# New data for Rinodina flavosoralifera

# Mireia GIRALT, Pieter P. G. van den BOOM, Tor TØNSBERG and John A. ELIX

**Abstract:** Well-developed ascospores of *Rinodina flavosoralifera* have been observed for the first time. The ascospores are described and illustrated and new data on the chemistry and distribution of this species are provided. New chorological data on *Rinodina disjuncta* are also included.

Key words: Canary Islands, chemistry, distribution, morphology, *Physciaceae*, rare lichens

#### Introduction

Rinodina flavosoralifera was first described by Tønsberg (1992). Although it has since been collected several times no specimen has been reported to have well-developed ascospores, an important character for distinguishing species of Rinodina and for segregating genera in the Physciaceae. The study of well-developed, fertile specimens from the Canary Islands has revealed that this species has typical Polyblastidium-type ascospores, an ascospore type previously known only in the genus Heterodermia in the Physciaceae. A brief description and drawings of the ascospores are presented here.

#### Methods

The specimens were examined by standard techniques using stereoscopic and compound microscopes. Ascospores were measured mounted in distilled water and ascus tip structures were observed after pretreatment with K and washing with distilled water before the

application of Lugol's iodine. The chemical constituents were identified by high performance liquid chromatography (HPLC) according to methods described by Elix *et al.* (2003).

### The Species

## Rinodina flavosoralifera Tønsberg

Sommerfeltia 14: 289–290 (1992); type: Norway, Hordaland, Bergen, Store Milde, Fana Folkehøgskole, 20–40 m, on a mossy trunk of *Prunus avium*, 1988, *T. Tønsberg* 11375 (BG—holotype!).

(Fig. 1)

Thallus morphology is totally in accordance with previous descriptions (Tønsberg 1992; Mayrhofer & Moberg 2002; Giavarini et al. 2009).

Apothecia lecanorine, 0.3–0.5 mm diam., sessile, constricted at the base. Thalline margin entire, thick, rather prominent, persistent, concolorous with the areolae. Proper margin visible as a ring within the thalline margin. Disc flat, brown, ± covered by a whitish pruina. Excipulum thallinum 55-70 µm thick laterally, expanded below to 80–120 μm. Cortex paraplectenchymatous, nearly indistinct to 10 µm wide laterally, expanded below to 45-60(-100) µm, I-. Excipulum proprium colourless, 10-20 µm wide laterally, expanded above to 30-60(-70) µm. Hymenium hyaline, up to 180 µm high. Epihymenium pale brown inspersed with crystals (pruina). Hypothecium

M. Giralt: Departament de Bioquímica i Biotecnologia (Àrea de Botànica), Facultat d'Enologia de Tarragona, Universitat Rovira i Virgili, Marcel·lí Domingo s/n, 43007, Tarragona, Spain. Email: mireia.giralt@urv.cat P. P. G. van den Boom: Arafura 16, 5691 JA, Son, The Netherlands.

T. Tønsberg: Museum of Natural History, University of Bergen, Allégt. 41, P.O. Box 7800, N-5020 Bergen, Norway.

J. A. Elix: Research School of Chemistry, Building 33, Australian National University, Canberra, ACT 0200, Australia.

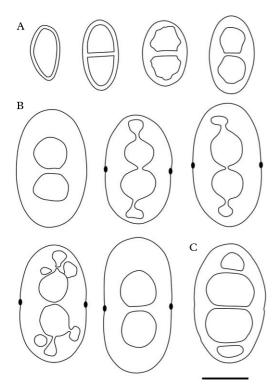


FIG. 1. Ascospore ontogeny and ascospore variability of *Rinodina flavosoralifera* (*P. & B. v.d. Boom* 42446, hb. v.d. Boom). A, Type A ontogeny and young colourless ascospores; B, mature ascospores of *Pachysporaria*-type and of *Polyblastidium*-type showing one or more sporoblastidia arising from each end of the two main lumina and a well developed torus; C, overmature, 3-septate ascospore. Scale = 10 µm.

hyaline, 40-60 µm deep. Paraphyses with pale brown apical cells, slightly enlarged, 2-3 µm wide. Asci Lecanora-type (Rambold et al. 1994), 8-spored. Ascospores colourless when immature,  $12-17 \times 7-10 \mu m$ ; mature ascospores brown,  $22-30 \times 11-17$  µm, Pachysporaria-type at first but usually developing sporoblastidia and then becoming Polyblastidium-type (Kurokawa sporoblastidia arising from each end of the two main lumina, rarely from the lateral parts (Fig. 1B & C); overmature ascospores 3-septate (Fig. 1C) or even submuriform; walls not ornamented, torus well developed, ontogeny of type A (Giralt 2001) (Fig. 1A).

