

## NEW MONOPHLETIC BRANCHES OF THE TELOSCHISTACEAE (LICHEN-FORMING ASCOMYCOTA) PROVED BY THREE GENE PHYLOGENY

S. Y. KONDRATYUK<sup>1,2</sup>, L. LŐKÖS<sup>3</sup>, D. K. UPRETI<sup>4</sup>, S. NAYAKA<sup>4</sup>, G. K. MISHRA<sup>4</sup>  
S. RAVERA<sup>5</sup>, M.-H. JEONG<sup>2</sup>, S.-H. JANG<sup>2</sup>, J. S. PARK<sup>2</sup> and J.-S. HUR<sup>2</sup>

<sup>1</sup>M. H. Kholodny Institute of Botany, Tereshchenkivska str. 2, 01004 Kiev, Ukraine  
E-mail: ksyu\_net@ukr.net

<sup>2</sup>Korean Lichen Research Institute, Sunchon National University  
Sunchon 540-742, Republic of Korea; E-mail: jshur1@sunchon.ac.kr

<sup>3</sup>Department of Botany, Hungarian Natural History Museum  
H-1431 Budapest, Pf. 137, Hungary; E-mail: lokos.laszlo@nhmus.hu

<sup>4</sup>CSIR-National Botanical Research Institute

Rana Pratap Marg, Lucknow-226001 Uttar Pradesh, India; E-mail: upretidknbri@gmail.com

<sup>5</sup>Dipartimento di Bioscienze e Territorio, Università degli Studi del Molise  
C. da Fonte Lappone, I-86090 Pesche (IS), Italy

(Received 10 August, 2016; Accepted 5 December, 2016)

Seventeen robust monophyletic branches newly discovered in the phylogenetic tree of the Teloschistaceae after separate nrITS, nrLSU and mtSSU, as well as combined phylogenetic analysis are proposed to consider as the following separate genera: *Dijigiella* S. Y. Kondr. et L. Lőkös gen. nov. for the *D. kaernefeltiana* group, *Elixjohnnia* S. Y. Kondr. et J.-S. Hur gen. nov. for the *Sirenophila jackelixii* group, *Fominiella* S. Y. Kondr., D. Upreti et J.-S. Hur gen. nov. for the *F. tenerifensis* group; *Gintarasiella* S. Y. Kondr. et J.-S. Hur gen. nov. for *Caloplaca aggregata*, *Hanstrassia* S. Y. Kondr. gen. nov. for the *Elenkiniana lenae* group, *Harusavskia* S. Y. Kondr. gen. nov. for *H. elenkinianoides* sp. n., *Huriella* S. Y. Kondr. et D. Upreti gen. nov. for *H. loekoesiana* sp. n., *Ikaeria* S. Y. Kondr., D. Upreti et J.-S. Hur gen. nov. for *Caloplaca aurantiellina*, *Klauderriella* S. Y. Kondr. et J.-S. Hur gen. nov. for the *Variospora thallincola* group, *Laundonia* S. Y. Kondr., L. Lőkös et J.-S. Hur gen. nov. for the *Gyalolechia flavovirescens* group, *Lazarenkoiopsis* S. Y. Kondr., L. Lőkös et J.-S. Hur gen. nov. for *Caloplaca ussuriensis*, *Nevilleiella* S. Y. Kondr. et J.-S. Hur gen. nov. for the *Caloplaca marchantii* group, *Opeltia* S. Y. Kondr. et L. Lőkös gen. nov. for the *Caloplaca neobaltistanica* group, *Oxneriopsis* S. Y. Kondr., D. Upreti et J.-S. Hur gen. nov. for the *Caloplaca oxneri* group, *Teuvoahntiana* S. Y. Kondr. et J.-S. Hur gen. nov. for the *Caloplaca rugulosa* group, *Tomnashia* S. Y. Kondr. et J.-S. Hur gen. nov. for the *Polycauliona rosei* group, and *Xanthapychia* S. Y. Kondr. et S. Raveri gen. nov. for the *Seirophora orientalis* group.

Hitherto missing molecular data on three gene sequences of the type species of the genera *Seirophora* and *Sirenophila* are completed within this study.

Six new to science species (*Dijigiella kaernefeltiana* S. Y. Kondr. sp. n., *D. subaggregata* S. Y. Kondr. et Kärnefelt sp. n., *Fominiella tenerifensis* S. Y. Kondr., Kärnefelt, A. Thell et T. Feuerer sp. n., *Hanstrassia jaeseounhurii* S. Y. Kondr., Ch.-H. Park et L. Lőkös sp. n., *Harusavskia elenkinianoides* S. Y. Kondr., X. Y. Wang, S.-O. Oh et J.-S. Hur sp. n., *Huriella loekoesiana* S. Y. Kondr. et D. Upreti sp. n.) are described, compared with closely related taxa.

A total of 34 new combinations for genera mentioned above are proposed.

Key words: phylogenetic analysis, phylogenetic tree, Teloschistaceae, three gene phylogeny

## INTRODUCTION

The taxonomy of the Teloschistaceae has undergone a radical change during recent years, including the creation of a large number of new genera, based mainly on molecular phylogeny (Arup *et al.* 2013a, Fedorenko *et al.* 2012, Gaya *et al.* 2012, Kondratyuk *et al.* 2013b, 2014a, b, 2015a, c, d, 2016c, d). Hence the number of genera in the Teloschistaceae has increased from 10 in the late 1990s (Kärnefelt 1989) to currently 80 (Arup *et al.* 2013a, Kondratyuk *et al.* 2013b, 2014a, c, 2015a, c, d, 2016c, d, Søchting *et al.* 2014a, b), and the family is divided in four subfamilies, Brownlielloideae, Caloplacoideae, Teloschistoideae and Xanthorioideae (Gaya *et al.* 2012, Arup *et al.* 2013b, Kondratyuk *et al.* 2015d).

New data on ITS1/ITS2 nrDNA, 28S nrLSU, and 12S mtSSU sequences for representatives of all four subfamilies of the Teloschistaceae accumulated during 2015 and 2016 were found to illustrate a number of new robust monophyletic branches within the phylogenetic tree of the Teloschistaceae.

The aim of this paper is to provide descriptions of newly discovered robust monophyletic branches, which are supported as separate ITS1/ITS2, nrLSU and mtSSU as combined phylogenetic analysis. Totally 17 robust monophyletic branches newly discovered in the phylogenetic tree of the Teloschistaceae after as separate nrITS, nrLSU and mtSSU, as combined phylogenetic analyses are proposed to consider as separate genera. Descriptions of six new to science species belonging to these genera are provided below, too.

## MATERIALS AND METHODS

More than 1,000 Teloschistaceae specimens, collected in 2014–2016 and deposited in the Korean Lichen Research Institute, Sunchon National University, South Korea (KoLRI), as well as some duplicates in the Hungarian Natural History Museum (BP) and the Lichen Herbarium in the M. H. Kholodny Institute of Botany of National Academy of Sciences of Ukraine (KW-L) have been examined using standard microscopical techniques, and hand-sectioned under a dissecting microscope (Nikon SMZ 645; Nikon, Tokyo, Japan). Anatomical characters were observed using a Nikon Eclipse E200 microscope and a Zeiss Scope, complemented with a digital camera AxioCam ERc 5s. Sections of apothecia were tested with water, K and IKI (10% potassium iodide).

Total DNA was extracted directly from the thalli according to Ekman (1999) and was purified with DNeasy Plant Mini Kit (QIAGEN, Germany). The nuclear ribosomal RNA gene region including the internal transcribed spacers 1 and 2 and the 5.8S subunit (ITS) was amplified using the primers ITS1F (Gardes and Bruns 1993) and ITS4 (White *et al.* 1990), the 28S LSU using

the primer LR5 (Vilgalys and Hester 1990), and the 12S mtSSU using the primers mtSSU1-mtSSU3R and mtSSU2R (Fedorenko *et al.* 2009, 2012).

The amplification was done using a Takara JP/TP600 PCR machine (Takara Bio Inc., Japan). One initial cycle of 5 min at 94 °C was followed by 30 cycles of the following steps: 30 seconds at 94 °C, 39 seconds at 57 °C and 1 min at 72 °C. Amplifications were ended with a final cycle at 72 °C for 10 min. PCR products were then sent to the sequencing facilities of the Genotech Cooperation, Seoul, South Korea, for cleaning and sequencing. The sequencing was carried out using the fluorescent marker BigDye and an ABI 3730xl sequencing machine (Applied Biosystems, Carlsbad, CA, USA).

The consensus sequence was aligned with all related species sequences retrieved from the GenBank database (Appendix). The consensus sequences were then deposited into GenBank under the accession numbers KY614390–KY614518. Phylogenetic analysis was performed using the ITS region and LSU gene of nrDNA and 12S SSU mtDNA sequences of 166 fungal taxa retrieved from the GenBank database and the 29 lichen-forming fungi investigated in this study. Sequence alignment was conducted in BioEdit and a phylogenetic tree was generated by the maximum parsimony (MP), minimum evolution (ME), and maximum likelihood (ML) analysis methods. Analyses were conducted using PAUP 4.0b10 on a Macintosh platform (Swofford 2003), and in Mega 5.0 (Tamura *et al.* 2011) with the number of bootstrap trials set to 1,000.

Our taxon sampling consists of 52 taxa of the Xanthorioideae (Fig. 1), 72 taxa of the Caloplacoideae (Fig. 2), 48 taxa of the Teloschistoideae (Fig. 3) and about 74 taxa of the Browlielloideae (Fig. 4) with *Brigantiae ferruginea* as outgroup (Appendix).

Totally 129 sequences on nrDNA and mtDNA are for the first time submitted to GenBank for 58 specimens of 29 taxa.

## RESULTS AND DISCUSSIONS

### *Current stage of the phylogenetic tree of the Teloschistaceae*

Phylogenetic trees of the teloschistoid, caloplacoid and xanthorioid lichens are presented in Figures 1–3 including respectively members of the sub-families Teloschistoideae, Caloplacoideae and Xanthorioideae only. In this case all genera, which were described and discussed before, are presented only by type species, while new groups, discussed in this paper are presented by larger number (between three and five) specimens/species.

The aim of the phylogenetic tree of the subfamily Brownlielloideae was also to show position of those taxa, which are in somewhat out position to main clades and subphylas of the subfamilies mentioned, while it includes

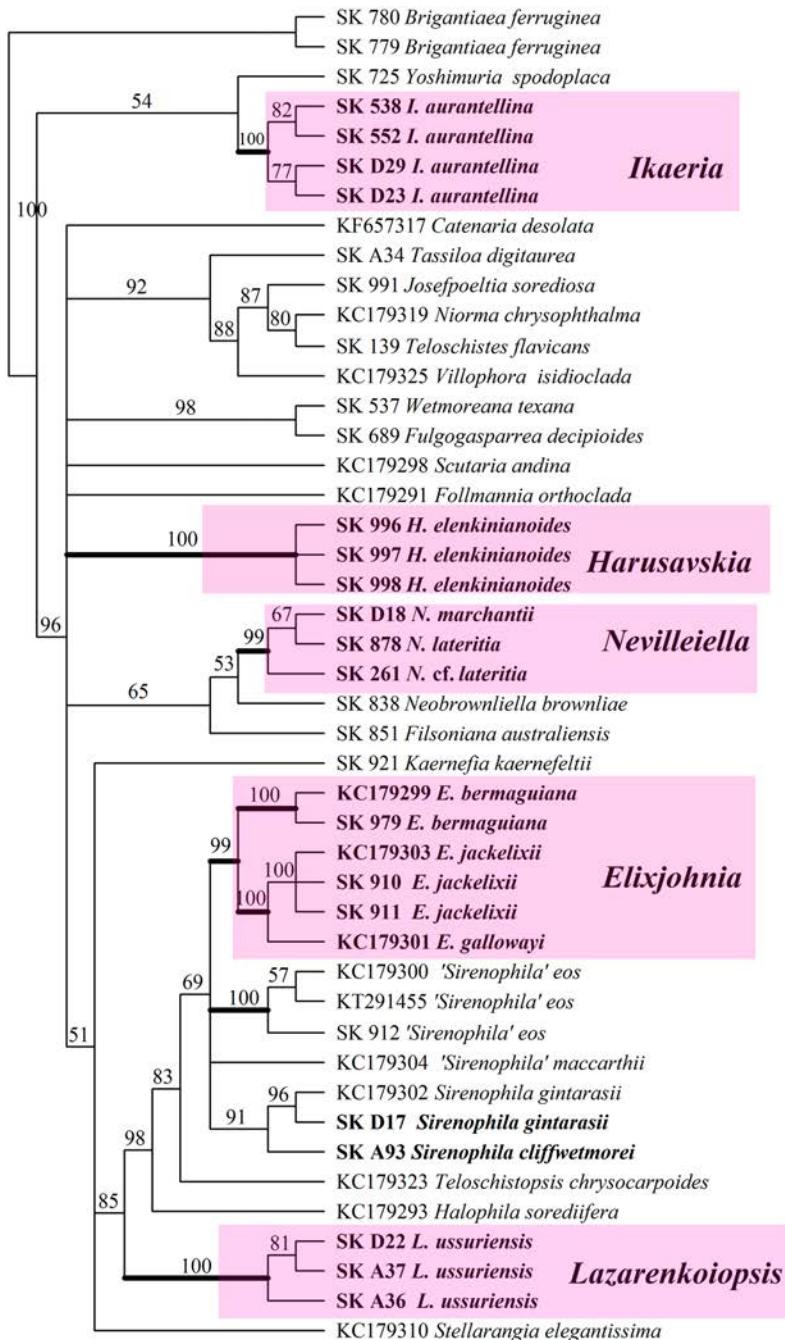


Fig. 1. Phylogenetic analysis of representatives of the subfamily Teloschistoideae after combined data set

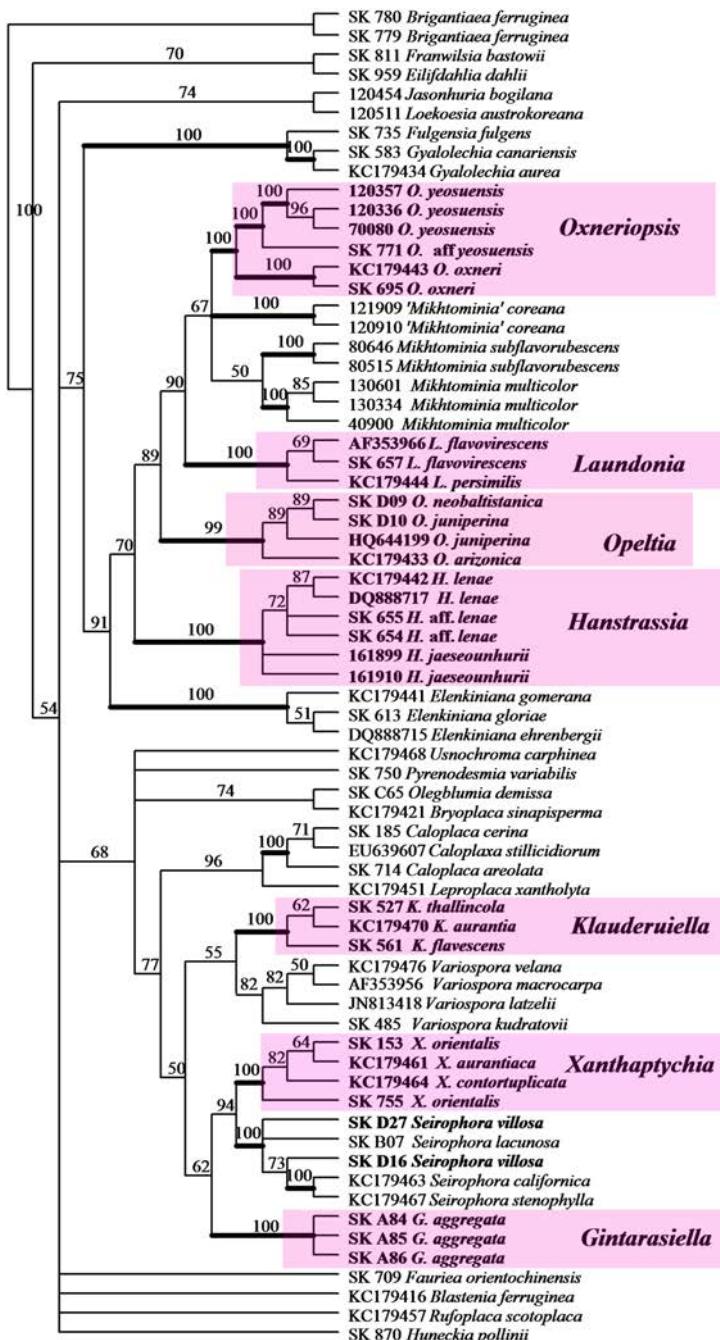


Fig. 2. Phylogenetic analysis of representatives of the subfamily Calopacoideae after combined data set

somewhat limited list of taxa of the Xanthorioideae, Caloplacoideae and Teloschistoideae. Main point of that three was to analyse the position of such genera as *Franwilsia*, *Eilifdahlia* and *Huneckia* from the Caloplacoideae, genera *Honeggeria*, *Schackletonia*, and *Solitaria* from the Xanthorioideae and some others being in somewhat out position to subfamilies mentioned.

In general from current phylogenetic tree of the Teloschistaceae we can make conclusion that position of the major part of genera within the subfamilies Teloschistoideae and Xanthorioideae is more or less stable. Only a few new monophyletic branches were added to these subfamilies (i.e. 5 genera to Teloschistoideae and 3 genera to Xanthorioideae).

Twenty-five monophyletic branches of the subfamily Teloschistoideae form single phylum, in which there are the following clades: the *Teloschistes* s.l. clade with 8 branches, the *Follmannia* s.l. clade with 4 branches, the *Filsoniana* s.l. clade with 3 branches, the *Sirenophila*–*Teloschistopsis*–*Halophila* clade with 5 or 6 branches, as well as two genera i.e. *Kaernefia* and the genus *Stellarangia*, which are positioning in sister position to the *Sirenophila*–*Teloschistopsis*–*Halophila* clade.

New branches, i.e. the newly proposed genus *Ikaeria* is positioning in sister position to the genus *Yoshimuria* in the *Teloschistes* s.l. clade, the newly proposed genera *Harusavskia* and *Nevilleiella* are in the *Filsoniana* s.l. clade, and two newly proposed genera *Elixjohnia* and *Lazarenkoiopsis* are positioned in the *Sirenophila*–*Teloschistopsis*–*Halophila* branch (Fig. 1).

The subfamily Caloplacoideae at current stage includes 2 large subphyla, i.e.: the *Gyalolechia* s.l. subphylum with 9 monophyletic groups belonging mainly to the *Mikhtomia* s.l. clade, and genera *Jasonhuria*, *Loekoesia* and *Gyalolechia* s. str. as outgroups to the *Mikhtomia* s.l. clade, and the *Caloplaca* s.l. subphylum with 14 monophyletic groups, which mainly belong to the *Caloplaca* s.l. clade, and three genera, i.e.: *Blastenia*, *Fauriea* and *Rufoplaca* forming the *Blastenia* s.l. clade.

The genera *Huneckia*, *Franwilsia*, and *Eilifdahlia* are positioned in the out position to the both subphyla of the Caloplacoideae mentioned.

The newly proposed genera *Hanstrassia*, *Laundonia*, *Opeltia*, and *Oxneriopsis*, are members of the *Mikhtomia* s.l. clade, while *Klauderuiella*, *Xantaptychia* and *Gintarasiella* – are members of the *Caloplaca* s.l. clade (Fig. 2).

There are two large subphyla within the subfamily Xanthorioideae (Fig. 3). Eighteen genera/monophyletic groups of this subfamily form the *Xanthoria* s.l. subphylum among which the genera *Flavoplaca* and *Calogaya* are positioned in the outermost position. Seventeen monophyletic groups form the *Xanthomendoza* s.l. subphylum among which *Cerothallia* and *Austroplaca* appeared to be in the outermost position. Additionally to these 35 branches mentioned above the genera *Schackletonia* and *Honeggeria* are positioned in out position to both subphyla mentioned above. Level of support of 34 monophyletic branches was discussed before (Kondratyuk et al. 2014c, 2015e). Three

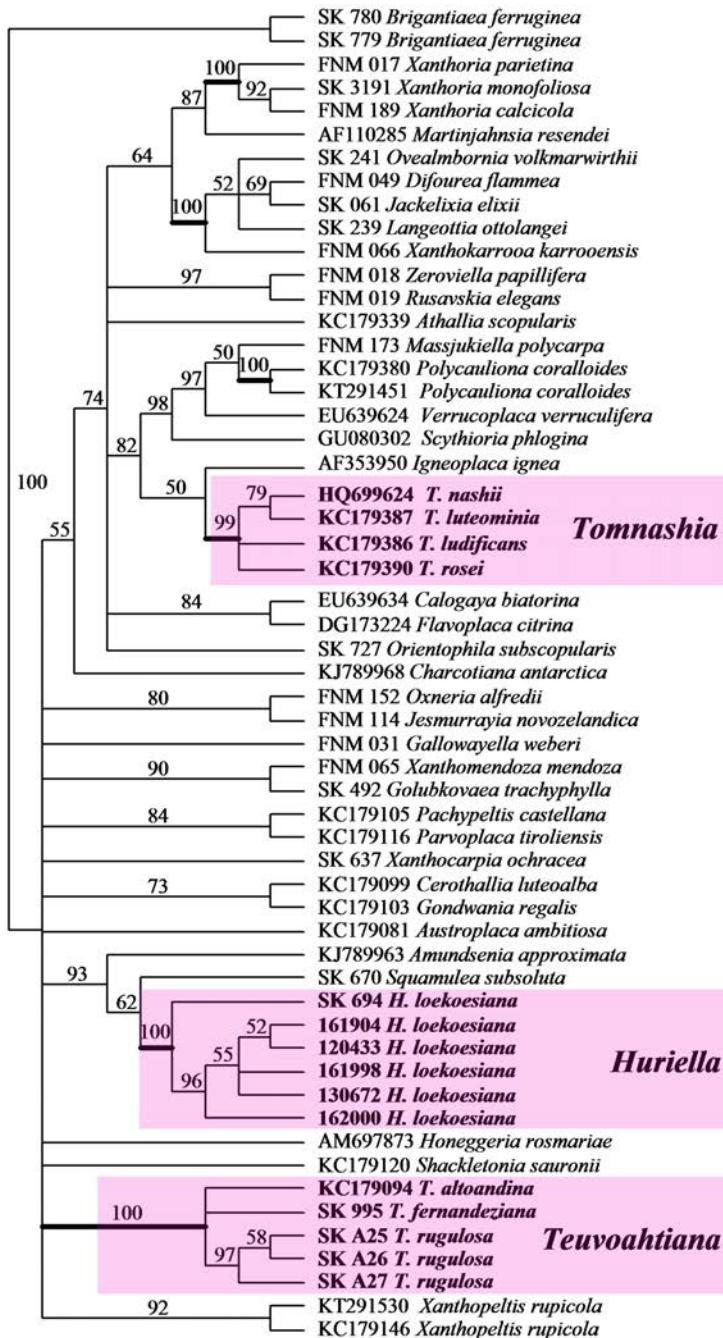


Fig. 3. Phylogenetic analysis of representatives of the subfamily Xanthorioideae after combined data set

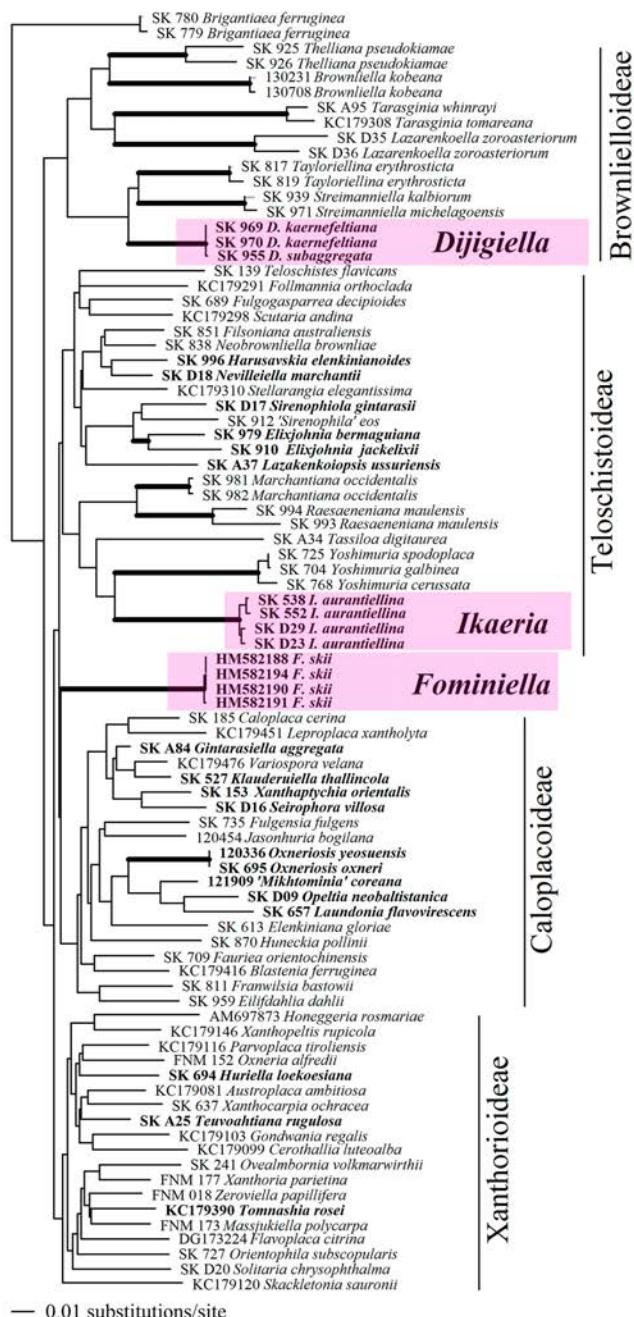


Fig. 4. Phylogenetic analysis of representatives of the subfamily Brownlielloideae after combined data set

groups, i.e.: *Huriella*, *Teuvoahtiana*, and *Tomnashia* are discussed below (in this paper). *Huriella* and *Teuvoahtiana* are members of the *Xanthomendoza* s.l. subphylum (i.e. of the *Xanthomendoza* s.l. clade and the *Xanthopeltis* s.l. clade, respectively), while *Tomnashia* is a member of the *Xanthoria* s.l. subphylum and the *Polycauliona* s.l. clade (Fig. 3).

The most unstable situation is with the subfamilies Brownlielloideae and Caloplacoideae and a number of taxa, which appeared to be in out position to all known subfamilies of the Teloschistaceae. Seven newly proposed genera are added to the subfamily Caloplacoideae (Fig. 3). The subfamily Brownlielloideae itself is almost with the same number of genera as it was in its original description (i.e. only one genus *Dijigiella* is added) (Fig. 4). However, the level of support of this branch is very low. The genus *Ikaeria* is positioned within the Caloplacoideae (compare Figs 1 and 4) and the genera *Fominiella* and *Solitaria* are in the out position to the subfamily Caloplacoideae, while the genus *Solitaria* was positioned in the subfamily Xanthorioideae before (Arup *et al.* 2013a, Kondratyuk *et al.* 2014c).

The new tendency that the subfamily Brownlielloideae will be divided into two or more separate subphylla with additional new members of this subphylum is found. However, new divisions of the Brownlielloideae can be observed only, when all monophyletic branches of the Teloschistaceae (as well as some still not analysed here) are included into analyses. In such case phylogenetic analysis is becoming more and more time consuming and in need of another computer facilities.

