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Oribatid mites of conventional and organic vineyards in the Valencian Community, Spain

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ABSTRACT

In this study the oribatid mite communities of conventional and organic vineyards in the Valencian Community (Spain) were compared. The soil samples were collected in El Poble Nou de Benitatxell in autumn 2014 and spring 2015 from four sites, treated as replicates, each including a conventional vineyard, an organic vineyard, and a control (natural habitat, i.e. in plots 1-3 an abandoned vineyard, in plot 4 an area never used in agriculture). Two parallel samples were collected in each vineyard from a zone between vine rows, driven by a tractor (Tr), a zone between vines (Vi), the border of the vineyard (Bo) and from a control, making a total of 112 samples. In total 3,225 oribatid mites were obtained represented by 59 species. No differences were found in density of Oribatida between the conventional, organic vineyards and the control, but the species diversity was higher in the control than in the vineyards. In the vineyards the density and species number of the oribatid mites were highest between vines (the average from all vineyards and both seasons was 4,400 individuals per 1 m², 15 species), followed by the border of the vineyards (2,800 individuals per 1 m², 14 species) and were lowest between vine rows (400 individuals per 1 m², 6 species). The species diversity of Oribatida was higher in autumn than in spring, while the density followed this pattern only in the vineyards, but not in the control. In the vineyards *Oribatula excavata* dominated ($D = 25$), followed by *Minunthozetes quadriareatus* and *Passalozetes africanus* ($D = 18$ and 14, respectively), while in the control these species were not abundant. In the control the most abundant species was *Oppiella subpectinata* ($D = 28$), followed by *Eremulus flagellifer* ($D = 20$). *Podoribates longipes* and *Steganacarus boulfekhari* are reported for the first time in Spain. To conclude, the oribatid mites did not benefit from the organic cultivation of the vineyards, probably because they are tolerant to herbicides used in the conventional systems but sensitive to mechanical cultivation of soil, which was even more intense in organic vineyards than in the conventional ones.

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Keywords Oribatida, conventional, organic, vineyards, tillage

Introduction

Spain is the country with the largest area of vineyards in the world (with more than 1 million ha, i.e. 14% of world's total vineyard area) (OIV 2016). It also leads in the organic viticulture, with

the largest area of organic vineyards (MAGRAMA 2015). The main organic wine producing regions in Spain are Castilla La Mancha (4,095 ha), Murcia (3,722 ha), València (2,086 ha) and Cataluña (1,127 ha) (Fabeiro *et al.* 2007).

The main difference between conventional and organic (also called ecological) agriculture is the use of pesticides and chemical fertilizers in conventional farming and prohibition of these activities in organic systems. The organic production relies on crop rotation, fixation of natural nitrogen, biologically active soil, recycled farm manure and crop residues, and on biological or mechanical weed and pest control (Mäder *et al.* 2002; Bengtsson *et al.* 2005). In Spain in organic vineyards some insecticides and fungicides are allowed (azadirachtin, sulfur, and sometimes, when the weather conditions are optimal for some fungi, copper calcium sulfate), while in conventional vineyards different pesticides are used (including chlorpyrifos, azoxistrobin, mancozeb).

The biological activity of soil is affected by oribatid mites (Walter and Proctor 1999), which are one of the most abundant arthropod group in the organic horizons, also in vineyards (Suzuki 1979), where they play an important role in decomposition of organic matter, nutrient cycling and soil formation (Behan-Pelletier 1999). Additionally, some species can possibly serve in the pest control. For example, the pathogenic fungus, *Rhizoctonia solani* that attacks grapevine roots (Walker 1992), can be controlled by the oribatid species *Protoribates agricola* and *Schelorbates azumaenis* (Nakamura *et al.* 1991; Enami and Nakamura 1996).

In undisturbed agroecosystems the density of Oribatida easily reaches several thousand individuals per 1 m² and 20-50 species (Behan-Pelletier 1999). These mites are more abundant in grasslands, with the densities up to several hundred thousand individuals per 1 m². In the agricultural fields the density of Oribatida is about ten times lower, mainly because of the cultivation practices that reduce the density of the soil fauna (Niedbala 1980). Among these practices, especially mechanical works on the soil, like tillage, ploughing, disking, etc. or using too high amounts of fertilizers or some chemicals, can have negative effects on the oribatid communities (summarized by Behan-Pelletier 1999).

The Oribatida of vineyards are poorly studied in comparison to other agroecosystems, like meadows, pastures, orchards or arable fields. Some studies on the mites from vineyards have been conducted in Germany (Jörger 1991) and Brazil (Johann *et al.* 2014), but the Oribatida were treated only as a group. More detailed studies on these mites from vineyards have been carried out in Japan (Suzuki 1979), Azerbaijan (Samedov *et al.* 1987), Italy (Nannelli and Simoni 2002) and India (Acharya and Basu 2014), but they were based only on adults, and none of them included organic vineyards.

Most studies showed a positive effect of organic farming on density (96 of 117 studies; i.e. 82%) and species richness (53 of 63 studies; i.e. 84%) of plants and animals (Bengtsson *et al.* 2005). Regarding microarthropods, most studies on the effect of organic and conventional agriculture have been carried out in the annual cultures (e.g. Bettiol *et al.* 2002; Van Leeuwen *et al.* 2015) and more soil microarthropods were found in organically managed farming systems than in conventional ones. Oribatida reacted positively to organic management, together with Uropodina (Badejo *et al.* 2004). However, in apple orchards that were permanent crop, there was no significant difference in density of oribatid mites between the organic and conventional management (Doles *et al.* 2001). Based on these results we hypothesized that organic cultivation of vineyards vs. conventional one has no impact on the oribatid density and species richness.

