Effect of plant products on the mycelial growth and conidial germination of Colletotrichum gloeosporioides causing anthracnose disease of mango fruits

K. PRABAKAR, P. MUTHULAKSHMI, T. RAGUCHANDER AND V.K. PARTHIBAN Department of Plant Pathology, Tamil Nadu Agrl. University, Coimbatore - 641 003, Tamil Nadu.

Abstract : The effect of plant extracts on the mycelial growth and conidial germination of Colletotrichum gloeosporioides causing anthracnose disease of mango fruits tested under in vitro conditions. Twenty six plant extracts belonging to 21 families were tested for the effect on mycelial growth and conidial germination of the pathogen. The plant extracts of Adenocalyma alleaceum and Bougainvillea spectabilis were most effective in inhibiting the mycelial growth completely upto tenth day. The conidial germination was recorded at different intervals upto 24 h. As observed in mycelial growth there was no conidial germination in Adenocalyma alleaceum and Bougainvillea spectabilis.

Key words: Mango, anthracnose, biocontrol, plant products.

ntroduction

Mango (Mangifera indica L.) is a major ruit crop in India and other tropical countries. n India, twenty different genera of fungi are nown to attack mango fruits during post harvest andling (Pathak, 1980) of which anthracnose aused by Colletotrichum gloeosporioides was he most important one (Snowdon, 1990). Anthracnose has been reported to cause heavy lamage in mango to the extent of 6.0 to 15.0 er cent in different parts of India. (Tandon, 967). Fungicides are primary means of controlling ost harvest diseases and they have recently ome under special scrutiny as posing potential oncogenic risks when applied to processed foods. Biological control offers an alternative for the hemical control of post harvest diseases of fruits which includes use of plant products and antagonistic organisms. Keeping this in view, the present study was taken up to find out effective plant products for the management of mango anthracnose.

Materials and Methods

Efficacy of plant extracts

Twenty six plant extracts were collected for evaluating their antifungal activities against C gloeosporioides both for spore germination and mycelial growth at Plant Pathology lab, TNAU. Among them, two species found effective were selected and used for further studies.

Preparation of plant extracts (Cold water extract) Fresh leaf / fruit / flower materials of

nlant species were used for extraction. They

were first washed with distilled water, then ground in a pestle and mortar by adding sterile water at the rate of 1:1 W/V and filtered through muslin cloth. This formed the standard plant extract solution (100%). The above extracts were diluted to 10 per cent by adding sterile The plant extracts so prepared were heated to 40-50°C for 10 min. to avoid contamination (Jagannathan and Narasimhan, 1987).

Effect of plant extracts on mycelial growth

The inhibitory effect of the plant extracts on mycelial growth was assayed by poisoned food technique (Bagchi and Das, 1968).

Effect of plant extracts on conidial germination

The conidia of the pathogen used in the study were harvested separately by flooding with sterile water and scrapping the culture with a glass rod (Montgomery and Moore, 1938).

Results and Discussion

Several plant products were found to inhibit the growth of pathogens hence an experiment was conducted to screen the plant products against C. gloeosporioides. Twenty six plant extracts belonging to 21 families were tested for the mycelial growth and conidial germination of the pathogen.

All the plant products tested reduced the mycelial growth significantly compared to control (Table 1). Among them three were highly effective (more than 75 per cent reduction) two were effective (50 to 75 per cent reduction) twelve were moderately effective (25 to 50 per cent reduction) and nine were less effective (0 to 25 per cent reduction). The two plant extracts Adenocalyma alleaceum and Bougainvillea spectabilis were most effective and inhibited the mycelial growth completely till the tenth day. While in Piper cubabe the mycelial growth was inhibited till the sixth day. The growth was completely inhibited upto 4 days in the plant extracts Abutilon indicum, Acalypha indica, Agiratum coninjoides, Allium cepa (leaf), Boerhaavia diffusa, Ocimum sanctum, Terminalia cephula and Tridax procumbens.

