# Species of Colletotrichum and Glomerella Pathogenic to Tomato Fruit

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#### **ABSTRACT**

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Eleven species of Colletotrichum and Glomerella and one subspecies of Colletotrichum were inoculated into detached, ripening tomato fruit. Colletotrichum sp. 1, Colletotrichum sp. 2, C. coccodes, C. dematium, C. dematium var. truncata, C. destructivum, C. gloeosporioides, C. graminicola, C. trichellum, Glomerella sp., G. glycines, and G. gossypii were pathogenic. C. falcatum and C. trifolii were nonpathogenic. In general, isolates of Colletotrichum sp. 1, C. gloeosporioides, C. dematium, and G. glycines were most aggressive. Differences in virulence among isolates occurred within G. glycines, Colletotrichum sp. 1, and C. gloeosporioides, with the greatest variability occurring in the latter species. C. dematium var. truncata, C. graminicola, C. trichellum, G. glycines, and G. gossypii are reported as pathogenic to tomato fruit for the first time.

Additional key words: anthracnose, barnyardgrass, big spurge, blue verbena, cocklebur, cowpea, johnsongrass, morningglory, okra, redroot pigweed, ryegrass, spotted spurge, three-seeded mercury, watermelon

Anthracnose is a major disease of fresh market, home garden, and processing tomatoes (2,4,10). It is most serious on processing types, which are allowed to ripen in the field (1,4). The latter condition is conducive to wounding of fruit, which predisposes them to infection

Although Colletotrichum coccodes (Wallr.) Hughes is often cited as the primary causal fungus of anthracnose (1,2,9,10), several other anthracnose fungi have been isolated from tomato fruit and their pathogenicity demonstrated in vitro (3,5,8). At least five species of anthracnose fungi have been isolated by us from ripe processing tomatoes in Mississippi.

There is little information on the relative pathogenicity and host range of fungi capable of inciting anthracnose of tomato fruit. This study was conducted to determine the relative ability of 12 anthracnose fungi isolated from various crops and weeds to incite anthracnose when inoculated into tomato fruit.

## MATERIALS AND METHODS

Isolation and identification of fungi. Sources of Colletotrichum and Glomerella isolates used for inoculation of tomato fruit are presented in Table 1. Fungi from

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0191-2917/82/12115303/\$03.00/0 © 1982 American Phytopathological Society tomato and apple were isolated from fruit lesions; others were isolated from leaf spots. Sections from the margins of diseased leaf and fruit tissue were surfacesterilized with 1% sodium hypochlorite, plated on Difco potato-dextrose agar (PDA), and incubated at 22 C. Species of Colletotrichum and Glomerella growing from plated material were identified or, to induce sporulation and facilitate identification, were cultured on sections of sterilized soybean stems in test tubes, on V-8 juice agar, or on PDA. Descriptions of Colletotrichum or Glomerella species reported by von Arx (13), Mordue (6,7), Sutton (11), and Tiffany and Gilman (12) were utilized for species determination.

Inoculation of tomato fruit. Sections of broom straw approximately 5 mm long and 1 mm in diameter were placed on 5-day-old PDA cultures of each fungus, incubated at 22 C for 1 wk, and used as

Ripe tomato fruits from the market, selected for uniformity in size and maturity, were surface-sterilized in 0.5% sodium hypochlorite for 1 min, rinsed three times in tap water, and allowed to dry. Six replicate fruits were inoculated per fungus by inserting infested straws into fruit using a sterile forceps, after which points of inoculation were covered with petrolatum. Fruits similarly treated with noninfested straws served as controls. The treatments and controls were completely randomized and incubated in the dark at 28 C. This experiment was conducted three times.

Disease rating. Disease severity was evaluated 3 days after inoculation using the following visual rating scale based on lesion diameter: 1 = no anthracnose lesion,  $2 = lesions \le 5 \text{ mm}$ , 3 = lesions

6-10 mm, 4 = lesions 11-15 mm, and 5 =lesions > 15 mm.

Isolation of fungi from inoculated fruit. Three days after inoculation, the fruit epidermis surrounding the point of inoculation was swabbed with 95% ethanol, peeled back with a sterile forceps, and a small portion of tissue aseptically removed from beneath and plated on PDA. The plated tissue was incubated at 22 C for 1 wk, during which fungi growing from it were identified.

## RESULTS AND DISCUSSION

All fungal isolates except C. falcatum and C. trifolii were pathogenic (Table 1) and reisolated from lesions on inoculated fruit. C. dematium var. truncata, C. graminicola, C. trichellum, G. glycines, and G. gossypii are recorded as pathogenic to tomato fruit for the first time. Thirty-one of the pathogenic isolates incited lesions rating a disease index of 4, 13 an index of 3, and five an index of <3. Additionally, two isolates of Colletotrichum sp. 1 (W2-6 and BG-3) and one of Glomerella sp. (VV-1) tested but excluded from the statistical analysis were highly pathogenic. Disease indexes for fruit infected with W2-6, BG-3, and VV-1 were 4.7, 4.7, and 4.8, respectively.