## DESCRIPTIONS OF TAXA

### *Dijigiella* S. Y. Kondr. et L. Lőkös, *gen. nov.*

MycoBank nr.: MB 819634.

*Similar to the genus Marchantiana of the Teloschistoideae, but differs in positioning in the Brownlielloideae of the Teloschistaceae.*

Type species: *Dijigiella kaernefeltiana* S. Y. Kondr.

Thallus small or indistinct, from endophloedal to corticolous, whitish grey to grey, continuous or more or less cracked to areolate, mainly distinct owing to bright yellow or yellow-orange apothecia, in some taxa densely aggregated in large groups.

Apothecia lecanorine to zeorine or biatorine, disc yellow to yellow-orange, dull brownish yellow to dull yellow-orange; true excipulum scleroplectenchymatous or "textura intricata" with matrix to paraplectenchymatous with ma-

trix, paraphyses broom-like, richly branched in the uppermost portion; ascospores 8-spored; ascospores bipolar, hyaline.

**Ecology:** Growing on bark of trees.

**Distribution and species content:** It hitherto contains two Australian taxa, while several taxa will probably be added in future.

**Etymology:** It is named after the well-known New Zealand lichenologist David J. Galloway in recognition of his contribution to the lichen flora of the Southern Hemisphere.

**Taxonomic notes:** After phylogenetic analysis this genus is positioned in the subfamily Brownlielloideae of the Teloschistaceae. From molecular data hitherto available the genus *Dijigiella* includes two species, i.e. *D. kaernefeltiana* and *D. subaggregata*.

A number of sorediate Australian taxa are still in progress of extracting DNA and getting sequences, and hopefully some of them will be added to this genus.

***Dijigiella kaernefeltiana* S. Y. Kondr., spec. nova**  
(Fig. 5)

MycoBank nr.: MB 819635.

*Similar to Marchantiana occidentalis, but differs in having greyish-greenish or greyish-whitish thallus, in having zeorine or lecanorine apothecia with bright orange disc and true exciple, in having scleroplectenchymatous or "textura intricata" true exciple, in having somewhat smaller ascospores, as well as in the lack of ascomatic acid, ascomatatic and 7-O-methylascomatatic acid.*

**Type:** Western Australia: Wicherina, E. of Geraldton, on the road to Mullewa, growing on dry shrubs together with *Streimanniella michelagoensis*, and species of the genera *Caloplaca*, *Buellia*, *Teloschistes* and *Candelariella*. Lat.: 28° 41' 46.0" S; Long.: 114° 56' 40.1" E. Coll.: Kärnefelt, I. (20042002), 07.01.2004 (LD1238056 sub *Streimanniella michelagoensis* – holotype).

Thallus of very small to 2–3(–5) mm across spots scattered among other lichens, i.e. *Streimanniella michelagoensis* thalli, greyish-whitish, from very thin consisting of very small ca 0.1–0.2 mm across irregular microareoles to rather thick among apothecia, and with rather larger size of "areoles" to 0.4–0.5 mm across, and easily distinct owing to bright yellow or yellow-orange discs or own margin of apothecia. Hypothallus not observed.

Apothecia (0.15–)0.3–0.6 mm diam. and to 0.18–0.2 mm thick in section; lecanorine or zeorine, if lecanorine with grey or grey-greenish thalline margin and yellow-orange or dull yellow-orange disc, if zeorine, own margin concolourous with disc dull yellow-orange observed; usually regularly



Fig. 5. *Dijigiella kaernefeltiana* (holotype), general habit. Scale 0.5 mm. (Photo of S. Kondratyuk)

rounded and distant, rarely aggregated and somewhat irregular, distinctly attenuated at the basis and uplifted above substrate level; in section true exciple to (10–)20–30 µm thick in the uppermost and the lower lateral portions, scleroplectenchymatous or “textura intricata” with matrix and radiating hyphae luminas, (10–)20–30 µm thick in basal portions paraplectenchymatous with matrix(?), cell lumina to 4 µm diam. (or cells to 7–9 µm across); thalline exciple to 50–70 µm thick, cortical layer absent or only of algal plectenchyma; hymenium to 50–60 µm high; paraphyses very richly branched at the tips, broom-like, to 3–4 µm diam. at the tips; subhymenium (50–)70–80 µm thick, straw yellow or to straw-brownish; ascospores 8-spored, with (2–4–)8 adult bipolar and young ascospores in the same ascus; ascospores bifusiform with somewhat attenuated ends, narrowly ellipsoid, (7.5–)8–13(–14) × 4.5–6 µm in water, and (7–)8–13(–15) × (4–)5–7(–8) µm in K, (in K becoming more or less widely ellipsoid), ascospores septum (2–)2.5–4 µm wide in water and (2–)3–6 µm wide in K.

**Chemistry:** Thalline cortical layer and algal plectenchyma of thalline exciple, and epihymenium K+ dirty crimson purple in places.

**Etymology:** Species is named after the well-known Swedish lichenologist Ingvar Kärnefelt (Lund, Sweden) in recognition of his contribution to lichenology, and who has kindly provided for us an opportunity to participate in a field trip to Australia in 2003–2004 and provided his earlier collections in our disposal.

**Distribution:** So far known from type locality in Western Australia.

**Taxonomic notes:** *Dijigiella kaernefeltiana* is similar to *Marchantiana occidentalis* (Elix, S. Y. Kondr. et Kärnefelt) S. Y. Kondr., Kärnefelt, A. Thell, Elix, J. Kim, A. S. Kondr. et Hur after having dull greenish-greyish or greenish greyish-whitish thallus, measurements of ascospores and having paraphyses richly branched in the upper portion and densely twisted above ascii, but differs in having greyish-greenish or greyish-whitish thallus (vs. brown to dark brown, greenish areoles immersed in bark), and in having zeorine or lecanorine apothecia with bright orange disc and true exciple (vs. biatorine), in having scleroplectenchymatous or “textura intricata” true exciple (vs. pseudo-prosoplectenchymatous in lateral portion and paraplectenchymatous in basal portion), as well as in the lack of ascomatic acid, ascomatatic and 7-O-methyl-ascamatatic acid, and in having somewhat shorter ascospores (8–13 × 4.5–6 µm vs. (9–)11–13(–14) × (4–)4.5–6(–7) µm).

*Dijigiella kaernefeltiana* often growing side by side with *Streimanniella michelagoensis* from which it differs in having lighter and thicker (with somewhat subconvex areoles) thallus (vs. dark brownish-greenish grey to dark brownish grey, smooth or slightly cracked to indistinct), in having much larger and brighter light yellow-orange apothecia (0.3–0.6 mm vs. ca 0.2–0.3 mm diam., lecanorine apothecia with blackish grey or greenish-greyish brown thalline

margin and somewhat dull yellow own margin and dull yellowish-blackish or yellowish, bluish-greyish to dull dark brownish orange disc), in having always hyaline ascospores of the same measurements (vs. bipolar ascospores becoming greyish or greyish-blackish at overmature, somewhat similar to *Rinodina* type ascospores), as well as in the lack of neochloroemodin (see also Kondratyuk *et al.* 2009). In type collection there is one more crustose member of the Teloschistaceae with rather thick to worthy, grey thallus. Unfortunately it was in insufficient amount for the further study. However, *Dijigiella kaernefeltiana* differs from this taxon in having lighter brownish-whitish-greyish thallus and in having much brighter dull yellow or dull yellow-orange disc and own margin of apothecia.

*Dijigiella kaernefeltiana* is similar to '*Caloplaca*' *ulcerosa* Coppins et P. James, but differs in having K<sup>+</sup> cortical layer of thallus or thalline exciple and algal plectenchyma in both as well as in the lack of whitish or greyish soredia, and in the lack of pustule-like immersed/ulcer-like soralia. Unfortunately status of '*Caloplaca*' *ulcerosa* is still uncertain from molecular point of view. After ITS phylogeny it is positioned in the Caloplaceoideae, but data on nrLSU and mtSSU are still missing for this taxon.

***Dijigiella subaggregata* S. Y. Kondr. et Kärnefelt, spec. nova  
(Fig. 6)**

MycoBank nr.: MB 819636.

*Similar to Gintarasiella aggregata, but differs in having thinner thallus, in having smaller and plane apothecia, in having hymenium and subhymenium without oil, in having smaller ascospores, in having wider ascospore septum, and in corticolous habit.*

Type: Australia: Victoria, Camperdown, in the central part of town, Wpt. 64, on bark of *Ulmus*, growing together with *Caloplaca hannesheretii*, and species of the genera *Opegrapha*, *Hyperphyscia*, *Phaeophyscia*, *Lecanora* and *Physcia*. Lat.: 38° 14.37' S; Long.: 143° 08.57' E. Coll.: Kärnefelt, E. I. (996401), 21.01.1999 (LD 1275974 – holotype).

Thallus (0.25–)0.5–1.5(–2.5) cm across, usually indistinct, endophloedal or very thin, whitish or whitish-greyish, well contrasting to darker aggregated apothecia in the centre of thallus, becoming somewhat greyish or dirty white among apothecia or to whitish yellow in peripheral zone probably yellow tinge owing to numerous young apothecia; seen as very dense aggregations of dull yellow or dull brownish yellow apothecia. Hypothallus absent.

Apothecia (0.15–)0.3–0.6 mm diam./across, rather small but easily distinct because often aggregated in very dense aggregations (to 200 apothecia



Fig. 6. *Dijigiella subaggregata* (holotype), general habit. Scale 1 mm. (Photo of S. Kondratyuk)

per thallus), initially regularly rounded, immersed into substrate, then becoming sessile and aggregated in large aggregations, biatorine with lighter bright or pale yellow own margin to 40(–50) µm thick, at initial stages well distinct, somewhat arising disc level, later irregularly developed to disappearing in densely pressed, irregular apothecia; disc more or less plane, dull dark yellow to dull brownish yellow, matt, without pruina; in section biatorine to zeorine, where thalline exciple to 30–40 µm thick developed only on underside; true exciple to 20(–40) µm thick in the uppermost lateral and to 30 µm thick in the lower lateral portion, and not developed or only to 7 µm thick in basal portion, more or less paraplectenchymatous; hymenium to 55–60 µm high; epiphymenium to 15 µm thick, dark orange, paraphyses with 1 or 2 uppermost cells distinctly widened to 3–4(–5) µm diam. towards the tips; subhymenium to 20–30(–40) µm thick, hyaline, without oil; asci 8-spored, often very variegating in measurements within the same ascus; ascospores rather small, but with rounded ends, mainly widely ellipsoid, widened at the septum, 7–10(–12) × (4–)5–6(–6.5) µm in water and (8–)9–13(–15) × (5–)6–8(–10) µm in K, septum (2–)3–4(–4.5) µm wide in water and (3–)3.5–5(–7) µm wide in K.

Chemistry: Epiphymenium and outer layers of the true exciple K+ crimson purple, somewhat washing out in solution.

Etymology: It is named after similarities to *Gintarasiella aggregata*.

Distribution: So far known only from type collection, Victoria, southern Australia.

Taxonomic notes: *Dijigiella subaggregata* is similar to *Gintarasiella aggregata*, after having thallus dominated almost entirely by clustered apothecia, but differs in having thinner thallus (vs. rather pulvinate), in having smaller and plane apothecia (0.3–0.6 mm vs. 0.3–1 mm wide, becoming convex in the oldest apothecia), in having hymenium and subhymenium without oil (vs. hymenium and subhymenium especially heavily inspersed with oil droplets or irregular oil aggregations), in having smaller ascospores (7–10 × 5–6 µm vs. 10–13.5 × 5–6 µm), in having wider ascospore septum (3–4 µm vs. 2–3 µm wide), and in corticolous habit.

*Dijigiella subaggregata* is similar to *Athallia cerinelloides* (Erichsen) Arup, Frödén et Søchting, but differs in having endophloedal or very indistinct thallus (vs. light grey, film-like), in having apothecia densely aggregated in large groups, and in having shorter ascospores (7–10 × 5–6 µm vs. 9–13 × 5–7 µm).

*Dijigiella subaggregata* is similar to *Cerothallia luteoalba* (Turner) Arup, Frödén et Søchting, but differs in having wider ascospore septum (3–4 µm vs. 1–1.5 µm wide), while ascospores are almost the same (7–10 × 5–6 µm vs. 8–12 × 3–6 µm).

*Dijigiella subaggregata* is similar to '*Caloplaca*' *aegatica* Giralt, Nimis et Poelt (see also comments below the genus *Ikaeria*), but differs in having smaller as-

cospores ( $7–10 \times 5–6 \mu\text{m}$  vs.  $10–15 \times 7–9 \mu\text{m}$ ), and narrower ascospore septum ( $3–4 \mu\text{m}$  vs.  $5–8 \mu\text{m}$  wide), in having endophloedal thallus (vs. greyish, dirty grey, grey-brown), and in having lighter apothecia (vs. orange to dark orange zeorine, with bright golden margin and disappearing thalline margin).

*Dijigiella subaggregata* is similar to *Fominiella skii* (Khodos., Vondrák et Šoun) S. Y. Kondr., D. Upreti et J.-S. Hur, but differs in having lecanorine to zeorine apothecia (vs. zeorine to biatorine) and larger (0.3–0.6 mm vs. 0.2–0.4 mm diam.) apothecia, in having smaller ascospores ( $7–10 \times 5–6 \mu\text{m}$  vs.  $9–11 \times 4.5–5.5 \mu\text{m}$ ) and narrower ascospore septum ( $3–4 \mu\text{m}$  vs.  $4.5–5.7 \mu\text{m}$  wide), as well as in distribution (vs. the Canary Islands).

### *Elixjohnia* S. Y. Kondr. et J.-S. Hur, *gen. nov.*

MycoBank nr.: MB 819637.

Similar to genus *Sirenophila* of the subfamily *Teloschistoideae*, but forming separate robust monophyletic branch.

Type species: *Elixjohnia jackelixii* (S. Y. Kondr., Kärnefelt et A. Thell) S. Y. Kondr. et J.-S. Hur.

Thallus distinctly zoned initially with brownish or dirty greenish yellow sterile circles of dull, smooth, continuous peripheral zones or without zonations; usually very thick towards the centre or evenly continuous; smooth and entire or distinctly areolate, whitish, yellowish or greyish to bright red or reddish orange; upper surface cracked and eroded, sometimes with isidia-like structures; medulla often visible through the numerous cracks. Protothallus well developed or absent. Apothecia common, small, zeorine (in some species at least initially), lecanorine or biatorine; disc brownish orange or yellowish brown, with a reddish tinge to dull orange, dark orange-red or scarlet; true exciple scleroplectenchyomatous; oil cells in paraphyses of *bermaguiana*-type (see Kondratyuk et al. 2007, 2009, 2012); asci 8-spored, but 2–4–6 mature bipolar ascospores together with aborted spores often present; ascospores narrowly to broadly ellipsoid; conidia broad ellipsoid to broad bacilliform, (2–)2–3 × 1.2–1.7  $\mu\text{m}$ .

Chemistry: It contains parietin (major).

Ecology: Species of the genus grow on exposed coastal rocks (quartzite, granite, basalt and dolerite), at or above the high tide level often together with each other (i.e. *E. jackelixii* and *E. gallowayi* often growing side by side), as well as together with species of the genera *Tarasginia whinrayi* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr., Kärnefelt, A. Thell, Elix, J. Kim, A. S. Kondr. et Hur, *T. tomareana* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr., Kärnefelt, A. Thell, Elix, J. Kim, A. S. Kondr. et Hur, *Sirenophila eos* (S. Y. Kondr. et Kärnefelt) Arup, Frödén et Søchting, *Jackelixia ligulata* (Körb.) S. Y. Kondr., Fedorenko, S. Sten-

roos, Kärnefelt et A. Thell, *Gondwania sublobulata* (Nyl.) S. Y. Kondr., Kärnefelt, Elix, A. Thell, J. Kim, M.-H. Jeong, N.-N. Yu, A. S. Kondratuk et J.-S. Hur, *H. cribrosa* (Hue) Søchting, Frödén et Arup and *Caloplaca conranii* S. Y. Kondr. et Kärnefelt.

Distribution: Species of this genus widely distributed in southern and southeastern Australia, Tasmania and New Zealand.

Etymology: Genus is named after the well-known Australian lichenologist and chemist Elix John Alan ('Jack') in recognition of his contribution to lichenology.

Taxonomic notes: After combined phylogenetic analysis the genus *Elixjohnia* is a member of the large *Sirenophila-Teloschistopsis-Halophila* subclade of the Teloschistoideae. This branch includes three species, i.e. *E. bermaguiana*, *E. jackelixii* and *E. gallowayi* after molecular data hitherto available.

Since 2013, when genus *Sirenophila* was described, situation with taxa of this branch has been problematic, because the genus *Sirenophila* was in fact segregated only on the basis of ITS phylogeny. Only ITS data were provided for the rather rare Australian taxon *S. gintarasii* (S. Y. Kondr. et Kärnefelt) Arup, Frödén et Søchting (Arup *et al.* 2013a), type species of the genus *Sirenophila*.

We were able to get 28S nrLSU and 12S mtSSU data for this taxon from isotype kept in KW-L and they are presented here (Appendix).

From phylogenetic analysis (Fig. 1) it is seen that the genus *Sirenophila* s. str. includes only *S. gintarasii* and *S. cliffwetmorei* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr., recently combines to this genus (Kondratyuk *et al.* 2015e). Status of *Sirenophila eos*, as well as of *S. maccarthii* is still questionable and should be repeatedly analysed, when more molecular data on this phylum will be available.

Two species of this genus, i.e.: *E. jackelixii* and *E. bermaguiana* were compared with *Tomnashia rosei* (see below), which belongs to the Xanthorioideae. Furthermore *E. bermaguiana* was compared also with *Caloplaca inconnexa* (Nyl.) Zahlbr. (supposedly the Xanthorioideae) and *Caloplaca nubigena* (Kremp.) Dalla Torre et Sarnth. (supposedly Caloplacoideae). Molecular data on *Caloplaca inconnexa* and *C. nubigena* are still missing. In original description *E. gallowayi* was compared only with *Sirenophila eos* (Kondratyuk *et al.* 2007).

### *Fominiella* S. Y. Kondr., D. Upreti et J.-S. Hur, *gen. nov.*

MycoBank nr.: MB 819638.

*Similar to genera Athallia of the Xanthorioideae, but differs in having different measurements of ascospores as well as in positioning in distant position from the Teloschistoideae, as well as in out position to both subfamilies Caloplacoideae and Teloschistoideae.*

Type species: *Fominiella tenerifensis* S. Y. Kondr., Kärnefelt, A. Thell et T. Feuerer.

Thallus thin, film-like, inconspicuous, whitish to light grey or pale yellowish. Hypothallus not developed.

Apothecia zeorine to lecanorine or biatorine, disc yellow-orange, own margin yellow, always paler than disc; thalline exciple white, greyish to yellow-white, soon disappearing or permanent, true exciple leptodermatous paraplectenchymatous; asci 8-spored; ascospores polarilocular, hyaline.

Chemistry: Parietin (major), emodin, parietinic acid, fallacial and teloschistin (trace).

**Etymology:** This genus is named after the well-known Ukrainian botanist Alexandr Valyljovych Fomin, specialists on cryptogamic plants, who is founder of several botanical institutions and periodicals in Georgia and Ukraine including recent institution known as M. H. Kholodny Institute of Botany, National Academy of Sciences of Ukraine.

**Distribution and species diversity:** So far genus includes two species, one of which, i.e.: *Fominiella skii* known from arid and semiarid conditions of southern and southeastern Europe and the Near East of Asia, and the second (*F. tenerifensis*) is known only from the Canary Islands.

**Taxonomic notes and phylogenetic affiliations:** After combined phylogenetic analysis the genus *Fominiella* is positioned in out position to all known subfamilies (Fig. 4). The genus includes two species, i.e.: *F. tenerifensis* and *F. skii* after molecular data hitherto available.

In original description (Vondrák *et al.* 2011) *Fominiella skii* was already compared with members of the following genera of the subfamily Xanthorioideae, i.e.: *Athallia* Arup, Frödén et Søchting (i.e.: *A. cerinelloides* (Erichsen) Arup, Frödén et Søchting), and *Xanthocarpia* (i.e.: *X. raesaenae* (Bredkina) S. Y. Kondr.), with which it is similar in having very reduced thallus. Reduction of the thallus has occurred independently several times in the evolution of the Teloschistaceae (Kärnefelt 1989; Vondrák *et al.* 2011).

On the basis of ITS phylogeny *Fominiella skii* was also included into the genus *Athallia* (Arup *et al.* 2013a). However, providing data on nrLSU and mtSSU has allowed showing its positioning in rather distant position from Xanthorioideae and being in somewhat out position to both subfamilies Calopacoideae and Teloschistoideae.

ITS sequence of the type specimen of *Fominiella skii* (HM582191) provided in original paper (Vondrák *et al.* 2011) is included into analysis. However, 28S nr LSU and 12S mtSSU were obtained from material of *F. tenerifensis* within this study only. Unfortunately we were not able to get ITS data on material investigated, while a number of special attempts were done. So this genus is in

urgent need of confirmation by molecular data (data on three or more genes) obtained from the same specimen (of the same species).

*Fominiella tenerifensis* S. Y. Kondr., Kärnefelt, A. Thell et T. Feuerer,  
*spec. nova*

MycoBank nr.: MB 819639.

*Similar to Fominiella skii, but differs in having well-developed thalline exciple, in having longer and wider ascospores, and in the lack of true exciple in lateral portion.*

Type: Spain, Canary Islands, Tenerife Island, Santiago del Teide, on bark of *Spartocytisus* shrubs. Lat.: 28° 17.631' N; Long.: 16° 48.989' W. Alt.: 920 m a.s.l. Coll.: Kondratyuk, S. Y. (20917), 15.01.2009 (KW-L – holotype, LD, BP, B – isotypes).

Thallus thin, film-like, inconspicuous or visible only around apothecia, whitish, light grey or pale yellowish. Hypothallus not developed.

Apothecia 0.2–0.35 mm diam. and to 0.12 mm thick in section, lecanorine, very small and usually aggregated in groups, true exciple not present in the uppermost lateral and lower lateral portions, to 10–15 µm thick in basal portion, leptodermatous paraplectenchymatous, cell lumina 3–5 µm diam./across; thalline exciple 50–60 µm thick with cortical layer to 15–20 µm thick, paraplectenchymatous, K–; hymenium 70 µm high; epihymenium K+ purple; paraphyses almost not widened towards the tips, with uppermost cells to 3–3.5 µm diam. in water (and to 2–3 µm diam. in K), richly broom-like branched (better seen in K), with oil droplets, but very indistinct or badly seen owing to small measurements, to 2–3 µm diam. (better seen in K); subhymenium 20–30 µm thick; asci 8-spored; ascospores elongated ellipsoid, 12–13(–15) × (5.5–)6–7 µm in water and 12–14(–15) × (5–)6–7 µm in K; septum 5–6(–7) µm wide in water and (6–)7–8 µm wide in K.

Chemistry: Epihymenium and outermost portions of true exciple K+ purple.

Ecology: It grows on barks of twigs of shrubs often associated with *Ikaeria aurantiellina*.

Etymology: It is named after type collection, Tenerife Island of Spanish Canary Islands.

Distribution: It is so far known from several places within the same Tenerife Island, Spain, Canary Islands.

Taxonomic notes: *Fominiella tenerifensis* is similar to *F. skii*, but differs in having well-developed thalline exciple, in having longer and wider ascospores (12–13(–15) × (5.5–)6–7 µm vs. (7.5–)9–11(–12.8) × (3.5–)4.6–5.9(–6.5)

$\mu\text{m}$ ) (while septum is almost the same (5–6(–7)  $\mu\text{m}$  vs. (3.5–)4.3–5.7(–6.5)  $\mu\text{m}$ )), and in the lack of true exciple in lateral portion.

*Fominiella tenerifensis* is similar to *Athallia cerinelloides* (Erichsen) Arup, Frödén et Søchting, but differs in having wider ascospore septum (5–6(–7)  $\mu\text{m}$  vs. 3–4  $\mu\text{m}$  wide), while ascospores are almost the same (12–13(–15)  $\times$  (5.5–)6–7  $\mu\text{m}$  vs. 9–13  $\times$  5–7  $\mu\text{m}$ ).

***Gintarasiella* S. Y. Kondr. et J.-S. Hur, *gen. nov.***

MycoBank nr.: MB 819640.

*The genus is characterised by a unique combination of the following characters among representatives of the subfamily Teloschistoideae, i.e.: pulvinate habit, with densely clustered apothecia that essentially obscure the thallus, a very densely inspersed hymenium and subhymenium and relatively small ascospores.*

Type species: *Gintarasiella aggregata* (Kantvilas et S. Y. Kondr.) S. Y. Kondr. et J.-S. Hur.

Thallus crustose, areolate, forming pulvinate patches, dominated almost entirely by clustered apothecia, yellow-orange. Apothecia from zeorine to biatorine, disc deep orange, somewhat darker than thallus, own margin colourous with disc; true exciple paraplectenchymatous in basal portion and composed of radiating more or less parallel hyphae in the lateral portions; hymenium and subhymenium heavily inspersed with oil droplets or irregular oil aggregations; asci 8-spored; ascospores polarilocular, ellipsoid, hyaline. Conidiomata and conidia not observed.

Chemistry: Thallus and all elements of apothecia K+ crimson purple.

Etymology: This genus is named after our friend and colleague, Dr Gintaras Kantvilas (Hobart, Tasmania), in acknowledgement of his enormous contributions to the taxonomy of the Tasmanian and Australian lichens, especially.

Distribution and species diversity: So far the genus includes only type species, which known from limestone outcrops of Kangaroo Island, Southern Australia.

Taxonomic notes and phylogenetic affiliations: After separate and combined phylogenetic analysis the genus *Gintarasiella* belongs to the subfamily Teloschistoideae, where it forms separate robust branch within the *Sirenophilina-Teloschistopsis-Halophila* subphylum as the most separate outgroup to this subphylum. From molecular data hitherto available the genus *Gintarasiella* includes only type species *G. aggregata*.

In original description the type species of this monotypic genus was already compared with members of the following genera of the subfam-

ily Xanthorioideae, i.e.: *Flavoplaca* (i.e.: *F. mereschkowskiana* (S. Y. Kondr. et Kärnefelt) Arup, Frödén et Søchting), *Cerothallia* (i.e. *C. yorkensis* (S. Y. Kondr. et Kärnefelt) Arup, Frödén et Søchting), and *Xanthocarpia* (i.e.: *X. jerramungupensis* (S. Y. Kondr., Kärnefelt et Elix) S. Y. Kondr., Kärnefelt, A. Thell, Elix, J. Kim, A. S. Kondr. et J.-S. Hur) (Kantvilas 2016).

From morphological point of view after having inspersed hymenium and subhymenium it can be also compared with the members of the genus *Franwilia* (subfamily Caloplacoideae), as well as after having pulvinate thallus with some *Calogaya* species (i.e. *C. lobulata* (Flörke) Arup, Frödén et Søchting known from bark of introduced trees in Tasmania) of the Xanthorioideae. However, it differs in having smaller ascospores and different chemistry, as well as in the positioning in the subfamily Teloschistoideae of the Teloschistaceae.

***Hanstrassia* S. Y. Kondr., gen. nov.**

Mycobank nr.: MB 819641.

*Similar to genus Elenkiniana, but differs in having mainly areolate and sorediate thallus, and in positioning in separate robust monophyletic branch within the Mikhtomia s.l. clade of the Caloplacoideae.*

Type species: *Hanstrassia lenae* (Søchting et G. Figueras) S. Y. Kondr.

