

Efficacy of bio-control agents with chemical fungicides against *Cercospora* leaf spot of Okra (*Abelmoschus esculentus* L.) Moench

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ABSTRACT

Okra (*Abelmoschus esculentus* L.) is one of the foremost vegetable crop grown during kharif as well as summer seasons. *Cercospora* leaf spot incited by *Cercospora* spp. is one of the emerging disease in all regions wherever okra is grown. *C. abelmoschica* causes sooty black, angular spots and cause severe defoliation common during humid seasons. An experiment was conducted to evaluate the efficacy of bioagents and chemicals viz., T₀ - Untreated control, T₁ Mancozeb (1%) + *Trichoderma* (4%), T₂ - Mancozeb (1%) + *Pseudomonas* (4%), T₃ Mancozeb (1%) + *Bacillus subtilis* (4%), T₄ - Mancozeb (1%) + *Trichoderma* (2%) + *Pseudomonas* (2%), T₅ - Mancozeb (1%) + *Pseudomonas* (2%) + *Bacillus subtilis* (2%), T₆ Mancozeb (1%) + *Bacillus subtilis* (2%) + *Trichoderma* (2%), T₇ - Mancozeb (1%) against *Cercospora* leaf spot of okra. Studies revealed that minimum disease intensity, Maximum plant height, maximum no. of branches per plant and Maximum no. of fruits was observed in T₄ - Mancozeb (1%) + *Trichoderma* (2%) + *Pseudomonas* (2%) and is hereby considered as the best treatment.

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Keywords: Mancozeb, *Trichoderma*, *Pseudomonas*, *Bacillus*.

1. INTRODUCTION

Okra (*Abelmoschus esculentus* L.) Moench is one of the most widely known species of the family Malvaceae and an economically important vegetable crop grown in tropical climate of temperature range between 25^o to 35^o c. The name "Okra" derives from one of the Nigero-Congolese group of languages. "Okra" originated in Ethiopia and was then propagated in North Africa. In India okra is grown in sub-tropical areas and it is commonly known as Bhendi.

Some studies are being developed targeting okra extract as a remedy to manage diabetes. Its ripe seeds are roasted, ground and used as a substitute for coffee in some countries. Mature pods and stems containing crude fibre are used in the paper industry. Okra seeds are a potential source of oil, which consists of linoleic acid up to 47.4% and polyunsaturated fatty acid essential for human nutrition. (Singh et al., 2014).

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Okra contains Potassium, Sodium, Magnesium and Calcium as principal elements in pods, which contains 17% seeds. Presence of Iron, Zinc, Manganese and Nickel also has been reported (Moyin-Jesu, 2007). Fresh pods are low in calories (20/100g), practically no fat, rich in fiber, and with several valuable nutrients. Okra seed is mainly composed of oligomeric catechins (2.5 mg g⁻¹ of seeds), while the mesocarp is mainly composed of hydroxycinnamic (0.2 mg g⁻¹) and quercetin derivatives (0.3 mg g⁻¹). Pods are rich in phenolic compounds with important biological properties like quercetin derivatives, catechin oligomers and hydroxycinnamic derivatives (Arapitsas, 2008).

Okra plant also contains many medicinal properties with it. But before using, it is very necessary to seek advice from a professional. The mucilage can be used as plasma replacement, helpful in washing away toxic substances from the body and have strongly demulcent action (Gemedé et al., 2015). In the treatment of syphilis infusion of root is used. In Nepal the juice of root is used in the boils, wound and cuts. It is used in the medication of catarrhal infections, dysuria and gonorrhoea. Other than this fibre present in okra has property of controlling blood sugar level in blood. Okra has nutrient that insure proper functioning of intestine. It is also effective in ulcer and joint healthiness. Due to its alkaline nature, it also guards the mucous membrane in the digestive system. Useful in curing of pulmonary inflammation, bowel irritation and sore throat (Kumar et al., 2013). Its fruit can be also used for the control of goitre due to high iodine content in it.

