

Microfungi associated with *Podocarpus* leaf litter in South Africa

P.W. Crous*, KA. Seifert¹ and R.F. Castañeda Ruiz²

*Department of Plant Pathology, University of Stellenbosch, Stellenbosch, 7600 Republic of South Africa

¹Centre for Land and Biological Resources Research, Research Branch, Agriculture and Agri-Food Canada, Ottawa, Ontario, K1A 0C6, Canada

²Instituto de Investigaciones Fundamentales en Agricultura Tropical 'Alejandro de Humboldt' Ministerio de la Agricultura, Calle 2 Esq. 1 Stgo. de las Vegas, Ciudad de la Habana, Cuba

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Nine microfungi are listed from leaf litter of *Podocarpus* spp. New species include *Chaetopsina mellitolunae* Crous & Seifert, *Rhinotrichella elegans* R.F. Castañeda & Crous, *Parasymptodiella podocarpi* Crous & Seifert, *Guignardia podocarpi* Crous and its probable anamorph *Phyllosticta podocarpi* Crous. A key is provided to distinguish the accepted species of *Parasymptodiella*. *Gyothrix verticiclada* (Goid.) S. Hughes & Piroz., which was found to be morphologically variable, is discussed in detail. New records for South Africa include *Camposporium antennatum* Harkn., *Dactylaria irregularis* de Hoog, *Endophragmiella boewei* (J.L. Crane) S. Hughes, and *Phaeoisaria clematidis* (Fuckel) S. Hughes.

Keywords: Follicolous fungi, *Chaetopsina mellitolunae*, *Guignardia podocarpi*, *Parasymptodiella podocarpi*, *Phyllosticta podocarpi*, *Rhinotrichella elegans*, *Podocarpus*, taxonomy, new species.

*To whom correspondence should be addressed.

Introduction

The foliicolous microfungi occurring on woody hosts in South Africa have been poorly studied. However, progress has been made on fungi occurring on genera such as *Eucalyptus* L'Hérit. and *Syzygium* Gaertn. in the Myrtaceae (Crous 1993; Crous & van der Linde 1993; Crous *et al.* 1995). On *Podocarpus* L'Hérit. ex Pers. species, which are endemic to South Africa, only a few foliar fungi have been recorded (Doidge 1950).

The aim of the present study was to collect leaf litter of the various *Podocarpus* spp. at Kirstenbosch and Stellenbosch botanical gardens, as well as from Knysna, where these trees occur in their natural environment. Three species were commonly encountered, namely *P. henkelii* Stapf ex Dallim. & Jacks., *P. elongatus* (Ait.) L'Hérit. ex Pers. and *P. latifolius* (Thunb.) R. Br. ex Mirb. Of these, leaf litter of *P. henkelii* proved to have hardly any microfungi, whereas litter of the other two were particularly rewarding. Several of the fungi isolated proved to be new records for South Africa, or apparently undescribed taxa. Four new species are described, and five listed as new records. Several fungi correlating closely with their original descriptions are merely listed, while other more variable taxa are discussed in detail.

Materials and Methods

Leaf litter samples were incubated in Petri dish moist chambers at 25°C on the laboratory bench, and examined at regular intervals for the presence of microfungi. Single-conidial isolates were made on 2% malt extract agar (MEA) (Oxoid), and plated onto fresh MEA and carnation-leaf agar (CLA) (Crous *et al.* 1992) plates, incubated at 25°C under near-ultraviolet light, and examined. Cardinal temperature requirements for growth were determined after 8 days at 5–35°C in 5° intervals, with three replicate plates per temperature. The experiment was repeated once. Mounts were prepared in lactophenol, and measurements were made at 1 000× magnification. Averages were derived from at least 30 observations, and the range is given in parentheses. Unless otherwise noted, all microscopic structures are hyaline with smooth, thin walls. Descriptions are based on living material from the natural substrata unless otherwise noted.

Taxonomy

Chaetopsina mellitolunae Crous & Seifert sp. nov. (Figure 1)

Mycelium consists in hyphis ramosis septates *in vitro*, 1.5–2.5 µm diam. Conidiophora setiformia, recta, parietibus crassis (basi usque ad 2 µm), apicem versus verrucosa, 8–16-septata, 180–360 µm longa, 3–4 µm lata ad septum subapicale, 6–10 µm lata ad septum basale, lutea, pallidiora vel hyalina supra regionem conidiogerae, 2–6 aggregata in stroma cellularum brunnearum; apicibus saepe fertilibus *in vitro*. Cellulae conidiogerae monophialidicae, rare polyphialidicae *in vitro*, ampulliformes, 8–13 × 3–4 µm, collis inconspicuis divergentibus vel cylindricis usque ad 2 µm longis; saepe in hyphis *in vitro* dispositae. Conidia non-septate, cylindrica, recta, in apice obtusa vel acuta, basim obtusa hilo parvo complanato, (11–)14(–17) × (1.5–)2(–3) µm, in massis mucosis aggregata.

Mycelium consisting of branched, septate, hyphae *in vitro*, 1.5–2.5 µm diam. Conidiophores setiform, straight, thick-walled (up to 2 µm at base), becoming verrucose at apex, 8–16-septate, 180–360 µm long, 3–4 µm wide at subapical septum, 6–10 µm wide at basal septum, yellow, becoming pale yellow to hyaline above conidiogenous region; most conidiophores with equivalent, appressed lateral branches near the mid-point, which sometimes branch again, giving rise to phialidic conidiogenous cells directly, or to branches that are cylindrical, ellipsoidal or globose and 4–6 µm long; conidiophores aggregated in groups of 2–6 on a stroma of brown cells on CLA; conidiophore apices frequently becoming fertile *in vitro*. Conidiogenous cells monophialidic, rarely polyphialidic *in vitro*, ampulliform; 8–13 × 3–4 µm, collarettes inconspicuous, divergent or cylindrical, up to 2 µm long; occurring singly, in pairs or whorls on branches, sometimes singly at the apex of the setae, or directly on mycelial hyphae *in vitro*. Conidia non-septate, cylindrical, straight, with apex obtuse to slightly acute, base obtuse with a small flattened hilum, (11–)14(–17) × (1.5–)2(–3) µm on CLA, aggregated in slimy masses.

Colonies on MEA attain a radius of 8–9 mm after 8 days at 20°C in the dark. Cultures are shiny with sparse aerial mycelium; initially white, turning orange. Cardinal temperatures for growth are minimum: above 5°C, optimum: 20°C, and maximum: below 35°C.

