

Discovery of *Cytospora* species associated with canker disease of tree hosts from Mount Dongling of China

Haiyan Zhu^{1,2,3}, Meng Pan¹, Jadson D.P. Bezerra⁴, Chengming Tian¹, Xinlei Fan¹

1 The Key Laboratory for Silviculture and Conservation of Ministry of Education, Beijing Forestry University, Beijing 100083, China **2** State Key Laboratory of Mycology, Institute of Microbiology, Chinese Academy of Sciences, Beijing 100101, China **3** College of Life Sciences, University of Chinese Academy of Sciences, Beijing 100049, China **4** Setor de Micologia, Departamento de Biociências e Tecnologia, Instituto de Patologia Tropical e Saúde Pública (IPTSP), Universidade Federal de Goiás, Rua 235, s/n, Setor Universitário, CEP: 7460505, Goiânia, Goiás, Brazil

Corresponding author: Xinlei Fan (xinleifan@bjfu.edu.cn)

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Abstract

Members of *Cytospora* encompass important plant pathogens, saprobes and endophytes on a wide range of woody hosts with a worldwide distribution. In the current study, we obtained seven representative isolates from six tree hosts of Betulaceae, Juglandaceae, Rosaceae, Tiliaceae and Ulmaceae in Mount Dongling of China. Based on morphological comparison and phylogenetic analyses using partial ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2* gene sequences, we identified two known species (*Cytospora leucostoma* and *C. pruinopsis*) and two novel species (*C. coryli* and *C. spiraeicola*). These results represent the first study on *Cytospora* species associated with canker disease from Mount Dongling of China.

Keywords

Cytosporaceae, phylogeny, taxonomy, wood-inhabiting fungi

Introduction

The genus *Cytospora* was established by Ehrenberg (1818) and belongs to Cytosporaceae, Diaporthales, Sordariomycetes (Wijayawardene et al. 2018, Fan et al. 2020). It is characterised by single or labyrinthine of pycnidial locules, filamentous conidiophores (enteroblastic and phialidic conidiogenous cells) producing hyaline, allantoid conidia

in the asexual morph; diaporthalean-like perithecia, clavate to elongate obovoid ascii with four or eight hyaline, allantoid ascospores in the sexual morph (Spielman 1983, 1985, Adams et al. 2005). Species of *Cytospora* contain important pathogens that cause stem canker and dieback disease on more than 100 species of woody and coniferous plants, thereby causing severe commercial and ecological damage and significant losses worldwide (Sinclair et al. 1987, Adams et al. 2005, 2006, Fan et al. 2014a, b, 2015a, b, Lawrence et al. 2018, Pan et al. 2018, Zhu et al. 2018a, Zhang et al. 2019). Previous *Cytospora* species and their related sexual morphs viz. *Leucostoma*, *Valsa*, *Val-sella* and *Valseutypella* were listed by old fungal literature without any living culture and sufficient evidence for their identification (Fries 1823, Saccardo 1884, Kobayashi 1970, Barr 1978, Sutton 1980, Gvritishvili 1982, Spielman 1983, 1985). Adams et al. (2005) revised the genus *Cytospora* from *Eucalyptus* with 28 species and accepted all sexual genera combined under *Valsa*, either as subgenera or species without additional infrageneric rank. Following the single-name for pleomorphic taxa, *Cytospora* (1818), the older asexual typified name was proposed as the recommended name against *Valsa* (1849), the younger sexual typified name (Fan et al. 2015a, b, Rossman et al. 2015).

Currently, 388 species epithets of *Cytospora* have been recorded in Index Fungorum (2020) (accessed 2 January 2020). However, Kirk et al. (2008) estimated approximately 110 species, but most of them lack herbarium materials, ex-type cultures and DNA sequence data.

Species identification criteria of *Cytospora* were previously carried out by the host-based method and morphology in China; however, these bases are unreliable due to the uninformative illustrations and descriptions, weak host specificity and overlapping morphological characteristics (Teng 1963, Tai 1979, Wei 1979). Recent studies have been able to use multiphase approaches to solve the taxonomy of *Cytospora* (Fan et al. 2014a, b, 2015a, b, Yang et al. 2015, Lawrence et al. 2016, Norphanphoun et al. 2017, Pan et al. 2018, Zhu et al. 2018a, Zhang et al. 2019). Fan et al. (2020) summarised 52 species of *Cytospora* associated with canker and dieback disease in China, using a six gene matrix (ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2*), of which 13 species were newly introduced.

Mount Dongling has high plant diversity in western Beijing, including more than 1,000 tree hosts (Ma et al. 1995). As more plant species were recorded in this region, the exploration of fungal diversity gradually increased as most fungi are often linked to particular host plants as pathogens or endophytes. Species of *Alternaria*, *Diaporthe*, *Leptostroma*, *Pestalotiopsis* and *Phoma* were the most commonly isolated endophytes from *Pinus tabuliformis* and later, an additional 38 endophytic taxa were identified from *Acer truncatum* from Mount Dongling (Guo et al. 2008, Sun et al. 2011). Further, pathogens belonging in Botryosphaerales have been identified from Mount Dongling, including five species from *Aplosporella*, *Botryosphaeria* and *Phaeobotryon* (Zhu et al. 2018b). Zhu et al. (2019) subsequently introduced six species of diaporthalean fungi residing in four families (viz. Diaporthaceae, Erythrogloeaceae, Juglanconidaceae and Melanconidaceae) from Mount Dongling. For the current understanding, many common host plants represent high fungal diversity causing canker and dieback disease in Mount Dongling. *Juglans mandshurica* and *J. regia* (Juglandaceae) were infected by *Botryosphaeria dothidea* (Botryosphaeriaceae), *Diaporthe eres*,

D. rostrata (Diaporthaceae) and *Juglanconis oblonga* (Juglanconidaceae). *Rhus typhina* (Anacardiaceae) was infected by *Aplosporella ginkgonis*, *A. javeedii* (Aplosporellaceae), *Phaeobotryon rhois* and *P. rhoinum* (Botryosphaeriaceae). *Quercus mongolica* (Fagaceae) was infected by *Dendrostoma donglinensis* (Erythrogloeaceae) (Zhu et al. 2018b, 2019).

During the course of cognitive practices to investigate forest pathogens that cause canker or dieback disease in Mount Dongling of China, seven *Cytospora* strains were obtained from six unrelated hosts, i.e. *Corylus mandshurica* (Betulaceae), *Juglans mandshurica* (Juglandaceae), *Prunus sibirica*, *Spiraea salicifolia* (Rosaceae), *Tilia nobilis* (Tiliaceae) and *Ulmus pumila* (Ulmaceae). Phylogenetic analyses inferred from combined ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2* gene regions were conducted to provide a multi-gene phylogeny for *Cytospora*, based on a large set of freshly collected specimens in Mount Dongling of China. Thus, the current study aims to clarify the systematics and taxonomy of *Cytospora* species with detailed descriptions and illustrations and compare it to known species in the genus.

Materials and methods

Sampling and isolation

Seven infected branches of six hosts were collected from Mount Dongling of China (Table 1). Sampled trees expressed general symptoms and signs of canker diseases including elongate, slightly sunken and discoloured areas in the bark, several prominent dark sporocarps immersed in bark, erumpent through the surface of bark when mature (Fig. 1). A total of seven isolates was established by removing a mucoid spore mass from conidiomata or ascomata of fresh material, spreading the suspension on the surface of 1.8 % potato dextrose agar (PDA) and incubating at 25 °C for up to 24 h. Single germinating spores were transferred on to fresh PDA plates. Specimens and isolates were deposited in the Key Laboratory for Silviculture and Conservation of the Ministry of Education in Beijing Forestry University (BJFU) and at the working Collection of X.L. Fan (CF), housed at the BJFU. Axenic cultures are maintained in the China Forestry Culture Collection Centre (CFCC).



Figure 1. Disease symptoms associated with *Cytospora* species. **A** *Corylus mandshurica* **B** *Spiraea salicifolia* **C** *Ulmus pumila* **D** *Prunus sibirica*.

Table I. Isolates and GenBank accession numbers used in the phylogenetic analyses of *Cytospora*.

