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Impact of the Invasive Argentine Ant in Citrus Agroecosystems: Effects on the Diversity and Frequency of Native Ant Species Foraging on Tree Canopy

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Simple Summary: We aimed at assessing the impact of the Argentine ant invasion on the native ant community in citrus ecosystems. We compared the Argentine ant's invaded and uninvaded citrus orchards in the south of Portugal, estimating species richness and the frequency of ant assemblages foraging on the tree canopy. The results suggest that the Argentine ant has a negative impact on the native ant community structure, markedly reducing the diversity and frequency of native species. This impact was more or less pronounced depending on the season. Possible implications for citrus pest management are discussed.

Abstract: The invasion of the Argentine ant, *Linepithema humile* (Mayr) (Hymenoptera, Formicidae) can alter the entire ecosystem with serious impacts on the native community structure (e.g., ant diversity) and processes (e.g., trophic interactions) leading to biodiversity loss and pest outbreaks. Most studies addressing these impacts have been conducted in natural or semi-natural areas, few are those conducted in agricultural ecosystems, such as citrus orchards. These are dominant agricultural ecosystems in Mediterranean landscapes. Furthermore, most studies have been conducted in a short span, not evidencing seasonal fluctuations. In this work, we assessed the ecological impact of the Argentine ant on the native ant communities in citrus orchards, in the region of Algarve, southern Portugal. By using principal response curve, we compared seasonal variation on ant assemblages in invaded and uninvaded citrus orchards foraging on tree canopy from a two-year sampling. The Argentine ant had a marked negative impact on the native ant community foraging on citrus canopy. In the uninvaded orchards, the native ant community had a rich assemblage composed of 16 ant species, in its majority (72%) controlled by the dominant species Lasius grandis Forel, Tapinoma nigerrimum (Nylander) and/or Pheidole pallidula (Nylander). In the invaded orchards, the native ant community was poorer and highly modified, mostly dominated by the Argentine ant (80%). Apparently, the only native ant species not affected by the presence of the Argentine ant was *Plagiolepis pygmaea* (Latreille). A significant negative effect was found between the proportion of infested trees by L. humile and the number of native ant species per orchard. Differences in the native ant community in the invaded and uninvaded orchards persisted over seasons and years. However, negative impacts were higher in the spring and summer, and less pronounced in the autumn. We discuss implications for citrus pest management.

Keywords: ants; Formicidae; invasive species; *Lasius grandis*; *Linepithema humile*; principal response curve; seasonal changes

1. Introduction

Ant invasions represent a worldwide concern and have been the subject of the largest number of publications on invasive insect studies in recent years [1]. Most of the introduced ant species are usually not detected and usually do not constitute a threat to native fauna. However, those species that become invasive are very successful and considered a serious threat to the world's native biodiversity [2,3]. The causes underlying the ecological success of invasive ants have been documented [4]. Apparently, it seems that a combination of characteristics such as omnivory, unicoloniality, absence of competitors and natural enemies makes them successful invaders [4]. The Argentine ant, *Linepithema humile* Mayr, appears to have these abilities [4]. Native to South America, this species easily spread and become established into new areas around the world [5]. It has been recognized that the Argentine ant can displace the native ant community leading to dramatic impacts on the ecosystems [4,6–10].

Understanding the ecological impacts of invaders and the mechanisms underlying invasion dynamics is of extreme importance to anticipate or mitigate their negative environmental impacts [11]. However, there are difficulties in performing experimental studies on the impact of invasive ants since researchers cannot ethically introduce invaders to uninvaded areas [12]. As such, the impacts of invaders have been quantified by comparing selected metrics between invaded and uninvaded areas [13–16]. Such comparisons can yield important insights into a wide variety of impacts associated with ant invasion [4].

To date, most studies on the impact of invasive ants have been conducted in natural or semi-natural areas [4,6,17]. Only a few were conducted in agricultural ecosystems, such as citrus orchards [18].

Citrus is an economically important irrigated crop in the Mediterranean region, representing 20% of the world's citrus production [19]. Ants are common insects foraging on citrus canopy [20,21], mainly due to the high diversity of honeydew-producing hemipteran in this crop, including aphids (Hemiptera, Aphidoidea), whiteflies (Hemiptera, Aleyrodoidea), and scale insects (Hemiptera, Coccoidea) [22–29]. The Argentine ant reaches high densities in Mediterranean citrus orchards, where it is considered a common pest, also facilitating the activity of some sap-sucking insect species [30].

The decreasing diversity and abundance of native ants resulting from ant invasions can give rise to a variety of direct and indirect effects on non-ant taxa [4]. This is of extremely high importance for integrated pest management (IPM), as the composition of ant communities may influence the pest status of insect species present in agricultural ecosystems. For example, the most recent update to the USA Road Map for IPM places a special emphasis on invasive species [31].

In a previous study on ant communities associated with citrus, in the region of Algarve (south of Portugal), the Argentine ant was detected in 33% of the sampled orchards within two subregions, and was completely absent from another subregion—Serra [21]. Since Algarve was the first Mediterranean area invaded by the Argentine ant, over 120 years ago [32,33] it is likely there was no limitation of time for its establishment in the region. Based on this assumption, Zina et al. [21] hypothesized that, among other factors, the absence of the Argentine ant in the Serra could be explained as the result of interspecific competition with dominant native ant species, such as *Lasius grandis* Forel, *Pheidole pallidula* (Nylander) and *Tapinoma nigerrimum* (Nylander).

In the present study, we aimed at assessing the impact of the Argentine ant invasion on the native ant community foraging on the tree canopy in citrus ecosystems. As an experimental approach, we compared the species richness and frequency of ant assemblages foraging on the tree canopy between invaded (treatment) and uninvaded (control) citrus orchards in the region of Algarve. It was assumed that differences between treatments on the structure of ant communities is a result of the

interaction between native ants and the Argentine ant. We further hypothesized that those interspecific interactions changed over the year. Autumn and winter must be a particular moment in which the Argentine ant is most vulnerable, as this species is known to have seasonal polydomy, a type of intrinsic nest relocation pattern in which colonies converge during the winter and spread among multiple nests in warmer periods [34–36]. Overall, we predict finding a higher number of ant species in uninvaded than invaded orchards, as well as a seasonal effect on this difference in species richness. This study was thus conducted during two years, with seasonal sampling, in order to identify the possible existence of seasonal patterns in ant species diversity and frequency.

Experimental studies and research focusing on the invasive ecology of ants and impacts on the native ant community are especially valuable contributions to agrobiodiversity and citrus pest management. Thus, we discuss the possible implications of the results for citrus pest management.

2. Materials and Methods

2.1. Study Area and Experimental Design

This study was conducted in Algarve, the main citrus producing region of Portugal. With a Mediterranean climate, the average annual temperature in the region is 15.5-17.5 °C, decreasing to 13 °C at 900 m altitudes. The maximal temperature in the summer can reach 30-35 °C, occasionally 40 °C [37].

Based on a previous study [21], we selected 24 commercial citrus orchards, distributed between Silves (37.177 N, 8.448 W) and Tavira (37.154 N, 7.652 W) (Figure 1), 12 of which were invaded by the Argentine ant. The orchards included 71% of sweet oranges, *Citrus sinensis* (L.) (cv. Valencia Late, Newhall, Navelina, Rhode, Navelate Lanelate, Salustiana), 13% lemons, *C. limon* Osbeck (cv. Eureka), 8% mandarins, *C. reticulata* (Blanco) (cv. Ortanique, Encore), and 8% mixed plots, with sweet oranges and mandarins. The average area per citrus plot was identical (p = 0.57) between invaded (1.46 ha \pm 1.08 SE) and uninvaded (1.73 ha \pm 1.25 SE) orchards. The nearest neighbour index among orchards was 0.88 (Z-Score = -1.14). Eventually, insecticide treatments under an IPM regime were applied to control key pests, such as the California red scale, *Aonidiella aurantii* (Maskell) and the Mediterranean fruit fly, *Ceratitis capitata* (Wiedemann). We assumed no differential effects of insecticide treatments between invaded and uninvaded orchards. Experimental evidence supports this assumption, as ant community structure has been shown to be insensitive to pesticide gradients (e.g., [38]).

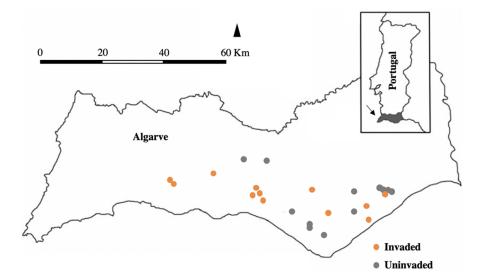


Figure 1. Map of the Algarve region with the invasion status of sampled orchards. The orange dots represent the invaded orchards by the Argentine ant, while grey dots represent uninvaded orchards.

