Colletotrichum species associated with Camellia anthracnose in China

Peng XJ, Wang QC, Zhang SK, Guo K* and Zhou XD*

State Key Laboratory of Subtropical Silviculture, College of Forestry and Biotechnology, Zhejiang A & F University, Hangzhou 311300, Zhejiang, China


Abstract

Species of Camellia oleifera, Ca. sinensis and Ca. japonica represent globally-important tree germplasm resource and have significant ecological and economic value. However, they have suffered from anthracnose caused by pathogenic fungi from the genus of Colletotrichum. To determine the diversity of Colletotrichum species associated with Camellia anthracnose in China, we collected infected leaves from three major cultivation provinces. A total number of 167 fungal strains resembling Colletotrichum were obtained, 41 from Ca. oleifera, 48 from Ca. sinensis, and 78 from Ca. japonica. Comparison of morphology and phylogenetic analyses based on six loci (ACT, CAL, CHS-1, GAPDH, TUB2 and ITS) of the representative isolates revealed that they fall in four species complexes and represent 15 known and one undescribed taxa. The species complexes are Colletotrichum gloeosporioides (C. aenigma, C. alienum, C. camelliae, C. fructicola, C. gloeosporioides, C. jiangxiense, C. pandanicola, C. siamense, C. wuxiense and C. puereense described here), C. acutatum (C. fioriniae and C. nymphaeae), C. boninense (C. boninense and C. karstii) and C. orchidearum (C. clivicola and C. plurivorum). Of these, C. fructicola was the most dominant species occurring on Ca. oleifera (68.3%), Ca. sinensis (50.0%) and Ca. japonica (29.49%). Pathogenicity test results indicated that C. camelliae is the most pathogenic to the three Camellia species. This study represents the first report of C. aenigma, C. alienum, C. boninense, C. camelliae, C. clivicola, C. fioriniae, C. fructicola, C. gloeosporioides, C. jiangxinse, C. karstii, C. nymphaeae, C. pandanicola and C. wuxiense causing anthracnose on Ca. japonica and C. pandanicola on Ca. sinensis. Knowledge gained in this study enlarges our understanding of Colletotrichum species causing anthracnose on three important Camellia trees and contributes to the disease management.

Keywords – 1 new species – Camellia japonica – fungal diversity – multi-gene phylogeny – pathogenicity

Introduction

Camellia is one of the most traditional and diverse genera within the family Theaceae. It includes at least 280 species, and they are mainly distributed in east Asia (Yang et al. 2011, 2019, Teixeira & Sousa 2021, Talhinhas & Baroncelli 2021). Certain Camellia species are of high commercial value. For example, tea-oil tree (Ca. oleifera) is China’s most important woody oil tree species (Quan et al. 2022). Up till now, its cultivation area has reached close to 5 million ha and the oil production from seeds is around 518,000 tons per year there (State-owned Forest Farms and Nurseries Station, State Forestry Administration of China 2022). Camellia sinensis belongs to the
tea group of *Camellia*, and is a vital beverage tree crop worldwide grown in more than 60 countries (Orrock et al. 2019). Tea leaves are rich in a variety of active substances such as alkaloids, steroids and polyphenols that are beneficial to human health (Anand et al. 2015). *Camellia japonica* is one of the top ten traditional famous flowers in China with great ornamental and medicinal value (Moon & Kim 2018, Zhang et al. 2018). These three *Camellia* species have thus been extensively studied due to their significant economic values (Yang et al. 2018, Teixeira & Sousa 2021, Lin et al. 2022, Zhao et al. 2022). Many species of *Colletotrichum* are fungal pathogens (Van der AA 1978, Liu et al. 2022). They can infect and cause anthracnose in different plant genera including *Camellia*, resulting in serious economic and ecological losses (Guarnaccia et al. 2018, Chen et al. 2021, Zhang et al. 2021, Yang et al. 2022). The disease is more severe and difficult to manage, especially under high temperature and humidity conditions (Zhou et al. 2007, Liu et al. 2009).

Taxonomy of *Colletotrichum* species was confusing as their morphology is similar and there was a lack of comprehensive study using a polyphasic approach to identify the species (Cannon et al. 2014, Talhinhas & Baroncelli 2021). Recently, comparisons of morphological characteristics and multi-gene sequence analyses have been applied to define the taxonomic status in *Colletotrichum*, and a total number of 280 species were clarified (Dowling et al. 2020, Liu et al. 2022). Previous studies reported that there are 11, 16 and three *Colletotrichum* species causing anthracnose on *Ca. oleifera*, *Ca. sinensis* and *Ca. japonica*, respectively (Table 1) (Liu et al. 2015, Hou et al. 2016, Orrock et al. 2019, Li & Li 2020, Wang et al. 2020, Zhang et al. 2021, Li & Li 2022, Peng et al. 2022, Chen et al. 2022). However, there is a lack of comprehensive study on *Colletotrichum* causing anthracnose on the three important *Camellia* species including diversity, occurrence and pathogenicity of pathogens. The aim of this study was thus to achieve these goals using a combined approach of fungal morphology and phylogenetic analyses of multi-locus sequences of ACT, CAL, CHS-I, GAPDH, TUB2, ITS (Weir et al. 2012, Wang et al. 2020, Zhang et al. 2021). The pathogenicity of the representative *Colletotrichum* isolates was further evaluated to contribute to disease management and germplasm selection.

**Table 1** *Colletotrichum* species previously reported from *Camellia* anthracnose.

<table>
<thead>
<tr>
<th>Host</th>
<th>Species</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Ca. oleifera</em></td>
<td><em>C. fructicola</em>, <em>C. camelliae</em>, <em>C. siamense</em>, <em>C. gloeosporioides</em>, <em>C. horii</em>, <em>C. boninense</em>, <em>C. kahavae</em>, <em>C. karstii</em>, <em>C. fioriniae</em>, <em>C. aenigma</em> and <em>C. aescynomenes.</em></td>
<td>Li &amp; Li 2020, Wang et al. 2020, Li &amp; Li 2022, Chen et al. 2022</td>
</tr>
<tr>
<td><em>Ca. sinensis</em></td>
<td><em>C. gloeosporioides</em>, <em>C. siamense</em>, <em>C. aenigma</em>, <em>C. fructicola</em>, <em>C. camelliae</em>, <em>C. acutatum</em>, <em>C. fioriniae</em>, <em>C. boninense</em>, <em>C. karstii</em>, <em>C. truncatum</em>, <em>C. cliviae</em>, <em>C. crassipes</em>, <em>C. jiangxiense</em>, <em>C. wuxiense</em>, <em>C. henanense</em> and <em>C. endophytic</em>.</td>
<td>Liu et al. 2015, Orrock et al. 2019, Zhang et al. 2021</td>
</tr>
<tr>
<td><em>Ca. japonica</em></td>
<td><em>C. siamense</em>, <em>C. aracearum</em> and <em>C. camelliae-japonica</em>.</td>
<td>Hou et al. 2016, Peng et al. 2022</td>
</tr>
</tbody>
</table>

**Materials & Methods**

**Sampling and fungal isolation**

Leaves with typical anthracnose symptoms of *Ca. oleifera*, *Ca. sinensis* and *Ca. japonica* plants were collected from the provinces of Zhejiang, Jiangxi, Yunnan and Shanghai in China during the period of 2020-2022. The diseased tissues with black acervuli on the surface were cut into pieces and incubated in moist chambers at 25 °C (Senanayake et al. 2020). The spore mass was collected using a sterile needle under a dissection microscope and the tissues without sporulation were cut into 4-5 mm² fragments and surface sterilized with 1% NaClO for 45 s, 75% ethanol for 1 min and rinsed three times in sterile water (Fu et al. 2019). They were transferred onto malt extract agar plates at 25 °C and purified. For further study, pure cultures were stored in 25% glycerol at -80 °C. The specimens were kept at culture collection of tree health division, Zhejiang A&F...
University, and ex-type living cultures were deposited in the China General Microbiological Culture Collection Centre (CGMCC), Beijing, China.