Species	Strain ¹	Host	Origin	GenBank accession numbers			
				ITS	LSU	act	rpb2
<i>Cytospora ailanthicola</i>	CFCC 89970 ^T	<i>Alanthus altheifolia</i>	Ningxia, China	MH933618	MH933653	MH933526	MH933494
<i>Cytospora amphiliformis</i>	MFLUCC 16-0583 ^T	<i>Sorbus intermedia</i>	Russia	KY417726	KY417760	KY417794	MH935565
	MFLUCC 16-0629	<i>Acer platanoides</i>	Russia	KY417727	KY417761	KY417795	NA
<i>Cytospora angulata</i>	CBS 144233 ^T	<i>Prunus dulcis</i>	California, USA	MG971853	NA	MG972002	NA
<i>Cytospora atracirrhiza</i>	CFCC 89615	<i>Juglans regia</i>	Qinghai, China	KR045618	KR045700	KF498673	KP310858
	CFCC 89616	<i>Juglans regia</i>	Qinghai, China	KR045619	KR045701	KF498674	KP310859
<i>Cytospora boilliniensis</i>	CFCC 50493 ^T	<i>Pinus armmandii</i>	Beijing, China	MH933619	MH933654	MH933527	NA
	CFCC 50494	<i>Pinus armmandii</i>	Beijing, China	MH933620	MH933655	MH933528	NA
<i>Cytospora berberidis</i>	CFCC 89927 ^T	<i>Berberis dasystachys</i>	Qinghai, China	KR045620	KR045702	KU710990	KU710948
	CFCC 89933	<i>Berberis dasystachys</i>	Qinghai, China	KR045621	KR045703	KU710991	KU710949
<i>Cytospora bungeana</i>	CFCC 50495 ^T	<i>Pinus bungeana</i>	Shanxi, China	MH933621	MH933656	MH933529	MH933593
	CFCC 50496	<i>Pinus bungeana</i>	Shanxi, China	MH933622	MH933657	MH933530	MH933594
<i>Cytospora californica</i>	CBS 144234 ^T	<i>Juglans regia</i>	California, USA	MG971935	NA	MG972083	NA
<i>Cytospora carbonacea</i>	CFCC 89947	<i>Ulmus pumila</i>	Qinghai, China	KR045622	KP310812	KP310842	KP310855
<i>Cytospora carpophrotri</i>	CMW 48981 ^T	<i>Carpobrotus edulis</i>	South Africa	MH382812	MH411216	NA	MH411212
<i>Cytospora celtidicola</i>	CFCC 50497 ^T	<i>Celtis sinensis</i>	Anhui, China	MH933623	MH933658	MH933531	MH933499
	CFCC 50498	<i>Celtis sinensis</i>	Anhui, China	MH933624	MH933659	MH933532	MH933500
<i>Cytospora centruroides</i>	MFLUCC 16-1206 ^T	<i>Sorbus domestica</i>	Italy	MF190122	MF190068	NA	NA
	MFLUCC 17-1660	<i>Sorbus domestica</i>	Italy	MF190123	MF190069	NA	NA
<i>Cytospora cerasasperma</i>	CFCC 89624	<i>Juglans regia</i>	Gansu, China	KR045645	KR045724	NA	KU710976
	CFCC 89625	<i>Juglans regia</i>	Gansu, China	KR045646	KR045725	NA	KU710977
<i>Cytospora ceratospernopsis</i>	CFCC 89626 ^T	<i>Juglans regia</i>	Shaanxi, China	KR045647	KR045726	KU711011	KU710978
	CFCC 89627	<i>Juglans regia</i>	Shaanxi, China	KR045648	KR045727	KU711012	KU710979
<i>Cytospora chrysosperma</i>	CFCC 89629	<i>Salix psammophila</i>	Shaanxi, China	KF765673	KF765689	NA	KF765705
	CFCC 89981	<i>Populus alba</i> subsp. <i>pyramidalis</i>	Gansu, China	MH933625	MH933660	MH933533	MH933597
<i>Cytospora eoryti</i>	CFCC 89982	<i>Ulmus pumila</i>	Tibet, China	KP281261	KP310805	KP310835	NA
<i>Cytospora cotini</i>	MFLUCC 14-1050 ^T	<i>Corylus mandshurica</i>	Beijing, China	MN854450	MN854661	MN850751	MN861120
	MFLUCC 15-0865 ^T	<i>Salix alba</i>	Russia	KX430142	NA	KX430144	NA
<i>Cytospora curvata</i>				KY417728	KY417762	KY417796	NA

Species	Strain ¹	Host	Origin	GenBank accession numbers			
				ITS	LSU	act	rpb2
<i>Cytospora davidianna</i>	CXY 1350 ^T	<i>Populus davidianna</i>	Inner Mongolia, China	KM034870	NA	NA	NA
	CXY 1374	<i>Populus davidianna</i>	Heilongjiang, China	KM034869	NA	NA	NA
<i>Cytospora elaeagni</i>	CFCC 89632	<i>Elaeagnus angustifolia</i>	Ningxia, China	KR045626	KR045706	KU710955	KU710918
	CFCC 89633	<i>Elaeagnus angustifolia</i>	Ningxia, China	KF765677	KF765693	KU710996	KU710919
<i>Cytospora elaeagnicola</i>	CFCC 52882	<i>Elaeagnus angustifolia</i>	Xinjiang, China	MK732341	MK732338	MK732344	KR045668
	CFCC 52883	<i>Elaeagnus angustifolia</i>	Xinjiang, China	MK732342	MK732339	MK732345	NA
<i>Cytospora erumpens</i>	CFCC 52884	<i>Elaeagnus angustifolia</i>	Xinjiang, China	MK732343	MK732340	MK732346	NA
	CFCC 50022	<i>Prunus padus</i>	Shanxi, China	MH933627	MH933661	MH933534	NA
<i>Cytospora eucaedipti</i>	MFLUCC 16-0580 ^T	<i>Salix × fragilis</i>	Russia	KY417733	KY417767	KY417699	KY417801
	CBS 144241	<i>Eucalyptus globulus</i>	California, USA	MG971907	NA	MG972056	NA
<i>Cytospora euonymicola</i>	CFCC 50499 ^T	<i>Euonymus kiautschovicus</i>	Shaanxi, China	MH933628	MH933662	MH933535	MH933570
<i>Cytospora euonymina</i>	CFCC 50500	<i>Euonymus kiautschovicus</i>	Shaanxi, China	MH933629	MH933663	MH933536	MH933598
	CFCC 89993 ^T	<i>Euonymus kiautschovicus</i>	Shaanxi, China	MH933630	MH933664	MH933537	MH933600
	CFCC 89999	<i>Euonymus kiautschovicus</i>	Shaanxi, China	MH933631	MH933665	MH933538	MH933601
<i>Cytospora fraxinigena</i>	MFLUCC 14-0868 ^T	<i>Fraxinus ornus</i>	Italy	MF190133	MF190078	NA	NA
	MFLU 17-0880	<i>Fraxinus ornus</i>	Italy	MF190134	MF190079	NA	NA
<i>Cytospora fagax</i>	CXY 1371	NA	NA	KM034852	NA	NA	NA
	CXY 1381	NA	NA	KM034853	NA	NA	NA
<i>Cytospora galactica</i>	CFCC 89620 ^T	<i>Juglans regia</i>	Qinghai, China	KR045628	KR045708	KU710997	KU710957
	CFCC 89621	<i>Juglans regia</i>	Qinghai, China	KR045629	KR045709	KU710998	KU710958
<i>Cytospora gigaspora</i>	CFCC 50014	<i>Juniperus procumbens</i>	Shanxi, China	KR045630	KR045710	KU710999	KU710959
	CFCC 89634 ^T	<i>Salix psammophila</i>	Shaanxi, China	KF765671	KF765687	KU711000	KU710960
<i>Cytospora granatii</i>	CBS 144237 ^T	<i>Punica granatum</i>	California, USA	MG971799	NA	MG971949	NA
<i>Cytospora hippophae</i>	CFCC 89639	<i>Hippophaë rhamnoides</i>	Gansu, China	KR045632	KR045712	KU711001	KU710961
	CFCC 89640	<i>Hippophaë rhamnoides</i>	Gansu, China	KF765682	KF765730	KU710962	KP310865
<i>Cytospora japonica</i>	CFCC 89956	<i>Prunus cerasifera</i>	Ningxia, China	KR045624	KR045704	KU710993	KU710953
	CFCC 89960	<i>Prunus cerasifera</i>	Ningxia, China	KR045625	KR045705	KU710994	KU710954
<i>Cytospora jaquinensis</i>	CBS 144235 ^T	<i>Populus deltoides</i>	California, USA	MG971895	NA	MG972044	NA
<i>Cytospora junipericola</i>	BBH 42444	<i>Juniperus communis</i>	Italy	MF190126	MF190071	NA	ME377579

