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RESEARCH ARTICLE

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Cytospora elaeagnicola sp. nov. Associated with Narrow-leaved Oleaster Canker Disease in China

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ABSTRACT

Cytospora is a genus including important phytopathogens causing severe dieback and canker diseases distributed worldwide with a wide host range. However, identification of Cytospora species is difficult since the currently available DNA sequence data are insufficient. Aside the limited availability of ex-type sequence data, most of the genetic work is only based on the ITS region DNA marker which lacks the resolution to delineate to the species level in Cytospora. In this study, three fresh strains were isolated from the symptomatic branches of Elaeagnus angustifolia in Xinjiang Uygur Autonomous Region, China. Morphological observation and multi-locus phylogenetic analyses (ITS, LSU, ACT and RPB2) support these specimens are best accommodated as a distinct novel species of Cytospora. Cytospora elaeagnicola sp. nov. is introduced, having discoid, nearly flat, pycnidial conidiomata with hyaline, allantoid conidia, and differs from its relatives genetically and by host association.

ARTICLE HISTORY

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KEYWORDS

Cytosporaceae; molecular phylogeny; new species; taxonomy

1. Introduction

The genus Cytospora contains important phytopathogens causing dieback and stem canker disease on multiple woody plants [1,2]. It was introduced by Ehrenberg in 1818 [3] and belonged to the family Cytosporaceae in Diaporthales [4]. This disease has globally caused great losses on ecologically and commercially important woody plants. Cytospora is characterized by the diaporthalean-like perithecial ascoma, clavate to elongate obovoid asci with allantoid, hyaline, aseptate ascospores in sexual state; and the single or labyrinthine locules, filamentous conidiophores, phialidic conidiogenous cells with allantoid, hyaline, aseptate conidia in the asexual state [2,5]. The asexual name Cytospora (1818) is an older name than all of the sexual synonyms Valsa (1849), Leucocytospora (1917), Leucostoma (1917), Valsella (1870) and Valseutypella (1919), and thus has the priority in nomenclature [2,6–8]. More than 610 species named Cytospora are listed at present in Index Fungorum (2019). However, the amount of species in Cytospora was with 110 estimated species [9]. Species criteria of Cytospora were previously based on host affiliations and morphology in China, however these bases are unreliable due to the uninformative

illustrations and descriptions, weak host specificity and overlapping morphological characteristics [10–12]. Recent studies have reported updated phylograms for the genus *Cytospora* on the basis of multigene phylogenetic analyses using ex-type or reference strains [6,7,13–15]. However, because availability of the extype sequence data is limited to few species, identification of a strain to species level is very difficult. Recently, only 14 new species were included to this genus [16].

Elaeagnus angustifolia is a drought-resistant tree that is grown as a major biomass energy source [17], and has high medicinal and ecological value as well [18]. Furthermore, during an investigation of phytopathogens in north of China, most E. angustifolia trees were observed to suffer from dieback and stem canker caused by Cytospora species. In the current representative three Cytospora were collected from Elaeagnus angustifolia in Xinjiang Uygur Autonomous Region, China. Multilocus phylogenetic analyses using combination of ITS, LSU, ACT and RPB2 sequences confirmed finding of a new species in Cytospora. In this paper, C. elaeagnicola sp. nov. is introduced, accompanied with descriptions, illustrations and comparison with other species in the genus.

2. Materials and methods

2.1. Sample collection and isolation

Fresh samples of Cytospora were collected from infected branches and stems of E. angustifolia during investigations of phytopathogens in Xinjiang Uygur Autonomous Region, China. The samples placed in paper bags were brought to the laboratory for processing and experimental purpose using the same methodology as in Fan et al. [14,15]. Single conidia were isolated by taking fruiting bodies and suspend the mucoid spore mass removed from conidiomata or ascomata in a drop of sterile water. The spore suspension from each sample was then spread over the surface of 1.8% potato dextrose agar (PDA) medium in a petri-dish and incubated at 25 °C. After 24 h, a single germinating conidium was transferred to a fresh PDA plate. Samples and isolates of the new species were deposited in the Museum of Beijing Forestry University (BJFC) and single-spore cultures in the China Forestry Culture Collection Center (CFCC).

