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Published in:
P L o S One

DOI:
10.1371/journal.pone.0073419

Publication date:
2013

Document Version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
Wynns, A. A., Eilenberg, J., & Jensen, A. B. (2013). Ascosphaera callicarpa, a new species of bee-loving fungus, with a key to the Genus for Europe. DOI: 10.1371/journal.pone.0073419
**Ascospaera callicarpa**, a New Species of Bee-Loving Fungus, with a Key to the Genus for Europe

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**Abstract**

We studied the bee specialist fungus *Ascospaera* in wild solitary bees to investigate the diversity of the genus in nature and the ecology of these fungi with their bee hosts. A new morphologically distinctive species was discovered which also has a unique nrITS sequence. This new species, here named *Ascospaera callicarpa*, is common on the larval feces of the solitary bee *Chelostoma florosomne* which nests in the *Phragmites* reeds of thatched roofs in Europe. Because collections of *Ascospaera* from wild bees are scarce and because little is known about the ecology and distribution of the majority of the species in the genus, a key to the species thus far reported for Europe is included.

**Introduction**

*Ascospaera* is a genus of 28 species of bee specialist fungi with a worldwide distribution in the temperate to tropical regions. The genus is remarkable for its host and habitat specificity with all species completing their entire life cycle within the nests of bees (Apoidea: Anthophila). *Ascospaera* was first discovered in the early 20th century in Europe after *A. apis*, the type species, was identified as the causative agent of a brood disease affecting honeybees [1,2]. This brood disease, known as chalkbrook, was later observed in a solitary bee in London [3]. *Ascospaera* is widely known as the chalkbrood fungus, although at least half of the species lead a saprotrophic rather than pathogenic lifestyle [4,5]. Saprotrophic *Ascospaera* species flourish on diverse substrates within the bee nest, for example on pollen provisions, on materials used by the bees to construct the nest and on larval feces [6,7]. Little is known about these saprotrophs which appear to live innocuously inside the brood cells of the bees. Consequently, the potential for research on the ecological and functional role of these fungi within the bee nest remains wide open.

*Ascospaera* is placed in Ascospaeraceae (Pezizomycotina: Eurotiomycetidae), a small family of ascomycetes primarily characterized by a unique fruiting body type called a spore cyst. Spore cysts are unicellular, cyst-like fruiting bodies that form from the expansion of a single cell called a nutriocyte [8]. The wall of a spore cyst is a double-layered membrane. Asci are free-floating and evanescent. Because of their anomalous fruiting bodies, the taxonomic affinities of *Ascospaera* and its relatives remained uncertain until ontological studies led C.F. Spiltoir and L.S. Olive [8] to confidently place them among the Ascomycota within Plectascales. This position was later confirmed by additional morphological study [9] and DNA sequenced-based phylogenies [10,11].

A distinguishing feature of *Ascospaera* is the presence of spore balls [8]. A spore ball is a compact aggregation of spores formed by groups of asci that are united by a single membrane [12]. The membrane surrounding a spore ball disintegrates and only remnants of it are sometimes observed in mature spore cysts [5]. Spore balls may contain as few as two to as many as several hundred ascospores [3,13]. The average number of ascospores per spore ball and the persistence of spore balls at maturity are meaningful taxonomic characters.

Pathogenic *Ascospaera* species afflict only the larval stage of bees. Typically diseased larvae die in the larval stage; however, in rare occurrences, larvae have been observed to enter pupation before being overcome by the fungus (Wynns pers. obs.). Pathogenic species of *Ascospaera* appear to be highly specialized fungi with ascospores typically germinating only when within the midgut of their host. Spore germination is followed by rapid hyphal growth, with the fungus consuming the larva from the inside out [14]. Two widespread pathogenic species, *Ascospaera aggregata* and *A. apis*, are of economic interest because of their potential to negatively impact populations of commercial pollinators, namely *Apis mellifera* L. and *Megachile rotundata* Fabricius [15,16].

Although *Ascospaera* lives in association with both solitary and social bees the majority of species (25 out of 28) were originally described from solitary bees. Within the nests of solitary bees *Ascospaera* grows on pollen provisions where an egg has failed to develop, on larval feces, on the surface of cocoons, within larvae, and on the diverse materials used by different bee species for brood cell construction [4,5,6]. Unlike their social relatives (e.g., honey bees), solitary bees lack adult-larva interaction, there is no nursing of the brood and no cooperative behavior (including social immunity) [17]. A consequence of no adult-larva contact and no nursing is that the brood is mass provisioned rather than progressively provisioned like their social counterparts; this means that once an egg hatches the larva has all the food it will need to...
complete development into an adult [17]. Following their flight and nesting period solitary bees overwinter in their individual brood cells with no activity until emergence the following spring or early summer. In this way solitary bee nests provide a relatively stable, undisturbed micro-environment that appears suitable for the growth of these specialised fungi.

