SUPPRESSION OF Fusarium solani IN VOLVED IN COTTON SEEDLING DAMPING – OFF BY Trichoderma SPP.

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ABSTRACT

Five isolates of *Trichoderma* spp., three belonging to *T. harzianum* and two belonging to T.viride were isolated from cotton seedlings infected with damping -off or from rotted roots of adult plants. The isolates were evaluated for biocontrol capacity against 14 isolates of Fusarium solani involved in seedling damping -off, of cotton under greenhouse conditions. Analysis of variance showed very highly significant effects of Trichoderma spp. isolates. F. solani isolates, and their interaction on preemergence damping- off, post-emergence damping-off, survival, plant height, and dry weight. Responses of F. solani isolates to antagonistic effects of Trichoderma isolates were different. Some isolates of F. solani exhibited response to all Trichoderma isolates, which siginifficantly increased the percentage of surviving seedlings, plant height and dry weight. Cluster analysis of Trichoderma spp. isolates based on antagonistic patterns showed that isolates were placed in two groups. The first group included isolates of T. harzianum, while the second group included isolates of T. viride. The results of cluster analysis of F. solani isolates, based on their response patterns to Trichoderma spp. Isolates. Placed the isolates in three distinct groups; however, grouping the isolates was not related to geographic origins. It is worth noting that isolates of Middle Egypt and Upper Egypt were placed in same group.

Keywords: Biological control, cotton, damping-off, *Gossypium barbadense* L., *F. solani* and *Trichoderma* spp.

ITTRODUCTION

Fusarium solani (Mart.) Sacc. is one of the most ubiquitous soil fungus and destructive plant pathogen of many of hosts (Booth,1971;Domsch et al.,1980). It is easily isolated from seeds. The degree of losses caused by F. solani has not been determined F. solani is one of the organisms contributing to the seedling disease complex of Gossypium spp. (Davis et al.,1981; Johnson, 1981). Disease symptoms include seed rot, pre-and potemergence damping-off, and seedling root rot which, individually or in combination, result in stand reductions and reduced seedling vigor that delays growth and maturity, F. solani caused significant reductions in emergence of cotton and increased root discoloration of surviving seedlings (Bastson and Trevathan, 1988). F. solani was isolated from cotton plants with severe foot rot in India during 1977, 1978 and 1980(Bharathudu and Rao,1982). Nelson and Windels(1992), This disease occurs as seed decay before germination and as pre-emergence damping-off (Abd-Elsalam et al 2007). Many studies demonstrated that some of the isolates of Trichoderma

spp. showed biocontrol potentiality against several micro-organtisms involved in cotton damping-off, and root-rot disease. Aly et al., 2000; Hanson,2000; Haq and Khan,2000 Xueling et al., 2003; Asran et al., 2005; Howell and Pukhaber, 2005, Aly et al., 2007and Asran-Imal et.al 2010). F. solani has been effectively controlled through Seed and soil treatment with Trichoderma virens preparations (Howell,1982 &1991 and Nelson, 1994). The objectives of this study were to: (1) isolate and to identify Fusarium spp. and Trichoderma spp. from cotton seedlings. (2) to evaluate the biocontrol capacity of isolated Trichoderma spp. against Fusarium spp. pathogenic to cotton seedlings.

MATERIALS AND METHODS

Isolation, purification and identification of Fusarium solani and Trichoderma spp. from cotton (Gossypium barbadense L.) roots :

Isolation, was made from samples collected from several localities in cotton producing areas in ten governorates, i.e Beheira, Dakahliya, Damietta, Gharbiya, ,kafr El-sheikh, Minufiya , sharkiya, Minya, Assiute, and Sohag (Table 1&2). Each sample consisted of 5 to 15 seedlings showing a variety of damping-off symptoms or rotted roots of 5 adult plants . Seedling or roots of adult plants were removed from soil and washed thoroughly under running tap water to remove any adhering soil . Small pieces (approximately 0.5 cm long) of necrotic root tissues were surface sterilizes with 10% clorox solution for 2 minutes, and washed several times with sterilized water. The surface sterilized pieces were then blotted dry between sterilized filter papers and plated on potato dextrose agar (PDA) medium amended with streptomycin sulphate or penicillin G and rose bengal to eliminate any bacterial contamination. The plates were incubated at 25±2° C for 3-7 days. Fusarium solani isolates were identified to species level according to Nelson et al. (1983) while, Trichoderma spp. isolates were identified to species level according to Rifai (1969). Identification of isolates to species level was kindly verified by Mycological Centre. Assiut University.

