

I.C. Senanayake^{1,2,3}, P.W. Crous⁴, J.Z. Groenewald⁴, S.S.N. Maharachchikumbura⁵, R. Jeewon⁶, A.J.L. Phillips⁷, J.D. Bhat^{8,9}, R.H. Perera³, Q.R. Li¹⁰, W.J. Li^{1,2,3}, N. Tangthirasunun^{11,12}, C. Norphanphoun³, S.C. Karunarathna^{1,2*}, E. Camporesi^{13,14,15}, I.S. Manawasighe¹⁶, A.M. Al-Sadi⁵, and K.D. Hyde^{1,2,3}

¹Key Laboratory for Plant Diversity and Biogeography of East Asia, Kunming Institute of Botany, Chinese Academy of Science, Kunming 650201, Yunnan, China; ²East and Central Asia, World Agroforestry Centre, Kunming 650201, Yunnan, China; ³Center of Excellence for Fungal Research, Mae Fah Luang University, Chiang Rai, Thailand; ⁴Westerdijk Fungal Biodiversity Institute, Uppsalalaan 8, 3584 CT Utrecht, The Netherlands; ⁵Department of Crop Sciences, College of Agricultural and Marine Sciences, Sultan Qaboos University, P.O. Box 34, Al-Khod 123, Omari, ⁶Department of Health Sciences, Faculty of Science, University of Mauritius, Reduit, 80837, Mauritius; ⁷Faculty of Sciences, Biosystems and Integrative Sciences Institute (BiolSI), University of Lisbon, Campo Grande, 1749-016 Lisbon, Portugal; ⁸Department of Botany, Goa University, Goa 403 206, India; ⁹No. 128/1-J, Azad Housing Society, Curca, P.O. Goa Velha 403108, India; ¹⁰Engineering Research Center of Southwest Bio-Pharmaceutical Resources, Ministry of Education, Guizhou University, Guiyang, Guizhou 550025, China; ¹¹Univ Paris Diderot, Sorbonne Paris Cité, Institut des Energies de Demain (IED), Paris 75205, France; ¹²Univ Paris Sud, Institut de Génétique et Microbiologie, UMR8621, Orsay 91405, France; ¹³A.M.B. Gruppo Micologico Forlivese, Antonio Cicognani, Via Roma 18, Forli, Italy; ¹⁴A.M.B. Circolo Micologico, Giovanni Carini, 314 Brescia, Italy; ¹⁵Società per gliStudiNaturalisticidella Romagna, 144 Bagnacavallo, RA, Italy; ¹⁶Institute of Plant and Environment Protection, Beijing Academy of Agriculture and Forestry Sciences, No. 9 of ShuGuangHuaYuanZhongLu, Haidian District, Beijing 100097, China

*Correspondence: S.C. Karunarathna, samanthakarunarathna@gmail.com

Abstract: Diaporthales is an important ascomycetous order comprising phytopathogenic, saprobic, and endophytic fungi, but interfamilial taxonomic relationships are still ambiguous. Despite its cosmopolitan distribution and high diversity with distinctive morphologies, this order has received relativelyiaceae, *Macrohilaceae, Melanconidaceae, Pseudoplagiostomaceae, Schizoparmaceae, Stilbosporaceae* and *Sydowiellaceae*. Taxonomic uncertainties among genera are also clarified and recurrent discrepancies in the taxonomic position of families within the *Diaporthales* are discussed. An updated outline and key to families and genera of the order is presented.

Key words: Multi-gene DNA phylogeny, New taxonomic arrangement, Phytopathogenic fungi, Sordariomycetes, Systematics.

Taxonomic novelties: New families: Apiosporopsidaceae Senan. Maharachch. & K.D. Hyde, Apoharknessiaceae Senan. Maharachch. & K.D. Hyde, Asterosporiaceae Senan. Maharachch. & K.D. Hyde, Auratiopycnidiellaceae Senan. Maharachch. & K.D. Hyde, Erythrogloeaceae Senan. Maharachch. & K.D. Hyde, Melanconiellaceae Senan. Maharachch. & K.D. Hyde, Prosopidicolaceae Senan. Maharachch. & K.D. Hyde, Prosopidicolaceae Senan. & K.D. Hyde, New genera: Marsupiomyces Senan. & K.D. Hyde, Microascospora Senan., Camporesi & K.D. Hyde, Phaeoappendicospora Senan., Q.R. Li & K.D. Hyde, Paradiaporthe Senan., Camporesi, & K.D. Hyde, Hyaliappendispora Senan., Camporesi & K.D. Hyde, Chiangraiomyces Senan. & K.D. Hyde; New species: Chiangraiomyces bauhiniae Senan. & K.D. Hyde, Coniella pseudokoreana Senan., Tangthir. & K.D. Hyde, Cytospora centrivillosa Senan., Camporesi & K.D. Hyde, Cytospora rosae Senan., Camporesi & K.D. Hyde, Gnomoniopsis agrimoniae Senan., Camporesi & K.D. Hyde, Hyaliappendispora galii Senan., Camporesi & K.D. Hyde, Marsupiomyces epidermoidea R.H. Perera, Senan., Bulgakov & K.D. Hyde, Marsupiomyces quercina Senan., Camporesi & K.D. Hyde, Melanconis italica Senan., Camporesi & K.D. Hyde, Microascospora rubi Senan., Camporesi & K.D. Hyde, Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, Melanconis italica Senan., Camporesi & K.D. Hyde, Microascospora rubi Senan., Camporesi & K.D. Hyde, Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, Microascospora rubi Senan., Camporesi & K.D. Hyde, Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, Microascospora rubi Senan., Camporesi & K.D. Hyde, Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, Microascospora rubi Senan., Camporesi & K.D. Hyde, Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, Microascospora rubi Senan., Camporesi & K.D. Hyde, Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, Microascospora fragariae (F

Available online 1 August 2017; http://dx.doi.org/10.1016/j.simyco.2017.07.003.

INTRODUCTION

The *Diaporthales* is a distinct order in the subclass *Diaporthomycetidae* (*Sordariomycetes*) and it includes pathogens, saprobes and endophytes, with no known coprophilous, hypersaprobes or mycophylic species (Barr 1978, Rossman *et al.* 2007, Vasilyeva *et al.* 2007, Maharachchikumbura *et al.* 2015, 2016). Taxa of this order inhabit a wide diversity of hosts and substrates, including most economically and ecologically important trees and crops, soil and living animal and human tissues (Barr 1978, Gryzenhout *et al.* 2006c). Species in *Diaporthales* form solitary or aggregated, immersed to erumpent, rarely superficial, orange, brown to black

perithecial ascomata, with short or long necks, that are located in stromatic tissues or substrates, with a centrum (or hamathecium) lacking or with few paraphyses (Alexopoulos & Mims 1978, Barr 1978, Castlebury *et al.* 2002). Asci are unitunicate with a conspicuous refractive ring (Hawksworth *et al.* 1995, Rossman *et al.* 2007). Ascospore morphology is diverse, ranging from short to elongate and aseptate or septate with hyaline or pigmented walls. The asexual morphs of *Diaporthales* are generally coelomycetous (Rossman *et al.* 2007), producing acervuli or pycnidial conidiomata, with or without a well-developed stroma. Conidiogenesis is phialidic or rarely annellidic and conidia are usually unicellular or 1-septate (Rossman *et al.* 2007).

Peer review under responsibility of Westerdijk Fungal Biodiversity Institute.

^{© 2017} Westerdijk Fungal Biodiversity Institute. Production and hosting by ELSEVIER B.V. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

Fungal taxa placed in "Diaporthaceae sensu lato" were divided into two groups (von Höhnel 1917) as "Eu-Diaportheen", to accommodate genera without allantoid ascospores and "Valseen" to accommodate genera with allantoid ascospores. Nannfeldt (1932) introduced the order Diaporthales to accommodate von Höhnel's Eu-Diaportheen group. Luttrell (1951) described Diaporthales as an order comprising species that have a "Diaporthe-type centrum" and "Endothia-type ascus". Chadefaud (1960) analysed characters of stromatic tissues in diaporthoid taxa and recognised families as Diaporthaceae or Cytosporaceae (= Valsaceae), Melanconidaceae and Gnomoniaceae. Wehmeyer (1975) classified the Diaporthales to include three families: Diaporthaceae, Gnomoniaceae and Cytosporaceae. Barr (1978) revised the order Diaporthales accepting Gnomoniaceae and Cytosporaceae in the suborder Gnomoniineae. Melanconidaceae and Pseudovalsaceae were accommodated in the suborder Melanconidineae. To differentiate genera, Barr (1978) used characters such as presence or absence of stromata, stromatic development and tissue types, the position of the perithecia and perithecial necks relative to the substrate, as well as ascospore shape; and Monod (1983) distinguished genera within Gnomoniaceae based on characters of the stromatic tissues, asexual morphs and ascospores. Three families were recognised in Diaporthales by Eriksson (2001), including Cytosporaceae, Melanconidaceae and Vialaeaceae. Based on analysis of LSU nrDNA sequence data, Castlebury et al. (2002) accepted Diaporthaceae, Gnomoniaceae, Melanconidaceae and Cytosporaceae in Diaporthales. Gnomoniaceae was revised by several recent studies and new taxa were introduced (Sogonov et al. 2008, Walker et al. 2010, 2012, Mejía1 et al. 2011). Castlebury et al. (2002) did not confirm Vialaeaceae as a family in Diaporthales and therefore excluded it from Diaporthales. Réblová et al. (2004) introduced Togniniaceae to this order based on small subunit (SSU) nrDNA; however, Mostert et al. (2006) concluded that its placement was ambiguous based on large subunit (LSU) nrDNA. Maharachchikumbura et al. (2015) excluded Togniniaceae from Diaporthales and accommodated it in Togniniales. Gryzenhout et al. (2006c) introduced the Cryphonectria-Endothia complex as the family Cryphonectriaceae. Sydowiellaceae and the Schizoparme-Pilidiella complex with the genus Coniella were introduced as Schizoparmaceae in Diaporthales (Rossman et al. 2007, Alvarez et al. 2016). Harknessiaceae was introduced into Diaporthales accommodating Harknessia with wuestneia-like sexual morphs (Crous et al. Pseudoplagiostomaceae was introduced 2012b). by Cheewangkoon et al. (2010) to accommodate Pseudoplagiostoma. Voglmayr & Jaklitsch (2014) resurrected Stilbosporaceae in Diaporthales based on phylogenetic analysis of LSU nrDNA sequence data and transferred the genera Stegonsporium and Stilbospora to this family. Macrohilaceae was introduced by Crous et al. (2015), based on an analysis of LSU nrDNA to accommodate Macrohilum. Suetrong et al. (2015) introduced Tirisporellaceae into Diaporthales; however, Jones et al. (2015) excluded this family from Diaporthales. Norphanphoun et al. (2016) introduced Lamproconiaceae to accommodate Lamproconium and Hercospora. Juglanconidaceae was introduced in the Diaporthales by Voglmayr et al. (2017). However, molecular data suggest that additional families still remain to be elucidated

(Gryzenhout *et al.* 2006c, Crous *et al.* 2012a, 2015, Voglmayr *et al.* 2017). Currently there are 14 families accepted in the *Diaporthales*.

Given the taxonomic discrepancies within *Diaporthales*, the present study uses a combined taxonomic approach based on morphology and DNA sequence analyses of the partial 28S nrDNA (LSU), the internal transcribed spacer regions and intervening 5.8S nrDNA (ITS), DNA-directed RNA polymerase II second largest subunit (*rpb2*), and translation elongation factor 1-alpha (*tef1*) gene regions to investigate phylogenetic relationships of all genera in *Diaporthales* to update their classification. All taxonomic novelties and present taxonomic families are redescribed and illustrated where necessary. We also present new data on each family to provide a better taxonomic understanding.

MATERIALS AND METHODS

Isolates and specimens

Specimens were collected from Germany, Italy, Russia, Thailand and the UK. They were placed in paper bags and collection details noted. Specimens were brought to the laboratory in Ziplock plastic bags and examined with a Motic SMZ 168 stereomicroscope. Rehydrated fruiting bodies were used to observe morphological characteristics of ascomata, asci, ascospores and other tissues and characters were photographed with a Canon 550D digital camera fitted to the Nikon ECLIPSE 80i compound microscope. Photomicrographs were arranged with Adobe Photoshop v. CS6 and all measurements were made with Tarosoft v. 0.9.0.7. Specimens were preserved and are deposited at the BBH and MFLU fungaria. Taxonomic novelties and descriptions were deposited in MycoBank (Crous et al. 2004), and new species were established using modern criteria and standards (Taylor et al. 2000, Seifert & Rossman 2010, Jeewon & Hyde 2016).

Sporocarps were removed from the substrate using a sterilised needle and placed in a few drops of sterilised distilled water on a sterilised cavity slide and a spore suspension was prepared as described in Chomnunti *et al.* (2014). Germinating ascospores were aseptically transferred to Petri dishes containing Potato Dextrose Agar (PDA) or Malt Extract Agar (MEA) (Crous *et al.* 2009). Colonies were photographed and characters were noted. Colony colour on PDA and MEA was determined with the colour charts of Rayner (1970). Living cultures are deposited at Mae Fah Luang University (MFLU) and the Westerdijk Fungal Biodiversity Institute (CBS) culture collections. Autoclaved pine needles were placed on water agar (WA) to observe conidiomatal development and sporulating (Crous *et al.* 2009).

Types and other relevant authentic specimens were loaned from accessible fungaria [New York State Museum (NY), Naturhistorisches Museum Wien (W), Swedish Museum of Natural History (S), Royal Botanic Gardens, Kew (K), Universität Wien (WU)]. A small part of the fungarium specimen was cut and rehydrated in water or 5 % KOH. Micro-morphological characters were observed from rehydrated ascomata and photography was done as previously described.

DNA extraction, PCR amplification and phylogeny

Fresh fungal mycelia grown on MEA for 4 wk at 18 °C was scraped from the colony margin and sometimes perithecial content of fresh specimens were used for genomic DNA extraction following the protocol outlined by Jeewon *et al.* (2002). PCR amplification and sequencing of the LSU nrDNA region using the primer pair LROR/LR5 (Vilgalys & Hester 1990, Rehner & Samuels 1994), ITS nrDNA region using primer pair ITS5/ITS4 (White *et al.* 1990), *rpb2* region using the primer pair fRPB2-5F/ fRPB2-7cR (Liu *et al.* 1999), and *tef1* region using primer pair EF1-728F/EF1-986R (Carbone & Kohn 1999) were performed.

Each amplification reaction contained 0.125 µL of 5 units/µL Ex-Tag DNA polymerase (TaKaRa), 2.5 µL of 10 × PCR buffer, 2 µL of 2 mM MgCl₂, 2.5 µL of 2 mM dNTPs, 1 µL of 0.2–1.0 µM primer. <500 ng DNA template and was adjusted with doubledistilled water to a total volume of 25 mL. Amplification reactions were performed in a thermal-cycler (BIORAD 1000™ Thermal Cycler, Bio-Rad Laboratories, Hercules, California). The temperature profile for both ITS nrDNA and LSU nrDNA 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 (Phillips et al. 2008). The temperature profile for the rpb2 was: initial denaturation at 94 °C for 120 s, 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 tef1 was: initial denaturation at 94 °C for 120 s, 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 PCR products with a DNA ladder were determined by electrophoresis at 120 V/cm for 20 min in 1 % agarose gel stained with ethidium bromide (0.5 mg/mL). The gel was visualised 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 condensed with DNASTAR Lasergene v. 7.1. 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).

Maximum likelihood analysis (ML) was performed by RAxMI 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 with 1000 replicates using the GTRGAMMAI model of nucleotide substitution, which was the best model predicted for the concatenated LSU nrDNA, ITS nrDNA, *rpb2* and *tef1* alignment by MrModeltest v. 2.3 (Nylander 2004).

For the Bayesian analyses (BI) of the individual loci and concatenated LSU nrDNA, ITS nrDNA, *rpb2* and *tef1* alignment, MrModeltest v. 2.3 (Nylander 2004) was used to determine the best nucleotide substitution model settings for MrBayes. A dirichlet state frequency was predicted for all four data partitions and GTR+I+G as best model for LSU nrDNA, ITS nrDNA, and *rpb2*; for *tef1* the best model was GTR+G. The heating parameter was set to 0.2 and trees were saved every 1 000 generations (Ronquist *et al.* 2012). The Markov Chain Monte Carlo (MCMC) analysis of four chains started in parallel from a random tree topology.

The maximum parsimony analysis (MP) was performed with PAUP v. 4.0b10 (Swofford 2003). Ambiguously aligned regions were excluded and all characters were unordered and given equal weight. Alignment gaps were treated as a fifth character state. Trees were inferred using the heuristic search option with TBR branch swapping and 100 random sequence additions. MaxTrees were set 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 (-In L) were calculated for trees generated under different optimality criteria. The robustness of the most parsimonious trees was evaluated by 1 000 bootstrap replications resulting from the maximum parsimony analysis, each with 10 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 different optimality criteria were significantly different.

Trees were viewed in FigTree v. 1.4.3 (Rambaut 2012). The final alignments and the trees obtained were deposited in TreeBASE (http://purl.org/phylo/treebase/phylows/study/TB2: S21148) and are available under study accession no. S21148.

RESULTS

To reveal the phylogenetic position of genera, families and genera *incertae sedis* within the order *Diaporthales*, a phylogenetic analysis was performed with LSU nrDNA, ITS nrDNA, *rpb2* and *tef1* sequence data. Sequences of representative species were selected from Maharachchikumbura *et al.* (2016), Norphanphoun *et al.* (2016), VogImayr *et al.* (2017) and supplemented with sequences from GenBank. The LSU nrDNA, ITS nrDNA, *rpb2*, *tef1* and combined data matrices contained 1423, 735, 1064, 427 and 3 652 characters with gaps, respectively. The alignment comprised 310 strains and *Eutypella* sp. (MFLUCC 16–1215) was selected as the outgroup.

The same concatenated alignment was subjected to phylogenetic analyses, including a Bayesian analysis, a maximum parsimony analysis and a maximum likelihood analysis. The concatenated sequence alignment contained 2027 parsimonyinformative characters, 385 were variable and parsimony uninformative and 1241 were constant. The parsimony analysis vielded the maximum of 1 000 equally most parsimonious trees (TL = 16 973 steps; CI = 0.278; RI = 0.728; RC = 0.202; HI = 0.722). The ML analysis yielded a tree with a likelihood value of In: -75295.054554 and the following model parameters: alpha: 0.368178; Π(A): 0.246723, Π(C): 0.249231, Π(G): 0.277805, and II(T): 0.226241. The Bayesian analysis lasted 72 151 000 generations (average standard deviation of split frequencies value = 0.016671) and the consensus trees and posterior possibilities were calculated from the 103 301 trees in each of the two run files, of which a total of 72 152 trees in each of the two run files, of which a total of 108 228 were sampled after discarding the first 25 % of generations for burn-in. The different data partitions contained 787, 529, 761 and 390 unique site patterns (LSU nrDNA, ITS nrDNA, rpb2 and tef1, respectively).

The phylogeny resulting from the analysis of combined gene sequence data is shown in Fig. 1. Overall, the topologies obtained from the different phylogenetic analyses were mostly

| Table 1. Details of the strains included for molecular and/or morphological study. | | | | | | | |
|--|-----------------|---------------|---------------------------------------|----------|-----------|-----------|----------|
| Fungal species | Culture | Specimen | Host/substrate | Gen | Bank acce | ssion num | nbers |
| | accession no. | voucher no. | | ITS | LSU | tef1-a | rpb2 |
| Alnecium auctum | CBS 124263 | WU 30206 | Alnus glutinosa | KF570154 | KF570154 | KF570200 | KF570170 |
| Ambarignomonia petiolorum | CBS 121227 | BPI 844274 | Liquidambar styraciflua | EU254748 | EU255070 | EU221898 | EU219307 |
| | CBS 116866 | BPI 843530 | Liquidambar styraciflua | EU199193 | AY818963 | _ | EU199151 |
| Amphilogia gyrosa | CBS 112922 | AFTOL-ID 1985 | Elaeocarpus dentatus | _ | FJ176889 | _ | FJ238374 |
| | YMJ 91123101 | HAST 91123101 | Elaeocarpus japonicus | EF026147 | _ | KC465404 | _ |
| | CMW 10470 | - | Elaeocarpus japonicus | _ | AY194108 | _ | - |
| Amphiporthe hranicensis | CBS 119289 | BPI 843515 | Tilia platyphyllos | EU199178 | EU199122 | _ | EU199137 |
| Anisogramma anomala | 529478 | - | Corylus avellana | EU683064 | EU683066 | _ | - |
| Anisogramma virgultorum | 529479 | - | Betula pendula | EU683062 | EU683065 | _ | - |
| Apiognomonia errabunda | AR 2813 | AR 2813 | Fagus sylvatica | DQ313525 | NG027592 | DQ313565 | DQ862014 |
| Apiognomonia veneta | MFLUCC 16-1193 | MFLU 17-0896 | Platanus acerifolia | MF190114 | MF190056 | _ | _ |
| | MFLUCC 17-1656 | MFLU 17-0896B | Platanus acerifolia | MF190115 | MF190057 | _ | - |
| | CBS 897.79 | Monod LAU | Platanus orientalis | _ | EU255195 | EU221910 | EU219259 |
| Apioplagiostoma populi | ID 858501 | ApLA2 | Populus tremuloides | KP637024 | _ | _ | _ |
| Apiosporopsis carpinea | CBS 771.79 | _ | Carpinus betulus | _ | AF277130 | _ | _ |
| Apiosporopsis sp. | Masuya 11Af2-1 | _ | Alnus firma | _ | AB669034 | _ | _ |
| Apoharknessia insueta | CBS 111377 | CPC 1451 | Eucalyptus pellita | JQ706083 | AY720814 | _ | _ |
| | CBS 114575 | CPC 10947 | Eucalyptus pellita | JQ706082 | AY720813 | _ | _ |
| Ascitendus austriacus | CBS 131685 | - | Decayed driftwood of Alnus glutinosa | - | AF261067 | - | JQ429257 |
| | CBS 102665 | - | Submerged stems of Fagus sylvatica | AF242263 | AF242263 | - | - |
| Asteroma alneum | CBS 109840 | _ | Alnus glutinosa | EU167609 | EU167609 | _ | _ |
| Asteroma sp. | Masuya 8Ah9-1 | _ | Alnus hirsuta | _ | AB669035 | _ | _ |
| Asterosporium asterospermum | - | MFLU 15-3555 | Fagus sylvatica | - | MF190062 | - | MF377615 |
| | KT 2125 | HHUF 30038 | Fagus crenata | _ | AB553743 | _ | _ |
| | CBS 112404 | _ | Fagus sylvatica | _ | AB553745 | _ | _ |
| | KT 2101 | HHUF 30037 | Fagus crenata | _ | AB553742 | _ | _ |
| | KT 2138 | HHUF 30039 | Fagus crenata | _ | AB553744 | _ | _ |
| Aurantioporthe corni | MNA 1003 | _ | Cornus alternifolia | KF495043 | KF495058 | _ | - |
| | SDS 1001 | - | Cornus alternifolia | KF495046 | KF495061 | _ | - |
| Aurantiosacculus acutatus | CPC 13704 | CBS H-20933 | Eucalyptus viminalis | JQ685514 | JQ685520 | _ | - |
| Aurantiosacculus eucalyptorum | CPC 13229 | - | Eucalyptus globulus | JQ685515 | JQ685521 | _ | - |
| Aurapex penicillata | CMW 11296 | - | Myrica faya | AY214315 | AY194090 | _ | - |
| | CMW 10032 | - | Miconia theaezans | AY214312 | AY194104 | _ | - |
| Auratiopycnidiella tristaniopsis | CBS 132180 | CBS H-20932 | Tristaniopsis laurina | JQ685516 | JQ685522 | - | - |
| Aurifilum marmelostoma | CBS 124930 | CMW 28288 | Terminalia ivorensis | FJ882856 | HQ730874 | _ | - |
| | CBS 124929 | PREM 60257 | Terminalia mantaly | FJ882855 | HQ730873 | _ | - |
| Brachysporium nigrum | MR 1346 | - | - | - | KT991662 | _ | KT991652 |
| Cainiella johansonii | Kruys 727 (UPS) | - | Dryas sp. | JF701922 | _ | _ | - |
| | Kruys 731 (UPS) | - | Dryas octopetala | JF701922 | JF701920 | _ | - |
| Calosphaeria pulchella | CBS 115999 | JF 3200 | Prunus sp. | EU367451 | AY761075 | _ | GU180661 |
| Celoporthe dispersa | CMW 9978 | _ | Syzygium cordatum | AY214316 | HQ730854 | HQ730841 | - |
| Celoporthe eucalypti | CMW 26913 | - | Eucalyptus EC48 clone | HQ730839 | HQ730865 | HQ730852 | _ |
| Chaetoconis polygoni | - | MFLU 17-0965 | Rumex acetosa | - | MF190063 | _ | - |
| | CBS 405.95 | _ | Polygonum sachalinense | - | EU754141 | - | - |
| Chapeckia nigrospora | CBS 125532 | BPI 863766 | <i>Betula</i> sp. | JF681957 | EU683068 | - | - |

| Table 1. (Continued). | | | | | | | |
|----------------------------|----------------|-----------------|-------------------------|-------------------------|----------|---------------|--------------|
| Fungal species | Culture | Specimen | Host/substrate | GenBank accession numbe | | | nbers |
| | accession no. | voucher no. | | ITS | LSU | tef1-a | rpb2 |
| Chiangraiomyces bauhiniae | MFLUCC 17-1669 | MFLU 17-0964 | Bauhinia sp. | MF190119 | MF190064 | MF377598 | MF377603 |
| | MFLUCC 17-1670 | CHUNI 81 | <i>Bauhinia</i> sp. | MF190118 | MF190065 | MF377599 | MF377604 |
| Chromendothia citrina | AR 3445 | _ | Quercus mongolica | _ | EU255074 | EU222013 | EU219342 |
| | CBS 109758 | BPI 747935 | Quercus mongolica | _ | AF408335 | _ | _ |
| Chrysocrypta corymbiae | CBS 132528 | CPC 19279 | Corymbia sp. | JX069867 | JX069851 | _ | _ |
| Chrysofolia barringtoniae | TBRC 5647 | SDBR-CMUENBA048 | Barringtonia sp. | KU948046 | KU948045 | - | - |
| Chrysofolia colombiana | CPC 24986 | CBS 139909 | Eucalyptus urophylla | KR476738 | KR476771 | - | _ |
| Chrysoporthe cubensis | CMW 14394 | _ | Eucalyptus sp. | JN942342 | JN940856 | GQ290137 | _ |
| Chrysoporthella hodgesiana | CMW 10641 | CBS 115854 | Tibouchina semidecandra | AY692322 | - | - | - |
| Coniella africana | CBS 114133 | CBS H-22706 | Eucalyptus nitens | AY339344 | AY339293 | KX833600 | KX833421 |
| Coniella australiensis | IMI 261318 | BPI 748425 | Leaf litter | AF408336 | AF408336 | KX833692 | KX833497 |
| Coniella crousii | NFCCI 2213 | AMH 9406 | Terminalia chebula | HQ264189 | - | - | - |
| Coniella fragariae | CBS 110394 | RMF 74.01 | Forest soil | KJ710463 | KJ710441 | KX833695 | KX833499 |
| | CBS 172.49 | STE-U 3930 | <i>Fragaria</i> sp. | AY339317 | AY339282 | AY339352 | - |
| Coniella koreana | CBS 143.97 | CBS H-22710 | - | KX833584 | AF408378 | KX833684 | KX833490 |
| Coniella pseudokoreana | MFLUCC 12-0427 | MFLU 13-0282 | - | MF190145 | - | - | - |
| | MFLUCC 17-1673 | MFLU 13-0282B | - | MF190146 | - | - | - |
| Coniella pseudostraminea | CBS 814.71 | IMI 233050 | <i>Fragaria</i> sp. | KX833582 | - | KX833682 | - |
| Coniella quercicola | CBS 283.76 | - | Eucalyptus nitens | AY339344 | AY339293 | AY339364 | - |
| Coniella straminea | CBS 149.22 | STE U 3932 | <i>Fragaria</i> sp. | AY339348 | AY339296 | AY339366 | KX833506 |
| Coniella tibouchinae | CPC 18511 | BECM1 | Tibouchina granulosa | JQ281774 | JQ281776 | JQ281778 | KX833507 |
| Coniella wangiensis | CPC 19397 | CPC 19397 | Eucalyptus sp. | JX069873 | JX069857 | KX833705 | KX833509 |
| Coryneum longipes | AR 3541 | BPI 872021 | Quercus cerris | - | EU683072 | - | - |
| Coryneum modonia | AR 3558 | BPI 749131 | Castanea sativa | - | EU683073 | - | - |
| Coryneum umbonata | AR 3897 | BPI 843585 | Quercus cerris | - | EU683074 | - | - |
| Corynym arausiaca | MFLUCC 13-0658 | MFLU 17-0875 | Quercus sp. | MF190120 | MF190066 | MF377574 | MF377609 |
| | MFLUCC 15-1110 | BBH 42437 | Quercus sp. | MF190121 | MF190067 | MF377575 | MF377610 |
| Crinitospora pulchra | CBS 138014 | CBS H-21729 | Mangifera indica | KJ710466 | KJ710443 | - | - |
| Cryphonectria parasitica | ATCC 38755 | - | Castanea dentata | AY141856 | EU199123 | EU222014 | - |
| | AFTOL ID 2123 | ATCC 38755 | - | - | - | DQ862033 | DQ862017 |
| Cryptodiaporthe aesculi | AR 3580 | BPI 748430 | Aesculus hippocastanum | EU199179 | AF408342 | - | EU199138 |
| | CBS 109765 | AFTOL-ID 1238 | Aesculus hippocastanum | - | DQ836905 | - | DQ836892 |
| Cryptometrion aestuescens | CMW 18790 | PREM 60249 | Eucalyptus grandis | GQ369458 | HQ730869 | - | - |
| | CMW 18793 | - | Eucalyptus grandis | GQ369459 | HQ730870 | - | - |
| Cryptosporella hypodermia | CBS 116866 | BPI 748432 | Ulmus minor | EU199181 | AF408346 | - | EU199140 |
| Cryptosporella suffusa | CBS 121077 | BPI 871231 | Alnus incana | EU199184 | EU199124 | - | EU199142 |
| Cytospora ambiens | ATCC 52280 | ATCC 52280 | Acer rubrum | AY347345 | AF277146 | - | - |
| Cytospora austromontana | Willow21 | - | - | KM669911 | - | KM669767 | - |
| Cytospora carbonacea | CFCC 50056 | - | Ulmus pumila | KP281263 | KP310809 | KP310852 | - |
| Cytospora cedri | CBS 196.50 | - | - | AF192311 | - | JX438575 | - |
| Cytospora centrivillosa | MFLUCC 16-1206 | MFLU 17-0887 | Sorbus domestica | MF190122 | MF190068 | - | MF377600 |
| | MFLUCC 17-1660 | BBH 42449 | Sorbus domestica | MF190123 | MF190069 | - | MF377601 |
| | - | MFLU 17-0999 | Sorbus domestica | MF190124 | MF190070 | - | - |
| Cytospora ceratosperma | AR 3426 | - | - | - | EU255209 | - | - |
| Cytospora chrysosperma | CFCC 89630 | - | Salix psammophila | KF765674 | KF765690 | - | KF765706 |
| Cytospora fraxinigena | MFLUCC 14-0868 | BBH 42442 | Fraxinus ornus | MF190133 | MF190078 | - | - |
| | - | MFLU 17-0880 | Fraxinus ornus | MF190134 | MF190079 | - | - |
| | | | | | | (continued of | n next page) |



| Table 1. (Continued). | | | | | | | | |
|------------------------|----------------|--------------|--------------------------|--------------------------|----------|----------|----------|--|
| Fungal species | Culture | Specimen | Host/substrate | GenBank accession number | | | bers | |
| | accession no. | voucher no. | | ITS | LSU | tef1-α | rpb2 | |
| Cytospora germanica | CXY 217 | _ | Populus sp. | JQ086564 | JX524618 | | _ | |
| Cytospora hippophaes | CFCC 89640 | - | Hippophae rhamnoides | KF765682 | KF765698 | KP310865 | KF765714 | |
| Cytospora junipericola | _ | BBH 42444 | Juniperus communis | MF190126 | MF190071 | MF377579 | _ | |
| | _ | MFLU 17-0882 | Juniperus communis | MF190125 | MF190072 | MF377580 | _ | |
| Cytospora mali | CFCC 50044 | - | Malus baccata | KR045637 | KR045717 | _ | _ | |
| Cytospora malicola | SXFX-V2 | - | Malus pumila | GU174579 | _ | JQ900335 | _ | |
| Cytospora melanodiscus | Jimslanding2 | - | Alnus tenuifolia | JX438621 | _ | JX438605 | _ | |
| | Worrall2b | - | Alnus tenuifolia | JX438620 | _ | JX438606 | _ | |
| Cytospora melnikii | MFLUCC 16-0635 | T 1104 | Populus nigra | KY417736 | KY417770 | _ | _ | |
| Cytospora nivea | CFCC 89643 | - | Salix psammophila | KF765685 | KF765701 | _ | KF765717 | |
| Cytospora punicae | CBS 199.50 | | Punica granatum | JX438622 | _ | JX438568 | - | |
| Cytospora quercicola | MFLUCC 14-0867 | BBH 42443 | Quercus sp. | MF190129 | MF190073 | - | - | |
| | _ | MFLU 17-0881 | Quercus sp. | MF190128 | MF190074 | - | - | |
| Cytospora ribis | CFCC 50027 | - | Ulmus pumila | KP281268 | KP310814 | KP310857 | - | |
| Cytospora rosae | MFLUCC 14-0845 | MFLU 17-0885 | Rosa canina | MF190131 | MF190075 | - | - | |
| | MFLUCC 17-1664 | BBH 42447 | Rosa canina | MF190130 | MF190076 | - | - | |
| Cytospora sacculus | CFCC 89625 | - | Juglans regia | KR045646 | KR045725 | KP310861 | - | |
| Cytospora salicina | MFLUCC 16-0637 | T-1017 | Salix fragilis | KY417751 | KY417785 | - | - | |
| | MFLUCC 16-1190 | MFLU 17-1655 | Cornus sanguinea | MF190132 | MF190077 | - | - | |
| Cytospora sordida | HMBF 159 | - | Juglans regia | KF225613 | KF225627 | - | - | |
| Cytospora sp. | CMON41 | - | Phaseolus vulgaris | JQ753989 | JQ754081 | - | - | |
| Cytospora translucens | CZ320 | - | - | FJ755269 | FJ755269 | - | - | |
| Diaporthe azadirachtae | TN 01 | - | Azadirachta indica | KC631323 | - | - | - | |
| Diaporthe cassines | CBS 136440 | CPC 21916 | Cassine peragua | KF777155 | KF777208 | KF777244 | - | |
| Diaporthe cynaroidis | 150e | - | Myrtus communis | KC959207 | - | - | - | |
| | CBS 122676 | CMW 22190 | Protea cynaroides | KC343058 | EU552122 | - | - | |
| Diaporthe decedens | CBS 114281 | UPSC 2957 | Corylus avellana | KC343059 | AF408348 | - | - | |
| Diaporthe eres | MFLUCC 17-1667 | T400 | Fraxinus pennsylvanica | MF190137 | MF190080 | MF377594 | - | |
| | MFLUCC 17-1668 | MFLU 17-0890 | Fraxinus pennsylvanica | MF190138 | MF190081 | MF377595 | - | |
| | MFLUCC 14-0862 | Т98 | Catalpa bignonioides | MF190135 | MF190082 | MF377596 | - | |
| | MFLUCC 17-1661 | MFLU 17-0889 | Catalpa bignonioides | MF190136 | MF190083 | MF377597 | - | |
| | AR 5193 | - | Ulmus sp. | KJ210529 | - | KJ210550 | - | |
| | PS57 | - | Glycine max | JF430494 | JF704176 | - | - | |
| Diaporthe eucalyptorum | MFLUCC 12-0306 | - | Leaf litter | KT459419 | - | KT459453 | - | |
| Diaporthe litoricola | MFLUCC 16-1195 | BBH 42436 | Stem of sea-shore plant | MF190139 | MF190086 | - | - | |
| | MFLUCC 17-1657 | MFLU 17-0894 | Stem of sea-shore plant | MF190140 | MF190087 | - | - | |
| Diaporthe maytenicola | CBS 136441 | CPC 21896 | Maytenus acuminata | KF777157 | KF777210 | - | - | |
| Diaporthe nobilis | Napa911 | - | - | KM669958 | - | KM669814 | - | |
| Diaporthe rudis | - | IT 1526 | Acer campestre | MF190141 | MF190088 | MF377576 | - | |
| | - | MFLU 17-0895 | Acer campestre | MF190142 | MF190089 | MF377577 | - | |
| | MFLUCC 16-1197 | BBH 42452 | Umbelliferous stem | MF190143 | MF190085 | - | - | |
| | MFLUCC 17-1658 | MFLU 15-2661 | Umbelliferous stem | MF190144 | MF190084 | - | - | |
| | LC6147 | - | Dendrobenthamia japonica | KY011890 | KY011864 | KY011901 | - | |
| | BPI 748231 | - | - | - | AF362560 | - | - | |
| | CBS 113201 | CBS H-7950 | Vitis vinifera | KC343234 | - | KC343960 | - | |
| Diaporthella corylina | CBS 121124 | BPI 871218 | Corylus sp. | KC343004 | - | - | - | |
| Diaporthella sp. | CN5 | - | Corylus avellana | KP205483 | - | KP205456 | - | |
| | CN13 | - | Corylus avellana | KP205484 | - | KP205457 | - | |
| Dicarpella dryina | ICMP 14042 | - | Quercus sp. | KC145909 | - | KC145954 | - | |

