# EFFECT OF INOCULATION WITH VESICULAR ARBUSCULAR MYCORRHIZA ON ROOT-ROT DISEASE INCIDENCE AND PLANT GROWTH OF TWO CULTIVARS OF BROAD BEAN

#### By

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#### **ABSTRACT**

Effects of soil infestation with vesicular arbuscular mycorrhizal (VAM) fungi, i.e. Glomus macrocarpum and G. australe, each alone or in combination with the root-rot infecting fungi, i.e. Fusarium solani (Mart.) Appel & Wollenew. or Rhizoctonia solani Kühn, in sterilized and unsterilized soils, on percentage of survived plants, root-rot disease severity index (DSI) and some growth characters of broad bean (Vicia faba L.) cultivars Giza 2 and Ribaya -40 (R-40), were investigated under greenhouse conditions.

Mycorrhizal plants grown in soils infested with F. solani or R. solani showed significant increases in percentage of survived plants and reductions in root-rot disease incidence compared with the nonmycorrhizal plants grown with pathogens only. The VAM-fungus, G. macrocarpum was more effective, in this respect, than G. australe and broad bean cultivar R-10 responded better than Giza 2 one especially in unsterilized soil. Root-rot disease incited by F. solani on cv. Giza 2 and R. solani on cv. R-40 were slightly but significantly reduced by introducing G. australe into soils infested with these pathogens. However, double infestation of unsterilized soil with both G. australe and R. solani resulted in significant increase in DSI on cv. Giza 2 compared with R. solani alone. Fusarium solani caused considerable increase in colonization intensity of roots by the VAM-fungus G. australe but not G. macrocarpum, while R. solani, in general decreased colonization indexes of both VAM-fungi. Inversed relationships were detected between colonization index of G. macrocarpum only and DSI particularly of F. solani. The lowest values of DSI of F. solani or R.

solani and highest colonization index of G. macrocarpum were associated with mycorrhizal plants of cv. R-40 in unsterilized soil.

Soil infestation with G. australe and F. solani or R. solani showed significant higher plant height, root length, shoot and root dry weight than in case of pathogen only. Height and root length of F. solani infected plants, in both soil types were not significantly different from those inoculated with both G. macrocarpum and F. solani. Results of this investigation indicate that, soil infestation with VAM-fungi cauld be promising for controlling root-rot diseases and/or improved growth of broad bean plants.

#### INTRODUCTION

Fusarium solani and R. solani caused severe damping-off and root-rot diseases and great reduction in growth characters of broad bean (Vicia faba L.) plants (Sirry et al., 1970; Nofal et al., 1982).

Several investigators reported that diseases caused-by soil-borne pathogen can be influenced by the action of mycorrhizae inside the root system and /or its surrounded area (Santoro and Casida, 1962; Harely and Smith, 1983 and Ahmed et al., 1994). In general, mycorrhizal plants suffer less damage due to root-pathogens (Dehne, 1982, Saleh and Ahmed, 1988). More recently, the vesicular-arbuscular mycorrhizal (VAM) fungi showed an important role in the biological control of tomato and sunflower bacterial crown gall disease (Abd El-Sayed, 1992) and root-rot of broad bean caused by R. solani (Ahmed et al., 1994). However, some reports indicate an increase in disease severity under the influence of VAM-fungi (Ross, 1972 and Davis and Menge, 1980). Mycorrhizal plants showed enhanced growth development especially under field condition, mainly, because improvement of nutrients uptake and may be due to providing further protection against soil-borne pathogen attacking their roots (Garcïa-Garrido and Ocampo, 1987 and Ahmed et al., 1994).

The present study was conducted under greenhouse conditions to investigate effects of two species of the VAM-fungi, i.e. Glomus macrocarpum and G. australe each alone or in combination with F. solani or R. solani on the disease incidence and growth characters of two cultivars of broad bean in sterilized and unsterilized soils. Relation between intensity of VAM-colonization and disease severity index was also investigated.

