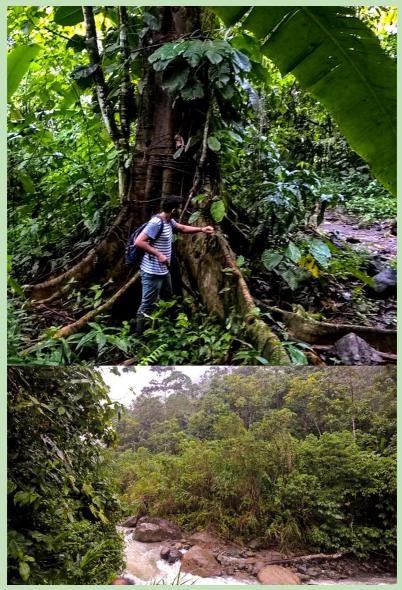
MYCOTAXON

THE INTERNATIONAL JOURNAL OF FUNGAL TAXONOMY & NOMENCLATURE

VOLUME 134 (1)

JANUARY–MARCH 2019



Type locality of *Coronosporidium ecuadorianum* gen. & sp. nov. (Sosa & al.— FIG. 1, p. 112)

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MYCOTAXON

Volume one hundred thirty-four (1) — table of contents

Nomenclatural novelties		vii	
Errata		viii	
Reviewers		ix	
2019 submission procedure		X	
From the Editor		xi	
History & Nomenclature			
North American mycology comes of ag	e:		
the 19th century participants and the	ir roles	Ronald H. Petersen	1
Ταχονομά			
Freddy Magdama, M	Ecuador ernando Espin Iarcos Vera, Si	Daynet Sosa, Noza, Lizette Serrano, Imón Pérez-Martinez, Sael F. Castañeda-Ruiz I	111
Trichoglossum tetrasporum, newly recor			
		IPAMOY CHAKRABORTY,	
Badal Kumar Da	atta, Panna D	as, Ajay Krishna Saha I	119
Perenniporia mopanshanensis sp. nov. fr	rom China		
	Chang-	-Lin Zhao & Xiang Ma	125
New records of Bilimbia and Toninia fro		Mei-jie Sun,	
		O WANG, LU-LU ZHANG	139
Anamylopsora altaica sp. nov from Nor			
	,	a Tumur, Shou-Yu Guo 🛛	147
Four Pyrenula species new to China	· ·	ig Fu, André Aptroot,	
Spadicoides matsushimae sp. nov., and	Zhong-Lian	ig Wang, Lu-Lu Zhang 🛛	155
Anisospadicoides gen. nov. for two aty	pical Spadicoi	des species Min Oiao.	
De-Wei Li, Ze-Fen Yu, KA			161
New records of Hymenoscyphus, Parasc			
<i>Scutellinia</i> for Turkey		Ali Keleş 1	169
Lemonniera yulongensis sp. nov. from Y	unnan, China	1	
, , ,		Lv, Bo Feng, Min Qiao I	177

VI ... MYCOTAXON 134(1)

Contributions to species of *Xylariales* in China—2. *Rosellinia pervariabilis* and *R. tetrastigmae* spp. nov., and a new record of *R. caudata* XIN XIE, LILI LIU, XU ZHANG, QINGDE LONG, XIANGCHUN SHEN, SARANYAPHAT BOONMEE, JICHUAN KANG, QIRUI LI 183 Catalog of *Penicillium* spp. causing blue mold on bulbs, roots, and tubers FRANK M. DUGAN & CARL A. STRAUSBAUGH 197 *Sulzbacheromyces yunnanensis*, a new record for Thailand NAKARIN SUWANNARACH, JATURONG KUMLA, KANITTA SATIENPERAKUL, WITCHAPHART SUNGPALEE, SUTHEERA HERMHUK, PIYAWAN SUTTIPRAPAN, KRIANGSAK SRI-NGERNYUANG & SAISAMORN LUMYONG 215 MYCOBIOTA (FUNGA) NEW TO THE MYCOTAXON WEBSITE A checklist of the non-gilled fleshy fungi (*Basidiomycota*)

of Kerala State, India (SUMMARY)

T.K. Arun Kumar, Anjitha Thomas, Krishnapriya Kuniyil, Salna Nanu, Vinjusha Nellipunath 221