The aims of this study were (1) to compare the oribatid communities in conventional and organic vineyards, (2) investigate the community structure of Oribatida in selected habitats in vineyards, (3) compare the dynamics of these mites in autumn and spring, and (4) improve the knowledge of the species diversity in the vineyards.

Materials and methods

Study sites

The study was carried out in El Poble Nou de Benitatxell, a village located in the Valencian Community, Spain (Figure 1). The village has an area of 12.65 km² and is situated on a hill of the elevation 159 m a.s.l. The climate of the area is clearly Mediterranean, with mild temperature in winter and higher in summer, with a characteristic period of drought in summer and higher rainfall in spring and autumn (Figure 2). The year 2014 was drier (annual rainfall 251 mm) than the year 2015 (518 mm). In the sampling seasons (autumn 2014 and spring 2015), the average temperature was 17°C and 18°C, respectively, and the average rainfall was 135 mm and 25 mm (MAGRAMA 2015).

The soils are calcareous, including deposition and lithology marls (“Tap” facies) and decarbonated soils (terra rossa) (Barbér Vallés and Moity Martín 2009). The morphology of the landscape is characterized by a strong anthropization, predominantly terraced with stone walls. Traditionally, these terraces were planted with cereals, legumes (beans) and Muscat grapes, but nowadays unirrigated vineyards with table grapes (Muscat of Alexandria variety) predominate (80% of the cultivated land, i.e. 368 ha; Statistics National Institute: Agrarian Census 2009).

Most vineyards have been cultivated in a conventional way, with the use of synthetic fertilizers and pesticides. But since farmers are becoming aware of problems caused by these substances in the ecosystems, they have started an initiative to change the conventional system into organic one. Thus, the crop receives water exclusively from the rain and the farmers

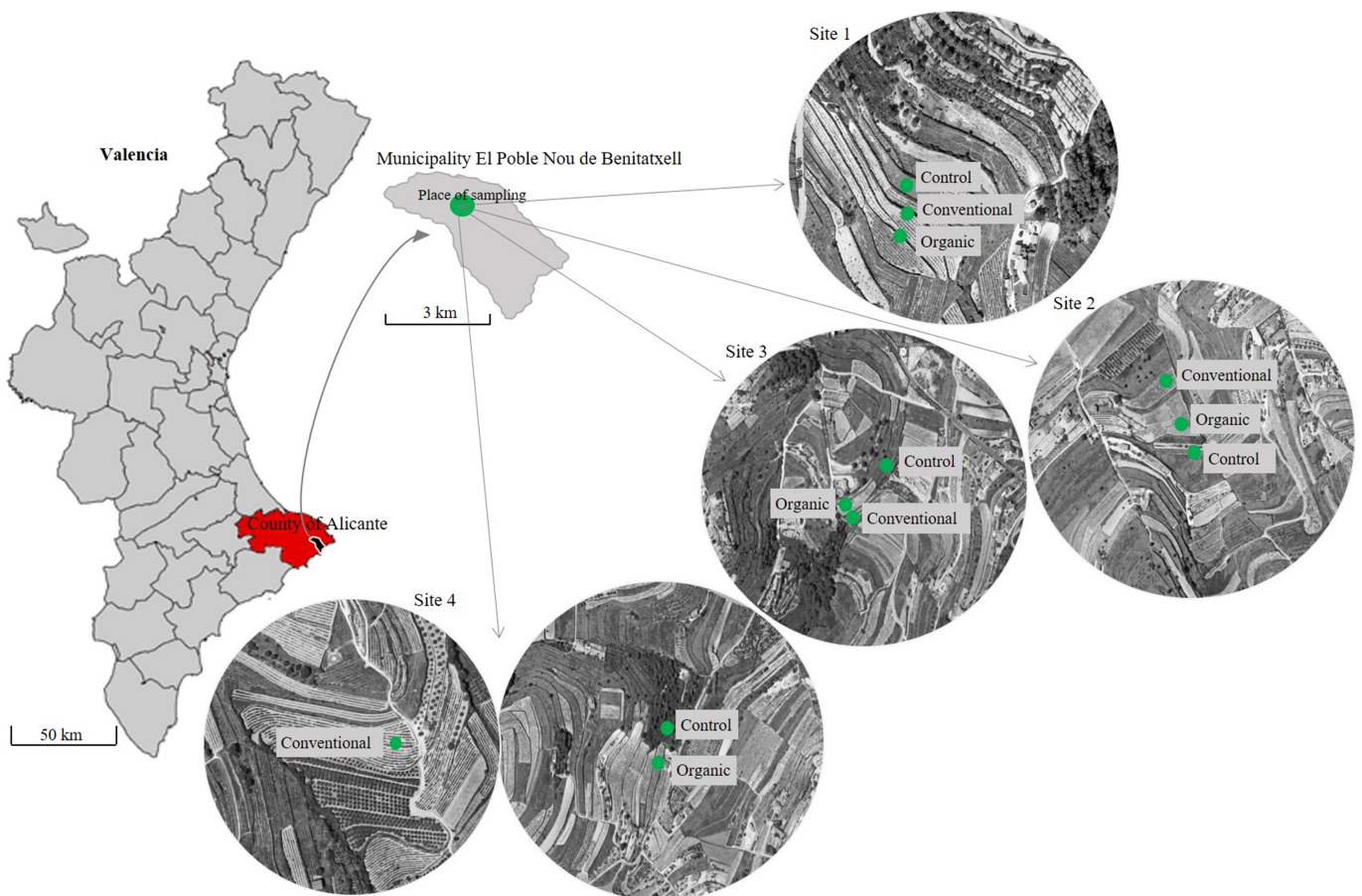


Figure 1 Locality of the study plots in El Poble Nou de Benitatxell.

maintain the traditional cultural techniques without mineral fertilizing and only some pesticides (azadirachtin, sulfur, and seldom, copper calcium sulfate). In Spain the normative says a farmer needs 4 years to get the organic national certification. The project also prioritizes fair trade, giving a fair price to the farmer and selling locally, thus reducing CO₂ emissions from the transport (Domingo *et al.* 2014).