| Table 1. (Continued). | | | | | | | | |
|------------------------------|----------------|---------------|-------------------------|-------------------|----------|---------------|--------------|--|
| Fungal species | Culture | Specimen | Host/substrate | GenBank accession | | | numbers | |
| | accession no. | voucher no. | | ITS | LSU | tef1-α | rpb2 | |
| | ICMP 14043 | _ | Quercus ilex | KC145858 | _ | KC145955 | _ | |
| Diplodina microsperma | CBS 114545 | CPC 2336 | Protea sp. | JN712461 | JN712525 | - | _ | |
| Discula destructiva | CBS 109771 | BPI 1107757 | Cornus nuttallii | EU199186 | AF408359 | - | EU199144 | |
| | MD 254 | BPI 1107741 | Cornus florida | AF429741 | AF429721 | AF429732 | - | |
| Disculoides eucalypti | CPC 17650 | - | Eucalyptus sp. | JQ685517 | JQ685523 | - | - | |
| Disculoides eucalyptorum | CBS 132184 | CPC 17648 | Eucalyptus viminalis | NR120090 | - | - | - | |
| Ditopella biseptata | - | MFLU 17-0884B | Alnus glutinosa | MF190147 | MF190091 | - | MF377616 | |
| | - | MFLU 17-0884 | Alnus glutinosa | MF190148 | MF190090 | - | MF377617 | |
| Ditopella ditopa | CBS 109748 | BPI 748439 | Alnus glutinosa | DQ323526 | EU199126 | - | EU199145 | |
| Ditopellopsis sp. | CBS 121471 | BPI 872061 | Clethra alnifolia | EU254763 | EU255088 | EU221936 | EU219254 | |
| Dwiroopa lythri | AR 3383 | BPI 747560 | Lythrum salicaria | - | AF408364 | - | - | |
| Endothia gyrosa | CMW 2091 | CRY 1515 | Quercus palustris | AF046905 | AY194114 | - | - | |
| Endothiella gyrosa | CMW 10436 | AFTOL-ID 1223 | Quercus sp. | AF452117 | - | - | _ | |
| Erythrogloeum hymenaeae | CPC 18819 | - | Hymenaea courbaril | JQ685519 | JQ685525 | - | - | |
| <i>Eutypella</i> sp. | MFLUCC 16-1215 | BBH 42446 | Alnus cordata | MF190165 | MF190112 | MF377578 | MF377618 | |
| Foliocryphia eucalypti | CBS 124779 | CPC 12494 | Eucalyptus coccifera | GQ303276 | GQ303307 | - | - | |
| Gnomonia gnomon | CBS 829.79 | Monod 267 LAU | Populus sp. | AY818957 | AY818964 | EU221905 | - | |
| | CBS 199.53 | _ | Corylus avellana | DQ491518 | AF408361 | EU221885 | EU219295 | |
| Gnomoniella fraxini | AR 3999 | BPI 843391 | Fraxinus americana | AY455814 | AY455818 | - | _ | |
| | AR 2793 | BPI 746411 | Fraxinus americana | AY455813 | AY455817 | - | _ | |
| Gnomoniopsis agrimoniae | MFLUCC 14-0844 | MFLU 17-0888 | Agrimonia eupatoria | - | MF190093 | MF377585 | _ | |
| | MFLUCC 17-1662 | BBH 42450 | Agrimonia eupatoria | - | MF190092 | MF377586 | _ | |
| Gnomoniopsis alderdunensis | CBS 125680 | BPI 879186 | Rubus parviflorus | GU320825 | _ | _ | _ | |
| Gnomoniopsis chamaemori | CBS 803.79 | Monod 345 LAU | Rubus chamaemorus | EU254808 | EU255107 | _ | _ | |
| Gnomoniopsis racemula | AR 3892 | BPI 871003 | Epilobium angustifolium | EU254841 | EU255122 | EU221889 | EU219241 | |
| Greeneria saprophytica | MFLUCC 12-0298 | MFLU 13-0255 | Syzygium cumini | KJ021933 | KJ021935 | - | - | |
| Greeneria uvicola | FI1 2007 | _ | Vitis sp. | HQ586009 | GQ870619 | - | _ | |
| | FI1 2008 | _ | Vitis sp. | HQ586010 | GQ870620 | _ | _ | |
| Hapalocystis berkeleyi | AR 3851 | _ | _ | _ | EU683069 | _ | _ | |
| | MFLUCC 13-0662 | IT 1187 | Platanus sp | _ | KP744486 | _ | _ | |
| Harknessia eucalypti | CBS 342.97 | _ | Eucalyptus regnans | AY720745 | AF408363 | _ | _ | |
| | CPC 13643 | _ | Eucalyptus regnans | JQ706089 | JQ706215 | _ | _ | |
| Harknessia karwarrae | CPC 10928 | _ | Eucalyptus botryoides | AY720748 | AY720841 | _ | _ | |
| Harknessia molokaiensis | CBS 114877 | _ | Eucalyptus robusta | AY720749 | AY720842 | _ | _ | |
| | CPC 19269 | _ | Eucalyptus cypellocarpa | JQ706127 | JQ706248 | _ | _ | |
| Harknessia weresubiae | CBS 113075 | _ | Eucalyptus sp. | AY720741 | AY720835 | _ | _ | |
| | CPC 5109 | _ | Eucalyptus sp. | AY720744 | AY720838 | _ | _ | |
| Hercospora tiliae | AR 3526 | _ | Tilia tomentosa | _ | AF408365 | _ | _ | |
| Holocryphia eucalypti | CBS 115852 | CMW 14545 | Eucalyptus sp. | JQ862840 | JQ862797 | JQ863037 | _ | |
| | CMW 7033 | PREM 56305 | Eucalyptus saligna | JQ862838 | JQ862795 | JQ863035 | _ | |
| Hyaliappendispora galii | MFLUCC 16-1208 | MFLU 17-0893 | Galium sp. | MF190150 | MF190095 | MF377587 | _ | |
| | MFLUCC 17-1761 | MFLU 17-0966 | Galium sp. | MF190149 | MF190094 | MF377588 | _ | |
| Hyalorostratum brunneisporum | A573 2b | ILL 40792 | _ | _ | HM191720 | _ | _ | |
| Immersiporthe knoxdaviesiana | CMW 37314 | PREM 60740 | Rapanea melanophloeos | JQ862770 | JQ862760 | _ | _ | |
| | CMW 37319 | PREM 60739 | Rapanea melanophioeos | JQ862765 | JQ862755 | _ | _ | |
| Juqlanconis iualandina | D96 | WU 35960 | Juglans regia | KY427145 | KY427145 | KY427214 | KY427195 | |
| J J | AR 3860 | WU 35959 | Juglanconis iuglandina | KY427149 | _ | KY427218 | KY427199 | |
| | CBS 121083 | BPI 843622 | Juglans regia | KY427148 | KY427148 | KY427217 | KY427198 | |
| Jualanconis oblonaa | MAFF 410216 | TFM FPH 2623 | Juglans ailanthifolia | KY427153 | KY427153 | KY427222 | KY427203 | |
| | | | g unununonu | | | (continued of | n next page) | |

| Table 1. (Continued). | Table 1. (Continued). | | | | | | | | |
|---------------------------------------|-----------------------|-----------------|-----------------------|-------------------------|---------------|----------------|-----------------|--|--|
| Fungal species | Culture | Specimen | Host/substrate | GenBank accession numbe | | | nbers | | |
| | accession no. | voucher no. | | ITS | LSU | tef1-α | rpb2 | | |
| | MAFF 410217 | TFM FPH 3599 | Juglans ailanthifolia | KY427154 | KY427154 | KY427223 | KY427204 | | |
| Juglanconis pterocaryae | MAFF 410079 | TFM FPH 3373 | Pterocarya rhoifolia | KY427155 | KY427155 | KY427224 | KY427240 | | |
| Lamproconium desmazieri | AR 3525 | BPI 748445 | <i>Tilia</i> sp. | - | AF408372 | - | _ | | |
| | MFLUCC 15-0870 | MFLU 15-1940 | <i>Tilia</i> sp. | KX430134 | KX430135 | MF377591 | MF377605 | | |
| | MFLUCC 14-1047 | MFLU 14-0780 | <i>Tilia</i> sp. | KX430132 | KX430133 | MF377592 | | | |
| | MFLUCC 15-0872 | MFLU 15-2111 | <i>Tilia</i> sp. | KX430139 | AF408372 | MF377593 | MF377606 | | |
| | MFLUCC 15-0873 | MFLU 15-2192 | <i>Tilia</i> sp. | KX430140 | KX430141 | _ | _ | | |
| | MFLUCC 15-0871 | MFLU 15-2037 | <i>Tilia</i> sp. | KX430136 | KX430137 | _ | _ | | |
| Lanspora coronata | AFTOL-ID 736 | JK 5839A | - | _ | U46889 | _ | DQ470899 | | |
| Lasmenia sp. | CBS 124122 | LMS 2011b | Nephelium lappaceum | GU797405 | JF838337 | _ | _ | | |
| | CBS 124123 | LMS 2011c | Nephelium lappaceum | GU797406 | JF838338 | _ | _ | | |
| | CBS 124124 | LMS 2011d | Nephelium lappaceum | JF838336 | JF838341 | _ | _ | | |
| | CBS 124125 | LMS 2011a | Nephelium lappaceum | GU797407 | JF838340 | _ | _ | | |
| Latruncellus aurorae | CBS 125526 | PREM 60348 | Galpinia transvaalica | HQ171209 | HQ171214 | _ | _ | | |
| | CBS 124904 | PREM 60349 | Galpinia transvaalica | GU726946 | HQ171213 | _ | _ | | |
| Leucostoma kunzei | ATCC 64881 | ATCC 64881 | Picea pungens | AY347320 | _ | JX438595 | _ | | |
| Luteocirrhus shearii | CBS 130776 | PERTH 08439362 | Banksia baxteri | KC197021 | KC197019 | _ | _ | | |
| | CBS 130775 | PERTH 08355312 | Banksia baxteri | KC197024 | KC197018 | _ | _ | | |
| Macrohilum eucalypti | CPC 10945 | _ | Eucalyptus sp. | DQ195781 | DQ195793 | _ | _ | | |
| ,, | CPC 19421 | CBS H-22279 | Eucalyptus piperita | KR873244 | KR873275 | _ | _ | | |
| Mamianiella coryli | _ | BPI 877578 | Corylus californica | EU254862 | _ | _ | _ | | |
| Marsupiomvces epidermoidea | _ | MFLU 15-2921 | Quercus robur | _ | MF190058 | _ | _ | | |
| · · · · · · · · · · · · · · · · · · · | _ | BBH 42451 | Quercus robur | _ | MF190059 | _ | _ | | |
| Marsupiomvces quercina | MFLUCC 13-0664 | MFLU 17-0876 | Quercus sp. | MF190116 | MF190061 | _ | _ | | |
| maroupromy ooo quoroma | MFLUCC 14-0566 | BBH 42438 | Quercus sp. | MF190117 | MF190060 | _ | _ | | |
| Mastigosporella anisophylleae | CPC 22461 | _ | Anisophyllea sp | KF779492 | KF777221 | _ | _ | | |
| | CBS 136421 | CBS H-21429 | Anisophyllea sp | NR137844 | _ | _ | _ | | |
| Mazzantia napelli | AR 3498 | BPI 748443 | Aconitum lycoctonum | _ | AF408368 | FU222017 | FU219345 | | |
| | AFTOL-ID 2126 | AR 3498 | _ | _ | _ | _ | DO862020 | | |
| Melanconiella chrysodiscosporina | MELUCC 17–1671 | MELU 16-1309 | Fagus sylvatica | ME190166 | _ | _ | _ | | |
| Melanconiella chrysomelanconium | _ | MFLU 17-0879 | Carpinus betulus | MF190167 | MF190113 | _ | MF377619 | | |
| Melanconiella ellisii | _ | BPI 878343 | Carpinus caroliniana | JQ926271 | _ | JQ926406 | JQ926339 | | |
| Melanconiella spodiaea | SPOD1 | WU 31854 | Carpinus betulus | JQ926301 | _ | - | JQ926367 | | |
| Melanconis alni | AR 3748 | BPI 872035 | Alnus viridis | FU199195 | FU199130 | _ | FU199153 | | |
| | AR 3500 | BPI 748444 | Alnus viridis | _ | AF408371 | FU221896 | EU100100 | | |
| Melanconis italica | MELUCC 17-1659 | MELU 15-1112 | Alnus cordata | ME190152 | MF190097 | _ | ME377602 | | |
| | MFLUCC 16-1199 | MFLU 17-0883 | Alnus cordata | MF190151 | MF190096 | _ | _ | | |
| Melanconis marginalis | AR 3442 | BPI 748446 | Alnus rubra | FI 1199197 | AF408373 | FI 1221991 | FU219301 | | |
| Melanconis stilbostoma | F01051 | _ | | AV57781/ | AV577813 | | | | |
| | E01051 | | | AV577811 | AV577810 | _ | _ | | |
| | AD 3548 | - | - Potula sp | AIJHOII | AT377010 | - | - | | |
| | AR 3540 | - BDI 7/823/ | Betula sp. | | AI 302307 | - EL1221886 | - EL 1210200 | | |
| Microascospora fragariae | 1 1 | DI 1 740234 | | - HM85/850 | | L0221000 | L0219299 | | |
| Microascospora Iragariae | 1.1 | - | - | | - | - | - | | |
| | 1.0 | _ | _ | | - | - | - | | |
| Microascospora rubi | 12 | | - Rubus ulmifolio | ME100154 | - ME100009 | - ME277501 | - ME377614 | | |
| ινιισι σαστοσμοί α ΤΟΝΙ | _ | BBH /2//2 | Rubus ulmifolio | ME100152 | ME100000 | ME277500 | ME277640 | | |
| Microthia havanansis | CMW/ 11208 | DDT 42440 | Fucelyntus solians | | AV104004 | WI 57750Z | | | |
| | 011111230 | | Lucarypius sanylia | TIZ14320 | 71134031 | - | - | | |

| Table 1. (Continued). | | | | | | | | |
|---------------------------------|---------------------------|-----------------|-----------------------|---------------|------------|----------------|---------------|--|
| Fungal species | Culture | Specimen | Host/substrate | Gen | Bank acce | ssion nun | nbers | |
| | accession no. voucher no. | | | ITS | LSU | tef1-a | rpb2 | |
| | CMW 38367 | _ | Psidium cattleianum | KJ027495 | _ | _ | _ | |
| Myrmecridium montsegurinum | JF 13180 | PRM 934684 | Fraxinus excelsior | KT991674 | KT991664 | _ | KT991654 | |
| Occultocarpon ailaoshanense | LCM 524.01 | BPI 879253 | Alnus nepalensis | JF779849 | JF779853 | _ | JF779856 | |
| | LCM 522.01 | BPI 879254 | Alnus nepalensis | JF779848 | JF779852 | JF779862 | JF779857 | |
| Ophiodiaporthe cyatheae | YMJ 1364 | HAST 1364 | Cyathea lepifera | JX570889 | JX570891 | KC465406 | JX570893 | |
| Ophiognomonia melanostyla | LCM 389.01 | BPI 879257 | Tilia cordata | JF779850 | JF779854 | _ | JF779858 | |
| Ophiognomonia vasiljevae | AR 4298 | BPI 877671 | Juglans nigra | EU254977 | EU255162 | EU221999 | EU219331 | |
| Ophiostoma gemellus | CMW 23059 | _ | Tarsonemus sp. | DQ821562 | DQ821533 | _ | _ | |
| Pachytrype princeps | Rogers s.n. | _ | - | _ | FJ532382 | _ | _ | |
| Pachytrype rimosa | FF1066 | _ | _ | _ | FJ532381 | _ | _ | |
| Papulosa amerospora | AFTOL-ID 748 | JK 5547F | _ | _ | DQ470950 | _ | DQ470901 | |
| Paradiaporthe artemisiae | MFLUCC 14-0850 | MFLU 12-2131 | Artemisia sp. | MF190155 | MF190100 | MF377583 | _ | |
| , | MFLUCC 17-1663 | BBH 42448 | Artemisia sp. | MF190156 | MF190101 | MF377584 | _ | |
| Phaeoacremonium aleophilum | CBS 631.94 | _ | Vitis vinifera | AF266647 | AB278175 | KF764643 | _ | |
| Phaeoacremonium vibratilis | CBS 117115 | BPI 2460 | Fagus sylvatica | KF764573 | DQ649065 | KF764645 | HQ878611 | |
| Phaeoappendispora thailandensis | MELUCC 13-0161 | MELU 17-0873 | Quercus sp. | MF190157 | MF190102 | _ | ME377613 | |
| | MELUCC 17-1762 | BBH 42435 | Quercus sp. | MF190158 | MF190103 | _ | ME377614 | |
| Phaeocytostroma ambiguum | CPC 17077 | _ | Zea mays | FR748041 | FR748100 | FR748073 | _ | |
| r naoooytootionia ambiguum | CPC 17078 | _ | Zea mays | FR748044 | FR748101 | FR748074 | _ | |
| Phaeodiaporthe appendiculata | CBS 123821 | WU 32449 | Acer campestre | KE570156 | KE570156 | _ | _ | |
| | CBS 123809 | WU 32448 | Acer campestre | KE570155 | KE570155 | _ | _ | |
| Phraamonorthe conformis | MELLICC 14-0567 | MELU 15-2662 | Alnus alutinosa | KU315388 | KU315389 | KU315391 | _ | |
| i magnioporato comorna | AR 3632 | BPI 748450 | Alnus rubra | _ | AF408377 | _ | _ | |
| Plagiostoma dilatatum | I CM 402 02 | BPI 878957 | Salix irrorata | GU367070 | _ | _ | GU367104 | |
| Plagiostoma ionesii | MELLICC 16-1189 | MELU 17-0878 | Limbelliferous stem | ME190159 | MF190104 | ME377589 | - | |
| r lagiottorna jonooli | MELUCC 17-1654 | BBH 42440 | Umbelliferous stem | MF190160 | MF190105 | ME377590 | _ | |
| Plagiostoma salicellum | CBS 109755 | BPI 843490 | Salix sp | EU255047 | _ | FU221912 | _ | |
| Plagiostoma salicicola | MELLICC 13-0656 | MELLI 17_0877 | Salix sp. | ME100161 | ME100106 | | _ | |
| r lagiostorna salicicola | MELLICC 17-1666 | BBH 42439 | Salix sp. | ME100162 | ME100107 | | | |
| Plaurocaras caprasa | CBS 372 69 | | Salix sp. | 1011 130 102 | AE2771/3 | | | |
| | AP /333 | - BDI 877710 | Salix sp. | - EU255060 | EL1255106 | - EU221031 | - EU210313 | |
| | CRS 006 70 | Monod 469/LALL | Salix sucheriss | EU255061 | EU255190 | EU221931 | EU219313 | |
| Pleuroceras tenellum | CBS 121082 | BDI 871050 | Acer rubrum | EU233001 | EU255197 | EU221902 | EU219311 | |
| Prosonidicola mevicana | CBS 113529 | CBS-H 79/8 | Prosonis alandulosa | AV720700 | | | E0100100 | |
| | CBS 113530 | C 158 | Prosopis glandulosa | AV720710 | | | | |
| Pseudoplagiostoma conumbiae | CPC 10287 | CBS H- 20057 | Conventia sp | 12060861 | - | | | |
| | CPS 124807 | CBS H 20303 | Eucolymbia sp. | GU073512 | CL 1073606 | - CLI073542 | - | |
| | CDC 1/161 | 00011-20303 | | CU073510 | CU073604 | CU073540 | - | |
| Decudenteriesterne eldii | CFC 14101 | - | | GU973510 | GU973004 | GU973540 | - | |
| rseudopiagiosionia oluli | CBS 113722 | - | | GU973535 | GU973010 | GU973503 | - | |
| Decudenteriesterne veriebile | CBS 124000 | CBS H-20300 | | GU973534 | GU973609 | GU973566 | - | |
| r seuuupiayiusiuilla Vallapile | | | Eucarypius globulus | RE0062E0 | GU3/3011 | GUSI JOOD | - | |
| Pusiulorinyces bambusicola | | | Dallipusa sp. | NF000/02 | NF0U0/03 | KL000122 | - | |
| Pyricularia potnriochioae | | UBS-H 21430 | | KF///186 | KF///238 | - | - | |
| Pyricularia oryzae | REA 8401 | | Opniuros exaltatus | км484916 | KM485022 | - | - | |
| Rossmania ukurunduensis | AK 3484 | BPI /4/566 | Acer ukurunduense | - | EU6830/5 | - | - | |
| Rostraureum tropicale | | PREM 5/519 | i erminalia ivorensis | AY16/436 | AY194092 | - | - | |
| | CMW 9975 | - | Terminalia ivorensis | AY167439 | - | - | - | |

(continued on next page)



| Table 1. (Continued). | | | | | | | |
|---|--------------------------|-----------------|-------------------------|---------------------------|---------------|----------|----------|
| Fungal species | Culture | Specimen | Host/substrate | GenBank accession numbers | | | bers |
| | accession no. | voucher no. | | ITS | LSU | tef1-α | rpb2 |
| Sillia ferruginea | AR 3440 | BPI 843619 | Corylus avellana | JF681959 | EU683076 | _ | _ |
| | CBS 126567 | BPI 843619 | Corylus avellana | JF681959 | EU683076 | - | - |
| Sirococcus tsugae | CBS 119626 | BPI 871167 | Tsuga mertensiana | EU199203 | EU199136 | EF512534 | EU199159 |
| | AR 4010 | - | Cedrus deodara | EF512478 | EU255207 | EU221928 | EU219289 |
| Sordaria fimicola | CBS 508.50 | - | Dung | AY681188 | AY681160 | - | DQ368647 |
| Stegonsporium acerophilum | CBS 117025 | WU 28050 | Acer saccharum | EU039982 | EU039993 | EU040027 | KF570173 |
| Stenocarpella macrospora | CBS 117560 | MRC 8615 | Zea mays | FR748048 | EU754219 | - | - |
| Stenocarpella maydis | CBS 117559 | MRC 8614 | Zea mays | FR748052 | DQ377937 | - | - |
| Stilbospora ellipsosporum | WJ 1840 | - | Carpinus betulus | - | AY616229 | - | - |
| Stilbospora macrosperma | CBS 121883 | - | Carpinus betulus | JX517290 | JX517299 | - | KF570196 |
| Sydowiella depressula | CBS 813.79 | - | Rubus sp. | EU552156 | - | - | - |
| Sydowiella fenestrans | CBS 125530 | BPI 843503 | Chamerion angustifolium | JF681956 | EU683078 | - | - |
| Sydowiella urticicola | MFLUCC 13-0665 | MFLU 13-0260 | Urtica dioica | - | MF190108 | - | - |
| | MFLUCC 17-1665 | BBH 42439 | Urtica dioica | - | MF190109 | - | - |
| Thyridium vestitum | AFTOL-ID 172 | OSC 100064 | - | - | AY544671 | - | DQ470890 |
| Tubakia seoraksanensis | CBS 127490 | - | Quercus mongolica | HM991734 | KP260499 | - | - |
| | BJFCCC140824-15 | - | - | KP260502 | KP260501 | - | - |
| Tubakia thailandensis | MFLUCC 12-0303 | MFLU 13-0260 | Decaying leaf | MF190163 | MF190110 | - | - |
| | MFLUCC 17-1672 | MFLU 13-0260B | Decaying leaf | MF190164 | MF190111 | - | - |
| Ursicollum fallax | CMW 18119 | PREM 58840 | Coccoloba uvifera | DQ368755 | EF392860 | - | - |
| Valsalnicola oxystoma | AR 4833 | BPI 884137 | Alnus viridis | JX519559 | JX519563 | - | - |
| | AR 5137 | BPI 884135 | Alnus tenuifolia | JX519561 | - | - | - |
| Valsella salicis | AR 3514 | BPI 748461 | Salix fragilis | - | EU255210 | EU222018 | EU219346 |
| Valseutypella multicollis Waydora typica | CBS 105.89 PDD 103894 | – PDD 103894 | Quercus ilex – | DQ243803 KF727412 | – KF727413 | - | - |

AFTOL: Assembling the Fungal Tree of Life culture collection; AMH: Ajrekar Mycological Herbarium, India; ATCC: American Type Culture Collection, Virginia, USA; BBH: National Science and Technology Development Agency, Thailand; BECM: British Empire and Commonwealth Museum, UK; BJFCCC: Beijing Forestry University, China; BPI: U.S. National Fungus Collections, Systematic Botany and Mycology Laboratory, USA; CBS: Westerdijk Fungal Biodiversity Institute, Utrecht, The Netherlands; CFCC: China Forestry Culture Collection Center, Beijing, China; CMW: Forestry and Agricultural Biotechnology Institute, University of Pretoria, South Africa; CPC: Culture collection of Pedro Crous, The Netherlands; FI: Museo di Storia Naturale dell'Universita, Italy; HAST: Herbarium, Biodiversity Research Center, Academia Sinica, Taiwan; HHUF: Hirosaki University, Japan; ICMP: International Collection of Micro-organisms from Plants, New Zealand; ILL: University of Illinois, USA; IMI: International Mycological Institute, Kew, UK; JF: Jonkershoek Forestry Research Centre, South Africa; LCM: University herbarium, Thailand; MFLUCC: Mae Fah Luang University Culture Collection, Thailand; MNA: Museum of Northern Arizona, USA; NFCCI: National Fungal Culture Collection, India; OSC: Oregon State University, Corvallis, USA; PERM: University of Perm, South Africa; PERTH: Western Australian Herbarium, Australia; PH: Academy of Natural Sciences of Drexel University, Philadelphia, PA; RMF: Rocky Mountain Herbarium, University of Wyoming, USA; SDSU: San Diego State University, USA; STE-U: University of Stellenbosch, Plant Pathology Department, South Africa; TBRC: Thailand Bioresource Research Center, Bangkok, Thailand; TFM: Forestry and Forest Products Research Institute, Matsunosato, Japan; UPSC: Fungal Culture Collection at the Botanical Museum, Uppsala University, Sweden; WU: Universitate Wien, Austria.

similar and the best scoring RAxML tree is illustrated here. The bootstrap support values of maximum likelihood analysis (MLB), maximum parsimony analysis (MPB) and Bayesian posterior probability scores (PP) are noted at the nodes. The separation of *Diaporthales* from other *Sordariomycetes* taxa is well-supported (MLB/MPB/PP = 100/96/1). The order separates into 21 familial clades with good support values and two genera *incertae sedis* clades labelled as 5 and 18.

Clade 1 is represented by *Gnomoniaceae* with moderate support values (MLB/MPB/PP = 60/-/0.9) and comprised Alnecium, Ambarignomonia, Amphiporthe, Anisogramma, Apiognomonia, Apioplagiostoma, Asteroma, Cryptodiaporthe, Cryptosporella, Discula, Ditopella, Ditopellopsis, Gnomonia, Gnomoniella, Gnomoniopsis, Mamianiella, Marsupiomyces, Occultocarpon, Ophiognomonia, Phragmoporthe, Plagiostoma,

Pleuroceras, Sirococcus and Valsalnicola. Anisogramma and Mamianiella formed a distinct clade with high support value. Morphologically they are similar and these genera appear to be congeneric. Therefore we synonymise Anisogramma under Mamianiella giving priority to the older name and its taxonomic stability. In addition, Mamianiella is nested in between Anisogramma species and this supports that both genera should be synonymised. Here we introduce one new genus and six new species to Gnomoniaceae. We introduce Plagiostoma salicicola based on morphology and phylogeny. Plagiostoma jonesii, the second species, was a fully-supported clade sister to Plagiostoma salicellum, P. dilatatum and P. salicicola. It is morphologically distinct from other species in Plagiostoma. Sequences of the asexual morph of Apiognomonia veneta (= Discula nervisequa) is included here and clustered with other Apiognomonia veneta strains. Ditopella biseptata is introduced based on phylogeny as well as morphology. Ditopella biseptata formed a fullysupported clade sister to *D. ditopa*. We introduce a new genus *Marsupiomyces* with two phylogenetically well-supported species, *M. quercina* and *M. epidermoidea*. Gnomoniopsis agrimoniae represents a new species distinct from other species of *Gnomoniopsis*.

Clade 2 is represented by *Melanconidaceae sensu stricto* with good support values (MLB/MPB/PP = 93/91/0.9) and it is considered as *Melanconidaceae*. Most genera listed under *Melanconidaceae* in recent publications are excluded from this family, based on morphology and available sequence data. We introduce a new *Melanconis* species as *M. italica*.

Clade 3 (MLB/MPB/PP = 100/92/1) represents the new family *Apiosporopsidaceae* which is introduced to accommodate a single genus, *Apiosporopsis*.

Clade 4 is represented by *Juglanconidaceae* (MLB/MPB/ PP = 99/94/1), which was recently introduced by Voglmayr *et al.* (2017) based on a fungal species isolated from *Juglans nigra*.

Clade 5 is represented by *Diaporthella* species with low bootstrap support and is considered here as *Diaporthales* genera *incertae sedis*, pending the availability of sequence data for the type species *Diaporthella aristata*.

The family *Cryphonectriaceae* is represented by Clade 6 which is phylogenetically poorly-supported (MLB/MPB/PP = 63/ –/0.9) but morphologically distinct from other families in *Diaporthales*. Almost all taxa in this family have molecular data.

Clade 7 is represented by the *Harknessiaceae*, which is phylogenetically poorly-supported (MLB/MPB/PP = 60/68/–). Species of *Harknessia* and wuestneia-like sexual morphs have been linked by morphological characteristics (Crous & Rogers 2001, Crous *et al.* 2012b), but the types of both genera have not been linked by molecular data. *Dwiroopa lythri* clusters basal to *Harknessia* species with low support values (MLB/MPB/PP = 60/68/–; Fig. 1).

Schizoparmaceae (Clade 8) is phylogenetically wellsupported (MLB/MPB/PP = 94/88/–). We introduce a new *Coniella* species as *Coniella* pseudokoreana and it clusters sister to *Coniella* straminea.

Clade 9 (MLB/MPB/PP = 91/71/1) represents the new family *Erythrogloeaceae* which is introduced in this study to accommodate *Chrysocrypta*, *Disculoides* and *Erythrogloeum*.

Clade 10 (MLB/MPB/PP = 93/-/1) currently encompassies 6 genera within the Melanconiellaceae. In particular, Melanconiella sisters to Microascospora where as Greeneria is nested in between Dicarpella and Tubakia. Microascospora is introduced here based on Microascospora rubi collected from Italy and Microascospora fragariae (= Sphaeronaemella fragariae) which was already placed in Microascales. Other genera were previously placed in Melanconidaceae, but phylogenetically they do not cluster with M. stilbostoma, which is the family type of the Melanconidaceae. Dicarpella and Tubakia form a distinct clade within this family to represent both genera as holomorphs. Greeneria saprophytica is distant from Greeneria uvicola, which is the type of this genus. We introduce a new Tubakia species as T. thailandensis. A new genus Microascospora based on M. rubi is introduced here and *M. rubi* strains have high support as a distinct species. Microascospora rubi forms a sister clade to Microascospora fragariae (= Sphaeronaemella fragariae) and high bootstrap support values confirmed it as a species. In addition. we include sequences of Melanconiella chrysodiscosporina and *M. chrysomelanconium* from recently collected specimens.

Clade 11 is represented by the monotypic family *Auratiopycnidiellaceae*, which is newly introduced in this study based on *Auratiopycnidiella tristaniopsis*, and it is morphologically and phylogenetically well-supported (MLB/MPB/PP = 95/–/1).

Clade 12 comprises the monotypic family *Pseudoplagiostomaceae* with full-support (MLB/MPB/PP = 100/100/1).

Clade 13 (MLB/MPB/PP = 100/68/1) represents Apoharknessiaceae to accommodate Apoharknessia and Lasmenia.

Clade 14 is represented by the family *Diaporthaceae* with good support (MLB/MPB/PP = 86/91/0.9). We also introduce several new genera such as *Chiangraiomyces* which is typified by *C. bauhiniae*; *Paradiaporthe*, typified by *P. artemisiae* and *Hyaliappendispora* typified by *H. galii. Paradiaporthe* is nested with reliable support between *Chiangraiomyces* and *Phaeocytostroma* while *Chiangraiomyces*, clusters sister to *Ophiodiaporthe*. *Hyaliappendispora* grouped sister to *Phaeodiaporthe*. Here we included several new isolates of *Diaporthe* species and *Diaporthe litoricola* forms a fully-supported clade sister to *Diaporthe maytenicola*. In addition, we include a new collection of *Diaporthe eres* and a new collection of *Diaporthe rudis*.

Clade 15 represents the family *Macrohilaceae* with full-support (MLB/MPB/PP = 100/100/0.9).

Clade 16 is represented by *Cytosporaceae* with good support (MLB/MPB/PP = 88/51/1). In addition to *Cytospora*, we include *Waydora* and *Pachytrype* in *Cytosporaceae* based on molecular data. Here we introduce five *Cytospora* species viz., *Cytospora centrivillosa*, *Cytospora fraxini*, *Cytospora junipericola*, *Cytospora quercicola*, and *Cytospora rosae*. *Cytospora centrivillosa* forms a distinct clade that is sister to *Cytospora melanodiscus* is morphologically quite different having 1-septate ascospores. Here we added sequences of *C. salicina* from freshly collected specimens.

Prosopidicolaceae (Clade 17) is introduced to accommodate *Prosopidicola mexicana*.

Phaeoappendicospora thailandensis (Clade 18) forms separate fully-supported (MLB/MPB/PP = 100/100/0.9) clade. It currently does not have a high affinity with any known family in *Diaporthales*, therefore we consider this species as *Diaporthales* genera *incertae sedis*.

Clade 19 represents the family *Stilbosporaceae* that is fullysupported (MLB/MPB/PP = 100/100/1). Although *Crinitospora* is morphologically different from *Stegonsporium* and *Stilbospora*, inclusion of this genus in the family is phylogenetically wellsupported.

Clade 20 comprises the family *Coryneaceae* and includes molecular data for *Coryneum arausiaca* collected from Italy. *Coryneum arausiaca* has high support (MLB/PP = 100/96/1) as a separate species.

Clade 21 represents the family *Sydowiellaceae* with very good support (MLB/MPB/PP = 98/79/1) and here we introduce a new species *Sydowiella urticicola*. *Sydowiella urticicola* clade received high support values.

Lamproconiaceae is represented by clade 22 and comprises *Lamproconium* and *Hercospora*.

The new family *Asterosporiaceae* (23) is introduced to accommodate *Asterosporium asterospermum*. This monogeneric family received high support (MLB/MPB/PP = 100/96/1) and is sister to the *Sydowiellaceae* and *Lamproconiaceae*.





Fig. 1. Consensus tree resulting from a maximum likelihood analysis of a combined LSU nrDNA, ITS nrDNA, *rpb2* and *tef1* sequence alignment for taxa of *Diaporthales* and other species in *Sordariomycetes*. Families are indicated in coloured blocks. RAxML bootstrap support values (MLB above 50 %), maximum parsimony bootstrap support values (MPB above 50 %) and Bayesian posterior probabilities (BPP above 0.90) are given at the nodes (MLB/MPB/BPP). The scale bar represents the expected number of changes per site. The tree is rooted to *Eutypella* sp (MFLUCC 16–1215). All the new sequences used in this study are in blue bold and type sequences are in black bold. The nodes that received maximum support (MLB/MPB/BPP = 100/100/1) are indicate by *



Fig. 1. (Continued).



Fig. 1. (Continued).



Fig. 1. (Continued).



Fig. 1. (Continued).



Fig. 1. (Continued).

Taxonomy

Diaporthales Nannf., Nova Acta R. Soc. Scient. Upsal. 8: 53. 1932.

Saprobic or pathogenic in plants, and animals, including humans or inhabiting soil. Sexual morph: Pseudostromata or ascostromata well-developed, poorly developed or absent, scattered, immersed or erumpent, solitary to aggregated, valsoid to diatrypoid, broadly elliptical, oval to circular from above, yellowish orange, pale brown, dark brown to black, some species turning purple or umber in KOH. Entostroma normally limited to the region near the perithecial walls, prosenchymatous, pale-coloured, and slightly differentiated from the surrounding bark tissue. Ectostromatic disc well- or poorly developed, subhvaline, vellowish white, pale brown, rarely dark brown to black, pulvinate, flat or slightly convex, orbicular, circular or somewhat irregular, with or without black zone or a crust consisting of fungus tissue, sclerotioid, coriaceous. Central column present or absent, if present beneath the disc more or less conical, comprising hyaline or pigmented hyphae mixed with a pigmented, cream, yellow, olive, brownish or grey, powdery amorphous substance. Ascomata perithecial, scattered, solitary or aggregated, immersed to erumpent, rarely superficial, globose to subglobose, sometimes circinate, arranged in a valsoid to diatrypoid configuration or single, coriaceous, sometimes with plate-like ornamentation around ostiole, black to brown, ostiolate, papillate. Papilla lacking or upright, long or short, one or more, central or eccentric, slanted to horizontal on host tissue, sometimes converging, with neck swollen at the tips, fuscous black to umber, ostiole with hyaline periphyses. Peridium thin or thick, comprising outer, dark, thickwalled, cells of textura angularis and inner, mostly small, hyaline, thin-walled, flattened cells of textura angularis. Hamathecium aparaphysate or comprising few broad cellular, filiform to cylindrical, septate to aseptate, branched to unbranched, hyaline paraphyses and sometimes parenchymatous cells attached at the base and asci dissolving at maturity. Asci generally 2-32-spored, unitunicate, ellipsoid, cylindrical, fusiform, clavate, oblong-clavate, broadly fusoid to cylindrical-fusoid, short pedicellate, apex blunt, usually with distinct, J- refractive ring. Ascospores overlapping

uniseriate, biseriate, partially biseriate to fasciculately arrange, ovoid, ellipsoid, oblong, fusoid, cylindrical, filamentous or allantoid, aseptate to multi-septate, rarely distoseptate, constricted or not at the septa, hyaline, olivaceous to brown, smooth- to sometimes ornamented walled, ends mostly rounded, rarely pointed, multi-guttulate, straight or curved, smooth- to sometimes ornamented walled to rarely ornamented, hyaline to dark brown. Appendages absent or present: if present, apical or basal, subulate, navicular or whip-shaped, smooth, hvaline, Asexual morph: Coelomycetous. Stroma present or absent, immersed to superficial, opening by irregular rupture, globose, subglobose to irregular, solitary to gregarious, orange, brown to dark brown, sometimes loculate. Conidiomata amphigenous, eustromatic, punctiform, pycnidial or acervular, sometimes pyriform in section and divided into compartments by bending of peridium, subcuticular, peridermal or subepidermal, brown to black or orange with dark brown border, sometimes with a central, well-developed, pale brown, pseudoparenchymatous layer, becoming thinner or absent at the margin of the conidiomata, sometimes with pale coloured, ectostromatic disc and central column or with radiate scutella. Scutella convex, membranous, brown, somewhat translucent, with a central hyaline or pale disc, giving rise to radiating hyphae, thick-walled cells radiating from a central point, rounded to pointed at the tips. Peridium comprising pale to dark brown cells of textura angularis to textura globulosa. Paraphyses present or absent. If present, hyaline, cellular, subcylindrical, branched or not, with obtuse apex, septate, constricted at septa. Conidiophores reduced to conidiogenous cells or arising from the upper most cells of basal and parietal tissue or under the developing scutellum, densely aggregated or few, filiform, fusiform, cylindrical to globose, simple or branched, septate or aseptate, sometimes septate only at the base, smooth, hyaline or hyaline at the top, pale brown at the base, sometimes dimorphic. Alpha conidiophores tightly aggregated, subcylindrical, branched in mid region, consisting of few supporting cells, giving rise to septate, ampulliform, cylindrical to irregular conidiogenous cells or paraphyses, straight to sinuous, septate, cylindrical, hyaline to pale brown, branched only at the base, smooth, formed from the innermost layer cells of the conidiomatal wall, sometimes with

terminal and lateral apex, with minute periclinal thickening and collarette. Beta conidiophores interspersed among alpha conidiophores, hyaline, subcylindrical, branched, septate. Conidiogenous cells lining the inner cavity of conidioma, enteroblastic to holoblastic, annellidic or phialidic, discrete or integrated, hyaline to olivaceous, smooth, lageniform, subcylindrical to ampulliform, with terminal truncate locus, simple or branched, proliferating several times percurrently near apex, with flaring collarettes or apex truncate, with minute periclinal thickening or terminal truncate locus. Conidia broadly ellipsoid, oval, obovoid, allantoid, fusoid to sigmoid, sinuate to slightly angular, hyaline to brown, hyaline when immature, becoming medium brown to dark brown at maturity, smooth-walled, guttulate, aseptate to septate or distoseptate, apex obtuse, base truncate with a visible scar or a flat protruding scar at the base, sometimes the apical and basal cell darker than other cells or with hyaline tip in apical cell, sometimes with or without a longitudinal germ slit, sometimes with marginal frill or becoming golden brown at germination, with solitary, brown, wavy germ tubes.

Notes: The order Diaporthales was introduced to accommodate "true" diaportheen taxa and Eriksson & Winka (1997) accommodated Diaporthales in Sordariomycetidae. Barr (1978), Monod (1983), Castlebury et al. (2002), Rossman et al. (2007), Maharachchikumbura et al. (2015, 2016), Rossman et al. (2015) and Voglmayr et al. (2017) clarified the taxonomic and phylogenetic concepts. Maharachchikumbura et al. (2015) introduced the subclass Diaporthomycetidae to accommodate the order Diaporthales. Morphologically and phylogenetically this is a wellsupported order comprising Apiosporopsidaceae, Apoharknessiaceae, Asterosporiaceae, Auratiopycnidiellaceae, Coryneaceae, Cryphonectriaceae, Cytosporaceae, Diaporthaceae, Erythrogloeaceae, Gnomoniaceae, Harknessiaceae, Juglanconidaceae, Lamproconiaceae, Macrohilaceae, Melanconidaceae, Melanconiellaceae, Prosopidicolaceae, Pseudoplagiostomaceae, Schizoparmaceae, Stilbosporaceae, and Sydowiellaceae.

Apiosporopsidaceae Senan., Maharachch. & K.D. Hyde, fam. nov. MycoBank MB821538. Facesoffungi number FoF03455. Clade 3.

Parasitic on living leaves and twigs. Sexual morph: Ascomata scattered, black, oval to almost spherical, immersed in the leaf tissue beneath a thin, well-developed clypeus, neck lacking or only slightly papillate, periphysate. *Peridium* comprises 5–6 outer layers of dark, thick-walled cells of *textura angularis* and inner, thin-walled, strongly flattened cells of *textura angularis*. *Hamathecium* aparaphysate. *Asci* 8-spored, unitunicate, short-pedicellate, apex blunt with J- apical ring. *Ascospores* 1–2-seriate, elliptical to fusoid, often slightly flattened on one side, unicellular, hyaline. Asexual morph: Coelomycetous. *Stroma* loculate, globose to irregular, sometimes with beaks. *Conidiogenous cells* phialidic, short to elongate, simple or branched. *Conidia* oblong or cylindrical to allantoid, 1-celled, hyaline.

Type genus: Apiosporopsis (Traverso) Mariani.

Type species: Apiosporopsis saccardoana Mariani.

Apiosporopsis carpinea (Fr.) Mariani, Atti Soc. ital. Sci. nat. (Modena) 50: 165. 1911. Facesoffungi number FoF03456. Fig. 2.

Basionym: Xyloma carpini Fr., Observ. mycol. (Havniae) 2: 363. 1818.

Illustration: For asexual morph see Potebnia (1910).