Species	Strain ¹	Host	Origin	GenBank accession numbers			
				ITS	LSU	act	rpb2
<i>Cytopora juniperola</i>	MFLU 17-0882 ^T	<i>Juniperus communis</i>	Italy	MF190125	MF190072	NA	MF377580
	CFCC 50501 ^T	<i>Juniperus przewalskii</i>	Sichuan, China	MH933632	MH933666	MH933539	MH933507
	CFCC 50502	<i>Juniperus przewalskii</i>	Sichuan, China	MH933633	MH933667	MH933540	MH933508
	CFCC 50503	<i>Juniperus przewalskii</i>	Sichuan, China	MH933634	MH933668	MH933541	MH933509
<i>Cytopora kantschavellii</i>	CXY 1383	<i>Populus maximowiczii</i>	Jilin, China	KM034867	NA	NA	NA
	CXY 1386	<i>Populus maximowiczii</i>	Chongqing, China	KM034867	NA	NA	NA
<i>Cytopora leucosperma</i>	CFCC 89622	<i>Pyrus brieschneideri</i>	Gansu, China	KR045616	KR045698	KU710988	KU710944
	CFCC 89894	<i>Pyrus brieschneideri</i>	Qinghai, China	KR045617	KR045699	KU710989	KU710945
<i>Cytopora leucostoma</i>	CFCC 50015	<i>Sorbus aucuparia</i>	Ningxia, China	KR045634	KR045714	KU711002	NA
	CFCC 50016	<i>Sorbus aucuparia</i>	Ningxia, China	MH820400	MH820408	NA	MH820404
	CFCC 50017	<i>Prunus cerasifera</i>	Ningxia, China	MH933635	MH933669	MH933542	MH933510
	CFCC 50018	<i>Prunus serrulata</i>	Gansu, China	MH933636	MH933670	MH933543	MH933511
	CFCC 50019	<i>Rosa helvetica</i>	Gansu, China	MH933637	MH933671	MH933544	NA
	CFCC 50020	<i>Prunus persica</i>	Gansu, China	MH933638	MH933672	MH933545	NA
	CFCC 50021	<i>Prunus salicina</i>	Gansu, China	MH933639	MH933673	MH933546	NA
	CFCC 50023	<i>Cornus alba</i>	Shanxi, China	KR045635	KR045715	KU711003	KU710964
	CFCC 50024	<i>Prunus pseudocerasus</i>	Qinghai, China	MH933640	MH933674	MH933547	MH933505
	CFCC 50467	<i>Betula platyphylla</i>	Beijing, China	KT732948	KT732967	NA	NA
	CFCC 50468	<i>Betula platyphylla</i>	Beijing, China	KT732949	KT732968	NA	NA
	CFCC 53140	<i>Prunus sibirica</i>	Beijing, China	MN854445	MN854656	MN850760	MN850746
	CFCC 53141	<i>Prunus sibirica</i>	Beijing, China	MN854446	MN854657	MN850761	MN850747
	CFCC 53156	<i>Juglans mandshurica</i>	Beijing, China	MN854447	MN854658	MN850762	MN850748
	MFLUCC 16-0574	<i>Rosa</i> sp.	Russia	KY417731	KY417764	KY417798	NA
	MFLUCC 16-0589	<i>Salix alba</i>	Russia	KY417732	KY417766	KY417800	NA
	MFLUCC 16-0628 ^T	<i>Salix × fragilis</i>	Russia	KY417734	KY417768	KY417802	NA
<i>Cytopora longistriolata</i>	CBS 144236 ^T	<i>Prunus domestica</i>	California, USA	MG971905	NA	MG971615	MG971764
<i>Cytopora longispora</i>	MFLUCC 17-0508 ^T	<i>Lumnitzeracervicola</i>	Thailand	MG975778	MH253461	MH253457	NA
<i>Cytopora lamniticervicola</i>	CFCC 50028	<i>Malus pumila</i>	Gansu, China	MH933641	MH933675	MH933548	MH933513
	CFCC 50029	<i>Malus pumila</i>	Ningxia, China	MH933642	MH933676	MH933549	MH933514
	CFCC 50030	<i>Malus pumila</i>	Shaanxi, China	MH933643	MH933677	MH933568	MH933524
	CFCC 50031	<i>Crataegus</i> sp.	Shaanxi, China	KR045636	KR045716	KU711004	KU710927
	CFCC 50044	<i>Malus baccata</i>	Qinghai, China	KR045637	KR045717	KU711005	KR045678

Species	Strain ¹	Host	Origin	GenBank accession numbers			
				ITS	LSU	act	rpB2
<i>Cytospora melnikii</i>	CFCC 89984	<i>Rhus typhina</i>	Xinjiang, China	MH933644	MH933678	MH933551	MH933515
	MFLUCC 15-0851 ^T	<i>Malus domestica</i>	Russia	KY417735	KY417769	KY417701	KY417803
	MFLUCC 16-0635	<i>Populus nigra</i> var. <i>italica</i>	Russia	KY417736	KY417770	KY417702	KY417804
<i>Cytospora nivea</i>	MFLUCC 15-0860	<i>Salix acutifolia</i>	Russia	KY417737	KY417771	KY417703	KY417805
	CFCC 89641	<i>Elaeagnus angustifolia</i>	Ningxia, China	KF765683	KF765699	KU711006	KU710967
	CFCC 89643	<i>Salix psammophila</i>	Shaanxi, China	KF765685	KF765701	NA	KU710968
	CBS 144248 ^T	<i>Olea europaea</i>	California, USA	MG971944	NA	MG972098	NA
<i>Cytospora palmi</i>	CXY 1276	<i>Cotinus coggygria</i>	Beijing, China	JN402990	NA	NA	KJ781296
	CXY 1280 ^T	<i>Cotinus coggygria</i>	Beijing, China	JN411939	NA	NA	KJ781297
<i>Cytospora parakantschavelii</i>	MFLUCC 15-0857 ^T	<i>Populus × sibirica</i>	Russia	KY417738	KY417772	KY417704	KY417806
	MFLUCC 16-0575	<i>Pyrus pyraster</i>	Russia	KY417739	KY417773	KY417705	KY417807
<i>Cytospora paripistariae</i>	CBS 144506 ^T	<i>Pistacia vera</i>	California, USA	MG971804	NA	MG971954	NA
<i>Cytospora parasitica</i>	MFLUCC 15-0507 ^T	<i>Malus domestica</i>	Russia	KY417740	KY417774	KY417706	KY417808
	XIAU 2542-1	<i>Malus</i> sp.	Xinjiang, China	MH798884	MH798897	NA	NA
<i>Cytospora paramtranslucens</i>	MFLUCC 15-0506 ^T	<i>Populus alba</i> var. <i>bolleiana</i>	Russia	KY417741	KY417775	KY417707	KY417809
	MFLUCC 16-0627	<i>Populus alba</i>	Russia	KY417742	KY417776	KY417708	KY417810
<i>Cytospora pistatiae</i>	CBS 144238 ^T	<i>Pastacia vera</i>	California, USA	MG971802	NA	MG971952	NA
<i>Cytospora planicola</i>	MFLU 17-0327 ^T	<i>Platanus hybrida</i>	Italy	MH253451	MH253452	MH253449	MH253450
<i>Cytospora platyclada</i>	CFCC 50504 ^T	<i>Platycladus orientalis</i>	Yunnan, China	MH933645	MH933679	MH933552	MH933560
	CFCC 50505	<i>Platycladus orientalis</i>	Yunnan, China	MH933646	MH933680	MH933553	MH933611
	CFCC 50506	<i>Platycladus orientalis</i>	Yunnan, China	MH933647	MH933681	MH933554	MH933612
<i>Cytospora platycladica</i>	CFCC 50038 ^T	<i>Platycladus orientalis</i>	Gansu, China	KT2222840	MH933682	MH933555	MH933613
	CFCC 50039	<i>Platycladus orientalis</i>	Gansu, China	KR045642	KR045721	KU711008	KU710973
<i>Cytospora plurivora</i>	CBS 144239 ^T	<i>Olea europaea</i>	California, USA	MG971861	NA	MG972010	NA
	CBS 144240 ^T	<i>Populus deltoides</i>	California, USA	MG971891	NA	MG972040	NA
<i>Cytospora populicola</i>	CFCC 89644 ^T	<i>Salix psammophila</i>	Shaanxi, China	KF765686	KF765702	KU711007	KU710969
<i>Cytospora populinina</i>	CFCC 50032 ^T	<i>Sorbus aucuparia</i>	Ningxia, China	MH933648	MH933683	MH933556	MH933614
<i>Cytospora populinopsis</i>	CFCC 50033	<i>Sorbus aucuparia</i>	Ningxia, China	MH933649	MH933684	MH933557	MH933615
<i>Cytospora pruinopis</i>	CFCC 50034 ^T	<i>Ulmus pumila</i>	Shaanxi, China	KP281259	KP281086	KU710970	KP310849
	CFCC 50035	<i>Ulmus pumila</i>	Jilin, China	KP281260	KP310807	KU710971	KP310850
	CFCC 53153	<i>Ulmus pumila</i>	Beijing, China	MN854451	MN854662	MN850752	MN850759
<i>Cytospora predappiensis</i>	MFLUCC 17-2458 ^T	<i>Platanus hybrida</i>	Italy	MG873484	MG873480	NA	NA

