# MYCOTAXON

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#### APRIL-JUNE 2022



*Cinereomyces wuliangshanensis* sp. nov. (Luo & Zhao— FIG. 2, p. 215)

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publication date for volume one hundred thirty-seven (1) MYCOTAXON for January–March 2022 (I–XII + 1–172) was issued on April 28, 2022

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Adustochaete yunnanensis C.L. Zhao [MB 841453], p. 266 Anapleurothecium clavatum J.S. Monteiro & R.F. Castañeda [IF557977], p. 231 Anapleurothecium leptospermi (J.A. Cooper) J.S. Monteiro & R.F. Castañeda [IF557979], p. 233 Andomyces Chuaseehar., Sri-indr. & Somrith. [MB823025], p. 252 Andomyces coronatus Chuaseehar., Sri-indr. & Somrith. [MB823026], p. 252 Chaetocapnodium zapotae L. Navarro, Salinas & Trigos [MB 837989], p. 182 Cinereomyces wuliangshanensis C.L. Zhao & K.Y. Luo [MB 836265], p. 214 Corynespora chinensis Jing W. Liu, Jian Ma, R.F. Castañeda & X.G. Zhang [IF 558664], p. 204 Ellisembia appendiculata J.W. Xia, R.Y. Liu & X.G. Zhang [MB 843925], p. 240 Erysiphe iranica Darsaraei, Khodap. & Pirnia [MB 842174] p. 276 Gangliostilbe wuzhishanensis J.W. Xia, R.Y. Liu & X.G. Zhang [MB 843815], p. 222 Nephromopsis awasthii G.K. Mishra, Nayaka & Upreti [MB 840586], p. 306 Passalora sicerariae Anu Singh, Bhartiya & P.N. Singh [MB 835143], p. 246 Podosporium simile J.S. Monteiro & R.F. Castañeda [IF557978], p. 233 Vesiculophora J.S. Monteiro & R.F. Castañeda [IF558008], p. 228 Vesiculophora diversiseptata J.S. Monteiro & R.F. Castañeda [IF558009], p. 229 Xylodon flocculosus C.L. Zhao [MB 840651], p. 194 Zasmidium sinense Jing W. Liu, Jian Ma, X.G. Zhang & R.F. Castañeda [IF 558665], p. 174

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#### Corrigenda

#### Mycotaxon 137(1)

p. 21, line 19 for: Additional examinations ... *Phaeocollybia P.* read: Additional examinations ... *P. spadicea* 

#### Corrigenda for Mycotaxon 137(2)

Cited below is an omission present in files submitted for PDF conversion in the current issue but not detected by the authors until after the paper had gone to press.

p. 311, line 23 AFTER THE LAST SENTENCE ADD: *Platismatia* is not closely related with the three other genera included in this study (Divakar et al. 2017).

#### Reviewers — volume one hundred thirty-seven (2)

The Editors express their appreciation to the following individuals who have, prior to acceptance for publication, reviewed one or more of the papers prepared for this issue.

Krishnendu Acharya D. Jayarama Bhat Uwe Braun Rafael F. Castaneda-Ruíz Maria Martha Dios Arun Kumar Dutta Daria Erastova Edit Farkas Patricia Oliveira Fiuza Yang Gao Yuehua Geng Danny Haelewaters Mei-Ling Han Hsiao-Man Ho Aamna Ishaq Sana Jabeen Makoto Kakishima Ali Keleş Munazza Kiran

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#### From the Editor-in-Chief

MYCOTAXON STYLE NUTS AND BOLTS—As this issue goes to press, we are in the process of editing a new set of instructions and manuscript templates to assist authors. In so doing, the supposedly helpful styles formatting of the previous MYCOTAXON template will be eliminated. This past year, although it became obvious that several authors had carefully prepared manuscripts using those styles, their texts arrived at the editorial desk looking nothing like what the authors had intended. There seems to be a growing incompatibility among different computer operating systems and word processing applications that is causing problems. (The fact that one manuscript left Nomenclature Editor Pennycook's computer in perfect Times New Roman but arrived on Editor-in-Chief Norvell's computer displaying several sections written entirely in the Greek alphabet alerted us that there was trouble brewing between Shaun's Times New Roman font family on his PC and Lorelei's Times font family on her Mac.)

