# Two new Clitocella species from North China revealed by phylogenetic analyses and morphological characters 

Ning Mao ${ }^{{ }^{*}}$, Jing-Chong Lv ${ }^{1^{*}}$, Yu-Yan Xu', Tao-Yu Zhao', Li Fan ${ }^{\text { }}$<br>I College of Life Science, Capital Normal University, Xisanhuanbeilu 105, Haidian, Beijing 100048, China<br>Corresponding author: Li Fan (fanli@mail.cnu.edu.cn)

Academic editor: Rui-Lin Zhao \| Received 4 January 2022 | Accepted 2 April 2022 | Published 13 April 2022
Citation: Mao N, Lv J-C, Xu Y-Y, Zhao T-Y, Fan L (2022) Two new Clitocella species from North China revealed by phylogenetic analyses and morphological characters. MycoKeys 88: 151-170. https://doi.org/10.3897/ mycokeys.88.80068


#### Abstract

Two new species of Clitocella are proposed based on morphological and phylogenetic investigations. Clitocella borealichinensis sp. nov. is closely related to C. orientalis but distinguished from the latter by its slightly smaller basidiospores and hyphae of pileipellis with pale brown to brown intracellular or parietal pigment. Clitocella colorata sp. nov. is closely related to C. popinalis and C. mundula in macromorphology but is differentiated from C. popinalis by its slightly smaller basidiospores and the difference in genetic profile, and from C. mundula by its relatively colorful pileus (white to yellowish white, grayish white to grayish brown, pink white). Phylogenetic analyses based on sequence data from five different loci (ITS, nrLSU, tef1, $r p b 2$ and $a t p 6$ ) support the taxonomic position of the two new species in the genus Clitocella. The illustrations and descriptions for the new taxa are provided.


## Keywords

Entolomataceae, multigene, phylogeny, taxonomy

## Introduction

The genus Clitocella Kluting, T.J. Baroni \& Bergemann (Entolomataceae, Agaricales), with C. popinalis (Fr.) Kluting, T.J. Baroni \& Bergemann as the type species, was established in 2014 (Kluting et al. 2014). The main characteristics of Clitocella are clitocyboid basidiomata, narrow and crowded, long-decurrent lamellae, central to eccentric stipe, thin-walled

[^0]$(<0.5 \mu \mathrm{~m})$ basidiospores with undulate pustules or minute bumps, clamp connections absent. (Baroni 1981; Kluting et al. 2014; Jian et al. 2020). Previous studies show that Clitocella is phylogenetically closely related to the genera Clitopilus (Fr. ex Rabenh.) P. Kumm. and Clitopilopsis Maire. Clitopilus differs from Clitocella in its longitudinally ridged basidiospore ornamentation, and Clitopilopsis in its basidiospores with thickened walls (0.5$0.9 \mu \mathrm{~m}$ ) and obscure irregular rounded angles of the basidiospores in polar view (Kluting et al. 2014; Baroni et al. 2020; Jian et al. 2020). There are 10 accepted species in Clitocella (Index Fungorum, http://www.Indexfungorum.org/; accessed date: 19 November 2021).

In China, the species diversity of Clitocella is scarce and only two species are described (Jian et al. 2020). Recently, several specimens of Clitocella were collected when we investigated the macrofungi in Shanxi province, North China. The morphological examination and phylogenetic analysis for these collections revealed that they represented three taxa of Clitocella, including two new species. The aim of this paper is to describe the new species and provide the DNA data to confirm the presence in China of a previously described species.

## Materials and methods

## Morphological studies

Collections were obtained and photographed in the field from Shanxi regions in China, and then dried in a fruit drier at $40-50^{\circ} \mathrm{C}$ and deposited in BJTC herbarium (Capital Normal University, Beijing, China). The sizes of basidiomata (pileal width) used in this study are as follows: small: $<30 \mathrm{~mm}$; medium-sized: $30-50 \mathrm{~mm}$; large: $>50 \mathrm{~mm}$. Standardised color values were obtained from ColorHexa (http://www. colorhexa.com/). Microscopic characters were observed in sections obtained from dry specimens mounted in $3 \% \mathrm{KOH}$, Congo Red, or Melzer's reagent (Dring 1971). For scanning electron microscopy (SEM), basidiospores were scraped from the dried gleba, placed onto double-sided tape that was mounted directly on the SEM stub, coated with platinum-palladium film of 8 nm thick using an ion-sputter coater (HITACHI E-1010), and examined with a HITACHI S-4800 SEM. The term " $[\mathrm{n} / \mathrm{m} / \mathrm{p}]$ " means n basidiospores from m basidiomata of p collections. Dimensions of basidiospores are given using the following format ' $(\mathrm{a}-\mathrm{)} \mathrm{~b}-\mathrm{c}(-\mathrm{d})$ ', where the range ' $\mathrm{b}-\mathrm{c}$ ' represents at least $90 \%$ of the measured values, and 'a' and ' d ' are the most extreme values. $\mathrm{L}_{\mathrm{m}}$ and $\mathrm{W}_{\mathrm{m}}$ indicate the average basidiospore length and width ( $\pm$ standard deviation) for the measured basidiospore, respectively. ' $Q$ ' refers to the length/width ratio of basidiospores in side-view; ' $\mathrm{Q}_{\mathrm{av}}$ ' refers to the average Q of all basidiospores $\pm$ standard deviation.

## DNA extraction, PCR amplification and DNA sequencing

Dried basidiomata were crushed by shaking for 45 s at $30 \mathrm{~Hz} 2-4$ times (Mixer Mill MM301, Retsch, Haan, Germany) in a 1.5 mL tube together with a 3 mm diam tungsten carbide ball. Total genomic DNA was extracted from the powdered basidiomata using

NuClean Plant Genomic DNA Kit (CWBIO, China), following the manufacturer's instructions. Primers ITS1F and ITS4 were employed for the ITS (White et al. 1990; Gardes and Bruns 1993), while LR0R and LR5 for nrLSU (Vilgalys and Hester 1990), EF1-983F and EF1-1953R for the tef1 (Rehner 2001), bRPB2-6F and bRPB2-7R2 for the rpb2 (Liu et al. 1999; Matheny 2005; Matheny et al. 2007), and ATP6-3 and ATP66r for the atp 6 (Kretzer and Bruns 1999; Binder and Hibbett 2003). Polymerase chain reactions (PCR) for ITS region, nrLSU region, tef1 gene, $r p b 2$ gene and atp 6 gene were performed in $25 \mu \mathrm{~L}$ reaction containing $2 \mu \mathrm{~L}$ DNA template, $1 \mu \mathrm{~L}$ primer $(10 \mu \mathrm{M})$ each, $12.5 \mu \mathrm{~L}$ of $2 \times$ Master Mix [Tiangen Biotech (Beijing) Co.], $8.5 \mu \mathrm{LddH} \mathrm{H}_{2} \mathrm{O}$.