DNA extraction, PCR amplification and sequencing

Fungal genomic DNA was extracted using an Ezup Column Fungal Genomic DNA Extraction Kit (Shenggong Bioengineering Co., LTD). Partial regions of six loci were amplified. The partial gene sequence of the actin (ACT) was amplified with primers ACT-512F/ACT-783R (Carbone & Kohn 1999), calmodulin (CAL) with primers CL1C/CL2C (Weir et al. 2012), chitin synthase (CHS-1) with primers CHS-79F/CHS-345R (Carbone & Kohn 1999), glyceraldehyde-3-phosphate dehydrogenase (GAPDH) with primers GDF1/GDR1 (Templeton et al. 1992), beta-tubulin (TUB2) with primers T1/Bt2b (Glass & Donaldson 1995) and internal transcribed spacer regions and intervening 5.8S nrRNA gene (ITS) with primers ITS4/ITS5 (White et al. 1990). The information on primer pairs used are listed in Table 2.

The PCR amplification was conducted as described by Weir et al. (2012). The PCR amplicons were purified and sequenced at the Qinke Biotech Company, Shanghai, China. The consensus sequences were obtained from forward and reverse sequences with the program SeqMan (DNASTAR, Madison, WI, USA). Sequences generated in this study were deposited in GenBank.

Table 2: Primers used for PCR amplification and sequencing.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer</th>
<th>Primer sequence (5'-3')</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>ITS</td>
<td>ITS1</td>
<td>TCCGTAGGTAACCTGCGG</td>
<td>White et al. (1990)</td>
</tr>
<tr>
<td></td>
<td>ITS4</td>
<td>TCCCTCGGTTATGGATATGC</td>
<td></td>
</tr>
<tr>
<td>ACT</td>
<td>ACT-512F</td>
<td>ATGTGCAAGGCGGTTTCGC</td>
<td>Carbone &amp; Kohn (1999)</td>
</tr>
<tr>
<td></td>
<td>ACT-783R</td>
<td>TACGAGTCTTCTGGCCCAT</td>
<td></td>
</tr>
<tr>
<td>TUB2</td>
<td>T1</td>
<td>AACATGCCGTAGATTGTAAGT</td>
<td>Glass &amp; Donaldson (1995),</td>
</tr>
<tr>
<td></td>
<td>Bt2b</td>
<td>ACCCTCAGTGTAGTACCTGGGC</td>
<td>O’Donnell &amp; Cigelnik (1997)</td>
</tr>
<tr>
<td>GAPDH</td>
<td>GDF</td>
<td>GCCGTCAAACGCCCTTCTGTA</td>
<td>Templeton et al. (1992)</td>
</tr>
<tr>
<td></td>
<td>GDR</td>
<td>GGGTGGAGTCGTAATCTGGACGATGT</td>
<td></td>
</tr>
<tr>
<td>CAL</td>
<td>CL1C</td>
<td>GAATTCAGGAGGCGCTTC</td>
<td>Weir et al. (2012)</td>
</tr>
<tr>
<td></td>
<td>CL2C</td>
<td>CTTCTGCATCATGACGGGTG</td>
<td></td>
</tr>
<tr>
<td>CHS-1</td>
<td>CHS-79F</td>
<td>TGCGGCAAGGAGTCTGGGAAAG</td>
<td>Carbone &amp; Kohn (1999)</td>
</tr>
<tr>
<td></td>
<td>CHS-345R</td>
<td>TGGAAGAACCCTCTGTGAGAGT</td>
<td></td>
</tr>
</tbody>
</table>

Phylogenetic analyses

Sequences of 167 representative isolates were assembled and blasted to investigate their phylogenetic relatives (Dissanayake et al. 2020). The sequences of different gene regions obtained from this study and ex-type cultures downloaded from GenBank were aligned using MAFFT v. 7 online servers (http://mafft.cbrc.jp/alignment/serve/index.html) (Katoh & Standley 2013) and manually adjusted in MEGA v. 7.06 (Kumar et al. 2016). Maximum Likelihood (ML) analysis was performed on the multi-locus alignment using PhyloSuite software v. 1.2.2 (Zhang et al. 2020). Clade stability was assessed in a bootstrap analysis with 1000 replicates. Concatenated analyses of ACT, CHS-1, GAPDH, TUB2 and ITS were conducted for the C. acutatum complex and C. orchidearum complex (Damm et al. 2012a, Liu et al. 2022), while sequences of ACT, CAL, CHS-1, GAPDH, TUB2 and ITS regions were concatenated for the analysis of the C. gloeosporioides complex and C. boninense complex (Damm et al. 2012b). PhyloSuite software was used to conduct the statistical selection of best-fit models of nucleotide substitution using the corrected Akaike information criterion (AIC).

The undescribed taxon and its most phylogenetically close neighbors were analyzed using the Genealogical Concordance Phylogenetic Species Recognition (GCPSR) model by performing a pairwise homoplasy index (PHI) test (Quaedvlieg et al. 2014). The PHI test was performed in SplitsTree v. 4.14.6 (Huson & Bryant 2006) to determine the recombination level within
phylogenetically closely related species using a six-locus concatenated dataset (ACT, CAL, CHS-1, GAPDH, TUB2, and ITS). If the resulting pairwise homoplasy index was below a 0.05 threshold ($\Phi_w < 0.05$), it was indicative of significant recombination in the dataset. The relationship between closely related species was visualized by constructing a split graph.

**Morphology**

Mycelial plugs (5 mm) of purified cultures were transferred from the growing edge of 5-d-old cultures to the centre of 9-mm-diameter malt extract agar (MEA, 2%) plates in triplicate at 25 °C. Colony diameters were measured from 3 to 6 days to calculate mycelial growth rates (mm/d). Colony color, size and density were also recorded.