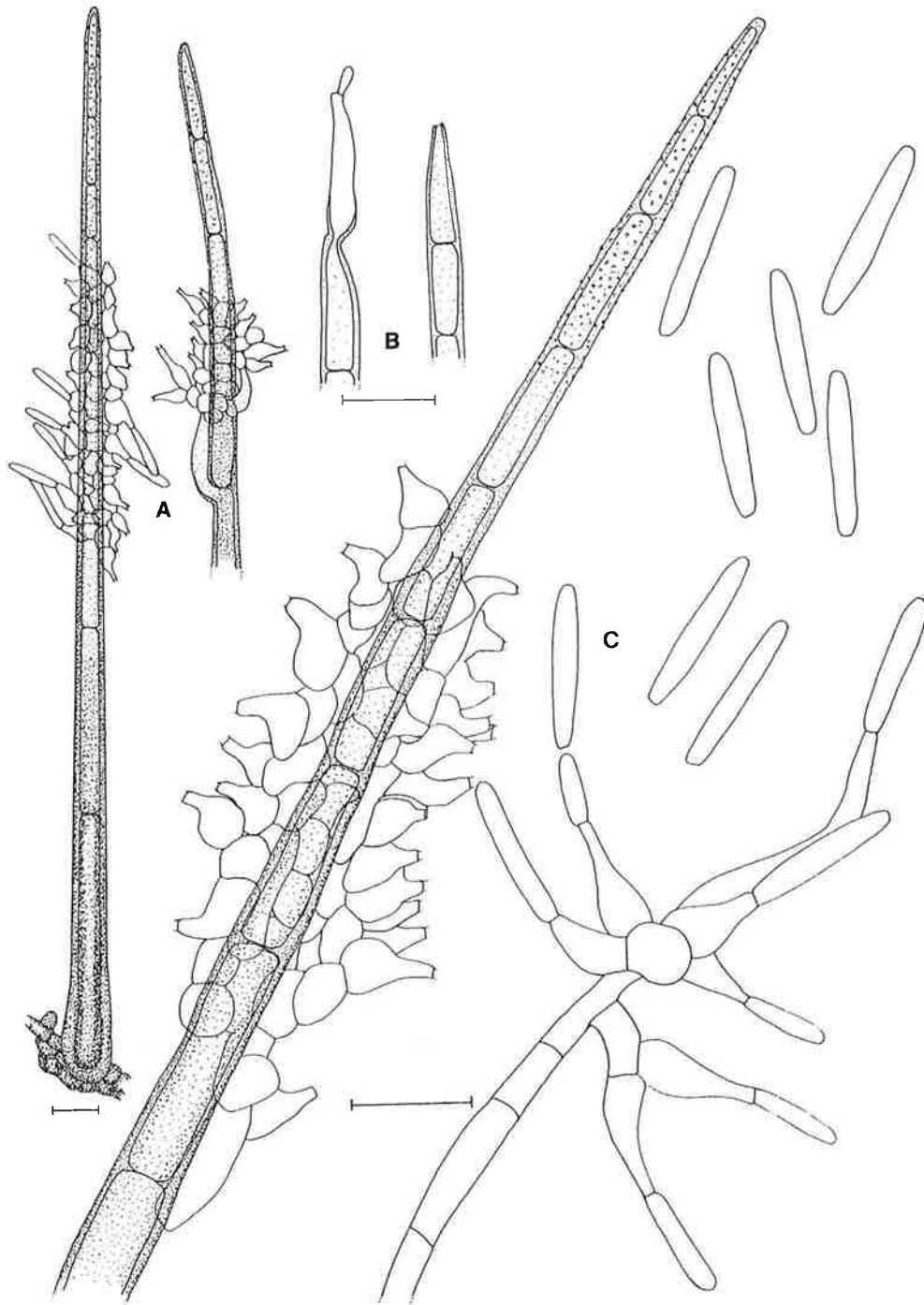


Figure 1 *Chaetopsina mellitolunae* (PREM 51901). A. Conidiophores *in vivo*. B. Setal apices becoming fertile *in vitro*. C. Conidia and conidiogenous cells on a conidiophore and hypha *in vitro* (bar = 10 µm).

Specimen examined: Southern Cape, Knysna, 'big tree', *Podocarpus elongatus* leaf litter, P.W. Crous & J.M. Crous, 9 Jan. 1995, PREM 51901 (holotype), DAOM 221070, culture STE-U 891.

Chaetopsina mellitolunae is morphologically similar to *C. nimbae* Ant. Rambelli, which was described from *Lophira alata* Banks ex Gaertn. collected in south-western Africa (Merli *et al.* 1992). The present collection closely matched the original description in its yellow conidiophores and conidial dimensions. An examination of the type specimen (ROBB 138 A) proved *C. nimbae* to be distinct from our collection. Conidiophores of *C. mellitolunae* are generally straight, not curved, and the conidiogenous cells are not restricted to only one side of the conidio-

phore. Furthermore, the conidiophores are lighter in colour and taper to more bluntly rounded apices, whereas those of *C. nimbae* are yellow-brown (*in vivo*), and have more acute apices. Phialides and conidiophore branches are also slightly larger, and cultures are generally lighter in colour on potato-dextrose agar than reported for *C. nimbae*. Conidiogenous cells are either mono- or polyphialidic, and in culture on CLA some setal conidiophores have fertile setal apices, as reported for *C. fulva* Ant. Rambelli by Merli *et al.* (1992).

***Guignardia podocarpi* Crous sp. nov. (Figure 2)**

Mycelium immersum, consists in hyphis septatis, ramosis, laevibus, mediobrunneis, 5–8 µm diam. Ascocarpi sparsi, immerse, sub-