Production of Fusarium solani inoculum used for soil infestation:

Substrate for growth of isolates was prepared in 500-ml glass bottles. Each contained 50g of sorghum grains and 40 ml tap water. Contents of each bottle were autoclaved for 30 minutes .inocula were taken from one-week old PDA cultures and aseptically introduced into the bottles and allowed to colonize sorghum grains for three weeks.

Production of *Trichoderma* spp. inoculum used for seed treatment (seed coating):

Inocula of *Trichaderma* spp. isolates were prepared as previously mentioned; however antagonist-sorghum mixtures. were air –dried in the greenhouse. The dry mixtures were triturated to a fine powder in a blender (Papaviza and Lewis,1981).

Interaction between *Trichoderma* spp. and *F. solani* isolates under greenhouse conditions :

Autoclaved clay loam soil was placed on a greenhoused bench and infested with inoculum of each *F. solani* isolate at a rate of 50g/kg soil. After thoroughly mixing, infested soil was dispensed into 15-cm –diameter clay

pots. In the control treatment, soil was infested with sorghum powder at the same rate. Slightly moistened seeds were treated with powdered inoculum of each *Trichoderma* isolate, at rate of 10g/ kg seeds and thoroughly shaken in plastic bags before being planted at the rate of 10 seeds/pot of *F. solani* ifested soil. Pots (4 for each treatment) were randomly distributed on a greenhouse bench under the temperature/ regime of 29-±5°C. Preemergence damping- off was recorded 15 days after planting, postemergence damping-off, survivals, plant height (cm/plant) and dry weight (mg/plant) were recorded 45 days after planting .

Table 1. Geographic origins of *Trichoderma* spp. used in the study

Isolate No.	Geographic origin	Trichoderma spp.
1	Dakahliya	T. harziaunum
2	Sharkiya	T . harziaunum
3	KafrEl-sheikh	T . harziaunum
4	Gharbiya	T . viride
5	Beheira	T . viride

Table 2.Geographic origins of Fusarim solani isolates used in the study

Isolate No.	Governorate	Previous Crops	Host cultivar
1	Beheira	Egyptian clover	Giza 70
2	Kafr El-sheikh	Egyptian clover	Giza 89
3	Dakahliya	Egyptian clover	Giza 86
4	Minufiya	Onion	Giza 86
5	Dakahliya	Faba bean	Giza 86
6	Beheira	Egyptian clover	Giza 70
7	Sharkiya	Egyptian clover	Giza 89
8	Kafr El-sheikh	Egyptian clover	Giza 89
9	Damietta	Faba bean	Giza 88
10	Gharbiya	Faba bean	Giza 89
11	Minya	Faba bean	Giza 83
12	Sharkiya	Pea	Giza 89
13	Sohag	Onion	Giza 83
14	Assiute	Pea	Giza 83

Statistical andalysis of data:

The experimental design of the present study was a randomized complete block design with four replicates. Analysis of variance (ANOVA) of the data was performed with the MSTAT-C statistical package. Least significant difference (LSD) was used to compare treatment means. Percentage date was transformed into arcsine angles before carrying out the ANOVA to produce approximately constant variance. Cluster analysis was performed with the software package SPSS 6.0.

RESULTS

In vivo evaluation of Trichoderma spp. antagonism against F. solani isolates:

ANOVA (Table 3) showed that *Fusarium solani* isolates and *Trichoderma* isolates was highly significant source of variation in all the tested parameters, the interaction was highly significant source of variation in all the tested parameters except plant height.