Saprobic on over-wintered plants. Sexual morph: Clypeus 70-140 µm wide, 50-70 µm high, slight, prosenchymatous. Ascomata 112-250 µm diam, 140-170 µm high, globose or depressed, immersed, usually hypophyllous, apapillate, apex rounded with plane pore or short papillate or conic. Peridium 10-20 um wide, comprising brown cells of textura thick-walled. angularis. Asci 40-75 × 8-14 µm, 8-spored, unitunicate, cylindrical, sessile, apical ring bilobed, distinct, shallow. Ascospores 10-15 × 3.5-6.5 µm, overlapping uniseriate, ellipsoid, ovoid or fusoid, straight or often inequilateral, guttulate, hyaline, aseptate. Asexual morph: Conidiomata acervular, superficial, black, coriaceous. Conidiophores reduced to conidiogenous cells. Conidiogenous cells 5-10 µm long, conical, wide, aseptate, hyaline. Conidia 12-15 × 8-9 µm, oblong to ellipsoid, hyaline, aseptate, with two small guttules (description of asexual morph from Potebnia 1910).

Material examined: Austria, Sonntagberg, New Rosenau, July, on leaves of Carpinus betulus (Betulaceae), P.P. Strasser, IMI 11662.

Notes: Traverso (1907) erected Apiosporopsis as a subgenus of *Guignardia* to accommodate *Guignardia carpinea* and *G. veneta* based on their distinct morphological characters. Mariani (1911) raised Apiosporopsis to generic rank describing *A. saccardiana* as a third species. Von Höhnel (1917) proposed Sphaerognomonia to accommodate Apiosporopsis carpinea. Reid & Dowser (1990) evaluated this genus and proposed Apiosporopsis as the correct name for Sphaerognomonia, retaining the type species as Apiosporopsis carpinea. Index Fungorum (2017) and MycoBank (2017) list another two species of Apiosporopsis as *A. saccardoana* and *A. coronillae*.

Apiosporopsis carpinea was recorded only on over-wintered living leaves. Gloeosporium robergei was reported as the asexual morph of A. carpinea (Potebnia 1910, Treigien & Markovska 2007). However, there are no molecular data to prove this. Gloeosporium robergei was reported as the causal agent of bud mortality and twig cankers on Ostrya virginiana (Sinclair & Hudler 1980). Sequences of this species (CBS 617.72 and CBS 738.68) placed the genus in the Diaporthales, but not in the Gnomoniaceae or Melanconidaceae (Sogonov et al. 2008). The molecular analysis of this study revealed that Apiosporopsis species formed a separate, well-supported clade (Fig. 1, Clade 3). Morphologically this clade is distinct from other families of Diaporthales having ascospores with pseudo-septate, sharply pointed ends, sessile unitunicate asci with a bilobed apical ring, and apapillate, immersed ascomata. Hence, we introduce the family Apiosporopsidaceae to accommodate these species.

Apoharknessiaceae Senan., Maharachch. & K.D. Hyde, fam. nov. MycoBank MB821881. Facesoffungi number FoF03457. Clade 13.

Endophytic, saprobic or pathogenic. Sexual morph: Undetermined. Asexual morph: *Conidiomata* stromatic or eustromatic, subepidermal to immersed, solitary to gregarious, subglobose to irregular, unilocular, pale brown. *Conidiomata wall* outer layer composed of thin-walled, pale brown cells of *textura*



Fig. 2. Apiosporopsis carpinea (IMI 11662). A. Packet of the herbarium specimen. B. Herbarium specimen. C. Ascomata on substrate. D, E. Vertical section of ascomata. F–I. Asci. J–M. Ascospores. Scale bars: C = 500 μm, D, E = 50 μm, F–M = 10 μm.

angularis, inner layer pale yellow to hyaline. Conidiophores reduced to conidiogenous cells or hyaline, septate, cylindrical, and sparingly branched. Conidiogenous cells holoblastic, cylindrical, lageniform to ampulliform, hyaline, smooth, invested in mucus. Conidia obclavate, conical, aseptate, pale brown, with a longitudinal band on the flat surface, thick and smooth-walled, guttulate, with short hyaline apiculus, with small globule of mucus on base or obtuse apex with a scar at the base.

Type genus: Apoharknessia Crous & S.J. Lee.

Type species: Apoharknessia insueta (B. Sutton) Crous & S.J. Lee.

Notes: Apoharknessia displays similar morphological characters to *Harknessia* but differs in having a hyaline, apical apiculus. Nag Raj (1993) listed *Mastigonetron*, as a synonym for *Harknessia*. *Mastigonetron* is typified by *M. fuscum* (= *H. insueta*). However, this species has a *Wuestneia* sexual morph, *W. fusca*, and it does not cluster with other *Harknessia* species. Therefore, *Apoharknessia* was introduced to accommodate *H. insueta* (Lee *et al.* 2004). The genus *Apoharknessia* presently accommodates two species (Crous *et al.* 2017).

Lasmenia species cause rachis necrosis, flower abortion and necrotic spots on leaves of *Nephelium lappaceum*. Several *Lasmenia* species associated with tropical fruits as pathogens have been isolated. DNA-based studies report a close affinity of *Lasmenia* to *Cryphonectriaceae* (Serrato-Diaz *et al.* 2011). Lasmenia was introduced in 1886 without designating any type species and *L. balansae* was selected as the lectotype species by von Höhnel (1910). There are 12 species recorded under *Lasmenia* in Index Fungorum (2017). *Lasmenia* species are reported as the causative agents of rachis necrosis, flower abortion, fruit rot, and leaf spots on *Nephelium lappaceum* (Serrato-Diaz *et al.* 2011). A few species have been transferred to *Lasmeniella*, but some species remain doubtful.

Phylogenetic analysis in the present study indicates that *Apoharknessia* and *Lasmenia* clearly belong to the *Diaporthales* in a well-supported clade (Fig. 1, Clade 13). However, the sequences of *Lasmenia* which are included in this study are not of a known species and given the sparse taxa in this family, any affinity between the two genera can not be ascertained.

Hence, we introduce a new family *Apoharknessiaceae* to accommodate these two genera. Morphologically species of this clade are distinct from other families of *Diaporthales* in having eustromatic to stromatic pycnidial conidiomata, blastic or phialidic conidiogenesis and ellipsoid to conical conidia with a longitudinal band on the flat surface or small globule of mucus at the base.

Apoharknessia insueta (B. Sutton) Crous & S.J. Lee, Stud. Mycol. 50: 240. 2004. Facesoffungi number FoF03458.

Illustration: See Lee et al. (2004).

Foliicolous forming bleached spots or saprobic on various substrates. Sexual morph: Undetermined. Asexual morph: *Conidiomata* stromatic, subepidermal to immersed, solitary to gregarious, subglobose to irregular, unilocular, pale brown. *Conidiomata wall* outer layer composed of thin-walled, pale brown cells of *textura angularis*, inner layer pale yellow to hyaline. *Conidiophores* reduced to conidiogenous cells. *Conidiogenous cells* $5-15 \times 4-6 \ \mu m (\overline{x} = 9 \times 4.8 \ \mu m)$, lageniform to ampulliform, hyaline, smooth, invested in mucus. *Conidia* $10-12 \times 7.5-9 \ \mu m (\overline{x} = 10.5 \times 8 \ \mu m)$, conical, aseptate, brown, with a longitudinal band on the flat surface, thick and smoothwalled, guttulate, with short hyaline apiculus, with small globule of mucus on base. *Basal appendage* 2 × 1–1.5 \ µm, often gelatinising and resulting in a minute marginal frill on the truncate base of the conidia (description based on Nag Raj 1993).

Notes: Apoharknessia was introduced and typified by *Apoharknessia insueta* and it clustered distant from *Harknessia sensu stricto* (Clade 7) (Lee *et al.* 2004). *Apoharknessia* is morphologically similar to *Harknessia* but distinct in having a hyaline apical apiculus in conidia and cultures on oatmeal or malt extract agar not forming fluffy aerial mycelium. In addition, it grows within the medium and sporulates directly on hyphae without forming conidiomata. Crous *et al.* (2017) introduced a new species as *Apoharknessia eucalyptorum*.

Asterosporiaceae Senan. Maharachch. & K.D. Hyde, fam. nov. MycoBank MB821539. Facesoffungi number FoF03459. Clade 23.

Endophytic or saprobic on *Betulaceae*, *Fagaceae*, *Juglandaceae* and *Sapindaceae*. Sexual morph: Undetermined. Asexual morph: *Conidiomata* acervular, subepidermal, erumpent at maturity, solitary, or occasionally confluent, unilocular, dark brown to black. *Conidiomata wall* composed of thin-walled,

brown cells of *textura angularis*. *Conidiophores* cylindrical, branched at the base, septate, hyaline to pale brown. *Conidiogenous cells* holoblastic, cylindrical, unbranched, integrated, determinate, hyaline to pale brown, smooth. *Conidia* terminal, transversely distoseptate, consisting of four arms, with reduced lumina, brown, smooth-walled.

Type genus: Asterosporium Kunze.

Type species: Asterosporium hoffmannii Kunze.

Notes: A molecular phylogenetic analysis based on SSU nrDNA, LSU nrDNA, ITS nrDNA and beta-tubulin positions Asterosporium species within Sordariomycetes (Tanaka et al. 2010). Wijayawardene et al. (2016) showed that Asterosporium species are related to Diaporthales forming a sister clade to species in Sydowiellaceae based on combined ITS nrDNA and LSU nrDNA sequence analyses. In this study, Asterosporium species are positioned in Diaporthales (Fig. 1, Clade 23) and constitute a well-supported sister clade to Svdowiellaceae and Lamproconiaceae. Morphologically, Asterosporium species are distinct from other members of Diaporthales in having star-like, brown conidia. Hence, we introduce a novel family Asterosporiaceae to accommodate Asterosporium species. We illustrate Asterosporium asterospermum collected from Italy.

Asterosporium asterospermum (Pers.) Hughes, Canad. J. Bot. 36: 738. 1958. Fig. 3.

Basionym: Stilbospora asterosperma Pers. [as 'asterospora'], Syn. meth. fung. (Göttingen) 1: 96. 1801.

Saprobic on twigs and branches of *Fagus sylvatica*. Sexual morph: Undetermined. Asexual morph: *Conidiomata* 2–2.5 mm high, 0.8–1 mm diam ($\bar{x} = 2.1 \times 0.86$ mm, n = 15), acervular, subepidermal, erumpent at maturity, solitary, or occasionally confluent, unilocular, dark brown to black. *Conidiomata wall* 25–30 µm ($\bar{x} = 29$, n = 20), composed of thin-walled, brown cells of *textura angularis*. *Conidiophores* 30–35 µm high, 5–8 µm wide ($\bar{x} = 29 \times 7$ µm, n = 20), cylindrical, branched at the base, septate, hyaline to pale brown. *Conidiogenous cells* 70–100 µm high, 4–7 µm wide ($\bar{x} = 80 \times 5$ µm, n = 20), holoblastic, cylindrical, unbranched, integrated, determinate, hyaline to pale brown, smooth-walled. *Conidia* 65–75 × 90–115 µm ($\bar{x} = 68 \times 100$ µm, n = 20), terminal, transversely distoseptate, consisting of four arms, with reduced lumina, brown, smooth-walled.

Specimen examined: Italy, Forlì-Cesena Province, Santa Sofia, near Passo la Calla, on dead branch of *Fagus sylvatica (Fagaceae)*, 29 Sep. 2012, E. Camporesi, IT 805, MFLU 15-3555, HKAS 92536.

Notes: Asterosporium was introduced and typified by Asterosporium asterospermum (= Stilbospora asterosperma and Asterosporium hoffmannii) and there are five species listed Fungorum А. Index (2017), namely acerinum, in asterospermum, A. attenuatum, A. hoffmannii and Α. A. strobilorum. However, only A. asterospermum has DNA sequence data in GenBank. There are no records for the sexual morph of Asterosporium (Tanaka et al. 2010). Species of this genus are associated with twigs and stems of overwintered plants as endophytes.



Fig. 3. Asterosporium asterospermum (MFLU 15–3555). A. Conidiomata on host substrate. B, C. Vertical section of conidiomata. D–H. Different stages of conidiogenesis. I–M. Conidia. Scale bars: A = 1 mm, B = 400 µm, C = 50 µm, D–H = 20 µm, I–M = 30 µm.

Auratiopycnidiellaceae Senan., Maharachch. & K.D. Hyde, fam. nov. MycoBank MB821540. Facesoffungi number FoF03460. Clade 11.

Foliicolous. Sexual morph: Undetermined. Asexual morph: *Conidiomata* amphigenous, pycnidia, globose, orange on leaves with dark brown border. *Peridium* comprises pale brown cells of *textura angularis. Paraphyses* hyaline, cellular, subcylindrical, branched or not, with obtuse apex, septate, constricted at septa. *Conidiophores* reduced to conidiogenous cells. *Conidiogenous cells* hyaline, smooth, lageniform to ampulliform, with terminal truncate locus, thick-walled, sometimes appearing to proliferate percurrently. *Conidia* ellipsoid, smooth, solitary, median 1-septate, constricted at septum, apex obtuse, base truncate, thickened, at times with marginal frill, becoming golden brown at germination with solitary, brown, wavy germ tubes.

Type genus: Auratiopycnidiella Crous & Summerell.

Type species: Auratiopycnidiella tristaniopsidis Crous & Summerell.

Notes: Crous *et al.* (2012a) described *Auratiopycnidiella* as a genus with subepidermal, orange, pycnidial conidiomata, forming hyaline, holoblastic conidiogenous cells, with or without a thickened scar and hyaline, ellipsoid, 1-septate conidia having a thickened hilum or minute marginal frill. Crous *et al.* (2012a)

reported that the genus is phylogenetically distant to *Melanconidaceae* based on LSU nrDNA sequence data and treated this genus as *Diaporthales* genera *incertae sedis* pending the availability of more molecular data. A megablast search of NCBI's GenBank nucleotide database using the calmodulin, ITS nrDNA, and beta-tubulin sequences retrieved sequence similarities with *Harknessiaceae* and *Cryphonectriaceae* (Crous *et al.* 2012a). Our phylogenies generated herein indicate that *Auratiopycnidiella* forms a single branch which is phylogenetically distinct from all other included families (Fig. 1, Clade 11) and hence we introduce *Auratiopycnidiellaceae* to accommodate *Auratiopycnidiella. Auratiopycnidiella* currently comprises a single species with a single isolate.

Auratiopycnidiella tristaniopsidis Crous & Summerell [as '*tristaniopsis*'], Persoonia 28: 69. 2012. Facesoffungi number FoF03461.

Illustration: See Crous et al. (2012a).

Foliicolous. Sexual morph: Undetermined. Asexual morph: *Conidiomata* up to 200 μ m diam, amphigenous, pycnidia, globose, orange on leaves with dark brown border, with irregular central opening. *Peridium* up to 25 μ m thick, comprising 4–7 layers of pale brown cells of *textura angularis*. *Paraphyses* hyaline, cellular, subcylindrical, branched or not, with obtuse apex, 2–6-septate, constricted at septa *Conidiophores* 10–25 × 3–6 μ m, reduced to

conidiogenous cells. *Conidiogenous cells* hyaline, smooth, lageniform to ampulliform, with terminal truncate locus, thick-walled, sometimes appearing to proliferate percurrently. *Conidia* $13-15 \times 5-5.5 \,\mu$ m, ellipsoid, smooth, solitary, medially 1-septate, constricted at septum, obtuse at apex, truncate at base, thickened at times with marginal frill, hyaline becoming golden brown during germination with solitary, brown, wavy germ tubes 90° to the long axis of the spore (description based on Crous *et al.* 2012a).

Notes: Auratiopycnidiella was introduced and typified by *Auratiopycnidiella tristaniopsis.* This is a monotypic genus comprising only the type species, *A. tristaniopsis. Auratiopycnidiella tristaniopsis* forms leaf spots on its host species. Morphologically this taxon shows some similarities to taxa of the *Cryphonectriaceae* in having orange conidiomata. However, phylogenetically it is distinct from *Cryphonectriaceae*.

Coryneaceae Corda, Icon. fung. (Prague) 3: 36. 1839. Clade 20. *Synonym: Pseudovalsaceae* M.E. Barr, Mycol. Mem. 7: 151. 1978.

Saprobic on dead wood or pathogenic. Sexual morph: Stromata solitary, erumpent, comprising pseudoparenchymatous cells. Ectostromatic disc well or poorly developed, brown to black, comprising small cells of textura prismatica cells. Ascomata perithecial, arranged in valsoid configuration, immersed, aggregated, globose to subglobose, coriaceous, brown to black, papillate, ostiolate. Papilla upright, central, broad, sometimes converging, comprising brown cells of textura porrecta. Peridium comprising outer, thick-walled, brown cells of textura angularis and inner, thick-walled, hvaline, compressed cells of textura angularis. Hamathecium comprising broad, cellular, septate paraphyses, attached to base, longer than asci. Asci 8-spored, unitunicate, ellipsoid to cylindrical, thin-walled, pedicellate, apex rounded with a J- apical ring. Ascospores overlapping uni- to biseriate, hyaline or initially hyaline, brown at maturity, irregularly fasciculate, ellipsoid, fusoid or elongate, 1-3-septate, often distoseptate, end cells pale brown or hyaline, sometimes end cells pointed, straight or curved not constricted at the septa, guttulate, smooth-walled. Asexual morph: Coelomycetous. Conidiomata acervular, solitary, erumpent through the outer periderm layers of host or immersed, scattered, surface tissues above slightly dome-shaped. Conidiomatal wall composed of thin-walled, vertically arranged dark brown cells of textura angularis. Conidiophores branched at the base or not, cylindrical to globose, septate or aseptate, hyaline or hyaline at the apex, pale brown at the base. Conidiogenous cells terminal, hyaline, annellidic, cylindrical, sometimes with setulose apical appendages. Conidia hyaline to dark brown, curved, broadly fusiform to cylindrical or clavate, smooth-walled, 4-6-distoseptate, sometimes the apical and basal cell darker than other cells with hyaline tip in apical cell.

Type genus: Coryneum Nees.

Type species: Coryneum umbonatum Nees.

Notes: The family *Coryneaceae* (Fig. 1, Clade 20) was introduced by Corda (1839) based on *Coryneum*. However, Barr (1978) introduced the family *Pseudovalsaceae* based on *Pseudovalsa lanciformis*, which is the sexual morph of *Coryneum umbonatum*. Hence *Pseudovalsaceae* must be synonymised under *Coryneaceae* giving priority to the older name. Rossman *et al.* (2015) protected the earliest name *Coryneum* (1816) over *Pseudovalsa* (1863) and conserved *Coryneum umbonatum* as the type species. This family comprises fungal taxa with upright, erumpent perithecia and central beaks. However, many genera previously included in *Coryneaceae* have been placed in various other families (Castlebury *et al.* 2002) and the only genus remaining in the family is *Coryneum*.

Coryneum arausiaca (Fabre) Senan., Maharachch. & K.D. Hyde, **comb. nov.** MycoBank MB821543. Facesoffungi number FoF03462. Fig. 4.

Basionym: Pseudovalsa arausiaca Fabre, Sphér. Vaucl.: 56. 1883.

Saprobic on branches of Quercus sp. Sexual morph: Stromata comprising loosely packed, black, hyphae mostly around the neck. Ascomata 600-700 µm high, 315-365 µm diam $(\overline{x} = 640 \times 340 \text{ µm}, \text{ n} = 20)$, immersed, 5–10 aggregated in one group, visible only as ostiolar opening through cracks in bark, valsoid, globose, brown to black, papillate, ostiolate, ostiole periphysate, periphyses hyaline, long. Peridium 25-50 µm $(\overline{x} = 45 \ \mu m, n = 10), 10-15$ layers of thick-walled, brown-walled cells of textura angularis and papilla comprising brown cells of textura porrecta. Paraphyses 5–10 μ m (\overline{x} = 7.6 μ m, n = 20), few, hyaline, septate, attached at base, longer than asci. Asci $145-155 \times 25-30 \ \mu m$ ($\overline{x} = 146 \times 25.6 \ \mu m$, n = 10), 8spored, unitunicate, clavate, short pedicellate, apically rounded, narrow, J-, without an obvious apical ring. Ascospores $70-90 \times 6.5-8.5 \ \mu m$ ($\overline{x} = 77 \times 7.5 \ \mu m$, n = 10), 2-3-seriate, broadly ellipsoidal, ends pointed, 1-3-septate, not constricted at the septa, hyaline, guttulate, smooth-walled. Asexual morph: Conidiomata acervular, 1-1.3 mm wide, 0.5-0.55 mm high $(\overline{x} = 1.1 \times 0.51 \text{ mm}, n = 20)$, solitary, erumpent through the outer periderm layers of host, scattered, surface tissues above slightly domed. Conidiomatal wall 100-150 μ m (\overline{x} = 135 μ m, n = 20), composed of thin-walled, vertically arranged, dark brown cells of textura epidermis. Conidiophores 20-35 µm long, 4-7 µm wide $(\overline{x} = 30 \times 6 \mu m, n = 20)$, branched at the base, cylindrical, septate, hyaline at the top, pale brown at the base. Conidiogenous cells 4–7 μ m long, 4.5–6 μ m wide (\overline{x} = 6 × 5 μ m, n = 20), formed from the apical cell of the conidiophore, holoblastic, cylindrical, hyaline. Conidia 42-56 × 13-16 µm $(\overline{x} = 48 \times 14 \ \mu m, n = 20)$, curved, broadly fusiform to fusiformcylindrical or clavate (rather variable in form), dark brown, smooth-walled, 4-6-disto-septate, with apical and basal cells darker than other cells, apical cell with a hyaline tip, truncate and black at base.

Culture characteristics: Ascospores germinating on MEA within 12 h and germ tubes produced from both ends, fast growing on MEA at 25 °C, after 1 wk reaching 3 cm diam, white, cottony, margin wavy, superficial, slightly effuse, radially striated, edges with more aerial mycelium than centre.

Specimens examined: **Italy**, Province of Forli-Cesena, Civitella di Romagna, Pian di Spino, on branch of *Quercus* sp. (*Fagaceae*), 25 Mar. 2013, E. Camporesi, IT 1144, (**neotype designated here** MFLU 14–0796, cultures exneotype, MFLUCC 13–0658); Province of Forli-Cesena, Civitella di Romagna, Pian di Spino, on branch of *Quercus* sp. (*Fagaceae*), 16 Feb. 2015, E. Camporesi, IT 1144A, **paraneotype** HKAS83943, cultures ex-paraneotype, MFLUCC 15–1110.



Fig. 4. Coryneum arausiaca (MFLU 14–0796). A, B. Ascostromata on substrate. C, D. Vertical section of ascoma. E. Peridium. F. Periphyses. G. Paraphyses. –J. Asci. K–S. Ascospores. T. Conidiomata on substrate. U. Vertical section of conidiomata. V. Conidiophores, conidiogeneous cells with conidia. W–ZB. Conidia. Scale bars: A, B = 500 µm, C, D, S = 100 µm, E–G, K, Q, U–Z = 10 µm, H–J = 20 µm, R = 1 mm, T = 50 µm.

Notes: We have re-collected and neotypified *Pseudovalsa arausiaca. Pseudovalsa arausiaca* has immersed, globose perithecia in a valsoid configuration with broadly ellipsoidal, 1–3septate, hyaline ascospores. The neotype is morphologically identical to *Pseudovalsa arausiaca* described by Fabre (1883). However, we could not locate the type specimens and assume that they are lost. Fortunately, we obtained fresh material from the same host genus and location. Therefore, a neotype is designated here with sequence data. Rossman *et al.* (2015) protected *Coryneum* over *Pseudovalsa arausiaca* as *Coryneum arausiaca.* Both sexual and asexual morphs of *Coryneum arausiaca* were obtained from the same specimen as well as cultures which indicate a holomorph connection. We illustrate both sexual and asexual morphs of *Coryneum arausiaca* and the combined gene analysis of LSU nrDNA, ITS nrDNA, *rpb2* and *tef1* shows the distinct placement of *C. arausiaca* within *Coryneaceae* (Fig. 1, Clade 20).

Cryphonectriaceae Gryzenh. & M.J. Wingf., Mycologia 98: 246. 2006. Clade 6.

Saprobic or pathogenic in forest trees and economic crops. Sexual morph: Ascostromata scattered, immersed or erumpent, aggregated, oval to circular from above, comprising two layers, upper layer of yellowish orange to pale brown cells, purpling in



Fig. 5. Cryphonectria parasitica (NY 01293321). A. Herbarium packet. B. Herbarium specimen. C. Ascostromata on substrate. D. Horizontal section of ascostroma. E. Vertical cross section of ascoma. F–I. Asci. J–M. Ascospores. N. Horizontal cross section of conidiomata. O. Vertical cross section of conidioma. P, Q. Conidia attached to the conidiogenous cells and conidiophore. R. Conidia. Scale bars: C, D = 1 mm, F–I, P–R = 10 μm, E, Q, O = 100 μm, N = 200 μm, J–M = 5 μm.

KOH and inner layer of hyaline cells, mixed with plant cells. Ascomata immersed, aggregated, several in one stroma, globose to subglobose, fuscous black to umber, with long neck, or ostiolar canal sometimes immersed in stromatic tissues, or superficial, necks covered in umber stromatic tissue of textura porrecta, inner wall of the necks or ostiolar canal covered with hyaline, filamentous periphyses. Peridium comprising inner layer of small, hyaline cells of textura angularis and outer layer of small, brown cells of textura angularis. Hamathecium comprising a few cellular paraphyses and parenchymatous cells, attached at the base and asci dissolving at maturity. Asci 8-spored, unitunicate, cylindrical-fusoid to clavate, pedicellate, with distinct, Jrefractive ring. Ascospores overlapping uniseriate to biseriate, ellipsoid, fusoid to cylindrical, aseptate to multi-septate, not constricted at the septa, hyaline, sometimes brown, smoothwalled. Asexual morph: Coelomycetous. Conidiomata occurring as a part of ascomata as conidial locules or solitary structures, uni- to multi-loculate, pyriform, subglobose to pulvinate, necks absent or present, if present, with one to several attenuated necks, superficial or semi-immersed, orange to fuscous-black. Conidiophores cylindrical, aseptate, hyaline, sometimes reduced to conidiogenous cells. Conidiogenous cells lining the

inner cavity of the conidiomata, phialidic, sometimes within flattened bases, ampulliform, inconspicuous, with attenuated or truncate apices, hyaline, smooth. *Conidia* minute, sometimes both micro- and macro-conidia present, sigmoid, broadly ellipsoid to fusoid, obovoid-cylindrical to allantoid, aseptate, hyaline.

Type genus: Cryphonectria (Sacc.) Sacc. & D. Sacc.

Type species: Cryphonectria parasitica D. Sacc.

Notes: Cryphonectriaceae (Fig. 1, Clade 6) is mostly a pathogenic family comprising some of the world's most important tree pathogens (Vermeulen *et al.* 2011). Cryphonectriaceous species are saprobes, endophytes and phytopathogens. They cause cankers, blights and dieback of economically important plants and forest trees. Castlebury *et al.* (2002) recognised the *Cryphonectria-Endothia* complex (a precursor to the *Cryphonectriaceae*) as a separate clade in *Diaporthales* based on analysis of LSU nrDNA sequence data. *Cryphonectriaceae* was formally established by Gryzenhout *et al.* (2006c) to accommodate the *Cryphonectria-Endothia* complex and other allied genera when analysing LSU nrDNA sequence data of fungal taxa in Diaporthales. Species of this family can be distinguished from other families of Diaporthales by orange stromatic tissues, which turn purple in KOH and yellow in lactic acid. Initially Amphilogia, Chrysoporthe, Cryphonectria, Endothia and Rostraureum were placed in the family (Gryzenhout et al. 2006c). Subsequently, several additional genera were added to the family, some associated with serious canker or foliar diseases, namely: Aurantiosacculus, Aurapex, Aurifilum, Celoporthe, Chromendothia, Chrysocrypta, Chrvsofolia, Crvptometrion, Diversimorbus, Foliocrvphia, Holocryphia, Immersiporthe, Lasmenia, Latruncellus, Luteocirrhus, Mastigosporella, Microthia, Prosopidicola and Ursicollum (Vasilyeva 1993, Gryzenhout et al. 2006a, b, Nakabonge et al. 2006, Begoude et al. 2010, Gryzenhout et al. 2010, Vermeulen et al. 2011, Crous et al. 2012a, Chen et al. 2013, Crane & Burgess 2013, Crous et al. 2013).

Endothia (1849) is typified by E. gyrosa and the asexual morph of this genus was reported as an Endothiella species (Barr 1978). However, Endothiella is congeneric with Cryphonectria and Endothiella eucalypti is the asexual morph of type species of Cryphonectria, C. eucalypti (Jackson 2003). Endothiella (1906) is based on the type species, Endothiella gyrosa, now placed in Cryphonectria as C. decipiens (Gryzenhout et al. 2009). Barr (1978) observed several specimens of Cryphonectria and Endothia and she used stromatic configuration and ascospore characters to differentiate these two genera. According to Barr (1978), Cryphonectria has a valsoid configuration of perithecia in prosenchymatous stromata and ellipsoid or ovoid, 1septate ascospores, while Endothia has a diatrypoid configuration of perithecia in pseudoparenchymatous stromata and allantoid, unicellular ascospores. Based on these characters, most Endothia species have been moved to Cryphonectria and the generic name Endothia was restricted to the species with a diatrypoid configuration of the perithecia and allantoid, unicellular ascospores. Combined analysis of LSU nrDNA, ITS nrDNA, rpb2 and tef1 sequence data in the present study shows Cryphonectriaceae is not well-supported (Fig. 1, Clade 6). Phylogenetic analyses of this study also place Cryphonectria and Endothia as two separate genera, as well as Chrysocrypta (Fig. 1, Clade 9), Lasmenia (Fig. 1, Clade 13) and Prosopidicola (Fig. 1, Clade 17) outside of Cryphonectriaceae. Hence, currently this family comprises Amphilogia, Aurantioporthe, Aurantiosacculus, Aurapex, Aurifilum, Celoporthe, Chromendothia, Chrysofolia, Chrysoporthe, Chrysoporthella, Cryphonectria, Cryptometrion, Diversimorbus, Endothia, Foliocryphia, Holocryphia, Immersiporthe, Latruncellus, Luteocirrhus, Mastigosporella, Microthia, Rostraureum and Ursicollum.

Cryphonectria parasitica (Murrill) M.E. Barr, Mycol. Mem. 7: 143. 1978. Facesoffungi number FoF03463. Fig. 5. *Basionym: Diaporthe parasitica* Murrill, Torreva 6: 189. (1906).

Pathogenic on branches of *Castanea dentata*. Sexual morph: *Ascostromata* 6.5–1 mm diam ($\bar{x} = 0.8$ mm, n = 20), comprising erumpent to superficial, orange epistromatic portion and immersed, hyaline, parenchymatous portion. *Ascomata* 650–715 µm high, 210–220 µm diam ($\bar{x} = 685 \times 216$ µm, n = 20), perithecial, immersed, globose to subglobose, with black to brown ostiole, ostiolar canal slender, covered with orange to fuscous-black stromatic tissue. *Hamathecium* aparaphysate, comprising parenchymatous tissues. *Asci* 20–35 × 5–8 µm ($\bar{x} = 28 \times 6.4$ µm, n = 20), 8-spored, unitunicate, fusiform to cylindrical base with small pedicel, apex oblong. Ascospores $5-6 \times 2-2.5 \ \mu m$ ($\overline{x} = 5.5 \times 2.2 \ \mu m$, n = 20), overlapping uni- or biseriate, hyaline, ellipsoid to fusiform, 1-septate. Asexual morph: Conidiomata 250–300 μm high, 180–200 μm diam ($\overline{x} = 280 \times 185 \ \mu m$, n = 20), eustromatic, erumpent, pyriform to pulvinate, orange to fuscous black, occurring in the same stroma as perithecia. Conidiophores $3-4 \times 1-1.5 \ \mu m$ ($\overline{x} = 3.4 \times 1.1 \ \mu m$, n = 20), cylindrical, unbranched, hyaline. Conidiogenous cells $2.5-5 \times 0.5-1 \ \mu m$ ($\overline{x} = 3.3 \times 0.8 \ \mu m$, n = 20), phialidic, simple or branched. Conidia $1.8-2.5 \times 0.5-1 \ \mu m$ ($\overline{x} = 2.1 \times 0.9 \ \mu m$, n = 20), hyaline, minute, allantoid to cylindrical, aseptate.

Materials examined: USA, New York. Bronx Co. Bronx. North of Botanical Museum, Bronx Park, on *Castanea dentata (Fagaceae)*, 26 Nov. 1905, W.A. Murrill (holotype 01293321, as *Diaporthe parasitica*, NY).

Notes: American chestnut blight, caused by *Cryphonectria parasitica*, destroyed American chestnut trees in the USA and Canada at the end of the 19th century. Scientists believed *Cryphonectria parasitica* arrived from north-east Asia in the late 19th century and they discovered that Japanese and Chinese chestnut varieties showed resistance to *C. parasitica*. Spores of this fungus are highly resistant to unfavourable environmental conditions and they can be produced at any time of year when conditions are suitable. The fungus can exist as a saprobe and a parasite. Mycelium can survive more than 10 mo in dried bark and soil (Hepting 1974). Conidia and ascospores of *C. parasitica* are sometimes forcibly ejected and spread in wind and rain. Spores of *Cryphonectria parasitica* are also dispersed by beetles and birds. In addition to chestnut species, some oak species and Chinquapin also are infected by *Cryphonectria parasitica*.

Cytosporaceae Fr. [as '*Cytisporei*'], Syst. orb. veg. (Lundae) 1: 118. 1825. Clade 16.

Synonym: Valsaceae Tul. & C. Tul. [as 'Valsarum'], Select. fung. carpol. (Paris) 1: 180. 1861.

Pathogenic or saprobic on plant tissues. Sexual morph: Stromata well or poorly developed. Ectostroma circular or irregular, usually well developed in the upper regions. Entostroma normally limited to the region near the perithecial walls. Ascomata perithecia, immersed to erumpent, solitary or 6-10 ascomata aggregated in valsoid configuration, globose to oblong, coriaceous, black to brown, with long neck swollen at the tips, ostiolate. Ostiole periphysate, open through the neck. Peridium thin, comprising outer, 4-6 layers of, dark brown, thick-walled, cells of textura angularis and 5-7 layers of, inner, small, hyaline, thin-walled, cells of textura angularis. Hamathecium comprising few, hyaline paraphyses limited only at young stage. Asci unitunicate, 8spored, clavate, short-pedicellate, apex round, with apical ring. Ascospores uni- to biseriate, unicellular or rarely bicellular, allantoid or ellipsoid, hyaline, smooth-walled. Asexual morph: Stromata uniloculate, black, circular in shape. Locule composed of numerous inter connecting chambers arranged radially or irregularly within a continuous mass of ectostromatic tissue, one conidioma per locule. Conidiomata pyriform in section, brown, divided into compartments by bending of peridium. Peridium consists of brown, 5-7 layers of textura angularis cells. Conidiophores reduced to conidiogenous cells. Conidiogenous cells arising from conidiomatal wall, phialidic, simple or branched, hyaline, cylindrical. Conidia unicellular, allantoid, hyaline, smooth-walled.

Type genus: Cytospora Ehrenb.

Type species: Cytospora chrysosperma (Pers.) Fr.

Notes: The Cytosporaceae (Fig. 1, Clade 16) comprises phytopathogenic species and saprobes. Most Cytospora species are plant pathogens and cause cankers and dieback of many hardwoods and coniferous trees, as well as rarely on herbaceous plants. Generally, Cytospora cankers are known as valsa-canker, Leucostoma-canker or perennial canker (Farr et al. 1989). Cytospora species have been reported as highly virulent and destructive pathogens on Prunus and Populus trees (Biggs 1989, Kepley & Jacobi 2000). A few Cvtospora species are considered as facultative wound parasites that attack damaged or weakened plants. Maharachchikumbura et al. (2015, 2016) listed 13 genera under this family as Amphicytostroma, Chadefaudiomyces, Cryptascoma, Cytospora, Ditopellina, Durispora, Harpostroma, Hypospilina, Kapooria, Leptosillia, Maculatipalma, Pachytrype, and Paravalsa. However, the type species of Amphicytostroma, A. tiliae is the asexual morph of the type species of Amphiporthe, A. hranicensis, and these generic names are synonyms (Sutton 1980). Amphiporthe is more widely used than Amphicytospora and it seems best to protect the former (Rossman et al. 2015). However, Amphiporthe belongs in Gnomoniaceae (Sogonov et al. 2008; Fig. 1, Clade 1) and we exclude this genus from Cytosporaceae. Rossman et al. (2015) proposed to use Cytospora (1818) rather than Valsa (1825), Valsella (1870), Leucostoma (1917), Valseutypella (1919), or Leucocytospora (1927). Xenotypa is a genus in Gnomoniaceae and typified by Xenotypa aterrima. This genus is characterised by having solitary or aggregated, erumpent, globose, papillate ascomata with allantoid to cylindrical, unicellular, hyaline ascospores. Morphologically this is similar to Paravalsa and Valsella. Ananthapadmanaban (1990) described the relationship between Xenotypa and Paravalsa, accommodating Paravalsa in Valsaceae. However, many of the fungal taxa listed in Maharachchikumbura et al. (2015) do not share similar morphological characters and it is necessary to restrict this family to Cytospora sensu-lato. Cytospora, Valsella, Leucostoma, Valsa and Pachytrype have sequence data in accessible data bases. Hence, we suggest to accommodate Cytospora, Paravalsa, Pachytrype, Waydora and Xenotypa in Cytosporaceae. However, the Cytospora sensu-lato complex still needs to be resolved using high resolution genes as it seems to comprise several genera.

Cytospora centrivillosa Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821567. Facesoffungi number FoF03464. Fig. 6.

Etymology: Name based on two Latin words "*centrum*" and "*villos*" meaning hamathecium comprising filiform paraphyses.

Saprobic on dead branch of *Sorbus domestica*. Sexual morph: *Stromata* poorly developed, comprising loosely packed parenchymatous cells, black. *Ascomata* 550–725 µm high, 160–215 µm diam ($\overline{x} = 611 \times 190 \mu$ m, n = 20), aggregated, immersed, globose to subglobose, dark brown, coriaceous, ostiolate, papillate. *Papilla* 285–430 µm high, 90–130 µm diam ($\overline{x} = 340 \times 101 \mu$ m, n = 20), long, central or asymmetrically located, wall thick, internally covered by hyaline periphyses. *Peridium* comprises brown, thick-walled cells of *textura angularis*. *Asci* 75–85 × 15–19 µm ($\overline{x} = 79 \times 18 \mu$ m, n = 20), 8-spored, unitunicate, clavate to fusiform, without apical ring and pedicel. *Ascospores*

16–20 × 4–6 μm (\bar{x} = 17 × 5 μm, n = 20), biseriate, allantoid, hyaline, smooth. Asexual morph: Coelomycetous. *Conidiomata* on MEA appears as pale yellow, slimy heads of conidial mass, immersed, black. *Conidiophores* 6.5–8 × 3–3.5 μm (\bar{x} = 7.4 × 3.1 μm, n = 20), cylindrical, unbranched, hyaline. *Conidiogenous cells* 10–13.5 × 1–2 μm (\bar{x} = 11.7 × 1.6 μm, n = 20), cylindrical, tapering towards the apices, bearing single conidia at each tip, hyaline. *Conidia* 4–6 × 1–1.5 μm (\bar{x} = 5.1 × 1.1 μm, n = 20), eguttulate, allantoid, aseptate, hyaline.

Culture characteristics: Colonies growing on MEA attenuated 1 cm incubated at 18 °C within 4 d, fast growing, circular, flat, entire, white, thin, tightly attached to the media, mycelia clots arrange radially from centre to margin.

Specimens examined: Italy, Province of Forlì-Cesena, Predappio, Monte Mirabello, on dead and aerial branch of *Sorbus domestica* (*Rosaceae*), 1 Oct. 2014, E. Camporesi, IT 2132 (holotype MFLU 17–0887, isotype BBH 42449, culture ex-type MFLUCC 16–1206); Province of Forlì-Cesena, Predappio, Monte Mirabello, on dead and aerial branch of *Sorbus domestica* (*Rosaceae*), 13 Oct. 2014, E. Camporesi, IT 2132B, MFLU 17–0999, culture MFLUCC 17–1660.

Note: Cytospora centrivillosa is morphologically and phylogenetically distinct from other species in Cytospora and our analysis results in a distinct clade with full support (Fig. 1, Clade 16).

Cytospora fraxinigena Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821568. Facesoffungi number FoF03465. Fig. 7.

Etymology: Named after the host genus Fraxinus.

Saprobic on dead branch of *Fraxinus omus*. Sexual morph: *Stromata* poorly developed, comprising loosely packed parenchymatous cells, black. *Ascomata* 350–500 × 150–230 µm ($\overline{x} = 429 \times 189 \mu$ m, n = 20), immersed in stromatic tissues, globose to subglobose, dark brown, coriaceous, ostiolate, papillate. *Papilla* 185–200 × 60–95 µm ($\overline{x} = 193 \times 79 \mu$ m, n = 20), long, central, wide, thick-walled, internally covered by hyaline periphyses. *Peridium* comprises brown, thick-walled cells of *textura angularis*. *Asci* 26–33 × 6.2–7.5 µm ($\overline{x} = 30 \times 6.7 \mu$ m, n = 20), 8-spored, unitunicate, clavate to fusiform, without apical ring and pedicel. *Ascospores* 5.5–7.5 × 1.5–2 µm ($\overline{x} = 6.4 \times 1.7 \mu$ m, n = 20), biseriate, allantoid, hyaline, smooth. Asexual morph: Not observed.