#### MATERIALS AND METHODS

### Isolation of the causal organism(s):

The used cultures of root-rot infecting fungi, i.e. F. solani and R. solani were isolated from rotted roots of broad bean plants. Purification of the isolated fungi was carried out using hyphal tip and/or monosporic culture techniques, then identified according to Nelson et al. (1983) and Parameter and Whitney (1970).

## Inoculum preparation and Pathogenicity test:-

The inocula were prepared by growing each of *Fusarium solani* (*Mart.*) Appel & Wollenew. and R. solani (Kuhn.) on sterilized sorghum grain medium in glass bottles for 2 weeks at 28 C. A clay loam soil with pH 7.5, unsterilized or sterilized by autoclaving at 15 lb/in² for two hours, was infested by the prepared inocula separately at rate of 5.0% of the soil weight. The infested soil was potted in plastic pots (\$\phi\$ 20 cm), each containing 1.5 kg infested soil. Pots were watered on alternate days and incubated under greenhouse conditions for 7 days to maintain equal distribution of the fungal inoculum. Sterilized, un-inoculated sorghum grain medium was used in control pots. Surface sterilized seeds ( with 0.1% mercuric chloride solution for 2 min) of Giza 2 and Ribaya-40 (R-40) broad bean cvs., were planted at the rate of 6 seeds/pot. Four pots, were used for each particular treatment.

#### Disease Assessment:-

Percentage of pre-emergence damping-off was calculated 15 days from sowing. While, after 60 days from sowing, plants were carefully removed, washed currently with tap water, then examined for root-rot symptoms. Determination of the root-rot disease severity index (DSI) was carried out, based on a scale from 0 (non-visible damage) to 5 (completely destroyed roots) according to Salt (1982). Also, plant height, root length, fresh and dry weights of both shoots and roots were estimated at the same time.

#### Mycorrhizal Studies:-

These experiments were carried out under greenhouse conditions to investigate effects of adding vesicular-arbuscular mycorrhizae (VAM) to sterilized or unsterilized soil not infested or infested with *F. solani* or *R. solani* just before sowing on percentage of survived plants and root-rot disease incidence as well as on plant growth of two broad bean cultivars. All experimental necessities were accomplished as mentioned before.

Two different species of the VAM i.e. Glomus macrocarpum and G. australe were used in this study. Each of them was kindly provided by Dr. Gendia, H., Dept. of Hort., Fac. of Agric., Moshtohor. The VAM-fungi were propagated, separately, on onion plants using the propagation technique described by Al-Fassi et al. (1990). After three months from onion cultivation, the mycorrhizal roots of onion bulbs together with its adjacent soil were collected and used for mycorrhizal infestation. Plastic pots (\$\phi\$ 20 cm), each containing 1.5 kg soil prepared as above described were used. The inocula of the VAM was added at rate of 10 g/pot (modified from Fares, 1986). Pots without mycorrhizal infestation were served as control. Root-rot disease severity index (DSI), percentage of survived plants, plant height, root length, dry weight

of shoots and roots were determined 60 days after sowing as mentioned before. This work was repeated for two seasons (1991 and 1992). All data obtained were statistically analyzed by calculating the least significant difference (L.S.D.) at the 5% level according to Little and Hills (1975).

