EFFICIENCY OF SOME BACTERIAL STRAINS FOR CONTROLLING LIMB ROT DISEASE OF PEANUT GROWN IN A SANDY SOIL

MOUSSA, LOBNA A. 1, EBTSAM M. MORSY 1, ABEER A. SHALTOUT² AND SOHEIR S. FAHMY¹

- 1. Soils, Water and Environment Res. Inst., ARC, Giza, Egypt.
- 2. Plant Pathology Res. Inst., ARC, Giza, Egypt.

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Abstract

Four biocontrol agents, Bacillus subtilis, Bacillus Pseudomonas synxantha amyloliquefaciens, Brevibacterium otitidis were tested individually and/or in combined mixtures for suppression of Rhizoctonia solani on peanut. All tested bacteria moderately inhibited the growth of R. solani. B. amyloliquefaciens was the best for inhibiting growth. In a field experiment, combined mixture of the four tested bacteria completely suppressed incidence disease also combined mixtures of B. amyloliquefaciens and Brevibacterium otitidis or Pseudomonas synxantha and Brevibacterium otitidis gave same result. Dehydrogenase and nitrogenase activities, root nodulation and peanut biomass yield were also determined. All tested strains increased plant parameters, however B. amyloliquefaciens and B. subtilis were the most effective.

INTRODUCTION

Peanut (*Arachis hypogaea* L.) growers must protect their crop from foliar and soil-born diseases to optimize profits and maintain high yields (Franke *et al.*, 1999).

One of the most common soil-born pathogen is Rhizoctonia solani AG-4. It induced many diseases in peanut including seed decay, pre- and post- emergence damping-off, hypocotyls and root necrosis, peg rot, limb and root rot. Bell and Summer (1984). Control of Rhizoctonia limb rot in peanut is based on an integrated management approach using crop rotation, proper fertilization, irrigation management, chemical and biological control. In response to environmental and health concerns about extended use of pesticides, there is a growing interest for finding alternative control approaches for use in integrated pest management (IPM) strategies for crop diseases. Biological control of plant diseases has been suggested as alternative control methods (Cook, 1993). Bacillus subtilis and Fluorescent pseudomonas, isolated from the rhizosphere, are known to suppress several soil-born diseases caused by phytopathoenic fungi and promote plant growth (Mosa et al., 2003). Yoshida et al. (2001) demonstrated capability of B. amyloliquefaciens to inhibit mulberry anthracnose and secretion of several antifungal compounds. Brevibacterium otitidis has been joined as a bacterial biocontrol agent to antagonize two of powerful soil-born pathogenic fungi (Macrophoamina phaseolina and Sclerotium rolfsii) causing charcoal

rot and southern blight of many crops (Moussa *et al.,* 2006). Morsy (2005) succeeded to prove the role of *Pseudomonas synxantha* as antagonistic bacterium against *R. solani* and *F. solani* in the rhizosphere of tomato plants.

MATERIALS AND METHODS

Materials

- Peanut (Arachis hypogaea L.) seeds, cv. Giza 6 were provided from Horticulture Research Institute, ARC, Giza, Egypt.
- Sandy soil at Ismailia Agric. Res. Station, ARC, was used during summer season under sprinkler irrigation. Mechanical and chemical characteristics of the experimental soil are shown in Table1.

Table 1. Some mechanical and chemical characteristics of the experimental soil

A) Mechanical characteristics						
Particle size distribution (%)						CaCO₃
Coarse sand	Fine sand	Silt	Clay	Textural class	(%)	(%)
31.82	61.61	1.22	5.35	Sandy	0.44	1.42

EC		Sol	uble cati	ons (me	s (meq/l) Soluble anions (meq/l)				/D	
dS/m	pН	Ca ²⁺	Mg ²⁺	Na ⁺	K ⁺	CO ₃ =	HCO ₃	CI.	SO ₄ =	SAR

- Rhizolex T (tolcofs-methyl thiram) was used as chemical fungicide.
- Biocontrol agents: Four of the potential biocontrol agents against limb rot and wilt disease were examined for their interaction with the pathogenic fungus *R. solani*.
 Two of them were isolated from Egyptian soil: *B. subtilis* and *Pseudomonas synxantha* (Morsey, 2005), *B. amyloliquefaciens* and *Brevibacterium otitidis* (Moussa *et al.*, 2006).
- Pathogenic fungus: Rhizoctonia solani was isolated from the experimental soil (Ismailia Agric. Res. Station, ARC). It was identified on the basis of cultural and microscopic characteristics. Pathogencity of the strain toward peanut was estimated (Sneh et al., 1991).