The effect of plant extracts on the conidial germination of C gloeoyporioides was studied at different intervals and the results are presented

in (Table 2) The data show that all the plant extracts tested significantly reduced the conidial germination over control. The conidial germination was recorded at different intervals upto 24 h. As observed in mycelial growth there was no conidial germination in Adenocalyma alleaceum and Bougainvillea spectabilis The Piper cutbabe extract followed the above two extracts in inhibiting the conidial germination.

Among the twenty six plant extracts screened against C. gloeosporioides, Adenocalyma alleaceum and Bougainvillea spectabilis were very effective in controlling the growth, conidial germination and disease development. Several workers have shown the possibility of controlling the post harvest diseases using plant products. Ark and Thompson (1959) effectively protected the peaches from brown rot caused by Monilinia fructicola

Table 1. Effect of plant extracts on the mycelial growth of C. gloeosporioides

	Ra	Per cent			
Plant extracts		P	reduction over control		
	4	6	. 8	10	
Abutilon indicum (leaf)	0.0i	171	40 ^k	551m	37.37
Acalypha indica (leaf)	0.0	101	37 ^m	52 ⁿ	41.11
Adenocalyma alleaceum (leaf)	00i	0.0 ⁿ	0.0	001	100.0
Aegle marmelous (leaf)	001	12 ^m	24 ^p	399	55.41
Agiratum coninjoides (leaf)	004	18 ¹	41	561	36.58
Allium cepa (leaf)	00i	10 ⁿ	229	31'	65.23
Anderographis paniculatus (leaf)	09 ^h	19h	461	57 [±]	34.99
Azadirachta indica (leaf)	20 ⁱ	31°	64°	83°	6.00
Azadirachta indica (seed)	125	27#	51h	641	27.52
Boerhaavia diffusa (leaf)	00,	16k	40k	55 ^m	38.05
Bougainvillea spectabilis (leaf)	001	00°	004	001	100.00
Casuarina equisetifolia (leaf)	13E	30 ^d	5218	69 ^t	21.55
Catheranthus roseus (leaf)	19 ^d	31°	61 ^d	774	12.80 .
Catheranthus roseus (flower)	12fg	28°	53°	66 ^g	24.92
Eucalyptus globulus (leaf)	24ªb	38"	67ab	86 ^d	1.81
Ipomea cornea (leaf)	15°	29°	55°	72*	18.80
Lawsonia alba (leaf)	12fg	28 ^r	51gh	65h	26.39
Mirabilis jalaba (leaf)	10h	24€	52gh	63 ¹	28.99
Nerium odorum (leaf)	001	17)	38 ⁿ	56lm	36.58
Ocinum sanctum (leaf)	001	00°	12 ^r	211	75.88
Piper cubabe (fruit)	23 ^b	37*	66b	86 ^d	1.81
Prosopis juliflora (leaf)	001	16k	35°	51°	42.58
Terminalia chebula (fruit)	001	141	32°	46b	47.59
Tridax procumbens (leaf)	18 ^d	32°	60 ^d	76ª	13.59
Vitex negundo (leaf)	21°	33 ^b	64°	831	5.66
Zizyphus jujuba (leaf) Control	25*	38*	68*	88*	

^{*} Mean of three replications

In a column, means followed by a common letter are not significantly different at the 5% level by DMRT

