

Ant community richness and composition across a gradient from *Eucalyptus* plantations to secondary Atlantic Forest

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Abstract: Secondary forests and exotic tree plantations are expanding across tropical landscapes. However, our current understanding of the value of these human-dominated forest landscapes for invertebrate biodiversity conservation is still very poor. In this paper, we use the leaf-litter ant fauna to assess invertebrate diversity in one commercially managed *Eucalyptus* plantation (four years old), two abandoned plantations of different regeneration ages (16 and 31 years), and one neighboring secondary Atlantic Forest in Southeastern Brazil. There was a clear gradient in species richness from the secondary forest to the managed *Eucalyptus* plantation; richness and diversity peaked in secondary forest and in the older regenerating *Eucalyptus* plantation. Significantly more species were recorded in secondary forest samples than in *Eucalyptus* plantations, but *Eucalyptus* plantations had a similar level of richness. Furthermore, a non-metric multidimensional scaling analysis revealed clear differences in species composition between the younger managed *Eucalyptus* plantation (understory absent) and habitats with sub-developed or developed understory. *Eucalyptus* plantations were characterized by an assemblage of widespread, generalist species very different from those known to occur in core forest habitats of southeastern Brazil. Our results indicate that while older regenerating *Eucalyptus* plantations can provide habitat to facilitate the persistence of generalist ant species, it is unlikely to conserve most of the primary forest species, such as specialized predators, Dacetini predators, and nomadic species.

Keywords: Formicidae, habitat modification, eucalipt plantations, diversity.

SUGUITURU, S.S., SILVA, R.R., SOUZA, D.R., MUNHAE, C.B. & MORINI, M.S.C. **Riqueza e composição das comunidades de formigas em um gradiente de plantio de *Eucalyptus* a Mata Atlântica secundária.** *Biota Neotrop.* 11(1): <http://www.biotaneotropica.org.br/v11n1/pt/abstract?inventory+bn00811012011>.

Resumo: As florestas secundárias e plantações de espécies exóticas estão se expandindo nas paisagens tropicais. No entanto, nossa compreensão sobre o valor destas florestas para a conservação da biodiversidade de invertebrados ainda é incipiente. Neste trabalho, usamos a fauna de formigas de serapilheira para avaliar a diversidade desses insetos entre três florestas de *Eucalyptus*, sendo uma comercial (quatro anos de idade) e duas abandonadas em diferentes idades de regeneração (16 e 31 anos) e uma área de Mata Atlântica secundária. A riqueza total foi mais alta na floresta secundária e nos plantios de *Eucalyptus* abandonados há mais tempo. A densidade de espécies na floresta secundária foi significativamente maior quando comparado as plantações de *Eucalyptus*, mas não difere entre eucaliptais; análise de ordenação revelou diferenças na composição de espécies entre as plantações de *Eucalyptus* com subbosque ausente e com subbosque desenvolvido ou em desenvolvimento. Ainda, foi constatada uma sobreposição acentuada entre amostras de serapilheira das florestas de eucaliptos abandonadas há mais tempo e a floresta secundária. Em geral, plantações de eucalipto foram caracterizadas pela presença de espécies generalistas e de ampla distribuição. Nossos resultados indicam que embora o subbosque de plantações de eucaliptos com maior idade de regeneração suporte um conjunto relativamente alto de espécies generalistas de formigas, é improvável que eucaliptais conservem a maioria das espécies de florestas primárias, especialmente predadores especializados, Dacetini e espécies nômades.

Palavras-chave: Formicidae, modificação de habitats, plantações de eucalipto, diversidade.

Introduction

The Atlantic Forest once covering more than a million squared kilometers, was reduced to 12% of the original cover (Ribeiro et al. 2009) and the total protected area is only 1.62% (Laurance 2009). The Atlantic Forest is recognized as one of the most highly threatened biodiversity hotspot of the world (SOS Mata Atlântica 1998, Myers et al. 2000), with high number of endemic species (Martini et al. 2007, Carnaval et al. 2009, Metzger et al. 2009).