Insects **2020**, 11, 785 4 of 17

2.2. Ant Sampling and Identification

Seasonal samplings (summer, autumn, spring) were carried out during two years, between July 2012 and May 2014, in the same 20 trees per orchard. Trees were first selected at haphazardly along 4–5 line transects, each tree separated from the other by at least 15 m to avoid spatial correlation. Line transects covered the central part of the studied citrus plots, as a grid, in order to consider within orchard variability. Samplings were carried out between 17 July and 24 August, 17 October and 15 November, in 2012; between 25 March and 26 May, 18 July and 23 August, 15 November and 30 November, in 2013; and between 19 May and 28 May, in 2014. For each sampling period, the number of sampled plots in each date was similar between invaded and uninvaded orchards. The method used was direct search by hand collection, considered the most efficient technique to estimate the diversity of ant species [39,40]. This method has proved effective in other ant studies conducted in Mediterranean citrus groves [20,21,41]. It records the presence of species inhabiting a habitat element and allows listing the ant fauna in relatively short time by experienced collectors [42]. The presence of ants in the tree canopy was determined by visual observation of the trunk, shoots, leaves, fruits and flowers, along the tree canopy perimeter up to 1.70 m height, during 60 s per tree, following Zina et al. [21]. This time is sufficient to determine the presence of ants in a tree by ant experts. Whenever ants were detected, a sample of specimens was collected and preserved in 96% alcohol, within Eppendorf tubes, for species identification. The samples were stored in the Entomology Laboratory of the School of Agriculture, University of Lisbon. The collected ant specimens were observed under magnification (640×) and identified at the species level based on Collingwood and Prince [43] and Gómez and Espadaler [44].

2.3. Data Analysis

We built a generalized linear mixed model (GLMM) with the Poisson distribution using a log link function to explain observed variation in ant diversity. We used the orchards as subjects with repeated measures, the year was considered a random variable, and the season and the treatment as fixed factors, with three and two levels, respectively. Pairwise comparisons of estimated marginal means were used to estimate significant differences in ant species richness between invaded and uninvaded orchards, and among seasons. The GLMM was fitted using SPSS [45].

Pearson's correlation tests were performed by season and overall to assess the relationship between the proportion of infested trees per orchard and the number of native ant species in those orchards.

Multivariate statistical analysis was used to describe the effects of Argentine ant's invasion at the community level. The principal response curve (PRC) method was applied to study the effect of the invasion by the Argentine ant on the native ant community, by comparing the native ant species over the sampling seasons in invaded and uninvaded orchards, using packages Vegan for R version 3.3.1 software for Windows [46], with data on occurrences per orchard. Monte Carlo permutation tests [47] were performed to test the significance of the first axis and the significance of the PRC deviations for each sampling season. PRC is based on redundancy analysis (RDA), adjusted for overall changes in community response over time, in relation to the control [48]. This is an interpretive method allowing a quantitative interpretation of the effects headed for the species level enabled by scoring the species weight, accounting for deviances [47].

3. Results

3.1. Ant Communities

In total, 10,930 individuals comprising 18 ant species, 10 genera, and three subfamilies were collected in the sampled orchards (Table 1). Of these, 55% were collected in the Argentine ant's invaded orchards, and 45% in the uninvaded orchards (Table 1). Overall, ants were present in all sampled orchards and in 81% of the sampled trees (89% and 72%, in invaded and uninvaded orchards, respectively). Sixteen ant species were identified in uninvaded orchards, while only 10 species were observed in the orchards invaded by the Argentine ant. In the first case, five ant species, i.e., *L. grandis*

Insects 2020, 11, 785 5 of 17

(41% of the specimens), *Plagiolepis pygmaea* (Latreille) (22%), *T. nigerrimum* (12%), *P. pallidula* (10%), and *P. schmitzii* Forel (7%) represented 92% of the collected specimens, whereas in the case of invaded orchards, 99% of the specimens belonged to *L. humile* (80%) and *P. pygmaea* (19%) (Figure 2).

Table 1. Ant species collected in Argentine ant invaded and uninvaded citrus orchards of Algarve, Portugal.

Subfamily	Nı	umber of	Specime	ens	Number of Orchards in Wh Each Ant Species Was Fou				
		vaded		aded		vaded	Invaded		
Ant Species	Orchards (N = 12)			nards = 12)		hards = 12)	Orchards $(N = 12)$		
	N	%	N	%	N	%	N	%	
Dolichoderinae									
Linepithema humile	0	0	5153	86.37	0	0	12	100	
Tapinoma madeirense	22	0.44	0	0	2	16.67	0	0	
Tapinoma nigerrimum	835	16.82	2	0.03	8	66.67	1	8.33	
Tapinoma simrothi	32	0.64	7	0.12	6	50.00	1	8.33	
Formicinae									
Camponotus gestroi	0	0	1	0.02	0	0	1	8.33	
Camponotus lateralis	6	0.12	0	0	1	8.33	0	0	
Camponotus micans	25	0.50	0	0	1	8.33	0	0	
Camponotus sylvaticus	12	0.24	3	0.05	4	33.33	1	8.33	
Formica cunicularia	24	0.48	0	0	2	16.67	0	0	
Lasius grandis	2099	42.28	0	0	10	83.33	0	0	
Plagiolepis pygmaea	1159	23.35	783	13.12	10	83.33	12	100	
Plagiolepis schmitzii	224	4.51	3	0.05	7	58.33	2	16.67	
Myrmicinae									
Aphaenogaster senilis	4	0.08	0	0	2	16.67	0	0	
Crematogaster auberti	41	0.83	4	0.07	2	16.67	1	8.33	
Crematogaster scutellaris	33	0.66	1	0.02	4	33.33	1	8.33	
Crematogaster sordidula	56	1.13	9	0.15	3	25.00	1	8.33	
Messor barbarus	4	0.08	0	0	3	25.00	0	0	
Pheidole pallidula	388	7.82	0	0	8	66.67	0	0	
Total Number of Specimens (N)	4964		5966						
Species Richness (S)	16		10						

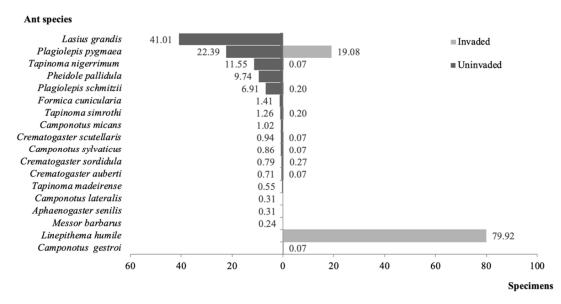


Figure 2. Frequency distribution (%) of ant species in Argentine ant invaded and uninvaded orchards.

Insects 2020, 11, 785 6 of 17

A significant negative correlation (at the 0.05 level) was found between the magnitude of invasion by L. humile (i.e., the proportion of infested trees) and the number of native ant species per orchard (Pearson Correlation (PC) = -0.42, p < 0.001; Appendix A). Individual correlations by season showed the same trend for summer (PC = -0.47, p = 0.019; Appendix A) and spring (PC = -0.46, p = 0.026; Appendix A), but not for autumn (PC = -0.17, p = 0.425; Appendix A). When L. humile was present, native ants were found only in very low levels (up to nine individuals). Most species were found together with L. humile in just one orchard (Crematogaster scutellaris (Olivier), C. sordidula (Nylander), C. auberti Emery, Camponotus sylvaticus (Olivier), C. gestroi Emery, T. nigerrimum, and T. simrothi Krausse). Plagiolepis schmitzii co-occurred with the Argentine ant in two orchards. An exception was found for P. pygmaea, which was observed in all invaded orchards (Figure 2, Table 1). Eight ant species, i.e., Aphaenogaster senilis Mayr, C. lateralis (Olivier), C. micans (Nylander), Formica cunicularia Latreille, L. grandis, Messor barbarus (Linnaeus), P. pallidula, and T. madeirense Forel appeared only in uninvaded orchards, whereas C. gestroi was only observed in an invaded orchard (Figure 2, Table 1).

Overall, the results on the GLMM showed that ant species richness is significantly affected by treatment ($F_{(1,138)} = 17.90$, p < 0.001) (Appendix B). The average number of ant species per orchard in uninvaded orchards (2.98 ± 0.23 ; maximum = 7) was significantly (p < 0.001) higher than in invaded ones (1.77 ± 0.17 ; maximum = 4) (Figure 3A). We present the analyses of all pairwise comparisons in Appendix C.

