THE SYNNEMATOUS ANAMORPH OF *EXIDIA SACCHARINA* (AURICULARIALES, BASIDIOMYCOTA): MORPHOLOGY, CONIDIOGENESIS AND ASSOCIATION WITH BARK BEETLES*

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Abstract. The paper describes a synnematous anamorph of *Exidia saccharina*, its natural occurrence in the field, and the ultrastructure of conidiogenesis. Conidiogenesis was revealed as annellidic by transmission electron microscopy. The anamorph was found rarely in galleries of the bark beetles *Ips typographus* and *Pityogenes chalcographus* (Coeleoptera, Scolytinae) in the bark of *Picea abies* in Germany. The anamorph–teleomorph connection was confirmed by morphological comparison of the anamorph from bark beetle galleries with the anamorph grown from basidiospores on natural medium. The newly discovered synnematous growth and annellidic conidiogenesis are similar to those of *Gloeosynnema* species (basidiomycetous hyphomycetes), which differ by details of septal clamp connections.

Key words: basidiomycetes, mitosporic fungi, synnemata, Europe

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INTRODUCTION

Anamorphs of Auriculariales usually are characterized by hyaline, one-celled, mostly allantoid conidia which are produced from germinating basidiospores or mycelia derived from basidiospores. The anamorph of Exidia glandulosa (Bull.) Fr. is illustrated as an example in Figure 1 (from dead branch; GERMANY. Baden-Württemberg, Tübingen, Schönblick, 22 Febr. 1998, leg. R. Kirschner 308, FR). The conidiogenous cells have sometimes been called phialide-like (Ingold 1982a, b, 1984, 1992, 1995), but the exact mode of conidiogenesis is usually not evident by light microscopy and has not been investigated with transmission electron microscopy. Brefeld (1888) found considerable morphological differences among anamorphs of certain species of Auriculariales in culture, but this approach has not been used consistently for monographs of this group. Production of secondary spores and conidia from germinating basidiospores can be observed often on freshly collected basidiomata of Auriculariales, but the anamorphs alone are rarely found in nature (Olive 1952; Ingold 1991), or if they occur their association with a teleomorph has not been recognized. Recently an anamorph with a relationship to the Hyaloriaceae (Auriculariales) was described from material on a dead branch in Taiwan and in culture as Helicomyxa everhartioides R. Kirschner & Chee J. Chen (Kirschner & Chen 2004). Another anamorph was found on dead bamboo in China (Kirschner et al., in press). Both species form superficial pustulate or cupulate conidiomata on dead plant material, and their asexual propagules probably are dispersed by rain. Another habit connected to a cryptic habitat hidden beneath tree bark and probable dispersal by insects is demonstrated in this paper.

MATERIALS AND METHODS

Bark of *Picea abies* (L.) Karst. with galleries of the bark beetles *Crypturgus pusillus* (Gyllenhal), *Ips typographus* L., and *Pityogenes chalcographus* (L.) was

^{*} This paper is dedicated to Professor Tomasz Majewski on the occasion of his 70th birthday.

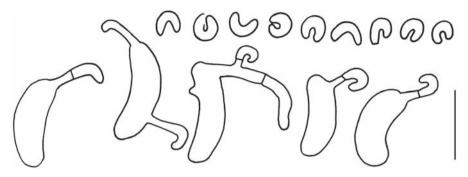


Fig. 1. Basidiospores of *Exidia glandulosa* (Bull.) Fr. germinating with allantoid conidia, an example considered typical for the majority of Auriculariales (*leg. R. Kirschner 308*). Scale bar = $10 \mu m$.

