

Research Article

# Morphology and multigene phylogeny reveal three new species of *Samsoniella* (Cordycipitaceae, Hypocreales) from spiders in China

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#### Abstract

The genus Samsoniella was erected based on orange cylindrical to clavate stromata, superficial perithecia and conidiophores with Isaria-like phialides and to segregate them from the Akanthomyces group. In this study, based on morphological features and multigene (SSU, LSU, TEF, RPB1 and RPB2) phylogenetic analysis six Samsoniella species parasitizing spiders were collected in China. Three of them belong to known species S. alpina, S. erucae and S. hepiali. Three new species S. anhuiensis sp. nov., S. aranea sp. nov. and S. fusiformispora sp. nov. are illustrated and described. They are clearly distinct from other species in Samsoniella occurring in independent subclades. Furthermore, among the four insect-pathogenic fungi specimens collected from similar sites, three of them were identified as the new species described below. Our study significantly broadens the host range of Samsoniella from Insecta to Arachnida, marking a noteworthy expansion in understanding the ecological associations of these fungi. Additionally, the identification of both mononematous and synnematous conidiophores in our study not only expands the knowledge of Samsoniella species but also provides a basis for future research by comparing the ecological significance between these conidiophore types. In conclusion, our study enhances the understanding of Samsoniella diversity, presenting a refined phylogenetic framework and shedding light on the ecological roles of these fungi in spider parasitism.

Key words: Araneogenous fungi, Isaria-like, Samsoniella, taxonomy

# Introduction

The genus *Isaria* Pers. was established by Persoon (1794) with *I. farinosa* (Pers.) Fr. as the type species (Hodge et al. 2005). *Isaria* is characterized by the formation of branched synnemata that give rise to flask-shaped phialides produced in whorls. For a considerable period, *Isaria* has been considered the asexual morph of *Cordyceps sensu stricto*, a classification within the family Cordycipitaceae, which encompasses numerous species featuring pallid or brightly pigmented, fleshy stromata (Sung et al. 2007; Maharachchikumbura et al. 2015). Samson (1974) transferred some species including *I. farinosa* to *Paecilomyces* Bainer



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(1907). However, Hodge et al. (2005), based on morphological and molecular phylogenetic studies, moved Paecilomyces farinosa back to Isaria re-establishing the type as Isaria farinosa (Holmsk.) Fr. Most of the insect-pathogenic mesophilic Paecilomyces species in sect. Isarioidea of Samson (1974) were transferred to Isaria (Luangsa-ard et al. 2004, 2005; Gams et al. 2005). Nonetheless, Kepler et al. (2017) proposed the rejection of the genus Isaria due to the polyphyletic distribution of Isaria species. Recently, molecular phylogenetic analysis, has shown that some Isaria-like fungi are distributed in the genus Akanthomyces of the family Cordycipitaceae, forming monophyletic branches and are closely related to the genus Akanthomyces. Mongkolsamrit et al. (2018) established this phylogenetic branch as a new genus Samsoniella Mongkols., Noisrip., Thanakitp., Spatafora & Luangsa-ard. They accommodated three species of Lepidoptera entomopathogenic fungi in the genus; S. alboaurantia (G. Sm.) Mongkolsamrit, S. aurantia Mongkolsamrit and S. inthanonensis Mongkolsamrit. The three species have orange cylindrical to clavate stromata, superficial perithecia and orange conidiophores with Isaria-like phialides and hyaline conidia.

Over the past seven years, there has been extensive research on the species diversity within the genus Samsoniella, possibly driven by the significant medical and ecological value associated with certain species in the genus. In a follow-up study, Wang et al. (2020a) documented nine new species within the genus Samsoniella. Specifically, Paecilomyces hepiali Chen, formerly misconstrued as the asexual counterpart of Ophiocordyceps sinensis, demonstrated the ability to produce Isaria-like phialides. The perplexing taxonomic status of P. hepiali prompted taxonomists to reconsider its classification. Wang et al. (2020a) determined that the most suitable systematic position for P. hepiali is within the genus Samsoniella. Consequently, they proposed the new taxonomic combination S. hepiali for this species. Subsequently, Chen et al. (2020) described three additional species of Samsoniella. Furthermore, phylogenetic analysis led to the repositioning of strains previously identified as I. farinosa. Notably, strains CBS 240.32 and CBS 262.58 were integrated into the genus Samsoniella and redesignated as S. alboaurantia (Mongkolsamrit et al. 2018; Chen et al. 2021). Similarly, strains OSC 111005 and OSC 111006 were reassigned to S. farinosa Wang (Wang et al. 2020b). More recently, Chen et al. (2021, 2022, 2023), Wang et al. (2022), Wang et al. (2023) and Crous et al. (2023) contributed descriptions of fifteen additional novel Samsoniella species. Consequently, the genus Samsoniella now comprises a total of thirty-one recognized species.

We carried out a series of surveys for spider pathogenic fungi in China. A total of seven spider cadavers infected by *Samsoniella* were collected and isolated. Based on morphological and molecular phylogenetic analyses, three were identified as *S. alpina*, *S. erucae*, and *S. hepiali*. However, the other four strains represented four new species, which are described here as *S. anhuiensis* sp. nov., *S. aranea* sp. nov. and *S. fusiformispora* sp. nov. Among the four insect-pathogenic fungi specimens collected from the same sites, three of them were identified as the new species described below. Our study enhances the understanding of *Samsoniella* diversity, presenting a refined phylogenetic framework and shedding light on the ecological roles of these fungi in spider parasitism.