In the Southeastern Brazil extensive Atlantic Forest areas had been replaced by managed tree plantations. The exotic *Eucalyptus* trees were introduced in Southern Brazil in 1868 to increase the production of railroad ties (Lima 1993). Presently, they are used mainly in the pulpwood industry and for energy production in ironworks. Plantations have increased rapidly within Brazil, where the coverage of *Eucalyptus* planted for the pulpwood industry has increased 17-fold since 1980 (Sociedade... 2006). It is estimated that Brazil has the largest land area of *Eucalyptus* plantations in the world; in Southeastern Brazil alone they occupy about 611,500 ha (Kronka 2005). The value for wildlife of extensive monocultures, as well as of areas of native regeneration, is surprisingly poorly understood (Kanowski et al. 2005). Our current knowledge is insufficient to predict whether these habitats could help to conserve tropical forest species for the future (Gardner et al. 2007, 2008). However, they are likely to become a dominant feature in the tropical forest landscapes (Wright & Muller-Landau 2006). Most studies have been restricted to a few well-studied taxa (birds and mammals) in tropical regions and landscapes dominated by sites with similar regeneration ages since abandonment (Machado & Lamas 1996, Barlow et al. 2007b). However, studies along gradients of landscape modification could be important for understanding response patterns of the invertebrate fauna to forest modification (Summerville & Crist 2002, Pinkus-Rendón et al. 2006), and the use of ecological indicators to assessment of forest disturbance has been suggested (McGeoch & Chown 1998, Uehara-Prado et al. 2009).

Ants are routinely used as indicators of ecological change and ecosystem dynamics (Underwood & Fisher 2006) because they are ubiquitous, highly abundant, diverse, and respond to ecosystem disturbance. Ants are important predators, competitors, and mutualists in most terrestrial habitats (Hölldobler & Wilson 1990), with effects on soil structure (McKey et al. 2010) and process which affect the flow of energy and material in ecosystems (Folgarait 1998).

In most habitats, plant communities determine the physical structure of the environment and have considerable influence on the distribution of animal species at a range of spatial scales (Tews et al. 2004, Lassau et al. 2005). Habitat structure is an important determinant of tropical ant community composition (Lassau & Hochuli 2004), and there is a strong association between ant species richness and composition and a more complex habitat structure in regenerating forests (Dunn 2004). The wildlife value of plantations can be highly dependent on levels of understory regeneration (Stallings 1991, Mitra & Sheldon 1993). Older unmanaged *Eucalyptus* plantations with dense understories in Southeastern Brazil can support relatively high numbers of bird species (Willis 2003). Studies that found a high diversity in tree plantations show that the latter are associated with mature stands that are floristically similar to natural forests (Duran & Kattan 2005).

In the present study, we compared the leaf-litter ant diversity in human-modified Atlantic Forest areas in Southeastern Brazil. We contrasted the leaf-litter ant diversity among naturally regenerated Atlantic Forest and abandoned *Eucalyptus* plantations. The abandoned plantation forests represent a gradient of Atlantic Forest regeneration and are composed of different tree species and different types of

understory vegetation. Our hypothesis is that the ant community richness and composition differs between forests across a gradient from *Eucalyptus* plantations to secondary Atlantic Forest. We expected that 1) the species richness of leaf-litter ant fauna is linked to structure of the understory vegetation at each site, and 2) the similarity in species composition between secondary forest and eucalypt plantations increases with increase regeneration age.

Material and Methods

1. Study area

The study sites are located in the Parque das Neblinas (23° 44' 51" S and 46° 08' 39" W), Bertioga county, State of São Paulo, Southeastern Brazil. Four adjacent study sites of similar size (40 ha) were chosen, close to primary Atlantic Forest:

- A secondary Atlantic Forest area (SF): The understory is developed, characterized by herbaceous plants, shrubs, and tree canopies > 1.5 m.
- Four year-old managed *Eucalyptus grandis* (Hill ex Maiden) plantation (Eg4): The plantation is characterized by the absence of an understory. The native understory vegetation is periodically cleared or suppressed, either by manual removal or herbicidal treatment.
- Sixteen year-old *Eucalyptus saligna* Smith plantation (Es16): This area was commercially managed and was abandoned in 1988. The understory is under-developed, characterized by herbaceous plants, shrubs, and trees > 1 m;
- Thirty-one year-old *Eucalyptus grandis* (Hill ex Maiden) plantation (Eg31): This plantation was commercially managed and was abandoned in 1973. The understory is developed, characterized by herbaceous plants, shrubs and tree canopies > 1 m.

