøberg 2003; Søchting & Arup 2018; Tønsgaard *et al.* 2021), but it has also been found in Sweden, Italy, France, Germany, Austria and the British Isles. Based on morphology, secondary chemistry and ITS data, it was identified from the Falkland Islands where it grew in similar habitats to those in the Northern Hemisphere, namely very thin twigs of dwarf shrubs. It is the first proven record from the Southern Hemisphere but it may have a much wider distribution.

Specimen studied. **Falkland Islands:** **East Island:** Gipsy Bay, 51.6765°S, 57.8101°W, 26 m, heathland, dead *Empetrum* twigs, 2018, U. Søchting 12604.

The molecular phylogeny (Fig. 2), combined with distribution and secondary chemistry, place *Marchantiana pyramus* in a sister position to *Marchantiana epibrya*, which is combined here into the genus *Marchantiana* as:

***Marchantiana epibrya* (Kantvilas & Søchting) Søchting & Arup comb. nov.**

MycoBank No: MB 848017

Basionym: *Caloplaca epibrya* Kantvilas & Søchting, *Kanunnah* 6, 110 (2013) (MycoBank No.: MB 805211).

Based on the molecular data (Fig. 1), *Caloplaca queenslandica* is combined into *Marchantiana* as:

***Marchantiana queenslandica* (Kalb, S. Y. Kondr., Elix & Kärnefelt) Arup & Søchting comb. nov.**

MycoBank No.: MB 848018

Basionym: *Caloplaca queenslandica* Kalb *et al.*, *Australas. Lichenol.* 66, 35 (2010) (MycoBank No.: MB 548337).

***Marchantiana pyramus* Søchting & Arup sp. nov.**

MycoBank No.: MB 848019

Thallus crustose, inconspicuous; similar to *Marchantiana ramulicola*, but apothecia orange and C+ red due to dominance of the anthraquinone emodin; ascospores polardiblastic, $12 \times 5.5 \mu\text{m}$, septum $5 \mu\text{m}$; on twigs in Patagonia.

Type: Chile, XII Región de Magallanes y de la Antártica Chilena, Seno Almirantazgo, Ainsworth Bay, 54.420°S , 69.562°W , 1 m, dead *Chilotrimum*, 6 December 2009, Søchting 11272.2a (C—holotype; LD—isotype).

(Fig. 6)

Thallus inconspicuous or smooth, thin and whitish.

Apothecia regular, biatorine, mostly few together and dispersed, sessile, 0.3–0.4 mm diam.; disc flat, orange, only slightly darker than the exciple, with coarse epipsamma; proper margin regular, orange, often brownish near the disc, eventually



Figure 6. *Marchantiana pyramus* (U. Søchting 10178). Scale = 0.5 mm. In colour online.

sometimes almost black at the surface, 30–50 μm thick, almost level with disc; true exciple fan-shaped, laterally up to c. 40 μm , close to surface with tissue of elongated cells, $3 \times 10 \mu\text{m}$; hymenium c. 60 μm ; paraphyses simple to poorly branched, 1.5–2 μm thick, terminal cells up to 3 μm thick. Asci clavate, 8-spored. Ascospores polardiblastic, $(10)12 \pm 1.2(15) \times (4.5)5.3 \pm 0.4(6) \mu\text{m}$, length/width ratio 2.3 ± 0.3 , septum $(3.5)4.8 \pm 0.6(6) \mu\text{m}$, length/septum ratio 2.6 ± 0.4 ($n = 22$).

Chemistry. Chemosynndrome named here as E3: emodin 20–60%, with oxidation products citreorosein 30–50%, emodinol 20–30% and emodinic acid 15–40%. Thallus K–; apothecia K+ purple and C+ dark reddish.

Etymology. The epithet commemorates the mythological person Pyramus, appearing in Ovid's classic play 'Metamorphoses' with his beloved Thisbe, whose name is used in this paper for a new *Austroplaca* species, very often growing intricately together with *M. pyramus* (see below and Fig. 3).

