

# The effects of land-use change on arthropod richness and abundance on Santa Maria Island (Azores): unmanaged plantations favour endemic beetles

Seline S. Meijer · Robert J. Whittaker ·  
Paulo A. V. Borges

Received: 4 March 2010 / Accepted: 9 August 2010 / Published online: 3 September 2010  
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**Abstract** We study how endemic, native and introduced arthropod species richness, abundance, diversity and community composition vary between four different habitat types (native forest, exotic forest of *Cryptomeria japonica*, semi-natural pasture and intensive pasture) and how arthropod richness and abundance change with increasing distance from the native forest in adjacent habitat types in Santa Maria Island, the Azores. Arthropods were sampled in four 150 m long transects in each habitat type. Arthropods were identified to species level and classified as Azorean endemic, single-island endemic (SIE), native, or introduced. The native forest had the highest values for species richness of Azorean endemics, SIEs and natives; and also had highest values of Azorean endemic diversity (Fisher's alpha). In contrast, the intensive pasture had the lowest values for

endemic and native species richness and diversity, but the highest values of total arthropod abundance and introduced species richness and diversity. Arthropod community composition was significantly different between the four habitat types. In the semi-natural pasture, the number of SIE species decreased with increasing distance from the native forest, and in the exotic forest the abundance of both Azorean endemics and SIEs decreased with increasing distance from the native forest. There is a gradient of decreasing arthropod richness and abundance from the native forest to the intensive pasture. Although this study demonstrates the important role of the native forest in arthropod conservation in the Azores, it also shows that unmanaged exotic forests have provided alternative habitat suitable for some native species of forest specialist arthropods, particularly saproxylic beetles.

S. S. Meijer · R. J. Whittaker  
Biodiversity Research Group, Oxford University Centre  
for the Environment, South Parks Road,  
Oxford OX1 3QY, UK

S. S. Meijer · P. A. V. Borges (✉)  
Azorean Biodiversity Group (CITA-A), Departamento  
de Ciências Agrárias, Universidade dos Açores, 9701-851  
Terra-Chã, Angra do Heroísmo, Portugal  
e-mail: pborges@uac.pt

R. J. Whittaker  
Center for Macroecology, Evolution and Climate, Department  
of Biology, University of Copenhagen Universitetsparken  
15, Copenhagen, Denmark

*Present Address:*

S. S. Meijer  
UCD Forestry, Agriculture & Food Science Centre,  
University College Dublin, Belfield, Dublin 4, Ireland

S. S. Meijer  
World Agroforestry Centre, Nairobi, Kenya

**Keywords** Arthropods · Azores · Conservation  
biogeography · Endemic species · Landscape matrix ·  
Land use

## Introduction

Humans are altering natural landscapes at an accelerating rate (Vitousek et al. 1997; Ramankutty and Foley 1999; Foley et al. 2005). Large-scale transformation and clearing of land have put ecosystems and biodiversity at serious risk (Tilman et al. 2001). The loss of natural habitats has led many species to become at risk of extinction (Pimm et al. 1995; Brook et al. 2003; Hanski et al. 2007) and some have claimed that habitat loss is causing a mass extinction of species (e.g. Myers 1992; Wilson 1992). Due to the time-lag between habitat loss and extinction (Janzen 1986; Heywood et al. 1994), it is expected that many more

species will go extinct in the future as a result of land conversion that has already taken place (Tilman et al. 1994; Brooks and Balmford 1996; Brooks et al. 1997, 1999a, b, 2002; Cowlshaw 1999; Grelle et al. 2005; Helm et al. 2006; Triantis et al. 2010).

As natural habitats get increasingly fragmented by human activities, it is important to understand the effects of this fragmentation on the distribution and survival of species in order to inform conservation strategies. The importance of protected areas in the conservation of biodiversity has been generally accepted (Margules and Pressey 2000; Bruner et al. 2001). This has been supported by classical ecological theories such as the theory of island biogeography (MacArthur and Wilson 1963, 1967) and metapopulation dynamics (Hanski 1998). However, these theories tend to view natural habitat patches as islands in a sea of uninhabitable matrix, which may be an unnecessarily pessimistic assumption in many cases (Bhagwat et al. 2008). As a result, the conservation of species has focussed mainly on the establishment of protected areas and the landscape outside of reserves has often been overlooked (Ricketts 2001; Whittaker and Fernández-Palacios 2007; Wiens 2007; Bhagwat et al. 2008).

Since further expansion of the existing network of protected areas might be constrained by competing claims for other land uses (Bhagwat et al. 2008) and their effectiveness in representing species diversity has been questioned (Rodrigues et al. 2004; Chape et al. 2005), it is important to start understanding which habitats besides natural vegetation can support viable populations of species. Various recent studies have examined the role of the landscape surrounding reserves and have indicated the importance of high quality matrix environments in supporting biodiversity (Gascon et al. 1999; Ricketts 2001; Vandermeer and Carvajal 2001; Hughes et al. 2002; Perfecto and Vandermeer 2002; Sekercioglu et al. 2002; Watson et al. 2005; Bhagwat et al. 2008). This switch of attention away from fragments to the matrix has been referred to as ‘countryside biogeography’ (Daily et al. 2001, 2003) or ‘reconciliation ecology’ (Rosenzweig 2003).

The Azorean archipelago has undergone drastic changes in land use since the first settlers arrived 600 years ago. Originally, the islands were covered mostly by *Laurisilva* forests. However, these have been reduced to small fragments restricted to higher altitudes (Borges et al. 2005a, 2006). The islands are now mainly occupied by non-native habitats, such as intensive pastures for cattle, areas for agricultural crop production, forest patches of exotic plant species, and abandoned agricultural land and pastures covered with invasive plants (Martins 1993). Since further increasing the protected area network and the area of the native forest presents many practical challenges, it is important to understand how the changes in land use have

affected the Azorean biodiversity, and which habitats outside the original native forest are able to support indigenous species.

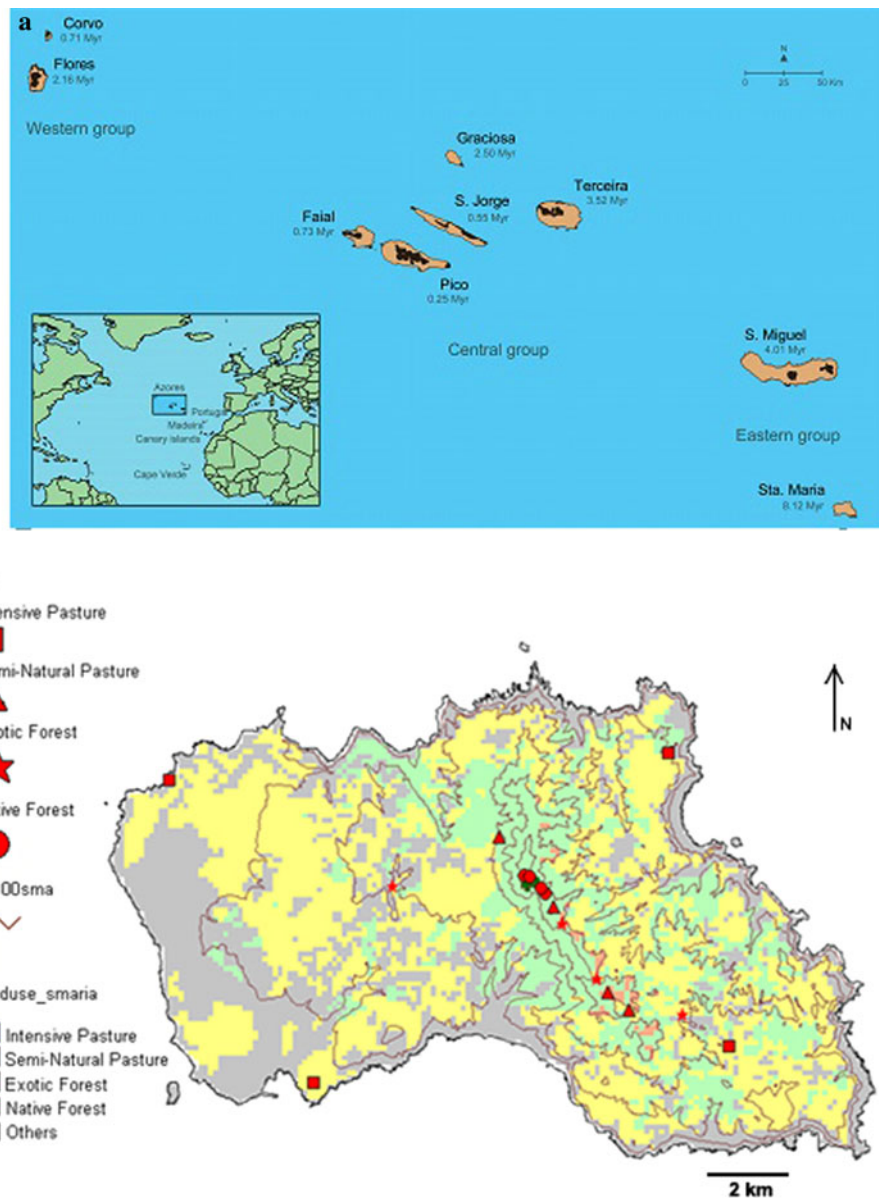
This study investigates how changes in land use have affected endemic, native and introduced arthropod richness and abundance on Santa Maria Island in the Azores. Santa Maria is the oldest island of the Azores, dating back 8.12 Myr (Abdel-Monem et al. 1975), which has resulted in higher numbers of endemic and native species compared to the other islands (Borges and Brown 1999; Borges and Hortal 2009). Arthropods were sampled using pitfall traps in four different habitat types (native forest, exotic forest, semi-natural pasture and intensive pasture) and classified as endemic (including the endemics from the target studied island, i.e. single-island-endemics—SIE), native non-endemic, or introduced. Native non-endemic species (“native” for simplicity) arrived in the Azores naturally and occur outside of the Azores as well. Introduced species are species that have been brought to the Azores by human activities. Where there was doubt over the colonisation status of a species, it was classified as native. In this study the following hypotheses will be tested: (i) species richness, abundance and diversity are highest in the native forest and lowest in the intensive pasture for Azorean endemic and native species, (ii) they are highest in the intensive pasture and lowest in the native forest for introduced species; (iii) the community composition of arthropods is different between the four habitat types; and (iv) an increase in distance to the native forest will result in a decrease in endemic, SIE and native arthropod richness and abundance in the other habitat types.

## Materials and methods

### Study area

This study was conducted in the Azores, an archipelago of nine islands situated in the North Atlantic (Fig. 1a). The islands are situated near the Mid-Atlantic Ridge and can be divided into three groups extending for 615 km: the western group (Flores and Corvo), the central group (Faial, Pico, São Jorge, Graciosa and Terceira) and the eastern group (São Miguel and Santa Maria). They are volcanic islands of recent origin, the youngest island is Pico (300,000 years) and Santa Maria is the oldest island (8.12 Myr) (Abdel-Monem et al. 1975; Borges and Hortal 2009). Santa Maria covers an area of 97 km<sup>2</sup> and the highest point is Pico Alto, with an elevation of 587 m. The temperate oceanic climate is strongly influenced by the proximity of the ocean and is characterised by high levels of humidity and small temperature fluctuations throughout the year. The average temperature on Santa Maria ranges between

**Fig. 1** Map of the region, showing **a** the nine islands of the Azores. Adopted from Borges et al. (2009); and **b** the four habitat types on Santa Maria and the position of the 16 transects (Land-use data extracted from DROTRH 2008)



14.3 and 17.5°C, with lowest temperatures in February and highest temperatures in August (Borges and Brown 1999). The average rainfall differs between the dry western part (752 mm year<sup>-1</sup>) and wet eastern part (1,386 mm year<sup>-1</sup>). January and February are generally the wettest months and August the driest (Borges and Brown 1999).

