RESEARCH ARTICLE

The efficacy of bio consortium of against *S. rolfsii* under *in vitro* conditions disease of chickpea

Rakesh Gurjar, A.R. Wasnekar, Mahesh Kumar Mimrot and Yashowardhan Singh

SUMMARY

A experiment was conducted in 2019-20 *Rabi* cropping season to obtain information on the incidence of chickpea diseases. The investigation was entitled the efficacy of bio consortium of against *S. rolfsii* under *in vitro* conditions disease of chickpea was conducted, Department of Plant pathology JNKVV. The experiment was laid out in Completely Randomized Design (CRD) concept comprising nine treatment combinations with three replications in vitro condition Dual culture technique was employed to test the efficacy of various bio consortiums. The maximum growth inhibition of *Sclerotium rolfsii* was recorded with *T. viride* + *T. harzianum* + *P. fluorescens* (65.74%). It is also found that treatment *T. viride* + *T. harzianum* + *P. fluorescens* are more efficient than other treatments according to the germination percentage, pre-emergence mortality, post-emergence mortality, phenotypic parameter and disease incidence.

Key Words: Bio consortium, S. rolfsii under in vitro conditions

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hickpea (*Cicer arietinum* L.) is a major legumes crop grown worldwide and ranks second in the global farming. It belongs to the family *Fabaceae*,

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Mahesh Kumar Mimrot, Department of Plant Pathology, Swami Keshwanand Rajasthan Agricultural University, Bikaner (Rajasthan) India sub family *Papilionaceae*. Chickpea is a rich source of protein (20 to 25 %) and also enriches soil fertility by biological nitrogen fixation. In India, it is grown over an area 95.39 lac hectares, with an annual production of 90.75 lac tonnes and yield was 951kg/ha. However, the total area and production of chickpea in MP is 35.90 lac hectare and 45.95 lac tonnes respectively, having productivity of 1056kg/ha (Anonyms, 2017). Chickpea collar rot is most serious and challenging disease which cause severe yield losses (upto 60-70%) under favourable conditions (Nene, 1984). The disease causes damage on root and stem of plant. The pathogen produces sclerotia which overwinter in soil and on plant debris besidesit can survive in a long period causing disease in the following season (Punja, 1985). Drying of plants with

foliage turned slightly yellow before death, scattered throughout the field is an indication of collar rot infection. The disease generally appears within two weeks of sowing and the younger plants collapse but older ones turn yellow and may dry without collapsing. The younger plants exhibit clear rotting at the collar region. The rotten portion is often covered with white mycelia strands of *S. rolfsii*. Thus, the control of the disease is very difficult. Various methods for controlling such disease have been investigated including the use of resistant varieties, chemical control, plant volatile compounds, plant extracts and biological control (Kumar and Tripathi, 1991; Dubey *et al.*, 2007 and El-Mougy *et al.*, 2007).

Management of plant disease through biological control has been considered as a viable alternative method as against the use of chemical pesticide and cultural practices. Different mode of action of bio control active micro-organism in controlling fungal plant disease include hyper-parasitism, predation, antibiosis, cross protection, competition for site and nutrient and induced resistance, the present investigation is mainly focussed to sort out the most effective organic amendments and antagonists for management of collar rot (*Sclerotium rolfsii*) disease of chickpea.

MATERIAL AND METHODS

The presentinvestigation was conducted to check the efficacy of bio consortium and oil cake against S. rolfsii under in vitro conditions. The pathogen was isolated from infected gram seedlings by hyphal tip method of fungal isolation. Identification of Sclerotium rolfsii were done by morphological characters formed white mat of hyaline mycelium with formation of initially white sclerotia which later turned into brown hard structure. Sclerotia were black, varied from spherical to irregular in shape and measured 80 to 85 µm in diameter. Pycnidial production was not observed in culture plates. Required bio-inoculants and oil cake Trichoderma viride, Trichoderma hazianum, Trichoderma aureoviride and Pseudomonas fluorescens, respectively were obtained from Microbes Research and Production Canter, JNKVV Jabalpur (M.P.).