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Diseases play a vital role in yield losses of the crop. Among them, fungi are one of the most important and prevalent pathogens which attack the crops from seedling to harvesting stage. Some of the fungal diseases that attack are *Cercospora* leaf spot (*Cercospora abelmoschi*), damping-off (*Pythium sp.* and *Rhizoctonia sp.*), powdery mildew (*Oidium sp.*), southern blight (*Sclerotium rolfsii*), verticillium wilt (*Verticillium albo-atrum*) and *alternaria leaf spot* (Raid and Palmateer, 2006).

Among the fungal diseases *Cercospora* leaf spot of bhendi incited by *Cercospora* is one of the most economically important in all regions where ever bhendi is grown. In India, two species of *Cercospora* produce leaf spots on bhendi. *C. malayensis* causes brown, irregular spots and *C. abelmoschi* causes sooty black, angular spots. Both the leaf spots cause severe defoliation and are common during humid seasons. Now a days, this disease incited by *C. abelmoschi* becomes more severe in southern transition zone of Karnataka. Initially the disease symptoms observed on the lower surface of the leaves as indistinct spots in the form of olivaceous specks. Later on, light brown to grey mouldy growth of the fungus covered the entire lower surface. The infected leaves ultimately dry and defoliate. The disease progress upward from lower leaves and infects stem and fruits and produces similar symptoms. (Naik et al., 2017).

Cercospora produce a perylene quinone toxin called cercosporin which is non-selective, affecting bacteria and fungi unless these produce protective antioxidants such as carotenoids. Morphology of the pathogen of genus *Cercospora* was first described by Frensius (1863). Etymologically the generic name means a fungus has obclavate (tail shaped) spores. Sporulation occurs at temperature range 8-24°C, where mature spores sporulate after 14 to 24 hours.

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For the management of *Cercospora* leaf spot of okra from many years, many have been relied on chemicals and this resulted in many undesirable problems. Now a day's tremendous use of chemicals in agriculture has resulted in growing concern of both public health and environment hazards thus, emphasis is now on judicious use of bio-agents, botanicals and organics for management of the plant diseases which is less costly, nontoxic and doesn't affect public health and

environment. Fungicides are also effective in managing this disease as such their use in the management strategy cannot be ruled out but their indiscriminate use should be avoided. There is need to incorporate alternative control components that are effective in field. Considering the above-mentioned facts, a study was conducted, entitled, "**Efficacy of bio control agents with chemical fungicides against Cercospora leaf spot of Okra (*Abelmoschus esculentus* L.) Moench**" with the following objectives :-

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1. To evaluate the effect of **bioagents** bio-agents on *Cercospora* disease intensity in okra.

2. MATERIAL AND METHODS

The experiment was conducted at the research plot of the Department of Plant Pathology and Central Research Field, Sam Higginbottom University of Agriculture Technology and Sciences, Prayagraj during the *Kharif* season 2022. The selected site was uniform, cultivable with typical sandy loam soil having good drainage.

Table 1. The treatment details.

S.No	Treatments	Treatment Details
1.	T0	Control
2.	T1	Mancozeb (1%) + <i>Trichoderma harzianum</i> (4%)
3.	T2	Mancozeb (1%) + <i>Pseudomonas fluorescens</i> (4%)
4.	T3	Mancozeb (1%) + <i>Bacillus subtilis</i> (4%)
5.	T4	Mancozeb (1%) + <i>Trichoderma harzianum</i> (2%) + <i>Pseudomonas fluorescens</i> (2%)
6.	T5	Mancozeb (1%) + <i>Pseudomonas fluorescens</i> (2%) + <i>Bacillus subtilis</i> (2%)
7.	T6	Mancozeb (1%) + <i>Bacillus subtilis</i> (2%) + <i>Trichoderma harzianum</i> (2%)
8.	T7	Mancozeb (1%)

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Disease severity scale of *Cercospora* leaf spot

Disease intensity was recorded as grades in five randomly selected plants in each plot at different times that is before spraying, 15 days after the first spray and 15 days after the second spray as per the scale of **Farrag (2011)** which is given below.