Four study sites, which are included in this local project of changing the table grapes production into organic system, were selected. They were treated as replicates and each included a conventional vineyard, an organic vineyard and a natural habitat (control), that was in plots 1-3 an abandoned vineyard and in plot 4 the area never used in agriculture. Their locality was respectively: site 1 - 38°44'16.28" N, 0°7'33.80" E; 38°44'16.20" N, 0°7'32.35" E; 38°44'17.52" N, 0°7'33.35" E, site 2 - 38°44'15.59" N, 0°8'22.83" E; 38°44'13.17" N, 0°8'24.57" E, and 38°44'11.74" N, 0°8'24.97" E), site 3 - 38°43'38.10" N, 0°8'51.19" E; 38°43'38.34" N, 0°8'50.50" E; 38°43'40.63" N, 0°8'54.48" E, and site 4 - 38°43'5.94" N, 0°8'17.71" E; 38°43'50.83" N, 0°7'48.42" E, 38°43'52.24" N, 0°7'48.48" E. The vines in sites 1-4 were planted respectively in the years 1992, 1967, 1980 and 1987, but since 2012 all of them have been cultivated in organic system.

In the conventional system of management (Table 1) the farmers apply herbicides to kill weeds before spring (the active ingredient is glyphosate 36%) and they use chemical fertilizers with different formulations. In organic vineyards they do not apply any herbicide, but from the second tilling on, an implement that cuts weeds between vines is added. Since 2013 these plots are fertilized with sheep manure. Finally, uncultivated plots do not show any intervention or modification, being occupied by spontaneous vegetation.

In total, 40 plant species were recorded in all plots during a 6-month study (March-August). The number of species was similar in conventional and organic vineyards and in the natural habitat (Table 2), while the plants density and plant cover were highest in the natural habitat, followed by the organic vineyards, and were lowest in conventional vineyards.

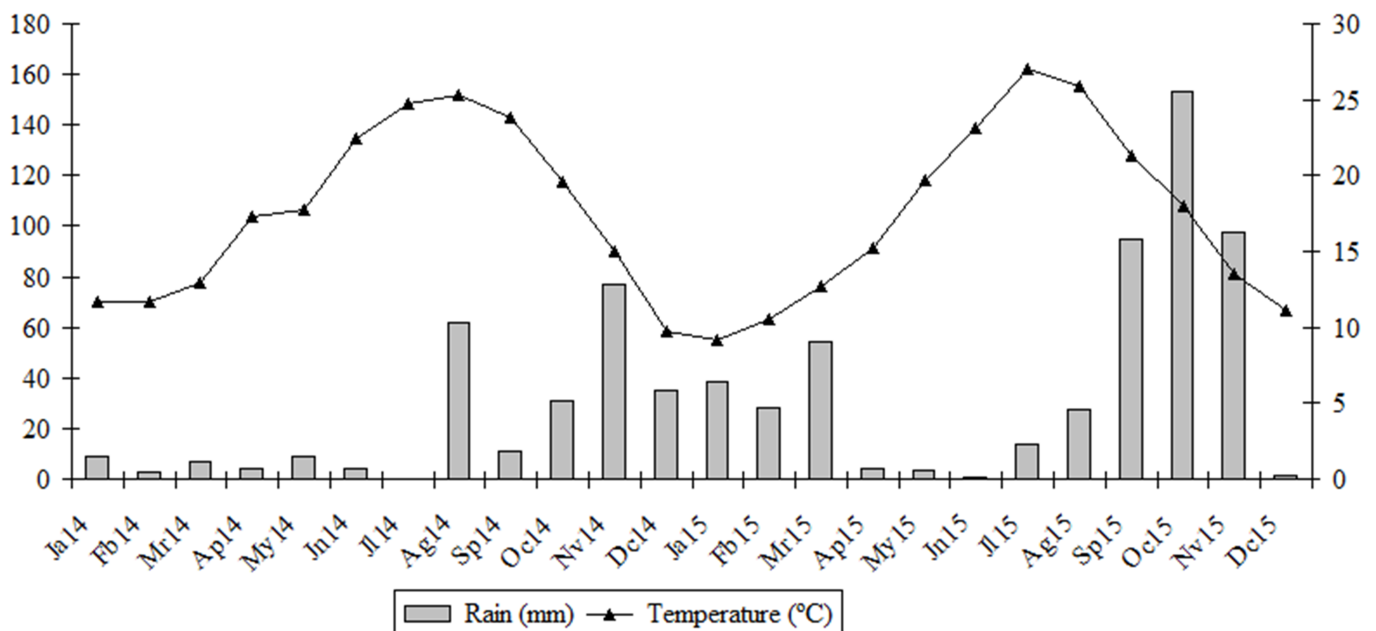


Figure 2 Mean temperature (°C) and sum of precipitation (mm) in all months of years 2014 and 2015 in El Poble Nou de Benitatxell (data from MAGRAMA 2015).

Table 1 Cultivation practices in vineyards in El Poble Nou de Benitatxell.

