SOIL MACROFAUNA IN COVER CROPS OF FIGS GROWN UNDER ORGANIC MANAGEMENT

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ABSTRACT: Soil fauna plays an important role in organic management through their effects on soil organic decomposition, nutrient mineralization, and amelioration of the soil's physical properties. This work evaluates the density and diversity of the soil macrofauna under types of cover plants in areas cultivated with *Ficus carica* L. under organic management. The soil macrofauna was collected in 0.25×0.25 m areas, down to a soil depth of 0.3 m, and at the surface layer. The treatments consisted of bahiagrass living mulch (*Paspalum notatum*), siratro living mulch (*Macroptilium atropurpureum*), and bahiagrass mulch. The highest macrofauna density and the lowest diversity were observed in bahiagrass, of which 80% were represented by ants, thus characterizing the soil under this cover crop as showing the lowest functional diversity and quality. Key words: *Ficus carica* L, soil fauna, soil quality indicators

MACROFAUNA EDÁFICA SOB COBERTURAS DO SOLO NO CULTIVO DO FIGO SOB MANEJO ORGÂNICO

RESUMO: A fauna do solo tem papel importante em sistemas orgânicos, através dos seus efeitos na decomposição da matéria orgânica, mineralização de nutrientes e condicionamento físico do solo. Este trabalho avaliou a densidade e a diversidade da macrofauna edáfica sob tipos de cobertura do solo em áreas cultivadas com *Ficus carica* L. sob manejo orgânico do solo. A macrofauna foi coletada em áreas de 0,25 × 0,25 m e amostraram-se as camadas de material vegetal e solo até a profundidade de 0,3 m, nos meses de março e setembro de 2001. Os tratamentos constaram de cobertura viva com grama batatais (*Paspalum notatum*), cobertura viva de siratro (*Macroptilium atropurpureum*) e cobertura morta formada por palha de grama batatais. A mais alta densidade da macrofauna e a mais baixa diversidade foi observada em grama batatais, sendo mais de 80% dessa fauna de formigas, o que caracteriza o solo sob influência desta cobertura como o de mais baixa diversidade funcional e qualidade, comparado às outras coberturas.

Palavras-chave: Ficus carica L, fauna do solo, indicadores de qualidade do solo

INTRODUCTION

Soil organic management represents a very promising alternative for the sustainable production of foods, satisfying a demand that comes from all sectors of society. The use of legume plants as living mulches for the soil fits well within this context, since it is a technique that contributes toward maintaining soil organic matter and incorporating atmospheric N, in addition to protecting the soil against erosive processes. However, little is known about the impact sustained by the soil fauna as a consequence of the use of these cover plants under organic management.

The soil fauna populations can be markedly affected by soil cover, as the availability of energy associated with the existence of new habitats favorable to colonization increases, thus influencing density and diversity of nearly all groups in this particular fauna (Tian et al., 1999). Both the soil's microorganisms and fauna can

modify the physical and chemical properties of the soil (Pankhurst & Lynch, 1994), and assume an essential role in the soil processes under organic management. This aspect has been considered as one of the key processes for maintaining the structure and fertility of tropical soils (Lavelle et al., 1993).

Among the animals that make up the soil fauna, the edaphic macrofauna comprises the largest invertebrates dwelling in it, with a body length longer than 10 mm (Swift et al., 1979). The community structure of the soil macrofauna can act as an indicator of environment-soil changes (Lavelle, 1997). Maybe the approach that contributes the most for an understanding of the regulatory ability of the soil fauna in different ecosystems and the expected consequences when one or more groups are excluded consists in analyzing the composition and importance of certain functional groups (Costa, 2002).

The macrofauna has a potential to become a quality indicator, since it is visible to the naked eye, in addi-

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tion to being responsive to management. However, knowledge has to be produced about the groups of organisms that occur under the various types of organic management, their type of response, and the role played by them.

The objective of this work was to evaluate the soil macrofauna community under different soil covers in fig trees growing under organic management, identifying their composition, diversity, and abundance, as a first step toward obtaining bioindicators for these systems in the future.

MATERIAL AND METHODS

A fig orchard (*Ficus carica* L.), variety Roxo de Valinhos, was implemented in a Typic Palehumult in March 2000 in an area that belongs to SIPA (Sistema Integrado de Produção Agroeocológica – Agroecological Production Integrated System, Almeida et al., 1999), in the municipality of Seropédica, RJ (43°41'10" W and 22°45'30" S, at 33 m above sea level), using organic production methods.