Nomenclatural novelties and typifications proposed in Mycotaxon 134(1)

Anamylopsora altaica Ahat, A. Abbas, S.Y. Guo & Tumur [FN 570571], p. 150 Anisospadicoides R.F. Castañeda, Qiao & Z.F. Yu [MB 827953], p. 162 Anisospadicoides macrocontinua (Matsush.) R.F. Castañeda, Qiao & Z.F. Yu [MB 827954], p. 165 Anisospadicoides macroobovata (Matsush.) Qiao, Z.F. Yu & R.F. Castañeda [MB 827955], p. 165 Coronosporidium R.F. Castañeda, Quevedo & D. Sosa [IF 554557], p. 113 Coronosporidium ecuadorianum R.F. Castañeda, Quevedo & D. Sosa [IF 554558], p. 113 Lemonniera yulongensis Z.F. Yu [MB 827810], p. 178 Perenniporia mopanshanensis C.L. Zhao [MB 827580], p. 132 Rosellinia pervariabilis Q.R. Li & J.C. Kang [MB 827512], p. 188 Rosellinia tetrastigmae Q.R. Li & J.C. Kang [MB 828157], p. 190 Spadicoides matsushimae R.F. Castañeda & D.W. Li [MB 809530], p. 165

VIII ... MYCOTAXON 134(1)

Errata from previous volumes

VOLUME 133(2)

Front cover, bottom line FOR: 133(#) READ: 133(2)

VOLUME 133(4)

p. 729, line 3 (masthead) FOR: pp. 729 READ: p. 729

Back cover (online edition only):

REPLACE single bottom line by (Kaur & al.—Figs 2,3; p. 678) Maninder Kaur, Artist

Back cover (printed hard copy only): after bottom line ADD: Maninder Kaur, ARTIST

Reviewers — volume one hundred thirty-four (1)

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.

Meredith Blackwell	Troy McMullin
Rafael F. Castañeda-Ruiz	Josiane Santana Monteiro
Dinushani Anupama Daranagama	Karen K. Nakasone
Edit Farkas	Lorelei L. Norvell
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Michael Loizides	İbrahim Türkekul
Patinjareveettil Manimohan	Jan Vondrák
Eric H.C. McKenzie	Xiu-Guo Zhang

x ... Mycotaxon 134(1)

2019 Mycotaxon submission procedure

Prospective MYCOTAXON authors should download the MYCOTAXON 2019 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 2019 MYCOTAXON journal or weblist 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 & illustration 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. They email 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 2019 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.

The PDF proof and bibliographic & nomenclatural index entries are sent to all coauthors for final inspection. After PDF processing, the *Editor-in-Chief* corrects ONLY PDF editorial/conversion and index entry errors; corrections of all other errors are listed in the ERRATA of a subsequent issue for no charge. Authors will pay fees for mycobiota uploads, optional open access, and correction of major author errors to the *Business Manager* <subscriptions@mycotaxon.com> at this time.

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FROM THE EDITOR-IN-CHIEF

ON TIME! (FOR A CHANGE)—Loyal MYCOTAXON authors and readers are undoubtedly weary of the constant stream of editorial apologies following yet another issue released 1–2 months AFTER the end of a March, June, September, or December quarter. We are equally tired of apologizing to you. Thus it is with immense pleasure that we announce the release of MYCOTAXON 134(1) in MARCH! To ensure future timely journal delivery, we [i] urge authors to send their manuscripts to the EDITOR-IN-CHIEF immediately after receiving their final review from the NOMENCLATURE EDITOR and [ii] set a June 15 closure deadline for 134(2), the 2019 April–June issue. Submissions received after that date will be scheduled for the following (July–September) issue.