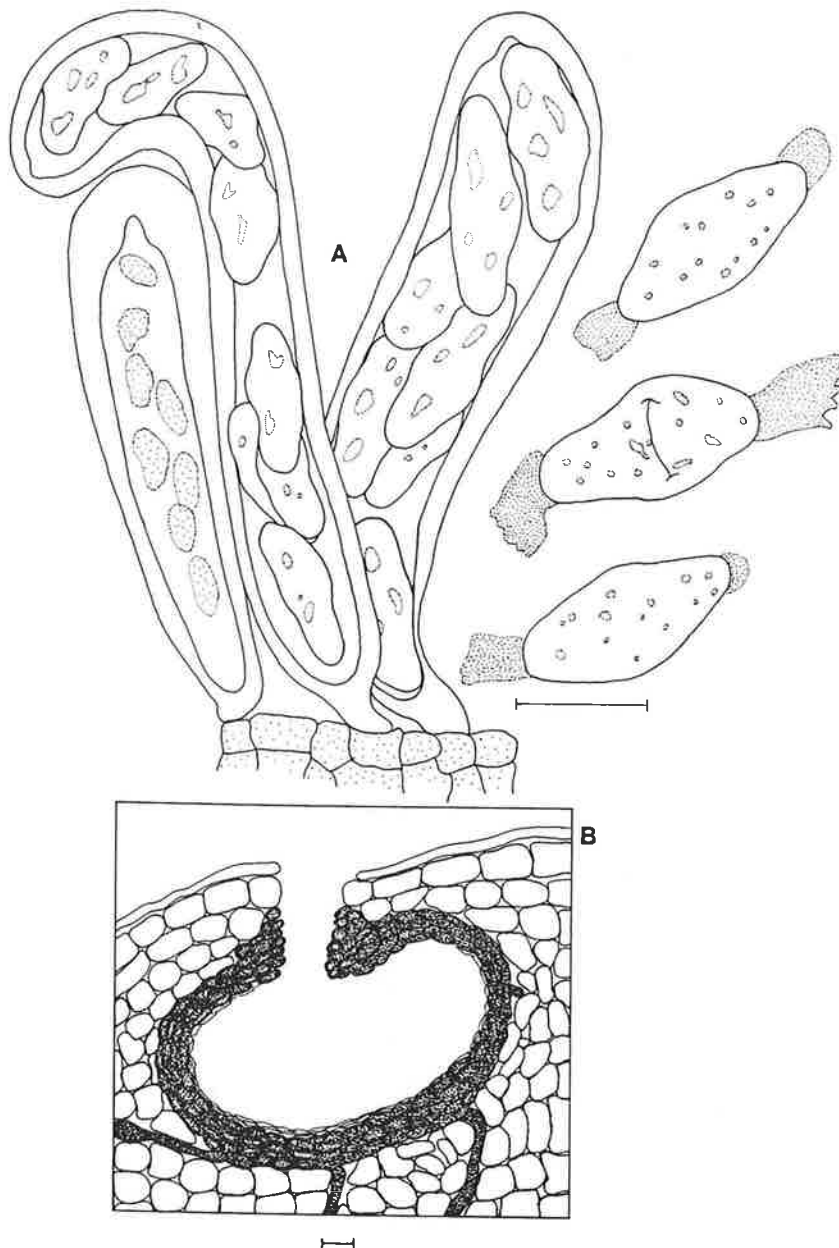


Figure 2 *Guignardia podocarpi* (PREM 51902). A. Bitunicate asci and guttulate ascospores with mucous caps. The horizontal line in the second ascospore is a fold in the exterior cell wall, presumably induced during slide preparation. B. Vertical section through a pseudothecium (bars = 10 μ m).

globosi, usque ad 200 μ m diam. et 150 μ m alti, inter pycnidia, atrobrunnei, solitarii, uniloculares, collo prominenti; parietes 3–6 cellulis crassis, ex textura angulare medio vel atro brunnea, 10–20 \times 5–6 μ m. Asci clavati vel cylindrici, bitunicati, 8-sporei, 60–85 \times 14–18 μ m. Ascosporae hyalinae, guttulatae, unicellulares, (19–)20(–23) \times (7–)8(–9) μ m, fusiformes vel ellipsoideae, latiores in mediano, in apice obtusae appendice gelatina exhibentes.

Mycelium immersed, consisting of septate, branched, medium-brown hyphae, 5–8 μ m diam. Ascocarps sparse, immersed, subepidermal, subglobose, up to 200 μ m diam. and 150 μ m in height, intermixed amongst pycnidia, dark brown, solitary, uni-locular with a prominent neck; wall consisting of 3–6 layers of *textura angularis*, cells 10–20 \times 5–6 μ m, medium to dark brown. Asci clavate to cylindrical bitunicate, 8-spored, 60–85 \times 14–18 μ m. Ascospores evenly distributed in asci, hyaline, guttulate, unicellular, (19–)20(–23) \times (7–)8(–9) μ m, fusiform-ellipsoidal, wider in middle, guttulate, ends obtuse with polar gelatinous appendages.

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, Jun. 1994, PREM 51902 (holotype).

As far as we could establish, no taxa of this group have been described from *Podocarpus*. Pseudothecia of *G. podocarpi* occurred on several leaves in close association with pycnidial conidiomata of a *Phyllosticta* species and a *Leptodothiorella* Höhn. microconidial synanamorph. In his review of the genus, van der Aa (1973) accepted that different *Phyllosticta* species are generally associated with different hosts. As *Phyllosticta* spp. are well established anamorphs of *Guignardia* (Bissett 1986a, 1986b), this species is subsequently described below as the suspected anamorph of *G. podocarpi*.

***Phyllosticta podocarpi* Crous sp. nov.** (Figure 3)

Conidiomata pycnidialia, dispersa, immersa, subepidermalia, erumpentia, subglobose, solitaria, unilocularia, usque ad 60 μ m diam.,

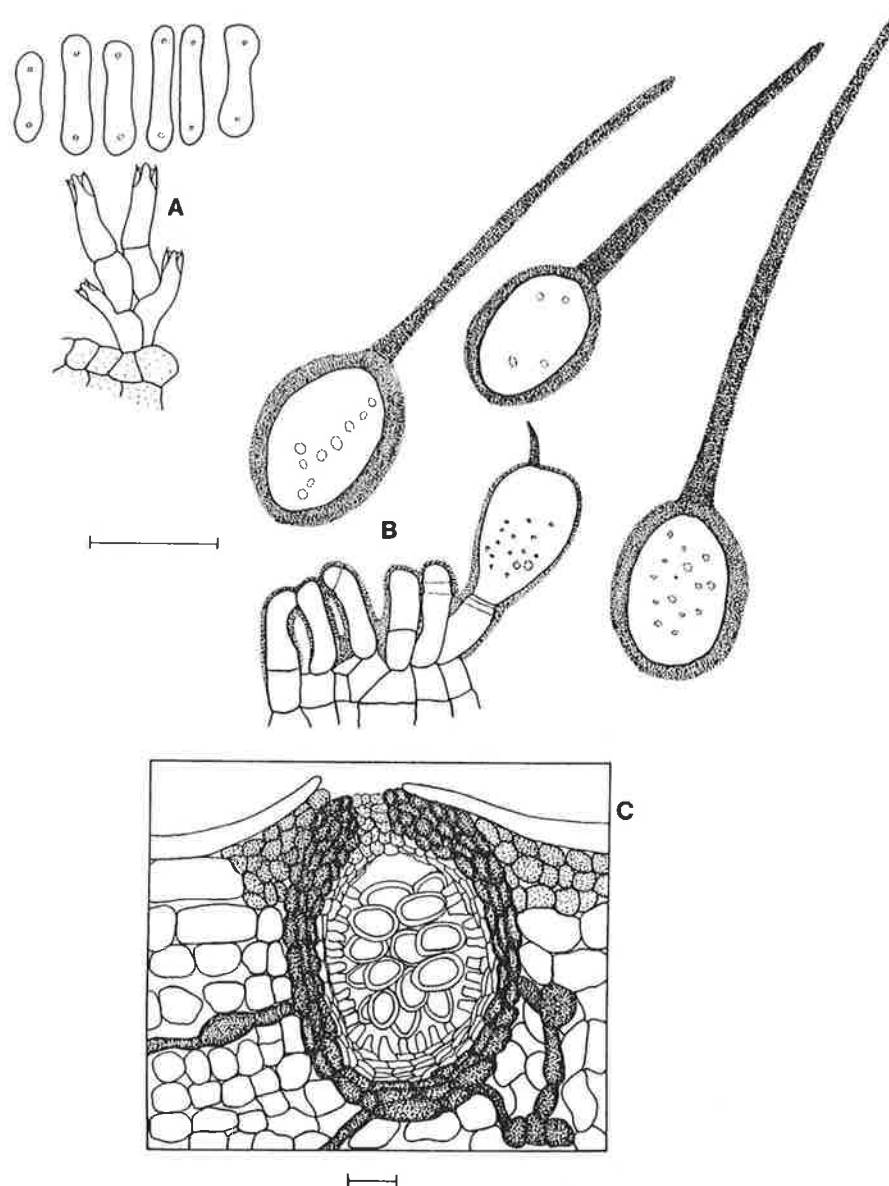


Figure 3 *Phyllosticta podocarpi* (PREM 51903). A. Microconidia and conidiophores. B. Macroconidia and conidiogenous cells. C. Vertical section through a conidioma (bars = 10 μm).

