

A new species and a new record in the *Agaricales* from Sri Lanka

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Abstract: Sri Lanka is one of the global biodiversity hotspots, and the island is characterized by rich and unique assemblage of fungi. However, the fungi of the country are poorly known, with less than 10% of the taxa having been described thus far. Studies of macrofungi are comparatively more numerous than those of microfungi due to relative importance. Nevertheless, most studies have been based only on morphological characterization of taxa. Several morpho-molecular based studies have been recently carried out, and these have yielded novel taxa and new records of macrofungi from Sri Lanka. The study reported herein focused on the macrofungi (mushrooms) in the wet zone of Sri Lanka. All fresh samples were collected from three districts (Galle, Kalutara and Matara) that represent two provinces (Southern, and Western). The samples were investigated based on both morphological and molecular analyses. The new species, *Candolleomyces ruhunensis*, is described from the family *Psathyrellaceae* and *Crepidotus striatus*, new to Sri Lanka, from *Crepidotaceae*. Sequence data from the ITS gene region were used to construct maximum likelihood (ML) and Bayesian inference analyses. The morphology of each taxon is illustrated with pictures, drawings and full descriptions.

INTRODUCTION

Agaricales is one of the best known groups of fungi to science; its taxa have been recorded as saprotrophs, symbionts and parasites. They occur in diverse habitats from the arctic to the tropics (Laala *et al.* 2018; Luna-Fontalvo *et al.* 2021). *Agaricales* are characterized by a fleshy fruiting body with an umbrella and a laminar, porous, or venous hymenophore. The order encompasses 44 families that accommodate more than 16,000 taxa (Wijayawardene *et al.* 2022; Luna-Fontalvo *et al.* 2021). *Agaricales* also include numerous macro-delicacies rich in carbohydrates, proteins and vitamins B and D which are good for human health. Several members of the genera *Pleurotus*, *Agaricus*, *Macrocybe*, and *Termitomyces* have both medicinal and edible properties (Hobbs 1996; Naeem *et al.* 2020; Kumar *et al.* 2021). On the other hand, several taxa such as *Amanita phalloides* and *Mycena polygramma* contain mushroom toxins that are highly detrimental to humans (Yagan *et al.* 2020; Luna-Fontalvo *et al.* 2021).

Several studies (Berkley & Broom 1870, 1871, 1873; Petch 1908a, b, 1910, 1913, 1915a, b, 1916a, b, c, 1917a, b, 1919, 1922, 1923, 1924a, b, c, d, 1925; Petch & Bisby 1950; Cesati 1879; Höehnel 1908, 1909, 1914) have been reported on the *Agaricales* of Sri Lanka, and these include 513 taxa from 50 genera. Species of *Agaricus*, *Entoloma*, *Hygrophorus*, *Inocybe*, *Lepiota*, *Marasmius*, and *Pleurotus* have been identified most frequently, based only on morphology (Karunarathna *et al.* 2012a, b). Most of those historical collections are deposited in the Royal Botanical Gardens, Peradeniya, Sri Lanka, and the Herbarium at Kew. Nevertheless, no reliable, detailed statistics have yet been generated on the assemblage of *Basidiomycetes* from Sri Lanka (Karunarathna *et al.* 2012).

The present paper describes a novel taxa, *Candolleomyces ruhunensis*, of *Candolleomyces*, and a new record of *Crepidotus striatus* reported from the wet zone of Sri Lanka. These taxa have been investigated based on evidence from morphology and phylogenetic analyses. Both maximum likelihood (ML) and Bayesian analysis were carried out using the ITS and LSU gene regions to determine their phylogenetic placements. Macro and micro-morphological characteristics are described with the aid of images and detailed descriptions including ecology and distribution.

MATERIALS AND METHODS

Collection sites

Specimens of mushrooms were collected from different areas of the country's wet zone, which include three districts (Matara, Galle and Kalutara) of two provinces (Southern and Western). The sampling areas are located in the southwestern portion of Sri Lanka, which receives a mean annual rainfall of over 2,500 mm, with a strong contribution of the south-west monsoons. The mean annual temperature ranges from 28–33°C while the relative humidity ranges from 85–91% (Department of Meteorology of Sri Lanka 2022). The specimens were collected from home gardens, agricultural lands, bare lands, roadsides, and public places.

Sample Collection

Fresh basidiomata were collected from 2020 and 2021. Specimens were photographed in the natural habitat and after being separating from the habitat, using a Can-on XD5 digital camera. Collected basidiomata were cleaned as much as possible to remove soil or attached debris and wrapped separately in aluminum foil to prevent spore contamination, damage, and mixing up of basidiomata of different taxa. Specimens were taken to the laboratory for morphological studies.

Morphological Studies

Macro-morphological characteristics such as size, shape, and structure of the pileus and stipe were recorded. The colour terminology used for macro-morphological identification follows Kornerup & Wanscher (1978). The specimens were dried with a portable dryer at 40°C for 24–48 hours and sealed in zip-lock plastic bags containing silica gel as a desiccant to control humidity. All the herbarium specimens were deposited in the Herbarium of Kunming Institute of Botany (HKAS), Kunming, China.

Micro-morphological observations of dry specimens were carried out with free hand sections. Slides were prepared with distilled water. In addition, 3–5% KOH, Congo red, and 3–5% NH₄(aq) were used to check some

morphological characters where necessary. Morphological characters were observed and photographed using a compound light microscope (Nikon Model Eclipse Ci-s) attached to a Canon 550D digital camera. The measurements were taken with the Tarosoft (R) Image Frame Work pro-gram, while images used for figures were processed with Adobe Photoshop CS3 (Version 15.0.0, Adobe®, San Jose, CA, USA) extended version 10.0 (Adobe Systems, San Jose, CA, USA).