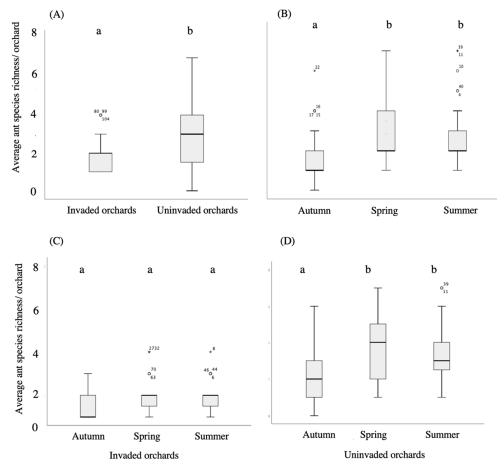


Figure 3. Box plots representing the average ant species' richness per orchard in function of the invasion of the Argentine ant (**A**) and the season (**B**) in invaded (**C**) and uninvaded (**D**) citrus orchards. Boxes show interquartile ranges (25th and 75th percentiles), middle lines are medians, whiskers are non-outlier ranges beyond the boxes, circles are the outliers and asterisks are the extreme outliers. Different letters show significant differences between invasion treatment groups and among seasons by the fitted generalized linear mixed model (GLMM).

Insects 2020, 11, 785 7 of 17

3.2. Seasonal Patterns

Ants were observed in all seasons and orchards, except for autumn 2012 and 2013, in which ants were observed in about 92% (11 out of 12) and 83% (10 out of 12) of the invaded and uninvaded orchards, respectively. Overall, the results on the GLMM showed that ant species richness is significantly affected by the season ($F_{(2,138)} = 5.50$, p = 0.005) (Appendix B). The average number of ant species per orchard was significantly lower in autumn (1.73 ± 0.17) than in spring (2.69 ± 0.25; p = 0.002) and summer (2.60 ± 0.25; p = 0.005) (Figure 3B). No significant differences were found between spring and summer (p = 0.80), both in invaded (Figure 3C) and uninvaded orchards (Figure 3D). We present the analyses of all pairwise comparisons in Appendix C.

Overall, ant frequency showed a seasonal pattern in both invaded and uninvaded orchards, with a maximum in spring/summer and a minimum in autumn (Figure 4). Still, the variation among seasons was more evident in non-invaded orchards, compared to those in which *L. humile* was present. The seasonal pattern also registered some variation in function of ant species. The level of variation among seasons was apparently lower for the most dominant species in each treatment, i.e., *L. grandis* and *L. humile* in uninvaded and invaded orchards, respectively (Figure 4). In addition, the seasonal peaks of activity varied among ant species. *Lasius grandis*, *T. nigerrimum* and *P. schmitzii* were more frequent in the spring, while the frequency of *L. humile*, *P. pygmaea* and *P. pallidula* was highest in the summer and that of *T. simrothi* in autumn. *Plagiolepis pygmaea* exhibited the same seasonal pattern and similar levels of activity in both invaded and uninvaded orchards.

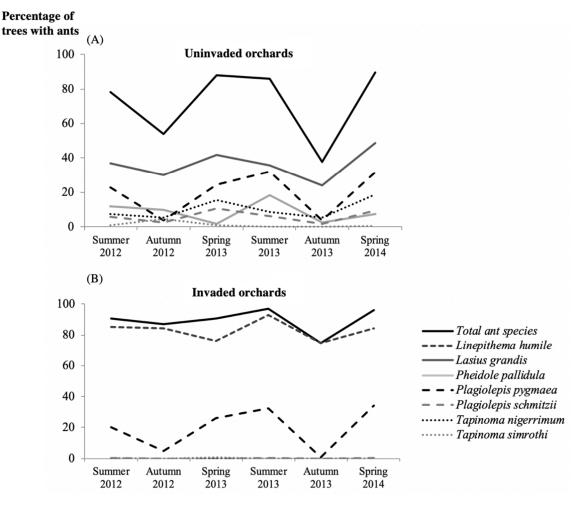


Figure 4. Seasonal variation in the percentage of trees with the most frequent ant species by season in the uninvaded (**A**) and invaded orchards (**B**). In each season, 240 trees were observed by modality.

PRC analysis revealed that the occurrence of native ants in uninvaded orchards was significantly higher (p = 0.001) than in the orchards in which the Argentine ant was present, in all sampling dates, during the two years' study (Figure 5). Still, differences were higher in spring and summer than in autumn. *Lasius grandis* was the major contributor to the differences between invaded and uninvaded orchards, followed by *T. nigerrimum*, *P. pallidula*, and *P. schmitzii*.

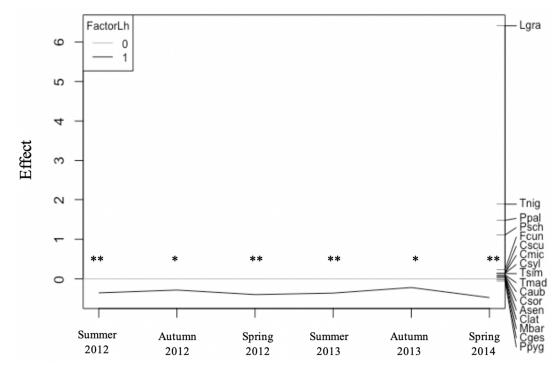


Figure 5. Principal response curves (PRC) representing the effects of the invasion by the Argentine ant on native ants foraging on citrus tree canopy. The left y axis represents deviances from the control (uninvaded orchards). Significant deviances based on Monte Carlo permutation tests are marked with one (p < 0.05) or two asterisks (p < 0.001). The right side of the figure represents ant species weight, accounting for the deviances of the PRC. The first axis explains 96% of the variance of species—environment. Legend: 0—native ant community (gray reference line); 1—ant community affected by the Argentine ant invasion (black line); Lgra—Lasius grandis, Tnig—Tapinoma nigerrimum, Ppal—Pheidole pallidula, Psch—Plagiolepis schmitzii, Fcun—Formica cunicularia, Cscu—Crematogaster scutellaris, Cmic—Camponotus micans, Csyl—Camponotus sylvaticus, Tsim—Tapinoma simrothi, Tmad—Tapinoma madeirense, Caub—Crematogaster auberti, Csor—Crematogaster sordidula, Asen—Aphaenogaster senilis, Clat—Camponotus lateralis, Mbar—Messor barbarus, Cges—Camponotus gestroi, Ppyg—Plagiolepis pygmaea.

4. Discussion

The evidence that invasive ants often become highly abundant in their introduced range and can outnumber native ants [4] is consistent with our results. We showed that the Argentine ant has a negative impact on the native ant community structure and that this effect is more or less pronounced depending on the season period. In addition, community processes are also likely to be affected by the Argentine ant invasion. The direct impacts on native ants modify networks and indirectly affect a variety of regulating and supporting services, disrupting ecosystem processes, such as trophic-based interactions, often leading to pest outbreaks [49]. We will thus focus our discussion on Argentine ants' impacts on community structure (e.g., ant diversity and frequency), seasonal dynamics, and possible implications for citrus pest management.

Insects 2020, 11, 785 9 of 17

4.1. Ant Community Structure

Ant communities of Mediterranean citrus orchards have been studied by several authors [18,20,21,41,50–54]. However, almost no studies addressed the impact of invasive species on the native ant community in this agroecosystem. Our results evidenced a negative impact of the Argentine ant on the native ant assemblages foraging on citrus canopy. Similar negative impacts have been reported in natural and forest ecosystems [6,8,17,55].

In invaded areas, the abundance of native ants can be reduced by over 90% ([4] and references therein) In our study, the orchards invaded by the Argentine ant showed a 44% and 76% reduction in the number and frequency of native ant species, respectively, compared to uninvaded orchards. A 60% decrease in native ant's biodiversity was also reported by Menke et al. [17], in California riparian woodlands.

Overall, the uninvaded orchards showed a far more complex and richer ant community, composed of 16 native ant species, mainly dominated by *L. grandis*, *T. nigerrimum* and *P. pallidula* (72%). On the other hand, in invaded orchards, in which the ant community was dominated by the Argentine ant (80%), the native ant community was poorer and highly modified, limited to few species. As a result, mean ant species richness per orchard was lower in invaded orchards, compared to uninvaded ones. This pattern supports the 'dominance-impoverishment rule', according to which ant communities dominated by behaviourally dominant species are associated with low ant species richness [7,56]. Recently, Arnan et al. [56] suggested that this rule only applies to invaded communities, and not to native ones.