Table3. Analysis of variance of the effect of *Trichoderma* spp. Isolates, Fusarim solani isolates and their interaction on cotton seedling disease variables (cv. Giza 92) under greenhouse conditions

Parameter and Source of variation	D.f.	M.S.	F .value	P > F								
Pre-emergence damping off												
Replication ^a	3	72.613	0.892	0.479								
Trichoderma isolates (T)	5	728.907	8.326	0.000								
Fusarium isolates (F)	13	347.844	3.93	0.000								
TXF	65	173.652	1.984	0.000								
Error	249	87.542										
F	Post-emerge	ence damping off										
Replication	3	134.312	0.835	0.476								
Trichoderma isolates(T)	5	2052.318	12.754	0.000								
Fusarium isolates(F)	13	1333.071	8.284	0.000								
TXF	65	532.078	3.307	0.000								
Error	249	160.919										
	S	urvival										
Replication	3	222.763	1.848	0.139								
Trichoderma (T)	5	661.191	5.487	0.000								
Fusarim (F)	13	512.579	4.253	0.000								
TXF	65	250.142	2.076	0.000								
Error	249	120.510										
	Plai	nt height										
Replication	3	186.813	1.461	0.226								
Trichoderma isolates (T)	5	310.605	2.429	0.004								
Fusarium isolates(F)	13	836.455	6.543	0.000								
TXF	65	141.386	1.106	0.290								
Error	249	127.847										
	Dry	weight										
Replication	3	19058.670	1.600	0.190								
Trichoderma isolates (T)	5	46262.888	3.883	0.000								
Fusarium (F)	13	171182.567	14.369	0.000								
TXF	65	11913.563	1.836	0.000								
Error	249	11913.563										

^a Replication is random, while each of *Trichoderma* . isolates and *Fusarium solani* isolates are fixed.

Table 4 showed that *Trichoderma* isolates were the most important source of variation in pre-and post emergence damping –off, and survival, while *Fusarium* isolates were the most important source at variation in plant height and dry weight. The interaction was always the least important source of variation. Due to the significant interaction of *Trichoderma* isolate× *Fusarium* isolate in pre emergence damping –off, an interaction LSD was

calculated to compare between Trichoderma isolates and the control for each Fusarium isolate; these comparisons showed that the difference was not the same for each Fusarium isolate for example, the difference between T. harzianum and the control was significant for Fusarium isolate No. 3, while it was nonsignificant for Fusarium isolate no. 2, another example is the non significant effect of T. harzianum no.1 on Fusarium no. 11 and the significant effect on Fusarium no. 12. The difference between Trichoderma isolates may vary depending on Fusarium isolate. For example, the difference between T4 and T5 was not significant on Fusarium no.1, while this difference was significant on fusarium no.5. the same conclusions hold true for post emergence damping-off, survival, and dry weight, where Trichoderma x Fusarium interaction was significant. Due to the lack of significant Fusarium. × Trichoderma interaction in the case of plant height, the general means were used to compare between the effects of Trichoderma isolates on plant height. These comparisons showed that all the tested Trichoderma isolates significantly increased plant height regardless of Fusarium isolates.

Table 4. Relative contribution of *Trichoderma* isolates, *Fusarium solani* isolates and their interaction to variation in cotton seedling disease variables (cv. Giza 92) under greenhouse conditions

Source of	Relative contribution a to variation in										
variation	Pre- emergence damping-off	Pot – emergence damping-off	Survival	Plant height	Dry weight						
Trichoderma isolates(T)	55.09	50.65	40.15	21.05	17.91						
Fusarium isolates (F)	26.29	32.90	31.13	56.70	66.25						
TxF calculated	13.13	13.13	15.19	9.58	8.35						

^a Colculated as percentage of squares of the explained (model) variation .

The comparisons between *Trichoderma* isolates and control within isolates of F. solani (Table 8) revealed that efficiency of the tested Trichoderma isolate in increasing the plant height of surviving seedlings was varied from one isolate to another. Thus, (T2) was effective against F. solani isolate No. 3 and ineffective against No.9 and 14 Trichoderma (T1) was effective against F. solani isolate No.1 and ineffective against F. solani isolate No. 13 F. solani isolate responded differrntly to the application of Trichoderma isolates for example F. solani Nos. 1,2,3,6 & 7 were highly responsive to Trichoderma isolates, while isolates Nos. 10,13& 14 had no response to any Trichoderma isolate. The majority of Trichoderma isolates showed significant effects no dry weight of surviving seedling Table (8). T. viride isolates significantly increased dry weight of seedlings regardless of F. solani isolate, while, T. harzianum showed various significant effects. On the other hand, Trichoderma isolates did not show significant effects in improving dry weight of seedling in the case of F. solani isolates Nos. 9,13 and 14. Fusarium isolates showed variable effects of plant height.

