

Interaction between endomycorrhizae and rhizosphere fungi in soils of Iraq

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Endomycorrhizal and rhizospheric fungi were recovered from soil samples collected from the root of plants in Iraq. The relations between the two fungal populations were investigated.

Key words: endomycorrhizae, rhizosphere fungi, fungi populations.

INTRODUCTION

The mutualistic association between plants and mycorrhizal fungi is beneficial. The main role of mycorrhizae is to supply the essential nutritional elements from soil to the plant roots and improving the growth of host plants (H a r l e y 1989). In additions, mycorrhizae have an impact on the soil surrounding plant roots (I n g h a m, M o l i n a 1991). Nevertheless, the interaction between mycorrhizal fungi and soil microorganisms including bacteria, protozoa, nemaodes and pathogenic fungi have received considerable attention (S y l v i a 1990; I n g h a m, M o l i n a 1991). Mycorrhizae protect plants from nematodes infection (K e l l a m, S c h e n c k 1980; S h a r m a J o h r i, G i a n i n a z z i 1992) affecting the bacterial populations in soil (A m e s, R e i d, I n g h a m 1984) and reducing the pathogenicity of parastic fungi (Z a m b o l i m, S c h e n c k 1993; G i o v a n n e t t i, A v i o, S a l u t i n i 1991). However, little information is available regarding the interaction between endomycorrhizae and rhizospheric fungi in soil and only few cases have been reported (D a n i e l s, M e n g e 1980; S c h e n c k 1981; K u c e y 1987).

In this study an attempt was made to demonstrate the relationship between endomycorrhizae and rhizospheric fungi in soils of five selected host plants in Iraq.

MATERIALS AND METHODS

Soil samples were taken from root zones (from a depth of 10-15 cm) of five host plant species, namely: *Cordia myxia* L., *Ficus carica* L., *Lawsonia inermis* L., *Punica granatum* L., and *Vitis vinifera* L. growing in Southern Iraq. A total of ninety soil samples were taken bimonthly between January and November 1994. At each collecting time, 100 g soil sample per plant was placed in a plastic bag and brought to the laboratory for fungal spore isolation. Soil properties were determined such as: pH (6.6-7.5), conductivity (4.5-9.8 mmohs), moisture (7.8-12.8%) and soil texture (silt 60.1%, clay 35.3, sand 1.6).

Twenty grams of each soil sample were processed to recover spores of endomycorrhizal fungi using wet-sieving technique following Koske, Halverson (1981). Spores enumeration was made according to Giovannetti, Mosse (1980). Triplicates were used for each soil sample per collection. For isolation of rhizosphaeric fungi, the dilution plating method was conducted as described by Warcup (1950); 20 grams of soil were diluted in 100 ml of distilled water and subsequently 1 ml of soil suspension was transferred into a Petri-dish to which Czapek agar medium was added. Fungal colonies were surveyed and counted after 4-5 days after being incubated at 20°C. Triplicate were made for soil samples.

RESULTS

A total of 17 species of endomycorrhizal fungi recovered from soils collected from the five host plants (Table 1). The number of species found was as follows: 9 species in samples of *Cordia myxia*, 11 species with *Ficus carica* and with *Vitis vinifera*, 12 species with *Lawsonia inermis* and with *Punica granatum*. Out of them four taxa namely *Acaulospora bireticulata*, *A. scrobiculata*, *A. trappei*, *Endogone incrassata* and *Glomus fasciculatum* were common for all the plants. The total spore number of endomycorrhizae varied in the collecting time and in the plants predominant species were *Glomus fasciculatum*, *G. leptotichum*, and *Acaulospora laevis* in the soils of the host plants (Table 1). Seasonal variations in spore number of endomycorrhizae was observed. High total spore number was detected in January-March for all the soil samples, except for *Lawsonia inermis*. A low total spore number was noted in September (Table 1). Among the host plants *Cordia myxia* rendered the highest spores number while *Vitis vinifera* showed the lowest spore number of endomycorrhizae.

