Biological Control of Chickpea Damping-off and Root Rot Diseases G. A. H. Alareny., A. Mahran., Heidi. I. G. Abo-Elnaga., M. S. Mohamed Plant pathology Faculty of Agriculture Assiut University.

Abstract:

Twenty three fusarial isolates of 3 species e.g. Fusarium oxysporum, F. solani and F. moniliforme which isolated from diseased chickpea collected from different fields located at Assiut governorate were tested for their pathogenicity to chickpea. All these isolates caused pre and post emergence damping-off and root rot. F. oxysporum isolate (No.35) gave the highest percentage of infection. Isolates of Trichoderma harzianum, Pseudomonas fluorescence and Bacillus subtilis which isolated from chickpea rhizosphere inhibited growth of F. oxysporum, F. solani and F. moniliforme, in vitro and reduced pre and post emergence damping-off and root rot under greenhouse conditions. The selected antagonists varied in their inhibitory effect on radial growth of the tested pathogen and the reduction of the diseases.

Received on: 12/12/2013 Accepted for publication on: 22/1/2014

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Introduction:

Chickpea (Cicer arietinum L) is considered the seven crops in Egypt. The cultivated area in Egypt reached in year 2011renged from 12000 to 20000 Fadden. Most of this area was cultivated in upper Egypt especially Assiut governorate.

Damping-off and root caused by Fusarium oxysporum, F. solani and F. moniliforme, is considered sever disease in chickpea in Egypt as well as all the world(Noher et al., 2009; Sumanti et al., 2009; Khetarpal et al., 2009; Singh et al., 2010; Igbal et al., 2010 and Meki et al., 2011) .Using fungicides is consider harmful for human and environment (Meki et al., 2011) Therefore we use biological control to control diseases caused by Fusarium oxysporum, F. solani and F. moniliforme. Which consider safe and cheaper. Using Trichoderma harzianum which considered is one of the efficient biocontrol agents that commercially produced to prevent development of several soil pathogenic fungi(Kaushal and Sood , 2008; Alam et al., 2009 and Anand 2009) A different and Reddy., mechanism has been suggested as being responsible for their biocontrol activity which includes mycoparsasitism antibiosis competition for neutrant and space and serration of chitinolytic enzymes (Harman ,2000). Antifungal metabolites produced by T. harzianum, Pseudomonas fluorescence and Bacillus subtilis have been investigated for their antagonistic and fungal properties (Gowily et al., 1995; Saikia et al., 2003; Goel et al., 2002 and Mujeebur et al., 2004).

Materials and Methods:

Isolation and identification of pathogenic isolates:

Forty fungal isolates were collected from different chickpea fields located at Assiut governorate. Isolates were isolated from infected chickpea plants showing damping-off and root rot according the methods described by (Hajieghrari, 2009). Isolated fungi were purified using single spore technique and identified on basis of morphological and cultural characteristic according to (Domsh et al., 2007). And confirmed by Assiut University, Mycological Center (AUMC)

Pathogenicity tests:

Twenty three Fusarium isolates were tested for their pathogenicity using Giza3 chickpea cultivar under greenhouse conditions during (2005/2006) growing season. Pathogen inocula were prepared on barley medium as described by (Singh et al., 1997). Sterilized barley grains were inoculated with 5 mycelial blocks (0.5-mm in diameter) 5-days old. The inoculated flasks were kept under room temperature for 3 weeks. The flasks were shaken with hand every alternate day. The autoclaved pots (30cm in diameter) filled with autoclaved soil and inoculated with fungal inocula at the rate of 50g/kg of soil. Five pots were used as replicates. Ten seeds of chickpea Giza3 cultivar were surface sterilized by dipping in 3% sodium hypochlorite solution for 3 min. and washing with sterilized water and seeded in every pot containing infested soil. Pre and postemergence damping- off were recorded after 15 and 35 days, respectively. Root rot was determined after 60 days from planning using disease index.

Disease index:

Root system discoloration index was determined for external discoloration according to the index described by (Achenbach and Jennifer 1996). Readings were converted to discoloration index using

Disease severity
$$\% = \frac{0.4+1.B+2.C+3.D+4.E}{4T} \times 100$$

Where A, B, C, D and E are the number of the plants corresponding to the numerical grade 0, 1, 2, 3, and 4, respectively and (4T) is the total number of plants (T) multiplied by the maximum disease grade 4, Where, T=A+B+C+D+E

The grade of external discoloration:

The numerical grade of external root system suggested by Achenbach and Jennifer (1996) with modification were used

0= No infection

- 1=1- 25% root tissue exhibiting discoloration
- 2= 26-50% root tissue exhibiting discoloration
- 3= 51-75% root tissue exhibiting discoloration
- 4=More than 75% root tissue exhibiting discoloration or plant died.

Source of antagonists:

Isolation of Trichoderma harzianum

T. harzianum isolates were isolated from chickpea rhizosphere using method described by (Mujeebur et al., 2004). Antagonistic isolates were purified using single spore isolation and identified on basis of morphological and culture characteristic according to (Domsh et al., 2007). The identification was confirmed by Assiut University, Mycological Center (A.U.M.C).

Isolation of Bacillus subtilis.

B. subtilis isolates were isolated from chickpea rhizosphere using method described by (Hervas et al., 1998). and identified according to their morphological and biochemical activities described by (Bergey's 1978).

Source of *Pseudomonas fluores*cence isolates.