Tabel 5. Effect of Trichoderma isolates, Fusarium solani isolates and their interaction on pre-emergence damping -off of cotton

seedlings (cv.Giza 92) under greenhouse conditions

occumings (ovicine oz) andor greenine dec centarients												
	T.		T.		T.		1	Г.		T.		
	harz	zianum	harzianum		harzianum		vir	ide	vi	ride	control	
	(T1)	(T2)		(T3)		(T4)		(1	Γ5)		
Fusarium solani		b						Trono		Trono	·	Trans-
Joiann	%	Trans-	%	Trans-	0/	0/	0/	Trans-	0/	Trans-	0/	
	70	forme	70	formed	%	%	%	forme	%	forme	%	forme
		d						d		d		d
1	10 ^a	15.86	17.5	24.53	225	27.33	32.5	34.72	30	3305	27.5	81.39
2	30	32.53	17.5	21.58	20	26.19	27.5	30.87	30	32.83	32.5	33.75
3	15	19.92	0.0	0.0	15	19.55	15	22.50	22.5	26.41	35.0	36.0
4	37.5	37.51	27.5	31.05	15	22.50	27.5	30.80	27.5	31.39	27.5	31.39
5	42.5	40.39	22.5	28.22	30	33.05	17.5	24.16	30	32.31	55	47.88
6	30	33.05	32.5	34.50	32.5	33.75	10	15.86	35	36	30	32.53
7	45	42.05	17.5	17.89	22.5	28.22	5	6.64	35	36	27.5	31.55
8	35	36.06	25	29.89	22.5	28.22	22.5	27.86	30	32.53	37.5	43.49
9	37.5	37.75	35	36.06	27.5	30.87	30	32.31	27.5	3139	50	44.94
10	32.5	34.34	30	32.31	20	23.25	45	42.05	30	32.67	42.5	40.61
11	27.5	31`.39	25	29.36	22.5	27.86	20	25.08	22.5	27.86	25	29.36
12	17.5	21.58	10	15.86	27.5	31.39	25	29.14	37.5	37.51	35	35.94
13	27.5	31.39	22.5	28.22	30	33.05	25	29.89	30	33.05	37.5	33.75
14	27.5	31.39	37.5	37.51	42.5	40.61	22.5	28.22	22.5	38.22	25	29.36

^a Mean of four replicates.

LSD (transformed date) for :

Trichoderma isolates (T)..... = 3.50 ($P \le 0.05$) Fusarium isolate (F)=5.34($P \le 0.05$) Interaction $T \times F$ = 13.1($P \le 0.05$)

Tabel 6. Effect of Trichoderma isolates, Fusarium solani isolates and their interaction on post-emergence damping -off of cotton seedling (cv.Giza 92) under greenhouse conditions

