

Efficacy of fungicides and bio-pesticide against the *Sclerotinia sclerotiorum* causing Sclerotinia rot of Mustard

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Abstract: Two methods were used for evaluating of fungicides. In case of in vitro condition all the fungicides, and bio pesticide were significantly superior over control in inhibiting the growth of the pathogen Vitavax (0.1 %), Bavistin (0.1%) and Benomyl (0.1 %) each were the most effective as they completely inhibited (100%) the growth of pathogen. Whereas Neemark (0.5%) was noted as least effective bio pesticide which inhibited the growth of fungus only upto 44.44 %. Ridomyl (0.2 %) and Chlorothalonil (0.2 %) were found to be the next best in superiority in inhibiting the growth of the pathogen. The spraying of Vitavax (0.1%) at an interval of 10 days was found more effective under field condition as is evident from its corresponding yield in both the years. The next best fungicide Bavistin (0.1%) was found statistically at par with Vitavax (0.1%) in respect of reduction in disease severity and enhancement of seed yield. However, Neemark (0.5%) proved to be the least effective under field condition also. Benomyl @ 0.1% ranked 3rd in effectiveness which was significantly superior with other tested fungicides such as Indofil M-45 (0.25), Ridomyl and Chlorothalonil in respect of decreasing the disease severity but was at par in respect of seed yield.

keywords: disease, pathogen, pesticides, sclerotinia sclerotiorum, brassica juncea

I. Introduction

Indian mustard [Brassica juncea (L.) Czern. & Coss.] Commonly known as “rai” is most important oil yielding crucifer crop belongs to family Brassicaceae. The rapeseed - mustard group accounts for 24% of the total oil seed production of which Indian mustard alone covers about 70%. The crop is grown extensively in Rajasthan, Maharashtra, U.P., Gujarat and Punjab. Brassica juncea is erect, much branched, 3' to 6' high annual plant with slender and tapering root. The stem branches from the axil of the fourth or fifth leaf upward. Lower leaves petiolated, green, sometimes with a whitish bloom, ovate to obovate, variously lobed with toothed, scalloped or frilled edges, lyrate-pinnatisect, with 1-2 lobes or leaflets on each side and a larger sparsely sectose, terminal lobe. Upper leaves sub-entire, short stalked, 30-60 mm long, 2-3.5 mm wide, the fruit is siliqua. The pods are bi-locular with a false septum between two halves. The oil content varied from 37 to 47 per cent in different species of rapeseed-mustard. In India, mustard ravaged by several major diseases viz. Alternaria blight (Alternaria brassicae), White rust (Albugo candida), Powdery mildew (Erysiphe cruciferarum) and Sclerotinia rot (Sclerotinia sclerotiorum) which influences the quantity and quality of yield. The pathogen affects many crops in India, particularly rapeseed-mustard and has become a widespread and destructive in mustard growing parts (Ghasolia et al., 2004) and take a heavy toll of yield (Chauhan et al., 1992). In mustard growing areas, this disease led to complete crop failure, as the disease incidence has been recorded up to 80 per cent in some parts of Punjab and Haryana states (Kang and Chahal, 2000; Sharma et al., 2001).

II. Materials and methods

The studies were conducted during rabi season in department of plant pathology and Student Instructional Farm of Narendra Deva University of Agriculture and Technology, Kumarganj, Faizabad. Six fungicides belonging to different groups and one bio-pesticide were bio-assayed against the pathogen under laboratory condition to find out their relative efficacy in inhibiting the growth of the pathogen by the “Food poison technique”, (Schmitz, 1930). Requisite quantity of each fungicide and bio-pesticide was incorporated in 2.0 per cent Potato Dextrose Agar medium, thoroughly mixed by shaking, prior to pouring in Petri dishes. The medium was allowed to solidify and then inoculated with 5.0 mm disc of inoculum from 3 days old culture of the pathogen. The inoculum was placed in the centre of each Petri dish. The fungal discs were reversed so that the pathogen could come in direct contact with the medium. Three replications were kept for each treatment. The efficacy of various fungicides and bio-pesticide were observed by measuring the radial growth of the fungal colony in mm. The per cent inhibition over control was calculated by following formula given by Bliss (1934).

$$\text{Per cent inhibition over control} = \frac{C-T}{C} \times 100$$

Where, C = Growth of fungus in control.
T = Growth of fungus in treatment

For evaluating the efficacy of various, fungicides and bio-pesticide as spray against the disease, the experiment was conducted in the field. The mustard cv. Varuna was sown in plots during both the years. Crop was inoculated artificially with 5 mm discs of inoculums from 3 days old culture of the pathogen. The first spraying of, fungicides and bio-pesticide was given as soon as the disease was noticed and subsequent two sprayings were followed at 10 days interval. The controlled plots were sprayed with water only. The observations on disease severity were recorded after 10 days of the last spraying and yield data were also recorded separately in each treatment after the harvest of the crop.