90 μm alta; paries consistens in 3–4 stratis cellularum brunnearum texturae angularis et strato interiore cellularum hyalinarum complanatarum. Cellulae conidiogerae cylindricae, 7–12 \times 3–5 μm proliferationibus 1–3 inconspicuis percurrentibus ad apicem. Conidia unicellularia, guttulata, late ellipsoidea vel subglobosa, (10–)14(–)17 \times (8–)9(–)10 μm tegmentis mucosis persistentibus ca. 1 μm crasses; appendicibus apicalibus 10–40 μm longis, base ca. 1.5–2 μm diam., acuta obtusa. Microconidia expulsa in cirris albis ex conidiomatibus immersis subepidermalibus pycnidialibus. Microconidiophora ramosa, 0–2-septata, 10–25 \times 3–4 μm . Cellulae microconidiogerae cylindricae, prominenter crassae periclinaliter, 7–11 \times 3–3.5 μm . Microconidia bacillaria apicibus obtusis tumidis, unicellularia, (6–)10(–)11 \times (2–)2.5(–)3 μm .

Conidiomata pycnidial, scattered, immersed, subepidermal, becoming erumpent, subglobose, solitary, unilocular, up to 60 μm diam, and 90 μm high; wall composed of 3–4 layers of brown cells of *textura angularis*, 7–20 \times 4–7 μm , and an inner layer of flattened cells. Conidiogenous cells cylindrical, 7–12 \times 3–5 μm , with 1–3 inconspicuous percurrent proliferations at the apex. Conidia unicellular, guttulate, broadly ellipsoidal to sub-

globose, (10–)14(–)17 \times (8–)9(–)10 μm with persistent mucous coats, ca. 1 μm thick; apical appendages 10–40 μm long, ca. 1.5–2 μm diam. at the base, tapering to an acutely rounded apex. Microconidia exuding as white cirri from immersed, subepidermal pycnidial conidiomata. Microconidiophores irregularly branched, 0–2-septate, 10–25 \times 3–4 μm . Microconidiogenous cells cylindrical, with prominent periclinal thickening, 7–11 \times 3–3.5 μm . Microconidia bacillar with swollen, obtuse ends, unicellular, (6–)10(–)11 \times (2–)2.5(–)3 μm .

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, Jun. 1994, PREM 51903 (holotype).

Parasymphodiella podocarpi Crous & Seifert sp. nov. (Figure 4)

Mycelium ex hyphis septatis, ramosis, laevibus, olivaceis vel brunneis, 2–4 μm diam compositum. Conidiophora mononemata, macronemata, non ramosa, cylindrica, recta, parietibus crassis, basim brunnea et subtiliter verrucosa, pallidiora et laeviora versus regionem conidiogeraem versus basim 10–14 μm lata, apicem 6–11 μm lata, regione fertili terminantia, 95–270 μm longa et 8–14 μm

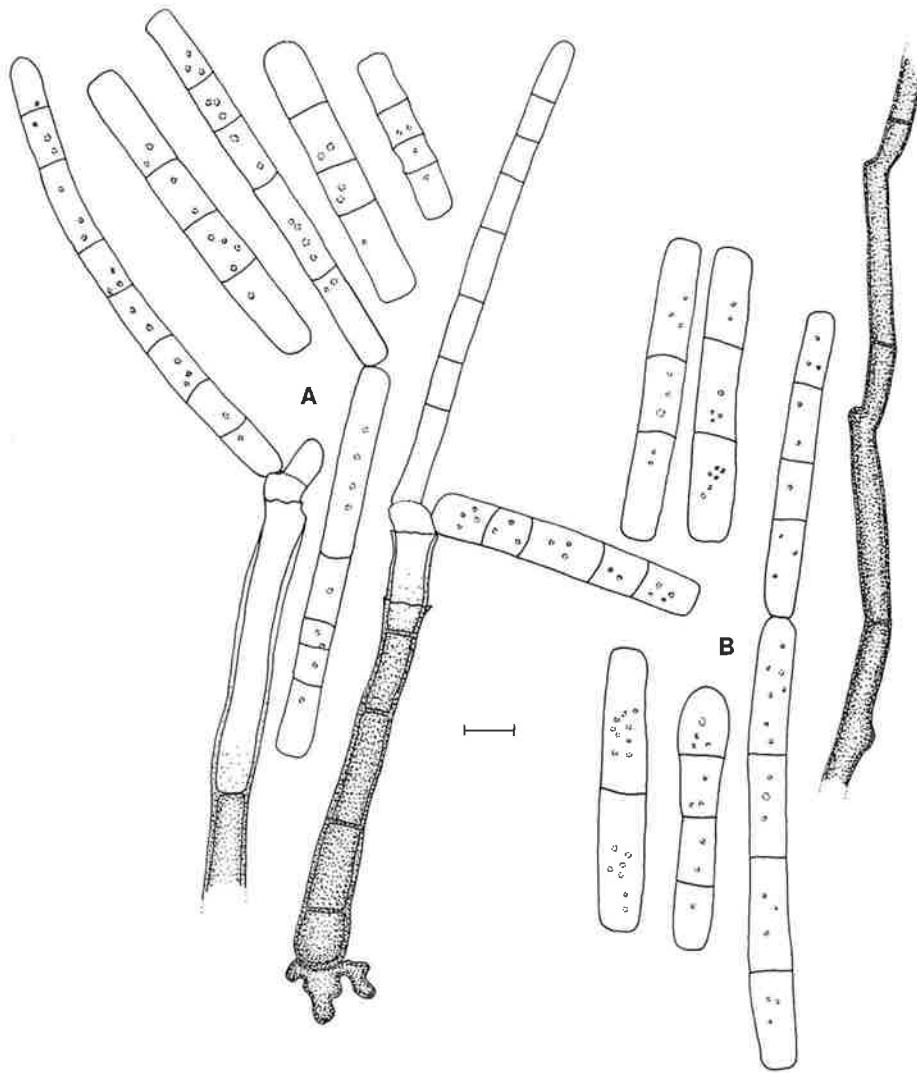


Figure 4 *Parasymptodiella podocarpi* (PREM 51904). Conidiophores and conidia *in vivo* (A), and *in vitro* on CLA (B) (bar = 10 μ m).

lata, 4–11-septata (*in vivo*), catenas longas formantia; conidiophora usque ad 1 000 μ m longa *in vitro*. Cellulae conidiogerae terminales, in conidiophoris incorporatae, indeterminatae, irregulariter sympodiales, 25–65 \times 5–6 μ m pallide brunneae vel hyalinae, 5–65 μ m inter locos conidiogeres. Conidia holothallica, sicca, catenulata, guttulata, recta vel plus minusve curvata, apicem et basim conidiorum intercalarium truncata, conidiis apicalibus apicibus obtusis et basim truncatis, (0–)3(–7)-septatis, (30–)62(–140) \times (6–)7(–9) μ m *in vivo*, 1(–3)-septatis, (30–)45(–70) \times (5–)8(–14) μ m *in vitro*.