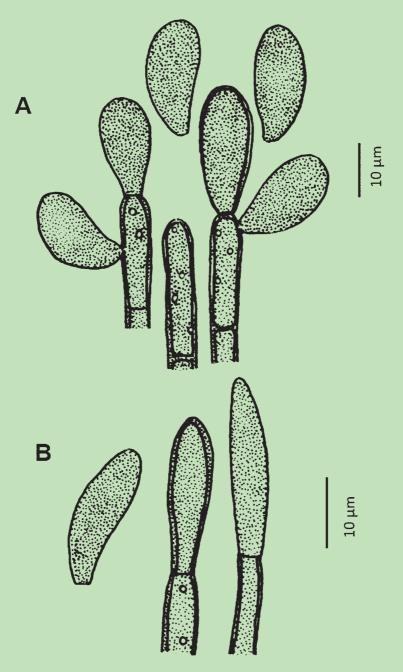
MYCOTAXON 134(1) presents 14 papers by 64 authors representing 10 countries and peer reviewed by 30 expert reviewers. The issue opens with Ronald H. Petersen's 110-page lively and welcome historical treatise on 19th century mycologists in North America. This provides us the opportunity to introduce two new journal divisions—HISTORY & NOMENCLATURE and TAXONOMY to supplant our former 'RESEARCH PAPERS' designation. ["Biodiversity & ecology" still find their home under the MYCOBIOTA/FUNGAE summaries of papers posted on the MYCOTAXON website, here announcing a checklist of fleshy non-gilled fungi from Kerala uploaded in February.]

The 2019 January–March MYCOTAXON also proposes TWO new genera (*Anisospadicoides* with two species from PERU and *Coronosporidium* from ECUADOR) and SEVEN species new to science representing *Anamylopsora*, *Lemonniera*, *Perenniporia*, and *Rosellinia* from CHINA; *Coronosporidium* from ECUADOR; and *Spadicoides* from PERU.

New species range extensions represent ascomycetes *Hymenoscyphus*, *Parascutellinia*, and *Scutellinia* for TURKEY and *Trichoglossum* for INDIA, ascolichens *Bilimbia*, *Pyrenula*, *Toninia* for mainland CHINA, and the basidiolichen *Sulzbacheromyces* for THAILAND.

Plant pathologists especially will welcome the paper covering *Penicillium* causing blue mold on bulbs, roots, and tubers that tabulates where to find references to descriptions, culture methodologies, phylogenies, and hosts for each species.

Warm regards, Lorelei Norvell (*Editor-in-Chief*) 22 March 2019



Anisospadicoides macrocontinua gen. & sp. nov. (Qiao & al.— Fig. 1, p. 163) RAFAEL F. CASTAÑEDA-RUÍZ, artist

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Perenniporia mopanshanensis sp. nov. from China

Chang-Lin Zhao^{1,2*} & Xiang Ma¹

¹ College of Biodiversity Conservation and Utilization & ² Key Laboratory for Forest Resources Conservation and Utilization in the Southwest Mountains of China, Ministry of Education, Southwest Forestry University, Kunming 650224, P.R. China

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ABSTRACT—A new poroid, white-rot, wood-inhabiting fungal species, *Perenniporia mopanshanensis*, is proposed based on morphological and molecular characters. This species from Yunnan Province, China, is characterized by resupinate basidiomes with a cream to buff to straw pore surface, dimitic hyphal system with strongly dextrinoid, unbranched, interwoven skeletal hyphae, and ellipsoid, non-truncate basidiospores ($5.5-6.5 \times 4-5 \mu m$) with hyaline, distinctly thick, smooth, strongly dextrinoid walls. Phylogenetic analyses of ITS+nLSU sequences showed that *P. mopanshanensis* is a distinct taxon in the *Perenniporia* sensu stricto clade and is sister to *P. bannaensis*.

KEY WORDS—molecular phylogenetics, *Polyporaceae*, taxonomy, *Truncospora*, wood-rotting fungi

Introduction

Perenniporia Murrill (*Polyporaceae, Polyporales*) is a large cosmopolitan genus characterized by poroid basidiomata with thick-walled, ellipsoid to distinctly truncate basidiospores that are cyanophilous and with variable dextrinoid reactions (Ryvarden 1991). The *Perenniporia* hyphal system is di- or trimitic: the generative hyphae with clamp connections and the skeletal hyphae with walls that are cyanophilous and variably dextrinoid (Decock & Stalpers 2006). About 100 species are accepted in the genus (Gilbertson & Ryvarden 1987; Hattori & Lee 1999; Núñez & Ryvarden 2001; Dai & al. 2002, 2011; Cui & al. 2007; Xiong & al. 2008; Choeyklin & al. 2009; Decock 2011, 2016; Zhao & al. 2013; Ryvarden & Melo 2014; Jang & al. 2015; Ji & al. 2017).