collected in Germany in 1994–2000. For details see the section on specimens examined; for general collection strategies see Kirschner (1998, 2001). Bark beetle galleries were immediately examined with a dissecting microscope. In order to obtain cultures, conidia were transferred from conidiophores in the galleries to a natural medium (1.5% water agar containing autoclaved pieces of twigs of *Picea abies*) with a flamed needle. Parts of a basidioma of *Exidia saccharina* (Alb. & Schwein.) Fr. (*leg. R. Kirschner 88* and *333*) were fixed under the lids of Petri dishes and allowed to disperse basidiospores onto 1.5% water agar containing autoclaved pieces of

twigs of *Picea abies* or 2% malt extract agar. Polyspore cultures were deposited under accession numbers F 3274 (ex-teleomorph), F 3275 and F 3468 (both ex-anamorph) in the culture collection of the Botanical Institute of the University of Tübingen, Germany. Dried specimens were deposited in the Herbarium Senckenbergianum in Frankfurt am Main, Germany (FR). Transmission electron microscopy (TEM) was used to study material from a gallery of an unidentified bark beetle in *Picea abies*, Germany, Bad Waldsee, 7 Apr. 1995, *leg. R. Kirschner 11*, using methods described in Kirschner *et al.* (2001a). For microscopic examination, specimens were mounted



Fig. 2. Synnemata of the anamorph of *Exidia saccharina* (Alb. & Schwein.) Fr. in a gallery of the bark beetle *Pityogenes chalcographus* in *Picea abies* (Germany, Bad Waldsee, 12 Sept. 1994, *leg. R. Kirschner*). Scale bar = 1 mm.

in water or 5–10% aqueous KOH, with and without 1% aqueous phloxine staining.

RESULTS

Anamorph of Exidia saccharina (Alb.

& Schwein.) Fr.

Figs 2–4

The anamorphic fungus described and illustrated below from material found in bark beetle galleries was indistinguishable from the anamorph derived from the polysporous cultures of the teleomorph of *E. saccharina* (*leg. R. Kirschner 88*, Fig. 5) on natural medium. For synonyms of the teleomorph see Neuhoff (1935) and Wojewoda

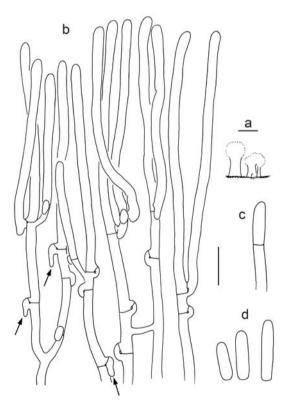


Fig. 3. Anamorph of *Exidia saccharina* (Alb. & Schwein.) Fr. isolated from conidiophores from a gallery of *Ips typographus* in *Picea abies*, Germany, Bad Waldsee, 19 Oct. 1995, *leg. R. Kirschner 10*, on autoclaved spruce twig in water agar. a – habit sketches of synnemata, b – conidiophores. Clamps with retrorse elongations (arrows), c – apex of conidiogenous cell forming a conidium, d – conidia. Scale bars: a = 100 μ m, b–d = 10 μ m.

(1981). Septal ultrastructure revealed dolipores with continuous parenthesomes.

Conidiophores on autoclaved spruce twigs and in bark beetle galleries predominantly synnematous, but also mononematous, erect, hyaline, sparsely and irregularly branched and septate, $120-150 \times 2-3$ µm. Clamps at the septa often with a retrorse projection which in some cases is delimited by a septum. One to three conidiogenous cells arising terminally from branches of the conidiophore, cylindrical, $50-63 \times 2-3 \mu m$. Mode of conidiogenesis unclear when viewed by LM, but visible by TEM as annellidic, because longitudinal sections through conidiogenous cells show lateral wall remnants of previous conidium dehiscence. Conidia accumulating in a slimy mass on the apex of the conidiogenous cells, hyaline, one-celled, smooth, cylindrical with truncate base and rounded apex, cytoplasm strongly granulate, $10-16 \times 3 \ \mu m$.

