THE OCCURRENCE OF ARBUSCULAR MYCORRHIZAL FUNGI OF THE PHYLUM GLOMEROMYCOTA IN ISRAELI SOILS

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ABSTRACT

In December 1997 and June-July 2000, 49 and 113 rhizosphere soil and root mixtures were collected, respectively, to determine the occurrence of arbuscular mycorrhizal fungi (AMF) of the phylum Glomeromycota in different sites of Israel. Except for five samples taken from under cultivated plants, all the others came from under Ammophila arenaria and Oenothera drummondii colonizing sand dunes adjacent to the Mediterranean Sea. After a continuous cultivation of the mixtures in pot trap cultures with Plantago lanceolata as the plant host up to 2006 and their examination at least twice a year, spores of AMF were found in 41 and 103 cultures with the 1997 and 2000 soil and root mixtures, respectively. The spores represented 30 species and 8 undescribed morphotypes in 7 genera of the Glomeromycota. The AMF most frequently found in Israeli soils were Glomus aurantium and G. constrictum, followed by G. coronatum, G. gibbosum, an undescribed Glomus 178, and Scutellospora diporpurascens. Up to 2001, 21 species of AMF were known to occur in Israel, and this paper increases this number to 33, of which 11 are new fungi for this country. Moreover, four species, G. aurantium, G. drummondii, G. walkeri and G. santium, were recently described as new for science based on spores isolated from Israeli soils. Additionally, the general distribution in the world of the formally described species found in Israel was presented.

KEY WORDS: arbuscular mycorrhizal fungi, Glomeromycota, Israeli soils, occurrence.

INTRODUCTION

One of the most frequently occurring and widely distributed soil microorganisms in the world are arbuscular mycorrhizal fungi (AMF) of the phylum Glomeromycota. Until quite lately, they have been considered to associate with ca. 80% of plants of the Earth (Gianinazzi and Gianinazzi-Pearson 1986). However, the use of specific study methods, including, e.g., cultivation of rhizosphere soil samples and/or root fragments in pot trap cultures as well as molecular analyses of roots, has showed that AMF also frequently coexist with plants of, e.g., of the families Brassicacese and Chenopodiaceae, earlier generally considered to be non-mycorrhizal (Harley and Harley 1987, 1990). This induced the Committee of the International Bank of Glomeromycota to express the supposition that “The majority of plants, strictly speaking, do not have roots; they have mycorrhizas” (http://www.kent.ac.uk/bio/beg/english-homepage.htm).

Literature data indicate that AMF play a vital role in the life of plants. They increase, e.g., (1) the growth and nutrition of plants due to increased absorptive area of their roots by extraradical hyphae extending up to 10 cm outside the roots (Bieleski 1973), (2) the rate of succession (Janos 1980) and competitiveness of plants (Allen and Allen 1984; Fitter 1977), and (3) pollen production (Lau et al. 1995). Additionally, AMF (1) influence plant phenology (Allen and Allen 1986), (2) improve structure and stability of the soil through binding sand grains into aggregates by extraradical mycorrhizal fungi (Koske 1975), (3) aid occurring plants by transferring nutrients from better nourished plants to those of a poorer condition by hyphal bridges (Newman 1988), (4) protect plants against pathogens and nematodes (Bagyaraj 1984; Schönbeck 1978), as well as (5) alleviate the harmful effects of soil toxic substances (Turnau and Haselwander 2002). However, the range of the improvements listed above highly depends on a species or a strain of the fungus associated with a given plant (Stahl and Christensen 1991).

Both the spore abundance of AMF and the species diversity of their spore populations may be highly influenced by, e.g., (1) a plant species and its age (Dalpé 1989; Gemma et al. 1989; Koske and Gemma 1997), (2) the composition of non-mycorrhizal soil microbiorganisms (Lee and Koske 1994), (3) the amount and composition of both organic and inorganic chemical substances (Koske and Gemma...
1997; Rose 1988), (4) pH (Porter et al. 1987), (5) soil compaction and its humidity (Koske and Halvorson 1981; Read 1989), as well as (6) above- and underground temperature (Koske 1987).

A few studies of the occurrence of AMF have been conducted in Israel to date. Dodd and Krikun (1984) found five species and one morphotype, and Haas and Menge (1990) nine species. Following the cultivation of field-collected rhizosphere soil and root mixtures in pot trap cultures, Błaszkowski et al. (2001b) revealed 17 species and 8 undescribed morphotypes. Among the described found species, 9 were new for this country. However, Błaszkowski et al. (2001b) examined only 49 soil and root samples collected from three sites and all but five were taken from dunes of the Mediterranean Sea adjacent to Tel-Aviv. Moreover, all the identified fungi were isolated only from the first cycle pot trap cultures containing the soil and root mixtures collected. Meanwhile, the recognition of the species diversity of AMF of a given area highly depends on the intensity of its sampling due to the aggregated occurrence of spores of these fungi (St. John and Koske 1988) and their seasonal sporulation (Gemma et al. 1989). Additionally, the absence of spores in the field or in early generations of trap cultures does not mean the absence of the arbuscular fungus inside the roots of the sampled plant. Gazea et al. (1992) concluded that sporulation of AMF begins after root colonization level is built up to a threshold value. In Stutz and Morton’s (1996) investigations, 75% of the spores recognized after three propagation cycles were not detected in the first cycle. Therefore, (1) in the year 2000, next soil and root samples were collected from many sites of Israel and (2) the occurrence of AMF in the sites sampled was determined based on examination of many generations of trap cultures with soils and roots collected in both 1997 and 2000.

The aim of this paper is to present all AMF found in Israel to date.

MATERIALS AND METHODS

Study sites

In 1997, most rhizosphere soil and root samples were collected from dunes of the Mediterranean Sea extending from the Kibbutz Shefayim to Tel-Aviv. Two samples were taken near Berscheva in northern Israel, two from a field located in the Negev Desert near the Volcanic Centre, and one from a meadow situated in the Jordan Valley. In 2000, all soil and root samples came from under Ammophila arenaria (L.) Link growing in different sites distributed along the bank of the Mediterranean Sea.