2.2. Morphology observation

Samples were observed on infected plant tissues including the structure and size of fruiting bodies. The photographs of the macro-morphological characteristics were recorded using a Leica stereomicroscope (M205 FA) while the micro-morphological observations were determined under a Leica compound microscope (DM 2500) with differential interference contrast (DIC). Over 20 fruiting bodies were sectioned, both vertically and horizontally, and 50 conidia were selected randomly to get the measurement of their length and width. Cultural characteristics, including the colony characters and the production of pigment of isolates on PDA incubated at 25 °C in the dark were recorded, after 3, 7, and 30-days growth [19].

2.3. 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 approach [20]. The ITS region was amplified with the primers ITS1 and ITS4 [21]; the LSU region with LR0R and LR7 [22]; the partial ACT region with ACT512F and ACT783R [23] and the RPB2 region with RPB2-5F and fRPB2-7cR [24]. The PCR amplicons were estimated visually by electrophoresis in 2% agarose gels. Fragments were sequenced in both directions using the respective primers and the BigDye Terminater

v.3.1 Cycle Sequencing Kit (Applied Biosystems; Foster City, CA). Sequences were joined and quality was examined with Seqman v.7.1.0 in the DNASTAR lasergene core suite software (DNASTAR Inc.; Madison, WI).

2.4. DNA sequence analysis

Sequences based on ITS region and the combined dataset (ITS, LSU, ACT and RPB2) were aligned using MAFFT v.6 [25] and edited manually using MEGA6 [26], and some characters were excluded from both ends of the alignments to approximate the size of our sequences to those included in the dataset.

MP analysis was carried out by using PAUP v.4.0b10 with a heuristic search option of 1000 random-addition sequences with a tree bisection and reconnection (TBR) as the branch swapping algorithm [27]. Zero length branches were collapsed, whereas all equally parsimonious trees were saved. Stability of the clade was assessed with a bootstrap analysis of 1000 replicates [28]. Other measures calculated parsimony scores were tree length (TL), consistency index (CI), retention index (RI) and rescaled consistency (RC) [27]. ML analysis was carried out by using RAxML v.7.2.8 with a GTR+G+I model of site substitution, including estimation of gamma-distributed rate heterogeneity and a proportion of invariant sites [29]. And the branch support from MP and ML analyses was evaluated with a bootstrapping method of 1000 replicates [28].

BI analysis employing a Markov Chain Monte Carlo (MCMC) algorithm was performed using in MrBayes v.3.1.2 with the inverse gamma rates (GTR+I+G) nucleotide substitution model, which was selected based on the AIC criterion, using MrModeltest v.2.3 [30,31]. Two MCMC chains were run from random trees for 1,000,000 generations, and trees were sampled every 100th generation, resulting in 10,000 total trees. The first 25% of trees were discarded as the burn-in phase of the analysis and the Bayesian posterior probabilities (BPP) were calculated using the remaining 7500 trees [32].

In all analyses, *C. elaeagnicola* was selected as a distinct and new grape. Phylograms were examined in Figtree v.1.3.1 [33]. Novel sequence data was deposited in GenBank (Table 1), the multilocus sequences alignment file was deposited in TreeBASE (www.treebase.org) accession S24181 and the taxonomic novelty was deposited in MycoBank.

3. Results

3.1. Phylogeny

The ITS sequences of the three isolates of Cytospora from E. angustifolia were aligned with



Table 1. Isolates and GenBank accession numbers used in this study.