SPECIMENS EXAMINED

Exidia saccharina anamorph: In galleries of Pitvogenes chalcographus in bark of Picea abies: GER-MANY. Baden-Württemberg, near Bad Waldsee, ca 600 m a.s.l., 12 Sept. 1994, leg. R. Kirschner (Fig. 2), not deposited; in abandoned galleries of an unidentified bark beetle in bark of Picea abies: GERMANY. Baden-Württemberg, near Bad Waldsee, ca 600 m a.s.l., 7 Apr. 1995, leg. R. Kirschner 11 (Fig. 4), FR; in galleries of Ips typographus in bark of Picea abies: GERMANY. Baden-Württemberg, near Bad Waldsee, ca 600 m a.s.l., 19 Oct. 1995, leg. R. Kirschner 10 (Fig. 3), FR, culture F 3275 in Tübingen; in abandoned bark beetle galleries secondarily colonized by Crypturgus pusillus in bark of felled stem of Picea abies, GERMANY. Bavaria, Oberjoch, Iseler, ca 1200 m a.s.l., 26 Sept. 2000, leg. R. Kirschner 756, FR, culture F 3468 in Tübingen.

Exidia saccharina teleomorph: On felled stems of *Pinus sylvestris*: GERMANY. Hessen, Darmstadt-Eberstadt, Malcher Schneise, *ca* 100 m a.s.l., 9 Nov. 1996, *leg. R. Kirschner* 88, FR, LSU rDNA sequence available under GenBank accession number AF291323 (Weiß & Oberwinkler

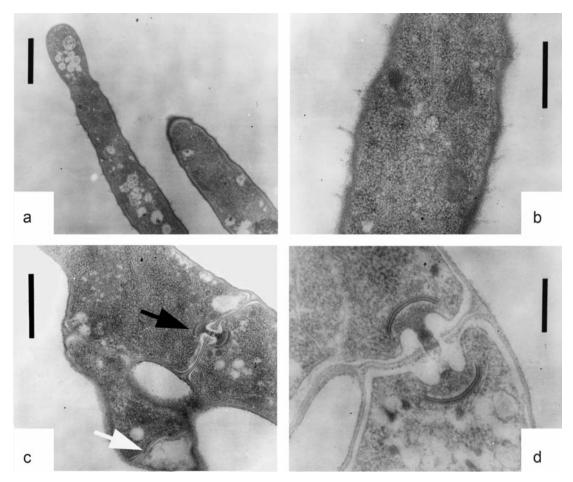


Fig. 4. Ultrastructure of the anamorph of *Exidia saccharina* (Alb. & Schwein.) Fr. from a gallery of an unidentified bark beetle in *Picea abies*, Germany, Bad Waldsee, 19 Oct. 1995, *leg. R. Kirschner 11*. a – longitudinal section through the apex of a conidiogenous cell with a developing conidium, b – longitudinal section through subapical part of a conidiogenous cell showing lateral wall remnants of previous conidium dehiscence indicating annellidic conidiogenesis, c – longitudinal section through a clamp connection showing the dolipore with parenthesomes in the regular septum (black arrow) and the retraction septum delimiting the retrorse, plasma-free elongation of the clamp (white arrow), d – median section through a dolipore with continuous parenthesomes. Scale bars: $a = 1.5 \mu m$, $b = 0.5 \mu m$, $c = 0.75 \mu m$, $d = 0.25 \mu m$.

2001); S of Darmstadt-Eberstadt, *ca* 100 m a.s.l., 11 Apr. 1998, *leg. R. Kirschner 330, 331, 333* (on three stems), FR, culture F 3274 derived from *R. Kirschner 333* deposited in Tübingen.

DISCUSSION

Brefeld (1888) described the conidia of *E. sac-charina* in culture as cylindrical, measuring 10–15 \times 3–5 µm, and accumulating in heads. Comparing these conidia to the curved conidia in most other