Dual culture technique:

To test the efficacy of antagonistic fungus, twenty ml of sterilized melted PDA was plated in Petri plates and allowed to solidify. Mycelial discs measuring 5 mm diameter from three-day old cultures of both fungal antagonist and the test pathogen were pieced at equidistant on sterile petri plate containing PDA medium.

To test efficacy of antagonistic bacterium 4cm line was streaked at one side of the plate. On the opposite side to the antagonist, mycelia disc measuring 5 mm diameter from four-day old culture of test pathogen was placed on sterile petri plate containing PDA medium.

The petri plates with pathogen inoculated at one end alone, served as control. The petri plates were then incubated at 28±2°C. Four replications were maintained in each treatment. Growth of antagonists, pathogen and zone of inhibition of the pathogen in control plate. Per cent inhibition of mycelia growth over control was calculated by using the formula given by Vincent (1947):

$$I = \frac{C - T}{C} \times 100$$
where

I = Per cent inhibition in growth of test pathogen

C = Radial growth (mm) in control

T = Radial growth (mm) in treatment.

RESULTS AND DISCUSSION

The results obtained from the present investigation as well as relevant discussion have been summarized under following heads:

Evaluation of bio consortium against radial growth of *Sclerotium rolfsii*:

To compare the growth rate different consortiumon radial growth of Sclerotium rolfsii was studied and observation have been presented in Table 1 and Plate 2. In dule culture test, each of all tested consortium differentially limited the growth of the test pathogen and overgrew the pathogen colony when compared to control. Minimum radial growth (30.83 mm) of pathogen with maximum per cent inhibition of radial growth (65.74 %) was recorded in T. viride + T. harzianum + P. fluorescens followed by Trichoderma viride + Trichoderma harzianum wich recorded radial growth of 32.37 mm with radial growth inhibition of 64.04 per cent which were statistically at par with each other. Consortium T. viride + Pseudomonas fluorescens, T. viride + T. harzianum + T. aureoviride, T. harzianum+ Pseudomonas fluorescens, T. viride + T. aureoviride and T. harzianum + T. aureoviride inhibited radial growth of target pathogen by 61.30, 60.92, 59.07, 55.41 and 51.81 per cent, respectively. Maximum radial growth 47.43 mm and minimum inhabitation 47.30 per cent was recorded in T. viride + T. harzianum + T.



Palate 1: Collection isolation and identification of Sclerotium rolfsi

Table 1: Evaluation of bio consortium against on radial growth of S. rolfsii						
Sr. No.	Name of treatment	Radial growth of pathogen (mm)	Per cent growth inhibition			
T_1	Trichoderma viride + Trichoderma harzianum	32.37	64.04			
T_2	T. viride + T. aureoviride	40.13	55.41			
T_3	T. viride + Pseudomonas fluorescens	34.83	61.30			
T_4	$T.\ harzianum+T.\ aureoviride$	43.37	51.81			
T_5	T. harzianum + Pseudomonas fluorescens	36.83	59.07			
T_6	T. viride + T. harzianum + P. fluorescens	30.83	65.74			
T ₇	T. viride + T. harzianum + T. aureoviride	35.17	60.92			
T_8	T. viride + T.harzianum + T.aureoviride + Pseudomonas	47.42	47.20			
	fluorescens	47.43	47.30			
T ₉	Control	90.00	-			
	S.E. ±	0.33	-			
	C.D. (P=0.05)	0.98	-			

aureoviride + *P. fluorescens*. The test pathogen showed 90.00 mm growth after 120 hours of incubation

In dual culture test, each of all tested fungal and bacterial consortium limited the growth of the pathogen and overgrew the pathogen colony when compared to the control. In consortium treatment, maximum per cent inhibition of *Sclerotium rolfsii* was observed in *T. viride* + *T. harzianum* + *P. fluorescens* (65.74) similarly result reported by Singh *et al.* (2013) in which combination of *Trichoderma* spp. and *Pseudomonas* spp. to assess the synergistic effect of compatible isolates for plant growth promotion and management of *S. rolfsii*. In dual culture, in consortium treatment, maximum inhibition of *Sclerotium rolfsii* was observed in *T. viride* + *T. harzianum* + *P. fluorescens* (65.74%).