The shape, color and size of conidia, conidiophores and ascospores were observed using a Carl Zeiss Imager A2 microscope after ten days of cultivation. Sizes of 30 conidia were measured unless no spores were produced. Appressoria were induced by dropping a conidial suspension ($10^6$ conidia/mL; 50 μL) on a concavity slide containing moistened filter papers with distilled sterile water, and then incubating at 25 °C in the dark. After 24 h to 48 h, the conidial appressoria formed at the ends of germ tubes were measured (Tan et al. 2022).

**Pathogenicity tests**

Sixteen representative isolates of *Colletotrichum* species were selected for the pathogenicity test on detached leaves of *Ca. oleifera*, *Ca. sinensis* and *Ca. japonica* plants under controlled conditions. The pathogenicity test was conducted with an aliquot of spore suspension ($1.0 \times 10^6$ conidia per mL) on detached healthy leaves of three hosts in eight replicates each. The healthy-looking leaves were collected and washed three times with sterile water, and air-dried on sterilized filter paper. The leaves are inoculated using the wound/drop inoculation method (Aiello et al. 2015, Cristóbal-Martínez et al. 2017). The 10 μL spore suspension was inoculated on the left side of a leaf after wounding once by pin-pricking with a sterilized needle (insect pin, 0.5 mm diam), an equal amount of sterile water on the right side of the same leaf in parallel as a control. The inoculated leaves were kept in a growth chamber with 85% relative humidity at 26 °C with a 12/12 h light/dark photoperiod. The length of lesions was measured 14 days after inoculation (Bhunjun et al. 2021).

**Results**

**Fungal isolation**

Two hundred thirty-eight isolates resembling *Colletotrichum* were obtained and preliminarily identified based on morphology and ITS sequence data. According to the different sampling sites and hosts, 167 representative isolates were chosen for further detailed analyses. Among them, 41 isolates were from *Ca. oleifera*, 48 from *Ca. sinensis* and 78 from *Ca. japonica*. Our result indicated that *C. fructicola* represents the dominant taxon causing anthracnose in these three-plant species (Fig. 1). The occurrence of other *Colletotrichum* species was summarized in Table 3. In short, we obtained five *Colletotrichum* species from *Ca. oleifera* anthracnose, 14 from *Ca. sinensis* and 14 from *Ca. japonica*, respectively. Among them, *C. clivicola* and *C. aegima* were isolated only from *Ca. japonica*, while *C. puerense* and *C. plurivorum* only from *Ca. sinensis*.

**Table 3** *Colletotrichum* species previously reported from *Camellia* anthracnose.

<table>
<thead>
<tr>
<th>Species</th>
<th>Occurrence rate (%) on <em>Ca. oleifera</em></th>
<th>Occurrence rate (%) on <em>Ca. sinensis</em></th>
<th>Occurrence rate (%) on <em>Ca. japonica</em></th>
</tr>
</thead>
<tbody>
<tr>
<td><em>C. fructicola</em></td>
<td>68.3</td>
<td>50.0</td>
<td>29.5</td>
</tr>
<tr>
<td><em>C. siamense</em></td>
<td>14.6</td>
<td>8.3</td>
<td>23.1</td>
</tr>
<tr>
<td><em>C. camelliae</em></td>
<td>2.4</td>
<td>8.3</td>
<td>11.5</td>
</tr>
<tr>
<td><em>C. fioriniae</em></td>
<td>12.2</td>
<td>2.1</td>
<td>7.7</td>
</tr>
<tr>
<td><em>C. wuxiense</em></td>
<td>2.4</td>
<td>2.1</td>
<td>1.3</td>
</tr>
</tbody>
</table>
Table 3 Continued.

<table>
<thead>
<tr>
<th>Species</th>
<th>Occurrence rate (%) on Ca. oleifera</th>
<th>Occurrence rate (%) on Ca. sinensis</th>
<th>Occurrence rate (%) on Ca. japonica</th>
</tr>
</thead>
<tbody>
<tr>
<td>C. alienum</td>
<td>-</td>
<td>2.1</td>
<td>6.4</td>
</tr>
<tr>
<td>C. boninense</td>
<td>-</td>
<td>2.1</td>
<td>1.3</td>
</tr>
<tr>
<td>C. clivicola</td>
<td>-</td>
<td>-</td>
<td>1.3</td>
</tr>
<tr>
<td>C. jiangxiense</td>
<td>-</td>
<td>4.2</td>
<td>1.3</td>
</tr>
<tr>
<td>C. karstii</td>
<td>-</td>
<td>2.1</td>
<td>3.8</td>
</tr>
<tr>
<td>C. aenigma</td>
<td>-</td>
<td>-</td>
<td>1.3</td>
</tr>
<tr>
<td>C. pandanicola</td>
<td>-</td>
<td>2.1</td>
<td>3.8</td>
</tr>
<tr>
<td>C. nymphaeae</td>
<td>-</td>
<td>4.2</td>
<td>1.3</td>
</tr>
<tr>
<td>C. plurivorum</td>
<td>-</td>
<td>4.2</td>
<td>-</td>
</tr>
<tr>
<td>C. gloeosporioides</td>
<td>-</td>
<td>4.2</td>
<td>6.4</td>
</tr>
<tr>
<td>C. puerense</td>
<td>-</td>
<td>4.2</td>
<td>-</td>
</tr>
</tbody>
</table>

Figure 1 – Occurrence of Colletotrichum species causing Camellia anthracnose.

Phylogenetic analyses

The 167 representative isolates from three Camellia plant species were subjected to multilocus phylogenetic analyses with concatenated ACT, CAL, CHS-1, GAPDH, TUB2, and ITS sequences for those belonging to the species complexes of C. gloeosporioides and C. boninense, and concatenated ACT, CHS-1, GAPDH, TUB2, and ITS sequences for C. acutatum and C. orchidearum species complexes. The results showed that isolates clustered together with 15 known species in four Colletotrichum species complexes, including gloeosporioides (143 isolates), acutatum (15), orchidearum (3), and boninense (6) (Figs 2–4). In the phylogenetic tree constructed for the isolates in the C. gloeosporioides complex, 143 isolates clustered in ten clades corresponding to C. aenigma (1), C. alienum (6), C. camelliae (14), C. fructicola (75), C. gloeosporioides (7), C. jiangxiense (3), C. pandanicola (4), C. siamense (28), C. wuxiense (3). Noticeably two isolates (YNS-22, YNS-30) clustered distantly from any known species in the complex, is herein described as a new taxon, namely C. puerense based on the guidelines established in Chethana et al. (2021) and Jayawardena et al. (2021) (Fig. 2). From 15 isolates in the C. acutatum species complex, 12 isolates grouped in the clade of C. fioriniae, and three isolates...
were clustered with *C. nymphaeae*. From three isolates in the *C. orchidearum* complex, two isolates clustered with *C. plurivorum* and one with *C. clivicola* (Fig. 3). From *C. boninense* complex, four isolates clustered with *C. karstii*, and two with *C. boninense* (Fig. 4).