## **Materials and methods**

## Sample collection, isolation and morphological observations

The majority of spider specimens infected by fungi were collected from all over China. Four specimens were collected from the Jingting Mountains National Forest Park, Anhui Province, southeastern China. Four specimens were collected from the Jinggang Mountains National Nature Reserve, Jiangxi Province, southeastern China. One specimen was collected from the Maiji National Forest Park, Gansu Province, northwestern China. One specimen was collected from the Yaoluoping National Forest Park, Anhui Province, southeastern China, and one specimen was collected from the Wanfo Mountains, Anhui Province, southeastern China. Several insect specimens infected by fungi were collected from sites similar to those where spider specimens were collected. The collections were noted and photographed in the field, then carefully deposited in plastic boxes and returned to the laboratory. Fungal cultures were isolated from fresh conidia or mycelia from spider cadavers. Pure cultures were established and incubated on fresh potato dextrose agar (PDA) plates and grown at 25 °C for 2 weeks. The fresh structures of specimens and isolated strains were mounted in water for measurements and lactophenol cotton blue solution for microphotography following Wang et al. (2020a). Features such as size and shape of conidia, colony color in culture, were made from squash mounts and sections made from fresh specimen and culture grown on oatmeal agar (OA, Difco), PDA and one quarter strength SDAY (SDAY/4, Difco) (Bischoff et al. 2009). The color of the cultures was characterized using the Naturalist's Color Guide (Smith 1975). Microscopic observations were made from squash mounts and sections made from fresh material using a ZEISS Axiolab 5 microscope. All samples and strains studied here were deposited in the Research Center for Entomogenous Fungi (RCEF) of Anhui Agricultural University.

## DNA extraction, PCR amplification and sequencing

Total genomic DNA was extracted from cultured mycelia with CTAB method (Liu et al. 2001), then stored in -20 °C. Two gene regions, namely the small subunit ribosomal RNA (SSU) and large subunit ribosomal RNA (LSU) were sequenced from the cell nuclei, and three protein coding genes, translation elongation factor-1a (TEF) and the largest and second largest subunits of RNA polymerase II (RPB1 and RPB2) were used in this study. The SSU and LSU were amplified with NS1/NS4 (White et al. 1990) and LROR (Vilgalys and Hester 1990)/LR7(Hopple 1994). The TEF with 983F/2218R (Rehner and Buckley 2005), RPB1 with CRPB1/RPB1-Cr (Castlebury et al. 2004) and RPB2 with fRPB2-7CR /fRPB2-5F (Liu et al. 1999) were amplified. PCR reactions of the five nuclear loci were carried out in 25 µL reaction mixture containing 12.5 µL 2× Tag Plus MasterMix (CoWin Biosciences, Beijing, China), 1 µL of each primer (10 µM), 1.5 µL of template DNA (1–2 ng) and 9  $\mu$ L of sterile water. PCR cycle conditions were as previously described (Sung et al. 2007). PCR products were purified and sequenced by Sangon Company (Shanghai, China). The resulting sequences were checked manually, then submitted to GenBank.

## Sequence alignment and phylogenetic analyses

The sequences in this study were uploaded to BLAST and searched in the GenBank database to determine probable taxa. DNA sequences generated in this study were assembled and edited using version 6.0. DNASTAR. Generated SSU, LSU, *TEF*, *RPB1* and *RPB2* sequences were aligned with those published by Chen et al. (2020) and Wang et al. (2020a) and others downloaded from Gen-Bank were used as a dataset of taxa in *Samsoniella* and closely related *Samsoniella* groups (Table 1). Sequences of the genus *Akanthomyces (A. aculeatus* HUA772 and HUA 186145) were chosen as the outgroup. Multiple sequence alignment was conducted with MAFFT 7.3.13 (Katoh and Standley 2013). The final sequence alignment of the combined dataset was used for analyses using Maximum Likelihood (ML) and Bayesian Inference (BI) to infer their phylogenetic relationships.

Phylogenetic inference was done according to Maximum Likelihood (ML) using RAxML version 8 (Stamatakis 2014) and Bayesian Inference (BI) using MrBayes v.3.2 (Ronquist et al. 2012). For the ML analysis, we used the GTR-CAT model for all partitions, in accordance with recommendations in the RAx-ML manual against the use of invariant sites and 1000 rapid bootstrap replicates. The GTR+I+G model was selected by MrModeltest 2.2 (Darriba et al. 2012) as the best nucleotide substitution model for the Bayesian analysis. Four MCMC chains were executed simultaneously for 2000,000 generations, sampling every 100 generations. Finally, phylogenetic trees were visualized using the Interactive Tree of Life (iTOL) (https://itol.embl.de) online tool (Letunic and Bork 2016).