DNA Extraction, PCR Amplification, and Sequencing

The genomic DNA of each specimen was extracted from dried samples using a Bio-spin Fungus Genomic DNA Extraction Kit (Bioer Technology Co., Ltd., Hangzhou, P.R. China). The nuclear ribosomal internal transcribed spacer (nrITS) and Large subunit (nrLSU) regions were amplified using the primer pair ITS5/ITS4 and LROR/LR5 (White *et al.* 1990; Vilgalys & Hester 1990). The amplification process was carried out for total volume of 25 µL comprising 1.0 µL of template DNA, 9.5 µL of double-distilled water, 1.0 µL of each primer, and 12.5 µL of 2× Power Taq PCR Master Mix. The latter consisted of a premixed, ready to use solution that included 0.1 Units/ µL Taq DNA polymerase, 500 µm of dNTP mixture each (dATP, dCTP, dGTP, and dTTP), 20 mM of Tris-HCl pH 8.3, 100 mM KCl, 3 mM of MgCl₂, stabilizer, and enhancer. During the polymerase chain reaction (PCR), each sample underwent 35 cycles according to the following settings: denaturation (95°C, 30 s), annealing (52°C, 30 s), extension (72°C, 1 min), and final extension (72°C, 10 min). Amplified products were con-firmed on a 1% agarose gel electrophoresis stained with ethidium bromide. The amplified PCR fragments were sent to a commercial sequencing provider (Beijing Bai Mai Hui Kang Biological Engineering Technology Co., P.R. China). The nucleotide sequence data were deposited in GenBank.

Sequence Alignment and Phylogenetic Analyses

The Basic Local Alignment Search Tool (BLAST) of the National Center for Biotechnology Information (NCBI) was used for the preliminary identification of the sequences data. Sequences were assembled using Seqman (Swindell & Plasterer 1997). The closely related sequences for each taxon were obtained from Genbank with consideration of recent literature. Sequences were automatically generated with MAFFT v. 7 (Katoh & Standley 2013) and terminal ends of sequences and ambiguous regions were trimmed manually using BioEdit v. 7.0.5.2 (Hall 1999) and excluded from the dataset. The phylogenetic web tool “ALTER” was used to convert sequence alignment from FASTA to PHYLIP for RAxML analysis and from FASTA to NEXUS format for Bayesian analysis (Glez-Peña *et al.* 2010). Maximum likelihood (ML) analysis was performed using RAxML-HPC2 on XSEDE (8.2.10) in CIPRES Science Gateway V. 3.3 with 1000 separate runs using the GTR+I+G model of evolution iterations. The optimal nucleotide substitution models used for Bayesian analysis were independently selected for each locus under the Akaike information criterion (AIC). Bayesian analysis was conducted with MrBayes v. 3.1.2 (GTR+I+G model) to evaluate posterior probabilities (PP) by Markov chain Monte Carlo sampling (BMCMC). Sequences for each strain were aligned using Clustal X (Thompson *et al.* 1997). Six simultaneous Markov chains were run for 2,000,000 generations, and trees were sampled every 1000 generations, resulting in 2000 trees. The first 25% of the trees, representing the burning phase of the analyses, were discarded, while the remaining 75% of the trees were used to calculate PP in the majority rule consensus tree. The ITS sequence data were analyzed using ML and Bayesian analyses. Trees were displayed in FigTree v1.4.0 (Rambaut 2012) and then copied to Microsoft PowerPoint 2013 and converted to jpeg files using Adobe Photoshop CS3 Extended 10.0 (Adobe Systems, San Jose, CA, USA).

RESULTS

Trees of similar topologies were constructed in ML and Bayesian analyses and the ML tree was taken to present the results. The results of our phylogenetic analyses show that *C. ruhunensis* is a new species, in accordance with its morphological characteristics, supported with high bootstrap values in combined phylogram of ITS and LSU: 94% in ML and 0.96 in PP (Figure 3).

TAXONOMY

Candolleomyces ruhunensis A.N. Ediriweera, P. Voto, S.C. Karunarathna & J. Kumla sp. nov. (Figure 1-2)



Figure 1. *Candolleomyces ruhunensis* (HKAS123158, holotype): (a-c) mature basidiomes; scale bars = 0.5 cm

Typus: Sri Lanka, Southern province, Matara, premises of the University of Ruhuna; saprotrophic on soil with decaying leaf litter and woody debris in a wetland in tropical climate; 2nd January 2021, A.N. Ediriweera, Herb. HKAS123158; GenBank ITS accession number: ON685315; Index Fungorum IF 555865.

Etymology. The specific epithet derives from “Ruhuna”, where the specimen was first detected from premises of the University of Ruhuna, Sri Lanka.

Diagnosis: *Candolleomyces ruhunensis* is characterized by 2–3.5 cm wide pileus, fugacious veil, lamellae not darkening with age, elliptic to oval, spores of (6.12) 6.65–8.30 (8.51) × (4.21) 4.43–5.94 (6.48) μm , basidia of 12–22 × 5–10 μm , utriform to subcylindraceous or sublageniform cheilocystidia of 23–28.2 × 7.1–8 μm , absence of pleurocystidia.

Macroscopic characters

Basidiomata small.

Pileus (primordia not observed) 20 – 35 mm across, paraboloid to convex when young, broadly convex to campanulate at maturity, margin thin, slightly eroded or slightly plicate and striate; surface bald, smooth, deep orange (6A8) to corn yellow (4A5) or light brown (6D8), brownish grey (6F8) at centre, chocolate brown (6F4) when dried; veil of fugacious, dispersed, white (1A1) to yellowish white (1A3) flocci up to center and around the margin, also scarcely appendiculate.

Lamellae crowded, with three tiers of lamellulae (2, 4 and 5 mm), adnate, thin, 2 mm height, yellowish white (2A2), dull yellow (3B3), not darkening with age; edge equal.