Section					GenBank accession numbers		
Cardiolifornis	Species	Strain	Host	ITS	LSU	ACT	RPB2
Condition	C. abyssinica	CMW 10181 ^T	Eucalyptus globulus	AY347353	_	-	_
C ampulliformis MFLUCC 16-0838 ¹ Sorbus intermedia KY417720 KY417760 KY417690 KY417760	C. abyssinica	CMW 10178			_	_	_
C ampulliformis MFLUCC 16 06259 Acer platanoides KY417726 KY417763 KY170648 KY417763 KY417763 KY170648 KY417763 KY417634 KY417763 KY417763 KY417763 KY417763 KY417763 KY417763 KY417763 KY417763 KY417763 <th< td=""><td></td><td></td><td></td><td></td><td>_</td><td>_</td><td>_</td></th<>					_	_	_
C absorbinhata CFCC 89015 Juglams regula RS045019 RR045700 KF498673 KU710946 C autorichnata CHW 679527 Berberis Garystory and RS045020 RR045702 KU710940 KU710940 C berberdis CFC 599273 Berberis Garystory and RS045020 RR045702 KU710940 KU710940 C berberdis CFC 59923 Berberis Garystory and RS045020 RR045702 KU710950 KU710940 C berberdigh CEC 59933 Berberis Garystory and RS045020 RR045702 KU710950 KU710940 C berseypora CBS 1168117 Euclybrus garandis Kereliconis AF192211 - - - - C carborica CFC 20947 Curborico dellos ME192315 - - MF31000 - MF31000 KU710950 - MF31000 - MF31000 - MF31000 - MF37700 - - - MF37700 - - - - - - - - - - - - -	•						
C. carcinorhanda (NEC 88616 Juglares regio (NB 473761 S7498674 KU171094 NJ - C. berberlois (FCC 89923 Berberis disystectyla (R8045201 K8045702 KU1710990 KU1710940 C. berberlois (FCC 89923 Berberis disystectyla (R804521 K8045702 KU1710990 KU1710940 C. berberlois (FCC 89923 Berberis disystectyla (R804521 K8045702 KU1710940 KU1710940 C. berberlois (FCC 89923 Berberis disystectyla (R804521 K8045702 KU1710940 KU1710940 C. berberlope (GS 114692 Eucolyptus globulus (R804522 K910812 K910812 KU1710940 C. berberlope (FCC 89947 Ulmus pumila (R804522 K910812	•		•				
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C. berkeleyi			3		-	-	-
C berkeleyi StarfordTi ^V Euchyptus globulus A7473730 — — — C brwispona CB\$ 116839 Euchyptus grandis AF193231 — — — C carbonacea CFCC 89947 Ullius pumila AF1923215 — — — C carbonacea CFC 89947 Ullius pumila AF192315 — — — — C carpobrati CKW 49861 CWW 49861 CWW 49861 CWW 49861 AW 5070 — MF190122 MF190068 — MF190122 CWP 19009 — MF190122 MF19009 — MF190123 MF19009 — MF190123 MF19009 — MF1937700 CW 2000 Chrysoperma CFCC 89981 Popular solution MF190123 MF190099 MF1903550 MF1933560 CW 19033560 MF1933560 MF1933560 MF1933560 MF1933560 MF1933560 MF1933560 CW 1903360 MF1933560 MF1933560 MF1933560 MF1933560 CW 1903360 CW 190320 CW 190320 CW 190320 CW 190320<	C. berberidis	CFCC 89927 ^T	Berberis dasystachya	KR045620	KR045702		KU710948
Ceberkley UCB Veraphras glandis AF147349 - - -					KR045703	KU710991	KU710949
Commission	,				_	-	_
Corbonation		•			_		_
Carophonice					_	_	_
Cedri	•				KP310812	KP310842	KU710950
Centrivillosa MFLUCC 11-1060 Sorbus domestica MF190122 MF190069 — MF377601 C. chrysosperma C.F.CC 98029 Solar psammaphila KF765673 KF765689 KF765721 KF765705 KF7656891 Populus alba subsp. pyramidalis KF765673 KF7656891 KF765721 KF765705 KF77705 K	C. carpobroti	CMW 48981 ^T	•	MH382812	MH411216	-	_
Centrivillosa			-		_	-	_
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C davidiana CXY 1350 ^T Populus davidana KM034869 — <td>C. cotini</td> <td></td> <td>33,3</td> <td></td> <td></td> <td></td> <td></td>	C. cotini		33,3				
C davidinan CXY 1374 Populus davidlana KM034869 —					KY417762	KY417694	KY417796
C distoflomis CMW 6509¹ Eucalyptus globulus AY347348 - - - - C. disciformis CMW 6750 Eucalyptus globulus AY3473734 -			•		_	_	_
C disciformis CMW 6750 Eucalptus grandis AY347359 — <td></td> <td></td> <td></td> <td></td> <td>_</td> <td>_</td> <td>_</td>					_	_	_
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(continued)

Table 1. Continued.