The organic production methods refer to those preconized in Instrução Normativa n° 07 (Normative Instruction no. 7), dated 17 May 1999, by Ministério da Agricultura (Brazilian Agriculture Ministry). Thus, no agrochemicals were used to control plant diseases, although different biological agents and mixtures such as Bordeaux and lime-sulfur could be used. With regard to the use of corrective amendments and fertilizers of industrial origin, only those allowed in the regulations contained in the normative instruction mentioned above were utilized; no synthetic fertilizers were used.

The experiment area had been previously cultivated with sugarcane for four years. The area was tilled by means of one plowing and two harrowings, with incorporation of plant residues. Lime was applied in the entire area, in addition to Thermophosphate as a source of phosphorus and micronutrients. Organic fertilization consisted of 30 litters per pit (10 t ha⁻¹ dry matter) rotted cattle manure; 1/3 were placed in the bottom of the pit and 2/3 were homogeneized with dirt and returned to the pit, with the horizons inverted. The initial fertilization for orchard formation was performed by broadcasting the fertilizers on a circle with a 50 cm radius; applications consisted of 4.5 kg bird manure per plant, 250 g Thermophosphate per plant (68 kg ha⁻¹ P), and 1.2 kg timber ash per plant (83 kg ha⁻¹ K). Bordeaux mixture was applied on the surface and inside the fig tree crowns in order to control plant rust.

Siratro was planted five days after the fig trees were planted, at a density of 30 seeds m⁻¹, distributed in furrows spaced at 0.5 m; bahiagrass was planted as seedlings at a row spacing of 0.3×0.3 m. The mulch was obtained by mowing the turf in areas of the Federal Ru-

ral University of Rio de Janeiro campus and of Embrapa Agrobiologia. In 2001, the addition of residues represented by mulch was 25 t ha⁻¹ in February and 32 t ha⁻¹ in August, in order to constantly maintain a layer of at least 0.2 m in height. Bahiagrass was maintained under a cutting management performed by means of hoeings, at a height of approximately 0.05 m from the surface, at intervals of ca. 60 days. Siratro management consisted of one cutting in the beginning of 2000 and another in October 2001. The climate data in the study region for the year 2001 are presented in Figure 1. These data were provided by the "Ecologia Agrícola" (Agricultural Ecology) Weather Station of Pesagro-Rio/INMET, located at km 47 in the former Rio-São Paulo Road (22°45' S and 43°41' O, at an altitude of 33 meters), in Seropédica/RJ, Brazil.

The experimental design was organized as random blocks, in a $3 \times 2 \times 2$ factorial scheme, with three replicates. The three treatments consisted of bahiagrass (*Paspalum notatum*) living mulch (perennial grass with a rhizomatous habit); siratro (*Macroptilium atropurpureum*) living mulch (herbaceous perennial legume with a voluble habit); and mulch consisting of bahiagrass straw. The macrofauna and plant covers were evaluated in two seasons: March and September 2001, representing the rainy and dry seasons, respectively.

The TSBF (Tropical Soil Biology and Fertility) method, described by Anderson & Ingram (1993), was used to sample the soil macrofauna. Three samplings were performed at each plot of the experiment area, totaling nine sampling points. A 0.25×0.25 m area was delimited at each point, and layers of plant material and soil were sampled down to a depth of 0.3 m.

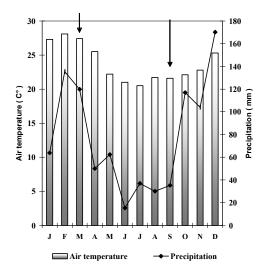


Figure 1 - Air temperature (°C) and precipitation (mm) for the year 2001. Arrows indicate the months when soil macrofauna samplings were taken.

The macrofauna individuals longer than 10 mm were removed manually and stored in containers with 70% alcohol. These were later counted and identified under a stereoscopic microscope, at the level of major taxonomic group. The term *group* was used in the soil macrofauna study, meaning either a family, or a class or order, with the objective of comprising a set of individuals with a similar life form.

The communities were characterized based on the following parameters: a) density, number of individuals per square meter; b) Shannon Diversity Index (H), calculated by the formula: H = -" pi.log pi; where pi = ni/N; ni = density of each group, N = " of densities of all groups (Magurran, 1988).

