

Article

A New Species and a New Record of Bysoid Arthoniaceae (Lichenized Ascomycota) from Southern China

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Abstract: This paper illustrates two species in the lichen-forming family Arthoniaceae from southern China, including a new species of *Herpothallon*, *H. fibrosum* L.L. Liu & Lu L. Zhang and a new record of *Cryptothecia*, *C. striata* G. Thor for China. *Herpothallon fibrosum* has fluffy, cylindrical pseudoisidia, like a bundle of fiber, and psoromic acid and confluent acid are present. Furthermore, the new record of *Cryptothecia striata* has been identified by morphological, anatomical, chemical, and molecular studies. The systematic position of the two species was clarified by the molecular sequence data from the small subunit of the mitochondrial ribosomal DNA (mtSSU). Detailed taxonomic descriptions, chemical characters, comparisons, and discussion of the characteristics of similar species are provided for the two species; the relationship between *Cryptothecia* and *Herpothallon* is also discussed here.

Keywords: bysoid thalli; *Cryptothecia*; *Herpothallon*; molecular phylogeny

1. Introduction

The family Arthoniaceae (Arthoniales, Arthoniomycetes, Ascomycota) contains over 700 lichenized species in 20 genera [1], and it is a major component of the lichen flora of many forest types, especially in the tropical and subtropical regions [2,3]. Up to now, 58 species of 11 genera in Arthoniaceae have been reported from China, including five species of *Cryptothecia* and 17 species of *Herpothallon* [4–7].

Morphologically, *Cryptothecia* and *Herpothallon* are closely related. Both have conspicuous bysoid crusts with trentepohlioid photobionts and asci not developing inside true ascomata. And species of *Herpothallon* were, until 2009, included within *Cryptothecia* but have been segregated by Aptroot et al. [8]; the thallus of *Cryptothecia* species is usually more tightly attached to the substrate than that of *Herpothallon*, most described species in *Cryptothecia* are fertile and without soredia and pseudoisidia; the species of *Herpothallon* typically have abundant pseudoisidia or other propagules for vegetative dispersal and are mostly sterile. Only one species, *H. fertile*, is known to produce isolated asci within pseudoisidia and ascospores with straight, not curved septa (as is characteristic for *Cryptothecia*) [8–14]. In addition, pycnidia are developed in some species inside or at the top of pseudoisidia, such as *H. isidiatum* Jagadeesh Ram & G. P. Sinha and *H. tomentosum* L.L. Liu & Lu L. Zhang [6,11]. However, in our research, we frequently encountered sterile crustose lichens that lack distinctive characteristics for accurate taxonomic classification. Adding many sterile species to *Herpothallon* or any other genus simply would make this genus even more of a dustbin. In this case, the genotypic characteristics assume great significance alongside the morphological and chemical traits. Aptroot et al. [15] described these unknown sterile species in the genera that are phylogenetically closest in their recent work. This work offers us valuable guidance in the nomenclature of sterile tropical Arthoniaceae.

In southern China, there are abundant subtropical to tropical evergreen resources. This habitat is favorable for the lichens of Arthoniaceae. Since 2020, we have conducted several investigations on Arthoniaceae in southern China and found that this family exhibits a high level of species richness, especially in the vicinity of flowing water sources, such as streams



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and river valleys within forested areas. During our research, a new species of *Herpothallon* and a new record of *Cryptothecia* for China were discovered; we used mitochondrial small subunit (mtSSU) rDNA data to construct a phylogenetic tree to show the position of these two taxa and to support the delimitation of these species in the family Arthoniaceae. Our study found that the ITS sequence, typically considered crucial for distinguishing between lichen species, is essentially identical in *Cryptothecia* and *Herpothallon* and cannot be used as a standard for distinguishing between species within these two genera.

2. Materials and Methods

2.1. Investigation of Lichen Specimens

The specimens studied were collected in Fujian, Guangdong, Guizhou, Yunnan, Zhejiang Provinces, and Guangxi Zhuang Autonomous Region, China, and are preserved in the Lichen Section of the Botanical Herbarium, Shandong Normal University, Jinan, China (SDNU). The voucher numbers refer to Table 1.