P.fluorescence (isolate No. 1) was obtained from bacterial collection of Institute of Plant Protection University of Georg-August, Gottingen, Germany. (Isolates No .2 and 3) obtained from Agriculture Research Center, Ministry of Agriculture Giza Egypt.

The antagonistic effect of *Tricho-derma* harzianum against *Fusarium* isolates.

The antagonistic capability of three fungal isolates isolated from chickpea rhizosphere was tested against the tested pathogenic fungi *in vitro*. The highly pathogenic isolates of *F.oxysporum* (Isolates No.12, 16 and 35), *F. solani* (Isolates No.4, 8 and 14) and *F. moniliforme*. (Isolates No.22, 24 and 32) were selected for this study. Petri dishes (9cm in diameter) each containing PDA medium (pH7) were inoculated with 5-mm equal disks of *T. harzianum* obtained from 4 days old cultures grown on PDA medium at 25+1°C.

A disc (5- mm in diameter) of F. oxysporum, F. solani and F. moniliforme, was inoculated at equal distance of the opposite side of Petri dish. Plates inoculated with pathogenic fungi only were used as control and five replicates were used for each test. The inoculated plates were incubated at $25+1^{\circ}C$.

Observation on antagonism and or mycoparasitism of the tested fungi recorded when the growth of the pathogenic fungi completely covered the plate surface in control.

Culture filtrate (nonvolatile) and early volatile metabolites testes:

Culture filtrates of three T. harzianum isolates showed over growth upon pathogen mycelium were tested by growing fungi in conical flasks (250 ml) each contained 100 ml of Czapek's solution liquid medium at 25±1°C.After 14 days incubation period, mycelia mats were discarded by filtration through filter papers and culture filtrates were sterilized by passing through Sitz filter. Sterilized culture filtrates were added to autoclaved Potato Dextrose Agar (PDA) medium to get 10% concentration (v/v) before dispensing medium in Petri dishes, when the temperature of the medium was about 50. Medium of PDA without addition of culture filtrates of antagonists was used as control. Disk 5-mm. in diameter, from 5days old cultures of each pathogenic fungus was transferred to the center of dishes and incubated 25±1°C. Three replicates were used for each treatment. After 6 days incubation period, linear growth of tested pathogens was recorded and percentage of inhibition was calculates as

% of growth reduction = $\frac{Growth \text{ in control- growth in treat-}}{ment \times 100}$ Growth in control

Antagonistic effect of *P. fluorescence* and *B. subtilis* isolates against the causal pathogens of *Fusarium in vitro*.

Bacterial suspension was prepared from 24h old cultures of *P. fluorescence* or *B. subtilis* and stroked at the center of Petri dish containing (PDA) medium, then two disks (5mm in diameter) from 4 days old culture of the causal pathogens were inoculated at equal distance of

the opposite side of Petri dish and three replicates were used. When the pathogenic fungi covered the plate surface of control without bacterial treatment, the distance between the edge of bacterial colonies and the fungus was measured.

Biological control of chickpea damping-off and root rot by *T. harzianum* under greenhouse conditions.

The effect of *T. harzianum*, on control of chickpea damping-off and root rot disease was carried out under greenhouse conditions during growing seasons (2007/2008) and (2008/2009). Clay loam soil with 1% organic maters was used. Completely randomize design was accomplished in greenhouse. Inocula of pathogens and *T. harzianum* were prepared as above mentioned described in pathogenicity tests.

Isolates of F. oxysporum (No.35), F. solani (No.4), F. moniliforme (No.22) and isolates of T. harzianum (1, 2 and 3) were used in this study.

Pots (30-cm in diameter) were filled with sterilized clay loam soil. *T. harzianum* inoculum was added at the same time of soil infestation with pathogens inoculum at rate of 50g/kg. Pots seeded with 10 seeds of Giza 3 chickpea cultivar. Pots were irrigated and fertilized as recommended, five pots were used as replicates and five replicates without treatment were used as control. Pre and post emergence damping-off were recorded after 15 and 35 days, respectively. Disease index of root rot determined 60 days after planting.

Biological control of chickpea damping-off and root rot by *P. fluorescence* or *B. subtilis* under greenhouse conditions.

The effect of three isolates of each *P. fluorescence* (No. 1,2 and 3)

and B. subtilis (No. 1,2and 3) on control chickpea damping-off and root rot were carried out under greenhouse conditions at Assiut University during growing seasons (2007/2008) and (2008/2009). Pots 30cm in diameter containing autoclaved soil were infested with the pathogenic isolates of F. oxysporum (No.35), F. solani (No.4) and F. moniliforme (No.22) 7days before planting at rate of 50g/kg .Inocula of pathogen were prepared as above mentioned described in pathogenicity tests. Bacterial suspension of P. fluorescence and B. subtilis was prepared by growing them in 250 ml conical flasks, each containing 100 ml of Martins medium at $27\pm2^{\circ}$ C. After 7 days of incubation period, the obtained cultures were

Statistical analysis:

All data were subjected to statistical analysis and means were compared using LSD test Gomez and Gomez, (1984).