Mycelium consisting of septate, branched, smooth, olivaceous to brown hyphae, 2–4 μ m diam. Conidiophores mononematous, macronematous, unbranched, cylindrical, straight, thick-walled (up to 1.5 μ m), dark brown and finely verrucose at the base, becoming smoother and lighter brown towards the conidiogenous region, 10–14 μ m wide at the base, 6–11 μ m wide at the apex, terminating in a smooth fertile region, 95–270 μ m long and 8–14 μ m wide, 4–11-septate (*in vivo*), giving rise to chains of conidia; conidiophores up to 1 000 μ m in length *in vitro*. Conidiogenous cells terminal, integrated, indeterminate, irregularly sympodial, 25–65 \times 5–6 μ m, light brown to hyaline, with 5–65 μ m between conidiogenous loci. Conidia holothallic, dry, catenate, guttulate, straight to slightly curved, cylindrical, apex and base of intercalary conidia truncate, apical conidia with obtuse apices and truncate bases, (0–)3(–7)-septate, (30–)62(–140) \times (6–)7(–9) μ m *in vivo*, 1(–3)-septate, (30–)45(–70) \times (5–)8(–14)

μ m *in vitro*; conidial chains appear sinuous as the conidia are developing.

Colonies on MEA attain a radius of 4–6 mm after 6 days at 15°C in the dark. Cultures are diffuse, spreading, with irregular margins of black fascicles of hyphae, extending beneath the agar surface, forming swollen, brown, chlamydospore-like cells, 12–25 \times 8–15 μ m; colony centres are dark brown to black with sparse aerial mycelium. Cardinal temperatures for growth are minimum: below 10°C, optimum: 15°C, and maximum: below 30°C.

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, Aug. 1994, PREM 51904 (holotype), DAOM 221069, culture STE-U 790, IMI 364301.

Of the remaining described species of *Parasymptodiella* Ponnappa with conidia devoid of septal plugs, *P. podocarpi* is most similar to *P. elongata* Crous, M.J. Wingf & W.B. Kendr. (1995), *P. minima* J.L. Crane & Schokn. (1982), *P. clarkii* B. Sutton (1978) and *P. longispora* (Tokum. & Tubaki) Tokum. However, it can easily be distinguished from *P. elongata* (20–40 \times 6–12 μ m; 0–2-septate), *P. minima* (11.5–14.5 \times 1.5–2 μ m; 3-septate), and *P. clarkii* (15–19 \times 2.5–3 μ m; 3-septate) which all have smaller conidia. In conidial dimensions, *P. podocarpi* is most similar to *P. longispora*, which produces 1(–3)-septate conidia *in vitro*, 32–68 \times 9–15 μ m (Tokumasu & Tubaki 1983), rather similar to

those we report for *P. podocarpus*. In culture, conidia of the type strain (CBS 544.84) produced 1–2(–5)-septate conidia, 35–130 × 6–9 μm. Chlamydospore-like structures were also observed to occur in the conidiophores (as nodal swellings) and in the mycelium. The growth rate of *P. longispora* at 25°C is 6 cm in 10 days, whereas that of *P. podocarpus* is reported to be 1–3 mm after 10 days at this temperature. Furthermore, although *P. longispora* produces abundant chlamydospores on malt agar (Tokumasu & Tubaki 1983), *P. podocarpus* only exhibited sparse chlamydospore formation, embedded in the agar of old cultures. Sutton *et al.* (1982) reported the presence of a *Stylaspergillus* B. Sutton *et al.* synanamorph for *P. laxa* (Subram & Vittal) Ponnappa. Tokumasu (1987) reported a similar *Stylaspergillus* anamorph from needles of *Pinus* in Japan that also had *P. longispora* present, but he was unable to confirm the connection in culture because conidia of the former anamorph did not germinate. We have not observed such a synanamorph in our cultures or specimens of *P. podocarpus*.

Discrepancies between conidiophore proliferation, conidial septation and conidial dimensions observed *in vitro* and *in vivo* for *P. podocarpus* bring to light problems in describing species from cultures alone. Although pure cultures can be grown under standardized conditions and thus make careful comparisons possible, there is a tendency for variation to occur in culture for some fungi. In many fungi with phragmoconidia, for example, conidia produced in culture have fewer septa than those produced in nature. In the cercosporoid complex, however, conidia again tend to be longer and develop numerous septa in culture (Crous

et al. 1989). In contrast to *P. longispora*, cultures of *P. podocarpus* had shorter conidia and longer conidiophores than observed in nature.

A key to distinguish the species presently accepted in *Parasymptodiella* is provided below.

Key to species

1. Conidia 3-septate, less than 20 μm in length 2
 Conidia if 3-septate, longer than 20 μm 3
2. Conidia 11.5–14.5 × 1.5–2 μm *P. minima*
 Conidia 15–19 × 2.5–3 μm *P. clarkii*
3. Conidia 3-septate with punctiform septal plugs, 18–50 × 6–8 μm *P. laxa*
 Conidia without septal plugs 4
4. Conidia 3–4-septate, 31–33 × 2 μm *P. africana*
 Conidia 0–multi-septate, more than 5 μm wide 5
5. Conidia 0–2-septate, 20–40 × 6–12 μm *P. elongata*
 Conidia with 1–5 septa on MEA, up to 70 μm long 6
6. Conidia 32–68 × 9–15 μm *in vivo*, 35–130 × 6–9 μm *in vitro*, forming numerous chlamydospores in vegetative hyphae, 23–46 × 10–25 μm, growing profusely at 25°C, occurring on *Pinus* *P. longispora*
 Conidia 30–140 × 6–9 μm *in vivo*, 30–70 × 5–14 μm *in vitro*, chlamydospores sparse, embedded in agar, 12–25 × 8–15 μm, growth stunted at 25°C, occurring on *Podocarpus*. *P. podocarpus*