Table 1. Specimens and sequences used for phylogenetic analyses. Newly generated sequences are in bold. * = outgroup.

| Taxon | Locality | GenBank Accession no. | Voucher |
|---------------------------------------|--------------|-----------------------|-------------------------------|
| <i>Arthonia calcarea</i> | Sweden | KJ850974 | Thor 11/6a (UPS) |
| <i>Arthonia radiata</i> | Sweden | KJ850968 | Frisch 10/Se29 (UPS) |
| <i>Arthothelium orbilliferum</i> | Norway | KY983977 | TRH-L-15449 |
| <i>Crypthonia palaeotropica</i> | Uganda | KJ850961 | Frisch 11/Ug457 (UPS) |
| <i>Cryptophaea phaeospora</i> | DR Congo | KX077541 | Van den Broeck 5809 (BR) |
| <i>Cryptothecia</i> sp. 1 | Rwanda | EU704053 | Ertz 8472 (BR) |
| <i>Cryptothecia</i> sp. 2 | Uganda | KJ850955 | Frisch 11/Ug18 (UPS) |
| <i>Cryptothecia</i> sp. 3 | Uganda | KJ850956 | Frisch 11/Ug194 (UPS) |
| <i>Cryptothecia striata</i> 1 | China | PP302048 | 20230938 (SDNU) |
| <i>Cryptothecia striata</i> 2 | China | PP302049 | 20233925 (SDNU) |
| <i>Cryptothecia subnidulans</i> 1 | Réunion | KJ850952 | v.d.Boom 40613 (hd v.d. Boom) |
| <i>Cryptothecia subnidulans</i> 2 | Guyana | KJ850953 | Joensson Guyana 6a (UPS) |
| <i>Herpothallon inopinatum</i> | Mexico | KJ850964 | Rudolphi 12 (UPS) |
| <i>Herpothallon kigeziense</i> | Uganda | KF707644 | Frisch 11/Ug26 (UPS) |
| <i>Herpothallon rubrocinctum</i> | Mexico | KF707643 | Rudolphi 5 (UPS) |
| <i>Herpothallon</i> sp. 1 | Uganda | KF707645 | Frisch 11/Ug401 (UPS) |
| <i>Herpothallon</i> sp. 2 | Brazil | OR544590 | Aptroot40528 |
| <i>Herpothallon echinatum</i> 1 | China | OQ676528 | 20220048 (SDNU) |
| <i>Herpothallon echinatum</i> 2 | China | OQ676540 | 20211610 (SDNU) |
| <i>Herpothallon fibrosum</i> 1 | China | OQ676530 | 20220088 (SDNU) |
| <i>Herpothallon fibrosum</i> 2 | China | OQ676541 | 20222323 (SDNU) |
| <i>Herpothallon fibrosum</i> 3 | China | OQ676536 | 20220374 (SDNU) |
| <i>Herpothallon glaucescens</i> | China | OQ676531 | 20220069 (SDNU) |
| <i>Herpothallon lilacinum</i> | China | OQ676532 | 20220090 (SDNU) |
| <i>Herpothallon tomentosum</i> | China | OQ676538 | 20220565 (SDNU) |
| <i>Inoderma nipponicum</i> | Japan | KP870146 | Frisch 12Jp227 (TNS) |
| <i>Leprantha cinereopruinosa</i> | Poland | MG207692 | Kukwa 17127 & Lubek (BR) |
| <i>Myriostigma candidum</i> 1 | Gabon | EU704052 | Ertz 9260 (BR) |
| <i>Myriostigma candidum</i> 2 | Uganda | KJ850959 | Frisch 11/Ug125 (UPS) |
| <i>Myriostigmaminiatum</i> | Brazil | KP843606 | Silva T2A29 (ISE—epitype) |
| <i>Pachnolepia pruinata</i> | Sweden | KJ850967 | Frisch 11/Se34 (UPS) |
| <i>Snippocia nivea</i> | France | MG207695 | Ertz 17437 (BR) |
| <i>Stirtonia neotropica</i> | Brazil | KP843611 | Cáceres & Aptroot 11112 (ISE) |
| <i>Sporodophoron gossypinum</i> | Japan | KP870154 | Frisch 12Jp186 (TNS) |
| <i>Sporodophoron primorskiense</i> | Russia | KP870157 | Ohmura 10509 (TNS) |
| <i>Tylophoron moderatum</i> | DR Congo | JF830780 | Ertz 14504 (BR) |
| <i>Chiodecton natalense</i> * | Zambia | EU704051 | Ertz 6576 (BR) |