Ecology and distribution. *Marchantiana pyramus* grows on thin branches of shrubs and on twigs of *Nothofagus*. It is very common in southern Patagonia, both along the coast and inland. Often associated with *Austroplaca thisbe* (Fig. 3). In the Falkland Islands it grew on dead twigs of dwarf bushes. It has not been recorded from Tasmania or New Zealand.

Notes. *Marchantiana pyramus* shares its morphology and ecology with most of the other species in the genus *Marchantiana*, particularly with *M. asserigena*, which also occurs in the Northern Hemisphere.

Additional specimens studied. **Chile:** XII Región de Magallanes y de la Antártica Chilena: Parque Nacional Torres del Paine, walk from Lago Grey, 51.1181°S , 73.1419°W , branch of *Ribes*, 2008, M. Z. Søgaard 57b; 20 km S of Puerto Natales, twig of *Berberis*, M. Z. Søgaard 58; Provincia Ultima Esperanza, 20 km S of Puerto Natales, on twigs of *Berberis*, 2008, M. Z. Søgaard 27b, 28; Seno Otway, 100 m from the sea, 53.0936°S , 71.3365°W , 1 m, dead stems of *Empetrum*, 2018, U. Søchting 12667; Montes Admiral, 52.0111°S , 72.3682°W , dead *Berberis* twigs, 2018, Søchting 12698; Hosteria Cabo San Isidro, 53.7822°S , 70.9737°W , 2 m, dead *Ribes*, 2009, U. Søchting 11363, 11369, 11372; Beagle Channel, Cabo Hyades, Bahía Yendegaia, 54.9158°S , 68.7137°W , 1 m, dead twigs of *Nothofagus*, 2015, U. Søchting 12317; Seno Pia, 54.780°S , 69.590°W , 3 m, dead *Misodendrum* on *Nothofagus antarctica*, 2009, U. Søchting 11312a; Isla Navarino, Puerto Williams around the airport, 54.9269°S , 67.6177°W , 9 m, *Berberis* twig, 2015, U. Søchting 12241; 37 km W of Puerto Williams, Caleta Honde, 54.93°S , 68.28°W , 5 m, dead *Chilotrimum*, 2005, U. Søchting 10178; *ibid.*, 54.92°S , 68.23°W , 10 m, dead twigs of *Pernettya*, 2015, U. Søchting 12340.—**Falkland Islands:** East Falkland: Gypsy Cove, $51.674340^{\circ}\text{S}$, $57.808660^{\circ}\text{W}$, 10 m, *Bolax* on cliff top, 2015, A.M. Fryday 11279 (C, MSC). Pebble Island: maritime rocks and pebbles on shore, 51.3094°S , 59.6124°W , 40 m, dead *Empetrum* stem, 2018, U. Søchting 12654.

***Marchantiana ramulicola* Søchting & Arup sp. nov.**

MycoBank No.: MB 848020

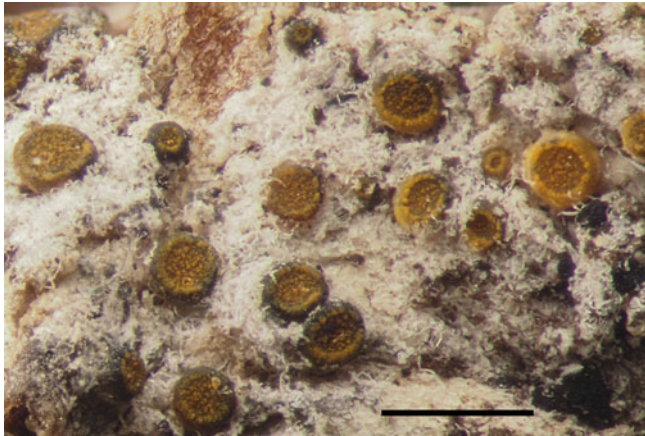


Figure 7. *Marchantiana ramulicola* (U. Søchting 11377). Scale = 0.5 mm. In colour online.

Thallus crustose, inconspicuous; similar to *Marchantiana pyramus* but apothecia more olive and C– due to chemosyndrome A dominated by parietin; ascospores polaridiblastic, $13 \times 8 \mu\text{m}$, septum $4.5 \mu\text{m}$; on twigs in Patagonia.