Cultivation practice	Method	Time	Conventional vineyards	Organic vineyards
Annual pruning of vines	Secateurs	January	Yes	Yes
Annual fertilizing	Fertilizer spreader	February	Yes (mineral)	Yes (sheep manure)
Tillage (to incorporate fertilizer)	Tractor	February	Yes	Yes
Weed control	Herbicide	April	Yes	No
Weed control	Tractor	April	No	Yes
Tillage	Tractor	May	Yes	Yes
Removing branches, clearing canopy	Manual	May	Yes	Yes
Tillage	Tractor	June	Yes	Yes
Collecting fruit	Manual	Mid-August-September	Yes	Yes

Sampling and mite analyses

Sampling was carried out in autumn (13 Nov. 2014) and spring (12 Jun. 2015). In each of the four sites two parallel samples (each of the area 950 cm², 10 cm deep) were collected with metal corer from seven plots: (1) natural habitat, treated as the control, (2) conventional vineyard, zone between vine rows, driven by a tractor (hereafter referred to as Tr), (3) conventional vineyard, zone between vines (Vi), (4) border of conventional vineyard (Bo), and organic vineyard from the same habitats, (5) Tr, (6) Vi and (7) Bo. In total 112 samples were collected (4 sites x 2 samples x 7 plots x 2 seasons).

The samples were extracted in Berlese funnels during 4 days at the Universitat Politècnica de València; the time of extraction was based on the observation that after 4 days no living mites were present in the samples. The mites were preserved in 70% ethanol and Oribatida were sorted out under stereomicroscope and determined, including the juveniles, at the UTP University of Science and Technology. The nomenclature of the Oribatida follows Weigmann (2006) and partly Subías (2004, 2015). The data presented in tables 3 and 5 and in Figure 3 refer to all mites, including adults and juveniles. Authors and dates of the taxonomic names are listed in Table 1.

Oribatid mite populations were characterized by the density (*A*), dominance (*D*) and constancy (*C*) indices, and oribatid mite communities were characterized by the number of species (*S*) and the Shannon (*H'*) diversity index (Odum 1982). The basic statistical descriptors included the minimum, maximum, mean values and standard deviation. For the other statistical analyses, the values were log-transformed $\ln(x+1)$ (Łomnicki 2010). Normality of the distribution was tested with the Kolmogorov-Smirnov test, while the equality of variance in different samples, with the Levene test. The assumption of normality or equality of variance was not met, so the non-parametric ANOVA rang Kruskal-Wallis was used and then, in case of significant differences between averages, the multiple comparison test between average ranks was used. The level of significance for all statistical tests was accepted at $\alpha = 0.05$. The statistical calculations mentioned above were carried out with STATISTICA 10.0 software.

Subsequently, based on the same log-transformed $\ln(x+1)$ data set the analysis of the community structure of the Oribatida with detrended correspondence analysis (DCA) was performed using MVSP 3.2 (Multi Variate Statistical Package, Kovach Computing Services;

Table 2 Vegetation characteristics in studied plots in El Poble Nou de Benitatxell (average data from 4 sites and from 6 vegetation analyses from March to August); *A* – density per 1 m², % of cover.

Species	Control		Conventional vineyards		Organic vineyards	
	<i>A</i>	%	<i>A</i>	%	<i>A</i>	%
<i>Anagallis arvensis</i> L.	0	0	0.6	0.1	0.6	0.4
<i>Arisarum vulgare</i> Targ.-Tozz.	0.3	0	0	0	0	0
<i>Avena fatua</i> L.	0.2	0	0.5	0	0.1	0.1
<i>Blackstonia perfoliata</i> (L.) Huds.	0.4	0.1	0	0	0	0
<i>Brachypodium retusum</i> Pers. (Beauv.)	26.3	35	0.3	0.3	0	0
<i>Bromus madritensis</i> L.	1	2.7	0.2	0	0.1	0
<i>Calendula arvensis</i> L.	0.1	0	0.4	0.1	4.7	1.2
<i>Centaurea cyanus</i> L.	0	0	0.3	0.7	0	0
<i>Convolvulus althaeoides</i> L.	0.1	1.3	0.7	0.1	2.3	0.9
<i>Conyza bonariensis</i> (L.) Cronquist	2.9	1	0	0	0.3	0.5
<i>Cynodon dactylon</i> (L.) Pers.	0	0	0.5	0.4	0	0
<i>Diplotaxis erucoides</i> (L.) DC.	0	0	0	0	0.1	0
<i>Elymus pungens</i> (Pers.) Melderis	0.1	1.2	1.3	0.4	0.1	0.3
<i>Erodium cicutarium</i> (L.) L'Hér. ex Aiton.	0.1	0	0.4	0.7	0.1	0.4
<i>E. malacoides</i> (L.) L'Hér.	0.1	0	0.3	0.7	1.5	2.9
<i>Euphorbia falcata</i> L.	0	0	0	0	0.3	0
<i>E. helioscopia</i> L.	1	0.1	0	0	0.3	0.1
<i>E. segetalis</i> L.	0.4	0.1	0.9	0.5	0.9	0.1
<i>Foeniculum vulgare</i> Mill.	0.3	0.5	0	0	0	0
<i>Fumaria officinalis</i> L.	0.3	0.1	0	0	0.6	0.2
<i>Galactites tomentosa</i> Moench	0	0	0.5	0.4	0	0
<i>Geranium rotundifolium</i> L.	0	0	0	0	0.7	2.7
<i>Hyparrhenia hirta</i> (L.) Stapf	2.4	8.5	0.5	0.3	0.2	0.4
<i>Lavatera cretica</i> L.	0	0	1.9	2.8	0	0
<i>Leucanthemum paludosum</i> (Poir.) Bonnet & Barratte	0	0	1.1	0.2	5.1	1.6
<i>Lotus ornithopodioides</i> L.	0.3	0.1	0.8	0.2	1	0.3
<i>Medicago minima</i> (L.) Bartal	0.7	0.5	0.6	1.6	2	2.9
<i>Mercurialis annua</i> L.	0	0	0	0	2.3	0.2
<i>Muscari neglectum</i> Guss. ex Ten.	0.1	0	0	0	0	0
<i>Oxalis pes-caprae</i> L.	3.8	2	6.8	4.3	4.6	4
<i>Picris echioides</i> L.	0.1	0	0.5	0.1	0.8	0.1
<i>Plantago albicans</i> L.	0.1	0	1	0.2	0	0
<i>P. lagopus</i> L.	0.2	0	0	0	0	0
<i>Reichardia tingitana</i> (L.) Roth	0	0	0.2	0.3	0.1	0.1
<i>Rhamnus alaternus</i> L.	0.1	0.1	0.1	0	0	0
<i>Rubia peregrina</i> L.	0.4	0.1	0	0	0	0
<i>Sonchus oleraceus</i> L.	1.3	0.2	1.3	0.5	2.3	0.7
<i>S. tenerrimus</i> L.	0.1	1	0.1	0	0.3	0
<i>Vicia pseudocracca</i> Bertol.	0.1	0	0	0	0.2	0
<i>V. sativa</i> L.	0.3	0	0	0	0.3	0.1
Total	43.3	54.7	21.7	14.7	31.3	20.2
Number of species		29		25		27