2.2. Morphological and Chemical Analyses

The morphological and anatomical characters were examined under a stereo microscope (COIC XTL7045B2) and a polarizing microscope (Olympus CX41). The thallus and medulla were tested with K (a 10% aqueous solution of potassium hydroxide), C (a saturated solution of aqueous sodium hypochlorite), P (a saturated solution of p-phenylenediamine in 95% ethyl alcohol), I (a 3% Lugol's iodine) for identification, and H₂SO₄ (a 10% sulfuric acid solution). Polarized light (pol) was used to locate crystals in the sections. The lichen substances were identified using standardized thin-layer chromatography techniques (TLC) with systems B' and C [16]. In this study, *Lethariella cladonioides* (Nyl.) Krog. containing atranorin and norstictic acid was used as the partition standard sample. Micrographs were taken under an Olympus SZX16 research stereo microscope and a BX61 research microscope with an Olympus DP72 CCD Microscope Digital Camera.

2.3. DNA Extraction, Amplification and Sequencing

Samples for molecular genetic analysis were taken from the clean growing portions of the thallus (e.g., prothallus hyphae, pseudoisidia, pseudoisidioid structure) of specimens, and pigmented or carbonized portions were removed as far as possible. The amount of lichen used depends on the thickness of the thallus; usually, 7 mm² was used.

Genomic DNA was extracted using the Sigma-Aldrich REDEExtract-N-Amp Plant PCR Kit (St. Louis, MO, USA) following the manufacturer's instructions, except only 30 µL of extraction buffer and 30 µL dilution buffer were used.

The following primers were used for PCR amplification: mrSSU1 and mrSSU3R [17]. The 50 µL PCR mixture consisted of 2 µL DNA, 2 µL of each primer, 25 µL 2 × Taq PCR MasterMix (Taq DNA Polymerase [0.1 unit/µL]; 3 mM MgCl₂; 100 mM KCl; 0.5 mM dNTPs; and 20 mM Tris-HCl [pH 8.3]) (Tiangen, Beijing, China) and 19 µL dd H₂O. PCR cycling conditions were 94 °C for 10 min, followed by 34 cycles of 95 °C for 45 s, 50 °C for 45 s, and 72 °C for 90 s, followed by a final extension at 72 °C for 10 min. Sequencing was performed by the company BioSune Biological Technology (Shanghai, China); an ABI 3730 XL sequencer was used to sequence both strands.

2.4. Phylogenetic Analysis

The mtSSU sequences were assembled and edited using SeqMan v.7.0 (DNASTAR packages). The assembled sequences were blasted using NCBI Blast (<http://www.ncbi.nlm.nih.gov/BLAST/>, accessed on 8 February 2024) to preliminarily identify which taxa they belong to and subsequently aligned with MAFFT v7.505 using E-INS-i algorithm and normal alignment mode [18]. The final alignment contained species of *Cryptothecia*, *Herpothallon*, and other genera in the Arthoniaceae. The species *Chiodecton natalense* Nyl. was chosen as the outgroup. In addition to our newly generated sequences, other related sequences were downloaded from GenBank and mostly from Frisch et al. [19] and Thiyagaraja et al. [20] (Table 1).

Phylogenetic relationships were inferred using maximum likelihood (ML) and Bayesian inference (BI). The best substitution models were estimated using ModelFinder v2.2.0 for the subsequent ML and BI analyses [21]. In ML analysis, TVM+F+I+G4 is the best-fit model for mtSSU; in BI analysis, GTR+I+G+F is the best-fit model. The ML analyses were performed by the IQ-TREE v2.2.0 with 1000 replicates as bootstrap analysis [22]. The BI analyses were performed by MrBayes v. 3.2.6 [23], using three Markov chains running for 10 million generations for the concatenated dataset. The trees were sampled every 100 generations, and the first 25% of the trees were discarded as burn-ins. Bootstrap support (BS) ≥ 70 and posterior probability (PP) ≥ 0.95 were considered significant support values. All of the above analysis software, except SeqMan v.7.0, operate within PhyloSuite v1.2.3 [24,25].