Results:

Data in Table (1) indicated that all tested isolates were able to infect chickpea plants of Giza 3 cultivar and causing root rot and damping-off diseases. The tested fungal isolates were varied in their

centrifuged for 5 minutes at 3000 r.p.m. After centrifugation, bacterial cells were re-suspended in sterile distilled water to give concentration of 1x10⁵ cell/ml.of the antagonistic isolates of P. fluorescence and concentration of $1x10^7$ cell/ml.of B. subtilis. The bacterial suspention was added to the soil 5 days before infestation with the pathogen. Ten sterilized chickpea seeds were planted in each pot. Five replicates were used for each individual treatment. Plants watered when necessary. Pre and post emergence damping-off were recorded after 15 and 35 days respectively and root rot determined after 60 days from planting. Pots containing non- infested soil mixed with 50g/kg. of sterilized barely medium were used as control.

virulence. In general, *F. oxysporum* isolate (No.35) caused the highest percentage of pre and postemergence damping-off (60%, 40%) respectively, followed by *F. oxysporum* isolates (No.16) (64%, 30%, and 92.5%). The lowest percentage of pre and post-emergence damping-off and root rot caused by *F. oxysporum* isolate (No. 40) (30%, 2%, and 31.25%).

Table (1) Pathogenic capability of twenty three Fusarium isolates on Giza (3) chickpea (cv) under greenhouse conditions during growing season (2005/2006)

No. of isolets	Fusarium isolates	Source of		rgence ng-off%	Disease index
1801618		fungal	pre	post	%
1	F. oxysporum	El-Ghanayim	56	12	65
2	F. oxysporum	El-Kussiah	52	12	62.5
5	F. oxysporum	El-Kussiah	28	10	35
11	F. oxysporum	Sahil- Selim	32	8	40
12	F. oxysporum	Sahil- Selim	56	18	72.5
16	F. oxysporum	Al-badary	64	30	92.5
20	F. oxysporum	Sahil- Selim	24	22	43.8
21	F. oxysporum	Abnoub	58	2	60
25	F. oxysporum	Assiut	. 58	8	63.8
26	F. oxysporum	Assiut	50	12	61.3
31	F. oxysporum	Sahil- Selim	52	8	60
34	F. oxysporum	Abnoub	38	4	41.5
35	F. oxysporum	Abo-Teg	60	40	**
36	F. oxysporum	Assiut	60	20	80
38	F. oxysporum	Assiut	50	4	52.5
39	F. oxysporum	Al-Fath	32	12	42.5
40	F. oxysporum	Al-Fath	30	2	31.3
4	F. solani	El-Kussiah	70	20	90
8	F.solani	El-Kussiah	28	6	32.5
14	F. solani	Abo-Teg	56	14	70
22	F. moniliforme	Sahil- Selim	70	10	80
24	F. moniliforme	Abo-Teg	52	6	55
32	F. moniliforme	Abo-Teg	66	20	83.8
	Control		0	0	0
	L.S.D. 0.05%		5.58	4.57	8.574

** : All plants dead

Effect of culture filtrates of T. harzianum on reduction of linear growth of pathogenic Fusarium isolates

Data presented in Table (2) indicated that all culture filtrates of tested *T. harzianum* which gave over growth upon *Fusarium* mycelium decreased significantly linear growth of tested *Fusarium* isolates. The greatest inhibition of growth caused when *T. harzianum* isolate (No. 2) tested with *F. solani* isolate (No.14) and (No.4) while the lowest inhibition caused by *T. harzianum* isolate (No.3) against *F. solani* (No.4), *F. moniliforme* (No.22) and (No.32).

Table (2) Effect of culture filtrates of *T.harzianum* on linear growth of pathogenic *Fusarium* isolates

Pathogenic	No. of	Bioagints	Linear growth	% Inhibition
<u>isolates</u>	isolate		(mm)	zone (mm)
		T. harzianum 1	64	35
F. oxysporum	12	T. harzianum 2	47	42
1. oxysporum	12	T. harzianum 3	- 58	41
		control	89	0.0
		T. harzianum 1	56	33
F. oxysporum	16	T. harzianum 2	63	26
1. oxysporum	10	T. harzianum 3	69	21
		control	89	0.0
		T. harzianum 1	64	25
E amonavir	35	T. harzianum 2	56	33
F. oxysporum	33	T. harzianum 3	69	21
		control	89	0.0
		T. harzianum 1	70	19
E!	1 , [T. harzianum 2	33	56
F. solani	4	T. harzianum 3	75	14
		control	89	0.0
		T. harzianum 1	36	53
		T. harzianum 2	50	39
F. solani	8	T. harzianum 3	64	25
		control	89	0.0
		T. harzianum 1	61	28
.		T. harzianum 2	28	61
F. solani	14	T. harzianum 3	36	53
		control	89	0.0
		T. harzianum 1	61	28
77 1716		T. harzianum 2	44	45
F.moniliforme	22	T. harzianum 3	67	22
		control	89	0.0
		T. harzianum 1	56	33
F.moniliforme		T. harzianum 2	50	49
, , , , , , , , , , , , , , , , , , ,	24	T. harzianum 3	61	28
		control	89	0.0
		T. harzianum 1	47	42
		T. harzianum 2	39	50
F.moniliforme	32	T. harzianum 3	67	22
	d de la companya de l	control	89	0.0

L.S.D at 5% 6.08 4.52

Antagonistic effect between P. fluorescence isolates and the tested Fusarium isolates in vitro.

Data presented in Table (3) indicated that all *P. fluorescence* isolates inhibited the growth of the tested *Fusarium* isolates in *vitro* compared with the control. In general *P. fluorescence isolate* (No.1and 3) more effective in reducing the growth of all tested pathogens in *vitro*.