			GenBank accession numbers			
Species	Strain	Host	ITS	LSU	ACT	RPB2
C. myrtagena	CBS 116843 ^T	Tibouchiina urvilleana	AY347363	_	_	-
C. nivea	MFLUCC 15-0860	Salix acutifolia	KY417737	KY417771	KY417703	KY417805
C. nivea	CFCC 89641	Elaeagnus angustifolia	KF765683	KF765699	KU711006	KU710967
C. nivea	CFCC 89643	Salix psammophila	KF765685	_	_	KU710968
C. palm	CXY1276_	Cotinus coggygria	JN402990	_	_	_
C. palm	CXY1280 ^T	Cotinus coggygria	JN411939	_	_	
C. parakantschavelii	MFLUCC 15-0857	Populus \times sibirica	KY417738	KY417772	KY417704	KY417806
C. parakantschavelii	MFLUCC 16-0575	Pyrus pyraster	KY417739	KY417773	KY417705	KY417807
C. parapersoonii	T28.1 ^T	Prunus persicae	AF191181	_	_	_
C. parasitica	MFLUCC 15-0507 ^T	Malus domestica	KY417740	KY417774	KY417706	KY417808
C. paratranslucens	MFLUCC 15-0506 ^T	Populus alba var. bolleana	KY417741	KY417775	KY417707	KY417809
C. paratranslucens	MFLUCC 16-0627	Populus alba	KY417742	KY417776	KY417708	KY417810
C. pini	CBS 197.42	Pinus Sylvestirs	AY347332	_	-	_
C. pini	CBS 224.52 ^T	Pinus strobus	AY347316	_	_	-
C. populina	CFCC 89644 ^T	Salix psammophila	KF765686	KF765702	KU711007	KU710969
C. predappioensis	MFLUCC 17-2458 ^T	Platanus sp.	MG873484	MG873480	_	_
C. pruinopsis	CFCC 500341	Ulmus pumila	KP281259	KP310806	KP310836	KU710970
C. pruinosa	CFCC 50035	Ulmus pumila	KP281260	KP310807	KP310837	KU710971
C. pruinosa	CFCC 50036	Syzygium aromaticum	KP310800	KP310802	KP310832	_
C. pruinosa	CFCC 50037	Syzygium aromaticum	MH933650	MH933685	MH933558	_
C. prunicola	MFLU 17-0995	Prunus sp.	MG742350	MG742351	MG742353	MG742352
C. quercicola	MFBBH 42443	Quercus sp.	MF190128	MF190074	_	_
C. quercicola	MFLUCC 14-0867	Quercus sp.	MF190129	MF190073	_	_
C. rhizophorae	MUCC302	Eucalyptus grandis	EU301057	-	-	-
C. ribis	CFCC 50026	Ulmus pumila	KP281267	KP310813	KP310843	KU710972
C. ribis	CFCC 50027	Ulmus pumila	KP281268	KP310814	KP310844	_
C. rosae	MFLU 17-0885 ^T	Rosa canina	MF190131	MF190075	-	_
C. rostrata	CFCC 89909 ^T	Salix cupularis	KR045643	KR045722	KU711009	KU710974
C. rostrata	CFCC 89910	Salix cupularis	KR045644	KR045723	KU711010	KU710975
C. rusanovii	MFLUCC 15-0853	Populus × sibirica	KY417743	KY417777	KY417709	KY417811
C. rusanovii	MFLUCC 15-0854 ¹	Salix babylonica	KY417744	KY417778	KY417710	KY417812
C. sacculus	CFCC 89624	Juglans regia	KR045645	KR045724	KM401888	KU710976
