

**Mycosphere 8(1): 172–217 (2017)** www.mycosphere.org

ISSN 2077 7019

# **Article**

Doi 10.5943/mycosphere/8/1/15 Copyright © Guizhou Academy of Agricultural Sciences

# Morphophylogenetic study of Sydowiellaceae reveals several new genera

# Senanayake IC <sup>1,2,3</sup>, Maharachchikumbura SSN <sup>4</sup>, Jeewon R<sup>5</sup>, Promputtha I <sup>6</sup>, Al-Sadi AM <sup>4</sup>, Camporesi E <sup>7,8,9</sup> and Hyde KD <sup>1,2,3</sup>

Senanayake IC, Maharachchikumbura SSN, Jeewon R, Promputtha I, Al-Sadi AM, Camporesi E, Hyde KD. 2017 – Morphophylogenetic study of Sydowiellaceae reveals several new genera. Mycosphere 8(1), 172–217, Doi 10.5943/mycosphere/8/1/15

#### **Abstract**

Sydowiellaceae is a poorly studied family of the order Diaporthales, comprising a collection of morphologically diversified taxa. Eleven genera have been previously listed under this family. In this study, we provide a DNA sequence-based phylogeny for genera of Sydowiellaceae based on analyses of a combined LSU, ITS, RPB2 and TEF sequence dataset to establish the boundaries within the family. We introduce, describe, and illustrate three new genera, *Italiomyces*, *Ranulospora*, and *Tenuiappendicula*, based on newly collected specimens. Another three new genera, *Alborbis*, *Breviappendix*, and *Paragnomonia* are introduced to accommodate *Cryptodiaporthe galericulata*, *Gnomonia rosae*, *G. rostellata*, *G. rubi*, and *G. fragariae*. *Alborbis galericulata* comb. nov., *Breviappendix rubi* comb. nov., *B. rosae* comb. nov., *B. rostellata* comb. nov. and *Paragnomonia fragariae* comb. nov. are proposed. Two new species, *Sydowiella centaureii* and *Sillia karstenii*, are introduced with descriptions and illustrations. *Winterella* is replaced with a new genus *Tortilispora* typified by *T. aurantiaca*. Three new combinations are proposed in *Tortilispora*. We synonymize *Stegophora* under *Lambro* based on morphological similarities. An updated key for genera of Sydowiellaceae is presented.

**Key words** – Alborbis, Breviappendix, Italiomyces, Paragnomonia, Ranulospora, Sillia karstenii, Sydowiella centaureii, Tenuiappendicula, Tortilispora.

Submitted 10 January 2017, Accepted 8 February 2017, Published 20 February 2017 Corresponding Author: Itthayakorn Promputtha – e-mail – ppam118@gmail.com

<sup>&</sup>lt;sup>1</sup>Key Laboratory for Plant Diversity and Biogeography of East Asia, Kunming Institute of Botany, Chinese Academy of Science, Kunming 650201, Yunnan, China

<sup>&</sup>lt;sup>2</sup>East and Central Asia, World Agroforestry Centre, Kunming 650201, Yunnan, China

<sup>&</sup>lt;sup>3</sup>Centre of Excellence for Fungal Research, Mae Fah Luang University, Chiang Rai, Thailand

<sup>&</sup>lt;sup>4</sup>Department of Crop Sciences, College of Agricultural and Marine Sciences, Sultan Qaboos University, P.O. Box 34, Al-Khod 123, Oman

<sup>&</sup>lt;sup>5</sup>Department of Health Sciences, Faculty of Science, University of Mauritius, Mauritius, 80837

<sup>&</sup>lt;sup>6</sup>Department of Biology, Faculty of Science, Chiang Mai University, Chiang Mai 50200, Thailand

<sup>&</sup>lt;sup>7</sup>A.M.B. Gruppo Micologico Forlivese, Antonio Cicognani, Via Roma 18, Forlì, Italy

<sup>&</sup>lt;sup>8</sup>A.M.B. Circolo Micologico, Giovanni Carini, 314 Brescia, Italy

<sup>&</sup>lt;sup>9</sup>Società per gli Studi Naturalistici della Romagna, 144 Bagnacavallo, RA, Italy

#### Introduction

Sydowiellaceae Lar. N. Vassiljeva was established to accommodate the genus *Sydowiella* Petr., which is typified by *S. fenestrans* (Duby) Petr., and a collection of morphologically diverse fungal taxa whose intergeneric relationships are still uncertain (Maharachchikumbura et al. 2015). Sydowiellaceae is a phylogenetically, well-supported, monophyletic family within Diaporthales (Kruys & Castlebury 2012), but is morphologically poorly studied. To date, Sydowiellaceae comprises eleven genera (Maharachchikumbura et al. 2015, 2016) and 43 species (Index Fungorum 2017).

The family Sydowiellaceae is generally characterized by taxa with solitary or aggregated, erumpent to immersed ascomata, with wide papilla, a peridium comprising an outer layer of brown-walled cells of *textura globulosa* and inner layer of hyaline cells of *textura angularis*, cylindrical to cylindro-clavate or broadly fusoid unitunicate asci, with a J-, sometimes bi-lobed apical ring, and hyaline or greenish-brown, 0–11-septate, ellipsoidal or filiform ascospores that sometimes have apical and basal appendages (Maharachchikumbura et al. 2016).

The family Sydowiellaceae also includes the genera Cainiella E. Müll. typified by C. johansonii (Rehm) E. Müll., Calosporella J. Schröt typified by C. innesii (Curr.) J. Schröt., Chapeckia M.E. Barr typified by C. nigrospora (Peck) M.E. Barr, Hapalocystis Auersw. ex Fuckel typified by H. berkeleyi Fuckel, Lambro Racib. typified by L. insignis Racib., Rossmania Lar. N. Vassiljeva typified by R. ukurunduense Lar. N. Vassiljeva, Stegophora Syd. & P. Syd. typified by S. ulmea (Fr.) Syd. & P. Syd., Sillia P. Karst. typified by S. ferruginea (Pers.) P. Karst., Uleoporthe Petr typified by U. orbiculata (Syd. & P. Syd.) Petr. and Winterella (Sacc.) Kuntze typified by W. aurantiaca (Wehm.) J. Reid & C. Booth (Maharachchikumbura et al. 2015, 2016). Due to morphological plasticity, the taxonomic affinities of most genera within Sydowiellaceae have been uncertain and they were referred to Diaporthales genera incertae sedis (Rossman et al. 2006). Despite morphological based taxonomic discrepancies, DNA sequence based phylogenies have consistently reported Sydowiellaceae to be a well-supported monophyletic family comprising the above-mentioned 2006, Rossman et al. 2007, Kruys & Castlebury Maharachchikumbura et al. 2015, 2016). Stilbospora Pers., Hendersonia E.J. Butler, and Dothiorella Sacc. have been reported as asexual morphs for Hapalocystis associated with ascomata on the natural substrate (Wehmeyer 1941, Barr 1978, Jaklitsch & Voglmayr 2004), While Glawe (1985) and Liu et al. (2015) reported a phoma-like asexual morph for H. berkeleyi in culture. Cylindrosporella Höhn. was reported as the asexual morph of Stegophora by Kobayashi (1970).

Members of Sydowiellaceae occur on herbaceous plants, dicotyledonous and hardwood trees, as saprobes, endophytes or pathogens. *Stegophora ulmea* was reported as a pathogen on living leaves of *Ulmus americana*, while *Sydowiella depressula* was reported on leaves and stem cankers of cultivated black berry (*Rubus fruticosus*) (Nordskog et al. 2003). *Sydowiella fenestrans* has been recorded on leaves and dead stems of various plants. *Cainiella johansonii* was reported on both living and dead leaves and petioles of *Dryas* species, while *Cainiella borealis* was found as a saprobe on dead branches of *Cassiope tetragona* (Kruys & Castlebury 2012). *Hapalocystis berkeleyi* forms stromatic fruiting bodies on dead woody plants. Collections of *Uleoporthe orbiculata* were recorded on living and fallen leaves. There are reports that *U. orbiculata* and many other taxa in this family may have biotrophic lifestyles (Cannon 2001, De Silva et al. 2016).

Species in Gnomoniaceae appear to be similar to Sydowiellaceae. Kruys & Castlebury (2012) showed that *Cryptodiaporthe galericulata*, *Gnomonia fragariae*, *G. rostellata*, and *G. rubi* (Gnomoniaceae) clustered in Sydowiellaceae based on ITS sequence analysis. Morocko (2006) also showed a close affinity of *Gnomonia rosae* to Sydowiellaceae. Voglmayr &

Jaklitsch (2014) introduced *Calosporella* for *Prosthecium innesii* and accommodated it as *Calosporella innesii*.

Here we provide an updated phylogeny based on DNA sequence analyses of taxa of Sydowiellaceae to clarify intergeneric relationships. In addition, we introduce and describe three new monotypic genera based on fresh collections, another three new genera based on combinations of existing species with molecular data and two new species in existing genera based on fresh collections. *Winterella* is replaced with a new genus *Tortilispora* typified by *T. aurantiaca*. *Stegophora* is synonymized under *Lambro* based on its morphology. An updated taxonomic key of genera in Sydowiellaceae is also provided.

#### **Materials & Methods**

# Specimen collection, morphological examination and photomicrography

Collections were made in Italy from 2013 to 2016. Specimens were placed in paper bags and collection details noted. Samples were brought to the laboratory and examined under a stereomicroscope to observe the characteristics of the fruiting bodies. Macromorphological characters were photographed using a digital camera fitted to the stereomicroscope. A few fruiting bodies were transferred to a drop of water mounted on a slide using fine needles or forceps for rehydration and they were crushed to observe internal structures. Cross sections of fruit bodies were hand cut with a razor blade and mounted in a drop of water on a slide. Morphological characteristics of ascomata/conidiomata, asci, ascospores/conidia and other tissues associated with ascomata/conidiomata were photographed using a digital camera fitted to ta compound microscope. Melzer's reagent, Indian ink and 5 % KOH were used where necessary to stain fungal structures. All microphotographs were arranged using Adobe Photoshop CS6 (v. 13.0.1.1) and all measurements were made with Tarosoft (R) Image Frame Work (v. 0.9.0.7). Specimens were preserved and deposited at Mae Fah Luang University herbarium (MFLU) and BIOTEC Bangkok Herbarium (BBH). Facesoffungi and Index Fungorum numbers were registered (Jayasiri et al. 2015, Index Fungorum 2017). New species were resolved following recommendations as outlined by Jeewon & Hyde (2016).

#### Single spore isolation

A few fruiting bodies were picked from substrates using a sterilized needle and placed in a few drops of sterilized distilled water on a sterilized cavity slide as detailed in Chomnunti et al. (2014). Fruiting bodies were crushed to separate the spores and obtain a spore suspension. The suspension was inoculated on to malt extract agar (MEA) and incubated at 18 °C. Germinating ascospores were aseptically transferred to new culture plates. Colonies were photographed and characters were noted. Living cultures are deposited at Mae Fah Luang University Culture Collection (MFLUCC) and the Thailand Bioresource Research Center (TBRC).

#### **Promoting asexual morphs**

Dried and sterilized grass leaves or pine needles were laid parallel to the colony margin of pure cultures which were grown on water agar (Phookamsak et al. 2015). Conidiomata produced on both water agar and sterilized pine needles were observed after few weeks of incubation at 18 °C. Morphological characteristics were examined as mentioned above.

#### Herbarium material examination and re-drawing

Types and other relevant authentic specimens were loaned from accessible herbaria (New York State Museum (NYS), Swedish Museum of Natural History (S),

Naturhistorisches Museum Wien (W), and International Mycological Institute (IMI)). A small part of the herbarium specimen was cut and rehydrated using water or 5 % KOH. Micromorphological characters were observed from rehydrated ascomata and photography was done as previously described. Some specimens were redrawn based on previously illustrated authentic specimens.

# DNA extraction, PCR amplification, sequencing and phylogenetic analyses

Scraped fresh fungal mycelia grown on MEA for 4 weeks at 18 °C, or internal content of ascomata or conidiomata were used for genomic DNA extraction using a modified protocol of Jeewon et al. (2002). PCR amplification and sequencing of large subunit ribosomal RNA (LSU) region using the primer pair LROR/LR5, internal transcribed spacer (ITS) regions using primer pair ITS4/ITS5, RNA polymerase II gene (RPB2) region using the primer pair fRPB2-5F/fRPB2-7cR, and translation elongation factor (TEF) region using primer pair 728F/986R were performed (Vilgalys & Hester 1990, White et al. 1990, Carbone & Kohn 1999, Liu et al. 1999).

Each PCR reaction contained 0.3 µl of TaKaRa Ex-Taq DNA polymerase, 12.5 µl of 2 × PCR buffer, 2.5 µl of dNTPs, 1 µl of forward and reverse primers, 1 µl of DNA template and was adjusted with 6.5 µl of double-distilled water to a total volume of 25 ml. Amplification reactions were performed in a thermal-cycler (BIORAD 1000TM Thermal Cycler, Bio-Rad Laboratories, Hercules, California). The temperature profile for both ITS and LSU was an initial denaturing step for 2 min at 94 °C, followed by 35 amplification cycles of denaturation at 94 °C for 60 s, annealing at 58 °C for 60 s and extension at 72 °C for 90 s and a final extension step of 72 °C for 10 min (Jeewon et al. 2004, Phillips et al. 2008). The temperature profile for the RPB2 was, initial denaturation at 94 °C for 2 min, followed by 35 amplification cycles of denaturation at 95 °C for 45 s, annealing at 57 °C for 50 s and extension at 72 °C for 90 s. (Liu et al. 1999). The temperature profile for the TEF was, initial denaturation at 94 °C for 2 min, followed by 35 amplification cycles of denaturation at 95 °C for 30 s, 58 °C for 50 s, 72 °C 60 s. (Glass & Donaldson 1995). All amplified PCR products were determined by electrophoresis at 90 V/cm for 40 min in 1% agarose gel stained with ethidium bromide (0.5 mg/mL). The gel was visualized under a UV transilluminator to estimate the fragment size. PCR products were purified and sequenced with both primers at the Sunbiotech Company, Beijing, China. Sequences were edited and assembled with DNASTAR Lasergene (v.7.1) to obtain consensus sequences. The sequences generated in this study were supplemented with additional sequences obtained from GenBank (Table 1) based on blast searches and published literature. Multiple sequence alignments were generated with MAFFT v. 7 (http://mafft.cbrc.jp/alignment/server/index.html) and the alignment was manually improved with BioEdit v. 7.0.5.2 (Hall 1999).

Phylogenetic analyses of the sequence data consisted of Maximum Likelihood, Bayesian inference and maximum parsimony. Maximum likelihood (ML) analysis was performed by RAxMl GUI v.1.3 (Stamatakis et al. 2008, Silvestro & Michalak 2012). The search strategy was set to rapid bootstrapping and the analysis was carried out using the GTRGAMMAI model of nucleotide substitution with 1000 replicates. The model of evolution was estimated by using MrModeltest 2.3 (Nylander 2004) and GTR+I+G was selected as the model for Bayesian analyses. Bayesian inference in MrBayes v. 3.2.1 (Ronquist et al. 2012) was performed with default settings, running four chains over 3 million generations and sampling each 100<sup>th</sup> tree. The first 6000 of the 3000000 saved trees were discarded and the consensus tree was based on the remaining 2994000 trees. Maximum parsimony analysis (MP) was performed using PAUP v. 4.0b10 (Swofford 2003). Ambiguously aligned regions were excluded and gaps were treated as a missing data. All characters were unordered and given equal weight. Trees were inferred using the heuristic

**Table 1.** Isolates used in the analyses and their GenBank and culture accession numbers. Newly generated sequences are in bold.

Taxa	Culture accession	TTC		ession numbers	and the
4	numbers	ITS	LSU	RPB2	TEF
Asterosporium asterospermum	KT2125	-	AB553743	-	-
Asterosporium asterospermum	CBS 112404	-	AB553745	-	-
Asterosporium asterospermum	KT2101	-	AB553742	-	-
Asterosporium asterospermum	KT2138	- LXX502472	AB553744	-	-
Breviappendix rubi	MFLUCC 16–1205	KY523473	KY523491	-	-
Breviappendix rubi	MFLUCC 16–1205A	KY523474	KY523492	-	-
Breviappendix rubi	MFLUCC 16–1205B	KY523475	KY523493	-	-
Cainiella borealis	Kruys 725 (UPS)	JF701921	-	-	-
Cainiella johansonii	Kruys 731 (UPS)	- IE:01065	JF701920	-	-
Calosporella innesii	AR 3639	JF681965	EU683071	-	-
Calosporella innesii	AR 3925	JF681966	-	-	-
Calosporella innesii	AR 3831	JF681964	-	-	-
Chapeckia nigrospora	CBS 125532	JF681957	EU683068	-	-
Cryphonectria macrospora	CBS 109764	EU199182	AF408340	EU220029	-
Cryphonectria parasitica	ATCC 38755	AY141856	EU199123	DQ862017	EU222014
Cryptodiaporthe galericulata	AR 3890	JF681967	-	-	-
Cryptodiaporthe galericulata	AR 4004	JF681968	-	-	-
Cryptodiaporthe galericulata	AR 3811	JF681969	-	-	-
Cryptodiaporthe galericulata	AR 4027	JF681970	-	-	_
Cytospora hippophaes	CFCC 89640	KF765682	KF765698	KF765714	KP310865
Diaporthe eres	AR 5193	KJ210529	-	-	KJ210550
Diaporthe eres	PS57	JF430494	JF704176	-	-
Gnomonia cf rosae	BPI 877452B	EU254935	EU255157	-	-
Gnomonia fragariae	CBS 146.64	EF212844	EF212856	-	-
Gnomonia fragariae	UN22	EF212833	EF212851	-	-
Gnomonia fragariae	F 5.6	EF212840	EF212855	-	-
Gnomonia fragariae	F 1.4	-	EF212853	-	-
Gnomonia fragariae	M1	EF212834	EF212852	-	-
Gnomonia fragariae	GF2	EF212832	EF212850	-	-
Gnomonia fragariae	SER1	EF212841	EF212854	-	-
Gnomonia fragariae	SER7	EF212843	-	-	-
Gnomonia fragariae	SER4	EF212842	-	-	-
Gnomonia fragariae	L56	EF212838	-	-	-
Gnomonia fragariae	O3	EF212837	-	-	-
Gnomonia fragariae	S12	EF212836	-	-	-
Gnomonia fragariae	S7	EF212835	-	-	_
Gnomonia gnomon	CBS 829.79	AY818957	AY818964	-	EU221905
Gnomonia gnomon	CBS 199.53	AY818956	AF408361	DQ470922	EU221885
Gnomonia rosae	CBS 850.79	EF212846	EF212858	-	_
Gnomonia rostellata	AR 3533	JF681962	-	_	_
Gnomonia rostellata	AR 3987	JF681963	_	_	_
Gnomonia rubi	CBS 102.72	EF212847	EF212859	_	_
Hapalocystis berkeleyi	AR 3851		EU683069	_	_
Hapalocystis berkeleyi	AR 3870	JF681958	-	_	_
Hapalocystis berkeleyi	MFLUCC 13–0662	-	KP744486	_	_
Hapalocystis occidentalis	WU 24705	_	AY616231	_	_
Hercospora tiliae	AR 3526	_	AF408365	_	_
Italiomyces centaureii	MFLU 16–2866	KY523478	KY523494	_	_
Italiomyces centaureii Italiomyces centaureii	MFLU 16–2867	KY523477	KY523495	_	_
Italiomyces centaureii Italiomyces centaureii	MFLUCC 14–0849	KY523477	KY523496	_	_
Lamproconium desmazieri	MFLUCC 14-0849 MFLUCC 15-0871	KX430136	KX430137	_	_
Lamproconium desmazieri Lamproconium desmazieri	MFLUCC 13-08/1 MFLUCC 14-1047	KX430130 KX430132	KX430137 KX430133	_	_
Lamproconium desmazieri Lamproconium desmazieri	MFLUCC 14–1047 MFLUCC 15–0870	KX430132 KX430134	KX430135	-	_
=				-	-
Lamproconium desmazieri Onkiognomonia rubi idaei	MFLUCC 15–0873	KX430140	KX430141	-	-
Ophiognomonia rubi-idaei	BPI 877638 MELLICC 12, 0702	EU254938	- VV502407	-	-
Ranulospora alnii	MFLUCC 13-0793	KY523479	KY523497	-	-
Ranulospora alnii	MFLUCC 13–0793A	KY523480	KY523498	-	-
Ranulospora alnii	MFLU 16-2868	KY523481	KY523499	-	-
Rossmania ukurunduensis	AR 3484	-	EU683075	-	-
Sillia italica	MFLU 16-0056A	KY523484	KY397949	-	-

**Table 1.** continued. Isolates used in the analyses and their GenBank and culture accession numbers. Newly generated sequences are in bold.