2. Sampling design

In each area, a 1,200 m long line transect was selected for the ant survey. Transects started 500 m from the forest edge and contained 25 points, each 50 m apart. At each point, two samples were collected, one 25 m to the left and one 25 m to the right of the point. At each of the 50 sample points, a 1 m² plot (quadrat) was established on the ground. All leaf-litter inside the plot was collected, sifted and put into a bag. The sifted material was brought to the field-laboratory and its fauna was extracted using mini-Winklers for 48 hours (Agosti & Alonso 2000, Bestelmeyer et al. 2000). Surveys were carried out during the rainy season and after rainy periods to enhance the likelihood of collecting cryptic and hypogaic ant species.

The material was initially identified to genera and then to morphospecies. Identification to species level was carried out by comparison with the Formicidae collection of the Museu de Zoologia da Universidade de São Paulo (MZUSP), and with pertinent literature. Classification follows (Bolton et al. 2006). Vouchers were deposited at the Universidade de Mogi das Cruzes (SP) and MZUSP.

We used leaf-litter depth as a surrogate for resource availability within the quadrats. In each sampling unit, the litter depth was measured at the center of quadrat by vertically inserting a ruler marked in 0.5 - cm increments into the topsoil.

3. Data analysis

In the analyses, we treated each sample as a statistical independent replicate. All samples come from points at least 50 m apart; as the home range of most leaf-litter ant species covers less than a few meters, the samples were considered independent. Species richness was defined as the number of ant species occurring in each litter

sample. Matrices of species by sites quadrats were constructed, and filled with occurrence data (presence = 1, absence = 0). One occurrence was defined as the presence of one species in one 1 m², no matter how many individuals were present.

We considered leaf-litter depth within the quadrats as a surrogate for resource availability for leaf-litter ant fauna. Leaf-litter depth is an appropriate measure because leaf-litter provides suitable places for nesting and is the resource base for litter-dwelling ants. We tested the hypothesis that understory age increases ant species richness by using the Kruskal–Wallis test, followed by multiple comparison test adjusted appropriately using “kruskalmc” function in pgirmess package (Giraudeau 2010). We evaluated the relationship between leaf-litter and species richness at each site using General Linear Models (GLM), with leaf-litter depth as an explanatory variable and species richness within quadrats as a response variable. Poisson errors were corrected for overdispersion (Crawley 2002).

Patterns of species richness among different forest types were compared using sample-based rarefaction curves constructed using EstimateS version 7.5.2 (Colwell 2005). Comparisons among areas were standardized by the number of occurrence, because we were interested in patterns of species richness and not in species density (Gotelli & Colwell 2001). Estimated species richness was calculated for each habitat using EstimateS version 7.5.2, using the mean of the three commonly employed incidence-based estimators for ant biodiversity studies (Chao2, Jack1 and Bootstrap2).

Ordination analysis (non-metric multidimensional scaling) based on a Bray–Curtis dissimilarity matrix, using presence-absence data to compare patterns of species composition and community structure, were implemented in the vegan package (Oksanen et al. 2009).

Results

We recorded a total of 37,486 leaf-litter ants in 200 samples from all habitats. These were assigned to 135 species and morphospecies representing 48 genera and 11 subfamilies. A total of 87 species were found in the secondary Atlantic Forest, 79 within one *E. grandis* plantation (31 years, developed understory), 76 in the *E. saligna* plantation (16 years, sub-developed understory) and 61 in the other *E. grandis* plantation (4 years, without understory) (Table 1). We identified 32 (23.5%), 8 (5.9%), 14 (17%) and 9 (6.62%) species, respectively, that were unique to each habitat. Thirty-two species (24%) were common to all habitats.