Type: Chile, XII Region de Magallanes y de la Antártica Chilena, Hosteria Cabo San Isidro, 53.7822°S , 70.9737°W , 2 m, dead twigs of *Chilotrimum*, 17 December 2009, U. Søchting 11377 (C—holotype; LD—isotype).

(Fig. 7)

Thallus inconspicuous or smooth, thin and greyish, on smooth bark with a blackish hypothallus.

Apothecia regular, biatorine, dispersed to aggregated, initially immersed, eventually sessile and restricted at the base, 0.2–0.4 mm diam.; *disc* flat, yellow to dark orange, with coarse epip-samma; *proper margin* regular, olive in outer part, with a brighter yellow tinge near the disc, c. $50 \mu\text{m}$ thick, initially sometimes urniform, inflated below and constricted above, prominent, with a sharp edge towards the disc; *true exciple* fan-shaped, laterally up to c. $40 \mu\text{m}$; *hypotheceum* very thin, c. $15 \mu\text{m}$; *hymenium* c. $60 \mu\text{m}$; *paraphyses* mostly simple, c. $1.5 \mu\text{m}$ thick, terminal cells slightly thickened, up to $3 \mu\text{m}$. *Asci* clavate, 8-spored. *Ascospores* $(10.5)13.1 \pm 1.3$ $(15) \times (6.5)7.8 \pm 0.8(10) \mu\text{m}$, length/width ratio 1.7 ± 0.2 , septum $(3.5)4.6 \pm 0.5(5) \mu\text{m}$, length/septum ratio 2.9 ± 0.4 ($n = 20$).

Chemistry. Chemosyndrome A of Søchting (1997): 6–10% teloschistin, 0–5% fallacinal, 0–4% parietinic acid and 85–90% parietin ($n = 3$).

Etymology. The epithet refers to the preferred growth on thin twigs.

Ecology and distribution. The species grows on thin, dead branches and twigs of, for example, *Chilotrimum* and *Fuchsia*. It is so far known only from two localities in southern Chile.

Additional specimens studied. **Chile:** XII Región de Magallanes y Antártica Chilena: Seno Almirantazgo, 54.432°S , 70.118°W , 1 m, dead twigs of *Chilotrimum*, 2009, U. Søchting 11295; Beagle Channel, Seno Pia, west arm, shore of fjord, 54.7897°S , 69.6946°W , 2 m from the sea, 1 m, twigs of *Chilotrimum* hanging over the sea,

2015, U. Søchting 12308; Hosteria Cabo San Isidro, 53.7822°S , 70.9737°W , 2 m, dead twigs, 2009, U. Søchting 11353, 11367.


Based on the molecular results *Caloplaca subpyracea* (Nyl.) Zahlbr. from New Zealand is combined into *Marchantiana*:

Marchantiana subpyracea (Nyl.) Søchting & Arup comb. nov.

Mycobank No.: MB 848016

Basionym: *Lecanora subpyracea* Nyl. *Lich. Nov. Zel.*, 59 (1888) (Mycobank No.: MB 389373); type: New Zealand, *sine loco*, 1867, Charles Knight (H–NYL 29847—lectotype selected here, MBT 10014522).