Species	Strain ¹	Host	Origin	GenBank accession numbers			
				ITS	LSU	act	rpb2
<i>Cytopora pruinosa</i>	CFCC 50036	<i>Syringa oblata</i>	Qinghai, China	KP310800	KP310802	KP310832	NA
	CFCC 50037	<i>Syringa oblata</i>	Qinghai, China	MH933650	MH933558	NA	KP310845
<i>Cytopora prunicola</i>	MFLU 17-0995 ^T	<i>Prunus</i> sp.	Italy	MG742350	MG742353	MG742352	MH933522
<i>Cytopora pruniae</i>	CBS 14x244	<i>Prunus granatum</i>	California, USA	MG971943	NA	NA	NA
<i>Cytopora quericola</i>	MFLU17-0881	<i>Quercus</i> sp.	Italy	MF190128	MF190074	NA	MG971798
	MFLUCC 14-0867 ^T	<i>Quercus</i> sp.	Italy	MF190129	MF190073	NA	NA
<i>Cytopora ribis</i>	CFCC 50026	<i>Ulmus pumila</i>	Qinghai, China	KP281267	KP310813	KU710972	KP310856
	CFCC 50027	<i>Ulmus pumila</i>	Qinghai, China	KP281268	KP310814	NA	KP310857
<i>Cytopora rosiae</i>	MFLU 17-0885	<i>Rosa canina</i>	Italy	MF190131	MF190076	NA	NA
<i>Cytopora rostrata</i>	CFCC 89909 ^T	<i>Salix capitularis</i>	Gansu, China	KR045643	KR045643	KU710974	KR045684
	CFCC 89910	<i>Salix capitularis</i>	Gansu, China	KR045644	KR045723	KU711010	KU710933
<i>Cytopora risanovii</i>	MFLUCC 15-0853	<i>Populus × sibirica</i>	Russia	KY417743	KY417777	KY417709	NA
	MFLUCC 15-0854 ^T	<i>Salix babylonica</i>	Russia	KY417744	KY417778	KY417710	NA
<i>Cytopora salicaceum</i>	MFLUCC 15-0861	<i>Salix × fragilis</i>	Russia	KY417745	KY417779	KY417711	NA
	MFLUCC 15-0509 ^T	<i>Salix alba</i>	Russia	KY417746	KY417780	KY417712	NA
	MFLUCC 16-0576	<i>Populus nigra</i> var. <i>italica</i>	Russia	KY417741	KY417775	KY417707	NA
	MFLUCC 16-0587	<i>Prunus cerasus</i>	Russia	KY417742	KY417776	KY417708	KY417813
<i>Cytopora salicicola</i>	MFLUCC 15-0866	<i>Salix alba</i>	Russia	KY417749	KY417783	KY417715	NA
	MFLUCC 14-1052 ^T	<i>Salix alba</i>	Russia	KU982636	KU982635	NA	NA
<i>Cytopora salicina</i>	MFLUCC 15-0862 ^T	<i>Salix alba</i>	Russia	KY417750	KY417784	KY417716	NA
	MFLUCC 16-0637	<i>Salix × fragilis</i>	Russia	KY417751	KY417785	KY417717	NA
<i>Cytopora schulzeri</i>	CFCC 50040	<i>Malus domestica</i>	Ningxia, China	KR045649	KR045728	KU711013	KU710980
	CFCC 50042	<i>Malus asiatica</i>	Qinghai, China	KR045650	KR045729	KU711014	KU710981
<i>Cytopora sibirae</i>	CFCC 50045 ^T	<i>Sibiraea angustata</i>	Gansu, China	KR045651	KR045730	KU711015	KU710982
	CFCC 50046	<i>Sibiraea angustata</i>	Gansu, China	KR045652	KR045731	KU711015	KU710983
	CFCC 50047	<i>Syphnolobium</i> <i>japonicum</i>	Shanxi, China	KR045653	KR045732	KU711017	KU710984
	CFCC 50048	<i>Magnolia grandiflora</i>	Shanxi, China	MH820401	MH820394	MH820409	MH820397
	CFCC 89598	<i>Syphnolobium</i> <i>japonicum</i>	Gansu, China	KR045654	KR045733	KU711018	KU710985

Species	Strain ¹	Host	Origin	GenBank accession numbers			
				ITS	LSU	act	rpB2
<i>Cytospora sphaericola</i>	CFCC 89595 ^T	<i>Syphnolobium japonicum</i> var. <i>pendula</i>	Gansu, China	KR045655	KR045734	KU711019	KU710986
	CFCC 89596	<i>Syphnolobium japonicum</i> var. <i>pendula</i>	Gansu, China	KR045656	KR045735	KU711020	KU710987
<i>Cytospora sphaeriopsis</i>	CFCC 89600 ^T	<i>Syphnolobium japonicum</i>	Gansu, China	KR045623	KP310804	KU710992	KU710951
<i>Cytospora sorbi</i>	MFLUCC 16-0631 ^T	<i>Sorbus aucuparia</i>	Russia	KY417752	KY417786	KY417718	KY417820
<i>Cytospora sorbicola</i>	MFLUCC 16-0584 ^T	<i>Acer pseudoplatanus</i>	Russia	KY417755	KY417789	KY417721	KY417823
	MFLUCC 16-0633	<i>Cotoneaster melanocarpus</i>	Russia	KY417758	KY417792	KY417724	KY417826
<i>Cytospora spinaceae</i>	CFCC 50049 ^T	<i>Spinacea salicifolia</i>	Gansu, China	MG708759	MG707643	MG708196	MG708199
	CFCC 50050	<i>Spinacea salicifolia</i>	Gansu, China	MG707860	MG707644	MG708197	MG708200
<i>Cytospora spinicola</i>	CFCC 53138 ^T	<i>Spiraea salicifolia</i>	Beijing, China	MN854448	MN854659	NA	MN850749
	CFCC 53139	<i>Tilia nobilis</i>	Beijing, China	MN854449	MN854660	NA	MN850750
<i>Cytospora tamariicola</i>	CFCC 50507	<i>Rosa multiflora</i>	Yunnan, China	MH933651	MH933686	MH933559	MH933525
	CFCC 50508 ^T	<i>Tamarix chinensis</i>	Yunnan, China	MH933652	MH933687	MH933560	MH933523
<i>Cytospora tamnifolia</i>	MFLUCC 14-1057 ^T	<i>Betula pubescens</i>	Russia	KT459411	KT459412	KT459413	NA
<i>Cytospora thailandica</i>	MFLUCC 17-0262 ^T	<i>Xylocarpus moluccensis</i>	Thailand	MG975776	MH253463	MH253459	MH253455
	MFLUCC 17-0263 ^T	<i>Xylocarpus moluccensis</i>	Thailand	MG975777	MH253464	MH253460	MH253466
<i>Cytospora tibouchinae</i>	CPC 26333 ^T	<i>Tibouchina semidecandra</i>	France	KX228284	KX228335	NA	NA
<i>Cytospora transluens</i>	CXY 1351	<i>Populus davidiana</i>	Inner Mongolia, China	KM034874	NA	NA	NA
<i>Cytospora ulmi</i>	MFLUCC 15-0863 ^T	<i>Ulmus minor</i>	Russia	KY417759	NA	NA	NA
<i>Cytospora vinacea</i>	CBS 141585 ^T	<i>Vitis interspecific</i> hybrid 'Vidal'	USA	KX256256	NA	NA	KX256277
<i>Cytospora viticola</i>	CBS 141586 ^T	<i>Vitis vinifera</i> 'Cabernet Franc'	USA	KX256239	NA	NA	KX256260
<i>Cytospora xylocarpi</i>	MFLUCC 17-0251 ^T	<i>Xylocarpus granatum</i>	Thailand	MG975775	MH253462	MH253458	MH253454
<i>Diaporthe vaccinii</i>	CBS 160.32	<i>Vaccinium macrocarpon</i>	USA	KC343228	NA	JQ807297	NA

Abbreviations: **BBH**: BIOTEC Bangkok Herbarium, National Science and Technology Development Agency, Thailand; **CBS**: Westerdijk Fungal Biodiversity Institute (CBS-KNAW Fungal Biodiversity Centre), Utrecht, The Netherlands; **CCRC**: China Forestry Culture Collection Centre, Beijing, China; **CMW**: Culture collection of Michael Wingfield, University of Pretoria, South Africa; **CPC**: Culture collection of Pedro Crous, The Netherlands; **MFLUCC**: Mae Fah Luang University herbarium, Thailand; **XJAU**: Xijiang Agricultural University, Xinjiang, China; **NA**: not applicable. All the new isolates used in this study are indicated in bold type and the strains from generic type species are marked by a superscript (T).

Morphological analysis

Species identification was based on morphological features of the ascomata or conidiomata from infected host materials and micromorphology, supplemented by cultural characteristics. Microscopic photographs (structure and size of stromata; structure and size of ectostromatic disc and ostioles) were captured using a Leica stereomicroscope (M205 FA) (Leica Microsystems, Wetzlar, Germany). Microscopic observations (shape and size of conidiophores, ascii and conidia/ascospores) were determined under a Nikon Eclipse 80i microscope (Nikon Corporation, Tokyo, Japan), equipped with a Nikon digital sight DS-Ri2 high definition colour camera, using differential interference contrast (DIC) illumination. The Nikon software NIS-Elements D Package v. 3.00, Adobe Bridge CS v. 6 and Adobe Photoshop CS v. 5 were used for the manual editing. More than 10 conidiomata/ascomata, 10 ascii and 30 conidia/ascospores were measured by Nikon software NIS-Elements D Package v. 3.00 to calculate the mean size/length and respective standard deviations (SD). Colony diameters were measured and the colony features were described using the colour charts of Rayner (1970).

DNA extraction, PCR amplification and sequencing

Fungal mycelium grown on the cellophane of PDA was scraped for the extraction of genomic DNA following a modified CTAB method (Doyle and Doyle 1990). The primers and PCR conditions are listed in Table 2. DNA sequencing was performed using an ABI PRISM 3730XL DNA Analyser with a BigDye Terminator Kit v.3.1 (In-vitrogen, USA) at the Shanghai Invitrogen Biological Technology Company Limited (Beijing, China). The DNA sequences, obtained from forward and reverse primers, were combined using SeqMan v. 7.1.0 in the DNASTAR Lasergene Core Suite software (DNASTAR Inc., Madison, WI, USA).

Phylogenetic analyses

The current isolates were initially identified as *Cytospora* species, based on both morphological observations and BLAST results. To clarify their further phylogenetic position, an analysis, based on the combined six genes (ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2*), was performed to compare *Cytospora* species from the current study with other strains in GenBank. *Diaporthe vaccinii* was selected as the outgroup in all analyses. Subsequent alignments for each gene were generated using MAFFT v.7 (Katoh and Standley 2013) and manually adjusted using MEGA v. 6 (Tamura et al. 2013). Ambiguously aligned sequences were excluded from analysis. Reference sequences were selected, based on ex-type or ex-epitype sequences available from recently published literature (Fan et al. 2014a, b, 2015a, b, 2020, Yang et al. 2015, Lawrence et al. 2016, Norphanphoun et al. 2017, Zhu et al. 2018a, Zhang et al. 2019, Fan et

Table 2. Genes used in this study with PCR primers, primer DNA sequence, optimal annealing temperature and corresponding references.