In dule culture test, each of all tested consortium differentially limited the growth of the test pathogen and overgrew the pathogen colony when compared to control. Minimum radial growth (30.83 mm) of pathogen with maximum per cent inhibition of radial growth (65.74 %) was recorded in T. viride + T. harzianum + P. fluorescens followed by Trichoderma viride + Trichoderma harzianumwich recorded radial growth of 32.37 mm with radial groth inhibition of 64.04 per cent which were statistically at par with each other. Consortium T. viride + Pseudomonas fluorescens, T. viride + T. harzianum + T. aureoviride, T. harzianum+ Pseudomonas fluorescens, T. viride + T. aureovirideand T. harzianum + T. aureovirideinhibited radial growth of target pathogen by 61.30, 60.92, 59.07, 55.41 and 51.81 per cent, respectively. Maximum radial growth 47.43 mm and minimum inhabitation 47.30 per cent was recorded in T. viride + T. harzianum + T.

aureoviride + *P. fluorescens*. The test pathogen showed 90.00 mm growth after 120 hours of incubation.

Effect of bio consortium against collar rot (*Sclerotium rolfsii*) disease incidence on variety JG-12:

Germination percentage:

Among the treatments minimum germination per cent 76.67 per cent was observed in T_8 (T. viride + T. harzianum + T. aureoviride + Pseudomonas fluorescens) followed by 77.78 per cent in T_4 (T. harzianum + T. aureoviride), 80.00 per cent in T_2 (T. viride + T. aureoviride), 81.11 per cent T_5 (T. harzianum + T seudomonas fluorescens), 84.44 per cent in T_7 (T. viride + T. harzianum + T. aureoviride), 86.67 per cent in T_3 (T. viride + T seudomonas fluorescens), 87.78 per cent in T_1 (T irichoderma viride + T irichoderma harzianum) and highest germination 91.11 per cent was recorded in T_6 (T. viride + T. harzianum + T. fluorescens) as compare to control.

Pre-emergence mortality:

Maximum pre-emergence mortality 23.33 per cent was recorded in T_8 (T. viride + T. harzianum + T. aureoviride + Pseudomonas fluorescens) followed by 22.22 per cent in T_4 (T. harzianum + T. aureoviride), 20.00 per cent in T_2 (T. viride + T. aureoviride), 18.89 per cent in T_5 (T. harzianum + Pseudomonas fluorescens), 15.56 per cent in T_7 (T. viride + T. harzianum + T. aureoviride), 13.33 per cent in T_3 (T. viride + Pseudomonas fluorescens), 12.22 per cent in T_1 ($Trichoderma\ viride + Trichoderma\ harzianum$). Minimum pre-emergence mortality 8.89 per cent was

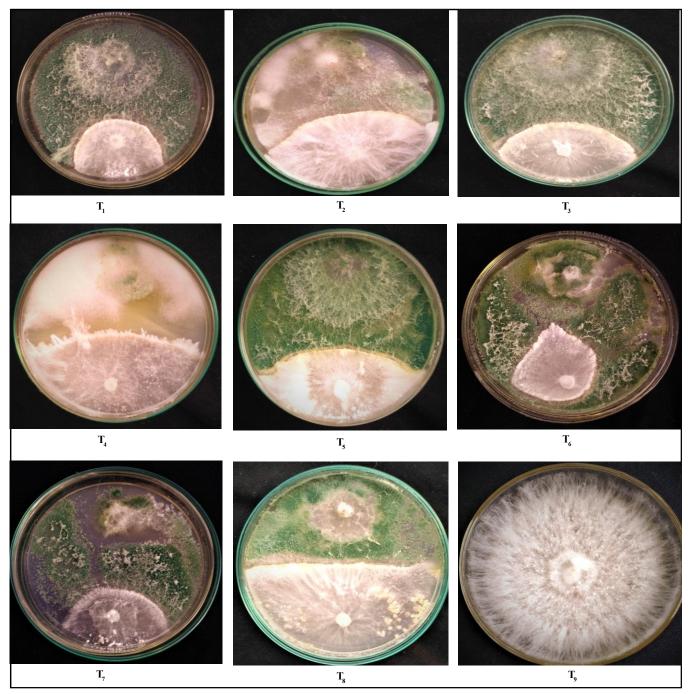


Plate 2: In-vitro efficacy of Bio-consortium against Sclerotium rolfsii after 120 hours