**Figure 2** – Phylogenetic tree using sequences of 143 *Colletotrichum* isolates from *Ca. oleifera*, *Ca. sinensis* and *Ca. japonica* leaves in the *C. gloeosporioides* complex. The species *C. boninense* (CBS 123755) was selected as an outgroup. The tree was built using concatenated sequences of the ACT, CAL, CHS-1, GAPDH, TUB2, and ITS gene regions. Bootstrap values >50% (1000 replication) are given at the nodes. Ex-type or reference cultures are in bold. Colored blocks indicate clades containing isolates from *Camellia* plants.
Morphological investigation

Cultures of the representative isolates were selected to observe their morphological characteristics. Few differences in colony morphology were obviously observed among the 16 species (Figs 6–21). When grown on MEA, all selected isolates firstly developed white colonies. They became gray-green to black after 7 days except those belonging to *C. acutatum* complex usually appear pink. Abundant orange conidial masses were often observed on the center of the colonies of *C. fructicola*, *C. gloeosporioides*, *C. siamense*, *C. karstii*, *C. fioriniae* and *C. nymphaeae*. A significant difference in growth rate across 16 *Colletotrichum* species was observed (Table 4). The average mycelial growth rate of *C. gloeosporioides* was the fastest, with an average of 11.87 ± 0.4 mm/d. The average mycelial growth rate of *C. fructicola* was 13.6 ± 0.6 mm/day (n = 11), followed by *C. siamense* (11.0 ± 0.9 mm/d), *C. alienum* (10.7 ± 0.6 mm/d) and *C. fructicola* (10.6 ± 0.6 mm/d), while *C. fioriniae* and *C. nymphaeae* in the *C. acutatum* complex showed the slowest growth rate with 8.2-8.3 mm/d. However, there were no significant differences in the conidial size among the species.
Figure 2 – Continued.

Taxonomy

Based on morphology and multi-locus sequence data, a total of 167 obtained isolates were assigned to 16 Colletotrichum species. Of these, one species represents an undescribed taxon and is described below. Species descriptions were also provided for those existing ones according to the complex they reside.


Materials examined – China, Shanghai City, on leaves of *Ca. japonica*, Mar. 2022.

Notes – *Colletotrichum aenigma* belongs to the gloeosporioides species complex and has been recorded as a pathogen on *Ca. sinensis* in China (Wang et al. 2016). This is the first report of *C. aenigma* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 70–71 mm in 6 d at 25 °C with entire margin, aerial mycelium sparse, surface white. The conidia are bigger than those of the ex-type isolate (ICMP 18608) 14.4–16.1 × 5.1–6.1 μm, mean ± SD = 15.3 ± 0.8 × 5.6 ± 0.5 μm, L/W ratio = 2.7 vs 12–16.5 × 5–7.5 μm, mean = 14.5 × 6.1 μm, L/W ratio = 2.4 of *C. aenigma*.


Materials examined – China, Zhejiang province, Fu yang City, on leaves of *Ca. japonica*, Mar. 2022; Zhejiang province, Lin’an City, on leaves of *Ca. japonica*, Sep. 2021; Zhejiang province, Lin’an City, on leaves of *Ca. sinensis*, Dec. 2021.

Notes – *Colletotrichum alienum* belongs to the gloeosporioides species complex. The species is known from many important crops such as *Camellia sinensis*, *Diospyros kaki* and *Grevillea* sp.
(Weir et al. 2012, Liu et al. 2013). This is the first report of *C. alienum* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 65–67 mm in 6 d at 25 °C with entire margin, aerial mycelium dense, surface greyish-green and reverse dark grey. Conidia (14.8–17.6 × 4.8–5.6 μm, mean ± SD = 16.2 ± 0.8 × 5.2 ± 0.8 μm, L/W ratio = 3.1) are slightly shorter than those of the ex-type isolate (ICMP 12071) 15.5–17.5 × 5–5.5 μm, mean = 16.5 × 5.0 μm, L/W ratio = 3.3 of *C. alienum*.

**Colletotrichum boninense** Moriwaki, Toy. Sato & Tsuki, Mycoscience 44: 48. 2003

Materials examined – China, Yunnan province, Pu’er City, on leaves of *Ca. sinsensis* and *Ca. japonica*, Mar. 2022.

Notes – *Colletotrichum boninense* belongs to the boninense species complex. This is the first report of *C. boninense* causing anthracnose on *Ca. japonica* in China. Colonies diam on MEA attained 55–58 mm in 6 d at 25 °C, aerial mycelium sparse, surface light pink. Conidia hyaline, smooth walled, cylindrical, one end rounded and one end slightly acute, 11.5–15.5 × 6.5–7.8 μm, mean ± SD = 14.2 ± 1.2 × 6.5 ± 0.5 μm, L/W ratio = 2.2. The L/W ratio measured here is smaller than described (2.3).

Figure 3 – Phylogenetic tree using sequences of 15 *Colletotrichum* isolates in *C. acutatum* complex and three in *C. orchidearum* complex. The species *Monilochaetes infuscans* (CBS 869.96) was selected as an outgroup. The tree was built using concatenated sequences of the ACT, CHS-1, GAPDH, TUB2, and ITS gene regions. Bootstrap values > 50% (1000 replication) are given at the nodes. Ex-type cultures are in bold. Colored blocks indicate clades containing isolates from *Camellia* plants.
Figure 4 – Phylogenetic tree using sequences of 6 Colletotrichum isolates in *C. boninense* complex. The species *C. gloeosporioides* (IMI 356878) was selected as an outgroup. The tree was built using concatenated sequences of the ACT, CAL, CHS-1, GAPDH, TUB2, and ITS gene regions. Bootstrap values > 50% (1000 replication) are given at the nodes. Ex-type cultures are in bold. Colored blocks indicate clades containing isolates from Camellia plants.

![Phylogenetic tree](image)

Figure 5 – PHI test of *C. puerense* and phylogenetically related isolates. PHI test value (Φw) < 0.05 indicates significant recombination within the dataset.

![PHI test](image)


Materials examined – China, Yunnan province, Pu’er City, on leaves of *Ca. japonica*, Mar. 2022; Jiangxi province, Ji’an City, on leaves of *Ca. japonica*, Dec. 2021; Zhejiang province,
Zhoushan City, on leaves of *Ca. japonica* and *Ca. oleifera*, Dec. 2021; Yunnan province, Pu’er City, on leaves of *Ca. sinensis*, Dec. 2021.

Notes – *Colletotrichum camelliae* belongs to the kahawae clade within the gloeosporioides complex. The species was first described by Massee on *Ca. sinensis* anthracnose from Sri Lanka (Willis 1899). Our study represents the first report of *C. camelliae* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 66–67 mm in 6 d at 25 °C with entire margin, aerial mycelium sparse, surface dark white, cottony, reverse pale-yellow. Conidia hyaline, smooth walled, cylindrical with obtuse ends, 14.2–17.2 × 5–6.5 μm, mean ± SD = 15.7 ± 1.5 × 5.8 ± 0.7 μm, L/W ratio = 2.7. Appressoria brown to black, round or oval.


Materials examined – China, Shanghai City, on leaves of *Ca. japonica*. Mar. 2022.