The only monographic work on *Ascosphaera* [5] focused on collections from an important commercial pollinator in Canada, the alfalfa leafcutting bee *Melissodes retundata*. While limited in scope, this monograph, which included the first key to the genus, remains the most useful and comprehensive reference for the identification of *Ascosphaera* species. Given the importance of wild pollinators and their increasing role in buffering the loss of honeybee pollination services [18] a more complete monograph with an updated key to these bee-specialist fungi is much needed.

Seven of the 28 described species of *Ascosphaera* are currently known from Europe. Here we describe a new species from Denmark occurring in the nests of the wild solitary bee *Collostoma florisonne* L. To stimulate interest and to facilitate the identification of *Ascosphaera* species so far known from Europe, we provide a key and descriptions for these species. Cumulative host reports and species distributions are also included with the hope that this information will result in additional records for these under-collected fungi.

**Materials and Methods**

**Morphological study**

Descriptions of spore cysts and ascospores were made from observations of spore cysts mounted in water on a glass slide. Measurements and light photomicrographs were made on an Olympus AX70 Provis light microscope and Olympus SZX16 dissecting microscope. Herbarium acronyms follow those of Index Herbariorum [19].

**Culture and isolation**

Attempts to isolate and culture the fungus were made by placing spore cysts and hyphae on three different solid agar media: malt agar with 20% dextrose (MY20), V8® agar with 2% yeast extract (V8YE), and malt extract agar (MEA). To induce spore germination spore suspensions were prepared from spore cysts placed in a modified V8 spore germination broth [20] and exposed to CO₂ as described in Wynns et al. [13].

**Molecular study**

Genomic DNA was obtained by plucking 5–10 spore cysts and grinding them inside a 1.5 ml Eppendorf tube. DNA was isolated using the Qiagen DNeasy Plant Mini Kit (Qiagen, Germany) using the standard protocol and eluted in two separate 50–100 μl fractions to avoid over-dilution.

We sequenced the entire nuclear ribosomal ITS region (ITS1-5.8S-ITS2) for *A. callicarpa* sp. nov. Genomic DNA was amplified using ITS1F and ITS4 primers [21]. PCR reactions were performed for a final 50 μl volume containing 29.8 μl of sterile deionized water, 5 μl of *Taq* polymerase reaction buffer (Sigma®), 1.0 μl 10 mM dNTPs, 3.0 μl 25 mM MgCl₂, 0.2 μl *Taq* DNA polymerase (Sigma®), 5.0 μl each 10 μM primer and 1 μl of genomic DNA template. PCR was performed on a Biometra® thermocycler (Whatman) under the following conditions: step 1) 1 min at 95°C, 2) 45 sec at 95°C, 3) 40 sec at 52°C, 4) 1 min 30 sec at 72°C, 5) return to step 2 30 times, 6) final step of 10 min at 72°C. Samples were kept at 4°C until electrophoresis was performed on a 1% agarose TAE gel and visualized with EZvision One® (Amresco). PCR reactions were cleaned using Qiaquick® PCR purification kit (Qiagen) and sent to Eurofins MWG Operon AG (Ebersberg, Germany) for sequencing. The nucleotide sequence was assembled using BioEdit [22] and subjected to a BLASTn search in GenBank.

**Nomenclature**

The electronic version of this article in Portable Document Format (PDF) in a work with an ISSN or ISBN will represent a published work according to the International Code of Nomenclature for algae, fungi, and plants, and hence the new names contained in the electronic publication of a *PLOS ONE* article are effectively published under that Code from the electronic edition alone, so there is no longer any need to provide printed copies. In addition, new names contained in this work have been submitted to MycoBank from where they will be made available to the Global Names Index. The unique MycoBank number can be resolved and the associated information viewed through any standard web browser by appending the MycoBank number contained in this publication to the prefix http://www.mycobank.org/MB/. The online version of this work is archived and available from the following digital repositories: PubMed Central, LOCKSS.

**Results and Discussion**

**Culture and isolation**

Despite repeated attempts, we were unable to obtain in vitro mycelial growth or induce ascospore germination of *Ascosphaera callicarpa*.

**Molecular study**

An ITS sequence was obtained for *A. callicarpa* (GenBank accession: JX070046). A BLASTn search of the ITS sequence revealed a highest sequence-similarity to other *Ascosphaera* species.