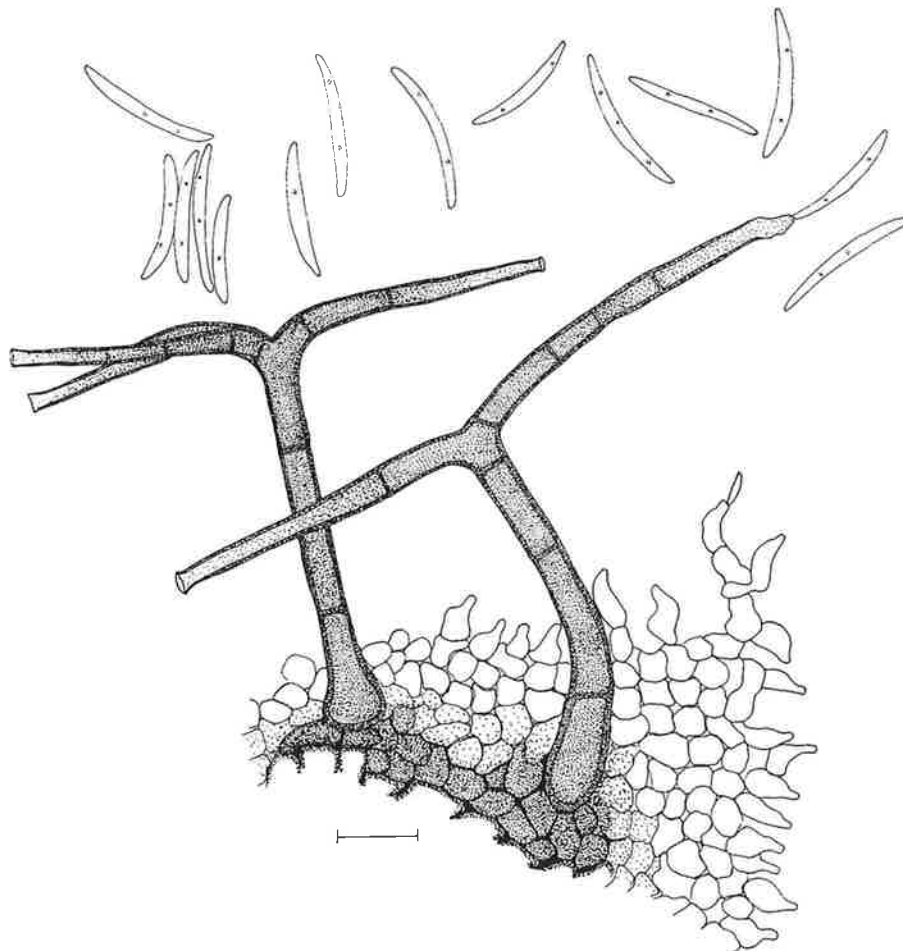


Figure 5 Setae, conidia and conidiogenous cells of *Gyrothrix verticiclada* *in vitro* on MEA (PREM 51906) (bar 10 = μm).

Gyrothrix verticiclada (Goid.) S. Hughes & Piroz., Can. J. Bot. 9: 42 (1971) (Figures 5, 9–13)

Peglionia verticiclada Goid., *Malpighia* 34: 7 (1935).

Mycelium immersed and superficial, consisting of branched, septate, hyaline to brown hyphae, 1.5–3 μm diam.; forming a large stroma consisting of smooth, brown, isodiametric cells on which the setae and conidiogenous cells are situated. The stroma is embedded in the host tissue (Hughes & Pirozynski 1971), or is formed beneath the agar surface in culture, giving colonies a dark brown to black appearance; strains lose some of their ability to form stromatic tissue with subsequent subculturing. Setae straight, erect, thick-walled, dark brown, becoming lighter brown in the upper two cells, occurring singly, or arranged in tight clusters, 50–80 μm tall (measured from base to below primary branch), 3.5–6 μm wide (at first basal septum), 4–7-septate, seldomly unbranched, frequently branching at the same locus to form 2–6 lateral branches nearly equal in length, 10–50 \times 4–6 μm , 1–3-septate, primary branches sometimes branch dichotomously to form secondary branches, 17–35 \times 3.5–4 μm , 1–3-septate; branch apices obtuse when immature, becoming swollen and fertile via small denticles or bumps, finally bursting open to appear like collarettes typical of phialides of *Dictyochaeta* Speg. spp. Conidiogenous cells smooth, olivaceous, irregular, straight, or geniculate-sinuuous, ampulliform to lageniform, 5–10 \times 3–4 μm *in vivo*, 10–23 \times 3–4.5 μm *in vitro*, giving rise to conidia via inconspicuous annellations. Conidia forming in a slimy mass around the base of setae (rarely at branch apices), hyaline, falcate, tapering to blunt apices, non-septate with a few minute guttules, (15–)17(–21) \times (1.5–)2 μm

Colonies on 2% malt-extract agar (MEA) attain a radius of 8–11 mm after 8 days at 25°C in the dark. Cultures are dark brown to black with little aerial mycelium on MEA. Cardinal temperatures for growth are minimum: below 10°C, and maximum: below 30°C.

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongus* leaf litter, P.W. Crous, Aug 1994, PREM 51906, DAOM 221067, IMI 364302, STE-U 786-788.

This fungus was originally described in Italy from leaf litter of *Laurus nobilis* L. and *Prunus cerasus* L. by Goidanich (1935), who erected a new genus *Peglionia* Goid. with *P. verticiclada* as type. Verona and Benedek (1967) commented that this fungus closely resembled species of *Gyrothrix* (Corda) Corda and *Circinotrichum* Nees. In 1963 two collections of this fungus were obtained from *Knightsia excelsa* R. Br. in New Zealand. In a subsequent study, Hughes & Pirozynski (1971) reduced *Peglionia* to synonymy with *Gyrothrix*, and also introduced a new name for this species as *G. verticiclada* (Goid.) S. Hughes & Piroz. The latter decision was chiefly based upon the branched setae present in both *Peglionia* and *Gyrothrix*, but that are lacking in *Circinotrichum*. In these genera, conidia are formed on conidiogenous cells that are situated around the base of the setae. The exact mode of conidiogenesis is unclear, but minute annellations can be seen at the apices of conidiogenous cells in *Circinotrichum* and *Gyrothrix* (Ellis 1971) (Figure 7). Furthermore, Castañeda and Kendrick (1990) introduced the genus *Selenodriella* Castañeda & W.B. Kendr. for species with minute denticles at the apices of their conidiogenous cells (Figure 8). In *G. verticiclada*, however, we could distinguish indistinct annellations at the apices of conidiogenous cells (Figures 9–13). Conidia were observed to be borne in whorls of 2–6 at the apices of conidiogenous cells. In a scanning electron microscopy study of the conidiogenesis of *Gyrothrix circinata* (Berk. & M.A. Curtis) S.

Hughes, Nakagiri and Ito (1991) illustrated the same arrangement of conidia. Conidiogenous cells were shown to have a collarette, with several denticle-like structures situated within the apex of the conidiogenous cell. These illustrations suggest, therefore, that the first conidium is probably produced holoblastically, and that the flat-tipped structures may be a compressed form of percurrent proliferation. The proliferation period of ontogeny is reduced, giving rise to inconspicuous scars on the apex within the collarette. Conidia then develop laterally to each other, and are borne in whorls as also seen in *G. verticiclada*. Onofri (1995) showed a similar mode of conidiogenesis in a culture of *Circinotrichum maculiforme* Nees, where the first conidium is produced holoblastically. Additional conidia are produced enteroblastically, with several loci forming laterally on the enteroblastic wall of the conidiogenous cell, appearing as small, flat-tipped scars in the illustrations of Nakagiri and Ito (1991) for *G. circinata*.

A closer examination of the apices of the dichotomously branching setae of *G. verticiclada* showed them to frequently become swollen, and to be open at maturity, appearing like a giant phialide. When young material is studied, however, small denticles are observed at the branch apices, to which conidia are attached in clusters. With age, these apices become swollen to the outside, whereupon they burst, appearing like an open phialide with a flared collarette, somewhat resembling that of the genus *Dictyochaeta*.

