

Amanita ochroterrea and *Amanita brunneiphylla* (Basidiomycota), one species or two?

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Abstract

Davison, E.M. *Amanita ochroterrea* and *Amanita brunneiphylla* (Basidiomycota), one species or two? *Nuytsia* 21(4): 177–184 (2011). *Amanita ochroterrea* Gentilli ex Bas and *A. brunneiphylla* O.K.Miller are robust, macroscopically similar mushrooms described from the south-west of Western Australia. According to the protologue of *A. brunneiphylla*, the main difference between them is the presence (in *A. ochroterrea*) or absence (in *A. brunneiphylla*) of clamp connections. However, in the current study abundant clamp connections have been observed in the holotype and paratypes of *A. brunneiphylla*. As other microscopic characters are indistinguishable, *A. brunneiphylla* is synonymised with *A. ochroterrea*, and an expanded description presented.

Introduction

Amanita species are large, conspicuous mushrooms with a worldwide distribution. They are readily recognized to the generic level in the field, but the majority are difficult to separate into species solely on their appearance. Microscopic characters are usually needed before collections can be confidently identified. The most commonly used microscopic characters are spore size and shape, their response to Melzer's iodine, the presence of clamp connections in the basidiome especially at the base of basidia, the structure of the subhymenium and underlying lamella trama, the pileipellis, and the structure of the universal veil on the pileus. All of these characters are used to separate species and construct diagnostic keys (Bas 1969; Reid 1979; Tulloss 1994; Wood 1997).

In 1953 Gentilli described several collections of *Amanita* spp. from Kings Park, Perth, Western Australia which included two large specimens with an earthy buff cap, buff lamellae and pale buff spore print that he called *A. preissii* (Fr.) Sacc. forma *ochroterrea* Gentilli (Gentilli 1953). This forma was not validly published since it was not accompanied by a Latin description or diagnosis. Gentilli sent these specimens to Bas who considered them to be so distinctive that he described them as a new species, *A. ochroterrea* Gentilli ex Bas (Bas 1969; MB308574).

Following a visit to the south-west of Western Australia in 1989, Miller described *A. brunneiphylla* O.K.Miller (MB358165), a robust species with a dull white cap, light brown lamellae and a pale yellow spore print. He recognised that this was similar to *A. ochroterrea*, but separated the two species on the presence (in *A. ochroterrea*) or absence (in *A. brunneiphylla*) of clamp connections, together with small differences in spore size and colour of the lamellae (Miller 1991).

This paper reports a re-examination of the holotype and paratypes of *A. brunneiphylla* held at the Western Australian Herbarium (PERTH). These are compared with the published description of *A. ochroterrea*. Additional collections named as *A. ochroterrea* in PERTH, and macroscopically similar specimens held in private collections in Western Australia have also been examined. As a result, *A. brunneiphylla* is synonymised with *A. ochroterrea* and an expanded description is provided.

Materials and methods

Methodology follows that of Tulloss (2000, 2008). Colours are from the Royal Botanic Garden, Edinburgh (1969) and colour codes in the form of 'p3A2' are from Korerup and Wanscher (1978). Dried material was rehydrated in 10% NH₄OH or 3% KOH and stained with 1% Congo Red. Biometric variables for spores follow Tulloss and Lindgren (2005), i.e. 'L = the average spore length computed for one specimen examined and the range of such averages, L' = the average spore length computed for all spores measured, W = the average spore width computed for one specimen examined and the range of such averages, W' = the average spore length computed for all spores measured, Q = the ratio of length/breadth for a single spores and the observed range of the ratio of length/breadth for all spores measured, Q = the average value of Q computed for one specimen examined and the range of such averages, Q' = average value of Q computed for all spores measured'.

Comparison of *Amanita ochroterrea* and *A. brunneiphylla*

The macroscopic descriptions of *A. ochroterrea* and *A. brunneiphylla* are similar; however, the protologue of *A. brunneiphylla* states that the flesh is 'firm, dull white tinted grey at the base' (Miller 1991). An image (E567) of a paratype (PERTH 007565259, *O.K. Miller* OKM 23747, as reproduced in Figure 1A, shows that the flesh is tinted brown, however, this may have resulted from drying. Neither Gentilli (1953) nor Bas (1969) comment on the colour of the context.