Auriculariales, he considered this characteristic important enough to propose the genus *Ulocolla* for the holomorph. This concept was not followed by later researchers (e.g., Neuhoff 1935). The germination of basidiospores and production of cylindrical or slightly curved conidia in culture were also reported by Neuhoff (1935) and Wojewoda (1981), but with smaller sizes reported for the conidia ($5-7 \times 1-2 \mu m$). Brefeld (1888) used water and liquid media for his observation of conidium development. Neuhoff (1935) probably observed

conidia developing from hyphae of germinating basidiospores on teleomorph specimens from nature. It is unclear whether the conidial dimensions differ because of different environmental or genetic conditions. Neuhoff (1935) also found different sizes of basidiospores among different collections of the same species. The different sizes of basidiospores and conidia might indicate some genetically fixed variability to be tested by mating experiments and DNA analyses. The occurrence of synnematous conidiophores with clamped stipe hyphae in nature or in culture was not observed by previous authors. This finding resulted only from field collection and cultivation on natural media (Kirschner 1998).

Anamorphs of other species of Auriculariales could be distinguished from the anamorph of E. saccharina based on the literature. Straight or nearly straight cylindrical conidia have been demonstrated clearly for only a few other Exidia species, namely E. badioumbrina (Bres.) Neuhoff, E. cartilaginea S. Lundell & Neuhoff, and E. recisa (Ditmar) Fr. (Neuhoff 1935; Pilát 1957; Ingold 1995). However, these species are restricted to angiosperms and the conidia are smaller, up to 6 µm long (Pilát 1957; Ingold 1995). As far as known, the conidia of the majority of Auriculariales s.str. are allantoid: Auricularia Bull. ex Juss. spp. (Brefeld 1888; Ingold 1982a), Ductifera sucina (Möller) K. Wells (Möller 1895), E. glandulosa (Bull.) Fr. (Brefeld 1888; Ingold 1982b), E. pithya Fr. (Neuhoff 1935; Pilát 1957; Wojewoda 1981), E. thuretiana (Lév.) Fr. [Neuhoff 1935, as E. albida (Huds. ex Louds.) Bref.; Ingold 1995], E. truncata Fr. (Neuhoff 1936), E. umbrinella Bres. (Pilát 1957), Exidiopsis cerina Möller (Möller 1895), Exidiopsis ciliata Möller (Möller 1895), Exidiopsis effusa (Bref.) Möller (Brefeld 1888; Ingold 1992), Exidiopsis glabra Möller (Möller 1895), Heterochaete hirneoloides (Pat.) K. Wells (Möller 1895, as Exidiopsis tremellispora Möller), H. shearii (Burt) Burt (Kisimova-Horovitz et al. 1997), H. verruculosa (Möller) Bodman (Möller 1895, as Exidiopsis verruculosa Möller), Myxarium grilletii (Boud.) D. A. Reid [Neuhoff 1935, as Exidia grilletii (Boud.) Neuhoff], M. laccatum (Bourdot & Galzin) D. A. Reid (Kisimova-Horovitz et al. 2000) and M. nucleatum Wallr. (Brefeld 1888, as Exidia corrugativa Bref. - synonymy according to Neuhoff 1935; Ingold 1984). In some of these cases, conidiogenous cells were described as phialide-like. As demonstrated here for the anamorph of E. saccharina, conidiogenesis is annellidic with percurrent proliferations not visible by LM but only by electron microscopy, in which longitudinal sections through conidiogenous cells show lateral wall remnants of previous conidium dehiscence. Phialidic development with apical replacement wall-building indicated by apical wall-thickening in the conidiogenous cell (Minter et al. 1983) has been confirmed in Basidiomycota only for spermatia in spermogonia of rust fungi (Mims et al. 1976). The clear distinction between phialidic and annellidic conidiogenesis, difficult to detect with LM but obvious with TEM, supported segregation of the genus Leucogloea R. Kirschner with annellidic conidium development, and basidiomycetous affinities from Pleurocolla Petr. with phialidic conidiogenesis (Kirschner 2004). Within ascomycetous anamorphs, detailed studies of conidiogenesis based on electron microscopy have led to consider-

