

Microarthropod communities related with biological soil crusts in a desert scrub in northwestern Mexico

Comunidades de microartrópodos relacionadas con costras biológicas de suelo en un matorral desértico en el noroeste de México

Jaramar Villarreal-Rosas¹, José G. Palacios-Vargas^{2⊠} y Yolanda Maya³

¹Universidad Autónoma Metropolitana, Unidad Xochimilco. Calz. del Hueso 1100, Col. Villa Quietud, 04960, México, D. F., México. ²Departamento de Ecología y Recursos Naturales, Facultad de Ciencias, Universidad Nacional Autónoma de México. Av. Universidad 3000, Col. Copilco, Covoacán, 04510, México, D. F.

³Centro de Investigaciones Biológicas del Noroeste. Av. Instituto Politécnico Nacional 195, Col. Playa Palo de Santa Rita Sur, 23096 La Paz, Baja California Sur, México.

🖂 troglolaphysa@hotmail.com

Abstract. In arid ecosystems, biological soil crusts closely interact with microarthropod communities. Together, both communities play one of the most important environmental services: decomposition of organic matter. In a desert scrub in the southern Baja California Peninsula of Mexico, microarthropod communities were correlated to biological soil crusts and the way soil properties influence distribution of the microarthropods. Twenty five soil samples were taken from 3 site types: without crusts (10), with crusts (10), and eroded surfaces (5). Microarthropods were extracted; specimens were identified to family level and feeding groups were identified. Of the 4 682 microarthropods within 40 taxa, Prostigmata had the greatest richness. The lack of plant coverage at eroded sites seems to affect micro-environmental conditions, so that no microarthropods were found at these sites and biological soil crusts were simple in structure. Among desert scrub, biological soil crusts were complex in structure, and edaphic properties were more favorable for microarthropods to thrive. Specific dissimilarities in community structure of microarthropods for each microhabitat were related to feeding preferences of each taxa.

Key words: Acari, Collembola, edaphic properties, microflora.

Resumen. En los ecosistemas áridos, las costras biológicas de suelo junto con las comunidades de microartrópodos dan lugar a uno de los servicios ambientales más importantes: la descomposición de la materia orgánica. La relación entre comunidades de microartrópodos y costras biológicas de suelo, así como la influencia de las propiedades edáficas en la distribución de microartrópodos fueron estudiadas en un matorral desértico en el sur de la península de Baja California. Se tomaron 25 muestras de suelo de 3 sitios: sin costras (10), con costras (10) y superficies erosionadas (5). Se extrajeron los microartrópodos, que fueron identificados hasta familia y se identificaron hábitos alimenticios. Se encontraron 4 682 microartrópodos en 40 taxa, Prostigmata con la mayor riqueza. La falta de cobertura vegetal en los sitios erosionados parece afectar las condiciones micro-ambientales, por lo que no fueron encontrados microartrópodos en estos sitios y las costras biológicas encontradas fueron simples. En el resto del matorral, las costras biológicas eran complejas en su estructura y las propiedades edáficas fueron más favorables para el desarrollo de microartrópodos. Se relacionaron las diferencias específicas dentro de la estructura de la comunidad de microartrópodos para cada microhábitat con los hábitos alimenticios de cada taxa.

Palabras clave: Acari, Collembola, propiedades edáficas, microflora.

Introduction

Over more than 70% of the surface of the world is arid and semiarid soils, where vascular plants are widely dispersed or absent; there, a highly specialized microorganism community of biological soil crusts (BSC), composed of cyanobacteria, algae, microscopic fungi, lichens, and mosses predominates (Belnap, 2001a). These organisms grow inside or on the surface of the upper layer of the soil; during its development, polysaccharides secreted by cyanobacteria and algae, together with filaments of lichens and moss, adhere to soil particles (Coleman et al., 2004). Consequently, BSC stabilize and reduce the susceptibility of soil to water and wind erosion. Furthermore, BSC supply a source of nitrogen for desert soils, which can support germination and establishment of vascular plants (Belnap, 2001a). Flat and rough types of BSC have been identified from a mountainous range of southern Baja California Peninsula (Maya et al., 2002).

According to Neher et al. (2009), there is a close relationship between the porous structure of BSC and the edaphic fauna; this microhabitat of microarthropods also supplies its main source of food. In turn, microarthropods support the bacteria and fungi by supplying detritus (Palacios-Vargas, 1983), as well as disseminating bacteria and fungi spores, and lichen fragments and soredia (Steinberger, 1991). Hence, microarthropods are important regulators of the bacterial and fungal populations of soil. Since microarthropods are relatively sedentary, they reveal soil conditions better than mobile microfauna (Olfert et al., 2002); hence, they can be used as bio-indicators of the health of desert soils (Sandor and Maxim, 2008).

