

IN VITRO STUDIES ON THE EFFECT OF DIFFERENT FUNGICIDES AGAINST MYCELIAL GROWTH OF Sclerotium rolfsii, THE CAUSAL AGENT OF STEM ROT IN GROUNDNUT

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ABSTRACT

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Groundnut is an important oil seed crop and is regarded as 'king of oil seeds'. Among several biotic stresses of groundnut, stem rot caused by *Sclerotium rolfsii* causes up to 80 per cent yield loss. Saprophytic nature, broad host range and ability to produce resting structures make this pathogen difficult to control. In this experiment, ten fungicides *viz.*, hexaconazole, azoxystrobin, tebuconazole, pyraclostrobin, carbendazim, carboxin, validamycin, chlorothalonil, mancozeb and copper oxy chloride were tested against the mycelial growth of the pathogen at four concentrations i.e. 500 1000, 1500 and 2000 ppm. Among these, hexaconazole, tebuconazole, azoxystrobin and carboxin showed 100 per cent mycelial inhibition at all the concentrations tested, whereas carbendazim proved to be least effective in inhibiting the mycelial growth of *S. rolfsii*, followed by copper oxychloride and validamycin.

KEYWORDS: Groundnut, fungicides, mycelial inhibition, Sclerotium rolfsii

INTRODUCTION

Groundnut which is popularly regarded as the 'King of oil seeds' is one of the most important oil seed crops grown in India. It is rich in oil and protein content. It is also good source of minerals and vitamins. India is one of the largest producers of groundnut in the world. Due to increasing population and decreasing area under cultivation, more production must be obtained from limited area. Soil borne diseases are one of the major factors that limit groundnut productivity and production. Among the soil borne pathogens, *Aspergillus niger* Van Tieghem, *Sclerotium rolfsii* Sacc, and *Rhizoctonia bataticola* Taub are the major limiting pathogens.

More than 500 species of plants are being attacked by *Sclerotium rolfsi*, of which, the most common hosts are legumes, crucifers, and cucurbits (Punja, 1985). It is a soil borne pathogen and can survive at all climatic conditions through resting structures called sclerotia. Production of round and brown color sclerotia is the characteristic feature of this pathogen. The pod yield loss due to stem rot disease is 10-25 per cent, but under severe diseased conditions yield losses may range to up 80 per cent (Rodriguez-Kabana *et al.*, 1975). Patil and Rane (1982) reported yield loss up to 10 to 50 per cent due to this disease. Adiver (2003) reported the yield loss of 15-70 per cent in groundnut is due to leaf spot, rust and stem rot singly or in combination. Chemical control can be suggested in those fields where stem rot disease is the major limiting factor in groundnut production. The efficacy of the chemical used depends on method of application, number of sprays, time of sprays and the dosages of appropriate chemicals used. Hence, in the present study, efforts were made to study the effect of already existing chemicals and new molecules on mycelial growth of the fungus.

MATERIAL AND METHODS

Efficacy of ten fungicides *viz.*, hexaconazole, azoxystrobin, tebuconazole, pyraclostrobin, carbendazim, carboxin, validamycin, chlorothalonil, mancozeb and copper oxy chloride were tested against the mycelial growth of the pathogen at four different concentrations i.e. 500 ppm, 1000 ppm, 1500 ppm and 2000 ppm. The details of fungicides, their recommended dosage and the concentrations tested in the present study were given in Table 1.

Poisoned food technique was employed to evaluate the efficacy of these chemicals. Stock solution of 30,000 mg a.i. ml⁻¹ was prepared for each fungicide. From the stock solution, required quantity was pipetted out and added to required quantity of molten PDA medium in conical flasks to obtain final test concentration. Twenty ml of fungicide amended medium was poured in each pre-sterilized Petri plate. A mycelial disc of 5 mm diameter

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was cut from 15 day old culture of *S. rolfsii* using sterile cork borer under aseptic conditions and placed at the centre of each poisoned plate. Control was maintained by placing fungal discs in plates containing untreated (non poisoned) medium. The experiment was laid out in CRD with four replications per treatment .The inoculated Petri plates were incubated at $28 \pm 2^{\circ}$ C in BOD incubator.

The data on radial growth of the fungal mycelium was recorded 4 days after inoculation. Per cent inhibition of mycelial growth of test pathogen over control was calculated by using the formula mentioned below. The data were analysed statistically using CRD.

$$I = C - T / C \times 100$$

where, I = Per cent mycelia inhibition of test pathogen, C = Radial growth (mm) in monocultured check, T = Radial growth (mm) in treatment plates.