Collection of soil and root samples, establishment of trap cultures, and extraction of spores of AMF

Rhizosphere soils and roots of sampled plants were collected from a depth of 5-30 cm using a small garden shovel. About 100-200 cm³ samples were placed in plastic bags. After their transfer to a laboratory in Poland, they were first stored at 4°C for ca. one month and then used to establish trap cultures. Trap cultures were established to initiate sporulation of AM fungal species rarely sporulating in the field and species that did not produce spores at the time of collection of the field samples. The growing substrata of the trap cultures was the field-collected material mixed with an autoclaved coarse-grained sand coming from maritime dunes adjacent to Świnoujście (pH 6.7; 12 and 26 mg L⁻¹ P and K, respectively; Błaszkowski 1995). The mixtures were placed into 9×12.5-cm plastic pots (500 cm³) and densely seeded with Plantago lanceolata L. Plants were grown in a greenhouse at 15-30°C with supplemental 8-16-h lighting provided by one SON-T AGRO sodium lamp (Philips Lighting Poland S.A.) placed 1 m above pots. The maximum light intensity was 180 μE m⁻² s⁻¹ at pot level. Plants were watered 2-3 times a week. No fertilizer was applied during the growing period. Trap cultures were first harvested four months after plant emergence and then every ca. 6 month until 2006. After each harvest, the cultures were reseeded with P. lanceolata. Spores were extracted by wet sieving and decanting (Gerdemann and Nicolson 1963).

Microscopy survey

Morphological properties of spores and their wall structures were determined based on observation of at least 100 spores mounted in polyvinyl alcohol/lactic acid/glycerol (PVLG; Omar et al. 1979) and a mixture of PVLG and Melzer’s reagent (1:1, v/v). Spores were crushed to varying degrees by applying pressure to the cover slip and then stored at 65°C for 24 h to clear their contents from oil droplets. These were examined under an Olympus BX 50 compound microscope equipped with Nomarski differential interference contrast optics. Microphotographs were recorded on a Sony 3CDD color video camera coupled to the microscope.

Terminology of spore structure is that suggested by Stürmer and Morton (1997) and Walker (1983). Spore colour was examined under a dissecting microscope on fresh specimens immersed in water. Nomenclature of fungi and plants is that of Walker and Trappe (1993) and Mirek et al. (1995), respectively. The authors of the fungal names are those presented at the URL web page http://www.indexfungorum.org/AuthorsOfFungalNames.htm. Specimens were mounted in PVLG on slides and deposited in the Department of Plant Pathology, University of Agriculture, Szczecin, Poland. Newly described species were deposited in the herbarium at Oregon State University in Corvallis, Oregon, USA.

Colour microphotographs of spores and mycorrhizae of the formally described species can be viewed at the URL http://www.agroar.szczecin.pl/~jblaszkowski/.

RESULTS AND DISCUSSION

General data

In the years 1997 and 2000, 49 and 113 rhizosphere soil and root samples were collected, respectively, in different sites of Israel. In 1997, except for two samples taken under Capitium annuum L. cultivated near Berscheva in northern Israel, two from under Lycopersicon esculentum Mill. grown in the Negev Desert near the Volcanic Center, and one from under Festuca rubra L. s. s. growing on an irrigated meadow in the Jordan Valley, all the other soil and root mixtures came from under Oenothera drummondii Hook. Colonizing sand dunes of the Mediterranean Sea extending up to ca. 10 km from Tel-Aviv (Table 1). In 2000, all the...
TABLE 1. Plants examined and numbers* of soil and root samples in which the occurrence of arbuscular mycorrhizal fungi was investigated.

<table>
<thead>
<tr>
<th>Plant species</th>
<th>Locality</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ammophila arenaria (L.) Link</td>
<td>2090-2203</td>
</tr>
<tr>
<td>Oenothera drumondii Hook</td>
<td>1177-188, 1190-1215, 1217-1222, 1224, 1225</td>
</tr>
<tr>
<td>Capsicum annuum L.</td>
<td>1189, 1216</td>
</tr>
<tr>
<td>Lycopersicon esculentum Mill.</td>
<td>1223</td>
</tr>
</tbody>
</table>

* the numbers correspond with the numbering system used by the first author of this paper.

samples collected came from under Am. arenaria growing in different dune sites adjacent to the Mediterranean Sea.

After a continuous cultivation of the rhizosphere soil and root samples collected in pot trap cultures and their examination at least twice a year, spores of AMF were found in 41 and 103 cultures with the 1997 and 2000 soil and root mixtures, respectively. The spores represented a total of 30 species and 8 undescribed morphotypes belonging to 7 genera of the Glomeromycota (Table 2). No representative of the genera Gigaspora and Paraglomus was found. The 1997 cultures hosted 20 species and 5 morphotypes in 5 genera, and those of 2000 20 species and 4 morphotypes in 5 genera. The fungi highly dominating in Israeli soils were members of the genus Glomus (76.1% of all the representatives of the Glomeromycota revealed) with 15 species and 5 morphotypes and 14 species and 4 morphotypes recognized in the 1997 and 2000 cultures, respectively.

The arbuscular fungi most frequently found in trap cultures containing rhizosphere soil and root mixtures collected in 1997 were G. constrictum and an undescribed Glomus 178 (present in at least 20% cultures), followed by Arch. trappei, G. aurantium, G. claroideum, an undescribed Glomus 130, Pac. scintillans, and Scu. diparurescens (found in 10-20% cultures; Table 2). The 2000 cultures most frequently hosted G. aurantium and G. constrictum, then G. coronatum, G. gibbosum, Glomus 178, and Scu. diparure-
sens.

Taking into account the results of the frequency of occurrence of AMF in cultures of both 1997 and 2000, the arbuscular fungi most frequently found in the Israeli soils examined were G. aurantium and G. constrictum, followed by G. coronatum, G. gibbosum, Glomus 178, and Scu. di-
parurescens (Table 2).

Of the fungal species identified in trap cultures with soils and roots collected in 1997, only Arch. trappei, G. constrictum, G. coronatum, G. geosporum, G. microcarpum, G. mossea, and G. sinuosum had earlier been recorded in Israel (Dodd and Krikun 1984; Haas and Menge 1990). Although not revealed by the authors of this paper in trap cultures with the 1997 soils, other AMF found by Dodd and Krikun (1984) and Haas and Menge (1990) were Ac. larvum Gerd. & Trappe, G. calvisporum (Trappe) R.T. Almeida & N.C. Schenck, G. fasciculatum, and G. macrocarpum Tul. & C. Tul.

The species of AMF not revealed in the first cycle of trap cultures with soils and roots sampled in 1997 but found to sporulate in next generations of the same cultures were D. spurca, Scu. fulgida, Scu. pelliculida, and Scu. persica.

TABLE 2. Species of arbuscular mycorrhizal fungi found in Israel.