Therefore, until we post the 2022 author guides on our MYCOTAXON website, we urge everyone to refer to the sample manuscript or a recent MYCOTAXON publication for formatting suggestions. Use the current blank templates (all of which are sized for the MYCOTAXON-size page) for all the required body-, legend-, and table-text files, but do not apply any built-in 'styles' or introduce your own author-defined character or paragraph styles to your document. Use instead the paragraph formatting menu built into your word-processing application. We hope to have the new guidelines available shortly!

FONT-FAMILY REMINDER—Remember that the only font families permitted in a Mycotaxon manuscript are the serif **TIMES** and san-serif **Arial** families. Authors should begin any manuscripts intended for Mycotaxon using only those fonts to prevent the sudden and highly dismaying resurrection of an 'alien' font at press time.

NO INTERVENING SPACE IN °C—AN EDITORIAL LABOR-SAVING DEVICE! For several years, we have advised inserting a space between the degree symbol ('o') and the temperature abbreviation ('C' or 'F'). Given the number of times scientific papers refer to temperature, it does not seem to make much sense to devote so much room to the lowly space. We feel that the degree symbol is analogous to the percent sign and that neither should be separated from its corresponding abbreviation. During editorial processing, the separate parts XII ... MYCOTAXON 137(2)

frequently fall on separate lines, requiring valuable editorial time to reunite. Henceforth, we ask all of you to join our Two-Elements-Together-Movement: 20°C and 100%.

The information above might not be of high scientific interest to readers, but your Editor-in-Chief thanks you!

The 2022 April–June MYCOTAXON offers 21 contributions by 89 authors (representing 17 countries) as revised by 38 expert reviewers and the editors.

With 13 titles, the NEW TAXA section proposes TWO new genera (*Andomyces* from Thailand & *Vesiculophora* from Brazil) and 15 species new to science representing *Adustochaete*, *Cinereomyces*, *Corynespora*, *Ellisembia*, *Gangliostilbe*, *Xylodon*, and *Zasmidium* from CHINA; *Anapleurothecium*, *Podosporium*, *Vesiculophora* from BRAZIL; *Andomyces* from THAILAND; *Chaetocapnodium* from MEXICO; *Erysiphe* from IRAN; and *Nephromopsis* and *Passalora* from INDIA. We also offer one new combination in *Anapleurothecium* from Brazil.

The NEW RANGES/HOSTS section contains six titles. New species range extensions are reported for [ascomycetes] *Elaphomyces* for TURKEY; [basidiomycetes] *Anthracoidea* for Russia and *Lactocollybia* for PAKISTAN & ASIA; [myxomycetes] *Diderma, Lamproderma, Lepidoderma, Meriderma,* and *Physarum* for the French and Spanish PYRENEES and *Fuligo & Stemonitis* for RUSSIA; AND [zygomycetes] *Coemansia* for BRAZIL & SOUTH AMERICA.

MYCOTAXON 137(2) also provides identification keys to species in all cetrarioid lichen genera (*Cetraria, Melanelia, Nephromopsis, Platismatia*) and species in India as well as to species in *Adustochaete, Anapleurothecium, Gangliostilbe*, and *Coemansia.* Papers providing conclusions supported by sequence analyses cover five new species representing *Adustochaete, Chaetocapnodium, Cinereomyces, Erysiphe* and *Xylodon* and one range extension in *Lactocollybia.* 

We also pleased to announce the posting on our MYCOBIOTA website of two new annotated species lists, which cover 1871 Indian cercosporoid fungi in INDIA and 1619 basidiomycetes collected from Grosseto Province in ITALY. Our issue concludes with book reviews of THE HIDDEN KINGDOM OF FUNGI (Seifert 2022) and THE BOLETES OF CHINA: TYLOPILUS S.L. (Chun & Yang 2021).

Warm regards,

Lorelei L. Norvell (*Editor-in-Chief*) 14 July 2022

#### 2022 Mycotaxon submission procedure

Prospective MYCOTAXON authors should download the MYCOTAXON 2022 guide, review & submission forms, and MYCOTAXON sample manuscript by clicking the 'file download page' link on our INSTRUCTIONS TO AUTHORS page before preparing their manuscript. This page briefly summarizes our '4-step' submission process.