PCR reactions were implemented as follows: an initial denaturation at $94{ }^{\circ} \mathrm{C}$ for 5 min , then to 35 cycles of the following denaturation at $94^{\circ} \mathrm{C}$ for 30 s , annealing at $52^{\circ} \mathrm{C}$ for 45 s (ITS), 60 s (nrLSU), $72{ }^{\circ} \mathrm{C}$ for 1 min ; and a final extension at $72^{\circ} \mathrm{C}$ for 10 min . Amplification of rpb2 and tef1 sequences followed Kluting et al. (2014), which entailed a touchdown protocol: an initial incubation of $94^{\circ} \mathrm{C}$ for 5 min ; 12 cycles of $94{ }^{\circ} \mathrm{C}$ for $1 \mathrm{~min}, 67^{\circ} \mathrm{C}$ for 1 min , decreasing $1^{\circ} \mathrm{C}$ each cycle, and $72^{\circ} \mathrm{C}$ for $1.5 \mathrm{~min} ; 36$ cycles of $94^{\circ} \mathrm{C}$ for $45 \mathrm{~s}, 55^{\circ} \mathrm{C}$ for 1 min , and $72^{\circ} \mathrm{C}$ for 1.5 min ; and a final extension period at $72^{\circ} \mathrm{C}$ for 7 min . Sequences of the atp 6 were amplified with a cycling protocol of $95^{\circ} \mathrm{C}$ for 5 min , followed by 40 cycles at $95^{\circ} \mathrm{C}$ for $30 \mathrm{~s}, 42^{\circ} \mathrm{C}$ for 2 min , and $72^{\circ} \mathrm{C}$ for 1 min , and a final extension at $72^{\circ} \mathrm{C}$ for 10 min . The PCR products were sent to Beijing Zhongkexilin Biotechnology Co. Ltd. for purification, sequencing, and editing. Validated sequences were deposited in the NCBI database (http://www.ncbi.nlm.nih. gov/). Other sequences of Clitocella and related species were mainly selected from those used by previous studies (Kluting et al. 2014; Vizzini et al. 2016; Baroni et al. 2020; Jian et al. 2020). The accession numbers of all sequences employed are provided in Table 1.

## Phylogenetic analyses

The combined nrLSU-rpb2-tef1-atp 6 dataset and ITS dataset were compiled to identify new species and to investigate their phylogenetic position in Clitocella. For the combined nrLSU-rpb2-tef1-atp 6 dataset, Clitopilopsis albida S.P. Jian \& Zhu L. Yang was chosen as outgroups for individual (nrLSU, rpb2, tef1, atp6) or combined analyses (Jian et al. 2020). For ITS dataset Mycena pura (Pers.) P. Kumm. was selected as outgroup taxon (Baroni et al. 2020). The sequences of each marker (ITS, nrLSU, rpb2, tef1, atp6) were independently aligned in MAFFT v.7.110 (Katoh and Standley 2013) under default parameters. Ambiguously aligned sites were identified by Gblocks v.0.91b (Castresana 2000; using default options except "Allowed Gap Positions" = half) with default parameters (For ITS: 1137, nrLSU: 180, rpb2: 611, tef1: 166, atp6: 25 position were deleted). The software BioEdit 7.0.9 (Hall 1999) was used to manually check the aligned sequences. To examine the conflict among topologies with maximum likelihood (ML), separate single-gene analyses were conducted. Sequences were then concatenated. The ITS alignment can be found on Suppl. material 5. For the combined analyses, a partitioned mixed model was used by defining the sequences of nrLSU, rpb2, tef1, and atp 6 as four independent partitions and each gene was separately estimated by different model parameters. Maximum Likelihood (ML) and Bayesian Inference (BI) analyses were conducted on the resulting concatenated dataset.

Table I. Specimens used in molecular phylogenetic studies and their GenBank accession numbers. Newly generated sequences are in bold.

| Species | Voucher | Locality | GenBank accession No. |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | ITS | nrLSU | $r p b 2$ | tef1 | atp 6 |
| Catathelasma ventricosum | DAOM221514 | USA | KP255469 | - | - | - | - |
| Clitocella colorata | BJTC FM1593 | China | OL966940 | - | - | - | - |
| Clitocella colorata | BJTC FM1594 | China | OL966941 | - | - | - | - |
| Clitocella colorata | BJTC FM1891 | China | OL966944 | OL966955 | OL989914 | OL989918 | OL989924 |
| Clitocella colorata | BJTC FM1892 | China | OL966945 | OL966956 | OL989915 | OL989919 | OL989925 |
| Clitocella colorata | BJTC FM1952 | China | - | OL966958 | OL989916 | OL989920 | OL989926 |
| Clitocella fallax | CBS 605.79 | - | AF357018 | - | - |  |  |
| Clitocella fallax | CBS 129.63 | - | AF357017 | AF223166 | EF421018 | - | - |
| Clitocella fallax | K(M): 116541 | Spain | - | - | KC816938 | KC816847 | KC816769 |
| Clitocella fallax | O-F88953 | Norway | - | - | KC816936 | KC816845 | KC816767 |
| Clitocella fallax | 25668 OKM | USA | - | - | KC816937 | KC816846 | KC816768 |
| Clitocella fallax | ME Noordeloos 1997173 | Italy | - | GQ289209 | GQ289275 | - | - |
| Clitocella fallax | ME Noordeloos 200367 | Slovakia | - | GQ289210 | GQ289276 | - | - |
| Clitocella mundula | 7161 TJB | USA | - | - | KC816952 | KC816862 | KC816782 |
| 'Clitocella mundula' | O-F19454 | Norway | - | - | KC816954 | KC816864 | KC816784 |
| Clitocella mundula | O-F71544 | Norway | - | - | KC816950 | KC816860 | KC816780 |
| 'Clitocella mundula' | AFTOL-ID 521 | USA | - | - | KC816953 | KC816863 | KC816783 |
| Clitocella mundula | 7115 TJB | USA | - | - | KC816951 | KC816861 | KC816781 |
| Clitocella mundula | K(M): 164736 | UK | - | - | KC816949 | KC816859 | KC816779 |
| 'Clitocella mundula' | K(M) : 49620 | UK | - | - | KC816948 | KC816858 | KC816778 |
| Clitocella mundula | HMJAU 7274 | China | - | MN065724 | MN148161 | MN166272 | MN133781 |
| Clitocella mundula | HMJAU 7275 | China | - | MN065723 | MN148160 | MN166271 | MN133780 |
| Clitocella mundula | HMJAU 27014 | China | - | MN065722 | MN148159 | MN166270 | MN133779 |
| Clitocella borealichinensis | BJTC FM1618 | China | OL966942 | OL966946 | OL989912 | - | OL989922 |
| Clitocella borealichinensis | BJTC FM1781 | China | OL966943 | OL966957 | OL989913 | OL989917 | OL989923 |
| Clitocella orientalis | HKAS 75548 | China | MN061333 | MN065727 | MN148164 | MN166275 | MN133784 |
| Clitocella orientalis | HKAS 75664 | China | MN061332 | MN065726 | MN148163 | MN166274 | MN133783 |
| Clitocella orientalis | HKAS 77899 | China | - | MN065725 | MN148162 | MN166273 | MN133782 |
| Clitocella orientalis | HKAS 78876 | China | MN061334 | MN065729 | MN148166 | MN166277 | MN133786 |
| Clitocella orientalis (Holotype) | HKAS 78763 | China | - | MN065728 | MN148165 | MN166276 | MN133785 |
| Clitocella orientalis | BJTC FM1539 | China | - | OL966947 | OL989911 | OL989921 | - |
| Clitocella popinalis | HBJU-550 | India | KU561066 | - | - | - | - |
| Clitocella popinalis | CBS 481.50 | UK | FJ770397 | - | - | - | - |
| Clitocella popinalis | KA12-1717 | Korea | KR673647 | - | - | - | - |
| Clitocella popinalis | RA802-3b | USA | MK217434 | - | - | - | - |
| Clitocella popinalis | $\begin{gathered} \text { Smith-2018 iNaturalist } \\ \# 17340579 \end{gathered}$ | USA | MK573922 | - | - | - | - |
| Clitocella popinalis | K(M): 143166 | UK | - | - | KC816971 | KC816878 | KC816796 |
| Clitocella popinalis | K(M): 167017 | UK | - | - | KC816972 | KC816879 | KC816797 |
| Clitocella popinalis | O-F63376 | Norway | - | - | KC816974 | KC816880 | KC816799 |
| Clitocella popinalis | 6378 TJB | Switzerland | - | - | KC816976 | KC816882 | KC816801 |
| Clitocella popinalis | O-F105360 | Norway | - | - | KC816975 | KC816881 | KC816800 |
| Clitocella popinalis | K(M): 146162 | UK | - | - | KC816970 | KC816877 | KC816795 |
| 'Clitocella popinalis' | MC2-TRENT | Italy | - | - | KC816973 | - | KC816798 |
| 'Clitocella popinalis' | ME Noordeloos 9867 | Austria | - | GQ289213 | GQ289280 | - | - |
| Clitocella popinalis | TB6378 | USA | - | AF261285 | GU384654 | - | - |
| Clitocella. Mundula | HMJAU 7275 | China | MN061331 | - | - | - | - |