<table>
<thead>
<tr>
<th>Fungal species</th>
<th>Frequency of occurrence (%)</th>
<th>1997</th>
<th>2000</th>
<th>Mean</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acarospora paulineae</td>
<td>4.1</td>
<td>6.2</td>
<td>5.5</td>
<td></td>
</tr>
<tr>
<td>Archaeospora trappei</td>
<td>14.3</td>
<td>4.3</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diversispora sparsa</td>
<td>4.1</td>
<td>1.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Entrophospora infrequens</td>
<td>1.8</td>
<td>1.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. arenatum</td>
<td>2.0</td>
<td>0.9</td>
<td>1.2</td>
<td></td>
</tr>
<tr>
<td>G. aurantium</td>
<td>18.4</td>
<td>47.8</td>
<td>38.9</td>
<td></td>
</tr>
<tr>
<td>G. caledonum</td>
<td>2.0</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. claroideum</td>
<td>10.2</td>
<td>3.1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. constrictum</td>
<td>24.5</td>
<td>48.7</td>
<td>41.4</td>
<td></td>
</tr>
<tr>
<td>G. coronatum</td>
<td>8.2</td>
<td>16.8</td>
<td>14.2</td>
<td></td>
</tr>
<tr>
<td>G. corymbiforme</td>
<td>2.7</td>
<td>1.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. drummondi</td>
<td>3.5</td>
<td>2.5</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. etunicatum</td>
<td>2.0</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. fasciculatum</td>
<td>0.9</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. geosporum</td>
<td>4.1</td>
<td>1.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. gibbosum</td>
<td>2.0</td>
<td>11.5</td>
<td>8.6</td>
<td></td>
</tr>
<tr>
<td>G. intraradices</td>
<td>4.1</td>
<td>6.2</td>
<td>5.6</td>
<td></td>
</tr>
<tr>
<td>G. microcarpum</td>
<td>4.1</td>
<td>0.9</td>
<td>1.9</td>
<td></td>
</tr>
<tr>
<td>G. mosseae</td>
<td>4.1</td>
<td>6.2</td>
<td>5.6</td>
<td></td>
</tr>
<tr>
<td>G. pastulatum</td>
<td>0.9</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. sinuosum</td>
<td>2.0</td>
<td>1.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. trimeatus</td>
<td>2.0</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. walkeri</td>
<td>0.9</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G. xanthium</td>
<td>4.1</td>
<td>7.1</td>
<td>6.2</td>
<td></td>
</tr>
<tr>
<td>Glomus 126</td>
<td>8.2</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glomus 130</td>
<td>10.2</td>
<td>3.1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glomus 139</td>
<td>2.0</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glomus 149</td>
<td>2.0</td>
<td>1.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glomus 163</td>
<td>5.3</td>
<td>3.7</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glomus 165</td>
<td>0.9</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glomus 178</td>
<td>20.4</td>
<td>15.0</td>
<td>16.7</td>
<td></td>
</tr>
<tr>
<td>Pasticora franciscana</td>
<td>0.9</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pasticora scintillans</td>
<td>14.3</td>
<td>0.9</td>
<td>4.9</td>
<td></td>
</tr>
<tr>
<td>Scutellopsora diparurescens</td>
<td>18.4</td>
<td>13.3</td>
<td>14.8</td>
<td></td>
</tr>
<tr>
<td>S. fulgida</td>
<td>0.9</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>S. pelliculida</td>
<td>1.8</td>
<td>1.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>S. persica</td>
<td>2.7</td>
<td>1.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Scutellopsora 179</td>
<td>0.6</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

In summary, up to 2001, 21 species of AMF were known to occur in Israel (Błaszkowski et al. 2001b; Dodd and Krikun 1984; Haas and Menge 1990). This paper increases this number to 33, of which D. spurca, E. infrequens, G. aurantium, G. corymbiforme, G. drummondi, G. pastulat-
rum, G. trimeatus, G. walkeri, Pac. franciscana, Scu. fulgida, Scu. pelliculida, and Scu. persica were recorded for the first time in this country. Moreover, G. aurantium, G. drummondi, G. walkeri, and G. xanthium have recently been described as new species for science from spores isolated from soils of, e.g., Israel (Błaszkowski et al. 2004; Bła-
szkowski et al. 2006).

The finding of spores of AMF in 83.7% and 91.2% of trap cultures with rhizosphere soil and root mixtures of plants of different sites of Israel collected in the years 1997 and 2000, respectively, confirms earlier suggestions of many authors that members of the phylum Glomeromycota belong to the most commonly occurring soil microorga-
nism in the world. However, literature data indicate that many species of AMF frequently found in the field sporulate rarely or not at all in pot cultures grown in a greenhouse. The reasons may be (1) inappropriate water and air relations (Anderson et al. 1984), pH (Porter et al. 1987), temperature (Koske 1987), and organic content (Koske and Gemma 1997) of pot cultures to initiate spore germination and/or to attain a minimum root colonization level needed to trigger sporulation (GazeY et al. 1992), (2) the lack or shifts in the structure of populations of non-mycorrhizal soil microorganisms reinforcing both spore germination of AMF and colonization of roots of their plant hosts (Bagyaraj 1984), and (3) the incompatibility of the AM fungal species x host plant arrangement used (Read 2002). Thus, the field samples collected by the authors of this paper could have still contained spores not listed here. Therefore, further studies are needed to reveal species of AMF sporulating in the field conditions of Israel.

The marked predominance of members of the genus Glomus in the spore populations of AMF isolated from trap cultures with Israeli soils was not a surprise. Glomus spp. are the most frequently found arbuscular fungi in different regions of the world and fastest adapt to a wide range of physical, chemical, and biological soil conditions (Błaszkowski 1993a; Smith and Read 1997). However, the high plasticity and productivity of Glomus spp. in pot cultures may have suppressed or even eliminate other species of the Glomeromycota functioning in the field. This may explain the exceptionally low proportion of members of the genera Acaulospora and Scutellospora and the lack of species of the genera Gigaspora and Paraglomus in the spore populations of AMF isolated. Gigaspora and Scutellospora spp. prefer warm sandy soils and commonly occur in maritime dunes (Błaszkowski 1993a; b; Koske 1987). Therefore, the recognition of both the real species diversity of communities of AMF and the natural proportions of components of these communities in sites considered in the studies presented here will need the determination of the occurrence of these fungi in both field-collected soil and root samples, as well as in trap cultures with different host plants.