The phylogenetic trees generated were visualized with FigTree v. 1.4.2 [26].

3. Results and Discussion

A total of five mtSSU sequences were newly generated from five specimens. We constructed ML and BI topologies based on mtSSU sequences and 32 additional sequences found on GenBank, mostly from Frisch et al. [19] and Thiagaraja et al. [20]. The phylogenetic trees obtained from ML and BI analyses exhibited consistent topologies; therefore, we present only the ML tree (Figure 1).

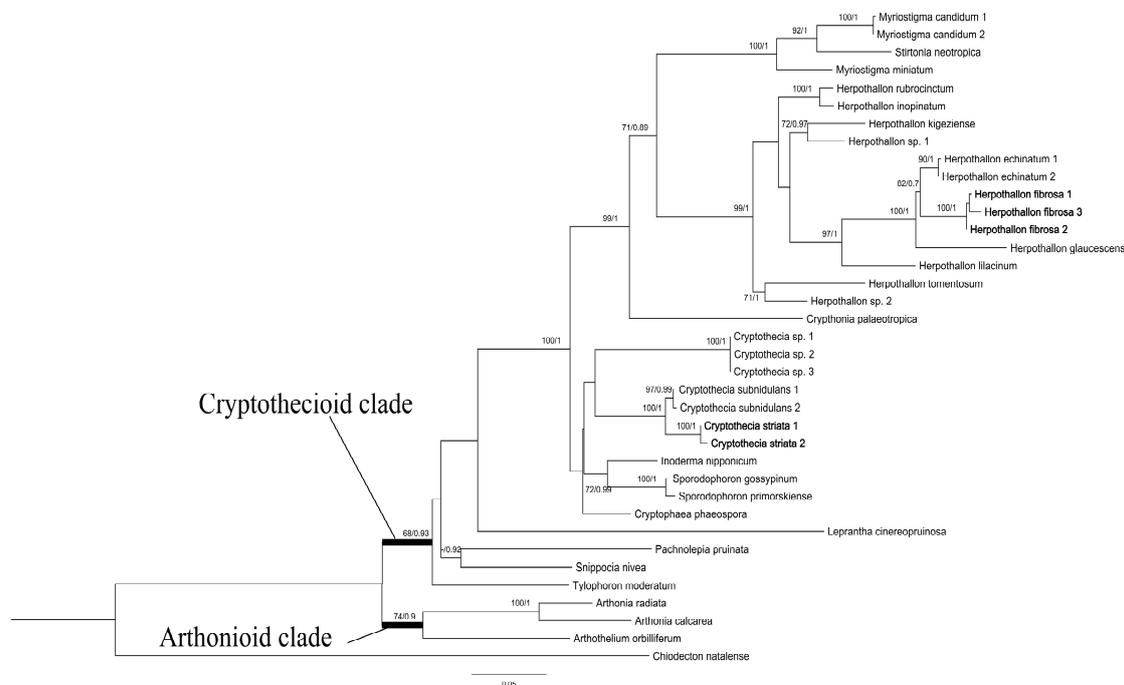


Figure 1. Phylogenetic tree generated from maximum likelihood (ML) analysis based on mtSSU sequences. ML bootstrap values (left) and Bayesian posterior probabilities (right) are indicated at the nodes. New species and records from China are shown in bold.

The topology of the phylogenetic tree, divided into the Arthonioid and Cryptothecioid clades, was largely congruent with that of earlier studies [19,20]. All species of *Herpothallon* and *Cryptothecia* belong to the Cryptothecioid clade. The newly identified *Herpothallon* specimens formed a single clade, represented by a bootstrap support of 100 and a posterior probability of 1 for the branch (Figure 1). Based on the differences in phylogeny and morphology compared to other species, which are described in detail below, it is classified as a new species named *Herpothallon fibrosum* (refer to Taxonomy below). The new record, *Cryptothecia striata*, is phylogenetically close to *C. subnidulans*, the type of the genus; they all have gray thallus and 1-spored asci, but *C. striata* exhibit radiating ridges on the thallus and contain gyrophoric acid and lecanoric acid as secondary metabolites, which distinguish it from *C. subnidulans* [9,27].