**Key to European species of Ascosphaera**

1. Wall of spore cyst smooth; ascospores cylindrical with rounded ends, (3.1–) 3.9×1.6(–2.0) μm; saprotroph.... *A. callicarpa* A.A. Wynns (Fig. 1)
   1) Wall of spore cyst with minute or conspicuous dark spots; ascospores not cylindrical; saprotroph or pathogen.................2
   2) 1. Wall of spore cyst smooth; ascospores cylindrical with rounded ends, (3.1–) 3.9×1.6(–2.0) μm; saprotroph.... *A. callicarpa* A.A. Wynns (Fig. 1)
   3) 2. Ascospores broadly sub-falcate, with a tendency to be trigoval when view on-end, 1.9–3.5×0.6–0.9 μm; saprotroph.... *A. tenax* Skou (Figs 2A–B)
   4) 2. Ascospores sub-falcate or trigoval in cross section 3
   5) 3. Ascospores always >2 μm wide, 4–7.9×2.3–6.5 μm, ellipsoid to broadly ellipsoid; spore balls not persistent; saprotroph.... *A. atrata* Skou & K. Hackett (Fig. 2C)
   6) 3. Ascospores otherwise; spore balls persistent; pathogen or saprotroph .................................................................4
   4) 4. Spore cysts not exceeding 125 μm diameter ...............5
   5) 4. Spore cysts mostly exceeding 125 μm diameter ..........6
   6) 5. Ascospores 3.0–5.0×1.3–1.8 μm; at least some ascospores and spore balls with attached granules; spore cyst wall brown with small spots visible at low magnification; saprotroph.... *A. fumicola* Skou (Fig. 3)
   7) 5. Ascospores 2.1–3.9×1.1–1.7 μm; ascospores and spore balls always without granules; spore cyst wall pale greenish to yellowish brown, with nearly smooth walls, minute spots visible at high magnification; obligate parasite, cause of chalkbrood disease of honeybees.... *A. apis* (Maasen ex Claussen) L.S. Olive & Spiltoir (Figs 2F–G)
   8) 6. Ascospores often ≥400 μm diameter, forming a dense layer beneath the cuticle of bee larvae with chalkbrood disease;
ascospores 3.4–5.9×1.3–2.6 μm, ellipsoid, sub-cylindrical or allantoid; obligate pathogen... A. aggregata Skou (Figs. 4A–B, E–F)

6. Spore cysts mostly less than and not exceeding 400 μm in diameter, developing on aerial hyphae above the cuticle of larvae with chalkbrood disease or growing saprotrophically on the cocoon, feces or leaf lining of a brood cell ......................... 7

7. Ascospores (2.4–) 2.8–4.0 (−5.0)×1.0–1.8 (−2.0) μm..... A. major (Prokschl & Zobl) Skou (Figs. 2D–E)

7. Ascospores 3.5–6.5×1.7–3.5 μm .......... A. proliperda Skou (Figs. 4C–D,G)

Taxonomy

Ascosphaera aggregata Skou, Friesia 11: 64, 1975.

Type: DENMARK.THRU, on larvae of Osmia rufa L., J.P. Skou s.n. (holotype, cf).

Fig. 1. Ascosphaera callicarpa. A) habitat. Phragmites reeds and female Chelostoma florisomne returning with pollen for her brood. B) fecal pellet of C. florisomne larva covered with spore cysts; pale spore balls are visible through the transparent spore cyst wall. C) close-up of spore cyst showing spore balls and smooth, unornamented spore cyst wall. D) spore balls. E) bacilliform ascospores. B, photographed from A.A. Wynns 5166; C–E from A.A. Wynns 5166. Scale bars: B = 200 μm, C = 50 μm, D = 15 μm, E = 10 μm.

doi:10.1371/journal.pone.0073419.g001

Ascosphaera callicarpa sp. nov.

Description. Mating system unknown but possibly homothallic [4]. Pathogenic. Infected larvae swollen, black, and filled with a solid core of pale buff mycelium. Ascomata black to dark brown spore cysts produced below surface of larval cuticle in a crowded continuous layer [23] or scattered and appearing as small individual boils [24], 280–750×130–290 μm, spherical or conical and faceted from being tightly packed beneath the larval cuticle; wall light reddish brown to black, minutely punctate. Spore balls pale brown to yellowish brown with small brown granules attached to surface, 9–25 μm diameter, mostly persistent. Ascospores ellipsoid to sub-cylindrical or allantoid, 3.4–5.9×1.3–2.6 μm. Culture on V8YE with moderate growth after 14 days, low, pale buff with a darker, brownish centrum with age, occasionally producing nutriocytes on aerial hyphae.
Figure 2. Light microphotographs of *Ascosphaera tenax*, *A. atra*, *A. major*, and *A. apis*. *Ascosphaera tenax* A) ascospores. B) punctate spore cyst wall.; *A. atra* C) broken spore cyst with ascospores.; *A. major* D) ascospores with attached granules. E) close-up of spore cyst wall.; *A. apis* F) ascospores. G) detail of pale spore cyst wall with minute spots. A–B photographed from holotype; C from ARSEF 693; D from A.A. Wynns 5170; E from A.A. Wynns 5175; F from A.A. Wynns 5174. Scale bars A = 5 μm, B–C = 10 μm, D = 5 μm, E–F = 10 μm.
doi:10.1371/journal.pone.0073419.g002