Unfortunately, from the terrestrial ecosystems, soil comprises one of the less studied resources (Coleman et al., 2004), both in its biodiversity and its internal processes. Of the microarthropods only ~10% have been examined and probably only 10% of the species described (André et al., 2002). The basis of distribution of the edaphic fauna and the way it interacts and develops are far from being understood. As a result, the importance of the biota in the soil processes are commonly underestimated (Coleman et al., 2004), regardless of the importance of the services provided to humanity and the rest of the biota. Few studies have been made regarding interactions between biological soil crusts and microarthropod communities. It is remarkable that in North America, studies of microarthropods in arid lands have focused on the Chihuahuan and Mojave Deserts and not the Sonoran Desert. The main objective was to analyze the influence of some edaphic parameters in microarthropod communities related to BSC in a desert scrub.

Material and methods

Study site. The study was conducted at the Northwest Biological Research Center (CIBNOR) reserve, located on an alluvial plain dominated by desert scrub vegetation; the common shrubs include Jatropha cinerea, J. cuneata, Prosopis articulata, Bursera microphylla, Fouquieria diguetti, Cyrtocarpa edulis and the cardon cactus Pachycereus pringlei. Summers are hot and arid, with occasional tropical storms bringing most of the rain. There is only one rainy season, from August to February with 2 peaks in September and January, with the greatest precipitation in summer; winter storms provide <10% of the annual total (León de la Luz et al., 1996).

Biological soil crusts are distributed in patches on the soil surface. Based on appearance, they are designated as "flat" or "rough" crusts; rough crusts contain very conspicuous lichens; in contrast, flat crusts are dominated by cyanobacteria. Flat crusts were found on eroded soils; rough crusts on scrubland soils. Stereoscopic and brightfield microscopy show that filamentous cyanobacteria are dominant in both crusts, most species belonging to *Microcoleus*, as well as many species of the genera *Scytonema* and *Nostoc*, which are nitrogen fixers.

Sampling. Field work was performed in September and October 2011. From a satellite image of 22 July 2009, we identified 25 sites in this area, of which 10 sites had BSC, 10 sites were without BSC, and 5 sites were eroded. To collect microarthropods, a sample of litter and the upper layer of the soil were excavated (together with the crusts, if present). At 10 sites (5 with and 5 without crusts) a soil sample (0-10 cm deep) was collected for analysis. From each sample, a volume of 500 mL of soil (or soil + litter) was placed in a box. At each site, the surrounding vegetation and the type of BSC (flat or rough) was described. The following soil properties were measured: pH, electrical conductivity, total dissolved solids, soluble phosphorus, organic matter, calcium, magnesium, total nitrogen, sodium, potassium, soil texture, bulk density, pore space, temperature, and relative humidity (the 2 last in the field). Microarthropod extraction and identification. Microarthropods were extracted from the soil samples using Berlese-Tullgren funnels (Palacios-Vargas and Mejía, 2007). Voucher specimens of each family were mounted in Hoyer's fluid on glass slides for initial identification and subsequently archived (Palacios-Vargas and Mejía, 2007). Only colembolans (Christiansen and Bellinger, 1998) and mites (Balogh and Balogh, 1988; Kethley, 1990; Walter et al., 2009) were identified to family. Feeding groups were assigned to each Acari family based on the feeding behavior (McDaniel, 1979; Neher et al., 2009; Walter et al., 2009). The groups were: predators (nematode and other microarthropods), phytophages, microphytophages (fungi and algae eaters), and saprophytes.

Data analysis. Kruskall-Wallis analysis was used to determine differences between the edaphic parameters in both conditions ("soil with crusts" and "soil without crusts"). For each condition, the diversity, evenness, and dominance were estimated. For diversity, the Shannon index (H') was used; evenness was obtained by dividing the value of Shannon's diversity index by the logarithm of the number of taxa, and dominance was obtained with the Berger-Parker index (Magurran, 1988). To assess differences in richness between crust types, Whittaker's similarity index was used (Arellano and Halffter, 2003).

Multiple regression tests identified the association between edaphic parameters and abundances of each taxon. For these correlations, only taxa that were present in at least 3 of the sample sites, each with more than 3 individuals were used. Kruskall-Wallis analysis and multiple regression tests were processed with software (Statistica 6.0, StatSoft, 1995). Indexes were obtained with additional software (PAST 2.04; Hammer et al., 2001).

Results

Description of the area. The total annual precipitation in 2009 was 215.9 mm, while a total of 0.15 mm was recorded until the sampling period in September-October 2011, the time of sampling. Dryness contributed to the very low quantity of litter found, around 1 cm thick, when most abundant. The vegetation was leafless and non-flowering. Most of the study area had well developed fleshystemmed scrub, but eroded sites lacked native plant cover, with scattered introduced buffel grass (*Cenchrus ciliaris*) as the dominant species (Fig. 1). Flat BSC was found underneath the buffel grass, but no microarthropods were captured from samples collected at eroded sites. In the remaining study area, where desert scrub prevailed, rough BSC were common (Fig. 2), but less abundant where the plant coverage was dense, probably as a consequence of the reduction of radiation.