	T.Harzianum		T.Harzianum		T.harzianum		<i>T.</i> v	⁄iride	<i>T.</i> v	riride	control		
Fusarim	(Τ1)	(T	2)	(]	(T3)		(T 4)		Γ5)	Control		
solani	%	Trans- b formed	%	Trans- formed	%	%	%	Trans- formed	%	Trans- formed	%	Trans- formed	
1	27.5 ^a	31.63	67.5	55.24	47.5	43.57	50	45	50	45	45	42.13	
2	22.5	28.32	32.5	34.76	62.5	52.24	42.5	40.69	37.5	34.76	50	45	
3	57.5	49.31	32.5	34.76	65	53.37	52.5	46.43	60	50.77	32.5	34.76	
4	30	33.21	32.5	34.76	52.5	46.43	67.5	55.24	55	47.87	55	47.87	
5	20	26.56	17.5	24.73	52.5	46.43	37.5	37.76	57.5	49.31	42.5	40.69	
6	25	30	42.5	40.69	37.5	37.76	32.5	34.76	50	45	70	56.79	
7	25	30	65	53.37	42.5	40.69	37.5	37.76	55	47.87	57.5	49.3	
8	52.5	46.43	60	50.77	47.5	43.57	32.5	34.76	40	39.23	42.5	40.69	
9	42.5	40.69	60	50.77	52.5	46.43	35	36.27	57.5	49.31	37.5	37.76	
10	52.5	46.43	62.5	52.24	77.5	61.68	55	47.87	70	56.79	52.5	46.43	
11	45	42.13	65	53.37	75	60	62.5	52.24	52.5	46.43	75	60	
12	35	36.27	70	56.79	52.5	46.43	50	45	52.5	46.43	55	47.87	
13	72.5	58.37	65	53.37	57.5	49.31	45	42.13	45	42.13	62.5	52.24	
14	67.5	55.24	57.5	49.31	50	45	45	42.13	37.5	37.76	65	53.37	

^a Mean of four replicates.

LSD (transformed date) for:

Trichoderma isolate (T)..... ...= 4.107 ($P \le 0.05$) Fusarium isolate (F)=6.27($P \le 0.05$) Interaction $T \times F$ = 15.37($P \le 0.05$)

^bPercentage date were transformed into drcsine angels before carrying out the analysis of variance to produce approximately constant variance.

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Trichoderma isolates were divided into two main groups based on their antagonism pattern. The first group included the three isolates of T. harzianum and the 2^{nd} group included the two isolates of T. viride (Fig.1) Fusarium isolates were divided into three groups based on their response pattern to the antagonism of Trichoderm isolates. Each group included isolates from different governorates.

Tabel 7.Effect of *Trichoderma* isolates, *Fusarium solani* isolates and their interaction on survival of cotton seedling (cv.Giza 92) under greenhouse conditions

	T.Har	rzianum	T.Harzianum		Tharzianum		T.viride		T.viride		control		
Fusarium	(T1)		(T2)		(T3)		(T 4)		(.	T5)	Control		
solani	%	Trans- b formed		Trans- formed	%	%	%	Trans- formed	%	Trans- formed	%	Trans- formed	
1	62.5 ^a	52.24	15	22.79	30	33.21	17.5	24.73	20	26.56	27.5	31.63	
2	47.5	43.57	50	45	17.5	24.73	30	33.21	32.5	34.76	17.5	24.73	
3	30	33.21	57.5	49.31	20	26.56	32.5	34.76	17.5	24.73	32.5	34.76	
4	32.5	34.76	40	39.23	32.5	34.76	2.5	9.10	17.5	24.73	17.5	24.73	
5	35	36.27	60	50.77	17.5	24.73	45	42.13	12.5	20.70	2.5	9.10	
6	25	30	25	30	30	33.21	57.5	49.31	15	22.79	0.0	0.0	
7	30	33.21	17.5	24.73	35	36.27	57.5	49.31	10	18.44	15	22.79	
8	12.5	20.70	15	22.79	30	33.21	45	42.13	30	33.21	10	18.44	
9	12.5	20.70	5	12.92	20	26.56	35	36.27	15	22.79	12.5	20.70	
10	15	22.79	7.5	15.89	2.5	9.10	0.0	0.0	2.5	9.10	5	12.92	
11	27.5	31.63	10	18.44	2.5	9.10	17.5	24.73	25	30	0.0	0.0	
12	27.5	31.63	20	26.56	17.5	24.73	25	30	10	18.44	5	12.92	
13	0.0	0.0	7.5	15.89	12.5	20.70	30	33.21	25	30	0.0	0.0	
14	5	12.92	5	12.92	2.5	9.10	32.5	34.76	40	39.23	7.5	15.89	

^a Mean of four replicates.

LSD (transformed date) for isolate of:

^bPercentage date were transformed into drcsine angels before carrying out the analysis of variance to produce approximately constant variance.