Thallus saxicolous or terricolous, areolate, effigurate or only weakly lobate at the margin, yellowish grey, pale yellow, ochre to orange or brownish yellow, often with a whitish pruina; areoles thick; soralia scarce or numerous, labriform, usually marginal, yellowish to ochre, usually brighter than the thallus; cortical layer pseudoprosoplectenchymatous or scleroplectenchymatous; medulla dense, consisting of interwoven hyphae, without a clear orientation.

Apothecia lecanorine to zeorine, few and dispersed; disc flat, later sometimes rather convex, deep orange to brownish, sometimes with whitish pruina at initial stages; true exciple prosoplectenchymatous; asci 8-spored; ascospores polarilocular. Conidia bacilliform,  $3\text{--}3.5 \times 1\text{--}1.25 \mu\text{m}$ .

Chemistry: Anthraquinone fragilin and depsidones caloploicin and vicanicin (major), and anthraquinones parietin, emodin, and depsidone isofulgidin in small concentrations.

Ecology: It grows on limestone and calcareous schist of vertical cliffs in dry continental conditions or on sandy soil in mountainous deserts of the Asian continent.

Etymology: It is named after the well-known Estonian lichenologist Hans Trass (02.05.1928–14.02.2017) in recognition of his contribution to lichenology and to geobotany, as well as for his contribution to Eurasian lichen flora.

**Distribution and species diversity:** Type species *Hanstrassia lenae* is known from northeastern Asia from the Altai Region in Siberia and the Mongolian People's Republic to Yakutia, while *H. jaeseounhurii* described here is so far known only from the type locality of China. Taxonomic status of material previously identified as *H. lenae* from Russia is in need of further clarifying.

**Taxonomic notes and phylogenetic affinities:** After combined phylogenetic analysis the genus *Hanstrassia* is positioned in separate robust monophyletic branch within the *Mikhtomia* s.l. branch, while after ITS phylogeny it was placed in the genus *Gyalolechia* s.l. (Arup et al. 2013a), and to the genus *Elenkiniana* (Kondratyuk et al. 2014c).

It is interesting to note, that in the first ITS phylogenetic tree published in the paper with original description of *H. lenae* it was positioned in sister position to the *Laundonia flavovirescens* (Søchting and Figueras 2007), and now the same position is confirmed by three gene phylogeny (Fig. 2).

***Hanstrassia jaeseounhurii* S. Y. Kondr., Ch.-H. Park et L. Lőkös, spec. nova  
(Figs 7–8)**

MycoBank nr.: MB 819642.

*Similar to Hanstrassia lenae, but differs in having much larger thalline areoles, in having only areolate thallus, in having very scarce soredia and in having scleroplectenchymatous cortical layer of thallus.*

Type: China. Qinghai Province, between Jingtai and Zhongwei (close to Yellow River), along 201 Provincial Road, on soil, growing together with *Candelariella* sp. Lat.: 37° 25' 19.78" N; Long.: 104° 34' 50.05" E; Alt.: 1,686 m a.s.l. Coll.: Hur, J.-S. and Park, Ch.-H. (CH 160016), 17.06.2016 (KoLRI 041972 – holotype); the same locality and collectors, on soil, growing together with *Candelariella* sp., *Megaspora* sp. and *Catapyrenium* sp. (CH 160015) (KoLRI 041972 sub *Candelariella* – isotype).

Thallus to 2–3(–5) cm across, distinctly areolate, without lobes in peripheral position, brownish green-yellow, with whitish pruina towards the peripheral portion; areoles distant, scattered and regularly rounded, to (0.2–)0.5–1 mm diam./across in peripheral portion, and much larger to (0.5–)1–3(–4) mm across, in the centre, seem to be overlapping by marginal portions; sorediate. In section thalline areoles (0.3–)0.4–0.8 mm thick, while together with slightly brownish medullar layer (may be including overgrowing old portions?) to 1–1.6 mm thick; cortical layer 50–75(–100) µm thick, yellowish in upper portion or throughout, scleroplectenchymatous, cell lumina to 3–4(–5) µm diam., often with oxalate crystals to (5–)7–15(–20)[–40] µm across; K+ purple, while

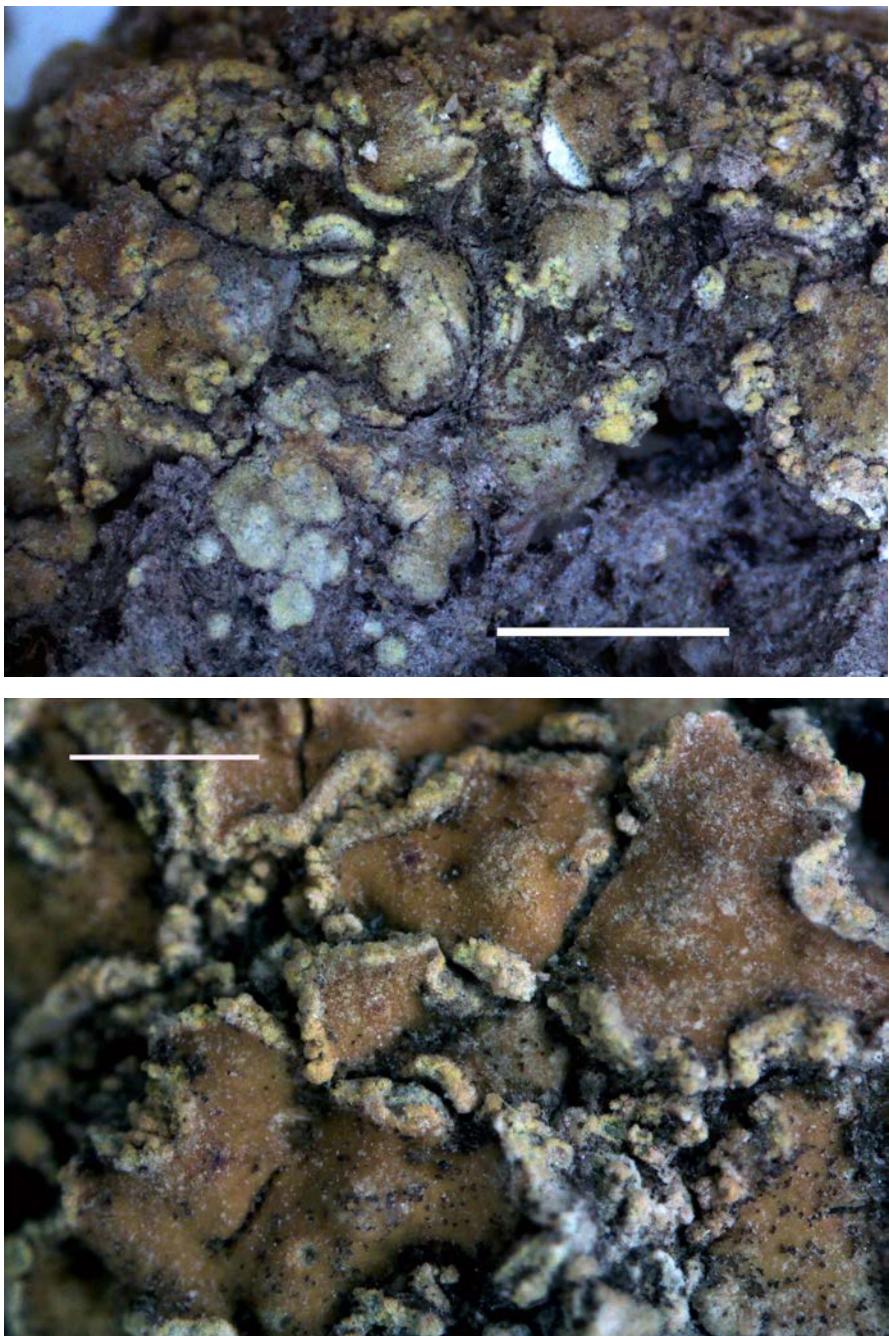


Fig. 7. *Hanstrassia jaeseounhurii* (holotype), general habit. Scale 2 mm (top) and 1 mm (bottom). (Photo of S. Kondratyuk)

portions with crystals K+ blackish purple in places; epinecral portions (better seen in K) to 20(–25) µm thick, in places dirty brownish; algal layer 50–70 (–150) µm thick, in places consisting of clusters, not entire, algal cells to 10–22.5 µm diam./ across; medulla white, compact (without hollows) to 0.4–1.4 mm thick, where upper portion to 0.4–0.5 mm thick below algal layer purely white, while lower portion (to 0.7–1 mm thick) straw yellowish or somewhat pale brownish, probably owing to dust/soil particles, usually hardly seen, owing to numerous crystals not dissolving in K. Soralia marginal, somewhat lip-like; soredia/schizidia usually rather scarce, 0.06–0.12(–0.2) mm diam./across, or rarely in larger aggregations, irregular, light yellow or lemon yellow (well contrasting to darker thallus). Apothecia, conidiomata, and conidia unknown.

**Chemistry:** Cortical layer of thallus K+ purple, while portions with numerous crystals K+ to blackish purple in places.

**Ecology:** Growing on soil among mosses and together with species of the genera *Candelariella*, *Placopyrenium*, *Collema* and *Megaspora*.

**Etymology:** It is named after Prof. Jae-Seoun Hur (Suncheon, Republic of Korea), who has collected the type material and has kindly provided it for our study.

**Distribution:** So far known only from the type locality in Inner Mongolia, Northern China, Asia.

**Taxonomic notes:** *Hanstrassia jaeseounhurii* is similar to *H. lenae*, but differs in having much larger thalline areoles (1–3(–4) mm vs. ca 1 mm diam./across), in having only areolate thallus (vs. sometimes slightly radially elongated at the periphery of the thallus and narrower, i.e. thallus is effigurate, but only weakly lobate at the margin), in having very scarce and much larger (0.06–0.12(–0.2) mm diam./across vs. 25–40(–50) µm diam.) soredia and in having scleroplectenchymatous (vs. prosoplectenchymatous) cortical layer of thallus.

*Hanstrassia jaeseounhurii* is similar to *Fulgensia desertorum* (Tomin) Poelt in having schizidia and crustose thallus often dispersed into elements, but differs in having distinctly areolate (vs. continuous crustose) and larger (2–3(–5) cm vs. 0.75–1.6 cm across) thallus, in having larger thalline areoles (1–3(–4) mm vs. 0.4–0.7 mm across), in the lack of lobate peripheral portion, in having much thicker cortical layer, in having distinct algal zone and medulla, as well as in positioning in separate monophyletic branch (vs. it is positioned in the *Gyalolechia* s.l. clade).

After lack of apothecia and non-lobate peripheral portion, and more or less warty squamulose thallus *Hanstrassia jaeseounhurii* is similar to *Fulgensia bracteata* subsp. *deformis* (Erichsen) Poelt, but differs in having distinctly areolate thallus (vs. warty crustose to squamulose), in having much larger areoles (1–3(–4) mm vs. only in peripheral zone distinct as ca 0.2–0.5 mm across), and in having scarce and smaller (0.06–0.12(–0.2) mm vs. 0.2–0.4 mm across) schizidia.

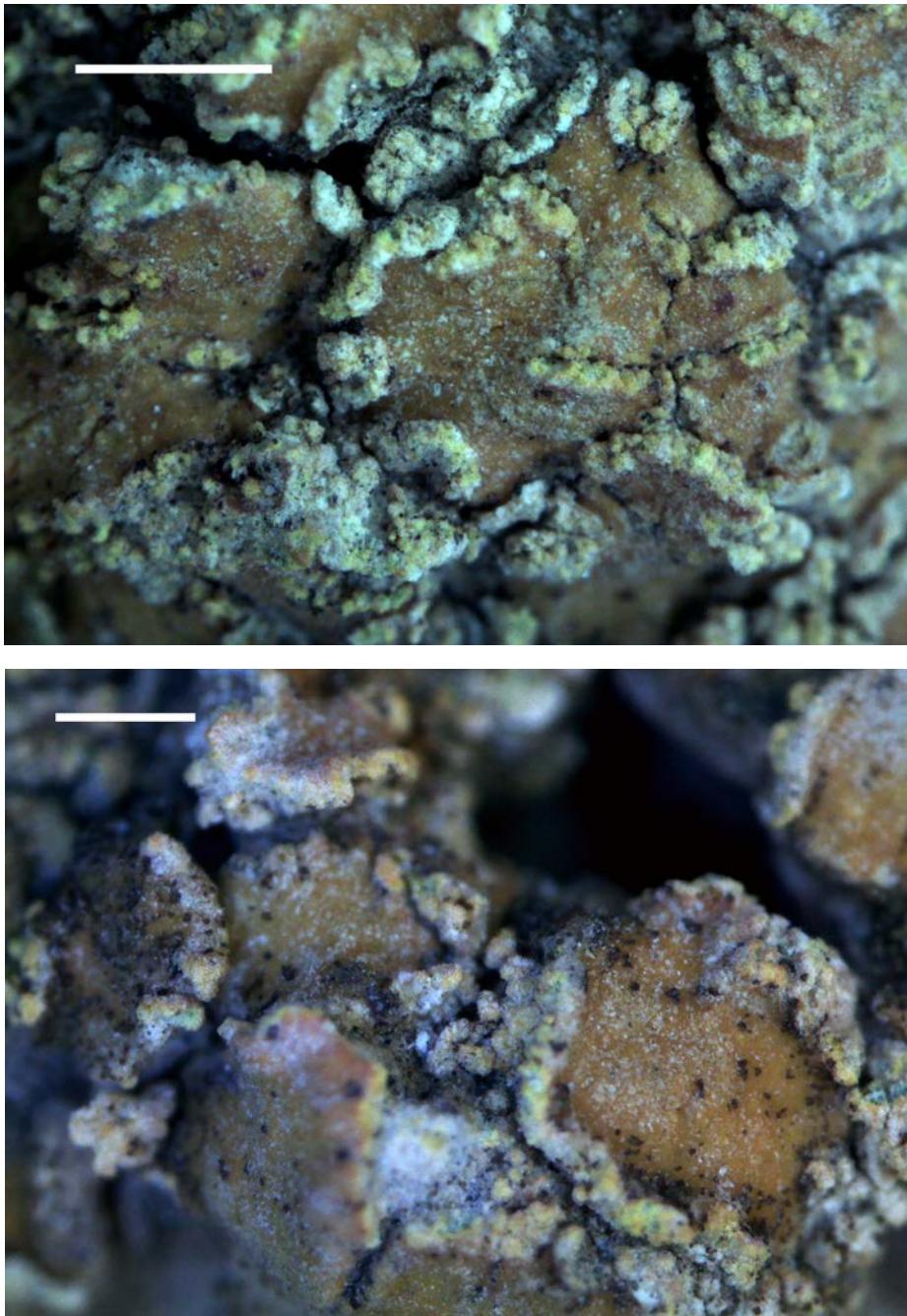


Fig. 8. *Hanstrassia jaeseounhurii* (holotype), enlarged portions with scarce schizidia. Scale 1 mm (top) and 0.5 mm (bottom). (Photo of S. Kondratyuk)

After having naked medulla in marginal soralia *Hanstrassia jaeseounhurii* is similar to *Megaspora rimisorediata* Valadbeigi et A. Nordin, growing on calcareous soil in desert conditions of Eurasia, but differs in having brownish green-yellow (vs. whitish or white grey) thallus and in having much larger (0.06–0.12(–0.2) mm vs. 0.05–0.07 mm) and lemon yellow (vs. blackish or bluish grey) soredia, as well as in having bright yellow medulla in naked portions (vs. medulla white).

***Harusavskia* S. Y. Kondr., gen. nov.**

MycoBank nr.: MB 819643.

*Similar to the genus Rusavskia, but differs in being distinctly areolate in the centre of the thallus, in the lack of lower cortical layer, in having narrower thalline lobes, in having ascospores with halo and in having very indistinct ascospore septum.*

Type species: *Harusavskia elenkinianoides* S. Y. Kondr., X. Y. Wang, S.-O. Oh et J.-S. Hur.

Thallus areolate in the centre and distinctly lobate, with well-developed lobes within peripheral zone; yellow-brownish orange to brownish yellow-orange in peripheral portions matt or somewhat shiny with rather scarce pseudocyphellae, while in the centre pseudocyphellae numerous forming eroded upper surface, whitish-brownish yellow-orange.

Apothecia lecanorine, where thalline exciple yellowish-brownish orange and disc mainly plane, dark brownish brick-orange; true exciple “textura intricata” or “textura globosa”, with very thin cell walls; cortical layer of thalline exciple paraplectenchymatous; subhymenium with numerous oil droplets; asci 8-spored; ascospores seem to be simple, septum mainly not developed (not seen in water) or only juvenile stage of septum seen, with well distinct halo to 1–1.2(–1.5) µm wide in K.

Chemistry: Thallus K+ purple. Epiphytum and cortical layer of thalline exciple K+ bright crimson purple washing in liquid; cortex of thalline section K+ bluish, violet- or ink-purple.

Ecology: Growing on siliceous rock.

Distribution: So far known only from the type locality in Chile, South America.

Etymology: Genus *Harusavskia* named after ascospores having “halo”, and after similarities with *Rusavskia elegans*, to which this material was originally included.

Taxonomic notes and phylogenetic affiliations: After combined phylogenetic tree this unique monotypic genus is positioned within the *Filsoniana*

clade of the subfamily Teloschistoideae (Fig. 1). From all representatives of the Teloschistaceae this genus differs in having ascospores with halo.

*Harusavskia elenkinianoides* S. Y. Kondr., X. Y. Wang, S.-O. Oh  
et J.-S. Hur, *spec. nova*  
(Fig. 9)

Mycobank nr.: MB 819644.

*Similar to Elenkiniana gomerana, but differs in having shorter and much wider thalline lobes, in having somewhat larger apothecia with lower cortical layer, in having much thicker true exciple especially in basal portion, and in having "textura intricata" or "textura globosa" of true exciple, in having higher hymenium, in having larger ascospores, and in having narrower ascospore septum, as well as in having ascospores with halo.*

Type: Chile. 'Laguna del Maule', Maule, on rock. Lat.: 35° 57' 25.0" S; Long.: 70° 34' 23.9" W; Alt.: ca 1,887 m alt. Coll.: Wang, X. Y., Oh, S.-O. and Hur, J.-S. (CL 120324), 30.01.2012 (KoLRI 014493 – holotype).

Thallus to several cm across, areolate in the centre and distinctly lobate, with well-developed lobes within peripheral zone to 2.5–3.5 mm wide; central portions more or less evenly areolate, areoles 0.4–0.8 mm across and to 0.2–0.3 mm thick, irregular; lobes to 2–3.5 mm long and (0.4–)0.8–1.2 mm wide towards the centre and widened towards the tips to 1–2 mm wide, dissected into secondary lobes to 0.7–1.5 mm long and to 0.7 mm wide (and to 0.4–0.5 mm thick); yellow-brownish orange to brownish yellow-orange in peripheral portions, matt or somewhat shiny with rather scarce pseudocyphellae, while towards the centre number of pseudocyphellae becoming larger/higher to totally eroded upper surface, whitish-brownish yellow-orange; thalline lobes convex, along the lobe ends in peripheral zone seen that the lower portion of lobe white.

Apothecia 0.4–1.3(–1.5) mm diam., and 0.5–0.6 mm thick in section; at first immersed into thalline lobes, then uplifted on thalline warts in the centre, but not constricted at the basis; usually very compactly aggregated covering almost the whole central part, and areoles of central part hardly recognise, because they are completely covered by apothecia; lecanorine to zeorine, where thalline exciple yellowish-brownish orange and disc mainly plane, dark brownish brick-orange. In section true exciple to 40–70(–120) µm thick in the uppermost lateral portion, to 30–70(–80) µm thick in lower lateral portion and (20–)30–60(–150) µm thick in basal portion, intermediate type between "textura intricata" and "textura globosa", with very thin cell walls, where cell

lumina to 2–3(–4)  $\mu\text{m}$  across; thalline exciple to 100–150(–170)  $\mu\text{m}$  thick with cortical layer to 15  $\mu\text{m}$  thick, paraplectenchymatous (in K to 30  $\mu\text{m}$  thick of lax paraplectenchyma, cells to 5–7  $\mu\text{m}$  across); algal cells to 10–11  $\mu\text{m}$  diam.; hymenium 100–120  $\mu\text{m}$  high; epihymenium to 25  $\mu\text{m}$  thick in K more or less becoming darker; paraphyses gradually becoming wider towards the tips to 4–6  $\mu\text{m}$  diam. in water (and distinctly widened towards the tips to 4–5  $\mu\text{m}$  diam. in K); subhymenium to 70–110(–300)  $\mu\text{m}$  thick, with numerous oil droplets to 5–6  $\mu\text{m}$  diam.; ascii 8-spored, uniseriate; ascospores seem to be simple (see below), straight or somewhat curved, almost cylindrical (10.5–)11–16(–17)  $\times$  (4.5–)5–6.5(–7)  $\mu\text{m}$  in water and (10–)12–15(–16)  $\times$  (4–)5–7  $\mu\text{m}$  in K (measurements mainly with halo); ascospore wall in water to 0.5  $\mu\text{m}$  thick, and with halo well distinct to 1–1.2(–1.5)  $\mu\text{m}$  wide in K (while not seen in water); septum mainly not developed (not seen in water) or only juvenile stage of septum observed in K, to 1.5–2(–2.5)  $\mu\text{m}$  wide (septum developed only at spore wall and not developed in the centre).

**Chemistry:** Thallus K+ purple. Epihymenium and cortical layer of thalline exciple K+ bright crimson purple washing in liquid; cortex of thalline section K+ bluish, violet- or ink-purple.

**Ecology:** It grows on siliceous rock.

**Distribution:** So far known only from the type locality in Chile, South America.

**Etymology:** Species is named after similarities to the Canary Islands endemic *Elenkiniana gomerana*.

**Taxonomic notes:** This taxon is characterised by the following features: lobate peripheral zone, lobes mainly matt and slightly lighter at the tips, yellowish-brownish or yellow-brownish orange; areoles in the centre richly covered by pseudocyphellae and upper surface seem to be heavily eroded and whitish-yellowish brown in contrast to dark brick-orange discs of apothecia, which are very abundant, almost completely covered some parts in the centre; ascospores mainly simple (only very rarely juvenile septum seen); very soft tissue in the true exciple (well developed, but concave), as well as presence of halo in ascospores.

*Harusavskia elenkinianoides* is similar to the Canary Islands endemic *Elenkiniana gomerana* (J. Steiner) S. Y. Kondr., Kärnefelt, A. Thell, Elix, J. Kim, A. S. Kondr. et J.-S. Hur in having areolate in the centre and lobate in peripheral zone and the same colouration thallus, in having numerous pseudocyphellae in the centre and more or less immersed into the thallus apothecia, but differs in having shorter (2–3.5 mm vs. to 8 mm long after Steiner 1911), and much wider (0.8–1.2 mm wide towards the centre and to 1–2 mm wide towards the tips, vs. 0.3–0.7(–1) mm wide) thalline lobes, in having somewhat larger apothecia (0.4–1.3(–1.5) mm vs. 0.5–0.7(–1.2) mm diam.), in having lower cortical layer (to 15(–30)  $\mu\text{m}$  vs. 37–65  $\mu\text{m}$  after Llimona and Werner 1975, and 16–28

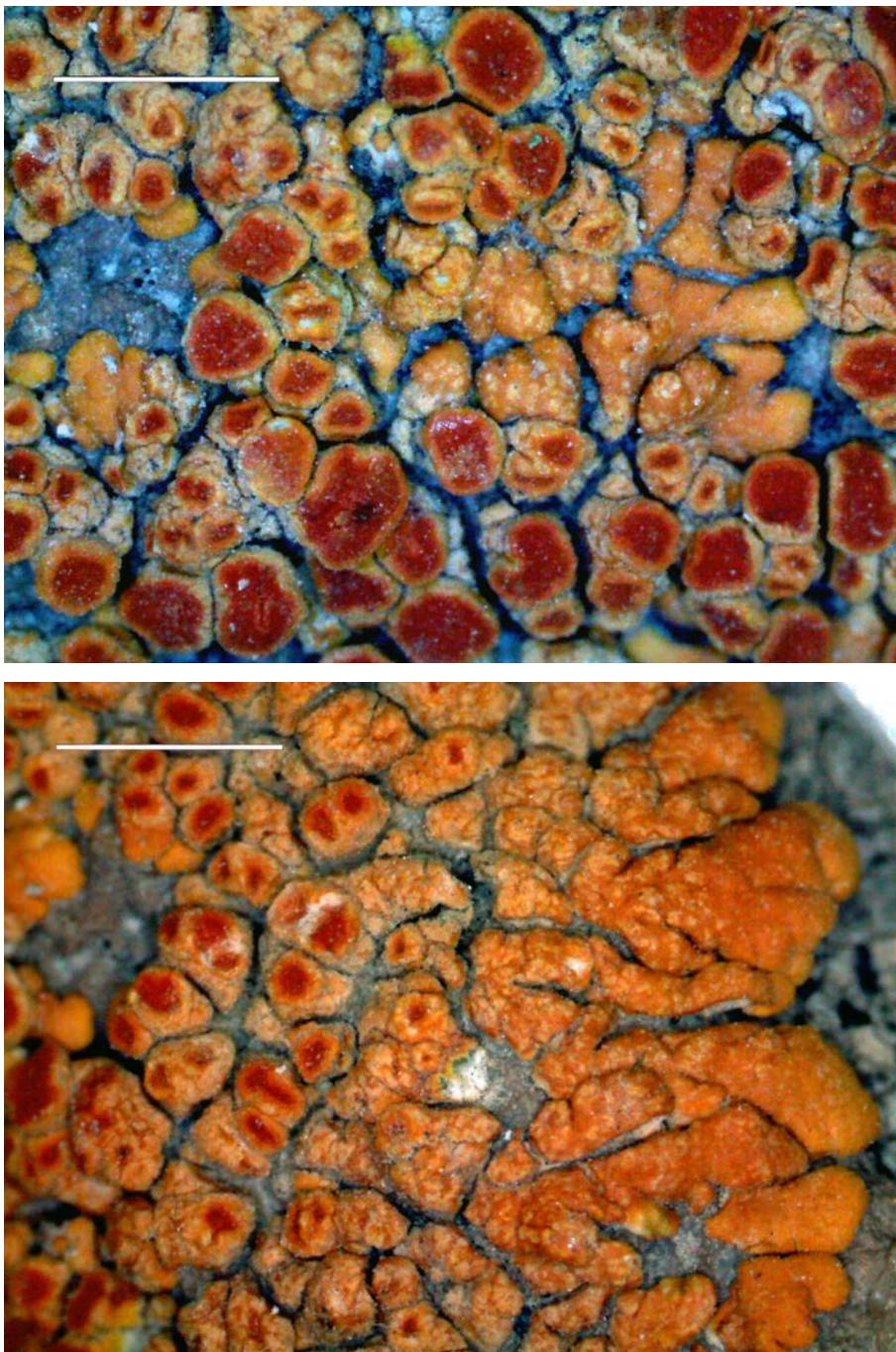


Fig. 9. *Harusavskia elenkinianoides* (holotype). Scale 2 mm. (Photo of S. Kondratyuk)

µm after Steiner 1911), in having much thicker true excipic especially in basal portion (30–60(–150) µm vs. 25 µm thick), and in having “textura intricata” or “textura globosa” of true excipic (vs. of parallel hyphae), in having higher hymenium (100–120 µm vs. 60–90 µm high), in having larger ascospores ((10.5)–11–16(–17) × (4.5)–5–6.5(–7) µm vs. 7.5–9.9–12.5 × 4–5.5–6 µm after Llimona and Werner 1975, and 7.5–11 × 5.5–6.5 µm after Steiner 1911), and in having narrower ascospore septum (1.5–2 µm vs. 2–4 µm wide), as well as in having ascospores with halo.