Stipe 40 – 55 × 3 mm, slender, cylindrical, equal, fragile, hollow, white (1A1) or pale yellow (1A3), slightly squamulose towards the base; annulus absent.

Context very thin (2 mm or less), pale yellow (1A4) to light yellow (4A5). Odour indistinct, taste not tested.

Microscopic characters

Basidiospores (n = 40) (6.12) 6.65 – 8.30 (8.51) × (4.21) 4.43 – 5.94 (6.48) μm , on average 7.39 × 5.04 μm , Q = (1.31) 1.37 – 1.59 (1.64), on average 1.46, in face view elliptic to oval, base rounded, in side view adaxially flattened to indistinctly subphaseoliform, thick-walled, smooth, hyaline to sub-hyaline in water and in NH_4 , inamyloid, prominent with hyaline to sub-hyaline oil drops; germ pore absents or an indistinct callus.

Basidia 12.0 – 22.0 × 5.0 – 10.0 μm , clavate, 4-spored, hyaline.

Pleurocystidia absent.

Cheilocystidia 23.0 – 28.2 × 7.1 – 8.0 μm , utriform to sometimes subcylindraceous or sublageniform, apex obtuse to broadly rounded, hyaline, abundant; paracystidia small to sometimes voluminous, numerous.

Pileipellis composed of a 2–3 layers paraderm of subglobose, weakly pigmented cells 30.3 – 41.0 μm broad.

Clamp connections present.

Ecology and habitat: saprotrophic, gregarious, growing on soil mixed with decaying coconut leaf litter and woody debris in wet humus soil in a wet soil in tropical climate.

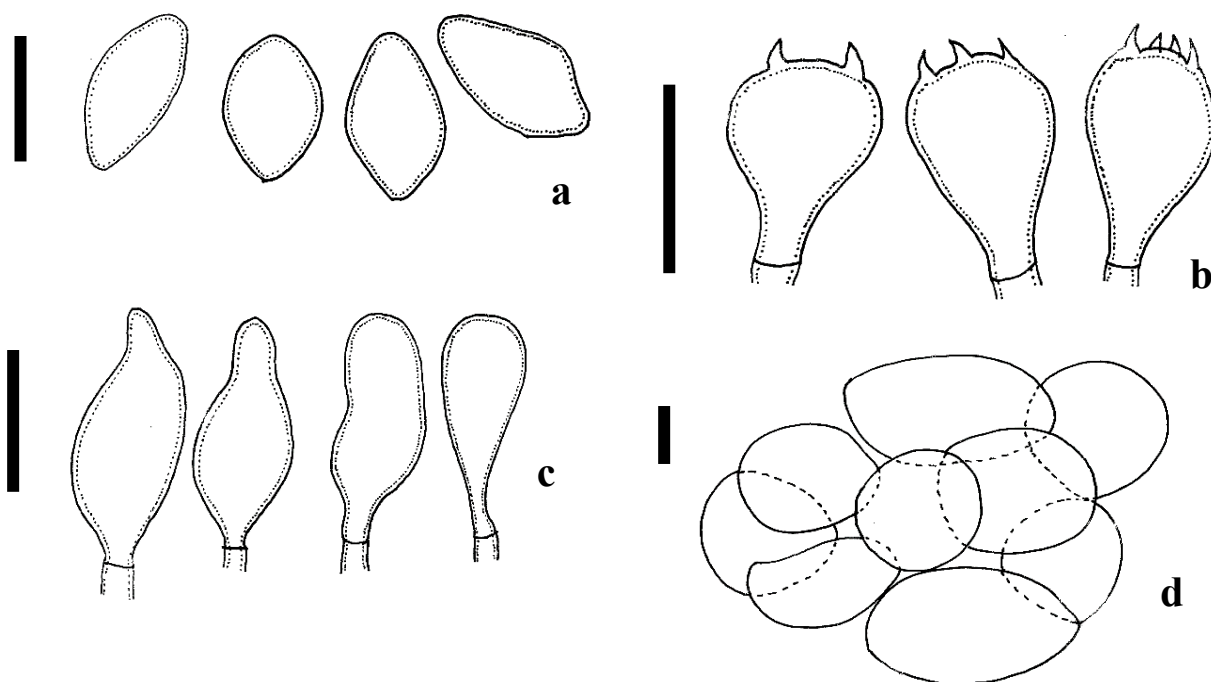


Figure 2. *Candolleomyces ruhunensis* (HKAS123158, holotype): (a) basidiospores; (b) basidia; (c) cheilocystidia; (d) pileipellis. Scale bars: (a) = 5 μm , (b) = 10 μm , (c) = 20 μm , (d) = 10 μm