The present data showed that 17 species of rhizospheric fungi inhabited in the soils of the plants studied (Table 2). Among the recovered fungi, 10 species were present in the soil sample of *C. myxia*, 12 species – *F. carica*, 13 – *P.*

granatum and 14 species — *L. inermis* and *V. vinifera*. Among these taxa seven species, i.e., *Alternaria alternata*, *Aspergillus fumigatus*, *A. niger*, *Cladosporium herbarum*, *Cunninghamella echinulata*, *Fusarium moniliforme* (and *Penicillium* sp.) were common for the plant soils. The dominating fungi were: *Aspergillus niger*, *Cladosporium herbarum* (and *Penicillium* sp.) in all soil samples and through out the collecting period. Temporal variation in the total number of fungal isolates was observed (Table 2). The maximum number of isolates was encountered in September, except for *Lawsonia inermis*, and the lowest number of isolates was noted in March, except for *Cordia myxia*.

Data, regarding the total number of isolates of endomycorrhizae and rhizosphere fungi, were presented according to the dates of collection for each plant (Fig. 1). The general seasonal variation trend of endomycorrhizal fungi showed that the highest population density was in March and the lowest in September. A reverse relationship was observed for the rhizospheric fungal population, i.e. maximum population density was noted in September and minimum in March (Fig. 1).

DISCUSSION

The markedly varied species composition and population density of endomycorrhizal and rhizospheric fungi can be related to the nature of the host plants. It has been stated that host plants play a role in the presence or absence of endomycorrhizae, which is, in part, due to root exudation process (I n g h a m, M o l i n a 1991). Consequently, any changes in root exudation would affect the fungal colonization and their populations. The low density in the population of endomycorrhizae in September is possible associated with the low amount of root exudates produced by plants at the senescent growth stage. These observations are in agreement with other studies (G e m m a, K o s k e, C a r r e i r o 1989; M c G e e 1989; O k e e f, S y l v i a 1991; S y l v i a, W i l l i a m s 1992). These studies showed that high and low spore production by endomycorrhizae occur at early and late plant growth stages.

The present study also demonstrated that the endomycorrhizal association developed more frequently, e.g., in *Glomus fueganum* which was more abundant in the soil of the host *C. myxia*, *Glomus fasciculatum* in the soil of *F. carica*, *Glomus aggregatum* seemed to have a wider range of occurrence in many plants. This is in agreement with the finding of (H e t r i c k, B l o o m 1983; H e t r i c k, B l o o m 1986).

On the other hand, it appeared that soils of the selected plants harboring diverse taxa of rhizosphere fungi. Nevertheless, seasonal variations in rhizospheric fungal isolates can be related to different biotic and abiotic factors (P u g h 1980; D o m s c h et al. 1980; C h r i s t e n s e n 1989). The high

T a b

Mean spore number (per 10 g of soil) of endomycorrhizal fungal (ENMF)

ENMF	<i>C. myxia</i>						<i>L. inermis</i>					
	Jan.	Mar.	May	July	Sept.	Nov.	Jan.	Mar.	May	July	Sept.	Nov.
<i>Acaulospora biradiculata</i> Roth. et Trappe	11	19	16	.	.	10	8	.	.	10	5	.
– <i>laevis</i> Gerd. et Trappe	25	4	.	9
– <i>scrobiculata</i> Trappe	14	.	9	10	2	.	.	.	13	8	.	7
– <i>digitata</i> Walker et Trappe	9	.	.	.	15	.
– <i>trappei</i> Ames et Linder	.	.	6	.	.	.	5	.	5	.	.	.
<i>Glomus aggregatum</i> Schenck et Smith	19	19	20	31	.	.	.	10	.	22	10	.
– <i>albidum</i> Walkers et Rhod.
– <i>deserticola</i> Trappe, Bloss et Menge	10	.	9	.	15	22	.	.	15	.	.	10
– <i>fasciculatum</i> (Thaxter) Gerd. et Trappe	51	40	15	.	.	10	10	18	.	.	15	10
– <i>fragile</i> (Berk. et Br.) Trappe et Gerd.	8	.	.	.
– <i>fuegianum</i> (Speg.) Trappe et Gerd.	.	5	12	23	33	33	.	.	10	.	5	.
– <i>fulvum</i> (Berk. et Br.) Trappe et Gerd.
– <i>leptotrichum</i> Schneck	23	6	15	14	8
– <i>microaggregatum</i> Koske, Gemma et Olexis	16	.	30	.	.
– <i>mosseae</i> (Nicol. et Gerd.) Gerd. et Trappe
– <i>occultum</i> Walker	.	10	2	.	14	6	.	.	7	.	.	3
– <i>reticulatum</i> Bhatt. et Mukerji
Total spores	128	99	104	78	72	71	32	44	83	84	73	59
	552						365					