*Harusavskia elenkinianoides* is similar to the epiphytic Chilean taxon *Caloplaca chilensis* S. Y. Kondr., Kärnefelt, Frödén et Arup, but differs in having larger thallus (several cm vs. 8–15(–25) mm diam.), being areolate in the centre, in having much wider thalline lobes (0.8–1.2 mm vs. 0.2–0.4 mm wide), in having pseudocyphellae, in having wider paraphysis tips (4–6 µm vs. to 3 µm diam.), in the lack of rhizine-like structures on underside and prothallus, in having smaller and nonstipitate apothecia (0.4–1.3 mm vs. 0.4–2.7 mm diam., stipe 0.3–1 mm long), in having much narrower ascospore septum (1.5–2 µm vs. 4–6(–7.5) µm wide), as well as in saxicolous habit, and having ascospores with halo.

Unfortunately molecular data on *Caloplaca chilensis* are still not available, while data on 6 genes of *Caloplaca chilensis* were mentioned in literature (Gaya et al. 2012). However, in fact these data belong to another taxon of the genus *Follmannia*.

After having convex thalline lobes with white sides along the lobe ends in the peripheral zone, as well as after having indistinctly whitish in the centre thallus *Harusavskia elenkinianoides* is similar to *Calogaya mogoltanica* (S. Y. Kondr. et Kudratov) S. Y. Kondr., Kärnefelt, Elix, A. Thell, J. Kim, M.-H. Jeong, N.-N. Yu, A. S. Kondratuk et J.-S. Hur known from limestone outcrops of Central Asia (Tadzhikistan and Iran). However, *Harusavskia elenkinianoides* differs in having brownish orange, epruinose (vs. dirty greenish yellow pruinose) thallus, smooth in the centre (vs. papillate and verruculose in the centre), in having thinner thalline lobes (0.4–0.5 mm vs. to 1–2(–3) mm thick), which well delimitated by wide cracks between each others (vs. indistinct in the centre of thallus and only seen as upper portion of convex surface of cortical layer), in having pseudocyphellae (vs. pruinose but without pseudocyphellae), in having smaller (mainly to 1 mm vs. 0.8–1.8 mm diam. not constricted at the basis zeorine apothecia (lecanorine, with thick thalline margin, attenuated at the basis), in having “textura intricata” or “textura globosa” type of true excipic (vs. well-developed paraplectenchymatous with well-developed matrix) in having shorter ascospores (12–15 × 5–7 µm vs. 15–18 × 6–7 µm) (Kondratyuk and Kudratov 2003).

In contrast to *Rusavskia elegans* (Link) S. Y. Kondr. et Kärnefelt (to which this material was originally included), it has distinctly areolate central portion of thallus, and lobes do not have well-developed lower cortex, as well as ascospores have halo, while septum is not well distinct.

***Huriella* S. Y. Kondr. et D. Upreti, *gen. nov.***

Mycobank nr.: MB 819645.

*Similar to the genus *Squamulea* of the subfamily *Teloschistoideae*, but differs in having areolate (not squamulose!) thallus, smaller zeorine apothecia, in having narrower ascospores, as well as in positioning in separate robust monophyletic branch in the *Amundsenia-Squamulea* clade.*

Type species: *Huriella loekoesiana* S. Y. Kondr. et D. Upreti.

Thallus crustose, small, but often forming larger aggregations, areolate (not squamulose!), dull yellow-brownish or yellowish green. Cortical layer paraplectenchymatous. Hypothallus not observed.

Apothecia small, seem to be biatorine, but zeorine in section, 1(–3–4) per areole, disc dull yellow to bright yellow (especially at initial stages) to dark brownish yellow; own margin concolourous with disc; true exciple paraplectenchymatous; asci 8-spored; ascospores polarilocular, small, widely ellipsoid with rounded ends. Conidiomata and conidia not seen.

Chemistry and ecology see below in species description.

Distribution and species diversity: So far monotypic genus, while possibly includes complex of species. *H. loekoesiana* is known from numerous localities in South Korea, Eastern Asia.

Etymology: It is named after Prof. Jae-Seoun Hur (Suncheon, Republic of Korea), who has kindly supported our study of the Teloschistaceae.

Taxonomic notes and phylogenetic affiliations: After combined phylogenetic analysis, as well as after separate ITS and mtSSU analyses the genus *Huriella* is always positioned in a sister position to the genera *Squamulea* and *Amundsenia* (or between the *Amundsenia-Squamulea* and the *Xanthocarpia* clades, if limited number of genera are included into the phylogenetic analysis) as separate robust monophyletic branch, while the level of support of the whole *Amundsenia-Squamulea* clade is much lower.

After molecular data hitherto available the genus *Huriella* includes only the type species, while we expect in future recognising a complex of taxa. One more species very similar to *H. loekoesiana* is in preparation for official publication.

***Huriella loekoesiana* S. Y. Kondr. et D. Upreti, spec. nova  
(Fig. 10)**

MycoBank nr.: MB 819646.

*Similar to Squamulea micromera, but differs in having areolate (not squamulose!) thallus, in having larger thalline areoles, in having larger zeorine apothecia and in having narrower ascospores.*

Type: Republic of Korea. Gangwon-do: Jeongseon-gun, Buk-myeon, Mt Baenon, on rock, growing together with *Laundonia flavovirescens* agg., *Rinodina* sp. Lat.: 37° 26' 34.9" N; Long.: 128° 45' 21.3" E; Alt.: ca 748 m a.s.l. Coll.: Wang, X. Y., Jeon, H. S., Lü, L. and Ryu, J. A. (100803), 28.05.2010 (KoLRI 012491 – holotype).

Thallus 2–4(–10) mm across, but forming larger aggregations to 5–10 cm across, crustose, areolate (not squamulose!), dull yellow-brownish, yellow-greenish; areoles (0.2–)0.3–0.8(–1.0) mm across, from regularly rounded to irregular, somewhat distant and scattered, with more or less even surface, becoming thinner towards the edges (not uplifted towards the edges!). Thallus in section to (60–)70–100(–120) µm thick; cortical layer to 10–15 µm thick, outermost portion somewhat brownish-yellowish in places, paraplectenchymatous, cell lumina to 5–7 µm diam./across; algal layer to 40–70 µm thick; medulla not developed or ca 10–30 µm thick below the algal layer in places. Hypothallus not observed.

Apothecia 0.2–0.4(–0.5) mm diam., to 0.2 mm thick in the section; 1(–3–4) per areole, immersed in thallus at first then sessile, seem to be biatorine, but zeorine in section; disc dull yellow to bright yellow, especially at initial stages (lighter than thallus) to dark brownish yellow at overmature, own margin bright or light yellow, concolourous with disc of young apothecia; in section zeorine, true exciple to 40–50 µm wide in the uppermost lateral portion, consisting of rounded cells to 5–6(–7) µm diam./across, to 10–15 µm thick in lower lateral portion and to 25–30 µm thick in basal portion, paraplectenchymatous, cell lumina 3–5 µm diam./across; thalline exciple to 40–50 µm thick, with well-developed cortical layer to 10–15 µm thick, paraplectenchymatous; cell rounded 5–7 µm diam./across; algal layer below true exciple entire or in clusters, algal cells to 13–20(–23) µm diam.; hymenium to 50–60 µm high; paraphyses distinctly swollen towards the tips to 5(–6) µm diam.; subhymenium 20–30 µm thick, hyaline, without oil; asci 8-spored; ascospores widely ellipsoid with rounded ends, (8.5–)9–11(–12) × (4.5)5–6 µm in water and becoming especially wider to (8–)10–12(–13) × 6–7.5(–8) µm in K; septum (3.5–)4–5(–6) µm wide in water and 4–6(–7) µm wide in K. Conidiomata and conidia not seen.

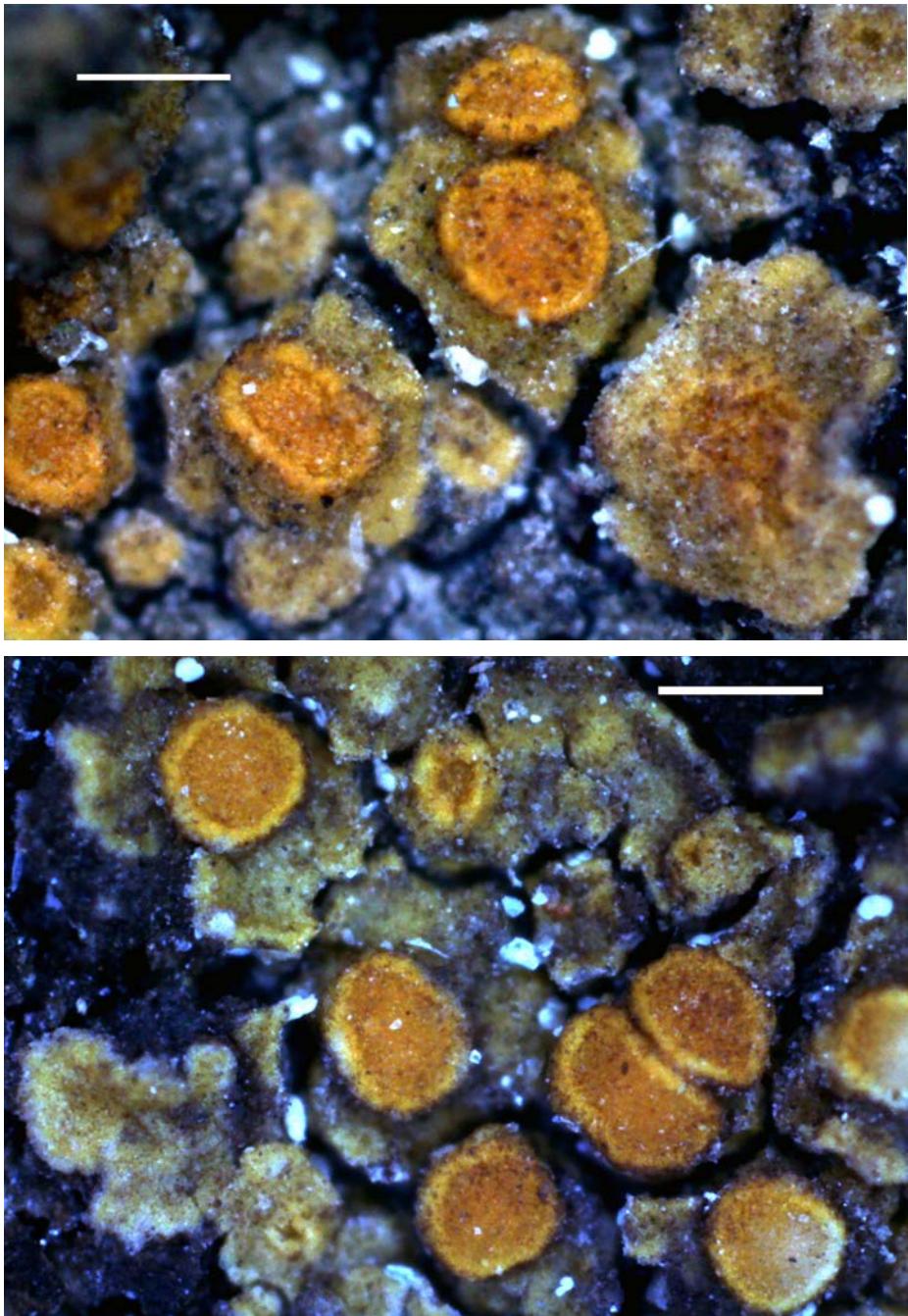


Fig. 10. *Huriella loekoesiana* (holotype), general habit. Scale 0.5 mm. (Photo of S. Kondratyuk)

**Chemistry:** Cortical layer of thallus, epihymenium and outer portion of true exciple and thalline exciple K+ purple; outer brownish portions of thalline cortical layer K+ dark brown purple; substances not studied.

**Ecology:** It grows on siliceous rocks growing often together with *Laundonia flavovirescens* agg., *Rinodina* sp.

**Distribution:** Known from scattered localities throughout the Republic of Korea, Eastern Asia.

**Etymology:** Species is named after our good friend and colleague, well-known Hungarian lichenologist László Lőkös (BP, Budapest) in recognition of his contribution to our recent knowledge on Korean lichen flora.

**Taxonomic notes:** *Huriella loekoesiana* is characterised by very small light yellow discs of apothecia, slightly lighter of (dull brownish yellow) colour of thallus, by areolate thallus with more or less rounded scattered areoles and small and especially narrow ascospores.

*Huriella loekoesiana* is similar to *Squamulea micromera* (Hue) S. Y. Kondr., L. Lőkös et J.-S. Hur, but differs in having areolate thallus (vs. squamulose), in having larger thalline areoles ((0.2–)0.3–0.8(–1.0) mm vs. 0.2–0.3 mm diam./across), in having larger zeorine apothecia (0.2–0.4(–0.5) mm vs. 0.2–0.3(–0.4) mm diam., biatorine), in having thinner true exciple in basal portion (25–30 µm vs. 40–60 µm thick), in having thinner subhymenium (20–30 µm vs. 50–60 µm thick), and in having narrower ascospores ((8.5–)9–11(–12) × (4.5–)5–6 µm vs. 9–12 × 6–7(–7.5) µm).

*Huriella loekoesiana* is similar to *Squamulea subsoluta* (Nyl.) Arup, Søchting et Frödén, the type species of genus *Squamulea* Arup, Søchting et Frödén, but differs in having smaller thalline areoles (not squamules, 0.3–0.8(–1.0) mm vs. 0.1–2.5 mm wide); in having smaller apothecia (0.2–0.4(–0.5) mm vs. 0.1–0.6(–1.0) mm diam.), in having narrower ascospores ((8.5–)9–11(–12) × (4.5–)5–6 µm vs. 9.5–12.5(–14) × 5.5–7 µm), and in having wider ascospore septum (4–5 µm vs. 3–4 µm wide).

It should be emphasised that taxonomic status of *Squamulea micromera* is waiting for confirmation by molecular data from freshly collected specimens. In general, the type specimen of *S. micromera* may be considered within the variation of *S. subsoluta* (Nyl.) Arup, Søchting et Frödén (sensu Wetmore 2003). However, we prefer to keep *S. micromera* as separate taxon until molecular characters of this material will not be checked especially.

There is one more taxon, very similar to *Huriella loekoesiana*, but differing in having smaller ascospores and other anatomical characters, which is probably also the member of the genus *Huriella* is in preparation.

Other specimens studied selected: Republic of Korea. Chungcheongbuk-do, Danyang-gun, Danseong-myeon, Daejam-ri, 584 Road, near Sunam Valley, on rock growing together with *Phaeophyscia* and *Endocarpon*. Lat.: 36° 54' 20.8" N; Long.: 128° 18' 42.9" E; Alt.:

ca 207 m a.s.l. Coll.: Lee, B. G. (152763), 05.09.2015 (KoLRI 037750 sub *Huriella*). – Gyeongsangbuk-do, Ulleung-do Island, Ulleung-gun, Seo-myeon, Namyang-ri, Turtle Rock (Mt 18-1), seashore rocks, on basalt. Lat.: 37° 27' 36.5" N; Long.: 130° 51' 27.9" E; Alt.: 2 m a.s.l. Coll.: Kondratyuk, S. Y. and Lőkös, L. (161904), 10.07.2016. (KoLRI 40141 sub *Candeliella*). – Gyeongsangbuk-do, Ulleung-do Island, Ulleung-gun, Ulleung-eup, Dodong-ri, Seaside Trail near Dodong Port, on steep siliceous rocks, growing together with *Squamulea squamosa*. Lat.: 37° 28' 59.9" N; Long.: 130° 54' 40.7" E; Alt.: 20 m a.s.l. Coll.: Kondratyuk, S. Y. and Lőkös, L. (161998), 11.07.2016. (KoLRI 40236 sub *Squamulea squamosa*); the same locality, (162000), (KoLRI 40238 sub *Squamulea squamosa*). – Jeju-do, Seogwipo-si, Eongsan-eup, Goseong-ri, Seopjicoji, on rock growing together with *Buellia stellulata*, and *Agonimia cavernicola*. Lat.: 33° 19' 21.0" N; Long.: 126° 50' 49.03" E; Alt.: ca 69 m a.s.l. Coll.: Gagarina, L. (140363), 19.06.2014 (KoLRI 022729 sub *Buellia stellulata*); the same locality and collector, (140355), (KoLRI 022723). – Jeju-do, Seogwipo-si, Sangjeong-dong, Yeongtto waterfall, on rock, growing together with *Acarospora ulleungdoensis* [Note: first time recorded from Cheju-do Island]. Lat.: 33° 16' 01.7" N; Long.: 126° 29' 49.00" E; Alt.: ca 210 m a.s.l. Coll.: Joshi, Y. and So, J.-U. (140595), 19.06.2014 (KoLRI 022989). – Jeju-do, Jeju-si, Gujwa-eup, Gimnyeong-ri, Gimnyeong Beach, seaside, on rock, growing together with *Leptogium* sp., *Endocarpon* sp., *Pyrenopsis* sp. Lat.: 33° 33' 53.91" N; Long.: 126° 45' 47.66" E; Alt.: ca 10 m a.s.l. Coll.: Hur, J.-S., Wang, X. Y., Ryu, J. A. and Hur, J. Y. (090020), 19.04.2009 (KoLRI 009784). – Jeollanam-do, Yeosu-si, Nam-myeon, Dumo-ri, Jickpo coast, Geumo-do, on rock. Lat.: 34° 30' 46.08" N; Long.: 127° 44' 16.04" E; Alt.: ca 30 m a.s.l. Coll.: Jayalal, U., Park, J. S. and Ryu, J. A. (120433), 26.04.2012 (KoLRI 015423 sub *Squamulea subsoluta*). – Jeollanam-do, Yeosu-si, Nam-myeon, Simjang-ri, Geumo-do, on rock, growing together with *Thelidium japonicum*. Lat.: 34° 28' 59.20" N; Long.: 127° 48' 15.48" E; Alt.: ca 4 m a.s.l. Coll.: Halda, J. P. (160440), 10.06.2016 (KoLRI 038585 sub *Thelidium japonicum* (det. by Halda, J. P.)). – Jeollanam-do, Sinan-gun, Haui-myeon, Unggok-ri, Haui-do, seaside, on rock growing together with *Orientophila subscopularis*. Lat.: 34° 36' 07.07" N; Long.: 126° 00' 52.02" E; Alt.: ca 20 m a.s.l. Coll.: Oh, S.-O., Park, J. S. and Woo, J.-J. (130672), 28.06.2013 (KoLRI 019017 sub *Orientophila subscopularis*).

*Ikaeria* S. Y. Kondr., D. Upreti et J.-S. Hur, *gen. nov.*

Mycobank nr.: MB 819647.

*Similar to Mikhtomia of the subfamily Caloplacoideae, but differs in having zeorine or lecanorine apothecia, and in forming separate robust monophyletic branch in the Teloschistoideae.*

Type species: *Ikaeria aurantiellina* (Harm.) S. Y. Kondr., D. Upreti et J.-S. Hur.

Thallus crustose, continuous, yellowish or greyish or whitish yellow. Hypothallus sometimes present.

Apothecia zeorine or lecanorine, disc orange to brownish orange, true exciple of "textura intricata" or somewhat scleroplectenchymatous, asci 8-spored, ascospores polaribilocular.

Chemistry: Not studied.

Ecology: Growing on bark of twigs of shrubs or trees.

Distribution and species diversity: So far this genus is monophyletic and known only from the Canary Islands.

Etymology: The genus is named after the well-known Swedish lichenologist Ingvar Kärnefelt (Lund, Sweden) in recognition of his contribution to lichenology, and who has kindly provided for us an opportunity to participate in field trips to Australia and to the Canary Islands and provided his earlier collections in our disposal.

Taxonomic notes and phylogenetic affiliations: After combined phylogenetic analysis the genus *Ikaeria* is positioned in the subfamily Teloschistoideae. From molecular data hitherto available the genus is monotypic including only type species *I. aurantiellina*, known from the Canary Islands and southwestern Europe.

*Ikaeria aurantiellina* is similar to *Mikhtomia* of the subfamily Caloplaeoidae (or to *Leutrouitia* of the Leutrouitiaceae), but differs in having zeorine or lecanorine apothecia, bipolar ascospores and in forming separate robust monophyletic branch in the Teloschistoideae.

*Ikaeria aurantiellina* is similar to taxa of the *Mikhtomia* s.l. subphylum, especially of the genera *Laundonia*, *Opeltia*, etc., but differs in positioning in somewhat out position to both Caloplaeoidae and Teloschistoideae after ITS, nrLSU and mtSSU analysis, as well as after combined phylogenetic analysis.

This genus often forms common branch with the genus *Yoshimuria*, but level of support of this branch is rather low to make conclusion that they are congeneric. In contrast, both genera, *Ikaeria* and *Yoshimuria*, have the highest level of support if they are presented by the same number of specimens (not shown in Figs 1–4).

Data on unique position of the genus *Ikaeria* were already obtained several years ago. However, these data were obtained from different specimens. Several attempts to get three gene sequences from the same specimen were attempted, but they were unsuccessful. So this genus, the same as genus *Fominella* (see above) is in urgent need of confirmation by molecular data obtained from the same specimen.

After having broom-like paraphyses often becoming brownish towards the tips the genus *Ikaeria* is similar to the Australian '*Caloplaca*' *marchantiorum* S. Y. Kondr. et Kärnefelt (Kondratyuk *et al.* 2009) (supposed member of the Teloschistoideae or Brownielloideae). Molecular data on '*Caloplaca*' *marchantiorum* are still not available, while several attempts to extract DNA were attempted.

Van den Boom and Etayo (2006) argue that the name *Caloplaca aegatica* Giralt, Nimis et Poelt, known from western Europe, might be a later synonym of *Ikaeria aurantiellina*.

*Klauderuiella* S. Y. Kondr. et J.-S. Hur, *gen. nov.*

Mycobank nr.: MB 819648.

Similar to *Variospora* of the subfamily *Caloplacoideae*, but differs in having lobate portions in peripheral portion of thallus, in having lecanorine apothecia, and in forming separate robust monophyletic branch.

Type species: *Klauderuiella thallincola* (Wedd.) S. Y. Kondr. et J.-S. Hur.

Thallus distinctly lobate (at least in peripheral portions, while areolate in the centre); often large size, yellow to orange, sometimes central and peripheral zone differently coloured (various shades of orange colour), matt, epruinose or rarely pruinose; lobes plane or convex; cortical layer "pseudo" paraplectenchymatous (*sensu* Clauzade and Roux 1985).

Apothecia lecanorine to zeorine; true excipie scleroplectenchymatous with well-developed matrix in basal portion and to paraplectenchymatous in lateral portions; subhymenium with oil droplets or greyish owing to numerous crystals; ascospores often widened in equatorial portion to rhombic or citriform.

Ecology: Growing especially on calcium containing rocks.

Distribution: Members of this genus are known as widely distributed in the Northern Hemisphere.

Etymology: It is named after the well-known French lichenologist Claude Roux (Klaŭdo Roux in Esperanto) (1945–) in recognition of his enormous contribution to lichenology. He also excellently illustrated the *Klauderuiella thallincola* and *Klauderuiella aurantia* groups in the well-known work "Likenoj de Okcidenta Eŭropo. Illustrata determinlibro." (1985), as well as he is a great patriot of the Esperanto language.

Taxonomic notes and phylogenetic affiliations: In current phylogenetic tree the genus *Klauderuiella*, i.e.: the *Variospora thallincola* group has the highest level of support after combined three gene phylogeny in contrast to very low support of the *Variospora-Bryoplaca* subclade, where it is positioned (see Fig. 2). The latter subclade (i.e.: the *Variospora-Bryoplaca* subclade) is probably still heterogeneous.

After molecular data hitherto available the genus *Klauderuiella* includes three species, i.e.: *K. thallincola*, *K. aurantia* and *K. flavesrens*.

From morphological point of view the genus *Klauderuiella* has very good (distinct) character – lobate in peripheral portion of the thallus in contrast to crustose members of the genus *Variospora* s. str. (i.e.: *V. velana* (A. Massal.) Arup, Søchting et Frödén, *V. latzelii* (Servit) S. Y. Kondr., *V. macrocarpa* (Anzi) Arup, Søchting et Frödén, *V. dolomitica* (Hue) Arup, Søchting et Frödén and *V. kudratovii* (S. Y. Kondr., B. Zarei-Darki et Hur) S. Y. Kondr., Kärnefelt, A. Thell, Elix, J. Kim, A. S. Kondr. et Hur).

The *Klauderuiella* is somewhat analogous to genera *Flavocitrina* and *Calogaya* in the Xanthorioideae and *Filsoniana* in the Teloschistoideae. However, the genus *Klauderuiella* differs in having different type of paraplectenchymatous tissue in cortical layer, as well as in having typical bipolar ascospores (without thickening at poles).

Status of *Variospora latzelii* and *V. australis* hitherto included in this genus *Variospora* only after ITS phylogeny should be confirmed by data on nrLSU and mtSSU.

Possibly one more monophyletic group will be segregated from the *Variospora* s.l. branch of the Caloplacoideae in future.

***Laundonia* S. Y. Kondr., L. Lőkös et J.-S. Hur, gen. nov.**

MycoBank nr.: MB 819649.

*Similar to the genus Mikhtomia of the subfamily Caloplacoideae, but forming separate robust monophyletic branch.*

Type species: *Laundonia flavovirescens* (Wulfen) S. Y. Kondr., L. Lőkös et J.-S. Hur.

Thallus crustose, continuous or areolate, margin thinning at edge, without elongated lobes; yellowish grey, yellowish orange, tan orange to greenish yellow or greenish orange, pruinose, without vegetative propagules or sorediate; prothallus absent or present, black.

Apothecia of medium size, lecanorine or biatorine, disc dark orange, reddish orange or brownish orange; true excipic consisting of radiating hyphae to "textura intricata" in basal portion; asci 8-spored; ascospores polaripolucar, hyaline.

Unfortunately data on conidia of type species and *L. persimilis* (see Wetmore 2007) not provided, while description of conidiomata present in descriptions.

Chemistry: Parietin, fallacinal, emodin, and teloschistin. Chemistry of *L. persimilis* was not especially investigated, however, C+ reaction may illustrate the presence of gyrophoric or lecanoric acids.

Ecology: It grows on wood or bark.

Etymology: It is named after the well-known British lichenologist Jack Rodney Laundon (1934–) who has made important contribution to taxonomy of this species complex.

Distribution: Type species, i.e. *Laundonia flavovirescens* is widely distributed in the Northern Hemisphere (see notes below), while *L. persimilis* is known only from western North America.

Taxonomic notes and phylogenetic affiliations: After combined phylogenetic analysis the genus *Laundonia* is positioned in out position to the other

genera of the *Mikhtomia* clade of the Caloplacoideae (Fig. 2), and it includes two species, i.e.: *L. flavovirescens* and *L. persimilis*.

In premolecular era it was thought that *Laundonia flavovirescens* is a rather common taxon distributed worldwide (see Wetmore 2007). However, when we have started extraction DNA and further sequencing appeared material morphologically similar to *Laundonia flavovirescens* to belong to many other genera like *Yoshimuria* S. Y. Kondr., Kärnefelt, Elix, A. Thell et Hur, *Mikhtomia* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur (subfamily Caloplacoideae), *Orientophila* Arup, Søchting et Frödén (subfamily Xanthorioideae) (Kondratyuk *et al.* 2016d), while *Laundonia flavovirescens* is rather rare itself.

