

Morphology and Pathogenicity of *Calonectria floridana*, *Calonectria kyotensis*,
and *Calonectria uniseptata*

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ABSTRACT

The perfect and imperfect states of *Calonectria floridana*, *C. kyotensis*, and *C. uniseptata* were compared morphologically and in pathogenicity tests with an isolate of *Cylindrocladium scoparium*. The three species of *Calonectria* reacted similarly to 20 hosts in leaf and root pathogenicity tests, but differed from *Cylindrocladium scoparium* in their pathogenicity to roots of three species

of *Lupinus* and three cultivars of peach. They were morphologically indistinguishable, and all produced an imperfect state identified as *Cylindrocladium floridanum*. On the basis of priority, *Calonectria kyotensis* is correctly assigned as the perfect state of *Cylindrocladium floridanum*.

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In April 1967, *Cylindrocladium floridanum* Sob. & Seymour was designated as a new species occurring on peach roots in Florida (5), and in March 1969, its perfect state was described as *Calonectria floridana* Sob. (4). In January 1968, *Calonectria kyotensis* Terashita was described with a *Cylindrocladium* conidial state from leaves of *Acacia dealbata* (6), and in April 1968, *Calonectria uniseptata* Gerlach was described as the perfect state of *Cylindrocladium scoparium* Morgan (2). Because of the apparent similarity of the three descriptions, it was decided to compare the perfect and imperfect states of the three species of *Calonectria* morphologically and in pathogenicity tests with an isolate of *Cylindrocladium scoparium* used in previous studies (1, 3, 4, 5).

MATERIALS AND METHODS.—Inocula for leaf pathogenicity tests were prepared by suspending conidia scraped from 10-day-old cultures grown on rehydrated Difco potato-dextrose agar (PDA, 39 g/liter of distilled water) in 10-ml portions of distilled water. The resulting suspensions were filtered through a single thickness of cheesecloth, adjusted to contain 20,000 conidia/ml, blended for 30 sec after adding

Triton B-1956 (active ingredient, 77% modified phthalic glyceryl alkylid resin) at a rate of 0.05 ml/20 ml of suspension, and sprayed on the leaves of test plants (Table 1). Mixtures containing 0.05 ml of Triton B-1956/20 ml of distilled water were sprayed on the leaves of control plants. All plants were maintained in a mist chamber for 24 hr at 28 C and 95 to 100% relative humidity after inoculation. Results were recorded 7 days after inoculation.

Inocula for root pathogenicity tests were prepared by comminuting 10-day-old cultures grown on PDA in 30-ml portions of distilled water. Sixty-five ml of the resulting suspensions were mixed into the top 3 cm of soil in each pot. Pots containing control plants received a like quantity of autoclaved inoculum. All plants were established in 6-inch clay pots containing soil fumigated with methyl bromide at a rate of 454 g/137.2 cm³. Results of root pathogenicity tests were recorded 4 weeks after the soil was infested. Plants in both studies were maintained in a greenhouse where temperatures varied from 17 C at night to 35 C during the day. The hosts used are listed in Table 1.

Transfers from the type cultures of *C. floridana*, *C. kyotensis* (ATCC-18834), and *C. uniseptata* were

TABLE 1. Number of plants with leaves and roots infected by *Calonectria floridana*, *C. kyotensis*, *C. uniseptata*, and *Cylindrocladium scoparium*