126 ... Zhao & Ma

Recently, phylogenetic studies of *Perenniporia* based on sequences of the internal transcribed spacer (ITS) region and the large subunit nuclear ribosomal RNA gene (nLSU) have revealed several well-supported clades that could be recognized as distinct genera (Robledo & al. 2009, Zhao & al. 2013). Robledo & al. (2009) demonstrated that *Perenniporia* was phylogenetically related to *Abundisporus* Ryvarden, *Hornodermoporus* Teixeira, *Perenniporiella* Decock & Ryvarden, and *Truncospora* Pilát, while Zhao & al. (2013) have shown that *Perenniporia* itself is polyphyletic. Several new species of *Perenniporia* have been described based on ribosomal DNA sequences (Zhao & Cui 2013, Jang & al. 2015, Ji & al. 2017)

During research on polypore diversity in southern China, an undescribed species of *Perenniporia* was found. We present morphological and molecular phylogenetic evidence that support the recognition of *P. mopanshanensis* as a new species.

Materials & methods

The specimens studied are deposited at the herbarium of Southwest Forestry University, Kunming, China (SWFC). Basidiomatal descriptions are based on field notes. Special colour terms follow Petersen (1996). Anatomical observations were obtained from the dried specimens and made using a light microscope following Dai (2010). The following abbreviations were used: KOH = 5% potassium hydroxide, CB = cotton blue, CB – a cyanophilous, IKI = Melzer's reagent, IKI – = both inamyloid 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 (a/b) = number of spores (a) measured from given number (b) of specimens.

The Omega EZNA HP Fungal DNA Kit was used to obtain genomic DNA from dried specimens according to the manufacturer's instructions with some modifications. A 30 mg sample from a dried fungal specimen was ground to powder with liquid nitrogen. The powder was transferred to a 1.5 ml centrifuge tube, suspended in 0.4 ml of lysis buffer, and incubated in a 65°C water bath for 60 min. After that, 0.4 ml phenol-chloroform (24:1) was added to each tube, and the suspension was shaken vigorously. After centrifugation at 13,000 rpm for 5 min, 0.3 ml supernatant was transferred to a new tube and mixed with 0.45 ml binding buffer. This mixture was transferred to an adsorbing column (AC) for centrifugation at 13,000 rpm for 0.5 min. Then, 0.5 ml inhibitor removal fluid was added in AC for a centrifugation at 12,000 rpm for 0.5 min. After washing twice with 0.5 ml washing buffer, the AC was transferred to a clean centrifuge tube, and 100 ml elution buffer was added to the middle of adsorbed film to elute the genomic DNA. ITS region was amplified with primer pairs ITS5 and ITS4 (White & al. 1990). Nuclear LSU region was amplified with primer pairs LR0R and LR7 (http://www.biology.duke.edu/fungi/mycolab/primers.htm). The ITS region

was amplified by initial denaturation at 95°C for 3 min, then 35 cycles of 94°C for 40 s, 58°C for 45 s, and 72°C for 1 min with a final extension of 72°C for 10 min. The nLSU region was amplified by 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, ending with 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).

ITS nLSU A. sclerosetosus MUCL 14138 FJ411101 FJ393868 Robledo & al. 2009 A. violaceus MUCL 35116 FJ411104 FJ393872 Robledo & al. 2009 Donkioporia expansa MUCL 35116 FJ411104 FJ393872 Robledo & al. 2013 Hornodermoporus Cui 6625 HQ876603 HQ654114 Zhao & al. 2013 MUCL 41677 FJ411092 FJ393859 Robledo & al. 2009 MUCL 41678 FJ411093 FJ393860 Robledo & al. 2009 Microporellus MUCL 45229 FJ411106 FJ393874 Robledo & al. 2009 Violaceocinerascens MUCL 45229 FJ411106 FJ393874 Robledo & al. 2013 Perenniporia africana Cui 8674 KF018120 KF018129 Present study Cui 8676 KF018120 KF018129 Present study Dai 12396 JQ001855 JQ001847 Zhao & al. 2013 P aridula Dai 12396 JQ001855 JQ001844 Zhao & al. 2013 Cui 8562 JQ291727 Zhao & al. 2013 Lao & al. 2013 Cui 8562	Species name	SAMPLE NO.	GenBank A	GenBank accession no.	
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P. medulla-panis MUCL 49581 FJ411088 FJ393876 Robledo & al. 2009 MUCL 43250 FJ411087 FJ393875 Robledo & al. 2009		Cui 5605	JN048760	JN048780	Zhao & Cui 2013
MUCL 43250 FJ411087 FJ393875 Robledo & al. 2009	P. macropora	Zhou 280	JQ861748	JQ861764	Zhao & Cui 2013
	P. medulla-panis	MUCL 49581	FJ411088	FJ393876	Robledo & al. 2009
Cui 3274 JN112792 JN112793 Zhao & al. 2013	-	MUCL 43250	FJ411087	FJ393875	Robledo & al. 2009
		Cui 3274	JN112792	JN112793	Zhao & al. 2013