Species of the genus *Laundonia* are very similar to members of the genus *Mikhtomia* in its current sense, especially in the Eastern Asian region. Many species of the latter genus (i.e. *Mikhtomia*), as well as the genus *Oxneriopsis* were segregated from the so-called *Caloplaca flavovirescens* group.

However, genus *Laundonia* differs in having zeorine or lecanorine apothecia, in the lack of vegetative propagules, and in positioning in the outermost position to the genus *Mikhtomia* of the *Mikhtomia* s.l. clade.

It can be similar to some species of the genus *Tomnashia* of the Xanthorioideae (see below), but differs in having lecanorine mainly orange to dark orange apothecia (without reddish tinge), as well as in preference to non-calcareous rocks, while occasionally growing on calcareous rocks, too.

From morphological point of view the rare, Eastern Asian, recently described taxon *Caloplaca kedrovopadensis* S. Y. Kondr. et J.-S. Hur (Kondratyuk *et al.* 2014b, 2016a) can be also candidate in being member of the genus *Laundonia*. However, molecular data are still missing for this taxon.

Earlier species of this genus were included mainly after ITS phylogeny to the genus *Gyalolechia* s.l. (Arup *et al.* 2013a), which was characterised by dominance of fragilin and presence of smaller amounts of parietin and emodin (often/sometimes with additional depsidones). However, as it was later shown the *Gyalolechia* s. str. branch including *G. aurea* and *G. canariensis* has the highest level of support, but this branch is in out position to the *Mikhtomia* s.l. clade (Kondratyuk *et al.* 2014c, and Fig. 2 in this paper). The genus *Laundonia* forms separate robust monophyletic branch within the *Mikhtomia* s.l. clade.

### *Lazarenkoiopsis* S. Y. Kondr., L. Lökö et J.-S. Hur, *gen. nov.*

MycoBank nr.: MB 819650.

*Similar to genus Solitaria of the subfamily Xanthorioideae, but differs in having distinctly cracked-areolated, much thicker thallus, numerous oil droplets in the hymenium, "textura intricata" tissue in the true exciple, not swollen towards the tips of paraphyses, as well as in having higher concentrations of fragilin and detectable*

*quantity of emodin, 7-chloro-emodin, erythroglauzin, 7-chloroparietinic acid, physcoin bysanthrone, physcoin 9-anthrone and physcoin 10-anthrone, as well as in positioning in the Caloplacoideae.*

Type species: *Lazarenkoiopsis ussuriensis* (Oxner, S. Y. Kondr. et Elix) S. Y. Kondr., L. Lőkös et J.-S. Hur.

Thallus rather thick in the centre and very thin at the periphery, smooth to distinctly cracked by rather broad cracks, upper surface whitish, dull greyish white to grey, greenish grey-brown or dull greenish yellow with well-contrasting bright yellow, dull yellow or dull brownish yellow soralia. Soralia initially rounded or irregular, usually at the margins of areoles, then becoming confluent and spectacular elongated, fissure-like. Sorediose mass convex, yellow to greenish yellow or dull brownish yellow.

Apothecia zeorine or lecanorine, disc brown to dark brown, own margin dull yellow to almost colourless; cortical layer of thalline exciple paraplectenchymatous; true exciple of "textura intricata" or poorly developed scleroplectenchyma with cell lumina of 1–1.5  $\mu\text{m}$  diam.; asci 8-spored; ascospores broadly ellipsoid to almost spherical or elongated and fusiform.

Chemistry: It contains fragilin (major), parietin (major/minor), emodin and physcoin bysanthrone (minor), and 7-chloroemodin, erythroglauzin, 7-chloroparietinic acid, physcoin 9-anthrone, and physcoin 10-anthrone (trace).

Etymology: It is named after the well-known Ukrainian bryologist Andrij Sozontovych Lazarenko (1901–1979) for his contribution to the bryoflora of the Eastern Asian region.

Ecology: It grows on various deciduous trees, especially on *Acer pseudoboldianum*, growing together with *Oxneriopsis oxneri* and *Caloplaca cerina*.

Taxonomic notes and phylogenetic affiliations: *Lazarenkoiopsis ussuriensis* is similar to *Solitaria chrysophthalma* (Degel.) Arup, Søchting et Frödén, but differs in having a much thicker, distinctly cracked-areolated thallus, numerous oil droplets in the subhymenium, more or less "textura intricata" tissue in the true exciple (scleroplectenchymatous in *S. chrysophthalma*), paraphyses not swollen towards the tips (4–6.5  $\mu\text{m}$  wide in *S. chrysophthalma*), a thinner exciple, as well as higher concentrations of fragilin and detectable quantity of emodin, 7-chloroemodin, erythroglauzin, 7-chloroparietinic acid, physcoin bysanthrone, physcoin 9-anthrone, and physcoin 10-anthrone, as well as in positioning in the Teloschistoideae (Fig. 1). The dimensions of the ascospores and ascospore septa are very similar for these two species.

*Lazarenkoiopsis ussuriensis* regularly grows together with *Oxneriopsis oxneri* (S. Y. Kondr. et Søchting) S. Y. Kondr., D. Upreti et J.-S. Hur, so it often appears that the same thallus produces both isidia and soredia. The thalli of both species (especially after long storage in herbaria) are the same colour.

However, when growing side by side, *O. oxneri* differs by characteristic ascending thalline fragments exposing the white medulla, while upper surface of *Lazarenkoiopsis ussuriensis* is always entire and medulla not apparent (except for soralia). *Caloplaca cerina* (Ehrh. ex Hedw.) Th. Fr. is also sometimes associated with *Lazarenkoiopsis ussuriensis*, but differs by its bright yellowish discs with a greyish thalline margin (in contrast to the darker brown discs in *L. ussuriensis*), as well as in lacking of soralia.

Suggestion that “*Caloplaca oxneri* would appear to be closely related to *C. ussuriensis*, but differs in having phyllidia and schistidia rather than well-developed soralia and convex sorediose mass” (Kondratyuk *et al.* 1996) is not supported by current molecular data. These taxa are members of the different robust monophyletic branches, i.e. the *Oxneriopsis* and *Lazarenkoiopsis* of the subfamilies Calopacoideae and the Teloschistoideae, respectively.

*Nevilleiella* S. Y. Kondr., et J.-S. Hur, *gen. nov.*

MycoBank nr.: MB 819651.

*Similar to genus Filsoniana of the subfamily Teloschistoideae, but differs in having crustose thallus and biatorine apothecia, as well as forming separate robust monophyletic branch.*

Type species: *Nevilleiella marchantii* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr. et J.-S. Hur.

Thallus crustose, developing almost spherical pustule-like formations or comprised of discrete, rather scattered distant areoles (which do not form entire crust), brownish yellow to brownish orange, sometimes with whitish pruina; areoles very variable, often convex, warty to almost spherical, sometimes with schizidial-like formations, cortical layer palisade paraplectenchymatous. Apothecia biatorine, sometimes becoming zeorine; disc and own margin rusty brownish orange or brownish orange, concolourous with thallus; true exciple paraplectenchymatous with distinct matrix in basal portion; subhymenium with oil droplets; ascospores polarilocular with almost invisible septum mounted in water (better seen in K). Conidia narrowly bacilliform, 3–4 × 0.8–1 µm.

Chemistry: Parietin (major), fallacial (submajor/minor), parietinic acid, teloschistin (minor/trace) present.

Ecology: It forms crust on clayey soil in saline areas, open localities or in mallee areas or on siliceous rocks along the coast and rough pasture, heathland and dry sclerophyll forest of Australian continent.

Etymology: This genus is named after Dr Neville Marchant, who kindly assisted us during our field work in Western Australia.

**Distribution and species diversity:** The genus includes two species rather widely distributed in Australia.

**Taxonomic notes and phylogenetic affiliations:** After combined phylogenetic analysis the genus *Nevilleiella* is positioned within the *Filsoniana* clade of the Teloschistoideae (Fig. 1).

The genus *Nevilleiella* is similar to genus *Filsoniana* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur of the subfamily Teloschistoideae, but differs in having crustose thallus and biatorine apothecia, as well as forming separate robust monophyletic branch.

The type species *Nevilleiella marchantii* was mentioned as the closest to the member of the genus *Filsoniana* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur, i.e. to *F. scarlatina* in the original publication (Kondratyuk et al. 2007). It was also compared with another Australian member of the same genus *Filsoniana* i.e.: *F. rexfilsonii* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur, but species of the genus *Nevilleiella* differs in the lack of oil droplets and oil cells in the paraphyses, and in having a very thin cortical layer, as well as much smaller, uniformly sized, ascospores in an ascus.

The taxonomic status of the second member of this genus, i.e. *Nevilleiella lateritia* is considered as rather questionable (Kantvilas 2016). Furthermore *Nevilleiella lateritia* considered as possible conspecific taxon with the *Filsoniana scarlatina* and *F. rexfilsonii*. However, molecular data confirm that these species have very good support as different species and that these taxa are members of two separate robust monophyletic branches (see also Kondratyuk et al. 2013b).

### *Opeltia* S. Y. Kondr. et L. Lőkös, gen. nov.

MycoBank nr.: MB 819652.

*Similar to the genus Blastenia, but differs in having zeorine and rusty brown apothecia, in having paraplectenchymatous true exciple, in having hymenium and especially subhymenium inspersed with oil, and in forming separate robust monophyletic branch in the Mikhtomia s.l. clade.*

Type species: *Opeltia neobaltistanica* (S. Y. Kondr. et J.-S. Hur) S. Y. Kondr. et L. Lőkös.

Thallus crustose, areolate to subsquamulose, margin abrupt at edge, without elongated lobes; whitish to dirty white, weakly yellowish whitish in places or yellowish orange, without asexual propagules or sorediate; protallus absent. Apothecia biatorine, zeorine to biatorine or lecanorine; disc dark reddish orange to rusty brown; true exciple paraplectenchymatous in basal portion to mesodermatous paraplectenchymatous with well-developed

matrix; hymenium and especially subhymenium inspersed with oil; ascospores 8-spored, ascospores polarilocular, often very varying in shape.

Chemistry: Parietin and emodin are recorded, while chemistry of all species of this genus is not studied in the same extent.

Ecology: It grows on bark of *Juniperus*, growing together with species of the genera *Candelariella*, *Caloplaca*, *Lecidella*, *Rinodina*, *Phaeophyscia*, and *Physconia*.

Etymology: The genus is named after the well-known German lichenologist Josef Poelt, in recognition of his enormous contribution to lichenology.

Distribution and species diversity: Two taxa, including the type species *O. neobaltistanica*, and *O. juniperina*, are known from Eurasia, while one species (*O. arizonica*) is known from southern North America.

Taxonomic notes and phylogenetic affiliations: After combined analysis the genus *Opeltia* is a member of the *Mikhtomia* s.l. clade of the Caloplacoidae, but forming separate robust monophyletic branch (Fig. 2). After molecular data hitherto available the genus *Opeltia* includes three species, i.e.: *O. arizonica*, *O. juniperina* and *O. neobaltistanica*.

After having yellow-orange areolate thallus it is similar to species of genus *Laundonia* (i.e. *L. flavovirescens*), while mainly growing on bark (vs. on rock), and with the exception of *O. arizonica* apothecia do not have visible thalline margin.

As it was mentioned earlier species of this genus were included mainly after ITS phylogeny to the genus *Gyalolechia* s.l. (Arup *et al.* 2013a), which was characterised by dominance of fragilin and presence of smaller amounts of parietin and emodin (often/sometimes with additional depsidones). However, as it was later shown the *Gyalolechia* s. str. branch including *G. aurea* and *G. canariensis* has the highest level of support, but this branch is in out position to the *Mikhtomia* s.l. clade (Kondratyuk *et al.* 2014c). The genus *Opeltia* forms separate robust monophyletic branch within the *Mikhtomia* s.l. clade.

***Oxneriopsis* S. Y. Kondr., D. Upreti et J.-S. Hur, gen. nov.**

Mycobank nr.: MB 819653.

*Similar to the genus Mikhtomia s. str., but differs in having more brownish orange to dark brown orange or even blackish brown-orange discs of apothecia and in forming separate robust monophyletic branch.*

Type species: *Oxneriopsis oxneri* (S. Y. Kondr. et Søchting) S. Y. Kondr., D. Upreti et J.-S. Hur.

Thallus crustose, continuous to cracked or areolated, often forming vegetative propagules, phyllidia or schizidia; greyish or greenish grey, while por-

tions with phyllidia and schizidia bright yellow or greenish yellow to dull yellow.

Apothecia lecanorine to zeorine, disc yellow, yellow-orange to orange-brown, dark brown or blackish brown; thalline margin as usual bright yellow; hymenium inspersed with oil; asci 8-spored; ascospores polarilocular, hyaline. Conidia bacilliform,  $2.5\text{--}3(3.5) \times 0.8\text{--}1 \mu\text{m}$ .

Chemistry: Fragilin (major) recorded for some taxa, however, chemistry of all taxa is not studied especially.

Ecology: It grows on bark of broad-leaved trees.

Distribution and species diversity: So far the genus includes Eastern Asian and North American taxa.

Etymology: It is named after the well-known Ukrainian lichenologist Alfred Mykolajovych Oxner (1898–1973) in recognition of his contribution to the Eurasian lichen flora.

Taxonomic notes and phylogenetic affinities: After ITS phylogeny species of genus *Oxneriopsis* were included into the genus *Gyalolechia* s.l. (Arup *et al.* 2013a). However, after combined phylogenetic analysis, based on nrITS, nrLSU and mtSSU sequences, the genus forms separate robust monophyletic branch within the *Mikhtomia* s.l. clade.

After molecular data hitherto available status of such species as *Oxneriopsis oxneri* and *O. yeosuensis* is confirmed within the genus. However, one more taxon (mentioned as *O. aff yeosuensis*) (see Fig. 2 and Appendix) is still under special revision and its status will be discussed separately.

It should be emphasised that within the genera *Oxneriopsis* and *Mikhtomia* we have the same tendency in forming phyllidia and schizidia, and two species *Oxneriopsis oxneri* and *Mikhtomia subflavorubescens* are sometimes morphologically totally the same. In this situation some specimens of *Mikhtomia subflavorubescens* were previously identified as “*Caloplaca oxneri*” (see Kondratyuk *et al.* 2014a) and superfluous combination “*Mikhtomia oxneri*” was proposed (Kondratyuk *et al.* 2014a). However, we should emphasise that combination was based on incorrectly identified specimens of *Mikhtomia subflavorubescens* and “*M. oxneri*” should be taken out from the usage. Correct place for this taxon is the *Oxneriopsis* branch. Fortunately Arup with colleagues (Arup *et al.* 2013a) have provided data on isotype specimens of the *Caloplaca oxneri*, and it allowed to clarify situation with real position of this taxon. After molecular data *Oxneriopsis oxneri* is very distinct, and appeared to be rather rare in South Korea for example (only a few specimens were confirmed as *Oxneriopsis oxneri*). At the same time *Mikhtomia subflavorubescens* is extremely common in this country, for which we have data more than 10 specimens, and almost half of specimens previously were incorrectly identified as “*M. oxneri*”. However, they all are confirmed to be *Mikhtomia subflavorubescens* after molecular phylogeny.

A separate paper on the molecular characters of the members of the genus *Mikhtomia* is still in progress (Kondratyuk *et al.* in prep.), however, some preliminary results are included here as proposals to new combinations (see below).

Unfortunately molecular data are still absent on such rare recently described taxon as '*Caloplaca*' *taranii* S. Y. Kondr., S. I. Tchabanenko, I. Galanina et L. Yakovczenko (Kondratyuk *et al.* 2013a), which differs in forming confluent soredious mass, but in general very similar to *Oxneriopsis oxneri*. May be the future involving this taxon in molecular phylogeny of the Teloschistaceae will confirm its status inside of the *Oxneriopsis* branch.

After morphological data we would expect that another Eastern Asian taxon '*Caloplaca*' *kiewkaensis* Yakovczenko, Galanina et S. Y. Kondr. (Kondratyuk *et al.* 2011), may also belong to this genus. However, we still do not have molecular data on this rare taxon.

It should be emphasised that in general the problem of molecular phylogeny of the *Mikhtomia* s.l. branch is that we do not have molecular data on the type species of the genus *Mikhtomia*, i.e. *M. gordejevii*, which was described in the 1920s and type specimen is very old. This species should be recollected in the type locality for the further extraction DNA and getting sequences for inclusion in the phylogenetic tree. In that case we will understand better importance of morphological and chemical characters for taxonomy of all genera of the *Mikhtomia* s.l. branch.

***Teuvoahtiana* S. Y. Kondr. et J.-S. Hur, gen. nov.**

Mycobank nr.: MB 819654.

*In having lobate, areolate and squamulose representatives in the same robust monophyletic branch *Teuvoahtiana* is similar to the Australian genus *Filsoniana* of the Teloschistoideae, but differs in positioning in the subfamily Xanthorioideae.*

Type species: *Teuvoahtiana rugulosa* (Nyl.) S. Y. Kondr. et J.-S. Hur.

Thallus crustose, areolate to squamulose, with rarely elongated in peripheral zone areoles to distinctly lobate; areolae and lobes usually very thick, convex, yellow-orange, dull yellowish brown to dull orange-brown, owing to numerous apothecia often dull reddish orange-brown; cortical layer paraplectenchymatous to mesodermatous paraplectenchymatous.

Apothecia of medium size to rather large, often densely aggregated, lecanorine to zeorine, disc plane, yellowish, roseus to pinkish orange-brown; true exciple of "textura intricata", asci (1-)2-4-6-8 spored, ascospores polaribilocular.

Chemistry is not studied in all members of this branch.

**Ecology:** It grows on non-calcareous rocks.

**Etymology:** It is named after the well-known Finnish lichenologist Teuvo Ahti (H, Helsinki, Finland) in recognition of his contribution to lichenology and especially to development of lichenological investigation in the South American continent.

**Distribution:** So far all members of this genus are distributed only within the South American continent.

**Taxonomic notes:** After combined phylogenetic analysis it is positioned in the Xanthorioideae as separate robust monophyletic branch or in some cases as sister branch to the *Xanthopeltis* branch (Fig. 3). The genus includes three South American species after molecular data hitherto available.

*Teuvoahtiana rugulosa* (Nyl.) S. Y. Kondr. et J.-S. Hur and *T. altoandiana* (Malme) S. Y. Kondr. et J.-S. Hur, as well as *T. fernandeziana* (Zahlbr.) S. Y. Kondr. et J.-S. Hur were compared to each other in original descriptions as the closest or the most similar taxa (Malme 1926, Zahlbruckner 1917). Molecular data entirely confirm their close relations.

Among taxa mentioned we have had difficulties only with identification of specimen of *Teuvoahtiana fernandeziana*, because our specimen differs in having somewhat thicker, somewhat *Toninia*-like thallus. However, as far details on ascospores and apothecia are within the *Teuvoahtiana fernandeziana* diagnosis (Malme 1926, Zahlbruckner 1917) we still hesitate to produce new taxon name. However, further molecular studies should clarify if position of this species in the genus *Teuvoahtiana* is correct, and if identification of specimen cited will be supported by further molecular data.

There are only a few genera in the Teloschistaceae, where we have distinctly lobate and crustose (elobate) taxa in the same monophyletic branches, i.e. the Australian genus *Filsoniana* (Teloschistoideae) and genera *Athallia*, *Gondwania*, *Orientophila* and *Teuvoahtiana* from the Xanthorioideae. In this case the distinctly lobate taxa *Filsoniana australiensis*, *Athallia scopularis*, *Orientophila subscopularis*, *O. yakjidoensis*, *O. jungakimae* and *Teuvoahtiana altoandina* are members of genera, where major portion of species diversity of the genera mentioned is represented by crustose representatives. In the genus *Gondwania* the majority of species is lobate taxa, while crustose lichens are also present (Kondratyuk *et al.* 2014c).

However, it should be mentioned that last data on *Orientophila* shows that this genus is more heterogeneous and more diverse as it was thought before (Arup *et al.* 2013a, Kondratyuk *et al.* 2013b, 2016d) and probably in the future the genus *Orientophila* will contain larger number of crustose species, or it will be divided into two or more separate branches. Special revision of this genus is also in progress (Kondratyuk *et al.* 2017).

Additionally to *Teuvoahtiana rugulosa*, *T. altoandina* and *T. fernandeziana* there are also undescribed taxa as '*Caloplaca* sp. 2' and '*Caloplaca* sp. 4' (sensu

Arup *et al.* 2013a), which belong to this monophyletic branch. However, they are still not legally described (not shown in the tree) unfortunately.

Arup with colleagues (Arup *et al.* 2013a) have recognised only *Caloplaca altoandina* and 4 undescribed taxa (mentioned as *Caloplaca* sp. 1–4), which were in sister position to the genus *Xanthopeltis*. After data of these authors monophyletic branch with *Xanthopeltis* has had high level of support, while *C. altoandina* did not have good support based on their matrix at that time. Taxa mentioned as *Caloplaca* sp. 1 and *Caloplaca* sp. 3 were included in this branch only after ITS phylogeny, while data on nrLSU and mtSSU were (and still are) missing.

Our data confirm previous data that *Teuvoahtiana* is closely related to the genera *Xanthopeltis* and *Austroplaca*. However, on another side, our data do not confirm previous data that *Austroplaca* is positioned in one clade with *Gondwania* and *Cerothallia*. The latter two genera are in somewhat out position to *Xanthomendoza* s.l. subphylum of the Xanthorioideae after our phylogenetic analysis.

***Tomnashia* S. Y. Kondr. et J.-S. Hur, *gen. nov.***

Mycobank nr.: MB 819655.

*Similar to genus Blastenia of the subfamily Caloplacoideae, but differs in having often yellow to yellowish orange or apricot orange thallus and mainly orange apothecia, as well as forming separate robust monophyletic branch in the Xanthorioideae.*

Type species: *Tomnashia rosei* (Hasse) S. Y. Kondr. et J.-S. Hur.

Thallus crustose, immersed into calcareous rock or absent, to continuous or areolate, white, grey, greenish yellow or yellowish orange to orange or apricot orange, in some taxa (*T. ludificans*) with a waxy translucent appearance, without asexual propagules. Prothallus absent, or yellow-orange if present.

Apothecia small biatorine, disc orange to reddish brown; true exciple consisting of radiating hyphae, in basal portion of “textura intricata”; asci 8-spored; ascospores polarilocular, hyaline.

Chemistry: Parietin, fallacinal, emodin, teloschistin and parietinic acid.

Ecology: It grows on calcareous rock, on soil and on non-calcareous rocks.

Eymology: It is named after the well-known American lichenologist Thomas H. Nash III (1945–) in recognition of his contribution to lichenology, especially of the North American lichen flora.

Distribution: Species of this genus are hitherto known only from North America, where mainly distributed in the southwestern part of the continent.

The genus *Tomnashia* similarly to the genera *Polycauliona* s. str. and *Igneoplaca* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur of the *Polycauliona* s.l. subphylum has rather limited distribution in coastal southwestern North America, while the other genera of the *Polycauliona* s.l. subphylum, like *Mass-*

*jukiella* S. Y. Kondr., Fedorenko, S. Stenroos, Kärnefelt, Elix, J.-S. Hur et A. Thell, *Verrucoplaca* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur, *Scythioria* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur are characterised by much wider distribution in the Northern Hemisphere or in the world.

Taxonomic notes and phylogenetic affiliations: After combined phylogenetic analysis the genus *Tomnashia* is positioned in the outermost position among monophyletic groups of the *Polycauliona* s.l. subclade (i.e. *Verrucoplaca*, *Igneoplaca*, *Massjukiella*) of the Xanthorioideae, which has rather low level of support (see tree Xanthorioideae), and in the most distant from the *Polycauliona* s. str. (i.e. *P. coralloides*) branch (Fig. 3).

From other members of the *Polycauliona* s.l. subphylum *Tomnashia* differs in having crustose, continuous, from immersed and indistinct to distinctly areolate thallus without elongated areoles at the margin, as well as mainly bright orange to reddish brown biatorine apothecia.

In having biatorine apothecia may be similar to the members of the genera *Blastenia* A. Massal. and *Eilidahlia* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur, as well as of *Laundonia* S. Y. Kondr., L. Lőkös et J.-S. Hur and *Mikhtomia* S. Y. Kondr., Kärnefelt, Elix, A. Thell et J.-S. Hur of the subfamily Caloplacoideae, but the genus *Tomnashia* is positioned in separate robust monophyletic branch of the Xanthorioideae.

Additionally to *Tomnashia rosei*, *T. nashii*, *T. luteominia* and *T. ludificans* this branch includes also '*Polycauliona* sp. 37' (after Arup et al. 2013a), but this taxon is still not described (and not included in the final tree).

### *Xanthaptychia* S. Y. Kondr. et S. Ravera, *gen. nov.*

MycoBank nr.: MB 819656.

*Similar to the genus Seirophora, but differs in having scleroplectenchymatous tissue in thallus and cortex of thalline margin of apothecia, and in distribution mainly in high altitude of mountainous regions of northern Eurasian or North American, as well as in forming separate robust monophyletic branch.*

Type species: *Xanthaptychia orientalis* (Frödén) S. Y. Kondr. et S. Ravera.

Thallus foliosus to subfruticose or caespitose to pulvinate, usually rosette-forming, small; lobes dorsiventral, horizontally orientated and three different portions (main lobes, secondary lobules and terminal portions observed (see also Kondratyuk et al. 2013c)), or overlapping and often ascending, branching mostly irregular to anastomosing, rarely podetium-like; upper surface whitish grey, often brownish grey partly brownish yellow to yellow, lower side whitish grey; tomentum often well developed on upper surface; in

section cortical layer scleroplectenchymatous; lower cortex absent or developed only on small portions; some taxa with various vegetative propagules.

Apothecia laminal, lecanorine, usually very large, to 5(–10) mm diam., disc yellow, orange to reddish orange or brownish orange; cortex scleroplectenchymatous; asci 8-spored, polarilocular with narrow septa, hyaline; conidia narrowly bacilliform, 3–3.8 × 0.9–1 µm. [Note: data of Frödén and Litterski (2005) on conidia bacilliform to narrowly ellipsoid, (3–)3.5–4(4.5) × (1–)1.4–1.8(–2) µm for *Xanthaptychia orientalis* are very doubtful and they are need in verifying].

Chemistry: Parietin (major) and low concentrations of emodin, fallacinal, teloschistin, parietinic acid and erythroglauclin.

Ecology: Corticolous species growing on *Picea schrenkiana*, *Ephedra canisetina*, *Acer pubescens*, *A. regalis*, *Rhamnus sintesii*, *Pistacia vera*, *Sageretia laetevirens*, *Amygdalus buharica*, and species of the genera *Populus*, *Juniperus* and *Acer* in montane belt between altitudes 1,100–2,100 m.

Etymology: It is named after similarities with members of the genus *Anaptychia* of the Physciaceae (i.e. lack of lower cortical layer) and xanthorioid lichens of the Teloschistaceae (in having foliose thallus).

Distribution: Northern Hemisphere with tendency to high altitudes of mountainous regions or polar latitudes of Eurasia and North America. Soredious taxa *Xanthaptychia contortuplicata* and *X. blumii* supposed to have wider distribution than the esoredious species *X. orientalis*. However, the latter two taxa are rather recently described and we still have limited data on their distribution (see Kondratyuk *et al.* 2013c, 2015b, 2016b). In the Mediterranean region species of the genera *Xanthaptychia* and *Seirophora* (i.e.: *Xanthaptychia orientalis* and *Seirophora villosa*) are completely allopatric.