RESULTS AND DISCUSSION

Mycelial growth of S. rolfsii varied in its sensitivity to the fungicides tested at varied concentrations. It was most sensitive to fungicides tebuconazole, azoxystrobin, carboxin, hexaconazole and showed 100 per cent inhibition at all the concentrations tested. Similar results were obtained by Johnson et al. (2008) by evaluation of five fungicides hexaconazole, propiconazole, mancozeb, chlorpyriphos and quinalphos against S. rolfsii in vitro and reported that hexaconazole at a concentration of 1000, 1500 and 2000 ppm and propiconazole at a concentration of 500, 750 and 1000 ppm completely inhibited the growth of S. rolfsii. Similarly, Perez et al. (2009) evaluated the in vitro reaction of fungicides to S. rolfsii and showed that tebuconazole inhibited mycelial growth and sclerotial production. Further, these triazoles are also effective against leaf spot disease (Adiver and Anahosur, 1995) preventing defoliation, which forms substrate for the growth of the pathogen.

The fungus was least sensitive to the fungicide carbendazim at all the concentrations tested and the difference in the mycelial growth was significant. Copper oxy chloride was less effective in inhibiting the mycelial growth (20.73%) at 1000 ppm and the sensitivity has increased to 48.10% with increase in the concentration to 2500 ppm. But, one more non-systemic fungicide, chlorothalonil was highly effective (100%) in inhibiting the mycelia growth at concentrations of 1500 ppm and above, while the inhibition was 54.76 per cent at 1000

ppm. The efficacy of antifungal antibiotic, validamycin in inhibiting the mycelial growth is higher (80.70%) at 1000 ppm and the sensitivity decreased with decrease in concentration and the differences are significant. Among all the fungicides tested, carbendazim proved to be least effective in inhibiting the mycelial growth of *S. rolfsii* followed by copper oxychloride and validamycin (Table 2 and Fig. 1). Less efficacy of carbendazim was also reported by Nagamani *et al.* (2018).

The efficacy of fungicides in inhibiting the mycelial growth is as follows: Tebuconazole = azoxystrobin = carboxin = mancozeb = hexaconazole > chlorothalonil > pyraclostrobin > validamycin > copperoxychloride > carbendazim.

Prabhu and Hiremath (2003) and Arunasri *et al.* (2011) reported that the triazole groups of fungicides *viz.*, hexaconazole, propiconazole, difenconazole were highly inhibitory to the growth of *S. rolfsii*.

From the above findings and in comparison with earlier research, it can be predicted that there is no much variability in the pathogen. As the pathogen is soil borne, seed treatment with effective chemical is practical, soil application being expensive, especially in developing country like India where small land holdings prevail. However, fungicide control of foliar diseases has some potential effect on control of stem and pod rots. Finally, integration of cultural practices with potential biological and effective chemicals may offer effective solution for management of the disease rather than using chemicals alone and the present study helps in identification of effective chemical with its right dosage.

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S. No.	Chemical	Active ingredient	Recommended dose (%)	Concentration tested (ppm)	
	Fungicide				
1.	Carbendazim	50 WP	0.2	500, 1000, 1500, 2000	
2.	Pyraclostrobin	20 EG	0.2	500, 1000, 1500, 2000	
3.	Azaxystrobin	23 SC	0.2	250, 500, 750, 1000	
4.	Hexaconazole	5 EC	0.2	500, 1000, 1500, 2000	
5.	Tebuconazole	25.9 EC	0.1	500, 1000, 1500, 2000	
6.	Validamycin	3 L	0.1	250, 500, 750, 1000	
7.	Carboxin	Carboxin 37.5 + Thiram 37.5 SD	0.2	500, 1000, 1500, 2000	
8.	Chlorothalonil	75 WP	0.3	1000, 1500, 2000, 2500	
9.	Mancozeb	75 WP	0.3	1000, 1500, 2000, 2500	
10.	Copper oxychloride	50 WP	0.25	1000, 1500, 2000, 2500	

Table 1.	Details (of fungicides	employed in	the study

Table 2. Effect of fungicides on the mycelial growth of S. rolfsii after 4 days of incubation

S. No	Fungicide	Trade name	Concentration (ppm)	Mycelial growth (cm)	Per cent inhibition over control (%)
1	Copperoxychloride	Dhanucop	1000	3.57	20.73
	(50% WP)	•			(27.06**)
			1500	3.10	31.10
					(33.88)
			2000	2.77	38.47
					(38.31)
			2500	2.33	48.10
					(43.89)
2	Mancozeb (75% WP)	Dithane M-	1000	0.00	100.00
		45			(90.00)
			1500	0.00	100.00
					(90.00)
			2000	0.00	100.00
					(90.00)
			2500	0.00	100.00
					(90.00)
3	Chlorothalonil (75% WP)	Daconil	1000	2.03	54.76
					(47.71)
			1500	0.00	100.00
					(90.00)
			2000	0.00	100.00
					(90.00)
			2500	0.00	100.00
					(90.00)

Cont...

Table 2. Cont...