Locus	Definition	Primers	Primer DNA sequence (5'-3')	Optimal annealing temp (°C)	References of primers used
ITS	internal transcribed spacer of ribosomal RNA	ITS1	TCCGTAGGTGAAACCTGGGG	51	White et al. 1990
		ITS4	TCCTCCGCTTTGATATGC		
LSU	large subunit of ribosomal RNA	LROR	ACCCGCTGAACCTTAAGC	55	Vilgalys and Hester 1990
		LR7	TACTACCACCAAGATCT		
act	actin	ACT-512F	ATGTGCAAGGCCGGTTTCGC	61	Carbone and Kohn 1999
		ACT-783R	TACGAGTCCTTCTGGCCCAT		
rpb2	RNA polymerase II second largest subunit	RPB2-5F	GA(T/C)GA(T/C)(A/C)G(A/T) GATCA(T/C)TT(T/C)GG	52	Liu et al. 1999
		RPB2-7cR	CCCAT(A/G)GCTTG(T/C)TT(A/G) CCCAT		
tef1a	translation elongation factor 1-alpha	EF1-668F	CGGTCACTTGATCTACAAGTGC	55	Alves et al. 2008
		EF1-1251R	CCTCGAACTCACCAGTACCG		
tub2	beta-tubulin	Bt2a	GGTAACCAAATCGGTGCTGCTTG	55	Glass and Donaldson 1995
		Bt2b	ACCCTCAGTGTAGTGACCCTTGGC		

al. 2020) (Table 1). Phylogenetic analyses were performed with PAUP v.4.0b10 for the maximum parsimony (MP) method (Swofford 2003), MrBayes v.3.1.2 for the Bayesian Inference (BI) method (Ronquist and Huelsenbeck 2003) and RAxML for the maximum likelihood (ML) method (Stamatakis 2006).

A partition homogeneity test (PHT) with heuristic search and 1,000 replicates was performed using PAUP v.4.0b10 to test the discrepancy amongst the ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2* sequence datasets in reconstructing phylogenetic trees. MP analysis was performed using a heuristic search option of 1,000 random-addition sequences with a tree bisection and reconnection (TBR) branch swapping algorithm (Swofford 2003). The branches of zero length were collapsed and all equally parsimonious trees were saved. Clade stability was assessed with a bootstrap analysis of 1,000 replicates (Hillis and Bull 1993). Other parsimony scores, such as tree length (TL), consistency index (CI), retention index (RI) and rescaled consistency (RC), were calculated (Swofford 2003). ML analysis was performed with the GTR + G + I model of site substitution following recent studies (Zhu et al. 2018a), including estimation of gamma-distributed rate heterogeneity and a proportion of invariant sites using PhyML v. 3.0 (Guindon et al. 2010). The branch support was evaluated with a bootstrapping method of 1,000 replicates (Hillis and Bull 1993). BI analysis was performed using a Markov Chain Monte Carlo (MCMC) algorithm with Bayesian posterior probabilities (Rannala and Yang 1996). A nucleotide substitution model was estimated by MrModetest v.2.3 (Posada and Crandall 1998) and a weighted Bayesian analysis was considered. Two MCMC chains were run from random trees for 1,000,000 generations and trees were sampled each 100 generations. The first 25% of trees were discarded as the burn-in phase of each analysis and the posterior probabilities (BPP) were calculated

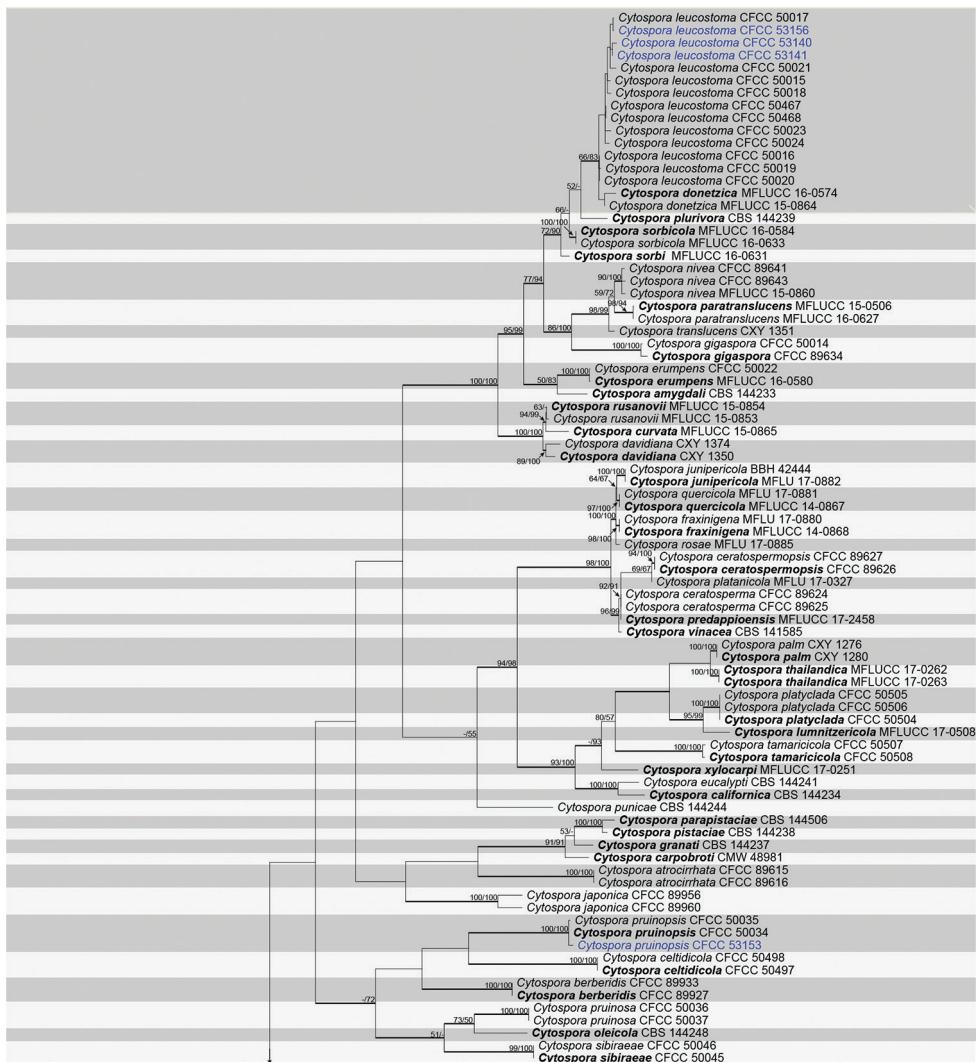


Figure 2. Phylogram of *Cytospora*, based on combined ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2* genes. The MP and ML bootstrap support values above 50% are shown at the first and second positions, respectively. Thickened branches represent posterior probabilities above 0.95 from the BI. Ex-type strains are in bold. Strains from the current study are in blue.

to assess the remaining trees (Rannala and Yang 1996). The branch support from MP and ML analysis was evaluated with a bootstrapping (BS) method of 1,000 replicates (Hillis and Bull 1993). Phylogenograms were plotted in Figtree v. 1.4.4 (<http://tree.bio.ed.ac.uk/software/figtree>) and edited in Adobe Illustrator CS6 v.16.0.0 (<https://www.adobe.com/cn/products/illustrator.html>). Novel sequences, generated in the current study, were deposited in GenBank (Table 1) and the aligned matrices, used for phylogenetic analyses, were submitted in TreeBASE (www.treebase.org; study ID S25564).