1—PEER REVIEW: Authors first contact peer reviewers (two for journal papers; three for mycobiota/fungae) before sending them formatted text & illustration files and the appropriate 2022 MYCOTAXON journal or mycota reviewer comment form. Experts return revisions & comments to BOTH the *Editor-in-Chief* <editor@mycotaxon.com> and authors. ALL co-authors MUST correct and *proof-read* their files before submitting them to the *Nomenclature Editor*.

2—NOMENCLATURAL REVIEW: Authors email all ERROR-FREE text, tables, legends, and graphics in separate files to the *Nomenclature Editor* <PennycookS@ LandcareResearch.co.nz>. Place first author surname + genus + 'MYCOTAXON' on the subject line, and (required) attach a completed SUBMISSION FORM. The Nomenclature Editor will (i) immediately assign the accession number and (ii) after a few weeks return his notes and suggested revisions to the author(s) and *Editor-in-Chief*.

3—FINAL SUBMISSION: All coauthors thoroughly revise and proof-read files to prepare error-free text and images ready for immediate publication. Poorly formatted copy will be rejected or returned for revision. E-mail the final manuscript to the *Editor-in-Chief* <editor@mycotaxon.com>, adding the accession number to the message and all files, which include a (i) revised 2022 submission form, all (ii) text files and (iii) jpg images, and (iv) FN, IF, or MB identifier verifications for each new name or typification. The *Editor-in-Chief* acknowledges submissions within two weeks of final submission but requests authors to wait at least 14 days before sending a follow-up query (without attachments).

4—FINAL EDITORIAL REVIEW & PUBLICATION: The *Editor-in-Chief* conducts a final grammatical and scientific review and returns her editorial revisions to all expert reviewers and coauthors for final author approval. Author-approved files are placed in the publication queue.

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*Podosporium simile* sp. nov. (Monteiro & al.— FIG. 3, p. 234)

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## *Xylodon flocculosus* sp. nov. from Yunnan, China

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ABSTRACT —A new corticioid fungal species, *Xylodon flocculosus*, is described from China based on morphological and ITS+LSU sequence analyses.

KEY WORDS — Honghe county, *Hymenochaetales*, *Schizoporaceae*, taxonomy, wood-rotting fungi

#### Introduction

The corticioid genus *Xylodon* (Pers.) Gray, with *X. quercinus* (Pers.) Gray as the type, is widespread. Its species are primarily wood decomposers causing white rot of both angiosperms and gymnosperms (Eriksson & Ryvarden 1976, Yurchenko & Wu 2014), although a few *Xylodon* taxa have also been collected on brown rotten spruce stumps, palms or palm tree inflorescences, bamboo, and ferns (Burdsall & al. 1981, Langer 1994, Nordén & al. 1999, Kotiranta & Saarenoksa 2000, Boidin & Gilles 2003, Hjortstam & al. 2005, Xiong & al. 2010, Jo & al. 2019). The genus is characterized by resupinate basidiomata with hymenophores that are smooth, tuberculate, grandinioid, odontioid, coralloid, irpicoid or poroid; a monomitic hyphal system with clamp connections on generative hyphae; cystidia that are bladder-like, bottle-shaped, and capitate to subulate; suburniform basidia; and globose to ellipsoid to cylindrical basidiospores (Gray 1821, Bernicchia & Gorjón 2010). Index Fungorum (http://www.indexfungorum.org) lists 192 specific and infraspecific names in *Xylodon*, but currently accepted number of species in the genus is about 90 (Wu 1990, 2000, 2001, 2006; Xiong & al. 2009, 2010; Dai 2011, 2012; Lee & Langer 2012; Yurchenko & al. 2013; Yurchenko & Wu 2014; Zhao & al. 2014; Chen & al. 2016, 2018; Kan & al. 2017a,b; Riebesehl & Langer 2017; Wang & Chen 2017; Shi & al. 2019; Ma & Zhao 2021). About 22 *Xylodon* species have been found and described in China (Wang & Chen 2017, Shi & al. 2019, Ma & Zhao 2021).