Table 2. Disease intensity and description.

Disease rating/grade	Description
0	No disease
1	Noticeable spotting with some defoliation (<25%)
3	Spotting heavy with significant defoliation (<50%)
5	Very heavy leaf spotting with severe defoliation (<75%)
7	Numerous spots on few remaining leaves and very heavy defoliation (<90%)
9	Very few remaining leaves covered with spots and nearly complete defoliation (<95%)

3.5 Disease intensity (%)

Percentage of Disease intensity will be recorded at 60, 75 and 90 days after incidence of *Cercospora* leaf spot. Percentage of Disease intensity will be calculated in accordance with following formula. The disease will be visually assessed in all the plots at weekly interval from first appearance of disease for each treatment. For each plot the number of infected okra plants will be counted and expressed as a percentage of the total number of okra plants in that plot. The mean percentage disease incidence for each treatment will be obtained from the three replications. The data will be further statistically analyzed.

Disease intensity (%) formula was given by **Wheeler (1969)**. It is calculated by using the following formula:

$$\text{Disease intensity (\%)} = \frac{\text{Sum of all disease ratings}}{\text{Total no. of ratings} \times \text{Maximum disease groups}} \times 100$$

Statistical analysis

The data obtained from the field experiment were statistically analyzed by following the standard procedures (**Panse and Sukhatme, 1989**). The percentage values were converted to arcsine values wherever required.

Analysis of variance:

The analysis of variance was worked out to test the significance of F and t-tests. It was carried out as followed according to procedure of RBD analysis for each character. The total variance and degree of freedom were partitioned into three components viz., replications, treatments and error. Analysis of variance was done under the fixed effect model given below: Let us suppose that there are 'k' treatments applied to 'r' number of replications. These can be represented by the symbols as follows:

Analysis of Variance was done under the fixed effect model given below

Let us suppose that there are 'k' treatments applied to 'r' number of replications. These can be represented by the symbols as follows

Table 3. Analysis of Variance

Source of Variation	d.f.	S.S.	M.S.S.	F(cal)	F(tab) at 5%	Result
Dueto Replication	r-1	R.S.S	M.S.S.R =RSS r-1	M.S.S.R M.E.S.S	F (r-1)	S/NS
Dueto Treatments	t-1	T.S.S.	T.S.S t-1 = M.T.S.R	M.T.S.S M.E.S.S	F(r-1)(t-1)	N/NS
Dueto error	(r-1) (t-1)	E.S.S	S.S.E. (r-1)(t-1)	-	f(t-1),(r-1),(t-1)	-
Total	(rt-1)	T.S.S.	-	-	-	-



Fig 1. Overview of Spraying



Fig 2. Overview of Disease Infested Leaves

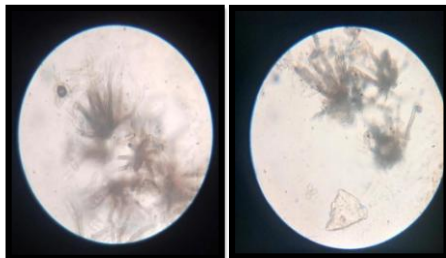


Fig 3. OVERVIEW OF MICROSCOPIC VIEW OF *Cercospora* sp.

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Table 4. Effect of treatments on Disease Intensity of *Cercospora* leaf spot of okra at 60,75 and 90 DAS