The nMDS ordination show distinct sample clusters, in special, the immature eucalypt plantation (Eg4). However, there were a large amount of overlap in sample composition among secondary forest and older eucalypt plantations (Es16 and Eg31; Figure 1) meaning that a number of secondary forest species can be found in areas of eucalypt plantations with an understory of native shrubs. Species generally considered rare and cryptic occurred in the studied areas irrespective to habitat; these were: *Adelomyrmex* sp. (secondary forest), *Amblyopone armigera* Mayr (Eg4 and Eg31), *A. elongata* (Santschi) (secondary forest), *Brachymyrmex micromegas* Emery (secondary forest), *Cylindromyrmex brasiliensis* Emery (Es16), *Basiceros* sp.2 (Eg4 and Eg31), *Phalacromyrmex fugax* Kempf (Eg31), and *Typhlomyrmex* sp.2 (secondary forest and Eg31) (Table 1).

The accumulated richness curves did not approximate the level of the asymptote (Figure 2), indicating that the sample size was not large enough to sample the expected species richness. The mean of the three estimators (Chao2, Jack 1 and Bootstrap) yielded an extrapolated maximum of 108.04 species for the secondary forest, 93.61 for the *E. grandis* plantation (31 years), 117.41 for the *E. saligna* plantation (16 years), and 71.26 species for the other *E. grandis* plantation (4 years).

There was no relationship between leaf-litter depth and ant species richness in any of the study sites (SF: t-value = -0.355, p = 0.724; Eg31: t-value = -0.381, p = 0.705; Es16: t-value = 0.254, p = 0.800; Eg4: t-value = 1.039; p = 0.304). We did not find significant differences in leaf-litter depth among sites (Kruskal–Wallis = 4.1024, df = 3, p-value = 0.2506). However, there was a significant effect of understory structure on leaf-litter ant species richness at the quadrat level (Kruskal–Wallis = 42.8354, df = 3, p-value = 0.000). Significantly more species were recorded in secondary forest than in *Eucalyptus* plantations (p < 0.01), but *Eucalyptus* plantations had a similar level of richness (p > 0.01).

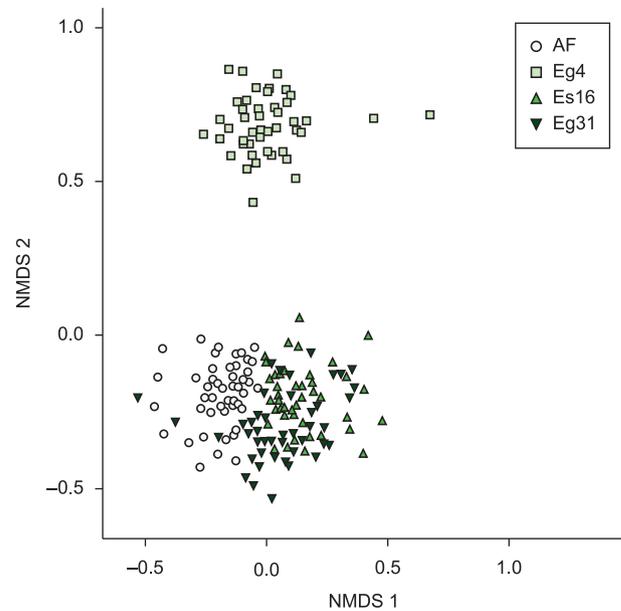


Figure 1. Non-metric multidimensional scaling (NMDS) ordination for leaf-litter ant communities sampled in four forests. Ordination analyses are based on presence-absence dissimilarity matrices (Bray–Curtis). SF: secondary forest; Eg4: 4 year-old *Eucalyptus grandis*; Es16: 16 year-old *Eucalyptus saligna*; and Eg31: 31 year-old *Eucalyptus grandis*.