Taxa	Culture accession		GenBank accession numbers		
	numbers	ITS	LSU	RPB2	TEF
Sillia italica	MFLU 16-0056B	KY523485	-	-	-
Sillia ferruginea	CBS 126567	JF681959	EU683076	-	-
Sillia karstenii	MFLU 16-2864	KY523482	KY523500	KY501636	-
Sillia karstenii	MFLU 16-2865	KY523483	KY523501	-	-
Stegonsporium acerophilum	CBS 117025	EU039982	EU039993	KF570173	-
Stilbospora macrosperma	CBS 121883	JX517290	JX517299	KF570196	-
Sydowiella centaureii	MFLU 16-2860	KY523486	KY523502	-	-
Sydowiella centaureii	MFLU 16-2858	KY523487	KY523503	-	-
Sydowiella depressula	CBS 813.79	-	EU683077	-	-
Sydowiella fenestrans	CBS 125530	JF681956	EU683078	-	-
Sydowiella urticola	MFLUCC 13-0665	-	KY523504	-	-
Tenuiappendicula alnicola	MFLU 16-1265	KY523488	KY523505	-	-
Tenuiappendicula alnicola	MFLU 16-2865	KY523489	KY523506	-	-
Tenuiappendicula alnicola	MFLUCC 16-1452	KY523490	KY523507	-	-
Valsa ambiens	CFCC 89622	KR045698	AF362564	KU710944	KU710911
Winterella aurantiaca	AR 4022	JF681960	-	-	-

search option with TBR branch swapping and 1000 random sequence additions. MaxTrees were set up to 1000, branches of zero length were collapsed and all multiple parsimonious trees were saved. Tree length (TL), consistency index (CI), retention index (RI), rescaled consistency index (RC), homoplasy index (HI), and log likelihood (-ln L) (HKY model) were calculated for trees generated under optimality criteria. The robustness of the most parsimonious trees was evaluated by 1000 bootstrap replications resulting from maximum parsimony analysis, each with ten replicates of random step-wise addition of taxa (Felsenstein 1985). The Kishino–Hasegawa tests (Kishino & Hasegawa 1989) were performed to determine whether the trees inferred under optimality criteria were significantly different. Trees were figured in Treeview (Page 1996). The final alignments and the trees obtained were deposited in TreeBASE (http://purl.org/phylo/treebase/phylows/study/TB2:S20545?x-access-code=e3bdc851b5ecb55a1daa3054ff455627&format=html) and are available under study accession no. S20545.

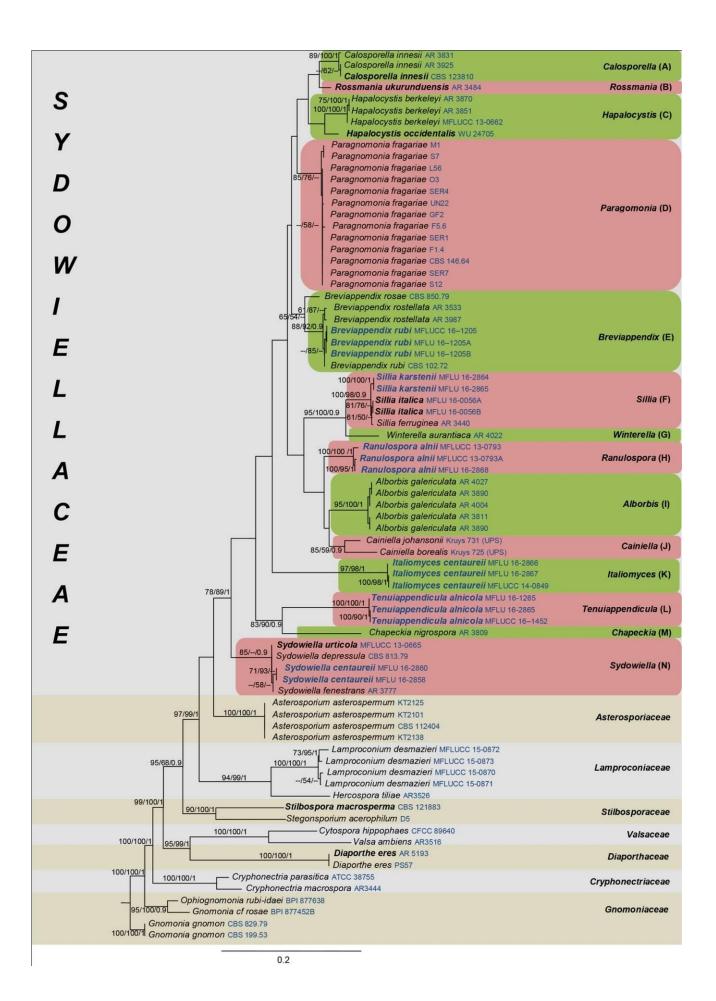
#### **Results and Discussion**

The phylogenetic analyses were carried out using combined LSU, ITS, RPB2 and TEF sequence data from taxa of Diaporthales as listed in Table 1. Taxon sampling was based on GenBank blast searches and recent publications (Morocko 2006, Kruys & Castlebury 2012, Maharachchikumbura et al. 2015, 2016). The final multiple alignment data comprised 76 taxa and 3283 characters including gaps. Overall, the topologies obtained from the different phylogenetic methods and parameters were congruent and the best scoring RAxML tree is illustrated together with bootstrap support values (TL = 3526, CI = 0.664, RI = 0.853, HI = 0.336, RC = 0.567) in Fig. 1. Maximum likelihood analysis based on 1000 bootstrap replicates, yielded a tree with the likelihood value of ln: -19847.102701 with the following model parameters: alpha: 0.779450,  $\Pi(A)$ : 0.238610,  $\Pi(C)$ : 0.265774,  $\Pi(G)$ : 0.286696, and  $\Pi(T)$ : 0.20892.

The separation of Sydowiellaceae from other diaporthalean taxa (Fig. 1) is well-supported and Sydowiellaceae comprises 14 clades (Clades A-N) representative of the generic placement. The type genus *Sydowiella* (Clade N) forms a moderately-supported clade within the family and here we introduce a new species *S. centaureii* which is morphologically and phylogenetically distinct. The phylogeny indicates that *Tenuiappendicula* (Clade L) constitutes an independent monophyletic clade with high statistical support, while *Chapeckia* 

(Clade M) has a sister relationship to Tenuiappendicula, but is phylogenetically distinct (Fig. 1). Chapeckia and Tenuiappendicula species have pseudostromata with very thin peridia and 1-septate, oval ascospores. The asexual morph of Tenuiappendicula is illustrated here and ITS, LSU sequences derived from conidiomatal DNA are similar to the sexual morph sequences. A new genus Italiomyces (Clade K) is introduced, with I. centaureii as the type species. Italiomyces differs from other genera in Sydowiellaceae in having mostly biguttulate, rarely 4-guttulate ascospores, with minute, spine-like apical appendages at both ends. The new genus Alborbis (Clade I) is introduced to accommodate Cryptodiaporthe galericulata which is morphologically and phylogenetically distinct from the type species of Cryptodiaporthe. This taxonomic circumscription of C. galericulata was suggested by Kruys & Castlebury (2012). Hence, we introduce Alborbis to accommodate Cryptodiaporthe galericulata with a new combination as Alborbis galericulata. Alborbis is sister to Cainiella (Clade J) and the new genus Ranulospora (Clade H), typified by R. alnii, is introduced as sister to Alborbis and Cainiella. Sillia (Clade F) and Tortilispora (Clade G) both have cylindrical asci containing long, filiform ascospores with pointed ends. Tortilispora, however, has a diaporthoid configuration of ascomata with 4-spored asci and septate paraphyses, while Sillia has a diatrypoid configuration of ascomata with 8-spored asci and aseptate paraphyses (Tibpromma et al. 2017). This morphological segregation is also supported in the phylogeny (Fig. 1). Reid & Booth (1987) studied Tortilispora (= Winterella), but further collections are needed together with support from molecular data as our assumption is based on only a single species. A new Sillia species, S. karstenii is introduced, as it constitutes a highly-supported subclade and is phylogenetically apart from other species (Fig. 1). The new species also differs from other Sillia species in having a very poorly developed stroma and ascospores with pointed ends (Fig. 15). A new genus Breviappendix (Clade E) is introduced to accommodate Gnomonia rubi and G. rostellata which were previously accommodated in Gnomoniaceae. Breviappendix forms a distinct clade in the family with moderate support. Breviappendix rosae, B. rostellata and B. rubi are introduced as new combinations for Gnomonia rosae, G. rostellata and G. rubi respectively and Breviappendix rubi is designated as the type species of the genus. A new genus Paragnomonia (Clade D) is introduced to accommodate Gnomonia fragariae which was previously placed in Gnomoniaceae. The placement in Sydowiellaceae was also noted by Mejía et al. (2011). Calosporella (Clade A) forms a well-supported separate clade as a wellresolved genus. This monotypic genus was recently revisited by Voglmayr & Jaklitsch (2014). Calosporella is sister to Rossmania (Clade B) which is represented by R. ukurunduensis. Hapalocystis (Clade C) does not have any support and Hapalocystis occidentalis is basal to all other Hapalocystis berkeleyi strains. Hapalocystis berkeleyi and H. occidentalis are morphologically similar, but differ primarily in spore ornamentation and size of appendages. Our phylogeny also supports the concept that Hapalocystis occidentalis can be considered as a separate species from H. berkeleyi based on morphological differences mentioned and phylogenetic support for this has been reported in other studies (Jeewon et al. 2002).

**Figure 1** – Consensus tree resulting from a maximum likelihood analysis of combined LSU, ITS, RPB2 and TEF sequence alignment of genera of Diaporthales. genera in Sydowiellaceae are indicated in pink and green blocks while families are indicated in ash-coloured blocks. Maximum parsimony bootstrap support (MPB above 50%), RAxML bootstrap support (MLB above 50%), and Bayesian posterior probabilities (PP above 0.9) are given at the nodes (MPB/MLB/PP). The scale bar represents the expected number of changes per site. The tree is rooted to *Gnomonia gnomon* (CBS 829.79; CBS 199.53). Ex-type strains are in bold and new strains are in blue bold.



# **Taxonomy**

**Sydowiellaceae** Lar.N. Vassiljeva, Pirenomits. Lokuloaskomits. Severa Dal'nego Vostoka (Leningrad): 210 (1987)

Saprobic on dead branches of deciduous trees or parasitic on both living and dead leaves and petioles of temperate shrubs. Sexual morph – Stromata present or absent. If erumpent through cracks, roughly circular, pseudoparenchymatous prosenchymatous with few ascomata in a stroma, erumpent, sometimes with simple to complex stellate splits in the substrate, with poorly to well-developed black lines around the stroma and internal eustromatic tissues with thick-walled, glassy, hyaline cells of textura globulosa to textura angularis around the basal regions of ascomata and dark brown-walled cells of textura globulosa to textura angularis around the ostioles, sometimes becoming rouge in 10 % KOH. If pseudostromatic, pseudostromata pustulate on surface, circular, containing up to 20 ascomata. Ectostromatic disc erumpent, conspicuous, white to light grey, becoming dark at maturity, fusoid to circular, containing 1–20 distinctly protruding ostioles. Entostroma crumbly, brownish. Ascomata scattered, solitary or aggregated, erumpent to immersed, globose to subglobose, coriaceous, brown to black, ostiolate, papillate. Papilla wide, erumpent from the substrate, internal wall covered by hyaline, filamentous periphyses. Peridium comprising outer layer of brown-walled cells of textura angularis to globulosa and inner layer of, hyaline cells of textura angularis. Hamathecium comprising cellular or filamentous, hyaline, septate or aseptate paraphyses or aparaphysate. Asci 8-spored, unitunicate, cylindrical to cylindro-clavate or broadly fusoid, short pedicellate, apex blunt, with J-, apical ring or without apical ring, sometimes apex wide and flat, with multi-layered, J-, apical ring. Ascospores overlapping uniseriate, biseriate to triseriate, kidney-shaped, filamentous, oval or ellipsoidal, hyaline or initially hyaline, becoming brown with age, 0–11septate, slightly constricted or not constricted at the septum, thick and smooth-walled, sometimes multi-guttulate, sometimes hyaline, with tapering bipolar appendages or appendages basal only. Asexual morph - Coelomycetous. Conidiomata pycnidia, superficial to erumpent, aggregated 3-5 in a single group, globose, orange to brown, conidiomatal wall thick-walled, with orange cells of textura angularis. Conidiophores annelidic, branched, hyaline, short, few conidiogenous cells arising from a single conidiophore, attached to conidiomatal wall. Conidiogenous cells cylindrical, hyaline, bottle-shaped, septate, pointedends, phialidic. Conidia ellipsoid or cylindroid, unicellular or multi-septate, hyaline or brown, smooth-walled.

Type genus – *Sydowiella* Petr.

Notes — Barr (1978) placed *Sydowiella* in the family Pseudovalsaceae (subfamily Pseudovalsoideae, tribe Ditopelleae) based on its morphological distinctiveness. There have been also a few suggestions to place *Sydowiella* in Gnomoniaceae. However, Monad (1983), did not accept the placement of *Sydowiella* in Gnomoniaceae because of absence of stromatic tissues. Vasilyeva (1987) introduced the family Sydowiellaceae based on *Sydowiella fenestrans* (Duby) Petr. and this family was accommodated in the order Ceratostomatales. Rossman et al. (2007) included *Chapeckia*, *Hapalocystis*, *Rossmania*, *Sillia*, and *Stegophora* in Sydowiellaceae. Sogonov et al. (2008) noted a close affinity of *Lambro* and *Uleoporthe* with this family, while Kruys & Castlebury (2012) analyzed ITS sequence data for genera in the family. The phylogenetic position of *Cainiella* in Sydowiellaceae was resolved and *Winterella* was also included in the family based on molecular data. *Calosporella* was introduced and placed in Sydowiellaceae to accommodate *Prosthecium innesii* by Voglmayr & Jaklitsch (2014).

#### List of accepted genera and species in Sydowiellaceae

Based on Index Fungorum 2017, Maharachchikumbura et al. (2015, 2016), Mejia et al. (2008), MycoBank (2017), Tibpromma et al. (2017)

Alborbis Senan. & K.D. Hyde

Alborbis galericulata (Tul. & C. Tul.) Senan. & K.D. Hyde

Breviappendix Senan. & K.D. Hyde

Breviappendix rosae (Fuckel) Senan. & K.D. Hyde

Breviappendix rostellata (Fr.) Senan. & K.D. Hyde

Breviappendix rubi (Rehm) Senan. & K.D. Hyde

Cainiella E. Müll.

Cainiella borealis M.E. Barr

Cainiella johansonii (Rehm) E. Müll.

Calosporella J. Schröt.

Calosporella innesii (Curr.) J. Schröt.

Chapeckia M.E. Barr

Chapeckia nigrospora (Peck) M.E. Barr Chapeckia ribesia (Sacc. & Scalia) M.E. Barr

Hapalocystis Auersw. ex Fuckel

Hapalocystis berkeleyi Auersw. ex Fuckel

Hapalocystis bicaudata Fuckel

Hapalocystis corni (Wehm.) M.E. Barr

Hapalocystis corticalis (Schwein.) M.E. Barr

Hapalocystis occidentalis Jaklitsch & Voglmayr

Hapalocystis ulmi (Wehm.) M.E. Barr

Hapalocystis vexans Speg.

*Italiomyces* Senan., Camporesi & K.D. Hyde

*Italiomyces centaureii* Senan., Camporesi & K.D. Hyde

Lambro Racib.

= Stegophora Syd. & P. Syd Lambro insignis Racib.

Lambro symploci G.C. Zhao & R.L. Zhao

Paragnomonia

Paragnomonia fragariae (Kleb.) Senan. & K.D. Hyde

*Ranulospora* Senan., Camporesi & K.D. Hyde

Ranulospora alnii Senan., Camporesi & K.D. Hyde

Rossmania Lar.N. Vassiljeva

Rossmania aculeata (Petch) Lar.N. Vassiljeva

Rossmania ukurunduensis Lar.N. Vassiljeva Sillia P. Karst.

Sillia albofusca Höhn.

Sillia biformis Rick

Sillia celastrina R. Rao

Sillia ferruginea (Pers.) P. Karst.

Sillia italica de Silva, Camporesi & K.D. Hyde

Sillia kamatii Tilak, S.B. Kale & S.V.S. Kale

Sillia karstenii Senan., Camporesi & K.D. Hyde

Sillia theae Hara

*Tenuiappendicula* Senan., Camporesi & K.D. Hyde

Tenuiappendicula alnicola Senan., Camporesi & K.D. Hyde

Tortilispora (Sacc.) Senan. & K.D. Hyde

= *Cryptosporella* Sacc.

Tortilispora albofusca (Cooke & Ellis)

Senan. & K.D. Hyde

Tortilispora aurantiaca (Wehm.) Senan. & K.D. Hyde

Tortilispora cinctula (Cooke & Peck)

Senan. & K.D. Hyde

# Key to genera within the family Sydowiellaceae

1. Štromata conspicuous, well-developed	7
1. Stromata absent, inconspicuous, poorly developed	
2. Ascomata valsoid, stromatic tissues do not form any colour with 10% KOH	
2. Ascomata diatrypoid, stromatic tissues form dull red with 10% KOH	
3. Ascospores oval to fusoid-oval, 1-septate, hyaline or hyaline to brown	
3. Ascospores fusiform to ellipsoid with long filiform basal cell, 2-septate, hyaline	
4. Ascomata solitary	_
4. Ascomata aggregated	
5. Asci 4-spored	
5. Asci 8-spored	
6. Ascospores apiosporous	
6. Ascospores non-apiosporous	
7. Ascospores oval, hyaline to brown, not constricted at the septa	
7. Ascospores fusoid, hyaline, constricted at the septa	
8. Ascospores non-appendaged, globules at the center of each cell	
8. Ascospores appendaged, globules at the septa of the spore	-
9. Ascospores oval, short fusoid to ellipsoidal	
9. Ascospores long fusoid cylindrical	
10. Ascospores 1-septate	
10. Ascospores multi-septate	
11. Ascospores with long strip-like appendages	
11. Ascospores with short, appendages	
12. Ascospores ellipsoid to cylindrical, usually 4-guttulate	
12. Ascospores fusiform, usually multi-guttulate	
13. Ascospores hyaline to brown, ellipsoidal with broadly rounded ends, 1–3-septate, constricted at the septa	<u> </u>
13. Ascospores hyaline, fusoid to oblong, 3–4-eusepta, not constricted at the septa	
14. Ascospores 0–5-septate	<del>-</del>
14. Ascospores 6–11-septate	Rossmania

### *Sydowiella* Petr., Annls mycol. 21(1/2): 30 (1923)

Saprobic or pathogenic on plants. Sexual morph – Ascomata solitary, scattered, erumpent to immersed, globose, brown, papillate, ostiolate. Papilla short and wide, comprising brown, elongated cells. Peridium comprising an inner layer of thin-walled, hyaline cells of *textura angularis* and outer layer of thick-walled, brown cells of *textura angularis*. Hamathecium comprising cellular, septate, hyaline paraphyses. Asci 8-spored, unitunicate, cylindrical, short-pedicellate, apex blunt, with a J-, apical ring. Ascospores overlapping uniseriate, oval to ellipsoid, 1-median septate, hyaline. Asexual morph – Undetermined.

Type species – Sydowiella fenestrans (Duby) Petr.

Basionym – Sphaeria fenestrans Duby

Notes — *Sydowiella* is characterized by solitarily, erumpent ascomata without stromatic tissues, a melanconis-like hamathecium, and 2-celled ascospores (Kobayashi 1970). The type species *Sydowiella fenestrans* was illustrated in Maharachchikumbura et al. (2016). *Sydowiella* comprises ten species (Index Fungorum 2017): *S. ambigua* (Mouton) Munk, *S. azukiae* I. Hino & Katum., *S. dakotensis* Petr., *S. depressula* (P. Karst.) M.E. Barr, *S. fenestrans* (Duby) Petr., *S. indica* Bilgrami & Rizwi, *S. juncina* Spooner, *S. stellatifolii* Marinc. et al. and *S. uticola* Senan. et al. Here we introduce an additional species.