Figure 3. Light microphotographs of *Ascosphaera fimicola*. A) two opaque, iridescent spore cysts still attached to hyphae. B–C) close-up of spore cyst showing maculate wall, D) ellipsoid ascospores with a few small granules attached to their surface. E) spore ball. A, photographed from A.A. Wynns 5167; B–C, E from A.A. Wynns 5130; D from J.P. Skou s.n. (paratype). Scale bars: A = 500 μm, B = 20 μm, C = 10 μm, D = 5, μm E = 15 μm.
doi:10.1371/journal.pone.0073419.g003
Ascosphaera callicarpa sp. nov.

(A.A.Wynns pers. obs.) and immature spore cysts below agar surface [4].

Ecology and distribution. *Ascosphaera aggregata* is an obligate pathogen with a preference for bees belonging to the family Megachilidae. This species has a broad distribution, with reports from both North America and Europe. Like *A. apis*, the distribution of *A. aggregata* is probably closely tied to the exchange and transport of bees (e.g. *M. rotundata*) for the pollination of commercial crops. In Europe *A. aggregata* is known from Denmark [23], Germany [present study], Spain [23,25] and Sweden [present study]. Attempts to isolate and grow *A. aggregata* in culture often results in the co-isolation of another pathogenic species, *A. proliperda*. Although *A. proliperda* and *A. aggregata* can be difficult to separate based on microscopic morphological features their growth in culture is strikingly different (Figs. 4B,D; see also *A. proliperda* for further discussion on its co-occurrence with *A. aggregata*).

Additional specimens examined. CANADA. ALBERTA: Brooks, on larvae of *M. rotundata*, 1988, J Jakobsen s.n. (C). DENMARK. THUR: Svendborg Kommune, on Osmia rufa (= O. bicornis L.), 1974, J.P. Skou s.n., paratype (C); On O. rufa, 1974, J.P. Skou s.n., holotype, (C); ZEALAND: Frederikssund Kommune, Slangerup, organic apple orchard, on larva of O. rufa, 2010, A.A. Wynns 5152; Roskilde Kommune, Roskilde, on larva of O. rufa, 2010, A.A. Wynns 5144; Taastrup Kommune, Taastrup, Højbakkegaard, on larvae of O. rufa, 2010, A.A. Wynns 5143 (C). USA. NEVADA: On Megachile pacifica (Paxzer) (= M. rotundata), 1975, K. Hackett s.n. paratype (C). SPAIN: On *M. rotundata*, 1972, J.P. Skow s.n., paratype (C); on *M. rotundata*, 1973, J.P. Skow s.n. paratype (C).

*Ascosphaera apis* (Maasen ex Clausen) L.S. Olive & Spiltoir in Spiltoir & Olive, Mycologia 47: 242, 1955.

=Persiciis aps= Maasen ex Clausen, Mitt. biol. BundAnst. Ld-u. Forstw.10: 470. 1921.

**Figs. 2F–G**
Description. Mating system heterothallic. Pathogenic. Infected larvae shrunken, pale buff, covered by a weft of hyphae, with or without the production of ascocarps. Ascocarps greenish (immature) to black (mature) spore cysts produced on aerial hyphae above the larval cuticle, 40–119 µm in diameter; wall pale greenish to yellowish-brown, nearly smooth with minute punctae at high magnification. Spore balls hyaline to pale yellowish, without granules, 7–20 µm in diameter, mostly persistent. Ascospores ellipsoid to sub-allantoid, 2.1–3.9 x 1.1–1.7 µm. Culture on SDA with rapid growth after 2–6 days, white with abundant production of spore cysts when both mating strains are present.

See Skou [7], Bissett [5], and Aroustine & Murray [26] for additional descriptions.