C. sacculus	CFCC 89625	Juglans regia	KF225616	KM401887	KM401889	- W/417014
C. salicacearum	MFLUCC 15-05091	Salix alba	KY417746	KY417780	KY417712	KY417814
C. salicacearum	MFLUCC 15-0861	Salix \times fragilis	KY417745	KY417779	KY417711	KY417813
C. salicacearum	MFLUCC 16-0587	Prunus cerasus	KY417748	KY417782	KY417714	KY417816
C. salicicola	MFLUCC 15-0866	Salix alba	KY417749	KY417783	KY417715	KY417817
C. salicicola	MFLUCC 14-1052 ¹	Salix alba	KU982636	KU982635	KU982637	-
C. salicina	MFLUCC 15-08621	Salix alba	KY417750	KY417784	KY417716	KY417818
C. salicina	MFLUCC 16-0637	$Salix \times fragilis$	KY417751	KY417785	KY417717	KY417819
C. schulzeri	CFCC 50040	Malus domestica	KR045649	KR045728	KU711013	KU710980
C. schulzeri	CFCC 50042	Malus asiatica	KR045650	KR045729	KU711014	KU710981
C. sibiraeae	CFCC 500451	Sibiraea angustata	KR045651	KR045730	KU711015	KU710982
C. sibiraeae	CFCC 50046	Sibiraea angustata	KR045652	KR045731	KU711015	KU710983
C. sophorae	CFCC 50047	Styphnolobium japonicum	KR045653	KR045732	KU711017	KU710984
C. sophorae	CFCC 89598	Styphnolobium japonicum	KR045654	KR045733	KU711018	KU710985
C. sophoricola	CFCC 89596	Styphnolobium japonicum	KR045656	KR045735	KU711020	KU710987
C. sophoricola	CFCC 89595 ^T	Styphnolobium japonicum var.	KR045655	KR045734	KU711019	KU710986
C. sorbi	MFLUCC 16-0631 ^T	Sorbus aucuparia	KY417752	KY417786	KY417718	KY417820
C. sorbicola	MFLUCC 16-0584 ¹	Acer pseudoplatanus	KY417755	KY417789	KY417721	KY417823
C. sorbicola	MFLUCC 16-0633	Cotoneaster melanocarpus	KY417758	KY417792	KY417724	KY417826
C. spiraeae	CFCC 50049 ¹	Spiraea salicifolia	MG707859	MG707643	MG708196	MG708199
C. spiraeae	CFCC 50050	Spiraea salicifolia	MG707860	MG707644	MG708197	MG708200
C. tanaitica	MFLUCC 14-1057 ^T	Betula pubescens	KT459411	KT459412	KT459413	_
C. tibouchinae	CPC 263331	Tibouchina semidecandra	KX228284	KX228335	-	_
C. translucens	CXY1351	Populus davidiana	KM034874	_	-	_
C. ulmi	MFLUCC 15-0863 ^T	Ulmus minor	KY417759	_	_	_
C. valsoidea	CMW 4309 ^T	Eucalyptus grandis	AF192312	-	_	_
C. valsoidea	CMW 4310	Eucalyptus grandis	AF192312	-	_	_
C. variostromatica	CMW 6766 ¹	Eucalyptus globulus	AY347366	-	_	_
C. variostromatica	CMW 1240	Eucalyptus grandis	AF260263	_	_	_
C. vinacea	CBS 141585 ¹	Vitis interspecific	KX256256	_	_	_
C. viticola	CBS 141586 ¹	Vitis vinifera	KX256239	_	_	_
Diaporthe vaccinii	CBS 160.32	Vaccinium macrocarpon	KC343228	_	JQ807297	_