The original natural vegetation of the island is *Laurisilva* forest, a sub-tropical evergreen broadleaf laurel type forest, which now only covers a small part of the island (Fig. 1b). The small fragment of *Laurisilva* forest on Santa Maria covers about 9 ha (Gaspar et al. 2008) and the overstorey consists of the broadleaf species *Laurus azorica* and *Picconia azorica*, the short-leaf species *Erica azorica*, and the shrub *Vaccinium cylindraceum*, all of them Azorean endemics. In addition, this small fragment is now being invaded by the exotic invasive plants *Pittosporum*

*undulatum* and *Hedychium gardnerianum*, which have both become abundant. The *Laurisilva* forests have been reduced since humans arrived on the islands in the fifteenth century. The rate of land conversion increased significantly after the Second World War, when the Government reforested parts of the islands with *Cryptomeria japonica* plantations and reinforced the development of cattle-raising and great amounts of land were converted into pastures (Martins 1993). The vegetation on Santa Maria is now dominated by exotic forest (*Cryptomeria japonica*, *Eucalyptus* spp., *Acacia* spp. and *Pittosporum undulatum*), semi-natural pasture (not stocked year-round and with relatively low levels of fertilisation and introduced grass species) and intensive pasture (stocked year-round and with high levels of fertilisation and introduced grass and legume species).

## Site selection

On the island of Santa Maria, four sites were selected in each of the four main habitat types: native forest, exotic forest (*Cryptomeria japonica* plantations), semi-natural pasture and intensive pasture, resulting in a total of 16 sites (Fig. 1b). The sites were selected so as to try and sample most of the island. However, the native forest is restricted to higher elevations in the central area of the island and reduced to a 9 ha fragment, whereas the intensive pastures tend to be located at lower elevations closer to the sea. The sites in the native forest of Pico Alto were sampled in August 1997 and June 2004 (Borges et al. 2005a; Gaspar et al. 2008), and data were collected in the other three habitat types in June 2009 (Appendix 1).

## Arthropod sampling

For the arthropod sampling the protocol followed since 1998 by Project BALA (see Borges et al. 2000, 2005a, Gaspar et al. 2008) was applied. At each site, 30 pitfall traps were used to capture arthropods along a transect of 150 m length. The pitfall traps were plastic cups with a top diameter of 42 mm and 78 mm deep. The traps were dug into the soil so that the rim of the cup was at the same level as the soil surface. Half of the traps were filled with 60 ml of an attractive solution (Turquin) and the other half with 60 ml of anti-freeze liquid with a small proportion of ethylene glycol. A few drops of liquid detergent were added to both solutions to reduce surface tension. A white plastic plate was placed 5 cm above the traps with two pieces of wire to protect the trap from rainfall. The traps were placed 5 m apart, starting the transect with a Turquin trap and alternating with ethylene traps. The traps were left in the field for 2 weeks and then the samples were taken to the laboratory. The specimens were sorted into morpho-species by para-taxonomists and then into species by a senior taxonomist (P. A. V. Borges). Due to non-availability of taxonomic expertise, Acari, Collembola, Hymenoptera and Diptera were excluded from this study. Based on expert opinions, the species were classified into three colonisation groups (see also Borges et al. 2005a): endemic (including the single-island endemics, or SIE), native (i.e. native but non-endemic species) and introduced species.

## Data analysis

To test the first hypothesis, species richness, abundance, dominance and diversity were compared between the four habitat types (native forest, exotic forest, semi-natural pasture and intensive pasture) for Azorean endemic, native, introduced and total species. Due to the low replication of transects ( $n = 4$ ) within each habitat type, tests of normality

could not be performed and the data were analysed using non-parametric Kruskal–Wallis tests in PASW Statistics 17. In transect T07 in the semi-natural pasture, 10 pitfall traps were lost as a result of trampling by cows, and therefore the abundance numbers in each transect were converted to average abundances by dividing the total abundance in the transect by the number of pitfall traps in that transect (20 traps for T07 and 30 for all other transects). It was assumed that the 20 pitfall traps in T07 were enough to give a representative number for species richness, diversity and dominance. The Berger–Parker Index was used as a measure of dominance and Fisher’s Alpha as an indicator of diversity. The total number of species expected in each habitat type was calculated using a first-order Jack-knife estimator, selected because it is considered generally robust and insensitive to scale effects (see Hortal et al. 2006). For each habitat type, the observed number of species was divided by the expected number of species to obtain a completeness index. A value above 0.75 is generally accepted as representing a complete sampling effort (see also Borges and Brown 2003). Richness and abundance data for SIEs were also analysed with Kruskal–Wallis tests. SIEs are important species for conservation purposes since they are, by definition, restricted to Santa Maria Island.

Differences in species composition between the habitat types were compared with an analysis of similarities (ANOSIM) for Azorean endemic, native, introduced and total species composition to test the second hypothesis (Clarke 1993). The  $R$  statistic value ranges between 1 and  $-1$  and indicates the degree of difference between the groups, with a value of 1 signifying completely distinct communities. Detrended correspondence analyses (DCA) were carried out to illustrate the difference in species composition between the four habitat types for each group of Azorean endemic, native, introduced and total species (Oksanen and Minchin 1997). A DCA was used to remove the arch effect that was observed when using a correspondence analysis (analysis not shown). The number of axis rescalings was four and rare species were not down-weighted. The ANOSIM and DCA were carried out in Community Analysis Package 4.0 (Henderson and Seaby 2007).

To assess the fourth hypothesis, that the increasing distance from the native forest results in a decrease in the richness and abundance of arthropod species in the adjacent habitat types, the distance to the native forest for each transect was correlated with the number of species and the number of individuals for Azorean endemic, SIE, native, introduced, and total species. The distance of each transect to the native forest was calculated using Google Maps. The intensive pastures were excluded from this analysis as they were not surrounding the native forest and were located at lower elevations. The data for the exotic forest and semi-natural pasture were analysed together (eight sites) as well as



separately (two groups of four sites). Non-parametric Spearman rank correlation (available in PASW Statistics 17) was used as the sample size was again too low to test for normality.

**Results**

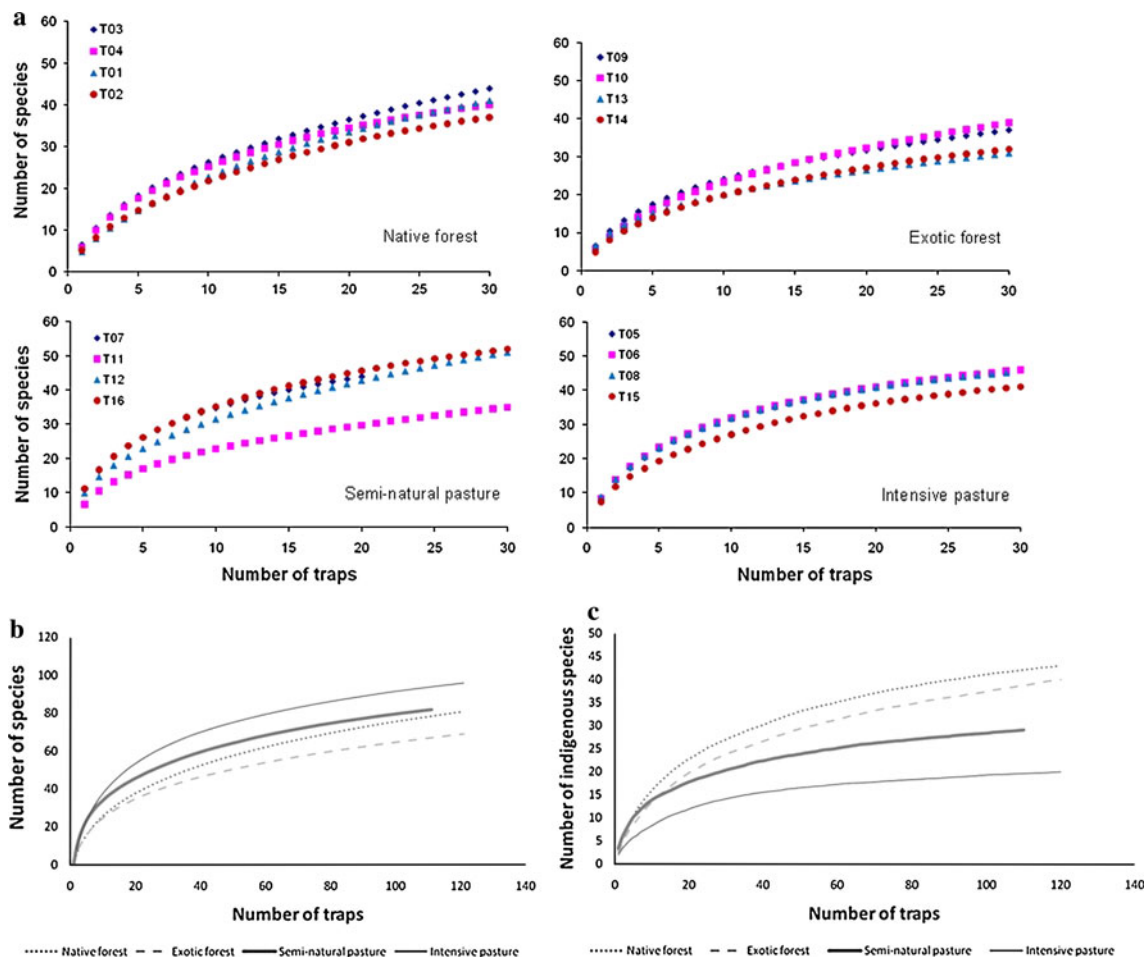
In this study, a total of 12,513 arthropod specimens were identified to species level in the four habitat types on Santa Maria (see Appendix 2). These specimens belonged to 188 different species, of which 25 were considered endemic to the Azores (including 9 SIEs), 58 were native to the archipelago (excluding the endemics) and 105 were introduced species.

The species accumulation curves for each habitat type are starting to level off, with the intensive pastures closest to asymptote (Fig. 2a). This indicates that most of the species that occur in this habitat type have been sampled in

this study, while a small number remain to be recorded in the other habitats. The same pattern can be observed for the completeness index of each habitat type, which was over or almost at the 0.75 level, indicating that about enough species were sampled to get a representative assessment of the composition and relative richness of each habitat (Appendix 3).