0	Otracia Na		Ge	nBank accession I	No.	
Species	Strain No.	SSU	LSU	TEF	RPB1	RPB2
Akanthomyces aculeatus	HUA772	KC519368	KC519370	_	_	-
A. aculeatus	HUA186145 <sup>™</sup>	MF416572	MF416520	MF416465	_	-
A. cf. coccidioperitheciatus	NHJ 5112	EU369109	EU369043	EU369026	EU369066	-
A. coccidioperitheciatus	NHJ 6709	EU369110	EU369042	EU369025	EU369067	EU369086
A. farinosa	CBS541.81	MF416606	MF416553	_	MF416655	-
A. lecanii	CBS101247	AF339604	AF339555	DQ522359	DQ522407	DQ522466
A. muscarius	CBS 143.62	KM283774	KM283798	KM283821	KM283841	KM283863
Beauveria bassiana	ARSEF1564 <sup>⊤</sup>	-	-	HQ880974	HQ880833	HQ880905
B. brongniartii	ARSEF 617 <sup>™</sup>	-	-	HQ880991	HQ880854	HQ880926
	BCC 16585	-	JF415967	JF416009	JN049885	JF415991
B. staphylinidicola	ARSEF 5718	EF468981	EF468836	EF468776	EF468881	-
Cordyceps farinosa	CBS111113	AY526474	MF416554	GQ250022	MF416656	GU979973
C. militaris	OSC 93623	AY184977	AY184966	DQ522332	DQ522377	AY545732
Isaria sp.	spat 09-050	MF416613	MF416559	MF416506	MF416663	MF416457
	spat 09-051	MF416614	MF416560	MF416507	MF416664	MF416458
Samsoniella alboaurantium	CBS 240.32	JF415958	JF415979	JF416019	JN049895	JF415999
	CBS 262.58	-	-	MF416497	MF416654	MF416448

Table 1. Species, strain numbers, accession numbers and origins of *Samsoniella* and related taxa used in this study, new sequences were shown in bold.

Creation	Canain Na		Ge	nBank accession I	No.	
Species	Strain No.	SSU	LSU	TEF	RPB1	RPB2
S. alpina	YFCC 5818	MN576753	MN576809	MN576979	MN576869	MN576923
	YFCC 5831	MN576754	MN576810	MN576980	MN576870	MN576924
S. alpina	RCEF0643	_	-	OM482385	_	-
S. anhuiensis	RCEF2830	OM268843	OM268848	OM483864	OM751889	_
	RCEF2590	OR978313	OR978316	OR966516	OR989964	_
S. antleroides	YFCC 6016	MN576747	MN576803	MN576973	MN576863	MN576917
	YFCC 6113	MN576748	MN576804	MN576974	MN576864	MN576918
S. aranea	RCEF2831	OM268844	OM268849	OM483865	OM751882	OM802500
	RCEF2868	OM268845	OM268850	OM483866	OM751883	OM802501
	RCEF2870	OR978314	OR978317	OR966517	OR989965	OR989966
S. aurantia	TBRC 7271 <sup>⊤</sup>	_	MF140728	MF140846	MF140791	MF140818
	TBRC 7273	_	-	MF140844	_	MF140816
S. cardinalis	YFCC5830	MN576732	MN576788	MN576958	MN576848	MN576902
	YFCC 6144	MN576730	MN576786	MN576956	MN576846	MN576900
S. cristata	YFCC6021	MN576735	MN576791	MN576961	MN576851	MN576905
	YFCC6023	MN576736	MN576792	MN576962	MN576852	MN576906
S. coccinellidicola	YFCC8772	ON563166	ON621670	ON676514	ON676502	ON568685
	YFCC8773	ON563167	ON621671	ON676515	ON676503	ON568686
S. coleopterorum	A19502	_	_	MT642602	MT642603	MN101587
S. duyunensis	DY09162	_	OQ363114	OQ398146	_	_
	DY07501	_	OR263307	OR282780	OR282773	OR282776
	DY07502	_	OR263427	OR282781	_	OR282777
S. erucae	KY11121	_	ON502835	ON525425	_	ON525424
	KY11122	_	ON502822	ON525427	_	ON525426
S. erucae	RCEF2595	OM268842	OM268847	OM483863	OM751888	-
	RCEF2592	_	_	OR966518	_	_
S. farinosa	OSC111005	DQ522558	DQ518773	DQ522348	DQ522394	_
	OSC111006	EF469127	EF469080	EF469065	EF469094	_
S. farinospora	YFCC8774	ON563168	ON621672	ON676516	ON676504	ON568687
·	YFCC9051	ON563169	ON621673	ON676517	ON676505	ON568688
S. fusiformispora	RCEF5406	OM268846	OM268851	OM483867	OM751890	_
·	RCEF2588	OR978312	OR978315	OR966515	_	_
S. guizhouensis	KY11161	_	ON502830	ON525429	_	ON525428
5	KY11162	_	ON502846	ON525431	_	ON525430
S. haniana	YFCC8769	ON563170	ON621674	ON676518	ON676506	ON568689
	YFCC8770	ON563171	ON621675	ON676519	ON676507	ON568690
	YFCC8771	ON563172	ON621676	ON676520	ON676508	ON568691
S. hepiali	YFCC 5823	MN576745	MN576801	MN576971	MN576861	MN576915
	YFCC 5828	MN576744	MN576800	MN576970	MN576860	MN576914
S. hepiali	RCEF1481	OL854202	_	OM482386	_	_
S. hymenopterorum	A19521	_	_	MN101588	MT642603	MT642604
	A19522	_	_	MN101591	MN101589	MN101590
S. inthanonensis	TBRC 7915	_	MF140725	MF140849	MF140790	MF140815
S. kunmingensis	YHH16002	MN576746	MN576802	MN576972	MN576862	MN576916
S. lanmaoa	YFCC6148 <sup>™</sup>	MN576733	MN576789	MN576959	MN576849	MN576903
	YFCC6193	MN576734	MN576790	MN576960	MN576850	MN576904