Table 4 Mean values \pm SD and range of density (A , in 10^3 individuals/m²), number of species (S) and Shannon (H') index of Oribatida in studied plots in El Poble Nou de Benitatxell in spring (Sp) and autumn (Au); Tr – zone between rows driven by a tractor, Vi – zone between vines, Bo – border; SD = standard deviation, H, p – the result of Kruskal-Wallis nonparametrical analysis of variance (ANOVA)

Parameter		Control	Conventional			Organic			ANOVA rang	
			Tr	Vi	Bo	Tr	Vi	Bo	Kruskal-Wallis	
									H	p
Adults	Au	5.5 ^{abc} \pm 9.2	1.0 ^{ac} \pm 0.8	6.9 ^{bc} \pm 4.8	3.9 ^{ab} \pm 1.7	0.3 ^c \pm 0.5	5.8 ^{ab} \pm 4.3	4.1 ^{abc} \pm 3.2	26.23	0
	A ($\bar{x} \pm$ SD, range)	0.5-28.0	0.2-2.6	2.5-16.2	2.2-6.8	0.0-1.5	0.3-11.2	0.0-9.7		
	Sp	4.6 ^a \pm 6.1	0.3 ^{ab*} \pm 0.6	2.2 ^{a*} \pm 1.8	0.9 ^{ab*} \pm 0.8	0.1 ^{bc} \pm 0.1	0.3 ^{ac*} \pm 0.4	0.9 ^{ac*} \pm 1.7	23.38	0
	A ($\bar{x} \pm$ SD, range)	0.0-18.3	0.0-1.7	0.0-5.4	0.2-2.6	0.0-0.4	0.0-1.3	0.0-4.4		
Juveniles	Au	1.4 ^{ac} \pm 2.0	0.0 ^b \pm 0.0	1.0 ^{ac} \pm 1.0	0.5 ^{abc} \pm 0.7	0.0 ^{ab} \pm 0.0	1.2 ^c \pm 0.9	0.6 ^{abc} \pm 0.7	25.39	0
	A ($\bar{x} \pm$ SD, range)	0.0-6.1	0.0-0.0	0.0-2.5	0.0-1.7	0.0-0.1	0.0-2.6	0.0-2.1		
	Sp	0.3 ^a \pm 0.3	0.0 ^{ab} \pm 0.04	0.1 ^{ab*} \pm 0.2	0.2 ^{ab} \pm 0.3	0.0 ^b \pm 0.0	0.0 ^{ab*} \pm 0.0	0.1 ^{ab*} \pm 0.2	27.09	0
	A ($\bar{x} \pm$ SD, range)	0.0-0.8	0.0-0.1	0.0-0.6	0.0-0.8	0.0-0.0	0.0-0.1	0.0-0.6		
Total	Au	6.9 ^{ac} \pm 11.1	1.0 ^{ab} \pm 0.8	7.8 ^c \pm 5.6	4.4 ^{ac} \pm 1.9	0.3 ^{bd} \pm 0.5	7.0 ^{ac} \pm 4.9	4.7 ^{acd} \pm 3.8	27.01	0
	A ($\bar{x} \pm$ SD, range)	0.6-34.1	0.2-2.6	2.6-17.8	2.2-7.5	0.0-1.6	0.3 \pm 13.9	0.0-11.8		
	Sp	4.9 ^a \pm 6.4	0.3 ^{ab*} \pm 0.6	2.3 ^{a*} \pm 1.9	1.1 ^{ab*} \pm 0.9	0.0 ^{bc} \pm 0.1	0.4 ^{ac*} \pm 0.5	1.0 ^{ac*} \pm 1.9	24.36	0
	A ($\bar{x} \pm$ SD, range)	0.0-19.1	0.0-1.7	0.0-6.0	0.3-2.7	0.0-0.4	0.0-1.4	0.0-5.0		
S	Au	27	10	21	17	7	21	17		
	Sp	13	4	10	15	2	6	7		
H'	Au	2.501	1.707	2.433	2.064	1.969	2.243	1.925		
	Sp	1.685	0.726	1.401	2.459	0.562	1.578	1.463		

* – significant differences between autumn and spring; ^{a,b,c,d} – differences between study plots; mean values with the same letter are not significantly different, at $p \leq 0.05$

In the same vineyards of El Poble Nou de Benitatxell as studied here, the abundance of thrips (Thysanoptera), eulophids (Eulophidae), aphelinids (Aphelinidae) and aphidiins (Aphidiinae) did not differ between both types of cultivation (Mesguer Cervera 2014), while the abundance of mesostigmatic mites was higher in conventional vineyards than in organic ones (García-Parra 2015).