Although invasive ants displace many native ant species, some are often able to persist ([4] and references therein, [55]). This is apparently the case of *P. pygmaea*, the only native ant species not affected by the presence of the Argentine ant. In our study, this species occurred in about 92% of the sampled orchards, with a similar frequency and seasonal pattern, in both invaded and uninvaded orchards. Similar observations were reported in other studies for *P. pygmaea* [57,58], and another species of the same genus, *P. schmitzii* [20]. *Plagiolepis pygmaea* is a common and not aggressive small species, which is tolerated by other ants, due to its submissive behaviour, allowing the coexistence with highly dominant species, such as *L. humile* [21,43,57]. In fact, species that coexist with *L. humile* are apparently small-sized species that can go unnoticed [6,55].

Invasive ants also compete with native ants indirectly via exploitative competition [4]. The native dominant species L. grandis and P. pallidula were only observed in the uninvaded orchards, and T. nigerrimum was only detected episodically in one of the invaded orchards. These results may support the hypothesis that the absence of the Argentine ant from citrus orchards in some areas of Algarve is, at least in part, related to interspecific competition with dominant native ant species. They may resist the invasion by the Argentine ant and limit its dispersion within the region. Competitive mechanisms underlying the displacement of native species by invasive ants may involve colony-level battles including the use of physical aggression by workers and nest raiding [4,59,60]. Colonies of L. humile, L. grandis and T. nigerrimum engage in aggressive and deadly battles (V. Zina, unpublished observation). Based on laboratory experiments, both L. grandis and T. nigerrimum showed to be strong competitors of L. humile [61] (V. Zina, unpublished observation). Other studies suggested that these dominant native species are able to resist and prevent the spread of the Argentine ant when its population is at low densities and/or the abiotic conditions are unfavourable to the invasion ([62] and references therein). Nevertheless, the outcome of the interaction between the Argentine and native dominant ants may be influenced by other factors, such as favourable habitat conditions and food resources availability [63,64]. Additional experimental data are needed, such as to test competition exclusion hypothesis, to confirm the hypothesis of biotic resistance.

It is known that the Argentine ant is able to secure the majority of food resources in areas where it meets native ants [60]. Additionally, the access to a carbohydrate-rich diet, such as hemipteran honeydew may allow invasive ants to feed workers at a high rate, making possible the maintenance

of high dynamic densities, the defence of absolute territories, and the further monopolization of resources [65].

Our work has been carefully designed on the basis of an extensive, replicated approach to produce reliable ecological information associated with ant invasions of citrus ecosystems. Nevertheless, as the sampling was focused on ant species foraging on tree canopy, we may have underestimated possible interactions with other ant species, such as hypogaetic ants (e.g., *Solenopsis* spp., *Hypoponera* spp.). However, it is known that the Argentine ant has a small impact on those species [6,17,66]. Moreover, we were particularly interested in understanding the possible interactions with ant species that may establish trophic relationships with citrus insect pests, which are expected to be limited to citrus canopy.

4.2. Seasonal Patterns

The impact of the Argentine ant on the native ant community in citrus orchards was shown to be seasonally dependent, with stronger differences between invaded and uninvaded orchards, and in spring and summer, in comparison with autumn. A seasonal effect was also reported by other authors [35,67]. However, a different pattern to that registered by us was reported by Heller et al. [35]. These authors observed that the impact of the Argentine ant on native ants in a Northern California reserve was greater in the autumn than in the spring, and that invasive and native ants overlapped more often in the spring. Apparent seasonal differences in the impact of the Argentine ant may be related with differences in the seasonal dynamics of food resources, such as hemipteran honeydew, which is habitat dependent. For example, a high diversity of hemipteran species excreting honeydew is commonly associated with citrus crops, including aphids, whiteflies and scale insects, and the population density of these insect pests is maximal in spring and summer [20]. Therefore, it is expected that the abundance of a carbohydrate-rich diet for the Argentine ant will be higher in those seasons, in the case of citrus, but not necessarily in other habitats, with a low diversity and abundance of hemipteran species.

Both the population dynamics of invasive and native species are expected to vary over time [68]. We found that season had a significant influence on the distributions of all the ants present in the orchards. However, this seasonal pattern registered some variation in their frequency depending on the ant species. For example, the dominant native species *L. grandis* established itself early in the year, reaching their peaks in the spring. In contrast, the Argentine ant showed the highest frequency in the summer. These different seasonal patterns are possibly related to the seasonal abundance of their preferred honeydew sources present in citrus canopy. In fact, *L. grandis* is known as an aphid/whitefly-tender species (e.g., *Aphis spiraecola* Patch, *Aleurothrixus floccosus* (Maskell)) [20,21,69], that are more abundant in springtime in parallel with citrus flushing, whereas the Argentine ant is more commonly associated with mealybugs and coccids (e.g., *Planococcus citri* (Risso), *Coccus hesperidum* Linnaeus) [21], with a population build-up occurring in the summer.

Invasion impacts can increase or decrease in magnitude over time [17]. The Argentine ant exhibits differences in population density along the year. The abundance of the Argentine ant in summer is related to the increase in the spatial extent of the colony, with many small dispersed nests, expanding the foraging range, while in autumn the ants return to the old winter nests and the spatial extent of the colony contracts [35,70]. Then, they move from nest aggregation in the winter (colony contraction), to nest dispersion in summer (colony expansion). It has been suggested that in the summer, when the Argentine ant is more dispersed and active, encounters with native ants will be less frequent [35]. However, with their peak abundance in the summer, we believe that major ecological impacts are expected to occur. In our work, Argentine ants' invasion varied in magnitude among orchards and consequently in the impact on native ant species. A negative correlation was significant in the summer and spring but not in the autumn. Menke et al. [17] suggested that variation in invader's abundance may be due to site-specific factors. In citrus orchards, it is likely to depend on the nutritional needs and resources abundance, yet a consequence of the season [64].

4.3. Implications for Citrus Pest Management

Some ant species are considered citrus pests [30], because they may originate direct damage on citrus plants, by feeding on leaves, shoots and buds, such as *T. nigerrimum* [71]. More frequently, ants' pest status on citrus is due to their mutualistic relationship with honeydew-producing hemipterans and consequent negative impact on the natural enemies of these sap-sucking insect pests. The disruption of biological control of hemipteran insect pests by ants has been reported by different authors [20,29,72,73]. The impact of ants in the levels of parasitism and predation of citrus pests is apparently dependent on ant species and population density [54,74,75].

The changes in native fauna driven by invasive species can have cascading consequences on the ecosystems services they provide [49]. Invasive and native ant species may preferentially explore different sources of honeydew. For example, Zina et al. [21] observed a positive correlation between Argentine ant and mealybugs and other scales insects (e.g., C. hesperidum), as well as between L. grandis and aphids and whiteflies, and between T. nigerrimum and I. purchasi. Different associations were reported by other authors, such as *P. pallidula* and *L. grandis* with mealybugs [20]. Such interactions among different honeydew-producing hemipterans may be of general importance for the Argentine ant's successful invasion in citrus orchards. There is a need to better understand the preferential associations between ants and honeydew-producers on citrus. Considering that ants may disrupt the biological control of citrus pests and that they may differ in their preferences for foraging in different honeydew sources (e.g., whiteflies, mealybugs, aphids) in the citrus canopy, we may expect that the different composition of ant assemblages in a citrus orchard will represent a different risk of insect pest outbreaks. For example, we would expect that the presence of the Argentine ant will favour the population build up and will increase the risk of outbreaks of the citrus mealybug, P. citri, a major pest of citrus crops in the Mediterranean area [30]. Besides differences in honeydew preference, different aggressive behaviour and potential for disrupting biological control, as well as different ability to build up large populations among ant species may also influence pest outbreak risks. Furthermore, ants have been recently reported as possible vectors of citrus diseases [76], and this role may also differ among ant species. Therefore, knowledge on the structure and composition of ant communities in citrus orchards, and the influence of invasive alien species, such as the Argentine ant, on native ant species, is of relevance for decision making in citrus pest management. Novel approaches of effective and selective ant control (e.g., prey-baiting [77]) may contribute to improve the biological control of citrus pests.

5. Conclusions

The results obtained in the present study provide strong evidence that the structure and composition of the native ant community foraging on the citrus canopy are affected by the presence of the Argentine ant, with a significant reduction in the diversity and frequency of native ant species in invaded orchards. The level of impact of the Argentine ant was season dependent, with a higher impact registered in spring and summer, in comparison with autumn, supporting our hypothesis that the dominance of the Argentine ant is expected to be higher in warmer periods of the year.