Colletotrichum sp. 1, a falcate-spored fungus, was provisionally identified as C. dematium. We refer to it by the former name to distinguish it from C. dematium and C. dematium var. truncata, whose identities are not in doubt. According to von Arx (13), C. trichellum, C. graminicola, and C. dematium are the only falcate-spored Colletotrichum species. Colletotrichum sp. 1 is neither trichellum nor graminicola, and although it can be referred to as dematium using von Arx's broad concept of this species, it differs from dematium in growth rate, size and shape of conidia, and morphology of appressoria (Roy, unpublished). Thus, it appears that there is ample justification for distinguishing between Colletotrichum sp. 1 and C. dematium. A comparative study of the morphology and pathogenicity of these two fungi is currently being conducted.

In general, isolates of Colletotrichum sp. 1, C. gloeosporioides (= G. cingulata (Stonem.) Spauld. & Schrenk), C. dematium, and G. glycines were the most aggressive (Table 1). C. destructivum and two isolates of Colletotrichum sp. 1 (JG-2 and JG-3) were least aggressive. Colletotrichum sp. 2, C. graminicola, C. trichellum, and G. gossypii did not differ

in pathogenicity. C. dematium was more aggressive than C. trichellum and two isolates of C. dematium var. truncata (T-2 and T-3). These data indicate that even though C. coccodes is emphasized in screening for resistance to anthracnose •(1,2), other species may be of equal or

greater importance and should, as Barksdale (3) cautioned, be considered in screening programs. In addition, the relative pathogenicity of species should be considered in establishing priorities for such programs.

Within some fungal species, statistically

significant differences in virulence occurred among isolates, occasionally even among isolates from the same host, and the magnitude of these differences varied among species. Such differences which occurred within Colletotrichum sp. 1, C. gloeosporioides, and G. glycines,

Table 1. Relative ability of Colletotrichum and Glomerella isolates from various hosts to incite anthracnose lesions when inoculated on tomato fruit in vitro