Culture characteristics: Colonies growing on MEA attenuated 1 cm incubated at 18 °C within 7 d, moderate fast growing, irregular, flat, undulate, white, woolly, loosely attached to the media.

Specimen examined: Italy, Province of Forlì-Cesena, Santa Sofia, near Corniolo, dead branch of *Fraxinus ornus* (*Oleaceae*), 6 Dec. 2013, E. Camporesi, IT 1562 (holotype MFLU 17–0880, isotype BBH 42442, culture ex-type MFLUCC 14–0868).

Notes: Cytospora fraxinigena forms a distinct clade which is sister to *Cytospora cedri* and *Cytospora rosae* (Fig. 1, Clade 16). Morphologically, *Cytospora fraxinigena* differs from those species in having slightly horizontal necks closely arranged at apex and hamathecium without paraphyses.

Cytospora junipericola Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821569. Facesoffungi number FoF03466. Fig. 8.



Fig. 6. Cytospora centrivillosa (MFLU 17–0887). A, B. Stromata on substrate. C, D. Vertical cross section of ascomata. E. Peridium. F–H. Asci. I. Paraphyses. J. Ascospores. K. Conidiomata on MEA. L–N. Conidiogenous cells, conidiophores, conidia. O. Conidia. Scale bars: A = 500 μm, B = 200 μm, C, D = 50 μm, E = 5 μm, I = 20 μm, F–H, J, L–O = 10 μm, K = 500 μm.

Etymology: Named after the host genus Juniperus.

Saprobic on dead branch of *Juniperus* sp. Sexual morph: *Stromata* poorly developed, comprising loosely packed parenchymatous cells, black. *Ascomata* 630–700 µm high, 150–250 µm diam ($\bar{x} = 670 \times 170$ µm, n = 20), immersed in stromatic tissues, globose to subglobose, dark brown, coriaceous, ostiolate, papillate. *Papilla* 300–500 µm high, 45–65 µm diam ($\bar{x} = 440 \times 58$ µm, n = 20), long, central, wide, thick-walled, internally covered by hyaline periphyses. *Peridium* comprises brown, thick-walled cells of *textura angularis*. *Asci* 30–35 × 5.5–7 µm ($\bar{x} = 32 \times 6$ µm, n = 20), 8-spored, unitunicate, clavate to fusiform, without apical ring and pedicel. *Ascospores* 5–10 × 1–2 µm ($\bar{x} = 7 \times 1.5$ µm, n = 20), biseriate, allantoid, hyaline, smooth. Asexual morph: Not observed.

Culture characteristics: Colonies growing on MEA attenuated 1 cm incubated at 18 °C within 7 d, moderate fast growing, irregular, flat, undulate, greenish ash, woolly, curled, loosely attached to the media.

Specimen examined: Italy, Province of Forlì-Cesena, Santa Sofia, near Cabelli, dead branch of *Juniperus communis* (*Cupressaceae*), 13 Jan. 2014, E. Camporesi, IT 1643 (holotype MFLU 17–0882, isotype BBH42444).

Notes: Cytospora junipericola forms a distinct clade that is sister to *Cytospora quercicola* with high bootstrap support (Fig. 1, Clade 16). Morphologically *Cytospora junipericola* produces tightly packed aggregated ascomata in poorly developed stromatic tissues. Papilla are asymmetrically located and only the ostiolar openings are close together.



Fig. 7. Cytospora fraxinigena (MFLU 17–0880). A. Ascomata on substrate. B. Vertical cross section of ascoma. C. Peridium. D–F. Asci. G–J. Ascospores. Scale bars: B = 100 μ m, C = 20 μ m, D–F = 10 μ m, G–J = 5 μ m.

Cytospora quercicola Senan., Camporesi, & K.D. Hyde, **sp. nov.** MycoBank MB821570. Facesoffungi number FoF03467. Fig. 9.

Etymology: Named after the host genus Quercus.

Saprobic on dead branch of *Quercus* sp. *Stromata* poorly developed, spread around the papilla, black. *Ascomata* 550–725 µm high, 160–215 µm diam ($\bar{x} = 611 \times 190$ µm, n = 20), scattered, aggregated, immersed, globose to subglobose, dark brown, coriaceous, ostiolate, papillate. *Papilla* 285–430 µm high, 90–130 µm diam ($\bar{x} = 340 \times 101$ µm, n = 20), long, central or asymmetrically located, papilla close to each other when open to host surface. *Peridium* comprises brown, thick-walled cells of *textura angularis*. *Asci* 75–85 × 15–19 µm ($\bar{x} = 79 \times 18$ µm, n = 20), 8-spored, unitunicate, clavate to fusiform, without apical ring and pedicel. *Ascospores* 16–20 × 4–6 µm ($\bar{x} = 17 \times 5$ µm, n = 20), biseriate, allantoid, hyaline, smooth.

Culture characteristics: Colonies growing on MEA becoming 1 cm within 7 d incubated at 18 °C, circular, flat, smooth colony with white mycelium, mycelia loosely attached to the substrate.

Specimen(s) examined: Italy, Province of Forli-Cesena, Santa Sofia, near Camposonaldo, on dead branch of *Quercus* sp. (*Fagaceae*), 10 Dec. 2013, E. Camporesi, IT 1568 (holotype MFLU 17-0881, isotype BBH 42443, culture extype MFLUCC 14-0867).

Notes: The *Cytospora quercicola* clade is fully-supported by the multi-gene phylogenetic analyses (Fig. 1, Clade 16). This species is sister to *Cytospora junipericola*.

Cytospora rosae Senan., Camporesi, & K.D. Hyde, **sp. nov.** MycoBank MB821571. Facesoffungi number FoF03468. Fig. 10.

Etymology: Named after the host genus Rosa.

Saprobic on Rosa canina. Sexual morph: Stromata restricted to around the ostiolar neck, black. Ascomata 235-255 µm high, 130–150 μ m diam (\overline{x} = 240 × 140 μ m, n = 20), solitary to rarely aggregated, scattered, immersed, globose, brown, coriaceous, ostiolate, papillate. Papilla 127-140 µm high, 70-90 µm diam $(\overline{x} = 135 \times 87 \ \mu m, n = 20)$, straight or curved, long, brown, internally covered by hyaline periphyses, wall comprising elongated, thick-walled cells. Peridium 16-23 μ m diam (\overline{x} = 20 μ m, n = 20), comprising outer, thick-walled, brown cells of textura angularis and inner, compressed, thick-walled, hyaline cells of textura angularis. Hamathecium comprising septate, hyphae-like, hyaline, 1.5–2.7 μ m diam (\overline{x} = 2.5 μ m, n = 20) paraphyses. Asci $20-23 \times 3.2-3.7 \ \mu m$ ($\overline{x} = 21 \times 3.7 \ \mu m$, n = 20), unitunicate, 8spored, clavate, short-pedicellate, apex round, with apical ring. Ascospores $4.2-6.3 \times 1-1.5 \ \mu m$ ($\overline{x} = 5.5 \times 1.3 \ \mu m$, n = 20), unito biseriate, unicellular, allantoid, or ellipsoid, hyaline, smoothwalled. Asexual morph: Conidiomata 100-200 µm diam



Fig. 8. Cytospora junipericola (MFLU 17–0882). A. Ascomata on substrate. B. Vertical cross section of ascoma. C. Peridium. D–G. Asci. H. Ascospores. Scale bars: A = 500 μm, B = 100 μm, C = 20 μm, D–H = 10 μm.

 $(\overline{x} = 150 \ \mu\text{m}, n = 20)$, solitary to aggregate, immersed, pyriform to subglobose, multi-loculate, black, coriaceous, ostiolate, papillate, peridium folded into centrum. *Pycnidial walls* 4–7 μ m diam ($\overline{x} = 6 \ \mu\text{m}, n = 20$), comprising small, thick-walled, brown cells of *textura angularis*. *Conidiophores* 8–12 × 1.5–2.5 μ m ($\overline{x} = 11 \times 2 \ \mu\text{m}, n = 20$), cylindrical, shorter than conidiogenous cells, branched, hyaline. *Conidiogenous cells* 10–15 × 1–1.5 μ m ($\overline{x} = 12 \times 1.2 \ \mu\text{m}, n = 20$), phialidic, cylindrical, tapering towards the apices, bearing single conidia at each tip. *Conidia* 3–5 × 0.5–1 μ m ($\overline{x} = 2 \times 1 \ \mu\text{m}, n = 20$), aguttulate, elongated to allantoid, slightly curved, aseptate, hyaline.

Culture characteristics: Colonies growing on MEA attained 2 cm within 7 d incubated at 18 °C, filamentous, flat, filiform, middle blackish ash, margin off white, cottony, tiny mycelium clots arrange radially from centre to margin.

Specimen(s) examined: Italy, Province of Forlì-Cesena, Galeata, near Passo delle Forche, on dead branch of *Rosa canina* (*Rosaceae*), 15 Apr. 2014, E. Camporesi, IT 1814 (holotype MFLU 17-0885, isotype BBH 42447, cultures extype MFLUCC 14–0845; Province of Forlì-Cesena, Galeata, near Passo delle Forche, on dead branch of *Rosa canina* (*Rosaceae*), 4 Jan. 2016, E. Camporesi, IT 1814 (paratype MFLU 15–3596, cultures ex-paratype MFLUCC 17–1664).

Notes: Combined ITS nrDNA, LSU nrDNA, *rpb2* and *tef1* sequence data in the current study shows that *Cytospora rosae*



forms a distinct clade with high bootstrap support, basal to *Cytospora fraxinigena* (Fig. 1, Clade 16). Morphologically, *Cytospora rosae* has unique characters of solitary ascomata and small asci with septate, wide, hyaline, hyphae-like paraphyses.

Cytospora salicina Norphanphoun *et al.*, Mycosphere 8: 80. 2017. Fig. 11.

Saprobic on twigs and branches of Cornus sanguinea. Sexual morph: Undetermined. Asexual morph: Stromata appear as black pinhead spots surrounding by yellow to pale brown tissues on the substrate, immersed, rosette, labyrinthine, pale brown to black, 1-5 pycnidia in a stroma, comprising loosely packed, pale brown cells of textura globosa, ostiole. Papilla narrow, short, internally covered by periphyses, converged, black, furfuraceous. Pvcnidial locules multi-chambered, subdivided by invaginations of common pycnidial walls. Conidiomata 530-600 µm high, $600-870 \ \mu m$ diam ($\overline{x} = 570 \times 705 \ \mu m$, n = 20), solitary to aggregate, immersed, pyriform to subglobose, black, coriaceous, ostiolate, papillate, peridium folded into centrum. Papilla 200–300 μ m high, 60–95 μ m diam (\overline{x} = 210 × 80 μ m, n = 20), internally covered by hyaline filiform periphyses. Pycnidial walls 7–11 μ m diam (\overline{x} = 9.2 μ m, n = 20), comprising small, thickwalled, brown cells of textura angularis, separates from



Fig. 9. Cytospora quercicola (MFLU 17–0881). A. Ascomata on substrate. B. Cross section of ascoma. C. Peridium. D–G. Asci. H. Ascospores. Scale bars: A = 200 μm, B = 100 μm, C = 20 μm, D–H = 10 μm.

stromata at maturity. *Conidiophores* 9–15 × 1.5–2.5 µm (\overline{x} = 11.4 × 2 µm, n = 20), cylindrical, shorter than conidiogenous cells, branched, hyaline. *Conidiogenous cells* 10–20 µm high, 1–1.5 µm diam (\overline{x} = 16 × 1.3 µm, n = 20), phialidic, cylindrical, tapering towards the apices, bearing single conidia at each tip. *Conidia* 4.5–6 × 0.5–1.5 µm (\overline{x} = 5 × 1.3 µm, n = 20), eguttulate, elongated to allantoid, slightly curved, aseptate, hyaline.

Culture characteristics: Colonies growing on PDA attenuated 2 cm incubated at 18 °C within 10 d, circular, flat, entire, white, thin, slightly aerial mycelia, loosely attached to the media.

Specimen examined: Russia, Rostov Region, Krasnosulinsky District, Donskoye forestry, Kabanya Balka (Boar gully), twigs and branches of *Cornus sanguinea* subsp. *australis* (*Cornaceae*), 27 Oct. 2015, T.S. Bulgakov, R1111, MFLU 17–0891, living culture MFLUCC 16–1190.

Notes: Cytospora salicina was introduced by Norphanphoun *et al.* (2017) from Russia causing canker on *Salix* sp. However, we collected this specimen from Russia associated with twigs and branches of *Cornus sanguinea*. *Cytospora salicina* is closely related to *C. chrysosperma*, *C. melnikii*, and *C. sordida* (Fig. 1, Clade 16). Diaporthaceae Höhn. ex Wehm., Am. J. Bot. 13: 638. 1926. Clade 14.

Pathogenic, endophytic or saprobic on terrestrial and rarely submerged plants. Sexual morph: Pseudostromata well- or poorly developed, pulvinate, erumpent, flat or slightly convex, orbicular, circular or somewhat irregular, sclerotioid, coriaceous, whitish to brownish black, with or without black zone or a crust consisting of fungus tissue, solitary or containing up to 10 ascomata in a stroma. Ectostromatic disk subhyaline to brown. Ascomata perithecial, immersed to erumpent, solitary or aggregated in a valsoid configuration, globose or compressed, coriaceous, black, ostiolate, papillate. Papilla short or long, erumpent, convergent, cylindrical to conical, black, internal wall covered by hyaline periphyses, composed of vertically arranged parenchymatous tissues. Peridium comprising outer layer of flattened, thick-walled, dark-brown cells of textura angularis and inner, hyaline, thin-walled cells of textura angularis. Hamathecium comprising septate, unbranched, cylindrical paraphyses. Asci 8-spored, unitunicate, clavate, oblong-clavate to broadly fusoid, sessile, with a distinct apical ring. Ascospores biseriate to partially biseriate, ellipsoid, oblong to fusoid, unicellular or 1-



Fig. 10. Cytospora rosae (MFLU 17–0885). A. Ascomata on substrate. B. Cross section of ascoma. C. Peridium. D–I. Asci. J. Paraphysis. K. Ascospores. L. Colony on MEA upper surface. M. Colony on MEA lower surface. N–P. Conidiomata on MEA. P. Horizontal cross section of conidioma. Q–R. Peridium with conidiogeneous cells, conidiophores and conidia. S. Conidia. Scale bars: A, P–N = 500 μm, B = 50 μm, C, I, Q–R = 10 μm, D–H, J, K, S = 5 μm, P = 100 μm, O = 1 mm.

septate, constricted at septum, with or without appendages at both ends, hyaline, dark brown, sometimes narrowly rounded ends and multi-guttulate, smooth-walled. Asexual morph: *Conidiomata* acervular or pycnidial, globose, initially immersed, erumpent at maturity, solitary, scattered, coriaceous, black, elongated ostiolar neck, sometime becoming multi-loculate with one to several clearly defined black necks extending above the stroma, often with yellowish, conidial mass extruding from ostiole. *Peridium* comprising 3–4 layers of pale brown cells of *textura intricata* to *textura angularis*. *Conidiophores* sometimes dimorphic. *Alpha conidiophores* tightly aggregated, subcylindrical, branched in mid region, consisting of 2–3 supporting cells, giving rise to septate, ampulliform, cylindrical to irregular conidiogenous cells or paraphyses, straight to sinuous, 1–5-septate, cylindrical, hyaline to pale brown, branched only at the base, smooth, formed from the inner most cell layers of the conidiomatal wall, sometimes terminal and lateral, apex with minute periclinal thickening and collarette. *Beta conidiophores* interspersed among alpha conidiophores, hyaline, subcylindrical, branched, 1–3-septate. *Alpha conidiogenous cells* enteroblastic, phialidic, cylindrical or subcylindrical, terminal and lateral, slightly tapering towards the apex or sometimes apex



Fig. 11. *Cytospora salicina* (MFLU 17–0891). A. Conidiomata on substrate. B. Horizontal cross section of conidioma. C, D. Vertical cross section of conidiomata. E. Conidiophore, conidiogenous cells arrangement. F. Peridium. G–M. Conidiophores, conidiogenous cells, conidia. Scale bars: A = 500 μ m, B = 200 μ m, C, D = 100 μ m, E, F = 15 μ m, G–M = 10 μ m, N = 5 μ m.

with minute periclinal thickening and collarette. Beta conidiogenous cells phialidic, integrated, terminal and lateral. Alpha conidia abundant, fusiform, ovate, subcylindrical to narrowly ellipsoid, straight or curved, occasionally irregular, smoothwalled, 0–2-septate, hyaline, base truncate to sub-truncate, apex obtuse, straight to curved, occasionally slightly sigmoid, pale to medium brown, with many guttules, sometimes short, hyaline, appendages at both ends. *Beta conidia* subcylindrical, fusiform to hooked, straight to slightly curved, aseptate, hyaline, smooth, base sub-truncate, sometimes widest in middle or in upper third, tapering to acutely rounded apex, truncate at base.

Type genus: Diaporthe Nitschke.

Type species: Diaporthe eres Nitschke.

Notes: The family Diaporthaceae (Fig. 1, Clade 14) comprises many endophytic and phytopathogenic fungal species (Udayanga et al. 2011) and it was introduced and accommodated in Diaporthales by von Höhnel (1917). Wehmeyer (1975) confined this family to Diaporthe and Mazzantia. However, Barr (1978) synonymised Diaporthaceae under Valsaceae. Castlebury et al. (2002) analysed LSU nrDNA sequence data of diaporthoid taxa and showed the distinct placement of Diaporthaceae in Diaporthales, forming a well-supported clade. Diaporthaceae previously comprised only Diaporthe (Phomopsis) and Mazzantia based on phylogenetic analysis (Castlebury et al. 2002). However, Lumbsch & Huhndorf (2010) included Apioporthella and Leucodiaporthe in this family. A LSU nrDNA sequences analysis by Lamprecht et al. (2011) indicates placement of Stenocarpella and Phaeocytostroma within Diaporthaceae. Pustulomyces was introduced based on a combined gene analysis of LSU nrDNA, SSU nrDNA and tef1 sequence data (Dai et al. 2014). Voglmavr & Jaklitsch (2014) confirmed the phylogenetic placement of Phaeodiaporthe in Diaporthaceae based on analysis of LSU nrDNA sequence data. Maharachchikumbura et al. (2015) listed Allantoporthe, Apioporthella. Clypeoporthella, Diaporthe, Diaporthella. Diaporthopsis, Leucodiaporthe, Mazzantia, Mazzantiella, Ophiodiaporthe and Pustulomyces as genera of Diaporthaceae. Rossman et al. (2015) synonymised Mazzantiella under Mazzantia based on greater usage of Mazzantia. The genus Clypeoporthella is based on C. brencklei, and a recently collected C. brencklei (BPI 843482) specimen was grown in culture and sequenced. DNA sequence data showed that C. brencklei clustered together with Diaporthe and it has a Phomopsis asexual morph. Thus, Clypeoporthella is considered as a synonym of Diaporthe (Sogonov et al. 2008). The genus Diaporthopsis was introduced to accommodate species that are similar to Diaporthe, with unicellular ascospores and was typified by D. angelicae. Molecular analysis of LSU nrDNA sequence data showed that D. angelicae clustered within the Diaporthe. In addition. Diaporthopsis angelicae has similar morphological characters of stromata, perithecia, and centrum to species of Diaporthe. Based on morphology and molecular data, Diaporthopsis was synonymised under Diaporthe (Castlebury et al. 2003, Gomes et al. 2013). The genus Diaporthella has aggregated perithecia within well-developed stromata and median, 1septate ascospores. Diaporthella corylina is strongly parasitic and causes dieback of Corylus stems. Morphologically Diaporthella corvlina shows similar characters to Anisogramma anomala. Anisogramma based on A. virgultorum is known to belong in the Gnomoniaceae (Castlebury et al. 2002, Vasilyeva et al. 2007). However, the LSU nrDNA, ITS nrDNA, rpb2 and tef1 combined gene analyses in the current study show (Fig. 1, Clade 5) the phylogenetic placement of Diaporthella is outside of Diaporthaceae and it does not show affinities with any families in Diaporthales. Hence Diaporthaceae comprises Allantoporthe, Apioporthella, Chaetoconis, Diaporthe, Leucodiaporthe, Mazzantia, Ophiodiaporthe, Phaeocytostroma, Phaeodiaporthe, Pustulomyces and Stenocarpella. Based on an LSU nrDNA phylogeny, Gao et al. (2017) showed Diaporthe sensu lato to be polyphyletic, including genera such as Mazzantia, Ophiodiaporthe, Pustulomyces, Phaeocytostroma, and Stenocarpella. In

the present study, we address this situation by proposing *Chiangraiomyces, Paradiaporthe, Hyaliappendispora* as new genera in *Diaporthaceae*. We collected and illustrate here several taxa in *Diaporthaceae* that are new to science or are poorly studied.

Chiangraiomyces Senan. & K.D. Hyde, **gen. nov.** MycoBank MB821544. Facesoffungi number FoF03469.

Etymology: Name related to the collection locality of Chiang Rai, Thailand.

Saprobic on dead wood. Sexual morph: Ascomata solitary, scattered, immersed to erumpent, globose to subglobose, coriaceous, black, papillate, ostiolate. Papilla long, internally covered by hyaline, periphyses. Peridium comprising outer, thick-walled, brown cells of textura angularis and inner, hyaline, thick-walled, compressed cells of textura angularis. Hamathecium comprising hyaline, aseptate, filamentous paraphyses. Asci unitunicate, 8-spored, fusiform, sessile to short pedicellate, with J-, funnel-shaped, apical ring. Ascospores biseriate to overlapping uniseriate, fusiform to ellipsoid, hvaline, smooth-walled, 1-septate, with two large guttules in the centre and two small guttules at the ends. Asexual morph: Conidiomata produced on PDA when incubated at 18 °C after 2 wk, pycnidial, globose, erumpent at maturity, black, coriaceous, short neck. Conidiomatal wall comprising pale brown, thick-walled cells of textura angularis. Conidiophores ampulliform, straight, branched, septate, hyaline, smooth. Conidiogenous cells phialidic, terminal, cylindrical, slightly tapering towards the apex. Hamathecium aparaphysate. Alpha conidia aseptate, hyaline, smooth, ovate to ellipsoidal, less in amount. Beta conidia fusiform to hooked, base subtruncate, aseptate, hyaline, smooth.

Type species: Chiangraiomyces bauhiniae Senan. & K.D. Hyde.

Chiangraiomyces bauhiniae Senan. & K.D. Hyde, **sp. nov.** MycoBank MB821545. Facesoffungi number FoF03470. Fig. 12.

Etymology: Name based on the host *Bauhinia*, from which it was collected.

Saprobic on Bauhinia sp. Sexual morph: Ascomata 200-300 µm high, $150-180 \ \mu m$ diam ($\overline{x} = 230 \times 240 \ \mu m$, n = 20), solitary, scattered, immersed to erumpent, globose to subglobose, coriaceous, black, papillate, ostiolate. Papilla 115-140 µm high, 75–90 μ m diam (\overline{x} = 130 × 85 μ m, n = 20), long, internally covered by hyaline, periphyses. Peridium 11-14 µm wide $(\overline{x} = 12.5 \ \mu m, n = 20)$, comprising outer, thick-walled, brown cells of textura angularis and inner, hyaline, thick-walled, compressed cells of textura angularis. Hamathecium 2.5-3 µm wide $(\overline{x} = 2.8 \ \mu m, n = 20)$, comprising hyaline, aseptate, filamentous paraphyses. Asci 75–90 × 12–13 μ m (\bar{x} = 78 × 12.5 μ m, n = 20), unitunicate, 8-spored, fusiform, with J-, funnel-shaped, apical ring, sessile to short pedicellate. Ascospores 17-18 × 3-4 µm $(\overline{x} = 17.8 \times 3.6 \mu m, n = 20)$, biseriate to overlapping uniseriate, fusiform to ellipsoid, hyaline, smooth-walled, 1-septate, with two large guttules in the centre and two small guttules at the ends. Asexual morph: Conidiomata 300–500 μ m diam (\overline{x} = 450 μ m, n = 20), produced on PDA when incubated at 18 °C after 2 wk, pycnidial, globose, erumpent at maturity, black, coriaceous, short neck. Conidiomatal wall comprising pale brown, thick-walled cells of textura angularis. Conidiophores 4-6 × 2-4 µm





Fig. 12. Chiangraiomyces bauhiniae (MFLU 17–0964). A, B. Ascomata on substrate. C. Cross section of ascoma. D. Peridium. E. Paraphyses. F–H. Asci. I–N. Ascospores. O. Conidiomata on MEA. P, Q. Alpha and beta conidiogeneous cells attached to conidiophores. R. Beta conidia. Scale bars: A, O = 500 μm, B = 200 μm, C = 100 μm, D–H, P–R = 10 μm, I–N = 5 μm.

 $(\overline{x} = 5 \times 3 \ \mu m, n = 20)$, ampulliform, straight, branched, septate, hyaline, smooth. *Conidiogenous cells* 7–10 × 2–3 μm $(\overline{x} = 8 \times 2.3 \ \mu m, n = 20)$, phialidic, terminal, cylindrical, slightly tapering towards the apex. *Hamathecium* aparaphysate. *Alpha conidia* 3–5 × 2–4 μm ($\overline{x} = 4.7 \times 3.3 \ \mu m, n = 20$), aseptate, hyaline, smooth, ovate to ellipsoidal, less in amount. *Beta conidia* 18–38 × 1.5–2 μm ($\overline{x} = 24 \times 1.7 \ \mu m, n = 20$), fusiform to hooked, base sub-truncate, aseptate, hyaline, smooth.

Culture characteristics: Colonies growing on MEA attained 1 cm within 7 d when incubated 25 °C, fast growing, circular, irregular, flat, white, forming aerial mycelia with hyphae loosely attached to the medium.

Specimen examined: **Thailand**, Chiang Rai, Mae Fah Luang University, near University President's house, on dead twigs of *Bauhinia* sp. (*Fabaceae*), I.C. Senanayake, 25 Dec. 2014, CHUNI 81 (**holotype** MFLU 17-0964, cultures extype MFLUCC 17–1669, MFLUCC 17–1670).

Notes: Chiangraiomyces bauhiniae has immersed, solitary ascomata, fusiform asci, with a J-, funnel-shaped, apical ring, and oval to fusiform ascospores with 2 large central guttules and 2 small marginal guttules. Phylogenetically, *Chiangraiomyces bauhiniae* forms a fully-supported clade that is sister to *Ophiodiaporthe cyatheae* (Fig. 1, Clade 14). Hence, we introduce *Chiangraiomyces* to accommodate this taxon.



Fig. 13. Paradiaporthe artemisiae (MFLU 17–0886). A. Ascomata on substrate. B. Cross section of ascoma. C. Peridium. D–G. Asci. H–L. Ascospores. Scale bars: A = 200 µm, B = 100 µm, C = 20 µm, D–G, H–L = 10 µm.

Paradiaporthe Senan., Camporesi & K.D. Hyde, gen. nov. MycoBank MB821546. Facesoffungi number FoF03471.

Etymology: The name reflects the morphological similarity to *Diaporthe*.

Saprobic on dead twigs of *Artemisia* sp. Sexual morph: *Ascomata* solitary, scattered, immersed, becoming erumpent when mature, globose to subglobose, black, coriaceous, ostiolate, papillate. *Papilla* periphysate with short, wide, prominent ostiole. *Peridium* thin at the base, gradually thickening towards the neck, comprising inner, hyaline, compressed, thin-walled cells of *textura angularis*

and outer, thick-walled, brown cells of *textura angularis*. *Hama-thecium* aparaphysate. *Asci* 8-spored, unitunicate, fusiform to clavate, sessile, apex rounded with a J-, apical ring. *Ascospores* biseriate, fusiform with pointed ends, medianly 1-septate, hyaline, smooth-walled. Asexual morph: Undetermined.

Type species: Paradiaporthe artemisiae Senan., Camporesi & K.D. Hyde.

Paradiaporthe artemisiae Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821547. Facesoffungi number FoF03472. Fig. 13.


Etymology: The name reflects the host genus Artemisia.

Saprobic on dead twigs of Artemisia sp. Sexual morph: Ascomata 280-300 µm high, 180-200 µm wide (x = 290 × 190 µm, n = 10), solitary, scattered, immersed, becoming erumpent when mature, globose to subglobose, black, coriaceous, ostiolate, papillate. Papilla 135-138 µm high, 110-140 µm wide, $(\overline{x} = 136 \times 115 \ \mu m, n = 10)$, comprising filiform, hyaline periphyses with short, wide, prominent ostiole. Peridium 8-13 µm $(\overline{x} = 10 \text{ µm}, \text{ n} = 10)$, thin at the base, gradually thickened towards the neck, comprising inner, hyaline, compressed, thin-walled cells of textura angularis and outer, thick-walled, brown cells of textura angularis. Hamathecium aparaphysate. Asci $45-60 \times 11-14 \ \mu m$ (x = 51 × 13.5 \ \mu m, n = 20) 8-spored, unitunicate, fusiform to clavate, sessile, apex rounded, with a J-, bilobed, apical ring. Ascospores 14-18.5 × 4-5 µm $(\overline{x} = 16 \times 4.2 \ \mu m, n = 20)$ biseriate to overlapping uniseriate, fusiform with two small globules at the ends and two large globules at the middle of spore, medianly 1-septate, hyaline, smooth-walled. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on MEA attained 1 cm within 7 d when incubated at 18 °C, irregular, circular, flat, woolly, white, mycelia loosely attached to the substrate.

Specimen examined: **Italy**, Province of Forlì-Cesena, Bagno di Romagna, Valbonella, on dead stem of *Artemisia* sp. (*Asteraceae*), E. Camporesi, 9 Jul. 2014, IT 1982 (**holotype** MFLU 17–0886, **isotypes** BBH 42448, cultures ex-type MFLUCC 14–0850, MFLUCC 17–1663).

Notes: Paradiaporthe artemisiae has erumpent, solitary ascomata with prominent, wide papilla. Morphologically, Paradiaporthe is similar to Diaporthe. However, Paradiaporthe artemisiae forms a distinct clade which is sister to Phaeocytostroma artemisiae (Fig. 1, Clade 14). Hence, we introduce Paradiaporthe as a new genus based on morphology and phylogeny.

Hyaliappendispora Senan., Camporesi & K.D. Hyde, **gen. nov.** MycoBank MB821548. Facesoffungi number FoF03473.

Etymology: Name reflects hyaline ascospores with long appendages.

Saprobic on dead stems. Sexual morph: Ascomata solitary to aggregate, immersed, globose to subglobose, black to brown, coriaceous, ostiolate, papillate. Papilla short, wide, internally covered by hyaline periphyses. Peridium comprising outer, dark brown, thick-walled cells of textura angularis and inner, thin-walled, hyaline, compressed cells of textura angularis. Hama-thecium comprising filiform, septate, hyaline paraphyses which are longer than asci. Asci 8-spored, unitunicate, cylindrical to fusiform, short pedicellate, apex rounded with a J- apical ring. Ascospores biseriate to overlapping biseriate, oval to ellipsoid, hyaline, medianly 1-septate, multiguttulate, with appendages. Appendages at both apical and basal ends, long, thread-like, covered by loose capsule. Asexual morph: Ceolomycetous. Sporulate on PDA at 20 °C after 1 mo, crowded at colony margin, appears at pale yellow bubbles when release the

conidial mass. *Conidiomata* globose, erumpent, black. *Peridium* comprising thick-walled, pale brown cells of *textura angularis*. *Conidiophores* ampulliform, septate, branched, hyaline. *Conidiogeneous cells* phialidic, terminal, cylindrical, elongate, hyaline. *Conidia* fusiform, unicellular, hyaline, smooth.

Type species: Hyaliappendispora galii Senan., Camporesi & K.D. Hyde.

Hyaliappendispora galii Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821549. Facesoffungi number FoF03474. Fig. 14.

Etymology: The name reflects the host genus Galium.

Saprobic on dead stem of Galium sp. Sexual morph: Ascomata $395-450 \ \mu m$ high, $180-200 \ \mu m$ wide ($\overline{x} = 419 \times 190 \ \mu m$, n = 10), solitary to aggregated, immersed, globose to subglobose, black to brown, coriaceous, ostiolate, papillate. Papilla 160-210 µm high, $100-185 \,\mu\text{m}$ wide ($\overline{x} = 172 \times 158 \,\mu\text{m}$, n = 10), short, wide, internally covered by hyaline periphyses. Peridium 15-25 µm wide ($\overline{x} = 20 \mu m$, n = 10), comprising outer, dark brown, thickwalled cells of textura angularis and inner, thin-walled, hyaline, compressed cells of textura angularis. Hamathecium comprising filiform, septate paraphyses $1.5-3.5 \,\mu\text{m}$ wide ($\overline{x} = 2 \,\mu\text{m}$, n = 10), which are longer than asci. Asci 110-125 × 20-25 µm $(\overline{x} = 116 \times 21 \ \mu m, n = 20)$, 8-spored, unitunicate, cylindrical to fusiform, short pedicellate, apex rounded with a J- apical ring. Ascospores $20-25 \times 7-10 \mu m$ ($\overline{x} = 22 \times 9 \mu m$, n = 20), biseriate to overlapping biseriate, oval to ellipsoid, hyaline, medianly 1septate, multiguttulate, with appendages. Appendages $6-11 \times 2-3 \ \mu m$ ($\overline{x} = 8 \times 2.3 \ \mu m$, n = 10), at both ends, long, thread-like, covered by loose capsule. Asexual morph: Ceolomycetous. Sporulate on PDA at 18 °C after 1 mo, crowded at colony margin, appears at pale yellow bubbles when release the conidial mass. Conidiomata globose, erumpent, black. Peridium comprising thick-walled, pale brown cells of textura angularis. Conidiophores $10-15 \times 1.5-2.5 \ \mu m$ ($\overline{x} = 13 \times 2.1 \ \mu m$, n = 10), ampulliform, septate, branched, hyaline. Conidiogeneous cells $8-16 \times 1.5-3 \ \mu m$ ($\overline{x} = 11 \times 2.5 \ \mu m$, n = 20), phialidic, terminal, cylindrical, elongate, hyaline. Conidia 7.5-9.5 × 1.5-2.5 µm $(\overline{x} = 8.3 \times 2.2 \,\mu\text{m}, n = 20)$, fusiform, unicellular, hyaline, smooth.

Culture characteristics: Colonies growing on PDA incubated at 18 °C attaining 1 cm diam within 14 d, irregular, undulate, umbonate, whitish ash clots with tightly arranged, short, aerial mycelium, erumpent, globose, pale brownish, with viscous droplets produced after 7 d, when colonies incubate further, conidiomata arised on culture media, concentrated at colony margin, appears as black, coriaceous bubbles at the beginning and later become yellow, slimy bubbles with conidial mass.

Specimen examined: Italy, Province of Arezzo, Quota, near Casuccia di Micheli, on dead stem of *Galium* sp. (*Rubiaceae*), E. Camporesi, 8 Jun. 2015, IT 2925 (holotype MFLU 15–2269, isotype BBH 42450, culture ex-type MFLUCC 16–1208).

Notes: Hyaliappendispora is morphologically distinct from other genera in Diaporthaceae in having biguttulate, uniseptate,

Fig. 14. Hyaliappendispora galii (MFLU 15–2269). A–C. Ascomata on substrate. D. Cross section of ascoma. E. Peridium. F. Paraphyses. G–J. Asci. K–N. Ascospores. O. Culture in upper surface view. P. Culture in lower surface view. Q. Conidiomata on PDA. R, S. Conidiogeneous cells, conidiophores and conidia. T. Conidia. Scale bars: A–C, Q = 200 µm, D = 100 µm, E = 20 µm, F–N, R–T = 10 µm.



hyaline ascospores with long filamentous apical and basal appendages and wall of the appendages makes a ring-like ornamentation at the proximal end. Phylogenetically *Hyaliappendispora galii* forms a fully-supported distinct clade that is sister to *Phaeodiaporthe* (Fig. 1, Clade 14).

Chaetoconis polygoni (Ellis & Everh.) Clem., Gen. fung. (Minneapolis): 176. 1909. Facesoffungi number FoF03475. Fig. 15.

Synonym: Amphorula polygoni (Ellis & Everh.) Petr., Sydowia 13: 181. 1959.

Saprobic on stem of Rumex acetosa. Sexual morph: Undetermined. Asexual morph: Conidiomata 175-250 µm high, $200-275 \,\mu\text{m}$ diam ($\overline{x} = 200 \times 250 \,\mu\text{m}$, n = 20), pycnidial, scattered, immersed to erumpent, globose to sub-globose, dark brown, unilocular or multilocular, ostiolate, papillate. Peridium 20-30 µm thick, comprising several layers of inner thin-walled, hyaline, compressed cells of textura angularis and outer, thick-walled, dark brown cells of textura angularis. Ostiole one or more, circular. Conidiophores $12-25 \times 2-3.5 \mu m$ ($\overline{x} = 20 \times 3 \mu m$, n = 20), hyaline, branched, septate, smooth, with acropleurogenous conidia, formed from the inner pycnidial wall cells. Conidiogenous cells $30-45 \times 9-11 \ \mu m$ ($\overline{x} = 32 \times 9.5 \ \mu m$, n = 20), enteroblastic, phialidic, determinate, integrated, cylindrical, hyaline, smooth, with minute channel and collarette. Conidia 35-50 × 4-5 um (x = 37 × 4.5 µm, n = 20), hyaline, 2-euseptate, continuous, base obtuse, apex extended into a filiform, cellular, unbranched appendage, thin-walled, smooth, guttulate, obclavate.

Specimen examined: **Germany**, on the edge of a mixed forest, 39 m asl, sandy, acid, fresh, mesotroph, on stem of *Rumex acetosa* (*Polygonaceae*), 9 May 2013, RK. Schumacher, CHUNI 73, MFLU 17–0965.

Notes: Chaetoconis polygoni has quite different morphological characteristics compared to other taxa in *Diaporthaceae*. Molecular analyses in this study showed that our collection clustered together with *C. polygoni* (CBS 405.95; Fig 1, Clade 14). However, we could not obtain a culture and therefore extracted DNA directly from the sporocarps. The sexual morph of *Chaetoconis polygoni* was reported as *Ceriospora polygonacearum* (Barney *et al.* 2006) which was assigned to *Sordariales* (Campbell *et al.* 2003) and later Senanayake *et al.* (2015) reassigned it to *Xylariales*. However, morphologically *Ceriospora polygonacearum* does not show any affinity to *Diaporthales*.

Diaporthe litoricola Senan., E.B.G. Jones & K.D. Hyde, **sp. nov.** MycoBank MB821550. Facesoffungi number FoF03476. Fig. 16.

Etymology: The name is based on the Latin words "litore" and "cola" meaning "beach-loving" since this fungus was collected from dead branches of beach plants.

Saprobic on dead stem of sea-shore plants. Sexual morph: Ascomata 800–900 µm high, 450–600 µm diam ($\overline{x} = 880 \times 475$ µm, n = 20), solitary, scattered, immersed, globose to subglobose, dark brown, coriaceous, ostiolate, papillate. *Papilla* 380–430 µm high, 110–140 µm diam ($\overline{x} = 420 \times 130$ µm, n = 20), conspicuous, long, black, with pale yellow apex, brown, unbranched seta in apex, internally covered by hyaline, filamentous periphyses. *Peridium* 7–12 µm wide, $(\overline{x} = 9.3 \ \mu m, n = 20)$, comprising several layers of compressed, thick-walled, olivaceous to brown cells of textura angularis. Hamathecium aparaphysate or sometimes with a few cellular paraphyses. Asci 80-90 × 11-12 μ m (\overline{x} = 87.5 × 11.1 μ m, n = 20), 8-spored, unitunicate, cylindrical, pedicellate, apex rounded, with bilobed, distinct apical ring. Ascospores $16-19 \times 4.5-5 \ \mu m$ (x = 18 × 4.8 μm , n = 20), biseriate, fusiform to ellipsoid, 1-septate, hyaline, guttulate. Asexual morph: Con*idiomata* 500–900 µm high, 800–1000 um diam $(\overline{x} = 880 \times 900 \text{ µm}, \text{ n} = 20)$, produced on PDA when incubated at 18 °C after 4 wk, pycnidial, globose, initially immersed, erumpent at maturity, black, coriaceous, elongated neck, often yellowish white, with conidial cirrus extruding from ostiole. Conidiomatal wall comprising pale brown, thick-walled cells of textura angu*laris.* Conidiophores $5-7 \times 4-7 \mu m$ ($\overline{x} = 6.3 \times 5.3 \mu m$, n = 20), ampulliform, straight to sinuous, unbranched, hyaline to olivaceous, smooth. Conidiogenous cells 14.5-21 × 1.8-2.8 µm $(\overline{x} = 17.3 \times 2.3 \,\mu\text{m}, n = 20)$, phialidic, terminal, cylindrical, slightly tapering towards the apex. Hamathecium aparaphysate. Alpha conidia $13-16 \times 2.8-3.8 \ \mu m$ ($\overline{x} = 14.7 \times 3.3 \ \mu m$, n = 20), aseptate, hyaline, smooth, ovate to ellipsoidal, base subtruncate, often biguttulate. Beta conidia 1.5-2 × 18-38 um $(\overline{x} = 1.7 \times 24 \mu m, n = 20)$, fusiform to hooked, base sub-truncate, aseptate, hyaline, smooth.