Yurchenko & Wu (2014) placed *Xylodon* in the *Xylodon-Schizopora-Palifer* clade based on nuclear DNA sequence studies of *Hyphodontia* s.l. In their studies based on morphological and molecular analyses Riebesehl & Langer (2017) proposed 16 new *Xylodon* combinations. Riebesehl & al. (2019), who accept 77 species in the genus, synonymised *Palifer* Stalpers & P.K. Buchanan and *Odontiopsis* Hjortstam & Ryvarden with *Xylodon*.

We encountered an undescribed taxon during our research on corticioid fungi in southern China. Morphological comparisons and internal transcribed spacer (ITS) and large subunit nuclear ribosomal RNA gene (nLSU) sequence analyses place the fungus in *Xylodon*. The taxon is proposed here as *Xylodon flocculosus*.

#### **Materials & methods**

The studied specimens have been deposited at the herbarium of Southwest Forestry University, Kunming, Yunnan Province, P.R. China (SWFC). Macromorphological descriptions are based on field notes. Colour terms follow Petersen (1996). Micromorphological data were obtained from the dried specimens and observed under a light microscope following Dai (2012). The following abbreviations are used: KOH = 5% potassium hydroxide, CB = Cotton Blue, CB- = acyanophilous, IKI = Melzer's reagent, IKI- = both non-amyloid and non-dextrinoid, L = mean spore length (arithmetic average of all spores), W = mean spore width (arithmetic average of all spores), Q = variation in the L/W ratios between the specimens studied, n = number of spores measured/number of specimens.

CTAB rapid plant genome extraction kit-DN14 (Aidlab Biotechnologies Co., Ltd.) was used to obtain genomic DNA from dried specimens, according to the manufacturer's instructions. The ITS region was amplified with primer pairs ITS5

	Sample	GenBank accession no.		- 2
Species		ITS	LSU	Reference
H. halonata	Mexico	MK575207	MK598738	Yurchenko & al. 2020
H. hastata	KHL14646	MH638232	MH638232	Viner & al. 2018
L. allantosporus	FR-0249548	KY800397	KY795963	Yurchenko & al. 2017
L. mascarensis	KAS-GEL 4833	KY800399	KY795964	Yurchenko & al. 2017
L. organensis	MSK 7247	KY800403	KY795967	Yurchenko & al. 2017
L. orientalis	GEL 3400	DQ340326	DQ340353	Yurchenko & al. 2017
X. apacheriensis	Wu 0910-58	KX857797	KX857822	Chen & al. 2017
X. asper	GEL 3257	EU583424	_	Yurchenko & al. 2020
X. astrocystidiatus	Wu 9211-71	JN129972	JN129973	Yurchenko & Wu 2014
X. australis	CANB869100	MT158715	MT158751	Fernández-López & al. 2020
X. borealis	Spirin 9416	MH317760	MH638259	Viner & al. 2018
X. brevisetus	KHL 12386	DQ873612	DQ873612	Larsson & al. 2006
X. bubalinus	Cui 12888	NR_154097	_	Wang & Chen 2017
X. chinensis	Wu 1407-105	KX857804	KX857811	Chen & al. 2017
X. cystidiatus	FR-0249200	MH880195	MH884896	Riebesehl & al. 2019
X. detriticus	Zíbarová 16.05.17	MH320794	_	Viner & al. 2018
X. exilis	MSK-F 7381	MH880196	MH884897	Riebesehl & al. 2019
X. filicinus	FR-0249797	MH880201	MH884901	Riebesehl & al. 2019
X. flocculosus	CLZhao 4544	MW980775	_	Present study
	CLZhao 18342 [T]	MW980776	MW980779	Present study
	CLZhao 18379	MW980777	MW980780	Present study
	CLZhao 18394	MW980778	MW980781	Present study
X. follis	FR-0249814	MH880204	MH884902	Riebesehl & al. 2019
X. hastifer	Ryvarden 19767	KY081801	_	Riebesehl & Langer 2017
X. heterocystidiatus	Wu 9209-27	JX175045	KX857821	Chen & al. 2017
X. hyphodontinus	LIP GG- MAR15-127	MH880208	MH884906	Riebesehl & al. 2019
X. kunmingensis	CLZhao 755	MK404530	_	Shi & al. 2019
X. lenis	Wu890714-3	KY081802	_	Riebesehl & Langer 2017
X. mollis	Wu 0808-32	JX175043	KX857820	Chen & al. 2017

 TABLE 1. Hastodontia, Lyomyces, and Xylodon species and sequences used in the phylogenetic analyses.