Edaphic properties. No significant differences were found regarding average temperature (26.4 °C) and (75.07%) relative humidity at sites with rough crusts. At sites without crusts, the averages were 24.26 °C and 72.85% RH. No significant differences were found between textural classes

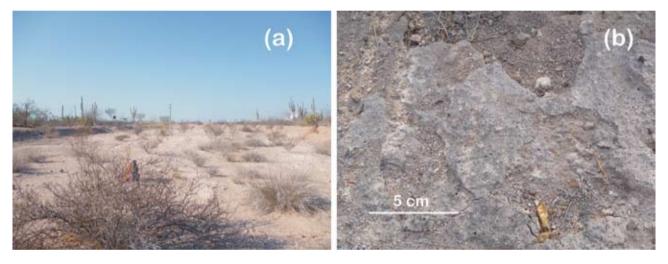


Figure 1. a), eroded site with individuals of *Cenchrus ciliaris*; b), flat biological soil crusts.

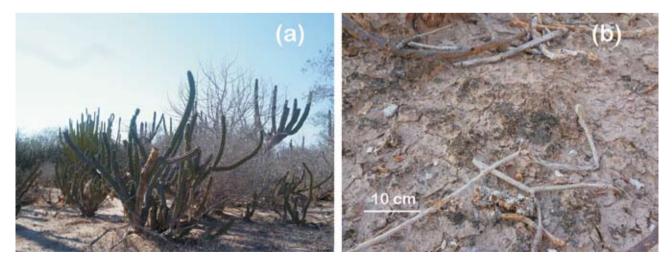


Figure 2. a), site where the scrub is well developed; b), rough biological soil crusts.

at both kinds of sites. Some slight differences occurred in the frequency of soil particle sizes. At sites without crusts, sand was more common than silt and clay, compared with sites with rough crusts (Table 1). At the latter sites, bulk density was less (1.8 g·cm⁻³ vs. 2.0 g·cm⁻³) and pore space was greater (32.1% vs. 22.3%). These results were significantly different (H= 10, df= 1, p= 0.0016). Regarding chemical parameters, nitrogen content was the only parameter that had significant differences between site types, with higher values from sites with rough crusts (0.10% vs. 0.04%, H= 10, df= 1, p= 0.0016). There were no significant differences between the values of OM and pH; however, the results suggest some variance that could be related to the presence of biological crusts (Table 1). *Edaphic fauna*. The 4 628 microarthropods were placed in 40 taxe 23 femiliae belanced to Apari 3 to Collombole

40 taxa, 23 families belonged to Acari, 3 to Collembola, and 12 to a miscellaneous group that contains specimens in Insecta and Aracnida (Table 2). Within Acari, the order with higher richness was Prostigmata with 20 families (1 306 specimens). In Oribatida, there were 4 families (1 020 specimens), and for Acaridae, there were 1 510 specimens in Astigmata; this family represented 32.6% of the specimens.

Among Collembola, the most abundant family was Isotomidae, with 27 specimens of a new species of *Folsomides*; it is remarkable that these specimens belonged to a new species (Palacios-Vargas and Villarreal-Rosas, 2013). It is worth mentioning that the 3 families of Collembola represent only 0.82% in all specimens. Amongst the remaining taxa, the orders with greatest abundance was Psocoptera (382 specimens), and Homoptera (186 specimens) (Table 2).

Regarding diversity, H' was 1.97 for the sites with crusts and 2.40 for the sites without crusts (t = -12.27, df =4537.6, p < 0.05). Diversity was higher at the sites without crusts despite the fact that the number of taxa was 34 in both site types. At sites without crusts, the number of individuals was lower (1 925 vs. 2 703). At sites without crusts, the communities have greater evenness, with values of 0.68 vs. 0.59. The results for the Berger-Parker index strengthen the finding that at the sites with crusts, there are fewer taxa with more individuals in each taxa, with values of 0.40 vs. 0.21. The taxa that may be influencing this result are: Aphelacarus acarinus (588 specimens), Acaridae (1 098 specimens), and Homoptera (158 specimens). The taxa richness per order is similar at sites with crusts and without crusts, but not abundances. Prostigmata was more abundant at sites without crusts (738 vs. 568). Oribatida, Astigmata, Collembola, and the miscellaneous group were more abundant at sites with crusts (Table 2).

Table 1. Physical and chemical properties of soil samples

The similarity value between the communities of both types of sites, as estimated by the Whittaker index was