ible 2. Effect of plant extracts on the mycelial growth of C. gloeosporioides

Plant extracts		Per cent reduction over control			
	6	12	18	24	
butilon indicum	10.00 (18.43) ^b	27.33 (31.52) ¹	46.33 (42.90)	55.00 (47.87) ^b	17.97
calypha indica	7.33 (15.71) ^a	25.00 (30.00) ¹	43.67 (41.36)	52.67 (46.53) ^b	40.60
denocalyma alleaceum	0.00 (3.18) ¹	0.00 (3.18) ⁿ	0.00 (3.18)	0.00 (3.18) ^b	100.00
egle marmelous	4.00 (11.54)*	15.67 (23.32) ¹	37.00 (37.46)	44.00 (41.55) ^b	50.38
giratum coninjoides	7.67 (16.08) ^d	40.33 (39.42) ^r	59.67 (50.58)	67.33 (55.14) ⁶	24.07
llium cepa	3.00 (9.97) ^b	12.67 (20.85)bk	33.33 (35.26)	40.00 (39.23) ^b	54.89
nderographis paniculatus	7.33 (15.71) ^d	25.00 (30.00) ¹	56.33 (48.64)	68.33 (55.75) ^b	22.94
zadirachta indica	10.00 (10.43) ^h	48.33 (44.04) ^c	70.33 (57.01)	84.00 (66.42) ^b	5.27
zadirachta indica	7.00 (15.34) ^d	42.33 (40.59) ^b	59.67 (50.58)	67.67 (55.35) ^b	23.68
oerhaavia diffusa	6.00 (14.18) ^{cf}	36.67 (37.27) ^s	49.33 (44.62)	53.67 (47.10) ^b	39.47
lougainvillea spectabilis	0.00 (3.18) ^j	0.00 (3.18) ⁿ	0.00 (3.18)	0.00 (3.18) ^b	100.00
Casuarina equisetifolia	10.33 (18.75) ^b	46.00 (42.71) ^d	59.67 (49.01)	69.33 (56.37) ^b	21.81
Catheranthus roseus	9.33 (17.79) ^{bc}	48.33 (44.04) ^b	70.00 (56.79)	85.33 (67.48) ^b	3.77
Catheranthus roseus	10.67 (19.07) ¹⁶	35.67 (36.67) ^b	58.00 (49.60)	66.00 (54.33) ^b	25.57
Eucalyptus globulus	11.00 (19.37)*	51.33 (45.76) ^b	70.00 (56.79)	86.67 (68.57) ⁵	2.26
pomea cornea	8.33 (16.78)°	45.67 (45.52) ^b	64.33 (53.33)	73.33 (58.91) ⁶	17.30
awsonia alba	6.67 (14.97)°	45.67 (45.52) ^b	63.33 (52.73)	70.33 (57.61) ^b	20.68
Mirabilis jalaba	7.67 (16.08) ^b	46.67 (43.09) ^b	64.00 (53.13)	72.00 (58.05) ¹	18.80
Nerium odorum	4.67 (12.48) ^b	40.00 (39.23) ¹	60.00 (50.77)	66.33 (54.53) ^h	25.20
Ocinum sanctum	5.33 (13.35)	28.00 (31.95) ^b	46.00 (42.71)	58.00 (49.60) ^b	34.59

Table 2 continues...

Plant extracts		Per cent reduction over control			
	1/11/2				
	6	12	18	24	4 61
Piper cubabe	2.33 (8.78) ^b	10.00 (18.43) ^b	24.33 (29.55)	28.00 (31.95) ^b	18.42
Prosopis juliflora	11.00 (19.37) ^h	50.33 (45.19) ^b	70.00 (56.79)	85.33 (67.48) ^b	3.77
Terminalia chebula	8.67 (17.12) ^b	30.00 (33.21) ^b	48.67 (44.24)	52.33 ⁻ (46.33) ^b	10.98
Tridax procumbens	8.00 (16.43) ^b	22.00 (27.97) ^b	39.00 (38.65)	48.67 (44.24) ^b	15.11
Vitex negundo	8.67 (17.12) ^b	46.67 (43.09) ^b	67.00 (54.94)	80.67 (63.92) ^b	9.02
Zizyphus jujuba	10.33 (18.75) ^b	49.67 (44.81) ^b	69.00 (56.17)	84.33 (66.68) ^b	4.90
Control	11.67 (19.98) ^b	52.33 (46.34) ^b	82.67 (65.41)	88.67 (70.33) ^b	

^{*} Mean of three replications

In a column, means followed by a common letter are not significantly different at the 5% level by DMR? The values in the parentheses are arcsine transformed values.

with deodorized garlic preparations. Sholberg and Shimizu (1991) reported the decay inhibition in strawberry and peach fruit by hinokitol, an antifungal compound derived from the trunk of Japanase cypress. Ghaouth and Wilson (1995) screened the 300 species of plants belonging to 43 families against *Bollytis cinerea*, among them five per cent showed fungicidal activity.

Conclusion

Among the twenty six plant extracts screened against C. gloeosporioides, Adenocalyma alleaceum and Bougainvillea spectabilis were very effective in controlling the growth and conidial germination.

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