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References

- Arup U, Søchting U and Frödén P (2013) A new taxonomy of *Teloschistaceae*. *Nordic Journal of Botany* **31**, 16–83.
- Arup U, Vondrák J and Halıcı MG (2015) *Parvoplaca nigroblastidiata*, a new corticolous lichen (*Teloschistaceae*) in Europe, Turkey and Alaska. *Lichenologist* **47**, 379–385.
- Bungartz F, Søchting U and Arup U (2020) *Teloschistaceae* (lichenized *Ascomycota*) from the Galapagos Islands: a phylogenetic revision based on morphological, anatomical, chemical, and molecular data. *Plant and Fungal Systematics* **65**, 515–576.
- Bungartz F, Søchting U and Arup U (2021) *Obscuropalca* gen. nov. – a replacement name for *Phaeoplaca*; *Teloschistaceae* (lichenized *Ascomycota*) from the Galapagos Islands. *Plant and Fungal Systematics* **66**, 240–241.
- Darriba D, Taboada GL, Doallo R and Posada D (2012) jModelTest 2: more models, new heuristics and parallel computing. *Nature Methods* **9**, 772.
- Degelius G (1964) Biological studies of the epiphytic vegetation of twigs of *Fraxinus excelsior*. *Acta Horti Gothoburgensis* **27**, 11–27.
- Degelius G (1978) Further studies of the epiphytic vegetation on twigs. *Botanica Gothoburgensis* **7**, 1–58.
- Döring H, Clerc P, Grube M and Wedin M (2000) Mycobiont-specific PCR primers for the amplification of nuclear ITS and LSU rDNA from lichenized ascomycetes. *Lichenologist* **32**, 200–204.
- Etayo J, Sancho LG, Gómez-Bolea A, Søchting U, Aguirres F and Rozzi R (2021) Catalogue of lichens (and some related fungi) of Navarino Island, Cape Horn Biosphere Reserve, Chile. *Anales Instituto de la Patagonia* **49**, 1–110.
- Gardes M and Bruns TD (1993) ITS primers with enhanced specificity for basidiomycetes – application to the identification of mycorrhizae and rusts. *Molecular Ecology* **2**, 113–118.
- Guindon S and Gascuel O (2003) A simple, fast, and accurate algorithm to estimate large phylogenies by maximum likelihood. *Systematic Biology* **52**, 696–704.
- Kondratyuk SY, Kärnefelt I, Thell A, Elix JA, Kim J, Kondratyuk AS and Hur J-S (2009) Contributions to the *Teloschistaceae*, with particular reference to the Southern Hemisphere. *Bibliotheca Lichenologica* **100**, 207–287.
- Kondratyuk SY, Kärnefelt I, Thell A, Elix JA, Kim J, Jeong M-H, Yu N-N, Kondratyuk AS and Hur J-S (2014) A revised taxonomy for the subfamily *Caloplacoideae* (*Teloschistaceae*, *Ascomycota*) based on molecular phylogeny. *Acta Botanica Hungarica* **56**, 93–123.
- Kondratyuk S, Lökös L, Kim JA, Kondratyuk AS, Jeong M-H, Jang SH, Oh S-O and Hur J-S (2015) Three new monotypic genera of the caloplacoid