4. Taxonomy

Herpothallon fibrosum L.L. Liu & Lu L. Zhang, sp. nov. (Figure 2).

Mycobank No: 852798.

Diagnosis: This species is easily recognized by the fluffy, cylindrical pseudoisidia like a bunch of wool and the presence of both psoromic and confluent acids.

Type: China, Guizhou Province: Tongren City, Xu Jia Ba town, Zhang Jia Gou, along the stream. 27°55'33.38" N, 108°1'59.13" E, on leaves, alt. 851 m, 11 June 2022, L.L. Liu, Y.X. Bi, Z.H. Jiang & D.C. Yan 20220088 (SDNU, Holotype).

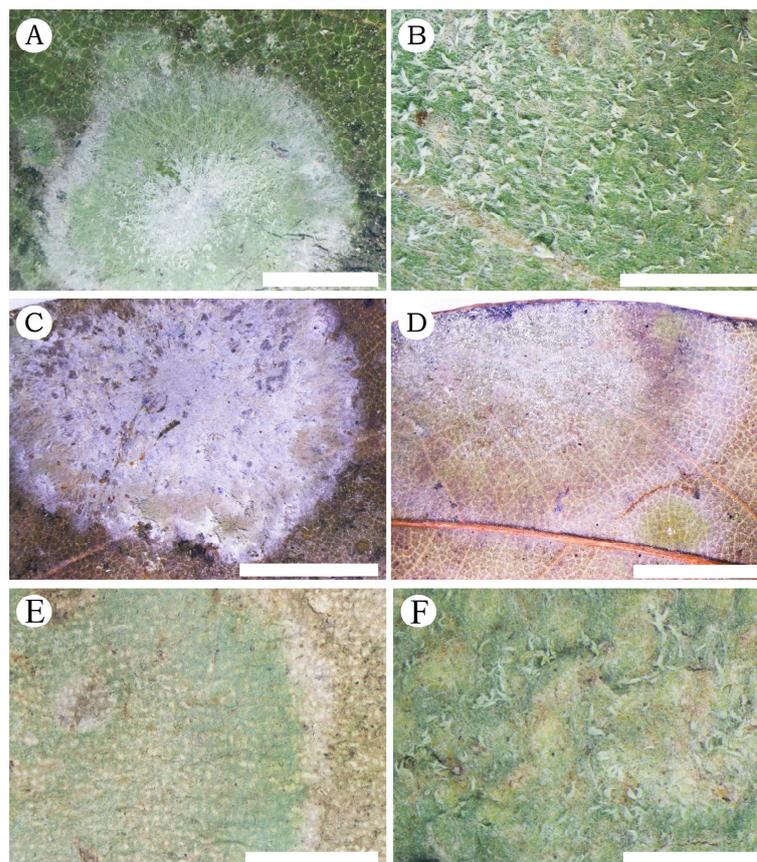


Figure 2. *Herpothallon fibrosum* (A,B) Thallus and pseudoisidia of fresh foliicolous specimen (20220088); (C,D) Thallus of foliicolous specimen after storage ((D): hypophyllous thallus) (20220088); (E,F) Thallus and pseudoisidia of fresh corticolous specimen (20220077) Scale bars: 2 mm (A); 800 μ m (B); 3 mm (C); 4 mm (D); 3 mm (E); 500 μ m (F).