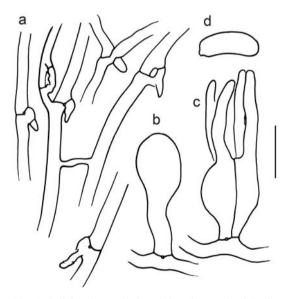


Fig. 5. Cellular characteristics of the teleomorph of *Exidia* saccharina (Alb. & Schwein.) Fr. (*leg. R. Kirschner 88*). a – hyphae with clamps with retrorse elongation ('spurred clamps'), b – young basidium, c – mature basidium, d – basidiospore. Scale bar = 10 μ m.

able taxonomic changes on the generic level in the former Leptographium complex (Wingfield et al. 1987). Conidiogenous cells of additional species of Auriculariales should therefore be investigated by electron microscopy. Helicomyxa everhartioides R. Kirschner & Chee J. Chen was described for a sporodochial anamorph with a close phylogenetic relationship to Hyaloriaceae and conidia developing from clamps (Kirschner & Chen 2004). Compared to the anamorph of Myxarium tremelloides (L. S. Olive) Wojewoda, for which separate conidiomata were also recorded from nature (Olive 1952), H. everhartioides might be closely related or identical, but the details of conidiophores and conidiogenesis of M. tremelloides are unknown (Olive 1952). Conidium development from clamps as in H. everhartioides appears to be unique within the Auriculariales.

The synnematous anamorph of E. saccharina is morphologically similar to species of Gloeosynnema Seifert & G. Okada, hitherto comprising the synnematous type species G. ochroleucum (Penz. & Sacc.) Seifert & G. Okada and the sporodochial species G. roseum Matsush. (Seifert & Okada 1988; Matsushima 1995). These species share the characters of hyaline or colored conidiomata, annellidic conidiogenesis, and clamps at the septa. The sporodochial G. roseum appears identical to the anamorph of Typhula micans (Pers.) Berthier (Agaricales) as described by Koske and Perrin (1971) and Berthier (1976). Clamps of Gloeosynnema species lack the retrorse projection typical for E. saccharina. Clamps with a posterior appendage, in some cases delimited by a retraction septum ('spurred clamp'), are a phylogenetically significant characteristic of the Auriculariales (Bandoni & Wells 1992). The spurred clamp is also illustrated for the teleomorph of E. saccharina (Fig. 5a). Other basidiomycetous synnematous anamorphs with annellidic conidiogenesis without clamps have been recorded for Basidiopycnis Oberw., R. Kirschner, R. Bauer, Begerow & Arenal (Atractiellales) and Stilbotulasnella Oberw. & Bandoni (Cantharellales) (Bandoni & Oberwinkler 1982; Oberwinkler et al. 2006). These anamorphs were not named separately by the original authors because they could be connected to the named teleomorph beyond any doubt. This practice is followed here for the distinct anamorph of *E. saccharina*. Though it would be nomenclaturally correct to create a new taxon, I consider it unnecessary for several reasons given in Kirschner and Oberwinkler (2009).

Exidia saccharina is known as a teleomorph from conifers with a preference for pine (Neuhoff 1935). Though bark beetle galleries in pine were also examined, the anamorph was found in this study only in spruce (Kirschner 1998). On these comparatively rare occasions, the anamorph appeared abundantly in the galleries but was never directly isolated from bark beetles. No teleomorph was found at the places where the anamorph of E. saccharina was collected. Exidia saccharina appears considerably less common than E. pithya, which can be found frequently on the bark of spruce after infestation by bark beetles (pers. observation). It may be that different environmental conditions are required for the development of both morphs of E. saccharina. As mentioned by Neuhoff (1935), basidiomata of this species in Europe appear very frequently in certain regions but are conspicuously rare in others. Although this paper adds a further species to the known basidiomycetous associates of bark beetles (e.g., Kirschner 2001; Kirschner et al. 2001a, b; Kirschner & Chen 2003; Kolařík et al. 2006; Oberwinkler et al. 2006), the complex ecological relationships, life cycles and systematic diversity of these fungi are not well understood, and more insights are to be expected from future work.