recorded in T_6 (*T. viride* + *T. harzianum* + *P. fluorescens*),) as compare to control.

Post-emergence mortality:

Maximum disease incidence of 24.48 and 22.77 percent was observed in T_8 (*T. viride* + *T. harzianum* + *T. aureoviride* + *Pseudomonas fluorescens*) and T_4 (*T. harzianum* + *T. aureoviride*), respectively.

Minimum disease incidence recorded in control.

Total mortality (%):

Maximum mortality of 90.17 per cent was receded in control (T_9) followed by 47.81 per cent was observed in T_8 (T. viride + T. harzianum + T. aureoviride + T. Pseudomonas fluorescens) and minimum mortality 17.37% was recorded in T_6 (T. viride + T. harzianum

+ P. fluorescens).

Effect of bio consortium on collar rot (*Sclerotium rolfsii*) disease and phenotypic parameters on variety JG-12:

Germination percentage:

All the treatment had higher germination percentage as compare to control. Among the treatments minimum germination per cent 78.60 per cent was observed in T_8 ($T. viride + T. harzianum + T. aureoviride + Pseudomonas fluorescens) followed by 79.77 per cent in <math>T_4$ (T. harzianum + T. aureoviride), 80.00 per cent in T_2 (T. viride + T. aureoviride), 83.50 per cent T_5 (T. harzianum + Pseudomonas fluorescens), 85.54 per cent in T_7 (T. viride + T. harzianum + T. aureoviride), 88.67 per cent in T_3 (T. viride + Pseudomonas fluorescens), 89.85 per cent in T_1 (Trichoderma viride + Trichoderma harzianum) and highest germination

94.40 per cent was recorded in T_6 (*T. viride* + *T. harzianum* + *P. fluorescens*).

Plants height (Shoot and Root):

Maximum shoot height 24. 33, 22.07, 20.07, 19.27, 16.37, 15.67, 14.23 and 13.30 cm were recorded in T_6 (T. viride + T. harzianum + P. fluorescens), T_1 ($Trichoderma\ viride + Trichoderma\ harzianum$), T_3 (T. $viride + Pseudomonas\ fluorescens$), T_7 (T. viride + T. harzianum + T. aureoviride), T_5 (T. $harzianum + Pseudomonas\ fluorescens$), T_2 (T. viride + T. aureoviride), T_4 (T. harzianum + T. aureoviride), T_8 (T. viride + T. harzianum + T. $aureoviride + Pseudomonas\ fluorescens$), respectively. As compare to control T_9 (control) which was 10.67 cm, respectively.

Maximum root height 10.37,10.27, 10.00, 9.60, 9.40, 9.10, 8.36 and 8.03 cm were recorded in T_6 (*T. viride* + *T. harzianum* + *P. fluorescens*), T_1 (*Trichoderma*

Table 2: Evaluation of bio consortium against on radial growth of S. rolfsii					
T.N.	Name of treatment	Radial growth of pathogen(mm)	Per cent growth inhibition		
T_1	Trichoderma viride + Trichoderma harzianum	32.37	64.04		
T_2	T. viride + T. aureoviride	40.13	55.41		
T_3	T. viride + Pseudomonas fluorescens	34.83	61.30		
T_4	T. harzianum + T . aureoviride	43.37	51.81		
T_5	T. harzianum + Pseudomonas fluorescens	36.83	59.07		
T_6	$T. \ viride + T. \ harzianum + P. \ fluorescens$	30.83	65.74		
T_7	T. viride + T. harzianum+ T.aureoviride	35.17	60.92		
T_8	T. viride + T.harzianum + T.aureoviride + Pseudomonas fluorescens	47.43	47.30		
T ₉	Control	90.00	-		
	S.E.±	0.33	-		
	C.D. (P=0.05)	0.98			