NOTES

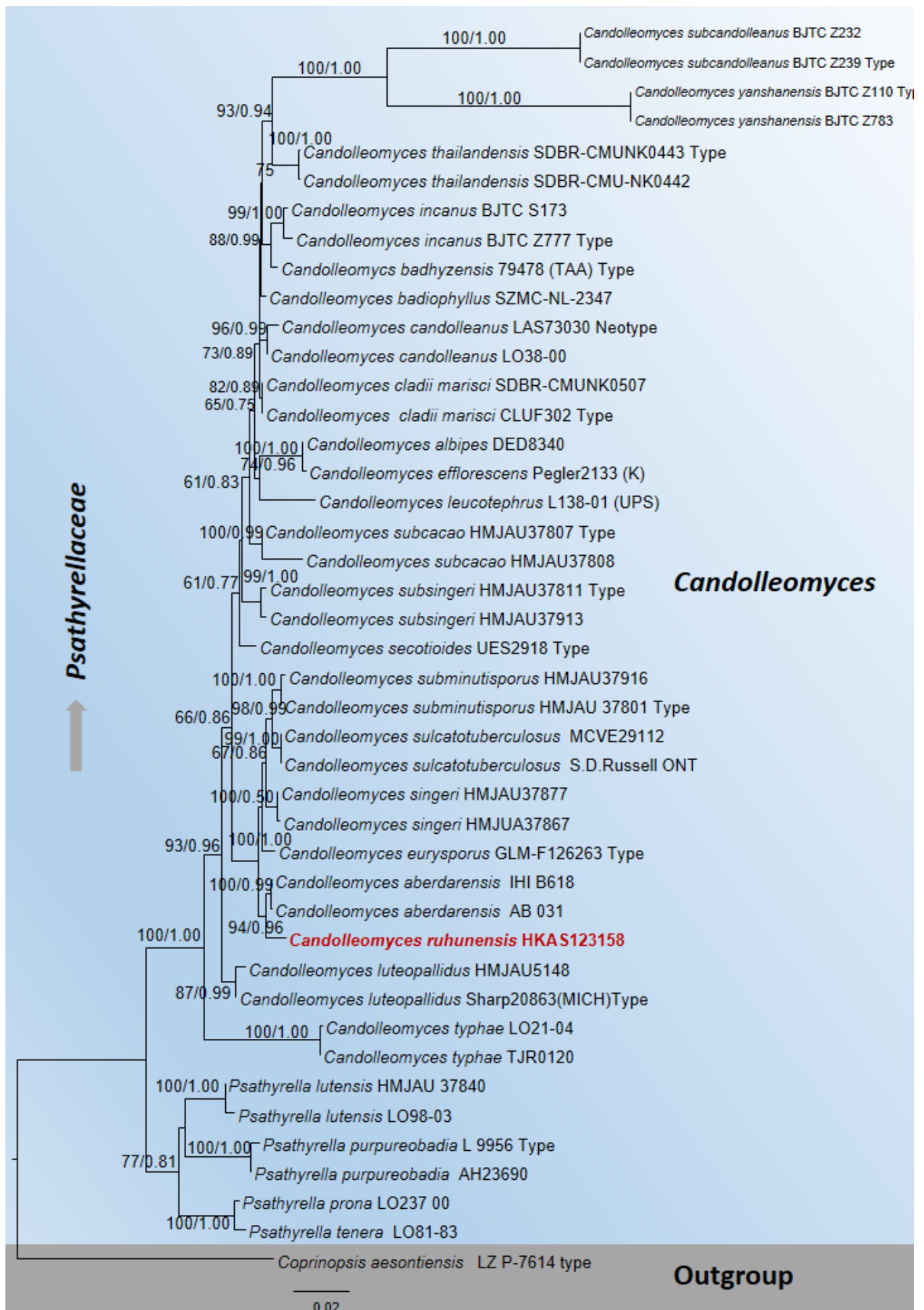
Candolleomyces ruhunensis is characterized by absence of pleurocystidia, which includes it in *Psathyrella* sect. *Spintrigerae* (Fr.) Konrad & Maubl. following a different systematics (e.g. Örstadius, Ryberg & Larsson 2015), hyaline, not phaseoliform spores mostly lacking a germ pore, distinctly utriform cheilocystidia, habitat on woody material in an equatorial climate, and a medium, not minute, habit. *Candolleomyces aberdarensis* A. Melzer, Kimani & R. Ullrich, described from Kenya (central Africa), shares several characteristics but differs in very minute habit and differently shaped spores [$7.5 - 8 (8.8) \times 4.4 - 5.0 \mu\text{m}$] having a higher quotient (1.6 – 1.75). This species is the closest taxon on a molecular basis with an identity between 97.94% – 98.09%. (see Figure 3 and Table 1).

The North American *Candolleomyces singeri* A.H. Sm. lacks truly differentiated cheilocystidia. *Candolleomyces sulcatotuberculosis* (J. Favre) Einhell, described from Europe, has a pileus distinctly striate and radially sulcate with age, and grows in moist to wet sites on debris of hygrophilous plants.

Candolleomyces cacao Desjardin & B.A. Perry and *Psathyrella atricastanea* (Murrill) A.H. Sm., from the equatorial African and American belt, have darker and ocellate pilei, and narrower spores.

The voucher *Psathyrella candolleana*, strain BRPCL, accession number MT658050, deposited by J. Xu, has a strong identity (99.27 – 99.41%, query cover 97 – 95%) with our materials. It was collected in China, associated with leaves of *Broussonetia papyrifera*, and may reveal a larger diffusion of *Candolleomyces ruhunensis* in Southeast Asia.

Figure 3 (below). RAxML analysis based on the ITS and LSU sequence data of *Candolleomyces ruhunensis*. Bootstrap support values for ML equal to or greater than 65%, and Bayesian posterior probabilities (BP) equal to or greater than 0.95 are given as ML/BP above the nodes. The tree is rooted with *Coprinopsis aesontiensis*. The newly obtained strain is indicated in red.