Table 1

species of five host plants throughout the study period

<i>F. carica</i>						<i>P. granatum</i>						<i>V. vinifera</i>					
Jan.	Mar.	May	July	Sept.	Nov.	Jan.	Mar.	May	July	Sept.	Nov.	Jan.	Mar.	May	July	Sept.	Nov.
8	.	.	.	10	.	12	.	6	7	10	.	10	10	.	14	.	6
.	8	.	13	.	21	7	20	20	28	.	14	.	.	13	10	9	13
.	17	27	.	7	20	10	15	12	.	.	10	.	6	8	12	.	6
.	.	.	.	8	9	3	.	.	.	19	25
.	20	16	30	25	4	.	.	13
.	7	.	6	16	.	.
.	8	10	28
.	8	4	7
21	29	15	7	.	.	18	21	17	28
11	.	.	4	.	.	9
.	2	4	15	13	15	.	2	13	11	.	.
7	11
24	19	11	.	21	.	.	10	8	.	31	14
.	.	1	12	4	.	7	10	.
.	.	.	4	16	10	10	21
.	9	5	.	.	.	5	5	5	.	.	.
.	10	.	.	16	6	.	5	.	10	.	4	6
71	104	70	40	62	68	84	94	92	103	70	74	46	55	59	70	42	69
411						517						339					

T a b

Mean spore number (per 10g of soil) of other rhizospheric fungal (ORF)

ORF	<i>C. myxia</i>						<i>L. inermis</i>					
	Jan.	Mar.	May	July	Sept.	Nov.	Jan.	Mar.	May	July	Sept.	Nov.
<i>Alternaria alternata</i> (Fr.) Keissl.	2	9	6	3	.	.
<i>Aspergillus fumigatus</i> Fres.	.	17	.	6	.	5	30	.	.	2	.	4
<i>A. niger</i> Tiegh.	.	.	22	58	10	42	79	15	17	45	40	.
<i>A. terreus</i> Thom	.	3	.	.	9	.	.	11
<i>A. versicolor</i> (Vuill.) Tirab
<i>Cladosporium herbarum</i> (Pers.) Link	1	3	6	10	69	.	.	.	13	.	24	.
<i>Cunninghamella chinolata</i> Thaxter	17	.	2	6	6	1	.	.
<i>Drechselara australiensis</i> Subram., Jain et Ellis (<i>Drechselara</i> sp. state of <i>Cochiobolus spicifer</i>)	6
<i>Fusarium moniliforme</i> Scheld.	8	4
<i>F. oxysporum</i> Link
<i>Humicola grisea</i> Traaen	10	.	.	2	5
<i>Mucor circinoid</i> Wehmer	4	3	.
<i>M. meahie</i> Cooney et Emers.	3	.	3	.	.
<i>Nigrospora oryzae</i> (Berk.) Petch
<i>Paecilomyces roseum</i> Bain. (<i>Penicillium</i> sp.)	.	7	25	21	.	.
<i>Stachybotrys atra</i> Corda	.	5	9	.	7	64	.	50	10	.	10	83
<i>Ulocladium botrytis</i> Preuss	1	.
Total spores	20	35	39	80	105	69	115	98	71	75	80	96

number of fungal isolates in September accounted for the predominant species: *Aspergillus niger*, *Cladosporium herbarum* (and *Penicillium* sp.). This is perhaps due to the fact that these species compete strongly with other fungi species in soil (P u g h 1980).