Table (3) Antagonistic effect between P. fluorescence isolates and the tested Fusarium isolates in vitro monitored as inhibition zone (mm)

Pathogenic isolates	No. of isolates	Bioagenets	Inhibition zone (mm)
E orange owner		P. fluorescence 1	21
F. oxysporum	12	P. fluorescence 2	4.9
		P. fluorescence 3	24
E companyone		P. fluorescence 1	24.2
F. oxysporum	16	P. fluorescence 2	19.2
		P. fluorescence 3	22.5
E orangeomanna		P. fluorescence 1	17.7
F. oxysporum	35	P. fluorescence 2	17.3
		P. fluorescence 3	17.3
F. solani		P. fluorescence 1	16.8
r. solani	4	P. fluorescence 2	6. 7
		P. fluorescence 3	21.3
F. solani		P. fluorescence 1	9.3
r. soluni	8	P. fluorescence 2	10.5
		P. fluorescence 3	15.3
F. solani	and the second	P. fluorescence 1	22.8
r. sount	14	P. fluorescence 2	6
		P. fluorescence 3	10. 7
F. moniliforme		P. fluorescence 1	11.2
1. montityorme	22	P. fluorescence 2	9.5
		P. fluorescence 3	15. 7
F. moniliforme		P. fluorescence 1	19.8
1. monnigorme	24	P. fluorescence 2	4.3
		P. fluorescence 3	15.2
F. moniliforme		P. fluorescence 1	16
1. monnyorme	32	P. fluorescence 2	8. 7
		P. fluorescence 3	19.2

L.S.D at 5%

2.9

Antagonistic Effect of B. subtilis isolates against the tested Fusarium isolates in vitro.

Data presented in Table (4) indicated that all *B. subtilis* isolates inhibited growth of all tested isolates of *Fusarium* in *vitro*. The inhibition zone between *B. subtilis* and the tested *Fusarium* was varied with *Fusarium* and *B. subtilis* isolates. Isolate of *B. subtilis* (No.2) with *F. oxysporum* isolates (No.16) and *F. solani* isolates (No.14) gave the highest inhibition zone (25 and 25.3 mm)

Table (4) Antagonistic Effect of B. subtilis isolates against the tested Fusarium isolates in vitro monitored as inhibition zone (mm)

Pathogenic isolates	No. of isolate	Biogenets	Inhibition zone (mm)
Farmanau		B. subtilis 1	15.5
F. oxysporum	12	B. subtilis 2	10
		B. subtilis 3	11.5
Fariananan		B. subtilis 1	14.7
F. oxysporum	16	B. subtilis 2	25
		B. subtilis 3	19.7
Forugnorum		B. subtilis 1	12.3
F. oxysporum	35	B. subtilis 2	21.8
		B. subtilis 3	18.7
F. solani		B. subtilis 1	13.7
r. solani	4	B. subtilis 2	10
		B. subtilis 3	20
F. solani		B. subtilis 1	9.2
1. Soluil	8	B. subtilis 2	9.3
		B. subtilis 3	8.5
F. solani		B. subtilis 1	17.2
r. sotani	14	B. subtilis 2	25.3
		B. subtilis 3	21.2
F. moniliforme		B. subtilis 1	10.5
1. monity of the	22	B. subtilis 2	10.2
		B. subtilis 3	2.8
F. moniliforme		B. subtilis 1	13
1. monningorme	24	B. subtilis 2	9.7
		B. subtilis 3	10
F. moniliforme		B. subtilis 1	6.8
1. monnyorme	32	B. subtilis 2	6.5
		B. subtilis 3	8.7

Effect of T. harzianum on chickpea damping-off and root rot caused by F.oxysporum, F. solani and F. moniliforme under greenhouse conditions during growing seasons (2007/2008) and (2008/2009).

Data in Table (5) indicated that in general treatment soil with three isolates of T. harzianum decreased significantly pre, post solani more than F. moniliforme. Trichoderma isolates varied in controlling chickpea damping-off and root rot with the variation of damping-off and root rot compared with control. Isolates of T. harzianum reduced damping-off and root rot caused by F.oxysporum, F. Fusarium isolates in the two seasons.

Table (5) Effect of T. harzianum on chickpea damping-off and root rot caused by F.oxysporum, F. solani and F. moniliforme under grcenhouse conditions during growing seasons (2007/2008) and (2008/2009)

	Fı	Fusarium oxysporum (No. 35)	n oxys	porum	(No. 3	5)	<i>I</i>	Fusarium solani (No. 4)	os un	lani (Vo. 4)		Fusc	ırium	monili	Fusarium moniliforme (No. 22)	No. 2	2)
	(20	(2007/2008)	(80	(20	(2008/2009)	(6)	(200	(2007/2008)	(8)	(200	(2008/2009)	(6	(200	(2007/2008)	(8)	(200	(2008/2009)	6)
Trichoderma	Damp	Damping-off		Dampi	Damping-off	rot	Dampi	Damping-off	101	Dampi	Damping-off	101	Damping-off	ig-off	101	Dampi	Damping-off	101
Isolates	%914	%120A	% root	%əл4	%1804	% root	%a14	%1204	% root	%a14	%180A	% root	%ə.14	%180d	% root	%ə14	%120A	1001 %
T. harzianum 1	52	36	85	52	34	83.8	64	8	71.3	64	10	72.5	89	-6	75.5.	99	∞	72.5
T. harzianum 2	48	30	75	44	32	73.8	09	15	73.3	99	12	75	99	8	72	65	8	70.5
T. harzianum 3	4	82	71.3	48	22	70	89	16	82.5	99	16	80.8	89	5	70	62	10	71.3
Pathogen control	09	40	*	09	40	*	70	20	06	70	20	06	70	10	80	70	10	08
Control	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
L.S.D at 5%	0	0	t	;	9	1	t C	(6	6		(,	,	t C	,	•	(
F%	7.59	67.7	7.76 2.11	7.11	.y	7.70	7:7	7.35	7.49	7.90	1.96	79.7	2.45	2.51	7.27	7.16	1.96	7.69
%L	2.54	2.31	2.86	2.15	1.88	2.80	2.39	2.40	2.56	2.12	1.90	2.65	2.44	2.21	2.57	2.16	1.96	2.68
F×T%	4.78	4.51	591	5.20	3.79	5.24	4.90	4.55	5.98	5.11	3.72	5.20	4.70	4.52	5.99	5.12	3.70	5.24
	1000	-						THE REAL PROPERTY AND ADDRESS OF THE PERSON NAMED IN COLUMN TWO IS NOT THE PERSON NAMED IN COLUMN TWO IS NAMED IN COLUMN TWO IS NAMED IN COLUMN TWO IS NAMED IN THE PERSON NAMED IN COLU		-								