Fungus		Source		
	Isolate no.	Common name	Scientific name	Disease rating
Colletotrichum sp. 1	BP-4	Sweet pepper	Capsicum annuum L.	5.0 a
C. gloeosporioides (Penz.) Sacc.	WM-1	Watermelon	Citrullus lanatus (Thunb.) Matsum. & Nakai	5.0 a
Glomerella glycines Hori	GG-1	Soybean	Glycine max (L.) Merr.	5.0 a
Colletotrichum sp. 1	OK-1	Okra	Abelmoschus esculentus (L.) Moench.	4.9 a
Colletotrichum sp. 1	W8-7	Big spurge	Euphorbia nutans L.	4.9 a
Colletotrichum sp. 1	MG-1	Morningglory	Ipomoea purpurea (L.) Roth	4.9 a
C. gloeosporioides	CB-6	Cocklebur	Xanthium pennsylvanicum Wallr.	4.9 a
Colletotrichum sp. 1	BD-1	Broadleaf dock	Rumex obtusifolius L.	4.9 a
Colletotrichum sp. 1	W1-1	Pigweed	Amaranthus sp.	4.9 a
Colletotrichum sp. 1	W4-2	Spotted spurge	Euphorbia maculata L.	4.9 a
Colletotrichum sp. 1	CF-1	Soybean	G. max	4.8 ab
Colletotrichum sp. 1	CB-4	Cocklebur	X. pennsylvanicum	4.8 ab
C. gloeosporioides	CB-5	Cocklebur	X. pennsylvanicum	4.8 ab
Colletotrichum sp. 1	WM-2	Watermelon	C. lanatus	4.8 abc
C. gloeosporioides	BP-1	Sweet pepper	C. annuum	4.8 abc
Colletotrichum sp. 1	W2-1	Redroot pigweed	Amaranthus retroflexus L.	4.7 abcd
C. dematium (Fr.) Grove	W8-6	Big spurge	E. nutans	4.7 abcd
G. glycines	OK3-A	Okra	A. esculentus	4.6 abcde
C. dematium	AF-4	Alfalfa	Medicago sativa L.	4.6 abcde
Colletotrichum sp. 1	CX-1	Tomato	Lycopersicon esculentum Mill.	4.6 abcdef
C. dematium	RC-1	Red clover	Trifolium pratense L.	4.6 abcdef
Colletotrichum sp. 1	CP-I	Cowpea	Vigna unguiculata (L.) Walp.	4.5 abcdef
G. gossypii Edg. G. glycines	GGS-3	Cotton	Gossypium hirsutum L.	4.2 bcdefg
s. glycines C. gloeosporioides	CP-6	Cowpea	V. unguiculata	4.2 cdefg
C. coccodes (Wallr.) Hughes	GTC-1	Soybean	G. max	4.1 defgh
C. gloeosporioides	RG-2 KZ-2	Ryegrass	Lolium multiflorum Lam.	4.1 efgh
C. gloeosporioides	GC-2	Kudzu	Pueraria lobata (Willd.) Ohwi.	4.1 efgh
C. graminicola (Ces.) Wilson	CG-1 <sup>t</sup>	Sweet pepper	C. annuum	4.1 efgh
C. gloeosporioides	AF-1	Alfalfa	M. sativa	4.1 efgh
Colletotrichum sp. 2	RG-1	Alfalfa	M. sativa	4.0 efghi
C. dematium Grove var.	KO-1	Ryegrass	L. multiflorum	4.0 efghi
truncata (Schw.) v. Arx	T-1	Soybean		
C. trichellum (Fr.) Duke	TC-5 <sup>u</sup>	Bamboo	G. max	3.9 fghi
C. gloeosporioides	RC-2	Red clover	Bambusa	3.8 ghi
C. coccodes	CX-3	Tomato	T. pratense	3.8 ghij
C. gloeosporioides	GC-3	Soybean	L. esculentum G. max	3.8 ghij
C. dematium var. truncata	T-3	Purple nutsedge		3.8 ghij
Colletotrichum sp. 1	CX-2	Tomato	Cyperus rotundus L.	3.8 ghij
. gloeosporioides	W3-2	Three-seeded mercury	L. esculentum	3.8 ghij
C. dematium var. truncata	T-2°	Soybean	Acalypha ostryaefolia L. G. max	3.7 ghij
Colletotrichum sp. 1	CT-4	Cotton	G. max G. hirsutum	3.7 ghij
. gloeosporioides	GC-1 <sup>w</sup>	Apple	Malus sylvestris Mill.	3.5 hij
. gloeosporioides	W3-1	Three-seeded mercury	A. ostryaefolia	3.4 ij
. gloeosporioides	W4-6	Spotted spurge	E. maculata	3.2 jk
. glycines	W8-4	Big spurge	E. nutans	3.2 jk 2.8 kl
. gloeosporioides	W6-1	Pigweed	Amaranthus sp.	2.8 kl 2.7 kl
'. destructivum O'Gara	CDS-2 <sup>x</sup>	Alfalfa	M. sativa	2.7 KI 2.6 I
Colletotrichum sp. 1	JG-3	Johnsongrass	Sorghum halepense (L.) Pers.	2.51
olletotrichum sp. 1	JG-2	Johnsongrass	S. halepense	2.5 I 1.8 m
. falcatum Went	CFC-3 <sup>y</sup>	Sugarcane	Saccharum officinarum L.	1.6 mn
. trifolii Bain & Essary	CTF-1 <sup>z</sup>	Alfalfa	M. sativa	1.0 mn 1.2 n
ontrol				1.2 n 1.0 n

Figures followed by the same letter are not significantly different (P=0.05) according to Duncan's multiple range test. Disease rating scale based on lesion diameter: 1 = no anthracnose lesions, 2 = lesions ≤5 mm, 3 = lesions 6-10 mm, 4 = lesions 11-15 mm, and 5 = lesions >15 mm.

<sup>1</sup> ATCC 11870.

<sup>&</sup>lt;sup>u</sup> ATCC 34168.

<sup>&</sup>lt;sup>v</sup> ATCC 18013.

<sup>\*</sup>Cultures obtained from L. E. Trevathan, Department of Plant Pathology and Weed Science, Mississippi State University.

<sup>\*</sup> ATCC 11869.

<sup>&</sup>lt;sup>y</sup> ATCC 12088.

<sup>&</sup>lt;sup>2</sup> ATCC 32358.

with the greatest variability occurring within C. gloeosporioides—indicate that choice of isolate may be an important consideration in screening for resistance.

Within pathogenic species, 26 isolates originated from 10 different crops (other than tomato) and an equal number from 14 different weeds. Some are newly recorded on certain hosts: Colletotrichum sp. 1 on watermelon, redroot pigweed, spotted spurge, big spurge, johnsongrass, cocklebur, morningglory, and barnyardgrass; C. gloeosporioides on spotted spurge and three-seeded mercury; C. coccodes on ryegrass; C. dematium on big spurge; Colletotrichum sp. 2 on ryegrass; G. glycines on okra, cowpea, and big spurge; and Glomerella sp. on blue verbena.

Our data suggest that numerous species of anthracnose fungiare potentially capable of infecting injured tomato fruit in the field. Further, they suggest that numerous crops and weeds, common in Mississippi and elsewhere, could serve as sources of inoculum. The extent to which these fungi occur on these hosts needs to be determined because it could have important implications in the epidemiology and control of anthracnose.

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