Culture characteristics: Colonies growing on PDA attained 1 cm diam within 7 d when incubated at 18 °C, flat, circular, smooth, white, slightly woolly, tightly attached to media, mycelial ends unbranched.

Specimen examined: UK, Hampshire, Eastney shore, on stem of undetermined sea-shore plant, 20 Mar. 2016, E.B.G. Jones, GJ 242 (holotype MFLU 17–0874, isotype BBH 42436, cultures ex-type MFLUCC 16–1195, MFLUCC 17–1657).

Notes: Diaporthe litoricola differs morphologically from *D. maytenicola* in having large, multi-guttulate ascospores, cylindrical asci, deeply immersed, long papillate, solitary ascomata and elongate, fusiform to cylindrical alpha conidia. Phylogenetically this fungus is closely related to *Diaporthe maytenicola*, *D. decedens* and *D. nobilis. Diaporthe litoricola* forms a moderately-supported clade in this study (Fig 1, Clade 14).

Diaporthe rudis (Fr.) Nitschke, Pyrenomycetes Germanici 2: 282. 1870. Facesoffungi number FoF03477. Fig. 17.

Saprobic on dead umbelliferous stems. Sexual morph: Clypeus appears as black, wide patches, forming a black mat on substrate connecting all the ascomata and spread around the individual ascomata. Ascomata 540-620 µm high, 250-275 µm wide $(\overline{x} = 590 \times 260 \ \mu m, n = 10)$, solitary or rarely aggregated, erumpent, globose to subglobose, black, coriaceous, ostiolate, papillate. Papilla 290-375 µm high, 75-95 µm wide $(\overline{x} = 330 \times 85 \ \mu m, n = 10)$, long, asymmetrically located, straight or curved, internally covered by hyaline periphyses, with apex of papilla pale brown, swollen, blunt, sometimes slightly covered by black, mycelial mat. *Peridium* 11–16 μ m wide (\overline{x} = 14 μ m, n = 10), comprising thick-walled, brown, compressed cells of textura angularis. Hamathecium aparaphysate. Asci 43-46 × 11-12 µm $(\overline{x} = 43 \times 11.6 \ \mu m, n = 10)$, 8-spored, unitunicate, clavate to fusiform, sessile, apex rounded, with a characteristic, bilobed, Japical ring. Ascospores $11-13 \times 3-4.5 \ \mu m$ ($\overline{x} = 12 \times 3.9 \ \mu m$, n = 10), biseriate, fusiform to elongate ellipsoid, 1-median septate,



Fig. 15. *Chaetoconis polygoni* (MFLU 17–0965). A. Conidiomata on substrate. B, D. Cross section of conidioma. C. Peridium. E–H. Conidiogenous cells attached to conidia. I–K. Conidia. Scale bars: A = 1 mm, B, D = 100 μ m, C, E–H = 50 μ m, I–K = 10 μ m.



Fig. 16. *Diaporthe litoricola* (MFLU 17–0874). **A–C.** Ascomata on substrate. **D.** Cross section of ascomata. **E.** Peridium. **F–I.** Asci. **J–L.** Ascospores. **M, N.** Conidioma. **O, P.** Conidiophores, conidiogenous cells and conidia arrangement. **Q.** Alpha conidia. **R, S.** Beta conidia. Scale bars: A = 500 μm, B–D, M, N = 200 μm, E = 50 μm, F–I, O, P = 20 μm, J–L, Q, S = 10 μm.



Fig. 17. Diaporthe rudis (MFLU 17–0895). A–C. Ascomata on substrate. D–E. Cross sections of ascoma. F. Peridium. G–J. Asci. K–P. Ascospores. Scale bars: A = 500 μ m, B, C = 200 μ m, D, E = 100 μ m, F = 20 μ m, G–P = 10 μ m.

with each cell containing two guttules, hyaline, smooth-walled. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on PDA attaining 2.5 cm diam within 10 d when incubated at 18 °C, circular, entire, flat, white, tightly attached to the media, aerial mycelia less or sparse, forming few, erumpent, globose, black, viscous droplets after 7 d.

Specimen examined: **UK**, Hampshire, Winchester, Whiteley, Botley Wood, on umbelliferous stem, 25 May 2016, E.B.G. Jones, GJ 301 (MFLU 17–0895, BBH 42452, living cultures MFLUCC 16–1197, MFLUCC 17–1658).

Notes: Diaporthe rudis was epitypified by Udayanga et al. (2014) based on morphology and phylogeny. Diaporthe rudis has a broad host range. This collection was obtained from umbelliferous woody stems and it forms very long, curved, narrow, papilla deeply immersed in substrate. They appear as pale yellow spots with black margins. Ostioles are blunt and covered by pale yellow cells. However, the base of the ascomata is immersed in deep layers of substrate. All *D. rudis* cluster together and phylogenetic affinities of these species are still unclear, but morphologically they are differing in terms of asci and ascospore morphology and size.

Diaporthe eres Nitschke, Pyrenomycetes Germanici 2: 245. 1870. Fig. 18.

Saprobic on stem of *Fraxinus pennsylvanica*. Sexual morph: Not observed. Asexual morph: Conidiomata 125-140 µm high, 265-300 μ m diam at base (\bar{x} = 135 × 280 μ m, n = 10), pycnidial, pyriform, initially immersed, erumpent at maturity, globose to pyriform, black, coriaceous, elongated neck, often with vellowish white, conidial cirrus extruding from ostiole. Conidiomatal wall 34–36 μ m diam (\overline{x} = 35 μ m, n = 10), parenchymatous, consisting of 4-7 layers of pale brown, thicktextura angularis. walled cells of Conidiophores $4-6 \times 4.5-8 \ \mu m$ ($\overline{x} = 4.6 \times 6.6 \ \mu m$, n = 20), ampulliform, straight to sinuous, unbranched, hyaline, smooth. Con*idiogenous cells* $8-14 \times 1.5-3 \ \mu m$ ($\overline{x} = 11.2 \times 2.2 \ \mu m$, n = 20), phialidic, terminal, cylindrical, slightly tapering towards the apex. Hamathecium aparaphysate. Alpha conidia $5.8-7.5 \times 2.5-3.5 \ \mu m$ (x = 6.4 × 2.8 \ \mu m, n = 10), aseptate, hyaline, smooth, ovate to ellipsoidal, base subtruncate, often biguttulate. Beta conidia not seen.

Culture characteristics: Colonies growing on MEA attenuated 2 cm within 10 d incubated at 18 °C, fast growing, entire, flat circular, white, with radially arranged minute mycelium clots later becoming creamy or pale yellow.

Specimen examined: Russia, Cotton Fabric urban micro district, on stem of *Fraxinus pennsylvanica* (Oleaceae), 14 May 2015, T.S. Bulgakov, T-400 (MFLU 15–2104, MFLU 17–0890, living cultures MFLUCC 17–1667, MFLUCC 17–1668).

Notes: Phylogeny depicts a close association between the two *D. eres* strains collected from *Fraxinus pennsylvanica*. In this study, *D. eres* has been treated as a "complex". It is noted herein that combined gene phylogeny also support such a complex as strains from different hosts/ regions are phylogenetically apart. We did not see beta conidia for this strain on the host or in culture.

Erythrogloeaceae Senan., Maharachch. & K.D. Hyde, fam. nov. MycoBank MB821551. Facesoffungi number FoF03478. Clade 9.

Foliicolous associated with leaf spots. Sexual morph: Undetermined. Asexual morph: Conidiomata epiphyllous, subepidermal, sometime eustromatic, acervular or subglobose, brown to black or yellow-orange, amphigenous, opening by irregular rupture, wall of 2-6 layers of orange-brown textura angularis, exuding slimy orange masses of conidia. Conidiophores reduced to conidiogenous cells. Conidiogenous cells lining the inner cavity of conidioma, hyaline to olivaceous, smooth, subcylindrical to ampulliform, tapering to a long, thin neck, at times apical part elongated into a long neck, proliferating several times percurrently near apex, with flaring collarettes, or apex truncate, with minute periclinal thickening. Conidia hyaline to olivaceous, smooth, guttulate or not, thin-walled, ellipsoid, fusoid, ovoid to somewhat obclavate, straight to curved, apex subobtuse, obtusely rounded, base truncate, with prominent marginal frill, or dimorphic, intermixed in same conidiomata. Macroconidia broadly ellipsoid to obovoid, hyaline, smooth, granular to guttulate, thick-walled, apex obtuse, base flattened, Microconidia hvaline, smooth, guttulate, fusoid-ellipsoid, acutely rounded at apex, truncate at base.

Type genus: Erythrogloeum Petr.

Type species: Erythrogloeum hymenaeae Gonz. Frag. & Cif. ex Petr.

Notes: Phylogenetic analyses from the current study based on combined LSU nrDNA, ITS nrDNA, *rpb2*, and *tef1* sequences showed that *Chrysocrypta* is basal to *Disculoides* and *Erythrogloeum* (Fig. 1, Clade 9). *Chrysocrypta* has previously been accommodated in the *Cryphonectriaceae* but the latter is distantly related (Fig. 1, Clade 6). Morphologically members of clade 9 depicts distinct characters in having epiphyllous acervuli, and subcylindrical to ampulliform conidiogenous cells. The sexual morphs of those taxa have not been reported. *Disculoides* was introduced and typified by *Disculoides eucalyptorum* (Crous *et al.* 2012a). *Disculoides eucalyptorum* was shown to be distinct from *Erythrogloeum hymenaeae*, which was sister to the *Greeneria-Melanconiella* complex based on rDNA sequence gene analyses (Crous *et al.* 2012a).

Chrysocrypta was introduced based on *Chrysocrypta corymbiae*, which was isolated from leaves of *Corymbia* species. *Chrysocrypta* is similar to *Foliocryphia* (*Cryphonectriaceae*), but is distinct in forming dimorphic conidia. Crous *et al.* (2012c) accommodated this taxon in *Cryphonectriaceae* based on morphology and rDNA sequence phylogeny. However, stromatic tissues of *Chrysocrypta corymbiae* do not turn purple with KOH, which is a basic characteristic of *Cryphonectriaceae*. In addition, DNA sequence data herein indicate that *Chrysocrypta corymbiae* does not belong in *Cryphonectriaceae*. Hence given the morphological distinctiveness and strongly supported clade (9), a new family *Erythrogloeaceae* is introduced to accommodate *Chrysocrypta, Disculoides* and *Erythrogloeum*.

Erythrogloeum hymenaeae Gonz. Frag. & Cif. ex Petr., Sydowia 7: 379. 1953. Facesoffungi number FoF03479. Fig. 19.

Foliicolous, associated with leaf spots. Sexual morph: Undetermined. Asexual morph: Conidiomata up to 250 µm



Fig. 18. Diaporthe eres (MFLU 15–2104). A, B. Conidiomata on substrate. C. Cross section of conidioma. D. Wall of conidioma. E-H. Conidiophore, conidiogenous cell attached to conidia. I, J. Conidia. Scale bars: A = 1 mm, B = 200 µm, C = 100 µm, D = 20 µm, E-H = 10 µm, I, J = 5 µm.

diam, acervular, epiphyllous, eustromatic, subepidermal, solitary, rupturing surface by irregular splits. *Peridium* comprises thin-walled cells of *textura angularis*. *Conidiophores* reduced to conidiogenous cells. *Conidiogenous cells* $5-10 \times 2.5-4 \mu$ m, hyaline, smooth, phialidic with periclinal

thickening, discrete, lageniform to cylindrical, lining the inner walls of cavity. *Conidia* 7–9 × 2.5–3 μ m, hyaline, smooth, guttulate or not, thin-walled, ellipsoid to ovoid, apex obtusely rounded, tapering to a truncate base (description based on Crous *et al.* 2012a).



Fig. 19. Erythrogloeum hymenaeae (F45467). A. Packet of specimen. B. Herbarium specimen. C-D. Conidiomata on substrate. E. Peridium in KOH. F-H. Conidiophores, Conidiogeneous cells and conidia. I. Conidia. Scale bars: C = 100 µm, D = 200 µm, E-G = 10 µm, H-I = 5 µm.

Specimen examined: Costa Rica, San José, on leaves of *Hymenaea courbaril*, Nov. 1929, H. Schmidt (F45468 syntype).

Notes: The monotypic genus *Erythrogloeum* comprises the type species *Erythrogloeum hymenaeae*, which is validly described based on *Phyllosticta hymenaeae* by Petrak (1953). *Erythrogloeum hymenaeae* is associated with a severe anthracnose of apical twigs and seedlings of *Hymenaeae* species (Ferreira *et al.* 1992). This fungus has been reported from in Brazil and Costa Rica.

Gnomoniaceae G. Winter [as 'Gnomonieae'], Rabenh. Krypt.-Fl., Edn 2 (Leipzig) 1.2: 570. 1886. Clade 1.

Saprobic on bark and leaves of overwintered plants. Sexual morph: *Stromata* lacking, or poorly to well-developed, scattered, erumpent, pustuliform with one or rarely two ascomata or valsoid, broadly elliptic to rounded, large. *Ectostromata* well-developed, brown to black, thick ectostromatic disc at perithecial necks. *Ascomata* immersed to erumpent, solitary or aggregated, globose to subglobose, black, coriaceous, thinwalled, with one or more long, central or eccentric necks with hyaline periphyses. *Peridium* comprising few layers of

brown, thick-walled cells of textura angularis. Hamathecium comprising few hyaline, septate, cellular paraphyses. Asci 8-32-spored, unitunicate, oval, fusiform to almost filiform, short pedicellate, with a distinct, J- apical ring. Ascospores biseriate, overlapping uniseriate to fasciculate, oval, fusiform, ovoid to subulate, small, unicellular to 1-septate, rarely multiseptate, ends mostly rounded, rarely pointed, appendages absent or subulate, navicular or whip-shaped, smooth. Asexual morph: Conidiomata acervulal or pycnidial, subcuticular, papillate or not, oblate to globose, black, thick-walled, with one chamber containing whitish conidial mass. Conidiophores simple, filiform to fusiform, annellation visible or invisible, densely branched. Conidiogenous cells usually phialidic, rarely with a few annellidic scars, irregular in shape, lageniform to cylindrical, gradually tapering to ends for one guarter to threequarters of their length, or abruptly narrowing to long neck at about half of the phialide length, or abruptly narrowing at apex, straight or curved, sometimes asymmetric swollen nodes, proliferating into other conidiogenous cells at basal or middle part. Conidia broadly ellipsoid to oval, sometimes obovoid, allantoid, occasionally curved or sinuate to slightly angular, hyaline, often unicellular.



Fig. 20. Plagiostoma salicicola (MFLU 17–0878). A. Ascomata on substrate. B. Ascomata distribution on bark epidermis (under surface). C. Cross section of ascoma. D. Papilla. E. Peridium. F–H. Asci. I–L. Ascospores. M. Culture on PDA, upper surface view. N. Culture on PDA, lower surface view. Scale bars: A, B = 500 μ m, C = 100 μ m, D = 50 μ m, E–L = 10 μ m.

Type genus: Gnomonia Ces. & De Not.

Type species: Gnomonia gnomon (Tode) J. Schröt.

Notes: Gnomoniaceae (Fig. 1, Clade 1) was introduced by Winter (1886). This family is characterised by immersed, rarely erumpent or superficial ascomata, without a stroma or aggregated with a rudimentary stroma. Species in *Gnomoniaceae* inhabit various hosts and substrates, including herbaceous plants, shrubs and trees as endophytes, pathogens and saprobes (Rossman *et al.* 2007, Walker *et al.* 2012). Pathogenicity of gnomoniaceous taxa is quite diversified, causing various diseases on plants. However, most gnomoniaceous species are restricted to overwintered plants in temperate and subtropical biomes.

Maharachchikumbura et al. (2015) accommodated 33 genera in *Gnomoniaceae*. Additionally, we introduce a new genus *Marsupiomyces* based on *M. quercina* and the second species *M. epidermoidea*. However, we exclude five genera from the family based on morphology and phylogeny and also included the additional genera *Mamianiella* and *Marsupiomyces* within this family. Hence, we accept 30 genera in this family: *Alnecium*, *Ambarignomonia*, *Amphiporthe*, *Anisomyces*, *Apiognomonia*, *Apioplagiostoma*, *Asteroma*, *Bagcheea*, *Cryptosporella*, Cylindrosporella, Diplacella, Ditopella, Ditopellopsis, Gloeosporidina, Gnomonia, Gnomoniella, Gnomoniopsis, Mamianiella, Marsupiomyces, Millerburtonia, Occultocarpon, Ophiognomonia, Phragmoporthe, Phylloporthe, Plagiostoma, Pleuroceras, Sirococcus, Spataporthe, Uniseta and Valsalnicola. Here we introduce, describe and illustrate new fungal taxa which belong to Gnomoniaceae.

Doubtful genera or genera excluded from *Gnomoniaceae*

Anisogramma was introduced and typified by Anisogramma virgultorum, and almost all characters of this genus are similar to Mamianiella. De Silva et al. (2009) analysed the phylogenetic relationship of Anisogramma species based on LSU nrDNA sequence data and reported its placement outside of Gnomoniaceae. Both Mamianiella and Anisogramma commonly occur on Corylus species. Combined LSU nrDNA, ITS nrDNA, rpb2 and tef1 gene analyses of the present study show that Mamianiella is nested in between Anisogramma species. Morphological comparison also reveals that both genera are characterised by Mamianiella and hence Mamianiella does not warrant generic status with high bootstrap support value (Fig. 1, Clade 1).



Mamianiella is an older name than Anisogramma. Therefore, we synonymise Anisogramma under Mamianiella giving priority to the older name. Mamianiella Höhn. was introduced and typified by *M. coryli*, (based on Sphaeria coryli) and Mamiania was introduced and typified by *M. fimbriata* (based on Sphaeria fimbriata). Von Arx & Muller (1954) suggested to retain both genera as one genus. However, Barr (1978) separated these two genera based on ascospore morphology as Mamianiella has unicellular ascospores, while Mamiania produces apiosporous ascospores. This is, however, not a strong character to differentiate these two genera, while almost all other characters are similar to each other. Hence, we synonymise Mamiania under Mamianiella giving priority to the older name Mamianiella.

Clypeoporthe, was reduced to synonymy in *Gnomonia* by Monod (1983). However, some species in this genus have eutypelloid configuration of ascomata in parenchymatous stromatic tissues. However, it is necessary to obtain DNA sequence data to resolve this genus. *Depazea* was typified by *D. frondicola* and it was assigned to *Mycosphaerellaceae* as *Sphaerulina frondicola* (Verkley *et al.* 2013). Hence, we exclude *Depazea* form *Gnomoniaceae*.

Phylloporthe, a plant parasitic, monotypic genus was introduced and is typified by *P. vernoniae*. There is no molecular data for *P. vernoniae* and it is not clear whether this genus belongs to *Gnomoniaceae* or not. Hence, we maintain this genus in *Gnomoniaceae* until molecular data for the type species are available.

Skottsbergiella was introduced and typified by *Skottsbergiella diaporthoides* which has large perithecia immersed in massive, externally crustose, pseudoparenchymatous stromata. Petrak (1971) assigned this genus to eutypoid fungi based on its stromatic consistency. This genus is morphologically similar to *Diaporthella*, which is placed in *Diaporthales incertae sedis* (Barr 1978). *Skottsbergiella diaporthoides* was renamed as *Diaporthe diaporthoides* and accommodated in *Diaporthaceae* (Barr 1978). Hence *Skottsbergiella* is not a valid genus.

Xenotypa is typified by X. aterrima. We observed a specimen of X. aterrima (as Hydnum aterrima, from S under accession no: F130640) on account of the elongated allantoid ascospores and solitary to aggregated ascomata, this taxon has closer affinity to Cytosporaceae than Gnomoniaceae. Hence, we exclude Xenotypa from Gnomoniaceae and include it in Cytosporaceae, until molecular data is available to confirm the placement.

Zythia is typified by *Z. resinae* which is synonymised under *Sarea resinae*. Molecular data demonstrate a placement of *Sarea resinae* within *Trapeliaceae* (*Baeomycetales*, *Ostropomycetidae*). Therefore, here we exclude *Zythia* from *Gnomoniaceae*. However, *Z. fragariae* shows an affinity to *Gnomoniaceae*. It is a common parasite on strawberry and Shipton (1967) reported *Zythia fragariae* as the asexual morph of *Gnomonia fragariae*. Walker *et al.* (2010) synonymised *Gnomonia fragariae* in *Gnomoniopsis* as *G. comari*. Hence *Zythia* is not considered to be a genus in *Gnomoniaceae*.

Plagiostoma salicicola Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821552. Facesoffungi number FoF03480. Fig. 20. *Etymology*: Based on the host genus *Salix* on which this fungus occurs and the Latin "-cola" which means loving.

Saprobic on dead twigs of Salix sp. Sexual morph: Stromata loosely packed comprising pseudoparenchymatous tissues. Ascomata 400-600 µm high, 250-400 um diam (\overline{x} = 580 × 300 µm, n = 20), perithecial, aggregated in groups of 3-10, immersed, oblate globose when moist and become convex with irregular dents around base of papilla when dry, coriaceous, black, ostiolate, papillate, Necks 420-700 µm long, 100-150 µm wide at base, 60-150 µm wide at apex, converged or not, eccentric to marginal, slightly curved. Asci $45-70 \times 10-20 \ \mu m$ ($\overline{x} = 62 \times 16 \ \mu m$, n = 20), 8-spored, unitunicate, fusiform, apex narrowly obtuse, sessile, with J- apical ring. Ascospores $15-25 \times 4-7 \ \mu m \ (\overline{x} = 17 \times 6 \ \mu m, \ n = 20)$, obliquely biseriate to fasciculate, ellipsoidal to fusiform, medianly 1-septate, constricted or not at the septum, ends rounded to tapering, with upper cell slightly wider than basal cell, hyaline. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on PDA attained 1 cm within 10 d incubated at 18 °C, circular, umbonate, undulate, white median region with ash outer margin, woolly, loosely attached to the substrate.

Specimen examined: **Italy**, Province of Trento, Val di Sole, near Croviana, on dead branch of *Salix* sp., 29 Jul. 2013, E. Camporesi, IT 1394 (**holotype** MFLU 17–0878, **isotype** BBH 42440, cultures ex-type MFLUCC 13–0656); Province of Trento, Val di Sole, near Croviana, on dead branch of *Salix* sp., 29 Jul. 2013, E. Camporesi, IT 1394 (**paratype** MFLU 15–2261, living cultures MFLUCC 17–1666).

Notes: Mejía et al. (2011) revisited the genus *Plagiostoma* and observed distinct grouping pattern of *Plagiostoma* species with expanded necks and species with cylindrical necks on *Salicaceae*. With species with expanded necks, *P. salicicola* is morphologically similar to *Plagiostoma dilatatum*. However, *P. dilatatum* has relatively small ascomata with short necks and long-pedicellate, cylindrical asci. The combined gene sequence analyses herein indicate a relationship of *P. salicicola* with other species of *Plagiostoma* separated with moderate support values, but sufficiently distinct of *P. dilatatum* (Fig. 1, Clade 1).

Plagiostoma jonesii Senan., & K.D. Hyde, **sp. nov.** MycoBank MB821553. Facesoffungi number FoF03481. Fig. 21.

Etymology: In honour of Prof. Gareth Jones, an eminent mycologist who collected this species.

Saprobic on umbelliferous stems. Sexual morph: Ascomata $380-420 \mu m$ high, $250-280 \mu m$ diam ($\overline{x} = 400 \times 270 \mu m$, n = 10), solitary or rarely aggregated, erumpent, globose to subglobose, black, coriaceous, ostiolate, papillate. Papilla 165-260 µm high, 70–100 μ m wide (\overline{x} = 200 × 80 μ m, n = 10), short, symmetrically or asymmetrically located, narrow at the base, widening towards the top, straight or curved, internally covered by hyaline periphyses. Peridium 15–25 μ m wide (\overline{x} = 18.5 μ m, n = 10), comprises thick-walled, brown, compressed cells of textura *porrecta.* Asci 40–50 × 8.5–9.5 μm (x = 48 × 8.8 μm, n = 10), 8spored, unitunicate, fusiform to clavate, apex with J-, bilobed, distinct apical ring, short pedicellate. Ascospores $12-14 \times 2.6-3.8 \,\mu m$ (x = 13 × 3.2 μm , n = 10), biseriate, fusiform to ellipsoid, hyaline, 1-septate, with two globules in each cell, with



Fig. 21. Plagiostoma jonesii (MFLU 17-0873). A. Ascomata on host surface. B. Cross section of ascoma. C. Peridium. D-G. Asci. H-L. Ascospores. Scale bars: A = 200 µm, B = 100 µm, C = 20 µm, D-L = 10 µm.

small spine-like appendages at both ends. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on PDA attenuated 1 cm within 7 d, incubated at 18 °C, fast-growing, flat, circular, smooth, less in aerial mycelia, white, tightly attached to the medium.

Specimen examined: UK, Sussex Ocidental, Arundel, river bank, on umbelliferous stem, 17 Feb. 2016, E.G.B. Jones, GJ 227 (holotype MFLU 17–0873, isotype BBH 42435, cultures ex-type MFLUCC 16–1189, MFLUCC 17–1654).

Notes: Plagiostoma jonesii is morphologically and phylogenetically distinct from other *Plagiostoma* species in having long, curved papilla arising out from the substrate appearing as spines and the opening is wider than base, ellipsoid to fusiform, 1septate, slightly or non-constricted ascospores with small, appendages. Our phylogeny shows that *Plagiostoma jonesii* is phylogenetically close to *P. salicellum* and *P. populinum*, but morphologically distinct from both species.

Gnomoniopsis agrimoniae Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821554. Facesoffungi number FoF03482. Fig. 22.

Etymology: Species epithet based on the host genus Agrimonia.

Saprobic on dead stems of *Agrimonia eupatoria*. Sexual morph: *Ascomata* 200–320 µm high, 245–400 µm diam ($\overline{x} = 273 \times 332$ µm, n = 20) solitary, scattered, erumpent, globose, black, coriaceous, ostiolate, papillate. *Papilla* 100–170 µm high,



Fig. 22. Gnomoniopsis agrimoniae (MFLU 17–0884). A, B. Ascomata on host substrate. C. Cross section of ascoma. D. Peridium. E–G. Asci. H. Ascospores. I. Culture growing on MEA, upper surface view. J. Culture growing on MEA, lower surface view. Scale bars: A = 100 μm, B = 200 μm, C = 50 μm, D = 10 μm, E–H = 5 μm.

70–105 µm diam ($\bar{x} = 160 \times 80$ µm, n = 20), short, comprising elongate brown cells of *textura porrecta*. *Peridium* 35–45 µm ($\bar{x} = 39$ µm, n = 10) comprising inner, hyaline, compressed cells of *textura angularis* and outer, brown, thick-walled, cells of *textura globosa*. *Asci* 28–32.5 × 5–5.5 µm ($\bar{x} = 30.5 \times 34.9$ µm, n = 20), 8spored, unitunicate, cylindrical to fusiform, short-pedicellate, apex obtuse with bilobed, J- apical ring. *Ascospores* 7–8 × 1.8–2.2 µm ($\bar{x} = 7.5 \times 2.1$ µm) overlapping uni- to biseriate, apiosporous, hyaline, uniseptate, smooth-walled. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on MEA becoming 2 cm within 7 d incubated at 18 °C, fast growing, circular, smooth mycelia concentrated at margins making a concave colony, off white, loosely attached to the substrate, wooly.

Specimen examined: Italy, Province of Forli-Cesena, near Santa Sofia, on dead stem of Agrimonia eupatoria (Rosaceae), 5 Apr. 2014, E. Camporesi, IT 1798 (holotype MFLU 17–0884, isotype BBH 42446, cultures ex-type MFLUCC 14–0844, MFLUCC 17–1662).

Notes: Gnomoniopsis agrimoniae has minute asci and ascospores compared to the other Gnomoniopsis species. Coriaceous, thick-walled ascomata and small apiosporous ascospores are prominent characters in this genus. Our combined gene analyses indicate a moderately supported phylogenetic distinction of *Gnomoniopsis agrimoniae* from other species with moderate support. Phylogeny analyses based on ITS sequence data following Walker *et al.* (2010) reported that *Gnomoniopsis agrimoniae* is distinct from other *Gnomoniopsis* species. *Gnomoniopsis* species are considered host specific and only *Gnomoniopsis agrimoniae* and *G. guttulata* are reported on *Agrimonia* species.

Apiognomonia veneta (Sacc. & Speg.) Höhn., Hedwigia 62: 47. 1920. Facesoffungi number FoF03483. Fig. 23.

Basionym: Laestadia veneta Sacc. & Speg., Michelia 1(no. 3): 351. 1878.

Pathogenic on living leaves of *Platanus acerifolia*. Sexual morph: Undetermined. Asexual morph: *Conidiomata* 180–200 µm high, 250–265 µm diam (\bar{x} = 188 × 260 µm, n = 10), acervular, irregularly round or oval, erumpent to immersed, solitary, scattered, conidiogenous layer covering the entire inner surface of acervular chambers and mostly in basal layer, yellowish-brown, initially developing under epidermis, then breaking through epidermis and forming thick whitish amorphous conidial masses. *Conidiophores* 10–15 × 2.5–4.5 µm (\bar{x} = 12 × 3.4 µm, n = 20), densely branched, ampulliform, hyaline. *Conidiogenous cells*



Fig. 23. Apiognomonia veneta (MFLU 15–3710). A–B. Conidiomata on host leaf surface. C. Cross section of conidioma. D–F. Conidiophore and conidiogenous cells attached to conidia. G, H. Conidia. I. Upper surface view of culture. J. Lower surface of culture. Scale bars: A, B = 1 mm, C = 100 μm, D–H = 10 μm.

14–21 × 2.7–3.5 µm (\overline{x} = 18 × 3 µm, n = 10), usually phialidic, rarely annellidic, lageniform to cylindrical, gradually tapering towards the apex, straight or curved, hyaline, smooth. *Conidia* 12–20 × 4–6 µm (\overline{x} = 15.6 × 5 µm, n = 10), broadly ellipsoid to oval, sometimes obovoid, occasionally curved or sinuate to slightly angular, hyaline, thick-walled, aseptate, guttulate.

Culture characteristics: Colonies growing on MEA attenuated 1 cm within 7 d, incubated at 18 °C, flat, circular, irregular, with circular ornamentations, margins concentrated with mycelial ends, white, rich in short aerial mycelia, loosely attached to the medium.

Specimen examined: Russia, Rostov region, Krasnosulinsky district, Donskoye forestry, lining-out nursery, on live leaves of *Platanus acerifolia (Platanaceae)*, 27

Oct. 2015, T.S. Bulgakov, R 1048, MFLU 15-3710, living cultures MFLUCC 16-1193, MFLUCC 17-1656.

Notes: Apiognomonia veneta is a common pathogen on *Plata-naceae*. Here we illustrate the asexual morph of *Apiognomonia veneta*. This is a common epifoliar pathogen. We could not obtain the sexual morph in culture or from the specimen.

Marsupiomyces Senan. & K.D. Hyde, gen. nov. MycoBank MB821555. Facesoffungi number FoF03484.

Etymology: Referring to the ascomata located in mycelial cavity not in stromatic tissues.

Saprobic on leaves of *Fagaceae*. Sexual morph: Appearing on the surface as black solitary swellings on the leaf surface. *Ascomatal cavity* pale in colour, tightly packed cells, forming a thin coating around ascomata. *Ascomata* solitary, scattered, immersed horizontally in the lower and upper leaf epidermis, globose to subglobose, coriaceous, black, ostiolate, papillate. *Papilla* long, asymmetrically located, slanted or on substrate, curved or erect. *Peridium* comprising thick-walled, brown, large cells of *textura globulosa* or *textura epidermoidea*. *Hamathecium* aparaphysate. *Asci* 8-spored, unitunicate, fusiform, with short, pointed pedicel, apex rounded with bi-lobed, distinct, apical ring. *Ascospores* uni- to tri-seriate, fusiform, cylindrical to elongate fusiform, straight or very slightly curved, 1-septate, hyaline, guttulate, smooth-walled. Asexual morph: Undetermined.

Type species: Marsupiomyces quercina Senan., Camporesi & K.D. Hyde.

Notes: Marsupiomyces is introduced and typified by *M. quercina*. Members of this genus occur on members of *Fagaceae*. Marsupiomyces comprises *M. quercina* and *M. epidermoidea*. Phylogenetically Marsupiomyces is closely related to Apioplagiostoma (Fig. 1, Clade 1). However, Apioplagiostoma differs from Marsupiomyces in having leaf lesions with dark purple to brown pigmentation, and apiosporous ascospores.

Marsupiomyces quercina Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821556. Facesoffungi number FoF03485. Fig. 24.

Etymology: Species epithet based on the host genus Quercus.

Saprobic on leaves of *Quercus*. Sexual morph: *Stromatic cavity* pale in colour, tightly packed cells, forming a thin, coating around ascomata. *Ascomata* 150–250 µm high 160–300 µm diam ($\bar{x} = 175 \times 200 \text{ µm}$, n = 10) solitary, scattered, immersed horizontally in the lower and upper leaf epidermis, globose to sub-globose, coriaceous, black, ostiolate, papillate. *Papilla* long, asymmetrically located, slanted or on substrate, curved or erect. *Peridium* 25–45 µm wide ($\bar{x} = 35 \text{ µm}$, n = 10), comprising thick-walled, brown, large cells of *textura globulosa*. *Hamathecium* aparaphysate. *Asci* 125–150 × 9–11 µm ($\bar{x} = 134 \times 9.8 \text{ µm}$, n = 30), unitunicate, 8-spored, fusiform, with short, pointed pedicel, apex rounded with bilobed, distinct, apical ring. *Ascospores* 15–21 × 6–8 µm ($\bar{x} = 17.4 \times 6.6 \text{ µm}$, n = 40), biseriate, cylindrical to elongate fusiform, 1-septate, hyaline, guttulate. Asexual morph: Undetermined.

Specimen examined: **Italy**, Province of Forlì-Cesena, San Paolo in Alpe, Santa Sofia, dead leaves of *Quercus* sp. (*Fagaceae*), 2 May 2013, E. Camporesi, IT 1214 (**holotype** MFLU 17–0876, **isotype** BBH 42438, cultures ex-type = - MFLUCC 14–0566, MFLUCC 13–0664).

Notes: The combined ITS nrDNA, LSU nrDNA, *rpb2* and *tef1* sequences analyses of this study shows that *Marsupiomyces quercina* forms a distinct clade which is sister to *Marsupiomyces epidermoidea* (Fig. 1, Clade 1).

Marsupiomyces epidermoidea R.H. Perera, Senan., Bulgakov & K.D. Hyde, **sp. nov.** MycoBank MB821557. Facesoffungi number FoF03486. Fig. 25.

Etymology: Fungal peridium comprising cells of *textura epidermoidea*.

Saprobic on dead leaves of Quercus robur. Sexual morph: Appearing on the surface as black solitary swellings on the leaf surface. Ascomatal cavity pale in colour, tightly packed cells, forming a thin, coating around ascomata. Ascomata 200-310 µm diam, depressed globose to irregular. Peridium 11-36 µm thick, comprising 3-8 layers of brown to hyaline cells of textura epidermoidea, outer cell laver brown to pale brown, inner cells hyaline, elongate. Asci 54–83 × 11–15 μ m (\overline{x} = 71 × 14 μ m, n = 25), 8-spored, unitunicate, clavate, apedicellate, with a J- refractive lying without paraphyses. apical ring, Ascospores $18-21 \times 3-3.6 \ \mu m$ (x = 19 × 3.4 μm , n = 30), uni- to tri-seriate, 1septate, not constricted at the septum, broadly fusiform, rounded at both ends, straight or very slightly curved, hyaline, guttulate, smooth-walled. Asexual morph: Undetermined.

Specimen examined: Russia, Rostov region, Shakhty city, Maisky, Cemetery Park, (47.6922302° E, 40.0925446° N), on dried leaf of *Quercus robur (Fagaceae)*, 21 Jun. 2015, T.S. Bulgakov, T 776 (holotype MFLU 15–2921, isotype BBH 42451).

Notes: Our new taxon, *Marsupiomyces epidermoidea* is a sister taxon to *Marsupiomyces quercina*, but sufficiently distinct. In addition, it is different from *Marsupiomyces epidermoidea* in having a distinct peridium comprising cells of *textura epidermoidea*.

Ditopella biseptata R.H. Perera, Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821558. Facesoffungi number FoF03487. Fig. 26.

Etymology: Species name refers to the ascospores that have two septa.

Saprobic on dead branch of Alnus glutinosa. Sexual morph: Stromata surrounding the perithecial necks, extending outward beneath the host periderm as a distinct clypeus, composed of dark brown thick-walled angular cells. Ascomata 500-900 um diam, immersed in the ectostroma, situated between the epidermis and the cortex of the host tissue, appearing as solitary swellings of the host epidermis, sometimes epidermis ruptures to expose the rounded apex of the ostiole, perithecial, depressed globose to oval, ostiolate. Ostiolar neck lined with thin-walled hyaline, septate periphyses. Peridium 44 µm thick, 2-layered, outer layer composed of angular, sometimes slightly compressed, dark brown, thick-walled cells of textura angularis, inner layer of elongate, hyaline, thin-walled, compressed cells of textura angularis, wider around the ostiole, composed of dark brown, thick-walled cells of textura angularis. Asci 63-90 × 15-19 µm (x = 79 × 18.3 µm, n = 10), 16- to 32-spored, elongate-ellipsoidal to clavate, apedicellate, with a J- refractive apical ring, lying without paraphyses. Ascospores $18-27 \times 3-4 \ \mu m \ (\overline{x} = 23.8 \times 3.6 \ \mu m, n = 30), multi-seriate, (1)$ 2(-3)-septate, not constricted at the septum, cylindrical to narrowly ellipsoidal, straight or very slightly curved, tapering slightly to bro2adly rounded ends, hyaline, guttulate, smoothwalled, with 2-polar appendages. Asexual morph: Undetermined.

Specimen examined: **Italy**, Province of Forlì-Cesena, Bagno di Romagna, near Lago Pontini, on dead branch of *Alnus glutinosa* (*Betulaceae*), 26 May 2014, E. Camporesi, IT 1891 (**holotype**, MFLU 15–2661).



Fig. 24. Marsupiomyces quercina (MFLU 17–0876). A. Ascomata on substrate. B. Stromatic cavity. C, D. Cross section of ascoma. E. Peridium. F. Apical ring. G–K. Asci. L–N. Ascospores. Scale bars: A, B = 500 μm, C = 200 μm, D = 50 μm, E = 5 μm, F, L–N = 10 μm, G–K = 20 μm.

Notes: Here we introduce a new species *Ditopella biseptata* based on phylogeny. *Ditopella biseptata* forms a distinct clade which is sister to *Ditopella ditopa* (Fig. 1, Clade 1). Morphologically *Ditopella biseptata* has 2-septa and minute appendages at both ends. We could not obtain a culture from this fungus and extracted DNA directly from the sporocarps.

Harknessiaceae Crous, Persoonia 28: 55. 2012. Clade 7.