#### Sydowiella centaureii Senan., Camporesi & K.D. Hyde, sp. nov.

Index Fungorum number – IF552715, Facesoffungi number – FoF 2834 Fig. 2

Etymology – Name referring to the host genus *Centaurea*.

Holotype – MFLU 16–2858

Saprobic on dead branches of *Centaurea* L. Sexual morph – Stromata pseudoparenchymatous, comprising outer dark brown to black line made by hyphal cells and inner light yellow to hyaline, large cells of *textura globosa*. Ascomata 450–510 µm high, 280–355 µm diameter ( $\bar{x}=504\times314$  µm, n = 20), scattered, solitary, immersed to erumpent, erect or slanted, globose to subglobose, coriaceous, black, papillate, ostiolate. Papilla 185–195 µm high, 110–120 µm wide ( $\bar{x}=190\times115$  µm, n = 20), internally covered with hyaline periphyses. Peridium 15–25 µm wide ( $\bar{x}=21$  µm, n = 20), comprising inner layer of hyaline, thin-walled, compressed cells of *textura angularis* and outer layer of dark brown, thick-walled, cells of *textura angularis*. Hamathecium comprising 4–7 µm ( $\bar{x}=6$  µm, n = 20) wide, septate, cellular paraphyses. Asci 85–110 × 10–15 µm ( $\bar{x}=102\times14$  µm, n = 20), 8-spored, unitunicate, cylindrical, short-pedicellate, with distinct, bilobed, J- apical ring. Ascospores 10–15 × 5–8 µm ( $\bar{x}=13\times6.8$  µm, n = 20), overlapping uniseriate, oval to widely fusiform, hyaline, 1-septate, slightly constricted at the septum, with a large guttule in each cell. Asexual morph – Undetermined.

Material examined – ITALY, Province of Forlì-Cesena, Fiumicello di Premilcuore, dead stem of *Centaurea* sp. (Asteraceae), 2 April 2014, Erio Camporesi, IT 1792 (MFLU 16–2858, holotype; *ibid* BBH, isotype).

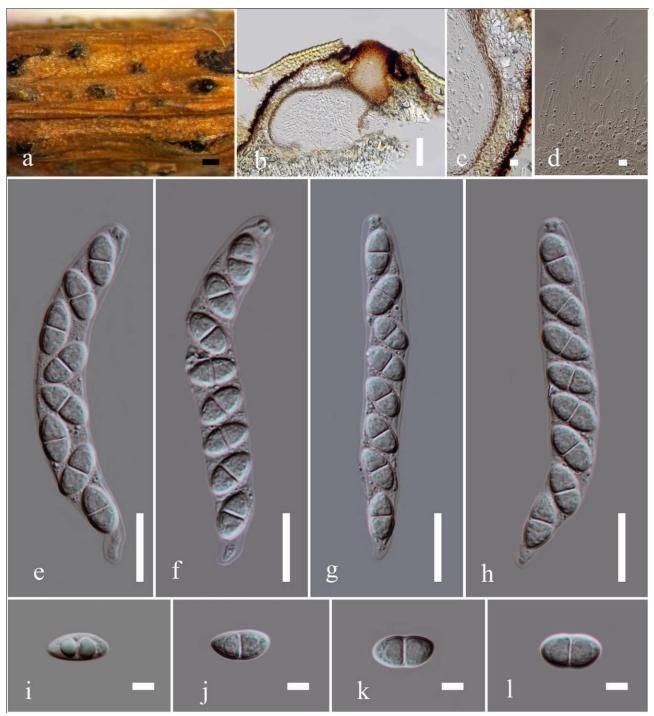
Notes – Sydowiella juncina and S. stellatifolii were collected on Juncus effusus L. and Brabejum stellatifolium L. respectively, and are possibly host-specific species. Sydowiella ambigua and S. azukiae form aggregated ascomata, while S. centaureii has solitary ascomata. Sydowiella centaureii differs from S. fenestrans and S. urticola in having large ascomata immersed within distinct stromatic tissues, surrounded by a layer of dark cells and small asci and ascospores (Maharachchikumbura et al. 2016, Index Fungorum 2017). However, we could not obtain a culture from this fungus and we extracted DNA directly from the ascomata.

#### Alborbis Senan. & K.D. Hyde, gen. nov.

Index Fungorum number – IF552717, Facesoffungi number – FoF 2835

Etymology – based on Latin words "alba" and "orbis", referring to the ascomata opening to the host surface through a whitish disk.

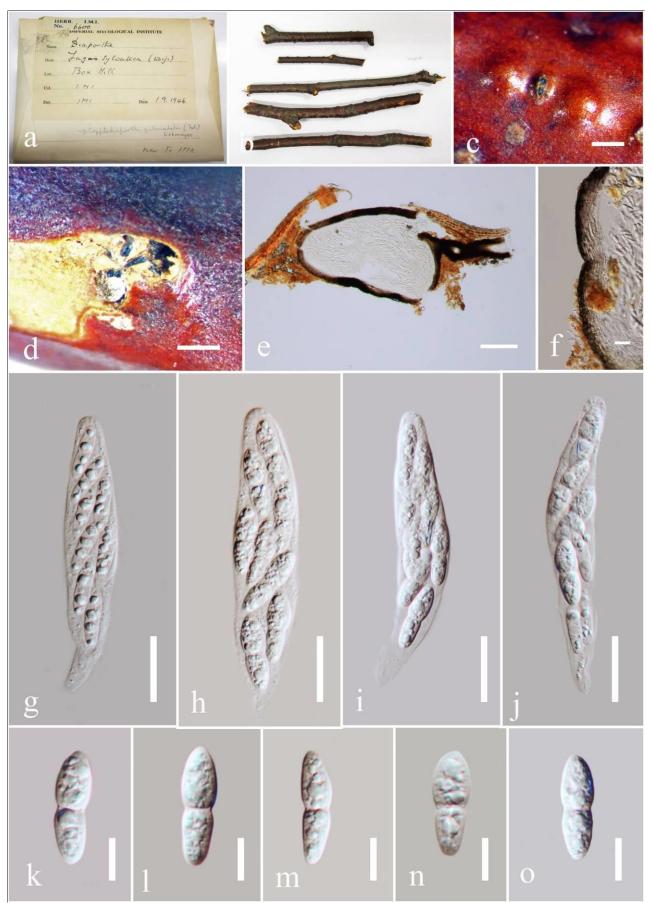
Saprobic on dead stems. Sexual morph – Ectostromatic disc appearing as pustulate swellings, small, whitish, usually containing several ascomata. Ascomata immersed, aggregated, globose to



**Figure 2** – *Sydowiella centaureii* (holotype) a Appearance of ascomata on substrate. b Vertical section of ascoma. c Peridium. d Paraphyses. e-h Asci. i-l Ascospores. Scale bars –  $a = 200 \mu m$ ,  $b = 100 \mu m$ ,  $c = 10 \mu m$ ,  $d = 5 \mu m$ , e-h =  $20 \mu m$ , i-l =  $5 \mu m$ .

subglobose, coriaceous, black, ostiolate. Ostiolar canal scarcely emergent, separate, opening to ectostromatic disc by a wide ostiole, internally covered by hyaline, filamentous periphyses. Peridium comprising thick-walled, dark brown cells of *textura angularis*. Asci 8-spored, unitunicate, cylindrical to clavate, apex rounded with J-, apical ring. Ascospores overlapping uniseriate to biseriate, fusiform, 1-septate, sometimes with apical and basal appendages. Asexual morph – Undetermined.

Type species – Alborbis galericulata (Tul. & C. Tul.) Senan. & K.D. Hyde



**Figure 3** – *Alborbis galericulata* (IMI 6600). a Herbarium packet. b Herbarium specimen. c Appearance of ascomata on host surface. d Horizontal section of ascomata through the ectostromatic disc. e Vertical cross section through the ascomata. f Peridium. g-j Asci. k-o Ascospores. Scale bars – c, d =  $500 \, \mu m$ , e =  $200 \, \mu m$ , f-j =  $20 \, \mu m$ , k-o =  $10 \, \mu m$ .

Alborbis galericulata (Tul. & C. Tul.) Senan. & K.D. Hyde, comb. nov.

Index Fungorum number – IF552718, Facesoffungi number – FoF 2836

Fig. 3

Basionym – Valsa galericulata Tul. & C. Tul., Select. fung. carpol. (Paris) 2: 203 (1863)

Holotype - NYBG 02929068

Saprobic on dead stems of *Fagus sylvatica* L. Sexual morph – Appearing on the surface as small, pustulate swellings, with a central stellate rupture of the periderm through a white, minute, elliptic to fusoid ectostromatic disk, usually containing 1–6 ascomata. Ascomata 600–950 µm high, 260–450 µm diameter, immersed, aggregated, globose to subglobose, coriaceous, black, ostiolate. Ostiolar canal 375–400 µm high, 80–90 µm wide ( $\bar{x}=390\times83$  µm, n = 20), scarcely emergent, separate, opening to host surface by wide ostiole, internally covered by hyaline, filamentous periphyses. Peridium 30–35 µm thick, comprising thick-walled, dark brown cells of *textura angularis*. Asci 90–110 × 17–20 µm ( $\bar{x}=100\times19$  µm, n = 20), 8-spored, unitunicate, cylindrical to clavate, apex rounded with J- apical ring, short-pedicellate. Ascospores 21–26 × 5–7 µm ( $\bar{x}=24\times6$  µm, n = 20), biseriate to triseriate, often 4-guttulate, ellipsoid to cylindrical, 1-septate, hyaline, slightly constricted at the septum lacking appendages. Asexual morph – Undetermined.

Material examined – UK, England, Surrey, Box Hill, on twigs of *Fagus sylvatica* L (Fagaceae), 1 September 1946, IMI 6600.

Notes – Cryptodiaporthe galericulata was introduced by Tulasne & Tulasne (1863) based on Valsa galericulata. Cryptodiaporthe galericulata has been reported as causing mild canker of Fagus species and saprobic on branches of Fagus sylvatica (Mejía et al. 2011). Wehmeyer (1933) mentioned that Cryptodiaporthe contains a heterogenous group of species. Barr (1978) accommodated Cryptodiaporthe galericulata in family Valsaceae (Tribe Valseae). However, the type species of Cryptodiaporthe is accommodated in Gnomoniaceae based on phylogeny. Mejía et al. (2011) noted that LSU based phylogeny of a fresh collection of C. galericulata (BPI 863767) from the USA, clustered in Sydowiellaceae rather than Gnomoniaceae. Kruys & Castlebury (2012) placed C. galericulata in Sydowiellaceae using ITS sequence data. Combined LSU, ITS, RPB2 and TEF sequence analysis in this study (Fig. 1) also places C. galericulata in Sydowiellaceae. Hence, we exclude C. galericulata from Cryptodiaporthe and introduce a new genus Alborbis to accommodate Cryptodiaporthe galericulata.

# Breviappendix Senan. & K.D. Hyde, gen. nov.

Index Fungorum number – IF552719, Facesoffungi number – FoF 2837

Etymology – based on Latin words "appendicibus" and "brevi" referring to the ascospores having short appendages.

Pathogenic on living stems forming cankers. Sexual morph — Ascomata solitary, scattered, erumpent to immersed, globose, coriaceous, brown, papillate, ostiolate. Papilla short, wide, comprising brown, elongated cells, internally covered by hyaline periphyses. Peridium comprising inner layer of thin-walled, hyaline, compressed cells of *textura angularis* and outer layer of thickwalled, brown, compressed cells of *textura angularis*. Hamathecium aparaphysate. Asci 8-spored, unitunicate, fusiform, sessile, apex blunt, with J- apical ring. Ascospores biseriate, fusiform to ellipsoid, 1-median septate, hyaline, ends with appendages. Asexual morph — Undetermined.

Type species – Breviappendix rubi (Rehm) Senan., Maharachch. & K.D. Hyde

#### Breviappendix rubi (Rehm) Senan., Maharachch. & K.D. Hyde, comb. nov.

Index Fungorum number – IF552720, Facesoffungi number – FoF 2838

Figs 4, 5

Synonymy – *Gnomonia tetraspora* var. *rubi* Rehm, Ascomyceten, fasc. 12: no. 596 (1880)

Gnomonia rubi (Rehm) G. Winter, Hedwigia 26(2): 62 (1885)

Epitype – MFLU 16–2861

Pathogenic on living stems of *Rubus ulmifolius* Schott. forming cankers. Sexual morph – Ascomata 660–760  $\mu$ m high, 520–535  $\mu$ m diameter ( $\bar{x} = 720 \times 530 \,\mu$ m, n = 15), solitary, scattered, erumpent to immersed, globose, coriaceous, brown, papillate, ostiolate. Papilla 380–410  $\mu$ m high, 130–220  $\mu$ m diameter ( $\bar{x} = 405 \times 210 \,\mu$ m, n = 15), comprising brown, elongate cells, internally

covered by hyaline periphyses. Peridium 20–35 µm wide ( $\bar{x} = 31$  µm, n = 15), comprising inner, thin-walled, hyaline, compressed cells of textura angularis and outer, thick-walled, brown, compressed cells of textura angularis. Hamathecium 9–11 µm wide at base ( $\bar{x} = 10$  µm, n = 15), 4-5 µm wide at the apex ( $\bar{x} = 4.8$  µm, n = 15), comprising thin-walled cellular, septate, paraphyses. Asci 35–45  $\times$  5–10 µm ( $\overline{x}$  = 41  $\times$  7.8 µm, n = 25), 4–(6)-spored, unitunicate, fusiform, sessile, apex blunt, with a J- apical ring. Ascospores  $10-15 \times 2.5-3.5 \, \mu \text{m}$  ( $\overline{x} = 13 \times 2.8 \, \mu \text{m}$ , n = 25), biseriate, ellipsoid, 1-median septate, slightly constricted at the septa, hyaline, smooth-walled, ends with small, pointed small, setose appendages 1.5-6 µm long. Asexual morph coelomycetous. Conidiomata on MEA sporodochial, developing in 14 days, with a golden-yellow, slimy, pycnidial, superficial, globoid conidial head 150–200  $\mu$ m wide/diameter ( $\bar{x} = 180 \,\mu$ m, n = 15). Conidiophores 7–11 µm high, 1–3 µm diameter ( $\bar{x} = 9 \times 1.6$  µm, n = 15), macronematous, cylindrical, short, septate, branched, hyaline, smooth, compactly packed. Conidiogenous cells 10-20  $\mu$ m high, 1–2.5  $\mu$ m diameter ( $\bar{x} = 14.8 \times 1.7 \mu$ m, n = 15), phialidic, discrete, vase-shaped, without collarette, hyaline, sometimes branched, aseptate, narrowing towards the apex. Conidia  $5.5-7 \times 1.5-2 \,\mu \text{m}$  ( $\bar{x} = 6.5 \times 1.8 \,\mu \text{m}$ , n = 15), lunar, widest in the centre, pointed at both ends, aseptate, smooth, hyaline, accumulated in a slimy, golden-yellow mass.

Material examined – ITALY, Province of Forlì-Cesena, Santa Sofia, near Corniolo, on dead branch of *Rubus ulmifolius* Schott (Rosaceae), 25 August 2014, Erio Camporesi, IT 2070 (MFLU 16–2861, epitype designated here), ex-epitype living culture, MFLUCC 16–1205, TBRC; *ibid* MFLU 16–2862, BBH).

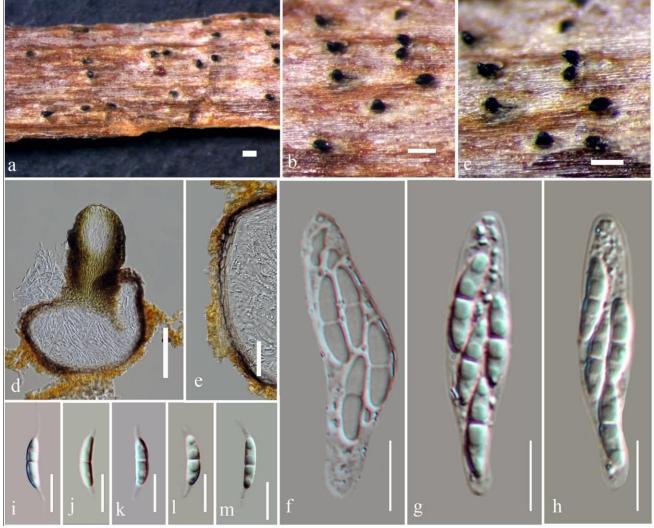
Culture characters – Colonies growing on MEA attaining 1 cm diameter within 7 days at 18 °C, rich in aerial mycelia, woolly, circular, flat, smooth, greyish white.

Notes – Gnomonia rubi was reported as the causal agent of severe stem cankers and dieback of various cultivated and wild Rubus species in both Europe and America (Ellis et al. 1984, Maas et al. 1989, Nordskog et al. 2003). Morphologically G. rubi and G. rostellata (Fr.) Wehm., have similar characters and some authors have synonymized G. rubi under G. rostellata (Monod 1983, Barr 1991). Barr (1978) reported G. rubi as the four-spored form of G. rostellata. Studies have placed Gnomonia rosae, G. rostellata and G. rubi out of Gnomoniaceae as they cluster together with Hapalocystis species (Morocko & Fatehi 2007). Kruys & Castlebury (2012) placed G. rubi in Sydowiellaceae based on ITS sequence data. In this study, we observed the same results and hence we introduced a new genus Breviappendix to accommodate G. rosae, G. rostellata and G. rubi and we proposed three new combinations as *Breviappendix rosae* (IF552798), *B. rostellata* (IF552799) and B. rubi. Breviappendix rubi is designated as the type species and we observed two specimens of Breviappendix rubi (= Gnomonia rubi) from IMI (IMI 252650, IMI 286320). Our fresh collection is morphologically identical to the prologue of Gnomonia rubi and the specimens we observed. However, we could not obtain the type specimen which is probably destroyed. The holotype was collected from Rubus fruticosus L (Rosaceae) in Switzerland and our fresh collection was collected from Rubus ulmifolius Schott in Italy. Considering the morphological similarity and close proximity of collections, we therefore designate our fresh collection as an epitype of Breviappendix rubi (sensu Ariyawansa et al. 2014) and both sexual and asexual morphs are illustrated herein.

*Breviappendix rosae* (Fuckel) Senan., Maharachch. & K.D. Hyde, *comb. nov*. Index Fungorum number – IF552798
Basionym – *Gnomonia rosae* Fuckel, Jb. nassau. Ver. Naturk. 23-24: 122 (1870) [1869-70]

*Breviappendix rostellata* (Fr.) Senan., Maharachch. & K.D. Hyde, *comb. nov.* Index Fungorum number – IF552799

Basionym – *Sphaeria rostellata* Fr., Observ. mycol. (Havniae) 1: 178 (1815)



**Figure 4** – *Breviappendix rubi* (MFLU 16–2861, epitype). a-c Appearance of ascomata on substrate. d Vertical section of ascoma. e Peridium. f-h Asci. i-m Ascospores. Scale bars – a-c =  $200 \, \mu m$ , d =  $100 \, \mu m$ , e =  $50 \, \mu m$ , f-m =  $10 \, \mu m$ .

#### Cainiella E. Müll., Sydowia 10(1-6): 120 (1957) [1956]

Facesoffungi number – FoF2839

Saprobic or parasitic on both living and dead leaves and petioles of temperate shrubs. Sexual morph – Ascomata scattered, solitary or in pairs, erumpent, globose to subglobose, coriaceous, black, ostiolate, papillate. Papilla wide, erumpent, internal wall covered by hyaline, filamentous periphyses. Peridium comprising a few layers of outer brown-walled cells of *textura globulosa* and inner layer of hyaline cells of *textura angularis*. Hamathecium comprising filamentous, hyaline, aseptate paraphyses. Asci few in number, 8-spored, unitunicate, cylindro-clavate, short pedicellate, apex wide, flat, with, J- apical ring. Ascospores overlapping uniseriate to biseriate, kidney-shaped or ellipsoidal, guttulate, initially hyaline, greenish or yellowish, becoming brown with age, 1-septate, with a thick and smooth epispore. Asexual morph – Undetermined.