These studies showed that the species of AMF most frequently occurring in the Israeli soils examined were G. auran
tium and G. constrictum, followed by G. coronatum, G. gibusum, an undescribed Glomus 178, and Scu. dipuqu-
rescens. Except for five soil and root samples taken from under cultivated plants, all the others came from dune sites extending along the bank of the Mediterranean Sea. Thus, the common occurrence of these species and their abundant sporulation in pot cultures indicate that they can be used in production of inoculum and, thereby, in protection of endangered or protected dune plants of Israel. Arbucular mycorrhizal fungi facilitate mycorrhizal plants to colonize a site, effectively exploit its resources, and, consequently, make them more competitive for light and space (Smith and Read 1997). Moreover, native autochthonous fungal species or strains usually are best adapted to actual soil and climatic conditions and used as inoculum are more successful (Stahl and Christensen 1991).

The distribution of arbuscular mycorrhizal fungi in Israeli soils and notes on their general occurrence

n=number of soil samples in which a particular fungal species was found. The numbers following are those of soil and roots samples given in Table 1 and correspond with the numbering system used by the first author of this paper.

1. Acaulospora paeiinae Błasz.

n=9: 1218, 1219, 2123, 2128, 2134, 2136, 2137, 2156, 2163.

The only other report of the occurrence of Ac. paeiinae in Israeli soils is that of Błaszkowski et al. (2001b) informing of its association with roots of O. drummondi.

Acaulospora paeiinae probably has a worldwide distribution. It has been found in many cultivated and uncultivated sites of Poland (e.g., Błaszkowski 1993a, b; Jansa et al. (2002) and Oehl et al. (2004) recovered spores of Ac. paeiinae from different soils of Switzerland. Koske et al. (1997) encountered this fungus among roots of Agrostis ca


Morton & D. Redecker

n=7: 1180, 1186, 1187, 1188, 1193, 1203, 1223.

In Israel, Arch. trappei has for the first time been recorded as Arch. trappei by Haas and Menge (1990) and then by Błaszkowski et al. (2001b).

Archaeospora trappei seems to occur in the whole world. Tadych and Błaszkowski (2000) and Błaszkowski et al. (2002) found this fungus in sand dune soils of the Słowiński National Park and the Błędowska Desert located in northern and southern Poland, respectively. This fungus has also been reported from, e.g., Australia, Brazil, Cuba, Japan, South Africa, Scotland, U.S.A. (Morton and Redecker 2001), Germany (Blaschke 1991), Switzerland (Oehl et al. 2005), and China (Gai et al. 2005). Błaszkowski (unpubl. data) isolated spores of Arch. trappei from many trap cultures with soils of Oman, Turkey, Cyprus, Italy, and France.

Spores of Arch. trappei may easily be omitted due to at least three reasons. First, they are very small and hyaline and, hence, difficult to see. Second, their wall consists of thin and delicate layers, which are easy to decompose by soil microorganisms. Third, sporulation of many AMF is seasonal (Gemma et al. 1989) and, therefore, their spores may be absent at the time of collection of field soils; most of the earlier investigations of the occurrence of AMF used only field-collected soil samples.

3. Diversispora spura (C.M. Pfeiff., C. Walker & Bloss)

n=1: 1197.

Diversispora spura has originally been discovered as G. spurae in a greenhouse bed of sand used for propagation of various ornamental plants cultivated in Arizona (Pfeiffer et al. 1996). This fungus has also been found in maritime dunes of Mexico (Pfeiffer et al. 1996), Hawaii (Koske and Gemma 1996), San Miguel Island, California (Koske, pers. comm.), as well as in different other natural and cultivated ecosystems of North America, Cuba and Africa (Kennedy et al. 1999; Stutz and Morton 1996; Stutz et al. 2000), and Poland (Błaszkowski et al. 2003; Iwaniuk and Błaszkowski 2004).
n=2: 2114, 2145.  
*Entrophospora infrequens* has originally been described as *G. infrequens* Hall from spores isolated from Long Bush located in New Zealand (Hall 1977). Ames and Schneider (1979) found identical spores in two celery fields in central California. However, they did not origin blastically at the end of a sporogenous hypha as in *Glomus* spp. but inside the neck of a sporiferous sacculus resembling that of fungi of the genus *Acaulospora*, whose spores develop laterally from such a sacculus. This was the base to erect a new genus, *Entrophospora* R.N. Ames & R.W. Schneid.

Apart from New Zealand and California, *E. infrequens* has also been recorded in many other regions of the U.S. (e.g., Koske and Gemma 1997; Koske and Halvorson 1989; Stahl and Christensen 1982), as well as in Canada (Boyetchko and Tewari 1993), Switzerland (Oehl et al. 2005), Poland (Błaszkowski 1993a, b), Taiwan (Wu and Chen 1986), and China (Gai et al. 2006). Additionally, Błaszkowski (unpubl. data) many times revealed this fungus in trap cultures with dune soils of Oman, Turkey, Cyprus, Italy, France, and Africa.

5. *Gnomus arenarium* Błaszk., Tadych & Madej  
n=1: 1210.  
This paper confirms the first Błaszkowski’s et al. (2001b) report of the presence of *G. arenarium* in Israeli dune soils.

*Gnomus arenarium* has been discovered in maritime dunes of the Baltic Sea adjacent to Świnoujście in northwestern Poland (Błaszkowski et al. 2001a).

n=62: 1203, 1210, 1211, 1215, 1219, 1220, 1221, 1222, 1224, 2093, 2102, 2103, 2106, 2116, 2117, 2120, 2122, 2125, 2126, 2127, 2130, 2132, 2134, 2135, 2136, 2137, 2140, 2143, 2146, 2148, 2149, 2151, 2152, 2158, 2161, 2166, 2168, 2171, 2172, 2173, 2174, 2175, 2176, 2178, 2180, 2182, 2183, 2184, 2185, 2186, 2187, 2188, 2189, 2190, 2192, 2193, 2194, 2195, 2197, 2198, 2200.

The holotype of *G. aurantium* has been selected from spores isolated from a one-species culture established from spores recovered from a trap culture with rhizosphere soil and roots mixture of *O. drummondi* colonizing sand dunes of the Mediterranean Sea located near Tel-Aviv (Błaszkowski et al. 2004). Additionally, this fungus has been found to occur among roots of *Am. arenaria* growing in dunes of Majorca, Spain, Calabrone, Italy (Błaszkowski et al. 2004), Turkey, Cyprus, Greece (Błaszkowski, unpubl. data), as well as in cultivated soils of Iran (Błaszkowski, unpubl. data).

7. *Gnomus caledonium* (Nicol. & Gerd.) Trappe & Gerd.  
n=1: 1225.  
The only earlier record of *G. caledonium* in Israeli soils is that of Błaszkowski et al. (2001b).