Ecology and distribution. Ascosphaera atra is an opportunistic pathogen of honeybees. Experimental trials showed A. apis is able to induce chalkbrood in the solitary bee M. rotundata [14]; however, A. apis is not known to live in association with solitary bees in nature. Reports of chalkbrood caused by A. apis in solitary bees before 1972 are most likely attributable to pathogenic species described after this time; e.g., A. major, A. aggregata or A. prolifera (the later two species are pathogenic specific to solitary bees). Originally described from Germany, A. apis is now known from all continents where honeybees are kept.

Additional specimens examined. USA. TEXAS: Weslaco, 26 Jun 2003, K.D. Murray s.n., ARSEF 7405 (+), 7406 (−).

Ascosphaera callicarpa Skou & K. Hackett, Friesia 11: 279, 1979.

Type: U.S.A. NEVADA, isolated from larva of Megachile pacifica with ragged-brood disease, 36836 (c), CBS 524.75 (holotype, c!).

Fig. 2C

Description. Mating system homothallic. Ascomata black, globose spore cysts, 30–140 µm; wall dark brown, punctate, punctae appearing as uniform dark circles often of variable size. Spore balls hyaline to pale yellowish brown, 8–17 µm diameter, evanescent. Ascospores ellipsoid to broadly ellipsoid, 4.7–9 x 2.3–6.5 µm, with or without small granules attached to the surface of the spore wall. Culture on SDA with moderate growth after 7 days, white to greyish-buff with abundant production of black spore cysts on aerial hyphae and on hyphae growing beneath the surface of the agar.

Ecology and distribution. Ascosphaera atra is a fast-growing saprotroph associated primarily with solitary bees. This species is typically found growing on pollen provisions. Less common substrates from which A. atra has been isolated include the surface of a diseased M. rotundata larva with chalkbrood caused by A. aggregata [27], from pollen within the gut of an otherwise healthy M. rotundata larva [7] and from the honey of A. mellifera [4]. Ascosphaera atra is the only species of the genus that has been found growing on plant material (grass silage) outside of the bee habitat [6]. Pathogenicity studies [14,27] demonstrated that A. atra is not a pathogen of solitary bees; however, Vojvodic et al. [28] concluded that it is a weak pathogen of honeybees. More work is needed to determine if A. atra is comparable to some of its bee- pathogen congeners e.g. A. aggregata and A. apis. The perceived pathogenicity of this species in honey bee larvae may be more closely tied to its rapid growth on suitable substrates. Ascosphaera atra is the most extensively studied saprotrophic species of Ascosphaera. This is reflected in the multiple reports from N. America [4,14], Europe [16], present study], New Zealand and Australia [4].

Additional specimens examined. Ascosphaera atra. AUSTRALIA. Peel: Waroona, A. mellifera honey, Nov 1994, D.L. Anderson 198, ARSEF 5147. CANADA. ALBERTA: Beaverlodge, Peace River region, from pollen in M. rotundata cells, Jan 1983, D. Farney s.n., DAOI 188981. USA. OREGON: Ontario. M. rotundata, Jun 1979, J.D. Vandenbog 6, ARSEF 693.

Ascosphaera callicarpa A.A. Wynns, sp. nov. [urn:lsid:indexfungorum.orgnames:518624]

Type. DENMARK. ZEALAND: Leje Kommune, Sagulanget (“Land of Legends”) Leje, Landbohusene, on fecal pellets of Chelostoma flaviscutentes nesting in the Phragmites reeds of thatched roof of shed behind 19th century cottage, 55°37’11”N; 11°22’13”, 2010, A.A.Wynns 5163 (holotype, c!).

Fig. 1

Description. Mating system unknown. Ascomata pale brown, semi-transparent and somewhat iridescent spore cysts (Fig. 1B), globose to subglobose 64–101 µm in diameter; wall smooth (Fig. 1C). Spore balls 10–16 µm in diameter, center grayish-brown to colorless, ascospores arranged spirally or not (Fig. 1C, D). Ascospores bacilliform, (3.1–)4.0 x 1.6–2.0 µm, colorless or slightly brownish (Fig. 1D–E); no attached granules. Mycelium sparse, white. No growth in culture on MY20, V8YE or MEA; no spore germination in V8 spore germination broth, either with or without the addition of carbon dioxide.

Ecology and distribution. Common in the nest reeds of the solitary bee C. flaviscutentes where it grows on the feral pellets of this bee. Although not definitely known, the distribution of A. callicarpa is probably closely tied to that of C. flaviscutentes. This fungus was not found in association with other bees, e.g. Osmia and Megachile, although these bees were observed nesting in the same Phragmites reeds as C. flaviscutentes. Ascosphaera callicarpa appears to be solely saprotrophic; it was not found in association with diseased bees or where a larva had failed to develop. Ascosphaera callicarpa is so far known only from the island of Zealand, Denmark.