In both types of vineyards the density of Oribatida and their species richness varied between three studied habitats (zone between vine rows, zone between vines, border of vineyard), which can be mainly explained by the sensitivity of these mites to the mechanical treatment of soil (Behan-Pelletier 1999). For example tillage reduced the density of Oribatida, but did not affect Mesostigmata, and increased the density of Prostigmata (Winter *et al.* 1990). Also in the vineyards of El Poble Nou de Benitatxell, the cultural practices did not affect the density of Mesostigmata, but their species diversity (García-Parra 2015). These differences between Oribatida and Mesostigmata can be explained by their different mobility; the predatory Mesostigmata move quickly and easily colonize new habitats, while Oribatida are slowly moving, 11-20 cm a day (Berthet and Gerard 1965), being more sensitive than the Mesostigmata to changes in their environment.

The density of Oribatida in the present study was similar to that reported from the vineyards in Italy (3,300-6,900 adult individuals per 1 m²; Nannelli and Simoni 2002), and seems to be typical for these ecosystems in the Mediterranean region. In other geographical regions, much higher densities of Oribatida were noted in vineyards; to a great extent this can be explained by different climatic conditions, especially higher humidity or colder climate. For example, in Japan, where grapes grow in monsoonal climate, with the annual rainfall above 1000 mm, including 800 mm in growing season, the density of Oribatida in vineyards was 57,000 individuals per 1 m² (Suzuki 1979). In Azerbaijan the density of Oribatida varied depending on the soil type between 40,000-422,000 individuals per 1 m² (Samedov *et al.* 1987). Although the wine regions of Azerbaijan are characterized by a low rainfall (250-600 mm), similarly to

Table 5 Mean values \pm SD and range of density (*A*, in 10^3 individuals/m²), dominance (*D*) and constancy (*C*) indices of Oribatida with *D* > 5 in studied plots in El Poble Nou de Benitatxell: Tr – zone between rows driven by a tractor, Vi – zone between vines, Bo – border; SD = standard deviation, H, *p* – the result of Kruskal-Wallis nonparametrical analysis of variance (ANOVA); symbols of species are explained in table 3.

Symbol of species		Autumn							ANOVA rang		
		Control	Conventional			Organic			Kruskal-Wallis	<i>H</i>	<i>p</i>
			Tr	Vi	Bo	Tr	Vi	Bo			
Efla	<i>A</i>	1.4 ^a \pm 2.6 0.0-7.5	0.0 ^a	0.8 ^a \pm 1.6 0.0-4.5	0.1 ^a \pm 0.2 0.0-0.4	0.0 ^a	0.0 ^a \pm 0.0 0.0-0.1	0.0 ^a \pm 0.0 0.0-0.1	11.68	0.07	
	<i>C</i>	50	0	37.5	25	0	12.5	12.5			
	<i>D</i>	19.9	0	9.9	1.8	0	0.2	0.3			
Gtar	<i>A</i>	0.1 ^{ab} \pm 0.1 0.0-0.3	0.0 ^a	1.3 ^b \pm 2.9 0.0-8.4	0.5 ^{ab} \pm 1.1 0.0-3.0	0.0 ^a	1.2 ^b \pm 1.4 0.0-3.9	0.0 ^{ab} \pm 0.0 0.0-0.1	25.68	0	
	<i>C</i>	50	0	87.5	25	0	75	12.5			
	<i>D</i>	1.5	0	16.1	10.7	0	16.8	0.3			
Hins	<i>A</i>	0.0 ^a	0.0 ^a \pm 0.0 0.0-0.1	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a \pm 0.0 0.0-0.1	5.09	0.53	
	<i>C</i>	0	12.5	0	0	0	0	12.5			
	<i>D</i>	0	1.3	0	0	0	0	0.3			
Lbur	<i>A</i>	0.0 ^a \pm 0.0 0.0-0.1	0.0 ^a \pm 0.0 0.0-0.1	0.1 ^a \pm 0.1 0.0-0.3	0.3 ^a \pm 0.4 0.0-1.0	0.0 ^a	0.5 ^a \pm 1.4 0.0-4.0	0.3 ^a \pm 0.6 0.0-1.7	9.54	0.14	
	<i>C</i>	12.5	12.5	37.5	50	0	12.5	37.5			
	<i>D</i>	0.2	1.3	0.8	7.2	0	7.2	5.6			
Mqua	<i>A</i>	0.5 ^a \pm 0.7 0.0-2.1	0.1 ^a \pm 0.1 0.0-0.4	1.0 ^a \pm 0.8 0.0-2.2	1.5 ^a \pm 2.5 0.0-7.2	0.0 ^a \pm 0.0 0.0-0.1	1.7 ^a \pm 1.8 0.0-4.7	1.2 ^a \pm 1.4 0.0-3.4	12.41	0.05	
	<i>C</i>	50	50	87.5	50	12.5	62.5	62.5			
	<i>D</i>	7	9	13.1	34	4.5	24.3	24.6			
Omin	<i>A</i>	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a \pm 0.0 0.0-0.1	0.1 ^a \pm 0.2 0.0-0.4	0.0 ^a	0.0 ^a	5.09	0.53	
	<i>C</i>	0	0	0	12.5	12.5	0	0			
	<i>D</i>	0	0	0	0.3	18.2	0	0			
Osub	<i>A</i>	1.9 ^a \pm 5.3 0.0-15.1	0.0 ^a	0.2 ^a \pm 0.4 0.0-1.0	0.0 ^a	0.0 ^a	0.3 ^a \pm 0.9 0.0-2.5	0.0 ^a	8.07	0.23	
	<i>C</i>	25	0	25	0	0	12.5	0			
	<i>D</i>	27.9	0	2.2	0	0	4.5	0			
Ori1	<i>A</i>	0.1 ^a \pm 0.2 0.0-0.4	0.1 ^a \pm 0.3 0.0-0.9	1.1 ^a \pm 2.0 0.0-4.4	0.0 ^a \pm 0.1 0.0-0.1	0.0 ^a \pm 0.0 0.0-0.1	0.1 ^a \pm 0.2 0.0-0.5	0.4 ^a \pm 0.6 0.0-1.4	5.67	0.46	
	<i>C</i>	37.5	12.5	25	25	12.5	12.5	50			
	<i>D</i>	1.7	11.5	13.6	0.6	4.5	0.9	8.9			
Oexc	<i>A</i>	0.1 ^a \pm 0.2 0.0-0.5	0.3 ^{ab} \pm 0.6 0.0-1.8	1.2 ^{ab} \pm 1.3 0.0-3.4	1.0 ^{ab} \pm 0.8 0.0-2.1	0.1 ^{ab} \pm 0.1 0.0-0.2	1.4 ^b \pm 1.3 0.0-3.7	1.5 ^{ab} \pm 2.3 0.0-6.8	16.52	0.01	
	<i>C</i>	25	87.5	87.5	75	37.5	75	75			
	<i>D</i>	1.1	34.6	14.8	22.4	22.7	19.8	32.7			
Ofri	<i>A</i>	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a \pm 0.0 0.0-0.1	0.0 ^a	6	0.42	
	<i>C</i>	0	0	0	0	0	12.5	0			
	<i>D</i>	0	0	0	0	0	0.19	0			
Pafri	<i>A</i>	0.1 ^a \pm 0.2 0.0-0.4	0.3 ^a \pm 0.4 0.0-0.8	1.0 ^a \pm 1.2 0.0-2.8	0.3 ^a \pm 0.3 0.0-0.9	0.0 ^a \pm 0.1 0.0-0.3	0.5 ^a \pm 0.5 0.0-1.4	0.7 ^a \pm 0.8 0.0-2.0	12.54	0.05	
	<i>C</i>	37.5	62.5	87.5	62.5	12.5	75	62.5			
	<i>D</i>	1.7	30.8	13.1	6.3	13.6	6.6	14.5			
Sspl	<i>A</i>	0.2 ^a \pm 0.3 0.0-1.5	0.1 ^a \pm 0.2 0.0-0.4	0.1 ^a \pm 0.3 0.0-0.8	0.0 ^a \pm 0.1 0.0-0.3	0.1 ^a \pm 0.2 0.0-0.6	0.0 ^a \pm 0.1 0.0-0.1	0.0 ^a \pm 0.0 0.0-0.1	3.77	0.71	
	<i>C</i>	37.5	25	37.5	12.5	12.5	37.5	12.5			
	<i>D</i>	2.3	6.4	1.7	0.9	27.3	0.6	0.3			