Species	Collection ID	ITS	LSU
<i>Candolleomyces aberdarensis</i>	AB 031	MH880928	MH880928
<i>C. aberdarensis</i>	IHI B618	MK421517	MK421517
<i>C. albipes</i>	DED8340	KX017209	
<i>C. badhyzensis</i>	79478 (TAA) Type	KC992883	KC992883
<i>C. badiophyllus</i>	SZMC-NL-2347	FN430699	FM876268
<i>C. cacao</i>	SFSU DED 8339 Type	NR148106	
<i>C. cacao</i>	FP1R4	KU847452	
<i>C. candolleanus</i>	LAS73030 Neotype	KM030175	KM030175
<i>C. candolleanus</i>	LO38-00	DQ389720	
<i>C. cladii-marisci</i>	CLUF302 Type	MK080112	
<i>C. cladii-marisci</i>	SDBR-CMUNK0507	MZ145228	
<i>C. effibulata</i>	LO37-96 Type	DQ389672	DQ389672
<i>C. efflorescens</i>	Pegler2133 (K)	KC992941	
<i>C. eurysporus</i>	GLM-F126263 Type	MT651560	MT651560
<i>C. incanus</i>	BJTC Z777 Type	ON042759	ON042766
<i>C. incanus</i>	BJTC S173	ON042760	ON042767
<i>C. leucotephrus</i>	LÖ138-01 (UPS)	KC992885	KC992885
<i>C. luteopallidus</i>	Sharp20863 (MICH) Type	KC992884	KC992884
<i>C. luteopallidus</i>	HMJAU5148	MG734736	MW301084
<i>C. ruhunensis</i>	HKAS123158	ON685315	to be submitted
<i>C. secotioides</i>	UES2918 Type	KR003281	KR003282
<i>C. singeri</i>	HMJUA37867	MG734718	MW301088
<i>C. singeri</i>	HMJAU37877	MW301073	MW301091
<i>C. subcacao</i>	HMJAU37807 Type	MW301064	MW301092
<i>C. subcacao</i>	HMJAU37808	MW301065	MW301093
<i>C. subcandolleanus</i>	BJTC Z239 Type	ON042755	ON042762
<i>C. subcandolleanus</i>	BJTC Z232	ON042756	ON042763
<i>C. subminutisporus</i>	HMJAU37801 Type	MW301066	MW301094
<i>C. subminutisporus</i>	HMJAU37916	MW301067	MW301095
<i>C. subsingeri</i>	HMJAU37811 Type	MG734715	MW301097
<i>C. subsingeri</i>	HMJAU37913	MG734725	MW301098
<i>C. sulcatotuberculosis</i>	MCVE29112	MF326002	
<i>C. sulcatotuberculosis</i>	S.D. Russell ONT	OP643311	
<i>C. trinitatensis</i>	TL9035 (C)	KC992882	KC992882
<i>C. trinitatensis</i>	ADK4162 (BR)	KC992886	KC992886
<i>C. yanshanensis</i>	BJTC Z783	ON042757	ON042764
<i>C. yanshanensis</i>	BJTC Z110 Type	ON042758	ON042765
<i>C. tuberculatus</i>	ADK3564	KC992934	KC992934
<i>C. tuberculatus</i>	Mushroom Observer # 315413	MH497604	
<i>C. subminutisporus</i>	HMJAU 37801 Type	NR_173318	
<i>C. subminutisporus</i>	X114	MK304138	
<i>C. typhae</i>	LO21-04	DQ389721	DQ389721

<i>C. typhae</i>	TJR0120	OK346575	
<i>C. thailandensis</i>	SDBR-CMU-NK0442	MZ145232	
<i>C. thailandensis</i>	SDBR-CMUNK0443 Type	MZ146874	
<i>Psathyrella lutensis</i>	LO98-03	DQ389685	DQ389685
<i>P. lutensis</i>	HMJAU 37840	MG734748	
<i>P. prona</i>	LO237 00	KJ939634	
<i>P. pseudogracilis</i>	LO172-02	DQ389675	DQ389675
<i>P. pseudogracilis</i>	CBS 166.47	MH856200	
<i>P. purpureobadia</i>	L 9956 Type	NR_119670	
<i>P. purpureobadia</i>	AH23690	KR261436	
<i>P. tenera</i>	LO81-83	KC992849	
<i>Coprinopsis aesontiensis</i>	LZ P-7614 type	NR_158339	KY554752

Table 1. Taxa information and GenBank accession numbers of ITS and LSU sequences of *Candolleomyces* specimens used in the molecular phylogenetic analysis.

Crepidotus striatus T. Bau & Y.P. Ge (Figures 4, 5)
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Figure 4. *Crepidotus striatus* (HKAS123160) (a,b): mature basidiomata in situ; scale bars = 2 cm

Macroscopic characters

Basidiomata small to medium, sessile.

Pileus fleshy, 2 – 4 mm of thickness, 10 – 32 mm across, hemispherical, pleurotoid or petaloid, occasionally with an umbo close to the attachment point; surface initially whitish (1A1-2), then butter yellow (4A5), yolk yellow (4B8) to brownish yellow (5C8), at maturity yellowish brown (5E8) close to the margin and in the middle, rust brown (6E8) at the attachment point, drying to light brown (6D8), smooth, slender, slightly pubescent, sticky, moist and slimy; margin slightly striate, plicate to sometimes lobed, sharp, concolorous with pileus surface.

Lamellae crowded, sinuate, dull yellow (3B3) to muddy brown and straw yellow (3B4) with maturity, lamellulae intermingled with 2-3 different lengths of 6, 2 and 2.2 mm, moist and fragile; edge smooth, even.

Stipe absent.

Context thin, 3 mm of thickness, whitish grey; odor distinct, taste not tested.

Microscopic characters

Basidiospores $7.5 - 8.4 \times 4.9 - 5.4 \mu\text{m}$, $Q = 1.30 - 1.55$, in front view elliptic, in side view adaxially flattened to sub-elliptic, light brown in water, thick-walled, with a guttule or yellowish to brown colour contents; germ pore visible, central, $1 - 2 \mu\text{m}$ of diam.

Basidia 4-spored and 2-spored, $16 - 22 \times 5.0 - 7.7 \mu\text{m}$, clavate to cylindraceous or subululiform, thin-walled, sub-hyaline to hyaline in water or 5% KOH, with a dark brown to light brown content.

Hymenium gelatinized. *Subhymenium* pseudoparenchymatous.

Hymenophoral trama regular, composed of thin-walled, hyaline, $2.5 - 4.0 \mu\text{m}$ wide hyphae.

