

SHORT NOTE

A. Rashid · T. Ahmed · N. Ayub · A. G. Khan

Effect of forest fire on number, viability and post-fire re-establishment of arbuscular mycorrhizae

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Abstract Forest fire can affect arbuscular mycorrhizal (AM) fungi by changing the soil conditions and by directly altering AM proliferation. We studied the effects of a severe forest fire at Margalla Hills near Islamabad on the number and viability of AM fungal propagules in the burnt soil and their role in the re-establishment of post-fire infection in colonized plants. Compared with a nearby control area, the burnt site had a similar number of total spores but a lower number of viable AM fungal propagules. The roots of the two most frequent species at the burnt site, *Dodonaea viscosa* and *Aristida adscensionis*, showed a gradual increase in percentage root length colonized by AM fungi in general and hyphal infection in particular. Our results indicate resumption of mycorrhizal activity following the fire, probably from AM hyphae in the roots of these dominant shrubs.

Key words Arbuscular mycorrhiza · Forest fire · Revegetation · MPN

Introduction

The importance of mycorrhiza, especially arbuscular mycorrhiza (AM) caused by Glomalean fungi, in the rehabilitation of severely disturbed ecosystems is well recognized (Miller 1979; Reeves et al. 1979; Khan 1981; Trappe 1987). The reduction and/or loss of infectivity of AM fungi as a result of natural or man-induced disturbance of ecosystems has been reported by various

researchers (Allen and Allen 1980; Powell 1980; Allen et al. 1984).

Annual forest fires are a typical feature of Margalla Hills near Islamabad, Pakistan. The role of fire in changing vegetation structure and soil characteristics has been the subject of a number of studies (Naveh 1973, 1975; Trabaud 1984, 1987; Johnson 1992; Kutiel and Shaviv 1993). Fire can dramatically change surface-soil characteristics and erosion rates (Anderson 1974; Amaranthus and McNabb 1984; Amaranthus 1989). Although several workers have reported significant effects of fire on soil microorganisms in general (Algren and Algren 1960; Perry et al. 1987), few studies have examined the effect of burning on AM fungi. The degradation of soil causes changes in the AM fungal population, reflected by a decrease in the number of AM fungal propagules (Allen et al. 1984; Day et al. 1987; Khan 1978) and hence a low inoculum potential (Khan 1988). Vilarino and Arines (1991) found that burning of vegetation had serious effects on both the spore population and the infection potential of AM fungi.

Our objectives were to determine the effects of forest fire on the number and viability of AM fungal propagules in affected areas of Margalla Hills, including post-fire re-establishment of AMF infection in the dominant shrubs of the burnt sites.

Materials and methods

The study was conducted on the south-facing slopes of Margalla Hills which constitutes the northwest boundary of Islamabad, Pakistan. The area of 12 605 ha is scrub type vegetation and the summer maximum temperature reaches 44 °C (Hoefflaken 1988). The vegetation on the burnt and the nearby control sites was identical before the fire and the sites experienced the same climatic conditions. In fact the demarcation of the sites was only possible after the fire. *Dodonaea viscosa* and *Artisia adscensionis* were among the dominant shrubs of both areas, covering 11.6 and 9.2%, respectively, of the control site and 2.3 and 4.3% of the burnt site.

Fire broke out at the end of July 1994 and plant and soil samples were collected in December 1994 and April 1995. At each

A. Rashid · T. Ahmad · N. Ayub
Department of Biological Sciences, Quaid-i-Azam University,
Islamabad, Pakistan

A. G. Khan (✉)
Department of Biological Sciences, University of Western
Sydney-Macarthur, P.O. Box 555,
Campbelltown NSW 2560, Australia
Fax: +46-203025; e-mail: a.khan@uws.edu.au

collection date, soil (sandy loam) samples, plants and their mycorrhizospheres were collected. Five uniformly spaced soil cores (4 cm wide × 10 cm deep) were taken from each site on each occasion and pooled to give two composite samples for extraction of AM fungal spores by wet sieving and decanting (Gerdeemann and Nicolson 1963). The composite soil samples were sieved with a 2-mm sieve to remove coarse debris and oven-dried (80°C overnight). Ca. 100 gm of dried soil sample was then thoroughly wetted for 1 h before sieving and decanting for AM fungal spore extraction.