Description: Thallus foliicolous, corticolous or saxicolous, up to 3 cm across, orbicular, sometimes flaking off, loosely to firmly appressed to the substrate, soft, felty to byssoid (sometimes arachnoid), dull, blue–green to greenish grey and greenish white in the inner part, rather thin, in sections up to 50 μ m thick, with many calcium oxalate crystals throughout the thallus (insoluble in KOH, forming colorless, needle-shaped crystals in 10% H₂SO₄). Hypothallus whitish, byssoid, composed of 1–3 μ m wide hyphae. Prothallus up to 3 mm broad, distinct, whitish, byssoid, composed of interwoven and radiating hyphae. Pseudoisidia are numerous, covered in the center of the thallus, loose, fluffy, cylindrical, like a bundle of fiber, sparsely branched, greenish white to whitish, 0.1–0.45 \times 0.02–0.1 mm. Photobiont *trentepohlia*, in short, irregular threads; cells yellowish green, 6–18 \times 5–10 μ m. Asci and pycnidia were not seen.

Chemistry: Thallus and prothallus K–, C–, P–, UV–, I+ orange–red in the medulla. TLC: psoromic acid (major), confluent acid (major).

Etymology: The epithet refers to the fluffy, cylindrical pseudoisidia, like a bundle of fiber.

Ecology and distribution: The new species was found growing on leaves of a tree next to a stream in Guizhou, high humidity in the growing environment.

Additional specimen examined: CHINA. Guizhou Province: Tongren City, Xu Jia Ba town, Zhang Jia Gou, along the stream, 851 m elev., 27°55′33.38″ N, 108°1′59.13″ E, on leaves, 11 June 2022, L.L. Liu, Y.X. Bi, Z.H. Jiang & D.C. Yan 20220083 (SDNU); Tongren City, Xu Jia Ba town, Zhang Jia Gou, along the stream, 851 m elev., 27°55′33.38″ N, 108°1′59.13″ E, on barks, 11 June 2022, L.L. Liu, Y.X. Bi, Z.H. Jiang & D.C. Yan 20220051, 20220077, 20220091, 20220092 (SDNU); Fujian Province: Zhangzhou City, Gongya Mountain Forest Park, near

Shenxiandong, 1224 m elev., 24°54'8.6" N, 117°25'33.76" E, on bark, 13 July 2022, L.L. Liu, J.X. Xue & L. Wang 20221040, 20221167, 20221216, 20221219, 20221259 (SDNU); Zhangzhou City, Gongya Mountain Forest Park, from Yingkemen to the "Precious *Taxus chinensis*", 1020 m elev., 24°54'28.05" N, 117°25'17.83" E, on the leaves of *Pseudosasa orthotropa* S.L. Chen et Wen, 13 July 2022, L.L. Liu, J.X. Xue & L. Wang 20221061, 20221062; Longyan City, Dongxiao National Forest Park, 621 m elev., 24°58'8.64" N, 117°1'11.98" E, on bark of a tree, 12 July 2022, L.L. Liu, J.X. Xue & L. Wang, 20220659 (SDNU); Xiamen City, Tianzhu Mountain Forest Park, the forest near Haowangjiao, 308 m elev., 24°35'40.26" N, 117°55'45.74" E, on bark, 11 July 2022, L.L. Liu, J.X. Xue & L. Wang 20220372, 20220374 (SDNU); Zhejiang Province: Lishui City, Jingning County, Wangdongyang Nature Reserve, 1299 m elev., on bark of tree, 2 December 2020, C.G. Zhao & L.L. Zhang 20222322, 20222323, 20222324, 20222325, 20222326, 20222327 (SDNU); Guangdong Province: Chaozhou City, Raoping County, Xiyuan Mountain tea farm, 830 m elev., on bark of tree, 16 July 2022, L.L. Liu, J.X. Xue & L. Wang 20220998, 20221000 (SDNU); Guangxi Zhuang Autonomous Region: Liuzhou City, Rongshui County, Jiuwan Mountain National Nature Reserve, 359 m elev., on bark of tree, 29 December 2020, L. Hu, P.F. Chen & Y.X. Bi, 20200325, 20200329, 20200331 (SDNU); Fangchenggang City, Shiwan Mountain National Nature Reserve, 753 m elev., on leaves of tree, 28 December 2020, L.L. Zhang et.al 20200237 (SDNU); Nanning City, Daming Mountain National Nature Reserve, 1210 m elev., on bark of tree, 30 December 2020, X. Zhang, 20200053 (SDNU); Nanning City, Daming Mountain National Nature Reserve, 1210 m elev., on rock, 30 December 2020, X. Zhang, 20200504 (SDNU).