Notes – *Colletotrichum clivicola* belongs to the orchidearum species complex. This is the first report of *C. clivicola* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 67–70 mm in 6 d at 25 °C, aerial mycelium sparse, surface dark white. Conidia hyaline, smooth walled, curved, tapering towards apex and base, 19.4–25.2 × 5.1–7.8 μm, mean ± SD = 22 ± 0.9 × 5.7 ± 0.3 μm, L/W ratio = 3.9. The conidia are slightly bigger than those of the ex-type (CBS 125375) 19.5–24.5 × 4.5–7 μm, mean = 21.8 × 5.7 μm, L/W ratio = 3.8 (Yang et al. 2009). Appressoria brown to dark brown, smooth, irregularly shaped.


Materials examined – China, Zhejiang province, Fu yang City, on leaves of *Ca. oleifera* and *Ca. japonica*, Mar. 2022; Zhejiang province, Lin’an City, on leaves of *Ca. sinensis*, Dec. 2021; Zhejiang province, Zhoushan City, on leaves of *Ca. japonica*, Dec. 2021; Zhejiang province, Lishui City, on leaves of *Ca. oleifera*, Sep. 2021; Jiangxi province, Ji’an City, on leaves of *Ca. japonica*, Dec. 2021; Yunnan province, Pu’er City, on leaves of *Ca. japonica*, Mar. 2022.

Notes – *Colletotrichum fioriniae* belongs to the acutatum species complex. This is the first report of *C. fioriniae* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 44–48 mm in 6 d at 25 °C, aerial mycelium sparse, surface light pink. Conidia hyaline, smooth walled, cylindrical, both ends slightly acute, 11.5–16.1 × 5.8–6.6 μm, mean ± SD = 14.5 ± 0.8 × 5.8 ± 0.5 μm, L/W ratio = 2.5. Appressoria brown to dark brown, smooth, rounded or irregularly shaped.


Materials examined – China, Zhejiang province, Lin’an City, on leaves of *Ca. japonica*, Sep. 2021; Zhejiang province, Lin’an City, on leaves of *Ca. sinensis*, Dec. 2021; Zhejiang province, Fu yang City, on leaves of *Ca. japonica* and *Ca. sinensis*, Mar. 2022; Zhejiang province, Quzhou City, on leaves of *Ca. oleifera*, Sep. 2021; Zhejiang province, Lishui City, on leaves of *Ca. oleifera*, Sep. 2021; Yunnan province, Pu’er City, on leaves of *Ca. sinensis*, Mar. 2022; Jiangxi province, Ji’an City, on leaves of *Ca. japonica*, Dec. 2021.

Notes – *Colletotrichum fructicola* belongs to the gloeosporioides complex. The species is the most dominant *Colletotrichum* species on *Camellia* with a total of 75 isolates. This is the first report of *C. fructicola* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 65–68 mm in 6 d at 25 °C, flat with entire margin, aerial mycelium dense, surface light green to gray-green. Conidia hyaline, smooth walled, cylindrical, ends rounded, 15–18.5 × 4.0–7.2 μm, mean ± SD = 15.2 ± 1.5 × 4.6 ± 0.9 μm, L/W ratio = 3.3. The conidia of *C. fructicola* are larger than those of ex-type (MFLU 090228) 9.7–14 × 3–4.3 μm, mean = 11.53 × 3.55 μm, L/W ratio = 3.2.


Notes – Colletotrichum gloeosporioides belongs to the gloeosporioides complex. This is the first report of C. gloeosporioides causing anthracnose on Ca. japonica in China.

Colonies diam on MEA attained 71–73 mm in 6 d at 25 °C, flat with entire margin, aerial mycelium dense, surface gray-green. Conidia hyaline, smooth walled, cylindrical, ends bluntly rounded, 14.2–19.3 × 5.2–7.2 μm, mean ± SD = 14.5 ± 1.5 × 5.4 ± 0.5 μm, L/W ratio = 2.7. Appressoria oval or irregular, brown to black, crenate or slightly lobed at edge.

Colletotrichum jiangxiense F. Liu. & L. Cai, Persoonia 35: 82. 2015

Materials examined – China, Yunnan province, Pu’er City, on leaves of Ca. sinensis and Ca. japonica, Mar. 2021.

Notes – Colletotrichum jiangxiense belongs to the gloeosporioides complex. This is the first report of C. jiangxiense causing anthracnose on Ca. japonica in China.

Colonies diam on MEA attained 70–71 mm in 6 d at 25 °C, flat with entire margin, aerial mycelium dense, surface white. Conidia hyaline, smooth walled, cylindrical, ends bluntly rounded, 13.2–19.5 × 4.5–6.2 μm, mean ± SD = 15.2 ± 1.2 × 5.2 ± 0.5 μm, L/W ratio = 2.9. The conidia are similar with those of the ex-type isolate (CGMCC 3.17363) 15.2 ± 1.0 × 5.2 ± 0.4 μm, mean = 15.2 × 5.2 μm, L/W ratio = 2.9 of C. jiangxiense.


Materials examined – China, Yunnan province, Pu’er City, on leaves of Ca. sinensis and Ca. japonica, Mar. 2022.

Notes – Colletotrichum karstii belongs to the boninense species complex. This is the first report of C. karstii causing anthracnose on Ca. japonica in China.

Colonies diam on MEA attained 65–67 mm in 6 d at 25 °C, aerial mycelium sparse, surface dark white. Conidia hyaline, smooth walled, cylindrical, one end rounded and one end slightly acute, 13.4–15.5 × 5.6–6.8 μm, mean ± SD = 14.8 ± 0.9 × 6.3 ± 0.3 μm, L/W ratio = 2.3. Appressoria brown to dark brown, smooth, rounded or irregularly shaped.

Colletotrichum nymphaeae Van der Aa, European Journal of Plant Pathology 84(3): 109-115. 1978


Notes – Colletotrichum nymphaeae belongs to the acutatum species complex. This is the first report of C. nymphaeae causing anthracnose on Ca. japonica in China.

Colonies diam on MEA attained 49–50 mm in 6 d at 25 °C, aerial mycelium sparse, surface light pink. Conidia hyaline, smooth walled, cylindrical, one end rounded and one end slightly acute, 14.8–17.6 × 4.8–5.6 μm, mean ± SD = 15.2 ± 0.9 × 5.3 ± 0.5 μm, L/W ratio = 2.8. Appressoria dark brown, smooth, rounded or irregularly shaped.

Colletotrichum pandanicola Tibpromma & K.D. Hyde, MycoKeys 33: 25. 2018

Materials examined – China, Zhejiang province, Lin’an City, on leaves of Ca. japonica, Sep. 2021; Shanghai City, on leaves of Ca. japonica, Mar. 2022; Yunnan province, Pu’er City, on leaves of Ca. sinensis. Sep. 2021.