The numbers of viable AM fungal propagules (spores, mycelium, infected root pieces) per 100 g of dried soil sample were determined by the most probable number (MPN) technique (Porter 1979, as modified by Woomey 1994), using pre-germinated red clover seedlings as the test plant. A tenfold dilution series of each mycorrhizosphere sample with acid-washed sand as diluent was planted with 6-day-old pre-germinated red clover seedlings in 50-ml vials with 1 seedling per vial and 5 replicates per dilution. The vials were maintained at field capacity by frequent watering to a constant weight and were randomized in trays in a glasshouse (temperature range 13–25°C and photoperiod 14 h). After 6 weeks, clover seedlings were harvested and their roots removed, washed and fixed in FAA (formalin, acetic acid, ethanol 5:5:90) and stored at 4°C. The MPN of AM fungal propagules for each inoculum were calculated according to Alexander (1965).

Roots of *Dodonaea viscosa* and *Aristida adscensionis* were collected by excavating the whole root systems of five plants per species. The roots were carefully washed to remove soil debris, cut into 1-cm pieces, fixed in FAA, and cleared (2.5% KOH) and stained (acid glycerol aniline blue) according to Koske and Gemma (1989).

The data was subjected to one-way ANOVA and χ^2 test.

Results and discussion

Our results indicate adverse effects of fire on the viability of AM fungal propagules (Table 1). The MPN method revealed a significantly lower number of viable AM fungal propagules at the burnt site than at the control site, suggesting that burning of the vegetation and subsequent loss of top soil by erosion reduced infectivity of AM fungal propagules. Moreover, direct effects of burning may have resulted in heat injury causing low viability and/or dormancy of AM fungal spores; the MPN bioassay may fail to detect such dormant spores. Dead AM fungal spores degrade slowly and the proportion of vital spores is often about half that of total spores (An et al. 1990). Similarly, the presence of propagules of AM fungi such as *Glomus intraradices*,

which produces spores inside roots and are not recovered in wet sieving, will be revealed in the MPN bioassay plants but are unlikely to be isolated by the wet-sieving technique (An et al. 1990). Reduced AM fungal spore viability at the burnt site in the present study may also be attributed to toxic substances released from organic matter as a result of fire (Raison 1979).

The total number of spores at the burnt site, viable or not, was not significantly different from the control-site 5 months after the fire (Table 1). These results are consistent with those of other workers (Vilarino and Arines 1991). At 10 months after the fire, rhizosphere soil from the control site did not contain a higher number of total AM fungal spores than at the burnt site. This contrasts with results of Dhillon et al. (1988), who recorded an increase in spore production with the reduced growth of little bluestem subjected to a prairie fire in the USA. No speciation differences in the AM fungal spore populations were noted (Table 2). In general the number of spores of *Glomus* sp. was greater than *Gigaspora gigantea* and *Gigaspora margarita* at both sites.

Examination of the roots of *Dodonaea viscosa* and *Aristida adscensionis* from the burnt site showed the gradual re-establishment of AM associations (Table 3). In December 1994, 4 months after the fire, the percentage root length colonized by AM fungi was significantly higher ($P < 0.05$) at the control site for all samples than at the burnt site for both plants studied. A significant increase in root length colonized by AM fungal hyphae in plants growing at the burnt site was found in April 1995, 10 months after the fire. It has been demonstrated by various workers that hyphal regrowth from root segments occurs more frequently and more rapidly than from spores, where infection is delayed by the time required for spores to germinate and pass through a highly branched phase before becoming infective (Powell 1976; Hayman and Stovold 1979; Hall 1988). Zakha et al. (1995) studied the mycorrhizal inoculum potentials in sugar maple forests and suggested that colonized root pieces and hyphal fragments are more likely to be propagules for infection than spores. Other researchers (Powell 1976; Abbott and Robson 1981) suggested that