TABLE 1. Species, specimens, and GenBank accession numbers of studied sequences

128 ... Zhao & Ma

P. mopanshanensis	CLZhao 5145 CLZhao 5152 CLZhao 2404 [T] CLZhao 2311	MH784912 MH784913 MH784911 MH784910	MH784916 MH784917 MH784915 MH784914	Present study Present study Present study Present study
P. nanlingensis	Cui 7620	HQ848477	HQ848486	Zhao & Cui 2013
-	Cui 7589	HQ848478	HQ848487	Zhao & Cui 2013
P. pyricola	Cui 9149	JN048762	JN048782	Zhao & Cui 2013
	Dai 10265	JN048761	JN048781	Zhao & Cui 2013
P. rhizomorpha	Dai 7248	JF706330	JF706348	Zhao & Cui 2012
	Cui 7507	HQ654107	HQ654117	Zhao & Cui 2013
P. russeimarginata	Yuan 1244	JQ861750	JQ861766	Zhao & Cui 2013
P. straminea	Cui 8718	HQ876600	JF706335	Zhao & Cui 2013
	Cui 8858	HQ654104	JF706334	Zhao & Cui 2013
P. subacida	Dai 8224	HQ876605	JF713024	Zhao & al. 2013
	Cui 3643	FJ613655	AY336753	Zhao & al. 2013
	MUCL 31402	FJ411103	AY333796	Robledo & al. 2009
P. subadusta	Cui 8459	HQ876606	HQ654113	Zhao & al. 2013
P. substraminea	Cui 10177	JQ001852	JQ001844	Zhao & al. 2013
	Cui 10191	JQ001853	JQ001845	Zhao & al. 2013
P. subtephropora	Dai 10962	JQ861752	JQ861768	Zhao & Cui 2013
	Dai 10964	JQ861753	JQ861769	Zhao & Cui 2013
P. tenuis	Wei 2783	JQ001858	JQ001848	Zhao & al. 2013
	Wei 2969	JQ001859	JQ001849	Zhao & al. 2013
P. tephropora	Cui 9029	HQ876601	JF706339	Zhao & Cui 2013
	Cui 6331	HQ848473	HQ848484	Zhao & Cui 2013
P. tibetica	Cui 9459	JF706327	JF706333	Zhao & Cui 2013
P. truncatospora	Cui 6987	JN048778	HQ654112	Zhao & Cui 2013
	Dai 5125	HQ654098	HQ848481	Zhao & Cui 2013
Perenniporiella chaquenia	MUCL 47647	FJ411083	FJ393855	Robledo & al. 2009
	MUCL 47648	FJ411084	FJ393856	Robledo & al. 2009
Pe. micropora	MUCL 43581	FJ411086	FJ393858	Robledo & al. 2009
Pe. neofulva	MUCL 45091	FJ411080	FJ393852	Robledo & al. 2009
Pe. pendula	MUCL 46034	FJ411082	FJ393853	Robledo & al. 2009
Pyrofomes demidoffii	MUCL 41034	FJ411105	FJ393873	Robledo & al. 2009
Truncospora detrita	MUCL 42649	FJ411099	FJ393866	Robledo & al. 2009
T. macrospora	Cui 8106	JX941573	JX941596	Zhao & al. 2013
T. ochroleuca	Dai 11486	HQ654105	JF706349	Zhao & al. 2013
	MUCL 39726	FJ411098	FJ393865	Robledo & al. 2009
	MUCL 39563	FJ411097	FJ393864	Robledo & al. 2009
T. ohiensis	Cui 5714	HQ654103	HQ654116	Zhao & al. 2013
	MUCL 41036	FJ411096	FJ393863	Robledo & al. 2009
Vanderbylia "delavayi"	Dai 6891	JQ861738	KF495019	Zhao & al. 2013
V. fraxinea	DP 83	AM269789	AM269853	Robledo & al. 2009
	Cui 7154	HQ654095	HQ654110	Zhao & al. 2013
	Cui 8885	HQ876611	JF706344	Zhao & al. 2013
	Cui 8871	JF706329	JF706345	Zhao & al. 2013
V. "robiniophila"	Cui 5644	HQ876609	JF706342	Zhao & al. 2013
	Cui 7144	HQ876608	JF706341	Zhao & al. 2013
	Cui 9174	HQ876610	JF706343	Zhao & al. 2013
V. vicina	MUCL 44779	FJ411095	AF518666	Robledo & al. 2009