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REFERENCES

- BANDONI R. J. & OBERWINKLER F. 1982. Stilbotulasnella: a new genus in the Tulasnellaceae. Canad. J. Bot. 60: 1875–1879.
- BANDONI R. J. & WELLS K. 1992. Clamp connections and classification of the Auriculariales and Tremellales. *Trans. Mycol. Soc. Japan* 33: 13–19.
- BERTHIER J. 1976. Monographie des Typhula Fr., Pistillaria Fr. et genres voisins. Bull. Mens. Soc. Linn. Lyon 45: 1–213.
- BREFELD O. 1888. Basidiomyceten. II. Protobasidiomyceten. Unters. Gesammtgeb. Mykol. 7: 1–178.
- INGOLD T. C. 1982a. Basidiospore germination and conidium development in *Auricularia. Trans. Brit. Mycol. Soc.* 78: 161–166.
- INGOLD T. C. 1982b. Basidiospore germination and conidium formation in *Exidia glandulosa* and *Tremella mesenterica*. *Trans. Brit. Mycol. Soc.* **79**: 370–373.
- INGOLD T. C. 1984. Myxarium nucleatum and its conidial state. Trans. Brit. Mycol. Soc. 82: 358–360.
- INGOLD T. C. 1991. The view of the active basidium in Heterobasidiomycetes. *Mycol. Res.* 95: 618–621.
- INGOLD T. C. 1992. The conidial state in *Exidiopsis effusa* and *Exidiopsis longispora*. Mycol. Res. 96: 932–934.
- INGOLD T. C. 1995. Types of reproductive cell in *Exidia* recisa and Sirobasidium intermediae. Mycol. Res. 99: 1187–1190.
- KIRSCHNER R. 1998. Diversität mit Borkenkäfern assoziierter filamentöser Mikropilze. Ph.D. Dissertation, Faculty for Biology, University of Tübingen, Germany.
- KIRSCHNER R. 2001. Diversity of filamentous fungi in bark beetle galleries in central Europe. In: J. K. MISRA & B. W. HORN (eds), *Trichomycetes and other fungal groups: Professor Robert W. Lichtwardt commemoration volume*, pp. 175–196. Science Publishers Inc., Enfield, New Hampshire, USA.
- KIRSCHNER R. 2004. Sporodochial anamorphs of species of *Helicogloea*. In: R. AGERER, M. PIEPENBRING & P. BLANZ (eds), *Frontiers in basidiomycote mycology*, pp. 165–178. IHW, Eching, Germany.
- KIRSCHNER R. & CHEN C.-J. 2003. A new record of *Rog-ersiomyces okefenokeensis* (Basidiomycota) from beetle galleries in pines in Taiwan. *Sydowia* 55: 86–92.
- KIRSCHNER R. & CHEN C.-J. 2004. Helicomyxa everhartioides,

a new helicosporous sporodochial hyphomycete from Taiwan with relationships to the Hyaloriaceae (Auriculariales, Basidiomycota). *Stud. Mycol.* **50**: 337–342.