Species	Strain No.		Ge	nBank accession I	No.	
Species	Strain No.	SSU	LSU	TEF	RPB1	RPB2
S. lepidopterorum	DL10071	-	-	MN101594	MN101592	MN101593
	DL10072	_	_	MT642606	_	MT642605
S. neopupicola	KY11321	-	ON502839	ON525433	_	ON525432
	KY11322	_	ON502833	ON525435	_	ON525434
S. pseudogunnii	GY407201	-	MZ827010	-	-	_
	GY407202	-	MZ831865	-	-	-
S. pseudotortricidae	YFCC9052	ON563173	ON621677	ON676521	ON676509	ON568692
	YFCC9053	ON563174	ON621678	ON676522	ON676510	ON568693
S. pupicola	DY101681	-	MZ827009	MZ855231	-	MZ855237
	DY101682	-	MZ827635	MZ855232	-	MZ855238
S. ramosa	YFCC6020 <sup>⊤</sup>	MN576749	MN576805	MN576975	MN576865	MN576919
S. sinensis	YFCC8766	ON563175	ON621679	ON676523	ON676511	ON568694
	YFCC8767	ON563176	ON621680	ON676524	ON676512	ON568695
	YFCC8768	ON563177	ON621681	ON676525	ON676513	ON568696
S. tiankengensis	KY11741	-	ON502838	ON525437	-	ON525436
	KY11742	-	ON502841	ON525439	-	ON525438
S. tortricidae	YFCC6013	MN576751	MN576807	MN576977	MN576867	MN576921
	YFCC6131	MN576750	MN576806	MN576976	MN576866	MN576920
S. vallis	DY07241	-	OR263306	OR282778	OR282772	OR282774
	DY07242	_	OR263308	OR282779	_	OR282775
	DY091091	-	OR263428	OR282782	-	-
	DY091092	-	OR263431	OR282783	-	-
S. winandae	TBRC17511	-	OM491231	OM687896	OM687901	OM687899
S. winande	TBRC17512	-	OM491232	OM687897	OM687902	OM687900
S. yunnanensis	YFCC 1527	MN576756	MN576812	MN576982	MN576872	MN576926
	YFCC 1824	MN576757	MN576813	MN576983	MN576873	MN576927

Boldface: data generated in this study.

## **Results**

## **Phylogenetic analysis**

To determine the phylogenetic relationship between these fungi and allied species from NCBI we constructed a phylogenetic tree based on Maximum Likelihood (ML) and Bayesian analysis, based on concatenated sequences of five genes included 89 taxa, comprising 4491 characters (SSU: 1047bp, LSU: 849 bp, *TEF*: 945bp, *RPB1*: 717 bp, *RPB2*: 933bp). The multi-gene phylogenetic tree consisted of four genera belonging to the family Cordycipitaceae, including *Akanthomyces, Beauveria, Cordyceps* and *Samsoniella*, with strong support (100%). Statistical support ( $\geq$ 75%/0.75) is shown at the nodes for ML bootstrap support/BI posterior probabilities and the strains' numbers are noted after each species' name (Fig. 1).

In the phylogenetic tree, *Samsoniella* species clustered in a clade easily distinguished from species of *Akanthomyces sensu stricto*, *Beauveria* and *Cordyceps*. Within the *Samsoniella* clade, the majority of *Samsoniella* species grouped together, while only two strains, named as *S. lepidopterorum*, formed a separate branch with a relatively far genetic distance. Furthermore, the



Tree scale: 0.05

Figure 1. Phylogenetic relationships between the genus *Samsoniella* and closely-related species, based on multigene dataset (SSU, LSU, *TEF*, *RPB1* and *RPB2*) for maximum likelihood/ Bayesian method. Note: The ML tree presented here, and the node support rate of the two methods is displayed on the branches. The maximum likelihood support values / Bayesian posterior probabilities value (≥75%/0.75) are shown, and bold lines mean support for the two analyses were 98%. The typical strain of the species is marked with the superscript "T"

seven spider- pathogenic strains (RCEF 0643, RCEF 1481, RCEF 2831, RCEF 2868, RCEF 2588, RCEF 2830, RCEF 2595) and four insect- pathogenic strains (RCEF2590, RCEF 2592, RCEF 2870, RCEF 5406) in this study are located on different branches of the Samsoniella clade. Strains RCEF 0643 and S. alpina were clustered in the same branch (MLBP=98, PP=1.00). Strain RCEF 2592 and RCEF 2595 were grouped with S. erucae clade (MLBP=95, PP=1.00). Strain RCEF 1481 was clustered in the same clade with S. hepiali (MLBP=100,PP=1.00). However, another seven strains formed three independent branches. S. fusiformispora (RCEF 5406 and RCEF 2588) formed a monophyletic group which closely clustered with S. hymenopterorum and S. farinosa with high bootstrap values. S. aranea (RCEF 2831 RCEF 2868, and RCEF 2870) clustered in an independent branch, which was phylogenetically close to S. yunnanensis (MLBP=100,PP=1.00). S. anhuiensis (RCEF 2830 and RCEF 2590) formed an independent sister branch with high support(MLBP=97, PP=0.97). Five-gene phylogenetic analyses suggested that RCEF 0643, RCEF 1481, RCEF 2592, and RCEF 2595 were known species. However, the other seven strains were three new species in Samsoniella.