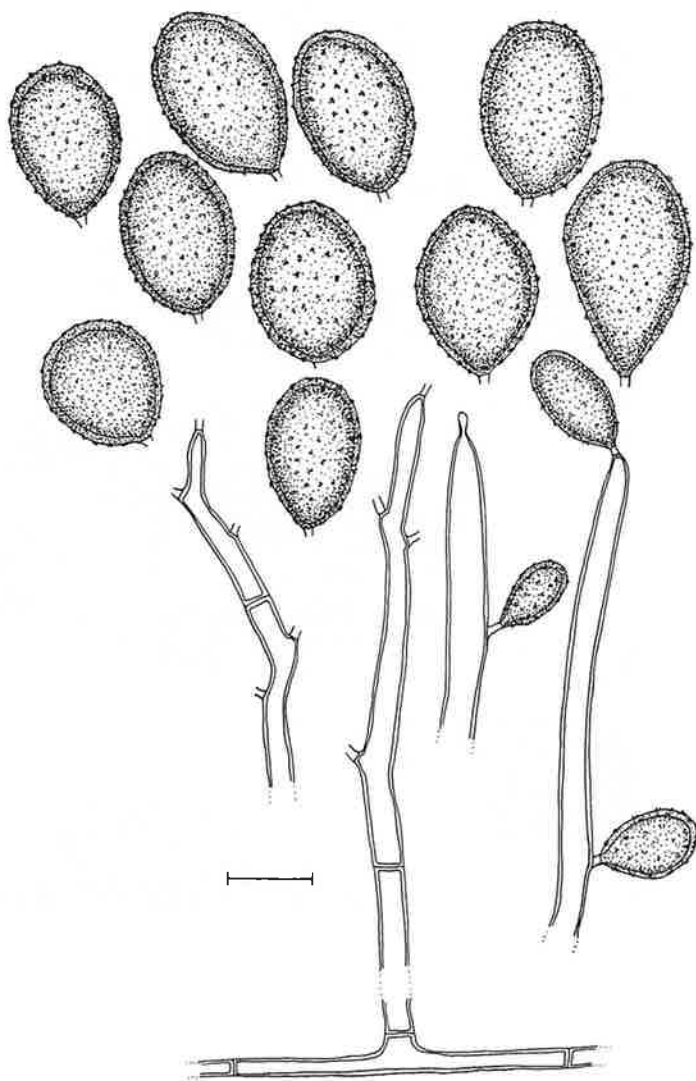
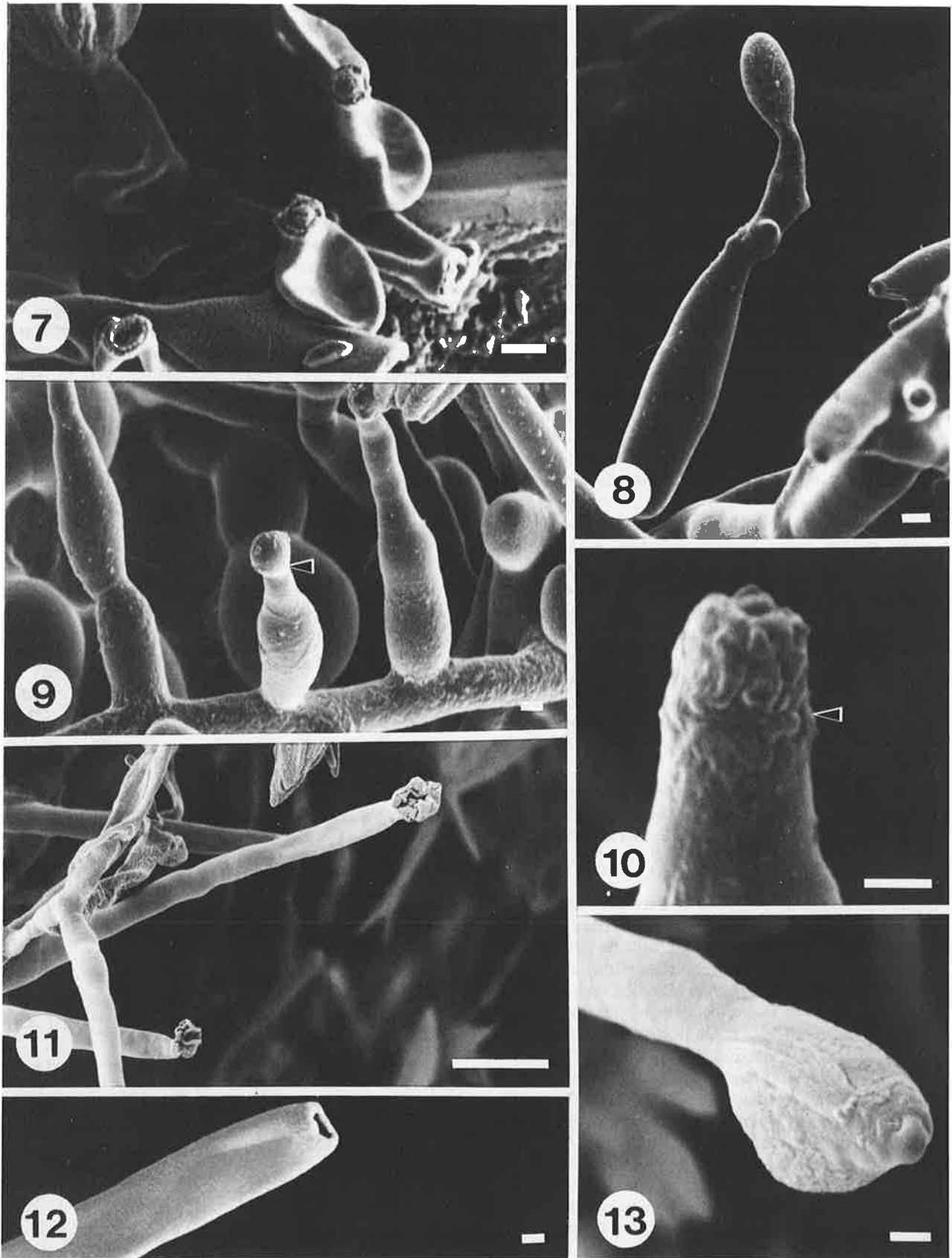
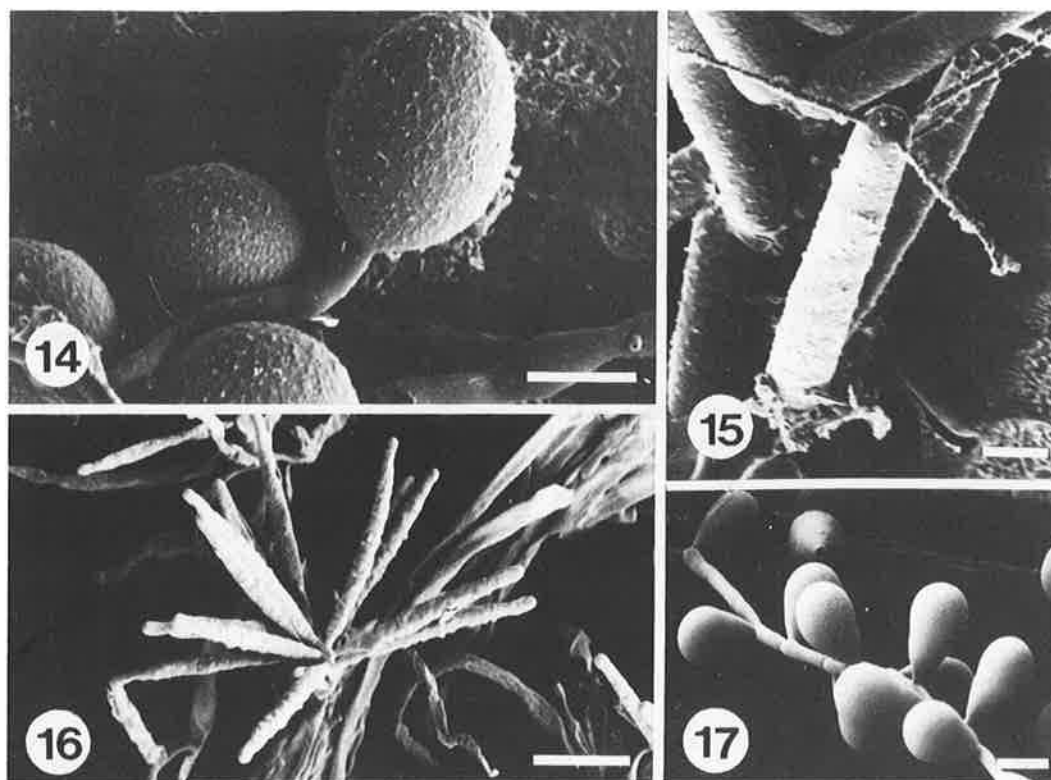