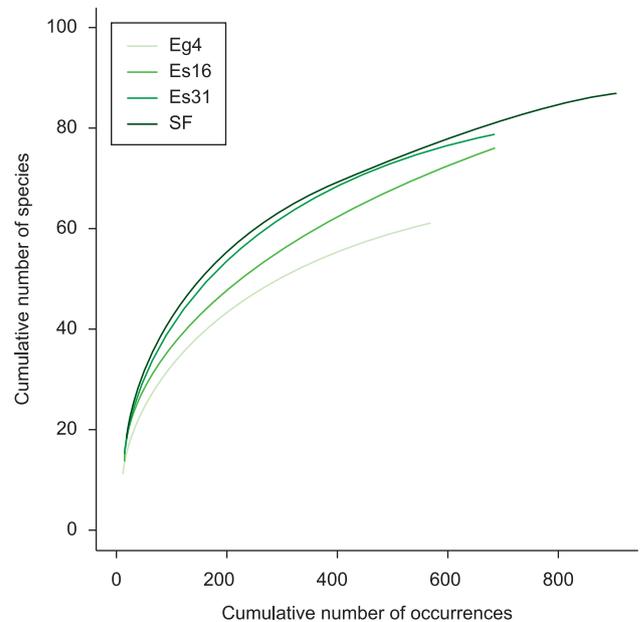


Figure 2. Species rarefaction curves of the number of species plotted against the number of occurrences of leaf-litter ant species. SF: secondary forest; Eg4: 4 year-old *Eucalyptus grandis*; Es16: 16 year-old *Eucalyptus saligna*; and Eg31: 31 year-old *Eucalyptus grandis*.

Table 1. Number of occurrences of each ant species collected in 50 m² samples of leaf-litter in the studied areas. Eg4: *Eucalyptus grandis* 4 years old; Es16: *Eucalyptus saligna* (16 years old); Eg31: *Eucalyptus grandis* (31 years old); and SF: secondary forest.

Subfamily/species	Eg4	Es16	Eg31	SF
Amblyoponinae				
<i>Amblyopone armigera</i> Mayr, 1887	2	-	6	-
<i>Amblyopone elongata</i> Santschi, 1912	-	-	-	5
<i>Prionopelta antillana</i> Forel, 1909	3	-	3	5
Cerapachinae				
<i>Cylindromyrmex brasiliensis</i> Emery, 1901	-	1	-	-
Dolichoderinae				
<i>Azteca</i> sp.2	-	1	-	-
<i>Linepithema neotropicum</i> Wild, 2007	17	35	10	18
<i>Linepithema iniquum</i> (Mayr, 1870)	-	-	1	2
<i>Tapinoma melanocephalum</i> (Fabricius, 1793)	-	1	-	2
Ecitoninae				
<i>Eciton burchelli</i> (Westwood, 1842)	-	-	-	7
<i>Eciton quadriglume</i> (Haliday, 1836)	-	1	1	-
<i>Labidus praedator</i> (Smith F., 1858)	1	-	-	-
Ectatomminae				
<i>Ectatomma edentatum</i> Roger, 1863	11	2	6	-
<i>Gnamptogenys continua</i> (Mayr, 1887)	-	1	1	6
<i>G. reichenspergeri</i> (Santschi, 1929)	-	-	2	1
<i>Gnamptogenys striatula</i> Mayr, 1884	5	19	18	11
<i>Gnamptogenys</i> sp.5	-	-	-	1
Formicinae				
<i>Camponotus (Tanaemyrmex)</i> sp.	-	1	-	-
<i>Camponotus</i> sp.6	-	-	1	-
<i>Camponotus</i> sp.8	-	-	-	1
<i>Camponotus</i> sp.10	-	1	-	-
<i>Brachymyrmex heeri</i> Forel, 1874	1	8	14	47
<i>Brachymyrmex incisus</i> Forel, 1912	-	25	12	2
<i>Brachymyrmex luederwaldti</i> Santschi, 1929	-	-	3	5
<i>Brachymyrmex micromegas</i> Emery, 1923	-	-	-	1
<i>Brachymyrmex pictus</i> Mayr, 1887	-	12	3	23
<i>Myrmelachista catharinae</i> Mayr, 1887	-	1	-	1
<i>Myrmelachista ruszkii</i> Forel, 1903	-	1	-	1
<i>Paratrechina fulva</i> (Mayr, 1862)	36	41	25	31
Heteroponerinae				
<i>Heteroponera dentinodis</i> (Mayr, 1887)	2	1	1	1
<i>Heteroponera dolo</i> (Roger, 1961)	-	-	1	-
<i>Heteroponera mayri</i> Kempf, 1962	-	11	21	25
Myrmicinae				
<i>Acromyrmex niger</i> (F. Smith, 1858)	10	3	4	8
<i>Acromyrmex rugosus rochai</i> Forel, 1904	2	-	-	-
<i>Acromyrmex</i> sp.7	1	-	-	-
<i>Apterostigma</i> sp.1	4	3	3	4
<i>Cyphomyrmex auritus</i> Mayr, 1887	-	1	-	-
<i>Cyphomyrmex strigatus</i> Mayr, 1887	-	-	-	1
<i>Mycetosoritis</i> sp.1	18	34	24	23
<i>Myrmicocrypta</i> sp.1	-	-	-	5
<i>Trachymyrmex (gr.septentrionales)</i> sp.4	-	1	5	-