#### Effect of root-rot pathogens on the VAM-colonization:-

Samples from root system of broad bean plants arised in the above experiment were collected and examined for VAMcolonization as described by Phillips and Hayman (1970). The roots were washed several times by tap water, cutted into small segments (1 cm long) and preserved in FAA solution (5% formaldehyde; 5% glacial acetic acid; 54% absolute alcohol and 36% distilled water). For microscopic preparation, the preserved roots of each treatment were washed several times by tap water to remove the preservative fluid. The roots were treated with 10% potassium hydroxide (KOH) in test tubes, then heated in water bath for 10 min. at 80-90 C. Root segments were then washed with tap water followed by 10% HCl. Trypan blue stain (0.5g l<sup>-1</sup>) was added to the root portions and heated again at 80-90 C for 5 min. Some of treated root segments (up to 20 portions/treatment) chosen at random were picked up and placed on glass slides to which few drops of fresh lactic acid were added. Then examined microscopically for mycorrhizal infection. Intensity of VAMcolonization expressed as an average number of the mycorrhizal structures i.e. vesicles and arbuscules, per root segment was calculated. Relationship between colonization intensity and rootrot disease severity index was also investigated.

#### RESULTS AND DISCUSSION

The obtained data (not shown) proved that the isolated fungi, Fusarium solani and Rhizoctonia solani, could infect broad bean and cause pre-emergence damping-off and root-rot diseases. Damping-off was relatively higher in sterilized than unsterilized soil, and on cv. R-40 than cv. Giza 2. R. solani, mostly, was more virulent than F. solani. It caused the greatest reduction in plant height, root length, fresh and dry weights of shoots and roots of diseased plants of both tested cvs., especially in sterilized soil, compared with healthy plants in control treatments. These results are in agreement with these reported by Sirry et al. (1970) and Nofal et al. (1982). They stated that, R. solani and/or F. solani reduced seed germination, plant height, fresh and dry weights of shoots and roots of broad bean plants.

# Effect of VA-mycorrhizal fungi on percentage of survival plants and root-rot disease severity:-

Data in **Table** (1) show clearly that, percentage of survived plants of cv. Giza 2 in unsterilized soil was significantly higher than cv. R-40. However, both cvs. survived better in sterilized soil without significant difference in between.

The lowest percentages of survived plants were obtained in soil inoculated with *R. solani* or *F. solani*. These figures were significantly raised by using VAM-pathogen combination. However, degree of improvement was depended on type of the combined partners. In this regard, percentage of survived plants in soil infested with *G. macrocarpum* and *R. solani* was significantly higher than the corresponding figure of *G. macrocarpum* and *F. solani* combination. Meanwhile, when *G. australe* was combined with these pathogens, it showed less effect. Soil infestation with *G. australe* and *F. solani* or *R. solani* resulted

in the highest and lowest significant increases in percentages of survived plants, respectively compared with those survived in soil infested by any of these pathogens alone.

Regarding root-rot-DSI, Table (1) proved that, DSI of F. solani and R. solani (averages) infections were reduced from 65% and 63% on non-mycorrhizal plants to 20.0% and 19.4%, when G. macrocarpum was combined with these root-rot infecting pathogens, respectively. However, G. australe, seems to be effective only against infection of F. solani as DSI were 48.1% and 60.7%, respectively. It is interest to state that DSI incited naturally (without artificial soil infestation) was significantly increased with G. australe (9.4%) as compared with G. macrocarpum or control treatments (6.3%).

Regardless of soil type, non-mycorrhizal plants of both tested cvs. were reacted similarly against F. solani infection, while, DSI of R. solani was significantly higher on cv. R-40 than cv. Giza 2. However, applying of G. macrocarpum in soils infested by F. solani or R. solani resulted in great reductions of DSI of these pathogens especially on R-40, the more susceptible cv., as compared with Giza 2, the less susceptible cv. The other VAMfungus, G. australe, was slightly effective only against infections caused by F. solani on cv. Giza 2 and R. solani on cv. R-40 as compared with G. macrocarpum. Variations in effectivities of these VAM-fungi in controlling root-rot disease incidence may be correlated with the manner by which it could colonized roots of broad bean plants. In this regard, data in Table (2) proved that, colonization index in roots of mycorrhizal plants inoculated with G. australe together with a given pathogen especially F. solani was increased considerably compared with those inoculated with G. australe only. R. solani-infection, in general, caused slight reduction in G. australe-colonization except in case of cv. R-40 in

Table 1: Effect of soil infestation with the VAM-fungi, G. macrocarpum and G. australe on root-rot disease severity index (DSI) and percentage of survived plants of two cultivars of broad bean cultivated in sterilized and unsterilized, infested (with F. solani or R. solani) or uninfested soils after 60 days from sowing.