Methods

Antibiosis of bioagents against phytopathogenic fungus

The antagonistic effect of the tested four biocontrol agents against *R. solani* was examined using agar plate inhibition zone technique. An explorative trial was made to study whether the biocontrol agents have the ability to grow and interact in the medium of the tested fungus. All the tested biocontrol agents were succeeded to grow

well on potato dextrose agar (PDA). A disc (5 mm. θ) of *R. solani* was inoculated at periphery of the plate against a steak of each tested bacterial strains at other periphery, which were firstly inoculated 24 hr before inoculation of the pathogenic fungus. Inoculated plates were incubated at 25 °C for 3 - 5 days. The growth and reduction in mycelial growth of the pathogenic fungus were calculated using the following equation according to Fokemma (1973):

Reduction% = $R_1 - R_2/R_1 \times 100$ where: R_1 = liner growth of control R_2 = liner growth of treatment

Field experiment

A field experiment was conducted in an attempt to practices controlling of limbrot disease biologically. The used experimental soil has a history of limb-rot disease causing by R.solani. The experimental layout comprised peanut (Arachis hypogaea L., Giza 6), Bradyrhizobium sp. (R617) added by seed coating, four biocontrol agents (B.subtilis, B. amyloliquefaciens, Pseudomonas synxantha and Brevibacterium otitidis), beside the fungicide (Rhizolex-T). To avoid the possibility of earlier negative interaction between the introduced Bradyrhizobium and biocontrol agents, they were alternately applied by two different ways, while biocontrol agents were applied as soil drench individually or in combined mixtures at rates of 1:1 after 15, 30 and 45 days of sowing. The fungicide (Rhizolex-T) was used with recommended dose (3 g/kg seeds). The NPK fertilizers were incorporated into soil as recommended dose 200 kg/fed of superphosphate (15 % P_2O_5), 45 N unit/fed as ammonium sulphate and 50 kg K/fed as potassium sulphate (48 % K_2O). Seeds were drilled in rows of 30 cm apart. The experimental design was a complete randomized block with three replicates.

After peanut maturity, shoots and roots were determined as dry weights. Pods and 100 seed weights as well as the shelling percent were recorded. Data obtained were subjected to the statistical analysis according to Snedecor and Cochran (1989), where mean values were compared using L.S.D. at 5 % level.

Biological determinations

Fungal count

The isolated fungi were counted and identified in the Mycology Res. and Plant Disease Survey Dept., Plant Pathology Res. Instit., ARC. The frequency of each individual fungal group was calculated as follows:

Fungi (%) = (No. colonies of each group/plate)/(No. total appeared fungal colonies) \times 100 (Martin, 1950).

Nitrogenase activity of peanut root nodules was determined by assaying acetylene reduction as described by Dilowarth (1970). In addition, dehydrogenase activity in the rhizosphere was assayed according to Thalmann (1967).

RESULTS AND DISCUSSION

Antagonistic efficacy of the tested bacteria

Four bacterial strains were evaluated for their potentiality to antagonize growth of the pathogenic fungus *Rhizoctonia solani*. Data presented in Table 2 show that all the tested bacterial strains succeeded to reduce radial growth of the pathogenic fungus. The most effective one was *Bacillus amyloliquefaciens* which gave 48.9 % reduction followed by *B. subtilis* (47.8 %). *Brevibacterium otitidis* and *Pseudomonas synxantha* resulted in reduction of the growth being 37.8 and 35.6 %, respectively in comparison with control. These findings are in agreement with those obtained by Yoshida *et al.* (2001) who attributed potentiality of *Bacillus amyloliquefaciens* St-RC-2 against *Colletotrichum dematium* (which cause mulberry anthracnose) to the antifungal compounds produced in the culture filtrate. Potentiality of *Ps. synxantha* and *Brevibacterium otitidis* to retard the fungal growth could be attributed to their antibioses towards the tested fungus.

Table 2. Bacterial bioagents effect on Rhizoctonia solani mycelia growth (in vitro).