Table 1. Continued...

Subfamily/species	Eg4	Es16	Eg31	SF
<i>Trachymyrmex oetkeri</i> Forel, 1908	2	-	1	-
<i>Basiceros disciger</i> (Brown & Kempf, 1960)	-	-	-	5
<i>Basiceros</i> sp.2	2	-	1	-
<i>Basiceros rugiferum</i> Mayr, 1887	24	3	4	29
<i>Basiceros stenognathum</i> (Brown & Kempf, 1960)	3	6	19	33
<i>Wasmannia</i> sp.3	42	38	6	19
<i>Procryptocerus</i> sp. pr. <i>schmalzi</i>	-	1	-	-
<i>Crematogaster</i> (<i>Orthocrema</i>) sp.1	-	3	-	6
<i>Crematogaster</i> sp.5	-	-	1	-
<i>Crematogaster</i> sp.8	1	-	-	-
<i>Acanthognathus ocellatus</i> Mayr, 1887	-	-	-	1
<i>Acanthognathus rudis</i> Brown & Kempf, 1969	-	1	3	-
<i>Strumigenys appretiata</i> (Borgmeier, 1954)	-	1	-	1
<i>Strumigenys crassicornis</i> Mayr, 1887	8	1	6	32
<i>Strumigenys denticulata</i> Mayr, 1887	48	48	41	23
<i>Strumigenys elongata</i> Roger, 1863	-	5	-	-
<i>Strumigenys schmalzi</i> Emery, 1906	2	-	2	4
<i>Strumigenys</i> sp.6	2	3	4	1
<i>Strumigenys</i> sp.12	12	-	3	6
<i>Strumigenys</i> sp.14	-	-	-	2
<i>Strumigenys</i> sp.15	1	7	10	-
<i>Strumigenys louisianae</i> Roger, 1863	-	3	4	7
<i>Adelomyrmex</i> sp.1	-	-	-	2
<i>Hylomyrma balzani</i> (Emery, 1894)	-	-	-	1
<i>Hylomyrma reitteri</i> (Mayr, 1887)	3	33	37	29
<i>Phalacromyrmex fugax</i> Kempf, 1960	-	-	1	-
<i>Pheidole</i> sp.4	1	-	-	-
<i>Pheidole</i> sp.5	-	-	-	8
<i>Pheidole</i> sp.6	-	-	-	3
<i>Pheidole</i> sp.7	45	48	45	44
<i>Pheidole</i> sp.8	-	-	-	1
<i>Pheidole</i> sp.9	1	-	2	-
<i>Pheidole</i> sp.12	-	7	-	-
<i>Pheidole</i> sp.13	1	8	3	32
<i>Pheidole</i> sp.14	-	-	3	2
<i>Pheidole</i> sp.15	1	-	-	29
<i>Pheidole</i> sp.16	2	2	19	-
<i>Pheidole</i> sp.17	-	-	-	1
<i>Pheidole</i> sp.18	-	2	-	-
<i>Pheidole</i> sp.19	1	-	1	-
<i>Pheidole</i> sp.20	7	7	6	1
<i>Pheidole</i> sp.21	-	11	25	2
<i>Pheidole</i> sp.22	3	3	-	-
<i>Pheidole</i> sp.23	2	3	12	-
<i>Pheidole</i> sp.24	4	-	-	-
<i>Pheidole</i> sp.26	-	-	-	1
<i>Pheidole</i> sp.27	2	1	-	-
<i>Pheidole</i> sp.28	21	12	14	19
<i>Pheidole</i> sp.30	3	10	5	4

Table 1. Continued...