All the new isolates used in this study are indicated in bold type and the strains from type materials are marked by an superscript (T).

available ITS sequences from related Cytospora species of published articles, resulting in an alignment containing 138 Cytospora ingroup strains and a total of 609 characters including gaps. In the alignment, 369 characters were constant, 72 variable characters were parsimony-uninformative and 168 characters were variable parsimonyand

informative. MP analyses generated 145 parsimonious trees, one of which is presented in Figure 1 (TL = 927, CI = 0.409, RI = 0.830, RC = 0.339).ML and BI analyses resolved results similar to the MP tree. C. elaeagnicola represented a monophyletic clade with overall high bootstrap support values (MP/ML/BI = 99/100/1; marked in blue in

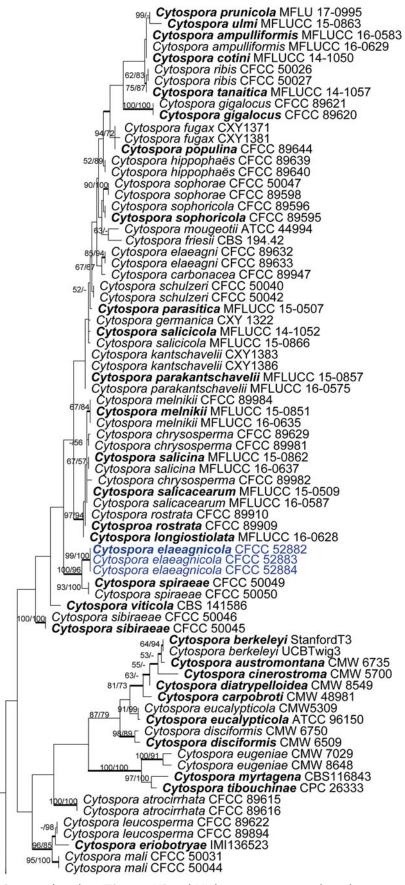


Figure 1. Phylogram of Cytospora based on ITS gene. MP and ML bootstrap support values above 50% are shown at the first and second position. Thickened branches represent posterior probabilities above 0.95 from BI. Ex-type strains are in bold. Strains in current study are in blue.

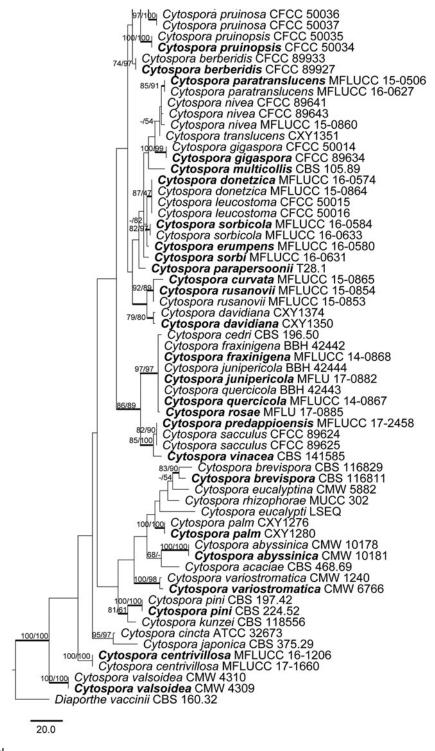


Figure 1. Continued

Figure 1). Subsequently, phylogenetic analyses were performed based on a concatenated alignment of ITS, LSU, ACT and RPB2 from published articles, comprised of 102 *Cytospora* ingroup strains with a total of 2207 characters including gaps. In the alignment, 1538 characters were constant, 104 variable characters were parsimony-uninformative and 565 characters were variable and parsimony-informative. MP analysis generated 105 parsimonious trees, one of which is presented in Figure 1 (TL = 2,350, CI = 0.412, RI = 0.827, RC = 0.341).

ML and BI analyses were similar to the MP tree. *Cytospora elaeagnicola* represented a monophyletic clade with full support values (MP/ML/BI = 100/100/1) (marked in blue in Figure 2).

3.2. Taxonomy

Cytospora elaeagnicola X.L. Fan sp. nov. Figure 3 Mycobank: MB830292.