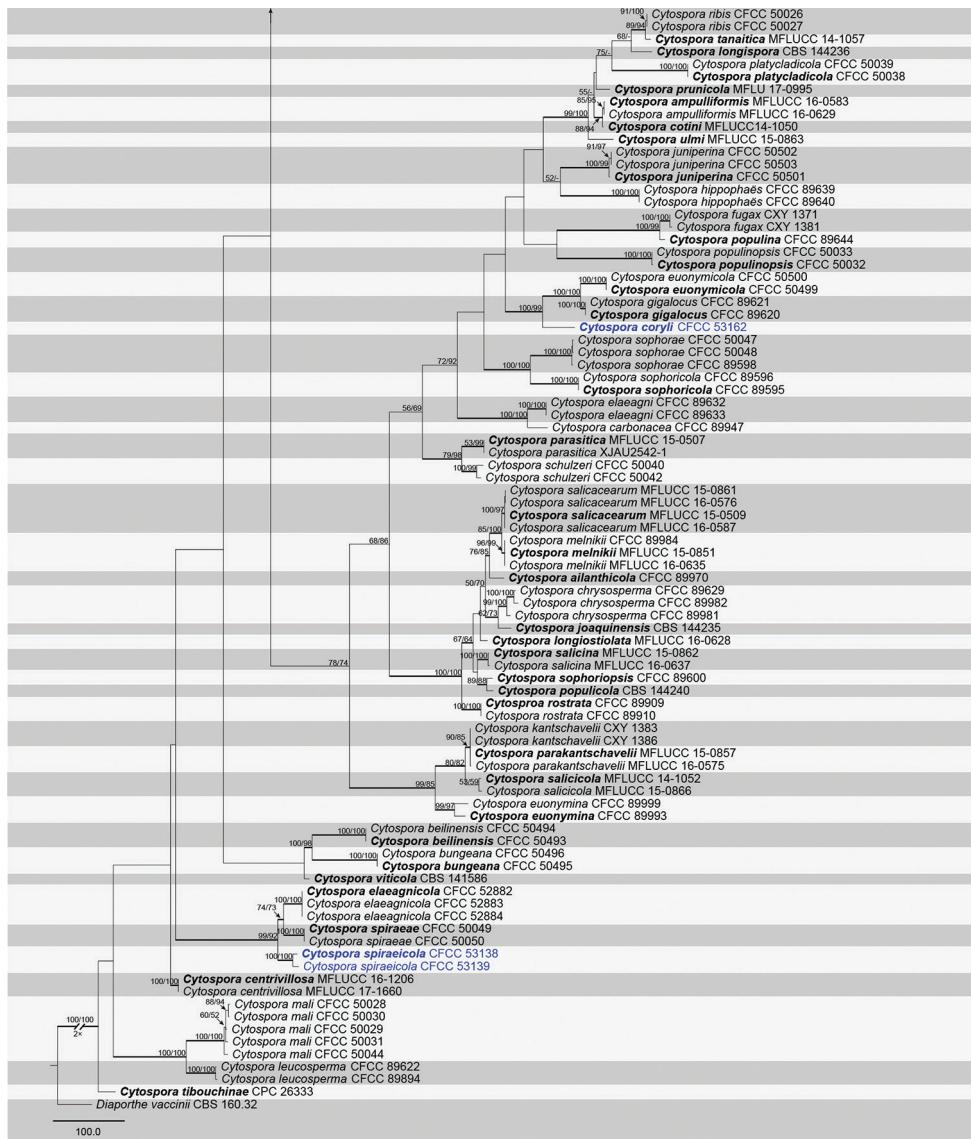


Figure 2. Continued.

Results

Phylogenetic analyses

A combined matrix of six gene sequences of *Cytospora* was considered. The combined alignments matrix (ITS, LSU, *act*, *rpb2*, *tef1-a* and *tub2*) included 172 accessions (seven from this study and 165 retrieved from GenBank) and counted 3,652 characters including gaps (665 characters for ITS, 525 for LSU, 337 for *act*, 730 for *rpb2*, 771 for *tef1-a*

and 624 for *tub2*), of which 2,067 characters were constant, 189 variable characters were parsimony-uninformative and 1,396 (38.22%) characters were variable and parsimony-informative. The MP analysis generated 100 parsimonious trees, the first tree of which is presented in Fig. 2 (TL = 8,029, CI = 0.345, RI = 0.804, RC = 0.278). Tree topologies of ML and BI analyses were similar to the MP tree. Based on the multi-locus phylogeny and morphology, seven strains were assigned to four species within *Cytospora coryli*, *C. leucostoma*, *C. pruinopsis* and *C. spiraeicola*, including two taxa which we describe here as new. The two isolates of *C. spiraeicola* formed a distinct and strongly supported clade (MP/ML/BI = 100/100/1) with close phylogenetic affinity to *C. elaeagnicola* and *C. spiraeae*. The strain of *C. coryli* from *Corylus mandshurica* shared a close relationship to *Cytospora euonymicola* and *C. gigalocus* with 100% MP, 99% ML, 0.99 BI supports.

Taxonomy

Cytospora coryli H.Y. Zhu & X.L. Fan, sp. nov.

Mycobank No: 833820

Fig. 3

Etymology. Named after the host genus on which it was collected, *Corylus*.

Holotype. CHINA, Beijing City, Mentougou District, Mount Dongling, Xiaolongmen Forestry Centre (115°27'07.00"E, 39°59'26.47"N), from branches of *Corylus mandshurica*, 17 Aug 2017, H.Y. Zhu & X.L. Fan, holotype CF 2019813, ex-type living culture CFCC 53162.

Description. Necrotrophic on branches of *Corylus mandshurica*. **Sexual morph:** not observed. **Asexual morph:** Conidiomata pycnidial, flat, immersed in the bark, scattered to gregarious, erumpent through the surface of bark, surrounded by conspicuous black stroma walls in the margin, with multiple locules. Conceptacle absent. Ectostromatic disc grey to black, discoid, circular to ovoid, 270–340 µm in diam., with one ostiole per disc. Ostiole grey to black, at the same or above level as the disc surface, inconspicuous. Locules numerous, subdivided frequently by invaginations with common walls, circular to irregular, 1550–1710 µm in diam. Conidiophores hyaline, branched at the base, in the middle, approximately cylindrical with the top end acute, 15.5–18.5 × 1–2 (av. = 17 ± 1.2 × 1.1 ± 0.2, n = 10) µm, sometimes reduced to conidiogenous cells. Conidiogenous cells enteroblastic, phialidic, sub-cylindrical to cylindrical, 7.5–14 × 1–2 (av. = 9.3 ± 1.7 × 1.4 ± 0.2, n = 10) µm. Conidia hyaline, allantoid, smooth, aseptate, thin-walled, 5–7 × 1–2 (av. = 5.6 ± 0.5 × 1.4 ± 0.2, n = 30) µm.

Culture characteristics. Cultures are initially white with hazel at the centre, growing fast up 9 cm in diam. after 3 days, becoming honey to hazel from the edge to centre after 7–10 days. In reverse, the cultures are the same as the upper colour after 3 days, becoming cinnamon from the edge to centre after 7–10 days. Colonies are flat, sparse at the centre and compact to the margin. Pycnidia distributed radially on colony surface.

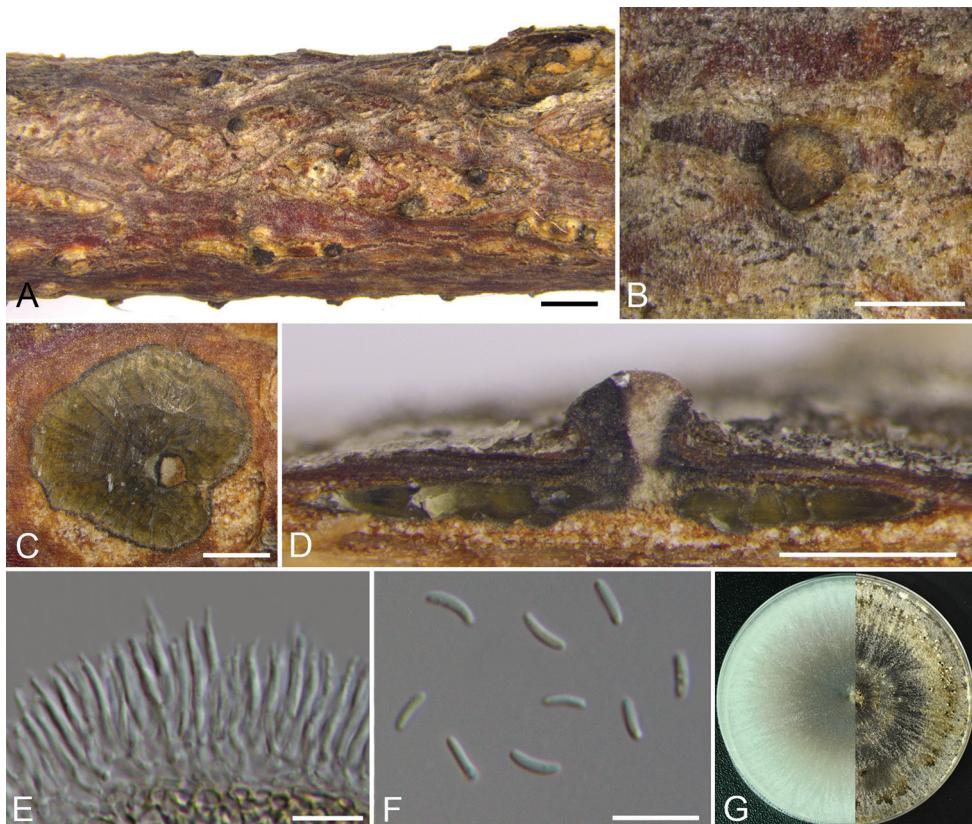


Figure 3. *Cytospora coryli* from *Corylus mandshurica* (CF 2019813). **A, B** habit of conidiomata on twig **C** transverse section of conidioma **D** longitudinal section through conidioma **E** conidiophores and conidiogenous cells **F** conidia **G** colonies on PDA at 3 days (left) and 30 days (right). Scale bars: 1 mm (**A**); 500 µm (**B–D**); 10 µm (**E, F**).

Habitat and distribution. Known from *Corylus mandshurica* in Mount Dongling, China.