**Species richness, abundance and diversity**

Total species richness and abundance were significantly different between the four habitat types on Santa Maria (Kruskal–Wallis test,  $H = 8.032, P = 0.045$  for species richness;  $H = 8.316, P = 0.040$  for species abundance, Table 1). Species richness differed significantly between the four habitat types for Azorean endemics, native and introduced species (Kruskal–Wallis test,  $H = 12.555, P = 0.006$ ;  $H = 10.296, P = 0.016$ ;  $H = 12.192, P = 0.007$ , respectively). As hypothesised, Azorean endemic and native



**Fig. 2** Species accumulation curves, showing **a** curves for the total number of species (1,000 randomisations) for each transect (each containing 30 pitfall traps) per habitat type (each containing 4 transects); **b** curves for the total number of species (1,000

randomisations) for each habitat type (each containing 4 transects with 30 pitfall traps) and **c** curves for the number of indigenous species (excluding the introduced species) (1,000 randomisations) for each habitat type (each containing 4 transects with 30 pitfall traps)

**Table 1** Mean ( $\pm 1$  SE) richness (number of species), abundance (number of individuals), dominance (1/Berger-Parker Index) and diversity (Fisher's alpha) values for Azorean endemic, native (excluding endemics), introduced and total species for the four habitat types and results of the Kruskal–Wallis tests

		Native forest	Exotic forest	Semi-natural pasture	Intensive pasture	<i>H</i>	<i>df</i>	<i>P</i>
Richness	Azorean endemic	10.0 (0.7)	5.3 (0.8)	4.0 (0.7)	1.5 (0.5)	12.555	3	<b>0.006</b>
	Native	15.8 (0.9)	14.0 (1.1)	13.3 (1.1)	7.5 (0.7)	10.296	3	<b>0.016</b>
	Introduced	14.8 (1.4)	15.5 (1.4)	28.3 (2.6)	35.8 (1.7)	12.192	3	<b>0.007</b>
	Total	40.5 (1.4)	34.8 (1.9)	45.5 (3.9)	44.8 (1.3)	8.032	3	<b>0.045</b>
Abundance	Azorean endemic	73.5 (25.0)	56.5 (34.6)	117.3 (52.1)	223.0 (111.0)	2.713	3	0.438
	Native	78.5 (8.3)	190.8 (48.5)	214.8 (71.1)	233.3 (154.4)	5.352	3	0.148
	Introduced	245.3 (64.8)	429.8 (117.0)	503.8 (141.6)	762.0 (135.1)	5.846	3	0.119
	Total	397.3 (82.7)	677.0 (124.3)	835.8 (171.6)	1218.3 (228.5)	8.316	3	<b>0.040</b>
Dominance	Azorean endemic	2.6 (0.3)	1.8 (0.4)	1.2 (0.1)	0.8 (0.3)	10.579	3	<b>0.014</b>
	Native	3.7 (0.1)	1.6 (0.3)	2.4 (0.5)	2.0 (0.7)	6.066	3	0.108
	Introduced	2.5 (0.1)	2.1 (0.3)	3.4 (0.7)	3.4 (0.9)	4.787	3	0.188
	Total	4.1 (0.2)	3.2 (0.8)	4.6 (0.5)	3.8 (0.9)	3.375	3	0.337
Diversity	Azorean endemic	3.6 (0.4)	2.2 (0.7)	0.9 (0.1)	0.3 (0.03)	10.975	3	<b>0.012</b>
	Native	6.1 (0.8)	3.7 (0.5)	3.8 (0.9)	2.0 (0.3)	11.184	3	<b>0.011</b>
	Introduced	3.5 (0.3)	3.2 (0.1)	6.7 (0.6)	8.0 (0.8)	12.044	3	<b>0.007</b>
	Total	11.7 (0.8)	8.0 (0.5)	10.5 (0.6)	9.7 (1.2)	7.013	3	0.071

In each habitat type, four 150 m long transects containing 30 pitfall traps were sampled. Probabilities in bold are significant at  $P < 0.05$ .

species richness were both highest in the forest habitats (native and exotic forest) and lowest in the semi-natural and intensive pasture, while the number of introduced species was lowest in the native forest and highest in the intensive pasture (Fig. 3). Total species abundance was highest in the intensive pasture and lowest in the native forest. The same trend was apparent when the species were separated into Azorean endemic, native and introduced species (Fig. 3), although these differences were not significant.

Interestingly, when the species accumulation curves are plotted with all the species as one curve for each habitat type (Fig. 2b), the curve for the native forest comes out below that of the two pasture types. However, when the same curves are plotted for just the indigenous species (so excluding the exotic species) the native and exotic forest come out above the curves for the pastures (Fig. 2c). This indicates that the greater richness of the pasture sites is due to the non-natives and the curve levels when they are excluded.

Species dominance was significantly different between the four habitat types for Azorean endemic species (Kruskal–Wallis test,  $H = 10.579$ ,  $P = 0.014$ ), with the intensive pastures being most dominated by a few species and the native forest having the least dominance (Table 1). Species diversity was not significantly different between the four habitat types for total species richness (Kruskal–Wallis test,  $H = 7.013$ ,  $P = 0.071$ ), but did differ significantly for Azorean endemic, native and introduced species (Kruskal–Wallis test;  $H = 10.975$ ,  $P = 0.012$  for Azorean endemic species,  $H = 11.184$ ,  $P = 0.011$  for native

species; and  $H = 12.044$ ,  $P = 0.007$  for introduced species). The native forest had the highest value of Fisher's alpha for Azorean endemic and native species, and the lowest for the intensive pasture. In contrast, introduced species diversity was highest in the intensive pasture, and lower in the forest habitats (Table 1).

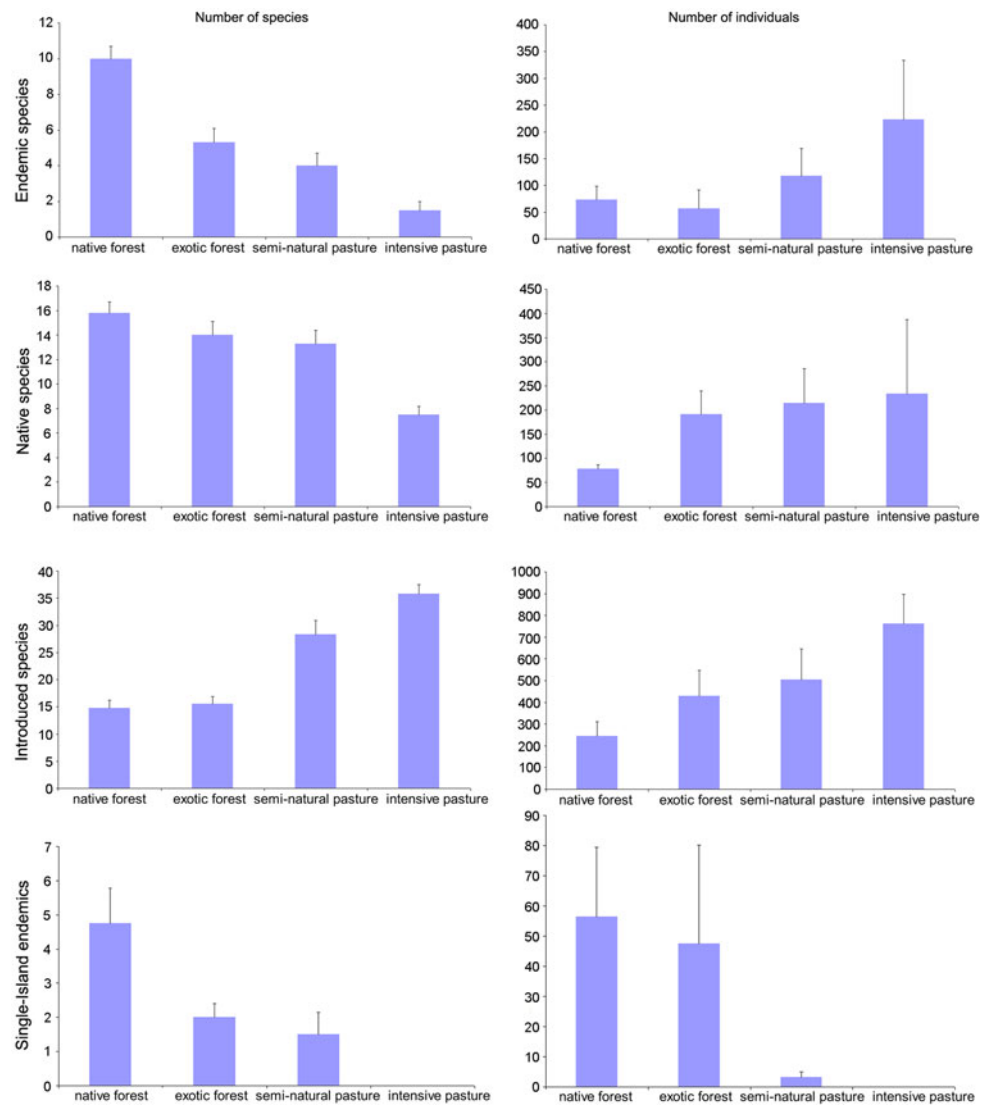
#### Single-island endemics

Of the 25 Azorean endemic species found in this study on Santa Maria, nine species are single-island endemics (SIEs), i.e. only found on Santa Maria (Table 2). The number of SIEs differed between the four habitat types (Kruskal–Wallis test;  $H = 11.465$ ,  $P = 0.009$ ), with the highest number found in the native forest and none in the intensive pasture (Fig. 3). The abundance of the single-island endemics was also significantly different between the four habitat types (Kruskal–Wallis test;  $H = 11.921$ ,  $P = 0.008$ ). The average abundance was highest in the native forest (Fig. 3), however, the highest abundance at transect level was recorded in one of the exotic forest transects, in which 145 individuals of the beetle *Tarphius pomboi* were found.

#### Species composition

In accordance with the third hypothesis, the total species composition of the arthropod communities was statistically different between the four habitat types (ANOSIM;

**Fig. 3** Mean ( $\pm 1$  SE) number of arthropod species and number of arthropod individuals per transect (four transects per habitat, each transect containing 30 pitfall traps) for Azorean endemic, native (excluding endemics), introduced and single-island endemic species found in the four habitat types



**Table 2** The number of single-island endemic species and the percentage single-island endemics out of the total number of Azorean endemic species in the four habitat types on Santa Maria

	Native forest	Exotic forest	Semi-natural pasture	Intensive pasture	Total
Azorean endemics	17	13	8	3	25
Single-island endemics	7	5	4	0	9
%	41.2	38.5	50.0	0.0	36.0

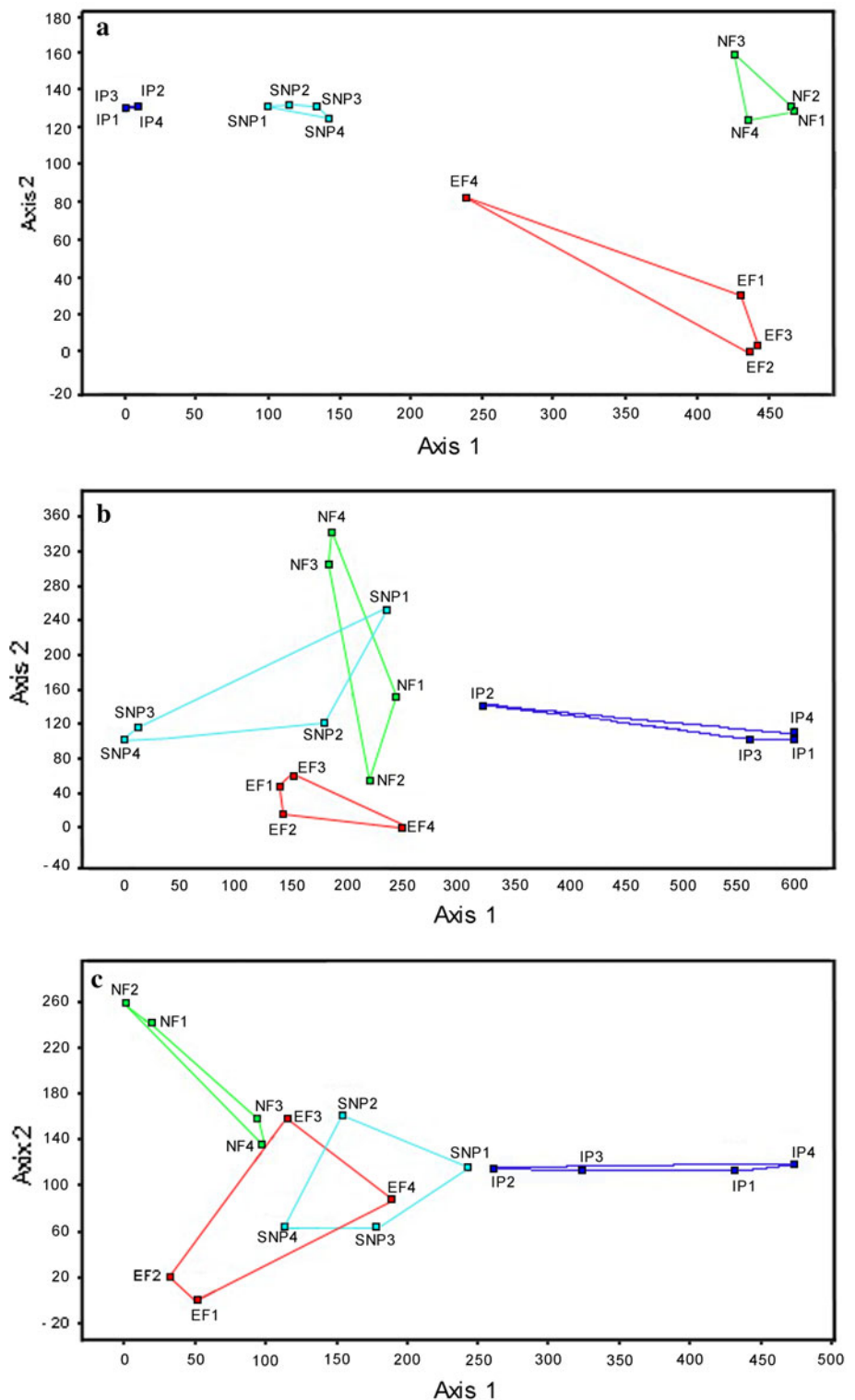
In each habitat type, four 150 m long transects containing 30 pitfall traps were sampled