Figure 6 Conidia and conidiophores of *Rhinotrichella elegans* (PREM 51905) (bar 10 = μm).



Figures 7–13 Conidiogenous cells and setae of *Gyrothrix* and *Selenodriella*. **7.** Pcurrent annellations at the apices of conidio-genous cells of a *Gyrothrix* sp. (bar = 1 μ m). **8.** Denticles at the apex of a conidiogenous cell of a *Selenodriella* sp. (bar = 1 μ m). **9–13.** *Gyrothrix verticillata* (PREM 51906). **9.** Outside cell layer (arrowed) that was involved in the holoblastic formation of the first conidium (bar = 1 μ m). **10.** Warty swelling developing at the setal apex (arrow) (bar = 1 μ m). **11.** Enlarged apical warty swellings on setae, that can also be fertile (bar = 10 μ m). **12.** Empty cell at setal apex without warty swelling (bar = 1 μ m). **13.** Protoplasm leaking from apical cell after discharge of warty swelling (bar = 1 μ m).



Figures 14–17 Conidia of various microfungi (bars = 10 μm). 14. *Rhinotrichella elegans* (PREM 51905). 15. *Camposporium antennatum* (STE-U 667). 16. *Dactylaria irregularis* (PREM 51908). 17. *Endophragmiella boewei* (PREM 51909).

Rhinotrichella elegans R.F. Castañeda & Crous sp. nov. (Figures 6, 14)

Mycelium copiosum, ex hyphis septatis, ramosis, laevibus, hyalinis, 1.5–2 μm diam. compositum. Conidiophora conspicua mononematosa, erecta, flexuosa, simplicia, interdum ramosa, cylindrica, septata, apice leviter geniculata, hyalina, basim 300–780 μm alta et 4–6.5(–8) μm crassa. Cellulae conidiogerae polyblasticae, terminales et intercalares, sympodialiter extendentes, in conidiophoris incorporatae, incoloratae, denticulatae, denticulis conspicuis, cylindricis, translucidis, 1–2 μm longis praeditae. Conidia obovata, interdum obpyriformia vel ellipsoidea, minime verrucosa vel levia, primo hyalina, tarde pallide brunnea vel dilute cinnabarina, unicellularia, acropleurogena, sicca, (19–)29(–37) \times (14–)18.5(–22) μm ad basim rotunda, appendice, truncato, translucido, conspicuo, 1–2.5 μm longo (reliquiis cellularum conidiogarium) praedita. Teleomorphosis ignota.

Mycelium abundant in culture, composed of septate, branched hyphae, 1.5–2 μm diam. Conidiophores undifferentiated, mononematous, erect, flexuous, mostly unbranched, cylindrical septate, slightly geniculate at the apex, colourless, 300–700 μm tall; 4–6.5(–8) μm wide at the base. Conidiogenous cells polyblastic, terminal and intercalary, proliferating sympodially, integrated, denticulate, with conspicuous, cylindrical denticles, 1–2 μm long. Conidia obovate, sometimes obpyriform to ellipsoid, finely verrucose to smooth-walled, initially hyaline, becoming pale brown to orange-red, thick-walled, 0-septate, acropleurogenous, dry, (19–)29(–37) \times (14–)18.5(–22) μm , base rounded but with a conspicuous, clear, truncate, hyaline appendage (remains of the conidiogenous cell), 1–2.5 μm long. Teleomorph unknown.

Colonies on MEA attain a radius of 18–30 mm after 8 days at 15°C in the dark. Cultures are floccose to cottony, initially white, turning orange. Cardinal temperatures for growth are minimum: below 5°C, optimum: 15°C, and maximum: below 30°C.

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, 31 Aug. 1993, PREM 51905 (holotype), DAOM 221068, culture STE-U 668.

The genus *Rhinotrichella* G. Arnaud ex de Hoog was erected by Arnaud (1953) without a Latin diagnosis, and was validated by de Hoog (1977) with *R. globulifera* de Hoog as type species. The latter has globose, smooth or finely verrucose, pale ochraceous conidia, 9–12 μm diam. Matsushima 1983 described an additional species, *R. macrospora* Matsush., with smooth, pale brown, globose conidia, 15–19 μm diam, and a synanamorph resembling a species of *Aspergillus* P. Micheli ex Link (not illustrated). Both these species are clearly distinct from *R. elegans*, which has larger conidia.

During the course of this study numerous other hyphomycetes were also isolated. As far as we could establish, several have not previously been recorded from South Africa. However, as their morphology closely corresponds with that of their respective descriptions, they are merely listed below.

Other new records

Camposporium antennatum Harkn., Bull. Calif. Acad. Sci. 1: 37–38 (1884) (Figure 15) (*vide* Ellis 1971).

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, 31 Aug. 1993, STE-U 667.

Dactylaria irregularis De Hoog, Stud. Mycol. 26: 124 (1985) (Figure 16)

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, 31 Aug. 1993, PREM 51908.

Endophragmiella boewei (Crane) S. Hughes, N.Z. J. Bot. 17: 147 (1979) (Figure 17) (fide Ellis 1976).

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, 31 Aug. 1993, PREM 51909, STE-U 670.

Phaeoisaria clematidis (Fuckel) S. Hughes, Can. J. Bot. 36: 795 (1958) (fide Ellis 1971).

Specimen examined: Western Cape Province, Stellenbosch, Botanical Garden, *Podocarpus elongatus* leaf litter, P.W. Crous, 26 Sept. 1994, PREM 51907, STE-U 813.

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