Our data also support the hypothesis that the interspecific competition with dominant native ant species, such as *L. grandis*, *P. pallidula* and *T. nigerrimum* may prevent the invasion by the Argentine ant in certain areas. However, the interaction between the Argentine and dominant ant species may be influenced by other factors, such as habitat conditions and food resources.

Future studies should aim at uncovering the mechanisms underlying the impacts of Argentine ant invasion in citrus orchards, namely through competitive interactions with dominant native species in the initial stages of invasion, and its associations with honeydew-producing hemipterans.

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Appendix A

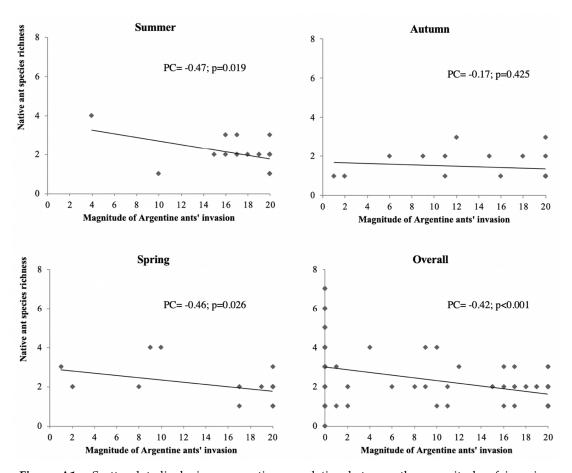


Figure A1. Scatterplot displaying a negative correlation between the magnitude of invasion, i.e., the percentage of infested trees by the Argentine ant, and the number of native ant species per orchard at summer, autumn, spring and overall. Pearson's Correlation (PC) and *p*-value are shown.

Appendix B

	Fixed E	ffects ^a		
Source	F	df1	df2	Sig.
Corrected Model	6.752	5	138	.000
Treatment	17.894	1	138	.000
Season	5.497	2	138	.005
Treatment * Season	.243	2	138	.785

Probability distribution: Poisson Link function: Log^a

a. Target: Ant richness

			Fi	xed Coe	fficients ^a				
					95% Confide	nce Interval	Exp	95% Confidence Exp(Coef	
Model Term	Coefficient	Std. Error	t	Sig.	Lower	Upper	(Coefficient)	Lower	Upper
Intercept	1.237	.1151	10.752	.000	1.010	1.465	3.445	2.744	4.326
Treatment=Invaded	566	.1889	-2.995	.003	939	192	.568	.391	.825
Treatment=Uninvaded	0 ^b								
Season=Autumn	487	.1779	-2.737	.007	839	135	.614	.432	.874
Season=Spring	.047	.1534	.307	.759	256	.350	1.048	.774	1.420
Season=Summer	0 ^b								
[Treatment=Invaded]* [Season=Autumn]	.163	.2870	.569	.570	404	.731	1.177	.668	2.076
[Treatment=Invaded]* [Season=Spring]	026	.2562	102	.919	533	.481	.974	.587	1.617
[Treatment=Invaded]* [Season=Summer]	0 ^b								
[Treatment=Uninvaded] *[Season=Autumn]	0 ^b								
[Treatment=Uninvaded] *[Season=Spring]	0 _p								
[Treatment=Uninvaded] *[Season=Summer]	0 ^b								

Probability distribution: Poisson Link function: Log^a

Figure A2. Results for the generalized linear mixed model (GLMM) of ant richness fitted by the Poisson distribution, with treatment, season and their interaction as fixed factors, the orchards as subjects with repeated measures and the year as a random variable.

a. Target: Ant richness

b. This coefficient is set to zero because it is redundant.