Notes: *Herpothallon fibrosum* is widely distributed in southern China and is unusual among species of *Herpothallon* in the fluffy, cylindrical pseudoisidia. The foliicolous specimens have thin thalli in sections up to 50 µm thick, usually epiphyllous, partly hypophyllous, and forming a hyphal network incompletely covering the substrate, and during the storage in the herbarium, their thallus turned beige to greyish white. The corticolous and saxicolous specimens have thick thalli in sections up to 200 µm thick.

The chemical composition of this species has not previously been reported for the genus, but the beige to greyish white thallus and similar form of pseudoisidia (20–60 µm diam.) can be seen at *Herpothallon aff. confluenticum* Aptroot & Lücking, it also produces confluentic acid, but without psoromic acid and calcium oxalate crystals, I Lugol's– and delimited by a compact brown prothallus [12]. The compact brown prothallus of *H. aff. confluenticum* mentioned in Bungartz et al. [12], it looks like there is another lichen around its thallus, so we suppose its compact brown prothallus may be caused by another lichen and thalli. *Herpothallon confluenticum* Aptroot & Lücking also contains confluentic acid as major; however, it has a dirty whitish prothallus and a whitish to brownish hypothallus, cylindrical pseudoisidia, partly branched at the tips, up to 0.6 × 0.2 mm [8].

Cryptothecia striata G. Thor, The Bryologist 32: 277 (1991) (Figure 3).

Type: USA. Florida: Sarasota Co. Myakka State Park, ca. 33 km ESE Sarasota, rather dry swamp forest, Thor 2200 (s, holotype; NY, isotype).

Description: Thallus corticolous, rarely saxicolous, up to 1–10.5 cm in diameter, ecorticate, firmly to loosely attached to the substrate, cottony, dull, greenish grey to whitish grey or pale green. Medulla white, with many calcium oxalate crystals (insoluble in KOH, forming colorless, needle-shaped crystals in 10% H₂SO₄). Hyphae 1.5–2 µm wide. Prothallus is usually distinct, thick, whitish byssoid, mainly composed of interwoven hyphae. Isidia not observed, but the thallus has granula isidia-like structures. Photobiont trentepohlioid, single or aggregated into bundles, cells yellowish green, 7–16.5 × 7–10 µm.

Ascigerous areas are generally delimited, whitish, and developing in the thallus center as white dots at first and then soon merging into distinctly radiating striae, rarely not forming striae. Asci hyaline, isolated or weakly aggregated, broadly pyriform to oblong or rarely globose, with a stalk, often covered by hyaline hyphae, 1-spored or very rarely 2-spored, 62–80 × 42–56 µm. Ascospores hyaline, ovoid to oblong, muriform, surrounded by abundant cytoplasm in the asci when mature, 54–80 × 21–42 µm. Pycnidia were not observed.

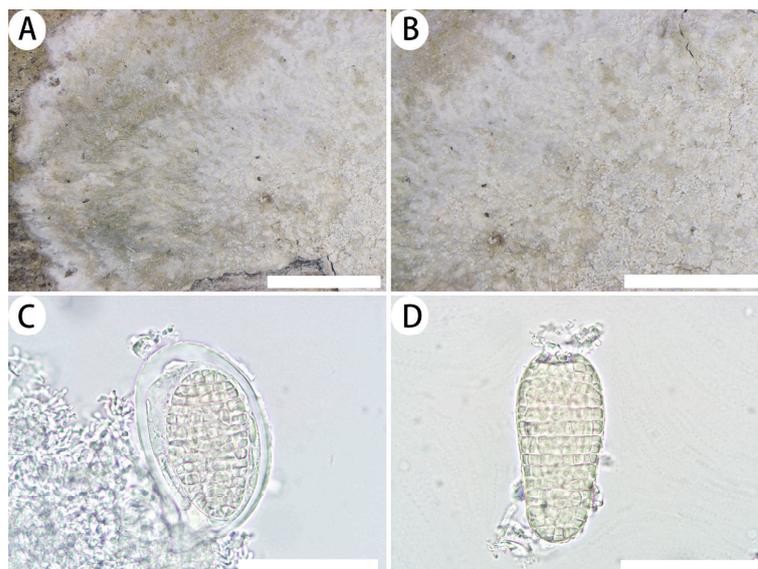


Figure 3. *Cryptothecia striata* (SDNU 20231073). (A) Thallus and prothallus. (B) Ascigerous areas. (C) Ascus. (D) Ascospores. Scale bars: 2 mm (A,B); 50 μ m (C,D).