Sampling site			Edap	Edaphic properties						J	Chemical properties	properti.	es			
	P_{-}	Particle size	ze	Textural classes	BD	Sd	Hd	MO	Ν	Ρ	Ex	Exchangeable cations	ble catio	su	EC	TDS
	Sand	Silt	Clay								Ca^{2+}	Mg^{2+}	Na^+	K^+		
		%			g.cm ⁻³	%		%	%			mg·kg		 1.	dS·m ⁻¹	mg·L ⁻¹
c1	85.5	~	6.5	sand	2.0	24.8	7.0	1.8	0.18	7.5	36.1	26.7	26.4	58.7	0.3	140.9
c2	55.5	28	16.5	sandy loam	1.7	34.5	7.0	2.3	0.09	10.5	36.1	46.2	32.5	52.8	0.3	159.3
c3	67.5	16	16.5	sandy loam	1.9	29.8	7.3	2.0	0.11	3.3	48.1	46.2	29.9	59.3	0.5	216.0
c7	59.5	20	20.5	sandy clay loam	1.7	34.9	7.5	1.6	0.07	6.4	20.0	41.3	21.5	27.7	0.3	139.8
c9	81.5	12	6.5	loamy sand	1.7	36.7	7.4	3.4	0.16	8.7	72.1	53.5	33.2	58.9	0.5	245.0
average	6.99	16.8	13.3		1.8	32.1	7.2	2.2	0.10	7.3	42.5	42.8	28.7	51.5	0.4	180.2
w2	53.5	24	22.5	sandy clay loam	2.1	22.6	7.3	1.3	0.05	6.3	68.1	38.9	23.2	40.9	0.4	191.6
w5	75.5	14	10.5	sandy loam	2.1	21.3	7.4	1.8	0.06	9.7	56.1	36.5	30.7	69.0	0.4	206.0
w8	89.5	3.3	7.2	loamy sand	2.1	21.5	7.4	1.1	0.03	3.7	80.2	34.0	14.5	31.0	0.4	182.0
6m	85.5	7.3	7.2	loamy sand	2.0	23.4	7.4	1.0	0.02	4.1	60.1	29.2	21.0	71.6	0.4	210.0
w10	83.5	8	8.5	loamy sand	2.0	22.9	7.2	1.7	0.03	12.0	28.1	26.7	26.6	54.5	0.3	146.6
average	77.5	11.32	11.18		2.1	22.3	7.3	1.4	0.04	7.2	58.5	33.1	23.1	53.4	0.4	187.2

		With	crust			Withc	out crust		Abundance/ family
	Abd	Density in 339.09 cm ²	Density in $1 m^2$	%	Abd	Density in 339.09 cm ²	Density in 1 m ²	%	
Prostigmata									
¹ Adamystidae	3	0.01	0.26	0.11	13	0.04	1.13	0.67	16
¹ Bdellidae	90	0.27	7.83	3.33	83	0.24	7.22	4.31	173
^{1, 2} Caeculidae	16	0.05	1.39	0.59	11	0.03	0.96	0.57	27
¹ Caligonellidae	13	0.04	1.13	0.48	27	0.08	2.35	1.40	40
^{1, 2} Camerobiidae	1	0.00	0.09	0.04				_	1
¹ Cheyletidae					1	0.00	0.09	0.05	1
^{1,2} Cunaxidae	7	0.02	0.61	0.26	2	0.00	0.09	0.10	9
² Dolichocybidae	4	0.02	0.01	0.20	1	0.01	0.09	0.10	5
	4	0.01							
*Eutrombidiidae			0.09	0.04	1	0.00	0.09	0.05	2
⁴ Linotetranidae	51	0.15	4.44	1.89	13	0.04	1.13	0.67	64
⁴ Nanorchestidae	6	0.02	0.52	0.22		—		—	6
Paratydeidae									
¹ Morphospecies 1	57	0.17	4.96	2.11	234	0.69	20.35	12.15	291
¹ Morphospecies 2	51	0.15	4.44	1.89	32	0.09	2.78	1.66	83
⁴ Penthaleidae				_	1	0.00	0.09	0.05	1
Pyemotidae									
² Pyemotes sp.	1	0.00	0.09	0.04					1
² Pygmephoridae	262	0.77	22.79	9.70	288	0.85	25.05	14.95	550
¹ Smarididae	1	0.00	0.09	0.04	200	0.01	0.17	0.10	3
	1	0.00	0.09	0.04	2	0.01	0.17	0.10	5
Tarsonemidae					1	0.00	0.00	0.05	1
² Tarsonemus sp.	—			_	1	0.00	0.09	0.05	1
Teneriffiidae									
⁴ Parateneriffia sp.	3	0.01	0.26	0.11	—			—	3
⁴ Tetranychidae				_	19	0.06	1.65	0.99	19
¹ Tydeidae	1	0.00	0.09	0.04	9	0.03	0.78	0.47	10
Richness	17				17				S21
Total	568	1.68	49.40	21.03	738	2.18	64.18	38.32	1,306
Oribatida									-,
Aphelacaridae									
² Aphelacarus acarinus	588	1.73	51.14	21.77	292	0.86	25.40	15.16	880
Aphelacarus acarinas	300	1.75	51.14	21.//					
Cymbaeremaeidae	_	_	_	_	1	0.00	0.09	0.05	1
² Scapheremaeus sp.									
² Gymnodamaeidae	—			—	2	0.01	0.17	0.10	2
Haplochthoniidae	40	0.12	3.48	1.48	97	0.29	8.44	5.04	137
² Haplochthonius sp.									
Richness	2				4				S4
Total	628	1.85	54.62	23.25	392	1.16	34.09	20.35	1,020
Astigmata					• / -		-		-,
^{2,3} Acaridae	1,098	3.24	95.49	40.65	412	1.22	35.83	21.39	1,510
Richness	1,070	5.24	JJ. T J	40.05	1	1.22	55.85	21.57	S1
	-	2.24	05 40	10 (5		1.22	25.09	21.20	
Total	1,098	3.24	95.49	40.65	412	1.22	35.98	21.39	1,510
Collembola									
Brachystomellidae	3	0.01	0.26	0.11	6	0.02	0.52	0.31	9
Brachystomella sp.									
Entomobryidae	2	0.01	0.17	0.07	_	_	_	_	2
Isotomidae									
Folsomides sp.	19	0.06	1.65	0.70	8	0.02	0.70	0.42	27
Richness	3	0.00	1.00	0.70	2	0.02	0.70	0.12	S3
Total	24	0.07	2.09	0.89	14	0.04	1.22	0.73	38
Thysanura	13	0.07	1.13	0.89	14	0.04	1.22	0.75	38 29
rnysanura	15	0.04	1.13	0.48	10	0.05	1.39	0.85	29