| Species | Voucher | Locality | GenBank accession No. |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | ITS | nrLSU | $r p b 2$ | tef1 | atp6 |
| Clitocella obscura | MK09051302 | Czech Republic | KX271753 | - | - | - | - |
| Clitocella prunulus | G.v. Zanen F96065 | - | KC885965 | - | - | - | - |
| Clitocella_termitophila | CORT014751 | Dominican <br> Republic | - | - | MN893319 | - | - |
| Clitopilus brunneiceps <br> (Holotype) | HKAS 104510 | China | - | MN065684 | MN148123 | MN166234 | MN133737 |
| Clitopilus yunnanensis (Holotype) | HKAS 104518 | China | - | MN065698 | MN148136 | MN166247 | MN133752 |
| Clitopilus. Amarus | A. d. Haan 98031 | - | KC885963 | - | - | - | - |
| Cltopilopsis albida (Holotype) | HKAS 104519 | China | - | MN065730 | MN148167 | MN166278 | MN133787 |
| Lyophyllum decastes | Sundberg091007a | Japan | HM572548 | - | - | - | - |
| Mycena pura | CBH371 | Denmark | KF913023 | - | - | - | - |
| Rhodocybe mellea | CORT013885 | Dominican <br> Republic | MN784992 | - | - | - | - |
| Rhodocybe mellea | JBSD127402 | Dominican Republic | MN784993 | - | - | - | - |
| Rhodocybe mellea | CORT014470 | Belize | MN784994 | - | - | - | - |
| Rhodocybe mellea | NYBG815044 | Costa Rica | MN784995 | - | - | - | - |

Maximum Likelihood (ML) was performed using RAxML 8.0.14 (Stamatakis et al. 2005; Stamatakis 2006, 2014) by running 1000 bootstrap replicates under the GTRGAMMAI model (for all partitions). Bayesian Inference (BI) analysis was performed with MrBayes v3.1.2 (Ronquist and Huelsenbeck 2003) based on the best substitution models $(G T R+I+G$ for ITS, GTR +I for $n r L S U, S Y M+G$ for $r p b 2, S Y M+I+G$ for $t e f 1$, and GTR+G for $\operatorname{atp} 6$ ) determined by MrModeltest 2.3 (Nylander 2004). Two independent runs with four Markov chains were conducted for 10 M generations under the default settings. Average standard deviations of split frequency (ASDSF) values were far lower than 0.01 at the end of the runs. Trees were sampled every 100 generations after burn-in ( $25 \%$ of trees were discarded as the burn-in phase of the analyses, set up well after convergence), and a $70 \%$ majority-rule consensus tree was constructed.

Trees were visualized with TreeView32 (Page 2001). Bootstrap values (BS) $\geq 70 \%$ and Bayesian Posterior Probability values (BPP) $\geq 0.95$ were considered significant (Hillis and Bull 1993; Alfaro et al. 2003).

## Results

## Phylogenetic analysis

Twenty-eight sequences were newly generated from our six collections in this study. Two datasets, nrLSU-rpb2-tef1-atp6 combined dataset and ITS dataset were compiled to investigate the phylogenetic position of these Clitocella species. For the combined dataset, the phylogenetic trees based on individual loci (including nrLSU, rpb2, tef1, atp $\sigma$ ) showed
the almost same major clades (Suppl. material 1-4: Figs S1-S4) as that of the combined dataset (Fig. 1). There was no strongly supported conflict between single gene phylogenies, except for the nrLSU phylogeny does not resolve Clitocella mundula and C.popinalis, while the atp 6 phylogeny does not resolve C. orientalis and the new species C. colorata. So here the


Figure I. Phylogeny derived from Maximum Likelihood analysis of the combined nrLSU-rpb2-tef1-atp6 dataset of Clitocella and related genera in the family Entolomataceae. Clitopilopsis albida was employed to root the tree as an outgroup. Numbers representing likelihood bootstrap support ( $\mathrm{BS} \geq 70 \%$, left) and significant Bayesian posterior probability ( $\mathrm{BPP} \geq 0.95$, right) are indicated above the nodes. New sequences are highlighted in bold.
combined dataset was used to infer the phylogenetic placement of Clitocella species. The final combined nrLSU-rpb2-tef1-atp6 dataset contained 2963 total characters ( 905 from nrLSU, 599 from rpb2, 1010 from tef1, 449 from atp 6 , gaps included) and included 40 samples of 11 taxa. The topologies of ML and BI phylogenetic trees obtained in this study were practically the same, therefore only the tree inferred from the ML analysis is shown (Fig. 1). Except for the species Clitocella termitophila T.J. Baroni \& Angelini, members of Clitocella in the dataset formed a monophyletic lineage with strong support (MLB $=98 \%$, BPP $=1.00$ ). Clitocella termitophila was sister to all other species of Clitocella but without strong support. Of our six collections, the sequences of a collection (BJTC FM1539) grouped in the clade C. orientalis S.P. Jian \& Zhu L. Yang, indicating it is identity with this species. The remaining specimens fell in two strongly supported clades, one comprised of two collections was described as the new species $C$. borealichinensis and another comprised of three collections was described as the new species C. colorata together with a collection from USA (AFTOL-ID 521) originally labelled as C. mundula. Clitocella colorata was sister to C. orientalis with strong support, implying C. colorata is closely related to C. orientalis. Clitocella borealichinensis further clustered with C. mundula and C. popinalis (Fr.) Kluting, T.J. Baroni \& Bergemann. One collection from Norway (O-F19454), which is labelled as Clitocella mundula, formed an independent clade.