I		•		%	Su	rviv	al		% Root-rot DSI						
So		oil Types	Unsterilized Soil			Sterilized Soil			Unsterilized Soil			Sterilized Se		Soil	
ł	CV.	Treatment	Con- troi	F. solani	R. solani	Con- trol	F. solani	R. solani	Con- trol	F. solani	R. solani	Con- trol	F. solani	R. solani	
4	Cira 2	Control G. macroca G. austarle		50.0	29.2 62.5 33.3	100 100 100	29.2 58.4 37.5	20.9 58.4 41.7	10.0 10.0 15.0	25.0	52.0 15.0 71.8	0.0 0.0 0.0	72.5 22.5 25.0	55.0 32.5 53.5	
	_	Control G. macroca G. austarle		66.7	33.3 62.5 45.8	100 100 100	20.9 58.4 62.5	16.7 50.0 37.5	15.0 15.0 22.5	52.5 7.5 60.0	65.0 15.0 60.0	0.0 0.0 0.0	82.5 25.0 57.5	80.0 15.0 55.0	
	L.S.D. at 5% for: Sur								val		Root	-rot	DSI		

L.S.D. at 5%	for:	Survival	Root-rot			
	Soil type (S)	N. S.	N. S.			
	Cultivar (C)	N. S.	N. S.			
	Treatment (T)	0.44	2.90			
	S×C	0.29	N. S.			
	$S \times T$	N. S.	N. S.			
	$C \times T$	N. S.	12.6			
	$S \times C \times T$	N. S.	N.S			

sterilized soil. These results are in agreement with Zamblin and Schenck (1983) working on *R. solani* of soybean and Kiran et al. (1987) on Fusarium-wilt of broad bean. In most cases DSI of a given pathogen on G. australe-mycorrhizal plants was affected proportionally with colonizing ability of G. australe. When the latter, VAM-fungus was introduced into sterilized soil, it could minimized infection of F. solani on cv. Giza 2 only up to 34.5% and its colonization index was increased up to 262.0%. The corresponding values, in unsterilized soil for the same cultivar and inoculation were 95.2% and 353.5% from those arised from F. solani and G. australe, each alone, respectively. As for cv. R-40, similar trend was also observed. These results indicated that both DSI and VAM-colonization index may be affected by the interaction between pathogen and VAM-fungus. In this respect,

Table 2:Colonization intensity and colonization index\* of the VAM-fungi G. macrocarpum and G. australe as affected by F. solani and R. solani, the root-rot pathogens on two broad bean cultivars.

So	oil Type	Uns	terilized	Soil	Sterilized Soil				
CV.	Treatment	Control	F. solani	R. solani	Control	F. solani			
Giza	G. macroc.	63.2 100.0	34.8 55.1	29.3 46.4	62.5 100.0	48.7 77.9	28.6 45.8		
2	G. australe %	40.9 100.0	144.6 353.5	32.3 79.0	48.2 100.0	126.3 262.0	46.2 95.9		
R-40	G. macroc.	43.0 100.0	51.2 119.1	66.0 153.5	56.4 100.0	41.9 74.3	39.0 69.1		
	G. australe %	51.9 100.0	160.7 309.6	41.5 80.0	42.2 100.0	94.2 223.2	46.6 110.4		

colonization intensity in case of VAM-pathogen combination x 100/ colonization intensity in VAM alone.