Lamella edge heteromorphous with scattered cheilocystidia.

Cheilocystidia $18.0 - 34.0 \times 9.5 - 17.6 \mu\text{m}$, slender to thickset, clavate, versiform to lageniform or cylindric-clavate, fusoid, often mucronate or with a long flexuous neck, hyaline, thin-walled. *Pleurocystidia* absent.

Pileipellis composed of a gelatinized cutis with thin-walled hyphae of $13.0 - 30.0 \times 4.0 - 10.0 \mu\text{m}$, yellowish in mass in ammonia. *Pileal trama* composed of thin-walled, hyaline, $6.1 - 10.0 \mu\text{m}$ wide hyphae,

Clamp connections not found.

Ecology and habitat: saprotrophic, caespitose, growing on a decaying huge woody trunk.

Collection examined: Sri Lanka, Southern province, Galle, Kanneliya; 29th August 2021, collectors G.P Pathirana and A.N. Ediriweera; Herbarium voucher code: HKAS123160; GenBank ITS Accession Number: ON704093.

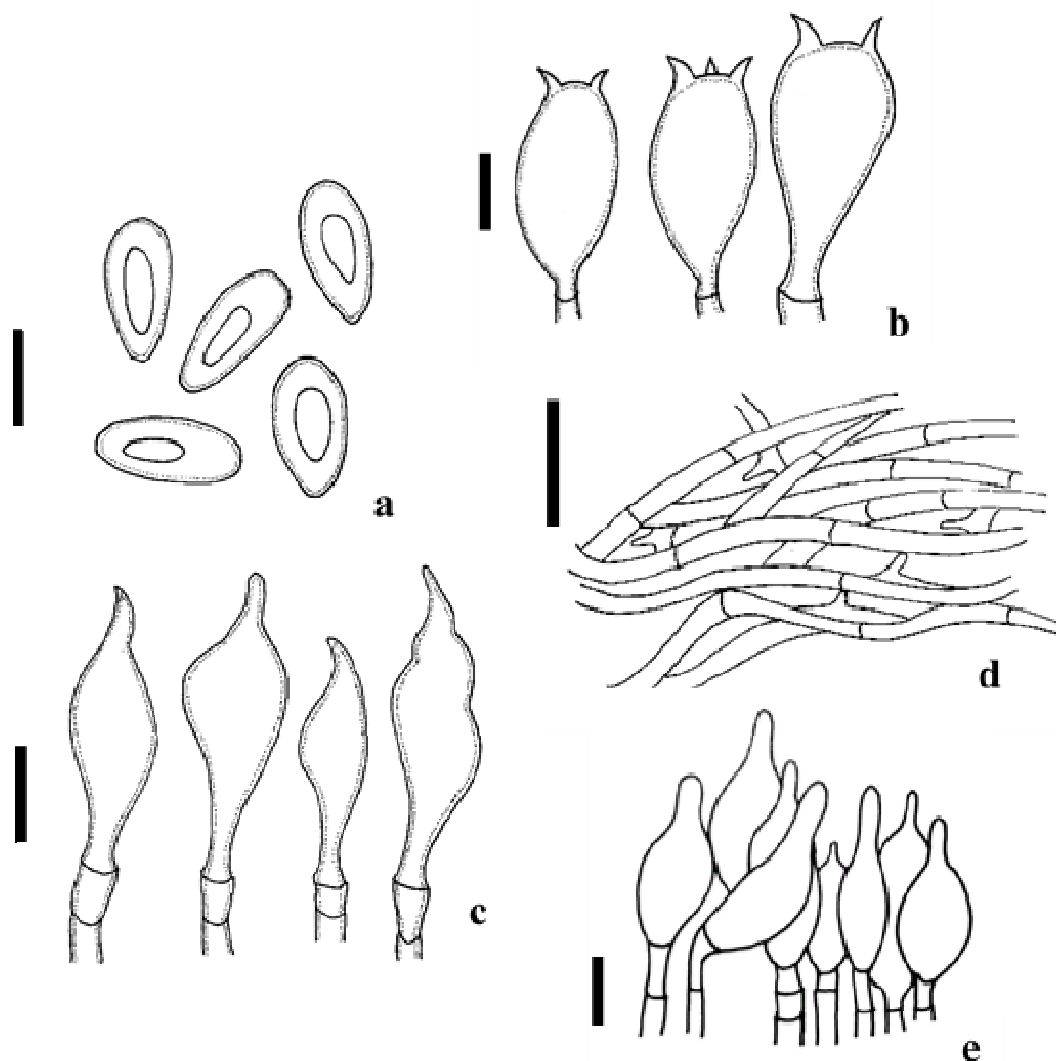
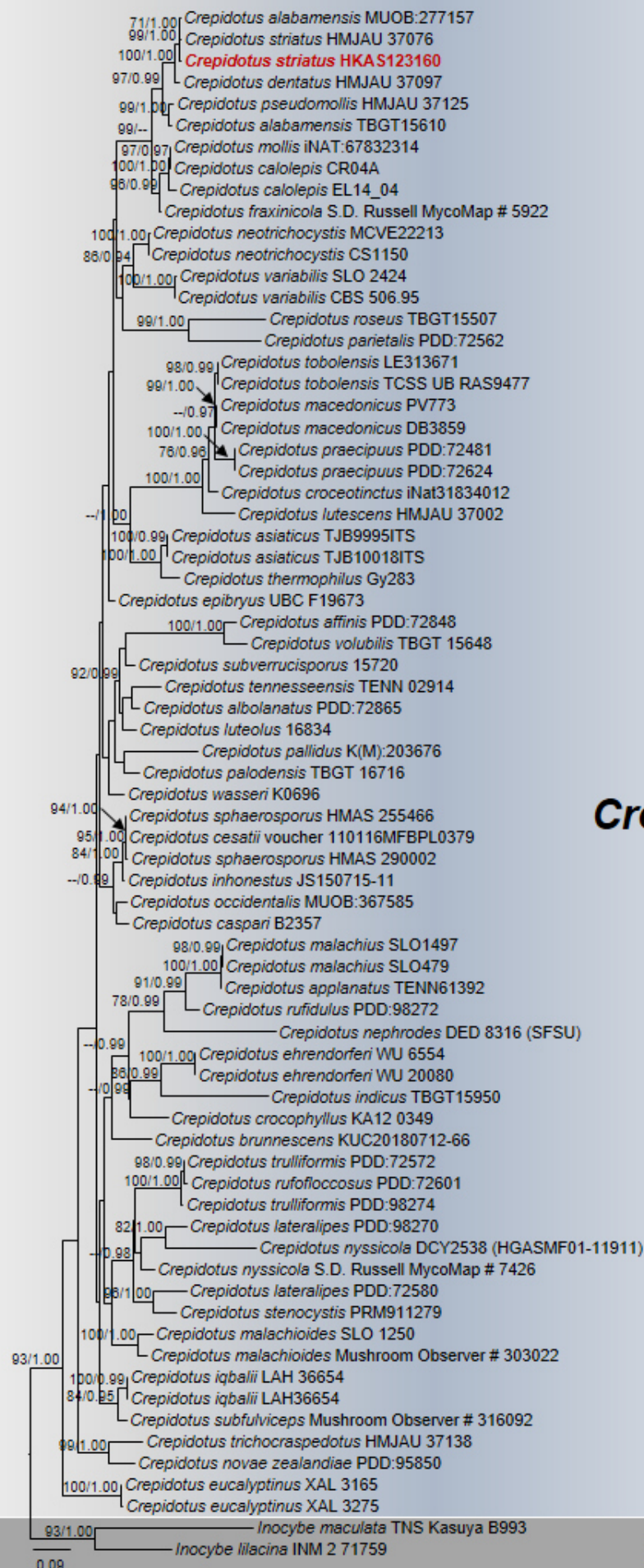