**: All plants dead

Effect of P. fluorescence on the disease incidence of chickpea damping-off and root rot caused by F.oxysporum, F. solani and F. moniliforme under greenhouse conditions during growing season (2007/2008) and (2008/2009).

during growing seasons(2007-2008) and (2008-2009). Data also indicated that P. fluorescence isolate(No3) more effective in reducing Data presented in Table (6) indicated that in general treatment soil with three isolates of *P. fluorescence* which inoculated with *F.* oxysporum, F. solani and F. moniliforme decrease significantly pre, post emergence damping-off and root rot compared with control pre, post-emergence damping-off and root rot caused by F. oxysporum, and F. solani compared with isolates of P. fluorescence (No1) and P. fluorescence (No.2).

Table (6) Effect of P. fluorescence on the disease incidence of chickpea damping-off and root rot caused by F.oxysporum, F. solani and F. moniliforme under greenhouse conditions during growing season (2007/2008) and (2008/2009)

American de la companya de la compa	Fu	sariun	ls/xx0 1	Fusarium oxysporum(No. 35)	(No. 3	5)		Fusar	Fusarium solani (No. 4)	lani (1	Vo. 4)		Fus	arium	moni	liform	Fusarium moniliforme(No. 22)	22)
	(20	(2007/2008)	(80	(20	(2008/2009)	(60	(20	(2002/2008)	(80	(20	(2008/2009)	(60	(20	(2007/2008)	(80	(20)	2008/2009	(60
P. fluorescence	Dampi	Damping-off	1	Dam _o	Damping- off	i	Damping- off	ping-	1	Damp off	Damping- off	'	Damping- off	oing- f		Dam 0	Damping- off	
_	%ə14	%1204	or % 101	%ə14	%1204	or % lor	%əz4	%1204	% ro roi	Pre%	%120 4	or % ior	Pre%	%1204	or % 101	Pre%	%1204	% ro roi
P. fluorescence 1	52	38	85	52	34	84.5	99	17	81.8	72	10	80.3	62	9	65	99	2	66.5
P. fluorescence 2	58	30	85	50	30	80	62	16	76.5	09	12	70.3	99	8	62.5	99	10	62.8
P. fluorescence 3	46	36	71.3	58	10	99	58	10	99	99	8	62.5	28	9	62.5	64	10	62.8
Pathogen control	09	40	*	09	40	*	70	20	06	70	20	06	70	10	80	70	10	08
Control	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
L.S.D at 5%																		
F%	1.69	1.55	2.47	1.55 2.47 1.89 1.39	1.39	2.39	1.64	1.65	2.41	1.87	1.36	2.39	1.64	1.65	2.41	1.87	1.36	2.39
T%	2.27	2.28	2.57	2.28 2.57 249 1.89 2.52	1.89	2.52	2.17	2.18	2.55	2.47	1.79	2.52	2.17	2.18	2.55	2.47	1.79	2.52
F×T%	4.43		5.19	4.49 5.19 4.96 3.68 5.05	3.68	5.05	4.33	4.36	4.36 5.09	4.94	3.58	5.05	4.33	4.36	5.09	4.94	3.58	5.05
L - L - T - T - T - X - X	1.2.1									The state of the s						and the same of th		

** : All plants dead

Effect of B.subtilis on disease incidence of chickpea damping-off and root rot caused by F.oxysporum, F. solani and F. moniliforme under greenhouse conditions during growing seasons (2007/2008) and (2008/2009).

damping-off and root rot compared with control. Data also indicated that in general B.subtilis isolate (No. 3) more effective in reducing root rot caused by F. oxysporum, F. solani and F. moniliforme compared with B.subtilis isolate (No. 1 and 2). Data presented in Table (7) indicated that treatment soil with three isolates of B. subtilis decrease significantly pre, post- emergence

Table (7) Effect of B.subtilis on disease incidence of chickpea damping-off and root rot caused by F.oxysporum, F. solani and F. *moniliforme* under greenhouse conditions during growing seasons (2007/2008) and (2008/2009)