Saprobic or pathogenic forming leaf spots. Sexual morph: Ascomata perithecial, solitary or aggregated, immersed,

globose, coriaceous, brown, papillate. *Papilla* emergent to depressed, wall comprising 3–5 layers of brown-walled cells of *textura angularis*. *Hamathecium* comprising hyaline, septate paraphyses. *Asci* 8-spored, unitunicate, cylindrical to clavate, short pedicellate, with J- apical ring. *Ascospores* unito biseriate, hyaline, ellipsoid to fusoid, aseptate, thick-walled, guttulate, smooth-walled. Asexual morph: Coelomycetous. *Conidiomata* eustromatic, pycnidial, scattered or aggregated, immersed, globose, coriaceous, with single or several locules, dark brown to black. *Peridium* comprising thin-walled, almost hyaline to brown cells of *textura angularis*. *Ostiole* wide,





Fig. 25. Marsupiomyces epidermoidea (MFLU 15–2921) A. Herbarium specimen. B, C. Appearance of ascoma on the leaf surface. D. Vertical section through ascoma. E. Peridium in surface view. F. Peridium. G-K. Asci. L-O. Ascospores. Scale bars: B = 1 mm, C = 500 μm, D = 100 μm, E = 50 μm, F-K = 20 μm, L-O = 10 μm.

central, surrounded by brown cells. *Conidiophores* lining the inner cavity or reduced to the basal layer, sometimes reduced to conidiogenous cells, sometimes septate, branched. *Conidiogenous cells* holoblastic, discrete, lageniform, sub-cylindrical to cylindrical, hyaline to pale yellow, smooth, producing macroconidia and sometimes microconidia from same conidiogenous cell, proliferating sympodially one or

several times. *Macroconidia* with a basal appendage, hyaline when young, brown at maturity, unicellular, although basal appendage separated by a septum thick-walled, smooth-walled, with or without pale and dark longitudinal bands, sometimes longitudinally striate, guttulate, basal appendage cellular, cylindrical to subcylindrical, hyaline, thin-walled, devoid of contents, apical appendage present or absent, if



Fig. 26. Ditopella biseptata (MFLU 15–2661). **A.** Herbarium specimen. **B, C.** Appearance of ascomata on host substrate. **D.** Vertical section through ascoma. **E.** Papilla. **G–J.** Asci (J in Melzer's reagent). **K–M.** Ascospores (M in Indian ink). Scale bars: C = 1 mm, D = 200 μm, E = 100 μm, F = 50 μm, G–J = 20 μm, K–M = 10 μm.

present elongated. *Microconidia* hyaline, oval to ellipsoid, aseptate, smooth-walled.

Type genus: Harknessia Cooke.

Type species: Harknessia eucalypti Cooke.

Notes: Harknessiaceae (Fig. 1, Clade 7) was introduced to accommodate Harknessia with its wuestneia-like sexual morph. Harknessia species, distributed in both tropical and temperate biomes, are associated with leaves and branches of host trees (Farr & Rossman 2001). Most pathogenic Harknessia species are associated with leaf spots, leaf tip dieback, leaf scorch and stem cankers (Crous *et al.* 1989, 1993, Yuan *et al.* 2000), but pathogenicity has not been properly studied (Crous *et al.* 2012b). Some saprobic species have also been isolated from asymptomatic plant tissues (Marincowitz *et al.* 2008, Crous *et al.* 2017). Twenty-one of the 60 species and seven of the 13 wuestneia-like sexual morphs have been linked to Harknessia asexual morphs

(Crous et al. 2012b, 2017). Ribosomal DNA sequence analysis of diaporthoid taxa revealed a distinct lineage for *Harknessia* sensu stricto within *Diaporthales* (Crous et al. 2012b). Crous et al. (2012b) introduced six novel species of *Harknessia* from *Eucalyptus* and phylogenetic relationships based on a multi-gene analysis of ITS nrDNA, calmodulin and beta-tubulin genes were provided for these species. However morphologically, *Dwiroopa lythri* has similar characters to *Harknessia* and phylogenetically it is moderately supported here. Hence, *Dwiroopa lythri* is accommodated within *Harknessiaceae* for now. Phylogeny herein, indicates support for the establishment of this family.

Harknessia eucalypti Cooke, Grevillea 9 (no. 51): 85. 1881. Facesoffungi number FoF03488. Fig. 27.

Saprobic on *Eucalyptus globulus* appearing as nearly circular, black distinct spots. Sexual morph: Undetermined. Asexual morph: *Conidiomata* $390-550 \mu$ m high, $400-600 \mu$ m diam, erumpent, scattered, pycnidial, unilocular, globose to





Fig. 27. Harknessia eucalypti (K (M) 195744). A. Packet of herbarium. B. Herbarium specimen. C. Cross section of conidioma. D–E. Conidia attached to the conidiogenous cells. F–K. Conidia. Scale bars: C = 100 μm, D–E = 20 μm, F–K = 10 μm.

subglobose, brown. *Peridium* 3–4 layers of brown cells of *textura angularis*. *Conidiophores* short, cylindrical, almost globose, branched, 1–2 layers, hyaline, mixed with peridium cells. *Conidiogenous cells* 8–13 × 4–6 µm, ampulliform, cylindrical, hyaline to brown. *Conidia* 11.5–15 × 8–9.5 µm ($\bar{x} = 13 \times 8.5 \mu$ m, n = 20), globose to ovoid with a truncate apiculate apex and an obtuse to blunt base, smooth, hyaline when young, brown at maturity, with longitudinal striations along the length of some conidia. *Basal appendages* 5–15 × 1.5–3 µm ($\bar{x} = 10 \times 2.5 \mu$ m, n = 20), hyaline, tubular, smooth, thin-walled, often collapsing.

Material examined: USA, California, on leaves of Eucalyptus globulus, Harkness 1280, isotype K (M) 195744.

Notes: Yuan & Mohammed (1997) observed the asexual morph of *Wuestneia epispora* from culture which was morphologically identical to *Harknessia eucalypti*, although this has not been proven based on sequences. *Harknessia* is associated with leaf spots, leaf tip dieback or leaf scorch, stem cankers and is also common on leaf litter (Crous *et al.* 1989, Marincowitz *et al.* 2008).

Juglanconidaceae Voglmayr & Jaklitsch, Persoonia 38: 142. 2017. Facesoffungi number FoF03489. Clade 4.

Synonym: Melansporellaceae C.M. Tian et al. Phytotaxa 305: 194. 2017.

Saprobic on dead corticated twigs and branches of Juglandaceae species. Sexual morph: Pseudostromata inconspicuous, ectostromatic disc pale yellow to pale brown, causing a more or less postulate bark surface. Central column more or less conical, beneath the disc. Ascomata surrounding the ectostromatic disc, with long, asymmetrical or symmetrical, lateral ostioles that emerge at the margin or within the ectostromatic disc, globose to subglobose, coriaceous, black. Hamathecium comprising hyaline paraphyses which deliquesce at maturity. Asci 8-spored, unitunicate, with a distinct apical ring, sessile. Ascospores hyaline, bicellular, with or without gelatinous appendages. Asexual morph: melanconium-like. Conidiomata acervular, with ectostromatic disc and central column. Conidiophores aseptate or few-celled, smooth, hyaline to brown. Conidiogenous cells annellidic, cylindrical, base swollen, hyaline to brown. Conidia ellipsoid to oval, brown, with gelatinous sheath. Conidial wall smooth on the outer surface, with inconspicuous to distinct irregular verrucae on the inner surface (description based on Voglmayr et al. 2017).

Type genus: Juglanconis Voglmayr & Jaklitsch.

Type species: Juglanconis juglandina (Kunze) Voglmayr & Jaklitsch.

Notes: Juglanconidaceae (Fig. 1, Clade 4) was introduced by Voglmayr et al. (2017) based on Melanconium juglandinum. This family comprises Juglanconis juglandina, J. oblonga, J. pterocaryae, and J. appendiculata. Juglanconidaceae is morphologically and phylogenetically distinct from other families of Diaporthales. Species in this family are mostly pathogenic on Juglandaceae tree species causing black pustular dieback disease (Graves 1923, Belisario 1999). Du et al. (2017) introduced a new family Melansporellaceae for Juglanconis species and here we synonymise Melansporellaceae under Juglanconidaceae.

Juglanconis juglandina (Kunze) Voglmayr & Jaklitsch, Persoonia 38: 144. 2017. Facesoffungi number FoF03490.

Illustration: See Voglmayr et al. (2017).

Saprobic on dead twigs and branches of Juglandaceae. Sexual morph: Pseudostromata 0.8-2 mm diam, typically inconspicuous, sometimes distinct, circular, slightly projecting, without perithecial bumps. Ectostromatic disc 0.5-1.2 mm diam, indistinct, circular or oblong, dark grey, brown or black, often covered by densely arranged ostioles, pulvinate. Central column yellowish to brownish grey. Entostroma indistinct. Ascomata 440-565 µm diam, perithecial, aggregated, immersed, globose to subglobose, coriaceous, black, arrange in various configurations. Asci 140-160 × 17-22 µm, 8-spored, unitunicate, clavate to fusoid, indistinct apical ring, with small narrow stalk. Ascospores 25-30 × 8-11 µm, uni- to irregularly biseriate, hyaline, inequilaterally ellipsoid or broadly fusoid, asymmetric, distinctly constricted at the septum, without appendages, upper cell mostly larger, with rounded to subacute end, lower cell subacute to narrowly rounded, multiguttulate, containing mostly one large and numerous small guttules per cell. Asexual morph: Conidiomata acervular, 1-4 mm diam, black, scattered or occasionally confluent, with central or eccentric stromatic column, at maturity covered by black discharged conidial masses, usually conspicuous. Conidiophores 25-35 µm high, 5-6.5 µm wide, cylindrical to lageniform, simple, rarely branched at the base, smooth, subhvaline to pale brown, Conidiogenous cells annellidic with distinct annellations, integrated. Conidia 20-25 × 12-15 µm, unicellular, hyaline when immature, brown to blackish when mature, broadly ellipsoid to broadly pipshaped, truncate with distinct scar at the base, multiguttulate, thick-walled, wall ornamented on the inside of the wall with irregular confluent verrucae and with gelatinous sheath.

Notes: Voglmayr *et al.* (2017) neotypified *Melanconium juglandinum* based on a freshly collected specimen due to misplacement or loss of the type specimen and poor condition of other authentic specimens. The conidiomata, conidiophores and conidia was nicely illustrated by Corda (1839) and the asexual morph is very common and conspicuous, while the sexual morph is infrequently found in fully-developed condition.

Lamproconiaceae C. Norphanphoun, T.C. Wen & K.D. Hyde, Phytotaxa 270: 94. 2016. Facesoffungi number FoF03491. Clade 22.

Pathogenic and saprobic on dead herbaceous branches. Sexual morph: Stromata prosenchymatous around perithecia, delimited

externally by greenish, blackened, dense pseudoparenchymatous zone, interior whitish, composed of interwoven hyphae mixed with substrate cells, 3-5 ascomata in a stroma. Ascomata perithecial, small, aggregated, scattered, globose to subglobose, pale to dark brown, coriaceous, ostiolate, papillate. Papilla converging and erumpent through stroma surface as single, large opening. Peridium comprises pale brown, compressed, cells of textura angularis. Asci 8-spored, unitunicate, cylindrical, short-stalked, Japical apparatus. Ascospores uniseriate, broadly ellipsoid, 1septate, not or slightly constricted at the septa, hvaline, smooth, Asexual morph: Conidiomata pycnidial, solitary, partly immersed in host tissue, uni- to multilocular or convoluted, dark blue or dark blackish brown, erumpent in the centre. Pycnidium thick-walled, thin at inner layer, hyaline or dark brown, comprising wall cells of textura angularis or textura intricata. Ostiole absent, dehiscence irregular. Paraphyses interspersed within conidiophores. Conidiophores filiform or cylindrical, pale-bluish or hyaline, septate, branched, smooth-walled, formed at the base of conidiomatal wall. Conidiogenous cells holoblastic, cylindrical to subcylindrical, each forming a single conidium at the apex, or annellidic, colourless to olivaceous, smooth-walled. Conidia fusiform, ellipsoid, thick-walled, contents granular, aseptate, bluish to glistening dark blue or hyaline, smooth-walled, produced in mucilage but without a distinct mucilaginous envelope or appendage.

Type: Lamproconium (Grove) Grove.

Type species: Lamproconium desmazieri (Berk. & Broome) Grove.

Lamproconium desmazieri (Berk. & Broome) Grove [as 'desmazieri'], British Stem- and Leaf-Fungi (Coelomycetes) (Cambridge) 2: 321. 1937. Facesoffungi number FoF03492. Fig. 28.

Pathogenic and saprobic on dead twigs and branches of lime trees (Tilia sp.). Sexual morph: Undetermined. Asexual morph: Conidiomata 0.8-1 × 0.4-0.55 mm, pycnidial, solitary, partly immersed in host tissue, uniloculate, dark blue, with a raised centre. Pycnidium 50-70 µm, with multi-layered wall, thin at inner layer, hyaline, wall cells of textura angularis. Paraphyses interspersed with conidiophores. Conidiophores 30-120 µm high, arising from the outermost wall layer at the basal of pycnidium, filiform or cylindrical, pale-bluish to hyaline, septate, branched, smooth-walled. Conidiogenous cells cylindrical to sub-cylindrical, annellidic, with flared periclinal thickenings in the collarette zone, colourless to oliva-8–10 smooth-walled. Conidia 22–28 x ceous, μm $(\overline{x} = 25.25 \times 9 \mu m, n = 30)$, fusiform, ellipsoid, infrequently slightly curved, aseptate, initially hyaline, bluish to glistening dark blue at maturity, narrowly rounded at ends, smooth-walled.

Material examined: **Russia**, Rostov region, Krasnosulinsky district, Donskoye forestry, artificial forest, on dead branches of *Tilia cordata (Tiliaceae)*, 21 May 2014, T. Bulgakov, MFLU 14–0780.

Notes: Melanconium desmazieri was reported as the asexual morph of Melanconis desmazieri from Tilia species (Petrak 1938). Grove (1937) re-circumscribed the species of Melanconium and postulated that Melanconium desmazieri differed from the type species of Melanconium in having 1-septate, bluish to glistening dark blue conidia. Therefore, Grove introduced a new genus Lamproconium to accommodate this taxon (Grove 1937, Sutton 1980), and Lamproconium desmazieri was placed in



Fig. 28. Lamproconium desmazieresi (MFLU 14–0780). A–C. Conidiomata on host. D. Vertical section of conidioma. E. Peridium and raised host. F. Apex of conidioma. G, H. Conidiogenous cells with attached conidia. I–M. Immature conidia. N. Mature conidium. Scale bars: A = 2 mm, B = 1 mm, C = 500 μm, D = 300 μm, E, F = 200 μm, G, H = 40 μm, I–N = 20 μm.

Diaporthales genera incertae sedis by Cannon & Minter (2014). Based on phylogenetic study, Norphanphoun *et al.* (2016) synonymised *Melanconis desmazieri* under *Lamproconium desmazieri* and introduced a new family *Lamproconiaceae* to accommodate *Lamproconium* and *Hercospora*. Morphologically *Lamproconiaceae* is distinct from other families of *Diaporthales* in having dark blue or dark blackish brown pycnidial conidiomata and fusiform to ellipsoid, aseptate, bluish to glistening dark blue or hyaline conidia. The sexual morph is reported only for *Hercospora*. Combined gene analysis of LSU nrDNA, ITS nrDNA, *rpb2*, and *tef1* shows that *Lamproconiaceae* is a distinct family that is sister to *Sydowiellaceae* (Fig. 1, Clade 22). Macrohilaceae Crous, IMA Fungus 6: 180. 2015. Clade 15.

Pathogenic forming leaf spots. Sexual morph: Undetermined. Asexual morph: Coelomycetous. *Conidiomata* pycnidial, immersed, becoming erumpent, medium brown, globose. *Conidiophore* reduced to conidiogenous cells. *Conidiogenous cells* lining the inner cavity, pale brown, cylindrical, proliferating percurrently near the apex. *Conidia* solitary, medium to dark brown, ovoid, smooth, guttulate, medianly septate, apex obtuse, base truncate with a visible scar.

Type genus: Macrohilum H.J. Swart.

Type species: Macrohilum eucalypti H.J. Swart.

Notes: The family Macrohilaceae was introduced and typified by Macrohilum (Crous et al. 2015) and its taxonomic placement in Diaporthales has been reported based on LSU nrDNA sequence data. In this study, our concatenated analysis on LSU nrDNA, ITS nrDNA, *rpb2* and *tef1* also indicates that the Macrohilum eucalypti strains cluster together with high support and belong to the Macrohilaceae (Fig. 1, Clade 15). Macrohilaceae differs from other families of Diaporthales in having single, dark brown, guttulate, thick-walled, medianly septate, oval conidia with obtuse apex and truncate base (Crous et al. 2015). This monotypic family comprises only a single species commonly associated with leaf spots of *Eucalyptus*.

Macrohilum eucalypti H.J. Swart, Trans. Br. mycol. Soc. 90: 288. 1988. Facesoffungi number FoF03493.

Illustration: See Crous et al. (2015).

Pathogenic forming leaf spots. Sexual morph: Undetermined. Asexual morph: *Conidiomata* immersed, becoming erumpent, medium brown, globose, to 300 µm diam. *Conidiogenous cells* lining the inner cavity, pale brown, cylindrical, finely roughened, proliferating percurrently near the apex, $10-15 \times 3-5$ µm. *Conidia* solitary, medium to dark brown, ovoid, smooth, guttulate, developing a single, dark brown, supra-median septum, thick-walled, frequently constricted at the septum, apex obtuse, base truncate and protruding, with a visible scar, 2-3 µm wide, $15-20 \times 10-12$ µm (description based on Crous *et al.* 2015).

Notes: Crous *et al.* (2015) epitypified *Macrohilum eucalypti* using an Australian specimen collected from *Eucalyptus piperita*. Although a New Zealand isolate (CPC 10945) differed from the Australian ex-epitype isolate (CPC 19421) by four base pairs in the ITS nrDNA, Crous *et al.* (2015) did not propose this isolate as a new species pending collection of more material.

Melanconidaceae G. Winter [as '*Melanconideae*'], Rabenh. Krypt.-Fl., Edn 2 (Leipzig) 1.2: 764. 1886. Clade 2.

Saprobic or pathogenic on plants. Sexual morph: *Pseudostromata* well-developed, obvious, erumpent. *Ectostromatic disc* surrounded by bark or not, yellowish white, ostiolar canal opening around the disc. *Ascomata* perithecial, arranged as circles around the ectos-tromatic disc, oblique or horizontal, globose to subglobose, coriaceous, black, with long, periphysate, lateral ostiolar canals. *Peridium* comprising outer, thick-walled, brown cells of *textura globosa* to *textura angularis* and inner, thick-walled, flat, hyaline cells of *textura angularis*. *Hamathecium* comprising wide, hyphaelike, paraphyses, deliquescent at maturity. *Asci* 8-spored,

unitunicate, oblong to fusiform, short pedicellate, with distinct, Japical ring. *Ascospores* overlapping uni- to biseriate, hyaline, ellipsoid, 1-septate. Asexual morph: coelomycetous, melanconiumlike. *Conidiomata* acervular, scattered, solitary, superficial, black, coriaceous. *Conidiophores* branched at the base, septate, *Conidiogenous cells* annellidic, cylindrical. *Conidia* hyaline to brown, ellipsoid or subglobose, smooth-walled, thick-walled.

Type genus: Melanconis Tul. & C. Tul.

Type species: Melanconis stilbostoma (Fr.) Tul. & C. Tul.

Notes: The family Melanconidaceae was introduced by Winter (1886) to accommodate species having yellowish-white ectostromatic discs surrounding ascomata arranged in a circle. Members of this family are plant pathogens causing disease on as economic plant species. well as saprobes. Maharachchikumbura et al. (2016) listed 24 genera under this family based on morphology, following Lumbsch & Huhndorf (2010). However, most genera do not have any DNA sequence data, except Dicarpella, Melanconiella, Melanconis, Melanconium, and Prosthecium. Voglmayr & Jaklitsch (2014) synonymised Prosthecium under Stilbospora and included it in Stilbosporaceae. Crous et al. (2012b) have linked more than half of known wuestneia-like species to Harknessia species accommodating it in Harknessiaceae. Based on morphological and phylogenetic evidence, Castlebury et al. (2002) and Rossman et al. (2007) reported that this family is monogeneric with Melanconis and its asexual morph Melanconium. However, Rossman et al. (2015) synonymised Melanconium under Melanconis based on the poor phylogenetic resolution of Melanconium species and poor host-specificity. Phylogenies generated in this study position Dicarpella and Melanconiella (both in Melanconiellaceae; Fig. 1, Clade 10) outside the Melanconidaceae (Fig. 1, Clade 2). Considering the lack of molecular data, diverse ecological strategies and variable morphology, the family Melanconidaceae is restricted to Melanconis sensu stricto. Hence, we exclude all genera listed in Maharachchikumbura et al. (2016) from this family except Melanconis.

Melanconis apiocarpum and *M. marginale* have been reported from leaf spots of *Alnus* species in Canada, England, and Switzerland (Sieber *et al.* 1991). *Melanconium juglandinum* causes black pustular dieback of *Juglans* species in Europe and was consistently isolated from diseased twigs and branches of Persian walnut trees (*Juglans regia*), proving to be a virulent pathogen (Belisario 1999).

Melanconis italica Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821560. Facesoffungi number FoF03494. Figs 29, 30.

Etymology: Species epithet based on the country where the fungus was collected, Italy.

Saprobic on *Alnus cordata*. Sexual morph: *Pseudostromata* poorlydeveloped, erumpent. *Ectostromatic disc* 500–600 µm diam, surrounded by bark or not, yellowish-white, causing a coarse bark surface, inverted conical, ostioles open into margin and rarely middle of the disc. *Ascomata* 0.90–1 mm high, 0.4–0.5 mm diam ($\overline{x} = 0.98 \times 0.47$ mm, n = 20), perithecial, oblique, globose to subglobose, coriaceous, black, with long periphysate, lateral ostiolar canal. *Hamathecium* comprising wide, hyphae-like, hyaline, septate



Fig. 29. Melanconis italica (MFLU 17–0879). A. Stromata on substrate. B–D. Horizontal cross section of stroma. E. Vertical section of ascoma. F. Peridium. G. Paraphyses. H–K. Asci. L–O. Ascospores. Scale bars: A = 500 μm, B-D = 200 μm, E = 100 μm, F = 50 μm, G–O = 10 μm.

4–12 µm wide ($\overline{x} = 9 \mu m$, n = 20) paraphyses. *Peridium* 15–28 µm diam ($\overline{x} = 21 \mu m$, n = 10), comprising thick-walled, brown cells of *textura angularis*. *Asci* 80–92 × 11–14 µm ($\overline{x} = 13 \times 8.5 \mu m$, n = 20), 8-spored, unitunicate, cylindrical, short pedicellate, with distinct, J-apical ring. *Ascospores* 18–21 × 2.8–4 µm ($\overline{x} = 19 \times 3.5 \mu m$), biseriate, hyaline, fusiform, 1-septate, slightly constricted at the septum, smooth-walled. Asexual morph: *Conidiomata* on MEA solitary, superficial, globose, appears as slimy bubbles of conidia mass, black. *Conidiophores* cylindrical, branched, thick-walled,

hyaline. *Conidiogenous cells* blastic, terminal or intercalary, bottle-shaped, narrowing towards the apex, hyaline, thick-walled. *Conidia* fusiform to ellipsoidal, aseptate, thick-walled, basal end pointed, apical end blunt, olivaceous.

Culture characteristics: Colonies growing on MEA attained 1 cm within 7 d incubation at 18 °C, flat, circular, smooth margin, white, tightly attached to the substrate, little aerial mycelia.



Fig. 30. Asexual morph of *Melanconis italica* (MFLUCC 16–1199). A, B. Conidiomata on PDA. C–F. Conidia attached to conidiogenous cells, and conidiophores. G. Conidia. Scale bars: A = 500 μm, B = 200 μm, C–F = 10 μm, G = 5 μm.

Specimen examined: Italy, Province of Forlì-Cesena, Fiumicello di Premilcuore, on dead branch of *Alnus cordata* (*Betulaceae*), 4 Dec. 2013, E. Camporesi, IT 1557 (holotype MFLU 17–0879, isotype BBH 42441, cultures ex-type, MFLUCC 16–1199, MFLUCC 17–1659).

Notes: Melanconis italica clusters in a clade with *M. alni* with high support (Fig. 1, Clade 2). Both *M. italica* and *M. alni* are associated with *Alnus* species. *Melanconis alnicola* is also reported from *Alnus* species. However, there are no DNA sequence data in GenBank for *Melanconis alnicola*. Morphologically, *M. alni* differs from *M. italica* in having short to long, hyaline, filiform appendages at both ends, and oval to ellipsoid ascospores. In contrast, *M. alnicola* has large, oval to ellipsoid ascospores $(25-45 \times 9-12 \ \mu m)$ and small asci $(50-60 \times 10-15 \ \mu m)$. Hence, a new species, *Melanconis italica*, is introduced to accommodate this taxon.

Melanconiellaceae Senan. & Maharachch., K.D. Hyde, fam. nov. MycoBank MB821561. Facesoffungi number FoF03495. Clade 10.

Synonym: Melanconiellaceae Locq., Mycol. gén. struct. (Paris): 210. 1984. (nom. inval., Art 39.1, Melbourne Code).

Phytopathogenic or saprobic. Sexual morph: *Stromata* present or absent. If present; *Pseudostromata* inconspicuous, erumpent, pale or dark coloured ectostromatic disc or pulvillus causing a more or less pustulate bark surface. *Ectostromatic disc* convex, flat to concave, surrounded by bark or not. *Central column* beneath the disc more or less conical, comprising hyaline or pigmented hyphae mixed with a pigmented, cream, yellow, olive, brownish or grey, powdery amorphous substance. *Perithecia* sometimes epiphyllous without stromatic tissues and immersed in host substrate, inconspicuous or appearing as rounded bumps beneath the bark surrounding the ectostromatic disc, oblique or horizontal, scattered or often arranged in a circle around the

central column, with long lateral ostioles that converge at the margin of the central column. Ostioles emerging in various positions in the ectostromatic disc, cylindrical. Peridium comprising dark, thick-walled cells of textura angularis. Hamathecium aparaphysate or comprising broad, hyaline paraphyses. Asci 2-8-spored, unitunicate, cylindrical-clavate, oblong or fusoid, with a J- distinct apical ring, tapering below to a short, narrow pedicel. Ascospores hyaline, yellowish or brown, oblong, fusoid or ellipsoid, 0-1-septate, septa central or slightly eccentric, slightly constricted or not, smooth, with or without short, blunt appendages and sometimes with a narrow gelatinous sheath. Asexual morph: Conidiomata acervular or pycnidia, punctiform, subcuticular, immersed or erumpent, sometimes with a central, well-developed, pale brown, pseudoparenchymatous layer becoming thinner or absent at the margin of the conidiomata, multiloculate, sometimes papillate, sometimes with pale coloured, ectostromatic disc and central column or with radiate scutella. Scutella convex, membranous, brown, somewhat translucent, with a central hyaline or pale disc, giving rise to radiating hyphae, thick-walled cells radiating from a central point, rounded to pointed at the tips. Conidiophores reduced to conidiogenous cells or branched, sometimes septate only at the base, few-celled, smooth, hyaline to pale brown, sometimes short, forming under the developing scutellum. Pseudoparaphyses filiform. Conidiogenous cells annellidic or phialidic. Conidia initially hyaline becoming brown, ellipsoid, obovoid, subglobose, ovoid or oblong, thick-walled, smooth to finely verrucose, with or without distinct hyaline sheath, each with a truncate base and obtuse to bluntly pointed apex, sometimes somewhat granular, sometimes with inconspicuous to conspicuous basal hilum, with or without distinct hyaline sheath or frill.

Type genus: Melanconiella Sacc.

Type species: Melanconiella spodiaea (Tul. & C. Tul.) Sacc.



Notes: The phylogenetic analyses of this study showed that *Greeneria*, *Melanconiella*, and *Dicarpella* (previously placed in *Melanconidaceae*), *Tubakia* (previously placed as *Diaporthales incertae sedis*), *Sphaeronaemella fragariae* (previously placed in *Microascales incertae sedis*) and *Microascospora* gen. nov. forms a distinct clade with moderate support, which we recognise as Melanconiellaceae (Fig. 1, Clade 10).

The genus Greeneria was introduced based on G. fuliginea (Scribner & Viala 1887) and was synonymised under Melanconium (Cavara 1889) as Melanconium fuligineum. Later van der Aa (1973) accommodated this genus in Phyllosticta as P. ampelicida. Punithalingam (1974) renamed this taxon as Greeneria uvicola providing a detailed description and illustration. A LSU nrDNA sequence analysis by Farr & Rossman (2001) showed the phylogenetic placement of Greeneria uvicola outside of Melanconidaceae, but within Diaporthales. Analyses in this current study showed the phylogenetic placement of G. uvicola and G. saprophytica within Melanconiellaceae (Fig. 1, Clade 10). However, G. saprophytica does not show a very close affinity to G. uvicola. Greeneria uvicola is one of the most common pathogens causing various diseases in grapes (Navarrete et al. 2009). Greeneria lacks a known sexual morph (Zhang & Blackwell 2001) and it differs from other diaporthalean asexual morphs in having holoblastic conidiogenesis, producing phialidic conidiogenous cells in acervuli or pycnidia, and pale brown conidia (Barr 1978). The fungus overwinters on stem lesions, mummified berries, leaves, and tendrils. It is known to attack several species of Vitis including V. aestivalis, V. labrusca, V. rotundifolia, and V. vinifera.

The genus *Dicarpella* is based on *Dicarpella bina* and the asexual morph of this genus was reported as *Tubakia* (Belisario 1991). *Tubakia* is typified by *Tubakia japonica*. The type species of these two genera are not linked to each other. However molecular data linked *Tubakia* and *Diplacella* together and a few *Diplacella* species have *Tubakia* asexual morphs (Sogonov *et al.* 2008). *Tubakia* is more commonly encountered compared with *Dicarpella* and it is also a more widely used name than *Dicarpella*. The phylogenetic analyses in this study indicate a plausible relationship of *Dicarpella dryina* and *Tubakia* are congeneric without analysing sequence data of the type species. Thus, here we retain *Dicarpella* and *Tubakia* as two separate genera until sequence data becomes available.

VogImayr et al. (2012) reviewed the genus *Melanconiella* based on herbarium material and recently collected specimens. The morphological and phylogenetic distinctness of *Melanconiella* from *Melanconis* was discussed. The generic type of *Melanconiella* was confirmed as *M. spodiaea*. Phylogenetic analysis in this study showed the distinct placement of *Melanconiella* within this new clade (Fig. 1, Clade 10).

A new genus *Microascospora* is introduced to this family based on *Microascospora rubi*. Phylogenetically *Sphaeronaemella fragariae* did not cluster with other *Sphaeronaemella* species and it forms a clade with *Microascospora rubi*. Hence *Sphaeronaemella fragariae* is excluded from *Sphaeronaemella* and accommodated in *Microascospora* as *M. fragariae*. However, *Melanconiellaceae* was originally invalidly published (Art. 39.1, Melbourne) by Locquin (1984). Hence *Melanconiellaceae* is herewith validated to accommodate *Dicarpella*, *Greeneria*, *Melanconiella*, *Microascospora* and *Tubakia*.

Melanconiella chrysodiscosporina Voglmayr & Jaklitsch, Fungal Diversity 57: 14. 2012. Facesoffungi number FoF03496. Fig. 31.

Saprobic on dead branch of Fagus sylvatica. Sexual morph: Pseudostromata indistinct, irregular or circular outline. Ectostromatic disc minute, circular, narrowly fusoid to oblong, yellow or grevish brown, central column vellow. Entostroma comprising subhyaline to yellowish hyphae. Ascomata 1.2-1.3 mm high, 0.3-0.5 mm diam ($\bar{x} = 1.25 \times 0.45$ mm, n = 20), immersed, aggregated, globose to subglobose, coriaceous, black, arranged in valsoid configuration. Papilla 600-950 µm high, 90-130 µm diam $(\overline{x} = 800 \times 117 \ \mu m, n = 10)$, long, asymmetrical or symmetrical, black, converging at upper region and make a common canal to open out, internally covered by hvaline periphyses. Peridium 14–17 μ m diam (\overline{x} = 15 μ m, n = 10), comprising outer few layers of thick-walled, brown, compressed cells of textura angularis and inner thick-walled, hyaline, compressed cells of textura angularis. Asci 85–100 × 13–17 μ m (\overline{x} = 95 × 15 μ m, n = 20), 8-spored, unitunicate, cylindrical to fusoid, with J- distinct apical ring, sessile or with short pedicel. Ascospores $17-20 \times 6-9 \mu m$ ($\overline{x} = 17.6 \times 7.5 \mu m$, n = 20), uni- or biseriate, broadly ellipsoid, not constricted at the septum, ends broadly rounded, hyaline, medianly 1-septate, multiguttulate with one large and numerous small guttules per cell, wall swelling and sometimes thicken and stuffed at the septum. Asexual morph: discosporina-like. Conidiomata 140-180 µm high, 490–600 µm diam (\bar{x} = 150 × 507 µm, n = 20), visible as darker spots marginated by a distinct dark brown to blackish marginal zone, with a central stromatic column, at maturity covered by whitish discharged conidial masses. Conidiophores 6-10 × 5-7.5 µm $(\overline{x} = 7.4 \times 6 \mu m, n = 20)$, few layers, cubic, thick-walled, hyaline. Conidiogenous cells $9-12 \times 1-2 \mu m$ ($\overline{x} = 11 \times 1.8 \mu m$, n = 20), phialidic, conical, base enlarged, narrowing towards the apex, thickwalled, hyaline. Conidia 10-12 × 4-6 μ m (\overline{x} = 10.8 × 5.2 μ m, n = 20), ellipsoid, oblong or cylindrical, with two large and numerous small guttules, hyaline, with gelatinous sheath.

Specimen examined: **Italy**, Province of Forlì-Cesena, Bagno di Romagna, near Riofreddo, on dead branches of *Fagus sylvatica (Fagaceae*), 14 Aug. 2016, E. Camporesi, IT 3066, MFLU 17–0893, living culture MFLUCC 17–1671.

Notes: Melanconiella chrysodiscosporina was introduced by Voglmayr et al. (2012). This fungus was mostly found in the summer season from late spring to autumn (Voglmayr et al. 2012). The holotype and other authentic specimens were collected from dead branches of *Carpinus betulus* (*Betulaceae*). However, we collected this specimen from dead branches of *Fagus sylvatica* (*Fagaceae*). This is the first host record of *Melanconiella chrysodiscosporina* on *Fagus sylvatica*.

Melanconiella chrysomelanconium Voglmayr & Jaklitsch, Fungal Diversity 57: 16. 2012. Facesoffungi number FoF03497. Fig. 32.

Saprobic on branches of *Carpinus betulus*. Sexual morph: Not observed. Asexual morph: melanconium-like. *Conidiomata* 0.4–1 mm diam, visible as blackish spots with central or eccentric ostiolar opening, pycnidial, epidermal to



Fig. 31. *Melanconiella chrysodiscosporina* (MFLU 17–0893). A. Herbarium specimen. B. Vertical cross section of ascoma. C. Peridium. D–F. Asci. G–J. Ascospores. K. Conidiomata on substrate. L. Vertical cross section of conidioma. M. Conidiophores, conidiogenous cells and conidia. N–Q. Conidia. Scale bars: A = 1 mm, B, L = 100 μm, C = 20 μm, D–J, M = 10 μm, K = 200 μm, N–Q = 5 μm.

subepidermal, globose to subglobose, black, coriaceous, at maturity covered by black discharged conidial masses. Ostiole present, pointed. Conidiomatal wall composed of thin-walled, brown cells of textura angularis. Conidiophores reduced to conidiogenous cells, arising from the uppermost layer of cells of the basal stromatic pycnidial wall. Conidiogenous cells 7–18 µm high, 2–6 µm diam ($\overline{x} = 14.4 \times 4 \mu m$, n = 20), annellidic, hyaline, cylindrical, thick-walled, determinate, integrated, with flared collarette and periclinal wall-thickening. Conidia 13–20 × 7–11 µm ($\overline{x} = 15 \times 8.5 \mu m$, n = 20), dark brown, broadly ellipsoid to globose, circular in outline, slightly

truncate at base, as eptate, multiguttulate with 1–2 large and numerous small guttules, thick-walled, with distinct gelatinous sheath, smooth.

Specimen examined: Italy, Province of Forlì-Cesena, Via Nenni, Forlì, on dead aerial branches of *Carpinus betulus* (*Betulaceae*), 2 Jan. 2015, E. Camporesi, IT 1622, MFLU 17-0966.

Notes: Melanconiella chrysomelanconium is morphologically similar and phylogenetically related to *M. chrysodiscosporina* (VogImayr *et al.* 2012). The combined gene analysis of this study



Fig. 32. Melanconiella chrysomelanconium (MFLU 17–0966). A, B. Conidiomata on substrate. C. Cross section of conidioma. D–F. Conidiophores, conidiogenous cells and attached conidia. G–J. Conidia. Scale bars: B = 200 µm, C = 500 µm, D–F = 20 µm, G–J = 10 µm.

illustrates its phylogenetic relationship to other *Melanconiella* species (Fig. 1, Clade 10).

Microascospora Senan. & K.D. Hyde, gen. nov. MycoBank MB821562. Facesoffungi number FoF03498.

Etymology: Name based on small ascospores (<20 µm in length).

Saprobic on dead stems. Sexual morph: *Ascomata* scattered, solitary, immersed, globose to subglobose, brown, coriaceous, papillate, ostiolate. *Papilla* narrow, long, straight or curved,

comprising thick-walled, brown, compressed cells of *textura* angularis, internally covered by hyaline periphyses. *Peridium* comprising thick-walled, brown, somewhat compressed cells of *textura angularis*. *Hamathecium* aparaphysate. *Asci* 8-spored, unitunicate, clavate to fusiform, J- apical ring, attached to base without a pedicel. *Ascospores* overlapping biseriate, ellipsoid to fusiform, hyaline, aseptate, with two large fat globules at ends, appendages long, filiform to wavy, hyaline. Asexual morph: Undetermined.



Fig. 33. *Microascospora rubi* (MFLU 15–1112). A. Herbarium specimen. B. Vertical section of ascoma. C. Peridium. D–F. Asci. G–J. Ascospores. K. Upper side of culture growing on MEA. L. Lower side of culture growing on MEA. Scale bars: A = 200 µm, B = 100 µm, C = 25 µm, D–F = 10 µm, G–J = 5 µm.

Type species: Microascospora rubi Senan., Maharachch. & K.D. Hyde.

Microascospora rubi Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821563. Facesoffungi number FoF03499. Fig. 33.

Etymology: Name based on host genus Rubus.

Saprobic on dead stems of Rubus ulmifolia. Sexual morph: Ascomata 250-290 × 205-255 µm (x = 269 × 230 µm), scattered, solitary, immersed, globose to subglobose, brown, coriaceous, papillate, ostiolate. Papilla 115-155 µm high, 55-67 µm diam ($\overline{x} = 139 \times 65 \mu m$, n = 10), narrow, long, straight or curved, comprising thick-walled, brown, compressed cells of textura angularis, internally covered by hyaline periphyses, Peridium $10-25 \ \mu m$ ($\overline{x} = 18 \ \mu m$), comprising thick-walled, brown, large, somewhat compressed cells of textura angularis. Hamathecium aparaphysate. Asci 68-70 × 15-18 µm (x = 69 × 16 µm, n = 20), 8-spored, unitunicate, clavate to fusiform, J- apical ring, attached to base without a pedicel. Ascospores 14-19 × 5-7 µm $(\overline{x} = 17 \times 6 \mu m, n = 20)$, overlapping biseriate, ellipsoid to fusiform, hyaline, aseptate, with two large fat globules at ends, appendages long, filiform to wavy, hyaline. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on MEA slow growing, becoming 1 cm within 10 d at 18 °C, circular, umbonate, irregular

margin, cream to olivaceous, cotton-like, loosely attached to the substrate.

Specimen(s) examined: Italy, Province of Forlì-Cesena, Bagno di Romagna, Ridracoli, on dead branch of *Rubus ulmifolia* (*Rosaceae*), 24 Jan. 2014, E. Camporesi, IT 1675 (holotype MFLU 15-1112, isotype BBH 42445).

Notes: A new genus *Microascospora* is introduced based on *M. rubi*. This genus is morphologically and phylogenetically distinct from other genera in *Melanconiellaceae* having small ascospores (<20 μ m length) with wavy, filiform long appendages, and immersed, solitary ascomata with wavy papilla.

Microascospora fragariae (F. Stevens & Peterson) Senan., Maharachch. & K.D. Hyde, **comb. nov.** MycoBank MB821631. Facesoffungi number FoF03500.

Basionym: Sphaeronaemella fragariae F. Stevens & Peterson, Phytopathology 6: 258. 1916.

Notes: The multi-gene sequence analysis in this study shows that Sphaeronaemella fragariae (Fig. 1, Clade 10) does not have any affinities to the type species of Sphaeronaemella, S. helvellae (incertae sedis in Microascales). Sphaeronaemella fragariae forms a well-supported clade that is sister to Micro-ascospora rubi in Melanconiellaceae. Hence, we exclude Sphaeronaemella fragariae from Sphaeronaemella and accommodate this taxon in Microascospora and propose a new combination as Microascospora fragariae.



Fig. 34. Tubakia thailandensis (MFLU 13–0260). A. Herbarium specimen. B. Conidiomata on the host surface. C. Vertical section of pycnothyrium. D. Top view of radiate scutellum and conidiogenous cells with developing conidia. E–H. Conidiogenous cells with developing conidia stained with lactophenol cotton blue. I–K. Conidia. L. Conidia stained with lactophenol cotton blue. M. Germinating conidium. N–O. Colonies on PDA from top. P–Q. Colonies on PDA from reverse. Scale bar: C = 50 µm, D, F–M = 10 µm, E = 5 µm.