## Taxonomy

Samsoniella anhuiensis T. Wang, Ming J. Chen & B. Huang, sp. nov. MycoBank No: 849801 Fig. 2

**Etymology.** Named after the location Anhui Province where the species was originally collected.

**Typification.** CHINA. Anhui Province: Xuancheng City, the Jingting Mountains National Forest Park, on a spider attached to a leaf, 15 March 2006, Mingjun Chen & Xueqiu Zhao, holotype XC20060315-06. Sequences from strain RCEF2830 and RCEF2590 have been submitted to GenBank with accession numbers. RCEF2830: SSU = OM268844; LSU = OM268849; *TEF* = OM483865; *RPB1* = OM751889. RCEF2590: SSU = OR978313; LSU = OR978316; *TEF* = OR966516; *RPB1* = OR989964.

**Description.** Sexual morph: Undetermined. Asexual morph: Isaria-like. Synnemata arising from the whole body of spider, white, flexuous, multiple, fleshy, up to 12 mm long, with terminal branched, white conidia produced from the branches of synnemata, powdery and floccose (Fig. 2A). Conidiophores arising from the aerial and prostrate hyphae, solitary and verticillate. Phialides in whorls of 2-5,  $5.0-15.2 \times 1.5-2.3 \mu m$ , smooth-walled, with basal portion swollen to ellipsoidal, tapering into a distinct neck,  $1.8-5.2 \times 0.8-1.2 \mu m$ . Conidia in chains, spherical to elliptical, aseptate, hyaline,  $2.1-3.2 \times 1.3-2.2 \mu m$ .

**Culture characteristics.** Colonies on 1/4 SDAY, attaining a diam 38–42 mm in 14 d at 25 °C. Colonies white, with smooth and neat edge, with high mycelial density at the centrum (Fig. 2B). Reverse pale yellow to yellowish, appears flesh pink at 30 d. Hyphae smooth, septate, hyaline,  $1.5-2.3 \mu m$  width. Erect conidiophores usually arising from aerial hyphae, with phialides in whorls of two to three or occasionally with solitary phialides along the hyphae. Phialides basal portion cylindrical, tapering to a distinct neck,  $4.8-16.0 \mu m$  long,  $1.4-2.0 \mu m$ 



**Figure 2.** Samsoniella anhuiensis **A** fungus on spider **B** colony on SDAY/4 **C** colony on PDA **D**, **F** conidiophores structure and conidia on SDAY/4 **E**, **G** conidiophores structure and conidia on PDA. Scale bars: 15 mm (**B**, **C**); 10 μm (**D**–**G**).

basal width and 0.6–1.0  $\mu$ m distinct neck width. Conidia in (Fig. 2D), smoothwalled, hyaline, spherical to elliptical, ovoid, occasionally pointed at both ends, 2.4–3.2 × 1.5–2.1  $\mu$ m (Fig. 2F). Chlamydospores and synnemata not observed.

Colonies on PDA, 39–41mm diameter in 14 d at 25 °C, white. The central part of the colony is raised and appears light yellowish (Fig. 2C). Reverse yellowish in the center. Hyphae smooth, septate, hyaline, with septum and branches, 1.5–2.8  $\mu$ m width, with phialides in whorls of two to five. Phialides basal portion cylindrical, tapering to a distinct neck, (7-)8–11.5(-13)  $\mu$ m long, 1.3–2.2  $\mu$ m basal width and 0.5–0.8  $\mu$ m distinct neck width (Fig. 2E). Conidia in chains, 1-celled, smooth-walled, hyaline, fusiform, elliptical, to obovate, 2–3(-3.5) × 1–2.5  $\mu$ m (Fig. 2G).

Habitat. Occurring on spider attached to the upperside of tree leaf.

**Notes.** Samsoniella anhuiensis was easily identified as belonging to Samsoniella based on the phylogenetic analyses (Fig. 1). Based on the combined multigene dataset, *S. anhuiensis* has an independent branch and has a close relationship with *S. tiankengensis*. However, colonies of *S. tiankengensis* exhibit a faster growth rate on PDA compared to *S. anhuiensis*, displaying white to light pink colonies with a light yellowish reverse. In contrast, colonies of *S. anhuiensis* appear light yellowish and take on a flesh-pink hue at 30 days on 1/4 SDAY, with a yellowish center in reverse. Notably, *S. anhuiensis* distinguishes itself from S. tiankengensis through the presence of larger spherical, elliptical to ovoid conidia (Table 2).