Subtilis Damping-off 5 Continuous		F	Fusarium oxysporum (No. 35)	ls/xo 1	orum (No. 35			Fusar	Fusarium solani (No. 4)	lani (N	5.4)		Fu	sarium	monil	iforme	Fusarium moniliforme (No. 22)	
Damping-off Sign Damping-off Sign Damping-off Sign Damping-off Sign Damping-off Sign	, ,	(20	02//200	8)	(20	08/200	9)	(2)	007/200)8)	(20	08/200	(6	(20	07/200	(8	(20	08/200	(
Property Property	B.subtilis	Damp	ing-off	10	Dampi	ng-off	1 0	Dampi	ing-off	10	Damp	ing-off	10	Damp	mg-off	10	Damp	ing-off	10
46 28 72.5 46 32 75 60 14 72.5 65 12 75.3 67 6 73.8 65 8 54 24 75 42 32 72.5 62 10 71.3 56 16 72.3 65 5 70 66 5 70 66 5 63.3 55 6 5 70 66 5 63.3 55 6 5 66 5 63.3 55 6 5 66 5 63.3 65 5 70 60 70 <	isolates	Pre%	Post %		Pre%	120A %		Pre%			Pre%			Pre%			Pre%	%180A	
54 24 75 42 32 72.5 62 10 71.3 56 16 72.3 65 5 70 66 5 70 66 5 63.3 55 6 5 70 66 5 63.3 55 6 5 70 66 5 63.3 55 6 3 6 6 7 6 7 6 7 6 7 7 6 7 7 7 7 7 7 7 7 7 7	B. subtilis 1	46	28	72.5	46	32	75	09	14	72.5	65	12	75.3	<i>L</i> 9	9	73.8	65	8	73.8
38 24 61.3 36 30 63.8 60 12 71.3 68 12 70 60 5 63.3 55 6 60 40 ** 70 20 90 70 20 90 70 10 80 70 10 0.0	B. subtilis 2	54	24	75	42	32	72.5	62	10	71.3	99	16	72.3	65	5	70	99	5	70.3
60 40 ** 60 40 ** 70 20 90 70 20 90 70 10 80 70 10 10 80	B. subtilis 3	38	24	61.3	36	30	63.8	09	12	71.3	89	12	70	09	~	63.3	55	9	09
0.0 0.0 <th>Pathogen control</th> <td>09</td> <td>40</td> <td>* *</td> <td>09</td> <td>40</td> <td>* *</td> <td>70</td> <td>20</td> <td>06</td> <td>70</td> <td>20</td> <td>06</td> <td>70</td> <td>10</td> <td>80</td> <td>70</td> <td>10</td> <td>80</td>	Pathogen control	09	40	* *	09	40	* *	70	20	06	70	20	06	70	10	80	70	10	80
1.60 1.51 2.87 2.01 1.59 2.09 1.65 2.44 2.25 1.66 2.59 1.94 1.91 2.48 1.97 1.57 2.21 2.24 2.27 2.39 2.67 1.99 2.72 2.37 2.39 2.95 2.57 1.90 4.41 4.45 5.39 4.99 3.88 5.25 4.46 5.18 4.99 3.78 5.15 4.53 4.57 5.39 4.94 3.80	Control	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
1.60 1.51 2.87 2.01 1.59 2.59 2.09 1.65 2.44 2.25 1.66 2.59 1.94 1.91 2.48 1.97 1.57 2.21 2.24 2.77 2.26 1.99 2.77 2.37 2.39 2.95 2.57 1.90 4.41 4.45 5.39 4.99 3.88 5.25 4.53 4.46 5.18 4.99 3.78 5.15 4.57 5.39 4.94 3.80	L.S.D at 5%																		
2.21 2.24 2.77 269 1.99 2.72 2.47 2.28 2.59 2.67 1.99 2.72 2.37 2.39 2.95 2.57 1.90 4.41 4.45 5.39 4.99 3.88 5.25 4.53 4.46 5.18 4.99 3.78 5.15 4.53 4.57 5.39 4.94 3.80	F%	1.60	1.51	2.87	2.01	1.59	2.59	2.09	1.65	2.44	2.25	1.66	2.59	1.94	1.91	2.48	1.97	1.57	2.49
4.41 4.45 5.39 4.99 3.88 5.25 4.53 4.46 5.18 4.99 3.78 5.15 4.53 4.57 5.39 4.94 3.80	Τ%	2.21		2.77	269	1.99	2.72	2.47	2.28	2.59	2.67	1.99	2.72	2.37	2.39	2.95	2.57	1.90	2.62
	F×T%	4.41	4.45	5.39		3.88	5.25	4.53	4.46	5.18	4.99	3.78	5.15	4.53	4.57	5.39	4.94	3.80	5.15

**: All plants dead

Discussion:

Damping-off and root-rot caused by F. oxysporum, F. solani and F. moniliforme were important diseases of chickpea plants. Twenty three Fusarium isolates were tested on chickpea Giza 3 (cv). All these isolates caused pre and post emergence damping-off and root- rot. F. oxysporum gave the highest percentage of infection. According to the available literature these pathogens were isolated from chickpea (Singh et al., 1997; Soregaon and Ravikumar 2010 and Meki et al., 2011) Three isolates of T. harzianum showed over growth upon tested pathogenic Fusarium isolates and their culture filtrates inhibited the growth of the tested Fusarium isolates in vitro .These results are in accordance with those obtained by (El-Nashar et al., 2001; Atef- Najwa 2008; Hajieghrari et al., 2008; Waheed and Khilare 2010 and Gajera et al., 2012). They reported T. harzianum treatment reduce the mycelia growth of the pathogenic fungi due to the rapid growth of T. harzianum which colonized medium surface and substrate or reduced hyphal coils over the interaction zone or produced chitins and β -(1-3)- gluconase .Soil treatment with T. harzianum reduced chickpea damping-off and root- rot diseases under greenhouse conditions such results in accordance with those reported by (Nawar-Lubna 2007; Kucuk et al., 2007; Atef-Najwa 2008; Haggag-Wafaa et al., 2011 and EL-Bramawy and EL-Sarag (2012). In vitro, isolates of P. fluorescence and B. subtilis inhibited linear growth of tested Fusarium isolates. The antagonists varied in their inhibitory effect on linear growth of the tested pathogen .Under greenhouse condition addition of P. fluorescence and B. subtilis to soil reduced incidence