The microscopic characters of the holotype and paratypes of *A. brunneiphylla* do not differ from the type description of *A. ochroterrea* (Table 1). Basidiospore dimensions, amyloid reaction, size of basidia, and shape of lamella edge cells in all mature collections are similar to those of the type description of *A. ochroterrea*. Clamp connections are present and abundant in all of these collections; thus the original description of *A. brunneiphylla* is neither supported by the holotype nor the paratypes.

These observations did not detect any significant differences between these taxa. As a result, *A. brunneiphylla* is synonymised with *A. ochroterrea*. An expanded description of *A. ochroterrea* is provided.

Expanded description of *Amanita ochroterrea*

Amanita ochroterrea Gentilli ex Bas, *Persoonia* 5: 505–506, figures 278–281 (1969). *Amanita preissii* (Fr.) Sacc. f. *ochroterrea* Gentilli, *W. Austral. Naturalist* 4: 30, figure 3 (1953), (*nom. inval.*, Art. 36.1). *Type*: Perth, Kings Park, Western Australia, June 1953, *J. Gentilli s.n. (holo: L)*. (MB308574).

Amanita brunneiphylla O.K.Miller, *Canad. J. of Bot.* 69: 2694 (1991). *Type*: Murdoch University campus, Western Australia, 7 May 1989, O.K. & H.H. Miller; E.M. & P.J.N. Davison OKM 23621 (*holo*: PERTH 07587473, image E 511). (MB358165).

Basidiome small to very large (Figures 1A, B). *Pileus* 35–170 mm diam, up to 15 mm thick, hemispheric when young, becoming more or less plane with a depressed centre with age, cream, pale buff to pale olivaceous buff (p1A2–p2B4–p4B3), margin of the pileus appendiculate, non-sulcate, no surface staining or bruising reaction. *Universal veil on pileus* (Figure 1C) adnate, forming a soft thin crust over the whole pileus, sometimes with small floccose warts in the centre, sometimes with thick felted angular flattened warts in the centre, cream, pale grey olivaceous, pale vinaceous buff, pale buff to pale olivaceous buff (p1A2–p3B5–p4B3). *Lamellae* close, free to narrowly adnate, 5–15 mm broad, ventricose, buff, olivaceous buff, hazel (p1B2–p5C4–6), drying buff to snuff brown (p5C4–F7), often with two tiers of plentiful lamellulae, the shorter truncate, the longer attenuate, lamella margin lighter in colour and slightly fimbriate. *Stipe* length (bottom of pileus context to top of bulb) 60–115 mm; width at mid-stipe 14–37 mm, more or less equal, solid, pale olivaceous buff, pale buff (p2B4–p4B3), furfuraceous or covered in mealy scales below the partial veil. *Partial veil* superior, descendent, soft, fugacious, initially greenish cream darkening to pale olivaceous buff (p1A2–B2). *Bulb* 55–72 × 28–44 mm, initially ovoid, narrowing with age, olivaceous buff, encrusted with sand. *Remains of universal veil* at the base of the stipe soft ridges and scales, in some specimens forming girdles; loose patches often remaining in the soil. *Flesh* cream, straw, to pale olivaceous buff (p1A2–p2B5) in both pileus and stipe, sometimes darkening on exposure to air. *Smell* mild and earthy when young, stronger when older. *Spore deposit* cream (p3A2–3) to buff.



Figure 1. *Amanita ochroterrea*. A – showing the brownish colour of freshly exposed flesh (paratype of PERTH 07565259, O.K. Miller OKM 23747); B – *Amanita ochroterrea* (PERTH 08334897, B-S 168); C – surface of pileus (PERTH 08059632, E.M. & P.J.N. Davison EMD 6-2008). Photographs by N.L. Bougher (A), L.I. Little (B) and E.M. Davison (C)

Table 1. Comparison of the type descriptions of *Amanita ochroterrea* and *A. brunneiphyllo* with observations from holotype and paratype collections of *A. brunneiphyllo*.