<b>Table</b> 2. Morph	ological comparison of thre	e new species	with other	related Samsoniella :	species (Wang et al. 2022).			
				Morphological cha	Iracteristics			
Species	Synnemata (mm)	Conidiophores (µm)	Colony growth rate (mm)(14d, 25 °C)	Phialide	Phialides size (μm)	Conidia (µm)	Hosts/ substrates	Reference
S. anhuiensis	white, flexuous, multiple, fleshy, up to 12, with terminal branched	1	39-41	verticillate, in whorls of 2 to 5	8.0-11.5 × 1.3-2.2, , wide (apex) 0.5-0.8, basal portion cylindrical to narrowly lageniform	Fusiform, spherical, to obovate 2.0-3.5 × 1.0-2.5	spider	this study
S. alpina	irregularly branched, 3–20 long, cylindrical or clavate stipes with white powdery heads	3.1-6.5 × 1.6-2.8	up to 40	verticillate on conidiophores, solitary or verticillate on hyphae	4.7-9.5 × 1.9-3.1, wide (apex) 0.5-1.1, basal portion cylindrical to narrowly lageniform	fusiform or oval 2.0-3.1 × 1.3-2.1	larvae of Hepialus baimaensis	Wang et al. 2020a
S. aranea	Synnemata not observed	1	34.5-36	verticillate, in whorls of 2 to 4	<ul><li>6.9-11.2 × 1.4-1.9, wide (apex)</li><li>0.5-0.9, basal portion cylindrical to narrowly lageniform</li></ul>	elliptical, fusiform 1.9–3.4 × 1.2–2.4	spider	this study
S. coleopterorum	Synnemata not observed	1	36-40	verticillate, in whorls of 2 to 4	5.4–9.7 × 1.2–1.8, a cylindrical to ellipsoidal basal portion	fusiform, ellipsoidal or subglobose 1.7–2.5 × 1.2–1.8	Snout beetle Curculionidae	Chen et al. 2020
S. erucae	branched or unbranched, fleshy	1	46-48	solitary or in groups of three	6.8 -13.7 × 1.1 -1.5 with a cylindrical or ellipsoidal basal portion and tapered into a short, distinct neck	fusiform to ellipsoidal 2.3–2.9 × 1.1–1.5	caterpillar Lepidoptera	Chen et al. 2022
S. fusiformispora	multiple, unbranched, 2–3 long	1	36.5-39	verticillate, in whorls of 2 to 5	7.4-16.0 × 1.3-1.9, wide (apex) 0.5-1.0, basal portion cylindrical to narrowly lageniform	fusiform 1.9-3.4 × 1.2-2.4	spider	this study
S. hepiali	branched or unbranched, 5-411ong	4.0-7.6 × 1.4-2.2	50-55	verticillate, in whorls of 2 to 5, solitary or opposite on hyphae	3.5-13.6 x 1.3-2.1, wide (apex) 0.5-1.0, basal portion cylindrical to narrowly lageniform	fusiform or oval 1.8–3.3 × 1.4–2.2	larvae of Hepialus armoricanus	Wang et al. 2020a
S. tiankengensis	branched or unbranched, fleshy	I	53-56	solitary or in groups of four	5.4–10.4 × 1.3–2.2, cylindrical or subellipsoidal basal portion and tapered into a short, distinct neck	ellipsoidal 2.3–2.8 × 1.6–1.8	pupa of Lepidoptera	Chen et al. 2022
S. yunnanensis	gregarious, flexuous, fleshy, 4.0–18.0 long, with terminal branches of $3-7 \times 1.0-2.0$	4.2-23.5 × 1.4-2.3	48-50	verticillate, in whorls of 2 to 7, usually solitary on hyphae	4.5-11.6 × 1.2-2.4, wide (apex) 0.6-1.0, basal portion cylindrical to narrowly lageniform	fusiform or oval 2.0–3.3 × 1.1–2.2	pupa of Limacodidae	Wang et al. 2020a

#### Samsoniella aranea T. Wang, Ming J. Chen & B. Huang, sp. nov.

MycoBank No: 849800 Fig. 3

Etymology. Referring to its host, spider, family Araneae.

**Typification.** CHINA. Anhui Province: Xuancheng City, the Jingting Mountains National Forest Park, on spiders, in the litter layer, 15 March 2006 and 27 April 2006, Mingjun Chen & Xueqiu Zhao, holotype XC20060427-06, ex-holotype XC20060315-12. Sequences from strains RCEF2868, RCEF2831 and RCEF 2870 have been submitted to GenBank with accession numbers: RCEF2868: SSU = OM268846; LSU = OM268851; *TEF* = OM483867; *RPB1* = OM751883; *RPB2* = OM802501. RCEF2831: SSU = OM268845; LSU = OM268850; *TEF* = OM483866; *RPB1* = OM751882; *RPB2* = OM802500. RCEF 2870: SSU = OR978314; LSU = O978317; *TEF* = OR966517; *RPB1* = OR989965; *RPB2* = OR989966.

**Description.** Sexual morph: Undetermined. Asexual morph: Isaria-like. Mycellium on the spider consisting of white, smooth, branched, septate,  $1.6-2.5 \mu m$ diam hyphae (Fig. 3A). Conidiophores solitary, arising from superficial hyphae, smooth, cylindrical, flexuous. Phialides verticillate, in whorl of 2–4, 5.0–12.6 × 1.2–2.3 µm, with basal portion swollen to ellipsoidal, tapering into a distinct neck,  $4.0-6.0 \times 0.8-1.0 \mu m$ . Conidia in chains, fusiform, aseptate, hyaline, 2.1–  $3.6 \times 1.5-2.4 \mu m$ .

**Culture characteristics.** Colonies on 1/4 SDAY, attaining a diam of 34.5–41.0 mm in 14 d at 25 °C, floccose, colonies white to cream-yellowish, with white smooth and neat edge (Fig. 3B), reverse light yellowish, sporulating abundantly. Hyphae smooth-walled, branched, hyaline, septate,  $1.5-2.3 \mu$ m wide. Conidiophores smooth-walled, cylindrical, verticillate,  $4.8-16.0 \times 1.4-2.0 \mu$ m. Phialides in whorls of two to four, usually solitary on hyphae, basal portion cylindrical, tapering to a distinct neck;  $5.1-16.9 \mu$ m long,  $1.3-2.1 \mu$ m wide at the base, and  $0.5-1.0 \mu$ m wide at the apex (Fig. 3D). Conidia in chains, smooth-walled, hyaline, elliptical, occasionally fusiform,  $1.9-3.5 \times 1.4-2.6 \mu$ m (Fig. 3G). Chlamydospores and synnemata not observed.