Table 5 Continued.

Symbol of species		Spring							ANOVA rang		
		Control	Conventional			Organic			Kruskal-Wallis	H	p
			Tr	Vi	Bo	Tr	Vi	Bo			
Efla	A	0.0 ^{a*}	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0	1	
	C	0	0	0	0	0	0	0			
	D	0	0	0	0	0	0	0			
Gtar	A	0.0 ^a ±0.0 0.0-0.1	0.0 ^a	0.1 ^{a*} ±0.2 0.0-0.4	0.0 ^a	0.0 ^a	0.0 ^{a*} ±0.0 0.0-0.1	0.0 ^a	11.42	0.08	
	C	12.5	0	37.5	0	0	12.5	0			
	D	0.3	0	3.4	0	0	3.6	0			
Hins	A	0.5 ^a ±1.0 0.0-2.4	0.0 ^a	0.0 ^a	0.0 ^a ±0.0 0.0-0.1	0.0 ^a	0.0 ^a	0.1 ^a ±0.4 0.0-1.0	11.5	0.07	
	C	37.5	0	0	12.5	0	0	12.5			
	D	11.1	0	0	1.1	0	0	13.3			
Lbur	A	0.2 ^a ±0.4 0.0-1.3	0.0 ^a ±0.0 0.0-0.1	0.5 ^a ±0.9 0.0-2.6	0.3 ^a ±0.6 0.0-1.3	0.0 ^a	0.2 ^a ±0.3 0.0-0.8	0	10.35	0.11	
	C	50	12.5	37.5	37.5	0	25	0			
	D	5	3.7	24	31	0	46.4	0			
Mqua	A	0.0 ^a ±0.1 0.0-0.1	0.0 ^{a*}	0.0 ^{a*} ±0.1 0.0-0.1	0.0 ^{a*}	0.0 ^a	0.0 ^{a*} ±0.1 0.0-0.2	0.0 ^{a*}	10.99	0.09	
	C	25	0	37.5	0	0	12.5	0			
	D	0.5	0	1.7	0	0	7.1	0			
Omin	A	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0	1	
	C	0	0	0	0	0	0	0			
	D	0	0	0	0	0	0	0			
Osub	A	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0.0 ^a	0	1	
	C	0	0	0	0	0	0	0			
	D	0	0	0	0	0	0	0			
Ori1	A	0.0 ^a	0.0 ^a	0.1 ^a ±0.2 0.0-0.6	0.0 ^a ±0.1 0.0-0.1	0.0 ^a	0.0 ^a	0.2 ^a ±0.5 0.0-1.4	10.86	0.09	
	C	0	0	25	37.5	0	0	25			
	D	0	0	4	3.4	0	0	20			
Oexc	A	0.0 ^a	0.0 ^{a*} ±0.0 0.0-0.1	0.0 ^{a*} ±0.1 0.0-0.1	0.0 ^{a*} ±0.0 0.0-0.1	0.0 ^a ±0.0 0.0-0.1	0.0 ^{a*} ±0.1 0.0-0.3	0.0 ^{a*} ±0.1 0.0-0.2	2.84	0.83	
	C	0	12.5	25	12.5	12.5	12.5	25			
	D	0	3.7	1.1	1.1	25	10.7	4			
Ofri	A	2.8 ^a ±6.2 0.0-18.0	0.0 ^b	0.0 ^b	0.0 ^b ±0.0 0.0-0.1	0.0 ^b	0.0 ^b	0.0 ^b	20.89	0.02	
	C	50	0	0	12.5	0	0	0			
	D	56.2	0	0	1.15	0	0	0			
Pafr	A	0.1 ^{ab} ±0.2 0.0-0.4	0.3 ^{ab} ±0.6 0.0-1.7	1.3 ^a ±1.4 0.0-3.5	0.1 ^{ab} ±0.1 0.0-0.3	0.0 ^b ±0.1 0.0-0.3	0.0 ^{a*} ±0.1 0.0-0.3	0.5 ^{ab} ±1.4 0.0-3.9	15.9	0.01	
	C	50	50	87.5	50	12.5	25	12.5			
	D	2.92	81.5	56.6	9.2	75	14.3	49.3			
Sspl	A	0.3 ^a ±0.5 0.0-1.5	0.0 ^a	0.0 ^a ±0.1 0.0-0.3	0.1 ^a ±0.0 0.0-0.1	0.0 ^a	0.0 ^a	0.0 ^a	11.63	0.07	
	C	37.5	0	12.5	12.5	0	0	0			
	D	6.1	0	1.71	1.15	0	0	0			