	<i>A. ochroterrea</i> type description (Bas 1969)	<i>A. brunneiphyllo</i> type description (Miller 1991)	<i>A. brunneiphyllo</i> holotype	<i>A. brunneiphyllo</i> PERTH 07587473 paratype	<i>A. brunneiphyllo</i> PERTH 07564465 paratype	<i>A. brunneiphyllo</i> PERTH 07565259 paratype
Basidiospores L × W (µm)	(10-)11-13 (-13.5) × 5-6.5 (20/1)	(8-)9-10.8 × 4.1-5	(8-)9-12 × 4.5-5 (20/1) from lamella	no spores, basidia immature	no spores, basidia immature	(8.5-)9.5-11(-11.5) × 4.5-6 (15/1) from lamella
Q	1.9-2.4, mean 2.1	1.8-2.3, mean 2.1	2.0-2.7, mean 2.2			1.7-2.3, mean 2.0
Amyloidy	amyloid	amyloid	amyloid			amyloid
Basidia L × W (µm)	50-55 × 11-12	34-38 × 7-10	38-45 × 9-10	basidia immature	basidia immature	30-50 × 9-11
Lamella edge cells (size in µm)	globose to clavate, 15-35 × 15-20	pyriform to clavate, 18-26 × 9-15	pyriform to clavate, up to 25 × 12	clavate to spherical, 15-40 × 10-15	clavate to pyriform, 20-28 × 14-20	globose, clavate to cylindric, <20 wide
Subhymenium	probably ramose	isodiametric cells	inflated ramose	ramose	ramose	ramose
Clamp connections	present at base of basidia and in pileipellis	none seen in any tissue	present and abundant in all tissues	present and abundant in all tissues	present and abundant in all tissues	Present and abundant in all tissues

Basidiospores (Figure 2A) [161/9/8] (8–) 9.5–13(–15) × (4–)4.5–6 (–6.5) μm, **L** = 10.1–11.7 μm; **L'** = 11.0 μm; **W** = 4.7–5.6 μm; **W'** = 5.2 μm; **Q** (1.67–) 1.82–2.44(–3.00), **Q** = 2.00–2.23; **Q'** 2.13) hyaline, colourless, thin-walled, smooth, amyloid, elongate to cylindrical, infrequently bacilliform, adaxially flattened, with apiculus sublateral, truncate, about 1 × 1 μm, with granular contents. *Pileipellis* difficult to delimit, merging into both universal veil and pileus context, not or slightly gelatinized at the centre, gelatinization of the hyphal walls in some specimens near the pileus margin; hyphae 2–10 μm wide, thin walled, hyaline, orientation mainly radial with some interweaving. *Pileus context* tissue yellow in NH₄OH, hyphae 6–30 μm wide, thin walled, hyaline, dominant, mainly with radial orientation; inflated cells up to 60 × 250 μm, thin walled, colourless. *Lamella trama* bilateral; width of central stratum 40–60 μm, hyphae 5–20 μm wide, no inflated cells seen; subhymenial base 25–60 μm wide, hyphae 4–20 μm wide, dominant orientation is initially about 30° from the vertical with the hyphae bending round in a smooth curve to the subhymenium, inflated cells up to 20 × 70 μm, infrequent; subhymenium (Figure 2B) 20–40 μm wide, ramose to inflated ramose, thin-walled; lamella margin cells (Figure 2C) globose, pyriform to clavate up to 10–15 × 25–40 μm. *Basidia* [132/7/7] (30–)35–60(–65) × (7.5–)9–11(–12.5) μm, thin-walled, yellow contents in NH₄OH, about 90 % four-spored about 10 % two-spored, sterigmata up to 5 μm long by 2 μm wide at the base; basal clamps present. *Universal veil on the pileus* (Figure 2E) comprising abundant mainly ellipsoidal to spherical, rarely pyriform to clavate, venticose to fusiform cells up to 70 × 150 μm, but most smaller, inflated cells terminal or in short chains; filamentous hyphae 5–10 μm wide, frequently branched, irregularly disposed, hyphae more abundant in the proximal part of the universal veil, while the inflated cells are more abundant in the distal part of the universal veil; some gelatinization of the walls of both filamentous and inflated hyphae evident in older specimens. *Universal veil on the stipe base* disordered tissue of terminal spherical and clavate cells up to 30 × 150 μm and filamentous hyphae 3–12 μm wide. *Stipe context* acrophysalidic, with acrophysalides up to 30 × 300 μm dominant, filamentous hyphae 3–13 μm wide with mainly axial orientation. *Partial veil* (Figure 2D) of dominant spherical to clavate or pyriform inflated cells 15–20 × 50 μm, but occasionally up to 30 × 300 μm, and infrequent, mainly radial, filamentous hyphae 2–8 μm wide. *Vascular hyphae* present but infrequent in all tissues, 2–15 μm wide, occasionally branched, thin walled, with yellow glassy contents in NH₄OH, frequently serpentine, no knots or concentrations of vascular hyphae noted in any tissue. *Clamp connections* present and frequent in all tissues.