Taxonomic notes and phylogenetic affiliations: After combined phylogenetic analysis the genus positioned in the *Seirophora* s.l. clade of the Caloplacoideae, where it is forming separate robust monophyletic branch (Fig. 2). From molecular data hitherto available the genus *Xanthaptychia* includes four species, i.e.: *X. blumii*, *X. aurantiaca*, *X. contortuplicata* and *X. orientalis*, while status of the other *Seirophora* species including *S. scorigena*, *S. tenera* and others is still waiting for clarifying.

Similar to the genus *Seirophora*, but differs in having scleroplectenchymatous tissue in thallus and cortex of thalline margin of apothecia (vs. prosoplectenchymatous), and in distribution mainly in high altitude of mountainous regions of northern Eurasian or North American (vs. Mediterranean region with preference to its western regions), as well as in forming separate robust monophyletic branch.

From 2013 the position of genus *Seirophora* in phylogenetic tree of the Teloschistaceae was somewhat questionable because in fact this genus was not

confirmed by three gene phylogeny. Only data on ITS nrDNA were provided for the type species of this genus, i.e. *S. villosa* (Arup *et al.* 2013a). Within our study it was special task to provide missing data on nrLSU and mtSSU data for *S. villosa*. After getting these data (see Appendix and Fig. 2) it became clear that the *Xanthaptychia* branch formed a separate robust monophyletic branch.

The type species of this genus, *Xanthaptychia orientalis* appeared to be described twice almost in the same time by Frödén and Litterski (2005) and one year later by Kondratyuk and Kudratov (2006) as *Xanthoanaptychia kotovii* S. Y. Kondr. et Kudratov (while manuscripts of both publications were submitted in 2005, in the same year). Description of the *Xanthoanaptychia kotovii* was based on specimens kept in KW-L and LE and it was somewhat wider and more detailed than in the earlier paper. In this paper type species of the genus *Xanthaptychia* was compared with *X. contortuplicata* and *Seirophora villosa*, as well as with *Massjukiella polycarpa* (Hoffm.) S. Y. Kondr., Fedorenko, S. Stenroos, Kärnefelt, Elix, J.-S. Hur et A. Thell (Xanthorioideae) and *Niorma chrysopthalma* (L.) S. Y. Kondr., Kärnefelt, Elix, A. Thell, N.-H. Jeong et J.-S. Hur (Teloschistoideae). Unfortunately data on conidia of the type species of the genus *Xanthaptychia* are especially different and they are in urgent need of repeated revision (see above).

In both original papers similarities and differences with *S. villosa* and *S. lacunosa* were emphasised, as well as *S. scorigena*. However, after segregation of the genus *Xanthaptychia* in separate genus all these characters are in need of the further revision.

Status of *Seirophora tenera* Frödén et Litterski is still waiting for clarifying, while some information was published that data on ITS region of *S. tenera* were obtained (Frödén and Litterski 2005). However, these data are still not available for wide access.

### New combinations

*Elixjohnia bermaguiiana* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr. et J.-S. Hur, comb. nova – MycoBank nr.: MB 819657 – Basionym: *Caloplaca bermaguiiana* S. Y. Kondr. et Kärnefelt, in Kondratyuk, Kärnefelt, Elix and Thell, Bibl. Lichenol. 95: 348 (2007).

*Elixjohnia gallowayi* (S. Y. Kondr., Kärnefelt et Filson) S. Y. Kondr. et J.-S. Hur, comb. nova – MycoBank nr.: MB 819658 – Basionym: *Caloplaca gallowayi* S. Y. Kondr., Kärnefelt et Filson, in Kondratyuk, Kärnefelt, Elix and Thell, Bibl. Lichenol. 95: 358 (2007).

*Elixjohnia jackelixii* (S. Y. Kondr., Kärnefelt et A. Thell) S. Y. Kondr. et J.-S. Hur, comb. nova – MycoBank nr.: MB 819660 – Basionym: *Caloplaca jackelixii* S. Y. Kondr., Kärnefelt et A. Thell, in Kondratyuk, Kärnefelt, Elix and Thell, Bibl. Lichenol. 100: 251 (2009).

*Fominiella skii* (Khodos., Vondrák et Šoun) S. Y. Kondr., D. Upreti et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819661. – Basionym: *Caloplaca skii* Khodos., Vondrák et Šoun, in Vondrák, Khodosovtsev, Šoun and Vondráková, Lichenologist 44(1): 83 (2011) (2012). ≡ *Athallia skii* (Khodos., Vondrák et Šoun) Arup, Frödén et Søchting (2013).

*Gintarasiella aggregata* (Kantvilas et S. Y. Kondr.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819697 – Basionym: *Caloplaca aggregata* Kantvilas et S. Y. Kondr., in Kantvilas, J. Adelaide Bot. Gard. 29: 56 (2016).

*Hanstrassia lenae* (Søchting et G. Figueras) S. Y. Kondr., *comb. nova* – MycoBank nr.: MB 819662 – Basionym: *Caloplaca lenae* Søchting et G. Figueras, Lichenologist 39(1): 8 (2007). ≡ *Gyalolechia lenae* (Søchting et G. Figueras) Søchting, Frödén et Arup, Nordic J. Bot. 31(1): 71 (2013).

*Ikaeria aurantiellina* (Harm.) S. Y. Kondr., D. Upreti et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819663 – Basionym: *Caloplaca aurantiellina* Harm., in Pitard and Harmand, Bull. Soc. bot. Fr. 58 (Mém. no. 22): 46 (1911).

*Klauderuiella aurantia* (Pers.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819664 – Basionym: *Lichen aurantius* Pers., Ann. Bot. (Usteri) 5: 14 (1794). ≡ *Variospora aurantia* (Pers.) Arup, Frödén et Søchting, Nordic J. Bot. 31(1): 76 (2013).

*Klauderuiella flavescens* (Huds.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819665 – Basionym: *Lichen flavescens* Huds., Fl. Angl.: 445 (1762). ≡ *Variospora flavescens* (Huds.) Arup, Frödén et Søchting, Nordic J. Bot. 31(1): 76 (2013).

*Klauderuiella thallincola* (Wedd.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819666 – Basionym: *Lecanora murorum* var. *thallincola* Wedd., Mém. Soc. natn. Sci. nat. Cherbourg 19: 274 (1875). ≡ *Variospora thal-lincola* (Wedd.) Arup, Frödén et Søchting, Nordic J. Bot. 31(1): 77 (2013).

*Laundonia flavovirescens* (Wulfen) S. Y. Kondr., L. Lőkös et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819667 – Basionym: *Lichen flavovirescens* Wulfen, Schr. naturf. Freunde, Berlin 8: 122 (1787). ≡ *Gyalolechia flavovirescens* (Wulfen) Søchting, Frödén et Arup, Nordic J. Bot. 31(1): 70 (2013).

*Laundonia persimilis* (Wetmore) S. Y. Kondr., L. Lőkös et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819668 – Basionym: *Caloplaca persimilis* Wetmore, Bryologist 107(4): 513 (2004). ≡ *Gyalolechia persimilis* (Wetmore) Søchting, Frödén et Arup, Nordic J. Bot. 31(1): 71 (2013).

*Lazarenkoiopsis ussuriensis* (Oxner, S. Y. Kondr. et Elix) S. Y. Kondr., L. Lőkös et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819695 – Basionym: *Caloplaca ussuriensis* Oxner, S. Y. Kondr. et Elix, Folia cryptog. Estonica 48: 21 (2011).

*Nevilleiella marchantii* (S. Y. Kondr. et Kärnefelt) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819670 – Basionym: *Caloplaca marchantii* S. Y. Kondr. et Kärnefelt, in Kondratyuk, Kärnefelt, Elix and Thell, Bibl. Lichenol. 95: 367 (2007).

*Nevilleiella lateritia* (Taylor) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819671 – Basionym: *Lecidea lateritia* Taylor, J. Bot (Hooker) 6: 149 (1847). ≡ *Caloplaca lateritia* (Taylor) Zahlbr., Cat. Lich. Univ. 7: 154 (1930).

*Opeltia arizonica* (H. Magn.) S. Y. Kondr. et L. Lőkös, *comb. nova* – MycoBank nr.: MB 819672 – Basionym: *Caloplaca arizonica* H. Magn., Bot. Notiser, p. 69 (1944). ≡ *Gyalolechia arizonica* (H. Magn.) Søchting, Frödén et Arup, Nordic J. Bot. 31(1): 70 (2013)

*Opeltia juniperina* (Tomin) S. Y. Kondr. et L. Lőkös, *comb. nova* – MycoBank nr.: MB 819673 – Basionym: *Caloplaca juniperina* Tomin, Bot. Materialy (Notul. System. E Sect. Cryptog. Inst. Bot. nomine V. L. Komarovii Acad. Sci. URSS, 11: 11(1953). ≡ *Gyalolechia juniperina* (Tomin) Søchting, Frödén et Arup, Nordic J. Bot. 31(1): 71 (2013).

*Opeltia neobaltistanica* (S. Y. Kondr. et J.-S. Hur) S. Y. Kondr. et L. Lőkös, *comb. nova* – MycoBank nr.: MB 819674 – Basionym: *Caloplaca neobaltistanica* S. Y. Kondr. et J.-S. Hur, in Kondratyuk, Lőkös, Farkas, Oh and Hur, Acta Bot. Hung. 57(1–2): 89 (2015).

*Oxneriopsis oxneri* (S. Y. Kondr. et Søchting) S. Y. Kondr., D. Upreti et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819675 – Basionym: *Caloplaca oxneri* S. Y. Kondr. et Søchting, in Kondratyuk, Søchting and Kärnefelt, Nat. Hist. Res. 4(1): 17 (1996). ≡ *Gyalolechia oxneri* (S. Y. Kondr. et Søchting) Søchting, Frödén et Arup, Nordic J. Bot. 31(1): 71 (2013).

*Oxneriopsis yeosuensis* (S. Y. Kondr. et J.-S. Hur) S. Y. Kondr., D. Upreti et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819696 – Basionym: *Caloplaca yeosuensis* S. Y. Kondr. et J.-S. Hur, in Kondratyuk, Lőkös, Zarei-Darki, Haji Moniri, Tchabanenko, Galanina, Yakovchenko, Hooshmand, Ezhkin and Hur, Acta Bot. Hung. 55(1–2): 52 (2013) nom invalid., validated here – MB 819676 – Type: South Korea, Jeollanam-do, Yeosu-si, Nam-myeon, Geumoh-do, Dumori, Jickpo coast, on rock, growing together with *Caloplaca diffluens*. Lat.: 34° 30' 45.00" N; Long.: 127° 44' 14.08" E; Alt.: 6 m a.s.l. Coll.: U. Jayalal et al. (120360), 26.04.2012. Holotype: KoLRI 015350; Isotypes: KoLRI 015346 (120357); KoLRI 015349 (120359); KoLRI 015325 (120336); KoLRI 015321 (120332); KoLRI 015369 (120380); KoLRI 015367 (120378); KoLRI 015359 (120370).

*Squamulea micromera* (Hue) S. Y. Kondr., L. Lőkös et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819677 – Basionym: *Lecanora micromera* Hue, Ann. Mycol. 13: 82 (1915). ≡ *Caloplaca micromera* (Hue) Zahlbr., Cat. Lich. Univers. 7: 157 (1931). – Type: [Japan], Mt Jizagatake prope Kofu, 07.1903 Urbain Faurie No 5651 (KYO 00031292 – isotype, designated here).

*Teuvoahhtiana altoandina* (Malme) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819678 – Basionym: *Callopisma altoandinum* Malme, Ark. Bot. 20A(9): 18 (1926). ≡ *Caloplaca altoandina* (Malme) Zahlbr., Cat. Lich. Univers. 7: 211 (1931).

*Teuvoahtiana fernandeziana* (Zahlbr.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819679 – Basionym: *Blastenia fernandeziana* Zahlbr., K. svenska Vetensk-Akad. Handl. 57(6): 46 (1917). ≡ *Caloplaca fernandeziana* (Zahlbr.) Follmann et Redón, Willdenowia 6(3): 448 (1972).

*Teuvoahtiana rugulosa* (Nyl.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819680 – Basionym: *Placodium rugulosum* Nyl., Annls Sci. Nat., Bot., sér. 4, 3: 153 (1855). ≡ *Caloplaca rugulosa* (Nyl.) Zahlbr., Cat. Lich. Univers. 7: 263 (1931).

*Tomnashia ludificans* (Arup) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819681 – Basionym: *Caloplaca ludificans* Arup, Bryologist 98(1): 107 (1995). ≡ *Polycauliona ludificans* (Arup) Arup, Frödén et Söchting, Nordic J. Bot. 31(1): 52 (2013).

*Tomnashia luteominia* (Tuck.) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819682 – Basionym: *Placodium luteominium* Tuck. (as ‘*luteominium*’), Lichens of California (Berkeley): 18 (1866). ≡ *Polycauliona luteominia* (Tuck.) Arup, Frödén et Söchting, Nordic J. Bot. 31(1): 52 (2013).

*Tomnashia nashii* (Nav.-Ros., Gaya et Hladún) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819683 – Basionym: *Caloplaca nashii* Nav.-Ros., Gaya et Hladún, Mycotaxon 79: 31 (2001). ≡ *Polycauliona nashii* (Nav.-Ros., Gaya et Hladún) Arup, Frödén et Söchting, Nordic J. Bot. 31(1): 53 (2013).

*Tomnashia rosei* (Hasse) S. Y. Kondr. et J.-S. Hur, *comb. nova* – MycoBank nr.: MB 819684 – Basionym: *Caloplaca rosei* Hasse, Bryologist 14: 102 (1911). ≡ *Polycauliona rosei* (Hasse) Arup, Frödén et Söchting, Nordic J. Bot. 31(1): 53 (2013).

*Variospora latzelii* (Servít) S. Y. Kondr., *comb. nova* – MycoBank nr.: MB 819685 – Basionym: *Blastenia latzelii* Servít (as ‘*latzeli*’), Hedwigia 74: 151 (1934). ≡ *Caloplaca latzelii* (Servít) Clauzade et Cl. Roux, Bull. Soc. bot. Centre-Ouest, Nouv. sér., num. spec. 7: 824 (1985).

*Xanthaptychia aurantiaca* (R. Br.) S. Y. Kondr. et S. Ravera, *comb. nova* – MycoBank nr.: MB 819686 – Basionym: *Borrera aurantiaca* R. Br., Suppl. to the Appendix of Cpt. Parrys Voyag., Nat. Hist.: CCCV (1824). ≡ *Seirophora aurantiaca* (R. Br.) Frödén, in Frödén et Lassen, Lichenologist 36(5): 295 (2004).

*Xanthaptychia blumii* (S. Y. Kondr. et Moniri) S. Y. Kondr. et S. Ravera, *comb. nova* – MycoBank nr.: MB 819785 – Basionym: *Seirophora blumii* S. Y. Kondr. et Moniri, in Kondratyuk, Lőkös, Tschabanenko, Moniri and Farkas, Acta Bot. Hung. 55(3–4): 275–349 (2013).

*Xanthaptychia contortuplicata* (Ach.) S. Y. Kondr. et S. Ravera, *comb. nova* – MycoBank nr.: MB 819688 – Basionym: *Parmelia contortuplicata* Ach., Syn. meth. lich. (Lund): 210 (1814). ≡ *Seirophora contortuplicata* (Ach.) Frödén, in Frödén and Lassen, Lichenologist 36(5): 297 (2004).

*Xanthaptychia orientalis* (Frödén) S. Y. Kondr. et S. Ravera, *comb. nova* – MycoBank nr.: MB 819689 – Basionym: *Seirophora orientalis* Frödén, in Frödén

and Litterski, *Graphis Scripta* 17(1): 22 (2005). = *Xanthoanaptychia kotovii* S. Y. Kondr. et Kudratov, Ukr. Botan. Zh. 63(3): 342 (2006).

*Xanthocarpia raesaenenii* (Bredkina) S. Y. Kondr., comb. nova – MycoBank nr.: MB 819690 – Basionym: *Caloplaca raesaenenii* Bredkina, Nov. sist. Niz. Rast. 23: 170 (1986). – Replaced synonym: *Placodium geophilum* Räsänen, Ann. bot. Soc. Zool.-Bot. fenn. Vanamo 12(1): 1 (1938), Nom. illegit., Art. 53.1 non *Placodium geophilum* Th. Fr. (1891).

## CONCLUSIONS

Seventeen newly discovered robust monophyletic branches proved by combined phylogenetic analysis based on ITS nr DNA, 28S nrLSU and 12S mtSSU, which are proposed to consider as separate genera are added to the sub-families Teloschistoideae (5), Caloplacoideae (7), Xanthorioideae (3) and Brown-lielloideae (1). There are also genera *Fominiella* and *Ikaeria*, which are positioned in out position to known subfamilies and possibly illustrate that additional sub-families of the Teloschistaceae still exist and are waiting for legal description.

\*

*Acknowledgements* – We are thankful to Ingvar Kärnefelt and Arne Thell (both Lund, Sweden) for kind help with checking some names, and comments to some taxa, to Anna Guttová (Bratislava, Slovak Republic), Lidia Yakovchenko (Vladivostok, Russia) and Gintaras Kantvilas (Hobart, Tasmania) for providing fresh collections, to curator of KYO herbarium for the loan of type specimens, and to Konstanze Bensch (The MycoBank Team, UK) for valuable nomenclatural comments and help. SK and LL are grateful to Prof. J.-S. Hur and Dr S.-O. Oh for assistance and support during visits to South Korea in 2014, 2015 and 2016. The project was supported by the Ministry of Education and Science of Ukraine (M/90-2015-285 and M/34-2016-285) and by the Korean Brain Pool Program (161S-4-3-1659) for SK, and the Korea National Research Resource Center Program, the Korean Forest Service Program (KNA 2012-2016) through the Korea National Arboretum, and (for LL) also by the Hungarian Scientific Research Fund (OTKA K81232).

## REFERENCES

- Arup, U. (2006): A new taxonomy of the *Caloplaca citrina* group in the Nordic countries, except Iceland. – *Lichenologist* 38(1): 1–20. <https://doi.org/10.1017/S0024282905005402>
- Arup, U. (2009): The *Caloplaca holocarpa* group in the Nordic countries, except Iceland. – *Lichenologist* 41(2): 111–130. <https://doi.org/10.1017/s0023282909008135>
- Arup, U. and Grube, M. (1999): Where does *Lecanora demissa* (Ascomycota, Lecanorales) belong? – *Lichenologist* 31: 419–430. <https://doi.org/10.1017/s0024282999000584>
- Arup, U., Søchting, U. and Frödén, P. (2013a): A new taxonomy of the family Teloschistaceae. – *Nordic J. Bot.* 31: 16–83. <https://doi.org/10.1111/j.1756-1051.2013.00062.x>
- Arup, U., Søchting, U. and Frödén, P. (2013b): Addendum to ‘A new taxonomy of the family Teloschistaceae’. *Nordic J. Bot.* 31: 256. <https://doi.org/10.1111/j.1756-1051.2013.00295.x>

- Clauzade, G. and Roux, C. (1985): Likenoj de Okcidenta Eŭropo. Illustrata determinlibro. – *Bull. Soc. bot. Centre-Ouest, Royan* 7: 1–893.
- Ekman, S. (1999): PCR optimization and troubleshooting, with special reference to the amplification of ribosomal DNA in lichenized fungi. – *Lichenologist* 31(5): 517–531. <https://doi.org/10.1017/s002428299000675>
- Fedorenko, N. M., Stenroos, S., Thell, A., Kärnefelt, I. and Kondratyuk, S. Y. (2009): A phylogenetic analysis of xanthorioid lichens (Teloschistaceae, Ascomycota) based on ITS and mtSSU sequences. – *Bibl. Lichenol.* 100: 49–84.
- Fedorenko, N. M., Stenroos, S., Thell, A., Kärnefelt, I., Elix, J. A., Hur, J. S. and Kondratyuk, S. Y. (2012): Molecular phylogeny of xanthorioid lichens (Teloschistaceae, Ascomycota), with notes on their morphology. – *Bibl. Lichenol.* 108: 45–64.
- Frödén, P. and Lassen, P. (2004): Typification and emendation of Seirophora Poelt to include species segregated from Teloschistes Norman. – *Lichenologist* 36: 289–298. <https://doi.org/10.1017/S002428290401432X>
- Frödén, P. and Litterski, B. (2005): Two new species of Seirophora from central Asia. – *Gravis Scripta* 17: 22–26.
- Gardes, M. and Bruns, T. D. (1993): ITS primers with enhanced specificity for basidiomycetes – application to the identification of mycorrhizae and rusts. – *Mol. Ecol.* 2: 113–118. <https://doi.org/10.1111/j.1365-294x.1993.tb00005.x>
- Gaya, E., Lutzoni, F., Zoller, S. and Navarro-Rosinés, P. (2003): Phylogenetic study of Fulgensia and allied Caloplaca and Xanthoria species (Teloschistaceae, lichen-forming Ascomycota). – *Amer. J. Bot.* 90(7): 1095–1103. <https://doi.org/10.3732/ajb.90.7.1095>
- Gaya, E., Navarro-Rosinés, P., Llimona, X., Hladun, N. and Lutzoni, F. (2008): Phylogenetic reassessment of the Teloschistaceae (lichen-forming Ascomycota, Lecanoromycetes). – *Mycol. Res.* 112(5): 528–546. <https://doi.org/10.1016/j.mycres.2007.11.005>
- Gaya, E., Högnabba, F., Holguín, Á., Molnár, K., Fernández-Brime, S., Stenroos, S., Arup, U., Søchting, U., van den Boom, P., Lücking, R., Sipman, H. J. M. and Lutzoni, F. (2012): Implementing a cumulative supermatrix approach for a comprehensive phylogenetic study of the Teloschistales (Pezizomycotina, Ascomycota). – *Mol. Phyl. Evol.* 63: 374–387. <https://doi.org/10.1016/j.ympev.2012.01.012>
- Gaya, E., Fernández-Brime, S., Vargas, R., Lachlan, R. F., Gueidan, C., Ramírez-Mejía, M. and Lutzoni, F. (2015): The adaptive radiation of lichen-forming Teloschistaceae is associated with sunscreening pigments and a bark-to-rock substrate shift. – *Proc. Nat. Acad. Sci. USA* 112(37): 11600–11605. <https://doi.org/10.1073/pnas.1507072112>
- Joshi, Y., Vondrák, J., Vondráková, O., Nguyen, T. T. and Hur, J.-S. (2011): Caloplaca allochroa (lichenized Ascomycetes), a new saxicolous lichen species from South Korea. – *Mycotaxon* 117: 261–267. <https://doi.org/10.5248/117.261>
- Kantvilas, G. (2016): A synopsis and key for the lichen genus Caloplaca (Teloschistaceae) on Kangaroo Island, with the description of two new species. – *J. Adelaide Bot. Gard.* 29: 53–69.
- Kärnefelt, I. (1989): Morphology and phylogeny in the Teloschistales. – *Cryptogamic Botany* 1: 147–203.
- Kondratyuk, S., Søchting, U. and Kärnefelt, I. (1996): Caloplaca oxneri (Teloschistaceae), a new lichen species from East Asia. – *Nat. Hist. Res.* 4: 17–20.
- Kondratyuk, S. Y., Elix, J. A., Galanina, I. A., Yakovchenko, L. S., Kärnefelt, I. and Thell, A. (2011): Four new Caloplaca species (Teloschistaceae, Ascomycotina). – *Folia Cryptog. Est.* 48: 17–23.
- Kondratyuk, S. and Kudratov, I. (2003): Three new Caloplaca species from Asia. – *Ukr. Bot. Zhurn.* 60(1): 65–69.