$R = 0.787$ ,  $P = 0.001$ ). The arthropod community in native forest has a larger proportion of Azorean endemic and native species compared to the other habitat types, whereas the semi-natural and intensive pasture have relatively more introduced species (Table 3). In addition, the

species composition was also different between the four habitat types for the three species groups (ANOSIM;  $R = 0.823$ ,  $P = 0.001$  for Azorean endemic species;  $R = 0.597$ ,  $P = 0.001$  for native species; and  $R = 0.602$ ,  $P = 0.001$  for introduced species), suggesting that the four habitat types have different Azorean endemic, native and introduced species. All pairwise comparisons were significant as well, except for Azorean endemic species composition in natural forest vs. exotic forest (ANOSIM;  $R = 0.146$ ,  $P = 0.157$ ) and for introduced species composition in exotic forest vs. semi-natural pasture (ANOSIM;  $R = 0.302$ ,  $P = 0.1$ ).

The differences in species composition between the four habitat types were further supported by the detrended correspondence analysis (DCA). The ordination of the sites is a product of the distribution of species across the sites and the first two axes represent the two most prominent compositional gradients inherent to the data (Fig. 4). For

**Fig. 4** Sites ordination plots by DCA for **a** endemic, **b** native and **c** introduced species composition. In each habitat type, four 150 m long transects containing 30 pitfall traps were sampled. *NF* native forest, *EF* exotic forest, *SNP* semi-natural pasture, *IP* intensive pasture



the Azorean endemic species, the habitat types were clearly clustered and each habitat type formed a separate group. For native and introduced species composition there was more overlap between the four habitat types, suggesting that these species are more uniformly spread. The first axis of the ordination separates the forests from the pastures for

Azorean endemic, native and introduced species (eigenvalues axis 1: 0.965; 0.972 and 0.741, respectively). The second axis separates the native forest from the exotic forest (eigenvalues axis 2: 0.277 for Azorean endemic species; 0.519 for native species and 0.245 for introduced species, Fig. 4).



**Table 3** The total number of Azorean endemic, native (excluding endemics) and introduced species in the four habitat types and the number and percentage of species that are unique to each habitat type

	Azorean endemic				Native				Introduced			
	NF	EF	SNP	IP	NF	EF	SNP	IP	NF	EF	SNP	IP
Total	17	13	8	3	28	27	22	18	32	28	49	67
Unique	9	4	1	1	9	1	3	5	6	1	7	28
% Unique	52.9	30.8	12.5	33.3	32.1	3.7	13.6	27.8	18.8	3.6	14.3	41.8

In each habitat type, four 150 m long transects containing 30 pitfall traps were sampled  
*NF* native forest, *EF* exotic forest, *SNP* semi-natural pasture, *IP* intensive pasture

Species uniqueness

From the 17 Azorean endemic species that are found in the native forest, nine species occur exclusively in the native forest and in none of the other habitat types (Table 3). This means that for at least 50% of the Azorean endemic species on Santa Maria, the native forest is the only place where they occur. The exotic forest has four out of 13 Azorean endemic species that are unique to this habitat type, whereas in the semi-natural pasture only one out of 8 Azorean endemics is restricted to this habitat type. There are three Azorean endemic species that occur on the intensive pasture, of which one is unique to this type of habitat.

For the native species, nine out of 28 species found in the native forest were not found outside of this habitat type, which represents about one-third of the native species found in the native forest (Table 3). In the exotic forest, just one out of 27 native species seems to occur solely in this habitat type, whereas the semi-natural pasture has three out of 22 native species that are unique. Surprisingly, the intensive pasture is home to five native species that occur exclusively in this habitat type, which is more than one-fourth of the total of 22 native species found in the intensive pastures.

For the introduced species, there is an opposite trend (Table 3). The native forest has only six introduced species restricted to this habitat type out of a total of 32. In the exotic forest there is only one introduced species that does not occur in any of the other habitat types. The semi-natural pasture has seven out of 49 species that only occur in

this habitat type. The intensive pasture has the highest proportion of unique introduced species, with 28 out of 67 species (40%) only occurring in this habitat type.

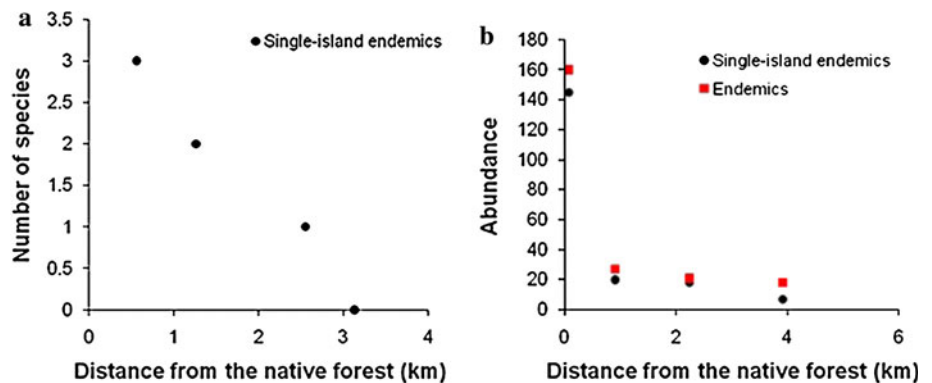
Distance from the native forest

There was no significant correlation between the distance from the native forest and species richness and abundance when the exotic forest and semi-natural pasture were grouped together. However, when these two habitat types were analysed separately, there were some significant correlations that support the third hypothesis. In the semi-natural pasture, the number of SIE species decreased with increasing distance from the native forest (Spearman correlation;  $r = -1, P < 0.01$ ; Fig. 5a). For the exotic forest, the abundance of both Azorean endemics and SIEs decreased with increasing distance from the native forest (Spearman correlation;  $r = -1, P < 0.01$ , Fig. 5b).

Discussion

Our findings support the proposition that land use change has significantly affected arthropod species richness and abundance on Santa Maria. As hypothesised, the numbers of Azorean endemic and native species were highest in the native forest and they were lowest in the highly modified intensive pasture. In addition, the intensive pastures had the

**Fig. 5** Relationship between distance to the native forest and **a** a number of SIE species in the semi-natural pasture and **b** abundance of endemic and SIE species in the exotic forest. In each habitat type, four 150 m long transects containing 30 pitfall traps were sampled



highest introduced species richness and abundance. Following some recent work confirming the relevance of native forests for the conservation of Azorean endemic arthropod biodiversity (e.g. Borges et al. 2005a, 2006, 2008; Ribeiro et al. 2005; Gaspar et al. 2008; Cardoso et al. 2009), our findings reinforce the important role of the native forest in arthropod conservation on the Azores, with 7 out of 9 sampled SIEs being found in the native forest and 50% of the sampled Azorean endemic species being confined to this habitat type. There seems to be a gradient of land-use change effects on arthropods, where an increase in land-use intensity results in more negative impact on the indigenous (i.e. endemic and/or native) arthropod community. These results are in agreement with other studies in the Azores (Borges and Wunderlich 2008; Borges et al. 2008; Cardoso et al. 2009) and elsewhere (Driscoll 2005; Schweiger et al. 2005; Jana et al. 2006; Rainio and Niemelä 2006; Acevedo and Restrepo 2008; Schmidt et al. 2008), that have demonstrated negative effects of such land-use change for several taxa.

Although not as good habitat as the native forest, the exotic forest and semi-natural pasture performed better than the intensive pasture, which is in accordance with results obtained for Terceira Island (Cardoso et al. 2009). These two habitat types harboured higher numbers of Azorean endemics and native species and had a community composition more similar to the native forest compared to the intensive pasture. Similarly, other studies have found that habitat types other than pristine forest, such as agricultural fields and secondary forest can support certain species and help to maintain biodiversity (Burel et al. 1998; Toth and Kiss 1999; Grill et al. 2005). Our results support the notion that the quality of the matrix is important in species conservation, as suggested by several other studies (Ricketts 2001; Vandermeer and Carvajal 2001; Perfecto and Vandermeer 2002; Watson et al. 2005; Prugh et al. 2008; Muriel and Kattan 2009).

The community composition of the arthropods differed between the four habitat types, with higher proportions of Azorean endemics in the forest habitats and more introduced species in the pastures. This is in agreement with the third hypothesis. When clustering the four habitat types according to species composition for each colonisation group, the clustering was most clear for the Azorean endemic species, suggesting that the habitat types harboured different endemic species. The overlap in species composition was most obvious for introduced species, which indicates that introduced species are able to occupy several habitat types on the island and spread easily. In doing so, introduced species form one of the major threats to terrestrial biodiversity in the Azores (Martins 1993; Borges et al. 2006, 2008, 2009).

Interestingly, the total abundance of Azorean endemic species was highest on the intensive pastures and not in the

native forest, which is not in accord with the first hypothesis. However, this was almost entirely the consequence of the great abundance of one beetle species, *Heteroderes azoricus*, of which 868 individual specimens were collected in the intensive pastures. Of the eight Azorean endemic species occurring in the semi-natural pasture, the spider *Pardosa acorensis* was also highly abundant, a pattern found in most of the Azorean islands (see also Borges and Wunderlich 2008; Cardoso et al. 2009). The number of Azorean endemic species inhabiting the pastures is low, yet these two generalist species are able to maintain populations with higher abundances than most endemic species in the native forest. The same pattern was found in a recent study on Terceira Island in the Azores (Cardoso et al. 2009), where these two species were found in high numbers in the pastures as well. Similar to our findings, Azorean endemic and native species richness on Terceira were also highest in the native forest and the intensive pasture had the highest number of introduced species. Cardoso et al. (2009) conclude that these two habitat types are the main drivers of the species composition of a particular site, with semi-natural pastures and exotic forests functioning as ‘connector habitats’ between natural forests. However, in the case of Santa Maria, exotic *Cryptomeria japonica* plantations seem to have a quite different role, accumulating a high density of endemic saproxylic beetles (see also below).

In addition to the endemic abundant and widespread *Heteroderes azoricus* and *Pardosa acorensis*, other endemic species are particularly important in Santa Maria, notably the nine SIE species exclusive to Santa Maria. The semi-natural pastures harboured four SIEs (*Tarphius pomboi*, *Tarphius rufonodulosus*, *Catops velhocabralli*, *Olisthopus inclavatus*), three of which are saproxylic beetle species. The exotic forest is home to five SIE saproxylic beetle species (*Tarphius serranoi*, *Tarphius pomboi*, *Tarphius rufonodulosus*, *Catops velhocabralli*, *Caulotrumpis parvus*), of which the species *Tarphius pomboi* was particularly abundant. This might be explained by the fact that this species feeds on wood-inhabiting fungi and is able to benefit from the decaying wood found in the exotic forest in Santa Maria, as has been shown for other saproxylic beetle species in northern Europe (Okland et al. 1996; Kaila et al. 1997; Martikainen et al. 2000; Jonsson et al. 2005).