*Gnomus caledonium* has a worldwide distribution and has frequently been among the dominating species of AMF of the sites examined (Błaszkowski 1993a). However, this fungus seems to prefer more fertile soils than maritime sand dunes.

n=5: 1181, 1184, 1185, 1191, 1201.

Although *G. claroideum* has many times been isolated from dune soils of Israel sampled in 1997 (Błaszkowski et al. 2001b), none of the samples collected in 2000 contained spores of this fungus.

The distribution of *G. claroideum* is worldwide and it has many times been found in both maritime sand dunes (Koske 1987; Mohankumar et al. 1988; Sylvia 1986) and other cultivated and uncultivated ecosystems (Błaszkowski 2003; Jansa et al. 2002; Schenck and Smith 1982; Vestberg et al. 2005; Zhang and Wang 1992).

9. *Glomus constrictum* Trappe  
n=66: 1178, 1180, 1182, 1183, 1187, 1202, 1208, 1218, 1219, 1220, 1222, 1223, 2090, 2091, 2092, 2093, 2105, 2112, 2120, 2122, 2125, 2133, 2134, 2136, 2139, 2140, 2141, 2144, 2146, 2148, 2149, 2151, 2152, 2153, 2154, 2158, 2160, 2161, 2163, 2166, 2167, 2168, 2170, 2172, 2173, 2174, 2175, 2176, 2177, 2178, 2179, 2181, 2183, 2185, 2186, 2188, 2191, 2192, 2193, 2194, 2195, 2196, 2197, 2199, 2200, 2201.

The first report of the occurrence of *G. constrictum* in Israel is that of Haas and Menge (1990), who revealed spores of this fungus in avocado orchard soils. Błaszkowski et al. (2001b) encountered *G. constrictum* in 12 Israeli soil samples, of which two came from under *L. esculentum*.

The presence of *G. constrictum* spores in almost half the trap cultures with the 2000 Israeli soils collected by the authors of this paper confirms conclusions of, e.g., Błaszkowski (2003) that this fungus occurs in the whole world and frequently dominates in the isolated spore populations of members of the Glomeromycota. In soils of Poland, *G. constrictum* has been first in respect of frequency of occurrence and ranked third considering the proportion of its spores in populations of all spores isolated from 332 soil samples collected from 113 sites (Błaszkowski 1993a). In maritime dunes, this fungal species has been found in, e.g., Quebec, New Brunswick and New Scotia, Canada (Dalpé 1989), U.S.A. (Koske 1987, 1988), Brazil (Stürmer and Bélée 1994), and Poland (Błaszkowski 1993b, 1994).

n=24: 1186, 1215, 1220, 2202, 2095, 2127, 2128, 2130, 2146, 2151, 2152, 2157, 2158, 2159, 2164, 2165, 2166, 2175, 2176, 2178, 2182, 2193, 2195, 2197.

Dodd and Krikun (1984; Dodd, pers. comm.) were the first who identified *G. coronatum* as *G. mosseae* in soils of Israel. Błaszkowski et al. (2001b) found this fungus in four cultures with rhizosphere soils and roots of *O. drummondi* growing near Tel-Aviv.

*Glomus coronatum* has originally been described from spores isolated from around roots of *Anacyclus radiatus* Loisel colonizing a maritime sand dune system near Folonica, Tuscan, Italy (Giovanetti et al. 1991). Błaszkowski (unpubl. data) extracted spores of this fungus from many cultures with maritime dune soils of Turkey, Cyprus, Italy, France, and Africa. The only records of this fungus in non-dune soils are those of Oehl et al. (2005) coming from the Upper Rine Valley extending between Basel (Switzerland), Freiburg (Germany), and Molhouse (France).

The lack of records of *G. coronatum* in ca. 3000 field soils representing different cultivated and uncultivated plants
growing in northern Europe (Błaszkowski, pers. observ.) suggests the occurrence of this fungus to be limited to regions of a warmer climate. According to Pirożyński (1968), temperature is the most important edaphic factor regulating the distribution of fungi in general.

   *n* = 3: 2164, 2183, 2191.

*Glomus corymbiforme* has originally been found in maritime sand dunes of the Baltic Sea adjacent to Świnoujście in north-western Poland (Błaszkowski 1995). Additionally, this fungus has occurred in dunes of the Mediterranean Sea located near Karabucak-Tuzla, Turkey (Błaszkowski, unpubl. data).

12. *Glomus drummondii* Błaszk. & C. Renker
   *n* = 4: 2092, 2104, 2111, 2131.

*Glomus drummondii* has recently been described based on spores recovered from trap cultures with rhizosphere soils of plants of maritime dunes of Cyprus, Greece, Poland, Spain, and Portugal (Błaszkowski et al. 2006). Thus, the maritime dunes of Israel are the next habitats of the presence of this fungus.

   *n* = 1: 1216.

In Israel, *G. etunicatum* has been associated with roots of *C. annuum* cultivated near Berscheva in 1997 (Błaszkowski et al. 2001b).

*Glomus etunicatum* is a common inhabitant of cultivated and uncultivated soils of different regions of the world (Błaszkowski 1993a).

   C. Walker & Koske
   *n* = 1: 2098.

Haas and Menge (1990) were the first to record *G. fasciculatum* in soils of Israel.

Literature data indicate that *G. fasciculatum* occurs in the whole world and is well adapted to different soils (Błaszkowski 2003; Walker and Koske 1987). However, according to Walker and Koske (1987), many reports of this fungus may have regarded other species of AMF due to its incomplete original description.

15. *Glomus geosporum* (Nicol. & Gerd.) C. Walker
   *n* = 2: 1188, 1189.

The first report of the occurrence of *G. geosporum* in Israel has been that of Haas and Menge (1990), who recovered spores of this fungus from among roots of *Persea americana* Mill. Later, Błaszkowski et al. (2001b) revealed *G. geosporum* associated with roots of *O. drummondii* and *C. annuum* growing in dunes near Tel-Aviv and a cultivated field near Berscheva in northern Israel.

Literature data suggest that *G. geosporum* occurs in the whole world and is adapted to conditions of different cultivated and uncultivated soils, including conditions of maritime dunes (e.g., Błaszkowski 1993a, b; Rose 1980).

   *n* = 15: 1182, 2106, 2107, 2108, 2118, 2119, 2125, 2131, 2154, 2155, 2157, 2160, 2161, 2176, 2187.