- Kondratyuk, S. and Kudratov, I. (2006): To revision of Middle Asian material of the 'Teloschistes brevior' complex (Teloschistaceae). – *Ukr. Bot. Zhurn.* **63**(3): 340–350.
- Kondratyuk, S., Kärnefelt, I., Elix, J. A. and Thell, A. (2007): New species of the genus Caloplaca in Australia. – *Bibl. Lichenol.* **95**: 341–386.
- Kondratyuk, S. Y., Kärnefelt, I., Elix, J. A. and Thell, A. (2009): Contributions to the Teloschistaceae, with particular reference to the Southern Hemisphere. – *Bibl. Lichenol.* **100**: 207–282.
- Kondratyuk, S. Y., Elix, J. A., Kärnefelt, I. and Thell, A. (2012): An artificial key to Australian Caloplaca species (Teloschistaceae, Ascomycota). – *Bibl. Lichenol.* **108**: 141–160.
- Kondratyuk, S. Y., Lőkös, L., Zarei-Darki, B., Haji Moniri, M., Tchabanenko, S. I., Galanina, I., Yakovchenko, L., Hooshmand, F., Ezhkin, A. K. and Hur, J.-S. (2013a): Five new Caloplaca species (Teloschistaceae, Ascomycota) from Asia. – *Acta Bot. Hung.* **55**: 41–60. <https://doi.org/10.1556/ABot.55.2013.1-2.4>
- Kondratyuk, S., Jeong, M. H., Yu, N. H., Kärnefelt, I., Thell, A., Elix, J. A., Kim, J., Kondratyuk, A. S. and Hur, J.-S. (2013b): Four new genera of teloschistoid lichens (Teloschistaceae, Ascomycota) based on molecular phylogeny. – *Acta Bot. Hung.* **55**: 251–274. <https://doi.org/10.1556/ABot.55.2013.3-4.8>
- Kondratyuk, S., Lőkös, L., Tschabanenko, S., Haji Moniri, M., Farkas, E., Wang, X. Y., Oh, S.-O. and Hur, J.-S. (2013c): New and noteworthy lichen-forming and lichenicolous fungi. – *Acta Bot. Hung.* **55**: 275–349. <https://doi.org/10.1556/ABot.55.2013.3-4.9>
- Kondratyuk, S. Y., Jeong, M. H., Yu, N. N., Kärnefelt, I., Thell, A., Elix, J. A., Kim, J., Kondratyuk, A. S. and Hur, J.-S. (2014a): A revised taxonomy for the subfamily Caloplaeoideae (Teloschistaceae, Ascomycota) based on molecular phylogeny. – *Acta Bot. Hung.* **56**: 93–123. <https://doi.org/10.1556/ABot.56.2014.1-2.10>
- Kondratyuk, S. Y., Lőkös, L., Tschabanenko, S., Skirina, I., Galanina, I., Oh, S.-O. and Hur, J.-S. (2014b): Caloplaca kedrovopadensis sp. nova and some new lichens from the Primorsky region, Russia. – *Acta Bot. Hung.* **56**: 125–140. <https://doi.org/10.1556/ABot.56.2014.1-2.11>
- Kondratyuk, S. Y., Kärnefelt, I., Thell, A., Elix, J. A., Kim, J., Jeong, M. H., Yu, N. H., Kondratyuk, A. S. and Hur, J.-S. (2014c): A revised taxonomy of the subfamily Xanthorioideae (Teloschistaceae, Ascomycota) based on molecular phylogeny. – *Acta Bot. Hung.* **56**: 141–178. <https://doi.org/10.1556/ABot.56.2014.1-2.12>
- Kondratyuk, S. Y., Kärnefelt, I., Thell, A., Elix, J. A., Kim, J., Kondratyuk, A. S. and Hur, J.-S. (2015a): Tassiloa, a new genus in the Teloschistaceae (lichenized Ascomycetes). – *Graptis Scripta* **27**: 22–26.
- Kondratyuk, S. Y., Lőkös, L., Farkas, E., Oh, S.-O. and Hur, J.-S. (2015b): New and noteworthy lichen-forming and lichenicolous fungi, 2. – *Acta Bot. Hung.* **57**(1–2): 77–141. <https://doi.org/10.1556/ABot.57.2015.1-2.10>
- Kondratyuk, S. Y., Lőkös, L., Kim, J. A., Kondratyuk, A. S., Jeong, M. H., Jang, S. H., Oh, S.-O. and Hur, J.-S. (2015c): Three new monotypic genera of the caloplacoid lichens (Teloschistaceae, lichen-forming Ascomycetes). – *Mycobiology* **43**: 195–202. <https://doi.org/10.5941/myco.2015.43.3.195>
- Kondratyuk, S. Y., Kärnefelt, I., Thell, A., Elix, J. A., Kim, J., Kondratyuk, A. S. and Hur, J.-S. (2015d): Brownlielloideae, a new subfamily in the Teloschistaceae (Lecanoromycetes, Ascomycota). – *Acta Bot. Hung.* **57**: 321–341. <https://doi.org/10.1556/034.57.2015.3-4.6>
- Kondratyuk, S. Y., Kim, J. A., Yu, N.-H., Jeong, M.-H., Jang, S. H., Kondratyuk, A. S., Zarei-Darki, B. and Hur, J.-S. (2015e): Zeroviella, a new genus of xanthorioid lichens (Teloschistaceae, Ascomycota) proved by three gene phylogeny. – *Ukr. Bot. J.* **72**(6): 574–584. <https://doi.org/10.15407/ukrbotj72.06.574>

- Kondratyuk, S. Y., Kärnefelt, I., Thell, A., Elix, J. A., Kim, J. A., Kondratyuk, A. S. and Hur, J.-S. (2015f): *Ovealmbornia reginae* (Teloschistaceae, Ascomycetes), a new xanthorhoid lichen from South Africa. – *Herzogia* **28**: 465–472. <https://doi.org/10.13158/heia.28.2.2015.465>
- Kondratyuk, S. Y., Lőkös, L., Halda, J. P., Haji Moniri, M., Farkas, E., Park, J. S., Lee, B. G., Oh, S.-O. and Hur, J.-S. (2016a): New and noteworthy lichen-forming and lichenicolous fungi 4. – *Acta Bot. Hung.* **58**(1–2): 75–136. <https://doi.org/10.1556/034.58.2016.1-2.4>
- Kondratyuk, S. Y., Lőkös, L., Halda, J. P., Upreti, D. K., Mishra, G. K., Haji Moniri, M., Farkas, E., Park, J. S., Lee, B. G., Liu, D., Woo, J. J., Jayalal, R. G. U., Oh, S.-O. and Hur, J.-S. (2016b): New and noteworthy lichen-forming and lichenicolous fungi 5. – *Acta Bot. Hung.* **58**(3–4): 319–396. <https://doi.org/10.1556/ABot.58.2016.3-4.7>
- Kondratyuk, S. Y., Lőkös, L., Kim, J. A., Kondratyuk, A. S., Jeong, M.-H., Jang, S. H., Oh, S.-O., Wang, X. Y. and Hur, J.-S. (2016c): *Fauriea*, a new genus of the lecanoroid caloplacoid lichens (Teloschistaceae, lichen-forming Ascomycetes). – *Acta Bot. Hung.* **58**(3–4): 303–318. <https://doi.org/10.1556/ABot.58.2016.3-4.6>
- Kondratyuk, S. Y., Lőkös, L., Kärnefelt, I., Thell, A., Elix, J. A., Oh, S.-O. and Hur, J.-S. (2016d): Three new Orientophila species (Teloschistaceae, Ascomycota) from Eastern Asia. – *Graphis Scripta* **28**(1–2): 50–58.
- Kondratyuk, S. Y., Lőkös, L., Halda, J. P., Roux, C., Upreti, D. K., Schumm, F., Mishra, G. K., Nayaka, S., Farkas, E., Park, J. S., Lee, B. G., Liu, D., Woo, J.-J. and Hur, J.-S. (2017): New and noteworthy lichen-forming and lichenicolous fungi 6. – *Acta Bot. Hung.* **59**(1–2): 137–260. <https://doi.org/10.1556/034.59.2017.1-2.7>
- Llimona, X. and Werner, R. G. (1975): Quelques lichens nouveau ou interessants de la Sierra de Gata (Almeria, SE de l'Espagne). – *Acta Phytotax. Barcinonensis* **16**: 1–32.
- Lutzoni, F., Pagel, M. and Reeb, V. (2001): Major fungal lineages are derived from lichen symbiotic ancestors. – *Nature* **411**(6840): 937–940. <https://doi.org/10.1038/35082053>
- Malme, G. O. A. (1926): Lichenes blasteniospori Herbarii Regnelli. – *Ark. Bot.* **20A**(9): 1–51.
- Martin, M. P. and Winka, K. (2000): Alternative methods of extracting and amplifying DNA from lichens. – *Lichenologist* **32**: 189–196. <https://doi.org/10.1006/lich.1999.0254>
- Søchting, U. and Figueras, G. (2007): *Caloplaca lenae* sp. nov. and other *Caloplaca* species with caloploicin and vicanicin. – *Lichenologist* **39**(1): 7–14. <https://doi.org/10.1017/s0024282907006299>
- Søchting, U., Søgaard, M. Z., Elix, J. A., Arup, U., Elvebakk, A. and Sancho, L. G. (2014a): *Catenaria* (Teloschistaceae, Ascomycota), a new Southern Hemisphere genus with 7-chlorocatenarin. – *Lichenologist* **46**: 175–187. <https://doi.org/10.1017/s002428291300087x>
- Søchting, U., Garrido-Benavent, I., Seppelt, R., Castello, M., Pérez-Ortega, S., De Los Ríos Murillo, A., Sancho, L. G., Frödén, P. and Arup, U. (2014b): *Charcotiana* and *Amundsenia*, two new genera in Teloschistaceae (lichenized Ascomycota, subfamily Xanthorioideae) hosting two new species from continental Antarctica, and *Austroplaca frigida*, a new name for a continental Antarctic species. – *Lichenologist* **46**: 763–782. <https://doi.org/10.1017/s0024282914000395>
- Steiner, J. (1911): Adnotationes lichenographicae. – *Österr. Bot. Zeitschr.* **61**: 177–183, 223–225.
- Swofford, D. L. (2003). PAUP\*, *Phylogenetic analysis using parsimony (\*and other methods)*. – Sunderland, Sinauer Associates, Massachusetts.
- Tamura, K., Peterson, D., Peterson, N., Stecher, G., Nei, M. and Kumar, S. (2011): MEGA5: molecular evolutionary genetics analysis using maximum likelihood, evolutionary distance, and maximum parsimony methods. – *Mol. Biol. Evol.* **28**: 2731–2739. <https://doi.org/10.1093/molbev/msr121>

- van den Boom, P. P. G. and Etayo, J. (2006): New records of lichens and lichenicolous fungi from Fuerteventura (Canary Islands), with descriptions of some new species. – *Cryptogamie, Mycol.* **27**(4): 341–374.
- Vilgalys, R. and Hester, M. (1990): Rapid genetic identification and mapping of enzymatically amplified ribosomal DNA from several *Cryptococcus* species. – *J. Bacteriol.* **172**(8): 4238–4246.
- Vondrák, J., Khodosovtsev, A., Šoun, J. and Vondráková, O. (2011): Two new European species from the heterogeneous *Caloplaca holocarpa* group (Teloschistaceae). – *Lichenologist* **44**(1): 73–89. <https://doi.org/10.1017/S0024282911000636>
- Vondrák, J., Šoun, J., Sogaard, M., Søchting, U. and Arup, U. (2010): *Caloplaca phlogina*, a lichen with two facies: an example of infraspecific variability resulting in the description of a redundant species. – *Lichenologist* **42**: 685–692. <https://doi.org/10.1017/S0024282910000435>
- Vondrák, J., Šoun, J., Vondráková, O., Fryday, A. M., Khodosovtsev, A. and Davydov, E. (2012): Absence of anthraquinone pigments is paraphyletic and a phylogenetically unreliable character in the Teloschistaceae. – *Lichenologist* **44**: 401–418. <https://doi.org/10.1017/s0024282911000843>
- Wetmore, C. M. (2003): The *Caloplaca squamosa* group in North and Central America. – *Bryologist* **106**(1): 147–156. [https://doi.org/10.1639/0007-2745\(2003\)106\[0147:tcsgin\]2.0.co;2](https://doi.org/10.1639/0007-2745(2003)106[0147:tcsgin]2.0.co;2)
- Wetmore, C. (2007): *Caloplaca*. – In: Nash III, T. H., Gries, C. and Bungartz, F. (eds): Lichen flora of the Greater Sonoran Desert Region. 3. Arizona State University, Lichens Unlimited, Tempe, pp. 179–220.
- White, T. J., Bruns, T., Lee, S. and Taylor, J. (1990): Amplification and direct sequencing of fungal ribosomal RNA genes for phylogenetics. – *PCR Protocols* **38**: 315–322. <https://doi.org/10.1016/b978-0-12-372180-8.50042-1>
- Zahlbrückner, A. (1917): Botanische Ergebnisse der schwedischen Expedition nach Patagonien und dem Feuerlande 1907–1909. VI. Die Flechten. – *Kongl. Svenska vetensk.-Akad. Handl.* **57**: 1–62.

## APPENDIX

Specimens included in the phylogenetic analysis with GenBank accession numbers. Newly submitted sequences are given in bold. Abbreviation: \* = this paper

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Amundsenia approximata</i>	Søchting <i>et al.</i> (2014)	KJ789964	KJ789972	KJ789974
<i>Amundsenia approximata</i>	Søchting <i>et al.</i> (2014)	KJ789963		
<i>Athallia holocarpa</i>	Arup (2009)	FJ346539		
<i>Athallia holocarpa</i>	Arup <i>et al.</i> (2013a)		KC179148	KC179478
<i>Athallia scopularis</i>	Arup <i>et al.</i> (2013a)	KC179339	KC179150	KC179480
<i>Austroplaca ambitiosa</i>	Arup <i>et al.</i> (2013a)	KC179081	KC179151	KC179481
<i>Blastenia ferruginea</i>	Arup <i>et al.</i> (2013a)	KC179416	KC179163	KC179493
<i>Brigantiae ferruginea</i>	Kondratyuk <i>et al.</i> (2013b); SK-779	KF264622		KF264684
<i>Brigantiae ferruginea</i>	Kondratyuk <i>et al.</i> (2013b); SK-780	KF264623		KF264685
<i>Brigantiae ferruginea</i>	*South Korea, 06.07.2012, Kondratyuk, S. Y. (121967), KoLRI 16922	KY614393		
<i>Brigantiae ferruginea</i>	*South Korea, 06.07.2012,			
	Kondratyuk, S. Y. (121971), KoLRI 16926	KY614394		
<i>Brigantiae ferruginea</i>	*South Korea, 06.07.2012, Kondratyuk, S. Y. (121981), KoLRI 16936	KY614395		
<i>Browniella kobeana</i>	Kondratyuk <i>et al.</i> (2015d); 120032, KoLRI	KT456212	KT456227	KT456242

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Browniella kobeana</i>	Kondratyuk et al. (2015d); 130231, KoLRI	KT456213	KT456228	KT456243
<i>Browniella kobeana</i>	Kondratyuk et al. (2015d); 130318, KoLRI	KT456214	KT456229	KT456244
<i>Bryoplaca sinapisperma</i>	Arup et al. (2013a)	KC179421		KC179495
<i>Calogaya biatorina</i>	Gaya et al. (2008)	EU639634		
<i>Calogaya aff. lobulata</i>	Kondratyuk et al. (2014c); SK-803	KJ133448	KJ133487	KJ133506
<i>Calogaya aff. lobulata</i>	Kondratyuk et al. (2014c); SK-804	KJ133449	KJ133488	KJ133507
<i>Calogaya aff. lobulata</i>	Kondratyuk et al. (2014c); SK-805	KJ133450	KJ133489	KJ133508
<i>Calogaya mogoltanica</i>	Kondratyuk et al. (2014c); SK-256	KJ133452	KJ133491	KJ133509
<i>Calogaya mogoltanica</i>	Kondratyuk et al. (2014c); SK-257	KJ133453	KJ133492	KJ133510
<i>Caloplaca cerina</i>	Fedorenko et al. (2009, 2012); FNM-185	EU681284		EU680863
<i>Caloplaca cerina</i>	Gaya et al. (2012)		JQ301549	
<i>Caloplaca pelodella</i>	Kondratyuk et al. (2013b); SK-714	KF264629		KF264689
<i>Caloplaca stillicidiorum</i>	Gaya et al. (2008)	EU639607		
<i>Catenarina desolata</i>	Söchting et al. (2014)	KF657317		KF657319
<i>Cerothallia luteoalba</i>	Arup et al. (2013a)	KC179099	KC179177	KC179511
<i>Cerothallia yarraensis</i>	Kondratyuk et al. (2014c); SK-890, isotype	KJ133454	KJ133493	KJ133511
<i>Charcotiana antarctica</i>	Söchting et al. (2014)	KJ789971	KJ789973	KJ789976
<i>Charcotiana antarctica</i>	Söchting et al. (2014)	KJ789968		
<i>Dijigiella kaernefeltiana</i>	*SK-969, Australia, 07.01. 2004, Kärnefelt, I. 20042002; LD 1238056, holotype		KY614396	KY614444 KY614475
<i>Dijigiella kaernefeltiana</i>	*SK-970, Australia, 07.01. 2004, Kärnefelt, I. 20042002; LD 1238056, holotype		KY614397	KY614445 KY614476
<i>Dijigiella subaggregata</i>	*SK-955, Australia: Victoria, 23.01.1999, Kärnefelt, I. 996401; LD 1275974, holotype	KY614398	KY614446	KY614477
<i>Dufourea flammea</i>	Arup et al. (2013)		KC179357	KC179183
<i>Dufourea flammea</i>	Fedorenko et al. (2009); FNM-083	EU681316		EU680898
<i>Dufourea flammea</i>	Fedorenko et al. (2009); FNM-049	EU681315		EU680897
<i>Eilifdahlia dahlii</i>	Kondratyuk et al. (2014a); SK-956	KJ021221	KJ021252	KJ021277
<i>Eilifdahlia dahlii</i>	Kondratyuk et al. (2014a); SK-959	KJ021318	KJ021253	KJ021279
<i>Eilifdahlia wirthii</i>	Kondratyuk et al. (2014a); SK-262	KJ021319	KJ021254	KJ021280
<i>Elenkiniana ehrenbergii</i>	Söchting and Figueras (2007)		DQ888715	
<i>Elenkiniana gloriae</i>	Kondratyuk et al. (2014a); SK-750	KJ021323		
<i>Elenkiniana gloriae</i>	Kondratyuk et al. (2014a); SK-611	KJ021321	KJ021256	KJ021282
<i>Elenkiniana gloriae</i>	Kondratyuk et al. (2014a); SK-613	KJ021322		KJ021283
<i>Elixjohnia bermaguiana</i>	Kondratyuk et al. (2013c, as <i>Sirenophila bermaguiana</i> ); SK-979			KF264706
<i>Elixjohnia bermaguiana</i>	Arup et al. (2013a, as <i>Sirenophila bermaguiana</i> ); isotype		KC179299	KC179245 KC179584
<i>Elixjohnia gallowayi</i>	Arup et al. (2013a, as <i>Sirenophila gallowayi</i> ); isotype		KC179301	KC179247 KC179586
<i>Elixjohnia jackelixii</i>	Kondratyuk et al. (2013c, as <i>Sirenophila jackelixii</i> ); SK-910		KF264655	KF264683 KF264707
<i>Elixjohnia jackelixii</i>	Kondratyuk et al. (2013b, as <i>Sirenophila jackelixii</i> ); SK-911			KF264708
<i>Elixjohnia jackelixii</i>	Arup et al. (2013a, as <i>Sirenophila jackelixii</i> )	KC179303	KC179248	KC179587
<i>Fauriea chujaensis</i>	Kondratyuk et al. (2016c); SK-D07, isotype	KX793097	KX793100	KX793103
<i>Fauriea orientochinensis</i>	Kondratyuk et al. (2016c); SK-709	KX793095	KX793098	KX793101
<i>Fauriea orientochinensis</i>	Kondratyuk et al. (2016c); SK-710	KX793096	KX793099	KX793102
<i>Filsoniana austroaliensis</i>	Kondratyuk et al. (2013c); SK-751	KF264631	KF264665	KF264691
<i>Flavoplaca citrina</i>	Arup (2006)	DG173224		
<i>Flavoplaca citrina</i>	Arup et al. (2013a)		KC179186	KC179521
<i>Flavoplaca aff. lutea</i>	Kondratyuk et al. (2014c); SK-840	KJ133456	KJ133494	KJ133514
<i>Flavoplaca aff. lutea</i>	Kondratyuk et al. (2014c); SK-841	KJ133457	KJ133495	KJ133515
<i>Flavoplaca aff. lutea</i>	Kondratyuk et al. (2014c); SK-843	KJ133458	KJ133496	KJ133516
<i>Follmannia orthoclada</i>	Arup et al. (2013a)	KC179291		

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Fominiella skii</i>	Vondrák <i>et al.</i> (2012), holotype	HM582191		
<i>Fominiella skii</i>	Vondrák <i>et al.</i> (2012)	HM582188		
<i>Fominiella skii</i>	Vondrák <i>et al.</i> (2012)	HM582194		
<i>Fominiella skii</i>	Vondrák <i>et al.</i> (2012)	HM582190		
<i>Fominiella tenerifensis</i>	*SK-D19, Spain: Tenerife, 15.01.2009, Kondratyuk, S. Y. 200916 KW-L, holotype	KY614447	KY614478	
<i>Franwilsia bastowii</i>	Kondratyuk <i>et al.</i> (2014a); SK-810	KJ021324	KJ021257	KJ021284
<i>Franwilsia kilcundaensis</i>	Kondratyuk <i>et al.</i> (2014a); SK-920	KJ021326	KJ021259	KJ021286
<i>Franwilsia renatae</i>	Kondratyuk <i>et al.</i> (2014a); SK-235	KJ021329		KJ021289
<i>Fulgensia cranfieldii</i>	Kondratyuk <i>et al.</i> (2014a); SK-983	KJ021333	KJ021262	KJ021292
<i>Fulgensia fulgens</i>	Kondratyuk <i>et al.</i> (2014a); SK-735	KJ021335		KJ021295
<i>Fulgensia poeltii</i>	Gaya <i>et al.</i> (2008)	EU639586		
<i>Fulgogasparrea decipiooides</i>	Kondratyuk <i>et al.</i> (2013b); SK-689	KF264644	KF264695	
<i>Fulgogasparrea decipiooides</i>	Kondratyuk <i>et al.</i> (2013b); SK-691	KF264643	KF264694	
<i>Fulgogasparrea decipiooides</i>	Arup <i>et al.</i> (2013a)	KC179333	KC179269	KC179608
<i>Gallowayella borealis</i>	Arup <i>et al.</i> (2013a)		KC179278	KC179617
<i>Gallowayella fulva</i>	Fedorenko <i>et al.</i> (2009); FNM-035	EU681352		
<i>Gallowayella hasseana</i>	Arup <i>et al.</i> (2013a)		KC179280	KC179619
<i>Gallowayella poeltii</i>	Fedorenko <i>et al.</i> (2009); FNM-111	EU681345		
<i>Gallowayella poeltii</i>	Arup <i>et al.</i> (2013a)		KC179282	KC179622
<i>Gallowayella poeltii</i>	Gaya <i>et al.</i> (2012)		JQ301583	
<i>Gallowayella weberi</i>	Fedorenko <i>et al.</i> (2009); FNM-031	EU681345		
<i>Gintarasiella aggregata</i>	*SK-A84, S Australia, 17.09.2012, Kantvilas, G. 476/12 HO 567239 – holotype	KY614390	KY614448	KY614479
<i>Gintarasiella aggregata</i>	*SK-A85, Australia: South Australia, 17.09.2012, Kantvilas, G. 476/12 KW-L ex HO 567239 – isotype			
<i>Gintarasiella aggregata</i>	*SK-A86, Australia: South Australia, 17.09.2012 Kantvilas, G. 476/12 KW-L ex HO 567239 – isotype	KY614391	KY614449	KY614480
<i>Golubkovia trachyphylla</i>	Kondratyuk <i>et al.</i> (2014c); SK-491	KY614392	KY614450	KY614481
<i>Golubkovia trachyphylla</i>	Kondratyuk <i>et al.</i> (2014c); SK-492	KJ133460		
<i>Golubkovia trachyphylla</i>	Kondratyuk <i>et al.</i> (2014c); SK-486	KJ133525		
<i>Golubkovia trachyphylla</i>	Arup <i>et al.</i> (2013a)	KJ133459	KJ133524	
<i>Gondwania cribrosa</i>	Arup <i>et al.</i> (2013a)	KC179283	KC179283	KC179623
<i>Gondwania regalis</i>	Arup <i>et al.</i> (2013a)	KC179102	KC179192	KC179526
<i>Gondwania sejongensis</i>	Arup <i>et al.</i> (2013a)	KC179103	KC179193	KC179527
<i>Gondwania sejongensis</i>	Kondratyuk <i>et al.</i> (2014c); SK-748, holotype	KJ133465	KJ133531	
<i>Gondwania sejongensis</i>	Kondratyuk <i>et al.</i> (2014c); SK-758	KJ133467		KJ133532
<i>Gyalolechia aurea</i>	Arup <i>et al.</i> (2013a)	KC179434	KC179196	KC179530
<i>Gyalolechia canariensis</i>	Gaya <i>et al.</i> (2008)	EU639587		
<i>Gyalolechia canariensis</i>	Kondratyuk <i>et al.</i> (2014a); SK-583	KJ021332		
<i>Haloplaca sorediella</i>	Arup <i>et al.</i> (2013a)	KC179293		
<i>Haloplaca suaedae</i>	Vondrák <i>et al.</i> (unpubl.)	HM582197		
<i>Hanstrassia jaeseounhurii</i>	*China, 17.06.2016, Park, Ch.-N., Hur, J.-S. (CH-160016) KoLRI 41972, holotype	KY614399		
<i>Hanstrassia jaeseounhurii</i>	*China, 17.06.2016, Park, Ch.-N., Hur, J.-S. (CH-160015) KoLRI 41971, isotype	KY614400		
<i>Hanstrassia lenae</i>	Söchting and Figueras (2007)	DQ888717		
<i>Hanstrassia lenae</i>	Arup <i>et al.</i> (2013a)	KC179442		
<i>Hanstrassia aff. lenae</i>	*SK-655, Russia, 16.09.2011, Yakovchenko, L. S., T-461 KoLRI	KY614401		KY614482
<i>Hanstrassia aff. lenae</i>	*SK-654, Russia, 16.09.2011, Yakovchenko, L. S., T-458 KoLRI	KY614402		KY614483

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Harusavskia elenkinianoides</i>	*SK-996, Chile, 12.12.2012, Oh, S.-O., Hur, J.-S., CL-120324 KoLRI 14493, holotype	KY614403	KY614451	KY614484
<i>Harusavskia elenkinianoides</i>	*SK-997, Chile, 12.12.2012, Oh, S.-O., Hur, J.-S., CL-120324 KoLRI 14493, holotype	KY614404	KY614452	KY614485
<i>Harusavskia elenkinianoides</i>	*SK-269, Chile, 12.12.2012, Oh, S.-O., Hur, J.-S., CL-120324 KoLRI 14493 holotype	KY614405	KY614453	KY614486
<i>Honeggeria rosmarieae</i>	Eichenberger <i>et al.</i> (unpubl., as <i>Xanthomendoza weberi</i> )	AM697873		
<i>Honeggeria rosmarieae</i>	Arup <i>et al.</i> (2013a, as <i>Xanthomendoza weberi</i> )	KC179145	KC179285	KC179625
<i>Huneckia pollinii</i>	Kondratyuk <i>et al.</i> (2014a); SK-3206	KJ021336	KJ021265	KJ021296
<i>Huneckia pollinii</i>	Kondratyuk <i>et al.</i> (2014a); SK-870	KJ021337	KJ021266	KJ021297
<i>Huneckia rheinigera</i>	Kondratyuk <i>et al.</i> (2014a); SK-3204	KJ021222		
<i>Huriella loekoesiana</i>	Kondratyuk <i>et al.</i> (2014c, as <i>Squamulea subsolata</i> ); SK-694	KJ133481		
<i>Huriella loekoesiana</i>	*South Korea, 24.04.2012, Jayalal, U. <i>et al.</i> (120433), KoLRI 15423	KY614406		KY614487
<i>Huriella loekoesiana</i>	*South Korea, 28.06.2013, Oh, S.-O. <i>et al.</i> (130672), KoLRI 19017	KY614407		KY614488
<i>Huriella loekoesiana</i>	*South Korea, 10.07.2016, Kondratyuk, S. Y., Lökös, L. (161904), KoLRI 40141	KY614408		
<i>Huriella loekoesiana</i>	*South Korea, 11.07.2016, Kondratyuk, S.Y., Lökös, L. (161998), KoLRI 40236	KY614409		
<i>Huriella loekoesiana</i>	*South Korea, 11.07.2016, Kondratyuk, S. Y., Lökös, L. (162000), KoLRI 40238	KY614410		
<i>Igneoplaca ignea</i>	Arup <i>et al.</i> (2013a)	KC179382	KC179219	KC179555
<i>Igneoplaca ignea</i>	Arup and Grube (1999)	AF353950		
<i>Ikaeria aurantiellina</i>	*SK-538, Spain, Tenerife, 15.01.2009, Kondratyuk, S. Y. 20916, KW-L	KY614411		KY614490
<i>Ikaeria aurantiellina</i>	*SK-552, Spain, Tenerife, 15.01.2009, Kondratyuk, S. Y. 20916, KW-L	KY614412		KY614491
<i>Ikaeria aurantiellina</i>	*SK-D29, Spain, Tenerife, 15.01.2009, Kondratyuk S. Y. 20916, KW-L	KY614413		KY614492
<i>Ikaeria aurantiellina</i>	*SK-D23, Spain, Tenerife, 15.01.2009, Kondratyuk S. Y. 20916, KW-L	KY614414		KY614493
<i>Ioplaca pindarensis</i>	Gaya <i>et al.</i> (2012)	JQ301672		
<i>Jackelixia angustata</i>	Fedorenko <i>et al.</i> (2009); FNM-064			EU680870
<i>Jackelixia angustata</i>	Arup <i>et al.</i> (2013)	KC179351		KC179180
<i>Jackelixia dissectula</i>	Arup <i>et al.</i> (2013)	KC179355	KC179182	KC179517
<i>Jackelixia elixii</i>	Fedorenko <i>et al.</i> (2009); FNM-061	EU681308		EU680877
<i>Jasonhuria bogilana</i>	Kondratyuk <i>et al.</i> (2015c); KoLRI 120454	KT220196	KT220205	KT220214
<i>Jasonhuria bogilana</i>	Kondratyuk <i>et al.</i> (2015c); KoLRI 120469	KT220197	KT220206	KT220215
<i>Jasonhuria bogilana</i>	Kondratyuk <i>et al.</i> (2015c); KoLRI 120641	KT220198	KT220207	KT220216
<i>Jasonhuria bogilana</i>	Kondratyuk <i>et al.</i> (2015c); KoLRI 120647	KT220199	KT220208	KT220217
<i>Jesmurraya novozelandica</i>	Arup <i>et al.</i> (2013a) (sub <i>Xanthomendoza novozelandica</i> )	KC179140		KC179621
<i>Jesmurraya novozelandica</i>	Fedorenko <i>et al.</i> (2009); FNM-114	EU681347		
<i>Jesmurraya novozelandica</i>	*SK-C82, New Zealand, Otago, 17.04.2010, D. J. Galloway 6000, KW-L			KY614454 KY614489
<i>Josefpoeltia sorediosa</i>	Kondratyuk <i>et al.</i> (2013b); SK-991	KF264645	KF264673	KF264696
<i>Kaernefia kaernefeltii</i>	Kondratyuk <i>et al.</i> (2013b); SK-921	KF264652	KF264680	KF264703
<i>Klauderuiella aurantia</i>	Arup <i>et al.</i> (2013a)	KC179470	KC179261	KC179600
<i>Klauderuiella aurantia</i>	Gaya <i>et al.</i> (2008)	EU639602		
<i>Klauderuiella aurantia</i>	Gaya <i>et al.</i> (2015)			KT291479