Table 2. Results of abundance (number of individuals), density (ind area⁻¹), richness (number of taxa), and percentage of edaphic microfauna. Only the feeding habits of the families belonging to Acari were included

		With	crust			Withc	Abundance/ family		
	Abd	Density in 339.09 cm ²	Density in 1 m ²	%	Abd	Density in 339.09 cm ²	Density in 1 m ²	%	
Araneae	1	0.00	0.09	0.04	1	0.00	0.09	0.05	2
Pseudoescorpionida	3	0.01	0.26	0.11	22	0.06	1.91	1.14	25
Coleoptera	10	0.03	0.87	0.37	5	0.01	0.43	0.26	15
Diptera	24	0.07	2.09	0.89	20	0.06	1.74	1.04	44
Embioptera	3	0.01	0.26	0.11	_	_	_	_	3
Hemiptera	_	_		_	2	0.01	0.17	0.10	2
Homoptera	158	0.47	13.74	5.85	28	0.08	2.44	1.45	186
Hymenoptera	3	0.01	0.26	0.11	4	0.01	0.35	0.21	7
Isoptera	7	0.02	0.61	0.26	51	0.15	4.44	2.65	58
Neuroptera	1	0.00	0.09	0.04	_	_	_	_	1
Psocoptera	162	0.48	14.09	6.00	220	0.65	19.13	11.42	382
Richness	11				10				S12
Total	385	1.14	33.48	14.25	369	1.09	32.09	19.16	754
Abundance per site type	2 703				1 925				4 628

Table 2. Continues

0.20, which implies that the communities are very similar, although only in the number of taxa. As stated earlier, both types of communities differ mainly in the number of individuals per taxa. Related to this, the feeding habits of the taxa appear to follow this pattern (Table 3), with the number of taxa in each habit similar, but with differences in abundance.

The families of predators were more abundant at sites without crusts (401 vs. 216). The microphytophages and phytophages were more abundant at sites with crusts (901 vs. 682 and 54 vs. 14, respectively). There were families containing 2 groups, predators and microphytophages, which were more abundant at sites with crusts (24 vs. 13). *Correlation analysis*. Only Linotetranidae, Bdellidae, and Pseudoescorpionida showed a significant correlation with a specific parameter (pore space, phosphorus, and temperature, respectively), when the total abundance (without discrimination between site types) was considered. Aphelacaridae, Linotetranidae, and Homoptera were positively correlated with pH and organic matter (Table 4) when total abundance was taken into account and when

only abundances recorded at the sites with rough crusts were considered. In contrast, they were not positively correlated when only abundance at the sites without crusts were considered.

Organic matter was the parameter with the highest number of positive correlations with abundance recorded at the sites with rough crusts, in which Linotetranidae, Pygmephoridae, Haplochtoniidae, and Homoptera had a p < 0.05. The second parameter with positive correlations was nitrogen content, with the same taxa and also at the sites with rough crusts (Table 4).

Discussion

Description of the sampling sites. During September and October, the scrub is usually phenologically very active (León de la Luz et al., 1996; Maya and Arriaga, 1996). However, this was not the case during the sampling period. The lack of foliage, flowers, and fruits, as well as small quantities of litter was a consequence of a prolonged drought.

Table 3. Richness and abundance of the recorded taxa according to the feeding behavior in which were classified

	Rough	h crusts	Withor	it crusts	Te	otal
	Richness	Abundance	Richness	Abundance	Richness	Abundance
Predators	7	216	8	401	8	617
Microphytophages	6	901	7	682	9	1583
Predators-microphytophages	3	24	2	13	3	37
Microphytophages-saprophytes	1	1098	1	412	1	1510
Phytophages	2	54	2	14	3	68

Table 4. Values obtained for the correlations between the edaphic parameters and the taxa. The following combination of variables was considered: 1), every edaphic parameter with every taxon, considering abundance in both site types; 2), each edaphic parameter with each taxon, setting aside abundance of each site type. No positive correlations were obtained when estimating each edaphic parameter with the total number of taxa.