Type species – Cainiella johansonii (Rehm) E. Müll.

#### Cainiella borealis M.E. Barr, Contr. Inst. bot. Univ. Montréal 73: 65 (1959)

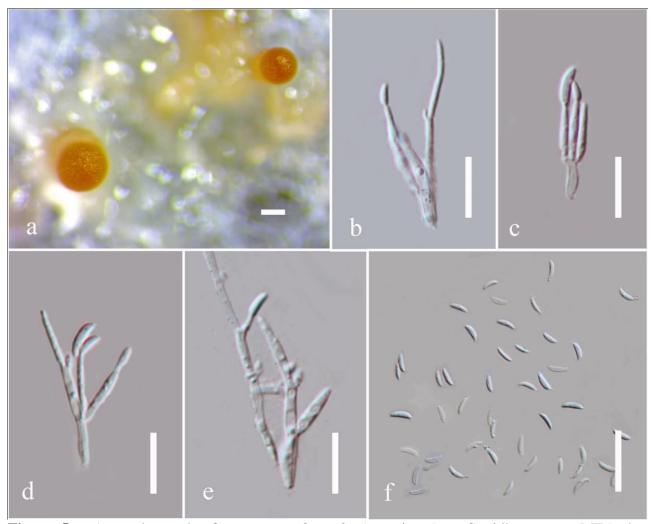
Facesoffungi number – FoF2840

Fig. 6

Saprobic or parasitic on both living and dead leaves and petioles of temperate shrubs. Sexual morph – Ascomata 775–1000  $\mu$ m high, 260–345  $\mu$ m diameter ( $\bar{x} = 846 \times 300 \, \mu$ m, n = 15), scattered, solitary, erumpent, globose to subglobose, coriaceous, black, ostiolate, papillate. Papilla 400–450  $\mu$ m high, 120–150  $\mu$ m diameter ( $\bar{x} = 415 \times 142 \, \mu$ m, n = 20), erumpent, internal wall

covered by hyaline, filamentous periphyses. Peridium 15–25  $\mu$ m wide ( $\bar{x}=21~\mu$ m, n = 20), comprising a few outer layers of brown-walled cells of *textura globulosa* and inner, hyaline, cells of *textura angularis*. Hamathecium comprising filamentous, hyaline, aseptate paraphyses. Asci 190–220 × 30–40  $\mu$ m ( $\bar{x}=204\times35~\mu$ m, n = 20), few in number, 8-spored, unitunicate, cylindro-clavate, short pedicellate, apex wide, flat with complex, J-, apical ring. Ascospores 30–35 × 17–20  $\mu$ m ( $\bar{x}=34\times19~\mu$ m, n = 20), biseriate, kidney-shaped, guttulate, hyaline, 1-septate, slightly constricted at the septum, with a thick epispore. Asexual morph – Undetermined.

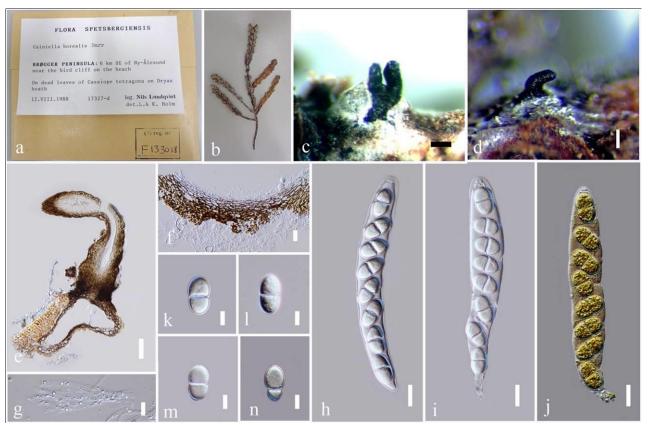
Material examined – NORWAY, Svalbard and Jan Mayen, Svalbard, Brøgger Peninsula: 6 km SE of Ny-Alesund near the bird cliff on the beach, on stems of *Cassiope tetragona* D. Don (Ericaceae), 12 August 1988, Nils Lundqvist, 17327-d, S-F133018.



**Figure 5** – Asexual morph of *Breviappendix rubi* (ex-epitype). a Conidiomata on MEA. b-e Conidiophores, conidiogenous cells and conidia. f Conidia. Scale bars –  $a = 100 \mu m$ , b-f =  $10 \mu m$ .

Notes – The genus *Cainiella* was introduced by Muller (1957) based on *Lizonia johansonii* Rehm. This genus is characterized by dark, immersed ascomata, with long-necks, deliquescing paraphyses, as well as asci with a J-, refractive ring and 1-septate ascospores. *Cainiella* comprises the type species *C. johansonii* (Rehm) E. Müll. and *C. borealis*. These taxa are associated with arctic dwarf shrubs and prostrate evergreen subshrubs. Initially, this genus was assigned to Sphaeriales (Muller 1956). However, Barr (1959) placed *Cainiella* in Xylariaceae when she introduced *C. borealis*. Subsequently *Cainiella* was placed in Amphisphaeriaceae (Muller & von Arx 1962), in Cainiaceae (Vasilyeva 1987, 1998) and in Hyponectriaceae (Kirk et al. 2008). Vasilyeva (1998) separated the two species, transferring *C. borealis* to *Sydowiella* (Sydowiellaceae) and *C. johansonii* to *Cainia* (Cainiaceae). However, Lumbsch & Huhndorf (2009) placed *Cainiella* in the family Sordariaceae.

Kruys & Castlebury (2012) reported that the taxonomic placement of *Cainiella* is in Sydowiellaceae based on LSU and ITS sequence data.



**Figure 6** – *Cainiella borealis* (S-F133018). a Herbarium packet. b Herbarium specimen. c, d Appearance of ascomata on substrate. e Cross section of ascoma. f Peridium. g Paraphyses. h, i Asci in water. j Ascus in Melzer's reagent. k-n Ascospores. Scale bars –  $c = 200 \, \mu m$ ,  $d = 200 \, \mu m$ , e =  $100 \, \mu m$ , f, h-j =  $20 \, \mu m$ , g, k-n =  $10 \, \mu m$ .

*Calosporella* J. Schröt., in Cohn, Krypt.-Fl. Schlesien (Breslau) 3.2(4): 442 (1897) [1908] Facesoffungi number – FoF2841

Saprobic on dead branches of deciduous trees. Sexual morph – Pseudostromata pustulate on surface, circular, containing up to 20 ascoma. Ectostromatic disc erumpent, conspicuous, white to light grey, becoming dark at maturity, fusoid to circular, possessing 1–20 distinctly protruding ostioles. Entostroma crumbly, brown. Ascomata immersed in stromata, aggregated, globose to subglobose, coriaceous, black, ostiolate, papillate. Papilla erumpent, cylindrical, black. Asci 8-spored, unitunicate, broadly fusoid, without or with inconspicuous pedicel, without a distinct apical ring. Ascospores bi- to triseriate, fusoid to oblong, 3–4-euseptate, hyaline, with tapering appendages at both ends. Asexual morph – Undetermined.

Type species – Calosporella innesii (Curr.) J. Schröt.

Calosporella innesii (Curr.) J. Schröt., in Cohn, Krypt.-Fl. Schlesien (Breslau) 3.2, 4: 442. 1897 (1908).

Basionym – Sphaeria innesii Curr., Trans. Linn. Soc. London 22: 281. 1858 ('1859').

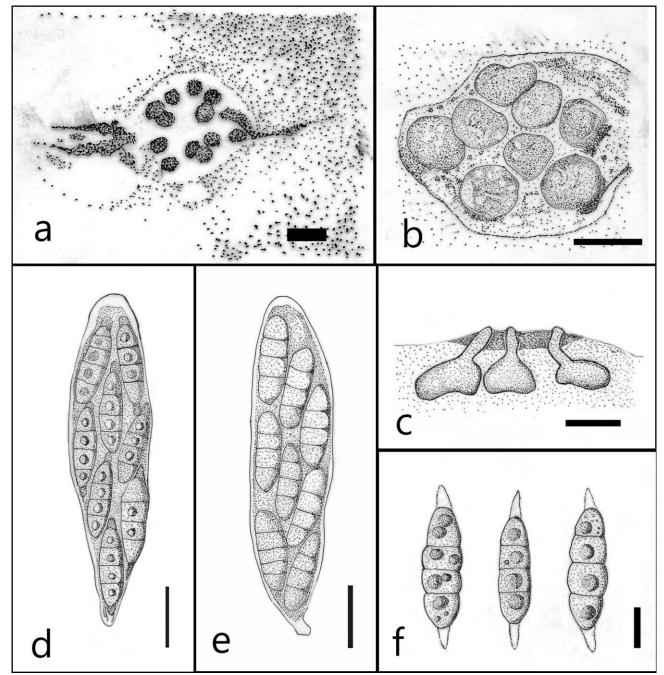
Facesoffungi number – FoF2842

Fig. 7

Epitype – WU 32161

Saprobic on dead branches of deciduous trees. Sexual morph – Pseudostromata 1–3 mm diameter, pustulate on surface, circular, containing up to 20 ascomata. Ectostromatic disc erumpent, conspicuous, white to light grey, becoming dark at maturity, fusoid to circular, possessing 1–20 distinctly protruding ostioles. Entostroma crumbly, brownish. Ascomata 400–600 µm diameter, immersed in stromata, aggregated, globose to subglobose, coriaceous, black, ostiolate, papillate.

Ostioles erumpent, cylindrical, black. Asci  $60-100 \times 16-22$  µm, 8-spored, unitunicate, broadly fusoid, without or with inconspicuous pedicel, lacking a distinct apical ring. Ascospores  $20-34 \times 6.5-10.5$  µm, bi- to triseriate, fusoid to oblong, 3-4-euseptate, not to slightly constricted at septa, hyaline, multi-guttulate, with rounded ends, hyaline tapering appendages at both ends, appendages 5-7 µm long, 3-4.5 µm wide at the base. Asexual morph – Undetermined (description based on Voglmayr & Jaklitsch 2014).



**Figure 7** – Calosporella innesii. (Redrawn from Voglmayr & Jaklitsch 2014). a Ascomata on substrate. b Horizontal cross section of ascoma. c Vertical cross section of ascoma. d, Matured ascus e. Over matured ascus. f Ascospores. Scale bars – a-c =  $500 \, \mu m$ , d, e =  $20 \, \mu m$ , f =  $10 \, \mu m$ .

Notes – Schröter (1897) erected the genus *Calosporella* as a replacement for *Calospora* Sacc. Saccardo (1883) listed 13 species under the genus without designating a generic type. However, the first two species, *C. platanoidis* (Pers.) Sacc. and *C. innesii* (Curr.) Sacc., are morphologically distinct (Saccardo 1883). Clements & Shear (1931) lectotypified *Calosporella* with *C. platanoidis*. However, Wehmeyer (1941) suggested the type specimen of its basionym, *Sphaeria platanoidis* 

Pers., is not congeneric with *C. platanoidis*, based on its morphology. Hence lectotypification of the genus *Calosporella*, with *C. platanoidis* could not be accepted. Therefore, Voglmayr & Jaklitsch (2014) typified *Calosporella* with *C. innesii* using a specimen collected from Austria associated with dead corticated branches of *Acer pseudoplatanus*. *Calosporella innesii* is characterized by hyaline, mostly 3-septate ascospores, with small apical appendages. It was transferred to *Prosthecium* by Wehmeyer (1941) based on phragmospores with apical appendages. However, in the prologue, Currey (1858) did not provide any details about the collection which was selected as the lectotype. Phylogenetic placement of *Calosporella* in Sydowiellaceae has been reported using ML analysis of the LSU sequence data from selected taxa in Diaporthales (Voglmayr & Jaklitsch 2014).

#### Chapeckia M.E. Barr, Mycol. Mem. 7: 164 (1978)

Facesoffungi number – FoF2843

Saprobic on branches of deciduous trees and shrubs. Sexual morph – Stromata pseudoparenchymatous, scattered on host surface, immersed, opening to the outside through cracks in the host epidermis, appearing as an aggregation of ostioles, rounded or elliptic, externally black, internally brown, composed of compact pseudoparenchymatous tissues, 3–9 ascomata in a stroma. Ascomata upright, deeply immersed in stromata. Peridium thin at the base, wider at base of neck. Hamathecium comprising cellular paraphyses. Asci 8-spored, unitunicate, oblong to ellipsoidal, short-pedicellate. Ascospores biseriate, oblong-ellipsoidal to oval, light to dark brown, 1-median septate. Asexual morph – Undetermined.

Type species – *Chapeckia nigrospora* (Peck)

#### Chapeckia nigrospora (Peck) M.E. Barr, Mycol. Mem. 7: 165 (1978)

Basionym – *Diatrype nigrospora* Peck, Ann. Rep. N.Y. St. Mus. nat. Hist. 33: 33 (1883) [1880]

Facesoffungi number – FoF2844 Holotype – NYS 2083 Fig. 8

Saprobic on branches of deciduous trees and shrubs. Sexual morph – Stromata 800–925  $\mu$ m wide ( $\bar{x}=900~\mu$ m, n = 10), pseudoparenchymatous, scattered on host surface, immersed, opening to outside through cracks in the host epidermis, appearing as an aggregation of ostioles, rounded or elliptic in shape, externally black, internally brown, composed of compact pseudoparenchymatous tissues, 3–4 or up to 9 ascomata in a stroma. Ascomata 360–455  $\mu$ m high, 160–225  $\mu$ m diameter ( $\bar{x}=408\times180~\mu$ m, n = 20), upright, deeply immersed in stromata, necks short and wide, flared at upper portion, globose. Peridium 10–13  $\mu$ m wide ( $\bar{x}=11~\mu$ m, n = 20), thin at the base, comprising 1–3 layers of brown-walled cells of *textura angularis* and base of neck comprising 6–8 layers of cells of *textura angularis*. Hamathecium comprising cellular paraphyses. Asci 40–55 × 10–15  $\mu$ m ( $\bar{x}=49\times12~\mu$ m, n = 20), 8-spored, unitunicate, oblong to ellipsoidal, apex slightly pointed, short-pedicellate. Ascospores 9.5–11 × 5–6  $\mu$ m ( $\bar{x}=10.7\times5.6~\mu$ m, n = 20), biseriate, oblong-ellipsoidal

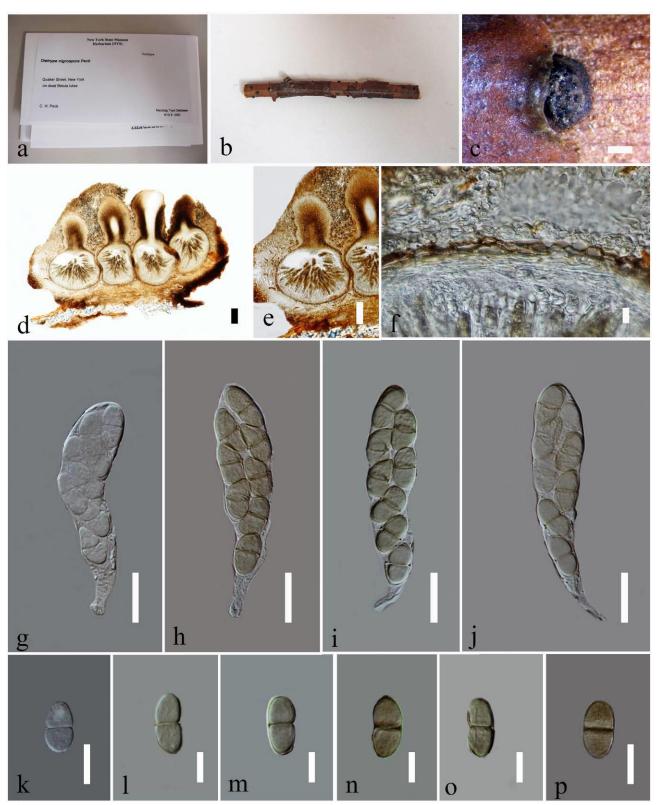
Material examined – USA, New York, Quaker Street, on bark of *Betula lutea*, C.H. Peck (NYS 2083, holotype).

dry, appendages narrow, short or elongate. Asexual morph – Undetermined.

to oval, hyaline when immature, light to dark brown at maturity, 1-median septate, slightly constricted at the septum, wall surrounded by narrow gel coating becoming irregularly rough when

Notes – *Chapeckia* was introduced by Barr (1978) based on *Diatrype nigrospora* and accommodated in Pseudovalsaceae. There are two species in the genus (Index Fungorum 2017) the type species and *C. ribesia* (Sacc. & Scalia) M.E. Barr.

Lumbsch & Huhndorf (2007) placed *Chapeckia* in Melanconidaceae while Rossman et al. (2007) placed it in Sydowiellaceae. Later, Lumbsch & Huhndorf (2010) also accommodated *Chapeckia* in Sydowiellaceae. De Silva et al. (2009) and Kruys & Castlebury (2012) analyzed sequence data for *Chapeckia* and agreed it belonged in Sydowiellaceae.



**Figure 8** – *Chapeckia nigrospora* (holotype). a, b Herbarium packet and specimen. c Appearance of stroma on host surface. d-e Cross section of ascomata in stroma. f Peridium. g-j Asci. k-p Ascospores. Scale bars –  $c = 500 \mu m$ , d,  $e = 100 \mu m$ ,  $f = 10 \mu m$ ,  $g-j = 20 \mu m$ ,  $k-p = 10 \mu m$ .

*Hapalocystis* Auersw. ex Fuckel, Fungi rhenani exsic., fasc. 6: no. 585 (1863) Facesoffungi number – FoF2845

Saprobic on woody bark. Sexual morph – Stromata forming a thin, weft of light brown hyphae around upper part of ascomata. Ascomata immersed, aggregated, subglobose to globose, coriaceous, black, ostiolate, papillate. Papilla short, black, cylindrical, periphysate. Peridium

comprising few layers of black, thick-walled, cells of *textura angularis*. Hamathecium comprising a few, septate, hyaline paraphyses attached to the base, longer than asci. Asci 4 or 8-spored, unitunicate, clavate to fusoid, with a short pedicel, and an inconspicuous flat, J-, refractive ring. Ascospores inequilaterally ellipsoidal, with broadly rounded ends, hyaline to dark brown, distoseptate, with short or long, hyaline strap-like appendages at both rounded ends. Asexual morph: Coelomycetous. Conidiomata pycnidia, superficial, aggregated in groups of 3–5, globose, orange to brown. Conidiomatal wall small, thick-walled, orange, comprising cells of *textura angularis*. Conidiophores annelidic, branched, hyaline, short, few conidiogenous cells arising from a single conidiophore, attached to conidiomatal wall. Conidiogenous cells cylindrical, hyaline, bottle-shaped, septate, ends pointed, phialidic. Conidia ellipsoid or cylindroid, unicellular or multiseptate, hyaline or brown.

