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DNA was extracted from 10 representative isolates, and a multilocus sequence analysis

amplified using corresponding primers. The obtained sequences were deposited in GenBank under the following accession numbers: ITS (LT745886, LT745887, and LT972205 to LT972210), ACT (LT976516 to LT976525), CHS-1(LT976833 to LT976842), GAPDH (LT976492 to LT976501), and TUB2 (LT976506 to LT976515). The deposited sequences showed high identity percentages (99 to 100%) with C. acutatum sequences present in GenBank (KX069826, KX069821 for ITS; JQ005839, KY171912 for ACT; MF979822, KY856133 for CHS-1; KX069805, KM252191 for GAPDH, and GU183311,

was carried out. Sequences of rDNA-ITS (ITS), partial actin (ACT), chitin synthase (CHS-1), glyceraldehyde-3-phosphate dehydrogenase (GAPDH), and  $\beta$ -tubulin-2 (TUB2) were

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GU183314 for TUB2). Phylogeny inferred from combined datasets of the five genes and *C. acutatum* species-complex sequences from GenBank revealed that all feijoa isolates were included in the *C. acutatum* subclade (100% branch bootstrap support value after 1,000 replicates). Koch's postulates were fulfilled by spraying a spore suspension (5 × 10<sup>4</sup> conidia/ml) from monoconidial colonies of 10 sequenced isolates on 50 leaves and inoculating 30 µl of the same spore suspension on 50 wounded mature feijoa fruit. As controls, 10 leaves and 10 mature feijoa fruit were inoculated with sterile distilled water. Following inoculation, leaves and fruits were maintained in a growth chamber at  $25 \pm 1^{\circ}$ C. Pathogenicity tests were carried out twice. Symptoms identical to those observed in the field appeared within 10 to 15 days on inoculated fruits and leaves and identified by both conidia morphology and molecular methods. The infection of feijoa plants by *C. acutatum sensu lato* has been previously reported in Auckland, New Zealand (Lardner et al. 1999). To our knowledge, this is the first report of *C. acutatum* on feijoa plants in Italy.



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