	<i>Cal. floridana</i>		<i>Cal. kyotensis</i>		<i>Cal. uniseptata</i>		<i>C. scoparium</i>	
	Leaves	Roots	Leaves	Roots	Leaves	Roots	Leaves	Roots
<i>Callistemon citrinus</i> Stapf	8/10	7/10	8/10	7/10	10/10	8/10	10/10	8/10
<i>C. rigidus</i> R. Br.	10/10	6/10	9/10	6/10	9/10	5/10	10/10	7/10
<i>Crotalaria spectabilis</i> Roth	6/10	0/10			8/10	1/10	10/10	3/10
<i>Eucalyptus camaldulensis</i> Dehnhardt	10/10	10/10	10/10	10/10	10/10	10/10	10/10	10/10
<i>E. grandis</i> J. E. Sm.	9/10	6/10			10/10	7/10	10/10	9/10
<i>E. robusta</i> J. E. Sm.	10/10	8/10	10/10	8/10	9/10	6/10	10/10	9/10
<i>E. rudis</i> Endl.	10/10	9/10	10/10	10/10	10/10	9/10	10/10	10/10
<i>E. saligna</i> J. E. Sm.	10/10	7/10	10/10	7/10	10/10	10/10	10/10	8/10
<i>E. tereticornis</i> J. E. Sm.	10/10	7/10	10/10	8/10	10/10	9/10	10/10	10/10
<i>Melaleuca leucadendra</i> L.	10/10	6/10	10/10	7/10	10/10	7/10	10/10	9/10
<i>Lupinus albus</i> L.	0/10	2/10		2/10	2/10	3/10	3/10	8/10
<i>L. angustifolius</i> L.	1/10	3/10		4/10	1/10	1/10	3/10	9/10
<i>L. luteus</i> L.	1/10	2/10		2/10	0/10	3/10	3/10	5/10
<i>Prunus persica</i> L.								
'Elberta'		10/10		8/10		8/10		3/10
'Nemaguard'		10/10		9/10		10/10		4/10
'Okinawa'		10/10		9/10		10/10		1/10
<i>Rhododendron obtusum</i> (Lindl.) Planch.								
'Rentschler's Pink'	9/10	3/10			10/10	2/10	10/10	4/10
'Valentine'	10/10	1/10	10/10	2/10	10/10	1/10	10/10	2/10
'White Christmas'	10/10	4/10	9/10	3/10	10/10	5/10	10/10	5/10
'White Water'	10/10	3/10	10/10	3/10	10/10	4/10	10/10	5/10

used in pathogenicity and morphological studies. The culture of *Cylindrocladium scoparium* was obtained from lesions on leaves of azalea collected at Fort Myers, Fla., in 1965.

Morphological comparisons of the *Cylindrocladium* state of *Calonectria floridana*, *C. kyotensis*, and *C. uniseptata* with *Cylindrocladium scoparium* were made from fructifications found in 10-day-old cultures on PDA. Sexual structures of the three species of *Calonectria* were compared after 21 days' growth on PDA.

RESULTS AND DISCUSSION.—Similar results were obtained with the imperfect states of *Calonectria floridana*, *C. kyotensis*, and *C. uniseptata* in leaf and root pathogenicity tests. They differed from the isolate of *Cylindrocladium scoparium*, however, in that they were highly virulent to roots of

three cultivars of peach and slightly virulent to roots of three species of *Lupinus*, whereas *C. scoparium* was slightly virulent to peach roots and highly virulent to roots of lupines. These results agree with previous comparative studies with *C. floridanum* and *C. scoparium* (3).

A comparison of the imperfect states of the three species of *Calonectria* revealed only minor differences, and suggested that they were the same (Table 2). Vesicles of the imperfect state of *C. uniseptata* were globose and typical of *Cylindrocladium floridanum*, rather than ellipsoidal as expected in isolates of *C. scoparium*. It was concluded that the imperfect states of the three species of *Calonectria* are *Cylindrocladium floridanum*, and that Gerlach (2) incorrectly identified the imperfect state of *Calonectria*

TABLE 2. Physical aspects of *Calonectria floridana*, *C. kyotensis*, and *C. uniseptata* after 10 days' growth on potato-dextrose agar

	<i>C. floridana</i>	<i>C. kyotensis</i>	<i>C. uniseptata</i>
Perithecia	Orange to reddish brown 335-510 × 310-405 μ	Orange to reddish brown 330-490 × 285-420 μ	Orange to reddish brown 310-480 × 300-425 μ
Asci	Clavate 83-145 × 14-22 μ	Clavate 74-136 × 14-21 μ	Clavate 79-133 × 13-19 μ
Ascospores	Mostly 1 septate 19-45 × 5.5-6.3 μ	Mostly 1 septate 18-49 × 5.5-6.3 μ	Mostly 1 septate 19-50 × 5.3-6.8 μ
Vesicles	Globose 8.2-17.9 μ	Globose 8.8-19.0 μ	Globose 8.2-20.4 μ
Conidia	Mostly 1 septate 34-55 × 3.5-4.9 μ	Mostly 1 septate 30-56 × 3.5-5.4 μ	Mostly 1 septate 36-47 × 3.5-4.8 μ

uniseptata as *Cylindrocladium scoparium*.

Examination of the sexual states of the three species of *Calonectria* showed no essential differences among them (Table 2). The paraphyses reported by Gerlach (2) for *C. uniseptata* were not observed, nor were they found in the other species. It was concluded, therefore, that *C. floridana*, *C. kyotensis*, and *C. uniseptata* are synonymous. Because *C. kyotensis* predates *C. floridana* and *C. uniseptata* by several months, *C. kyotensis* is correctly assigned as the perfect state of *Cylindrocladium floridanum*.

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