Table 3	Table 3: Effect of bio consortium against collar rot (Sclerotium rolfsii) disease incidence on variety JG-12						
T.N.	Combination	Germination	Pre emergence mortality	Post emergence mortality	Total mortality		
T_1	Trichoderma viride + Trichoderma harzianum	87.78	12.22	13.96	26.18		
T_2	T. viride + $T.$ aureoviride	80.00	20.00	20.74	40.74		
T_3	T. viride + Pseudomonas fluorescens	86.67	13.33	15.50	28.83		
T_4	$T.\ harzianum+T.\ aureoviride$	77.78	22.22	22.77	44.99		
T_5	T. harzianum + Pseudomonas fluorescens	81.11	18.89	20.50	39.39		
T_6	T. viride + T. harzianum + P. fluorescens	91.11	8.89	8.48	17.37		
T_7	T. viride + T. harzianum+ T.aureoviride	84.44	15.56	20.92	36.48		
T_8	T. viride + T. harzianum+ T.aureoviride + Pseudomonas fluorescens	76.67	23.33	24.48	47.81		
T ₉	Control	62.22	36.67	53.51	90.17		
	S.E. ±	1.92	1.99	2.83	2.93		
	C.D. (P=0.05)	5.76	5.97	8.48	8.77		

viride + Trichoderma harzianum), T_3 (T. viride + Pseudomonas fluorescens), T_7 (T. viride + T. harzianum + T. aureoviride), T_5 (T. harzianum + Pseudomonas fluorescens), T_2 (T. viride + T. aureoviride), T_4 (T. harzianum + T. aureoviride) and T_8 (T. viride + T. harzianum + T. aureoviride + Pseudomonas fluorescens), respectively. As compare to control T_9 (control) which was 6.43 cm, respectively.

Vigour index:

Maximum vigour index per cent of 3,274.21 and 2,906.31 was recorded in T_6 (*T. viride* + *T. harzianum* + *P. fluorescens*) and T_1 (*Trichoderma viride* + *Trichoderma harzianum*), as compare to control. Minimum vigour index per cent of 1,676.34 and 1,794.34

were observed in T_8 (*T. viride* + *T. harzianum* + *T. aureoviride* + *Pseudomonas fluorescens*) and T_4 (*T. harzianum* + *T. aureoviride*), respectively.

Effect of bio consortium on collar rot (*Sclerotium rolfsii*) disease incidence on variety JG-14:

Germination percentage:

Data presented in Table 5 indicated that germination percentage among treatment range from 77.78 to 92.22 per cent. All the treatment had higher germination percentage as compare to control. Among the treatments minimum germination per cent 77.78 per cent was observed in T_8 (*T. viride* + *T. harzianum* + *T. aureoviride* + *Pseudomonas fluorescens*) followed by 78.89 per cent in T_4 (*T. harzianum* + *T. aureoviride*),

Table 4: Effect of bio consortium on collar rot (Sclerotium rolfsii) disease and phenotypic parameters on variety JG-12						
T.N.	Combination	Germination	Shoot length	Root length	Vigour index	
T_1	Trichoderma viride + Trichoderma harzianum	89.85	22.07	10.27	2,906.31	
T_2	$T. \ viride + T. \ aureoviride$	80.00	15.17	9.10	1,941.33	
T_3	T. viride + Pseudomonas fluorescens	88.67	20.07	10.00	2,666.47	
T_4	T. harzianum $+$ $T.$ aureoviride	79.77	14.23	8.27	1,794.34	
T_5	T. harzianum + Pseudomonas fluorescens	83.50	16.37	9.40	2,151.22	
T_6	$T. \ viride + T. \ harzianum + P. \ fluorescens$	94.40	24.33	10.37	3,274.21	
T_7	T. viride + T. harzianum+ T.aureoviride	85.54	19.27	9.60	2,469.57	
т	T. viride + T. harzianum+ T.aureoviride + Pseudomonas	78.60	13.30	8.03	1,676.34	
T ₈	fluorescens					
T ₉	Control	70.32	10.67	6.43	1,202.93	
	S.E. ±	1.02	0.45	0.16	45.07	
	C.D. (P=0.05)	3.05	1.33	0.48	134.96	