Figure 5. *Crepidotus striatus* (HKAS123160). (a) basidiospores; (b) basidia; (c) cheilocystidia; (d) pileipellis; (e) lamellae edge. Scale bars: (a) = $5 \mu\text{m}$, (b) = $5 \mu\text{m}$, (c) = $6 \mu\text{m}$, (d) = $10 \mu\text{m}$, (e) = $10 \mu\text{m}$

Crepidotaceae



Crepidotus

Outgroup

Figure 6 (above). RAxML analysis based on the ITS sequence data of *Crepidotus striatus*. Boot-strap support values for ML equal to or greater than 65%, and Bayesian posterior probabilities (BP) equal to or greater than 0.95 are given as ML/BP above the nodes. The tree is rooted with *Inocybe lilacina* and *Inocybe maculata*. The newly obtained strain is indicated in red.

NOTES

Our new collection formed a distinct clade with *C. striatus*, *C. alabamensis* Murrill and *C. dentatus* T. Bau & Y.P. Ge with an high bootstrap (98% ML) support. The base pair comparison between them and our collection revealed a 0.62 %, 0.46 % and 2.0 % difference respectively (see Figure 6 and Table 2).

Crepidotus striatus, very recently described from China (Yu-Peng & Bau 2020), shares all qualitative characters with our collection: gelatinized pileus and gills, crowded gills, a striate pileus margin, basidia also 4-spored or mainly 2-spored, a tropical to sub-tropical habitat (the type of *C. striatus* was found in Nanning City, China, on the tropic of cancer), lageniform cheilocystidia, spore shape, and structure of the pileipellis. A slight deviation in the dimensional parameters of spores, basidia and cheilocystidia falls within a normal range of intraspecific quantitative variability in a recently described and still little known species.

Table 2. Taxa information and GenBank accession numbers of ITS sequences of *Crepidotus* specimens used in the molecular phylogenetic analysis.

Species	Collection ID	ITS
<i>Crepidotus affinis</i>	PDD:72848	KY827291
<i>C. alabamensis</i>	Muob:277157	OL142283
<i>C. alabamensis</i>	TBGT15610	MK459545
<i>C. albolanatus</i>	PDD 72865 type	NR_159820
<i>C. albolanatus</i>	PDD 72865 type	KY827292
<i>C. applanatus</i>	TENN61392 clone C5	FJ596805
<i>C. applanatus</i>	TENN61392 clone C4	FJ596804
<i>C. asiaticus</i>	TJB10018ITS paratype	MF077339
<i>C. asiaticus</i>	TJB9995ITS paratype	MF077337
<i>C. brunnescens</i>	KUC20180712-66	MN197898
<i>C. calolepis</i>	CR04A	KF879617
<i>C. calolepis</i>	EL14_04	FJ904178
<i>C. caspari</i>	B2357	MW722982
<i>C. cesatii</i>	110116MFBPL0379	MW554348
<i>C. croceotinctus</i>	iNat31834012	MN498116
<i>C. crocophyllus</i>	KA12-0349	KR673424
<i>C. dentatus</i>	HMJAU 37097 type	NR_173279
<i>C. ehrendorferi</i>	WU:6554 type	MH780918
<i>C. ehrendorferi</i>	WU:20080	MH780919
<i>C. epibryus</i>	UBC F19673	HM240524
<i>C. eucalyptinus</i>	XAL 3275	KT715782
<i>C. eucalyptinus</i>	XAL 3165	KT715780
<i>C. fraxinicola</i>	S.D. Russell MycoMap # 5922	MN906146
<i>C. fraxinicola</i>	S.D. Russell MycoMap 6297	MK560109
<i>C. indicus</i>	TBGT15950 paratype	MK370662
<i>C. inhonestus</i>	JS150715-11	KX963787