The ITS dataset comprised 27 samples of 11 taxa and 662 characters. The topology of phylogenetic trees based on the ITS dataset generated from ML and BI analyses were almost identical and only the tree inferred from the ML analysis is shown (Fig. 2). The sequences of the new species C. borealichinensis formed an independent and strong support branch, like that of multilocus phylogeny (Fig. 1), supporting it is a distinct species. The sequences of the new species C. colorata together with five sequences labelled as C. popinalis from India, South Korea, UK and USA formed an independent and strong support branch, indicating they represented a distinct species.

## Taxonomy

## Clitocella borealichinensis L. Fan \& N. Mao, sp. nov.

MycoBank No: 843689
Figs 3a, 4, 6a, b
Etymology. borealichinensis, referring to north China as the place of origin.
Holotype. China. Shanxi Province, Qinshui County, Lishan Mountain, $35^{\circ} 36.49^{\prime} \mathrm{N}, 112^{\circ} 11.7^{\prime} \mathrm{E}$, alt. $1150 \mathrm{~m}, 26$ July 2021 , on the ground in broad-leaved forest dominated by Quercus sp., N. Mao MNM340 (BJTC FM1781).

Diagnosis. Clitocella borealichinensis is characterized by its clitocyboid basidiomata, globose to subglobose, occasionally broadly ellipsoid basidiospores, the absence of hymenial cystidia and clamp connection, and usually growing in broad-leaved forests. It is most similar to C. orientalis but differs from it by the slightly smaller basidiospores, non-gelatinized hyphae of pileipellis and stipitipellis with pale brown to brown intracellular or parietal pigment.


Figure 2. Phylogeny derived from Maximum Likelihood analysis of the ITS sequences from Clitocella and related genera in the family Entolomataceae. Mycena pura was employed to root the tree as an outgroup. Numbers representing likelihood bootstrap support ( $\mathrm{BS} \geq 70 \%$, left) and significant Bayesian posterior probability ( $\mathrm{BPP} \geq 0.95$, right) are indicated above the nodes. New sequences are highlighted in bold.

Description. Basidiomata clitocyboid, small to medium-sized. Pileus $13-50 \mathrm{~mm}$ wide, low convex to plane convex when young, then slightly depressed at center; surface smooth, grayish white (\#f2f2f2) to pale white (\#ffffff), yellowish white (\#ffcd9a);


Figure 3. Basidiomata of Clitocella a Clitocella borealichinensis (BJTC FM1781, holotype) b-d Clitocella colorata (b BJTC FM1593 c BJTC FM1952 d BJTC FM1891, holotype) Scale bars: 10 mm (a-d). Photos by JingZhong Cao
margin incurved, non-striate; context thin pale white, $1.0-1.2 \mathrm{~mm}$ thick. Lamellae decurrent, grayish white (\#f2f2f2), pale yellow (\#fff3e7), crowded, edges smooth, thin and fragile, lamellulae numerous and concolorous with lamellae. Stipe 20-32× $2-8 \mathrm{~mm}$, central to eccentric, occasionally lateral, cylindrical to subcylindrical, equal or sometimes slightly tapering at base, pale white (\#fffff), smooth or tomentose, usually with white rhizomorphs. Odor unrecorded. Taste not recorded. Chemical color reaction: not reacting with $\mathrm{KOH} 3 \%$ at pileus of dried specimens.

Basidiospores $[60 / 2 / 2](3.8-) 4-5(-5.5) \times 3.8-4.5 \mu \mathrm{~m}, \mathrm{~L}_{\mathrm{m}} \times \mathrm{W}_{\mathrm{m}}=4.61( \pm 0.42)$ $\times 4.06( \pm 0.18), \mathrm{Q}=0.95-1.25\left(\mathrm{Q}_{\mathrm{av}}=1.13 \pm 0.10\right)$, hyaline, globose to subglobose, occasionally broadly ellipsoid in profile view, slightly angled in polar or face view with obscure minute pustules or bumps. Basidia $17-25 \times 5-6(-7) \mu \mathrm{m}$, clavate, hyaline, four spored, rarely two spored; sterigmata $2-4 \mu \mathrm{~m}$ long. Lamellar trama more or less regular, composed of 3-8 $\mu \mathrm{m}$ wide hyaline hyphae, subhymenium consisting of filamentous hyphal segments. Lamellae edges fertile. Pleurocystidia and cheilocystidia absent. Pileipellis a cutis composed of more or less radially, loosely arranged, non-gelatinized, smooth, cylindrical hyphae, $2-6 \mu \mathrm{~m}$ wide and with pale brown to brown intracellular or parietal pigment; terminal hyphae subcylindric, narrowly clavate, occasionally irregular, 3-5 $\mu \mathrm{m}$ wide; subcutis made up of subparallel, compactly arranged, thinwalled, hyaline, smooth, cylindrical hyphae, 3-6 $\mu \mathrm{m}$ wide; pileal trama composed of interwoven, cylindrical hyphae, 2.5-10 $\mu \mathrm{m}$ wide. Stipitipellis a cutis composed of


Figure 4. Microscopy of Clitocella borealichinensis a basidiospores basidia c pileipellis. Scale bars: $5 \mu \mathrm{~m}(\mathbf{a}) ; 10 \mu \mathrm{~m}(\mathbf{b}, \mathbf{c})$. Drawings by Ning Mao.


Figure 5. Microscopy of Clitocella colorata $\mathbf{a}$ basidiospores $\mathbf{b}$ basidia $\mathbf{c}$ pileipellis. Scale bars: $10 \mu \mathrm{~m}(\mathbf{a}, \mathbf{c})$; $5 \mu \mathrm{~m}$ (b). Drawings by Ning Mao.
parallel, compactly arranged, thin-walled, non-gelatinized, hyaline hyphae, 2.5-6 $\mu \mathrm{m}$ wide. Stipititrama composed of interwoven, hyaline, cylindrical hyphae, 3-10 $\mu \mathrm{m}$ wide. Caulocystidia absent. Clamp connections absent.

Habit. Scattered or in groups on soil in broad-leaved (Quercus) forest, Shanxi province, China.

Additional specimens examined. China. Shanxi province, Xia County, alt. 970m, 7 October 2020, N. Mao MNM172 (BJTC FM1618).