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Tr.no	Treatment	DISEASE INTENSITY		
		60DAS	75 DAS	90DAS
T0	Control	29.183 ^a	36.290 ^a	41.18 ^a
T1	Mancozeb (1%) + <i>Trichoderma harzianum</i> (4%)	21.033 ^d	27.403 ^e	34.44 ^c
T2	Mancozeb (1%) + <i>Pseudomonas fluorescens</i> (4%)	18.810 ^e	25.920 ^f	32.33 ^d
T3	Mancozeb (1%) + <i>Bacillus subtilis</i> (4%)	23.553 ^c	30.810 ^c	36.47 ^b
T4	Mancozeb (1%) + <i>Trichoderma harzianum</i> (2%) + <i>Pseudomonas fluorescens</i> (2%)	14.367 ^g	21.920 ^h	26.66 ^f
T5	Mancozeb (1%) + <i>Pseudomonas fluorescens</i> (2%) + <i>Bacillus subtilis</i> (2%)	15.703 ^f	23.847 ^g	29.33 ^e
T6	Mancozeb (1%) + <i>Bacillus subtilis</i> (2%) + <i>Trichoderma harzianum</i> (2%)	23.847 ^c	28.887 ^d	34.51 ^c
T7	Mancozeb (1%)	27.183 ^b	33.480 ^b	37.33 ^b
	C.D (5%)	1.009	1.044	1.867

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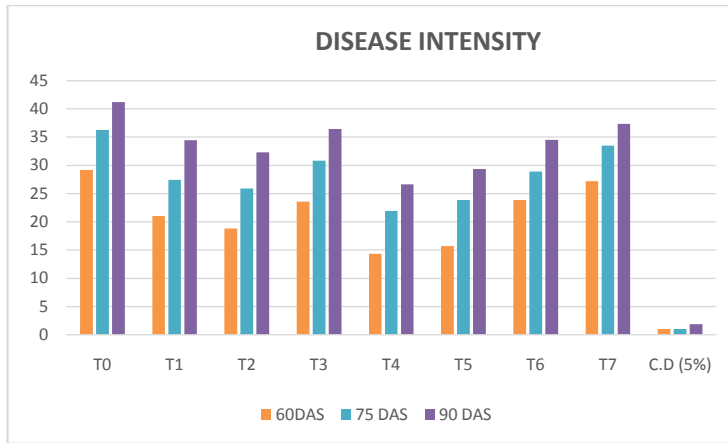


Fig 4.DISEASE INTENSITY

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4.1 Disease Intensity:

4.1.1 Disease Intensity at 60 DAS

The data presented in table 4 and depicted in figure 4 reveals that maximum Disease intensity ofokraat60DASwasrecordedinT4- Mancozeb (1%) + *Trichoderma*(2%) + *Pseudomonas*(2%) (14.36)followedby T5 - Mancozeb (1%) + *Pseudomonas*(2%) + *Bacillus subtilis*(2%) (15.70) and T2 - Mancozeb (1%) + *Pseudomonas*(4%) (18.81) followed by T₁ Mancozeb (1%) + *Trichoderma*(4%)(21.03) , T₆ Mancozeb (1%) + *Bacillus subtilis*(2%) + *Trichoderma*(2%)(23.553), T₃ Mancozeb (1%) + *Bacillus subtilis*(4%) (23.847)ascomparedtoT7-Mancozeb (1%) (27.18)andT0–untreatedcontrol-(29.18).Allthetreatmentsweresignificantover untreated control. Among the treatments (T₇ and T₄) were statistically non-significant to each other

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4.1.2 Disease Intensity at 75 DAS

The data presented in table 4 and depicted in figure 4 reveals that maximum Disease Intensity ofokraat75DASwasrecordedinT4- Mancozeb (1%) + *Trichoderma*(2%) + *Pseudomonas*(2%) (21.92)followedby T5 - Mancozeb (1%) + *Pseudomonas*(2%) + *Bacillus subtilis*(2%) (23.84) and T2 - Mancozeb (1%) + *Pseudomonas*(4%) (25.92) followed by T₁ Mancozeb (1%) + *Trichoderma*(4%)(27.40) , T₆ Mancozeb (1%) + *Bacillus subtilis*(2%) + *Trichoderma*(2%)(28.88), T₃ Mancozeb (1%) + *Bacillus subtilis*(4%) (30.81)ascomparedtoT7-Mancozeb (1%) (33.48)andT0–untreatedcontrol-(36.29).Allthetreatmentsweresignificantover untreated control.

