INTRODUCTION

The lichen genus *Pertusaria* DC. s.l. (Pertusariaceae, Ascomycota) is characterized by crustose, continuous to cracked thalli (in many taxa with isidia or soredia), usually large apothecia with open or perithecia-like discs, 1–8-spored asci with thick walls, and large and simple ascospores with thick, simple layered or laminate walls which are uniform, ornamented or striate. Secondary lichen metabolites are very diverse in *Pertusaria* and include depsides, depsidones fatty acids and xanthones (Dibben, 1980; Archer, 1993, 1997; Chambers et al., 2009). The genus is cosmopolitan, and its members are found from the Arctic and Antarctic to the tropics in both Hemispheres. About 800 species of *Pertusaria* s.l. are known worldwide, many of them with several varieties, which actually usually may represent habitat modification only and are unworthy of highlighting (Tobolewski, 1972; Dibben, 1980; Archer, 1993, 1997; Lumbsch & Nash, 2001; Archer & Elix, 2018 and literature cited therein). Nevertheless, there are still significant gaps in the distribution of many taxa in different regions of the world and many uncertainties related to taxonomy within the genus.

During a revision of the type materials of *Pertusaria* DC. in the Herbarium of the Turku University (TUR-V) collections named *Pertusaria atropallida* Vain. and *P. uralensis* Vain., which appeared to be synonyms of *P. coccodes* (Ach.) Nyl. The species *Pertusaria atropallida* was described from Näränkäväara Mt. (Kuusamo, Finland) as growing on the bark of fir (Wainio, 1881). According to the protologue, the species has ± rimose-cracked, yellowish-white or yellowish-pink thallus with smooth upper surface without zoned margin, isidia densely covering parts of the thallus, immersed apothecia with black discs, and simple ascospores measuring 30–80 × 24–40 µm. *Pertusaria uralensis* was reported only from Russia „prope Stationem Uralskaja in Montibus Uralensibus” (Vainio, 1928). The species is characterized by lacking isidia or soredia, and with 2–4 ostiolate apothecia, but without ascospores. Thallus reacts K+ yellow then turning red.

MATERIAL AND METHODS

The study is based on specimens deposited in TUR. Additional specimens from B, H, KRA, KRAM, SLTC, TRN, UGDA and UPS were studied for comparison (almost 140 specimens were examined). All samples were examined for anatomical, morphological and chemical characters. The morphology of the specimens was studied using a stereomicroscope. The following characters were examined: thickness, structure and colour of thallus, colour, shape and size of isidia and soredia. Anatomy was examined in handmade section or squash preparations mounted in water or KOH (in case of deformed ascospores in old specimens). Secondary metabolites were identified by thin-layer chromatography (TLC; solvents A and C) according to the methods proposed by Culberson & Kristinsson (1970), White & James (1985) and by Orange et al. (2001); metabolites reported refer to those detected in type material and other studied specimens.

Locality information is provided here in the same form as written on the original labels.
RESULTS & DISCUSSION

Two herbarium specimens (TUR-V 6743, 6744) were found to represent original material of *P. atropallida*. TUR-V 6743 is annotated “Fennia, Kuusamo, Näränkävaara” and has original note “e specim. orig.”, while TUR-V 6744 is annotated “Kuusamo, Näränkävaara” and is designated as “n. sp.” Information on labels and morphological characters correspond to the description in the protologue, but E. Vainio did not determine any specimen as a holotype, therefore lectotype should be selected (Art. 9.11; Turland et al., 2018). Because the specimen TUR-V 6744 is accompanied by original description with handwritten notes of E. Vainio, it should be designated as lectotype, while TUR-V 6743 as its duplicate (isolectotype). However, the situation is more complicated as the name *P. atropallida* is based on two organisms, because the original material includes sterile lichenized thalli and black perithecioid ascomata of a lichenicolous fungus, and both elements were apparently used for the species description.

The lichenized thallus present in the material is isidiate and contains norstictic acid as found by thin layer chromatography (TLC) and fits well the current concept of *P. coccodes* (Chambers et al., 2009). Meanwhile, the ascomatal structures described by Wainio (1881) clearly refer to lichenicolous fungus as no other immersed ascomata were detected. I sectioned four perithecioid ascomata, but no ascospores were found. Since no lichenicolous fungus growing on *Pertusaria* has similar size of ascospores as reported by Wainio (1881), the identity of the fungus species remains unknown.

As the material of *P. atropallida* belongs to more than one taxon, one has to be selected as the lectotype (Art. 9.11 and Art. 9.14; Turland et al., 2018). In order to keep the name in the genus *Pertusaria* has similar size of ascospores as reported by Wainio (1881), the identity of the fungus species remains unknown.

Vainio (1928) did not determine any specimen as a holotype in the protologue, and only locality was reported. Therefore lectotype has to be selected, because it is not possible to determine that at the time of describing the species E. Vainio had only one element upon which the validating description was based (Art. 9.3; Turland et al., 2018; see also McNeill, 2014).

The new synonymy of *P. coccodes* are as follows:


Lectotype (designated here): Finlandia, Kuusamo, Kuusamon pit., Näränkävaara, Kuusen kuorella kuusimetsässä, 1877, E. Wainio (TUR-V 6744!), thallus of *Pertusaria coccodes*; isolecototypes; TUR-V 6743!, H 9505289!, HBG!).


Lectotype (designated here): Rossia, Jugum Uralense, Uralskaja. Pichta, 1880, E. Wainio (TUR-V 6802!).
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REFERENCES