Ross (1972) found that Phytophthora root-rot of disease-susceptible but not disease-tolerant soybeans was increased by VA-mycorrhiza. This may have been because the tested Endogone spp. produced large vesicles in root cortex which could have caused some splitting of tissues, thereby facilitating entry of the pathogen. The obtained results are in agreement also with those reported by Zamblin and Schenck (1983) and Kiran et al. (1987). In case of mycorrhizal plants inoculated with G. macrocarpum and F. solani or R. solani, inversed relationship between DSI and colonization index was observed. For example, DSI of F. solani was minimized up to 14.3 % on G. macrocarpum-mycorrhizal plants of cv. R-40 grown in unsterilized soil. For the same combination, colonization index of G. macrocarpum was maximized up to 119.1%. On the other side, mycorrhizal plants of cv. Giza 2 infected with F. solani in unsterilized soil showed the maximum DSI and minimum colonization index. These results are in accordance with Al-Raddad (1991) who stated that, all tested Glomus spp. (seven isolates) reduced percentage

of Fusarium-infection in tomato and pepper roots at different rates. In fact, mycorrhizal fungi may changed the natural reaction of plants against their pathogen or may have no effect. The final reaction seems to be depending on several factors including soil conditions, host, pathogen, and the particular nature of the interaction between the host and the pathogen. In this point, Dehne (1982) reported that, mycorrhizal fungi, in some cases, reduced the disease incident due to pathogens causing morphological or physiological changes in the plant. Garcia-Garrido and Ocampo (1987) indicated also, that VAmycorrhizae could protect the plant against pathogenic organisms such as nematodes, fungi, bacteria and viruses. On the other hand, several investigators reported that, the mycorrhizal plants suffered more from, Verticillium-wilt on cotton (Davis et al., 1979), Phytophthora-rootrot on citrus (Davis and Mange, 1980). In addition, Zamblin and Schenck (1983) showed that, Glomus mosseae, the VAM-fungus, did not reduce significantly the disease index rating of R. solani on broad bean plants, but R. solani significantly reduced the percentage of root colonization by G. mosseae in autoclaved soil. The maintained successful in control of root-rot diseases by using VAM-fungi might be attributed to improved uptake of phosphorus, nitrogen and other mineral nutrients from soil as reported by Ross (1971) and Kucey and Paul (1983). In general, mycorrhizal fungi may conceivably afford protection to the roots by, utilizing surplus carbohydrate, thus reducing attractiveness of the roots to the pathogens (Zak, 1964); increasing in lignification of the cell wall (Dehne and Schonbeck, 1979) secreting antibiotic substances (Santoro and Casida, 1962) or by favoring, along with the root, protective rhizosphere organisms (Ahmed et al., Protection of the roots, against root-rot infection, by G. 1994). macrocarpum may not be limited to roots actually formed mycorrhizae. The protective substances which may be secreted by it

into root tissue during colonization process could be translocate to and benefit other root parts.

#### Effect of VAM-mycorrhizae on plant growth:

Data presented in Table (3) exhibited that, inoculation with both tested VAM-fungi increased plant height significantly compared with control treatment. Double inoculation with G. ausrale and F. solani resulted in the highest and significant increase in plant height compared with G. australe or F. solani alone. Similar increases in plant height was also obtained by combinations of R. solani with any of the tested VAM-fungi compared with R. solani alone. However, G. macrocarpum and F. solani combination has no significant effect on plant height compared with F. solani alone, but it was significantly decreased as compared with G. macrocarpum alone. Reduction was more pronounced on cv. R-40. These results were true in both sterilized and unsterilized soils. However, soil infestation with both G. macrocarpum and F. solani, in both types of soil, produced the highest reduction in plant height of cv. R-40. The same data proved that, root length was affected similarly as in plant height. However, G. australe combined with F. solani or R. solani produced significant improvement of root length of both tested cvs. in both soil types compared with the pathogen alone. These results could be supported by Al-Raddad and El-Saket (1991) who reported that, roots of good growth olive seedlings showed higher mycorrhizal structures, while week seedlings showed lower intensity of colonization. These are in agreement with the present work.