The fact that so many SIE forest-specialist beetles are found in both exotic forest and semi-natural pasture sites surrounding the native forest fragment of Santa Maria is a novel finding in the Azores. Patterns of endemic species distribution in other Azorean islands are not so positive, with most of the endemic species being restricted to native forest or having only sink populations in nearby human modified habitats (see Borges et al. 2008). The management practices

in Santa Maria in both exotic *Cryptomeria japonica* plantations and semi-natural pastures are obviously different compared to similar habitats on the other islands. In fact, particularly in *C. japonica* plantations, the management is less intensive or even almost absent, leading to a more diverse vegetation and high density of dead wood favourable for many dead-wood and fungi-eating endemic saproxylic beetles. The *C. japonica* plantations surrounding the Pico Alto native forest fragment were planted in the 1960s to catch water from the fog and improve water collection for human use. Consequently, the past and current management of these human-modified exotic forests has a positive effect on the diversity of endemic arthropods on Santa Maria. Despite the fact that Santa Maria has only 0.2% of the total area of native forest in the Azores (Gaspar et al. 2008), it is home to 57 endemic arthropod species (ATLANTIS database consulted in July 2009), which makes up 20% of the total number of Azorean endemic arthropod species. As such, Santa Maria can be considered as a ‘hotspot’ among the Azorean islands (see also Borges and Brown 1999; Borges et al. 2005a, b; Borges and Hortal 2009).

In accordance with the fourth hypothesis, our results showed also that with increasing distance from the native forest, the abundance of both Azorean endemics and SIEs decreased in the exotic forest and the number of SIE species decreased in the semi-natural pasture. Similar patterns of decreasing species numbers with increasing distance from natural forest have been found elsewhere (e.g. Klein et al. 2006). This suggests that the favorability of these two habitats is not general within the island, since their ability to support arthropods important to conservation depends on where they are located relative to the native forest. In addition, it is not clear what will be the long-term management of the *C. japonica* plantations surrounding the Pico Alto native forest fragment, casting some doubts on the successful conservation of saproxylic invertebrates on this island.

This study has some important implications for conservation management activities on Santa Maria. We clearly demonstrated the importance and uniqueness of the native forest for arthropod conservation and recommend that efforts should be made to protect the small patches of native forest that are left on the island. This small area of native forest is critical for the conservation of both archipelagic and single-island endemic species (Borges et al. 2005a). This study clearly showed that the exotic forest and semi-natural pasture are better habitats for these arthropods than the intensive pastures (see also Borges et al. 2008; Cardoso et al. 2009). Our results indicate that it would be better to replace disappearing native forest or abandoned fields with less intensively managed habitats such as exotic forest and semi-natural pasture, rather than intensive

pastures (see also Cardoso et al. 2009), although conservation of the original habitats is of course by far the best option.

The species accumulation curves suggest that not all arthropod species that occur in the different habitat types have been sampled in this study, and this is particularly true for the native forest for which the accumulation curve is only just about to start levelling off. The latter is probably due to the fact that a lower number of individuals was captured in the native forest, but this should not be considered a bias. The fact that not all arthropod species have been sampled is not a problem for this study, as the aim was not to obtain a complete list of species for Santa Maria, but to compare arthropod species richness and abundance between the different habitat types. This finding does suggest that further sampling is likely to find more species on Santa Maria, especially in the native forest. Moreover, it is important that the arthropods on the Azores are the subject of long-term studies, as the effects of native habitat that has been lost already might become pronounced in the future. Due to this so-called ‘extinction debt’ (Tilman et al. 1994), many more arthropods might be on the brink of extinction than current studies have shown (Triantis et al. 2010).

#### Implications for the conservation of S. Maria endemic arthropods

A regional strategy for the long-term persistence of SIEs from S. Maria is needed. We suggest: (i) it is important to monitor the abundance of SIEs in the different habitats over the following decades following the Long-Term Ecological Research (LTER) strategy (see e.g. Pereira and Cooper 2006); (ii) removal of *P. undulatum* and other plant invaders and its progressive replacement by native woody species; (iii) in addition, pastures and forest plantations surrounding the small area on native forest at Pico Alto should retain their current low-intensity management in such a way that they will be able to support species that will otherwise be driven to extinction in the long term.

This study demonstrates that the matrix quality in the form of less intensive agricultural management is important for the conservation of rare endemic saproxylic invertebrates in the Azores. The studied native forest fragment and part of the surrounding area is now a protected area named as Santa Maria Park using the IUCN Protected Areas Management Categories System (see Borges et al. 2005a). The Pico Alto area was classified as Category IV (protected area for the management of habitats and species), which implies some restrictions on human activities. Our results support the importance of maintaining the diverse matrix surrounding Pico Alto, to ensure the

continuing availability of wood in several stages of decay, essential for saproxylic arthropods.

**Acknowledgments** We are very thankful for the support in the fieldwork by the Ecoteca of St. Maria and to the Serviços de Desenvolvimento Agrário de St. Maria. We thank Fernando Pereira and João Moniz for the help with setting up the traps and collecting of the insects, and Vasiliki Orfanou and João Moniz for helping with the

sorting and identification of the arthropods. Enésima Mendonça is thanked for her help in preparing the map of Santa Maria, and Kostas Triantis for giving useful comments on the introduction. This study was supported by a Huygens Scholarship Programme (HSP) grant and a VSB Foundation grant to S. S. M. and financial support in the Azores was provided by project “Consequences of land-use change on Azorean fauna and flora—the 2010 Target” (Ref: Direcção Regional da Ciência e Tecnologia M.2.1.2/I/003/2008) to P. A. V. B.

## Appendix 1

See Table 4.

**Table 4** Information on all transects in the four habitat types on Santa Maria

Name	Transect	Sampled	Habitat	Longitude	Latitude	Elevation (m)
Pico Alto	T01	August 1997	Native forest	669927	4094384	530
Pico Alto	T02	August 1997	Native forest	669823	4094394	530
Pico Alto	T03	June 2004	Native forest	670257	4094043	479
Pico Alto	T04	June 2004	Native forest	670170	4094140	479
Aeroporto—Cabrestantes	T05	June 2009	Intensive pasture	662662	4096328	43
Pico do Facho	T06	June 2009	Intensive pasture	665588	4090185	52
Casas Velhas	T07	June 2009	Semi-natural pasture	671945	4091662	379
Fonte Jordão	T08	June 2009	Intensive pasture	673963	4090920	271
Pico Alto—L	T09	June 2009	Exotic forest	670185	4094194	464
Cruzamento Pico Alto	T10	June 2009	Exotic forest	670608	4093423	419
Pico Alto—PSN	T11	June 2009	Semi-natural pasture	670417	4093748	482
Fontinhas—PSN	T12	June 2009	Semi-natural pasture	671513	4092040	448
Fontinhas	T13	June 2009	Exotic forest	671308	4092293	419
St. Espírito	T14	June 2009	Exotic forest	673033	4091592	318
Ponta do Norte	T15	June 2009	Intensive pasture	672741	4096864	219
Alto da Nascente	T16	June 2009	Semi-natural pasture	669313	4095181	296

Long/Lat coordinates are presented in UTM's

## Appendix 2

See Table 5.

**Table 5** Taxonomic list of all the species found, with the indication of the abundance of each species in the four types of land-use

N/E/I	Code MF	Species	Class	Order	Family	NF	EF	SNP	IP
E	MF134	<i>Gibbaranea occidentalis</i> Wunderlich	Arachnida	Araneae	Araneidae	3		1	
E	MF17	<i>Pardosa acorensis</i> Simon	Arachnida	Araneae	Lycosidae	4	1	410	23
E	MF5	<i>Rugathodes acorensis</i> Wunderlich	Arachnida	Araneae	Theridiidae		6		
E	MF50	<i>Lepthyphantes acorensis</i> Wunderlich	Arachnida	Araneae	Linyphiidae	6			
E	MF526	<i>Neon acorensis</i> Wunderlich	Arachnida	Araneae	Salticidae		1		
I	MF1014	Gen. sp.	Arachnida	Araneae	Salticidae				2
I	MF1047	<i>Heliophanus kochi</i> Simon	Arachnida	Araneae	Salticidae				4
I	MF1100	<i>Zelotes tenuis</i> (C.L. Koch)	Arachnida	Araneae	Gnaphosidae				3
I	MF1071	<i>Chalcoscirtus infimus</i> (Simon)	Arachnida	Araneae	Salticidae				5
I	MF1118	<i>Micaria pallipes</i> (Lucas)	Arachnida	Araneae	Gnaphosidae				10

**Table 5** continued

N/E/I	Code MF	Species	Class	Order	Family	NF	EF	SNP	IP
I	MF122	<i>Clubiona terrestris</i> Westring	Arachnida	Araneae	Clubionidae	1		6	
I	MF127	<i>Steatoda grossa</i> (C.L. Koch)	Arachnida	Araneae	Theridiidae	1			
I	MF133	<i>Metellina merianae</i> (Scopoli)	Arachnida	Araneae	Tetragnathidae	3			
I	MF140	<i>Ero furcata</i> (Villers)	Arachnida	Araneae	Mimetidae	1	1		
I	MF21	<i>Tenuiphantes tenuis</i> (Blackwall)	Arachnida	Araneae	Linyphiidae	125	49	65	9
I	MF233	<i>Oedothorax fuscus</i> (Blackwall)	Arachnida	Araneae	Linyphiidae			378	664
I	MF234	<i>Erigone autumnalis</i> Emerton	Arachnida	Araneae	Linyphiidae			5	67
I	MF24	<i>Eperigone fradeorum</i> (Berland)	Arachnida	Araneae	Linyphiidae				17
I	MF250	<i>Xysticus nubilus</i> Simon	Arachnida	Araneae	Thomisidae			4	66
I	MF28	<i>Dysdera crocata</i> C. L. Koch	Arachnida	Araneae	Dysderidae	12	5	7	25
I	MF330	<i>Meioneta fuscipalpis</i> (C.L. Koch)	Arachnida	Araneae	Linyphiidae				32
I	MF333	<i>Ostearius melanopygius</i> (O. P.-Cambridge)	Arachnida	Araneae	Linyphiidae		1	1	4
I	MF334	Gen. sp.	Arachnida	Araneae	Linyphiidae			2	
I	MF34	<i>Erigone atra</i> (Blackwall)	Arachnida	Araneae	Linyphiidae				2
I	MF488	<i>Achaearanea acoreensis</i> (Berland)	Arachnida	Araneae	Theridiidae		1	5	4
I	MF489	<i>Lycosoides coarctata</i> (Dufour)	Arachnida	Araneae	Agelenidae				2
I	MF701	<i>Haplodrassus signifer</i> (C. L. Koch)	Arachnida	Araneae	Gnaphosidae				1
I	MF988	<i>Oecobius similis</i> Kulczynski	Arachnida	Araneae	Oecobiidae				6
I	MF998	<i>Zelotes aeneus</i> (Simon)	Arachnida	Araneae	Gnaphosidae				38
I	MF688	<i>Clubiona genevensis</i> L. Koch	Arachnida	Araneae	Clubionidae				30
N	MF117	<i>Lathys dentichelis</i> (Simon)	Arachnida	Araneae	Dictynidae	1			
N	MF2	<i>Tenuiphantes miguelensis</i> Wunderlich	Arachnida	Araneae	Linyphiidae	34	18		
N	MF20	<i>Palliduphantes schmitzi</i> (Kulczynski)	Arachnida	Araneae	Linyphiidae	9	15	2	
N	MF33	<i>Homalenotus coriaceus</i> (Simon)	Arachnida	Opiliones	Phalangiidae	61	24	139	11
N	MF103	<i>Chthonius tetrachelatus</i> (Preysler)	Arachnida	Pseudoscorpiones	Chthoniidae	2		5	
N	MF38	<i>Chthonius ischnocheles</i> (Hermann)	Arachnida	Pseudoscorpiones	Chthoniidae	1	6		
N	MF26	<i>Geophilus truncorum</i> Bergsøe & Meinert	Chilopoda	Geophilomorpha	Geophilidae				17
N	MF27	<i>Lithobius pilicornis pilicornis</i> Newport	Chilopoda	Lithobiomorpha	Lithobiidae		1		
N	MF1006	<i>Lithobius</i> sp.	Chilopoda	Lithobiomorpha	Lithobiidae				4
I	MF336	<i>Scutigera coleoptrata</i> (Linnaeus)	Chilopoda	Scutigermomorpha	Scutigeridae				42
N	MF468	<i>Haplobainosoma lusitanum</i> Verhoeff	Diplopoda	Chordeumatida	Haplobainosomatidae	3			
I	MF309	<i>Choneiulus palmatus</i> (Nemec)	Diplopoda	Julida	Blaniulidae	83		2	
I	MF9	<i>Ommatoiulus moreletii</i> (Lucas)	Diplopoda	Julida	Julidae	218	194	330	93
I	MF48	<i>Blaniulus guttullatus</i> (Fabricius)	Diplopoda	Julida	Blaniulidae	305	101		
I	MF62	<i>Brachyiulus pusillus</i> (Leach)	Diplopoda	Julida	Julidae	3	30	9	2
N	MF282	<i>Cylindroiulus propinquus</i> (Porat)	Diplopoda	Julida	Julidae	15	20	11	14
N	MF49	<i>Nopoiulus kochii</i> (Gervais)	Diplopoda	Julida	Blaniulidae	16	2		
N	MF53	<i>Proteroiulus fuscus</i> (Am Stein)	Diplopoda	Julida	Blaniulidae	26	8	1	
N	MF544	<i>Cylindroiulus latestriatus</i> (Curtis)	Diplopoda	Julida	Julidae	5			
I	MF71	<i>Oxidus gracilis</i> (C.L.Koch)	Diplopoda	Polydesmida	Paradoxosomatidae	20	82	33	1
N	MF37	<i>Polydesmus coriaceus</i> Porat	Diplopoda	Polydesmida	Polydesmidae	19	510	40	
N	MF59	<i>Zetha vestita</i> (Brullé)	Insecta	Blattaria	Polyphagidae	32	10	35	9
N	MF1113	<i>Loboptera decipiens</i> (Germar)	Insecta	Blattaria	Blattellidae				803
E	MF107	<i>Atlantocis gillerforsi</i> Israelson	Insecta	Coleoptera	Ciidae		2		
E	MF1098	<i>Olisthopus inclavatus</i> Istaelson	Insecta	Coleoptera	Carabidae			6	
E	MF128	<i>Caulotrups parvus</i> Israelson	Insecta	Coleoptera	Curculionidae	6	1		
E	MF138	<i>Athous pomboi</i> Platia & Borges	Insecta	Coleoptera	Elateridae	3			
E	MF244	<i>Alestrus dolosus</i> (Crotch)	Insecta	Coleoptera	Elateridae				1