Colonies on PDA, attaining a diam of 34.5-36 mm in 14 d at 25 °C, floccose, colonies white to cream-yellowish, with a white smooth and neat edge, forming radial folds from the center outwards (Fig. 3C). Reverse yolk yellowish, sporulating abundantly. Hyphae smooth walled, branched, hyaline, septate,  $1.5-2.6 \mu$ m wide. Conidiophores smooth – walled, cylindrical, verticillate. Phialides in whorls of two to four, usually solitary on hyphae, basal portion cylindrical, tapering to a distinct neck;  $6.9-11.2 \mu$ m long,  $1.4-1.9 \mu$ m wide at the base, and  $0.5-0.9\mu$ m wide at the apex (Fig. 3E). Conidia 1-celled, in chains, smooth-walled, hyaline, elliptical, occasionally fusiform,  $1.9-3.4 \times 1.2-2.4 \mu$ m (Fig. 3F).

Habitat. Occurring on spiders in the litter layer.

**Notes.** Samsoniella aranea was readily classified within the genus Samsoniella through phylogenetic analyses (Fig. 1). Analysis of the combined multigene dataset unveiled that *S. aranea* forms an independent branch and shares a close relationship with *S. yunnanensis*. However, notable distinctions were observed between the two species. Unlike *S. yunnanensis*, where synnemata arise from insect cocoons, synnemata of *S. aranea* were not observed. Additionally, distinct growth characteristics were noted, with colonies of *S. yunnanensis* exhibiting a faster growth rate on PDA compared to *S. aranea*.



**Figure 3.** Samsoniella aranea **A** fungus on spider **B** colony on SDAY/4 **C** colony on PDA **D**, **G** conidiophores structure and conidia on SDAY/4 **E**, **F** conidiophores structure and conidia on PDA. Scale bars: 15 mm (**B**, **C**); 10 μm (**D**–**G**).

Morphological differences were evident in the colonies on PDA, with *S. aranea* colonies being floccose, white to cream-yellowish, and having a yolk-yellowish reverse. On the other hand, colonies of *S. yunnanensis* were described as loose and hairy, appearing white with a reddish-brown reverse.

Samsoniella fusiformispora T. Wang, Ming J. Chen & B. Huang, sp. nov. MycoBank No: 849799 Fig. 4

Etymology. Referring to the typical fusiform conidia.

**Typification.** CHINA. Gansu Province: Tianshui City, Maiji National Forest Park, on a spider, underside of tree leaf, 22 September 2010, Wang Liming, holotype MJS20100922-21. Sequences from strain RCEF5406 and RCEF2588 submitted to GenBank with accession numbers. RCEF5406: SSU = OM268843; LSU = OM268848; *TEF* = OM483864; *RPB1* = OM751890. RCEF2588: SSU = OR978312; LSU = OR978315; *TEF* = OR966515.

**Description.** Sexual morph: Undetermined. Asexual morph: Isaria-like. Synnemata multiple, unbranched, arising from the whole body of spider, 3-6 mm long, Stipes cylindrical or clavate, 0.5-1.0 mm wide, pale yellowish, white conidia produced from the synnemaya and hyphal layer (Fig. 4A). Phialides verticillate, in whorl of 2-5,  $5.0-12.0 \times 1.9-2.8 \mu$ m, with basal portion swollen



**Figure 4.** Samsoniella fusiformispora **A** fungus on spider **B** colony on SDAY/4 **C** colony on PDA **D**, **F** conidiophores structure and conidia on SDAY/4 **E**, **G** conidiophores structure and conidia on PDA. Scale bars: 15 mm (**B**, **C**); 10 μm (**D**–**G**).

to ellipsoidal, tapering into a distinct neck, 2.3 -3.8 × 0.5–1.2  $\mu$ m. Conidia in chains, fusiform, aseptate, hyaline, 2.1–3.5 × 1.6–2.2  $\mu$ m.

**Culture characteristics.** Colonies on 1/4 SDAY fast-growing, 39.5–44 mm diameter in 14 d at 25 °C, colonies white edge to yellowish center, cottony (Fig. 4B), reverse yellow to orange-yellow, hyphae smooth – walled, branched, hyaline, septate,  $1.7-2.6 \mu$ m wide. Conidiophores smooth-walled, cylindrical, verticillate. Phialides in whorls of three to five, usually solitary on hyphae, basal portion cylindrical, tapering to a distinct neck;  $7.6-15 \mu$ m long,  $1.9-2.6 \mu$ m wide at the base, and  $0.7-1.2 \mu$ m wide at the apex (Fig. 4D). Conidia in chains, smooth-walled, hyaline, fusiform,  $2.1-3.6(-4.4) \times 1.8-2.2 \mu$ m (Fig. 4F). Chlamydospores and synnemata not observed. Size and shape of phialides and conidia similar in culture. Sexual state not observed.