Taxon	Edaphic		Value	
	parameter			
Without differ	entiation	R	df	р
Bdellidae	Р	0.69	1.8	0.028
Linotetranidae	Bd	0.65	1.8	0.040
	Pe	0.66	1.8	0.037
	OM	0.88	1.8	0.0006
	Ν	0.79	1.8	0.005
	Mg	0.64	1.8	0.042
Aphelacaridae	pĤ	0.68	1.8	0.030
Pseudoescorpionida	Temperature	0.50	1.8	0.027
Homoptera	ОM	0.82	1.8	0.003
	Ν	0.77	1.8	0.009
Rough cr	usts	R	df	р
Linotetranidae	OM	0.97	1.3	0.009
Paratydeidae	Р	0.95	1.3	0.013
morphospecies 1				
Pygmephoridae	OM	0.97	1.3	0.007
	Ν	0.89	1.3	0.038
Aphelacaridae	pН	0.90	1.3	0.038
Haplochthoniidae	ÔM	0.96	1.3	0.008
	Ν	0.90	1.3	0.033
Homoptera	OM	0.92	1.3	0.022
	Ν	0.93	1.3	0.020
	Ca	0.89	1.3	0.048
Without cr	rusts	R	df	р
Linotetranidae	Humidity	0.70	1.8	0.021
Paratydeidae	pH	0.91	1.3	0.033
morphospecies 1				
Acaridae	Clay	0.99	1.3	0.001
Haplochthoniidae	K	0.92	1.3	0.026
Psocoptera	Na	0.94	1.3	0.019
^	Κ	0.94	1.3	0.019
Diptera	EC	0.88	1.3	0.047
	TDS	0.88	1.3	0.049

All the environmental factors may affect the abundance of microarthropods. Arriaga and Maya (2007) describe the importance of large plants as shade providers, a factor that favors fungal activity in the litter of *J. cuneata* and *F. diguetti* and modulate ground temperatures during the day. In summer, temperature could fluctuate on bare ground more than 22 °C. At eroded sites, where the crusts were flat, there was not enough vegetation to provide moderate temperatures; here, microarthropods were not found. This type of crust has less diversity of cyanobacterial morphotypes, characteristics related with earlier successional stages. Furthermore, flat BSC have less organic matter and are found in disturbed areas (Belnap, 2001b).

The differences found between the site types "rough crusts" and "without crusts" regarding the soil properties may be related precisely to the presence or absence of BSC. Broadly speaking, the sites with a rough crust had more favorable soil conditions for microarthropod community development. The fungi hyphae and filamentous cyanobacteria, which are components of a rough crust, interweave the soil particles (Kieft, 1991; Maya et al., 2002); thus a higher nitrogen content derives from cyanobacteria that fix atmospheric nitrogen (Belnap, 2001a; Maya et al., 2002; Rivera Aguilar et al., 2004). Steinberger (1991) states that feces from microfauna may be a source of nitrogen in these crusts. Together, the lower bulk density results from greater pore space and higher N and OM content. Hence, sites with crusts have better soil structure. Although the soil crusts have higher levels of potassium and phosphorus (Rivera-Aguilar et al., 2004), significant differences in these elements were not found, nor in exchangeable cations. The absence of significant differences in these elements may be tied to the OM content as it serves as a substrate for retention of these cations (Knight, 1991).

Edaphic fauna description. At a broad level, there is a similarity among microarthropods in arid lands of North America, Chile, Africa, and Australia (Noble et al., 1996). Our survey found 7 Acari families in common with the 10 reported by Elkins and Whitford (1984). Other taxa were recorded by Santos and Whitford (1983), Cepeda and Whitford (1990), Noble et al. (1996), Kay et al. (1999). Most families collected are associated with arid environments.

A general pattern in arid lands is the dominance of Prostigmata over Oribatida (Cepeda and Whitford, 1990; Noble et al., 1996; Neher et al., 2009). Oribatid mites are commonly referred to as moss mites or beetle mites because they ingest litter and fungi directly and are commonly more abundant in soils rich in organic matter. In soils with little organic matter, as in arid lands, oribatid mites are replaced by prostigmatid mites, which generally feed by drilling and sucking the cytoplasm of plants, fungi, bacteria, or microfaunal cells.

Among Oribatida, Neher et al. (2009) found 23 families; only 4 were found in our study area. This low diversity could be attributed to the scant litter in the scrub, which is the main source of food for the oribatida. A family that seems to be well adapted to the arid conditions is Aphelacaridae, specifically *Aphelacarus* sp., which was abundant or common in several reports (Wallwork et al., 1986; Cepeda and Whitford, 1990; Noble et al., 1996; Neher et al., 2009).

Another common feature in these micro-environments is the low diversity of Collembola, a group that is usually found in moister conditions (Palacios-Vargas, 1991; Hopkin, 1997); however, some families in this group are adapted to arid conditions. One of these are Isotomidae (Suhardjono and Greenslade, 1994). This was the only family found in the Chihuahuan Desert by Neher et al. (2009) and the colembolan family that had the most individuals, of the 3 families of colembolans found in our study.