III. Results and discussions

Results presented in **Table-1** indicated that all the fungicides and bio-pesticide were significantly superior over control in inhibiting the growth of the fungal colony. Vitavax (0.1 %), Bavistin (0.1%) and Benomyl (0.1 %) were most effective as they completely inhibited (100%) the growth of pathogen. Whereas, bio-pesticide Neemark (0.5%) was noted as least effective and inhibited the growth of fungus only up to 44.44 %. Ridomyl (0.2 %) and Indofil M-45 (0.25 %) were found to be the next best in superiority in inhibiting the growth of the pathogen **Fig 1, 2 and 3**. The results presented in **Table-2** indicated that all the fungicides and bio-pesticide proved significantly effective in controlling the disease over control. The spraying of Vitavax (0.1%) at an interval of 10 days was found more effective under field condition as is evident from its corresponding yield in both the years. The next best fungicide Bavistin (0.1%) was found statistically at par with Vitavax (0.1%) in respect of reduction in disease severity and enhancement of seed yield. However, Neemark (0.5%) proved to be the least effective under field condition also. Benomyl @ 0.1% ranked 3rd in effectiveness which was significantly superior with other tested fungicides such as Indofil M-45, Ridomyl and Chlorothalonil in respect of decreasing the disease severity but was at par in respect of seed yield. On the basis of average yield of both the years and additional cost involved in each treatment, the benefit-cost ratio was calculated. It was recorded that Bavistin @ 0.1% was found most economical with benefit-cost ratio of 1:2.59 followed by Indofil M-45 @ 0.25% (1:1.57) and Vitavax @ 0.1% (1:1.38), respectively. Vitavax, was found most effective in controlling the disease and increasing the seed yield in comparison to others, but due to its high cost of product it ranked third after Dithane M-45 which was less effective. Similarly, Benomyl which ranked third in effectiveness was also lower in benefit-cost ratio and ranked fourth (1:0.90) due to its high product price. Among the chemicals, the benefit- cost ratio of Ridomyl was recorded lowest (**Table-3**), though it was at par with Indofil M-45 (0.25) in effectiveness. Neemark was recorded overall least effective with lowest benefit-cost ratio (1:0.21).

Concurrent with present findings i.e. much effectiveness of Vitavax (0.1%) against the inhibition of mycelia growth of *Sclerotinia sclerotiorum* was also reported by earlier workers time to time (Singh and Gangopadhyay,1984, Abdou et al. ,1982 and Roy and Saikia,1976) under laboratory conditions. It was observed that the performance of systemic fungicides was better in controlling the disease in comparison to non-systemic fungicides. Singh et al. (1994) and Singh et al. (2003) have also reported the effectiveness of these fungicides. Sharma, A.K. (1987) reported that Bavistin, Benomyl, Chlorothalonil and Indofil M-45 were effective against *Sclerotinia sclerotiorum* on pea crop.

Table 1. Efficacy of fungicides / bio-pesticide against the growth of *Sclerotinia sclerotiorum* under in-vitro

Fungicides/ bio-pesticide(%)	Average diameter of colony (mm)	Per cent inhibition over control
Bavistin (0.1)	0.0	100.0
Vitavax (0.1)	0.0	100.0
Chlorothalonil(0.2)	46.20	48.67
Indofil M-45 (0.25)	36.83	59.08
Ridomyl (0.2)	28.67	68.14
Benomyl (0.1)	0.0	100.0
Neemark (0.5)	50.00	44.44
Control	90.00	0.00
CD at 5%	3.56	

Table 2. Efficacy of fungicides / bio-pesticide against the *Sclerotinia sclerotiorum* under in vivo.

Fungicides/ biopesticide (%)	1 st year		2 nd year		Av. yield of both years
	Disease severity (%)	Yield (q/ha)	Disease severity (%)	Yield (q/ha)	
Bavistin (0.1)	8.00	10.83	9.33	10.33	10.58
Vitavax (0.1)	5.33	12.33	6.66	11.66	11.99
Chlorothalonil (0.2)	18.67	8.33	17.33	6.66	7.49
Indofil M-45 (0.25)	21.33	7.5	22.67	7.50	7.50
Ridomyl (0.2)	20.00	7.08	22.67	8.66	7.87
Benomyl (0.1)	12.00	9.58	13.33	9.41	9.49
Neemark (0.5)	24.00	5.83	28.67	5.83	5.83
Control	39.50	5.00	38.33	5.20	5.10
CD at 5%	5.10	2.65	4.80	2.66	2.85

Table.3 Economics of fungicides/ bio-pesticide.