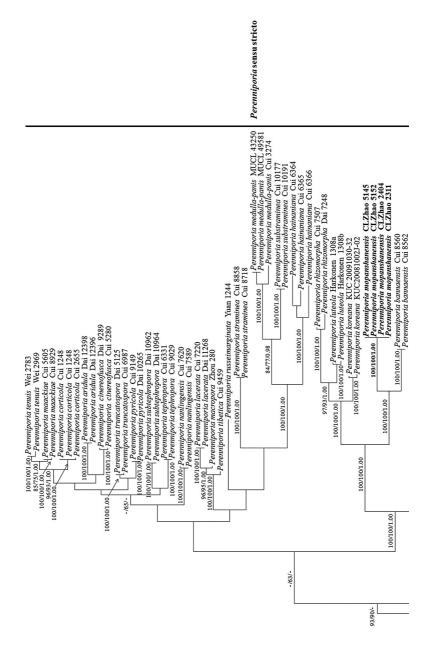
Sequencher 4.6 (GeneCodes, Ann Arbor, MI, USA) was used to edit the DNA sequence. Sequences were aligned in MAFFT 7 using the "G-INS-I" strategy (https://mafft.cbrc.jp/alignment/server/index.html) and manually adjusted in BioEdit (Hall 1999). The sequence alignment was deposited in TreeBase (submission ID 22900). *Donkioporia expansa* (Desm.) Kotl. & Pouzar and *Pyrofomes demidoffii* (Lév.) Kotl. & Pouzar were used as outgroup to root trees following Zhao & al. (2013) in the ITS+nLSU analyses.

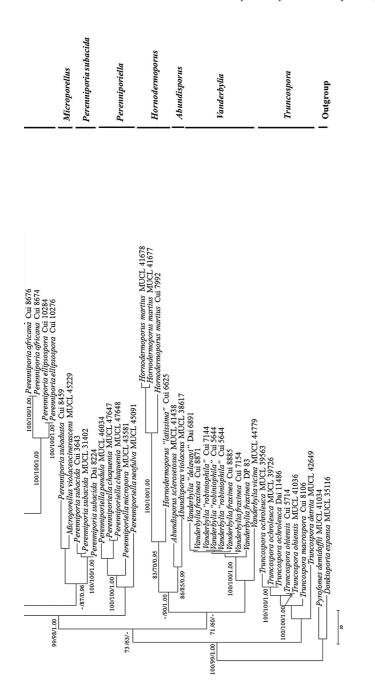
ITS+nLSU sequence analyses were performed using maximum parsimony, maximum likelihood, and Bayesian inference methods. Maximum parsimony (MP) analyses followed Song & al. (2016), and tree construction was performed in PAUP* version 4.0b10 (Swofford 2002). All characters were equally weighted, with gaps treated as missing data. Trees were inferred using the heuristic search option with TBR branch swapping and 1000 random sequence additions. Max-trees was set to 5000, branches of zero length were collapsed, and all parsimonious trees were saved. Clade robustness was assessed using bootstrap (BT) analysis with 1000 replicates (Felsenstein 1985). Tree length (TL), consistency index (CI), retention index (RI), rescaled consistency index (RC), and homoplasy index (HI) were calculated for each Maximum Parsimonious Tree (MPT) generated. Sequences were 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 replicates.