Collections examined. WESTERN AUSTRALIA: Murdoch University campus, 25 Apr. 1990, E.M. & P.J.N. Davison EMD 12-1990 (PERTH 08059640); Melville, 12 May 2008, E.M. & P.J.N. Davison EMD 6-2008 (PERTH 08059632); near Southern Cross, 23 June 1974, B. Dell & K. Elson s.n. (PERTH 00768804, UWA 1862); Swan View, 20 May 2005, K. Griffiths B-S 168 (PERTH 08334897); Murdoch University campus, 7 May 1989, O.K. & H.H. Miller, E.M. & P.J.N. Davison, OKM 23621 (PERTH 07587473, E 511 holotype of *A. brunneiphylla*); Kings Park, 21 May 1989, O.K. & H.H. Miller OKM 23660 (PERTH 07587562, E 529 paratype of *A. brunneiphylla*); Moore River, 23 May 1989, O.K. & H.H. Miller OKM 23720 (PERTH 07564465, E 560 paratype of *A. brunneiphylla*); Regans Ford, 30 May 1989, O.K. & H.H. Miller OKM 23747 (PERTH 07565259, E 567 paratype of *A. brunneiphylla*); Kings Park, 26 May 1982, anon. s.n. (PERTH 07607512, E 239).

Rejected collections. Corrigin shire, J. Catchpole, I.C. Tommerup & N.L. Bougher s.n., 8 July 1999 (PERTH 07658508, E 6235); Grass Patch, T.C. Daniell s.n., 18 Aug. 1984 (PERTH 00918326, UWA 2966) (green form); Gleneagle, R.N. Hilton s.n., 18 June 1975 (PERTH 00771961, UWA 2001); Walpole Nornalup National Park, K. Syme 29:87, 24 May 1987 (PERTH 07575270, UWA 3510).