The plant layer sampled in both seasons was chemically analyzed according to the method recommended by Bataglia et al. (1983) for N, P, and K determination, and was analyzed for Ca and Mg according to Claessen et al. (1997). The results are presented in Table 1.

The chemical characterization of the soil was performed during the March 2001 collection, according to Claessen et al. (1997). Soil moisture was determined in both collection seasons (March - M_i and September - M_i). The results are presented in Table 2.

The results were submitted to analysis of variance (ANOVA) using the SAS® (Statistical Analysis System) software. Tukey test was employed to make comparisons between means, at a 5%. Significant interactions were partitioned according to the factors involved, for treatments, seasons, and depths, whenever significance was found.

Table 1 - Nutrient contents of living mulch and senescent materials of siratro (S) and bahiagrass (B), and mulch (bahiagrass straw) (M). Means of two sampling seasons (March and September, 2001).

Plant cover	Ca	Mg	P	K			
	g kg ⁻¹						
S	17.2 a	4.0 a	1.4 ab	13.3 a			
В	4.6 b	2.2 a	0.9 b	4.7 b			
M	8.1 ab	4.6 a	1.5 a	10.7 ab			

Values followed by the same letter do not differ at 5% (Tukey test).

RESULTS AND DISCUSSION

Soil macrofauna density

The soil macrofauna groups Chilopoda, Oligochaeta, and Formicidae were found at higher densities in the soil (Figure 2). Araneae, however, showed a higher density in the plant layer.

Coleoptera, Diplopoda, and Oligochaeta were the most sensitive groups in relation to the season when evaluations were performed, and were more abundant in March (Table 3). Ant density in bahiagrass was similar to ant density in siratro (Table 3). Relatively to all other groups found, ants represented more than 80% of the macrofauna in bahiagrass, 58% in siratro, and 8% in mulch. It is possible that the low nutritional quality of the bahiagrass living mulch (Table 1) did not favor the abundance of other groups. On the other hand, ants, have a wide number of species that become adapted depending on the conditions of the environment (Fowler, 1998), which make them capable of maintaining their reproductive capacity, even when the environmental conditions are unfavorable (Bruyn, 1999). This wide occurrence, in association with the role they play in the soil make ants important, especially in agroecosystems under organic man-

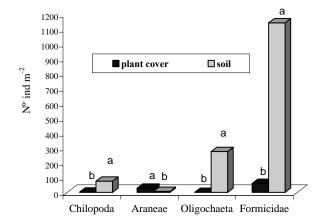


Figure 2 - Soil macrofauna density in plant layers and in soil cultivated with fig trees. (Values refer to means of sampling seasons and different plant cover types). For statistical analysis, values were transformed to log x+1. The same letters in each group do not differ at 5% (Tukey test).

Table 2 - Chemical characteristics of a Red-Yellow Argisol at a 0-0.3 m depth, under three soil covers (S = siratro, B = bahiagrass, and M = mulch).

Treatment	pH_{H2O}	$\mathbf{M}_{\scriptscriptstyle (i)}$	$\mathbf{M}_{\scriptscriptstyle{(\mathrm{f})}}$	Al	Ca+Mg	Ca	Mg	P	K
		%			mmolc dm ⁻³			mg dm ⁻³	
S	5.9 a	16 a	15 a	0.0 a	4.6 b	3.2 a	1.4 a	57 a	128 b
В	6.1 a	14 b	10 b	0.0 a	4.5 b	3.9 a	1.1 a	68 a	104 b
M	5.9 a	17 a	14 a	0.0 a	5.3 a	3.4 a	1.9 a	39 a	368 a

These values refer to March 2001, except for M_i and M_p , which refer to soil moisture measured in March and September 2001, respectively. Values followed by the same letter in the same column do not differ at 5% (Tukey test).

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Table 3 - Density of individuals in the soil macrofauna in a fig orchard area under different soil covers. (Mean values in the surface layers and in the soil).

Invertebrate groups	Evaluation seasons	Siratro	Bahiagrass	Mulch	Mean	
		No. ind m ²				
Coleoptera	March ^{1,2}	45	16	72	44 a	
	September	14	3	0	05 b	
	mean	29 AB	9 B	36 A		
Diplopoda	March	251	32	317	200 a	
	September	112	37	29	59 b	
	mean	181 A	35 B	173 A		
Gastropoda	March	25	6	0	10 a	
	September	75	6	27	36 a	
	mean	50 A	6 B	13 B		
Oligochaeta	March	222	131	237	196 a	
	September	139	75	24	79 b	
	mean	180 A	103 B	131 AB		
Formicidae	March	227	1323	16	522 a	
	September	1024	968	45	679 a	
	mean	625 A	1145 A	31 B		

¹For statistical analysis, values were transformed to log x+1.

agement, where the work they perform in biological processes is more expressive than the damage threat they pose to crops.