Note. Clitocella borealichinensis is easily confused with C. orientalis, C. obscura (Pilát) Vizzini et al. and C. pallescens Silva-Filho \& Cortez in morphology because they are all have white to grayish white pileus and decurrent lamellae. However, C. orientalis differs from C. borealichinensis by its viscid pileus and stipe when wet, gelatinized pileipellis and stipitipellis, and slightly larger basidiospores of (4-)4.5-6 $\times 4-5 \mu \mathrm{~m}$ (Jian et al. 2020). Clitocella obscura produce a distinctly reddish reaction when $3 \% \mathrm{KOH}$ is placed on the pileus surface (Baroni 1981; as Rhodocybe) while C. borealichinensis has not that kind of reaction. Clitocella pallescens differs C. borealichinensis by its pale grey to yellowish white stipe (Silva-Filho et al. 2018; Jian et al. 2020).

Clitocella mundula and C. popinalis clustered with C. borealichinensis in our multilocus phylogeny (Fig. 1), indicating they are phylogenetically closely related to each other. Morphologically, C. mundula differs from C. borealichinensis by its yellowish gray or brown to dark smoke gray pileus and slightly larger basidiospores of (4-)4.5-6(-6.5) $\times 4-5 \mu \mathrm{~m}$ (Jian et al. 2020), C. popinalis by its brown to grayish brown pileus, bigger basidiospores of $5.5-7-5-5.5 \mu \mathrm{~m}$, and its pileus surface produces a reddish reaction in 3\% KOH (Baroni 1981; as Rhodocybe). Moreover, DNA analysis also revealed that $C$. borealichinensis shared less than $91.10 \%$ similarity in tef1 sequence with $C$. mundula and $91.20 \%$ similarity with C. popinalis, supporting their separation.

## Clitocella colorata L. Fan \& N. Mao, sp. nov.

MycoBank No: 843690
Figs 3b-d, 5, 6c, d

Etymology. colorata, referring to the colorful pileus.
Holotype. China. Shanxi Province, Pu County, Wulushan Mountain, 36³3.2'N, $111^{\circ} 11.58^{\prime} \mathrm{E}$, alt. $1740 \mathrm{~m}, 28$ July 2021, on the ground in coniferous forest dominated by Pinus armandii Franch., N. Mao MNM292 (BJTC FM1891).

Diagnosis. Clitocella colorata is characterized by its clitocyboid basidiomata, relatively colorful pileus (white to yellowish white, grayish white to grayish brown, pink white), globose or subglobose to broadly ellipsoid basidiospores, hyphae of pileipellis with pale yellow to yellowish brown intracellular or parietal pigment, the absence of hymenial cystidia and clamp connection. It is most similar to C. popinalis and C. mundula but differs from C. popinalis by its slightly smaller basidiospores, only appearing in the forest and genetic profile, and from C. mundula by its colorful pileus (white to yellowish white, grayish white to grayish brown, pink white).

Description. Basidiomata clitocyboid, small to large. Pileus $20-62 \mathrm{~mm}$ wide, dry, convex to plano-convex, sometimes infundibuliform, with a shallow depression at the center; margin not striate, often enrolled or flat, sometimes slightly uplifted; surface white (\#ffffff) to yellowish white (\#ffffe7), grayish white (\#f2f2f2) to grayish brown (\#dba773), pink white (\#fff3f5); context white (\#ffffff) to grayish white (\#f2f2f2), 1.0-1.5 mm thick. Lamellae decurrent, white (\#ffffff) to yellowish white(\#fff3e7), becoming yellowish brown (\#e0b487) on drying, crowded, 1.0-2.0 mm deep, edges entire and concolorous, thin and fragile, lamellulae in 2-4 tiers


Figure 6. Basidiospores of species in Clitocella. Clitocella revealed by SEM a,b Clitocella borealichinensis c, d Clitocella colorata Scale bars: $3 \mu \mathrm{~m}(\mathbf{a}, \mathbf{b}) ; 5 \mu \mathrm{~m}(\mathbf{c}, \mathbf{d})$. Photos by Li Fan.
of varying lengths. Stipe $22-42 \times 4-10 \mathrm{~mm}$, central, cylindrical, equal, pale white (\#ffffff) to yellowish brown (\#e0b487), smooth, usually with white rhizomorphs. Odor unrecorded. Taste not recorded. Chemical color reaction: pileal surface of dried samples negative with $3 \% \mathrm{KOH}$.

Basidiospores $[100 / 5 / 2](3.8-) 4.5-5.5(-6.0) \times(3.5-) 4-4.8(-5.0) \quad \mu \mathrm{m}$; $\mathrm{L}_{\mathrm{m}} \times \mathrm{W}_{\mathrm{m}}=4.90( \pm 0.44) \times 4.29( \pm 0.35), \mathrm{Q}=1.00-1.25\left(\mathrm{Q}_{\mathrm{av}}=1.14 \pm 0.09\right)$; hyaline, globose or subglobose to broadly ellipsoid in profile view, slightly angled in polar or face view with obscure minute pustules or bumps. Basidia $20-30 \times(4.5-) 5-6.5 \mu \mathrm{~m}$, clavate, hyaline, with four spored, rarely two spored; sterigmata $2-3.5 \mu \mathrm{~m}$ long. Lamellar trama composed of subparallel, hyaline, cylindrical hyphae, 2.5-6 $\mu \mathrm{m}$ wide, subhymenium consisting of filamentous hyphal segments, $2-3.5 \mu \mathrm{~m}$ wide. Lamellae edges fertile. Pleurocystidia and cheilocystidia absent. Pileipellis a cutis composed of parallel, compactly arranged, non-gelatinized, smooth, cylindrical hyphae, $2-5 \mu \mathrm{~m}$ wide, with pale yellow to yellowish brown intracellular or parietal pigment; subcutis made up of interwoven, slightly loosely arranged, hyaline, smooth, cylindrical hyphae, 3-6.5 $\mu \mathrm{m}$ wide; pileal trama composed of parallel, compactly arranged, hyaline, cylindrical hyphae, 3-10 $\mu \mathrm{m}$ wide. Stipitipellis a cutis composed of parallel, compactly arranged, thin-walled, non-gelatinized, cylindrical hyphae, $2-5 \mu \mathrm{~m}$ wide, heavily
or moderately encrusted with brown pigment. Stipititrama composed of parallel, compactly arranged, hyaline, cylindrical hyphae, 3-7 $\mu \mathrm{m}$ wide. Caulocystidia absent. Clamp connections absent.

Habit. Scattered or in groups on soil or rotten wood in coniferous (Pinus) or broad-leaved (Quercus) forest, Shanxi province, China.

Additional specimens examined. China. Shanxi province, Pu County, Wulushan Mountains, alt. 1750m, 28 July 2021, N. Mao MNM293 (BJTC FM1892); Wenshui County, alt. 1760m, 30 July 2021, L. Fan CF1219 (BJTC FM1952); Xia County, alt. 931m, 6 October 2020, N. Mao MNM102 (BJTC FM1593); Xia County, alt. 931m, 6 October 2020, N. Mao MNM103 (BJTC FM1594).