Most of the mite families recorded in this study were microphytophages and microphytophages-saprophytes, similar to the findings of Neher et al. (2009). Mites with this preference were more abundant at sites with rough crusts. At sites without crusts, predator families were more abundant. This result suggests that the abundance of microarthropods is closely related to the resources provided in the rough crusts, either protective or feeding. Neher et al. (2009) state that species belonging to Aphelacaridae are strict micro-phytophages. This family was the most abundant in their study within the first 10 cm of soil, where the biomass of fungi and algae was higher. In our study, *A. acarinus* and Linotetranidae, which are strict phytophages, were more abundant at the sites with rough crusts.

Among the miscellaneous group, all orders are important in the equilibrium of the soil ecosystem, some of them containing genera that are known to be specific for arid lands. For example, some species of Thysanura survive in habitats with low water content since they absorb atmospheric moisture (McGavin, 2002).

Correlation analysis. In arid ecosystems, where biological processes are mainly controlled by physical and chemical factors more than biological interactions, it is believed that there is less species diversity (Nov-Meir, 1985). Similarly, Schnürer et al. (1985) mentions that the biomass of microorganisms in the soil is proportional to the organic matter content; hence, the mineral soils of arid regions generally support fewer microorganisms. This explains the low microarthropod abundance in the study area in comparison to other environments. Despite the low absolute densities of edaphic microarthropods in arid lands, desert soils can exhibit microarthropod communities that are relatively rich in species, that are, surprisingly well adapted to the stringency of the climate (Wallwork et al., 1984, 1986; Cepeda-Pizzaro and Whitford, 1989). The short length of the body of most of the specimens we captured and their particular shape suggest that some of them are ecomorphic, a phenomenon that implies changes in the morphological and anatomical features adapted to environmental conditions (Palacios-Vargas, 1980). Folsomides californicus (Palacios-Vargas and Villarreal-Rosas, 2013), the new species of collembolan found in

this area, can dehydrate and enter an inactive phase called cryptobiosis. This adaptation is not common in terrestrial invertebrates; however, it has been well described in *Folsomides angularis* (Belgnaoui and Barra, 1989).

Even if there are no positive correlations between edaphic properties and the taxa, Jenny (1980) states that biological activity depends on soil characteristics that positively influence water retention. The latter seems to occur at the sites with rough crusts, where total abundances are higher than at the sites without crusts. Some correlations suggest that the dynamics between edaphic parameters and abundances of taxa are different in each site type, even more so, since there were significant differences between each site type in 3 parameters: nitrogen content, density, and pore space. Indeed, the positive correlations were distinctly different between the 2 types of sites.

The relationship between the quantity of litter and organic matter for abundance and diversity of mites has been established by other authors, mainly in the Chihuahuan Desert (Santos and Whitford, 1983; Kamil et al., 1985; Steinberger and Whitford, 1985; Cepeda and Whitford, 1989; Noble et al., 1996). This relationship is supported in our study by the majority of positive correlations between litter abundance at the sites with rough crusts with the organic matter and nitrogen content. The importance of litter, together with other factors, is confirmed by the absence of microarthropods at eroded sites.

To determine if abiotic factors are influencing the structure of the microarthropod communities or if the feeding habits of the taxa are more important, a long term study covering several years on a monthly basis is recommended. In this way, the trigger effect of rainfall on soil biota could also be assessed.

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Literature cited

- André, H. M., X. Ducarme and P. Lebrun. 2002. Soil biodiversity: myth, reality or conning? Oikos 96:3-24.
- Arellano, L. and G. Halffter. 2003. Gamma diversity: derived from and a determinant of alpha diversity and beta diversity. An analysis of 3 tropical landscapes. Acta Zoológica Mexicana 90:27-76.