Data in Table (4) show that, dry weights of shoots and roots of both tested cvs., were clearly affected by the VAM-fungi and their combinations with root-pathogens. In most cases, *G. austarle* alone or combined with *F. solani* or *R. solani* gave the best results. However, *G. macrocarpum* combined with any of these pathogens produced the highest dry weight of shoots of R-40 cv. in unsterilized

Table 3: Effect of soil infestation with the VAM-fungi, G. macrocarpum and G. australe on plant height and root length of two cultivars of broad bean plants cultivated in sterilized and unsterilized, infested (with F. solani or R. solani) or uninfested soils after 60 days from sowing.

	·	Plant height (cm)							Root length (cm)						
So	il Types	Unsterilized Soil			Sterilized Soil		Unsterilized Soil			Sterilized Soil		ed			
cv.	Treatment	Cont-	F. solani	R. solani	Cont- rol	F. solani	R. solani	Cont-	F. solani	R. solani	Cont-	F. solani	R. solani		
Giza 2	Control G. macroc, G. australe		17.5 16.2 36.6	1	19.6 23.6 24.7	19.8 22.1 34.0	19.8 23.8 26.5	12.4 11.1 22.3	12.3	10.3 15.1 20.2			12.1 16.7 19.8		
92	Control G. macroc. G. australe		15.7 13.8 27.0	18.2 19.8 27.7		16.4		10.9 12.6 22.3	12.1 13.3 20.8	14.2 13.2 20.4	13.4 15.2 19.3	10.4			

L. S. D. at 5% for:	Plant height	Root length
Soil type (S)	N. S.	N. S.
Cultivar (C)	N. S.	N. S.
Treatment (T)	2.70	2.64
S × C	N. S.	N. S.
$S \times T$	N. S.	N. S.
$C \times T$	3.89	N. S.
$S \times C \times T$	5.28	5.28

soil only compared with its specific control. The same combinations on Giza 2 cv. in both types of soil and R-40 cv. in sterilized soil showed slight reduction in shoot dry weight compared with *G. macrocarpum* alone. As for root dry weight, the same data proved that it was affected similarly as in shoot dry weight. These results are supported by Khan (1973); Graham et al. (1976) and Kucey and Paul (1983). They reported that, their mycorrhizal plants grew much better and the dry matter of its shoots and roots were higher than the non-mycorrhizal plants. Also, Al-Raddad (1991) stated that, mycorrhizal tomato plants inoculated with *F. oxysporum* possessed a significantly higher root, shoot weight and plant height than those inoculated with *F. oxysporum* only. Pepper plants inoculated with *G. mosseae* showed higher fresh weight and

Table 4: Effect of soil infestation with the VAM-fungi, G. macrocarpum and G. australe on shoot and root dry weight of two cultivars of broad bean plants cultivated in sterilized and unsterilized, infested (with F. solani or R. solani) or uninfested soils after 60 days from sowing.

<u> </u>	Shoot dry weight(g)									Root-dry weight(g)							
Soil Types		Unsterilized Soil			Sterilized Soil			Unsterilized Soil			Sterilized Soil						
	Treatment		F. solani	R . solani	Control		R. solani	Control	F.	R	Control	F.	R				
0 Giza 2	G. australe	24.9 31.8 23.4	15.7 27.2 42.4	15.6 18.1 46.7	24.7 25.4 49.5	15.2 14.4 55.3	10.6 14.8 55.8	3.55 4.22 7.50	3.45 2.67 8.71	5.17 9.56	7.82 4.35 5.61	2.93 3.65 8.00	3.66				
R4	G. macroc. G. australe	31.1	62.5	14.9 56.6	24.0 21.8	19.3 13.6	11.8	4.10	3.36	3.32 5.10		3.55 4.43	3.82 3.34				

height than the mycorrhizal plants inoculated with *F. oxysporum* and plants inoculated with *F. oxysporum* alone. He added that, growth of pepper plants inoculated with *F. oxysporum* was not significantly different from plants inoculated with both VAM-fungi and *F. oxysporum*. In fact, the VA-mycorrhizal plants could absorb greater amounts of phosphorus and other nutrients from soil (Harley and Smith, 1983) and have more healthy roots when grown in infested soil with the pathogen (Davis and Menge, 1981), finally, this make them grew better than the non-mycorrhizal plants.