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

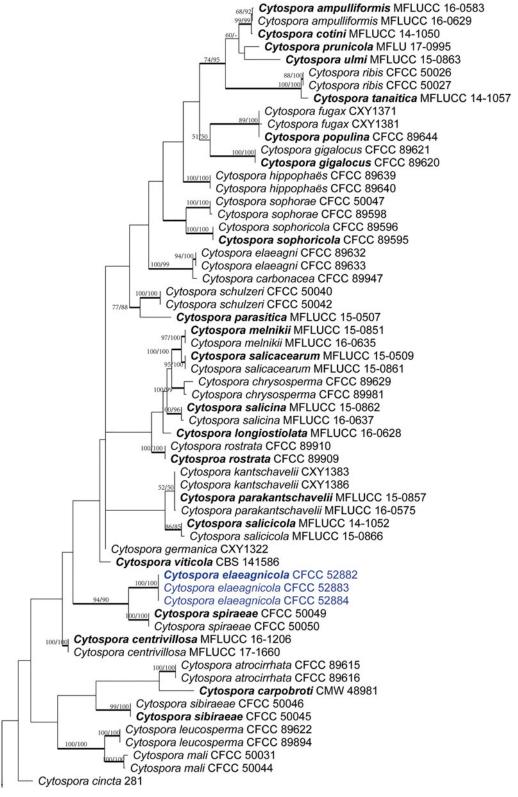


Figure 2. Phylogram of Cytospora based on combined ITS, LSU, ACT and RPB2 genes. MP and ML bootstrap support values above 50% are shown at the first and second position. Thickened branches represent posterior probabilities above 0.95 from Bl. Ex-type strains are in bold. Strains in current study are in blue.

Holotype: CF 20175831.

Host/Distribution: from branches of Elaeagnus angustifolia in China.

Descriptions: Asexual state: Conidiomata pycnidial, ostiolate, discoid, nearly flat, immersed in bark, scattered, producing black area on bark, erumpent through the surface of bark when mature. Locules multiple,

circular to ovoid, arranged irregularly with common walls, $(890-)905-1160(-1240) \mu m (\bar{x} = 1060 \pm 120 \mu m)$ n = 30) in diameter. Conceptacle absent. Ectostromatic disc iron grey to violaceous black, circular, disc dark, (160-)170-310(-350) µm ($\bar{x} = 240 \pm 60$ µm, n = 30) in diameter, with one ostiole in the centre of disc. Ostiole conspicuous, circular to ovoid, iron grey to violaceous

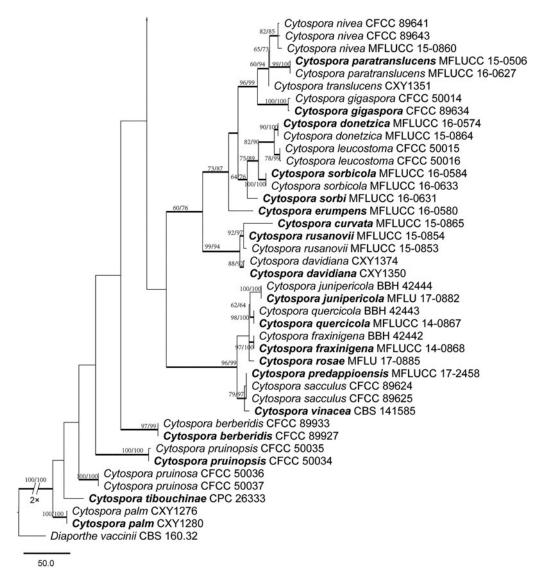


Figure 2. Continued

black at the same level as the disc, (48-)51-71(-78) µm ($\overline{x}=60\pm11$ µm, n=30) in diameter. Conidiophores hyaline, branched at base or not branched, thin walled, filamentous, (12-)13.5-19.5(-20) µm ($\overline{x}=16.5\pm3$ µm, n=30). Conidiogenous cells enteroblastic, phialidic. Conidia hyaline, allantoid, eguttulate, smooth, aseptate, thin-wall, $5.5-6.5(-7)\times(1-)1.5-2$ µm ($\overline{x}=6.1\pm0.4\times1.6\pm0.1$ µm, n=50). Sexual morph: not observed.

Culture characteristics: On PDA, cultures are white. The colony is flat, felt-like with a thick texture at the center with thin surrounding texture. Pycnidia are sparse, distributed irregularly on medium surface.

Materials examined: China, Xinjiang Uygur Autonomous Bole Mongol Autonomous Region, Prefecture, Provincial Road 202, 45°06'29.50"N, 82°33'32.82"E, from branches of Elaeagnus angustifolia, July 2017, C.M. Tian & X.L. Fan, deposited by X.L. Fan, holotype CF 20175831, ex-type living culture CFCC 52882; ibid. CF 20175832, living culture CFCC 52883; CF 20175833, living culture CFCC 52884.