Type species – *Hapalocystis berkeleyi* Auersw. ex Fuckel, Fungi rhenani exsic., fasc. 6: no. 585 (1863)

Hapalocystis berkeleyi Auersw. ex Fuckel, Fungi rhenani exsic., fasc. 6: no. 585 (1863)

Facesoffungi number – FoF2846

Fig. 9

Isotype – K(M) 195743

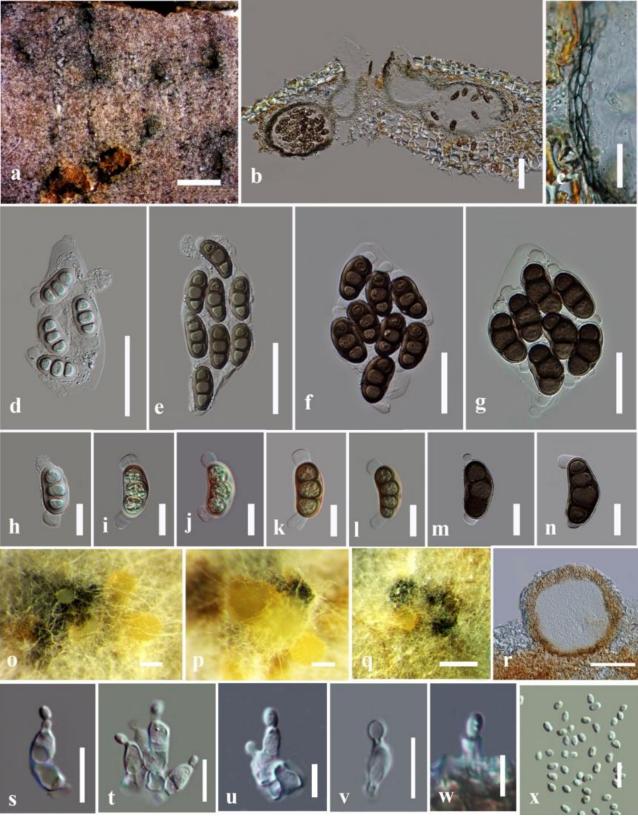
Saprobic on woody bark. Sexual morph – visible through dark cracks on the host surface. Stromata forming a thin, weft of light brown hyphae around upper part of ascomata. Ascomata 90–185 µm high, 206–302 µm diameter ( $\bar{x} = 150 \times 245 \text{ µm}$ , n = 15), immersed, aggregated, subglobose to globose, coriaceous, black, ostiolate. Ostiole 130–195 µm high, 90–113 µm wide ( $\bar{x} = 145 \times 105$  $\mu$ m, n = 10), short papillate, black, cylindrical, periphysate with hyaline, filamentous periphyses. Peridium 4–16 µm wide ( $\bar{x} = 10$  µm, n = 15), comprising 3–5-layers of black, thick-walled cells of textura angularis. Paraphyses 3–7 µm wide ( $\bar{x} = 6$  µm, n = 15), few, septate, hyaline, attached to the base, longer than asci. Asci 112–127  $\times$  46–80  $\mu$ m ( $\overline{x}$  = 121  $\times$  66  $\mu$ m, n = 20), 4–or 8-spored, unitunicate, clavate to fusoid, with a short pedicel, with an inconspicuous flat, refractive ring at the lower end of the thickened apical wall, apex narrow and blunted. Ascospores 34-42 × 12-18 um  $(\bar{x} = 37 \times 17 \mu m)$ , 2–5-seriate, inequilaterally ellipsoidal, with broadly rounded ends, initially hyaline, dark brown at maturity, (1-)-2-(-3)-septate, with cells of equal length, thick and smoothwalled, with short, hyaline, strap-like appendages situated at both rounded ends, having the same width as the ascospores. Asexual morph – Coelomycetous. Conidiomata 165–195 µm diameter ( $\bar{x}$ = 182  $\mu$ m, n = 15), pycnidial, superficial, aggregated, 3–5 in a group, globose, orange to brown. Conidiomata wall 7–13 µm wide ( $\bar{x} = 10 \mu m$ , n = 10), comprising 5–10 layers of small, thickwalled, orange cells of textura angularis. Conidiophores 1.5–2 µm high, 1–1.5 µm wide ( $\bar{x} = 2 \times$ 1.5  $\mu$ m, n = 10), branched, hyaline, short, few conidiogenous cells arising from a single conidiophore, attached to conidiomatal wall. Conidiogenous cells 10–14 µm high, 5–7 µm wide, cylindrical, hyaline, bottle-shaped, septate, pointed at ends, phialidic. Conidia 0.5–1.5 μm diameter  $(\overline{x} = 1 \text{ µm}, n = 20)$  ellipsoid, 1-celled, hyaline, smooth-walled.

Material examined – ITALY, Province of Forlì-Cesena [FC], Modigliana, Montebello, on branch of *Platanus hybrida* L. (Platanaceae), 14 April 2013, Erio Camporesi, IT 1187 (MFLU 14–0798).

Notes – *Hapalocystis* was established to accommodate two species, *H. berkeleyi* and *H. bicaudata* Fuckel (Fuckel 1863). Clements & Shear (1931) designated *H. berkeleyi* as the type species. Later this genus was synonymized under *Pseudovalsa* (Winter 1887, Ellis & Everhart 1892) and *Prosthecium* (Wehmeyer 1941).

Barr (1978) however excluded *Hapalocystis* from *Prosthecium* based on morphology and placed the genus in Melanconidaceae (Barr 1978, Lumbsch & Huhndorf 2007). Rossman et al. (2007) placed *Hapalocystis* in Sydowiellaceae based on its morphology. *Hapalocystis* now comprises *H. bicaudata H. corni* (Wehm.) M.E. Barr, *H. corticalis* (Schwein.) M.E. Barr, *H. mirabilis* Sorokīn, *H. occidentalis* Jaklitsch & Voglmayr, *H. ulmi* (Wehm.) M.E. Barr, and *H. berkeleyi* (*H. berkeleyi* var. *berkeleyi* and *H. berkeleyi* var. *kickxii* (Westend.) M.E. Barr and *H. vexans* Speg. (Index Fungorum 2017). The asexual morph of *Hapalocystis* was recorded as

*Stilbospora*, *Hendersonia* or *Dothiorella* (Jaklitsch & Voglmayr 2004). Glawe (1985) and Liu et al. (2015) reported a *Phoma*-like asexual morph for *H. berkeleyi* from culture.



**Figure 9** – *Hapalocystis berkeleyi*. a Appearance of ascomata on substrate. b Vertical cross section of ascomata. c Peridium. d–g Asci. h–n Ascospores. o–q Conidiomata. r Vertical cross section of conidioma. s–w Conidiophores, conidiogenous cells and conidia. x Conidia. Scale bars – a=1 mm, b=100 μm, c=20 μm, d-g=50 μm, h-n=20 μm, o-q=500 μm, g=100 μm, s-w=10 μm, x=5 μm.

Italiomyces Senan., Camporesi & K.D. Hyde, gen. nov.

Index Fungorum number – IF552721, Facesoffungi number – FoF 2847

Etymology – Based on the country where the fungus was collected.

Saprobic on dead stem of *Centaurea* L. Sexual morph – Ascomata solitary to rarely aggregated, scattered, immersed, globose to subglobose, black coriaceous, papillate, ostiolate. Papilla short, wide, emerging through cracks in the substrate, comprising small, thick-walled, cells of *textura angularis*, internally covered by hyaline, periphyses. Peridium comprising thick-walled, compressed, dark brown cells of *textura angularis*. Asci 8-spored, unitunicate, fusiform, short-pedicellate, apex blunt with bi-lobed apical ring. Ascospores overlapping uniseriate to biseriate, fusiform to narrowly oval, 1-septate, large, hyaline, with appendages. Asexual morph – Undetermined.

Type species – Italiomyces centaureii Senan., Camporesi & K.D. Hyde

Italiomyces centaureii Senan., Camporesi & K.D. Hyde, sp. nov.

Index Fungorum number – IF552722, Facesoffungi number – FoF 2848 Fig. 10

Etymology – Based on the host genus.

Holotype – MFLU 16–2866

Saprobic on dead stems of *Centaurea* L. Sexual morph – Ascomata 360–380 µm high, 405–415 µm diameter ( $\overline{x} = 375 \times 411$  µm, n = 15), solitary to rarely aggregated, scattered, immersed, globose to subglobose, black, coriaceous, papillate, ostiolate. Papilla 140–165 µm high, 150–185 µm diameter ( $\overline{x} = 160 \times 175$  µm, n = 15), emerging through cracks in the substrate, comprising small, thick-walled cells of *textura angularis*, internally covered by hyaline, periphyses. Peridium 15–25 µm wide ( $\overline{x} = 20$  µm, n = 15), comprising thick-walled, compressed, dark brown cells of *textura angularis*. Asci 90–110 × 15–20 µm ( $\overline{x} = 105 \times 17$  µm, n = 25), 8-spored, unitunicate, fusiform, short-pedicellate, apex blunt with bi-lobed apical ring. Ascospores 20–30 × 6.5–8.5 µm ( $\overline{x} = 26 \times 7.7$  µm, n = 25), overlapping uniseriate to biseriate, fusiform to narrowly oval, 1-septate, biguttulate, hyaline, with short, spine-like appendages at both ends. Asexual morph – Undetermined.

Culture characters – Colonies growing on MEA attaining 1 cm within 7 days when incubated at 18° C, with sparse aerial mycelia, forming minute mycelia tufts, powdery, flat, circular, smooth, white to off-white.

Material examined – ITALY, Province of Forlì-Cesena, Fiumicello di Premilcuore, on dead stem of *Centaurea* sp. (Asteraceae), 2 April 2014, Erio Camporesi, IT 1976 (MFLU 16–2866, holotype), ex-type living cultures MFLUCC 14–0849, *ibid* (BBH isotype), ex-type living culture, TBRC.

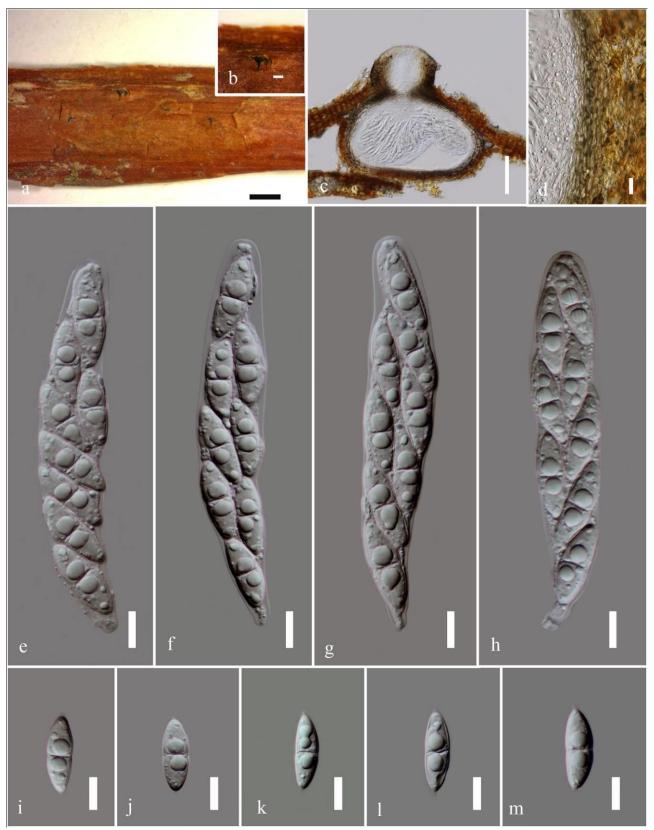
Notes – *Italiomyces* is distinct from other genera in Sydowiellaceae in its bi-guttulate or rarely 4-guttulate ascospores with minute, spine-like apical appendages. The guttule arrangement is unique for this genus and comprising two guttules located very close to each septum and smaller guttules at the ends of the spores. Furthermore, phylogenetically *Italiomyces* is distinct (Fig. 1) and hence we introduce the new genus.

Lambro Racib., Parasit. Alg. Pilze Java's (Jakarta) 2: 13 (1900)

= *Stegophora* Syd. & P. Syd., Annls mycol. 14(5): 364 (1916)

Index Fungorum number – IF552801 Facesoffungi number – FoF 2849

Pathogenic forming spots on living leaves. Sexual morph – Stromata pseudoparenchymatous to prosenchymatous, well-developed, black, with 3–12 ascomata in a stroma, forming a dark outer crust around the hyaline interior. Ascomata solitary, scattered, vertically or obliquely immersed within leaf tissue, globose to subglobose, coriaceous, black, with short neck and effuse clypeus-like hypostroma. Peridium membranaceous, outer layer comprising brown, thick-walled cells of *textura angularis* and inner layer of hyaline, thin-walled cells of *textura angularis*. Necks central or asymmetrically located, slightly protruding from either upper or lower leaf surface as blackish papillae, broader near the tip, internally covered by hyaline periphyses, wall comprising plectenchymatous, vertically elongate cells. Hamathecium aparaphysate. Asci 8-spored,



**Figure 10** – *Italiomyces centaureii* (holotype). a, b Herbarium specimen. c Vertical section of ascoma. d Peridium. e-h Asci. i-m Ascospores. Scale bars: a=1 mm, b=200  $\mu$ m, c=100  $\mu$ m, d-m =10  $\mu$ m.

unitunicate, clavate, tapering towards the base, apex thickened with J- apical ring. Ascospores overlapping uniseriate, apiosporous, elliptic to ovoid, tapering towards the base, hyaline. Asexual morph — Coelomycetous, Cylindrosporella-like. Appearing as greyish-brown to brown, circular or

irregularly-shaped. Conidiomata acervuli, scattered, solitary, erumpent. Conidiophores simple, short or obscure. Conidiogenous cells phialidic. Conidia hyaline, small, unicellular, filiform, fusiform, clavate or fusoid.

Type species – *Lambro insignis* Racib., Parasit. Alg. Pilze Java's (Jakarta) 2: 10 (1900)

*Lambro oharana* (Y. Nisik. & H. Matsumoto) E. Müll., in Müller & von Arx, Beitr. Kryptfl. Schweiz 11(no. 2): 778 (1962)

Facesoffungi number – FoF 2850

Fig. 11

Parasitic forming yellowish spots on living leaves, without distinct marginal zone, aggregated in a small black phyllachoroid scab, which is the cover of the conidial layer. Sexual morph – Stromata hypostromatic, pseudoparenchymatous, spread beneath ascomata, surrounded by clypeus in epidermal layer. Ascomata 370–480  $\mu$ m diameter, entirely immersed within leaf tissue, usually located as pairs, brown to light brown, globose to subglobose, with central or asymmetrically located short neck. Necks 120–180  $\mu$ m high, 100–110  $\mu$ m wide, slightly elevated from the upper leaf surface, parallel between paired ascomata, broader near the apex, with hyaline periphyses. Peridium plectenchymatous, light brown. Asci 43–56 × 7.5–10  $\mu$ m ( $\bar{x}$  = 48.8 × 8.8  $\mu$ m), 8-spored, clavate, tapering toward the base, thin-walled, with slightly visible apical ring at the thickened apex, irregularly loose in ascomata. Ascospores 12.5–15 × 5–6.5  $\mu$ m ( $\bar{x}$  = 13.4 × 5.5  $\mu$ m), irregularly biseriate, apiosporous, elliptic to ovoid, hyaline. Asexual morph – Coelomycetous. Conidiomata 0.45–0.5 mm diameter, acervuli, subepidermal, initially with thick, black cover, flat, round to elliptic. Conidiophores obscure. Conidiogenous cells phialidic. Conidia 4.5–8 × 1.5–2.5  $\mu$ m ( $\bar{x}$  = 5.4 × 1.5  $\mu$ m), clavate or oblong-ovoid, straight or curved, hyaline, unicellular (description is based on Kobayashi 1970).

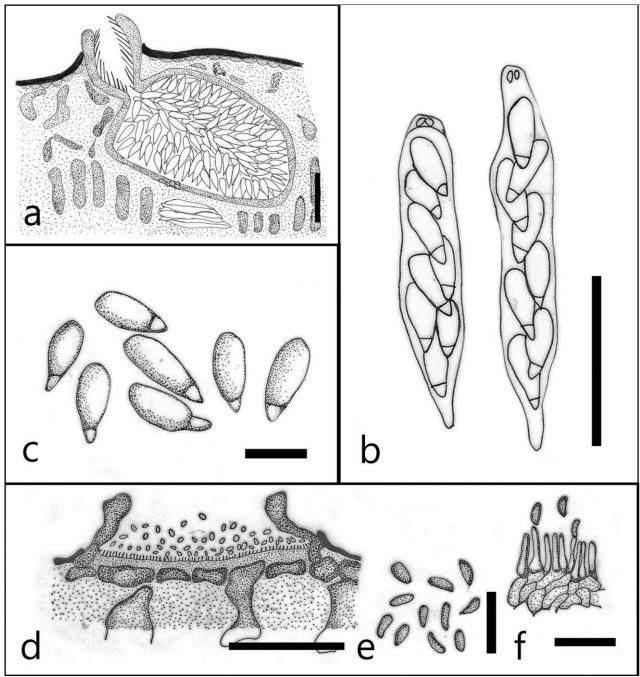
Notes – The genus *Lambro* is typified by *L. insignis*, which produces necrotic spots in leaves of *Sterculia subpeltata* Blume (Sterculiaceae) (Raciborski 1900). Müller & von Arx (1962) described and illustrated this genus and reported the morphological similarity with *Stegophora* and *Metadothella* Henn. species. *Lambro insignis* has a well-developed pseudoparenchymatous stromata completely enveloping 3–12 ascomata, and a dark outer crust around the hyaline interior region. However, the type species of *Stegophora*, *S. ulmea* has only subepidermal hypostroma (Müller & von Arx 1962). *Stegophora oharana* forms both a clypeus and hypostroma, which is limited to the epidermal layer of leaves. Sydow (1936) described *S. aenula* Syd. on *Ulmus davidiana* collected in China. Morphologically and biologically this species is identical to *S. oharana*. *Stegophora oharana* was transferred to *Lambro* based on its morphological similarity. Monod (1983) examined the type species, *L. insignis* and suggested that this species is related to *Stegophora*, thus it may belong in Sydowiellaceae. Hence, we synonymize *Stegophora* under *Lambro* giving priority to the older generic name *Lambro* and ignoring the small morphological differences between stromata. The asexual morph of *Lambro* is reported as a Cylindrosporella-like species.

# Paragnomonia Senan. & K.D. Hyde, gen. nov.

Index Fungorum number – IF552723, Facesoffungi number – FoF 2851

Etymology – In reference to the similarity to *Gnomonia*.

Pathogenic on plants. Sexual morph – Ascomata immersed, on roots and petioles, only neck visible when associated with host, solitary, globose to subglobose, black, coriaceous, papillate, ostiolate. Papilla central, straight to curved. Peridium comprising outer layer of large, brown cells of *textura angularis* and inner layer of compressed, hyaline cells of *textura angularis*. Asci fusiform to obovoid, with short pedicel and J- apical ring. Ascospores obliquely uniseriate, fusiform, straight to curved, 1-median-septate, not constricted at septum, with 4–6 guttules in each cell, sometimes more than 6-guttulate. Appendages apical and basal, hyaline. Asexual morph – Conidiomata tiny, pycnidia, solitary, immersed to erumpent, black. Conidiophores reduced to conidiogenous cells. Conidiogenous cells elongate, conical, unbranched, hyaline. Conidia oval to ellipsoid, aseptate, 2-guttulate, hyaline (description based on Alexopoulos & Cation 1952).



**Figure 11** – *Lambro oharana* (drawing based on Kobayashi 1970) a Vertical section of ascomata. b Asci. c Ascospores. d Vertical section of conidiomata. e Conidia. f Conidiogenous cells attached to conidia. Scale bars – a,  $d = 100 \, \mu m$ . b, c, e,  $f = 10 \, \mu m$ .

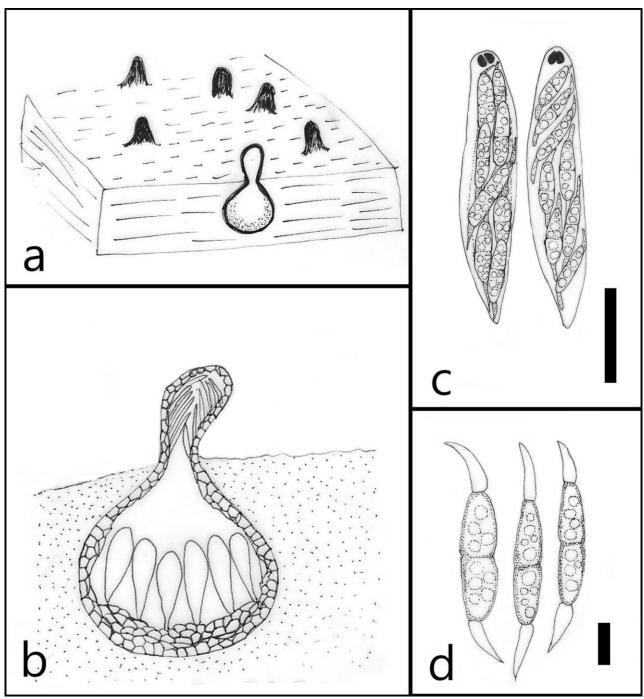
Paragnomonia fragariae (Kleb.) Senan. & K.D. Hyde, comb. nov.