Subfamily/species	Eg4	Es16	Eg31	SF
<i>Pheidole</i> sp.31	12	9	-	3
<i>Pheidole</i> sp.32	-	-	-	2
<i>Pheidole</i> sp.33	-	-	2	4
<i>Pheidole</i> sp.36	-	-	2	1
<i>Pheidole</i> sp.39	-	2	14	30
<i>Pheidole</i> sp.43	-	-	2	-
<i>Carebara</i> sp.1	-	-	-	1
<i>Megalomyrmex goeldii</i> Forel, 1912	-	1	-	-
<i>Megalomyrmex iheringi</i> Forel, 1911	-	4	4	6
<i>Monomorium pharaonis</i> (Linnaeus, 1758)	-	-	1	-
<i>Oxyepoecus venezyi</i> (Forel, 1907)	-	-	-	1
<i>Oxyepoecus</i> sp.2	-	6	6	4
<i>Solenopsis saevissima</i> (F. Smith, 1855)	1	-	-	-
<i>Solenopsis wasmannii</i> Emery, 1894	20	13	12	33
<i>Solenopsis</i> sp.2	42	44	35	48
<i>Solenopsis</i> sp.4	39	20	25	41
<i>Lachnomyrmex plaumanni</i> Borgmeier, 1957	4	17	9	14
<i>Lachnomyrmex</i> sp.1	-	1	1	-
Ponerinae				
<i>Anochetus altisquamis</i> Mayr, 1887	-	1	2	6
<i>Hypoponera</i> sp.1	24	34	30	42
<i>Hypoponera</i> sp.3	1	1	-	6
<i>Hypoponera</i> sp.4	20	4	16	4
<i>Hypoponera</i> sp.5	1	-	1	10
<i>Hypoponera</i> sp.6	-	3	10	12
<i>Hypoponera</i> sp.7	5	1	2	2
<i>Hypoponera</i> sp.8	33	41	48	35
<i>Hypoponera</i> sp.9	9	1	7	13
<i>Hypoponera</i> sp.10	-	-	-	1
<i>Hypoponera</i> sp.11	-	1	4	4
<i>Hypoponera</i> sp.12	-	-	1	1
<i>Leptogenys</i> sp.2	-	-	-	2
<i>Odontomachus affinis</i> Guérin, 1844	1	2	1	1
<i>Odontomachus meinerti</i> Forel, 1905	-	-	-	2
<i>Pachycondyla bucki</i> (Borgmeier, 1927)	-	-	1	-
<i>Pachycondyla constricta</i> (Mayr, 1884)	5	15	9	17
<i>Pachycondyla crenata</i> (Roger, 1961)	-	1	-	-
<i>Pachycondyla ferruginea</i> (F. Smith, 1858)	2	-	2	-
<i>Pachycondyla harpax</i> (Fabricius, 1804)	2	3	2	1
<i>Pachycondyla striata</i> Fr. Smith, 1858	-	1	-	-
<i>Pachycondyla</i> sp.9	-	1	-	-
<i>Pachycondyla</i> sp.10	2	-	-	-
<i>Typhlomyrmex</i> sp.2	-	-	1	4
Proceratiinae				
<i>Discothyrea neotropica</i> Bruch, 1919	5	-	-	-
<i>Discothyrea sexarticulata</i> Borgmeier, 1954	-	1	-	2
Pseudomyrmecinae				
<i>Pseudomyrmex gracilis</i> (Fabricius, 1804)	1	1	1	-
<i>Pseudomyrmex pallidus</i> (F. Smith, 1855)	-	-	2	-
Number of species	61	76	79	87

