

RESEARCH ARTICLE

Evaluation of efficacy of *Neem* oil, castor oil, carbendazim, *Trichoderma harzianum*, *Trichoderma viride* and *Pseudomonas fluorescens* against *Alternaria carthami*

■ D. Amrutha Gayathri and V. Krishna Rao

SUMMARY

The efficacy of two botanicals viz., *Neem* oil and castor oil, one fungicide i.e., carbendazim and three bioagents viz., *Trichoderma harzianum*, *Trichoderma viride*, *Pseudomonas fluorescens* were tested in vitro and in vivo against *Alternaria carthami* inciting leaf spot safflower leaf spot/blight. In vitro efficacy of botanicals and fungicide was evaluated by poison food technique against *Alternaria carthami*. In vitro efficacy of bioagents was evaluated by dual culture technique against *Alternaria carthami*. In in vitro evaluation of fungicide and botanicals carbendazim found to be most effective and showed maximum inhibition of mycelial growth (43.33%) followed by *Neem* oil (30.53%). Among the bioagents maximum inhibition of radial growth of the test pathogen was noticed in *P. fluorescens* (87.36 %) which was found on par with *T. viride* (86.22 %). Mycelial growth of test pathogen was inhibited to an extent of 81.08 per cent *T. harzianum*. In in-vivo evaluation, combined seed treatment with of *P. fluorescens* (10 g kg⁻¹ seed) + carbendazim (2 g kg⁻¹ seed)+ *Neem* oil (10 ml kg⁻¹ seed) was effective in controlling *Alternaria* leaf spot/blight.

Key Words : Botanicals, Bioagents, Safflower, Leaf spot/blight

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Safflower (*Carthamus tinctorius* L.) is one of the important oilseed crops of the world. It is popular not only for seed and oil, but also for its brightly

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coloured petals, high levels of linoleic acid (75-80%) in oil and amino acids (Nagaraj, 2009). *Alternaria* leaf spot/blight is one of the most serious diseases of safflower reducing yield by 50%. Infected seed are often smaller with reduced oil content. It mostly infects leaves, stems, heads and seeds. Plant extracts are known to possess antifungal properties (Nene and Thaplial, 1993). The presence of antifungal compounds in higher plants is well recognized and considered valuable for plant disease control. *Neem* has attracted special interest of scientists due to presence of variety of bioactive compounds of

Trichoderma sp. *Pseudomonas fluorescens* are commercially applied as bio control agents against many fungal pathogens. So keeping in view the present study was undertaken to know the efficacy of two botanicals and three bioagents for the control of *Alternaria carthami* inciting leaf spot safflower leaf spot/blight.

MATERIAL AND METHODS

Two botanicals viz., *Neem* oil, castor oil, one fungicide i.e., carbendazim were evaluated by poisoned food technique with three replications on potato dextrose agar (PDA) and incubated at $28 \pm 2^\circ\text{C}$ for seven days.

The fungal and bacterial antagonists were evaluated against the test pathogen *Alternaria carthami* in laboratory by dual culture technique. Petri dishes (90 mm) containing PDA was inoculated with 5 mm diameter mycelia disc of 7 days old culture of *Alternaria carthami* and fungal/bacterial antagonists at equal distance from periphery. Inoculated plates were incubated $28 \pm 2^\circ\text{C}$. Each treatment was replicated four times. After required period of incubation i.e., in the control plate growth reached 90 mm diameter, the radial growth of pathogen was measured. Per cent inhibition over control was assessed.

$$R_N \frac{C-T}{C} \times 100$$

where, R= Per cent inhibition

C = Radial growth of pathogen colony in control

T = Radial growth of pathogen colony in treatments

Under glass house studies the healthy seeds of safflower (cv. NIRA) were surface sterilized and artificially inoculated with the test pathogen by rolling the seeds in 10 days old sporulating culture grown on PDA. The inoculated seeds were kept for 8 h in Petri plates having moistened blotter papers. After incubation, inoculated seeds were treated separately by coating with potential botanicals and bioagents by imposing different treatments. Seeds from each treatment were then sown in pots (20 cm diameter) filled with sterilized soil @ five seeds per pot. Observations on pre emergence mortality, post emergence mortality, per cent seedling emergence were recorded after 35 days.

Treatment details :

Design: CRD

Replications: 4

Treatments: 8

T₁- Seed inoculation with test pathogen followed by seed treatment (10 g kg⁻¹seed) with potential bioagent

T₂- Seed inoculation with test pathogen followed by carbendazim seed treatment @ 2 g kg⁻¹seed.

T₃- Seed inoculation with test pathogen followed by seed treatment with neem oil @ 10ml kg⁻¹seed

$$T_4 = T_1 + T_2$$

$$T_5 = T_2 + T_3$$

$$T_6 = T_1 + T_3$$

$$T_7 = T_1 + T_2 + T_3$$

$$T_8 = \text{Inoculated control}$$

RESULTS AND DISCUSSION

The results are presented in Tables 1 to 4.

In *in vitro* evaluation, out of two botanicals tested *Neem* oil was found to be effective in inhibiting the mycelia growth (30.53%). Similar results were obtained by Ghewande (1989) and Usman *et al.* (1991) who reported the antifungal properties of neem based products.