Notes: Cytospora elaeagnicola is associated with canker disease of Elaeagnus angustifolia. The phylogenetic inferences resolved this species an individual clade both in ITS and combined multigene phylograms (Figures 1 and 2), which was closed spiraeae from Spiraea salicifolia. Morphologically, Cytospora elaeagnicola has obvious symptoms with black area on bark, and smaller conidia $(5.5-6.5 \times 1.5-2 \text{ vs. } 7-8 \times 2-2.5 \text{ } \mu\text{m})$ as compared with C. spiraeae; the cultures of C. elaeagnicola are white, differing from the cultures of C. spiraeae which becomes fawn after 7-10 days [34]. Considering the clearly distinction between these two species based on molecular phylogenetic position and on the host affiliation, Cytospora elaeagnicola is thus described as a novel species.

4. Discussion

In the current study, *C. elaeagnicola* sp. nov. was described from infected branches and twigs of *E.*

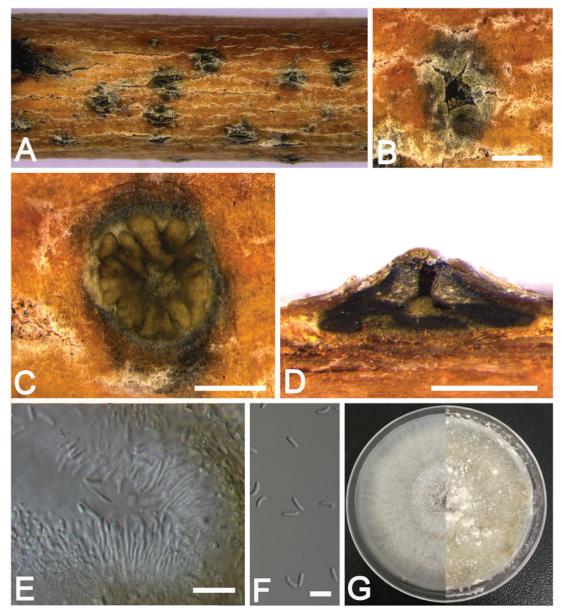


Figure 3. Morphology of Cytospora elaeagnicola from Elaeagnus angustifolia (CF 20175831). (A), (B) Habit of conidiomata on twig; (C) Transverse section of conidioma; (D) Longitudinal section through conidioma; (E) Conidiophores and conidiogenous cells; (F) Conidia; (G) Colonies on PDA after 3 d and 14 d (scale bars: $B-C=250 \,\mu m$, $D=200 \,\mu m$, $E=10 \,\mu m$, $F=5 \,\mu m$).

angustifolia in northwest region of China, an area that has undergone desertification at an alarming rate. Previously, Fan et al. [7] described C. elaeagni and C. nivea from E. angustifolia during the investigation of canker disease of three anti-desertification plants. Compared to C. elaeagnicola, C. elaeagni has smaller locules $(630-920 \mu m)$ with larger conidia $(6.3-9.3 \times 2-2.9 \,\mu\text{m})$ and dense cultures producing light brown pigment; C. nivea has obvious dark black conceptacle surrounding the conidiomata with larger conidia $(6.2-9.2 \times 1.7-2.4 \,\mu\text{m})$, and cultures producing dark green to black pigment [7]. These morphological deviations are in line with the combined phylogenetic analyses which resolved C. elaeagnicola as a separate, highly supported clade, both in the single ITS analyses and the concatenated analyses.

Cytospora species were previously identified by host association and morphological characteristics.

However, the uninformative illustrations and descriptions, overlapping morphological characteristics and low host-specificity have caused confusion in the identification of strains. Current study indicated more than one species of Cytospora are present on one host plant. In the future study, the taxonomy requires fresh collections from wide geographical ranges with comprehensive pathogenicity tests. Further studies are also needed in the clarification of the species diversity and in the understanding of their roles in plant diseases, especially for anti-desertification plants such as E. angustifolia in Northwestern China.

Disclosure statement

No potential conflict of interest was reported by the authors.

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