Index Fungorum number – IF552724, Facesoffungi number – FoF 2852

Figs 12, 13

= *Gnomonia fragariae* var. *fragariae* Kleb., Haupt- und Nebenfruchtformen der Ascomyzeten (Leipzig) 1: 285 (1918)

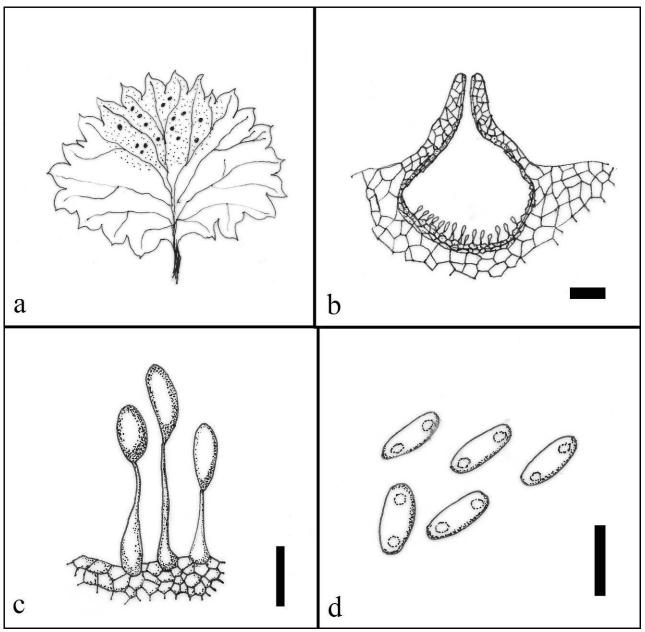
Pathogenic on plants. Sexual morph – Ascomata immersed on roots and petioles, papilla only visible on host, solitary, globose to subglobose, black, coriaceous, papillate, ostiolate. Necks central, straight to curved. Peridium comprising outer layer of large, brown cells of *textura angularis* and inner layer of compressed, hyaline cells of *textura angularis*. Asci  $25-35\times6-7~\mu m$ , fusiform to obovoid, with short pedicel and J- apical ring. Ascospores  $8-11\times2-4~\mu m$ , obliquely uniseriate, fusiform, straight to curved, 1-median septate, not constricted at the septum, with 4-6



**Figure 12** – *Paragnomonia fragariae* (drawing based on Morocko & Fatehi 2007). a Ascospores on substrate. b Vertical cross section of ascomata. c Asci. d Ascospores. Scale bars: c,  $d = 10 \mu m$ .

guttules, sometimes more than 6-guttulate. Appendages apical and basal, hyaline, elongate-conical. Asexual morph — Conidiomata tiny, pycnidial, solitary, immersed to erumpent, black. Conidiophores reduced to conidiogenous cells. Conidiogenous cells elongated, conical, unbranched, hyaline. Conidia oval to ellipsoid, aseptate, bi-guttulate, hyaline (description based on Alexopoulos & Cation 1952).

Notes – Paragnomonia fragariae is a poorly studied taxon, which was described by Klebahn (1918) from dead petioles and leaves of strawberry and causes strawberry decline in Germany. Klebahn (1918) considered this fungus as a saprobe of dead tissues of strawberry. Later, P. fragariae was recorded on dead tissues of pink barren strawberry (Potentilla micrantha), silver weed or goose grass (P. anserina), rock cinquefoil (P. rupestris), woodland strawberry (Fragaria vesca) and cultivated strawberry (Fragaria xananassa) collected in several localities of Switzerland



**Figure 13** – *Paragnomonia fragariae* (drawing based on Alexopoulos & Cation 1952) a Conidiomata on host leaf. b Cross section of conidioma. c Conidiophore, conidiogenous cells attached to conidia. d Conidia. Scale bars –  $b = 100 \, \mu m$ , c-d =  $5 \, \mu m$ .

(Bolay 1972). A severe strawberry petiole blight caused by *Gnomonia comari* was reported by van Adriechem and Bosher (1958) and is very similar to petiole blight caused by *P. fragariae*. However, *Gnomonia comari* did not cause any disease symptoms on roots, unlike *P. fragariae*. Morocko (2006) described the ascomata, asci and ascospores from cultures grown on artificial media and naturally infected plant tissues, with similar characteristics to *P. fragariae* (Klebahn 1918) and herbarium specimens collected by Bolay (1972) and Monod (1983). Morocko (2006) noted a great variation in length of the ascospore appendages.

Farr et al. (1989) and Maas (1998) suggested *P. fragariae* was a synonym of *Gnomonia comari* and Sogonov et al. (2008) placed *G. comari* in the newly introduced genus *Gnomoniopsis*. *Gnomoniopsis comari* has smaller asci and ascospores with a submedian septum and lacks appendages (Barr 1978). Morphologically *Gnomoniopsis comari* shows similarity to only *Gnomonia fragariae* var. *fructicola* (G. Arnaud) Fallahyan and not to *Gnomonia fragariae* var. *fragariae* Kleb. Morocko (2006) analyzed LSU and ITS sequence data of *Gnomonia* species and members of Diaporthales and showed that *G. fragariae* var. *fragariae* is phylogenetically

segregated from the generic type *G. gnomon*, and most other members of Gnomoniaceae. Morocko (2006) also suggested that *G. fragariae* f. *fragariae*, *G. rubi*, *G. rosae* and *Hapalocystis* species represents a genetically distinct group, possibly a new family, within Diaporthales.

Combined gene analyses of LSU, ITS, RPB2 and TEF sequence data in this study confirm the phylogenetic placement of *Gnomonia fragariae* var. *fragariae* in Sydowiellaceae. Hence, we exclude *Gnomonia fragariae* var. *fragariae* from *Gnomoniopsis comari* and introduce a new genus, *Paragnomonia* to accommodate *Gnomonia fragariae* var. *fragariae*.

# Ranulospora Senan., Camporesi & K.D. Hyde, gen. nov.

Index Fungorum number – IF552725, Facesoffungi number – FoF 2853

Etymology – Name referring to the tadpole-like shape of ascospores.

Saprobic on dead branches of deciduous plants. Sexual morph – Appearing as clusters of black, ostiolate necks which are erumpent. Pseudostromata comprising sparse, brown, fungal hyphae spreading around the ascomata. Ascomata immersed, aggregated, globose to subglobose, coriaceous, dark brown to black, with 6–8 aggregated ascomata, with long ostiolar necks. Peridium composed of 4–6 outer layers of thick-walled, brown cells of *textura angularis* and inner, hyaline, thick-walled cells of *textura angularis*. Hamathecium comprising few, septate, thin-walled paraphyses, which are longer than the asci. Asci 8-spored, unitunicate, cylindrical to fusiform, short pedicellate, apex rounded, with J- apical ring. Ascospores uniseriate or biseriate, fusiform to ellipsoid, with long basal appendage. Asexual morph – Undetermined.

Type species – Ranulospora alnii Senan., Camporesi & K.D. Hyde

# Ranulospora alnii Senan., Camporesi & K.D. Hyde, sp. nov.

Index Fungorum number – IF552726, Facesoffungi number – FoF 2854 Fig. 14

Etymology – Named after the host genus.

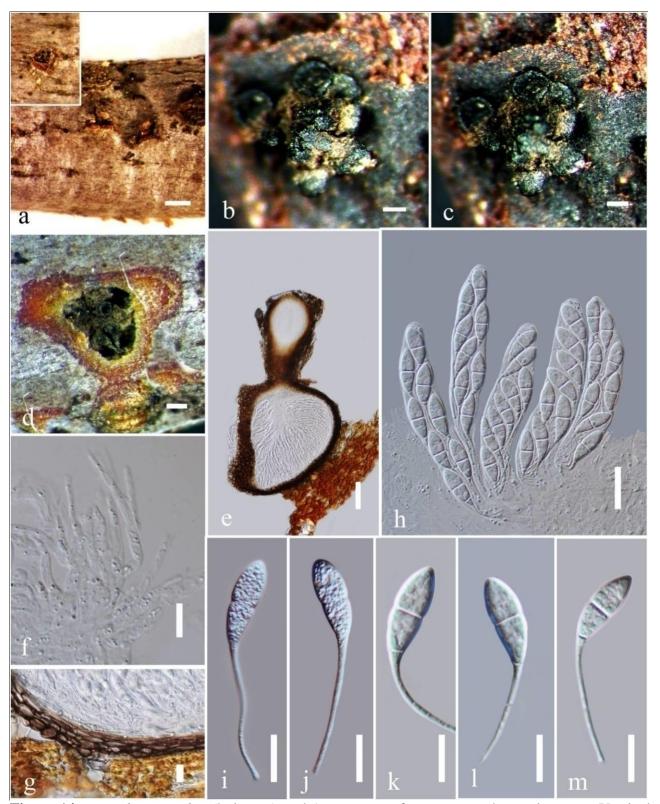
Holotype – MFLU 16–2868

Saprobic on dead branches of *Alnus incana* (L.) Moench. Sexual morph – Appearing as clusters of black, ostioles emerging through the cracked bark. Pseudostromata comprising sparse, brown, fungal hyphae spreading around ascomata. Ascomata 480–765 µm high, 280–345 µm diameter ( $\bar{x}=640\times315$  µm, n = 20), immersed, aggregated, globose to subglobose, coriaceous, dark brown to black, with 6–8 ascomata in an aggregation, papillate, with long ostiolar necks. Papilla 320–380 µm high, 160–260 µm wide ( $\bar{x}=349\times207$  µm, n = 20). Peridium 13–23 µm wide ( $\bar{x}=16.8$  µm, n = 20) composed of 4–6 outer layers of thick-walled, brown cells of *textura angularis* and inner, hyaline, thick-walled cells of *textura angularis*. Hamathecium comprising few, septate, hyaline, thin-walled paraphyses, longer than asci. Asci 85–120 × 16–22 µm ( $\bar{x}=101.7\times18.5$  µm, n = 20), 8-spored, unitunicate, cylindrical to fusiform, short pedicellate, apex rounded, with J- apical ring. Ascospores 45–55 × 7–8 µm ( $\bar{x}=48.6\times7.3$  µm, n = 20), uniseriate or biseriate, fusiform to ellipsoid, with long basal appendage, with 1 median and 1 basal septum, slightly constricted at the septa, smooth-walled, apical cell slightly wider than basal cell, appendage long filiform, straight or curved, with slight apical swelling, germ tubes emerging at the median septa and ascospores becoming olivaceous at germination. Asexual morph – Undetermined.

Culture characters – Colonies growing on MEA becoming 2 cm within 7 days, circular, flat to umbonate, margin filiform, dark green, tightly attached to the substrate, forming erect mycelium.

Material examined – ITALY, Province of Trento, Mezzana, Marilleva 900, on dead branch of *Alnus incana* (L.) Moench (Betulaceae), 15 August 2013, Erio Camporesi, IT 1415 (MFLU 16–2868 holotype; *ibid* BBH isotype); ex-type living cultures MFLUCC 13–0793, TBRC.

Notes – *Ranulospora alnii* was collected from bark of grey alder. This taxon appears to be a saprobe and several ascomata were grouped together with the necks combined by brown, fungal hyphae. The ascospore morphology is unique and we have not observed any taxon like this within Sydowiellaceae, although such appendages occur in other families (such as *Ceriospora* in Annulatascaceae (Campbell et al. 2013)). Phylogenetically this taxon forms a well-supported clade, which is sister to *Cainiella* and *Alborbis* (Fig. 1).

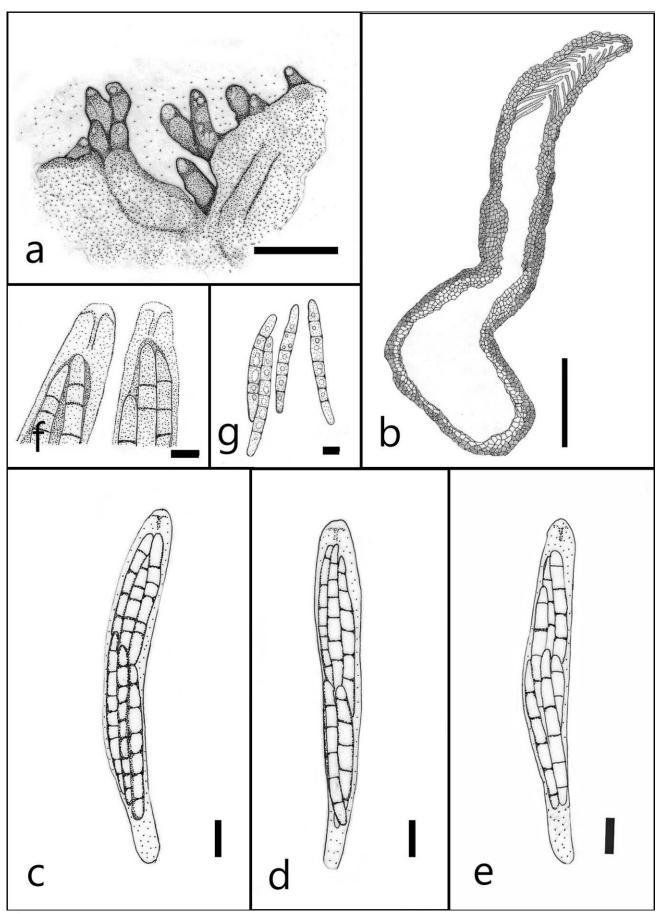


**Figure 14** – *Ranulospora alnii* (holotype). a-d Appearance of ascomata on host substrate. e Vertical section of ascoma. f Paraphyses. g Peridium. h Asci. i-m Ascospores with basal appendage. Scale bars –  $a = 500 \, \mu m$ , b-d =  $200 \, \mu m$ , d, e =  $100 \, \mu m$ , g, h =  $20 \, \mu m$ , f, i-m =  $10 \, \mu m$ .

Rossmania Lar.N. Vassiljeva, Mycoscience 42(4): 401 (2001)

Facesoffungi number – FoF 2855

Saprobic on dead woody bark. Sexual morph – Ascomata immersed under the bark, in clusters with convergent, long, stout erumpent necks, without an ectostromatic disc, globose, coriaceous, black. Necks wide, opening through the ostiole, internally covered by hyaline periphyses, comprising



**Figure 15** – *Rossmania ukurunduensis* (redrawn from Vasilyeva 2001). a Appearance of ascomata on substrate. b Vertical section of ascoma. c-e Asci. f Apical ring. g Ascospores. Scale bars – a=1 mm,  $b=200 \, \mu m$ , c-e =  $20 \, \mu m$ , f, g =  $10 \, \mu m$ .

brown cells of *textura porrecta*. Peridium thick, comprising outer, thick-walled, brown cells of *textura angularis* and inner thick-walled, hyaline cells of *textura angularis* at the base. Hamathecium aparaphysate. Asci 8-spored, unitunicate, clavate to cylindrical, short pedicellate, apex rounded, lacking an apical ring. Ascospores 4–5-seriate to fasciculate, long fusoid to cylindrical, slightly curved, 0–11-septate, hyaline to light yellow. Asexual morph – Undetermined.

Type species – Rossmania ukurunduensis Lar.N. Vassiljeva, Mycoscience 42(4): 401 (2001)

# Rossmania ukurunduensis Lar.N. Vassiljeva, Mycoscience 42(4): 401 (2001)

Facesoffungi number – FoF 2856

Fig. 15

Isotype – BPI 747566

Saprobic on dead bark of *Acer ukurunduense* Wall. Sexual morph – Ascomata immersed under the bark, with clusters of convergent, long, stout erumpent necks, without a ectostromatic disc, globose, coriaceous, black. Necks wide, opening through the ostiole, internally covered by hyaline periphyses, comprising brown cells of *textura porrecta*. Peridium thick, comprising outer, thick-walled, brown cells of *textura angularis* and inner thick-walled, hyaline cells of *textura angularis* at the base. Hamathecium aparaphysate. Asci 190–230 × 24–28  $\mu$ m, 8-spored, unitunicate, clavate to cylindrical, short pedicellate, apex rounded, lacking an apical ring. Ascospores 76–125 × 7–9  $\mu$ m, 4–5-seriate to fasciculate, long fusoid to cylindrical, slightly curved, 0–7-septate in Melzer's reagent, not constricted at the septa, hyaline, sometimes guttulate. Asexual morph – Undetermined.

Notes – Rossmania was introduced to accommodate taxa similar to Diaporthe, in its clusters of ascomata with long beaks and non-united ectostromatic disc, but differs in its septate elongate ascospores (Vasilyeva 2001). Therefore, superficially this genus is similar to Diaporthe or Phragmodiaporthe, but differs in having elongate Winterella-like ascospores. Rossmania comprises R. ukurunduensis and R. aculeata (Petch) Lar.N. Vassiljeva, the latter transferred from Phragmodiaporthe aculeata (Petch) Wehm., because it lacks true phragmospores (Vasilyeva 2001). Rossmania ukurunduensis differs from R. aculeata in having narrow, 5–7-septate ascospores, while R. aculeata has wider, 7–11-septate ascospores (Vasilyeva 2001). Rossmania ukurunduensis was collected in Russia associated with Acer ukurunduense, while R. aculeata was collected from Sri Lanka associated with Theaceae (Vasilyeva 2001). Vasilyeva (2001) placed Rossmania in Valsaceae. However, Rossman et al. (2007) and Lumbsch & Huhndorf (2010) placed Rossmania in Sydowiellaceae based on its morphology. The placement of Rossmania within Sydowiellaceae has been confirmed in phylogenetic studies (Fig. 1) (Kruys & Castlebury 2012, Maharachchikumbura et al. 2015). It differs from Winterella and Sillia by non stromatic ascoma and short cylindrical ascospores.

#### *Sillia* P. Karst., Bidr. Känn. Finl. Nat. Folk 23: 20, 159, 251 (1873)

Facesoffungi number – FoF 2857

Saprobic on dead stems of deciduous trees. Sexual morph – Stroma erumpent through cracks in bark, hemisphaerical, black on the surface, dark yellow internally, becoming rouge in 10 % KOH. Ascomata immersed in stromatic tissues, arising as several layers, globose, coriaceous, black, with cylindrical necks, protruding above the surface of the stroma. Peridium comprising thick-walled, brown cells of *textura angularis*. Asci 8-spored, unitunicate, narrowly clavate, short pedicellate, with a distinct small apical structure. Ascospores fasciculate, narrowly fusiform, slightly curved, hyaline, with 3–5-transverse septa, sometimes guttulate. Asexual morph – Undetermined.

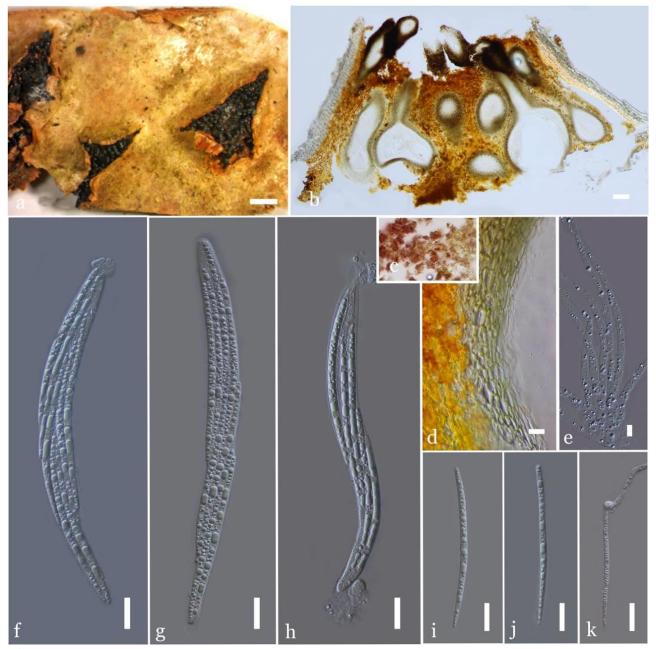
Type species – Sillia ferruginea (Pers.) P. Karst., Bidr. Känn. Finl. Nat. Folk 23: 159 (1873)

#### Sillia karstenii Senan., Camporesi & K.D. Hyde, sp. nov.

Index Fungorum number – IF552727, Facesoffungi number – FoF 2858

Fig. 16

Etymology – in honour of late Prof Petter Adolf Karsten, an eminent mycologist who introduced the genus *Sillia*.



**Figure 16** – *Sillia karstenii* (holotype). a Herbarium specimen. b Vertical section of stroma showing ascomata. c Stromatic tissues become umber in 10% KOH. d Peridium. e Paraphyses. f-h Asci. i-k Ascospores. Scale bars – a = 1 mm, b = 100  $\mu$ m, d, f-k = 20  $\mu$ m, e = 10  $\mu$ m.