All the bioagents viz., *Trichoderma harzianum*, *Trichoderma viride*, *Pseudomonas fluorescens* inhibited mycelia growth of the pathogen. *Pseudomonas fluorescens* inhibited maximum mycelia growth with a mean inhibition of (87.36 %) which was found at par with *T. viride* (86.22 %). Mycelial growth of test pathogen was inhibited to an extent of 81.08 per cent in *T. harzianum*. Similar results were obtained by Amaresh (2000) who reported that among fungi *T. viride* and *T. harzianum* overgrew and inhibited the growth of *A. helianthi*, while the bacterium *P. fluorescens* produced

Table 1 : Effect of botanicals on growth of *A. carthami* in vitro

Treatments	* Radial growth (mm)	Per cent inhibition
Neem oil (<i>Azadirachta indica</i>)	20.83	30.53 (33.50)
Castor oil (<i>Ricinus communis</i>)	26.2	12.66 (20.77)
Carbendazim	17	43.33 (41.14)
Control	40	0.00 (0.00)
C.D. (P=0.05)		3.696
S.E. ±		1.116

*Mean of three replications; Figures in parenthesis are angular transformed values

Table 2 : Effect of bioagents on growth of *A. carthami* in vitro

Treatments	* Radial growth (mm)	Per cent inhibition
<i>Trichoderma viride</i>	12	86.22 (66.192)
<i>Trichoderma harzianum</i>	15	81.02 (64.20)
<i>Pseudomonas fluorescens</i>	11.3	87.36 (69.17)
Control	90	0.00 (0.00)
C.D. (P=0.05)		1.803
S.E. ±		0.544

*Mean of three replications; Figures in parentheses are angular transformed values

Table 3 : Effect of seed treatment of botanicals/bioagents/fungicide on Pre emergence mortality of safflower cv. NIRA against *A. carthami* under glass house conditions

Sr. No.	Treatments	*Pre emergence mortality (%)	Per cent decrease over control
1.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed)	13.27 (21.35)	75.62
2.	Carbendazim (2 g kg ⁻¹ seed)	20.20 (26.69)	62.89
3.	Neem oil (10 ml kg ⁻¹ seed)	26.833 (31.17)	50.71
4.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed) + Carbendazim (2 g kg ⁻¹ seed)	5.08 (12.97)	90.66
5.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed) + Neem oil (10 ml kg ⁻¹ seed)	6.19 (14.40)	88.62
6.	Carbendazim (2 g kg ⁻¹ seed)+ Neem oil(10 ml kg ⁻¹ seed)	13.01 (21.12)	76.10
7.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed) + Carbendazim (2 g kg ⁻¹ seed)+ Neem oil (10 ml kg ⁻¹ seed)	1.20 (6.29)	97.97
8.	Control	54.44 (47.54)	-
	C.D. (P=0.05)	2.552	
	S.E. ±	0.844	

*Mean of three replications; Figures in parentheses are angular transformed values.

Table 4 : Effect of seed treatment of botanicals/bioagents/fungicide on Post emergence mortality of safflower cv. Nira against *A. carthami* under glass house conditions

Sr. No.	Treatments	*Post emergence mortality (%)	Per cent decrease over control
1.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed)	6.637 (14.91)	88.5
2.	Carbendazim (2 g kg ⁻¹ seed)	13.51 (21.55)	76.59
3.	Neem oil (10 ml kg ⁻¹ seed)	19.00 (25.82)	67.08
4.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed) + Carbendazim (2 g kg ⁻¹ seed)	6.67 (14.95)	88.44
5.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed) + Neem oil (10 ml kg ⁻¹ seed)	6.88 (15.19)	88.20
6.	Carbendazim (2 g kg ⁻¹ seed)+ Neem oil(10 ml kg ⁻¹ seed)	13.30 (21.38)	96.96
7.	<i>Pseudomonas fluorescens</i> (10 g kg ⁻¹ seed) + Carbendazim (2 g kg ⁻¹ seed)+ Neem oil (10 ml kg ⁻¹ seed)	0.753 (4.69)	99.43
8.	Control	57.73(49.44)	-
	C.D. (P=0.05)	2.287	
	SE (m)±	0.756	

*Mean of three replications; Figures in parentheses are angular transformed values.

maximum inhibition zone.

All the seed treatments were significantly superior in reducing the pre emergence and post emergence mortality seed treatment with combined treatment of with *Pseudomonas fluorescens* (10 g kg⁻¹ seed) + carbendazim (2 g kg⁻¹ seed) + neem oil (10 ml kg⁻¹ seed) resulted in high per cent reduction of pre and post emergence mortality (97.97 % and 99.43 %, respectively) followed by *P. fluorescens* (10 g kg⁻¹ seed) + carbendazim (2 g kg⁻¹ seed) when compared to control (54.44 %). The beneficial effect of seed treatments with bioagents and fungicides in minimizing the pre and post emergence mortality is in accordance with that of Govindappa *et al.* (2011) in safflower.

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