Etymology. The epithet callicarpa means with beautiful fruits, here referring to the spore cysts.

Preliminary conservation status. Ascosphaera callicarpa should be sought in other aggregations of C. flaviscutentes in thatched roofs throughout Europe in order to assess its conservation status. As a possible obligate associate of the bee C. flaviscutentes, the conservation of this fungus should be considered dependent on the conservation of its host.

Additional specimens examined. DENMARK. ZEALAND: Leje Kommune, Sagulanget Leje, Landbohusene, shed behind 19th century houses. All specimens on feral pellets of C. flaviscutentes nesting in the Phragmites reeds of the thatched roof, 2008, A.A.Wynns 5011, 5012, 5013, 5014, 5015, 5018, 5023, 5026, 5027, 5072, 5074, 5136, all specimens in c; 2011, A.A.Wynns 5166, 5168 (c); Soro Kommune, Kristiansminde, University of Copenhagen field station, east facing side of classroom building, growing on the feral pellets of C. flaviscutentes nesting in Phragmites reeds of the thatched roof, 2012, A.A. Wynns 5169, 5170 (c).

Morphological comparison of A. callicarpa with A. fimicola. Ascosphaera callicarpa most closely resembles A. fimicola Skou which also grows on the feral pellets of bees. This new species is distinguishable from A. fimicola by a pale brown, highly transparent fragile spore cyst (Fig. 1B–C) with a wall (Fig. 1C) that is not sculptured or maculate as in A. fimicola (Fig. 3B–C). The spore cysts of A. fimicola (Fig. 3A) are dark brown to pale brown, also somewhat iridescent, and it transparent, not as strikingly so as in A. callicarpa (Fig. 1B). The spores of A. fimicola are ellipsoid-fusiform (Fig. 3D) and often have small brown granules attached to their surface while the spores of A. callicarpa are bacilliform (Fig. 1E) without surface granules. Ascosphaera callicarpa grows on digested Ranunculus pollen voided by C. flaviscutentes. It is not clear if A. callicarpa grows on pollen collected from other plants since C. flaviscutentes is strictly oligolectic on Ranunculus species [29].

Ascosphaera fimicola Skou, Friesia 11: 68, 1975.
**Type:** DENMARK, THURO, on fecal pellets from larvae of *Osmia rufa*, J.F. Skou s. n., (holotype, c1).

**Fig. 3.**

**Description.** Mating system not known. Ascomata light to dark brown, somewhat iridescent, glistening spore cysts, (23–) 64–125 μm in diameter; wall brown, punctate, punctate minute and of uniform size. Spore balls yellowish, with small granules on the surface, (3–)10–15–20 μm in diameter, mostly persistent. Ascosporae ellipsoid to sub-allantoid, 3.0–5.0×1.3–1.9 μm with or without small granules attached to the spore wall. Mycelium on natural substrate noticeable, stringy, white and opaque. No growth in culture.

**Ecology and distribution.** *Ascosphaera fumiola* grows saprotrophically on the larval feces and cocoons of the solitary bee *Osmia bicornis* (syn. *O. rufa*) and was recently collected on the larval feces of *Caseton adiagatum* (Diptera: Drosophilidae) a cleptoparasite of this bee. Despite extensive collecting, *A. fumiola* was not found on the larval feces of the solitary bee *C. florisomne*. The composition of the pollen provisions of these bees may play a role in the absence or presence of *A. fumiola* in their nests. *Catoasporae florisomne* feeds exclusively on pollen from the plant genus *Ranunculus* (*Ranunculaceae*) [29] while *O. bicornis* often collects pollen from the plant family *Rosaceae* [30]. The last report of *A. fumiola* prior to our study was in 1975 [29]. We found that this species is more common than the previous few collections indicate. The known distribution of *A. fumiola* is restricted to Denmark but, like other species in the genus, this narrow distribution is most likely an artifact of under-collecting because of a more focused interest in the pathogens rather than the saprotrophs.

**Additional specimens examined.** DENMARK. ZELAND: Frederiks bund Kommune, Slang erup, organic apple orchard belonging to Verner Andersen, growing on pollen and feces of *O. bicornis*, 2010, A.A. Wynns 5150 (c); Lejre Kommune, Lejre Forsogscenter, growing on cocoon and between walls of leaf lining of healthy *Megachile* sp., 2008, A.A. Wynns 5039 (c); Roskilde Kommune, Roskilde, Gorderupvej 3, on leaf-lining of brood cell belonging to *Megachile*, 2010, A.A. Wynns 5173, on larval feces of *Megachile* sp. without disease, 2010, A.A. Wynns 5175.