MrModeltest 2.3 (Posada & Crandall 1998, Nylander 2004) was used to determine the best-fit evolution model for each data set for Bayesian inference (BI), which was calculated with MrBayes_3.1.2 using a general time reversible (GTR) 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 7 million generations (ITS+nLSU), and trees were sampled every 100 generations. The first 25% of the generations was discarded as burn-in. A majority rule consensus tree of all remaining trees was calculated. Branches that received bootstrap support for maximum likelihood (BS), maximum parsimony (BT), and Bayesian posterior probabilities (BPP) greater than or equal to 75% (BS, BT) and 0.95 (BPP) were considered significantly supported.

Molecular phylogeny

The ITS+nLSU dataset included sequences from 86 fungal specimens representing 46 species (TABLE 1). The dataset had an aligned length of 2100 characters, of which 1553 characters were constant, 86 variable and parsimony-uninformative, and 461 parsimony-informative. Maximum parsimony analysis yielded four equally parsimonious trees (TL = 1925, CI = 0.375, HI = 0.525, RI = 0.745, RC = 0.275). Best model for the ITS+nLSU dataset estimated and applied in the Bayesian analysis: GTR+I+G. Bayesian analysis and ML analysis produced similar topologies to MP analysis, with an average standard deviation of split frequencies = 0.006258 (BI).







132 ... Zhao & Ma

The ITS+nLSU phylogeny places *Perenniporia mopanshanensis* in the *Perenniporia* sensu stricto clade (FIG. 1), where it forms a strongly supported monophyletic lineage (BS = 100%; BT = 100%; BPP = 1) and is sister to *P. bannaensis* B.K. Cui & C.L. Zhao.

Taxonomy

Perenniporia mopanshanensis C.L. Zhao, sp. nov.

FIGS 2, 3

МусоВанк МВ 827580

Differs from *Perenniporia luteola* by its larger pores, its smaller basidiospores, and its unbranched skeletal hyphae.

TYPE: China. Yunnan Province: Yuxi, Xinping county, Mopanshan National Forestry Park, on an angiosperm trunk, 19 August 2017, C.L. Zhao 2404 (Holotype, SWFC 002404; GenBank MH784911, MH784915).

ETYMOLOGY: The specific epithet *mopanshanensis* (Lat.) refers to the locality (Mopanshan) of the type specimen.

BASIDIOMATA perennial, resupinate, without odor or taste when fresh, becoming corky upon drying, $\leq 15 \times 6$ cm, 8 mm thick at center. Pore surface cream when fresh, cream to buff to straw upon drying; pores round, 3–5 per mm; dissepiments thin, entire. Sterile margin narrow, cream, ≤ 0.5 mm wide. Subiculum cream, thin, each layer ≤ 0.5 mm thick. Tubes cream to buff, corky, each layer ≤ 2 mm long.

HYPHAL STRUCTURE dimitic; generative hyphae with clamp connections; skeletal hyphae strong dextrinoid, CB+; tissues unchanged in KOH.

SUBICULUM generative hyphae infrequent, hyaline, thin-walled, unbranched, $2-3 \mu m$ in diam.; skeletal hyphae dominant, hyaline, thick-walled with a narrow to wide lumen, unbranched, interwoven, $2-3.5 \mu m$ in diam.

TUBE generative hyphae infrequent, hyaline, thin-walled, unbranched, 1.5–3 μ m in diam.; skeletal hyphae dominant, hyaline, thick-walled with a narrow to wide lumen, unbranched, interwoven, 2–3.5 μ m. Plenty of crystals present among hyphae. Cystidia absent, fusoid cystidioles present, hyaline, thin-walled, 13–19 × 4.5–6.5 μ m; basidia barrel-shaped to clavate, with four sterigmata and a basal clamp connection, 14–20 × 7–12 μ m; basidioles dominant, mostly pear-shaped, but slightly smaller than basidia.

BASIDIOSPORES ellipsoid, non-truncate, hyaline, distinctly thick-walled, smooth, strong dextrinoid, CB+, $(5-)5.5-6.5(-7) \times 4-5(-5.5) \mu m$, L = 6.15 μm , W = 4.6 μm , Q = 1.35-1.42 (n = 120/4).

TYPE OF ROT: white rot.

ADDITIONAL SPECIMENS EXAMINED: CHINA. YUNNAN PROVINCE. Yuxi: Xinping county, Mopanshan National Forestry Park, on the angiosperm trunk, 19 August 2017,



FIG. 2. *Perenniporia mopanshanensis* (holotype, SWFC 002404). Basidiomata. Scale bar: 3 cm.