Discussion

Although the small spatial scale of plantations and second-growth forest in our study area did not enable us to implement a robust replicated design, our results suggest that the leaf-litter ant fauna responds to the age of regenerating *Eucalyptus* plantations. In general, the number of species present in *Eucalyptus* plantations varied according to the understory structure, being markedly different in the youngest *Eucalyptus* plantation. Despite the increase in average ant species richness at the sample level along the regeneration gradient, the secondary forest maintained the highest species richness (Soares et al. 1998, Schnell et al. 2003). The capacity of more of ant species to colonize *Eucalyptus* plantations is probably limited by the shortage of native plant species (Majer et al. 1984) and micro-habitats, such as leaf-litter accumulated around fallen logs (Andrew et al. 2000).

In our study, the site with assemblage most like those of the secondary forest reference site was the older *Eucalyptus* plantation (the sites share approximately 50% of the ant fauna). Although eucalypt plantations are intended primarily for the purpose of timber production, their ability to restore invertebrates biodiversity in modified landscapes has been evaluated in Australia (Schnell et al. 2003) and South America (Soares et al. 1998, Marinho et al. 2002, Fonseca & Diehl 2004, Barlow et al. 2007a, b). Strong evidences of a successional change in ant communities in response to tree growth within *Eucalyptus* plantations has been described (Schnell et al. 2003). Soares et al. (1998) also compared the ant fauna of a eucalypt plantation with that of a native secondary regrowth forest in Brazil and found that the secondary forest support a greater number of ant species than the eucalypt plantation. However, some studies have found different responses of the ant fauna to eucalypt plantation ages. For example, the richness of epigeic ant species was not related to age of eucalypt forests planted in sandbanks (Fonseca & Diehl 2004), and Marinho et al. (2002) demonstrated that most of the species found in native vegetation occur also in the eucalypt plantations.

We did not find a relationship between species richness and leaf-litter depth (used as a surrogate of resource availability for the leaf-litter ant fauna) in the *Eucalyptus* or secondary forest. Although the density of litter-dwelling ants increases with litter biomass (Kaspari 1996a, McGlynn et al. 2007), litter biomass can only account for a fraction of the variance in ant densities or ant species richness (Kaspari 1996b, McGlynn et al. 2009). In the litter, frequent disturbances, varying in scale from branch falls to stronger rainfalls, obscures the resource-density correlation (Kaspari 1996b). Further, studies on patch dynamics of ant colonies have showed that colonies may migrate on a weekly or monthly basis in response to predation (Kaspari 1996b).

It is well known that the physical and chemical structure of leaf-litter formed by *Eucalyptus* trees decrease decomposition by micro-organisms (Majer & Recher 1999), affecting the structure and dynamics of litter formation (Schneider 2003). Because the resource base of ants is supported by the detrital food web (Kaspari & Yanoviak 2008), we may expect a weaker correlative relationship between leaf-litter depth and ant species richness in *Eucalyptus* plantations.

Our data show the presence mainly of generalist species and of few species specialized (Brown Jr. 2000) in *Eucalyptus* plantations, such as specialized predators or hypogaic nomadic species of Ecitoninae. The simplified habitats created by management practices in commercial forests often favours generalist species and decrease the diversity of forest specialists such as litter-dwelling cryptic species (Andersen et al. 2009), which require a complex litter layer. Generalist ants are more likely to find sufficient resources within a patch, as compared to specialist species, which have limited mechanisms of dispersal and prefer locally limited resources (Pacheco et al. 2009).

It is interesting to mention the presence of the rarely collected *Phalacrotermes fugax* Kempf in the 31-years-old *Eucalyptus* site; until now, this species was known only from a few individuals in museum collections.

In sum, our results indicate that the ant species richness and composition changed along the regeneration gradient of *Eucalyptus* forests and in comparison to secondary Atlantic Forest. However, *Eucalyptus* and secondary forests are unlikely to conserve most primary forest species, such as groups of specialist predators and nomadic species, which show specialized biology. Furthermore, this study serves to highlight the potential importance of native understory vegetation for the conservation of leaf-litter ants. Further, our study suggest that examine the invertebrate ecological responses to disturbance regimes is vital if research is to inform and improve sustainable forest management practices.

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