of pre and post emergence dampingoff and root- rot of chickpea .The reduction of the diseases varied according to the pathogen and antagonistic isolates which used. These results agree with (Gowily et al., 1995; Inam et al., 2009; Priyanka, 2010; and Raiput et al., 2010). Pseudomonas fluorescence produce siderphore (iron chelating compounds) which strongly reacted with free Fe and inhibit the fungal mycelia growth and spore germination produce antibiotic substances produces cyanic acid (Abo-Alnaga- Heidi, 2002). Antifungal metabolites produced by bacteria like Pseudomonas fluorescence and Bacillus subtilis have been investigated for their antagonistic and fungal properties by (Gowily et al., 1995; Goel et al., 2002; Saikia et al., 2003; and Mujeebur et al., 2004).

References:

Abo- Alnaga- Heidi, I.G. (2002). Studies on root rot Disease of Soybean (Glycine Max.) in Upper Egypt. Ph.D. These Department Plant Pathology Faculty of Agriculture Assiut University Egypt pp. 145.

Achenbach, L.A. and P. Jennifer (1996). Use of RAPD markers as a diagnostic tool for the Identification of *Fusarium solani* isolates that cause soybean sudden death syndrome-Plant Disease, 80:1228-1232.

Alam, S., S. Zacharia, S. Simon, V. Choudhary, K. Sherzama and M. Naseem, (2009). Effect of *Trichoderma harzianum rifai* against chickpea wilt caused by *Fusarium oxysporum f. sp. ciceri* International Journal of Agriculture Environment Biotechnology. 2(4): 424-428.

Anand, S. and J. Reddy (2009). Biocontrol potential of *Tricho-*

- derma sp. against plant pathogens. International Journal of Agriculture Sciences. 1(2): 30-39. 20 ref.
- Atef-Nagwa. (2008). Bacillus subtilits and Trichoderma harzianum as wheat inoculants for Biocontrol of Rhizoctonia solani. Australian Journal of Basic and Applied Sciences 2(4):1411-1417.
- Bergey's, D.H. (1978). Bergerys Manual of Determinative Bacteriology. Seventh ed. Williams and Wilkins Co. Baltimore.1247 pp.
- Domsh, K.H., W.Gams, and T.H.Anderson (2007). Compendium of soil fungi. 2nd Edition IHW Verlage. Eching Germany. 672 pp.
- EL-Bramawy, M.A.S., and E.E.EL-Sarag (2012). Enhancement of seed yield and its components in some promising sesame lines using antagonism of *Trichoderma* spp. Against soil-borne fungal diseases.Int. J. Forest, Soil and Erosion 3: 148-154.
- EL-Nashar-Faten, Mehreshan. El-Mokadem, T.H.Abd-El-Moity, and H.A.M.Ammar (2001). Biological control of root-rot disease of wheat Egyptian Journal of Agricultural Research. 79(1).
- Gajera H.P., R. P. Bambharolia, S. V. Patel, T. J. Khatrani and B. A. Goalkiya (2012). Antagonism of *Trichoderma* spp. against *Macrophomina phaseolina*: Evaluation of coiling and cell wall degrading enzymatic activities J. Plant Pathol.Microb. 3:149.
- Goel, A. K., S. S. Sindhu, K. R. Dadarwal (2002). Stimulation of nodulation and plant growth of chickpea (Cicer arietinum L.)

- by *Pseudomonas spp.* antagonistic to fungal pathogens. Biology and Fertility of Soils. 36(6): 391-396.
- Gomez, K. A. and A. A. Gomez (1984). Statically Procedures for Agricultural Research. 2 nd Edn., John Wily and Sons Inc., New York, USA. 680.
- Gowily, A. M., G. L. Soliman and Abd El-Ghany(1995). Biological control of chickpea root rot caused by *Fusarium solani*. Annals of Agriculture Science, Moshtohor 33:1307-1315.
- Haggag-Wafaa. M., and M.S. Mohamed (2011). Biodiversity ,Biological and Molecular Investigations of Biocontrol by the Genus Hypocrea *Trichoderma* spp. European Journal of Scientific Research 65(2):281-292.
- Hajieghrari, B., M. Torabi-Giglou, M.R. Mohammadi, and Davari (2008). Biological potential of some Iranian *Trichoderma* isolates in the control of soil borne plant. African Journal of Biotechnology 7(8): 967-972.
- Hajieghrari, B. (2009). Wheat crown and root rotting fungi in Moghan area, Northwest of Iran. African Journal of Biotechnology 7(22): 6214-6219.
- Harman ,G. E. (2000). The myths and dogmas of biocontrol change in perception derived from research on *Trichoderma harzianum* strain T-22. Plant Dis., 84:377-393.
- Hervas, A., B. Landa, L. E. Datnoff and R. M. Jimenez-Diaz (1998)
 Effects of commercial and indigenous microorganisms of Fusarium wilt development in chickpea. Biological Control. 13(3):166-176.