Species	Sample	GenBank accession no.		D
		ITS	LSU	KEFERENCE
X. mollissimus	Yuan 4391	KY007518	_	Kan & al. 2017b
X. nesporii	MA:Fungi:79920	MT158717	MT158753	Fernández-López & al. 2020
X. niemelaei	Wu 1010-62	KX857799	KX857817	Chen & al. 2017
X. nongravis	GC 1412-22	KX857801	KX857818	Chen & al. 2017
X. nothofagi	PDD: 91630	GQ411524	_	Fukami & al. 2010
X. ovisporus	SFC20180822-22	MK992859	_	Lupala & al. 2019
X. paradoxus	KAS-JR06	MH880219	_	Riebesehl & al. 2019
X. pseudotropicus	Dai 10768	NR_154066	_	Zhao & al. 2014
X. quercinus	MA:Fungi:84446	MT158719	MT158755	Fernández-López & al. 2020
X. raduloides	KAS-JR26	MH880225	MH884910	Riebesehl & al. 2019
X. reticulatus	GC 1512-1	KX857808	KX857813	Chen & al. 2017
X. rhizomorphus	Dai 12354	KF917544	_	Zhao & al. 2014
X. rimosissimus	Ryberg 021031	DQ873627	_	Larsson & al. 2006
X. serpentiformis	TUB-FO 40688	MH880229	_	Riebesehl & al. 2019
X. spathulatus	MSK-F 12931	MH884914	MH880231	Riebesehl & al. 2019
X. subclavatus	TUB-FO 42167	MH880232	_	Riebesehl & al. 2019
X. subflaviporus	Wu 0809-76	KX857803	KX857815	Chen & al. 2018
X. subtropicus	Wu 9806-105	KX857807	KX857809	Chen & al. 2017
X. verecundus	KHL 12261	DQ873642	_	Larsson & al. 2006

and ITS4 (White & al. 1990), and the nLSU region was amplified with primer pairs LR0R and LR7 (http://lutzonilab.org/nuclear-ribosomal-dna). The ITS PCR protocol was: initial denaturation at 95 °C for 3 min; then 35 cycles at 94 °C for 40 s, 58 °C for 45 s, and 72 °C for 1 min; and a final extension of 72 °C for 10 min. The nLSU PCR protocol was: initial denaturation at 94 °C for 1 min; then 35 cycles at 94 °C for 30 s, 48 °C for 1 min, and 72 °C for 1.5 min; and a final extension of 72 °C for 10 min. The PCR products were purified and directly sequenced at Kunming Tsingke Biological Technology Limited Company. All newly generated sequences were deposited at GenBank (TABLE 1).

Sequencher 4.6 (https://www.genecodes.com) was used to edit the DNA sequence. Sequences were aligned using the "G-INS-I" strategy in MAFFT 7 (https://mafft.cbrc.jp/alignment/server/) and manually adjusted in BioEdit (Hall 1999). The sequence alignment was deposited in TreeBase (submission ID 28085). The outgroup *Hastodontia halonata* (J. Erikss. & Hjortstam) Hjortstam & Ryvarden

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FIG. 1. Maximum parsimony strict consensus tree illustrating the phylogeny of *Xylodon flocculosus* and related species in the residual polyporoid clade, based on ITS+nLSU sequences. Branches are labeled with maximum likelihood bootstrap >70%, parsimony bootstrap proportions >50% and Bayesian posterior probabilities >0.95.

and *H. hastata* (Litsch.) Hjortstam & Ryvarden was used to root the tree in the combined analyses following Shi & al. (2019) (FIG. 1).

Maximum parsimony (MP) analysis was applied to the combined dataset sequences. Approaches to phylogenetic analysis followed Zhao & Wu (2017), and the tree was generated in PAUP\* 4.0b10 (Swofford 2002). All characters were equally weighted, and gaps were treated as missing data. Trees were inferred using the heuristic search option with TBR branch swapping and 1000 random sequence additions. Max-trees were set to 5000, branches of zero length were collapsed, and all parsimonious trees were saved. Clade robustness was assessed using a bootstrap

(BT) analysis with 1000 replicates (Felsenstein 1985). Descriptive tree statistics tree length (TL), consistency index (CI), retention index (RI), rescaled consistency index (RC), and homoplasy index (HI) were calculated for each MP tree generated. Sequences were also analyzed using Maximum Likelihood (ML) with RAxML-HPC2 through the Cipres Science Gateway (www.phylo.org, Miller & al. 2009). Branch support (BS) for ML analysis was determined by 1000 bootstrap replicate.