Notes. *Cytospora coryli* is associated with canker disease of *Corylus mandshurica* in China. The only strain CFCC 53162 representing *Cytospora coryli* clusters as a single lineage and appears mostly related to *C. euonymicola* from *Euonymus kiautschovicus* and to *Cytospora gigalocus* from *Juglans regia* (Fan et al. 2015a, 2020) (Fig. 2). *Cytospora coryli* differs from *C. euonymicola* by its larger locules (1550–1710 vs. 1150–1400 µm) and larger conidia (5–7 × 1–2 vs. 4.5–5 × 1 µm) (Fan et al. 2020), *C. coryli* differs from *C. gigalocus* by its smaller locules (1550–1710 vs. 1630–2180 µm) with single ostiole (one to five ostioles in *C. gigalocus*) and the larger size of conidia (5–7 × 1–2 vs. 4.6–5.6 × 0.8–1.3 µm) (Fan et al. 2015a). Based on morphology and sequence data, we describe it as a new species.

***Cytopsora leucostoma* (Pers.) Sacc., *Michelia* 2: 264 (1881)**

Figs 4, 5

Sphaeria leucostoma Pers., Ann. Bot. 11: 23 (1794)*Valsa leucostoma* (Pers.) Fr., Summa Veg. Scand., Section Post. (Stockholm): 411 (1849)*Valsa persoonii* Nitschke, Pyrenomyc. Germ. 2: 222 (1870)*Leucostoma persoonii* (Nitschke) Höhn., Mitt. Bot. Inst. Tech. Hochsch. Wien 5: 78 (1928)

[Additional synonyms in Species Fungorum.]

Description. *Necrotrophic* on branches of Betulaceae, Juglandaceae and Rosaceae. **Sexual morph:** *Ascostromata* immersed in the bark, erumpent through the surface of bark, scattered, 950–2550 µm in diam., with 8–10 perithecia arranged circularly to irregularly. *Conceptacle* absent. *Ectostromatic disc* pale grey, fusoid, 600–2150 µm in diam., with 8–10 ostioles arranged irregularly per disc. *Ostioles* numerous, dark grey to black, at the same or above the level as the disc, concentrated, arranged irregularly in a disc, 60–120 µm in diam. *Perithecia* beige with a little black when mature, flask-shaped to spherical, arranged circularly to irregularly, 270–560 µm in diam. *Paraphyses* large, broad and cylindrical with 1–4 septa, 39–78 × 5.8–8.7 (av. = 50.6 ± 13.7 × 7 ± 0.8, n = 10) µm. *Asci* free, clavate to elongate ovoid, 35–45 × 6–8 (av. = 40.4 ± 3.3 × 6.9 ± 0.5, n = 10) µm, 8-spored. *Ascospores* uniseriate to biseriate, elongate-allantoid, thin-walled, hyaline, aseptate, 7–10 × 2–3 (av. = 8.3 ± 0.9 × 2.6 ± 0.2, n = 30) µm. **Asexual morph:** *Conidiomata* pycnidial, immersed in the bark, scattered, erumpent through the surface of bark, with multiple locules and a conspicuous central column. *Central column* beneath the disc more or less conical, brown. *Conceptacle* absent. *Ectostromatic disc* buff, discoid, circular to ovoid, 190–310 µm in diam., with 1–2 ostioles per disc. *Ostioles* grey to black, at the same or above the level as the disc surface, 60–65 µm in diam. *Locules* numerous, subdivided frequently by invaginations with common walls, circular to ovoid, 700–1000 µm in diam. *Conidiophores* hyaline, branched at the base or unbranched, approximately cylindrical, 8–14 × 1–2 (av. = 11.5 ± 1.8 × 1.4 ± 0.2, n = 10) µm, sometimes reduced to conidiogenous cells. *Conidiogenous cells* enteroblastic phialidic, sub-cylindrical to cylindrical, 7–11 × 1–2 (av. = 9 ± 1.4 × 1.5 ± 0.3, n = 10) µm. *Conidia* hyaline, elongate-allantoid, smooth, aseptate, 4.5–6 × 1–2 (av. = 5.4 ± 0.3 × 1.5 ± 0.2, n = 30) µm.

Culture characteristics. *Cultures* initially are white, growing fast up to 8 cm in diam. after 3 days and entirely covering the 9 cm Petri dish after 4 days, becoming greenish-olivaceous after 7–10 days and grey olivaceous after 30 days. In reverse, the cultures are the same as the upper colour after 7 days, becoming olivaceous grey to iron grey after 30 days. *Colonies* are flat with a uniform texture; sterile.

Habitat and distribution. Known from several species of Betulaceae, Juglandaceae and Rosaceae around the world.

Materials examined. CHINA, Beijing City, Mentougou District, Mount Dongling, Xiaolongmen Forestry Centre (115°26'47.36"E, 39°56'06.45"N), from branches of *Prunus sibirica*, 17 Aug 2017, H.Y. Zhu & X.L. Fan, CF 2019814, living culture CFCC



Figure 4. *Cytospora leucostoma* (Sexual morph) from *Prunus sibirica* (CF 2019814). **A, B** habit of ascocarps on twig **C** transverse section of ascocarp **D** longitudinal section through ascocarp **E** ascospores **F** ascus **G** ascospores **H** colonies on PDA at 3 days (left) and 30 days (right). Scale bars: 1 mm (**A**); 500 µm (**B–D**); 10 µm (**E–G**).

53140; *ibid.* CF 2019815, living culture CFCC 53141. CHINA, Beijing City, Mentougou District, Mount Dongling, Xiaolongmen Forestry Centre (115°29'20.52"E, 39°57'47.49"N), from branches of *Juglans mandshurica*, 17 Aug 2017, H.Y. Zhu & X.L. Fan, CF 2019809, living culture CFCC 53156.

Notes. *Cytospora leucostoma* is commonly associated with canker disease of Prunoideae of Rosaceae in China (Fan et al. 2020). Morphologically, our taxa are similar to previous descriptions of *C. leucostoma* in having multi-loculate pycnidial stromata with a conspicuous black conceptacle, producing elongate-allantoid, large conidia (4.5–6 × 1–2 µm) (Teng 1963, Zhuang 2005, Fan et al. 2020). The greenish-yellow of the cultures on PDA medium from *Juglans mandshurica* is similar to descriptions of those

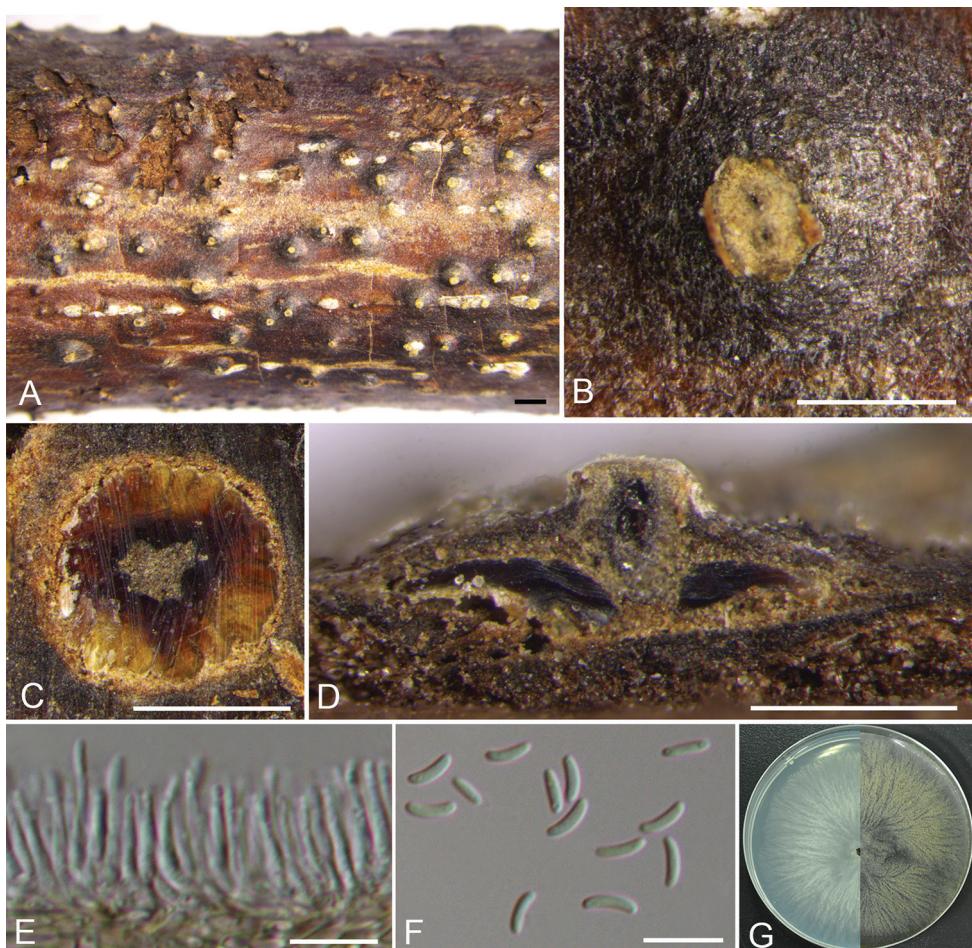


Figure 5. *Cytospora leucostoma* (Asexual morph) from *Juglans mandshurica* (CF 2019809). **A, B** habit of conidiomata on twig **C** transverse section of conidioma **D** longitudinal section through conidioma **E** coidiophores and conidiogenous cells **F** conidia **G** colonies on PDA at 3 days (left) and 30 days (right). Scale bars: 1 mm (**A**); 500 µm (**B–D**); 10 µm (**E, F**).

collected from Prunoideae (Fan et al. 2020). Multigene phylogenetic analyses supported the morphological results with high support values (ML/MP/BI = 100/100/1, Fig. 2). By combining morphology and the DNA data, our isolates collected from dead branches of *Prunus sibirica* and *Juglans mandshurica* belong to this species. The current study represents a new host record of *Juglans mandshurica*.