- lichens (*Teloschistaceae*, lichen-forming ascomycetes). *Mycobiology* **43**, 195–202.
- Kondratyuk S, Lökös L, Kim JA, Kondratiuk AS, Jeong M-H, Jang SH, Oh S-O, Wang XY and Hur J-S (2016) *Fauriea*, a new genus of the lecanoroid caloplacoid lichens (*Teloschistaceae*, lichen-forming ascomycetes). *Acta Botanica Hungarica* **58**, 303–318.
- Kondratyuk S, Lökös L, Upreti DK, Nayaka S, Mishra GK, Ravera S, Jeong M-H, Jang S-H, Park JS and Hur J-S (2017) New monophyletic branches of the *Teloschistaceae* (lichen-forming *Ascomycota*) proved by three gene phylogeny. *Acta Botanica Hungarica* **59**, 71–136.
- Kondratyuk S, Persson P-E, Hansson M, Mishra GK, Nayaka S, Liu D, Hur J-S and Thell A (2018) *Upretia*, a new caloplacoid genus (*Teloschistaceae*, lichen-forming *Ascomycota*) from India. *Cryptogam Biodiversity and Assessment Special Volume* (2018), 22–31.
- Kondratyuk S, Lökös L, Farkas E, Kärnefelt I, Thell A, Yamamoto Y and Hur J-S (2020) Three new genera of the *Teloschistaceae* proved by three gene phylogeny. *Acta Botanica Hungarica* **62**, 109–136.
- Larsen RV, Wolseley PA, Søchting U and Chimonides J (2009) Biomonitoring with lichens on twigs. *Lichenologist* **41**, 189–202.
- Lumbsch HT, Ahti T, Altermann S, Amo de Paz G, Aptroot A, Arup U, Bárcenas Peña A, Bawingan PA, Benatti MN, Betancourt L, et al. (2011) One hundred new species of lichenized fungi: a signature of undiscovered global diversity. *Phytotaxa* **18**, 1–127.
- Nguyen L-T, Schmidt HA, von Haeseler A and Minh BQ (2015) IQ-TREE: a fast and effective stochastic algorithm for estimating maximum-likelihood phylogenies. *Molecular Biology and Evolution* **32**, 268–274.
- Ott S, Schröder T and Jahns HM (2000) Colonization strategies and interactions of lichens on twigs. *Bibliotheca Lichenologica* **75**, 445–455.
- Ronquist F, Teslenko M, van der Mark P, Ayres DL, Darling A, Höhna S, Larget B, Liu L, Suchard MA and Huelsenbeck JP (2012) MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across a large model space. *Systematic Biology* **61**, 539–542.
- Shimodaira H (2002) An approximately unbiased test of phylogenetic tree selection. *Systematic Biology* **3**, 492–508.
- Søchting U (1997) Two major anthraquinone chemosyndromes in *Teloschistaceae*. *Bibliotheca Lichenologica* **68**, 135–144.
- Søchting U and Arup U (2018) *Marchantiana asserigena* comb. nov., a possible European immigrant from Australia. *Graphis Scripta* **30**, 115–120.
- Søchting U and Arup U (2021) *Austroplaca sibirica* (lichenized *Ascomycota*, *Teloschistaceae*) – a bipolar lichen with an austral origin documented by molecular data. *Graphis Scripta* **33**, 12–16.
- Søchting U and Fröberg L (2003) *Caloplaca asserigena*, the correct name for a small lichen with a unique chemistry. *Bibliotheca Lichenologica* **86**, 47–51.
- Søchting U and Sancho LG (2012) *Caloplaca magellanica* sp. nova, a southern Patagonian parasite on *Zahlbrucknerella*. *Bibliotheca Lichenologica* **108**, 215–220.
- Søchting U, Søgaard MZ and Elix JA (2014) *Catenarina* (*Teloschistaceae*, *Ascomycota*), a new Southern Hemisphere genus with 7-chlorocatenarin. *Lichenologist* **46**, 175–187.
- Søchting U, Søgaard MZ, Sancho LG, Frödén P and Arup U (2016) *Sirenophila ovis-atra*, a new species of maritime *Teloschistaceae* from the Southern Hemisphere. *Opuscula Philolichenum* **15**, 1–5.
- Søchting U, Søgaard MZ, Sancho LG and Arup U (2021) The lichen genus *Villophora* (*Teloschistaceae*, *Ascomycota*). *Lichenologist* **53**, 245–255.
- Tønsgaard T, Palice Z and Tindal E (2021) *Marchantiana asserigena*, a genus and species new to Norway. *Graphis Scripta* **33**, 67–78.
- Vilgalys R and Hester M (1990) Rapid genetic identification and mapping of enzymatically amplified ribosomal DNA from several *Cryptococcus* species. *Journal of Bacteriology* **172**, 4239–4246.
- Vondrák J, Frolov I, Košnar J, Arup U, Veselská T, Hahcı G, Maliček J and Søchting U (2020) Substrate switches, phenotypic innovations and allopatric speciation formed taxonomic diversity within the lichen genus *Blastenia*. *Journal of Systematics and Evolution* **58**, 295–330.
- White TJ, Bruns TD, Lee S and Taylor J (1990) Application and direct sequencing of fungal ribosomal DNA for phylogenetics. In Innis MA, Gelfand DH, Sninsky JJ and White TJ (eds), *PCR Protocols: a Guide to Methods and Applications*. San Diego: Academic Press, pp. 315–322.
- Wilk K, Pabijan M, Saluga M, Gaya E and Lücking R (2021) Phylogenetic revision of South American *Teloschistaceae* (lichenized *Ascomycota*, *Teloschistales*) reveals three new genera and species. *Mycologia* **113**, 278–299.
- Zhou S and Stanosz GR (2001) Primers for amplification of mtSSU rDNA, and a phylogenetic study of *Botryosphaeria* and associated anamorphic fungi. *Mycological Research* **105**, 1033–1044.
- Zoller S, Scheidegger C and Sperisen C (1999) PCR primers for the amplification of mitochondrial small subunit ribosomal DNA of lichen-forming ascomycetes. *Lichenologist* **31**, 511–516.