In Israel, *G. gibbosum* has for the first time been found in samples coming from under *O. drummondii* growing in dunes located near Tel-Aviv in 1997 (Błaszkowski et al. 2001b). Examination of the 2000 soil-root samples revealed this fungus to co-occur frequently with maritime dune plants of this country.

The original description of *G. gibbosum* has been prepared based on spores isolated from maritime dunes adjacent to Świnoujście in north-western Poland (Błaszkowski 1997). Błaszkowski (unpubl. data) also found this fungus in dunes of the Mediterranean Sea of Turkey, Cyprus, and Italy.

   *n* = 9: 1183, 1217, 2098, 2099, 2101, 2107, 2112, 2115, 2196.

The Błaszkowski’s et al. (2001b) finding of *G. intraradices* spores in two trap cultures with rhizosphere soils of *O. drummondii* was the first record of this fungus in Israel.

*Glomus intraradices* has frequently been recorded from maritime dunes of different regions of the world (see Błaszkowski et al. 2001b).

   *n* = 3: 1184, 1196, 2142.

In Israel, *G. microcarpum* has for the first time been found in soils of an avocado orchard (Haas and Menge 1990). Examination of trap cultures with rhizosphere soil and root samples of *O. drummondii* collected by the authors of this paper in 1997 and 2000 showed this fungus to be an inhabitant of dune sites of this country as well.

*Glomus microcarpum* is widely distributed in the world. It has earlier been encountered in many both cultivated and uncultivated sites (Błaszkowski 1993a), including maritime dunes of, e.g., the Baltic Sea, Poland (Błaszkowski 1993a, b, 1994), Madras, India (Mohankumar et al. 1988), and Italy (Puppi and Riess 1987).

19. *Glomus mossea* (Nicol. & Gerd.) Gerd. & Trappe
   *n* = 9: 1212, 1223, 2090, 2100, 2101, 2105, 2111, 2121, 2197.

The first report of the occurrence of *G. mossea* in Israel is that of Haas and Menge (1990), who found it associated with roots of cultivated *P. americana*. Later, Błaszkowski et al. (2001b) isolated spores of this fungus from under *L. esculentum* cultivated in the Negev Desert near The Volcanic Center. The studies presented here show that *G. mossea* also frequently co-occurred with maritime dune plants of Israel.

*Glomus mossea* is one of the most frequently recorded AMF in different regions of the world. In Poland, this species has been the second most frequently encountered AM fungus (Błaszkowski 1993a). However, it occurred three times more frequently in cultivated than uncultivated soils.

20. *Glomus pastulatum* Koske, Friese, C. Walker & Dalpé
   *n* = 1: 2144.

This paper is the first report of the presence of *G. pastulatum* in Israel. This fungus was revealed only in samples of dune soils collected in 2000.

*Glomus pastulatum* has originally been described from spores isolated from under *Am. breviligulata* Fern. colonizing maritime dunes of Rhode Island, U.S.A. (Koske et al. 1986). Additionally, this fungus has been identified in maritime dunes of Canada (Dalpé 1989), Poland (Błaszkowski...
1993a, b, 1994), and India (Kulkarni et al. 1997; Mohankumar et al. 1988). There is no literature report of the occurrence of *G. pustulatum* in non-dune sites.

n=1: 1177.
   The first record of *G. sinuosum* in Israel is that from the Negev Desert (Dodd and Krikun 1984). Later, Haas and Menge (1990) found this fungus under avocado, and Blaszkowski et al. (2001b) under *O. drummondii* growing in dunes near Tel-Aviv.
   *Glomus sinuosum* probably occurs in the whole world and is adapted to different soils, including dune soils (e.g., Almeida and Schenck 1990; Koske 1988; Wu 1993).

22. *Glomus trimurales* Koske & Halvorson
n=1: 1182.
   The only earlier finding of *G. trimurales* in Israel is that of Blaszkowski et al. (2001b). This fungus, characterized as *Glomus* 131, was revealed under *O. drummondii* growing in dunes near Tel-Aviv.
   *Glomus trimurales* has originally been characterized based on spores isolated from under *Abronia chamissonis* var. *bipinnatisepta* (Less.) Greene growing in maritime dunes of San Miguel Island (Koske and Halvorson 1989), although this fungus has earlier also been identified in maritime dunes of New Jersey, Maryland, and Virginia (Koske 1987).
   In Europe, *G. trimurales* has for the first time been encountered in dunes of the Baltic Sea adjacent to Świnoujście in north-western Poland (Blaszkowski et al. 2003).

23. *Glomus walkeri* Blaszk. & C. Renker
n=1: 2099.
   *Glomus walkeri* has recently been described based on spores isolated from a one-species culture established from spores isolated from a trap culture with a rhizosphere soil and root mixture of *O. drummondii* growing in dunes of the Mediterranean Sea near Tel-Aviv (Blaszkowski et al. 2006). Additionally, this fungus occurred in maritime dunes of Majorca, Spain and Calambrone, Italy (Blaszkowski et al. 2006).

n=10: 1220, 1222, 2092, 2108, 2120, 2122, 2126, 2130, 2198, 2200.
   Spores of *G. xanthium* were found in trap cultures with Israeli dune soils collected in both 1997 and 2000.
   *Glomus xanthium* has for the first time been revealed in a trap culture with a soil and root mixture taken under *Xanthium cl. spinosum* L. colonizing maritime dunes located near Verico in northern Greece (Blaszkowski et al. 2004). Later, this fungus has been encountered in dunes of the Mediterranean Sea adjacent to Karabucak-Tuzla, Turkey, Calambrone, Italy, and Majorca, Spain (Blaszkowski et al. 2004).
   Blaszkowski (unpubl. data) also found this fungus in dunes of Cyprus, France and Africa, as well as in cultivated soils of Iran. The lack of any finding of *G. xanthium* in ca. 3000 soil samples coming from northern Europe suggests it to be limited to regions of a warmer climate.

25. *Glomus 126*
   n=4: 1177, 1180, 1182, 1188.
   Spores of *Glomus 126* have been revealed in trap cultures with rhizosphere soils and roots sampled in both 1997 and 2000.
   The morphological characters of spores of this fungus have been presented previously (Blaszkowski et al. 2001b).

26. *Glomus 130*
   n=5: 1179, 1180, 1182, 1186, 1217.
   *Glomus 130* probably frequently occurs in dunes of the Mediterranean Sea of Israel as showed results of inspections of trap cultures established in both 1997 and 2000.
   The description and illustrations of morphological properties of *Glomus 130* spores have been presented by Blaszkowski et al. (2001b).