MrModeltest 2.3 (Nylander 2004) was used to determine the best-fit evolution model for each data set for Bayesian Inference (BI). BI was calculated with MrBayes 3.1.2 with a general time reversible (GTR+I+G) model of DNA substitution and a gamma distribution rate variation across sites (Ronquist & Huelsenbeck 2003). Four Markov chains were run for 2 runs from random starting trees for 1.4 million generations, with trees sampled every 100 generations. The first one-fourth generations were discarded as burn-in. A majority rule consensus tree was calculated for all remaining trees. Branches were considered as significantly supported if they received a maximum likelihood bootstrap value (BS) >70%, maximum parsimony bootstrap value (BT) >50%, or Bayesian posterior probabilities (BPP) >0.95.

#### Molecular phylogeny

The combined ITS+nLSU dataset included sequences from 49 fungal specimens representing 46 taxa. The dataset had an aligned length of 2617 characters, of which 960 characters were constant, 208 parsimony-uninformative and 428 parsimony-informative. MP analysis yielded 1 equally parsimonious tree (TL = 2617, CI = 0.380, HI = 0.620, RI = 0.481, RC = 0.183). The best-fit model for ITS+nLSU alignment estimated and applied in the BI was GTR+I+G, lset nst = 6, rates = invgamma; prset statefreqpr = dirichlet (1,1,1,1). BI analysis produced a similar topology with an average standard deviation of split frequencies = 0.009310.

The phylogenetic tree (FIG. 1) inferred from ITS+nLSU sequences includes 46 *Xylodon* species. *Xylodon flocculosus* formed a well-supported distinct lineage and was sister to *X. verecundus* (G. Cunn.) Yurchenko & Riebesehl.

#### Taxonomy

*Xylodon flocculosus* C.L. Zhao, sp. nov.

Figs 2, 3

MB 840651

Differs from *Xylodon verecundus* by its grandinoid hymenial surface and strongly encrusted cystidia.

TYPE: China. Yunnan Province: Honghe, Pingbian County, Daweishan National Nature Reserve, on a fallen angiosperm branch, 3 Aug 2019, CLZhao 18342 (Holotype, SWFC 0018342; GenBank MW980776, MW980779).

ETYMOLOGY: *flocculosus* (Lat.) refers to the flocculent hymenophore of the type specimen.



FIG. 2. *Xylodon flocculosus* (holotype, SWFC 0018342). Basidiomata. Scale bars: A = 1 cm; B = 1 mm.

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BASIDIOMATA annual, resupinate, soft, cottony, without odor or taste when fresh, becoming flocculent on drying,  $\leq 10 \times 3$  cm (length × breadth),  $\leq 200 \mu$ m thick. Hymenial surface grandinoid, aculei 15–20 per mm, 50–100  $\mu$ m long, white to pale buff when fresh, turn to buff upon drying. Margin sterile, white to pale buff,  $\leq 1$  mm wide.

HYPHAL STRUCTURE monomitic; generative hyphae clamped, hyaline, more or less interwoven, thick-walled, branched, 2–3  $\mu$ m in diameter; IKI–, CB–, tissues unchanged in KOH.

HYMENIUM Cystidia numerous in the aculei, strongly encrusted at the obtuse apex,  $22-55 \times 2.9-5.6 \mu m$ ; cystidioles absent; basidia barrel-shaped, with four sterigmata, basally clamped, slightly constricted in the middle to somewhat sinuous,  $11-20 \times 3.3-4.8 \mu m$ ; basidioles dominant, in shape similar to basidia, but slightly smaller.

BASIDIOSPORES ellipsoid, hyaline, thick-walled, smooth, IKI–, CB–, (4–)4.2–5.7 (-6)  $\times$  3.1–4.4(–4.6) µm, L = 4.92 µm, W = 3.57 µm, Q = 1.35–1.42 (n = 120/4). Type of rot: white rot.

