Effect of compost along with bio agents on root knot nematode (*Meloidogyne spp.*) in okra (*Abelmoschus esculentus* L., Walp)

ABSTRACT

Okra [Abelmoschus esculentus L., Moench] a native of South-Africa and commonly known as 'Bhindi' is an annual malvaceous vegetable crop, especially grown in tropical and subtropical climates. It is also called "perfect villager's vegetable". Root-knot nematodes are considered among the top five major plant pathogens and the first among the ten most important genera of plant parasitic nematodes in the world. Amongst nematodes, root-knot nematode (Meloidogyne incognita and Meloidogyne javanica) causes severe damage to okra. Trichoderma viride, Pseudomonas fluorescene and Rhizobium were tested under field conditions during Rabi 2020-2021 for their efficacy against the Root knot nematode and growth & nematode population. Among the treatments the maximum plant height(cm) at 60 DAS was recorded in T₆-Vermicompost+ spent mushroom compost+ *Trichoderma viride*+ Pseudomonas fluorescence + Rhizobium (51.50 cm) followed by T₅ - Vermicompost +spent mushroom compost+ Trichoderma viride + Rhizobium (44.45 cm) as compared to untreated check control T_0 – (25.13 cm). The maximum Root weight was recorded in T_3 – Vermicompost +spent mushroom compost + Rhizobium (9.90 gm) followed by T₁ -Vermicompost+ spent mushroom compost+ Trichoderma viride(9.65gm) as compared to untreated check control T₀-(10.25gm). The maximum Root length was recorded in T₆ vermicompost + spent mushroom compost+ Trichoderma viride + Pseudomonas fluorescence + Rhizobium (13.70gm) followed by T₄ -vermicompost + spent mushroom compost + Trichoderma viride + Pseudomonas fluorescence (13.35gm) as compared to untreated check control T_0 – (10.12). The minimum nematode intensity (%) at 60 DAS was recorded in T₆ - Vermicompost+ spent mushroom compost + Trichoderma viride + pseudomonas fluorescence + Rhizobium (17), followed by T₄ - Vermicompost +spent mushroom compost + Trichoderma viride + Pseudomonas fluorescence (25.50) as compared to untreated check control $T_0(57)$.

Keywords: Meloidogyne incognita, pseudomonas fluorescence, Rhizobium, Root-knot Nematode of Okra, Trichoderma viride.

1. INTRODUCTION

Okra [Abelmoschus esculentus L. Moench] a native of South- Africa and commonly known as 'Bhindi' is an annual malvaceous vegetable crop, especially grown in tropical and subtropical climates [1]. India ranks first in the world with 3.5 million tonnes

(70% of the total world production) of okra produced from over 0.35-million-hectare land[2]. In India, total grown in an area of 511 hectare having total production of 6219 million tons with the productivity of 12.0 million tons per hectare[3]. Dry seed contains 13-22% edible oi; and 2024% protein [4]. The production of okra also suffers from several pests (fruit borer, white flies, jassids, thrips, mites etc.) and pathogens (yellow vein mosaic, powdery mildew, root rot etc.) including nematodes [5]. Plant parasitic nematodes viz. *Meloidogyne incognita, Meloidogyne javanica, Rotylenchulus* spp., *Tylenchorhynchus* spp., *Hoplolaimus* spp., *Aphelenhusavenae, Nothotylenchus* spp., *Helicotylenchus* spp., *Hemicriconemoides* spp., *Longidorus* spp., *Paralongidorus* spp., *Trichodorus* spp., *Paratrichodorus* spp., *Ditylenchus* spp., *Tylenchus* spp., *Rotylenchulus* spp., and Xiphinema spp, were found associated with okra in India [6]. Root-knot nematode (*Meloidogyne incognita* and *Meloidogyne javanica*) causes severe damage to okra [7]. Severe attack of root-knot disease caused by *Meloidogyne* spp[8]. On okra and yield losses up to 27% [9]. Rootknot nematodes (RKN) are sedentary endo-parasite and is among the most damaging agricultural pests, attacking a wide range of crops [10].

2. MATERIALS AND METHODS

Keeping in view, the experiment was conducted in nematode infested soil at the courtyard of Department of Plant Pathology, SHUATS, Prayagraj, Uttar Pradesh during *Rabi* season in the year 2020-2021. The soil sample was collected from the infested field and processed in laboratory by following cobbs decanting and sieving technique followed by modified Baermann funnel technique to estimate the nematode population. Before laying out the experiment it was assured that the experimental field possess 2 larvae/gm of soil.

The selected field was dug up and the soil become pulverized and then whole location was divided into sub-plots and specified in randomized block design with six treatments viz., vermi compost @ 8 t/ha, spent mushroom compost @ 8 t/ha was used as basal application. These were incorporated into the soil by forming specific ridges according to the crop spacing and covered by thin layer of soil. The field was irrigated for fifteen days at regular intervals to enhance decomposition process. After fifteen days, seed treatment was done with Trichoderma @ 10 gm/kg, Pseudomonas @ 10 gm/kg, Rhizobium@ 5 gm/kg of seed where, vermi compost and spent mushroom compost was incorporated initially. Each treatment was replicated four times with plot size of $2.5 \times 1 \text{ m}^2$ each and local variety seed was sown with a spacing of $45 \times 30 \text{ cm}$. Root knots in the root system and plant growth parameters of okra was recorded at 30, 45, 60 days after sowing of the crop. The records have been subjected to th3e statistical analysis.