Notes. Morphologically, Clitocella colorata is easily confused with C. mundula and C. popinalis. However, according to Baroni (1981; as Rhodocybe), the pileus surface in C. mundula and C. popinalis can produce a reddish reaction in $3 \% \mathrm{KOH}$, whereas that is not exhibited in Clitocella colorata. The basidiospores of C. popinalis, $5.5-7 \times 5-5.5 \mu \mathrm{~m}$ (Baroni 1981; Kluting et al. 2014; Jian et al. 2020), are broader and longer than those of $C$. colorata $(4.5-5.5 \times 4-4.8 \mu \mathrm{~m})$. DNA analysis revealed that $C$. colorata shared less than $87.80 \%$ similarity in tef1 sequence with C. mundula and $86.10 \%$ similarity with C. popinalis, supporting their separation. Moreover, five ITS sequences (FJ770397, KR673647, KU561066, MK217434 and MK573922) labelled "C. popinalis" from India, Norway, South Korea, UK and USA are probably conspecific to the new species $C$. colorata as they clustered together with C. colorata in ITS tree (Fig. 2) and have more than $98.4 \%$ similarity in ITS region. However, these "C. popinalis" collections still need more other DNA regions and detailed morphology to support this view. One collection of " $C$. mundula," namely, AFTOLID 521 from Norway, should be re-identified C. colorata as it clustered together with C. colorata in the combined nrLSU-rpb2-tef1-atp6 tree (Fig. 1) and have more than $98.1 \%$ similarity in tef1 region. These showed that the new species C. colorata maybe have a wide geographical distribution. Although C. orientalis is sister to C. colorata with strong support, these two species have obvious differences in morphology. The pileus and stipe of C. orientalis are usually viscid when wet and have gelatinized pileipellis and stipitipellis. Clitocella colorata has non-gelatinized pileipellis and stipitipellis, and its pileus is more colorful and darker (Jian et al. 2020). DNA analysis revealed that C. colorata shared less than $95.80 \%$ similarity in tef1 sequence with C. orientalis and $90.20 \%$ similarity in ITS sequence. Moreover, C. colorata has a wider distribution range than $C$. orientalis, which is only distributed in China.

## Discussion

Three species of Clitocella are confirmed from Shanxi Province, north China in this study. Of them, C. colorata is the most commonly encountered species, which distributes across the provincial area and grows in almost all kinds of forest. Clitocella orientalis and Clitocella borealichinensis are probably limited in southern Shanxi province, and they usually occur in the Quercus spp. forests.

ITS gene is rarely used in the species classification of Clitocella in previous studies because it contains many ambiguous sites. In the contrast, the partial sequences of three protein-coding genes (the atp $6, r p b 2$ and tef1) are usually used to infer the phylogeny of Clitocella (Kluting et al. 2014; Baroni et al. 2020; Jian et al. 2020). However, we found that ITS, rpb2, and tef1 gene tree are similar to the combined (nrLSU-rpb2-tef1$\operatorname{atp} 6$ ) gene regions tree when we performed phylogenetic tree construction respectively using the ITS, nrLSU, rpb2, tef1 and atp6 gene of Clitocella (Fig. 2, Suppl. material $1-4$ : Figs $S 1-S 4)$. DNA analysis also showed that the intraspecific similarity of the ITS region is $\geq 98.4 \%$ and of tef1 gene is $\geq 98.1 \%$, the interspecific similarity of ITS region is $\leq 96.1 \%$ and of tef1 is $\leq 95.9 \%$ (Table 2, Table 3). But for the rpb2 gene, the intraspecific variation of $C$. mundula is more than the interspecific variation of C. colorata and C. orientalis (Table 4). Therefore, we consider that both the ITS and tefl may be more effective for the classification of Clitocella species.

Our molecular phylogenetic analysis (Fig. 1) revealed that one Norway collection O-F19454, which is labelled as Clitocella mundula, formed an independent clade, and it shared less than $93.40 \%$ similarity in tef1 sequence with other Clitocella species. These show that it probably represents a new species of Clitocella. The sequences of Clitocella fallax formed two or three (in rpb2 phylogeny) independent branches in our phylogenetic analyses (Fig. 2, Suppl. material 1-4: Figs S1-S4), and the similarity between the branches is less than $90.2 \%$ in tef1 sequence and $94.9 \%$ in $r p b 2$ sequence. These indicate that these specimens of $C$. fallax probably represented two or three species. The specimens of $C$. fallax should be therefore re-examined to resolve this taxonomic issue. Clitocella termitophila is not clustered in the genus Clitocella (Fig. 1). Moreover, in the rpb2 gene

Table 2. Interspecific variation and intraspecific variation of ITS in Clitocella species.

| Species | Number (n) | Intraspecific variation (\%) | Interspecific variation (\%) |
| :--- | :---: | :---: | :---: |
| Clitocella colorata | 9 | $<1.6 \%$ | $>3.9 \%$ |
| C. fallax | 3 | $<0.3 \%$ | $>11.8 \%$ |
| C. mundula | 1 | - | $>6.0 \%$ |
| C. borealichinensis | 2 | - | $>9.6 \%$ |
| C. obscura | 1 | - | $>6.6 \%$ |
| C. orientalis | 3 | $<0.9 \%$ | $>3.9 \%$ |

Table 3. Interspecific variation and intraspecific variation of tefl in Clitocella species.

| Species | Number (n) | Intraspecific variation (\%) | Interspecific variation (\%) |
| :--- | :---: | :---: | :---: |
| Clitocella colorata | 4 | $<1.9 \%$ | $>4.1 \%$ |
| C. fallax | 1 | - | $>9.8 \%$ |
| C. fallax | 2 | $<0.1 \%$ | $>9.8 \%$ |
| C. mundula $^{\text {b }}$ | 6 | $<0.3 \%$ | $>7.5 \%$ |
| C. mundula'c | 1 | - | $>4.7 \%$ |
| C. borealichinensis | 1 | - | $>8.4 \%$ |
| C. orientalis | 3 | $<0.1 \%$ | $>4.1 \%$ |
| C. popinalis | 7 | - | $>4.7 \%$ |

[^1]Table 4. Interspecific variation and intraspecific variation of $r p b 2$ in Clitocella species.

| Species | Number (n) | Intraspecific variation (\%) | Interspecific variation (\%) |
| :--- | :---: | :---: | :---: |
| Clitocella colorata | 4 | $<0.7 \%$ | $>1.7 \%$ |
| C. fallax ${ }^{\text {a }}$ | 1 | - | $>4.0 \%$ |
| C. fallax ${ }^{\text {b }}$ | 4 | $<0.1 \%$ | $>5.1 \%$ |
| C. fallax | - | $>4.0 \%$ |  |
| C. mundula | 1 | $<2.1 \%$ | $>4.9 \%$ |
| ' mundula | - | $>2.2 \%$ |  |
| C. borealichinensis | 6 | - | $>5.5 \%$ |
| C. orientalis | 1 | $<0.5 \%$ | $>1.7 \%$ |
| C. popinalis | 6 | $<0.4 \%$ | $>2.2 \%$ |
| C. termitophila | 9 | - | $>16.9 \%$ |

${ }^{a}$ represents voucher 256680 KM ; ${ }^{\text {b }}$ represents voucher O-F88953, K(M): 116541, CBS 129.63, ME Noordeloos 1997173; ${ }^{\text {c represents }}$ voucher ME Noordeloos 200367; ${ }^{\text {d represents voucher O-F19454. }}$
tree C. termitophila did not gather with Clitocella, Clitopilopsis or Clitopilus but formed a single branch (Suppl. material 2: Fig. S2). These indicate that Clitocella termitophila probably represents a potential taxonomic position rather than the species of Clitocella.