**Ascosphaera proliperda** Skou, Friesia 10: 15, 1972.

**Type:** DENMARK. ZELAND: Frederiksgaard Kommune, Frederiksgaard, in *Megachile centuncularis* larvae collected from the greenhouse of the Royal Veterinary and Agricultural University, J.P. Skou s.n., Jun 1967, CBS 687.71 (holotype, CBS H-6729, non vidi).

**Figs. 4C–D,G.**

**Description.** Mating system homothallic [5]. Pathogenic. Infected larva shrunken, covered by erect or low compact aerial hyphae bearing ascomata. Ascomata black spore cysts produced on tips of aerial hyphae above the larval cuticle, 60–250(–400) μm in diameter; wall dark-brown, appearing mottled from the confluence of very fine granules on the inner surface. Spore balls pale brown to sub-hyaline, 9–17(–25) μm, often with small brown granules on the surface. Asco spores sub-cylindrical or sub-allantoid, 3.5–6.5×1.7–3.5 μm, hyaline to sub-hyaline, with or without minute granules attached to the surface. Culture on MY20 with rapid growth after 7 days, white with abundant production of spore cysts.

**Ecology and distribution.** *Ascosphaera proliperda* causes chalkbrood in *Megachile centuncularis* [31,32] and *Megachile centuncularis* [7]. It is more often found growing saprotrophically on larval feces within the brood cells of *M. centuncularis* [7]. In the present study *A. major* was found growing on the larval feces and leaf material lining the brood cell of a species of *Megachile* and on the larval feces and pollen provisions of *O. bicornis*. The frequency of *A. major* as a cause of chalkbrood in honeybees is not known. Outwardly *A. apis* and *A. major* induce the same disease symptoms; therefore, the etiology of chalkbrood in honeybees should be carefully verified by morphological study of the fungus to distinguish infections by *A. apis* or *A. major* or to identify co-infection with both species. *Ascosphaera major* is known from N. America [33,34] and Europe. In Europe this species is reported from Switzerland [32], Austria [31] and Denmark [7,9].

**Additional specimens examined.** DENMARK. ZELAND: Frederiksbund Kommune, Slang erup, organic apple orchard belonging to Verner Andersen, growing on pollen and feces of *O. bicornis*, 2010, A.A. Wynns 5129 (c); on cocoons and feces of *O. rufa*, 2010, A.A. Wynns 5147, 5167 (c).

**Ascosphaera major** (Proksch & Zobl) Skou, Friesia 10:15, 1972.

**Type:** DENMARK. ZELAND: Glostrup, isolated from chalkbrood cells of *Megachile centuncularis*, CBS 696.71 (neotype, CBS H-9050, non vidi).

**=Perycystis apis** Maass ex Claussen var. major Proksch & Zobl in Proksch, Arch. Microbiol. 18: 200. 1953.

**=Ascosphaera apis** (Maassen ex Claussen) L.S. Olive & Spiltoir var. major (Proksch & Zobl) L.S. Olive & Spiltoir in Spiltoir & Olive, Mycologia 47: 243. 1955.

**Figs. 2D–E.**

**Description.** Mating system heterothallic. Ascomata dark brown to black, spore cysts, 60–150(–380) μm in diameter; wall greenish brown, with indistinct punctae or small granules attached to the inner surface, occasionally with larger crystalliferous brown precipitations with age. Spore balls hyaline to greyish-brown, (6–)14–10(–24) μm in diameter, usually with granules attached to the surface. Ascosporae suballantoid or bacilliform, (2.1–)2.8–4.0(–5.0)×1.0–1.8(–2.0) μm, at least some with small granules attached to the spore wall. Mycelium white to greyish-white. Culture on V8YE with moderate growth after 10 days, with abundant production of spore cysts when both strains are present.

**Ecology and distribution.** *Ascosphaera major* causes chalkbrood in *Apis mellifera* [31,32] and *Megachile centuncularis* [7]. It is more often found growing saprotrophically on larval feces within the brood cells of *M. centuncularis* [7]. In the present study *A. major* was found growing on the larval feces and leaf material lining the brood cell of a species of *Megachile* and on the larval feces and pollen provisions of *O. bicornis*. The frequency of *A. major* as a cause of chalkbrood in honeybees is not known. Outwardly *A. apis* and *A. major* induce the same disease symptoms; therefore, the etiology of chalkbrood in honeybees should be carefully verified by morphological study of the fungus to distinguish infections by *A. apis* or *A. major* or to identify co-infection with both species. *Ascosphaera major* is known from N. America [33,34] and Europe. In Europe this species is reported from Switzerland [32], Austria [31] and Denmark [7,9].