those in Spain, in many vineyards irrigation is used, and the average annual temperatures are lower than in Spain (10.5-15.5°C), that seems to provide more favorable conditions for oribatid mites.

In the control the density of Oribatida was similar in both seasons, while in the vineyards it varied, which can be explained by the cultivation practices that took place mainly in the spring, reducing the density of mites.

In the present study 59 species of Oribatida were found, and 57 of them were reported from the vineyards. Similar number of oribatid species was found in Italy (51 species; Nannelli and Simoni 2002) and Azerbaijan (61 species; Samedov *et al.* 1987), but was lower in Japan (24 species; Suzuki 1979) and India (16 species; Acharya and Basu 2014). Agricultural treatments of vineyards reduced the species richness of Oribatida, which is consistent with the conclusions of Meseguer Cervera (2014) and García-Parra (2015) who made similar observations on other arthropod groups in the same vineyards.

In the vineyards, *Oribatula excavata* dominated, followed by *Minunthozetes quadriareatus* and *Passalozetes africanus*. *Oribatula excavata* is a common species in Europe that is considered eurytopic, little sensitive to humidity conditions or organic matter content, and often found in cultivated fields (Pérez-Íñigo 1993) and in meadows (Weigmann 2006). *Minunthozetes quadriareatus* has a Mediterranean occidental distribution and is found in cultivated fields (Mingue *et al.* 1986). It is considered xerophilous, with preferences to cultivated soils (Pérez-Íñigo 1993), what explains its higher abundance in the vineyards than in the control. Similarly, *Passalozetes africanus* is a xerophilous species that is characteristic for very dry Mediterranean environment. It is tolerant of high temperatures and found in the areas without vegetation (Pérez-Íñigo 1993), which explains its highest dominance in the area between the vine rows, driven by a tractor, that was devoid of vegetation.

In the control, most abundant was *Oppiella subpectinata*, followed by *Eremulus flagellifer*. The former species is considered eurytopic, found in various habitats, but preferring higher content of organic matter (Weigmann 2006). This is why it was very abundant in the natural, undisturbed habitat, and in the vineyards was found exclusively in the areas between the vines, where the content of organic matter was higher than in other microhabitats. *Eremulus flagellifer* is a cosmopolitan species that requires a high humidity for development (Pérez-Íñigo 1997). This may explain why it was found only in the autumn, when precipitation was five-fold higher than in the spring.

In the vineyards in Italy (Nannelli and Simoni 2002), the most abundant was *Tectocepheus velatus*, which is eurytopic and easily colonizes new habitats. In the present study it also occurred in most plots, but in low numbers. In the vineyards in Japan, *Scheloribates latipes* was recorded in high number (Suzuki 1979), but in the present study it occurred in low numbers, while more common and more abundant was congeneric *S. barbatulus*. The presence of the species from genera *Scheloribates* and *Protoribates* in the vineyards is interesting, because they possibly play some role in control of pathogen fungi (Nakamura *et al.* 1991; Enami and Nakamura 1996).

A new species for Spain, *Podoribates longipes*, was found only in the control. It has a Holarctic distribution and was found in meadows (Migliorini *et al.* 2003), including salty ones (Weigmann 2006). *Steganacarus boulfekhari* was found only at the border of organic vineyard and has been known exclusively from Algeria, where it was recorded from many localities, mainly under different pine species (*Pinus halepensis* Mill., *P. pinaster* Aiton, *P. canariensis* C. Smith, *P. radiata* D. Don), oaks (*Quercus faginea* Lam.), eucalyptus (*Eucalyptus* L'Hér.), in evergreen bushes with *Erica arborea* L. and *Pistacia* L., and in orchards (Niedbała 2008).

In conclusion we can say that the oribatid mites do not benefit from organic cultivation of vineyards, probably because they are tolerant to the herbicides used in conventional vineyards, but are sensitive to the mechanical works on soil, which are more numerous in organic vineyards than in conventional ones. This study also supports the opinion (e.g. Dufлот *et al.* 2015 and included references) that natural habitats in the agricultural landscape are important zones, increasing its total species diversity.

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