Table 5: Effect of bio consortium on collar rot (Sclerotium rolfsii) disease incidence on variety JG-14					
T.N.	Combination	Germination	Pre emergence mortality	Post emergence mortality	Total mortality
T_1	Trichoderma viride + Trichoderma harzianum	88.89	11.11	12.49	23.60
T_2	T. $viride + T.$ $aureoviride$	81.11	18.89	19.17	38.06
T_3	T. viride + Pseudomonas fluorescens	87.78	12.22	15.19	27.42
T_4	T. $harzianum + T$. $aureoviride$	78.89	21.11	22.49	43.60
T_5	T. harzianum +Pseudomonas fluorescens	83.33	16.67	18.69	35.35
T_6	$T.\ viride + T.\ harzianum + P.\ fluorescens$	92.22	7.78	8.47	16.24
T ₇	T. viride + T. harzianum+ T.aureoviride	85.56	14.44	15.54	29.99
T_8	T. viride + T. harzianum+ T.aureoviride + Pseudomonas fluorescens	77.78	22.22	24.27	46.49
T ₉	Control	65.56	34.44	50.79	85.24
	S.E.±	1.77	2.32	2.60	2.36
	C.D. (P=0.05)	5.32	6.12	5.50	7.06

81.11 per cent in T_2 (T. viride + T. aureoviride), 83.33 per cent T_5 (T. harzianum + Pseudomonas fluorescens), 85.56 per cent in T_7 (T. viride + T. harzianum + T. aureoviride), 86.67 per cent in T_3 (T. viride + Pseudomonas fluorescens), 88.89 per cent in T_1 ($Trichoderma\ viride + Trichoderma\ harzianum$) and highest germination 92.22 per cent was recorded in T_6 (T. viride + T. harzianum + P. fluorescens).

Pre-emergence mortality

Post-emergence mortality:

Disease intensity, at flowering stage, among treatment varied from 8.47 to 24.27 as compared to control post-emergence mortality was 50.79 per cent. Maximum disease incidence of 24.27 and 22.49 per cent was observed in T_8 (T. viride + T. harzianum + T. aureoviride + T pseudomonas fluorescens) and T_4 (T harzianum + T aureoviride), respectively. Minimum

disease incidence of 8.47 was recorded in T_6 (*T. viride* + *T. harzianum* + *P. fluorescens*).

Total mortality (%):

Total mortality per cent among treatment ranges from 16.24 to 46.49 per cent as compared to control 85.24 per cent. Maximum mortality of 46.49 per cent was observed in T_8 (*T. viride* + *T. harzianum* + *T. aureoviride* + *Pseudomonas fluorescens*) and minimum mortality 16.24 per cent was recorded in T_6 (*T. viride* + *T. harzianum* + *P. fluorescens*).

Effect of bio consortium on collar rot (*Sclerotium rolfsii*) disease and phenotypic parameters on variety JG-14:

Germination percentage:

Data presented in Table 6 indicated that germination percentage among treatment range from 80.68 to 96.52 per cent. All the treatment had higher germination percentage as compare to control. Among the treatments minimum germination percent 80.68 per cent was observed in T_s (T. viride + T. harzianum + T. aureoviride + Pseudomonas fluorescens) followed by 82.55 per cent in T_A (*T. harzianum* + *T. aureoviride*), 84.45 per cent in T_2 (*T. viride* + *T. aureoviride*), 87.76 per cent T₅ (T. harzianum + Pseudomonas fluorescens), 89.35 per cent in T_7 (T. viride + T. harzianum + T. aureoviride), 90.70 per cent in T_3 (T. viride + Pseudomonas fluorescens), 91.85 per cent in T₁ (Trichoderma viride + Trichoderma harzianum) and highest germination 96.52 per cent was recorded in T_6 (T. viride + T. harzianum + P. fluorescens).