**Additional specimens examined.** DENMARK. ZELAND: Frederiksbund Kommune, Slang erup, organic apple orchard belonging to Verner Andersen, growing on pollen and feces of *O. bicornis*, 2010, A.A. Wynns 5150 (c); Lejre Kommune, Lejre Forsogscenter, growing on cocoon and between walls of leaf lining of healthy *Megachile* sp., 2008, A.A. Wynns 5039 (c); Roskilde Kommune, Roskilde, Gorderupvej 3, on leaf-lining of brood cell belonging to *Megachile*, 2010, A.A. Wynns 5173, on larval feces of *Megachile* sp. without disease, 2010, A.A. Wynns 5175.

**Ascosphaera proliperda** Skou, Friesia 10: 15, 1972.

**Type:** DENMARK. ZELAND: Frederiksbund Kommune, Frederiksbund, in *Megachile centuncularis* larvae collected from the green house of the Royal Veterinary and Agricultural University, J.P. Skou s.n., Jun 1967, CBS 687.71 (holotype, CBS H-6729, non vidi).

**=Ascosphaera proliperda** Skou, Zeesl. 687.71 (neotype, CBS H-6729, non vidi).

**=Ascosphaera proliperda** Skou, Zeesl. 687.71 (neotype, CBS H-6729, non vidi).

**=Ascosphaera proliperda** Skou, Zeesl. 687.71 (neotype, CBS H-6729, non vidi).

**=Ascosphaera proliperda** Skou, Zeesl. 687.71 (neotype, CBS H-6729, non vidi).

**=Ascosphaera proliperda** Skou, Zeesl. 687.71 (neotype, CBS H-6729, non vidi).
**Ascosphaera tenax** Skou & S.N. Holm, *Mycotaxon* 35: 212, 1989.

**Type**: DENMARK, NEKSELØ: Kalundborg Kommune, inside cocoons of *Megachile willughbiella*, 1985, J.P. Skou s.n., (holotype, C!).

**Description.** Mating system unknown. Ascomata lustrous black, less often dark brown, spore cysts, (33–)40–90(2105) μm diameter; wall dark brown, 1.5 μm thick, tough and leathery, smooth or minutely punctate. Spore balls hyaline, (7.7–)9–14(215.4) μm diameter. Ascospores sub-falcate, with a tendency to be trigonal when viewed on-end, 1.9–3.5×0.6–0.9 μm.

**Ecology and distribution.** *Ascosphaera tenax* grows saprotrophically on pollen provisions, larval feces and the inner side of cocoons of *Megachile willughbiella* and *M. rotundata*. Spore cysts are common beneath the inside of the leaf cap of *Megachile* cells. The last collections of *A. tenax* date from 1988, when the species was found growing in nearly half (18 out of 44) *M. willughbiella* cocoons examined [24]. *A. tenax* is known only from Denmark on the islands of Nekselø and Zealand. More focused collecting is needed to determine its real geographical range.

**Conclusion**

Our study is the first to provide a regional key to *Ascosphaera*. With the addition of *A. callicarpa* sp. nov., eight *Ascosphaera* species are now known from Europe (Table 1). Our collections of *A. fimicola* (see discussion under *A. fimicola*) from a dipteran cleptoparasite of *Osmia bicornis* add to the mounting evidence that, although undoubtedly a bee specialist, *Ascosphaera* is not restricted to bees; further evidence includes an isolated report of the saprotroph *A. atra* growing on grass [6] and molecular based identification of *Ascosphaera* DNA from *Eristalis* (Diptera: Syrphidae) and *Vespula* (Hymenoptera: Vespidae) species [37]. As previously suggested by Wynns [42], *Ascosphaera* should be sought outside the bee habitat in association with other pollenivorous insects and where high-sugar substrates are available. Reports of *Ascosphaera* in non-apoidean insects are quite possibly relevant for the control of chalkbrood in commercial bee pollinators since these insects may act as pathogen reservoirs or vectors of *Ascosphaera*. More frequent collections of *Ascosphaera* are needed to begin to grasp the diversity and ecology of these fungi in nature and to elucidate their potentially significant role within the bee habitat. Additional regional keys, such as the one provided here, may ease identification for the non-specialist and bring attention to the lesser-known species of both saprotrophs and pathogens.

**Acknowledgments**

We thank Holger Philipsen and Louise Lee Munk Larsen for field assistance, the Land of Legends (Sagnlandet Lejre) for kindly allowing us to collect bees from their thatched roofs and Justin Wynns.
Author Contributions
Conceived and designed the experiments: AAW JE AR. Performed the experiments: AAW. Analyzed the data: AAW. Wrote the paper: AAW.

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