27. *Glomus 139*
   n=1: 1219.
   Spores of *Glomus 139* occurred only in the 1997 cultures.
   The diagnostic characters of this fungus have earlier been presented (Blaszkowski et al. 2001b).

28. *Glomus 149*
   n=1: 1198.
   Spores single in the soil; hyaline to white; globose to subglobose; (38-150(65) μm diam (Fig. 1). *Spore wall* with two layers (layers 1 and 2). Layer 1 evanescent, at first smooth, then roughened, hyaline (0.5-0.8(-1.2) μm thick, rarely present in mature spores. Layer 2 laminate, smooth, hyaline to white, (2.3-6.2(-7.5) μm thick. Layers 1 and 2 not reacting in Melzer’s reagent.
   The species forming spores most similar to those of *Glo- mus 149* is *G. minutum* Blaszk. et al. Both fungi produce colourless spores of a similar size (Blaszkowski 2000). However, most spores of *G. minutum* occur in loose clusters usually associated with its host plant roots after their washing away from the soil, whereas those of *Glomus 149* always occurred singly in the soil. Additionally, the outermost spore wall layer of *Glomus 149* sloughs with age, and that of *G. minutum* is a permanent structure.

29. *Glomus 163*
   n=4: 2109, 2141, 2157, 2174.
   Spores formed in clusters of 10 to more than 50; hyaline; globose to subglobose; (40-865(-80) μm (Fig 2). *Spore wall* consists of three layers (layers 1-3; Fig. 3). Layer 1 evanescent, smooth to roughened, hyaline, (0.5-0.9(-1.3) μm thick. Layer 2 laminate, smooth, hyaline, (2.2-6.3(-7.0) μm thick. Layer 3 membranous, ca. 0.5-0.7 μm thick. Only layer 3 stains pinkish red in Melzer’s reagent (Fig. 3).
   The only other species whose spores occur in clusters and resemble in colour and size those of *Glomus 163* is *G. minutum* (Blaszkowski 2000). Examination of crushed spores in a mixture of PVLG and Melzer’s reagent readily separates these fungi. The spore wall of the latter species does not possess the innermost flexible spore wall layer of the former fungus.

30. *Glomus 165*
   n=1: 2102.
   Spores single in the soil; red brown; (100-150(-210) μm diam (Fig. 4). *Spore wall* composed of two layers (lay-
Fig. 1. Spores of *Glomus* 149 with spore wall layers 1 and 2 (sw1 and sw2). Fig. 2. Cluster of spores of *Glomus* 163. Fig. 3. Spore wall layers 1–3 (sw1–3) of *Glomus* 163 with layer 3 (sw3) stained in Melzer’s reagent. Fig. 4. Two-layered spore wall (sw1 and sw2) of *Glomus* 165. Fig. 5. Spores of *Glomus* 178. Fig. 6. Wall layers 1–4 (sw1–4) of a crushed spore of *Glomus* 178. Fig. 7. Spores of *Scutellospora* 179. Fig. 8. Spore wall layers 1 and 2 (sw1 and sw2) and germination wall layers 1 and 2 (gwl1 and gwl2) of *Scutellospora* 179. Figs. 1, 5, and 7: spores mounted in lactic acid; Figs 2–4, 6, and 8: spores in a mixture of PVLG and Melzer’s reagent. Bars: Figs 1, 2, 5=20 µm; Figs 3, 4, 6, 8=10 µm; Fig. 7=50 µm.

ers 1 and 2). Layer 1 evanescent, hyaline, (0.8–1.5)(2.1) µm thick (Fig. 4). Layer 2 laminate, red brown, (3.5–8.9 (-11.0) µm thick. *Subtending hypha* flared or slightly funnel-shaped, (14.0–21.5)(24.5) µm wide at the spore base, occluded by a curved septum continuous with the innermost lamina of spore wall layer 2.

The species of AMF most resembling *Glomus* 165 is *G. coronatum*. Both fungi form spores of a similar size and colour, as well as their wall consists of two layers of identical phenotypic properties (Błaszkowski 2003). The main character separating the two fungi is the width of their subtending hypha. The subtending hypha of *G. coronatum* is much wider (28.5–40.0 µm wide; Błaszkowski 2003) than that of spores of *Glomus* 165 (14.0–24.5 µm wide).
31. *Glomus 178*

n=30: 1177, 1179, 1180, 1182, 1188, 1194, 1201, 1202, 1211, 1217, 1218, 2117, 2124, 2132, 2137, 2139, 2144, 2153, 2155, 2160, 2163, 2170, 2175, 2178, 2183-2185, 2190, 2194, 2198.

Spores single in the soil; hyaline; globose to subglobose; (45.0)-65.5-(70.5) μm diam (Fig. 5). Spore wall consists of four layers (layers 1-4; Fig. 6). Layer 1 evanescent, hyaline, <0.5 μm thick, usually completely sloughed even in young spores. Layer 2 permanent, hyaline, 0.5-0.8 μm thick, frequently separating from layer 3. Layer 3 laminate, hyaline, (3.8-5.6)-(7.7) μm thick. Layer 4 flexible, hyaline, usually tightly adherent to the inner surface of layer 3. None of spore wall layers 1-3 reacts in Melzer’s reagent.

Of the described species of AMF, *Glomus 178* somewhat resembles *G. diaphanum* J.B. Morton & C. Walker, *G. lucidum* Blasz., and *Paraglomus occultum* (C. Walker) J.B. Morton & D. Redeker. Spores of all these fungi are glo- moid, hyaline, and their size range more or less overlaps (Błaszkowski 2003; Morton 2002). However, only *G. diaphanum* has an innermost flexible spore wall layer similar to that of *Glomus 178*. In contrast to the four-layered spore wall of *Glomus 178*, that of *G. diaphanum* consists of three layers, lacking the spore wall layer 2 of *Glomus 178*.

32. *Pacispora franciscana* Sieverd. & Oehl

n=1: 2145.

In this study, spores of *Pac. franciscana* have been found in one trap culture with a rhizosphere soil and roots of *Am. arenaria* sampled only in 2000.