Table 5 continued

N/E/I	Code MF	Species	Class	Order	Family	NF	EF	SNP	IP
E	MF540	<i>Heteroderes azoricus</i> Tarnier	Insecta	Coleoptera	Elateridae		2	42	868
E	MF64	<i>Tarphius serranoi</i> Borges	Insecta	Coleoptera	Zopheridae	108	5		
E	MF70	<i>Tarphius pomboi</i> Borges	Insecta	Coleoptera	Zopheridae	103	176	3	
E	MF73	<i>Metopthalmus occidentalis</i> Israelson	Insecta	Coleoptera	Lathridiidae	4			
E	MF76	<i>Tarphius rufonodulosus</i> Israelson	Insecta	Coleoptera	Zopheridae	1	7	3	
E	MF77	<i>Pseudechinosoma nodosum</i> Hustache	Insecta	Coleoptera	Curculionidae	10	3		
E	MF84	<i>Catops velhocabrali</i> Blas & Borges	Insecta	Coleoptera	Leiodidae		1	1	
E	MF85	<i>Tarphius depressus</i> Gillerfors	Insecta	Coleoptera	Zopheridae	6	20	3	
I	MF108	<i>Cryptamorphia desjardinsii</i> (Guérin-Méneville)	Insecta	Coleoptera	Silvanidae		1	21	
I	MF1097	Gen. sp.	Insecta	Coleoptera	Lyctidae		2		
I	MF1099	<i>Carpophilus</i> sp.	Insecta	Coleoptera	Nitidulidae			1	
I	MF1101	Gen. sp.	Insecta	Coleoptera	Cucujidae				40
I	MF1105	<i>Bruchus</i> sp.	Insecta	Coleoptera	Chrysomelidae				2
I	MF1106	<i>Sitona</i> sp.	Insecta	Coleoptera	Curculionidae				1
I	MF1110	Gen. sp.	Insecta	Coleoptera	Corylophidae			1	
I	MF1111	Gen. sp.	Insecta	Coleoptera	Curculionidae			2	2
I	MF1114	<i>Calamosternus granarius</i> (Linnaeus)	Insecta	Coleoptera	Aphodiidae				28
I	MF1117	Gen. sp.	Insecta	Coleoptera	Chrysomelidae				2
I	MF1120	<i>Notiophilus quadripunctatus</i> Dejean	Insecta	Coleoptera	Carabidae				1
I	MF113	<i>Typhaea stercorea</i> (Linnaeus)	Insecta	Coleoptera	Mycetophagidae	1	2	2	10
I	MF120	<i>Coccotrypes carpophagus</i> (Hornung)	Insecta	Coleoptera	Curculionidae	6		1	
I	MF142	<i>Cilea silphoides</i> (Linnaeus)	Insecta	Coleoptera	Staphylinidae		1		1
I	MF145	<i>Cryptophagus</i> sp.	Insecta	Coleoptera	Cryptophagidae	1		5	108
I	MF161	<i>Amara aenea</i> (De Geer)	Insecta	Coleoptera	Carabidae				2
I	MF162	<i>Atheta amicula</i> (Stephens)	Insecta	Coleoptera	Staphylinidae	2			
I	MF247	<i>Aleochara bipustulata</i> (Linnaeus)	Insecta	Coleoptera	Staphylinidae			1	7
I	MF266	<i>Chaetocnema hortensis</i> (Fourcroy)	Insecta	Coleoptera	Chrysomelidae			3	
I	MF268	<i>Carpophilus</i> sp.	Insecta	Coleoptera	Nitidulidae				1
I	MF271	<i>Anotylus</i> sp. 2	Insecta	Coleoptera	Staphylinidae			1	
I	MF275	<i>Agonum muelleri muelleri</i> (Herbst)	Insecta	Coleoptera	Carabidae			3	2
I	MF308	<i>Stelidota geminata</i> (Say)	Insecta	Coleoptera	Nitidulidae	2	824	115	56
I	MF341	<i>Philonthus politus politus</i> (Linnaeus)	Insecta	Coleoptera	Staphylinidae				1
I	MF342	<i>Cercyon</i> sp.	Insecta	Coleoptera	Hydrophilidae		1	1	3
I	MF344	<i>Sitona discoideus</i> Gyllenhal	Insecta	Coleoptera	Curculionidae			1	10
I	MF386	<i>Sitona puberulus</i> Reitter	Insecta	Coleoptera	Curculionidae			1	
I	MF395	<i>Psylliodes marcidus</i> (Illiger)	Insecta	Coleoptera	Chrysomelidae			10	1
I	MF45	<i>Anisodactylus binotatus</i> (Fabricius)	Insecta	Coleoptera	Carabidae			50	2
I	MF502	<i>Omosita colon</i> (Linnaeus)	Insecta	Coleoptera	Nitidulidae				1
I	MF51	<i>Paranchus albipes</i> (Fabricius)	Insecta	Coleoptera	Carabidae	8	258	128	97
I	MF52	<i>Cordalia obscura</i> (Gravenhorst)	Insecta	Coleoptera	Staphylinidae	1	1	6	6
I	MF523	<i>Sphenophorus abbreviatus</i> (Fabricius)	Insecta	Coleoptera	Dryophthoridae				2
I	MF541	<i>Hirticomus quadriguttatus</i> (Rossi)	Insecta	Coleoptera	Anthicidae				407
I	MF57	<i>Atheta atramentaria</i> (Gyllenhal)	Insecta	Coleoptera	Staphylinidae		2	7	
I	MF570	<i>Meligethes aeneus</i> (Fabricius)	Insecta	Coleoptera	Nitidulidae				1
I	MF61	<i>Epuraea biguttata</i> (Thunberg)	Insecta	Coleoptera	Nitidulidae	3	9	30	
I	MF65	<i>Sericoderus lateralis</i> (Gyllenhal)	Insecta	Coleoptera	Corylophidae		1	200	7
I	MF66	<i>Amischa analis</i> (Gravenhorst)	Insecta	Coleoptera	Staphylinidae	6	2		3
I	MF693	Gen. sp.	Insecta	Coleoptera	Curculionidae			1	