Table 6: Effect of bio consortium on collar rot (Sclerotium rolfsii) disease and phenotypic parameters on variety JG-14						
T. N.	Combination	Germination	Shoot length	Root length	Vigour index	
T_1	Trichoderma viride + Trichoderma harzianum	92.85	24.90	10.73	3,309.11	
T_2	T. $viride + T.$ $aureoviride$	84.45	16.37	9.20	2,159.88	
T_3	T. viride + Pseudomonas fluorescens	90.70	22.77	10.23	2,993.02	
T_4	T. harzianum + T. aureoviride	82.55	15.17	8.63	2419.17	
T_5	T. harzianum + Pseudomonas fluorescens	87.76	18.10	9.47	2,463.27	
T_6	T. viride + T. harzianum + P. fluorescens	96.52	26.37	11.53	3,658.07	
T ₇	T. viride + T. harzianum+ T.aureoviride	89.35	21.10	9.67	2749.31	
	T. viride + T. harzianum+ T.aureoviride +	80.68	15.20	8.27	1,893.21	
T_8	Pseudomonas fluorescens					
T ₉	Control	77.60	12.57	6.77	1,500.40	
	S.E. \pm	0.67	0.41	0.35	57.44	
	C.D. (P=0.05)	2.02	1.24	1.04	171.98	

Plants Height (Shoot and Root):

Data represented in Table 6 at the time of maturity so that shoot hight and root length was significantly increased in all the treatment except T_9 (control) which was 12.57 cm and 6.77 cm, respectively.

Maximum shoot height 26.37, 24.90, 22.77, 21.10, 18.10, 14.37, 16.17 and 15.20 cm were recorded in T_6 (T. viride + T. harzianum + P. fluorescens), T_1 ($Trichoderma\ viride + Trichoderma\ harzianum$), T_3 (T. $viride + Pseudomonas\ fluorescens$), T_7 (T. viride + T. harzianum + T. aureoviride), T_5 (T. $harzianum + Pseudomonas\ fluorescens$), T_2 (T. viride + T. aureoviride) and T_8 (T. viride + T. harzianum + T. aureoviride) and T_8 (T. viride + T. harzianum + T. $aureoviride + Pseudomonas\ fluorescens$), respectively.

Maximum root lenght 11.53,10.73, 10.23, 9.67, 9.47, 9.20, 8.63 and 8.27 cm were recorded in T_6 (T. viride + T. harzianum + P. fluorescens), T_1 (Trichoderma viride + Trichoderma harzianum), T_3 (T. viride + T2. harzianum + T3. (T4. viride + T5. harzianum + T5. (T5. T7. (T7. viride + T8. T9. (T8. viride + T9. T9. (T8. viride + T9. aureoviride), T9. (T9. viride + T9. aureoviride), T9. (T9. viride + T9. aureoviride + T9. T9. (T9. viride + T9. aureoviride + T9. T9. T9. (T8. viride + T8. T9. T9. T9. (T8. viride + T8. T9. T9. T9. (T8. viride + T9. T9. T9. (T9. viride + T9. T9. T9. (T9. viride + T9. viride +

Vigour index

Data presented in Table 6 indicated that vigour index (%) recorded at the time of maturity sowing that indicated that all the treatment has higher vigour index per cent as compared to T_9 (control). At the time of maturity, it varied from 1,893.21 to 3,658.07 as compare to 1,500.40 in T_9 (control). Maximum vigour index per cent of 3,658.07 and 3,309.11 was recorded in T_6 (T. viride + T. harzianum + P. fluorescens) and T_1 (T (T ichoderma

viride + Trichoderma harzianum). Minimum vigour index per cent of 1,893.21 and 2419.17 were observed in T_8 (T. viride + T. harzianum + T. aureoviride + Pseudomonas fluorescens) and T_4 (T. harzianum + T. aureoviride), respectively.

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