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4.1.3 Disease Intensity at 90 DAS

The data presented in table 4 and depicted in figure 4 reveals that maximum Disease Intensity ofokraat90DASwasrecordedinT4- Mancozeb (1%) + *Trichoderma*(2%) + *Pseudomonas*(2%) (26.66)followedby T5 - Mancozeb (1%) + *Pseudomonas*(2%) + *Bacillus subtilis*(2%) (29.33) and T2 - Mancozeb (1%) + *Pseudomonas*(4%) (32.33) followed by T₁ Mancozeb (1%) + *Trichoderma*(4%)(34.44) , T₆ Mancozeb (1%) + *Bacillus subtilis*(2%) +

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Trichoderma(2%)(34.51), T₃ Mancozeb (1%) + *Bacillus subtilis*(4%) (36.47) as compared to T₇-Mancozeb (1%) (37.33) and T₀-untreated control-(41.18). All the treatments were significant over untreated control. Among the treatments (T₈ and T₄), (T₇ and T₂) were statistically non significant to each other

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From this present study entitled "**Efficacy of bio control agents with chemical fungicides against *Cercospora* leaf spot of Okra (*Abelmoschus esculentus* L.) Moench**" based on the observations it can be concluded that the efficacy of combining readily available and ecologically safe bioagents with synthetic safe mancozeb fungicide for the management of *Cercospora* leaf spot of okra .

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From the critical analysis of the present findings, it can be concluded that after the application of all the treatments with three replications, T₄- Mancozeb (1%) + *Trichoderma*(2%) + *Pseudomonas*(2%) is the best treatment as it showed The **Disease Intensity of okra** at 60,75 and 90 DAS which was significantly increased by the use of Mancozeb (1%) + *Trichoderma*(4%) + *Pseudomonas*(4%) under Prayagraj Agro climatic conditions . Based on analysis T₄- Mancozeb (1%) + *Trichoderma*(2%) + *Pseudomonas*(2%) is recommended to control the *cercospora* leaf spot disease in Okra. The present findings were limited to one crop season *kharif* under the climatic conditions of Prayagraj , U.P. , therefore substantiate the present result more trails are required for further recommendations .

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Reference

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Arapitsas (2008). Identification and quantification of polyphenolic compounds from okra seeds and skins. *Food Chemistry*, 110:1041-1045.

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Faisal, M. and Tiwari, S. (2015). Comparison of bio-agents and botanicals with fungicides against tikka and anthracnose diseases of groundnut (*Arachis hypogaea* L.). *International Journal of Plant Protection*, 8(1): 45-48.

Gemedé, H. F., Ratta, N., Haki, G. D. and Beyene, A. Z. W. F. (2014). Nutritional Quality and Health Benefits of Okra (*Abelmoschus esculentus*): A review. *Global Journal of Medical Research*, 14(5):2249-4618.

Kumar, S., Jaiswal, S., Lal, A.A., Kumar, A. and Verma, A. (2017). Influence of natural products and bio-fungicide against tikka disease of groundnut caused by *Cercospora* spp. *The Pharma Innovation Journal*, 6 (3):213-216.

- Kumar, R., Meena, A. K., Kumar, V. and Singh, J. (2022)** Isolation of *Cercospora canescens* and management of *Cercospora* Leaf Spot (*Cercospora canescens*) of moth bean through botanicals. *Journal of Experimental Agriculture International*, **44(5):57-63**.
- Naik, G. R. and Jayalakshmi, K. (2017)**. Evaluation of fungicides against leaf spot of bhendi incited by *Cercospora abelmoschi* under field conditions. *International Journal of Chemical Studies*, **5(5):1210-1212**.
- Singh, P., Chauhan, V., Tiwari, B. K., Chauhan, S. S., Simon, S., Bilal, S. and Abidi, A. A. (2014)**. An overview on okra (*Abelmoschus esculentus*) and its importance as a nutritive vegetable in the world. *International Journal of Pharmacy and Biological Sciences*, **4(2)227-233**.

DEFINITIONS, ACRONYMS, ABBREVIATIONS

Here is the Definitions section. This is an optional section.

Term: Definition for the term