- KIRSCHNER R. & OBERWINKLER F. 2009. Supplementary notes on *Basidiopycnis hyalina* (Basidiomycota, Atractiellales) and its anamorph. *Mycotaxon* 109: 29–38.
- KIRSCHNER R., BAUER R. & OBERWINKLER F. 2001a. Colacosiphon: a new genus described for a mycoparasitic fungus. Mycologia 93: 634–644.
- KIRSCHNER R., BEGEROW D. & OBERWINKLER F. 2001b. A new Chionosphaera species associated with conifer inhabiting bark beetles. *Mycol. Res.* 105: 1403–1408.
- KIRSCHNER R., YANG Z.-L., ZHAO Q. & FENG B. 2010. Ovipoculum album, a new anamorph with gelatinous cupulate bulbilliferous conidiomata from China and with affinities to the Auriculariales (Basidiomycota). Fungal Diversity (in press, published online: DOI 10.1007/s13225–010– 0038–0).
- KISIMOVA-HOROVITZ L., OBERWINKLER F. & GÓMEZ P. L. D. 1997. Basidiomicetos resupinados de Costa Rica. Exidiaceae (Tremellales). *Rev. Biol. Trop.* 45: 1325–1347.
- KISIMOVA-HOROVITZ L., OBERWINKLER F. & GÓMEZ P. L. D. 2000. Basidiomicetos resupinados de Costa Rica. Myxariaceae s. Jülich, Sebacinaceae Wells & Oberw., y Tremellodendropsidaceae Jülich. *Revista Biol. Trop.* 48: 519–538.
- KOLAŘÍK M., SLÁVIKOVÁ E. & PAŽOUTOVÁ S. 2006. The taxonomic and ecological characterization of the clinically important heterobasidiomycete *Fugomyces cyanescens* and its association with bark beetles. *Czech Mycology* 58: 81–98.
- KOSKE R. E. JR. & PERRIN P. W. 1971. Basidiocarps, annelloconidia, and sclerotia in agar culture of *Pistillaria* (Clavariadelphaceae). *Canad. J. Bot.* **49**: 695–697.
- MATSUSHIMA T. 1995. *Matsushima Mycological Memoirs* No. 8. Published by the author, Kobe, Japan.
- MIMS C. W., SEABURY F. & THURSTON E. L. 1976. An ultrastructural study of spermatium formation in the rust fungus *Gymnosporium juniperi-virginianae*. Amer. J. Bot. 63: 997–1002.
- MINTER D. W., KIRK P. M. & SUTTON B. C. 1983. Thallic phialides. *Trans. Brit. Mycol. Soc.* 80: 39–66.
- MÖLLER A. 1895. Protobasidiomyceten. Botanische Mittheilungen aus den Tropen 8: 1–179 & 6 plates.
- NEUHOFF W. 1935. Die Pilze Mitteleuropas. Band IIa. Die Gallertpilze. 1. & 2. Lieferung: 1–28. Verlag von Dr. Werner Klinkhardt, Leipzig.
- NEUHOFF W. 1936. Die Pilze Mitteleuropas. Band IIa. Die Gallertpilze. 3. & 4. Lieferung: 29–48. Verlag von Dr. Werner Klinkhardt, Leipzig.
- OBERWINKLER F., KIRSCHNER R., ARENAL F., VILLARREAL M., RUBIO V., BEGEROW D. & BAUER R. 2006. Two new pyc-

nidial members of the Atractiellales: *Basidiopycnis hyalina* and *Proceropycnis pinicola*. *Mycologia* **98**: 627–649.

- OLIVE L. S. 1952. Taxonomic notes on Louisiana fungi. III. Additions to the Tremellales. *Mycologia* **43**(1951): 677–690.
- PILÁT A. 1957. Übersicht der europäischen Auriculariales und Tremellales unter besonderer Berücksichtigung der tschechoslowakischen Arten. Sborn. Nar. Muz. v Praze, Rada B, Prir. Vedy 13: 115–210.
- SEIFERT K. A. & OKADA G. 1988. *Gloeosynnema*, a new genus of synnematous hyphomycetes with basidiomycetous affinities. *Mycotaxon* 32: 471–476.
- WEISS M. & OBERWINKLER F. 2001. Phylogenetic relationships in Auriculariales and related groups – hypotheses derived from nuclear ribosomal DNA sequences. *Mycol. Res.* **105**: 403–415.
- WINGFIELD M. J., VAN WYK P. S. & WINGFIELD B. 1987. Reclassification of *Phialocephala* based on conidial development. *Trans. Brit. Mycol. Soc.* 89: 509–520.
- WOJEWODA W. 1981. Basidiomycetes (Podstawczaki), Tremellales (Trzęsakowe), Auriculariales (Uszakowe), Septobasidiales (Czerwcogrzybowe). In: S. DOMAŃSKI (ed.), *Mala flora grzybów.* 2: 1–408. Państwowe Wydawnictwo Naukowe, Warszawa – Kraków.

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