Pycnidia and conidia not seen in the material studied here. According to Tønsberg

(1992), the only pycnidium observed was similar to the apothecia in form and size and the conidia were rod–shaped,  $4-5 \times c$ . 1 µm.

Chemistry [v. d. Boom 42446]. Arthothelin [major], thiophanic acid [minor], 6-O-methylarthothelin [minor], 4,5-dichloronorlichexanthone [trace]. The latter two substances were not reported previously from this species.

Observations. As the morphology of the thallus, the chemistry and the immature ascospores are identical with that reported previously for this species, we are certain that the Canarian specimens belong to *R. flavosoralifera*. As a consequence, *R. flavosoralifera* can now be characterized by the yellowish thallus composed of sorediate areolae, the presence of the arthothelin chemosyndrome and the large *Pachysporaria*-type ascospores which develop sporoblastidia (*Polyblastidum*-type), have smooth walls and exhibit a very well-developed torus.

The *Polyblastidium*-type ascospore together with the presence of xanthones make *R. flavosoralifera* unique within the genus *Rinodina* and supports the uncertainty of its generic position as indicated by other authors (Mayrhofer & Moberg 2002; Giavarini *et al.* 2009).

According to the literature, typical Polyblastidium-type ascospores are only known from some species of the foliose genus Heterodermia Trevis. (cf. Kurokawa 1962; Poelt 1965). However, some other Rinodina species have been described with ascospores somewhat similar to the *Polyblastidium*-type, including R. subanceps (Nyl.) Zahlbr. (cf. Mayrhofer 1984), R. verruciformis Sheard (cf. Sheard & Mayrhofer 2002) and the R. dolichospora-group (cf. Giralt et al. 2009). As far as the chemistry is concerned, at present the only Rinodina species known to contain xanthones are R. thiomela (Nyl.) Müll. Arg. and R. xanthomelana Müll. Arg. from the Southern Hemisphere, the pantropical R. lepida (Nyl.) Müll. Arg. (all three contain thiomelin and various congeners, cf. Leuckert & Mayrhofer 1984; Trinkaus et al. 2002) and R. efflorescens Malme and R. sheardii Tønsberg (both containing secalonic acid A, see Elix & Tønsberg 1999) from the Northern Hemisphere. A further *Rinodina* species probably contains xanthones, namely *R. chrysomelaena* (Ach.) Tuck., since it has a citrine thallus which reacts C+ faint orange (Lendemer & Sheard 2006). However, none of these *Rinodina* species have *Polyblastidium*type ascospores.

Ecology and distribution. The specimen v. d. Boom 42446 was growing together with Rinodina disjuncta Sheard & Tønsberg, Frullania sp., Hypogymnia tavaresii D. Hawksw. & P. James, Micarea peliocarpa (Anzi) Coppins & R. Sant., Placynthiella dasaea (Stirt.) Tønsberg, Trapeliopsis granulosa (Hoffm.) Lumbsch and *Usnea* spp., on the main branches of Erica arborea, on a gently sloping hill with grassy undergrowth. Elsewhere trunks and branches of the same phorophyte have been found to support Evernia prunastri (L.) Ach., Heterodermia obscurata (Nyl.) Trevis., Parmotrema perlatum (Huds.) M. Choisy and P. reticulatum (Taylor) M. Choisy.

Rinodina flavosoralifera is known from Norway (Tønsberg 1992; Mayrhofer & Moberg 2002), from the western Pyrenees (van den Boom et al. 1995), from Alaska (Tønsberg 2002) and from England, Wales and Scotland (Coppins & O'Dare 1994; Giavarini et al. 2009). Previously R. disjuncta was known from Norway and the Pacific coast of North America, from Alaska to northern California (Tønsberg 1992; Sheard 1995; Mayrhofer & Moberg 2002).

Both *Rinodina flavosoralifera* and *R. disjuncta* show a western European – north-west North American disjunct distribution. Both are new records for the Canary Islands (*cf.* van den Boom *et al.* 2008) and for Africa and constitute the most southern known records for both species.

Additional specimens examined. Canary Islands: El Hierro: near top of mountain range, NNW of El Pinar, along road to El Pinar, c. 900 m S of crossing with road H1-1, SW of Montaña de la Fuente, rim of 'Fayal non mature Erica arborea (together with Rinodina disjuncta), 17°59.70′W; 27°44.00′N, 2009, P. & B. van den Boom 42446 (hb. v.d. Boom). Gran Canaria: 30–

40 m (vertical) uphill from country road GC-216 between Artenara village and Mt Tamadaba, 28°01.508′N; 15°39.791′W, 1170–1180 m, corticolous near base of trunks of *Pinus canariensis* in *P. canariensis* forest in steep N-facing slope, 2007, *T. Tønsberg* 37706 (to be distributed in Tønsberg, *Lich. isid. sored. crust. exs.*), 37707 (BG, fertile).

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