Notes – Colletotrichum pandanicola belongs to the gloeosporioides complex. This is the first report of C. pandanicola causing anthracnose on Ca. sinensis and Ca. japonica in China.
Colonies diam on MEA attained 62–65 mm in 6 d at 25 °C, flat with entire margin, aerial mycelium abundant, cottony, surface white. Conidia hyaline, smooth walled, cylindrical, ends rounded or one end slightly acute, 16.1–19.1 × 5–6.3 μm, mean ± SD = 17.3 ± 1.1 × 5.7 ± 0.4 μm, L/W ratio = 3.0.

**Colletotrichum plurivorum** Damm, Alizadeh & Toy. Sato, Studies in Mycology 92: 31. 2019

Materials examined – China, Yunnan province, Pu’er City, on leaves of *Ca. sinensis*. Mar. 2022.

Notes – *Colletotrichum plurivorum* belongs to the orchidearum species complex. Colonies diam on MEA attained 62–64 mm in 6 d at 25 °C, aerial mycelium sparse, surface dark white. Conidia hyaline, smooth walled, straight cylindrical, both ends bluntly rounded, 15.2–17.5 × 5.2–5.8 μm, mean ± SD = 15.8 ± 0.8 × 5.5 ± 0.3 μm, L/W ratio = 2.9. The conidia are slightly smaller than those of the ex-type (CBS 125474) 15–17 × 5.5 μm, mean = 16.0 ± 0.9 × 5.6 ± 0.1 μm, L/W ratio = 2.9 (Damm et al. 2019).

**Colletotrichum puerense** Peng XJ & Zhou XD, sp. nov.

Etymology – Referring to the tree host from which the fungus was isolated.

Type – China, Yunnan Province, Pu’er City, on leaves of *Ca. sinensis*, Mar. 2022. Sexual morphs were not observed. Asexual morphs developed on PDA. Vegetative hyphae were hyaline, smooth-walled, septate and branched. Setae not observed. Conidiomata acervular, yellow to light brown, conidiogenous cells were cylindrical to clavate, 9.5–15.2 × 3.5–5 μm, opening 1.5–2.5 μm. Conidia were smooth-walled, aseptate, cylindrical, both ends rounded, contents granular, 17–21 × 4–6 μm, mean ± SD = 19.1 ± 1.8 × 4.6 ± 0.6 μm, L/W ratio = 4.2. Appressoria were dark brown, irregular, often square to ellipsoid in outline, the margin lobate, 6–12 × 6–8 μm, mean ± SD = 8.9 ± 1.2 × 7± 0.5 μm, L/W ratio = 1.3.

Culture characteristics – Colonies diam on MEA attained 58–60 mm in 6 d at 25 °C with entire margin, aerial mycelium dense, surface grey-white with white margin; reverse pale yellow.

Notes – *Colletotrichum puerense* is phylogenetically closely related to *C. wuxiense* CGMCC 3.17894 and *C. wuxiense* JS1A44 (Fig. 2). However, *C. puerense* differs from the latter in ACT (with 98.83% sequence identity), CAL (100%), CHS-1 (99.52%), GAPDH (100%), TUB2 (99.31%) and ITS (99.62%) sequences. Moreover, *C. puerense* has larger conidia (YNS-22, 17–21 × 4–6 μm, mean ± SD = 19.1 ± 1.8 × 4.6 ± 0.6 μm) than those of *C. wuxiense* (mean = 16.9 × 5.5 μm). The PHI test (Φw = 1.0) detected no significant recombination between the isolates and *C. wuxiense* CGMCC 3.17894 and *C. wuxiense* JS1A44 (Fig. 5).


Materials examined – China, Zhejiang province, Fu yang City, on leaves of *Ca. oleifera* and *Ca. japonica*, Mar. 2022; Zhejiang province, Lin’an City, on leaves of *Ca. japonica*, Sep. 2021; Shanghai City, on leaves of *Ca. japonica*, Mar. 2022; Yunnan province, Pu’er City, on leaves of *Ca. sinensis*, Mar. 2022; Jiangxi province, Ji’an City, on leaves of *Ca. japonica*, Dec. 2021.

Notes – *Colletotrichum siamense* belongs to the gloeosporioides complex. The species has been recorded as a pathogen on a wide range of hosts (Weir et al. 2012) and is the dominant species on *Camellia* in this study.

Colonies diam on MEA attained 68–70 mm in 6 d at 25 °C, flat with entire margin, aerial mycelium dense, cottony, surface white. Conidia hyaline, smooth walled, cylindrical, ends rounded, 14.8–17.5 × 5.1–6.3 μm, mean ± SD = 15.5 ± 1.1 × 5.2 ± 0.5 μm, L/W ratio = 3.0. The conidia are bigger than those of the ex-type isolate (CGMCC 3.17363) 15.2 ± 1.0 × 5.2 ± 0.4 μm, mean = 15.2 × 5.2 μm, L/W ratio = 2.9 of *C. siamense*.


Notes – *Colletotrichum wuxiense* belongs in the gloeosporioides species complex. This is the first report of *C. wuxiense* causing anthracnose on *Ca. japonica* in China.

Colonies diam on MEA attained 62–63 mm in 6 d at 25°C, aerial mycelium dense, surface dark white or olive green. Conidia hyaline, smooth walled, cylindrical, ends rounded or one end slightly acute, 15.2–18.5 × 4.5–6.5 μm, mean ± SD = 15.5 ± 1.2 × 5.5 ± 0.5 μm, L/W ratio = 2.8. Appressoria dark-brown, irregular in shape.

Pathogenicity test

Sixteen representative strains of *Colletotrichum* species were selected for pathogenicity test on detached leaves of *Ca. oleifera*, *Ca. sinensis* and *Ca. japonica*. Under wounded conditions, all strains were pathogenic and infected leaves developed necrotic lesions at 14 dpi (Fig. 22). Diseased lesion lengths varied among the species and hosts (Fig. 23). There were significant differences (P < 0.05) of pathogenicity among the 16 species, using the test of least significant difference. Clearly, *C. camelliae* JXJ-20 showed the strongest pathogenicity on three *Camellia* plants, followed by *C. aenigma* SHJ-60, *C. jiangxiense* YNJ-17 and *C. alienum* FYJ-11, and the lesion diameters range from 5.12 mm to 25.67 mm. The other species such as *C. gloeosporioides* LAS-6, *C. pandanicola* SHJ-65, *C. siamense* LAJ-20, *C. wuxiense* LAJ-14, *C. puerense* YNS-22, *C. fiorinia* YNJ-1, *C. nympheae* YNS-42, *C. boninense* YNJ-4, *C. karstii* YNS-32, *C. clivicola* SHJ-58 and *C. plurivorum* YNS-6 showed weak pathogenicity with lesion diameters ranging from 3.06 mm to 1.35 mm. Noticeably, pathogenicity of the *C. camelliae* JXJ-20 to three *Camellia* detached leaves significantly varied. Species of *C. pandanicola* SHJ-65 and *C. clivicola* SHJ-58 were almost nonpathogenic to *Ca. sinensis* (1.33 mm–1.35 mm). No lesions were observed in controls.