Bacteria	Mean radial	growth (cm)
	Radial growth	Reduction %
B. amyloliquefaciens	4.6	48.9
Pseudomonas snyxantha	5.8	35.6
B. subtilis	4.7	47.8
Brevibacterium otitidis	5.6	37.8
Control	9.0	0.0

Efficiency of the bacteria on limb rot incidence on peanut

Data presented in Figure 1 show that all the tested bacterial strains as well as the fungicide Rizolex T succeeded to suppress the incidence disease on peanut plants. The most effective bacteria were *B. subtilis* and *B. amyloliquefaciens* which caused 84.1 % reduction in disease incidence followed by *Brevibacterium otitidis* (74.3 % reduction) and *Ps. synxantha* (56.6 % reduction).

Data also cleared that using a combined mixture of the four strains or mixtures of *Brevibacterium otitidis* with either *B. amyloliquefaciens* or *Ps. synxantha* completely suppressed the incidence disease on peanut plants (100 % reduction). These results are in harmony with those obtained by Yu and Sinclair (1996). On the other hand, Jetiyanon *et al.* (2003) indicated that *B. amyloliquefaciens* induced systemic resistance (ISR) in several plants against different plant pathogens. These (ISR) include lignification, peroxidase and superoxide dismutase production. Strains of *B. mojavensis*, *B. amyloliquefaciens*, *B. subtilis* and *Brevibacterium halotolerans* are antagonistic to the fungus *Fusarium moniliforme* (Bacon and Hinton 2002).

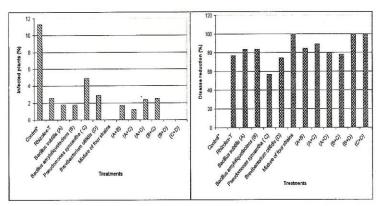


Fig. 1. Effect of the four tested bacteria on incidence of peanut limb rot caused by Rhizoctonia solani

Tested bacterial effect on beneficial fungi in the peanut rhizosphere

Effect of the tested bacteria on the frequency percentage of beneficial fungi such as *Trichoderma harzianum* and *T. hamatum* in the rhizosphere of peanut through growth stages was investigated. Data in Table 3 indicate that frequency of the two beneficial fungi was increased with adding the four tested bacteria compared to the control and use of (Rhizolex-T).

Also, it was shown that the most frequencies of T. harzianum were recorded due to B. subtilis and B. amyloliquefaciens. However, T. hamatum frequencies varied with the tested bacteria and growth stages. Also, the mixture of the four tested bacteria dramatically increased the frequency percentages of the two fungi compared with using either one of them. This was also shown when adding mixtures of Brevibacterium otitidis with each of B. amyloliquefaciens and Ps. Synxantha. Di Pietro (1995) mentioned that T. harzianum is known to produce extracellular cell wall degrading enzymes such as chitinase, β -1-3- glucanases and cellulases which are important features of mycoparasites for the colonization of their host fungi.

The obtained results of this test suggest that *Trichoderma* species found in peanut rhizosphere could be also having an antagonistic effect against the tested pathogen.

Table 3. Tested bacteria effect on beneficial fungi frequency in peanut rhizosphere.

		otal fun	aal		Frequer	ncy of be	eneficial	fungi 9	6	
Treatments	N 1000 113	Counts (cfu x 10 ³)			Trichoderma harzianum			Trichoderma hamatum		
	V	F	M	٧	F	M	٧	F	M	
Control										
Rhizolex -T	25	28	22	15	18	20	12	10	12	
Bacillus subtilis	5	10	10	0	10	10	0	10	20	
Bacillus amyloliquefaciens	12	15	10	25	26	30	17	20	10	
Pseudomonas synxantha	14	17	13	24	24	23	24	14	15	
Brevibacterium otitidis	12	16	11	16	25	27	16	18	18	
Mixture of four strains	13	15	10	15	20	20	15	18	10	
B. subtilis + B. amyloliquefaciens	10	12	9	30	33	33	20	16	22	
B. subtilis + Ps. synxantha.	14	16	10	28	30	30	14	18	20	
B. subtilis + Br. otitidis	8	10	10	25	25	30	12	20	20	
B. amyloliquefaciens + Ps.	10	13	11	20	23	27	10	15	18	
synxantha.	13	15	12	15	20	25	15	20	20	
B. amyloliquefaciens + Br. otitidis	13	15	12	15	20	25	15	26	25	
Ps. synxantha.+ Br. otitidis	12	14	10	25	28	30	16	21	20	
Before sowing		20		a va utes	15		V/V/V/See	10		

V= Vegetative stage

F= Flowering stage

M= Maturity

Effect of different bacterial strains on enzyme activity and dry weight of nodules in soil infected with *R. solani:*

Table 4 shows that all treatments inoculated with antagonistic bacteria remarkably recorded increases in dehydrogenase activity rather than the control (uninoculated treatment). Soil inoculated with *B. amyloliquefaciens*, *B. subtilis* and mixture of *B. subtilis* and *Ps. synxantha* gave the highest enzyme activity.