Although recently described, *Pac. franciscana* probably has a worldwide distribution. This fungus has originally been described from spores isolated from a grassland with olive trees growing in Umbra, Italy (Oehl and Sieverding 2004). The same mycologists also encountered this fungus in the High Alpines of Eastern Switzerland. Earlier, *Pac. franciscana* has probably been reported as the “white reticu- late spore” by Mosse and Bowen (1968) in Australia and the “white smooth-walled azygospore” in Libyan and the Negev Desert soils by El Ghami et al. (1976) and Dodd and Krikun (1984), respectively. In Poland, *Pac. franciscana* has for the first time been found associated with roots of *Lupinus luteus* L. cultivated in the north-west in 1985. Lat- ter, spores of this fungus have been isolated from under many cultivated and uncultivated plant species growing in different regions of Poland (Błaszkowski, unpubl. data). Additionally, Błaszkowski (unpubl. data) found *Pac. franciscana* spores to occur among roots of *Am. arenaria* colonizing sand dunes of the Mediterranean Sea located near Karabucak-Tuzla, Turkey.

33. *Pacispora scintillans* (S.L. Rose & Trappe) Sieverd. & Oehl

n=8: 1180, 1182, 1185, 1187, 1203, 1221, 1222, 2177.

The relatively frequent presence of spores of *Pac. scintillans* in trap cultures established in both 1997 and 2000 in- dicates this fungus to be rather a common inhabitant of soils of Israel.

*Pacispora scintillans* has originally been described as *G. scintillans* from spores recovered from under *Cereocarpus ledifolias* Nutt. growing in Oregon, U.S.A. (Rose and Trappe 1980). Later, this fungus has been found in other states of the U.S. (Halvorson and Koske 1987; Walker et al. 2004), Switzerland (Oehl et al. 2004), Germany, United Kingdom, Poland, Turkey and Australia (Walker et al. 2004), as well as in China (Gai et al. 2006), where it was associated with both cultivated and uncultivated plants.

34. *Scutellospora dipurpureascens* J.B. Morton & Koske

n=25: 1180, 1182, 1187, 1189, 1203, 1210, 1216, 1219, 1221, 2097, 2108, 2118, 2130, 2138, 2140, 2143, 2152, 2157, 2158, 2166, 2171, 2172, 2185, 2186, 2198.

The results of investigations of the authors of this paper indicate *Scu. dipurpureascens* to be a frequent component of spore populations of AMF of dune soils of Israel.

*Scutellospora dipurpureascens* has been discovered in a rhizosphere soil of *Festuca arundinacea* Schreb. growing in West Virginia (Morton and Koske 1988). This fungus has also dominated or has been one of the more frequently occurring species of AMF in both maritime and inland sand dunes of Poland (Błaszkowski et al. 2001b). Addition- ally, *Scu. dipurpureascens* has occurred in non-dune sites of Canada (Marcel et al. 1988), Mexico (Estrada-Tores et al. 1992), and the Netherlands (Griffioen 1994).

35. *Scutellospora fulgida* Koske & C. Walker

n=1: 2177.

Although *Scu. fulgida* has not been mentioned to occur in the Israeli soils collected in 1997 (Błaszkowski et al. 2001b), later examination of next generations of the same trap cultures revealed the presence of this species.

*Scutellospora fulgida* has originally been described based on spores isolated from under *Am. breviligulata* growing in maritime dunes of Virginia, U.S.A. (Koske and Walker 1986). Other sites of its occurrence cited in the literature are maritime dunes extending from New Jersey to Virginia (Koske 1987), maritime dunes of Florida (Sylvia and Will 1988), and soils of the south and east coasts of China (Gai et al. 2006). Recently, Błaszkowski (unpubl. data) found spores of *Scu. fulgida* in the Mediterranean Sea dunes adja- cent to Calambrone, Italy. Thus, the sites of occurrence of this fungus listed above suggest *Scu. fulgida* to be restricted to regions of a warmer climate.

36. *Scutellospora pellucida* (Nicol. & N.C. Schenck) C. Walker & F.E. Sanders

n=1: 2092.

Similarly as *Scu. fulgida*, *Scu. pellucida* has not sporula- ted in the first generation of trap cultures with Israeli soils sampled in 1997 (Błaszkowski et al. 2001b). However, fur- ther cultivation of these cultures and examination of trap cultures with the 2000 soils revealed this species to occur in dunes of the Mediterranean Sea of Israel.

*Scutellospora pellucida* has been discovered in cultivated soils of northern and central Florida (Nicolson and Schenck 1979). Literature data indicate that this fungus occurs in the whole world in both cultivated and non-dune and dune natural ecosystems (e.g., Oehl et al. 2004; Koske and Gen- ma 1997; Sieverding 1989; Saito and Vargas 1991; Błaszk-owski 1993a, b; Gai et al. 2006).

37. *Scutellospora persica* (Koske & C. Walker) C. Walker & F.E. Sanders

n=3: 2177, 2189, 2190.

*Scutellospora persica* is the third species not revealed in the first cycle of trap cultures with soils collected in 1997.
but this fungus was found to sporulate in both older cultures with the 1997 soils and the first generation of cultures representing the 2000 year.

The original description of *Scu. persica* has been prepa-
red based on spores isolated from a barrier dune located in
New Jersey, U.S.A., although this fungus has also occurred in coastal sand dunes of the east coast of the U.S.A. from
Massachusetts to Virginia (Koske and Walker 1985). Apart from the U.S., this fungal species has also been recorded in
dunes and other soils in Brazil (Grandi and Trufem 1991),
Italy (Puppi et al. 1986), Poland and Greece (Blaszkowski and Tadych 1997), as well as in China (Gai et al. 2006).

38. *Scutellospora* 179

Spores single in the soil; capsicum red to red brown;
250–400 µm diam (Fig. 7). Spores with two walls, a sora
wall and an inner germination wall. *Spore wall* consists
of two layers (layers 1 and 2; Fig. 8). Layer 1 permanent,
yellow brown, 0.5–0.9 µm thick, ornamented with small warts,
0.5–0.8 µm high. Layer 2 laminate, capsicum red to red
brown, 4.0–11.5 µm thick. *Germination wall* composed of
two hyaline layers (layers 1 and 2; Fig. 8). Layer 1 flexible,
0.5–0.8 µm thick, frequently adherent to layer 2. Layer 2
flexible to semi-flexible, 1.0–1.8 µm thick, staining pinkish
red in Melzer’s reagent.

Spores of *Scutellospora* 179 are most reminiscent of tho-
se of *Scu. persico* (Koske & C. Walker) C. Walker & F.E.
Sanders in size and in having a warty outer spore wall lay-
er forming the spore surface. The only character distin-
guishing the two fungi is colour of their spores. Spores of *Scu-
tellospora* 179 are darker (capsicum red to red brown) than
those of *Scu. persica* (sunflower yellow (4A7) to apricot
yellow; Blaszkowski 2003).

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