## Key to the species of Clitocella

1 Basidiomata clitocyboid........................................................................... 2

- Basidiomata pleurotoid ...................... C. termitophila* (Baroni et al. 2020)

2 Pileus surface gray, dark gray, pale yellow to yellowish brown, pigments present in pileipelli3

- Pileus surface almost white to pastel gray, pigments absent in pileipellis ..... 8
3 Basidiospores globose to subglobose. ..... 4
Basidiospores ellipsoid ..... 7
4 Pileus surface of dried samples with a positive KOH reaction ..... 5
Pileus surface of dried samples with a negative KOH reaction. ..... 6
5 Occurring in grassland systemsC. popinalis (Baroni 1981; Kluting et al. 2014; Jian et al. 2020)- Occurring in forested systems
$\qquad$
..............C. mundula* (Baroni 1981; Kluting et al. 2014; Jian et al. 2020)
6 Pileus color with pink tinges C. colorata*
- Pileus color without pink tinges C. borealichinensis*
7 Pileus color with yellow tinges, basidiospores small, $5-8 \times 3.5-5.5 \mu \mathrm{~m}$.
C. himantiigena (Silva-Filho et al. 2018)
- Pileus color without yellow tinges, basidiospores large, 7-9 $\times 5-6 \mu \mathrm{~m}$
$\qquad$C. ammophila (Contu 1999)
8 Basidiospores globose to subglobose or ovatae ..... 9
- Basidiospores amygdaliform to ellipsoid. ..... 11

[^2]| 9 | Basidia long, length > $40 \mu \mathrm{~m}$..........................C. nigrescens (Maire 1945) |
| :---: | :---: |
| - |  |
| 10 | Pileus infundibuliform to plano-convex, basidiospores $4-5 \times 3-4.5 \mu \mathrm{~m}$ $\qquad$ C. pallescens (Silva-Filho et al. 2018; Jian et al. 2020) |
| - |  |
| 11 |  |
| - |  |

## Acknowledgements

We extend our appreciation to Dr. J.Z. Cao for collecting specimens and providing valuable suggestions. The study was supported by the National Natural Science Foundation of China (No. 31750001) and the Beijing Natural Science Foundation (No. 5172003).

## References

Alfaro ME, Zoller S, Lutzoni F (2003) Bayes or bootstrap? A simulation study comparing the performance of Bayesian Markov chain Monte Carlo sampling and bootstrapping in assessing phylogenetic confidence. Molecular Biology and Evolution 20(2): 255-266. https://doi.org/10.1093/molbev/msg028
Baroni TJ (1981) A revision of the genus Rhodocybe Maire (Agaricales). Beih Nova Hedwigia 67: 1-194.
Baroni TJ, Angelini C, Bergemann SE, Lodge DJ, Lacey L, Curtis TA, Cantrell SA (2020) Rhodocybe-Clitopilus clade (Entolomataceae, Basidiomycota) in the Dominican Republic: New taxa and first reports of Clitocella, Clitopilus, and Rhodocybe for Hispaniola. Mycological Progress 19(10): 1083-1099. https://doi.org/10.1007/s11557-020-01619-y
Binder M, Hibbett DS (2003) Oligonucleotides. http://www.clarku.edu/faculty/dhibbett/Protocols_Folder/Primers/Primers.htm [accessed 18 Mar 2012]
Castresana J (2000) Selection of conserved blocks from multiple alignments for their use in phylogenetic analysis. Molecular Biology and Evolution 17(4): 540-552. https://doi. org/10.1093/oxfordjournals.molbev.a026334
Contu M (1999) Ecologia e tassonomia del genere Rhodocybe (Basidiomycetes, Entolomataceae) in Sardegna. Revista Catalana de Micologia 22: 5-14.
Contu M (2009) Studi sul genere Clitopilus (incl. Rhodocybe) 1. Prima segnalazione in Italia di Clitopilus blancii comb. nov., nuove raccolte di Clitopilus giovanellae, iconografia di Clitopilus carneolus comb. nov. e ulteriori nuove combinazioni. Bollettino AMER 77-78, 15-31.
Dring DM (1971) Techniques for microscopic preparation. In: Booth C (Ed.) Methods in microbiology, vol 4. Academic, New York, 98 pp. https://doi.org/10.1016/S0580-9517(09)70008-X

Gardes M, Bruns TD (1993) ITS primers with enhanced specificity for basidiomycetes application to the identification of mycorrhizae and rusts. Molecular Ecology 2(2): 113118. https://doi.org/10.1111/j.1365-294X.1993.tb00005.x

Hall TA (1999) BioEdit: a user-friendly biological sequence alignment editor and analysis program for Windows 95/98/NT. Nucleic Acids Symposium Series 41: 95-98.
Hillis DM, Bull JJ (1993) An empirical test of bootstrapping as a method for assessing confidence in phylogenetic analysis. Systematic Biology 42(2): 182-192. https://doi. org/10.1093/sysbio/42.2.182
Jian SP, Tolgor B, Zhu XT, Deng WQ, Yang ZL, Zhao ZW (2020) Clitopilus, Clitocella, and Clitopilopsis in China. Mycologia 112(2): 371-399. https://doi.org/10.1080/00275514.2 019.1703089