Data presented in Fig. 1 showed a close reverse relationship between endomycorrhizal and rhizosphere fungal populations. The negative relationship was marked throughout the growing season and was related to the plants. It is evident that the low population density of rhizosphere fungi was influenced by the increase in endomycorrhizal population. Such inverse relationship might be the result of depletion in root exudates due to mycorrhizal colonization (L a h e u r t e, B e r t h e l i n 1986), high competition of mycorrhizae with non-mycorrhizal fungi (G r a h a m, M e n g e 1982), or to the production of antifungal substances produced by mycorrhizae (I n g h a m, M o l i n a 1991) which may hinder the growth and populations of soilborn fungi.

Table 2

species of five host plants

<i>F. carica</i>						<i>P. granatum</i>						<i>V. vinifera</i>					
Jan.	Mar.	May	July	Sept.	Nov.	Jan.	Mar.	May	July	Sept.	Nov.	Jan.	Mar.	May	July	Sept.	Nov.
2	2	.	.	7	27	42	5	.
2	7	10	.	11	4	22	.	30	.	3	.	14
21	16	.	17	24	27	54	34	18	55	77	41	.	50	58	47	30	49
.	8	3	.	.
.	9
49	.	78	76	37	50	32	20	17	24	.	8	41	13	22	15	60	30
.	5	10	8	9	.	2	3	1	6	.	.	10	.	.	5	.	.
.
.	2	.
.	4	.	.	2	.	.	2	4
.	.	.	.	2	3	2	3
.	.	.	.	4	3	.	.	2	.	.	3	.
16	12	11	20	.	.	.	8	.
.	1	1	2
.	3
.
.	10	.	10	13	12	.	12	16	13	27	31	79	55	37	31	83	15
.	2
.	.	.	.	4	4
90	39	98	111	104	97	92	69	87	98	121	103	165	129	152	143	193	110

The present results indicate that the endomycorrhizae infested soilborn inhibit or reduce the activity of rhizosphere fungi. Few reports concerning the relationship between endomycorrhizae and soilborn fungi are available and only single examples were cited in literature. Daniels and Menge (1980) reported a negative interaction between *Glomus macrocarpum* and *Pythium* sp. in soil of wheat. Schenk and Kellam (1978) indicated that mycorrhizae reduced fungal pathogen attack on some plant crops. In addition, reduction of the soybean root-infecting fungi including *Fusarium solani* and *Rhizoctonia solani* was observed by Zambolim and Schenk (1983).

It is worth mentioning that among the isolated fungi which are known as common saprophytes, some species, however, were reported as plant pathogens (Domsch et al. 1980). These observations should lead to further studies regarding the introduction of endomycorrhizae into soil to reduce the negative effect of endomycorrhizae into soil to reduce the negative effect of other fungi.