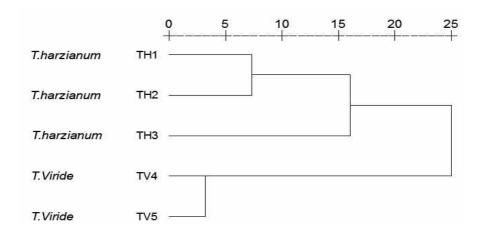
Table 8. Effect of *Trichoderma* isolates, *Fusarium solani* isolates and their interaction on plant height and dry weight of cotton seedlings (cv. Giza 92) under greenhouse conditions

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Fusarim		Plant height (cm/plant)							Dry weight (mg / plant)					
solani	(T1)	(T2)	(T3)	(T 4)	(T5)	control	Mean	(T1)	(T2)	(T3)	(T4)	(T5)	control	
1	13.01 ^a	24.62	12.31	24.68	22.76	19.95	19.56	204	301.5	314	296.3	179.5	186.8	
2	27.40	26.48	23.27	27.22	21.14	18.62	28.56	201	293.3	227.3	318.3	264	136	
3	26.33	31.01	23.15	33.14	25.67	20.56	26.64	286	314.3	281.3	244.5	162	152.3	
4	22.22	22.92	27.93	26.85	24.38	22.30	24.43	307.3	311.3	280.3	76.8	244.5	128.8	
5	23.52	19.95	21.19	29.22	24.33	13.13	21.89	278.3	272.3	234	279.5	174.3	45.8	
6	23.59	29.16	20.79	20.52	20.76	0.0	19.14	210.8	268.3	317	257.8	321.3	0.0	
7	22.33	20.57	7.13	32.61	21.86	10.89	19.23	280.8	294.3	265	201.3	155	143.3	
8	27.94	22.24	20.08	26.89	22.55	21.44	23.52	224.8	238.3	227.3	209.5	304.3	109	
9	17.08	13.05	22.01	23.42	21.84	6	17.23	321.5	164.5	219.8	294	163.3	118.5	
10	22.67	20.88	24.92	0.0	24.07	17.91	18.41	255.8	235	83.25	0.0	81.5	62.8	
11	23.81	25.41	26.09	15.69	23.68	18.69	22.23	183.5	87.5	81	240.5	309.3	0.0	
12	23.89	21.64	18.78	24.64	20.67	11.55	20.19	260.5	252.3	239.5	242.5	146.8	63.5	
13	0.0	19.64	22.54	23.99	0.0	21.33	14.58	0.0	98.5	249.8	283.8	234.5	0.0	
14	23.93	25.32	10.52	24.38	18.17	15.43	19.83	151	164.8	88.8	157.3	123.5	118.3	
Mean	21.27	23.06	20.05	23.80	20.85	15.56								

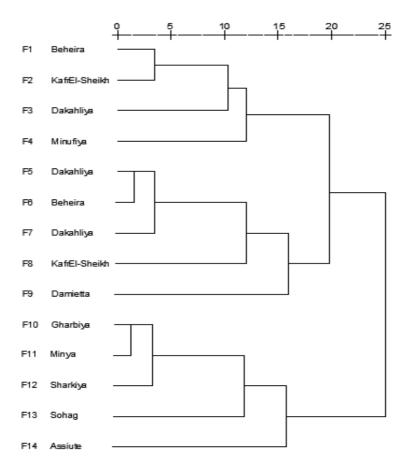
^a Mean of four replicates.

LSD for:

Trichoderma isolate (T)...... = 4.23 (P ≤ 0.05) *Fusarium* isolate (F).....=4.23 (P ≤ 0.05) Interaction $T \times F$ = 152.82 (P ≤ 0.05)



Fig(1) Phenogeram based on average linkage cluster analysis of antagonism patterns for 5 isolates of *Trichoderma* spp. against 14 isolates of *F. solani*.



Fig(2) Phenogeram based on average linkage cluster analysis of response pattern of 14 isolates of *Fusarium solani* to the antagonism of *Tricoderma* isolates.