- Inam, M. H., S. El-Hassan, S. Gowen and N. Javed (2009). Effects of two rhizobacterial isolates and neem cake application on control of chickpea wilt caused by *Fusarium oxysporum f.sp. ciceri*. Arab Journal of Plant Protection. 27(1):103-110.
- Iqbal, S. M., A. Ghafoor, A. Bakhsh, I. Ahmad and A. Sher (2010). Identification of resistant sources for multiple disease resistance in chickpea. Pakistan Journal of Phytopathology. 22(2): 89-94.
- Kaushal, R. P. and R. Sood (2008). Management of root rot in chickpea. Journal of Food Legumes. 21 (3):178-181.
- Khetarpal, R. K., S. Khetarpal, P. Kumar, M. Pal, D. Chand and S. Lata (2009). Effect of association of *Fusarium solani* and *F. moniliforme* on photosynthetic characteristics in chickpea (*Cicer arietinum L.*) Indian Phytopathology. 62(4): 484-487.
- Kucuk, C., M.Kivance, E.Kinaci, and G.Kinaci (2007). Biological efficacy of *Trichoderma harzianum* isolate to control some fungal pathogens of wheat (*Triticum aestivum*) in Turkey. Biologia 62: 283-286.
- Meki, S., S. Ahmed, and P.K Sakhuja (2011). Control of chickpea wilt Fusarium oxysporum f.sp. ciceri using Trichoderma spp. In Ethiopia. Archives of Phytopathology and Plant Protection. 44(5): 432-440.
- Mujeebur, R., K. Shahana, M. Khan and F. A. Mohiddin (2004). Biological Control of Fusarium wilt of chickpea through seed treatment with commercial formulation of Trichoderma har-

- zianum and Pseudomonas fluorescens. Phytopath Mediterr. 20-25.
- Nawar-Lubna, S.,(2007). Pathological and rhizospherical studies on root-rot disease of squash in Saudi Arabia and its control. African Journal of Biotechnology 6(3): 219- 226.
- Noher, A. M., M. Y. Abdalla, M. A. Abou Attia and, N. M. Abou Zeid (2009). Pathological and genetic variability among isolates of *Fusarium oxysporum f.* sp. ciceri causing wilt of chickpea in Egypt. Association Francaise de Protection des Planets, 9eme conference international sure les maladies des planets, Tours, France, 8 et 9 December. 143-151.
- Priyanka, J. M. (2010). Assessment of the potential of plant growth promoting *rhizobacteria Bacillus sp.*in enhancing symbiotic efficiency in chickpea with Mesorhizobium .Biochemical and Cellular Archives. 10(1): 57-62.
- Rajput, V.A., S.A. Konde, and M.R. Thakur, (2010). Evaluation of biogagents against chickpea wilt complex. Journal of Soils and Crops. 20(1): 155-158.
- Saikia, R., T. Singh, R. Kumar, J. Srivastava, A. K. Srivastava, K. Singh and D. K. Arora, (2003) Role of salicylic acid in systemic resistance induced by *Pseudomonas fluorescens* against *Fusarium oxysporum f. sp. ciceri* in chickpea. Microbiological Research. 158(3): 203-213.
- Singh, R. S., D. Singh, H. V. Singh and D. Singh(1997). Effect of fungal antagonists on the growth of chickpea plants and

- wilt caused by Fusarium oxysporum f. cp. ciceri. Plant Disease Research .12:103-107.
- Singh, R. K., Abul Hasan and R. G. Chaudhary2010). Variability in *Fusarium oxysporum f. sp. ciceri* causing vascular wilt in chickpea. Archives of Phytopathology and Plant Protection. 43 (10): 987-995.
- Soregaon, C. D. and R. L. Ravikumar, (2010). Marker assisted characterization of wilt resistance in productive chickpea genotypes. Electronic Journal of Plant Breeding. 1(4):1159-1163.
- Sumanti G., D. Chakraborti, R. K. Rangi, D. Basu, S. Das (2009). A molecular insight into the early events of chickpea (*Ccer arietinum L*) and *Fusarium oxysporum f. sp. ciceri* (race 1) interaction through cDNA-AFLP analysis. Phytopathology. 99 (11): 1245-1257.
- Waheed, M. A., and V.C.Khilare (2010). Biological control of mulberry root rot incited by species of *Fusarium*. Indian Journal of Sericulture 49(2): 218-219.

المكافحه البيولوجيه لمرض سقوط البادرات وعفن الجذور فى الحمص جمال على العريني ، عمرو مهران ، هايدي ابراهيم ابوالنجا ، محمد سامى محمد أمراض النبات – كلية الزراعة – جامعة أسيوط

الملخص:

تم عزل ثلاثة وعشرون عزله من فطر فيوزاريوم تشمل ثلاثة أنواع المصابه (Coxysporum, F. solani and F. moniliforme في أماكن مختلفة بمحافظه أسيوط قد سببت أمراض سقوط البادرات المفاجئ وعفن الجذور في أماكن مختلفة بمحافظه أسيوط قد سببت أمراض سقوط البادرات المفاجئ وعفن الجذور في الحمص وكانت عزله الفطر معربي المعمل والصوبه ووجد ان فطر Trichoderma harzianum أو راشحه لهم القدره على تثبيط النمو الطولي لجميع العزلات الفطريه الممرضه المختبره وكذلك بكتريا Pseudomonas النمو الطولي لجميع العزلات الفطريه المختبره ولوحظ في معظم الحالات أن معامله التربة باستخدام fluorescence and Bacillus subtilis T. harzianum, P. fluorescence and البادرات قبل وبعد الظهور وحدوث عفن الجذور وجميع الفطريات ألممرضه ألمختبره بالمقارنة بالكنترول تحت ظروف الصوبه خلال تجربتين عامي ۱۹۰۷ واختلفت هذه العزلات في قدرتها على المكافحه باختلاف العزلات الممرضة.