Chemistry: Thallus C+ red, K−, P−, UV+ pale grey–white; medulla and paraphysoids I+ blue. TLC: gyrophoric acid (major), lecanoric acid, traces of atranorin.

Ecology and distribution: *Cryptothecia striata* has previously been reported in the USA, Georgia, Ecuador, and India et al. [12,27,28] and is new to China. It was found growing on the bark of trees in a humid forest in Yunnan, Guangxi Zhuang Autonomous Region, and Fujian Province.

Additional specimen examined: CHINA. Yunnan Province: Jinghong City, Mengla County, Menglun Town, Xishuangbanna Tropical Botanical Garden, 21°55′35.23″ N, 101°15′15.09″ E, alt. 512 m, on the bark of trees, 4 March 2023, L.L. Liu et al. 20230938 (SDNU); same collection data for preceding, 21°55′43.17″ N, 101°15′39.08″ E, alt. 525 m, on the bark of trees, 5 March 2023, L.L. Liu et al. 20231073 (SDNU); Xishuangbanna Dai Autonomous Prefecture, Jinghong City, Wild Elephant Valley, 22°10′17.90″ N, 100°51′39.06″ E, alt. 732 m, on the bark of trees, 6 March 2023, L.L. Liu et al. 20233925 (SDNU); Guangxi Zhuang Autonomous Region: Nanning City, Shiwan Mountain, 21°53′58.25″ N, 107°54′04.41″ E, alt. 533 m, on the bark of trees, 27 December 2020, Lu L. Zhang et al. 20200189 (SDNU); same collection data for preceding, 21°53′58.04″ N, 107°54′23.54″ E, alt. 341 m, on the bark of trees, 28 December 2020, Lu L. Zhang et al. 20200230 (SDNU); Fujian Province: Nanping City, Jianou City, Fangdao Town, Wanmulin Nature Reserve, 27°3′0.43″ N, 118°8′35.51″ E, alt. 265 m, on the bark of trees, 15 April 2023, J.X. Xue & L. Wang 20231685 (SDNU); same collection data for preceding, 27°2′51.11″ N, 118°8′34.32″ E, alt. 300 m, on the bark of trees, 14 April 2024, Lu L. Zhang et al. 20231311 (SDNU).

Notes: According to Thor, Bungartz, and Mishra et al. [12,27,28], *Cryptothecia striata* is characterized by thallus having granula isidia-like structures and ascigerous areas merging into distinctly radiating striae. The Chinese material closely matches the description in Thor, Bungartz, and Mishra et al. [12,27,28]. *Cryptothecia striata* is similar to *C. punctosorediata* Sparrius in having greenish grey to whitish grey thallus and isidia-like structures, but *C. punctosorediata* has (6–)8-spored asci and smaller ascospores (40–50 \times 15–20 μ m) [29]. In addition, *Cryptothecia striata* is similar to *C. scripta* G. Thor in having isidia-like structures, 1-spored asci and gyrophoric acid and lecanoric acid as secondary metabolites, but the latter has globose isidia-like structures, shorter asci (30–40 μ m), and shorter ascospores (19–27 μ m) [9].

Author Contributions: Conceptualization, L.Z.; methodology, J.X. and L.Z.; software, J.X. and L.Z.; validation, L.L.; formal analysis, J.X., L.L. and L.Z.; investigation, L.L., J.X. and L.Z.; data curation, J.X. and L.Z.; writing—original draft preparation, L.L. and L.Z.; writing—review and editing, L.Z. and L.L.; visualization, J.X. and L.L.; supervision, L.Z.; project administration, L.Z.; funding acquisition, L.Z. All authors have read and agreed to the published version of the manuscript.

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Conflicts of Interest: The authors declare no conflicts of interest.

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