Table 4. Effect of different bacterial strains on enzyme activity and dry weight of nodules in soil infected with *R. solani*

	Enzyme activity							Decominh of and to		
Treatments	Dehydrogenase ug TPF/g dry soil/day			Nitrogenase u/ C₂H₄/plant/hr			Dry weight of nodules g/plant			
	45 *d	75 *d	100 *d	45 *d	75 *d	100 *d	45 *d	75 *d	100 *d	
Control	23.3	35.0	46.2	16.8	17.0	31.2	0.13	0.33	0.700	
Rizolex T	27.4	31.3	54.0	34.6	47.4	75.9	0.15	0.23	0.730	
Bacillus subtilis	66.9	53.5	56.4	63.4	67.9	208.3	0.17	0.52	0.800	
Bacillus amyloliquefaciens	59.1	51.2	57.1	78.6	164.4	433.2	0.16	0.49	0.832	
Pseudomonas synxantha	46.3	46.3	54.5	64.8	65.4	165.6	0.19	0.25	0.860	
Brevibacterium otitidis	20.3	55.4	54.2	113.5	141.2	175.9	0.18	0.45	0.763	
Mixture of four strains	22.2	49.5	48.9	87.2	213.6	227.0	0.17	0.84	0.900	
B. subtilis + B. amyloliquefaciens	36.9	42.8	64.1	103.6	137.2	113.4	0.18	0.47	0.780	
B. subtilis + Ps. synxantha	54.7	50.8	76.3	52.0	117.1	281.0	0.17	0.51	0.950	
B. subtilis + Brevi. otitidis	41.4	37.0	77.4	121.5	130.3	294.0	0.19	0.30	0.995	
B. amyloliquefaciens + Ps. Synxantha	28.8	57.0	49.3	46.7	48.4	200.9	0.16	0.37	0.874	
B. amyloliquefaciens + Brevl. otitidis	34.0	43.8	69.6	111.3	174.8	337.6	0.18	0.42	0.950	
Ps. Synxantha + Brevi. otitidis	38.2	45.3	61.4	79.9	102.7	344.0	0.20	0.45	0.880	

* After 45 days, 75 days and 100 days

LSD at 5% for:	Dehydrogenase	Nitrogenase	Dry weight of nodules
Treatment	17.3	50.8	0.16
Periods	9.37	24.8	0.11
TxP	30.1	88.1	0.27

The increase in DHA activity as a result of soil inoculation with antagonistic bacteria may be due to the synergistic effect between the native soil micro organisms and the introduced ones.

Bacterial inoculation increased nodulation as well as N_2 -ase activity but B. amyloliquefaciens, B. B. amyloliquefaciens + Brevi. otitidis and B. subtilis + Brevi. otitidis gave higher N_2 -ase activity and high dry weight of nodules than the other treatments. Bai et al. (2002) found that, soybean plants infested by three Bacillus strains resulted in enhancing nodulation and subsequent nitrogen fixation.

Influence of repeated inoculation with different bacterial strains on growth and yield

Data presented in Table 5 show that introduced antagonistic bacteria significantly increased the dry weights of peanut shoot and root. The increase percentages were 92.7 and 87.3 for shoot while they were 100 and 85.7 for root with *B. amyloliquefaciens* and *B. subtilis*, respectively. The combined mixtures effect of the four strains on increasing the dry weights were 76.6 and 71.4 % for shoot and root, respectively, compared with control and soil treated with the fungicide Rizolex-T. The obtained results indicate potentialities of antagonistic bacteria to suppress the plant pathogens and stimulated its growth.