<i>C. iqbalii</i>	LAH 36654 type	NR_173306
<i>C. iqbalii</i>	LAH 36654 type	MT973498
<i>C. lateralipes</i>	PDD:98270 isotype	KY827295
<i>C. luteolus</i>	16834	JF907963
<i>C. lutescens</i>	HMJAU 37002 type	NR_158400
<i>C. macedonicus</i>	PV773	MH780921
<i>C. macedonicus</i>	DB3859	MH780922
<i>C. malachioides</i>	SLO 1250 type	NR_132047
<i>C. malachioides</i>	Mushroom Observer # 303022	MK607561
<i>C. malachius</i>	SLO479	MF621033
<i>C. malachius</i>	SLO1497	MF621032
<i>C. mollis</i>	iNAT:67832314	MW676780
<i>C. neotrichocystis</i>	MCVE22213 type	MT055895
<i>C. neotrichocystis</i>	CS1150	OL672745
<i>C. nephrodes</i>	DED 8316 (SFSU)	KX017200
<i>C. novae-zelandiae</i>	PDD:95850	HQ533046
<i>C. nyssicola</i>	DCY2538 (HGASMF01-11911)	MZ413059
<i>C. occidentalis</i>	MUOB:367585	OK376745
<i>C. pallidus</i>	K(M):203676	MZ159614
<i>C. palodensis</i>	BGT16716 type	MH84489
<i>C. parietalis</i>	PDD:72562	KY827320
<i>C. praecipuus</i>	PDD:72481	KY827311
<i>C. praecipuus</i>	PDD:72624	KY827312
<i>C. pseudomollis</i>	HMJAU 37125 type	NR_173280
<i>C. roseus</i>	TBGT15507	MK567976
<i>C. rufidulus</i>	PDD 98272 type	NR_159823
<i>C. rufidulus</i>	PDD:98272 isotype	KY827319
<i>C. rufofloccosus</i>	PDD 72601 type	NR_159822
<i>C. rufofloccosus</i>	PDD:72601	KY827296
<i>C. sphaeorosporus</i>	HMAS 255466	MK966515
<i>C. sphaeorosporus</i>	HMAS 290002	MK966514
<i>C. stenocystis</i>	PRM911279	MF621030
<i>C. striatus</i>	HMJAU 37076 type	MH320742
<i>C. striatus</i>	HMJAU 37076 type	NR_173281
<i>C. striatus</i>	HKAS123160	ON704093
<i>C. subfulviceps</i>	Mushroom Observer # 316092	MH484051
<i>C. subverrucisporus</i>	MVCE15720	JF907961
<i>C. tennesseensis</i>	TENN 029144 type	NR_119720
<i>C. tennesseensis</i>	TENN 029144 type	FJ601806
<i>C. thermophilus</i>	Gy283	MF163183
<i>C. tobolensis</i>	LE313671	OL739885
<i>C. tobolensis</i>	TCSS UB RAS9477 paratype	MK522392
<i>C. trichocraspedotus</i>	HMJAU 37138 type	NR_173282

<i>C. trichocraspedotus</i>	HMJAU 37138 type	MH320743
<i>C. trulliformis</i>	PDD:72572 type	KY827297
<i>C. trulliformis</i>	PDD:98274 isotype	KY827298
<i>C. variabilis</i>	CBS 506.95	OL687124
<i>C. variabilis</i>	SLO 2424	MT055894
<i>C. volubilis</i>	TBGT15648 type	MH845231
<i>C. wasserii</i>	K0696 type	MW72298
<i>Inocybe lilacina</i>	INM 2 71759	KF680277
<i>I. maculata</i>	TNS Kasuya B993	KF680276

DISCUSSION

In the present study the new taxon *Candolleomyces ruhunensis* from *Psathyrellaceae* is introduced and a rare occurrence of the recently described *Crepidotus striatus* is reported. Based on a molecular biological evidence and on morphological differentiation (Hopple & Vilgalys 1999; Moncalvo *et al.* 2002; Matheny *et al.* 2006) *Candolleomyces* was recently separated from the genus *Psathyrella* inside which, in a different classification (Örstadius, Ryberg & Larsson 2015), it is treated as section *Spintrigerae*.

Several *Psathyrella* species have been reported from Sri Lanka which had not been identified to the species level (Ediriweera *et al.* 2014). Therefore *C. ruhunensis* is the first *Candolleomyces* species reported from Sri Lanka. Nevertheless, the highest number of *Candolleomyces* have been reported from China (Yan 2018; Bau and Yan 2021) which approximately amount to one third of the total number of *Candolleomyces* species reported worldwide.

Several species of *Crepidotus*, including *C. reversus* (Berk. & Broome) Sacc. and *C. citrinus* Petch, have been described and reported in previous studies from Sri Lanka (Pegler 1986). The report of *C. striatus* in the present study is the second ever finding of *C. striatus* after it was described from China (Yu-Peng & Bau 2020) in 2020 and first time to report from Sri Lanka.

The diversity of mushrooms in Sri Lanka is hardly explored even though the country is rich in diversity. Most familiar species with medicinal and culinary values naturally occur in home gardens, agricultural lands, forest borders and other disturbed areas, where they are known as a result of the traditional knowledge and experiences of villagers (Caron 1995; Rajapakse 2014; Hewage 2015). Thus, those species are known by their local names, which are highly synonymized based on macro-morphological characteristics and their habitats. Those locally known species are also just a handful of the island's total diversity. Hence, it is not fallacious to mention that Sri Lanka's diversity of mushrooms needs considerable more attention in order to identify its immense values and to derive the maximum output from this natural resource.

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