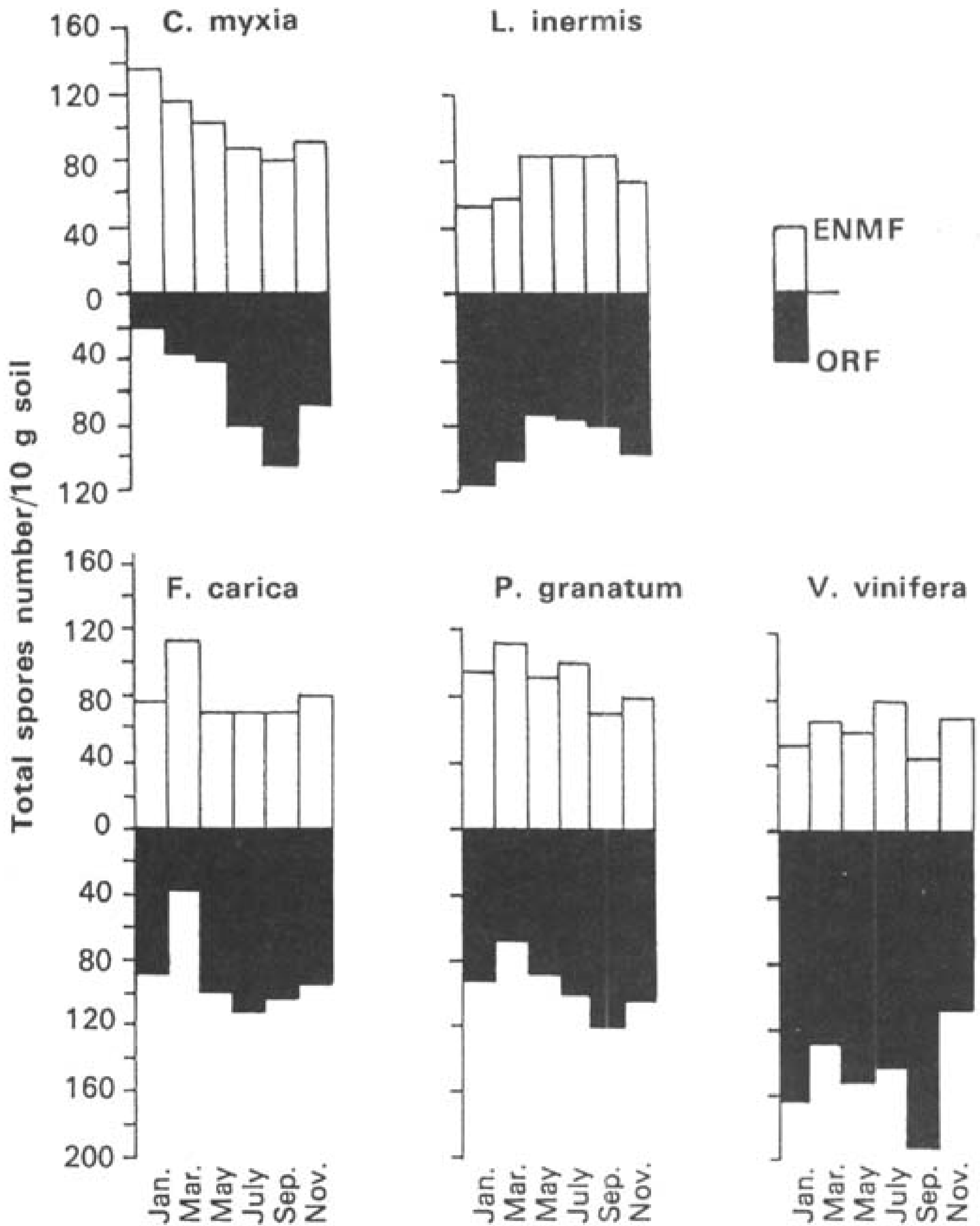


Fig. 1. Correlation between the total number of endomycorrhizal fungal (ENMF) spores and the total number of other rhizospheric fungal (ORF) spores in soil of five host plant species

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Interakcje grzybów mikoryzowych i ryzosferowych w glebach Iraku

S t r e s z c z e n i e

W próbkach gleby pobranej ze strefy korzeni pięciu roślin w Płd. Iraku wykryto obecność dziewiętnastu gatunków grzybów mikoryzowych i ryzosferowych. Największą liczbę endomikoryz stwierdzono w marcu, najmniejszą we wrześniu; odwrotny stosunek wystąpił w przypadku izolatów grzybów ryzosferowych. W pierwszym przypadku dominowały *Glomus fasciculatum*, *G. leptotrichum* i *Acaulospora laevis*, natomiast w drugim w glebie najwięcej było *Aspergillus niger*, *Cladosporium herbarum* oraz *Penicillium* sp. Przyczyną tak głębokich różnic pomiędzy tymi dwoma populacjami mogłaby być konkurencja między nimi, zasiedlanie korzeni przez gatunki oddziałujące niekorzystnie na grzyby i nie-mikoryzowe lub też przez wytwarzanie przez grzyby mikoryzowe substancji hamujących lub redukujących populacje gatunków ryzosferowych.