Appendix C

									Pairwis	e Contras	ts				
		Estimat	es			Treatment F	Palmuleo	Contrast					95% Confid	dence Interval	
			95% Conf	idence Interva	ıl	Contrasts	allwise	Estimate	Std. Error	t	df	Adj. Sig.	Lower	Upper	
Treatment	Mean	Std. Error	Lower	Upper		Invaded - U	ninvaded	-1.207	.286	-4.225	138	4.312E-5	-1.771	642	
Invaded	1.769	.169	1.46	2.13	7	Uninvaded -		1.207	.286	4.225	138	4.312E-5	.642	1.771	
Uninvaded	2.975	.230	2.55	3.46	7	The least sig	gnificant differe	ence adjusted si	gnificance lev	el is .05.					
Estimate	ed Mear	ıs 2: sea	ison						Pairwise	Contrast	s				
		Estimates				Season Pairv Contrasts	vise	Contrast Estimate	Std. Error	t	df	Adj. Sig.	95% Confide Lower	nce Interval Upper	
			95% Confide			Autumn – Sp	ring	955	.309	-3.090	138	.002	-1.567	344	
Season		Std. Error	Lower	Upper		Autumn – Su	ımmer	865	.306	-2.833	138	.005	-1.469	261	
Autumn	1.731	.196	1.383	2.166		Spring - Aut	umn	.955	.309	3.090	138	.002	.344	1.567	
Spring	2.686	.250	2.235	3.230		Spring - Sun	nmer	.090	.338	.266	138	.791	579	.759	
Summer	2.596	.245	2.154	3.130		Summer - A	utumn	.865	.306	2.833	138	.005	.261	1.469	
Summer	2.596	.245	2.154	3.130		Summer - S	pring	090	.338	266	138 138	.005 .791	.261 759	1.469 .579	
Summer	2.596	.245	2.154	3.130		Summer - S	pring		.338	266					
			2.154 eat * sea			Summer - S	pring	090	.338 gnificance leve	266	138				
		ns 3: Tre	at * sea			Summer - S	pring gnificant differe	090 ence adjusted si	.338 gnificance leve	266 el is .05.	138				nce Inter
		ns 3: Tre		son		Summer - S	pring	090 ence adjusted si	.338 gnificance leve	266 el is .05.	138			.579	
		ns 3: Tre Esti	eat * sea mates	SON 95% Confide		Summer - Sp The least sig	pring pnificant differe Season Pairw	090 ence adjusted si	.338 gnificance leve Pai	266 el is .05. irwise Cor	138 ntrasts	.791	759	.579 95% Confider	Uppe
Stimate Treatment	ed Mear	ns 3: Tre Estii Mean	eat * sea mates Std. Error	SON 95% Confide Lower	Upper	Summer - Sp The least sig	pring gnificant differe Season Pairw Contrasts	090 ence adjusted significant vise	.338 gnificance leve Pal Contrast Estimate	266 el is .05. irwise Cor	138 ntrasts	.791	759 Adj. Sig.	.579 95% Confider Lower	Uppe .1
Stimate Treatment	Season Autumn	Estin	eat * sea mates Std. Error .248	95% Confide Lower 1.002	Upper 2.000	Summer - Sp The least sig	oring gnificant differe Season Pairw Contrasts Autumn - Sp	090 ence adjusted significance adjusted si	.338 gnificance leve Pai Contrast Estimate 583	266 el is .05. irwise Cor Std. Error .378	138 ntrasts t -1.544	df 138 138	759 Adj. Sig.	.579 95% Confider Lower -1.329	Uppe .1
Estimate Treatment	ed Mear	Mean 1.415 1.998	eat * sea mates Std. Error .248 .297	95% Confide Lower 1.002 1.490	2.000 2.680	Summer - Sp The least sig	Season Pairw Contrasts Autumn - Sp Autumn - Su	090 ence adjusted significance adjusted significance adjusted significance and a signif	.338 gnificance levi Pal Contrast Estimate 583541	266 el is .05. irwise Cor Std. Error .378 .375	138 htrasts t -1.544 -1.443	df 138 138 138 138	759 Adj. Sig. .125 .151	.579 95% Confider Lower -1.329 -1.283	.1 .2 1.3
Estimate Treatment Invaded	Season Autumn Spring Summer	Mean 1.415 1.998 1.957	eat * sea mates Std. Error .248 .297 .293	95% Confide Lower 1.002 1.490 1.455	2.000 2.680 2.632	Summer - Sp The least sig	Season Pairw Contrasts Autumn - Sp Autumn - Su Spring - Autumn - Autumn - Su	090 ence adjusted significance adjusted si	.338 gnificance levi Pal Contrast Estimate 583 541 .583	266 el is .05. irwise Cor Std. Error .378 .375 .378	138 htrasts t -1.544 -1.443 1.544	df 138 138 138 138 138 138	759 Adj. Sig125 .151 .125	.579 95% Confider Lower -1.329 -1.283164	.1 .2 1.3
Estimate Treatment Invaded	Season Autumn Spring	Mean 1.415 1.998 1.957 2.117	eat * sea mates Std. Error .248 .297 .293 .305	95% Confide Lower 1.002 1.490 1.455 1.592	2.000 2.680 2.632 2.816	Summer - Sp The least sig	Season Pairw Contrasts Autumn - Sp Autumn - Su Spring - Autu Spring - Sum	090 ence adjusted si vise vise mmer umn nmer	.338 gnificance level Pal Contrast Estimate583541 .583	266 el is .05. irwise Cor Std. Error .378 .375 .378 .406	138 t -1.544 -1.443 1.544 .103	df 138 138 138 138 138 138	759 Adj. Sig125 .151 .125 .918	.579 95% Confider Lower -1.329 -1.283 164 761	Uppe .1 .2 1.3 .8 1.2
Estimate Treatment Invaded	Season Autumn Spring Summer	Estin Mean 1.415 1.998 1.957 2.117 3.611	eat * sea mates Std. Error .248 .297 .293 .305 .407	95% Confide Lower 1.002 1.490 1.455 1.592 2.890	2.000 2.680 2.632 2.816 4.512	Summer - Sp The least sig	Season Pairw Contrasts Autumn - Sp Autumn - Su Spring - Aut Spring - Sum Summer - Au	090 ence adjusted si vise wing mmer umn mer utumn pring	.338 gnificance levi Pai Contrast Estimate 583 541 .583 .042	266 el is .05. irwise Cor Std. Error .378 .375 .378 .406 .375	t -1.544 -1.443 1.544 .103	df 138 138 138 138 138 138 138 138	759 Adj. Sig125 .151 .125 .918	.579 95% Confider Lower -1.329 -1.283164761201	Uppe .1 .2 1.3 .8 1.2
Estimate Treatment Invaded	Season Autumn Spring Summer Autumn	Mean 1.415 1.998 1.957 2.117	eat * sea mates Std. Error .248 .297 .293 .305	95% Confide Lower 1.002 1.490 1.455 1.592	2.000 2.680 2.632 2.816	Summer - Si The least sig Treatment Invaded	Season Pairw Contrasts Autumn – Sp Autumn – Su Spring – Aut Spring – Summer – Al Summer – Al Summer – Sp	090 ence adjusted si vise vise immer umn immer utumn pring pring	.338 gnificance leve Pai Contrast 583 541 .583 .042 .541 042	266 el is .05. irwise Cor Std. Error .378 .375 .378 .406 .375 .406	138 htrasts t -1.544 -1.443 1.544 .103 103	df 138 138 138 138 138 138	759 Adj. Sig125 .151 .125 .918 .151 .918	.579 95% Confider Lower -1.329 -1.283 164 761 201 844	Uppe .1 .2 1.3 .8 1.2 .7
Estimate Treatment Invaded	Season Autumn Spring Summer Autumn Spring	Estin Mean 1.415 1.998 1.957 2.117 3.611	eat * sea mates Std. Error .248 .297 .293 .305 .407	95% Confide Lower 1.002 1.490 1.455 1.592 2.890	2.000 2.680 2.632 2.816 4.512	Summer - Si The least sig Treatment Invaded	Season Pairw Contrasts Autumn - Sp Autumn - Su Spring - Autu Spring - Summer - Au Summer - Sp Autumn - Sp	090 ence adjusted si vise viring mmer umn nmer utumn poring rring mmer	.338 gnificance leve Pai Contrast 583 541 .583 .042 .541 042	266 el is .05. irwise Cor Std. Error	138 htrasts t -1.544 -1.443 1.544 .103 103 -3.048	df 3 138 4 138 5 138 6 138 6 138 6 138 6 138 7 138 7 138	759 Adj. Sig125 .151 .125 .918 .151 .918	.579 95% Confider Lower -1.329 -1.283164761201844 -2.464	Uppe .1 .2 .1 .3 .8 .1.2 .753
Estimate Treatment Invaded	Season Autumn Spring Summer Autumn Spring	Estin Mean 1.415 1.998 1.957 2.117 3.611	eat * sea mates Std. Error .248 .297 .293 .305 .407	95% Confide Lower 1.002 1.490 1.455 1.592 2.890	2.000 2.680 2.632 2.816 4.512	Summer - Si The least sig Treatment Invaded	Season Pairw Contrasts Autumn - Sp Autumn - Su Spring - Autumn - Su Summer - Au Summer - Sp Autumn - Sp Autumn - Sp	090 ence adjusted si vise vise viring mmer umn nmer utumn poring viring mmer umn	.338 gnificance lew Pai Contrast Estimate 583 541 .583 .042 .541 042 -1.494	266 el is .05. irwise Cor Std. Error .378 .375 .378 .406 .490 .483	t -1.544 -1.443 -1.03 -3.048 -2.752	df 138 138 138 138 138 138 138 138 138 138	759 Adj. Sig125 .151 .125 .918 .151 .918 .003	.579 95% Confider Lower -1.329 -1.283164761201844 -2.464 -2.283	Uppe .1 .2 .1 .3 .8 .1.2 .75 .3 .2.4
	Season Autumn Spring Summer Autumn Spring	Estin Mean 1.415 1.998 1.957 2.117 3.611	eat * sea mates Std. Error .248 .297 .293 .305 .407	95% Confide Lower 1.002 1.490 1.455 1.592 2.890	2.000 2.680 2.632 2.816 4.512	Summer - Si The least sig Treatment Invaded	Season Pairw Contrasts Autumn - Sp Autumn - Su Spring - Autus Summer - Ar Summer - Sp Autumn - Su Autumn - Su Autumn - Su Spring - Autus	090 ence adjusted si	.338 gnificance lew Pai Contrast 5-541 5-83 541 042 042 -1.494	266 el is .05. irwise Cor Std. Error .378 .375 .378 .406 .375 .490	t -1.544 -1.443 -1.03 -3.048 -2.752 3.048	df 138 138 138 138 138 138 138 138 138 138	759 Adj. Sig125 .151 .125 .918 .151 .918 .003 .007	.579 95% Confider Lower -1.329 -1.283164761201844 -2.464 -2.283	.1. .2. 1.3 .8 1.2 .7 5 3 2.4

Figure A3. Results for the GLMM's estimated marginal means, showing the pairwise combinations of treatments (invaded/uninvaded orchards) and seasons (summer, autumn, spring).

References

- 1. Kenis, M.; Auger-Rozenberg, M.A.; Roques, A.; Timms, L.; Péré, C.; Cock, M.J.W.; Settele, J.; Augustin, S.; Lopez-Vaamonde, C. Ecological effects of invasive alien insects. *Biol. Invasions* **2009**, *11*, 21–45. [CrossRef]
- 2. McGlynn, T.P. The worldwide transfer of ants: Geographical distribution and ecological invasions. *J. Biogeogr.* **1999**, *26*, 535–548. [CrossRef]
- 3. Wetterer, J.K. Geographic origin and spread of cosmopolitan ants (Hymenoptera: Formicidae). *Halteres* **2015**, *6*, 66–78.
- 4. Holway, D.A.; Lach, L.; Suarez, A.V.; Tsutui, N.D.; Case, T.J. The causes and consequences of ant invasions. *Annu. Rev. Ecol. Syst.* **2002**, *33*, 181–233. [CrossRef]
- 5. Wetterer, J.K.; Wild, A.L.; Suarez, A.V.; Roura-Pascual, N.; Espadaler, X. Worldwide spread of the Argentine ant, *Linepithema humile* (Hymenoptera: Formicidae). *Myrmecol. News* **2009**, 12, 187–194.
- 6. Ward, P.S. Distribution of the introduced Argentine ant (*Iridomyrmex humilis*) in natural habitats of the lower Sacramento valley and its effects on the indigenous ant fauna. *Hilgardia* **1987**, *55*, 1–16. [CrossRef]
- 7. Hölldobler, B.; Wilson, E.O. *The Ants*; Springer: Berlin/Heidelberg, Germany, 1990.
- 8. Cammell, M.E.; Way, M.J.; Paiva, M.R. Diversity and structure of ant communities associated with oak, pine, eucalyptus and arable habitats in Portugal. *Insectes Soc.* **1996**, *43*, 37–46. [CrossRef]
- 9. Carpintero, S.; Reyes-López, J. The role of competitive dominance in the invasive ability of the Argentine ant (*Linepithema humile*). *Biol. Invasions* **2008**, *10*, 25–35. [CrossRef]
- 10. Gao, Y.; Reitz, S.R. Emerging themes in our understanding of species displacements. *Annu. Rev. Entomol.* **2016**, *6*2, 165–183. [CrossRef]
- 11. Rice, E.S.; Silverman, J. Propagule pressure and climate contribute to the displacement of *Linepithema humile* by *Pachycondyla chinensis*. *PLoS ONE* **2013**, *8*, e56281. [CrossRef]
- 12. Lach, L.; Hooper-Bùi, L.M. Consequences of ant invasions. In *Ant Ecology*; Lach, L., Parr, C., Abbott, K., Eds.; University Press: Oxford, UK, 2010; pp. 261–286.