Katoh K, Standley DM (2013) MAFFT multiple sequence alignment software version 7: Improvements in performance and usability. Molecular Biology and Evolution 30(4): 772-780. https://doi.org/10.1093/molbev/mst010
Kluting KL, Baroni TJ, Bergemann SE (2014) Toward a stable classification of genera within the Entolomataceae: A phylogenetic re-evaluation of the Rhodocybe-Clitopilus clade. Mycologia 106(6): 1127-1142. https://doi.org/10.3852/13-270
Kretzer AM, Bruns TD (1999) Use of atp6 in fungal phylogenetics: An example from the Boletales. Molecular Phylogenetics and Evolution 13(3): 483-492. https://doi. org/10.1006/mpev.1999.0680
Liu YJ, Whelen S, Hall BD (1999) Phylogenetic relationships among Ascomycetes: Evidence from an RNA polymerase II subunit. Molecular Biology and Evolution 16(12): 1799-1808. https://doi.org/10.1093/oxfordjournals.molbev.a026092
Maire R (1945) Études mycologiques. Fascicule 5. Bulletin de la Société d'Histoire Naturelle de l'Afrique du Nord 36: 24-42.
Matheny PB (2005) Improving phylogenetic inference of mushrooms with RPB1 and RPB2 nucleotide sequences (Inocybe; Agaricales). Molecular Phylogenetics and Evolution 35: 1-20. https://doi.org/10.1016/j.ympev.2004.11.014
Matheny PB, Wang Z, Binder M, Curtis JM, Lim YW, Nilsson RH, Hughes KW, Hofstetter V, Ammirati JF, Schoch CL, Langer E, Langer G, McLaughlin DJ, Wilson AW, Frøslev T, Ge ZW, Kerrigan RW, Slot JC, Yang ZL, Baroni TJ, Fischer M, Hosaka K, Matsuura K, Seidl MT, Vauras J, Hibbett DS (2007) Contributions of rpb2 and tef1 to the phylogeny of mushrooms and allies (Basidiomycota, Fungi). Molecular Phylogenetics and Evolution 43(2): 430-451. https://doi.org/10.1016/j.ympev.2006.08.024
Nylander J (2004) MrModeltest 2.2. Computer software distributed by the Evolutionary Biology Centre, University of Uppsala, Uppsala.
Page RD (2001) TreeView. Glasgow University, Glasgow.
Rehner $S$ (2001) Primers for elongation factor 1- $\alpha$ (tef1). [cited 2021 Nov 1] http://www.aftol. org/pdfs/EF1primer.pdf
Ronquist F, Huelsenbeck JP (2003) MrBayes 3: Bayesian phylogenetic inference under mixed models. Bioinformatics (Oxford, England) 19(12): 1572-1574. https://doi.org/10.1093/ bioinformatics/btg180

Silva-Filho AGS, Teixeira-Silva M, Cortez VG (2018) New species, new combination, and notes on Clitocella and Rhodocybe (Entolomataceae) from Paraná State, Brazil. Darwiniana 6: 58-67. https://doi.org/10.14522/darwiniana.2018.61.775
Stamatakis A (2006) RAxML-vI-HPC: Maximum-likelihood-based phylogenetic analyses with thousands of taxa and mixed models. Bioinformatics (Oxford, England) 22(21): 2688-2690. https://doi.org/10.1093/bioinformatics/bt1446
Stamatakis A (2014) RAxML version 8: A tool for phylogenetic analysis and post-analysis of large phylogenies. Bioinformatics (Oxford, England) 30(9): 1312-1313. https://doi. org/10.1093/bioinformatics/btu033
Stamatakis A, Ludwig T, Meier H (2005) RAxML-III: A fast program for maximum likelihoodbased inference of large phylogenetic trees. Bioinformatics (Oxford, England) 21(4): 456-463. https://doi.org/10.1093/bioinformatics/bti191
Vilgalys R, Hester M (1990) Rapid genetic identification and mapping of enzymatically amplified ribosomal DNA from several Cryptococcus species. Journal of Bacteriology 172(8): 4239-4246. https://doi.org/10.1128/jb.172.8.4238-4246.1990
Vizzini A, Baroni TJ, Sesli E, Antonín V, Saar I (2016) Rhodocybe tugrulii (Agaricales, Entolomataceae), a new species from Turkey and Estonia based on morphological and molecular data, and a new combination in Clitocella (Entolomataceae). Phytotaxa 267(1): 001-015. http://doi.org/10.11646/phytotaxa.267.1.1
White TJ, Bruns T, Lee S, Taylor J (1990) Amplification and direct sequencing of fungal ribosomal RNA genes for phylogenetics. In: Innis MA, Gelfand DH, Sninsky JJ, White TJ (Eds) PCR protocols: a guide to methods and applications. Academic Press, New York, 315-322. https://doi.org/10.1016/B978-0-12-372180-8.50042-1

## Supplementary material I

## Figure S1

Authors: Ning Mao, Jing-Chong Lv, Yu-Yan Xu, Tao-Yu Zhao, Li Fan
Data type: JPG file
Explanation note: Phylogeny derived from Maximum Likelihood analysis of the $n r L S U$ dataset of Clitocella and related genera in the family Entolomataceae. The bootstrap frequencies (> $70 \%$ ) is shown on the supported branches. New species are highlighted in red.
Copyright notice: This dataset is made available under the Open Database License (http://opendatacommons.org/licenses/odbl/1.0/). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.
Link: https://doi.org/10.3897/mycokeys.88.80068.suppl1

## Supplementary material 2

Figure S2
Authors: Ning Mao, Jing-Chong Lv, Yu-Yan Xu, Tao-Yu Zhao, Li Fan
Data type: JPG file
Explanation note: Phylogeny derived from Maximum Likelihood analysis of the rpb2 dataset of Clitocella and related genera in the family Entolomataceae. The bootstrap frequencies (> 70\%) is shown on the supported branches. New species are highlighted in red.
Copyright notice: This dataset is made available under the Open Database License (http://opendatacommons.org/licenses/odbl/1.0/). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.
Link: https://doi.org/10.3897/mycokeys.88.80068.suppl2

## Supplementary material 3

Figure S3
Authors: Ning Mao, Jing-Chong Lv, Yu-Yan Xu, Tao-Yu Zhao, Li Fan
Data type: JPG file
Explanation note: Phylogeny derived from Maximum Likelihood analysis of the tef1 dataset of Clitocella and related genera in the family Entolomataceae. The bootstrap frequencies (>70\%) is shown on the supported branches. New species are highlighted in red.
Copyright notice: This dataset is made available under the Open Database License (http://opendatacommons.org/licenses/odbl/1.0/). The Open Database License ( ODbL ) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.
Link: https://doi.org/10.3897/mycokeys.88.80068.suppl3

## Supplementary material 4

## Figure S4

Authors: Ning Mao, Jing-Chong Lv, Yu-Yan Xu, Tao-Yu Zhao, Li Fan
Data type: JPG file
Explanation note: Phylogeny derived from Maximum Likelihood analysis of the atp6 dataset of Clitocella and related genera in the family Entolomataceae. The bootstrap frequencies (> 70\%) is shown on the supported branches. New species are highlighted in red.
Copyright notice: This dataset is made available under the Open Database License (http://opendatacommons.org/licenses/odbl/1.0/). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.
Link: https://doi.org/10.3897/mycokeys.88.80068.suppl4

## Supplementary material 5

## ITS alignment

Authors: Ning Mao, Jing-Chong Lv, Yu-Yan Xu, Tao-Yu Zhao, Li Fan
Data type: PHY file
Explanation note: The ITS dataset comprised 27 samples of 11 taxa and 662 characters.
Copyright notice: This dataset is made available under the Open Database License (http://opendatacommons.org/licenses/odbl/1.0/). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.
Link: https://doi.org/10.3897/mycokeys.88.80068.suppl5


[^0]:    * These authors contributed equally to this work.

    Copyright Ning Mao et al. This is an open access article distributed under the terms of the Creative Commons Attribution License (CC BY 4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

[^1]:    ${ }^{\text {a }}$ represents voucher 25668 OKM ; ${ }^{\mathrm{b}}$ represents voucher O-F88953, K(M): 116541; ${ }^{\text {c represents voucher O-F19454 }}$

[^2]:    * Indicates the presence of molecular data.