The ant genera found were: *Crematogaster, Brachymyrmex* (which occurred especially in bahiagrass), *Solenopsis, Pheidole, Hypoponera* (especially in siratro), and *Mycocepurus*. This provides an indication of the relevance of this group to the soil fauna community, since it presents great diversity and becomes established in several habitats. When abundant, ants can modify the soil's physical structure by creating systems of galleries and chambers, influencing soil porosity, aeration, infiltration, and drainage (Lavelle & Spain, 2001). They also change soil texture and other soil properties through the horizontal and vertical transfer of materials of different particle sizes (Alvarado et al., 1981).

When both living mulches are compared, a higher density of Oligochaeta, Diplopoda, and Gastropoda can be found in the siratro living mulch than in bahiagrass (Table 3). Since Oligochaeta and Diplopoda are saprophytic, the higher density observed is probably associated with the greater food supply, organic matter quality, and higher soil moisture (Tables 1 and 2). Gastropods depend on calcium to build their shells, which represent a protection for their bodies (Burch & Pearce, 1990). This was probably a decisive factor why the siratro vegetation influenced the distribution of this group (Table 2).

In the mulch treatment, incorporation of residues amounted to 25 t ha⁻¹ in February and 32 t ha⁻¹ in August. With regard to the living mulches, the production

of siratro and bahiagrass residues was not quantified, but results obtained under similar edaphic-climatic conditions showed that bahiagrass yields about 4,000 kg ha⁻¹ in summer and 1,600 kg ha⁻¹ in the beginning of spring (Espíndola et al., 2001), and that siratro yields 3,100 kg ha⁻¹ when harvested in the dry season and 3,700 kg ha⁻¹ when harvested in the rainy season (Espíndola, 2001). However, evaluations performed in the experiment area of the fig orchard have shown that siratro presents substantial spontaneous leaf shedding, which represented around 1.45 t ha⁻¹ dry matter per month throughout 2002 (personal comm.).

Mulch cover increases earthworm populations in the soil, as compared with bare soil, and this effect is positively correlated with the quality of the mulch (Tian et al. 1999) although these factors are determinative for the occurrence of earthworms (Edwards & Lofty, 1982; Lee, 1985), exclusive colonization by *Pontoscolex corethrurus* (Muller) was observed, and no native species were found, such as those reported by Aquino (1999) to occur in a nearby secondary vegetation area.

Araneae and Chilopoda predators were found in all soil covers at low densities. These groups are related to more diversified habitats (Altieri et al., 1996; Odum, 1988).

Soil macrofauna diversity

Diversity determination is highly important because it can function as a bioindicator of community stability and describe the ecological dynamics of the commu-

²The same upper case letters in the same row or lower case letters in the same column do not differ at 5% (Tukey test).

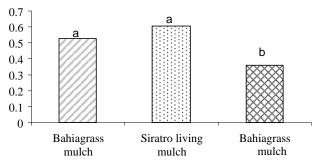


Figure 3 - Soil macrofauna diversity (Shannon index) in a fig orchard area under different soil covers (values refer to the mean between two sampling seasons). Same letters do not differ 5% (Tukey test).

nity and the impact of stress (Kennedy & Smith, 1995). No difference of Shannon index was detected between evaluation seasons (March and September). Therefore, the data presented refer to the mean of these collections in each treatment. Shannon index, which refers to both richness and density, was similar for mulch and siratro (Figure 3).

In spite of the high macrofauna density found in the bahiagrass treatment (1,306 individuals m²), the Shannon index was lower in this treatment in relation to the mulch and siratro treatments (Figure 3). The diversity values found in this experiment can be considered high, when compared to those found in other areas cultivated under the conventional system and under no-till, and are very close to the values found in forest soils (Aquino et al., 2000).

CONCLUSIONS

Using the soil macrofauna community as a whole worked better as a bioindicator than if a certain group from the fauna would have been selected for that purpose.

The macrofauna density and diversity indicated that the living mulch with bahiagrass encourages a more stressing and impacting environment for the community than soil covers consisting of siratro living mulch or bahiagrass straw mulch.

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