S.	Fungicide	Trade name	Concentration	Mycelial	Per cent inhibition
No.	-		<u>(ppm)</u>	growth (cm)	over control (%)
4	Hexaconazole	Trigger	500	0.00	100.00
	(5% EC)		1000	0.00	(90.00)
			1000	0.00	100.00
			1500	0.00	(90.00)
			1500	0.00	100.00
			••••	0.00	(90.00)
			2000	0.00	100.00
_	~	.		0.00	(90.00)
5	Carboxin	Vitavax	500	0.00	100.00
	(Carboxin 37.5% +			0.00	(90.00)
	Thiram 37.5%)		1000	0.00	100.00
					(90.00)
			1500	0.00	100.00
					(90.00)
			2000	0.00	100.00
					(90.00)
6	Tebuconazole	Torque	500	0.00	100.00
	(25.9% EC)				(90.00)
			1000	0.00	100.00
					(90.00)
			1500	0.00	100.00
					(90.00)
			2000	0.00	100.00
					(90.00)
7	Pyraclostrobin (20%	Insignia	500	1.43	68.07
	EG)				(55.57)
			1000	1.03	76.97
					(61.30)
			1500	0.47	89.57
					(71.15)
			2000	0.00	100.00
					(71.15)
8	Carbendazim (50 WP)	Dhanustin	500	4.43	1.46
-					(5.68)
			1000	4.13	8.10
					(16.40)
			1500	3.97	11.83
					(20.09)
			2000	3.87	14.03
			-000	2107	(21.93)
9	Azoxystrobin	Onestar	250	0.00	100.00
9	(23% SC)	Uncotai	200	0.00	(90.00)
			500	0.00	100.00
			500	0.00	(90.00)
			750	0.00	100.00
			750	0.00	(90.00)
			1000	0.00	100.00
			1000	0.00	(90.00)

Effect of different fungicides against Sclerotium rolfsii

Table 2. Cont...

S. No.	Fungicide	Trade name	Concentration (ppm)	Mycelial growth (cm)	Per cent inhibition over control (%)
10	Validamycin	Sheathmar	250	1.97	56.27
	(3% L)				(48.58)
			500	1.73	61.47
					(51.61)
			750	1.30	71.03
					(57.45)
			1000	0.86	80.70
					(63.94)
11	Control			4.50	0.00
					(0.00)
				C.D.	1.967
				SE(m)	0.698
				SE(d)	0.987
				C.V.	1.755

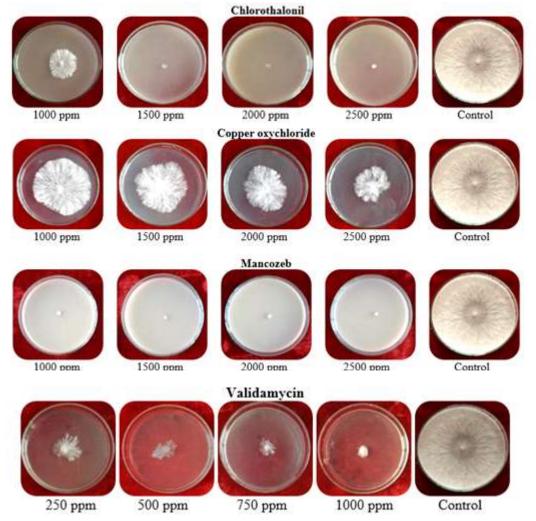


Fig. 1. Effect of different fungicides at different concentrations on mycelial growth of Sclerotium rolfsii

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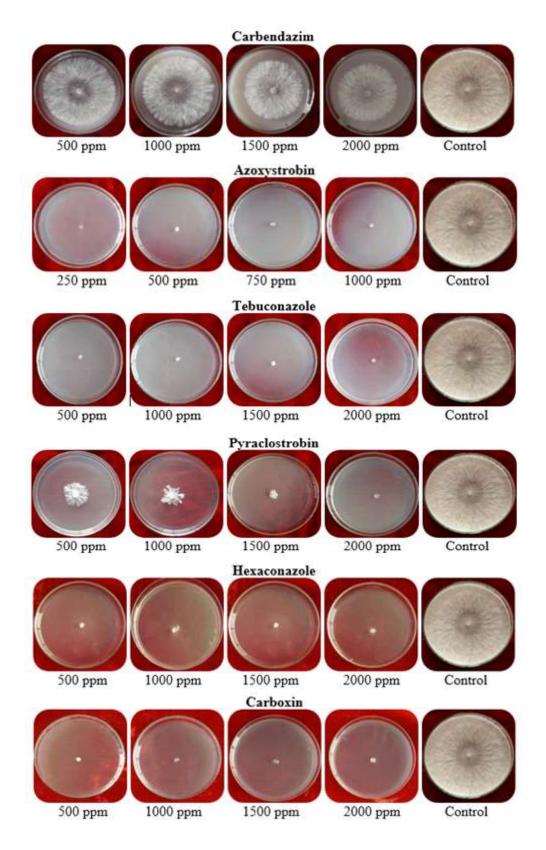


Fig. 1. Effect of different fungicides at different concentrations on mycelial growth of Sclerotium rolfsii

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