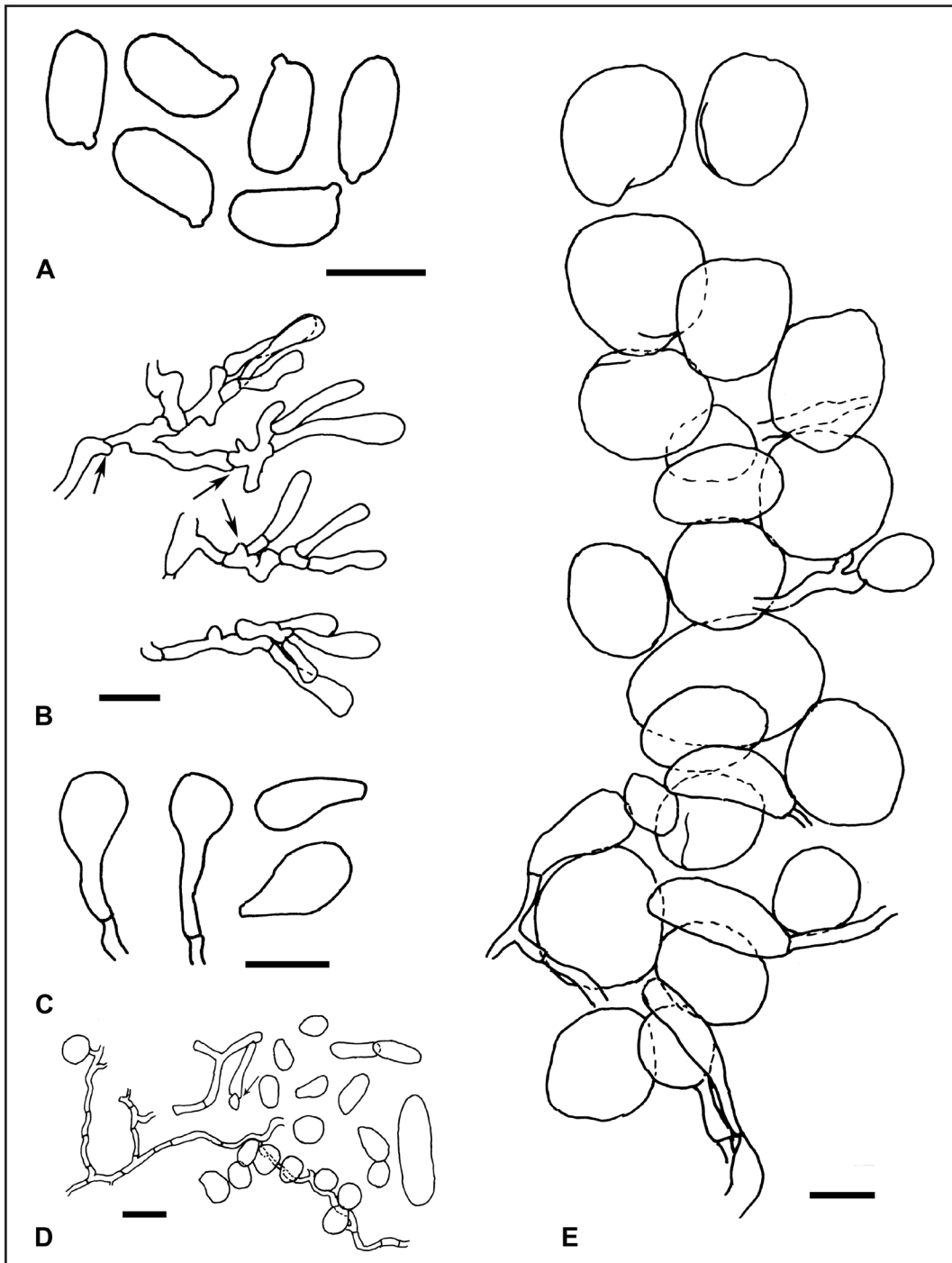


Figure 2. *Amanita ochroterrea*. A – spores from spore print; B – squash of young basidia and subhymenium, clamp connections indicated by arrows; C – lamella edge cells; D – cells from the partial veil, clamp connection indicated by an arrow; E – vertical section through the universal veil on the pileus, the proximal part is at the bottom. A, C, D, E (PERTH 08059632, *E.M.* & *P.J.N. Davison* EMD 6-2008); B (PERTH 07587562 *O.K. Miller* OKM 23660, paratype of *A. brunneiphylla*). Scale bars A = 10 μ m, B–E = 20 μ m.

These collections have been rejected for the following reasons: PERTH 07658508 has a saccate volva and probably resides in Section *Amidella*; PERTH 00771961 and PERTH 07575270 have been rejected because they have low *Q*: 1.64 and 1.72 respectively; PERTH 00918326 has a much wider central stratum and wider lamellae.

Distribution and habitat. Solitary or gregarious, in sandy soil in dry sclerophyll woodland and sand plain, often associated with *Eucalyptus marginata* Sm. and *Corymbia calophylla* (Lindl.) K.D.Hill & L.A.S.Johnson. *Amanita ochroterrea* is a distinctive species that is widely distributed in the south-west of Western Australia from the Moore River (31°00' S, 115°30' E) to Southern Cross (31°13' S, 119°18' E) although it does not appear to be common. It has not been recorded in South Australia (Grgurinovic 1997) or eastern Australia (Wood 1997).

Fruiting period. April to August.

Diagnostic features. Robust basidiomes with buff pileus, brown gills, yellowish spore print and clamp connections throughout.

Suggested common name. Brown-gilled amanita.

Notes. Bas (1969) placed *A. ochroterrea* in *Amanita* (subsection *Solitariae* Bas) *Stirps Grossa* in part because of the irregularly disposed remnants of the universal veil on the pileus. He commented, however, that these remnants were difficult to analyse. The collections examined here support Bas' placement of *A. ochroterrea* in *Stirps Grossa* because the basidia have clamp connections, *Q* is less than 2.2, the universal veil on the pileus forms a subfelted layer and is composed of irregularly disposed inflated cells intermixed with hyphae.

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