DISCUSSION

Three isolates of *T. harzianum* and two isolates of *T. viride* were evaluated *in vivo*, to assess their antagonistic potential against *Fusarium solani* implicated in seedling damping-off of cotton. The interaction between *Trichoderma* isolates and *Fusarium solani* isolates was very highly significant source of variation for most of the tested parameters except plant heigh. This interaction implies that a single isolate of antagonist can be highly effective against an isolate of *F.solani*, but may have only minimal effects on the other isolates of *F.solani*. Bell *et al* .(1982) reported similar results when they studied the *in vitro* antagonism of *Trichoderma* spp. against six fungal plant pathogens. The findings of the present study have an important bearing on antagonism testing methods. Isolates of *Trichoderma* spp.should be

tested against as many isolates of F. solani as possible, as this will improve the chance of identifying *Trichoderma* spp. isolates effective against several isolates of F. solani. The interaction also suggests that it may be more prudent to evaluate blends of *Trichoderma* isolates for wider application against more isolates of F. solani. In this investigation, the interaction between F. solani and the Trichoderma spp. isolates was evaluated under greenhouse conditions in a soil and at temperatures favourable for the growth of both F.solani and Trichoderma spp. Under field conditions, soil nutrients and temperatures during the different periods of cotton growing season may be more favourable for *F. solani* isolates or for *Trichoderma* isolates. Thus, the results of present work are not expected to be necessarily related to the degree of biological control that may be observed in the field, but should reflect the capacities and genetic variability of the Trichoderma isolates and of the various F.solani isolates to resist antagonism (Bell et. al., 1982). The application of cluster analysis has been suggested previously for assessing similarity and/or dissimilarity in gene—for-gene host-parasite relationships (Lebeda and Jendrulek, 1987, Priestley et al., 1984). The method was also used to express exactly the genetic similarity among 48 physiological races of Bremia lactucae Regel (Lebeda and Jendrulek, 1987) and '20 isolates of Macrophomina phaseolina (Omar, 2005 and Asran-Amal et.al 2010). In this study, cluster analysis divided the tested isolates of F.solani into groups based on their response pattern to Trichoderma isolates. However, grouping the isolates was not related to their geographic origin. On the other hand, the cluster analysis of Trichoderma spp. isolates based on their biocontrol capacity showed that grouping the isolates was related to their morphological taxonomy (Rifai, 1969). Thus, it seems reasonable to conclude that morphological variations among T. harzianum and T. viride, the basis of the genus taxonomy, may provide sufficient explanation for the variation in their biocontrol capacity against F. solani isolates.

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- تثبيط فطر فيوزاريوم سولانى المسبب لمرض موت بادرات القطن بواسطة نوعين من التريكوديرما
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عزلت خمس عزلات من فطر التريكودرما تنتمى الى النوعين هيرزيانم وفيريدى من بادرات قطن مصابة بمرض موت البادرات ومن جذور نباتات بالغة مصابة بمرض عفن الجذور. قيمت قدرة العزلات على المقاومة الحيوية لعدد ١٤ عزلة من فطر فيوزاريوم سولانى المسبب لمرض موت بادرات القطن، وذلك تحت ظروف الصوبة. اظهرت نتائج تحليل التباين ان عزلات التريكودرما وعزلات فيوزاريوم سولانى والتفاعل بينهما كانت جميعا مصادر عالية المعنوية للتباين في الصفات التالية: النسبة المئوية للبادرات الميتة قبل وبعد ظهورها فوق سطح التربة والنسبة المئوية للبادرات السليمة الباقية على قيد الحياة والوزن الجاف للبادرات في حين كان التفاعل غير معنوى في حالة طول البادرة.

اظهر التحليل العنقودى لعز لات التريكودرما بناء على قدرتها التضادية انقسامها الى مجموعتين رئيسيتين إحداهما تضم عز لات تريكودرما هيرزيانم والاخرى تضم عز لتى تريكودرما فيريدى. كذلك استخدم التحليل العنقودى لتقسيم عز لات فيوز اريوم سو لانى الى مجاميع بناء على استجابتها لعز لات التريكودرما فوجد ان العز لات انقسمت الى ثلاثة مجاميع رئيسية الا أن هذه المجاميع لم ترتبط بالأصول الجغر افية للعز لات