13. Oliveras, J.; Bas, J.M.; Casellas, D.; Gómez, C. Numerical dominance of the Argentine ant vs native ants and consequences on soil resource searching in Mediterranean cork-oak forests (Hymenoptera: Formicidae). *Sociobiology* **2005**, *45*, 1–16.

- 14. Walters, A.C. Invasion of the Argentine ants (Hymenoptera, Formicidae) in South Australia: Impacts on community composition and abundance of invertebrates in urban parklands. *Austral Ecol.* **2006**, *31*, 367–376. [CrossRef]
- 15. Rowles, A.D.; O'Dowd, D.J. Impacts of the invasive Argentine ant on native ants and other invertebrates in coastal scrub in south-eastern Australia. *Austral Ecol.* **2009**, *34*, 239–248. [CrossRef]
- 16. Devenish, A.J.M.; Gomez, C.; Bridle, J.R.; Newton, R.J.; Summer, S. Invasive ants take and squander native seeds: Implications for native plant communities. *Biol. Invasions* **2019**, *21*, 451–466. [CrossRef]
- 17. Menke, S.B.; Ward, P.S.; Holway, D.A. Long-term record of Argentine ant invasions reveals enduring ecological impacts. *Ecology* **2018**, *99*, 1194–1202. [CrossRef] [PubMed]
- 18. Cerdá, X.; Palacios, R.; Retana, J. Ant community structure in citrus orchards in the Mediterranean Basin: Impoverishment as a consequence of habitat homogeneity. *Environ. Entomol.* **2009**, *38*, 317–324. [CrossRef]
- 19. FAO. *Citrus Fruit Fresh and Processed. Statistical Bulletin 2016;* Food and Agriculture Organization of the United Nations: Rome, Italy, 2017.
- 20. Pekas, A.; Tena, A.; Aguilar, A.; Garcia-Marí, F. Spatio-temporal patterns and interactions with honeydew-producing Hemiptera of ants in a Mediterranean citrus orchard. *Agric. For. Entomol.* **2011**, 13, 89–97. [CrossRef]
- 21. Zina, V.; Branco, M.; Franco, J.C. Ant community associated to citrus canopy in Southern Portugal over a century after the invasion by the Argentine ant. *Phytoparasitica* **2017**, *45*, 183–200. [CrossRef]
- 22. Samways, M.J. Community structure of ants (Hymenoptera: Formicidae) in a series of habitats associated with citrus. *J. Appl. Ecol.* **1983**, *20*, 833–847. [CrossRef]
- 23. Queiroz, J.M.; Oliveira, P.S. Tending ants protect honeydew-producing whiteflies (Homoptera: Aleyrodidae). *Environ. Entomol.* **2001**, *30*, 295–297. [CrossRef]
- 24. Buckley, R.; Gullan, P. More aggressive ant species (Hymenoptera: Formicidae) provide better protection for soft scales and mealybugs (Homoptera: Coccidae, Pseudococcidae). *Biotropica* **1991**, 23, 282–286. [CrossRef]
- 25. Franco, J.C.; Silva, E.B.; Carvalho, J.P. Cochonilhas-Algodão (Hemiptera, Pseudococcidae) Associadas Aos Citrinos em Portugal; ISA Press: Lisbon, Portugal, 2000.
- 26. Gullan, P.J. Relationships with ants. In *Soft Scale Insects—Their Biology, Natural Enemies and Control*; Ben-Dov, Y., Hodgson, C.J., Eds.; Elsevier Science: Amsterdam, The Netherlands, 1997; pp. 351–377.
- 27. Bodenheimer, F.S. Citrus Entomology in the Middle East with Special References to Egypt, Iran, Irak, Palestine, Syria, Turkey; Dr. W. Junk Publishers: The Hague, The Netherlands, 1951.
- 28. Ben-Dov, Y. Relationships with ants. In *Armored Scale Insects—Their Biology, Natural Enemies and Control;* Rosen, D., Ed.; Elsevier: Amsterdam, The Netherlands, 1990; pp. 339–343.
- 29. Martínez-Ferrer, M.T.; Grafton-Cardwell, E.; Shorey, H.H. Disruption of parasitism of the California red scale (Homoptera: Diaspididae) by three ant species (Hymenoptera: Formicidae). *Biol. Control.* **2003**, *26*, 279–286. [CrossRef]
- 30. Franco, J.C.; Garcia-Marí, F.; Ramos, A.P.; Besri, M. Survey of the situation of citrus pest management in Mediterranean countries. *IOBC WPRS Bull.* **2006**, *29*, 335–346.
- 31. OPMP. A National Road Map for Integrated Pest Management. United States Department of Agriculture, Office of Pest Management Policy. 2018. Available online: https://www.ars.usda.gov/arsuserfiles/opmp/ipm%20road%20map%20final.pdf (accessed on 10 August 2020).
- 32. Martins, M.N. Une fourmi terrible envahissant l'Europe (*Iridomyrmex humilis* Mayr). *Brotéria Series Zoologia* **1907**, *6*, 101–102.
- 33. Coutinho, M.P. A "formiga-argentina" "*Iridomyrmex humilis*" Mayr "var. arrogans" Santschi. *B Minist. Agric.* **1929**, *11*, 95–116.
- 34. Robinson, E.J.H. Polydomy: The organization and adaptive function of complex nest system in ants. *Curr. Opin. Insect. Sci.* **2014**, *5*, 37–43. [CrossRef]
- 35. Heller, N.E.; Sanders, N.J.; Gordon, D.M. Linking temporal and spatial scales in the study of an Argentine ant invasion. *Biol. Invasions* **2006**, *8*, 501–507. [CrossRef]
- 36. McGlynn, T.P. The ecology of nest movement in social insects. *Annu. Rev. Entomol.* **2012**, *57*, 291–308. [CrossRef]

37. Koop, E.; Sobral, M.; Soares, T.; Woerner, M. *Os Solos do Algarve e as Suas Características*; Vista Geral Direcção Regional de Agricultura do Algarve: Faro, Portugal, 1989.