**Table 5** continued

N/E/I	Code MF	Species	Class	Order	Family	NF	EF	SNP	IP
I	MF703	<i>Nephus helgae</i> Fursh	Insecta	Coleoptera	Coccinellidae				36
I	MF72	<i>Ptenidium pusillum</i> (Gyllenhal)	Insecta	Coleoptera	Ptiliidae	1	1	3	
I	MF74	<i>Pseudophonus rufipes</i> (DeGeer)	Insecta	Coleoptera	Carabidae	1		2	1
I	MF758	Gen. sp.	Insecta	Coleoptera	Anthicidae		1		11
I	MF764	<i>Harpalus distinguendus</i> (Duftschmidt)	Insecta	Coleoptera	Carabidae				11
I	MF767	<i>Oligota parva</i> Kraatz	Insecta	Coleoptera	Staphylinidae			1	3
I	MF771	<i>Astenus lyonessius</i> (Joy)	Insecta	Coleoptera	Staphylinidae				3
I	MF777	Gen. sp.	Insecta	Coleoptera	Anthicidae				37
I	MF823	<i>Coproporus pulchellus</i> (Erichson)	Insecta	Coleoptera	Staphylinidae				1
I	MF828	<i>Carpophilus hemipterus</i> (Linnaeus)	Insecta	Coleoptera	Nitidulidae	1			1
I	MF885	<i>Oxytelus sculptus</i> Gravenhorst	Insecta	Coleoptera	Staphylinidae			2	3
I	MF888	Gen. sp.	Insecta	Coleoptera	Ptiliidae			2	
I	MF89	<i>Tachyporus chrysomelinus</i> (Linnaeus)	Insecta	Coleoptera	Staphylinidae	1	23	110	25
I	MF944	Gen. sp.	Insecta	Coleoptera	Laemophloeidae				1
I	MF99	<i>Phloeonomus</i> sp.3	Insecta	Coleoptera	Staphylinidae		3	2	
I	MF153	<i>Atheta</i> sp.2	Insecta	Coleoptera	Staphylinidae			3	
I	MF16	<i>Atheta fungi</i> (Gravenhorst)	Insecta	Coleoptera	Staphylinidae	147	21	123	
I	MF354	<i>Orthochaetes insignis</i> (Aubé)	Insecta	Coleoptera	Curculionidae			5	
I	MF673	<i>Mecinus pascuorum</i> (Gyllenhal)	Insecta	Coleoptera	Curculionidae				2
I	MF708	<i>Aridius</i> sp.	Insecta	Coleoptera	Lathridiidae			1	
N	MF102	<i>Pseudophloeophagus tenax</i> (Wollaston)	Insecta	Coleoptera	Curculionidae	3	6	1	
N	MF1086	Gen. sp.	Insecta	Coleoptera	Curculionidae		1		12
N	MF1104	<i>Rhyzobius litura</i> (Fabricius)	Insecta	Coleoptera	Coccinellidae				11
N	MF1115	<i>Psilothrix viridicoerulea</i> (Geoffroy)	Insecta	Coleoptera	Dasytidae				5
N	MF23	<i>Phloeonomus</i> sp.	Insecta	Coleoptera	Staphylinidae		1		
N	MF25	<i>Pseudophloeophagus aenopiceus</i> (Boheman)	Insecta	Coleoptera	Curculionidae		1		
N	MF262	<i>Rugilus orbiculatus orbiculatus</i> (Paykull)	Insecta	Coleoptera	Staphylinidae		1	204	3
N	MF68	<i>Sepedophilus lusitanicus</i> (Hammond)	Insecta	Coleoptera	Staphylinidae	1			
N	MF69	<i>Stilbus testaceus</i> (Panzer)	Insecta	Coleoptera	Phalacridae			4	1
N	MF78	<i>Anaspis proteus</i> (Wollaston)	Insecta	Coleoptera	Scraptiidae			10	
N	MF79	<i>Quedius curtipennis</i> Bernhauer	Insecta	Coleoptera	Staphylinidae		5	261	
N	MF82	<i>Proteinus atomarius</i> Erichson	Insecta	Coleoptera	Staphylinidae	2	4	5	
N	MF88	<i>Ocyopus olens</i> (Muller)	Insecta	Coleoptera	Staphylinidae		3		
N	MF94	<i>Anommatus duodecimstriatus</i> (Muller)	Insecta	Coleoptera	Bothrideridae		1		
N	MF98	<i>Placonotus</i> sp.1	Insecta	Coleoptera	Laemophloeidae		9	2	
N	MF15	<i>Ocys harpaloides</i> (Audinet-Serville)	Insecta	Coleoptera	Carabidae		23		
N	MF691	<i>Otiorynchus cribicollis</i> Gyllenhal	Insecta	Coleoptera	Curculionidae		85	1	2
N	MF360	Gen. sp.	Insecta	Coleoptera	Corylophidae				11
I	MF352	<i>Euborellia annulipes</i> (Lucas)	Insecta	Dermaptera	Anisolabididae	18	89	246	741
I	MF56	<i>Forficula auricularia</i> Linnaeus	Insecta	Dermaptera	Forficulidae	2	7	21	9
E	MF8	<i>Aphrodes hamiltoni</i> Quartau & Borges	Insecta	Hemiptera	Cicadellidae	30			
E	MF925	<i>Cixius azomariae</i> Remane & Asche	Insecta	Hemiptera	Cixiidae	3			
I	MF155	<i>Pseudacaudella rubida</i> (Borner)	Insecta	Hemiptera	Aphididae	2			
I	MF335	<i>Rhopalosiphum insertum</i> (Walker)	Insecta	Hemiptera	Aphididae			1	
I	MF501	<i>Loricula coleoptrata</i> (Fallén)	Insecta	Hemiptera	Microphysidae				1
I	MF521	<i>Brachysteles parvicornis</i> (A. Costa)	Insecta	Hemiptera	Anthocoridae				23
I	MF818	<i>Dysaphis plantaginea</i> (Passerini)	Insecta	Hemiptera	Aphididae			1	1
I	MF676	<i>Microplax plagiata</i> (Fieber)	Insecta	Hemiptera	Lygaeidae				119

**Table 5** continued

N/E/I	Code MF	Species	Class	Order	Family	NF	EF	SNP	IP
I	MF926	<i>Anoscopus albifrons</i> (Linnaeus)	Insecta	Hemiptera	Cicadellidae	2		54	45
N	MF1119	<i>Triatoma rubrofasciata</i> (De Geer)	Insecta	Hemiptera	Reduviidae				1
N	MF124	<i>Cyphopterus adscendens</i> (Herr.-Schaff.)	Insecta	Hemiptera	Flatidae	4			
N	MF232	<i>Acyrtosiphon pisum</i> Harris	Insecta	Hemiptera	Aphididae			11	
N	MF254	<i>Megamelodes quadrimaculatus</i> (Signoret)	Insecta	Hemiptera	Delphacidae			64	
N	MF326	<i>Buchananiella continua</i> (White)	Insecta	Hemiptera	Anthocoridae	1			
N	MF60	<i>Rhopalosiphonimys latysiphon</i> (Davidson)	Insecta	Hemiptera	Aphididae	8	1	20	
N	MF83	<i>Plinthisus brevipennis</i> (Latreille)	Insecta	Hemiptera	Lygaeidae	1	3	34	
N	MF890	<i>Euscelidius variegatus</i> (Kirschbaum)	Insecta	Hemiptera	Cicadellidae				6
N	MF101	<i>Geotomus punctulatus</i> (Costa)	Insecta	Hemiptera	Cydnidae	2			6
N	MF1096	Gen. sp.	Insecta	Hemiptera	Lygaeidae		2		
N	MF118	<i>Scolopostethus decoratus</i> (Hahn)	Insecta	Hemiptera	Lygaeidae	38	1		
N	MF119	Gen. sp.	Insecta	Hemiptera	Coccidae	1			
N	MF132	Gen. sp.	Insecta	Hemiptera	Coccidae	1			
N	MF156	Gen. sp.	Insecta	Hemiptera	Coccidae	8		6	
N	MF54	Gen. sp.	Insecta	Hemiptera	Margarodidae	10			
N	MF58	Gen. sp.	Insecta	Hemiptera	Coccidae	3			
E	MF12	<i>Cyclophora pupillaria granti</i> Prout	Insecta	Lepidoptera	Geometridae	2			
E	MF130	<i>Brachmia infuscatella</i> Rebel	Insecta	Lepidoptera	Gelechiidae	1			
E	MF19	<i>Argyresthia atlanticella</i> Rebel	Insecta	Lepidoptera	Yponomeutidae	3			
E	MF90	<i>Ascotis fortunata azorica</i> Pinker	Insecta	Lepidoptera	Geometridae	1			
I	MF1	Gen. sp.	Insecta	Lepidoptera	Tortricidae		6		
I	MF918	<i>Blastobasis</i> sp.3	Insecta	Lepidoptera	Blastobasidae	1			
N	MF10	<i>Mythimna unipuncta</i> (Haworth)	Insecta	Lepidoptera	Noctuidae	3	2	2	16
N	MF126	<i>Chrysodeixis chalcites</i> (Esper)	Insecta	Lepidoptera	Noctuidae	1			
N	MF220	<i>Noctua pronuba</i> (Linnaeus)	Insecta	Lepidoptera	Noctuidae			1	
N	MF365	Gen. sp.	Insecta	Lepidoptera	Pyalidae	1			
I	MF123	<i>Gryllus bimaculatus</i> (De Geer)	Insecta	Orthoptera	Gryllidae	2			3
I	MF245	<i>Eumodicogryllus bordigalensis</i> (Latreille)	Insecta	Orthoptera	Gryllidae				38
N	MF319	Gen. sp.	Insecta	Orthoptera	Gryllidae				1
E	MF184	<i>Elipsocus azoricus</i> Meinander	Insecta	Psocoptera	Elipsocidae		1		
N	MF121	<i>Ectopsocus briggsi</i> McLachlan	Insecta	Psocoptera	Ectopsocidae	1			
N	MF36	<i>Lachesilla greeni</i> (Pearman)	Insecta	Psocoptera	Lachesillidae	1			
I	MF135	<i>Nesothrips propinquus</i> (Bagnall)	Insecta	Thysanoptera	Phlaeothripidae	1			

In the first column *E* endemic from the Azores, *I* introduced species, *N* native non-endemic. Land-uses are coded as follows: *NF* native forest, *EF* exotic forest, *SNP* semi-natural forest, *IP* intensive pasture

### Appendix 3

See Table 6.

**Table 6** Completeness index (observed number of species divided by the expected number of species) for the four habitat types (four transects per habitat, each transect containing 30 pitfall traps)

	Native forest	Exotic forest	Semi-natural pasture	Intensive pasture
Observed nr of species	81	69	82	95
Expected nr of species	108.8	92.8	103.8	116.8
Completeness Index	0.74	0.74	0.79	0.81

The expected number of species was calculated using a first-order Jack-knife estimator