#### Holotype – MFLU 16–2864

Saprobic on dead branches of *Corylus avellana* L. Sexual morph – Stromata poorly developed, pseudoparenchymatous, comprising yellowish brown cells of *textura angularis*, turning to umber in 10% KOH. Ascomata 720–1200 µm high, 230–265 µm diameter ( $\bar{x}=1032\times255$  µm, n = 10), immersed to erumpent, gregarious, globose to subglobose, black, coriaceous, papillate, ostiolate. Neck 315–700 µm long, 50–140 µm wide ( $\bar{x}=432\times70$  µm, n = 10), symmetrical, erumpent through host surface, internally covered by hyaline periphyses, darker than peridium cells at the base. Peridium 35–50 µm thick ( $\bar{x}=47$  µm, n = 15), comprising thick-walled, brown, compressed cells of *textura angularis*. Hamathecium comprising thin-walled, septate, unbranched, hyaline paraphyses 6–10 µm wide ( $\bar{x}=8$  µm, n = 15). Asci 175–230 × 15–18 µm ( $\bar{x}=200\times17$  µm, n = 25), 8-spored, unitunicate, cylindrical, straight or somewhat curved, with a short obtuse pedicel. Ascospores 110–120 × 4.5–6 µm ( $\bar{x}=114\times5.2$  µm, n = 30), overlapping bi-seriate,

filiform with pointed ends, 5–6-septate when mature, guttulate, smooth-walled, ends forming a swelling when germinating. Asexual morph – Undetermined.

Material examined – ITALY, Province of Forlì-Cesena, near Bertinoro, dead branch of *Corylus avellana* L (Corylaceae), 30 December 2013, Erio Camporesi, IT 1611 (MFLU 16–2864, holotype, *ibid* BBH, isotype).

Notes – Winter (1887) accommodated *Sillia* species with hyaline scolecosporous ascospores in the genus *Melogramma* which has brown phragmospores. Therefore, Lindau (1897) separated them again and placed *Sillia* in Melogrammataceae. Wehmeyer (1926) considered *Sillia* to be similar to *Cryptospora* Tul. (= *Ophiovalsa*), which was placed in Diaporthaceae. Ascospores of *Sillia* are filiform or acicular, scolecosporous and multi-septate. *Ophiovalsa* has unicellular, long-cylindric-fusoid ascospores. However, the presence or absence of septa is not thought to be a significant character in scolecosporous spores (Tibpromma et al. 2017). Hence *Sillia* was accepted as closely related to *Ophiovalsa* in Diaporthaceae. Munk (1953, 1957) and Dennis (1960) also treated *Sillia* as a separate genus of Diaporthaceae. However, Barr (1978) included *Sillia* in family Gnomoniaceae (Tribe Endothieae). Later, Rossman et al. (2007) transferred *Sillia* in Sydowiellaceae based on its morphology. Several phylogenetic studies supported the taxonomic placement of *Sillia* in Sydowiellaceae (De Silva et al. 2009, Kruys & Castlebury 2012). The type species *Sillia ferruginea* has been reported from various countries in the northern hemisphere, including Austria, Bulgaria, Canada, Czechoslovakia, Denmark, England, France, Germany, Ireland, Poland, Russia, Spain, Sweden, Ukraine and the USA (Farr & Rossman 2006).

Sillia karstenii is the second species reported from Italy. There are six Sillia species listed in Index Fungorum (2017). Sillia theae Hara (on Camellia sinensis (L.) Kuntze in Japan), S. kamatii Tilak, S.B. Kale & S.V.S. Kale (on dead stem of Gymnema sylvestre (Retz.) Schult. in India), S. celastrina R. Rao (on dead stems of Celastrus paniculata Willd. in Brazil) and S. biformis Rick from Brazil. S. albofusca Höhn. has fused papilla and thus is quite distinct from S. karstenii. The type species S. ferruginea (Pers.) P. Karst., S. italica de Silva, Camporesi & K.D. Hyde and the new species S. karstenii were associated with Corylus species. Sillia karstenii also differs in having large ascomata with very poorly developed stromata. Phylogenetically, Sillia karstenii is relatively distinct from sequenced species with strong support and its establishment as a new species is justified.

#### Tenuiappendicula Senan., Camporesi & K.D. Hyde, gen. nov.

Index Fungorum number – IF552728, Facesoffungi number – FoF 2859

Etymology – Name comprising two Latin words "tenuis" and "appendicula", referring to the slender ascospore appendages.

Saprobic on branches Alnus cordata (Loisel.) Duby. Sexual morph - Pseudostromata parenchymatous, comprising thin-walled, light brown cells of textura globosa. Ascomata aggregated, immersed, globose to subglobose, black, coriaceous, ostiolate, papillate. Papilla slanted to horizontal in substrate, curved, asymmetrical, sometimes convergent, widening towards the apex, internally covered by hyaline, filamentous periphyses. Peridium comprising an outer, thick-walled, light brown layer of very few compressed layers of cells of textura angularis and inner thin-walled, hyaline layer of compressed cells of textura angularis. Hamathecium aparaphysate. Asci 8-spored, unitunicate, fusiform to clavate, with short pedicel, apex blunt, lacking apical ring. Ascospores biseriate to overlapping biseriate, oval, 1-septate, thick-walled. Appendages at both ends, slender, long, hyaline. Asexual morph - Coelomycetous. Conidiomata acervuli, scattered, solitary, immersed, subglobose, unilocular, dark brown. Conidiomatal wall multi-layered, comprising thickwalled, brown, small cells of textura angularis. Conidiophores branched at the base, cylindrical, septate, hyaline, formed from the upper pseudoparenchyma, smooth-walled. Conidiogenous cells holoblastic, terminal, discrete, with 1–2 percurrent proliferations, indeterminate, cylindrical, simple, hyaline, smooth-walled. Conidia cylindrical, narrowing towards ends, with small spine at upper end, (4)–5–(6)-septate, hyaline, sometimes guttulate, thick-walled.

Type species – Tenuiappendicula alnicola Senan., Camporesi & K.D. Hyde

Tenuiappendicula alnicola Senan., Camporesi & K.D. Hyde, sp. nov.

Index Fungorum number – IF552729, Facesoffungi number – FoF 2860

Figs 17, 18

Etymology – Name after the host genus.

Holotype – MFLU 16–1265A

Saprobic on branch of Alnus cordata (Loisel.) Duby. Sexual morph – Pseudostromata parenchymatous, comprising thin-walled, light brown cells of textura globosa. Ascomata 180–200  $\times$  50–80 µm ( $\bar{x}$  = 191  $\times$  78 µm, n = 20), aggregated, immersed, globose to subglobose, black, coriaceous, ostiolate, papillate. Papilla 85–115  $\times$  30–40 µm ( $\bar{x} = 93 \times 36$  µm, n = 15), slanted to horizontal in the substrate, curved, asymmetrical, sometimes convergent, widening towards the apex, internally covered by hyaline, filamentous periphyses. Peridium 12–16  $\mu$ m ( $\bar{x} = 14 \mu$ m, n = 15), comprising an outer layer of thick-walled, light brown, compressed cells of textura angularis and inner thin-walled, hyaline layer of compressed cells of textura angularis. Hamathecium 5–9  $\mu$ m ( $\bar{x} = 7.8 \mu$ m, n = 15), comprising cellular, wide, septate, thin-walled paraphyses. Asci 105–120  $\times$  25–35 µm ( $\bar{x}$  = 116  $\times$  33 µm, n = 25), 8-spored, unitunicate, fusiform to clavate, with short pedicel, apex blunt, lacking an apical ring. Ascospores  $25-30 \times 7-10 \mu m$  ( $\bar{x} = 27 \times 8.8 \mu m$ , n = 25), biseriate to overlapping biseriate, oval, 1-septate, constricted at septum, thick-walled, appendaged. Appendages slender,  $\log 20-30 \times 1.5-2.5 \ \mu m \ (\overline{x} = 27 \times 2 \ \mu m, \ n = 20)$ , one each at each end, hyaline. Asexual morph – Conidiomata  $30-45 \times 120-135 \, \mu m$  ( $\bar{x} = 35 \times 130 \, \mu m$ , n = 15), acervuli, scattered, solitary, immersed, subglobose, unilocular, dark brown. Conidiomatal wall 35-50 µm ( $\bar{x} = 37$  µm, n = 15), multi-layered, comprising thick-walled, brown, small cells of textura angularis. Conidiophores  $16-21 \times 2.5-6 \mu m$  ( $\bar{x} = 18 \times 4.7 \mu m$ , n = 15), branched at the base, cylindrical, septate, hyaline, formed from the upper pseudoparenchyma, smooth-walled. Conidiogenous cells  $14-16 \times 4.5-5.5 \, \mu m$  ( $\bar{x} = 15 \times 4.8 \, \mu m$ , n = 25), holoblastic, terminal, discrete, with 1–2 percurrent proliferations, indeterminate, cylindrical, simple, hyaline, smooth-walled. Conidia 35–45  $\times$  4.5–5.5 µm ( $\bar{x} = 40 \times 5$  µm, n = 25), cylindrical, narrowing towards the ends, with a small spine at upper end, (4) –5–(6)-septate, hyaline, sometimes guttulate, thick-walled.

Culture characters – Colonies growing on MEA attained 1 cm within 7 days when incubated at 18 °C, flat, circular, irregular, white to off-white, woolly, sometimes mycelium in tufts.

Material examined – ITALY, Province of Forlì-Cesena, Fiumicello di Premilcuore, on dead branch of *Alnus cordata* (Loisel.) Duby (Betulaceae), 24 April 2013, Erio Camporesi, IT 1199 (MFLU 16–1265A, holotype), ex-type living culture MFLUCC 16–1452; *ibid* BBH isotype, TBRC.

Notes — *Tenuiappendicula* forms a well-supported clade as a sister taxon to *Chapeckia*. *Tenuiappendicula* is characterized by 1-septate ascospores with long, thin appendages and asci without an apical ring. We observed the asexual morph of *Tenuiappendicula* associated with ascomata and we extracted genomic DNA from the conidiomata. LSU and ITS sequences derived from both sexual and asexual morphs are identical and hence the holomorph is illustrated.

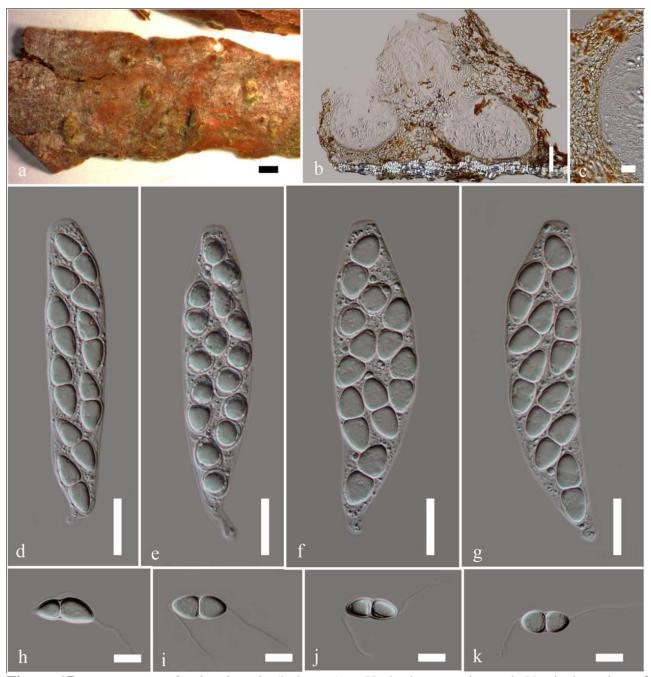
#### Tortilispora Senan. & K.D. Hyde, gen. nov.

Etymology – Name based on two Latin words "tortilis" and "spora" referring to the twisted ascospores.

Index Fungorum number – IF552802, Facesoffungi number – FoF 2863

Saprobic on dead twigs and branches. Sexual morph – Stromata forming angular, rather prominent pustules, with grey-brown to orange-brown ectostromatic discs, with 2–15 ascomata embedded in a well-developed ectostroma in a valsoid configuration. Ascomata immersed, aggregated, globose to subglobose, coriaceous, dark brown to black, ostiolate, papillate, paraphysate. Papilla black with periphyses. Peridium comprising thick-walled, dark brown to black, small cells of *textura angularis*. Asci 8-spored, unitunicate, cylindrical, sessile, apex rounded, with a J-, apical ring. Ascospores fasciculate, narrowly fusiform to long cylindrical, hyaline, transseptate. Asexual morph – Undetermined.

Type species – Tortilispora aurantiaca (Wehm.) Senan. & K.D. Hyde



**Figure 17** – *Tenuiappendicula alnicola* (holotype). a Herbarium specimen. b Vertical section of ascomata. c Peridium. d-g Asci. h-k Ascospores. Scale bars –  $a=500~\mu m$ ,  $b=100~\mu m$ ,  $c=20~\mu m$ ,  $d-g=20~\mu m$ ,  $h-k=10~\mu m$ .

Tortilispora aurantiaca (Wehm.) Senan. & K.D. Hyde, comb. nov.

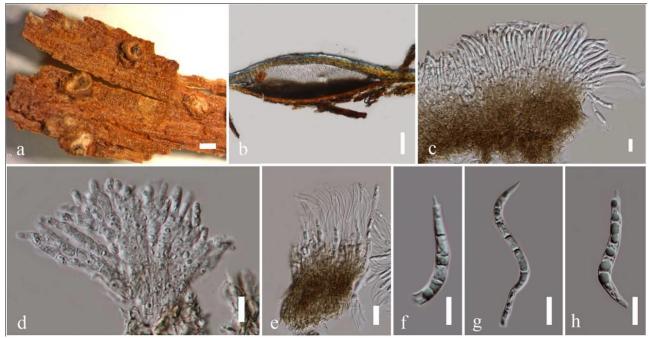
Index Fungorum number – IF552803, Facesoffungi number – FoF 2864 Fig. 19

Basionym – *Cryptospora aurantiaca* Wehm., Canadian Journal of Research, Section C 20(12): 589 (1942)

Holotype (part) – MICH 13964

Saprobic on dead twigs and branches. Sexual morph – Stromata 0.8–3 mm diameter, forming angular, rather prominent pustules, with grey-brown to orange-brown ectostromatic discs, immersed, scattered, solitary, black, parenchymatous, with 2–15 ascomata embedded in a well-developed ectostroma, which consists of hyaline to subhyaline, densely compact, thin-walled hyphae, with a valsoid configuration. Ascomata 675–875  $\mu$ m high, 225–300  $\mu$ m wide ( $\bar{x}=765\times275~\mu$ m, n = 25), immersed, aggregated, globose to subglobose, coriaceous, dark brown to black,

ostiolate, papillate. Papilla 375–500 µm high, 100–125 µm wide ( $\bar{x}=107\times466$  µm, n = 20), black, comprising parallel to interwoven, thick-walled, light to dark brown hyphae, internally covered by hyaline, filamentous periphyses. Peridium 10–15 µm wide ( $\bar{x}=13$  µm, n = 25), comprising thick-walled, dark brown to black, small cells of *textura angularis*. Hamathecium comprising broad, tapering band-like, septate paraphyses. Asci 90–100 × 11–13 µm ( $\bar{x}=93\times12$  µm, n = 25), 8-spored, unitunicate, cylindrical, sessile, apex rounded at apex, with a J- apical ring, comprising two refractive bodies. Ascospores 85–95 × 3–4 µm ( $\bar{x}=93\times3.5$  µm, n = 25), fasciculate, narrowly fusiform to long cylindrical, hyaline, 1–7-septate, smooth-walled. Asexual morph – Undetermined.



**Figure 18** – *Tenuiappendicula alnicola* (holotype). a Herbarium specimen. b Vertical section of conidioma. c, d Conidiophores, conidiogenous cells and conidia. e Broken conidial wall attached to conidiogenous cells. f-h Conidia. Scale bars –  $a = 500 \mu m$ ,  $b = 100 \mu m$ ,  $c-e = 10 \mu m$ ,  $f-h = 10 \mu m$ .

Material examined – CANADA, Nova Scotia, on *Alnus* sp. Mill. (Betulaceae), 09 July 1933, L.W. Wehmeyer, (1624), MICH (13964), holotype.

Notes – Winterella was accepted in Sydowiellaceae by Maharachchikumbura et al. (2016). There is, however, some confusion surrounding Winterella and we follow Mejia et al. (2008) and treat Ophiovalsa and Winterella as synonyms of Cryptosporella (Gnomoniaceae). Winterella albofusca, W. aurantiaca, and W. cinctula were excluded from Cryptosporella by Mejia et al. (2008) based on morphological and molecular data. LSU data (not shown) indicated that W. albofusca and W. aurantiaca belonged in the Diaporthales, although outside Gnomoniaceae. The above-mentioned species are characterized by multi-septate ascospores, and are placed in Sydowiellaceae in our phylogenetic analyses (Fig. 1). Since they do not cluster with Sillia species and are not typical of Rossmania species, genera with filiform ascospores, there is no obvious place to accommodate them. We thus introduce a new genus Tortilispora to accommodate these taxa with T. aurantiaca as the type species and we proposed T. albofusca (IF 552807), T. cinctula (IF552808) for other species.



**Figure 19** – *Tortilispora aurantiaca* (holotype). a, b Herbarium packet. d Herbarium specimen. c Ascomata on substrate. e Horizontal cross section of stromata. f, g Cross sections of ascomata. h Papilla. i Paraphyses. j Peridium. k-m Asci in water. n-q Ascospores. Scale bars: c,  $e = 500 \, \mu m$ , f-h =  $100 \, \mu m$ ,  $i = 20 \, \mu m$ , j-q =  $10 \, \mu m$ . Note – Rouge stromatic tissues in  $10 \, \%$  KOH, arrowed in g.

# Diaporthales genera, incertae sedis

In this section, we deal with *Uleoporthe* which appears to be atypical of Sydowiellaceae.

*Uleoporthe* Petr., Annls mycol. 39(4/6): 279 (1941) Facesoffungi number – FoF 2861 Saprobic or biotrophic on recently dead plant tissues. Sexual morph – Stromata roughly circular, on the surface of leaves. Ascomata surrounding the conidiomata, deeply immersed in stromatic tissues, globose to subglobose, ostiolate, papillate, with copious paraphyses. Peridium of strongly flattened, pale brown cells of *textura angularis* at the base and thick-walled, hyaline cells of *textura angularis* in the papilla region. Asci 8-spored, unitunicate, short-pedicellate to almost sessile, apex rounded. Ascospores overlapping uniseriate to biseriate, clavate, 2-celled, hyaline. Asexual morph – Conidiomata loculate, irregular in shape, ostiolate. Peridium comprising layers of brown, thin-walled cells of *textura epidermoidea*. Conidiophores poorly differentiated, thin-walled, ellipsoidal. Conidiogenous cells in small clusters or short branched chains of cells, irregular or narrowly flask-shaped, tapering towards the upper region, both percurrent and sympodial proliferation, minute, slightly flared collarettes where percurrent proliferation occurs. Conidia ellipsoidal to fusiform, hyaline, aseptate (description of asexual morph from Cannon 2001).