- 38. Chong, C.S.; Hoffmann, A.A.; Thomson, L.J. Commercial agrochemical applications in vineyards do not influence ant communities. *Environ. Entomol.* **2007**, *36*, 1374–1383. [CrossRef]
- 39. King, J.R.; Porter, S.D. Evaluation of sampling methods and species richness estimators for ants in upland ecosystems in Florida. *Environ. Entomol.* **2005**, *34*, 1566–1578. [CrossRef]
- 40. Gotelli, N.J.; Colwell, R.K. Quantifying biodiversity: Procedures and pitfalls in the measurement and comparison of species richness. *Ecol. Lett.* **2011**, *4*, 379–391. [CrossRef]
- 41. Martínez-Ferrer, M.T.; Campos-Rivela, J.M. Diversity, spatial distribution, and sampling for ant management decision-making in integrated pest management programs in citrus groves. *Entomol. Exp. Appl.* **2017**, 162, 251–260. [CrossRef]
- 42. Dunn, R.R.; Guénard, B.; Weiser, M.D.; Sanders, N. Geographic gradients. In *Ant Ecology*; Lach, L., Parr, C.L., Abbott, K.L., Eds.; Oxford University Press: Oxford, UK, 2010; pp. 38–58.
- 43. Collingwood, C.; Prince, A. A guide to ants of continental Portugal (Hymenoptera: Formicidae). *B Soc. Port. Entomol.* **1998**, *5*, 1–49.
- 44. Gómez, K.; Espadaler, X. Hormigas Ibéricas. 2007. Available online: http://www.hormigas.org (accessed on 15 November 2016).
- 45. IBM Corp. IBM SPSS Statistics for Windows, Version 25.0; IBM Corp: Armonk, NY, USA, 2017.
- 46. R Studio Team. RStudio: Integrated Development for R; RStudio Inc.: Boston, MA, USA, 2015.
- 47. Van den Brink, P.J.; Van den Brink, N.W.; Ter Braak, C.J.F. Multivariate analysis of ecotoxicological data using ordination: Demonstrations of utility on the basis of various examples. *Australas J. Ecotox.* **2003**, *9*, 141–156.
- 48. Van den Brink, P.J.; Ter Braak, C.J.F. Principal response curves: Analysis of time-dependent multivariate responses of biological community to stress. *Environ. Toxicol. Chem.* **1999**, *18*, 138–148. [CrossRef]
- 49. Del Toro, I.; Ribbons, R.R.; Shannon, L.P. The little things that run the world revisited: A review of ant-mediated ecosystem services and disservices (Hymenoptera: Formicidae). *Myrmecol. News* **2012**, *17*, 133–146.
- 50. Vanaclocha, P.; Monzó, C.; Gómez, K.; Tortosa, D.; Pina, T.; Castañera, P.; Urbaneja, A. Hormigas (Hymenoptera: Formicidae) presentes en el suelo de los cítricos de la provincia de Valencia. *Phytoma* **2005**, *171*, 14–24.
- 51. Alvis, L.; Garcia-Marí, F. Identification and abundance of ants (Hymenoptera: Formicidae) in citrus trees from Valencia (Spain). *IOBC WPRS Bull.* **2006**, 29, 111–116.
- 52. Urbaneja, A.; Garcia-Marí, F.; Tortosa, D.; Navarro, C.; Vanaclocha, P.; Bargues, L.; Castañera, P. Influence of ground predator on the survival of the Mediterranean fruit fly pupae, *Ceratitis capitata*, in Spanish citrus orchards. *BioControl* **2006**, *51*, 611–626. [CrossRef]
- 53. La Pergola, A.; Alicata, A.; Longo, S. Survey of the ants (Hymenoptera, Formicidae) in citrus orchards with different types of crop management in Sicily. *Control Citrus Fruit Crop. IOBC WPRS Bull.* **2008**, *38*, 233–237.
- 54. Calabuig, A.; Garcia-Marí, F.; Pekas, A. Ants in citrus: Impact on the abundance, species richness, diversity and community structure of predators and parasitoids. *Agric. Ecosyst. Environ.* **2015**, 213, 178–185. [CrossRef]
- 55. Wetterer, J.K.; Wetterer, A.L.; Hebard, E. Impact of the Argentine ant, *Linepithema humile* on the native ants of Santa Cruz Island, California. *Sociobiology* **2001**, *38*, 709–721.
- 56. Arnan, X.; Andersen, A.N.; Gibb, H. Dominance-diversity relationships in ant communities differ with invasion. *Glob. Chang. Biol.* **2018**, *24*, 4614–4625. [CrossRef]
- 57. Abril, S.; Gómez, C. Ascertaining key factors behind the coexistence of the native ant species *Plagiolepis pygmaea* with the invasive Argentine ant *Linepithema humile* (Hymenoptera: Formicidae). *Sociobiology* **2009**, *53*, 559–568.
- 58. Roura-Pascual, N.; Bas, J.M.; Hui, C. The spread of the Argentine ant: Environmental determinants and impacts on native ant communities. *Biol. Invasions* **2010**, *12*, 2399–2412. [CrossRef]
- 59. Human, K.G.; Gordon, D.M. Behavioral interactions of the invasive Argentine ant with native ant species. *Insectes Soc.* **1999**, *46*, 159–163. [CrossRef]
- 60. Holway, D.A. Competitive mechanisms underlying the displacement of native ants by the invasive Argentine ant. *Ecology* **1999**, *80*, 238–251. [CrossRef]
- 61. Blight, O.; Provost, E.; Renucci, M.; Tirard, A.; Orgeas, J. A native ant armed to limit the spread of the Argentine ant. *Biol. Invasions* **2010**, *12*, 3785–3793. [CrossRef]

62. Castro-Cobo, S.; Carpintero, S.; Reyes-López, J.L.; Sergio, F.; Angulo, E. Humans and scavenging raptors facilitate Argentine ant invasion in Doñana National Park: No counter-effect of biotic resistance. *Biol. Invasions* **2019**, *21*, 2221–2232. [CrossRef]

- 63. Way, M.J.; Cammell, M.E.; Paiva, M.R.; Collingwood, C.A. Distribution and dynamics of the Argentine ant *Linepithema* (*Iridomyrmex*) *humile* (Mayr) in relation to vegetation, soil conditions, topography and native competitor ants in Portugal. *Insect Soc.* **1997**, *44*, 415–433. [CrossRef]
- 64. Rust, M.K.; Reierson, D.A.; Paine, E.; Blum, L.J. Seasonal activity and bait preferences of the Argentine ant (Hymenoptera: Formicidae). *J. Agric. Urban Entomol.* **2000**, *17*, 201–212.
- 65. Davidson, D.W. Resource discovery versus resource domination in ants: A functional mechanism for breaking the trade-off. *Ecol. Entomol.* **1998**, 23, 484–490. [CrossRef]
- 66. Centorame, M.; Lancia, A.; Mori, E.; D'Eustacchio, D.; Fanfani, A. Could *Linepithema humile* (Hymenoptera, Formicidae) influence ant community composition? A preliminary study in a natural area in Italy. *Redia* **2017**, *100*, 89–94.
- 67. Sanders, N.J.; Barton, K.E.; Gordon, D.M. Long-term dynamics of the distribution of the invasive Argentine ant, *Linepithema humile*, and native ant taxa in northern California. *Oecologia* **2001**, *127*, 123–130. [CrossRef] [PubMed]
- 68. Parker, I.M.; Simberloff, D.; Lonsdale, W.M.; Goodell, K.; Wonham, M.; Kareiva, P.M.; Williamson, M.H.; Von Holle, B.; Moyle, P.B.; Byers, J.E.; et al. Impact: Toward a framework for understanding the ecological effects of invaders. *Biol. Invasions* 1999, 1, 3–19. [CrossRef]
- 69. Mestre, L.; Piñol, J.; Barrientos, J.A.; Espadaler, X. Differential ant exclusion from canopies shows contrasting top-down effects on community structure. *Oecologia* **2016**, *180*, 193–203. [CrossRef]
- 70. Diaz, M.; Abril, S.; Enriquez, M.L.; Gómez, C. Assessment of the Argentine ant invasion management by means of manual removal of winter nests in mixed cork oak and pine forests. *Biol. Invasions* **2014**, *16*, 315–327. [CrossRef]
- 71. Carvalho, J.P.; Ilharco, F.A.; Ferreira, M.A.; Carvalho, M.U.P. *Manual de Pragas e Sintomas do Ataque de Insectos e Ácaros em Citrinos*; Estação Agronómica Nacional: Oeiras, Portugal, 1999.
- 72. James, D.G.; Stevens, M.M.; O'Malley, K.J. The impact of foraging ants on populations of *Coccus hesperidum* L. (Hem., Coccidae) and *Aonidiella aurantii* (Maskell) (Hem., Diaspididae) in an Australian citrus grove. *J. Appl. Ent.* 1997, 121, 257–259. [CrossRef]
- 73. James, D.G.; Stevens, M.M.; O'Malley, K.J.; Faulder, R.J. Ant foraging reduces the abundance of beneficial and incidental arthropods in citrus canopies. *Biol. Control* **1999**, *14*, 121–126. [CrossRef]
- 74. Smith, D.; Beattie, G.A.C.; Broadley, R. Citrus Pests and Their Natural Enemies: Integrated Pest Management in Australia; Desert Oak Publishing Services: Brisbane, Australia, 1997.
- 75. Calabuig, A.; Garcia-Marí, F.; Pekas, A. Ants affect the infestation levels but not the parasitism of honeydew and non-honeydew producing pests in citrus. *Bull. Entomol. Res.* **2014**, *104*, 405–417. [CrossRef]
- 76. Benfradj, N.; Vettraino, A.M.; Tomassini, A.; Bruni, N.; Vannini, A.; Boughalleb-M'Hamdi, N. Citrus gummosis incidence and role of ants (*Lasius grandis*) and snails (*Helix aspersa*) as vectors of the disease in Tunisia. *For. Pathol.* **2018**, *48*, e12423. [CrossRef]
- 77. Buczkowski, G.; Mothapo, N.P.; Wossler, T.C. Let them eat termites—Prey-baiting provides effective control of Argentine ants, *Linepithema humile*, in a biodiversity hotspot. *J. Appl. Entomol.* **2018**, 142, 504–512. [CrossRef]

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