Table 1 Number of viable arbuscular mycorrhizal (AM) propagules and spores in burnt and control forest sites as determined by MPN and wet-sieving techniques on two sampling dates. Mean values in columns within a sampling followed by different letters are significantly different at $P < 0.05$ ($n = 2$)

Method	Site	December 1994	April 1995
MPN	Burnt	93 ^a	105 ^a
	Control	156 ^b	203 ^b
Wet-sieving technique	Burnt	311 ^a	322 ^a
	Control	298 ^a	382 ^a

Table 2 Numbers of spores of AM fungal species recovered per 100 g soil from burnt and control sites by wet sieving and decanting ($n = 10$). The data are mean results from 10 plants ± SD

Species	December 1994		April 1995	
	Control	Burnt	Control	Burnt
<i>Glomus</i> sp.	49 ± 13	37 ± 21	89 ± 17	71 ± 12
<i>Glomus fasciculatum</i>	60 ± 25	84 ± 32	67 ± 16	45 ± 19
<i>Glomus mosseae</i>	101 ± 26	98 ± 28	110 ± 31	94 ± 30
<i>Gigaspora gigantea</i>	50 ± 11	41 ± 19	83 ± 9	55 ± 21
<i>Gigaspora margarita</i>	9 ± 3	22 ± 12	12 ± 2	34 ± 8
<i>Acaulospora</i> sp.	18 ± 5	24 ± 7	7 ± 2	14 ± 3
Unknown	11 ± 7	5 ± 1	14 ± 1	9 ± 1
Total	298 ± 77	311 ± 54	382 ± 41	322 ± 38

Table 3 Mean \pm SD percentage root length colonization by AM fungi of plants from burnt and the control sites. Mean values in a line, within a sampling, followed by different letters are significantly different at $P < 0.05$ ($n = 10$)

Host	Form of infection	December 1994		April 1995	
		Control	Burnt	Control	Burnt
<i>Dodonaea viscosa</i>	Total	57 ^b \pm 8	23 ^a \pm 3	99 ^b \pm 13	47 ^a \pm 3
	Hyphal	35 ^b \pm 11	14 ^a \pm 3	61 ^b \pm 19	26 ^a \pm 7
	Vesicular	14 ^a \pm 6	8 ^a \pm 2	26 ^b \pm 9	16 ^a \pm 8
	Arbuscular	8 ^b \pm 2	1 ^a \pm 2	11 ^b \pm 4	5 ^a \pm 3
<i>Artisia adscensionis</i>	Total	43 ^b \pm 3	20.7 ^a \pm 2	45 ^b \pm 6	29 ^a \pm 5
	Hyphal	28 ^b \pm 5	15 ^a \pm 2	31 ^b \pm 6	14 ^a \pm 3
	Vesicular	12 ^b \pm 2	5 ^a \pm 2	11 ^a \pm 3	13 ^a \pm 2
	Arbuscular	3 ^b \pm 1	0 ^a \pm 1	3 ^a \pm 1	2 ^a \pm 1

infectivity of AM is probably related to the amount of nutrient reserves in fungal structures in roots, and hypothesized that intra-radical vesicles contribute greatly to inoculum potential. Mosse (1988) and Burggraaf and Beringer (1989) also reported that vesicles separated from roots retain their infectivity, and suggested that mycorrhizal root pieces are preferable to germinating spores.

To establish the relationships of mycorrhizal activity, time following fire and soil edaphic factors, both the dynamics of succession of vegetation and associated AM fungal populations on the Margalla Hills need to be examined further (Janos 1980). Mycorrhizal dependence and AM fungal propagule availability are prime determinants of early stages of community development and affect long-term plant dominance (Allen and Allen 1990; Molina et al. 1992).

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