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Klauderuiella flavescentis</i>	*SK-561, Spain, Tenerife, 16.01.2009, Kondratyuk, S. Y. 20925, KW-L	KY614416		KY614495
<i>Klauderuiella thallincola</i>	*SK-527, 18.08.1997, Kärnefelt, I. 970901 LD 1076480	KY614415		KY614494
<i>Langeottia ottolangei</i>	Kondratyuk et al. (2014c); SK-239	KJ133468	KJ133497	KJ133533
<i>Langeottia ottolangei</i>	Kondratyuk et al. (2014c); SK-240	KJ133469	KJ133498	KJ133534
<i>Langeottia ottolangei</i>	Kondratyuk et al. (2014c); SK-264	KJ133470	KJ133499	KJ133535
<i>Langeottia ottolangei</i>	Kondratyuk et al. (2014c); SK-265	KJ133471	KJ133500	KJ133536
<i>Laundonia flavovirescens</i>	*SK-657, Russia, 16.09.2011, Yakov- chenko, L. S., T-461, KoLRI	KY614417		KY614496
<i>Laundonia flavovirescens</i>	Arup and Grube (1999)	AF353966		
<i>Laundonia flavovirescens</i>	Arup et al. (2013a)		KC179198	KC179532
<i>Laundonia persimilis</i>	Arup et al. (2013a)	KC179444		
<i>Laundonia persimilis</i>	Vondrák et al. (unpubl.)	KT804978		
<i>Laundonia persimilis</i>	Vondrák et al. (unpubl.)	KT804979		
<i>Lazarenkoella zoroasteriorum</i>	Kondratyuk et al. (2015d); SK-A45	KT456215	KT456230	KT456245
<i>Lazarenkoella zoroasteriorum</i>	Kondratyuk et al. (2015d); SK-A51	KT456216	KT456231	KT456246
<i>Lazarenkoella zoroasteriorum</i>	Kondratyuk et al. (2015d); SK A55	KT456217	KT456232	KT456247
<i>Lazarenkoiopsis ussuriensis</i>	*SK A36, Russia, 03.08.2013, Kondratyuk, S. Y., 213R3 130130 KoLRI			KY614497
<i>Lazarenkoiopsis ussuriensis</i>	*SK A37, Russia, 03.08.2013, Kondratyuk, S. Y., 213R3 130130 KoLRI	KY614418	KY614455	KY614498
<i>Lazarenkoiopsis ussuriensis</i>	*SK D22, Russia, 03.08.2013, Kondratyuk, S. Y., 213R3 130130 KoLRI	KY614419	KY614456	KY614499
<i>Leproplaca oblitterans</i>	Arup et al. (2013a)	KC179449		KC179207
<i>Leproplaca xantholyta</i>	Arup et al. (2013a)	KC179451	KC179208	KC179542
<i>Leproplaca xantholyta</i>	Gaya et al. (2012)	JQ301670		JQ301565
<i>Loekoesia austrocoreana</i>	Kondratyuk et al. (2015c); KoLRI 120511	KT220200	KT220209	KT220218
<i>Loekoesia austrocoreana</i>	Kondratyuk et al. (2015c); KoLRI 120523	KT220201	KT220210	KT220219
<i>Loekoesia austrocoreana</i>	Kondratyuk et al. (2015c); SK-261	KT220202	KT220211	KT220220
<i>Marchantiana kalbiorum</i>	Kondratyuk et al. (2014a); SK-939	KJ021225	KJ023183	KJ021300
<i>Marchantiana occidentalis</i>	Kondratyuk et al. (2014a); SK-981	KJ021227	KJ021268	KJ021303
<i>Marchantiana occidentalis</i>	Kondratyuk et al. (2014a); SK-982	KJ021228	KJ021269	KJ021304
<i>Martinjahnzia resendei</i>	Martin and Winka (2000)	AF101285		
<i>Martinjahnzia resendei</i>	Arup et al. (2013a)		KC179290	KC179630
<i>Massjukiella polycarpa</i>	Arup et al. (2013a)	KC179389	KC179222	
<i>Massjukiella polycarpa</i>	Fedorenko et al. (2012); FNM-172			JN984146
<i>Massjukiella polycarpa</i>	Fedorenko et al. (2009); FNM-173	EU681333		EU680919
<i>'Caloplaca' allochroa</i>	Joshi et al. (2011)	HQ415800		
<i>Mikhtomia gordejevii</i>	Kondratyuk et al. (2014a); SK-80515 *South Korea, 06.07.2012, Kondratyuk, S. Y. (121959), KoLRI 16914	KJ021231		KJ021307
<i>Mikhtomia gordejevii</i>	*South Korea, 01.06.2012, collector unmentioned (120924), KoLRI 15954	KY614420	KY614457	
<i>Mikhtomia gordejevii</i>	Kondratyuk et al. (2014a); SK-80646	KY614421	KY614458	
<i>Mikhtomia multicolor</i>	Kondratyuk et al. (2014a); SK-A14 *South Korea, 28.06.2013, Oh, S.-O. et al. (130601), KoLRI 18946	KJ021232		KJ021308
<i>Mikhtomia multicolor</i>	*South Korea, 29.08.2004, Hur, J.-S. (40900), KoLRI 001689	KY614422	KY614459	
<i>Mikhtomia multicolor</i>	*SK A19, South Korea, 11.05.2012, Oh, S.-O. (120832), KoLRI 15829	KY614423	KY614460	
<i>Mikhtomia multicolor</i>	*South Korea, 28.06.2013, Oh, S.-O. et al. (130604), KoLRI 18949	KY614424		
			KY614461	

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Mikhtomia subflavorubencens</i>	Kondratyuk <i>et al.</i> (2014a as <i>M. oxnerii</i> ); SK-90117	KJ021233		KJ021311
<i>Mikhtomia subflavorubencens</i>	Kondratyuk <i>et al.</i> (2014a as <i>M. oxnerii</i> ); SK-90755	KJ021234		KJ021312
<i>Neobrowniella brownlieae</i>	Kondratyuk <i>et al.</i> (2013b); SK-831	KF264626	KF264661	KF264687
<i>Neobrowniella brownlieae</i>	Kondratyuk <i>et al.</i> (2013b); SK-838	KF264627	KF264662	KF264688
<i>Neobrowniella montisfracti</i>	Kondratyuk <i>et al.</i> (2013b); SK-230	KF264624	KF264659	
<i>Nevilleiella lateritia</i>	*SK-878, Australia, 10.01.1999, Kärnefelt, I. 990904, LD 1236736 *SK-261, Australia, 10.01.1999, Kärnefelt, I. 990904, LD 1236736	KY614426	KY614463	KY614501
<i>Nevilleiella lateritia</i>	*SK-D18, Western Australia, 08.01.2004, Kondratyuk, S. Y. 20444, KW-L – isotype Eichenberger <i>et al.</i> (unpubl.)	KY614427	KY614464	KY614502
<i>Nevilleiella marchantii</i>	Gaya <i>et al.</i> (2012)	KY614425	KY614462	KY614500
<i>Niorma chrysophthalma</i>	Kondratyuk <i>et al.</i> (2015c); SK-C65	AM292836		JQ301576
<i>Olegblumia demissa</i>	Arup <i>et al.</i> (2013a)	KT220203	KT220212	KT220221
<i>Opeltia arizonica</i>	*SK-D10, China, 01.08.2013, Oh, S.-O., Hur, J.-S., CH 130112 KoLRI	KC179433	KC179195	KC179529
<i>Opeltia juniperina</i>	Vondrák and Halici (unpubl.)	KY614429		KY614504
<i>Opeltia juniperina</i>	Joshi <i>et al.</i> (2011)	JN813383		
<i>Opeltia neobaltistanica</i>	*SK-D09, China, 01.08.2013, Oh, S.-O., Hur, J.-S., CH 130098 KoLRI	HQ644199		
<i>Orientophila loekoesii</i>	Kondratyuk <i>et al.</i> (2014c); SK-721	KY614428		KY614503
<i>Orientophila loekoesii</i>	Kondratyuk <i>et al.</i> (2014c); SK-692			KJ133540
<i>Orientophila loekoesii</i>	Kondratyuk <i>et al.</i> (2014c); SK-691			KJ133539
<i>Orientophila loekoesii</i>	Kondratyuk <i>et al.</i> (2014c); SK-690			KJ133538
<i>Orientophila subscopularis</i>	Arup <i>et al.</i> (2013a)			KJ133537
<i>Orientophila subscopularis</i>	Kondratyuk <i>et al.</i> (2014c); SK-727	KC179375		KC179546
<i>Orientophila subscopularis</i>	Kondratyuk <i>et al.</i> (2014c); SK-692	KJ133476		
<i>Orientophila subscopularis</i>	Kondratyuk <i>et al.</i> (2014c); SK-717	KJ133475		
<i>Ovealmbornia reginae</i>	Arup <i>et al.</i> (2013a, as <i>Ovealmbornia bonae-spei</i> ) (see Kondratyuk <i>et al.</i> 2015f)	KJ133474		
<i>Ovealmbornia volkmarwirthii</i>	Arup <i>et al.</i> (2013a, as <i>Ovealmbornia bonae-spei</i> ) (see Kondratyuk <i>et al.</i> 2015f)		KC179181	KC179516
<i>Ovealmbornia volkmarwirthii</i>	Kondratyuk <i>et al.</i> (2014c; 2015f); SK-241	KC179353		
<i>Ovealmbornia volkmarwirthii</i>	Kondratyuk <i>et al.</i> (2014c; 2015f); SK-242	KJ133477	KJ133501	KJ133541
<i>Ovealmbornia sp.</i>	Fedorenko <i>et al.</i> (2009, as <i>Ovealmbornia bonae-spei</i> ); FNM-099 (see Kondratyuk <i>et al.</i> 2015f)	KJ133478	KJ133502	KJ133542
<i>Oxneria alfredii</i>	Fedorenko <i>et al.</i> (2009); FNM-152	EU681319		EU680902
<i>Oxneria huculica</i>	Fedorenko <i>et al.</i> (2009, as <i>Oxneria fallax</i> ); FNM-199	EU681345		EU680933
<i>Oxneria huculica</i>	Gaya <i>et al.</i> (2012, as <i>Oxneria fallax</i> )			JQ301580
<i>Oxneria huculica</i>	Arup <i>et al.</i> (2013, as <i>Xanthomendoza fallax</i> )			KC179279
<i>Oxneria huculica</i>	Fedorenko <i>et al.</i> (2009, as <i>Oxneria fallax</i> ); FNM-022	EU681346		
<i>Oxneriopsis oxneri</i>	Arup <i>et al.</i> (2013, as <i>Caloplaca oxneri</i> )	KC179443		
<i>Oxneriopsis oxneri</i>	*SK-695, Russia, 17.09.2011			
<i>Oxneriopsis aff. oxneri</i>	Kondratyuk, S. Y. 21138, KW-L	KY614430		KY614505
<i>Oxneriopsis yeosuensis</i>	*SK-771, Russia, 17.09.2011	KY614431		KY614506
<i>Oxneriopsis yeosuensis</i>	Kondratyuk, S. Y. 21138, KW-L			
	*South Korea, 24.03.2007, Hur, J.-S. (70080), KoLRI 007067	KY614432	KY614465	KY614507

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Oxneriopsis yeosuensis</i>	*South Korea, 26.04.2012, Jayalal, U. et al. (120357), KoLRI 15346	KY614433	KY614466	KY614508
<i>Oxneriopsis yeosuensis</i>	*South Korea, 26.04.2012, Jayalal, U. et al. (120336), KoLRI 15325	KY614434	KY614467	KY614509
<i>Pachypeltis castellana</i>	Arup et al. (2013a)	KC179105		KC179547
<i>Parvoplaca tiroliensis</i>	Arup et al. (2013a)	KC179116	KC179216	KC179552
<i>Polycauliona coralloides</i>	Arup et al. (2013a)	KC179380	KC179218	KC179554
<i>Polycauliona coralloides</i>	Gaya et al. (2015)	KT291451	KT291539	KT291485
<i>Pyrenodesmia alociza</i>	Kondratyuk et al. (2014a); SK-747	KJ021239		KJ021313
<i>Pyrenodesmia chalybaea</i>	Gaya et al. (2012)	JQ301550		
<i>Pyrenodesmia teicholyta</i>	Vondrák et al. (2012)	JN641791		
<i>Pyrenodesmia teicholyta</i>	Arup et al. (2013a)		KC179176	
<i>Pyrenodesmia variabilis</i>	Gaya et al. (2003)	AY233224		
<i>Raeseneniana maulensis</i>	Kondratyuk et al. (2015d); SK-993, holotype	KT456218	KT456233	KT456248
<i>Raeseneniana maulensis</i>	Kondratyuk et al. (2014a); SK-994	KJ023182	KJ023184	
<i>Rufoplaca scotoplaca</i>	Arup et al. (2013a)	KC179457	KC179235	KC179573
<i>Rusavskia dasanensis</i>	Kondratyuk et al. (2015e); SK-702	KU056846		KU043372
<i>Rusavskia elegans</i>	Fedorenko et al. (2009); FNM-019	EU681336		EU680921
<i>Rusavskia elegans</i>	Arup et al. (2013a)		KC179238	KC179576
<i>Rusavskia sorediata</i>	Fedorenko et al. (2009); FNM-046, Finland	EU681335		EU680920
<i>Rusavskia sorediata</i>	Arup et al. (2013a)		KC179239	
<i>Scutaria andina</i>	Arup et al. (2013a)	KC179298	KC179242	KC179581
<i>Scythioria phlogina</i>	Arup (2006) (sub <i>Caloplaca phlogina</i> )	DQ173234		
<i>Scythioria phlogina</i>	Arup et al. (2013a, as <i>Polycauliona phlogina</i> )		KC179221	KC179557
<i>Scythioria phlogina</i>	Vondrák et al. (2010, as <i>Caloplaca scythica</i> )	GU080302		
<i>Seirophora californica</i>	Arup et al. (2013a)	KC179643		
<i>Seirophora lacunosa</i>	Kondratyuk et al. (2015c); SK-B07	KT220204	KT220213	KT220222
<i>Seirophora stenophylla</i>	Arup et al. (2013a)	KC179647		
<i>Seirophora villosa</i>	*SK-D27, Italy, 01.04.2015, Ravera, S. (specimens 1), KW-L	KY614435	KY614468	KY614510
<i>Seirophora villosa</i>	*SK-D16, Morocco, 25.03.2014, Guttová, A., KW-L ex BRA	KY614436	KY614469	KY614511
<i>Seirophora villosa</i>	Martin and Winka (2000)	AF098407		
<i>Shackletonia hertelii</i>	Arup et al. (2013a)	KC179118		KC179579
<i>Shackletonia sauronii</i>	Arup et al. (2013a)	KC179120	KC179241	KC179580
<i>Sirenophila cliffwetmorei</i>	*SK-A93, Australia, 15.09.2012, Kantvilas, G., 446/12 KW-L ex HO	KY614438	KY614471	KY614513
' <i>Sirenophila</i> ' eos	Kondratyuk et al. (2013b); SK-912	KF264656		
' <i>Sirenophila</i> ' eos	Arup et al. (2013a)	KC179300	KC179246	KC179585
' <i>Sirenophila</i> ' eos	Gaya et al. (2015)	KT291455	KT291542	KT291489
<i>Sirenophila gintarasii</i>	Arup et al. (2013a)	KC179302		
<i>Sirenophila gintarasii</i>	*SK-D17, Australia, 19.01.2004, Kondratyuk, S. Y., KW-L – isotype	KY614437	KY614470	KY614512
' <i>Sirenophila</i> ' maccarthyi	Arup et al. (2013a)	KC179304	KC179249	KC179588
<i>Solitaria chrysopthalma</i>	*SK-D20, Sweden, 27.01.2012, Arup, U., L 05051 LD 1109837	KY614439		KY614514
<i>Squamulea squamosa</i>	Arup et al. (2013a)	KC179125	KC179252	KC179591
<i>Squamulea subsoluta</i>	Arup (2006)	DQ173238		
<i>Squamulea subsoluta</i>	Arup et al. (2013a)		KC179253	KC179592
<i>Squamulea subsoluta</i>	Kondratyuk et al. (2014c); SK-670	KJ133480		
<i>Stellarangia elegantissima</i>	Arup et al. (2013a)	KC179310	KC179254	KC179593
<i>Streimanniella kalbiorum</i>	Kondratyuk et al. (2014a); SK-939	KJ021225	KJ023183	KJ021300
<i>Streimanniella michelagoensis</i>	Kondratyuk et al. (2014a); SK-971	KJ021226	KJ023185	KJ021301

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Streimannia seppeltii</i>	Kondratyuk <i>et al.</i> (2014a); SK-855	KJ021229	KJ023186	KJ021305
<i>Tarasginia whinrayi</i>	Kondratyuk <i>et al.</i> (2015d); SK-A95	KT456220	KT456235	KT456250
<i>Tarasginia whinrayi</i>	Kondratyuk <i>et al.</i> (2015d); SK-B02	KT456221	KT456236	KT456251
<i>Tassiloc digitarea</i>	Kondratyuk <i>et al.</i> (2015a); SK-A34	KP096222		KP096224
<i>Tayloriella erythrostictica</i>	Kondratyuk <i>et al.</i> (2015d); SK-817	KT456222	KT456237	KT456252
<i>Tayloriella erythrostictica</i>	Kondratyuk <i>et al.</i> (2015d); SK-814	KT456223	KT456238	KT456253
<i>Tayloriella erythrostictica</i>	Kondratyuk <i>et al.</i> (2015d) SK-819	KT456224	KT456239	KT456254
<i>Teloschistes flavicans</i>	Fedorenko <i>et al.</i> (2009, 2012); FNM-139	EU681363		EU680955
<i>Teloschistes flavicans</i>	Arup <i>et al.</i> (2013a)	KC179317	KC179255	KC179594
<i>Teloschistes flavicans</i>	Fedorenko <i>et al.</i> (2009, 2012); FNM-218	EU681362		JN984150
<i>Teloschistes flavicans</i>	Gaya <i>et al.</i> (2012)			JQ301578
<i>Teloschistopsis bonae-spei</i>	Arup <i>et al.</i> (2013a)	KC179322	KC179257	KC179596
<i>Teloschistopsis chrysocarpoidea</i>	Arup <i>et al.</i> (2013a)	KC179323		
<i>Teloschistopsis eudoxa</i>	Arup <i>et al.</i> (2013a)	KC179324	KC179258	KC179597
<i>Thelliana pseudokiamae</i>	Kondratyuk <i>et al.</i> (2015d); SK-925	KT456225	KT456240	KT456255
<i>Thelliana pseudokiamae</i>	Kondratyuk <i>et al.</i> (2015d); SK-926	KT456226	KT456241	KT456256
<i>Teuvoahrtiana altoandina</i>	Arup <i>et al.</i> (2013a, as 'Caloplaca' altoandina)	KC179094	KC179170	KC179503
<i>Teuvoahrtiana fernandeziana</i>	*SK-995, Chile, 27.01.2012, Wang, X. Y., Oh, S.-O., Hur, J.-S., CL-120121			
	KoLRI 14287	KY614440		KY614515
<i>Teuvoahrtiana rugulosa</i>	*SK-A25, Chile, 30.01.2012, Wang, X. Y., Oh, S.-O., Hur, J.-S., CL-120331 KoLRI 14500	KY614441	KY614472	KY614516
<i>Teuvoahrtiana rugulosa</i>	*SK-A26, Chile, 30.01.2012, Wang, X. Y., Oh, S.-O., Hur, J.-S., CL-120331 KoLRI 14500	KY614442	KY614473	KY614517
<i>Teuvoahrtiana rugulosa</i>	*SK-A27, Chile, 30.01.2012, Wang, X. Y., Oh, S.-O., Hur, J.-S., CL-120331 KoLRI 14500	KY614443	KY614474	KY614518
<i>Tommashia ludificans</i>	Arup <i>et al.</i> (2013a, as <i>Polycauliona ludificans</i> ); holotype		KC179386	
<i>Tommashia luteominia</i>	Arup <i>et al.</i> (2013a, as <i>Polycauliona luteominia</i> )	KC179387	KC179220	KC179556
<i>Tommashia nashii</i>	Vondrák <i>et al.</i> (unpubl., as <i>Caloplaca nashii</i> )			HQ699624
<i>Tommashia rosei</i>	Arup <i>et al.</i> (2013a, as <i>Polycauliona rosei</i> )	KC179390	KC179223	KC179559
<i>Usnochroma carpinea</i>	Arup <i>et al.</i> (2013a)	KC179468	KC179259	KC179598
<i>Usnochroma carpinea</i>	Gaya <i>et al.</i> (2008)	EU639595		
<i>Usnochroma carpinea</i>	Gaya <i>et al.</i> (2012)			JQ301548
<i>Usnochroma scoriphila</i>	Gaya <i>et al.</i> (2012)	JQ301664	JQ301560	
<i>Variospora kudratovii</i>	Kondratyuk <i>et al.</i> (2014a); SK-487	KJ021244	KJ021276	KJ023192
<i>Variospora kudratovii</i>	Kondratyuk <i>et al.</i> (2014a); SK-485	KJ021243	KJ021275	KJ023191
<i>Variospora kudratovii</i>	Kondratyuk <i>et al.</i> (2014a); SK-484	KJ021242	KJ021274	KJ023190
<i>Variospora latzelii</i>	Vondrák <i>et al.</i> (unpubl.)	JN813418		
<i>Variospora macrocarpa</i>	Arup and Grube (1999)		AF353956	
<i>Variospora velana</i>	Arup <i>et al.</i> (2013a)	KC179476	KC179265	KC179605
<i>Verrucoplaca verruculifera</i>	Arup <i>et al.</i> (2013a)	KC179404	KC179223	
<i>Verrucoplaca verruculifera</i>	Gaya <i>et al.</i> (2008)	EU639624		
<i>Verrucoplaca verruculifera</i>	Gaya <i>et al.</i> (2012)			JQ301564
<i>Villophora isidioclada</i>	Arup <i>et al.</i> (2013a)	KC179325	KC179266	KC179606
<i>Wetmoreana texana</i>	Kondratyuk <i>et al.</i> (2013b); SK-537	KF264657		KF264710
<i>Wetmoreana texana</i>	Kondratyuk <i>et al.</i> (2013b); SK-536	KF264658		KF264711
<i>Wetmoreana texana</i>	Arup <i>et al.</i> (2013a)	KC179337	KC179273	KC179612
<i>Xanthptychia aurantiaca</i>	Arup <i>et al.</i> (2013a, as <i>Seirophora aurantiaca</i> )	KC179461		
<i>Xanthptychia blumii</i>	Kondratyuk <i>et al.</i> (2015d, as <i>Seirophora blumii</i> ); SK-A65	KT456219	KT456234	KT456249

Species name	References / voucher details	ITS	LSU	mtSSU
<i>Xanthptychia contortuplicata</i>	Arup et al. (2013a, as <i>Seiophora contortuplicata</i> )	KC179464		
<i>Xanthptychia contortuplicata</i>	Gaya et al. (2015, as <i>Seiophora contortuplicata</i> )	KT291471		KT291522
<i>Xanthptychia contortuplicata</i>	Kondratyuk et al. (2014a, as <i>Seiophora contortuplicata</i> ); SK-775			KJ021316
<i>Xanthptychia orientalis</i>	Fedorenko et al. (2009, as <i>Seiophora orientalis</i> ); FNM-153	EU681287		
<i>Xanthptychia orientalis</i>	Kondratyuk et al. (2014a, as <i>Seiophora orientalis</i> ); SK-756	KJ021241		
<i>Xanthptychia orientalis</i>	Kondratyuk et al. (2014a, as <i>Seiophora orientalis</i> ); SK-755	KJ021240		KJ023189
<i>Xanthocarpia jerramungupensis</i>	Kondratyuk et al. (2014c); SK-142	KJ133484	KJ133503	KJ133543
<i>Xanthocarpia jerramungupensis</i>	Kondratyuk et al. (2014c); SK-917	KJ133485	KJ133504	KJ133544
<i>Xanthocarpia cf. jerramungupensis</i>	Kondratyuk et al. (2014c); SK-247	KJ133486	KJ133505	KJ133545
<i>Xanthocarpia ochracea</i>	Kondratyuk et al. (2014c); SK-637	KJ133483		
<i>Xanthocarpia ochracea</i>	Arup et al. (2013a)	KC179132	KC179277	KC179617
<i>Xanthokarrooa karrooensis</i>	Fedorenko et al. (2009); FNM-066	EU681320		EU680903
<i>Xanthomendoza mendozae</i>	Fedorenko et al. (2009); FNM-215	EU681349		
<i>Xanthomendoza mendozae</i>	Fedorenko et al. (2009); FNM-065			EU680938
<i>Xanthomendoza mendozae</i>	Fedorenko et al. (2009); FNM-069			EU680939
<i>Xanthomendoza mendozae</i>	Arup et al. (2013a)	KC179138	KC179281	KC179620
<i>Xanthopeltis rupicola</i>	Arup et al. (2013a)	KC179146	KC179286	KC179626
<i>Xanthopeltis rupicola</i>	Gaya et al. (2015)		KT291570	KT291530
<i>Xanthoria parietina</i>	Fedorenko et al. (2009, 2012); FNM-177	EU681289		EU680868
<i>Xanthoria parietina</i>	Gaya et al. (2012)		JQ301589	
<i>Xanthoria parietina</i>	Fedorenko et al. (2009, 2012); FNM-177	EU681289		EU680868
<i>Xanthoria parietina</i>	Gaya et al. (2012)		JQ301589	
<i>Xanthoria parietina</i>	Fedorenko et al. (2009, 2012); FNM-017	EU681288	EU680867	
<i>Xanthoria parietina</i>	Lutzoni et al. (2001)			AF356687
<i>Yoshimuria cerussata</i>	Kondratyuk et al. (2014a); SK-768	KJ021248		
<i>Yoshimuria galbina</i>	Kondratyuk et al. (2014a); SK-704			KJ023197
<i>Yoshimuria spodoplaca</i>	Kondratyuk et al. (2014a); SK-725	KJ021249		KJ023194
<i>Zeroviella coreana</i>	Kondratyuk et al. (2015e); SK-D06	KU056848		KU043373
<i>Zeroviella esfahanensis</i>	Kondratyuk et al. (2015e); SK-501	KU056849		KU043374
<i>Zeroviella papillifera</i>	Fedorenko et al. (2009); FNM-018	EU681331		EU680927
<i>Zeroviella papillifera</i>	Fedorenko et al. (2009); FNM-204	EU681330		EU680926
<i>Zeroviella ussurica</i>	Kondratyuk et al. (2015e); SK-D08, isotype	KU056857		KU043379