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#### تَاثَير الحقن بالميكورهيزا الحويصلية الشجيرية على النمو ودرجة الإصابة بالعفان الجذور في صنفين من الفول البلدي

في هذه الدراسة جرى حقن التربة المعقمة أو غير المعقمة تحت ظروف الصوبة بساحد فطريات الميكورهيزا الداخلية Glomus macrocarpum أو Glomus australe منفردا أو مع أي من مسببات عفن الجذور

solani أو Rhizoctonia solani بهدف التعرف على تأثير ذلك على حدوث الإصابة بأعفان الجذور وبعض صفات النمو في صنفي الفول البلدي جيزة ٢ وريباية ٤٠ وكذلك دراسة العلاقة بين شدة الإصابة بالمرض وكثافة إستعمار الجذور بفطريات الميكورهيزا المستخدمة وفيما يلي أهم النتائج التي تم التوصل إليها :-

- كاتت الإصابة بالمرض في التربة المعقمة أعلى منها في غير المعقمة ، وعلى الصنف ريباية ، ٤ أكثر منها على الصنف جيزة ٢ ، وفي معظم القياسات كان الفطر ريزوكتونيا أشد تأثيرا من الفطر Fusarium حيث سبب فضلا عن تأثيره الممرض نقصا شديدا في أطوال الجذور ووزنها الغض والجاف في كلا الصنفين تميزت النباتات النامية في تربة محقونة بأي من فطريات الميكورهيزا وأحد مسببات المرض بتحسن معنوي في كل من نسبة بقائها ودرجة مقاومتها للإصابة بأعفان الجذور مقارنة بتلك النامية في تربة محقونة بالمسبب المرضي فقط وفي مؤا الخصوص كان الصنف ريباية ، ٤ أفضل في استجابته من الصنف جيزة ٢ هذا الخصوص كان الصنف ريباية ، ٤ أفضل تأثيرا من G. australe على الصنف والفطر الى زيادة معنوية في شدة الإصابة بالفطر Rhizoctonia على الصنف جيزة ٢ في التربة غير المعقمة.
- أدى الفطر F. solani إلى زيادة كبيرة في كثافة إستعمار الميكور هيزا للجذور خاصة G. australe بينما سبب As. solani في معظم الأحيان تناقصا واضحا في كثافة الإستعمار بكل من فطري الميكور هيزا.
- لوحظت علاقة عكسية بين كل من معدل كثافة استعمار الجذور بالفطر G. ## macrocarpum وبين شدة إصابة هذه الجذور بمسببات المرض خاصة F. ## solani ومثال ذلك الصنف ريباية ٤٠ النامي في تربة غير معقمة ومحقونة ## بالفطر macrocarpum مع أي من مسببات المرض تحت الدراسة والذي ظهر عليه أعلى معدل استعمار ميكورهيزي مع أقل نسبة إصابة بالمرض مقارنة بنفس الصنف المحقون فقط بالمسببات المرضية .
- أدى إستخدام G. australe مع أي من المسببات المرضية المستخدمة إلى زيادة معنوية كبيرة في إرتفاع النبات وطول الجذور ، كذلك زيادة الوزن الجاف للجذور مقارنة باستخدام تلك المسببات المرضية كل على الفراد ، ومن ناحية أخرى لم تظهر هناك اختلافات معنوية في هذه القياسات عند استخدام كل من آخرى لم تظهر هناك اجتلافات معنوية معا مقارنة بالأخير منفردا.