Colonies on PDA, attaining a diam of 36.5-39 mm in 14 d at 25 °C, floccose, colonies white to yellowish, with high mycelial density at the centrum (Fig. 4C). Reverse pale yellowish edge to orang center. Hyphae smooth- walled, branched, hyaline, septate,  $1.5-2.5 \mu$ m wide. Conidiophores smooth – walled, cylindrical, verticillate. Phialides in whorls of two to five, usually solitary on hyphae, basal portion cylindrical, tapering to a distinct neck;  $7.4-16(-26) \mu$ m long,  $1.3-1.9(-2.4) \mu$ m wide at the base, and  $0.5-1.0 \mu$ m wide at the apex (Fig. 4E). Conidia 1-celled, in chains, smooth-walled, hyaline, fusiform,  $1.9-3.4 \times 1.2-2.4 \mu$ m (Fig. 4G).

Habitat. Occurring on spider attached to the underside of tree leaf.

**Notes.** Samsoniella fusiformispora was unequivocally identified as a member of the Samsoniella genus through phylogenetic analyses (Fig. 1) and was found to share a close relationship with *S. coleopterorum*. However, upon further investigation and comparison of the morphological characteristics of the three new species with other related *Samsoniella* species (Table 2), distinct differences emerged. Colonies of *S. fusiformispora* were noted to be white to yellowish, with a pale yellowish edge transitioning to an orange center in reverse. In contrast, colonies of *S. coleopterorum* were observed to be white, with a yellowish reverse.

## Discussion

The typical characteristics of *Samsoniella* were oval to fusiform conidia, bright red-orange stromata of the sexual morphs and synnemata of the asexual morphs (Chen et al. 2020). In this study, we present a phylogenetic investigation of cordycipitaceous *Isaria*-like fungi pathogenic on spiders. Combined with microscopic characteristics and phylogenetic analysis based on multi-locus sequence data, *S. fusiformispora*, *S. aranea* and *S. anhuiensis* were described and illustrated as new species in *Samsoniella*. It was found that the hosts of most reported *Samsoniella* species are Lepidoptera larvae or pupae, while the host of *S. coleopterorum* is a snout beetle (Curculionidae), and the host of *S. hymenopterorum* is a bee (Mongkolsamrit et al. 2018; Chen et al. 2020; Wang et al. 2020a). However, it should be noted that Wang et al. (2020a) described the host of *S. hymenopterorum* as being "Bee, family Vespidae". The family Vespidae are wasps, not bees. Our study has expanded the hosts of *Samsoniella* from Insecta to Arachnida.

Generally, the phialides of *S. fusiformispora* were longer and thinner than those of the closely-related *S. coleopterorum* while they also had bigger typical fusiform conidia with greater length to width ratio. In the ML and BI phylogenetic trees, *S. aranea* was inferred as a phylogenetic sister of *S. yunnanensis* with strong support (93%/1.00) and distinct from other related species in *Samsoniella*. The synnemata of *S. aranea* was not observed, but *S. yunnanensis* has gregarious, flexuous and fleshy synnemata arising from the limacodid cocoons (Wang et al. 2020a). Furthermore *S. yunnanensis* has smaller fusiform to oval conidia than *S. aranea* and the colonies on PDA grow faster than *S. aranea*. Similarly, *S. anhuiensis* was easily separated by the phylogenetic analyses with independent branches in the phylogenetic tree.

Kepler et al. (2017) found that sequences of *Isaria* sp. spat 09-050 and *Isaria* sp. spat 09-051 were firstly obtained, and two strains were clustered as the phylogenetic sister of *Isaria* spp. with 100 bootstrap proportion in the weighted parsimony (WP) analytic tree based on five genes (SSU, LSU, TEF, *RPB1* and *RPB2*), which was classified as *Akanthomyces* group. Then Wang et al. (2020a) constructed the multigene phylogenetic tree studied the new taxa of the family Cordycipitaceae and the new systematic position of the Chinese cordycipitoid fungus *Paecilomyces hepiali*. In this multigene phylogenetic tree, *Isaria* sp. spat 09-050 and *Isaria* sp. spat 09-051 were clustered in genus *Samsoniella* as sister group of *S. vallis* but in two independent branches. In this study, we obtained the same results. We convinced that *Isaria* sp. spat 09-050 and *Isaria* sp. spat

09-051 is an unpublished new species of the *Samsoniella*, should be revised to *Samsoniella* sp. spat 09-050 and *Samsoniella* sp. spat 09-051.

In this study, based on morphological characteristics and five loci phylogenetic analysis, *S. anhuiensis*, *S. aranea* and *S. fusiformispora* were separated from other *Samsoniella* species, which are described here as new species. The strain RCEF0643 was identified as *S. alpina*, the strain RCEF1481 was named as *S. hepiali*, and the strains RCEF2592 and RCEF 2590 was identified as *S. erucae*. Furthermore, our study significantly broadens the host range of *Samsoniella* from Insecta to Arachnida, marking a noteworthy expansion in understanding the ecological associations of these fungi. Additionally, the identification of both mononematous and synnematous conidiophores in our study not only expands the knowledge of *Samsoniella* species but also provides a basis for future research by comparing the ecological significance between these conidiophore types.

# **Additional information**

## **Conflict of interest**

The authors have declared that no competing interests exist.

## **Ethical statement**

No ethical statement was reported.

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## **Author contributions**

MC and BH conceived and designed the study. TW and MC wrote the manuscript, conducted the experiments, and analyzed the data. JL, and XC did a part of the experiments. ZI and NH edited the manuscript. MC and BH edited the manuscript and supervised the project.

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## **Data availability**

All of the data that support the findings of this study are available in the main text.

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