Type species – *Uleoporthe orbiculata* (Syd. & P. Syd.) Petr., Annls mycol. 39(4/6): 280 (1941)

Uleoporthe orbiculata (Syd. & P. Syd.) Petr., Annls mycol. 39(4/6): 280 (1941)

Basionym – Valsonectria orbiculata Syd. & P. Syd., Annls mycol. 14(1/2): 86 (1916)

Facesoffungi number – FoF 2862

Lectotype – W (E. Ule 3389)

Saprobic or biotrophic on recently dead plant tissues. Sexual morph - Stromata 3-7 mm diameter, roughly circular in external view, epigenous on leaves, occupying the upper epidermis and palisade layers of the leaf, erumpent, convex, initially with simple stellate splits, later becoming more complex and branched, mostly coalescing with yellow-brown host tissue at the surface, with poorly to well-developed black line around the stroma and internal eustromatic tissue with thick-walled, glassy hyaline cells of textura globulosa to textura angularis around the basal region of ascomata and dark brown cells of textura globulosa to textura angularis around the ostioles. Ascomata 435–450 µm high, 400–460 µm wide ( $\bar{x} = 445 \times 450$  µm, n = 15), surrounding the conidiomata, deeply immersed in stromatic tissues, solitary, globose to subglobose, coriaceous, light brown, ostiolate. Papilla 190–200 µm high, 80–90 µm wide ( $\bar{x} = 195 \times 85$  µm, n = 15), conical, slightly eccentric, hyaline, filamentous periphysate. Peridium 15–20 µm wide ( $\bar{x} = 19$  µm, n = 15), comprising 5-6 layers of strongly flattened, pale brown cells of textura angularis at the base and thick-walled, hyaline cells of textura angularis in the ostiolar region. Hamathecium comprising copious, very thin-walled, persistent paraphyses, swollen at the base and gradually tapering towards the apex. Asci  $60-80 \times 10-20 \, \mu m$  ( $\bar{x} = 65 \times 14 \, \mu m$ , n = 25), 8-spored, unitunicate, narrowly clavate, short-pedicellate to almost sessile, apex rounded, sometimes slightly attenuate. Ascospores  $16-20 \times 5-6.5 \mu m$  ( $\bar{x} = 18 \times 5.4 \mu m$ , n = 15), overlapping uniseriate to biseriate, clavate, upper cell ellipsoidal and the lower cylindric-ellipsoidal, with a median to slightly supramedian septum, thick-walled, hyaline, smooth-walled, each cell sometimes biguttulate. Asexual morph - Conidiomata loculate, irregularly-shaped, ostiolate. Peridium comprising 2-3 layers of brown, thin-walled cells of textura epidermoidea. Conidiophores poorly differentiated, thin-walled, ellipsoidal. Conidiogenous cells in small clusters or short branched chains of cells, irregular or narrowly flask-shaped, tapering towards the upper region, both percurrent and sympodial proliferation, minute, with slightly flared collarettes where percurrent proliferation occurs. Conidia ellipsoidal to fusiform, apex acute to obtuse, base narrowly truncate, thick-walled, hyaline, aseptate, smooth-walled.

Material examined – VENEZUELA, Miliquebe, Rio Cuquenau, in living leaves of *Cybianthus fulvopulverulentus* Mart. (Primulaceae), February 1910, E. Ule, 3389, W, lectotype.

Notes – The asexual morph of *Uleoporthe* was described by Sydow & Sydow (1916) connected to *Valsonectria orbiculata* (= *Uleoporthe orbiculata*). However, *Valsonectria* is now considered a member of the Bionectriaceae and its type species, *Valsonectria pulchella* Speg., is clearly not conspecific with *U. orbiculata* (Hawksworth et al. 1995). Muller & von Arx (1962) treated *Uleoporthe* as a synonym of *Phylloporthe*. However, *Uleoporthe* is characterized by a

Fig. 20



**Figure 20** – *Uleoporthe orbiculata* (holotype). a Herbarium packet. b-d Herbarium specimen and stromata. e Cross section of ascoma. f Peridium. g-i Asci. j-m Ascospores. n Conidiophores, conidiogenous cells attached to conidia. o Conidia. Scale bars – c = 1 mm,  $d = 500 \mu m$ ,  $e = 100 \mu m$ ,  $f = 10 \mu m$ , g - i, g - i

prosenchymatous stroma and hyaline, 2-celled ascospores, while *Phylloporthe* has a plectenchymatous to pseudoparenchymatous stroma. *Phylloporthe* also has strongly erumpent stromata composed mostly of fungal tissue, short pedicellate asci which are evanescent at the base, an aparaphysate centrum and periphysate ostioles (Cannon 2001). Petrak (1941) reported characters of the asexual morph of *Uleoporthe* as similar to *Phomopsis*. We could not observe the asexual morph from the lectotype. Barr (1978) did not include *Uleoporthe* as a synonym of *Phylloporthe* but placed *Uleoporthe* as a member of Gnomoniaceae (subfamily Stegophoroideae, tribe Stegophoreae). However, this was not accepted by Cannon (2001) based on the stromatic development and the small, acervular, spermatial asexual morph. Cannon (2001) placed *Uleoporthe* in Melanconidaceae, but this is problematic because the classification was developed based on exclusively temperate taxa, which may show different patterns of evolution to tropical taxa, and

fewer asexual morph characters (Cannon 2001). Lumbsch & Huhndorf (2010) and Maharachchikumbura et al. (2015) placed *Uleoporthe* in Sydowiellaceae. Sequence data for *Uleoporthe* is unavailable in GenBank to clarify the correct phylogenetic position. Thus, fresh collections and sequence data are needed to establish the true affinities of this taxon.

We examined the holotype of *Uleoporthe orbiculata*. We could not observe an apical ring in the ascus as it has a delicate wall. Morphologically, *U. orbiculata* is atypical of other taxa in Sydowiellaceae. *Uleoporthe orbiculata* is a leaf epiphyte forming circular, erumpent to superficial stromata, while other taxa form immersed stromata on bark. Therefore, we exclude *Uleoporthe* from Sydowiellaceae and place it in Diaporthales genera, *incertae sedis*, until sequence data is available.

#### Acknowledgements

K.D. Hyde thanks Kunming Institute of Botany for the award of Visiting Professor.

#### References

- Alexopoulos CJ, Cation D. 1952 *Gnomonia fragariae* in Michigan. Mycologia 44, 221–223
- Ariyawansa HA, Hawksworth DL, Hyde KD, Jones EBG. et al. (2014) Epitypification and neotypification: guidelines with appropriate and inappropriate examples. Fungal Diversity 69: 57–91
- Barr ME. 1959 Northern pyrenomycetes 1. Canadian eastern arctic. Contributions Institute Botany, University Montreal 73, 1–101
- Barr ME. 1978 The Diaporthales in North America: with emphasis on *Gnomonia* and its segregates. Mycologia Memoirs 7, 1–232
- Barr ME. 1991 Revisions and additions to the Diaporthales. Mycotaxon 41, 287-305
- Bolay A. 1972 Contribution à la connaissance de Gnomonia comari Karsten. Étude taxonomique, phytopathologique et recherches sur sa croissance in vitro. Bericht der Schweizerischen Botanischen Gesellschaft 81, 398–482
- Campbell J, Shearer CA, Crane JL, Fallah PM. 2003 A reassessment of two freshwater ascomycetes, *Ceriospora caudae-suis* and *Submersisphaeria aquatica*. Mycologia. 95:41-53.
- Cannon PF. 2001 Rediscovery and redescription of the genus *Uleoporthe* (Melanconidaceae). Fungal Diversity 7, 17–25
- Carbone I, Kohn LM. 1999 A method for designing primer sets for speciation studies in filamentous ascomycetes. Mycologia 91, 553 556
- Chomnunti P, Hongsanan S, Aguirre-Hudson B, Tian Q et al. 2014 The sooty moulds. Fungal Diversity 66, 1–36
- Clements FE, Shear CL. 1931 Genera of Fungi. 2nd ed. Wilson, New York, USA.
- Currey F. 1858 Synopsis of the fructification of the compound Sphaeriae of the Hookerian herbarium. The Transactions of the Linnean Society of London 22, 257–287
- De Silva H, Castlebury LA, Green S, Stone JK. 2009 Characterization and phylogenetic relationships of *Anisogramma virgultorum* and *A. anomala* in the Diaporthales (Ascomycota). Mycological Research 113, 73–81
- De Silva NI, Lumyong S, Hyde KD, Bulgakov T et al. 2016 Mycosphere Essays 9: Defining biotrophs and hemibiotrophs Mycosphere 7, 545–559
- Dennis RWG. 1960 Cryptogamic botany. In: A discussion on the biology of the southern cold temperate zone. Proceedings of the Royal Soc. Series B 152 (949), 539–540
- Ellis JB, Everhart BM. 1892 New species of fungi. Journal of Mycology 7 (2), 130–135
- Ellis MA, Kuter GA, Wilson LL. 1984 Fungi that cause cane cankers on thornless blackberry in Ohio. Plant Disease 68, 812–815
- Farr DF, Bills GF, Chamuris GP, Rossman AY. 1989 Fungi on Plants and Plant Products in the United States, APS Press, St. Paul, Minnesota

- Farr DF, Rossman AY. 2006 Fungal Databases, Systematic Mycology and Microbiology Laboratory, ARS, USDA. URL http://nt.ars-grin.gov/fungaldatabases/ [accessed 10–02–2017]
- Felsenstein J. 1985 Confidence limits on phylogenies: an approach using the bootstrap. Evolution 39(4), 783–791
- Fuckel KWGL. 1863 Fungi Rhenani exsiccati a L. Fuckel collecti. Hedwigia 2, 132–136
- Glass NL, Donaldson GC 1995 Development of primer sets designed for use with the PCR to amplify conserved genes from filamentous ascomycetes. Applied and Environmental Microbiology 61, 1323–1330
- Glawe DA. 1985 *Hapalocystis berkeleyi* and *Macrodiplodiopsis desmazieresii* in artificial culture. Mycologia 77, 880–886
- Hall TA. 1999 BioEdit: a user-friendly biological sequence alignment editor and analysis program for Windows 95/98/NT. Nucleic Acids Symposium Series 41, 95–98
- Hawksworth. DL, Kirk PM, Sutton BC, Pegler DN. 1995 Ainsworth& Bisby's Dictionary of the Fungi. 8th edition.CAB International, Wallingford
- Index Fungorum. 2017 (http://www.indexfungorum.org/Names/Names.asp
- Jaklitsch WM, Voglmayr H. 2004 *Hapalocystis occidentalis* a new species of Diaporthales from North America and a key to the species of *Hapalocystis*. Studies in Mycology 50, 229–234
- Jayasiri SC, Hyde KD, Ariyawansa HA, Bhat J et al. 2015 The Faces of Fungi database: Fungal names linked with morphology, molecular and human attributes. Fungal Diversity 74, 3–18
- Jeewon R, Hyde KD. 2016 Establishing species boundaries and new taxa among fungi: recommendations to resolve taxonomic ambiguities. Mycosphere 7 (11), 1669–1677
- Jeewon R, Liew ECY, Hyde KD. 2002 Phylogenetic relationships of *Pestalotiopsis* and allied genera inferred from ribosomal DNA sequences and morphological characters. Molecular Phylogenetics and Evolution 25, 378–392.
- Jeewon R, Liew, ECY, Hyde KD. 2004 Phylogenetic evaluation of species nomenclature of *Pestalotiopsis* in relation to host association. Fungal Diversity 17, 39–55.
- Kirk PM, Cannon PF, Minter DW, Stalpers JA. 2008 Dictionary of the Fungi. 10th ed. Wallingford, UK: CAB International. 1–784
- Kishino H, Hasegawa M. 1989 Evaluation of the maximum likelihood estimate of the evolutionary tree topologies from DNA sequence data. Journal of Molecular Evolution 29, 170–179
- Klebahn H. 1918 Hauptund Nebenfruchtformen der Askomuyzeten, Verlag von Gebruder Borntraeger, Leipzig
- Kobayashi T. 1970 Notes on new or little-known fungi inhabiting woody plants in Japan II. Transactions of the Mycological Society of Japan 11, 57–63
- Kruys A, Castlebury LA. 2012 Molecular phylogeny of Sydowiellaceae–resolving the position of *Cainiella*. Mycologia 104, 419–426
- Lindau G. 1897 Pyrenomycetineae, Laboulbeniineae. In: A. Engler & K. Prantl (eds) Die Natürlichen Pflanzenfamilien 1(1), 321–505
- Liu JK, Hyde KD, Jones EBG, Ariyawansa HA et al. 2015 Fungal diversity notes 1–110: taxonomic and phylogenetic contributions to fungal species. Fungal Diversity 72, 1–197
- Liu Y, Whelen S, Hall BD. 1999 Phylogenetic relationships among ascomycetes: evidence from an RNA polymerase II subunit. Molecular Biology and Evolution 16, 1799–1808
- Lumbsch HT, Huhndorf SM. 2009 Outline of Ascomycota 2009 Part 1. Myconet 14, 1–42
- Lumbsch HT, Huhndorf SM. 2010 Myconet Volume 14. Part One. Outline of Ascomycota 2009. Part Two. Notes on Ascomycete Systematics. Nos. 4751–5113. Fieldiana Life Earth Sciences 1, 1–64
- Lumbsch TH, Huhndorf SM. 2007 Outline of Ascomycota: 2007. Myconet (Chicago, USA: The Field Museum, Department of Botany) 13, 1–58
- Maas JL, Galleta GJ, Ellis MA. 1989 Cane canker diseases of thornless blackberry in eastern United States. Acta Horticulturae 262, 205–208

- Maas JL. 1998 Compendium of Strawberry Diseases, 2nd edn. APS Press, St. Paul, Minnesota. 38–39
- Maharachchikumbura SS, Hyde KD, Jones EBG, McKenzie EHC et al. 2015 Towards a natural classification and backbone tree for Sordariomycetes. Fungal Diversity 72, 199–301
- Maharachchikumbura SSN, Hyde KD, Jones EBG, McKenzie EHC et al. 2016 Families of Sordariomycetes. Fungal Diversity 79: 1–317.
- Mejía LC, Castlebury LA, Rossman AY, Sogonov MV, White JF. 2008 Phylogenetic placement and taxonomic review of the genus *Cryptosporella* and its synonyms *Ophiovalsa* and *Winterella* (Gnomoniaceae, Diaporthales). Mycological Research 112, 23–35
- Mejía LC, Castlebury LA, Rossman AY, Sogonov MV, White JF. 2011 A systematic account of the genus *Plagiostoma* (Gnomoniaceae, Diaporthales) based on morphology, host associations, and a four-gene phylogeny. Studies in Mycology 68, 211–235
- Monod M. 1983 Monographie taxonomique des Gnomoniaceae (Ascomycetes de l'ordre des Diaporthales). I. Beiheft Sydowia 9, 1–315
- Morocko I, Fatehi J. 2007 Molecular characterization of strawberry pathogen *Gnomonia fragariae* and its genetic relatedness to other *Gnomonia* species and members of Diaporthales. Mycological Research 111, 603–614
- Moročko I. 2006 Characterization of the strawberry pathogen *Gnomonia fragariae*, and biocontrol possibilities. Diss. (sammanfattning/summary) Uppsala: Sveriges lantbruksuniv., Acta Universitatis agriculturae Sueciae, 1652–6880
- Müller E, von Arx JA. 1962 Die Gattungen der didymosporen Pyrenomyceten. Beiträge zur Kryptogamenflora der Schweiz. 11(2), 1–922
- Müller E. 1956 Uber die neue sphaeriale Gattung Cainiella. Sydowia 10, 118–121
- Müller E. 1957 Haupt- und Nebenfruchtformen bei Guignardia Viala et Ravas. Beihefte zur Sydowia 1, 210–224
- Munk A. 1953 The system of the Pyrenomycetes. A contribution to a natural classification of the group Sphaeriales sensu Lindau. Dansk Botanisk Arkives 15, 1–163
- Munk A. 1957 Danish Pyrenomycetes. A preliminary flora. Dansk Botanisk Arkives 17, 1–491 MycoBank. 2017 http://www.mycobank.org/
- Nordskog B, Stensvand A, Heiberg N. 2003 Fungi Occurring on Aerial Constituents of Cultivated Blackberry (*Rubus fruticosus* L.) in Norway. Acta Agriculturae Scandinavica, Section B, Soil & Plant Science. 53, 21–28
- Nylander JAA. 2004 MrModeltest v2. Program Distributed by the Author. Evolutionary Biology Centre, Uppsala University
- Page RDM 1996 TREEVIEW: An application to display phylogenetic trees on personal computers. Computer Applications in the Biosciences 12, 357–358
- Petrak F. 1941 Mykologische Notizen XIV (nos 931-1000). Annales Mycologici 39, 251–349
- Phillips AJL, Alves A, Pennycook SR, Johnston PR et al. 2008 Resolving the phylogenetic and taxonomic status of dark-spored teleomorph genera in the Botryosphaeriaceae. Persoonia 21, 29–55
- Phookamsak R, Norphanphoun C, Tanaka K. et al. 2015 Towards a natural classification of Astrosphaeriella-like species; introducing Astrosphaeriellaceae and Pseudoastrosphaeriellaceae fam. nov. and Astrosphaeriellopsis, gen. nov. Fungal Diversity 74–143
- Raciborski M. 1900 Parasitische Algen und Pilze Java's. H. Teil, 1–46
- Reid J, Booth C. 1987 *Winterella*, the correct name for *Cryptospora* and *Ophiovalsa*. Canadian Journal of Botany 65, 1320–1342
- Ronquist F, Teslenko M, van der Mark P, Ayres DL et al. 2012 MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across a large model space. Systematic Biology 61, 539–42

- Rossman AY, Castlebury LA, Samuels GJ. 2006 Phylogeny and biodiversity of the Hypocreales and Diaporthales. Eighth International Mycological Congress, 21–25, August 2006, Cairns. Congress Handbook & Abstracts, Book 2
- Rossman AY, Farr DF, Castlebury LA. 2007 A review of the phylogeny and biology of the Diaporthales. Mycoscience 48, 135–144
- Saccardo PA. 1883 Sylloge Fungorum, vol. 2. Saccardo, Pavia, Italy.
- Schröter J. 1897 Pilze. In: Cohn F (ed), Kryptogamen-Flora von Schlesien 3.2. Kern, Breslau, Germany.
- Silvestro D, Michalak I. 2012 raxmlGUI: a graphical front-end for RaxML Organisms Diversity & Evolution 12, 335–337
- Sogonov MV, Castlebury LA, Rossman AY, Mejía LC, White JF. 2008 Leaf-inhabiting genera of the *Gnomoniaceae*, *Diaporthales*. Studies in Mycology 62, 1–79
- Stamatakis A, Hoover P, Rougemont J. 2008 A rapid bootstrap algorithm for the raxml web servers. Systematic Biology 57, 758–771
- Swofford DL. 2003 –. PAUP\*. Phylogenetic Analysis Using Parsimony (\*and Other Methods). Version 4.Sinauer Associates, Sunderland, Massachusetts.
- Sydow H, Sydow P. 1916 Weitere Diagnosen neuer Philippinischer Pilze. Annales Mycologici 14, 362–364
- Sydow H. 1936 Novae fungorum species XXIV. Annales Mycologici 34, 411–422
- Tibpromma S, Karunarathna SC, Hyde KD, Camporesi E et al. 2017 Fungal diversity notes 491–601: taxonomic and phylogenetic contributions to fungal taxa. Fungal Diversity (in prep)
- Tulasne ELR, Tulasne C. 1863 Selecta Fungorum Carpologia 2: i-xix, 319 pp., 34 tabs. Paris; Imperial. Typograph.
- Van Adriechem MCY, Bosher JE. 1958 Leaf blotch and petiole blight of strawberry caused by *Gnomonia fructicola*. Plant Disease Reporter 42, 772–775
- Vasilyeva L. 2001 Pyrenomycetes of the Russian Far East–Additions and corrections. 1. Rossmania ukurunduensis gen. et sp. nov. Mycoscience. 42, 399–401
- Vasilyeva LN. 1987 Pirenomitsety i Lokuloaskomitsety severa Dalbnego Vostoka: 1–255
- Vasilyeva LN. 1998 Nizshie Rasteniya, Griby i Mokhoobraznye Dalnego Vostoka Rossii. 4, 1–419
- Vilgalys R, Hester M. 1990 Rapid genetic identification and mapping of enzymatically amplified ribosomal DNA from several *Cryptococcus* species. Journal of Bacteriology 172, 4239–4246
- Voglmayr H, Jaklitsch WM. 2014 Stilbosporaceae resurrected: generic reclassification and speciation. Persoonia 33, 61–82
- Wehmeyer LE. 1926 A biologic and phylogenetic study of the stromatic Sphaeriales. American Journal of Botany 13, 575–645
- Wehmeyer LE. 1933 The British species of the genus *Diaporthe* Nits. and its segregates. Transactions of the British Mycological Society 17, 237–295
- Wehmeyer LE. 1941 A revision of *Melanconis*, *Pseudovalsa*, *Prosthecium* and *Titania*. University of Michigan Studies Scientific Series 14, 1–161
- White TJ, Bruns T, Lee S, Taylor J. 1990 Amplification and direct sequencing of fungal ribosomal RNA genes for phylogenetics. In: PCR Protocols: a guide to methods and applications. (Innis MA, Gelfand DH, Sninsky JJ, White TJ, eds). Academic Press, New York, USA, 315–322
- Winter G. 1887 Pilze, Ascomyceten. In: Rabenhorst's Kryptogamen-Flora von Deutschland, Oesterreich und der Schweiz [i–vi] 1(2), 1–928