**Figure 6** – Morphological characteristics of *C. aenigma*. a Front views of 6-d-old MEA culture. b Back views of culture. c-d Conidiophores. e-f Conidia. g-i Appressoria. a-i Isolate YNS-6. Scale bars = 10 μm.
Figure 7 – Morphological characteristics of *C. alienum*. a Front views of 6-d-old MEA culture. b Back views of culture. c-d Conidia. e-f Conidiophores. g-i Appressoria. a-i Isolate FYJ-11. Scale bars = 10 μm.

Figure 8 – Morphological characteristics of *C. boninense*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d Conidia. e Conidiophores. f-j Appressoria. Scale bars: d-j = 10 μm, c = 200 μm.
Figure 9 – Morphological characteristics of *C. camelliae*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d-e Conidia. f Conidiophores. g-j Appressoria. a-j Isolate JXJ-20. Scale bars: d-j = 10 μm, c = 200 μm.

Figure 10 – Morphological characteristics of *C. clivicola*. a Front views of 6-d-old MEA culture. b Back views of culture. c Ascomata. d Conidia. e-f Conidiophores. g-h Appressoria. a-h Isolate SHJ-58. Scale bars: d-h = 10 μm, c = 200 μm.
Figure 11 – Morphological characteristics of *C. fioriniae*. a, c Front views of 6-d-old MEA culture. b, d Back views of culture. e Conidiomata. f Conidia. g-h Conidiophores. i-j Appressoria. a-b, e, h Isolate YNJ-1. c-d, f-g, i-j Isolate DJO-36. Scale bars: f-j = 10 μm, e = 200 μm.

Figure 12 – Morphological characteristics of *C. fructicola*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d Conidia. e Conidiophores. f-i Appressoria. a-i Isolate LAS-24. Scale bars: d-i = 10 μm, c = 200 μm.
Figure 13 – Morphological characteristics of *C. gloeosporioides*. a, c Front views of 6-d-old MEA culture. b, d Back views of culture. e Conidiomata. f-g Conidia. h Appressoria. a-b, e, f Isolate LAJ-6. c-d, g-h LAS-6. Scale bars: f-h = 10 μm, e = 200 μm.

Figure 14 – Morphological characteristics of *C. jiangxiense*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d Conidia. e Conidiophores. f-i Appressoria. a-i Isolate YNJ-17. Scale bars: d-i = 10 μm, c = 200 μm.
**Figure 15** – Morphological characteristics of *C. karstii*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d-e Conidia. f Conidiophores. g-j Appressoria. a-j Isolate YNS-32. Scale bars: d-j = 10 μm, c = 200 μm.

**Figure 16** – Morphological characteristics of *C. nymphaeae*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d-e Conidia. f-g Conidiophores. h-i Appressoria. a-i Isolate YNS-42. Scale bars: d-i = 10 μm, c = 200 μm.
Figure 17 – Morphological characteristics of *C. pandanicola*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d-e Conidia. f Conidiophores. g Appressoria. a-g Isolate SHJ-65. Scale bars: d-g = 10 μm, c = 200 μm.

Figure 18 – Morphological characteristics of *C. plurivorum*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d Conidia. e-f Conidiophores. g-i Appressoria. a-i Isolate YNS-6. Scale bars: d-i = 10 μm, c = 200 μm.
Figure 19 – Morphological characteristics of *C. puerense*. a Front views of 6-d-old MEA culture. b Back views of culture. c Conidiomata. d Conidiophores. e Conidia. f-i Appressoria. a-i Isolate YNS-22. Scale bars: d-i = 10 μm, c = 200 μm.

Figure 20 – Morphological characteristics of *C. siamense*. a, c Front views of 6-d-old MEA culture. b, d Back views of culture. e Conidiomata. f Conidia. g Conidiophores. h-k Appressoria. a-b, e, i Isolate LAJ-20. c-d, f-h, j-k Isolate FYO-85. Scale bars: f-k = 10 μm, e = 200 μm.
Figure 21 – Morphological characteristics of *C. wuxiense*. a, c Front views of 6-d-old MEA culture. b, d Back views of culture. e Conidia. f Conidiophores. g-k Appressoria. a-b, e-i Isolate LAJ-14. c-d, j-k Isolate DJO-82. Scale bars: e-k = 10 μm.

Table 4 Morphology and lesion lengths of *Colletotrichum* causing on *Camellia* plants.

<table>
<thead>
<tr>
<th>Species</th>
<th>Conidial size</th>
<th>Mycelial growth diameter (mm/day)</th>
<th><em>Ca. oleifera</em> Lesion diameter on leaves (mm)</th>
<th><em>Ca. sinensis</em> Lesion diameter on leaves (mm)</th>
<th><em>Ca. japonica</em> Lesion diameter on leaves (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>C. aenigma</em></td>
<td>16.5 ± 0.5 × 5.1 ± 0.3</td>
<td>9.6 ± 0.4 ab</td>
<td>21.5 ± 2.5 b</td>
<td>2.6 ± 0.1 de</td>
<td>2.6 ± 0.6 de</td>
</tr>
<tr>
<td><em>C. alienum</em></td>
<td>16.0 ± 1.2 × 5.2 ± 0.7</td>
<td>10.7 ± 0.6 ab</td>
<td>2.0 ± 0.4 de</td>
<td>1.8 ± 0.2 de</td>
<td></td>
</tr>
<tr>
<td><em>C. camelliae</em></td>
<td>16.1 ± 1.5 × 5.3 ± 0.5</td>
<td>8.9 ± 0.4 ab</td>
<td>25.7 ± 3.7 a</td>
<td>4.4 ± 1.1 cde</td>
<td></td>
</tr>
<tr>
<td><em>C. fructicola</em></td>
<td>17.2 ± 1.3 × 6.2 ± 0.8</td>
<td>10.6 ± 0.6 ab</td>
<td>2.3 ± 0.1 de</td>
<td>2.2 ± 0.7 de</td>
<td></td>
</tr>
<tr>
<td><em>C. gloeosporioides</em></td>
<td>16.8 ± 0.5 × 6.3 ± 0.5</td>
<td>11.9 ± 0.4 a</td>
<td>1.5 ± 0.1 de</td>
<td>2.3 ± 0.3 de</td>
<td></td>
</tr>
<tr>
<td><em>C. jiangxiense</em></td>
<td>15.2 ± 1.2 × 5.2 ± 0.5</td>
<td>9.9 ± 0.5 ab</td>
<td>1.6 ± 0.1 de</td>
<td>1.8 ± 0.1 de</td>
<td></td>
</tr>
<tr>
<td><em>C. pandanicola</em></td>
<td>16.5 ± 0.8 × 5.2 ± 0.4</td>
<td>10.5 ± 0.3 ab</td>
<td>1.3 ± 0.1 de</td>
<td>2.0 ± 0.5 de</td>
<td></td>
</tr>
<tr>
<td><em>C. siamense</em></td>
<td>16.2 ± 0.7 × 5.9 ± 0.6</td>
<td>11.0 ± 0.9 ab</td>
<td>2.5 ± 0.7 de</td>
<td>1.6 ± 0.2 de</td>
<td></td>
</tr>
<tr>
<td><em>C. wuxiense</em></td>
<td>16.9 ± 1.1 × 5.5 ± 0.2</td>
<td>10.5 ± 0.2 ab</td>
<td>1.7 ± 0.2 de</td>
<td>2.8 ± 0.4 de</td>
<td></td>
</tr>
<tr>
<td><em>C. puerense</em></td>
<td>19.1 ± 1.8 × 4.6 ± 0.6</td>
<td>10.2 ± 0.5 ab</td>
<td>2.0 ± 0.1 de</td>
<td>2.3 ± 0.3 de</td>
<td></td>
</tr>
<tr>
<td><em>C. fioriniae</em></td>
<td>15.5 ± 1.2 × 5.2 ± 0.7</td>
<td>8.3 ± 0.4 b</td>
<td>2.1 ± 0.2 de</td>
<td>2.4 ± 0.1 de</td>
<td></td>
</tr>
<tr>
<td><em>C. nymphaeae</em></td>
<td>13.8 ± 1.1 × 6.0 ± 0.6</td>
<td>8.2 ± 1.0 b</td>
<td>2.5 ± 0.4 de</td>
<td>2.8 ± 0.6 de</td>
<td></td>
</tr>
<tr>
<td><em>C. boninense</em></td>
<td>12.5 ± 1.2 × 7.0 ± 0.4</td>
<td>10.2 ± 0.5 ab</td>
<td>1.9 ± 0.1 de</td>
<td>1.7 ± 0.1 de</td>
<td></td>
</tr>
<tr>
<td><em>C. karstii</em></td>
<td>14.5 ± 0.8 × 7.0 ± 0.5</td>
<td>9.8 ± 0.4 ab</td>
<td>1.8 ± 0.2 de</td>
<td>1.6 ± 0.1 de</td>
<td></td>
</tr>
<tr>
<td><em>C. clivicola</em></td>
<td>13.5 ± 0.8 × 6.0 ± 0.7</td>
<td>9.8 ± 0.3 ab</td>
<td>1.4 ± 0.1 e</td>
<td>2.0 ± 0.2 de</td>
<td></td>
</tr>
<tr>
<td><em>C. plurivorum</em></td>
<td>18.0 ± 1.2 × 5.8 ± 0.7</td>
<td>10.1 ± 0.6 ab</td>
<td>1.5 ± 0.1 de</td>
<td>1.9 ± 0.1 de</td>
<td></td>
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</tbody>
</table>
Figure 22 – Symptoms of 16 Colletotrichum species inoculated on leaves of Ca. oleifera, Ca. sinensis and Ca. japonica at 14th day.