Bai *et al.* (2002) pointed out that *B. subtilis* when applied as co-inoculation to soybean plant provided the most consistent increase in shoot and root weights. The application of antagonistic bacteria alone significantly increased the pod peanut yield up to 48.9, 44.9 and 38.9 % for *B. amyloliquefaciens, B. subtilis* and *Brervibacterium otitidis*, respectively, compared with the control and soil treated with Rizolex T. The combined mixture of *Ps. synxantha* + *Brevi. otitidis* also increased the pod yield of peanut by 40.8 %.

Table 5. Influence of repeated inoculation with different bacterial strains and fungicide on shoot, root biomass and yield of peanut plants

	Shoot	dry weight	Root dry weight		Pods yield		100 Seed	Shelling
Treatments	g/ plant	Increase %	g/ plant	Increase %	Ardeb/ fed.	Increase %	Weight (g)	%
Control*	44.9		1.4	-	14.9	-	92.5	64.5
Rizolex T	61.0	35.9	2.3	64.3	15.7	5.4	86.3	65.8
Bacillus subtilis	84.1	87.3	2.6	85.7	21.6	44.9	99.6	66.5
Bacillus amyloliquefaciens	86.5	92.3	2.8	100	22.2	48.9	98.7	67.1
Pseudomonas synxantha	68.3	52.1	2.5	78.6	18.2	22.1	96.1	64.6
Brevibacterium otitidis	73.8	54.4	2.4	71.4	20.7	38.9	98.2	64.9
Mixture of four strains	79.3	76.6	2.4	71.4	19.3	29.5	93.9	66.3
B. subtilis + B. amyloliquefaciens	68.2	51.4	2.3	64.3	18.1	21.5	95.6	66.9
B. subtilis + Ps. synxantha	61.2	36.3	2.7	92.9	18.3	22.8	93.7	66.3
B. subtilis + Brevi. otitidis	70.4	56.8	2.6	85.7	19.7	32.2	92.7	66.6
B. amyloliquefaciens + Ps. Synxantha	65.0	44.8	2.5	78.6	15.5	4.0	93.8	68.1
B. amyloliquefaciens + Brevi. Otitidis	61.5	35.6	2.2	57.1	18.1	21.5	93.7	67.1
Ps. Synxantha + Brevi. Otitidis	63.2	40.8	2.8	100	20.3	36.2	95.4	66.4
LSD at 0.05	18.4		0.4		5.3		6.4	

^{*} Uninoculated treatment

^{**} Soil treated with Rizolex T fungicide

This result is in harmony with that of Turner and Backman (1991) who recorded an increase in the yield of carrots (48 %), oats (33 %) and peanuts (37 %) when plants were inoculated with *B. subtilis*.

Concerning dry weights of 100 seeds and shelling percentages, data (Table 4) show significant increases in most of the treatments as compared with those of the control and treatment with Rizolex T.

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كفاءة بعض السلالات البكتيرية في المكافحة الحيوية لمرض عفن جذور الفول السوداني النامي في الأراضي الرملية

لبنى عبدالعزيز موسى ، إبتسام محمد مرسى ، عبير أحمد شلتوت ، سهير سيد محمد فهمى ا

- ١- معهد بحوث الأراضي والمياه والبيئة، مركز البحوث الزراعية، الجيزة، مصر
 - ٢- معهد بحوث أمراض النبات، مركز البحوث الزراعية، الجيزة، مصر

تم إختبار كفاءة أربعة سلالات بكتيرية في المكافحة الحيوية لفطر ريزوكتونيا سولاني الدى يصيب الفول السوداني وهي باسلس ساتلس، باسلس أميلوليكوفيشنس، سيدوموناس ساينسانسا، بريفيباكتريم أوتيتداس وأظهرت النتائج أن كل السلالات أدت إلى تثبيط نموالفطر وأظهر ميكروب باسلس أميلوليكوفيشنس أعلى نسبة تثبيط لنمو هذا الفطر. وفي تجربة حقلية بمحطة بحوث الأسماعيلية تم تقبيم كفاءة هذه السلالات كلاً على حدة وبخليط منهم حيث ظهرت كفائتهم في مكافحة المرض. وقد تم تقدير نشاط إنزيمي الدهيدروجينيز والنيتروجينيز وتكوين العقد الجذرية والمحصول المنتج من الفول السوداني. وأظهرت النتائج أن كل السلالات المختبرة أدت إلى زيادة دلائل النمو النباتية بينما كانت السلالاتين باسلس ساتلس، باسلس أميلوليكوفيشنس أكثرهما كفاءة.