## References

- Abdel-Monem AA, Fernandez LA, Boone GM (1975) K–Ar ages from the eastern Azores group (Santa Maria, São Miguel and the Formigas Islands). *Lithos* 8:247–254
- Acevedo MA, Restrepo C (2008) Land-cover and land-use change and its contribution to the large-scale organization of Puerto Rico's bird assemblages. *Diversity Distrib* 14:114–122
- Bhagwat SA, Willis KJ, Birks HJB, Whittaker RJ (2008) Agroforestry: a refuge for tropical biodiversity? *Trends Ecol Evol* 23:261–267
- Borges PAV, Brown VK (1999) Effect of island geological age on the arthropod species richness of Azorean pastures. *Biol J Linn Soc* 66:373–410
- Borges PAV, Brown VK (2003) Estimating species richness of arthropods in Azorean pastures: the adequacy of suction sampling and pitfall trapping. *Graellsia* 59:5–22
- Borges PAV, Hortal J (2009) Time, area and isolation: factors driving the diversification of Azorean arthropods. *J Biogeogr* 36:178–191
- Borges PAV, Wunderlich J (2008) Spider biodiversity patterns and their conservation in the Azorean archipelago, with description of new taxa. *Syst Biodivers* 6:249–282
- Borges PAV, Serrano ARM, Quartau JA (2000) Ranking the Azorean Natural Forest Reserves for conservation using their endemic arthropods. *J Insect Conserv* 4:129–147
- Borges PAV, Aguiar C, Amaral J, Amorim IR, André G, Arraiol A, Baz A, Dinis F, Enghoff H, Gaspar C, Ilharco F, Mahnert V, Melo C, Pereira F, Quartau JA, Ribeiro S, Ribes J, Serrano ARM, Sousa AB, Strassen RZ, Vieira L, Vieira V, Vitorino A, Wunderlich J (2005a) Ranking protected areas in the Azores using standardized sampling of soil epigeal arthropods. *Biodivers Conserv* 14:2029–2060
- Borges PAV, Cunha R, Gabriel R, Martins AF, Silva L, Vieira V, Dinis F, Lourenço P, Pinto N (2005b) Description of the terrestrial Azorean biodiversity. In: Borges PAV, Cunha R, Gabriel R, Martins AMF, Silva L, Vieira V (eds) A list of the terrestrial fauna (Mollusca and Arthropoda) and flora (Bryophyta, Pteridophyta and Spermatophyta) from the Azores. Direcção Regional de Ambiente and Universidade dos Açores, Horta, Angra do Heroísmo and Ponta Delgada, pp 21–68
- Borges PAV, Lobo JM, de Azevedo EB, Gaspar C, Melo C, Nunes LV (2006) Invasibility and species richness of island endemic arthropods: a general model of endemic vs. exotic species. *J Biogeogr* 33:169–187
- Borges PAV, Ugland KI, Dinis FO, Gaspar C (2008) Insect and spider rarity in an oceanic island (Terceira, Azores): true rare and pseudo-rare species. In: Fattorini S (ed) *Insect ecology and conservation*. Research Signpost, Kerala, pp 47–70
- Borges PAV, Amorim IR, Cunha R, Gabriel R, Martins AF, Silva L, Costa A, Vieira V (2009) Azores. In: Gillespie R, Clague D (eds) *Encyclopedia of Islands*. University of California Press, California, pp 70–75
- Brook BW, Sodhi NS, Ng PKL (2003) Catastrophic extinctions follow deforestation in Singapore. *Nature* 424:420–423
- Brooks T, Balmford A (1996) Atlantic forest extinctions. *Nature* 380:115
- Brooks TM, Pimm SL, Collar NJ (1997) Deforestation predicts the number of threatened birds in insular Southeast Asia. *Conserv Biol* 11:382–394
- Brooks TM, Tobias J, Balmford A (1999a) Deforestation and bird extinctions in the Atlantic Forest. *Anim Conserv* 2:211–222
- Brooks TM, Pimm SL, Kapos V, Ravilious C (1999b) Threat from deforestation to montane and lowland birds and mammals in insular South-east Asia. *J Anim Ecol* 68:1061–1078
- Brooks TM, Mittermeier RA, Mittermeier CG, da Fonseca GAB, Rylands AB, Konstant WR, Flick P, Pilgrim J, Oldfield S, Magin J, Hilton-Taylor C (2002) Habitat loss and extinction in the hotspots of biodiversity. *Conserv Biol* 16:909–923
- Bruner AG, Gullison RE, Rice RE, da Fonseca GAB (2001) Effectiveness of parks in protecting tropical biodiversity. *Science* 291:125–128
- Burel F, Baudry J, Butet A, Clergeau P, Delettre Y, Le Coeur D, Dubs F, Morvan N, Paillat G, Petit S, Thenail C, Brunel E, Lefeuvre JC (1998) Comparative biodiversity along a gradient of agricultural landscapes. *Acta Oecol* 19:47–60
- Cardoso P, Aranda SC, Lobo JM, Dinis F, Gaspar C, Borges PAV (2009) A spatial scale assessment of habitat effects on arthropod communities of an oceanic island. *Acta Oecol* 35:590–597
- Chape S, Harrison J, Spalding M, Lysenko I (2005) Measuring the extent and effectiveness of protected areas as an indicator for meeting global biodiversity targets. *Philos Trans R Soc Biol Sci* 360:443–455
- Clarke KR (1993) Non-parametric multivariate analyses of changes in community structure. *Aust J Ecol* 18:117–143
- Cowlishaw G (1999) Predicting the pattern of decline of African primate diversity: an extinction debt from historical deforestation. *Conserv Biol* 13:1183–1193
- Daily GC, Ehrlich PR, Sánchez-Azofeifa GA (2001) Countryside biogeography: use of human-dominated habitats by the avifauna of southern Costa Rica. *Ecol Appl* 11:1–13
- Daily GC, Ceballos G, Pacheco J, Suzán G, Sánchez-Azofeifa A (2003) Countryside biogeography of neotropical mammals: conservation opportunities in agricultural landscapes of Costa Rica. *Conserv Biol* 17:1814–1826
- Driscoll DA (2005) Is the matrix a sea? Habitat specificity in a naturally fragmented landscape. *Ecol Entomol* 30:8–16
- DROTRH (2008) Carta de ocupação do solo da região Autónoma dos Açores—projecto SUEMAC. Secretaria Regional do Ambiente. Direcção Regional do Ordenamento do território e dos Recursos Hídricos, Ponta Delgada
- Foley JA, DeFries R, Asner GP, Barford C, Bonan G, Carpenter SR, Chapin FS, Coe MT, Daily GC, Gibbs HK, Helkowski JH, Holloway T, Howard EA, Kucharik CJ, Monfreda C, Patz JA, Prentice IC, Ramankutty N, Snyder PK (2005) Global consequences of land use. *Science* 309:570–574
- Gascon C, Lovejoy TE, Bierregaard RO Jr, Malcolm JR, Stouffer PC, Vasconcelos HL, Laurance WF, Zimmerman B, Tocher M, Borges S (1999) Matrix habitat and species richness in tropical forest remnants. *Biol Cons* 91:223–229
- Gaspar C, Borges PAV, Gaston KJ (2008) Diversity and distribution of arthropods in native forests of the Azores Archipelago. *Arquipélago, Life Marin Sci* 25:1–30
- Grelle CEV, Alves MAS, Bergallo HG, Geise L, Rocha CFD, Van Sluys M, Caramaschi U (2005) Prediction of threatened tetrapods based on the species-area relationship in Atlantic Forest, Brazil. *J Zool Soc Lond* 265:359–364
- Grill A, Knoflach B, Cleary DFR, Kati V (2005) Butterfly, spider, and plant communities in different land-use types in Sardinia, Italy. *Biodivers Conserv* 14:1281–1300
- Hanski I (1998) Metapopulation dynamics. *Nature* 396:41–49
- Hanski I, Koivulehto H, Cameron A, Rahagalala P (2007) Deforestation and apparent extinctions of endemic forest beetles in Madagascar. *Biol Lett* 3:344–347
- Helm A, Hanski I, Pärtel M (2006) Slow response of plant species richness to habitat loss and fragmentation. *Ecol Lett* 9:72–77
- Henderson PA, Seaby RMH (2007) Community Analysis Package 4.0. Pisces Conservation Ltd. Available via [www.pisces-conservation.com](http://www.pisces-conservation.com)

- Heywood VN, Mace GM, May RM, Stuart SN (1994) Uncertainties in extinction rates. *Nature* 368:105
- Hortal J, Borges PAV, Gaspar C (2006) Evaluating the performance of species richness estimators: sensitivity to sample grain size. *J Anim Ecol* 75:274–287
- Hughes JB, Daily GC, Ehrlich PR (2002) Conservation of tropical forest birds in countryside habitats. *Ecol Lett* 5:121–129
- Jana G, Misra KK, Bhattacharya T (2006) Diversity of some insect fauna in industrial and non-industrial areas of West Bengal, India. *J Insect Conserv* 10:249–260
- Janzen DH (1986) The future of tropical ecology. *Ann Rev Ecol Syst* 17:305–324
- Jonsson BG, Kruys N, Ranius T (2005) Ecology of species living on dead wood—lessons for dead wood management. *Silva Fenn* 39:289–309
- Kaila L, Martikainen P, Punttila P (1997) Dead trees left in clear-cuts benefit saproxylic Coleoptera adapted to natural disturbances in boreal forest. *Biodivers Conserv* 6:1–18
- Klein AM, Steffan-Dewenter I, Tschamtk T (2006) Rain forest promotes trophic interactions and diversity of trap-nesting Hymenoptera in adjacent agroforestry. *J Anim Ecol* 75:315–323
- MacArthur RH, Wilson EO (1963) An equilibrium theory of insular zoogeography. *Evolution* 17:373–387
- MacArthur RH, Wilson EO (1967) The theory of island biogeography. Princeton University Press, Princeton
- Margules CR, Pressey RL (2000) Systematic conservation planning. *Nature* 405:243–253
- Martikainen P, Siitonen J, Punttila P, Kaila L, Rauh J (2000) Species richness of Coleoptera in mature managed and old-growth boreal forests in southern Finland. *Biol Conserv* 94:199–209
- Martins AMF (1993) The Azores—Westernmost Europe: where evolution can be caught red-handed. *Bol Mus Mun Funchal* 2:181–198
- Muriel SB, Kattan GH (2009) Effects of patch size and type of coffee matrix on Ithomiine butterfly diversity and dispersal in cloud-forest fragments. *Conserv Biol* 23:948–956
- Myers N (1992) The primary source: tropical forests and our future. W.W. Norton, New York
- Okland B, Bakke A, Hagvar S, Kvamme T (1996) What factors influence the diversity of saproxylic beetles? A multiscaled study from a spruce forest in southern Norway. *Biodivers Conserv* 5:75–100
- Oksanen J, Minchin PR (1997) Instability of ordination results under changes in input data order: explanations and remedies. *J Veg Sci* 8:447–454
- Pereira HM, Cooper HD (2006) Towards the global monitoring of biodiversity change. *Trends Ecol Evol* 21:123–129
- Perfecto I, Vandermeer J (2002) Quality of agroecological matrix in a tropical montane landscape: ants in coffee plantations in southern Mexico. *Conserv Biol* 16:174–182
- Pimm SL, Russell GJ, Gittleman JL, Brooks TM (1995) The future of biodiversity. *Science* 269:347–350
- Prugh LR, Hodges KE, Sinclair ARE, Brashares JS (2008) Effect of habitat area and isolation on fragmented animal populations. *Proc Natl Acad Sci* 105:20770–20775
- Rainio J, Niemelä J (2006) Comparison of carabid beetle (Coleoptera: Carabidae) occurrence in rain forest and human-modified sites in south-eastern Madagascar. *J Insect Conserv* 10:219–228
- Ramankutty N, Foley JA (1999) Estimating historical changes in global land cover: croplands from 1700 to 1992. *Glob Biochem Cycles* 13:997–1027
- Ribeiro SP, Borges PAV, Gaspar C, Melo C, Serrano ARM, Amaral J, Aguiar C, André G, Quartau JA (2005) Canopy insect herbivores in the Azorean Laurisilva forests: key host plant species in a highly generalist insect community. *Ecography* 28:315–330
- Ricketts TH (2001) The matrix matters: effective isolation in fragmented landscapes. *Am Nat* 158:87–99
- Rodrigues ASL, Andelman SJ, Bakarr MI, Boitani L, Brooks TM, Cowling RM, Fishpool LDC, da Fonseca GAB, Gaston KJ, Hoffmann M, Long JS, Marquet PA, Pilgrim JD, Pressey RL, Schipper J, Sechrest W, Stuart SN, Underhill LG, Waller RW, Watts MEJ, Yan X (2004) Effectiveness of the global protected area network in representing species diversity. *Nature* 428:640–643
- Rosenzweig ML (2003) Reconciliation ecology and the future of species diversity. *Oryx* 37:194–205
- Schmidt MH, Thies C, Nentwig W, Tschamtk T (2008) Contrasting responses of arable spiders to the landscape matrix at different spatial scales. *J Biogeogr* 35:157–166
- Schweiger O, Maelfait JP, Van Wingerden W, Hendrickx F, Billeter R, Speelmans M, Augenstein I, Aukema B, Aviron S, Bailey D, Bukacek R, Burel F, Diekötter T, Dirksen J, Frenzel M, Herzog F, Liira J, Roubalova M, Bugter R (2005) Quantifying the impact of environmental factors on arthropod communities in agricultural landscapes across organizational levels and spatial scales. *J Appl Ecol* 42:1129–1139
- Sekercioglu CH, Ehrlich PR, Daily GC, Aygen D, Goehring D, Sandi RF (2002) Disappearance of insectivorous birds from tropical forest fragments. *Proc Natl Acad Sci* 99:263–267
- Tilman D, May RM, Lehman CL, Nowak MA (1994) Habitat destruction and the extinction debt. *Nature* 371:65–66
- Tilman D, Fargione J, Wolff B, D'Antonio C, Dobson A, Howarth R, Schindler D, Schlesinger WH, Simberloff D, Swackhamer D (2001) Forecasting agriculturally driven global environmental change. *Science* 292:281–284
- Toth F, Kiss J (1999) Comparative analyses of epigeic spider assemblages in northern Hungarian winter wheat fields and their adjacent margins. *J Arachnol* 27:241–248
- Triantis KA, Borges PAV, Ladle RJ, Hortal J, Gaspar C, Dinis F, Mendonça E, Silveira LMA, Gabriel R, Cardoso P, Melo C, Santos AMC, Amorim IR, Ribeiro SP, Serrano ARM, Quartau JA, Whittaker RJ (2010) Extinction debt on oceanic islands. *Ecography* 33:285–294
- Vandermeer J, Carvajal R (2001) Metapopulation dynamics and the quality of the matrix. *Am Nat* 158:211–220
- Vitousek PM, Mooney HA, Lubchenco J, Melillo JM (1997) Human domination of Earth's ecosystems. *Science* 277:494–499
- Watson JEM, Whittaker RJ, Freudenberger D (2005) Bird community responses to habitat fragmentation: how consistent are they across landscapes? *J Biogeogr* 32:1353–1370
- Whittaker RJ, Fernández-Palacios JM (2007) Island biogeography: ecology, evolution, and conservation, 2nd edn. Oxford University Press, Oxford
- Wiens J (2007) The dangers of black-and-white conservation. *Conserv Biol* 21:1371–1372
- Wilson EO (1992) The diversity of life. Harvard University Press, Cambridge