Figure 23 – Lesion diameters on Ca. oleifera, Ca. sinensis and Ca. japonica leaves of 16 Colletotrichum species at 14th day after inoculation. Letters over the error bars indicate a significant difference at the $p = 0.05$ level.
Discussion

In this study, we employed morphological and multi-locus phylogenetic analyses to identify the Colletotrichum species associated with anthracnose from three important Camellia species in China. In total, we obtained a number of 167 isolates residing in four Colletotrichum species complexes including C. gloeosporioides, C. acutatum, C. boninense, and C. orchidearum. They represent 16 taxa with a newly described species, C. puereuse. Our study further confirmed that species of C. aenigma, C. alienum, C. boninense, C. camelliae, C. clivicola, C. fioriniae, C. fructicola, C. gloeosporioides, C. jiangxiense, C. karstii, C. nymphaeae, C. pandanicola and C. wuxiense can cause Ca. japonica anthracnose.

Species in the C. gloeosporioides complex such as C. fructicola, C. siamense, C. camelliae were most predominantly isolated from three Camellia plants. This is consistent with the previous studies of tea and tea-oil trees anthracnose (Liu et al. 2015, Zhu et al. 2015, Wang et al. 2016, Sun et al. 2021, Zhang et al. 2021). Other studies also revealed that C. gloeosporioides complex is the causal agent of anthracnose on a wide range of subtropical and tropical agricultural crops and tree species such as coffee, peach, pear, and chili (Liu et al. 2013, Cao et al. 2019, Jayawardena et al. 2020, Shi et al. 2021, Bhunjun et al. 2022, Tan et al. 2022). The broad host and geographic distribution of this Colletotrichum complex might be due to their higher genetic diversity to adopt to changing environmental conditions (McDonald & Linde 2002). The C. acutatum complex is the second major pathogen group confirmed from the present study. Colletotrichum fioriniae was obtained from three Camellia species with five isolates from Ca. oleifera, six from Ca. japonica and one from Ca. sinensis. The second species in the complex C. nymphaeae was isolated from Ca. japonica (one isolate) and Ca. sinensis (two isolates). Further, six isolates in C. boinense complex and three isolates in the C. orchidearum complex associated with Ca. sinensis and Ca. japonica anthracnose were identified. It is worth noting that C. acutatum and C. gloeosporioides complexes were recognized as frequently occurring endophytic fungi in Camellia (McDonald & Linde 2002, Liu et al. 2015). This implies that the members of these Colletotrichum complexes in Camellia may be converted from endophytes to pathogens (Carroll 1988, Wheeler et al. 2019). More importantly, the study described a previously unknown taxon causing Camellia anthracnose, namely C. puereuse from Ca. sinensis in Yunnan province. The possible reason is that the various climate environment there might be helpful promoting species diversity (Yang et al. 2004, Liu et al. 2021, Bhunjun et al. 2022).

Pathogenicity test using 16 representative Colletotrichum isolates of each obtained species showed that all species were pathogenic, and significant differences existed. Pathogenicity of C. camelliae JX-20, C. aenigma SHJ-60, C. alienum FYJ-11 were significantly stronger than other Colletotrichum species on three Camellia hosts. This is consistent with the previous results demonstrating that C. camelliae has strong pathogenicity on different plant hosts (Wang et al. 2016, Wang et al. 2020, Zhang et al. 2021). In our study, the pathogenicity of C. camelliae JX-20 to three plant hosts varied significantly. Species of C. aenigma SHJ-60, C. alienum FYJ-11, C. jiangxiense YNJ-17 were more pathogenic to Ca. oleifera while C. pandanicola SHJ-65 and C. clivicola SHJ-58 were least pathogenic to Ca. sinensis. The reasons behind this could be the differences in host resistance and fungal biological characters. Further, only one variety of each host plant was used and it is not enough to fully reflect the actual condition (Mo et al. 2018). Additional research using more genetic clones should be conducted to assess the pathogenicity of the pathogens.

In summary, this represents the first comprehensive investigation of Colletotrichum species occurring on three important plant species of Camellia, especially those from Ca. japonica. A total number of 16 Colletotrichum species were characterized and dominant taxa were determined. Our result disclosed that these fungal taxa are pathogenic to three plant hosts, and their pathogenicity varied. The knowledge gained and the diversity of Colletotrichum species in Camellia gathered provides a useful clue for resistant-germplasm selection and disease management.
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