ADDITIONAL SPECIMENS EXAMINED: CHINA. YUNNAN PROVINCE. Puer: JINGDONG COUNTY, Wuliangshan National Nature Reserve, on fallen angiosperm branch, 6 Oct 2017, CLZhao 4544 (SWFC 004544; GenBank MW980775). Honghe: PINGBIAN COUNTY, Daweishan National Nature Reserve, on fallen angiosperm branch, 3 Aug 2019, CLZhao 18379 (SWFC 018379; GenBank MW980777, MW980780) CLZhao 18394 (SWFC 018394; GenBank MW980778, MW980781).

#### Discussion

Earlier morphological and molecular studies (Yurchenko & Wu 2014, Riebesehl & al. 2019) strongly supported *Xylodon* as an independent genus in the *Xylodon-Schizopora-Palifer* clade. The newly described *X. flocculosus* is nested in *Xylodon* (FIG. 1) based on the combined ITS+nLSU sequence data (BS = 100%, BT = 100%, BPP = 1). The usefulness of the ITS region alone to delimit species in *Xylodon* is approaching its phylogenetic limits. Riebesehl & al. (2019) called for additional genetic markers in *Xylodon*. The present study, based on ITS and nLSU sequences, supports *X. flocculosus* within a distinct, well-supported monophyletic lineage. In ITS phylogenetic tree, *Xylodon flocculosus* was sister to *X. verecundus*. However, morphologically *X. verecundus* differs from *X. flocculosus* in its alutaceous hymenial surface, capitate cystidia, and narrower basidiospores (5 × 3 µm; Hjortstam & Ryvarden 1997).

Morphologically, *Xylodon flocculosus* resembles *X. australis* (Berk.) Hjortstam & Ryvarden, *X. rimosissimus* (Peck) Hjortstam & Ryvarden,

Xylodon flocculosus sp. nov. (China) ... 197



FIG. 3. *Xylodon flocculosus* (holotype, SWFC 0018342). A. Basidiospores; B. Basidia and basidioles; C. Encrusted cystidia; D. Section of hymenium. Scale bars: A = 5 μm; B–D = 10 μm.

and *X. tenellus* Hjortstam & Ryvarden in sharing a grandinoid hymenial surface. *Xylodon australis* differs in having hymenophore cracked into small polygons and larger basidiospores ( $6-7.5 \times 4-4.5 \mu m$ , Riebesehl & al. 2019);

*X. rimosissimus* is distinguished by its cream to beige, slightly orange hymenial surface and capitate cystidia (Langer 1994); and *X. tenellus* differs in its smaller globose basidiospores ( $4-4.2 \times 4.2-4.5 \mu m$ , Hjortstam & Ryvarden 2007).

*Xylodon fimbriatus* (Sheng H. Wu) Hjortstam & Ryvarden, nom. illeg. [ $\equiv$  *Lyomyces fimbriatus* (Sheng H. Wu) Riebesehl & Yurchenko (Yurchenko & al. 2020)], *X. papillosus* (Fr.) Riebesehl & al., and *X. subflaviporus* C.C. Chen & Sheng H. Wu also resemble *X. flocculosus* in having strongly encrusted cystidia. However, *X. fimbriatus* is distinguished by its limy-white hymenophore and capitate cystidia (Langer 1994), *X. papillosus* by its smooth or finely odontioid hymenophore (Rattan 1977), and *X. subflaviporus* by its poroid hymenophore with cream to straw-colored hymenial surface and apically encrusted cystidia (Chen & al. 2018).

*Xylodon archeri* (Berk.) Kuntze, *X. lenis* Hjortstam & Ryvarden, and *X. tuberculatus* (Kotir. & Saaren.) Hjortstam & Ryvarden are also reported with ellipsoid basidiospores. *Xylodon archeri* differs in its coralloid basidiomata with cinnamon-buff or buckthorn brown hymenial surface and presence of the capitate cystidia (Nakasone 2012), *X. lenis* in its pinkish-tinted ochraceous hymenophore and moniliform cystidia (Langer 1994), and *X. tuberculatus* in its smooth to papillose or tuberculate hymenophore and presence of gloeocystidia (Kotiranta & Saarenoksa 2000).

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