At 60 days after sowing the root knots in the roots of okra are identified. The galled roots were removed and washed thoroughly with sterile water. a gall is placed on the sterile slide using sterile forceps and teared using a sterile needle and examined under microscope. Eggs and female Meloidogyne were identified when observed under microscope. Mature females are swollen, melon like with elongate neck at anterior end, forms perineal patterns, short stylet with well-developed basal knobs, eggs laid in gelatinous matrix outside the body, tail absent. Males are vermiform, 1.5-2.0 mm long, basal knobs; oesophageal glands overlap intestine ventrally; tail elongate conoid with pointed tip.

3. RESULTS

The result presented in table 1 revealed that all the treatments were statistically significant and decreased the number of root knots in the roots of okra as compared to control. Among the bio agents used, the treatment T6- Vermicompost + SMC+ Trichoderma spp.+ Pseudomonas spp.+ Rhizobium (17) significantly decreased the root knots in the root system (57) in okra when compared to other bio agents. The treatments (T_6 , T_4), (T_2 , T_5) and (T_1 , T_3) are not significantly differ from each other.

Table 1: Effect of compost along with bio-agents on the number of root knot nematodes on the roots of okra at 60 DAS.

Tr no.	Treatments	Number of root knots
T_0	Control+ Vermicompost+ SMC	57
T_1	Vermicompost+ SMC+ <i>Trichoderma</i> spp. 50	
T_2	Vermicompost + SMC+ <i>Pseudomonas</i> spp 31	
T_3	Vermicompost+ SMC+ Rhizobium	51
T_4	Vermicompost + SMC+ <i>Trichoderma</i> spp.+ <i>Pseudomonas</i> spp.	26
T_5	Vermicompost + SMC+ <i>Trichoderma</i> spp.+ <i>Rhizobium</i>	34
T ₆	Vermicompost+ SMC+ <i>Trichoderma</i> spp.+ <i>Pseudomonas</i> spp.+ <i>Rhizobium</i>	17
	F- test	S
	S. E. (d) ±	1.91
	C. D. (5%)	3.93



Plate 1: Root knots in the root system of okra at 60 days after sowing.

The result presented in table 2 revealed that all the treatments were statistically significant and increased the plant growth parameters of okra. Among the Bio agents, the treatment T_6 -Vermicompost + SMC + Trichoderma spp. + Pseudomonas spp. + Rhizobium (19.01cm) significantly increased the plant height of okra. Among the Bio agents the treatments (T_6, T_4), (T_2, T_5) and (T_1, T_5) found non-significant to each other. The treatment T_6 -Vermicompost + SMC + Trichoderma spp. + Pseudomonas spp. + Rhizobium (37.20 cm) significantly increased the plant height of okra. Among the Bio agents the treatment (T_1, T_3) found non-significant to each other. The treatment T_6 -Vermicompost + SMC + Trichoderma spp. +

Table 2: Effect of bio agents along with bio agents on plant growth parameters of okra:

Treatments	Plant height (cm)	Root length (cm)	Root weight (gm)
T_0	25.13	10.12	10.25
T_1	35.27	11.50	9.65
T_2	41.85	12.62	8.93
T_3	32.35	11	9.90
T_4	44.45	13.35	8.43
T_5	38.35	12.20	9.13
T_6	51.50	13.70	8.25
F-test	S	S	S
S. E (d) \pm	0.75	0.29	0.40
C.D. (5%)	1.58	0.60	0.84

The result presented in table 2 revealed that all the treatments were statistically significant and decreased the root weight of cowpea as compare to control. The treatments T6-Vermicompost + SMC + Trichoderma spp.+ Pseudomonas spp.+ Rhizobium (8.25 gm) significantly decreased the root weight due to the less number of root knots in root system of okra The treatments (T_3,T_1) , (T_3,T_1,T_5) , (T_1,T_5,T_2) , (T_5,T_2,T_4) and (T_2,T_4,T_6) found non-significant to each other. Highest root weight was recorded in control T_0 - (10.25 gm) due to higher number of root knots in the root system of cowpea. The present investigation indicates that Vermicompost +

SMC + *Trichoderma* spp.+ *Pseudomonas* spp.+ *Rhizobium* into the soil used as an effective treatment for root knot nematodes and to develop eco-friendly strategy for the management of root knot nematodes of okra.

4. DISCUSSION

Probable reason for such finding may be due to the inhibitory impact of bio agents because of the nemato-toxic compounds present in the bio agents which help to reduce the severity of the nematodes in the soil and plants [11]. Application of Trichoderma had a significant effect on root knot nematode population and found toxic to *Meloidogyne* spp. due to Myco-parasitism mechanisms involved in the antagonisms of *Trichoderma*.

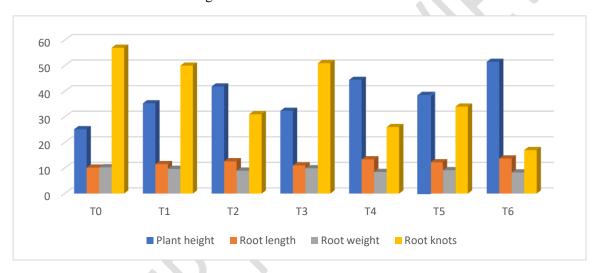


Fig 1: Effect of bio agents along with bio agents on plant growth parameters of okra.

5. CONCLUSION

In the present study on the basis of observation, it was found that for managing the root knots in the root system of okra, Vermicompost + SMC+ *Trichoderma* spp.+ *Pseudomonas* spp.+ *Rhizobium* (17) was significant in comparison to control (57). Hence from present study it can be concluded that Vermicompost + SMC+ *Trichoderma* spp.+ *Pseudomonas* spp.+ *Rhizobium* can be used effectively to reduce the root knots and to increase the plant growth of okra.

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