C.L. Zhao 2311 (SWFC 002311; GenBank MH784910, MH784914); 13 January 2018, C.L. Zhao 5145 (SWFC 005145; GenBank MH784912, MH784916); C.L. Zhao 5152 (SWFC 005152; GenBank MH784913, MH784917).

Discussion

The new species, *Perenniporia mopanshanensis*, is supported by phylogenetic analyses and morphological characters. In the ITS+nLSU analyses (FIG. 1), it forms a strongly supported monophyletic lineage (BS = 100%; BT = 100%; BPP = 1) where it forms a clade with *P. bannaensis*. However, morphologically *P. bannaensis* differs from *P. mopanshanensis* by its annual basidiocarps with buff-yellow to pinkish buff pore surface and smaller pores (6–8 per mm; Zhao & al. 2013).

Morphologically, the presence of non-truncate basidiospores is shared by several other species in *Perenniporia* sensu stricto: *P. africana* Ipulet & Ryvarden, *P. ellipsospora* Ryvarden & Gilb., *P. koreana* Y. Jang & J.J. Kim, *P. luteola* B.K. Cui & C.L. Zhao, *P. rhizomorpha* B.K. Cui & al., and *P. subacida* (Peck) Donk. A morphological comparison between *P. mopanshanensis* and these six species is presented in TABLE 2.

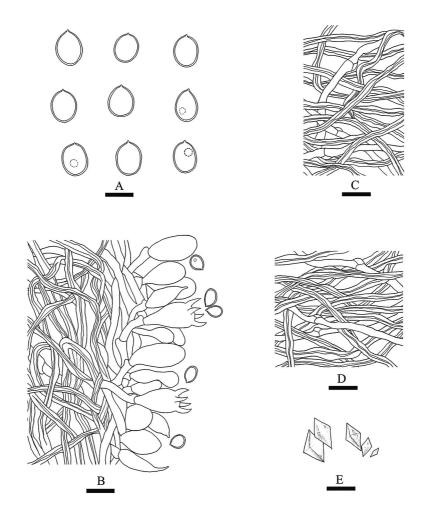


FIG. 3. *Perenniporia mopanshanensis* (holotype, SWFC 002404). A. Basidiospores; B. Section of hymenium; C. Hyphae from trama; D. Hyphae from subiculum; E. Rhombic crystal. Scale bars: $a = 5 \mu m$; $b-e = 10 \mu m$.

Polypores are an extensively studied group in *Basidiomycota* (Gilbertson & Ryvarden 1987, Núñez & Ryvarden 2001, Ryvarden & Melo 2014), but Chinese polypore diversity is still not well known, especially in the subtropics and tropics. The new species, *Perenniporia mopanshanensis*, was found in the Chinese subtropics, where many new taxa in the *Polyporales* and *Hymenochaetales* have

Species	Навіт	Pore surface	Pores / mm	Spores (µm)	Skeletal hyphae	Reference
africana	Annual	Pale orange to brown	6-8	4–5× 3–4	Branched	Ipulet & Ryvarden 2005
ellipsospora	Annual	Whitish to pale yellowish brown	3-4	4–5.5 × 3–4	Unbranched	Gilbertson & Ryvarden 1987
koreana	Annual	Grayish orange	5-6	6–7 × 3.9–5.2	Rarely branched	Jang & al. 2015
luteola	Perennial	Buff- yellow	4-6	6–7 × 5–5.5	Frequently branched	Zhao & Cui 2013
mopanshanensis	Perennial	Cream, buff to straw	3-5	5.5–6.5 × 4–5	Unbranched	This study
rhizomorpha	Annual	Yellow- buff to yellowish orange	4-6	5.3–6.5 × 4.1–5.2	Branched	Cui & al. 2007
subacida	Perennial	Ivory to yellowish	5-6	4.5–6 × 3.5–4.5	Unbranched	Decock & Stalpers 2006

TABLE 2. A comparison of Perenniporia species with non-truncate basidiospores

been described (Cui & Dai 2008; Cui & al. 2009, 2010, 2011; Du & Cui 2009; Li & Cui 2010; He & Li 2011; Jia & Cui 2011; Yu & al. 2013; Yang & He 2014; Chen & al. 2015). We anticipate that additional polypore taxa will be found in China after further investigation and molecular analysis.

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