***Cytospora pruinopsis* C.M. Tian & X.L. Fan, Mycological Progress 14(9): 74 (2015)
Fig. 6**

Description. See Yang et al. (2015).

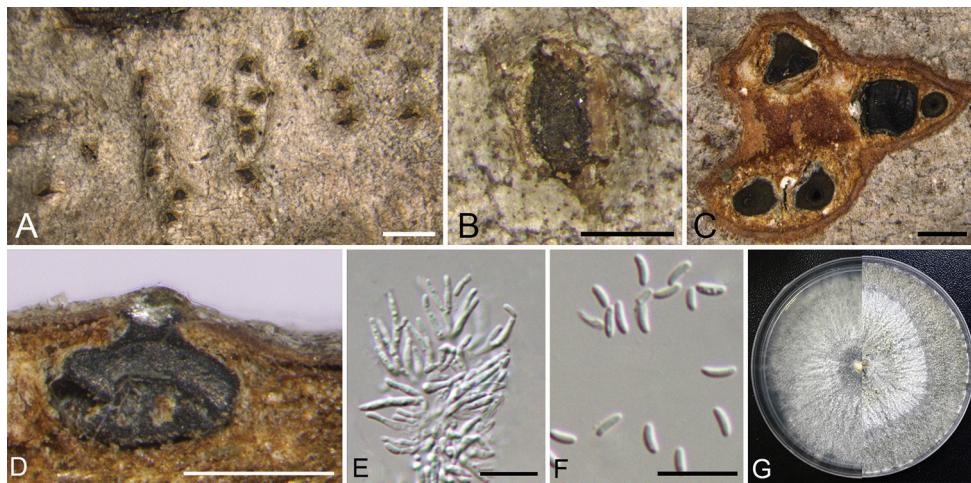


Figure 6. *Cytospora pruinopsis* from *Ulmus pumila* (CF 2019806). **A, B** habit of conidiomata on twig **C** transverse section of conidiomata **D** longitudinal section through conidioma **E** conidiophores and conidiogenous cells **F** conidia **G** colonies on PDA at 3 days (left) and 30 days (right). Scale bars: 1 mm (**A**); 250 µm (**B**); 500 µm (**C, D**); 10 µm (**E, F**).

Material examined. CHINA, Beijing City, Mentougou District, Mount Dongling, Xialongmen Forestry Centre (115°27'29.37"E, 39°56'47.49"N), from branches of *Ulmus pumila*, 22 Aug 2017, H.Y. Zhu & X.L. Fan, CF 2019806, living culture CFCC 53153.

Habitat and distribution. Known from *Ulmus pumila* in Northern China.

Notes. Yang et al. (2015) described *Cytospora pruinopsis* from cankers of *Ulmus pumila* in Shannxi Province of China. The strain CFCC 53153 clusters in a well-supported clade with high support value (MP/ML/BI = 100/100/1), based on combined multi-locus gene phylogenetic analyses (Fig. 2). Morphologically, it confirms *Cytospora pruinosa* in having a single locule and small conidia (2–4 × 1 µm) as per the descriptions of Yang et al. (2015). Phylogenetically, our isolates represent 6/771 nucleotide differences of *tef1-a* comparing with ex-type strains CFCC 50034 of *C. pruinosa*. Morphology and sequence data confirmed that our isolates represent this species.

Cytospora spiraeicola H.Y. Zhu & X.L. Fan, sp. nov.

Mycobank No: 833821

Fig. 7

Etymology. Named after the host genus on which it was collected, *Spiraea*.

Holotype. CHINA, Beijing City, Mentougou District, Mount Dongling, Xialongmen Forestry Centre (115°28'28.52"E, 39°55'49.42"N), from branches of *Spiraea salicifolia*, 17 Aug 2017, H.Y. Zhu & X.L. Fan, holotype CF 2019803, ex-type living culture CFCC 53138.



Figure 7. *Cytospora spiraeicola* from *Spiraea salicifolia* (CF 2019803). **A, B** habit of ascomata on twig **C** transverse section of ascoma **D** longitudinal section through ascoma **E** ascii and ascospores **F, G** ascus **H** ascospores **I** colonies on PDA at 3 days (left) and 30 days (right). Scale bars: 1 mm (**A, B**); 500 µm (**C, D**); 10 µm (**E–H**).

Description. Necrotrophic on branches of *Spiraea salicifolia* and *Tilia nobilis*. **Sexual morph:** Ascostromata immersed in the bark, erumpent through the surface of bark, scattered, with 3–5 perithecia arranged regularly, 660–890 µm in diam. Conceptacle absent. Ectostromatic disc pale grey, usually surrounded by tightly crowded ostiolar necks, quadrangular, 240–350 µm in diam., with 5–8 ostioles arranged regularly per disc. Ostioles numerous, dark grey to black, at the same or above the level as the disc, concentrated, arranged regularly in a disc, 25–40 µm in diam. Perithecia dark grey to black, flask-shaped to spherical, arranged circularly, 210–250 µm in diam. Paraphyses lacking. Ascii free, clavate to elongate, obovoid, 26–37 × 7.5–9 (av. = $33 \pm 2.5 \times 8.3 \pm 0.9$, $n = 10$) µm, 8-spored. Ascospores biseriate, elongate-allantoid, thin-walled, hya-

line, slightly curved, aseptate, $8.5\text{--}12 \times 2.5\text{--}3.5$ (av. = $10 \pm 1 \times 3 \pm 0.3$, n = 30) μm . **Asexual morph:** not observed.

Culture characteristics. Cultures are white, growing up to 4 cm in diam. with irregular margin after 3 days, covering the 9 cm Petri dish after 6 days, becoming vinaceous buff to hazel after 7–10 days. In reverse, the cultures are the same as the upper colour after 3 days, becoming isabelline to umber after 7–10 days. Colonies are felty with a heterogeneous texture, lacking aerial mycelium.

Habitat and distribution. Known from *Spiraea salicifolia* and *Tilia nobilis* in Mount Dongling, China.

Additional material examined. CHINA, Beijing City, Mentougou District, Mount Dongling, Xiaolongmen Forestry Centre ($115^{\circ}29'20.49''\text{E}$, $39^{\circ}57'47.43''\text{N}$), from branches of *Tilia nobilis*, 17 Aug 2017, H.Y. Zhu & X.L. Fan, CF 2019804, living culture CFCC 53139.

Notes. *Cytospora spiraeicola* is associated with canker disease of *Spiraea salicifolia* and *Tilia nobilis* in China, with characteristics similar to *Cytospora elaeagnicola* and *C. spiraeae* in phylogram (Fig. 2). Morphologically, it differs from *C. spiraeae* by the smaller perithecia (210–250 vs. 270–400 μm in diam.) and longer ascospores ($8.5\text{--}12 \times 2.5\text{--}3.5$ vs. $7\text{--}8 \times 2\text{--}2.5 \mu\text{m}$) (Zhu et al. 2018a). Phylogenetically, *C. spiraeicola* (CFCC 53138) differs from *C. elaeagnicola* (CFCC 52882) by ITS (8/665), *rpb2* (44/730), *tef1-a* (75/771) and *tub2* (42/624) and *C. spiraeae* (CFCC 50049) by ITS (4/665), *rpb2* (38/730), *tef1-a* (63/771) and *tub2* (44/624) (Zhu et al. 2018a, Zhang et al. 2019). Therefore, we describe it as a novel species.

Discussion

In the present study, seven specimens were collected from symptomatic branches and twigs associated with canker disease. Four *Cytospora* species were isolated from six tree hosts of Betulaceae, Juglandaceae, Rosaceae, Tiliaceae and Ulmaceae, which include two known species (*Cytospora leucostoma* and *C. pruinopsis*) and two novel species (*C. coryli* and *C. spiraeicola*). This study represents an investigation of *Cytospora* species associated with canker disease in Mount Dongling of China and included a comprehensive analysis of DNA sequence data to compare the novelties with known *Cytospora* species.

In a previous study, Zhu et al. (2018a) described *Cytospora spiraeae* from *Spiraea salicifolia* in Gansu Province of China during an investigation of forest pathogens of three hosts. Compared to the new species *Cytospora spiraeicola*, *C. spiraeae* has larger perithecia (270–400 vs. 210–250 μm in diam. and shorter ascospores ($7\text{--}8 \times 2.5\text{--}3.5 \times 8.5\text{--}12$ vs. $2\text{--}2.5 \mu\text{m}$). These morphological deviations are in line with the combined phylogenetic analyses which resolved *C. spiraeicola* as a unique lineage, highly supported. Besides this, the only strain of *C. coryli*, closely related to *C. euonymicola* and *C. gigalocus*, was distinguished by its different size of multiple locules and conidia (Fan et al. 2015a, 2020).

This study focused on *Cytospora* species in Mount Dongling of Beijing (China), which is considered as an attractive location with a high richness of fungal species (Guo et al. 2008, Zhu et al. 2018b, 2019). We hope that the descriptions and molecular data of *Cytospora* in this study could provide a resource for future studies in this genus and lay the foundation for the future canker disease caused by *Cytospora* species.

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