Tubakia thailandensis Senan., Tangthir. & K.D. Hyde, **sp. nov.** MycoBank MB821564. Facesoffungi number FoF03501. Fig. 34.

Etymology: Name based on the country from which this species was collected, Thailand.

Saprobic on dead leaves. *Conidiomata* 40–50 µm high, 50–75 µm diam, pycnothyria with radiate scutella, scattered to gregarious, superficial on the substratum. *Scutella* convex, brown to dark brown, thick-walled cells, radiating from a central point. *Conidiophores* short, forming under the developing scutella. *Conidiogenous cells* 5–10 × 2–4 µm, phialidic, with a minute collarette and wide periclinal thickening. *Conidia* 10–12.4 × 7.4–8.7 µm (\overline{x} = 11.3 × 8.1 µm, n = 20), globose to subglobose, smooth, hyaline, thick-walled.

Specimen examined: **Thailand**, Chiang Rai, Doi Mae Salong, on dead leaf, 2 May 2012, K. Wisitrassameewong, NTCL059 (**holotype** MFLU 13–0260, culture ex-type MFLUCC 12–0303).

Culture characteristics: Mycelium white when young, dark green, pale grey to black from above and reverse when aged, with medium mycelium, flat, rhizoid to irregular form, labate margin, and attaining a diam of 46 mm on PDA in 7 d at 27 °C.

Notes: Tubakia comprises seven species (Index Fungorum 2017, MycoBank 2017). Braun *et al.* (2014) presented a taxonomic key to the genus *Tubakia* and according to that key, this species is morphologically quite similar to "*Tubakia* sp." which has a small scutellum (40–80 µm diam.) and hyaline or subhyaline conidia (9–11 × 7–9 µm) collected from *Castanea henryi* in China. Therefore, we introduce this species as *Tubakia* species in having small (length < 15 µm), globose or subglobose, hyaline conidia, without microconidial development. Tubakia shares close phylogenetic affinities to *Greeneria saprophytica* (Fig. 1, Clade 10).

Prosopidicolaceae Senan. & K.D. Hyde, fam. nov. MycoBank MB821565. Facesoffungi number FoF03502. Clade 17.

Pathogenic on species of *Fabaceae*. *Conidiomata* pycnidial, rarely acervular, solitary or aggregated in a eustromatic stroma with one to several ostioles or astromatic, grey to black, erumpent to immersed. *Peridium* comprising grey-brown cells of *textura angularis*. *Conidiophores* reduced to conidiogenous cells or lining the whole inner layer of the wall, subcylindrical, branched, septate, straight to irregularly curved, base pale brown, becoming medium green-brown at apex. *Conidiogenous cells* mono- to polyphialidic, tightly aggregated, hyaline, smooth, ampulliform, subcylindrical to lageniform, prominent periclinal thickening, at times with percurrent proliferation. *Conidia* solitary, subhyaline to brown, smooth, guttulate, straight to variously curved, ellipsoid to fusoid-ellipsoid, apex obtuse, base truncate to bluntly round.

Type genus: Prosopidicola Crous & C.L. Lennox.

Type species: Prosopidicola mexicana Crous & C.L. Lennox.

Notes: Prosopidicolaceae is a monotypic family introduced here to accommodate *Prosopidicola* species. *Prosopidicolaceae* (Fig. 1, Clade 17) is phylogenetically not associated with any support to known families of *Diaporthales*, but is morphologically well-delineated. Species in this family are pathogens on

Fabaceae host plants. This family comprises *Prosopidicola albizziae* and *P. mexicana* (Lennox *et al.* 2004, Crous *et al.* 2016).

Prosopidicola mexicana Crous & C.L. Lennox, Stud. Mycol. 50: 191. 2004. Facesoffungi number FoF03503.

Illustration: See Lennox et al. (2004).

Pathogenic causing pod rot disease on Prosopidis glandulosae. Lesions 2-3 mm wide and up to 7 mm long, covering the pod, irregular, extending across the width of the pod, pale brown with a raised, dark brown margin. Conidiomata up to 250 µm diam, amphigenous, pycnidial, rarely acervular, scattered, immersed to erumpent, globose to subglobose, unilocular, black. Peridium up to 15 µm thick, consisting of 3-4 layers of brown cells of textura angularis. Conidiophores 5-50 µm high, 3-4 µm diam, lining the whole inner layer of the wall, subcylindrical, branched, 0-3septate, straight to irregularly curved, base pale brown, becoming medium greenbrown at apex. Conidiogenous cells $5-16 \times 3-4 \mu m$, phialidic when young, with prominent periclinal thickening and proliferating percurrently with age, subcylindrical to lageniform, green-brown, smooth when young, becoming medium to dark green-brown and warty with maturity, apex obtaining flared collarettes, rarely with two loci per conidiogenous cell. Conidia 10–15 × 4.5–5.5 µm, solitary, broadly ellipsoidal, medium brown, straight to slightly curved, rounded at the apex, tapering to a subtruncate base, with an inconspicuous dehiscence scar, smooth, thin-walled, aseptate (description based on Lennox et al. 2004).

Notes: Prosopidicola mexicana is the cause of a severe pod rot disease on *Prosopidis glandulosa*. It appears as black lesions surrounded by a dark brown margin. Lennox *et al.* (2004) revealed it to group closely to *Cryphonectriaceae*. However in the phylogenetic analyses generated in this study, it forms a distinct clade which is basal to *Cytosporaceae*.

Pseudoplagiostomataceae Cheew. *et al.*, as "*Pseudoplagios-tomaceae*", Fungal Diversity 44: 95. 2010. Clade 12.

Pathogenic on leaves, forming spots. Sexual morph: Ascomata solitary, scattered, immersed, slanted to horizontal on host tissue, globose or elliptical, black, coriaceous, papillate, ostiolate. Papilla short, internally covered with hyaline, filamentous periphyses. Peridium comprising a few layers of thick-walled, brown cells of textura angularis. Hamathecium lacking paraphyses. Asci 8spored, unitunicate, cylindrical, sessile, with J-, subapical ring. Ascospores overlapping uni- to biseriate, hyaline, fusiform to ellipsoid, 1-septate, with terminal, elongate, hyaline appendages. Asexual morph: Coelomycetous. Conidiomata acervular or pycnidial, brown. Peridium comprising small, brown cells of textura angularis. Conidiophores absent. Conidiogenous cells cylindrical to ampulliform, enteroblastic, percurrently proliferating with periclinal thickening and collarette. Conidia holoblastic, hyaline to brown, ellipsoid, unicellular, subglobose to broadly allantoid, with obtuse apex and a flat protruding scar at the base.

Type genus: Pseudoplagiostoma Cheew. M.J. Wingf. & Crous.

Type species: Pseudoplagiostoma eucalypti Cheew., M.J. Wingf. & Crous.

Notes: Pseudoplagiostomaceae was introduced by Cheewangkoon *et al.* (2010). *Pseudoplagiostomaceae* is similar to *Gnomoniaceae* (Fig. 1, Clade 1) based on morphological characters of its sexual morph, such as solitary, immersed, non stromatic ascomata with lateral beaks, asci with a distinct apical ring and 1-septate ascospores (Sogonov *et al.* 2008). However, in our phylogenetic analyses it formed a fully-supported clade (Fig. 1, Clade 12) sister to *Apoharknessiaceae*.

Pseudoplagiostoma eucalypti Cheew., M.J. Wingf. & Crous, Fungal Diversity 44: 98. 2010. Facesoffungi number FoF03504.

Illustration: See Cheewangkoon et al. (2010).

Pathogenic on leaves forming leaf spots. Sexual morph: Ascomata 130-150 µm high, 100-130 µm diam, perithecia, immersed in host tissue, slanted to horizontal, globose to elliptical, coriaceous, brown to black, papillate, ostiolate. Papilla 60-65 µm diam, erumpent, internal wall lined by hyaline periphyses. Peridium comprising few layers of thick-walled, brown cells of textura angularis. Hamathecium aparaphysate. Asci 65-70 × 11-13 µm, 8-spored, unitunicate, subcylindrical to long obovoid, with wedgeshaped, J-subapical ring, apex blunt and without a distinct pedicel. Ascospores 17-19 × 5-7 µm, overlapping uni- to biseriate, ellipsoid, tapering towards rounded ends, hyaline, median 1septate, widest at septum, with terminal, elongate, hyaline appendages. Asexual morph: Conidiomata 180-200 µm high, 170–190 µm diam, acervular to pycnidial, subcutical to epidermal. Peridium comprising small, brown cells of textura angularis. Conidiophores absent. Conidiogenous cells 8-12 × 2-4 µm, cylindrical to ampulliform, enteroblastic proliferation with periclinal thickening. Conidia 17-19 × 7-8 µm, holoblastic, ellipsoid, unicellular, with obtuse apex and a flat protruding scar at the base (description based on Cheewangkoon et al. 2010).

Notes: The monotypic family Pseudoplagiostomataceae was introduced by Cheewangkoon et al. (2010) to accommodate a cryptosporiopsis-like fungus isolated from Eucalyptus. The type species, P. eucalypti (as Cryptosporiopsis eucalypti) and two other new species, P. oldii and P. variabile, were isolated as foliar pathogens from Eucalyptus leaf spots. However, Cryptosporiopsis eucalypti was not closely related to the generic type of Cryptosporiopsis, C. nigra (Dermateaceae, Helotiales), and hence Cheewangkoon et al. (2010) introduced Pseudoplagiostoma to accommodate this taxon. Pseudoplagiostoma corymbiae (Crous et al. 2012c) and P. dipterocarpi (Suwannarach et al. 2016) were introduced to this genus from Corymbia sp. and Dipterocarpus tuberculatus respectively. The ascospore morphology, in particular, is distinct and morphologically this family differs from other families in the order in having astromatic, slanted to horizontal, globose ascomata with aparaphysate hamathecium, ascospores with terminal, elongate, hyaline appendages and a cryptosporiopsis-like asexual morph. Cheewangkoon et al. (2010) analysed LSU nrDNA sequence data of the order Diaporthales, to show the distinct placement of Pseudoplagiostomaceae with 100 % bootstrap support.

Schizoparmaceae Rossman, Mycoscience 48: 137. 2007. Clade 8.

Saprobic, parasitic or pathogenic on woody, herbaceous plants. Sexual morph: Ascomata perithecial, solitary, scattered, subepidermal, immersed to erumpent, becoming superficial, globose, coriaceous, brown to black, short papillate, ostiole with hyaline periphyses, plate-like ornamentation around ostiole. Peridium comprising thick-walled, brown-cells of textura angularis. Hamathecium aparaphysate. Asci 8-spored, unitunicate, ellipsoid to fusiform, sessile, with a J- apical ring. Ascospores biseriate, hyaline to becoming pale brown at maturity, ellipsoidal, aseptate, with or without mucoid caps. Asexual morph: Coelomvcetous. Conidiomata pycnidial, subepidermal, immersed to erumpent, unilocular, globose, slightly depressed globose to subglobose. Conidiophores densely aggregated, slender, subulate, simple or branched, hyaline, smooth, occasionally septate and branched at base, invested in mucus, developing from basal pad. Conidiogenous cells discrete, simple, subcylindrical, obclavate or lageniform, smooth, proliferating percurrently or with prominent periclinal thickening. Conidia ellipsoid, globose, napiform, fusiform or naviculate with a truncate base and an obtuse to apiculate apex, hyaline or olivaceous brown to brown, unicellular, broadly or narrowly ellipsoidal, apices tapering, with or without a longitudinal germ slit, with or without a mucoid appendage.

Type genus: Coniella Höhn.

Type species: Coniella pulchella Höhn.

Notes: The monogeneric family *Schizoparmaceae* (Fig. 1, Clade 8) was introduced to accommodate *Coniella* (= *Pilidiella*, *Schizoparme*). Species of the asexual *Pilidiella* have been more widely reported than *Schizoparme* (Farr & Rossman 2017) and thus, *Schizoparme* was synonymised under *Pilidiella* giving priority to the older name (Rossman *et al.* 2015). Although van Niekerk *et al.* (2004) treated *Coniella* and *Pilidiella* as two distinct genera, the generic boundaries of the former were recently expanded to include "hyaline to dark brown conidia", as Alvarez *et al.* (2016) reported that conidial pigmentation was lost or gained several times during the evolution of species within *Coniella*.

Coniella pseudokoreana Senan., Tangthir. & K.D. Hyde, **sp. nov.** MycoBank MB821542. Facesoffungi number FoF03505. Fig. 35.

Etymology: Somewhat similar to *Coniella koreana*, however phylogenetically distant from this species.

Saprobic on dead leaves. *Conidiomata* pycnidial, solitary to gregarious, globose, brown, unilocular, ostiolate, 85–130 µm high, 78–106 µm diam ($\bar{x} = 108 \times 92 \mu$ m, n = 10), immersed, with a central short ostiolar canal on each conidioma. *Conidiomata wall* 2–4-layered, 6–15 µm wide ($\bar{x} = 10 \mu$ m), with outer brown to dark brown layers composed of thick-walled cells of *textura angularis*, with inner pale brown layer composed of thin-walled cells of *textura angularis*, with inner pale brown layer composed of thin-walled cells of *textura prismatica*, except at the base which has a pulvinate convex giving rise to conidiophores or conidiogenous cells. *Conidiophores* 4–8 µm high, 1.5–4 µm wide, short, branched at the base, hyaline, smooth. *Conidiogenous cells* 5–10 µm high, 1.5–2 µm wide, holoblastic to enteroblastic, phialidic. *Conidia* 18–26 × 3–4 ($\bar{x} = 23 \times 3.6 \mu$ m) fusiform, navicular, with one side slightly curved and another straight, smooth, hyaline, conidium length/width ratio = 6.5:1.

Culture characteristics: Colonies attaining a diam of 4 cm on PDA after 5 d at 27 °C; surface white with medium to sparse mycelium, flat, irregular, undulate or wavy margin.



Fig. 35. Coniella pseudokoreana (MFLU 13–0282). A. Specimen on dead leaf. B. Conidiomata on host surface. C–D. Longitudinal section of a conidioma. E–H. Conidiogenous cells with developing conidia. I. Conidiogenous cells with developing conidia stained in lactophenol cotton blue. J. Conidia. K. Conidia stained with lactophenol cotton blue. L. Germinating conidium. M. Colonies on PDA from top. N. Colonies on PDA from reverse. Scale bar: C = 100 µm, D = 50 µm, E–L = 10 µm.

Specimen examined: Thailand, Phitsanullok, Tung Salang Luang, on decaying leaf, 18 Jun. 2012, N. Tangthirasunun, NTCL093 (holotype MFLU 13–0282, culture ex-type MFLUCC 12–0427).

Notes: Coniella pseudokoreana displays somewhat similar morphological characters to *C. koreana* and *C. castaneicola* in having linear, falcate, pale brown conidia (Alvarez *et al.* 2016). The colony morphology of *Coniella koreana* described in Alvarez *et al.* (2016) on PDA is similar to the colony morphology of our strain. However, conidiomatal morphology and size of the conidia are different. We collected *Coniella pseudokoreana* on a decaying leaf in Thailand. Our phylogeny reveals, *C. pseudokoreana* is distant from *Coniella koreana* and shares a sister taxon relationship to *C. straminea* (Fig. 1, Clade 8).

Stilbosporaceae Link [as '*Stilbosporei*'], Abh. Königl. Akad.-Wiss. Berlin 1824: 180. 1826, emend. Clade 19.

Saprobic on bark of trees and shrubs. Sexual morph: Pseudostromata inconspicuous, immersed. Ectostromatic disc absent or present, if present inconspicuous, pale brown, rarely dark brown. Entostroma prosenchymatous, pale coloured, slightly differentiated from the surrounding bark tissue. Ascomata loosely arranged as valsoid groups in a single layer, immersed, aggregated, globose to subglobose, coriaceous, black, ostiolate, papillate. Ostiole not obvious, convergent in groups. Hamathecium comprising filiform, aseptate, hyaline paraphyses. Asci 8spored, unitunicate, cylindrical, initially attached to the base, later floating in centrum, with J- refractive, apical ring. Ascospores overlapping uni- to biseriate, brown, ellipsoid to oblong, distoseptate. Asexual morph: Coelomycetous. Conidiomata stromatic, acervular with circular outline, epidermal, immersed to semi-immersed, brown, basal stroma of textura angularis to textura globulosa, with simple, septate, hyaline paraphyses and hyaline, unbranched cylindrical conidiophores. Conidiophores arising from the uppermost cells of basal and parietal tissue, unbranched, cylindrical, septate at only the base, hyaline, smooth, invested in mucus. Conidiogenous cells annellidic, discrete or integrated, cylindrical to lageniform, hyaline, smoothwalled, proliferating several times percurrently at apex. Conidia ellipsoid or oblong, with an obtuse apex and broad truncate base, sometimes 3-euseptate or distoseptate, with a hyaline sheath, hyaline to brown, thick-walled, smooth, sometimes with several, tubular, unbranched, filiform, flexuous, apical appendages.

Type genus: Stilbospora Pers.

Type species: Stilbospora macrosperma Pers.

Notes: The family Stilbosporaceae was introduced by Link (1826) to accommodate Prosthecium and its asexual morph. However, it is not a phylogenetically well-supported family and hence, Stilbosporaceae has been synonymised under several different families. Voglmayr & Jaklitsch (2014) resurrected the family Stilbosporaceae in Diaporthales based on a phylogenetic analysis of LSU nrDNA sequence data and accommodated the genera Stegonsporium and Stilbospora within the family, synonymising Prosthecium under Stilbospora. This decision is also supported by our multi-gene phylogeny (Fig. 1, Clade 19). The type species of Stilbospora, S. macrosperma has been linked to its asexual morph Prosthecium ellipsosporum, the generic type of Prosthecium

(VogImayr & Jaklitsch 2008). This genus comprises opportunistic or moderately phytopathogenic fungal species that cause branch dieback or twig blight of various plants. Maharachchikumbura *et al.* (2015) included Natarajania in *Stilbosporaceae* based on LSU nrDNA, SSU nrDNA, *tef1* and *rpb2* sequence data. However, in other analyses (not shown here), phylogenies also indicated a close association to the genera *Crinitospora*, *Stilbospora* and *Stegonsporium*. This is rather interesting as up to date, this is the only hyphomycetous taxon affiliated to the diaporthean taxa which are known to have coelomyceteous asexual morphs. The reliability of the deposited sequences as well as the identification of that taxon needs further investigation.

Stilbospora macrosperma Pers., Syn. meth. fung. (Göttingen) 1: 96 (1801). Facesoffungi number FoF03506. Fig. 36.

Saprobic on branches of Acer pseudoplatanus. Sexual morph: Pseudostroma comprising white, grevish to yellowish hyphae. µm hiqh. Ascomata 300-350 350-465 um diam $(\overline{x} = 325 \times 420 \ \mu m, n = 20)$, immersed, aggregated, globose to subglobose, coriaceous, ostiolate, papillate. Papilla cylindrical, pale brown, emerging from perithecial apices and merging separately with the stromatal disc, inconspicuous, often invisible on the bark surface. Peridium 20-40 μ m diam (\overline{x} = 32 μ m, n = 20), comprising thick-walled, brown, large, cells of textura angularis and hyaline, thick-walled, compressed cells of textura angularis around the base of papilla. Hamathecium comprising multiguttulate, hyaline, septate paraphyses. Asci 165-200 × 35-50 µm $(\overline{x} = 182 \times 42 \mu m, n = 20)$, 8-spored, unitunicate, clavate to ellipsoidal, thick-walled, very short pedicellate, apex containing a Jrefractive canal usually wider towards its base. Ascospores $40-50 \times 20-26 \ \mu m$ ($\overline{x} = 46 \times 22 \ \mu m$, n = 20), biseriate, ellipsoidal, oblong or rarely pyriform, with (3-)5-distosepta and sometimes 1, longitudinal, distoseptum, appendages on both ends projecting, subglobose, outer margin becoming diffuse. Asexual morph: Conidiomata 340-450 µm high, 450-460 µm diam $(\overline{x} = 410 \times 453 \ \mu m, n = 20)$, immersed, acervular, solitary, with circular outline, dark brown to black. Paraphyses 2.5-4 µm diam $(\overline{x} = 3.2 \ \mu m, n = 10)$, unbranched, aseptate, hyaline. Conidiophores reduced to conidiogenous cells. Conidiogenous cells $25-35 \times 7-10 \ \mu m$ (x = 31 × 9 μm , n = 20), holoblastic, cylindrical, septate, hyaline. Conidia 40-45 × 20-25 μ m (\overline{x} = 43 × 23 μ m, n = 20), pyriform, oval, ellipsoid or oblong, base truncate and hyaline, brown, with several distosepta, 1(-2)-longitudinal distosepta, with hyaline sheath.

Material examined: **Austria**, Wien, Landstraße, 3rd District, Botanical Garden of the University of Vienna (HBV), grid square 7864/1, on dead corticated branches of *Acer pseudoplatanus* (*Sapindaceae*), holomorph, 4 Feb. 2006, H. Voglmayr, D39 (**epitype** WU 28068).

Notes: Voglmayr & Jaklitsch (2014) epitypified the type species of *Stilbospora, Stilbospora macrosperma* and *S. macrosperma* was confirmed as the asexual morph of *Prosthecium ellipsosporum*, the generic type of *Prosthecium* (Voglmayr & Jaklitsch 2008). *Stilbospora* (1801) is older than *Prosthecium* (1852) and therefore *Stilbospora* has priority (Voglmayr & Jaklitsch 2014).

Sydowiellaceae Lar.N. Vassiljeva, Pirenomits. Lokuloaskomits. Severa Dal'nego Vostoka (Leningrad): 210. 1987. Clade 21.



Fig. 36. Stilbospora macrosperma (WU 28068). A. Herbarium specimen. B. Stromata on host substrate. C. Cross section of ascoma. D. Peridium. E–H. Asci. I–M. Ascospores. N. Paraphyses. O. Cross section of conidioma. P. Conidia attached to conidiogenous cells. Q–S. Conidia. Scale bars: C–D = 500 μ m, E, O, P = 100 μ m, F = 50 μ m, H–K = 20 μ m, G, L–M, Q–S = 10 μ m.

Saprobic or parasitic on plant matter. Sexual morph: Stromata well- or poorly developed, prosenchymatous, scattered, immersed to erumpent, appearing as an aggregation of ostioles, rounded or elliptic in shape, dark brown to black, composed of compact pseudoparenchymatous tissues, several ascoma in a stromata, some species turn umber in 5 % KOH. Ascomata solitary or aggregated, immersed or erumpent. globose to sub-globose, sometimes circinate, coriaceous, central or asymmetrically located ostiolar canal opens through an individual or converged ostiole, internally covered by filamentous, hyaline periphyses, sometime ostiolar opening wider than canal, black to brown. Peridium comprising a few layers of brown, thick-walled cells of textura angularis. Hamathecium comprising cellular, septate, branched, hyaline paraphyses. Asci 8-spored, unitunicate, cylindrical to sub-globose, short pedicellate, apex blunt with J- apical ring. Ascospores uni- to multi-seriate, filamentous, ellipsoid or long fusoid-cylindrical, 1-11-septate, hyaline, pale brown to dark brown, sometimes with apical and basal appendages, wall smooth. Asexual morph: Conidiomata sometimes stromatic, pycnidia, uniloculate, superficial, aggregated 3-5 in one group, globose, orange to brown. Conidiomatal wall comprising thick-walled, orange cells of textura angularis. Conidiophores elongate, branched, hyaline, few conidiogenous cells arising from one conidiophore, attached to conidiomatal wall. Conidiogenous cells cylindrical, hyaline, ampulliform, septate, ends pointed, phialidic. Conidia ovoid to ellipsoid, unicellular, hyaline, smooth-walled.

Type genus: Sydowiella Petr.

Type species: Sydowiella fenestrans (Duby) Petr.

Notes: The family Sydowiellaceae (Fig. 1, Clade 21) was established to accommodate the genus Sydowiella, which is typified by S. fenestrans. Members of this family occur on herbaceous plants, dicotyledonous and hardwood trees as saprobes, parasites and pathogens. Initially, most genera in this family were placed in Diaporthales incertae sedis (Rossman et al. 2006). However, DNA sequence data analyses of different gene regions of taxa in the family Sydowiellaceae proved it to be a well-supported, and its relationships to other families have been clarified (Rossman et al. 2007, Maharachchikumbura et al. 2015, Senanayake et al. 2017). Sydowiellaceae comprises the genera Alborbis, Breviappendix, Cainiella, Calosporella, Chapeckia, Italiomyces, Hapalocystis, Lambro, Paragnomonia, Ranulospora, Rossmania, Sillia, Sydowiella, Tenuiappendicula and Tortilispora (Senanayake et al. 2017). Here we introduce a new Sydowiella species as S. urticicola. Sydowiella urticicola produce solitary ascomata and ascospores containing a large guttule in each cell which clearly demarcates it from other species. Phylogenetically it is also distinct from other Sydowiella species.

Sydowiella urticicola Senan., Camporesi & K.D. Hyde, **sp. nov.** MycoBank MB821566. Facesoffungi number FoF03507. Fig. 37.

Etymology: Named after the host genus Urtica.

Saprobic on dead branches of deciduous plants. Sexual morph: Ascomata 290–325 µm high, 290–395 µm diam ($\bar{x} = 309 \times 314$ µm, n = 20), perithecia, astromatic, scattered, solitary, superficial to erumpent, globose to sub-globose, coriaceous, black, papillate, ostiolate. *Papilla* 115–150 µm, 155–205 μm diam (\overline{x} = 133 × 173 μm, n = 20), short, wide, internally covered by hyaline periphyses. *Peridium* 15–25 μm diam (\overline{x} = 19 μm, n = 20) comprising inner, hyaline, thick-walled, compressed, 1–3 layers of cells of *textura angularis* and outer, dark brown, thick-walled, rigid, 3–7 layers of cells of *textura angularis*. *Hamathecium* comprising wide, cellular, septate paraphyses. *Asci* 125–145 × 10–15 μm (\overline{x} = 136 × 14 μm, n = 20) 8-spored, unitunicate, cylindrical, short pedicellate, distinct, Japical ring. *Ascospores* 20–25 × 10–15 μm (\overline{x} = 22 × 12 μm, n = 20), overlapping uniseriate, ends blunted, hyaline, 1-septate, slightly constricted at the septum, with a large fat globule in each cell. Asexual morph: Undetermined.

Culture characteristics: Colonies growing on MEA becoming 2 cm within 5 d at 18 °C, fast growing, circular, umbonate, margin irregular, white, tightly attached to the substrate without spreading aerial mycelium.

Specimen examined: Italy, Province of Forlì-Cesena, Monte Fumaiolo, dead stem of *Urtica dioica (Urticaceae)*, 16 May 2013, N. Camporesi, IT 1268 (holotype MFLU 17–0877, isotype BBH 42439, cultures ex-type MFLUCC 13–0665, MFLUCC 17–1665).

Notes: Here we introduce a new *Sydowiella* species as *S. urticicola. Sydowiella urticicola* produces solitary ascomata and ascospores containing a large guttule in each cell. Phylogenetically this is distinct from other *Sydowiella* species (Fig. 1, Clade 21).

Diaporthales genera incertae sedis

Phaeoappendicospora Senan., Q.R. Li & K.D. Hyde, **gen. nov.** MycoBank MB821572. Facesoffungi number FoF03508. Clade 18.

Etymology: Name based on three Latin words "*phaeo*", "*appendicem*" and "*spora*" referring to the brown spores with appendages.

Saprobic on dead stems of *Fagaceae* species. Sexual morph: *Stromata* forming thin weft of pale brown hyphae around upper part of perithecia. *Ascomata* immersed, aggregated, subglobose to globose, coriaceous, black, ostiolate, papillate. *Papilla* black, cylindrical, lateral, periphysate. *Periphyses* hyaline, filamentous. *Peridium* comprising few layers of black, thick-walled, cells of *textura angularis*. *Hamathecium* comprising few, septate, hyaline paraphyses attached to the base, longer than asci. *Asci* 8-spored, unitunicate, ellipsoid, with a short pedicel, inconspicuous, flat, J-refractive ring at the lower end of the thickened apical wall, apex narrow and blunted. *Ascospores* ellipsoidal to ovoid, with broadly rounded ends, pale brown, 1-septate, not constricted at the septa, thick and smooth-walled, guttulate, with short, hyaline, rounded appendages at both ends. Asexual morph: Undetermined.

Type species: Phaeoappendicospora thailandensis Q.R. Li, Senan. & K.D. Hyde.

Phaeoappendicospora thailandensis Senan., Q.R. Li & K.D. Hyde, **sp. nov.** MycoBank MB821573. Facesoffungi number FoF03509. Fig. 38.

Etymology: The species epithet is based on the collection locality.



Fig. 37. Sydowiella urticicola (MFLU 17–0877). A–C. Ascomata on host surface. D. Cross section of ascoma. E. Peridium. F. Papilla. G–I. Asci. J–N. Ascospores. Scale bars: A = 500 µm, B, C = 200 µm, D, E = 100 µm, F–I = 50 µm, J–N = 20 µm. www.studiesinmycology.org



Fig. 38. Phaeoappendicospora thailandensis (MFLU 12–2131). A. Appearance of fungus on host substrate. B. Horizontal cross section of stroma. C. Vertical cross section of stromata. D. Peridium. E. Apical ring in Melzer's reagent. F–I. Asci. J–M. Ascospores. Scale bars: A = 1 mm, B = 500 µm, C = 100 µm, D = 25 µm, F–M = 10 µm.

Saprobic on dead stems of *Quercus*. Sexual morph: *Stromata* 2–4 mm diam ($\overline{x} = 3$ mm, n = 15), forming thin weft of pale brown hyphae around upper part of perithecia. *Ascomata* 180–230 µm high, 170–220 µm diam ($\overline{x} = 208 \times 198$ µm, n = 25), immersed, aggregated, subglobose to globose, coriaceous, black, ostiolate, papillate. *Papilla* black, cylindrical, lateral, periphysate. *Periph*yses hyaline, filamentous. *Peridium* 20–35 µm wide ($\overline{x} = 28$ µm, n = 20), comprising few layers of black, thick-walled cells of *tex-tura angularis*. *Hamathecium* comprising few, septate, hyaline paraphyses attached to the base, longer than asci. *Asci* 195–265 × 18.5–27 µm ($\bar{x} = 223 \times 22.5$ µm, n = 30), 8-spored, unitunicate, ellipsoid, with a short pedicel, inconspicuous, flat, J-refractive ring at the lower end of the thickened apical wall, apex narrow and blunted. *Ascospores* 26–34.5 × 11–13 µm
$(\overline{x} = 30 \times 12.5 \mu m, n = 30)$, ellipsoidal to ovoid, with broadly rounded ends, pale brown, 1-septate, not constricted at the septa, thick and smooth-walled, guttulate, with short, hyaline, rounded appendages at both ends. Asexual morph: Undetermined.

Culture characteristics: Colonies on PDA reaching 5 cm in 7 d at 25 °C, fast growing, circular, flat, smooth, whitish cream in upper, pale yellow in reverse.

Specimen examined: Thailand, Chiang Rai, near Khun korn water fall, on branch of *Quercus* sp. (*Fagaceae*), 25 Dec. 2012, Q.R. Li, TL19 (holotype MFLU 12–2131, culture ex-type MFLUCC 13–0161).

Notes: The newly introduced monotypic genus *Phaeoappendicospora* is typified by *P. thailandensis* and it comprises brown ascospores with guttules and appendages. *Phaeoappendicospora thailandensis* is associated with dead plant parts may be as saprobes. This fungus is morphologically similar to *Hapalocystis berkeleyi* in *Sydowiellaceae*. However, phylogenetically it does not show an affinity to any of the families in *Diaporthales* (Fig. 1, Clade 18). Hence, we accommodate this species in *Diaporthales* genera *incertae sedis*.

Key to families and genera of Diaporthales

| 1. | | Coelomycetous2 |
|----|-----|--|
| 1. | | Ascomycetous |
| 2. | | Conidia hyaline to olivaceous or bluish to glistening dark blue3 |
| 2. | | Conidia brown to dark brown6 |
| 3. | | Conidial wall thick, mostly specific on <i>Tiliaceae</i> host |
| 3. | | Conidial wall thin, mostly on Myrtaceae, Fabaceae host4 |
| 4. | | Conidia aseptata5 |
| 4. | | Conidia 1-septateAuratiopycnidiellaceae |
| 5. | | Conidiophores reduced to conidiogenous cells, conidia with acute apexErythrogloeaceae |
| | 5a. | Conidia olivaceousDisculoides |
| | 5a. | Conidia hyaline3b |
| | 5b. | Conidiomata acervular, sides of conidiomatal wall appearing dark brown to black cells of <i>textura angularis</i> , conidiogenous cells lageniform to cylindrical, conidia monomorphic |
| | 5b. | Conidiomata subglobose, conidiomatal wall uniformly comprises orange-brown cells of <i>textura angularis</i> conidiogenous cells ampulliform, conidia dimorphicChrysocrypta |
| 5. | | Conidiophores subcylindrical, branched, 0–3-septate, conidia with obtuse apexProsopidicolaceae |
| 6. | | Conidia ovoid, obclavate to conical7 |
| 6. | | Conidia transversely distoseptate, consisting of four armsAsterosporiaceae |
| 7. | | Conidial wall smooth on the outer surface, with inconspicuous to distinct irregular verrucae on the inner surfaceJuglanconidaceae |
| 7. | | Conidial wall without ornamentation8 |
| 8. | | Conidia ovoid, 1-septate |
| 8. | | Conidia obclavate to conical, aseptataApoharknessiaceae |
| | 8a. | Conidia obclavate, with a scar at the baseLasmenia |
| | | (continued on next page) |

(Continued).

| | 8a. | Conidia conical, with small appendage at base and apical apiculus |
|----|------|--|
| 9. | | Stromata well-developed10 |
| 9. | | Stromata absent or poorly developed |
| 10 |). | Stromatic tissues orange, becoming purple in KOHCryphonectriaceae |
| | 10a. | Asexual morph not reported, hamathecium paraphysate, ascospores brownChromendothia |
| | 10a. | Asexual morph reported, hamathecium aparaphysate, ascospores hyaline10b |
| | 10b. | Coelomycetes10c |
| | 10b. | Ascomycetes10i |
| | 10c. | Conidia sigmoid with obtuse to subobtuse apex and swollen, obtuse baseAurantiosacculus |
| | 10c. | Conidia cylindrical, ellipsoid to fusoid, occasionally allantoid |
| | 10d. | Conidiomata pulvinate10e |
| | 10d. | Conidiomata rostrate, pyriform to globose10g |
| | 10e. | Neck presentLuteocirrhus |
| | 10e. | Neck absent |
| | 10f. | Conidiomata superficial, pale to medium brown, conidiophores consisting of basal subglobular to angular cells, that branch irregularly, becoming cylindrical, transversely septate |
| | 10f. | Conidiomata immersed to erumpent, orange, conidiophores aseptate, occasionally with separating septa and branching |
| | 10g. | Conidiophores hyaline, cylindrical, delimited by septa or not, conidia cylindricalUrsicollum |
| | 10g. | Conidiophores reduced to conidiogenous cells, conidia fusoid-ellipsoid or allantoid10h |
| | 10h. | Conidia with apical appendageMastigosporella |
| | 10h. | Conidia without apical appendageChrysofolia |
| | 10i. | Conidiomata uniformly orange10j |
| | 10i. | Conidiomata uniformly brown to black, with or without orange necks |
| | 10j. | Conidiomata pulvinate to globose10k |
| | 10j. | Conidiomata conical, rostrate, pyriform or convex10q |
| | 10k. | Ascospores septate10I |
| | 10k. | Ascospores aseptate10p |
| | 101. | Ascostromata superficial, conidiomata paraphysate10m |
| | 10I. | Ascostromata immersed to erumpent, conidiomata aparaphysate 10n |
| | 10m. | Perithecia valsoidDiversimorbus |
| | 10m. | Perithecia diatrypoidMicrothia |
| | 10n. | Conidiomata usually more than 350 µm diam, ascospores with median septumCryphonectria |
| | 10n. | Conidiomata usually less than 350 µm diam, ascospores with median to submedian septum10o |
| | 10o. | Ascospores oval to ellipsoid, papilla parallel to each other and open individuallyCryptometrion |
| | 10o. | Ascospores fusiform, sometimes curved, papilla close to each other and converge at the apexAurantioporthe |
| | 10p. | Stromata strongly developed, large, erumpent, mostly superficial, numerous conidial locules, no paraphysesEndothia |
| | 10p. | Stromata small to medium, semi-immersed, few conidial locules or one convoluted locule, paraphyses present |
| | | (continued on next page) |

| (00111 | nucu). |
|--------|--|
| 10q. | Conidiomata with necks10r |
| 10q. | Conidiomata without necks10s |
| 10r. | Conidiomata with prominent, delimited neckLatruncellus |
| 10r. | Conidiomata with neck continuous with base, rostrate, white sheath of tissue surrounding perithecial necksRostraureum |
| 10s. | Conidiomata conical, uniformly orangeAmphilogia |
| 10s. | Conidiomata convex, with blackened ostiolar openingsAurifilum |
| 10t. | Conidiomata uniformly black when mature10u |
| 10t. | Conidiomata black with orange neckAurapex |
| 10u. | Conidiomata base tissue of <i>textura globulosa</i> when sectioned longitudinally, perithecial necks long and covered with dark tissue, emerging from orange stromaChrysoporthe |
| 10u. | Conidiomata base tissue prosenchymatous, apices of conidiomata can be orange to scarlet when young, perithecial necks short, orange to umber stromaCeloporthe |
| 10. | Stromatic tissues dark brown to black, not becoming purple in KOH11 |
| 11. | Perithecia with very long, narrow, wavy ostiolar neck opening to or around ectostromatic disc12 |
| 11. | Perithecia with medium to short, somewhat wide, straight ostiolar neck opening to host surface14 |
| 12. | Ectostroma conspicuous |
| 12. | Ectostroma inconspicuous13 |
| 13. | Ascospores distoseptataStilbosporaceae |
| 13a. | Conidia hyaline with several tubular, unbranched, filiform apical appendagesCrinitospora |
| 13a. | Conidia brown without any appendages13b |
| 13b. | Ascospores and conidia with three transverse eusepta, ellipsoid to oblong; asci without a refractive canal in the apexStilbospora |
| 13b. | Ascospores and conidia with more than three transverse distosepta, ascospores sometimes and conidia always with additional longitudina distosepta, ascospores ellipsoid to oblong, conidia mostly pyriform; asci with a cylindrical, slightly refractive canal in the apexStegonsporium |
| 13. | Ascospores not distoseptata |
| 13a. | Coelomycetous |
| 13a. | Ascomycetous |
| 13b. | Conidiomata subcuticular. acervularGreeneria |
| 13b. | Conidiomata epiphyllous or hypophyllous with radiate scutella |
| | Tubakia |
| 13c. | Astromatic with solitary ascomata13d |
| 13c. | Stromatic with aggregated ascomataMellanconiella |
| 13d. | Ascospores ellipsoid without appendagesDicarpella |
| 13d. | Ascospores fusiform with appendagesMicroascospora |
| 14. | Perithecia arranged in valsoid configuration15 |
| 14. | Perithecia not arranged in valsoid configuration16 |
| 15. | Conidiomata non-loculate and forming both alpha and beta conidia |
| 15a. | Coelomycetes15b |
| 15a. | Ascomycetes15e |
| 15b. | Conidia olivaceous15c |
| 15b. | Conidia brown15d |
| 15c. | Conidia aseptate, guttulate, elongate fusiform to sigmoidPustulomyces |
| | |

(Continued).