Particulars	Control	Bavistin (0.1%)	Vitavax (0.1%)	Indofil M-45 (0.25%)	Chlorothalonil (0.2%)	Benomyl (0.1%)	Ridomyl (0.2%)	Neemark (0.5%)
Amount of fungicides /bio-pesticide kg/ha	-	3	3	6	6	3	6	15
Cost of fungicides (Rs.)	-	2210	6000	1488	1500	6000	8700	3600
Sprayer charges (15Rs/days)	-	45	45	45	45	45	45	45
Labour charges (Rs. 58/day)	-	522	522	522	522	522	522	522
Total cost (Rs.)	-	2777	6567	2055	2067	6567	9267	4167
Av. Yield (q/ha)	5.20	10.33	11.66	7.50	6.60	9.41	8.66	5.83
Additional yield over control	-	5.13	6.46	2.30	1.40	4.21	3.46	0.63
Net income Rs. / ha	7280	7182	9044	3220	1960	5894	4844	882
C:B ratio	-	1:2.59	1:1.38	1:1.57	1:0.95	1:0.90	1:0.52	1:0.21

Sale price of mustard: Rs. 1400/q

References:

- [1]. Abdou, Y.A.M.; Fl. Sharkaway, T.A.; Osman, A.R. and Ragab, M.M. (1982). In vitro test for control of *Sclerotinia sclerotiorum*. Egyptian J. Phytopathology. 14: 87-92.
- [2]. Anonymous, 2011. Statistical Year Book India. Directorate of Economics and Statistics, Ministry of Agriculture, pp: 117.
- [3]. Bliss, C.I. (1934). The method of probits. Science. 79: 38.
- [4]. Chauhan, L.S., Singh, Jyoti and D.R.Chandra, 1992: In: proc. Natl. Symp. On management Microbes in Service of Mankind". Nov. 19-21,1992 at Univ. Allahabad, India. 65-66 pp (Abstr.).
- [5]. Chandel, S.R.S. (1993). A hand book of Agriculture statistics. Achal Prakashan Mandir, Kanpur, pp 558 .
- [6]. Ghasolia, R.P., Shivpuri, Asha and A.K. Bhargava, 2004: *Sclerotinia* rot of Indian mustard (*Brassica juncea*) in Rajasthan. Indian Phytopath.57: 76-79.
- [7]. Kang, I.S. and S.S. Chahal, 2000: Prevalence and incidence of white rot of mustard incited by *Sclerotinia sclerotiorum* in Punjab. Plant Dis. Res. 15: 232-233.
- [8]. Roy, A.K. and Saikia, V.N. (1976). White blight of mustard and its control. Indian J. of Agric.Sci. 46 (6): 274-277.
- [9]. Schmitz, H. (1930). A suggested toximetric method for wood preservative Indus. Engin Chem., Anlyt. Edi II. 361-363. Rev. Appl. Mycol. 10: 21-1931.
- [10]. Sharma, S., J.L. Yadav and G.R. Sharma, 2001: Effect of various agronomic practices on the incidence of white rot of Indian mustard caused by *Sclerotinia sclerotiorum* J.Mycol. Pl. Pathol 31: 83-84.
- [11]. Singh, N. and Grangopadhyay, S. (1984). Control of white rot seed cauliflower. Pesticides. 18: 23- 24.
- [12]. USDA., 2010. United States Department of Agriculture-Rapeseed are, yield and production Table No.15 [http:// www. Fas.usda.gov/psd online/ psd report. Asps](http://www.Fas.usda.gov/psd online/ psd report. Asps) (created on july 3, 2010).

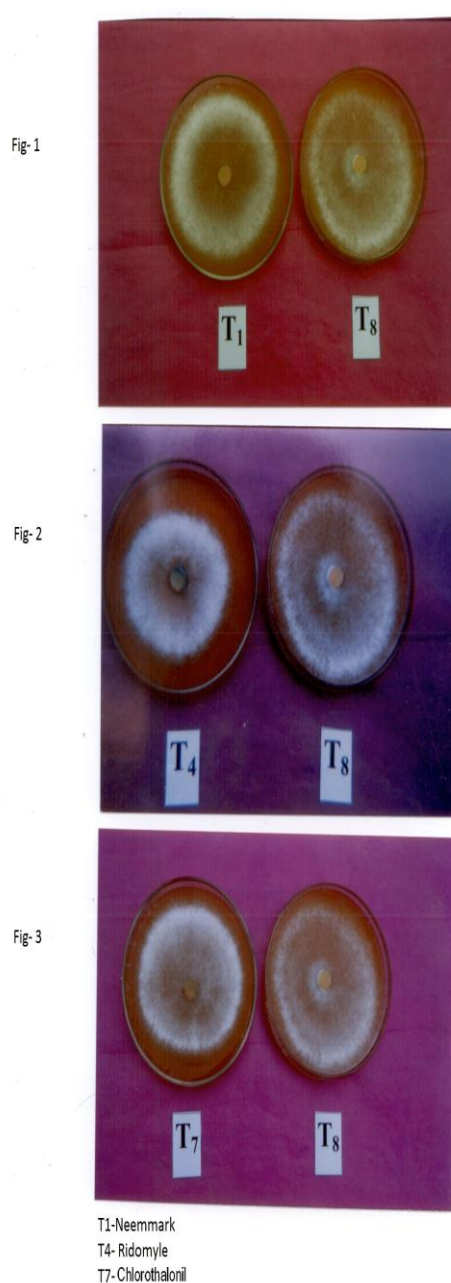


Fig.1,2,3:Effect of different fungicides against the pathogen (*Sclerotinia sclerotiorum*) under *in vitro*.