| 15d. | Conidia dimorphic, aseptate, ellipsoid to pyriformPhaeocytostroma |
|------|---|
| 15d. | Conidia monomorphic, uniseptate, subcylindrical to narrowly ellipsoid |
| 15e. | Ascospores brownPhaeodiaporthe |
| 15e. | Ascospores hyaline15f |
| 15f. | Ascospores aseptataMazzantia |
| 15f. | Ascospores septate15g |
| 15g. | Septa submedian, large cell usually 2-guttulate, small cell usually 1-guttulate |
| 15g. | Septa median, with or without guttules15h |
| 15h. | Ascospores with long slender, thread-like appendage at both ends |
| 15h. | Ascospores without appendages15i |
| 15i. | Papilla long-cylindrical, conidia globose to subglobose, multiguttulata |
| 15i. | Papilla short, conidia absent or if present; ovate to ellipsoidal, biguttulata15j |
| 15j. | Ascospores ovoid, not constricted at the septa15k |
| 15j. | Ascospores fusiform to elongate-ellipsoid, constricted at the septa 151 |
| 15k. | Asci form long, pointed apex by narrowing towards the apical ring Chiangraiomyces |
| 15k. | Asci form blunt apexLeucodiaporthe |
| 151. | Ascospores overlapping uniseriate, often with 4 guttules, larger guttules at the center and smaller ones at the ends |
| 151. | Ascospores biseriate, without guttulesAllantoporthe |
| 15m. | Ascospores fusiform, ends pointed, papilla short and wideParadiaporthe |
| 15m. | Ascospores elongate to elliptical, ends round, papilla long and narrowDiaporthe |
| 15. | Conidiomata loculate forming numerous interconnecting chambers arranged radially or irregularly with in ectostromatic tissues and without forming beta conidiaCytosporaceae |
| 15a. | CoelomycetesWaydora |
| 15a. | Ascomycetes15b |
| 15b. | Stromata inconspicuous, ascospores allantoids15c |
| 15b. | Stromata conspicuous, well-developed, ascospores fusiform |
| 15c. | Ascomata solitaryParavalsa |
| 15c. | Ascomata aggregatedXenotypa |
| 15d. | Perithecia in groups with convergent beaks; asci clavate to fusoid <i>Cytospora</i> |
| 15d. | Perithecia in groups with non-convergent beaks; asci more or less rectangularPachytrype |
| 16. | Conidia dark brown, broadly fusiform to cylindrical or clavate, 3-5-cellular, distoseptataCoryneaceae |
| 16. | Conidia hyaline, ellipsoid, unicellularSydowiellaceae |
| 16a. | Stromata conspicuous, well-developed16b |
| 16a. | Stromata absent, inconspicuous, poorly developed16d |
| 16b. | Ascomata valsoid, stromatic tissues do not turn any colour with 10 % KOH16c |
| 16b. | Ascomata diatrypoid, stromatic tissues form dull red with 10 % KOH |
| 16c. | Ascospores oval to fusoid-oval, 1-septate, hyaline or hyaline to brownChapeckia |

| 16c. | Ascospores fusiform to ellipsoid with long filiform basal cell, 2-septate, hyalineRanulospora |
|------|---|
| 16d. | Ascomata solitary16e |
| 16d. | Ascomata aggregated16i |
| 16e. | Asci 4-sporedBreviappendix |
| 16e. | Asci 8-spored16f |
| 16f. | Ascospores apiosporousLambro |
| 16f. | Ascospores non-apiosporous16g |
| 16g. | Ascospores oval, hyaline to brown, not constricted at the septa |
| 16g. | Ascospores fusoid, hyaline, constricted at the septa16h |
| 16h. | Ascospores non-appendaged, globules at the center of each cell Svdowiella |
| 16h. | Ascospores appendaged, globules at the septa of the spore |
| 16i. | Ascospores oval, short fusoid to ellipsoidal16j |
| 16i. | Ascospores long fusoid cylindrical16n |
| 16j. | Ascospores 1-septate16k |
| 16j. | Ascospores multi-septate16m |
| 16k. | Ascospores with long strip-like appendagesTenuiappendicula |
| 16k. | Ascospores with short, appendages16 |
| 161. | Ascospores ellipsoid to cylindrical, usually 4-guttulateAlborbis |
| 161. | Ascospores fusiform, usually multi-quttulateParagnomonia |
| 16m | Ascospores hyaline to brown, ellipsoidal with broadly rounded ends, 1–3-septate, constricted at the septa |
| 16m | Ascospores hyaline, fusoid to oblong, 3-4-eusepta, not constricted at the septa |
| 16n. | Ascospores 0–5-septate |
| 16n | Ascospores 6–11-septate Rossmania |
| 17 | Ascospores unicellular 18 |
| 17 | Ascospores multicellular 19 |
| 18. | Ascomata superficial to erumpent, conidia elongate ellipsoidal to fusiform without appendages |
| 18. | Ascomata immersed, conidia oval to globose with basal appendage |
| 19. | Perithecia with 2–3 necks opening on both sides of the substrate/leaf blade |
| 19. | Perithecia with one neck opening to upper side of the substrate/leaf blade |
| 20. | Papilla short, conidia with microcyclic conidiation Pseudoplagiostomataceae |
| 20. | Papilla long, conidia do not have microcyclic conidiation |
| | Gnomoniaceae |
| 20a. | Coelomycetes20b |
| 20a. | Ascomycetes |
| 20b. | Conidia hyaline20c |
| 20b. | Conidia brownUniseta |
| 20c. | Conidiomata pycnidia20e |
| 20c. | Conidiomata acervuli |
| 20d. | Conidia 1-septateSirococcus |
| 20d. | Conidia aseptateMillerburtonia |
| 20e. | Conidia filiform to fusiformAsteroma |
| 20e. | Conidia ellipsoidalCvlindrosporella |
| 20f. | |
| | |

| (continued | on | next | page) |
|------------|----|------|-------|
|------------|----|------|-------|

(Continued).

| Conta | nucu). |
|-------------|---|
| | Pseudostromata/ stromata absent; perithecia immersed in host tissues |
| 20f. | Pseudostromata/ stromata present; perithecia immersed in stromatic tissues20u |
| 20g. | Perithecial necks not parallel to substrate20h |
| 20g. | Perithecial necks parallel to substrate and not fused |
| 20h | Infected lesions distinct with dark number to brown normanization or |
| 2011. | blackish area with pale brown sharp margin |
| 20h. | Infected lesions indistinct20j |
| 20i. | Ascospores apiosporousApioplagiostoma |
| 20i. | Ascospores non-apiosporousDiplacella |
| 20j. | Ascospores slightly isthmoid with a median septum, often readily separate as part sporesPleuroceras |
| 20j. | Ascospores non-isthmoid, do not separate into part spores20k |
| 20k. | Appendages generally present20I |
| 20k. | Appendages generally absent200 |
| 201. | Perithecia mostly epiphyllous20m |
| 201. | Perithecia mostly hypophyllous20n |
| 20m. | Appendages ovoid to subulateAmbarignomonia |
| 20m. | Appendages cuneiform with diffuse ends or ovoid, subulate acicular |
| 20n. | Ascospores fusiform; arranged irregularly fasciculate or, obliquely in one longitudinal rowApiognomonia |
| 20n. | Ascospores oval to filiform; arranged unevenly parallel, irregularly multiseriate or obliquely uniseriate, occasionally evenly parallel Onligonomonia |
| 200 | Perithecia occurring on both sides of the bost leaf |
| 200. | Parithecia occurring on only upper or lower side of host leaf 20n |
| 200. 20n | Necke present |
| 20p. | Necks present |
| 20p. | Necks absent |
| 20q. | Spataporthe |
| 20q. | Bell-shaped to hemispherical chamber at base of neck absent |
| 20r. | Asci 32-spored;Ditopella |
| 20r. | Asci 8-spored |
| 20s. | Ascomata immersed in pale brownish, parenchymatous cellular cavity and ascomata easily separate from them <i>Marsupiomyces</i> |
| 20s. | Ascomata not immersed in any, parenchymatous cellular pockets |
| 20t. | Stromata blackGnomoniopsis |
| 20t. | Stromata grey, brownish, cream, yellowish white |
| 20u. | Ectostromatic disc presentDitopellopsis |
| 20u. | Ectostromatic disc absent |
| 20v. | Stromatic tissues grey to pale brown; on top of perithecia and cream yellow mycelium at bottom of peritheciaOccultocarpon |
| 20v. | Stromatic tissues black; do not form any mycelium at bottom of perithecia |
| 20w. | Ascospores septate20x |
| 20w. | Ascospores aseptate |
| 20x. | Cytoplasm of ascospores granular and divide into two parts with wide vacuous space forming diplastic polarity |
| 20x. | Cytoplasm of ascospores does not divide into any parts20v |
| 20y. | Ascospores apiosporous20z |
| | (continued on next page) |

| 20y. Ascospores non-apiosporous20aa |
|---|
| 20z. Ascospores form brown large cell and small hyaline cell at maturity |
| 20z. Ascospores form both cells hyaline at maturity |
| 20aa. Ascospores 3-septatePhragmoporthe |
| 20aa. Ascospores 1-euseptate |
| 20ab. Stromata immersed in host tissues in bark, ascospores fusiform |
| 20ab. Stromata erumpent to superficial on leaves, ascospores ovalPhylloporthe |
| 20ac. Ascospores oval to ellipsoidAmphiporthe 20ac. Ascospores allantoidValsalnicola |

List of accepted families and genera in Diaporthales

Apiosporopsidaceae Senan. et al. Apiosporopsis (Traverso) Mariani Apoharknessiaceae Senan. et al. Apoharknessia Crous & S.J. Lee Lasmenia Speg. Asterosporiaceae Senan. et al. Asterosporium Kunze Auratiopvcnidiellaceae Senan. et al. Auratiopycnidiella Crous & Summerell Coryneaceae Corda = Pseudovalsaceae M.E. Barr Coryneum Nees Cryphonectriaceae Gryzenh. & M.J. Wingf. Amphilogia Gryzenh. et al. Aurantiosacculus Dyko & B. Sutton Aurapex Gryzenh. & M.J. Wingf. Aurifilum Begoude et al. Celoporthe Nakab. et al. Chromendothia Lar.N. Vassiljeva Chrysofolia Crous & M.J. Wingf. Chrysoporthe Gryzenh. & M.J. Wingf. =Chrysoporthella Gryzenh. & M.J. Wingf. Cryphonectria (Sacc.) Sacc. & D. Sacc. Cryptometrion Gryzenh. & M.J. Wingf. Diversimorbus S.F. Chen & J. Roux Endothia Fr. Foliocryphia Cheewangkoon & Crous Holocryphia Gryzenh. & M.J. Wingf. Immersiporthe S.F. Chen et al. Latruncellus M. Verm. et al. Luteocirrhus C.F. Crane & T.I. Burgess Mastigosporella Höhn. Microthia Gryzenh. & M.J. Wingf. Rostraureum Gryzenh. & M.J. Wingf. Ursicollum Gryzenh. & M.J. Wingf. Cytosporaceae Fr. =Valsaceae Tul. & C. Tul.

(Continued).

Cytospora Ehrenb. Pachytrype Berl. ex M.E. Barr et al. Paravalsa Ananthap Waydora B. Sutton Xenotypa Petr. Diaporthaceae Höhn. ex Wehm. Allantoporthe Petr. Apioporthella Petr. Chaetoconis Clem. Chiangraiomyces Senan. & K.D. Hyde Diaporthe Nitschke Hyaliappendispora Senan. et al. Leucodiaporthe M.E. Barr et al. Mazzantia Mont. Ophiodiaporthe Y.M. Ju et al. Paradiaporthe Senan. et al. Phaeocytostroma Petr. Phaeodiaporthe Petr. Pustulomyces D.Q. Dai et al. Stenocarpella Syd. & P. Syd. Erythrogloeaceae Senan. et al. Chrysocrypta Crous & Summerell Disculoides Crous et al. Erythrogloeum Petr. Gnomoniaceae G. Winter Alnecium Voglmayr & Jaklitsch Ambarignomonia Sogonov Amphiporthe Petr. Anisomyces Theiss. & Syd. Apiognomonia Höhn. Apioplagiostoma M.E. Barr Asteroma DC Bagcheea E. Müll. & R. Menon Clypeoporthe Höhn. Cryptosporella Sacc. Cylindrosporella Höhn. Diplacella Syd. Ditopella De Not. Ditopellopsis J. Reid & C. Booth Gloeosporidina Petr. Gnomonia Ces. & De Not. Gnomoniella Sacc. Gnomoniopsis Berl. Marsupiomyces Senan. & K.D. Hyde Millerburtonia Cif. Occultocarpon L.C. Mejía et al. Ophiognomonia (Sacc.) Sacc. Phragmoporthe Petr. Phylloporthe Syd. Plagiostoma Fuckel Pleuroceras Riess.

Sirococcus Preuss Spataporthe Bronson et al. Uniseta Ciccar Valsalnicola D.M. Walker & Rossman Harknessiaceae Crous Dwiroopa Subram. & Muthumary Harknessia Cooke Juglanconidaceae Voglmayr & Jaklitsch Juglanconis Voglmayr & Jaklitsch Lamproconiaceae C. Norphanphoun et al. Hercospora Fr. Lamproconium (Grove) Grove Macrohilaceae Crous Macrohilum H.J. Swart Melanconidaceae G. Winter Melanconis Tul. & C. Tul. Melanconiellaceae Senan. et al. Dicarpella Syd. Greeneria Scribn. & Viala Melanconiella Sacc. Microascospora Senan. & K.D. Hyde Tubakia B. Sutton Prosopidicolaceae Senan. & K.D. Hyde Prosopidicola Crous & C.L. Lennox Pseudoplagiostomataceae Cheew. et al. Pseudoplagiostoma Cheew. et al. Schizoparmaceae Rossman DF et al. Coniella Höhn. Stilbosporaceae Link Crinitospora B. Sutton & Alcorn Stegonsporium Corda Stilbospora Pers. Sydowiellaceae Lar.N. Vassiljeva Alborbis Senan. & K.D. Hyde Breviappendix Senan. & K.D. Hyde Cainiella E. Müll Calosporella J. Schröt Chapeckia M.E. Barr Italiomyces Senan. et al. Hapalocystis Auersw. ex Fuckel I ambro Racib Paragnomonia Senan. & K.D. Hyde Ranulospora Senan. et al. Rossmania Lar.N. Vassiljeva Sillia P. Karst. Sydowiella Petr. Tenuiappendicula Senan. et al. Tortilispora (Sacc.) Senan. & K.D. Hyde

(Continued).

Caudospora Starbäck Chadefaudiomyces Kamat et al. Cryptascoma Ananthap. Cryptoleptosphaeria Petr. Cytomelanconis Naumov Dictyoporthe Petr. Ditopellina J. Reid & C. Booth Durispora K.D. Hyde Fremineavia Nieuwl. Hypodermina Höhn. Hypophloeda K.D. Hyde & E.B.G. Jones Kapooria J. Reid & C. Booth Keinstirschia J. Reid & C. Booth Lollipopaia Inderbitzin Macrodiaporthe Petr. Maculatipalma J. Fröhlich & K.D. Hyde Massariovalsa Sacc. Mebarria J. Reid & C. Booth Melanamphora Lafl. Melanconiopsis Ellis & Everh. Natarajania Pratibha & Bhat Phaeoappendicospora Senan. et al. Phraqmodiaporthe Wehm. Plagiophiale Petr. Plagiostigme Syd. Prostratus Sivan. et al. Pseudocryptosporella J. Reid & C. Booth Pseudothis Theiss. & Syd. Pseudovalsella Höhn. Rabenhorstia Fr. Savulescua Petr. Skottsbergiella Petr. Stioclettia Dennis Trematovalsa Jacobesco Uleoporthe Petr. Vismaya V.V. Sarma & K.D. Hyde Wehmeyera J. Reid & C. Booth Wuestneia Auersw. ex Fuckel Wuestneiopsis J. Reid & Dowsett

ACKNOWLEDGEMENTS

We thank Elias Jonk, Westerdijk Fungal Biodiversity Institute, for DNA isolation and sequencing of some of the included CBS cultures. Alan J.L. Phillips acknowledges the support from Biosystems and Integrative Sciences Institute (BioISI, FCT/UID/ Multi/04046/2013). Kevin D. Hyde thanks to National Research Council of Thailand (Mae Fah Luang University) for the grant "Biodiversity, phylogeny and role of fungal endophytes of Pandanaceae" (Grant No: 592010200112) and Thailand Research Fund (TRF) grant no RSA5980068 entitled "Biodiversity, phylogeny and role of fungal endophytes on above parts of *Rhizophora apiculata* and *Nypa fruticans*", the National Research Council of Thailand (Mae Fah Luang University) grant no 60201000201 entitled "Diseases of mangrove trees and maintenance of good forestry practice", and the Chinese Academy of Sciences, project number 2013T2S0030, for the award of Visiting Professorship for Senior International Scientists at Kunming Institute of Botany. Samantha C. Karunarathna thanks to

Diaporthales genera incertae sedis

Anisomycopsis I. Hino & Katum.

(continued on next page)



Yunnan Provincial Department of Human Resources and Social Security funded postdoctoral project (number 179122).

REFERENCES

- Alexopoulus CJ, Mims CW (1978). Introductory mycology, 3rd ed. John Wiley, New York, USA.
- Alvarez LV, Groenewald JZ, Crous PW (2016). Revising the Schizoparmaceae: Coniella and its synonyms Pilidiella and Schizoparme. Studies in Mycology 85: 1–34.
- Ananthapadmanaban D (1990). Paravalsa indica sp. nov. from India. Mycological Research 94: 275–276.
- Barney JN, Tharayil N, DiTommaso A, et al. (2006). The biology of invasive alien plants in Canada; Polygonum cuspidatum Sieb & Zucc [= Fallopia japonica (Houtt) Ronse Decr]. Canadian Journal of Plant Science 86: 887–905.
- Barr ME (1978). The *Diaporthales* in North America: with emphasis on *Gnomonia* and its segregates. *Mycologia Memoirs* **7**: 1–232.
- Begoude BAD, Slippers B, Wingfield MJ, et al. (2010). Botryosphaeriaceae associated with Terminalia catappa in Cameroon, South Africa and Madagascar. Mycological Progress 9: 101–123.
- Belisario A (1991). Dicarpella dryina sp. nov., teleomorph of Tubakia dryina. Mycotaxon 41: 147–155.
- Belisario A (1999). Cultural characteristics and pathogenicity of *Melanconium* juglandinum. European Journal of Forest Pathology **29**: 317–322.
- Biggs AR (1989). Integrated control of Leucostoma-canker of peach peach in Ontario. *Plant Disease* **73**: 869–874.
- Braun U, Bien S, Hantsch L, *et al.* (2014). *Tubakia chinensis* sp. nov. and a key to the species of the genus *Tubakia*. *Schlechtendalia* **28**: 23–28.
- Campbell J, Anderson JL, Shearer CA (2003). Systematics of Halosarpheia based on morphological and molecular data. Mycologia 95: 530–552.
- Cannon PF, Minter DW (2014). Lamproconium desmazieresii. IMI descriptions of fungi & bacteria: 1996. CABI Bioscience, Wallingford, UK: 1–3.
- Carbone I, Kohn LM (1999). A method for designing primer sets for speciation studies in filamentous ascomycetes. *Mycologia* **91**: 553–556.
- Castlebury LA, Farr DF, Rossman AY, et al. (2003). Diaporthe angelicae comb nov, a modern description and placement of Diaporthopsis in Diaporthe. Mycoscience 44: 203–208.
- Castlebury LA, Rossman AY, Jaklitsch WJ, et al. (2002). A phylogeny overview of the Diaporthales based on large subunit nuclear ribosomal DNA sequences. Mycologia 94: 1017–1031.
- Cavara F (1889). Matériaux de mycologie Lombarde. *Revue Mycologique* 11: 173–1913.
- Chadefaud M (1960). Les Végétaux non Vasculaires (Cryptogamie). In: *Traité de Botanique Systématique* (Chadefaud M, Emberger L, eds). Masson, Paris: 1–1018.
- Cheewangkoon R, Groenewald JZ, Verkley GJM, et al. (2010). Re-evaluation of *Cryptosporiopsis eucalypti* and cryptosporiopsis-like species occurring on *Eucalyptus* leaves. *Fungal Diversity* **44**: 89–105.
- Chen SF, Wingfield MJ, Roux J (2013). *Diversimorbus metrosiderotis* gen. et sp. nov. and three new species of *Holocryphia* (*Cryphonectriaceae*) associated with cankers on native *Metrosideros angustifolia* trees in South Africa. *Fungal Biology* **117**: 289–310.
- Chomnunti P, Hongsanan S, Aguirre-hudson B, et al. (2014). The sooty moulds. Fungal Diversity 66: 1–36.
- Corda ACI (1839). Coniomycetes Nees ab Esenb. *Icones Fungorum hucusque Cognitorum* **3**: 1–55.
- Crane C, Burgess TI (2013). *Luteocirrhus shearii* gen. sp. nov. (*Diaporthales*, *Cryphonectriaceae*) pathogenic to *Proteaceae* in the South Western Australian Floristic Region. *IMA Fungus* **4**: 111–122.
- Crous PW, Carris LM, Giraldo A, *et al.* (2015). The Genera of Fungi: fixing the application of the type species of generic names G 2: *Allantophomopsis, Latorua, Macrodiplodiopsis, Macrohilum, Milospium, Protostegia, Pyricularia, Robillarda, Rotula, Septoriella, Torula, and Wojnowicia. IMA Fungus* **6**: 163–198.
- Crous PW, Gams W, Stalpers JA, et al. (2004). MycoBank: an online initiative to launch mycology into the 21st century. *Studies in Mycology* **50**: 19–22.
- Crous PW, Knox-Davies PS, Wingfield MJ (1989). Newly-recorded foliage fungi of *Eucalyptus* sp. in South Africa. *Phytophylactica* **21**: 85–88.
- Crous PW, Rogers JD (2001). Wuestneia molokaiensis and its anamorph Harknessia molokaiensis sp. nov. from Eucalyptus. Sydowia 53: 74–80.
- Crous PW, Summerell BA, Alfenas AC, et al. (2012a). Genera of diaporthalean coelomycetes associated with leaf spots of tree hosts. *Persoonia* 28: 66–75.

- Crous PW, Summerell BA, Shivas RG, et al. (2012b). A re-appraisal of Harknessia (Diaporthales), and the introduction of Harknessiaceae fam. nov. Persoonia 28: 49–65.
- Crous PW, Summerell BA, Shivas RG, et al. (2012c). Fungal Planet description sheets: 107–127: molecular phylogeny and evolution of fungi. *Persoonia* 28: 138–182.
- Crous PW, Verkley GJM, Groenewald JZ, et al. (2009). Fungal biodiversity. CBS laboratory manual series: 1. Centraalbureau voor Schimmelcultures, Utrecht, Netherlands: 1–269.
- Crous PW, Wingfield MJ, Burgess TI, *et al.* (2017). Fungal Planet description sheets: 558–624. *Persoonia* **38**: 240–384.
- Crous PW, Wingfield MJ, Guarro J, et al. (2013). Fungal Planet description sheets: 154–213. Persoonia **31**: 188–296.
- Crous PW, Wingfield MJ, Nag Raj TR (1993). *Harknessia* species occurring in South Africa. *Mycologia* **85**: 275–280.
- Crous PW, Wingfield MJ, Richardson DM, et al. (2016). Fungal Planet description sheets: 400–468. Persoonia **36**: 316–458.
- Dai DQ, Wijayawardene NN, Bhat DJ, et al. (2014). Pustulomyces gen. nov. accommodated in Diaporthaceae, Diaporthales, as revealed by morphology and molecular analyses. Cryptogamie, Mycologie 35: 63–72.
- De Silva H, Castlebury LA, Green S, *et al.* (2009). Characterisation and phylogenetic relationships of *Anisogramma virgultorum* and *A. anomala* in the *Diaporthales* (Ascomycota). *Mycological Research* **113**: 73–81.
- Du Z, Hyde KD, Yang Q, et al. (2017). Melansporellaceae: a novel family of Diaporthales (Ascomycota). Phytotaxa 305: 191–200.
- Eriksson OE (2001). Outline of Ascomycota. Myconet 6: 1-27.
- Eriksson OE, Winka K (1997). Supraordinal taxa of Ascomycota. Myconet 1: 1–16.
- Fabre JH (1883). Essai sur les Sphériacées du département de Vaucluse [concl]. Annales des Sciences Naturelles Botanique 6: 31–43.
- Farr DF, Bills GF, Chamuris GP, et al. (1989). Fungi on plant and plant products in the United States. APS Press, St Paul, MN.
- Farr DF, Rossman AY (2001). Harknessia lythri, a new species on purple loosestrife. Mycologia 93: 997–1001.
- Farr DF, Rossman AY (2017). Fungal databases. Systematic Mycology and Microbiology Laboratory, ARS, USDA, Beltsville, MD. http://ntars-gringov/ fungaldatabases/.
- Felsenstein J (1985). Confidence limits on phylogenies: an approach using the bootstrap. *Evolution* **39**: 783–791.
- Ferreira FA, Demuner NL, Rezende DV (1992). Mancha de folha, des folha e antracnose do Jatobá (*Hymenaea* spp) causadas por *Erythrogloeum hymenaeae*. *Fitopatologia Brasileira* **17**: 106–109.
- Gao Y, Liu F, Duan W (2017). *Diaporthe* is paraphyletic. *IMA Fungus* 8: 153–187.
- Glass NL, Donaldson G (1995). Development of primer sets designed for use with PCR to amplify conserved genes from filamentous ascomycetes. *Applied and Environmental Microbiology* **61**: 1323–1330.
- Gomes RR, Glienke C, Videira SIR, et al. (2013). Diaporthe: a genus of endophytic, saprobic and plant pathogenic fungi. Persoonia **31**: 1–41.
- Graves AH (1923). The *Melanconis* disease of the butternut (*Juglans cinerea* L). *Phytopathology* **13**: 411–435.
- Grove WB (1937). British stem and leaf-fungi. Coelomycetes 2: 1-406.
- Gryzenhout M, Myburg H, Hodges CS, et al. (2006a). Microthia, Holocryphia and Ursicollum, three new genera on Eucalyptus and Coccoloba for fungi previously known as Cryphonectria. Studies in Mycology 55: 35–52.
- Gryzenhout M, Myburg H, Rodas CA, et al. (2006b). Aurapex penicillata gen. sp. nov. from native Miconia theaezans and Tibouchina spp. in Colombia. Mycologia 98: 105–115.
- Gryzenhout M, Myburg H, Wingfield BD (2006c). Cryphonectriaceae (Diaporthales), a new family including Cryphonectria, Chrysoporthe, Endothia and allied genera. Mycologia **98**: 239–249.
- Gryzenhout M, Tarigan M, Clegg PA, et al. (2010). Cryptometrion aestuescens gen. sp. nov. (Cryphonectriaceae) pathogenic to Eucalyptus in Indonesia. Australasian Plant Pathology 39: 161–169.
- Gryzenhout M, Wingfield BD, Wingfield MJ (2009). Taxonomy, phylogeny and ecology of bark-infecting and tree-killing fungi in the Cryphonectriaceae. APS Press, St Paul, Minnesota: 1–378.
- 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, et al. (1995). Ainsworth & Bisby's dictionary of the fungi, 8th ed. CAB International, Wallingford.
- Hepting GH (1974). Death of the American chestnut. *Journal of Forest History* **18**: 60–67.

Index Fungorum (2017). Retrieved 24 April 2017, from http://wwwindexfungorumorg/Names/Namesasp.

- Jackson T (2003). Occurrence and variation of Endothiella eucalypti in Eucalyptus globulus plantations of south-western Australia and the influence of some biotic and abiotic factors on the response of the host to the pathogen. Ph.D. dissertation. School of Biological Sciences and Biotechnology, Murdoch University Perth, Western Australia.
- Jeewon R, Hyde KD (2016). Establishing species boundaries and new taxa among fungi: recommendations to resolve taxonomic ambiguities. *Mycosphere* **7**: 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.
- Jones EBG, Suetrong S, Sakayaroj J, *et al.* (2015). Classification of marine Ascomycota, Basidiomycota, Blastocladiomycota and Chytridiomycota. *Fungal Diversity* **73**: 1–72.
- Kepley JB, Jacobi WR (2000). Pathogenicity of Cytospora fungi on six hardwood species. Journal of Aboriculture 26: 326–332.
- 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.
- Lamprecht SC, Crous PW, Groenewald JZ, et al. (2011). Diaporthaceae associated with root and crown rot of maize. IMA Fungus 2: 13–24.
- Lee S, Groenewald JZ, Crous PW (2004). Phylogenetic reassessment of the coelomycete genus *Harknessia* and its teleomorph *Wuestneia* (*Diaporthales*), and the introduction of *Apoharknessia* gen. nov. *Studies in Mycology* **50**: 235–252.
- Lennox CL, Serdani M, Groenewald JZ, et al. (2004). Prosopidicola mexicana gen. et. sp. nov., causing a new pod disease of Prosopis species. Studies in Mycology 50: 187–194.
- Link HF (1826). Entwurf eines phytologischen Pflanzensystems nebst einer Anordnung der Kryptogamen. Abhandlungen der königlichen Akademie der Wissenschaften zu Berlin aus dem Jahre 1824: 145–194.
- 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.
- Locquin MV (1984). Mycologie Générale et Structurale. Masson, Paris.
- 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.
- Luttrell ES (1951). Taxonomy of Pyrenomycetes. University of Missouri Studies 24: 1–120.
- Maharachchikumbura SS, Hyde KD, Jones EBG, et al. (2015). Towards a natural classification and backbone tree for Sordariomycetes. Fungal Diversity 72: 199–301.
- Maharachchikumbura SSN, Hyde KD, Jones EBG, et al. (2016). Families of Sordariomycetes. Fungal Diversity 79: 1–317.
- Mariani (1911). Pyrenomycetae, Sphaeriaceae, Guignardia. Atti della Societa Italiana di Scienze Naturali e del Museo Civico di Storia Naturale di Milano 50: 1–165.
- Marincowitz S, Crous PW, Groenewald JZ, et al. (2008). Microfungi occurring on *Proteaceae* in the fynbos. *CBS Biodiversity Series* **7**: 1–166.
- Mejía LC, Castlebury LA, Rossman AY, et al. (2011). A systematic account of the genus Plagiostoma (Gnomoniaceae, Diaporthales) based on morphology, hostassociations, and a four-gene phylogeny. Studies in Mycology 68: 211–235.
- Monod M (1983). Monographie taxonomique des *Gnomoniaceae* (Ascomycetes de l'ordre des *Diaporthales*) | Beih. *Sydowia* 9: 1–315.
- Mostert L, Groenewald JZ, Summerbell RC, et al. (2006). Taxonomy and pathology of Togninia (Diaporthales) and its Phaeoacremonium anamorphs. Studies in Mycology 54: 1–115.
- MycoBank (2017). http://wwwmycobankorg/defaultinfoaspx?Page=Home.
- Nag Raj TR (1993). Coelomycetous anamorphs with appendage-bearing conidia. Mycologue Publications, Waterloo, Canada: 1–1101.
- Nakabonge G, Gryzenhout M, Roux J, et al. (2006). Celoporthe dispersa gen. et sp. nov. from native Myrtales in South Africa. Studies in Mycology 55: 255–267.
- Nannfeldt JA (1932). Studien über die Morphologie und Systematik der nichlichenisierten inoperculaten Discomyceten. Nova Acta Regiae Societatis Scientiarum Upsaliensis Series 4(8): 1–368.
- Navarrete F, Abreo E, Bettucci L, *et al.* (2009). First report of *Greeneria uvicola* as cause of grapevine dead-arm dieback in Uruguay. *Australasian Plant Disease Notes* **4**: 117–119.
- Norphanphoun C, Doilom M, Daranagama DA, *et al.* (2017). Revisiting the genus *Cytospora* and allied species. *Mycosphere* **8**: 51–97.

- Norphanphoun C, Hongsanan S, Doilom M, et al. (2016). Lamproconiaceae fam. nov. to accommodate Lamproconium desmazieri. Phytotaxa 270: 89–102.
- Nylander JAA (2004). *MrModeltest*. Program distributed by the Author. Evolutionary Biology Centre, Uppsala University. v. 2.
- Petrak F (1938). Beiträge zur Kenntnis der Gattung Hercospora mit besonderer Berücksichtigung ihrer Typusart Hercospora tiliae (Pers) Fr. Annales Mycologici 36: 44–60.
- Petrak F (1953). Erythrogloeum nov gen, eine neue Gattung der Sphaeropsideen. Sydowia 7: 378–380.
- Petrak F (1971). Ergebnisse einer Revision der Grundtypen verscheidener Gattungen der Ascomyzeten und Fungi Imperfecti. Sydowia 24: 249–255.
- Phillips AJL, Alves A, Pennycook SR, et al. (2008). Resolving the phylogenetic and taxonomic status of dark-spored teleomorph genera in the *Botryos-phaeriaceae*. Persoonia 21: 29–55.
- Potebnia A (1910). Beiträge zur Micromycetenflora Mittel-Russlands (Gouv Kursk und Charkow). Annales Mycologici 8: 42–93.
- Punithalingam E (1974). Studies on Sphaeropsidales in culture II. Mycological Papers 136: 1–63.
- Rambaut A (2012). Fig.Tree. Tree figure drawing tool. v. 1.4.0. http://tree.bio.ed. ac.uk/software/figtree/.
- Rayner RW (1970). A mycological colour chart. CMI and British Mycological Society, Kew, UK.
- Réblová M, Mostert L, Gams W, et al. (2004). New genera in the Calosphaeriales: Togniniella and its anamorph Phaeocrella, and Calosphaeriophora as anamorph of Calosphaeria. Studies in Mycology 50: 533–550.
- Rehner SA, Samuels GJ (1994). Taxonomy and phylogeny of *Gliocladium* analysed from nuclear large subunit ribosomal DNA sequences. *Mycological Research* 98: 625–634.
- Reid J, Dowsettj A (1990). On Dicarpella, Sphaerognomonia and Apiosporopsis. Canadian Journal of Botany 68: 2398–2407.
- Ronquist F, Teslenko M, van der Mark P, et al. (2012). MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across a large model space. Systematic Biology 61: 539–542.
- Rossman AY, Adams GC, Cannon PF, et al. (2015). Recommendations of generic names in *Diaporthales* competing for protection or use. *IMA Fungus* 6: 145–154.
- Rossman AY, Castlebury LA, Samuels GJ (2006). Phylogeny and biodiversity of the Hypocreales and Diaporthales. In: 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.
- Scribner FL, Viala P (1887). Le Greeneria fuliginea, nouvelle forme de rot des fruits de lab vigne observee en Amärique. Comptes rendus Hebdomadaires des Séances de l'Académie des Sciences 105: 473.
- Seifert KA, Rossman AY (2010). How to describe a new fungal species. *IMA Fungus* 1: 109–116.
- Senanayake IC, Maharachchikumbura SSN, Hyde KD, et al. (2015). Towards unraveling relationships in Xylariomycetidae (Sordariomycetes). Fungal Diversity 73: 73–144.
- Senanayake IC, Maharachchikumbura SSN, Jeewon R, et al. (2017). Morphophylogenetic study of Sydowiellaceae reveals several new genera. Mycosphere 8: 172–217.
- Serrato-Diaz LM, Rivera-Vargas LI, Goenaga R, et al. (2011). First report of a Lasmenia sp causing rachis necrosis, flower abortion, fruit rot, and leaf spots on rambutan in Puerto Rico. Plant Disease 95: 1311–1313.
- Shipton PJ (1967). A fruit rot of strawberries caused by *Zythia fragariae*. *Plant Pathology* **16**: 123–125.
- Sieber TN, Sieber-Canavesi F, Petrini O, et al. (1991). Characterization of Canadian and European *Melanconium* from some *Alnus* species by morphological, cultural and biochemical studies. *Canadian Journal of Botany* 69: 2170–2176.
- Silvestro D, Michalak I (2012). raxmlGUI: a graphical front-end for RAxML. Organisms Diversity & Evolution 12: 335–337.
- Sinclair WA, Hudler GW (1980). Tree and shrub pathogens new or noteworthy in New York state. *Plant Disease* **64**: 590–592.
- Sogonov MV, Castlebury LA, Rossman AY, et al. (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.
- Suetrong S, Klaysuban A, Sakayaroj J, et al. (2015). Tirisporellaceae, a new family in the order Diaporthales (Sordariomycetes, Ascomycota). Cryptogamie, Mycologie 36: 319–330.

- Sutton BC (1980). The Coelomycetes Fungi imperfecti with pycnidia, acervuli and stromata. Commonwealth Mycological Institute, Kew, UK.
- Suwannarach N, Kumla J, Lumyong S (2016). *Pseudoplagiostoma dipterocarpi* sp. nov., a new endophytic fungus from Thailand. *Mycoscience* **57**: 118–122.
- Swofford DL (2003). PAUP* phylogenetic analysis using parsimony (*and other methods). Sinauer Associates, Sunderland, Massachusetts. Version 4.
- Tanaka K, Mel'nik VA, Kamiyama M, et al. (2010). Molecular phylogeny of two coelomycetous fungal genera with stellate conidia, *Prosthemium* and *Asterosporium*, on Fagales trees. *Botany* 88: 1057–1071.
- Taylor JW, Jacobson DJ, Kroken S, et al. (2000). Phylogenetic species recognition and species concepts in fungi. Fungal Genetics and Biology **31**: 21–32.
- Traverso GB (1907). Flora Italica Cryptogama Pars 1: Fungi Pyrenomycetae Sphaeriaceae: Allantosporae, Hyalosporae, Phaeosporae: 1. Società Botanica Italiana, Rocca S Casciano, Italy: 353–492.
- Treigien A, Markovskaja S (2007). Microscopic fungi on *Carpinus Betulus* in Lithuania 2. *Anamorphic Fungi Botanica Lithuanica* **13**: 45–50.
- Udayanga D, Castlebury LA, Rossman AY, *et al.* (2014). Insights into the genus *Diaporthe*: phylogenetic species delimitation in the *D. eres* species complex. *Fungal Diversity* **67**: 203–229.
- Udayanga D, Xingzhong L, McKenzie EHC, *et al.* (2011). The genus *Phomopsis*: biology, applications, species concepts and names of common pathogens. *Fungal Diversity* **50**: 189–225.
- Van der Aa HA (1973). Studies in Phyllosticta I. Studies in Mycology 5: 1-110.
- Van Niekerk JM, Groenewald JZ, Verkley GJ, et al. (2004). Systematic reappraisal of Coniella and Pilidiella, with specific reference to species occurring on Eucalyptus and Vitis in South Africa. Mycological Research 108: 283–303.
- Vasilyeva LN (1993). Chromendothia, a new genus of the family Hypocreaceae. Mikologiya i Fitopatologiya 27: 1–7.
- Vasilyeva LN, Rossman AY, Farr DF (2007). New species of the Diaporthales from eastern Asia and eastern North America. Mycologia 99: 916–923.
- Verkley GJM, Quaedvlieg W, Shin HD, et al. (2013). A new approach to species delimitation in Septoria. Studies in Mycology 75: 213–305.
- Vermeulen M, Gryzenhout M, Wingfield MJ, et al. (2011). New records of the Cryphonectriaceae from southern Africa including Latruncellus aurorae gen. sp. nov. Mycologia 103: 554–569.
- 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, Castlebury LA, Jaklitsch WM (2017). Juglanconis gen. nov. on Juglandaceae, and the new family Juglanconidaceae (Diaporthales). Persoonia 38: 136–155.
- Voglmayr H, Jaklitsch WM (2008). Prosthecium species with Stegonsporium anamorphs on Acer. Mycological Research 112: 885–905.
- Voglmayr H, Jaklitsch WM (2014). Stilbosporaceae resurrected: generic reclassification and speciation. *Persoonia* 33: 61–82.
- VogImayr H, Rossman AY, Castlebury LA, et al. (2012). Multigene phylogeny and taxonomy of the genus Melanconiella (Diaporthales). Fungal Diversity 57: 1–44.
- Von Arx JA, Müller E (1954). Die Gattungen der amerosporen Pyrenomyceten. *Beiträge zur Kryptogamenflora der Schweiz* **11**: 1–434.
- Von Höhnel FXR (1910). Fragmente zur Mykologie no. 538. Sitzungsberichte der Kaiserlichen Akademie der Wissenschaften in Wien 119: 632–633.
- Von Höhnel FXR (1917). Über die Benennung Stellung und Nebenfruchtformen von Sphaerella Fries. Berichte der Deutschen Botanischen Gesellschaft 35: 627–631.
- Walker DM, Castlebury LA, Rossman AY, et al. (2010). Systematics of genus Gnomoniopsis (Gnomoniaceae, Diaporthales) based on a three gene phylogeny, host associations and morphology. Mycologia 102: 1479–1496.
- Walker DM, Castlebury LA, Rossman AY, et al. (2012). Phylogeny and taxonomy of Ophiognomonia (Gnomoniaceae, Diaporthales), including twenty-five new species in this highly diverse genus. Fungal Diversity 57: 85–147.
- Wehmeyer LE (1975). The Pyrenomycetous Fungi. Mycologia Memoirs 6: 1–250.
- White T, Bruns T, Lee S, et al. (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, San Diego, California: 315–322.
- Wijayawardene NN, Hyde KD, Wanasinghe DN, et al. (2016). Taxonomy and phylogeny of dematiaceous coelomycetes. Fungal Diversity 77: 1–316.
- Winter G (1886). Fungi Australienses. Revue Mycologique Toulouse 8: 207-213.
- Yuan ZQ, Mohammed C (1997). Wuestneia epispora sp. nov. on stems of eucalypts from Australia. Mycological Research 101: 195–200.
- Yuan ZQ, Wardlaw T, Mohammed C (2000). Harknessia species occurring on eucalypt leaves in Tasmania, Australia. Mycological Research 104: 888–892.
- Zhang N, Blackwell M (2001). Molecular phylogeny of dogwood anthracnose fungus (*Discula destructiva*) and the *Diaporthales*. *Mycologia* **93**: 355–365.