- Arriaga, L. and Y. Maya. 2007. Spatial variability in decomposition rates in a desert scrub of northwestern Mexico. Plant Ecology 189:213-225.
- Balogh, J. and P. Balogh. 1988. Oribatid mites of the neotropical region I. *In* The soil mites of the world, Vol. 2, J. Balogh (ed.). Akademiai Kiadó, Budapest. p. 335.
- Belnap, J. 2001a. Microbes and microfauna associated with biological soil crusts. *In* Biological soil crusts: structure, function, and management, J. Belnap and O. L. Lange (eds.). Ecological Studies 150. Springer-Verlag, Berlin. p. 167-176.
- Belnap, J. 2001b. Comparative structure of physical and biological soil crusts. *In* Biological soil crusts: structure, function, and management, J. Belnap and O. L. Lange (eds.). Ecological Studies 150. Springer-Verlag, Berlin. p. 177-192.
- Belgnaoui, S. and J. A. Barra. 1989. Water loss and survival in the anhydrobiotic Collembola *Folsomides angularis* (Insecta). Revue d'Écologie et de Biologie du Sol 26:123-132.
- Cepeda-Pizarro, J. G. and W. G. Whitford. 1989. Species abundance distribution patterns of microarthropods in surface decomposing leaf-litter and mineral soil on a desert atershed. Pedobiologia 33:254-68.
- Cepeda, J. and W. G. Whitford. 1990. Microartrópodos edáficos del desierto Chihuahuense, al norte de México. Folia Entomológica Mexicana 78:257-272.
- Christiansen, K. and P. Bellinger. 1998. The Collembola of North America: north of the Rio Grande, a taxonomic analysis, Part 1. Grinnell College, Grinnell, IA. 1518 p.
- Coleman, D. C., D. A. Crossley Jr. and P. F. Hendrix. 2004. Fundamentals of soil ecology, second edition. Elsevier Academic Press, San Diego. 408 p.
- Elkins, N. Z. and W. G. Whitford. 1984. The effects of high salt concentration on desert soil microarthropod density and diversity. Southwestern Naturalist 29:239-241.
- Hammer, Ø., D. A. T. Harper and P. D. Ryan. 2001. PAST: Paleontological statistics software package for education and data analysis. Palaeontología Electrónica 4, 9. http://palaeoelectronica.org/2001_1/past/issue1_01.htm; last access: 07.III.2012.
- Hopkin, S. P. 1997. Biology of the springtails (Insecta: Collembola). Oxford University Press, New York. 223 p.
- Jenny, H. 1980. The soil resource: origin and behaviour. Springer-Verlag, New York. 377 p.
- Kamil, B. W., Y. Steinberger and W. G. Whitford. 1985. Soil microarthropods from the Chihuahuan Desert of New Mexico. Journal of Zoology (London) 205:273-286.
- Kay, F. R., H. M. Sobhy and W. G. Whitford. 1999. Soil microarthropods as indicators of exposure to environmental stress in Chihuahuan Desert rangelands. Biology and Fertility of Soils 28:121-128.
- Kethley, J. 1990. Acarina: Prostigmata (Actinedida). *In* Soil biology guide, D. L. Dindal (ed.). Wiley-Interscience, New York, p. 667-756.
- Kieft, T. L. 1991. Soil microbiology in reclamation. *In* Semiarid lands and deserts: soil resource and reclamation, J. Skujins (ed.). Marcel Dekker, New York. p. 209-256.

- Knight, W. G. 1991. Chemestry of arid region soils. *In* Semiarid lands and deserts: soil resource and reclamation, J. Skujins (ed.). Marcel Dekker, New York. p. 81-109.
- León de la Luz, J. L., R. Coria and M. Cruz. 1996. Fenología floral de una comunidad árido-tropical de Baja California Sur, México. Acta Botanica Mexicana 35:45-64.
- Magurran, A. E. 1988. Ecological diversity and its measurement. Princeton University Press, Princeton. 179 p.
- Maya, Y. and L. Arriaga. 1996. Litterfall and phenological patterns of the dominant verstory species of a desert scrub community in north-western Mexico. Journal of Arid Environments 34:23-35.
- Maya, Y., A. López-Cortés and A. Soeldner. 2002. Cyanobacterial microbiotic crusts in eroded soils of a tropical dry forest in the Baja California Peninsula, Mexico. Geomicrobiology Journal 19:505-518.
- McDaniel, B. 1979. How to know the mites and ticks. Wm. C. Brown, Dubuque, Iowa. 335 p.
- McGavin, G. C. 2002. Entomología esencial. Ariel, Barcelona. 352 p.
- Neher, D. A., S. A. Lewins, T. R. Weicht and B. J. Darby. 2009. Microarthropod communities associated with biological soil crusts in the Colorado Plateau and Chihuahuan Deserts. Journal of Arid Environments 73:672-677.
- Noble, J. C., W. G. Whitford and M. Kaliszweski. 1996. Soil and litter microarthropod populations from two contrasting ecosystems in semi-arid eastern Australia. Journal of Arid Environments 32:329-346.
- Noy-Meir, I. 1985. Desert ecosystem structure and function. *In* Hot deserts and arid shrublands, A. M. Evenari, I. Noy-Meir and D. W. Goodall (eds.). Elservier, Amsterdam. p. 93-103.
- Olfert, O., G. D. Johnson, S. A. Brandt and G. Thomas. 2002. Use of arthropod diversity and abundance to evaluate cropping systems. Agronomy Journals 94:210-216.
- Palacios-Vargas, J. G. 1980. Problemas taxonómicos en Collembola debidos a la variación intraespecífica. Anales de la Escuela Nacional de Ciencias Biológicas 23:85-95.
- Palacios-Vargas, J. G. 1983. Catálogo de los colémbolos mexicanos. Anales de la Escuela Nacional de Ciencias Biológicas 27:61-76.
- Palacios-Vargas, J. G. 1991. Manuales y guías para el estudio de microartrópodos II: Introducción a los insectos sin alas (Protura, Diplura, Collembola, Thysanura). Departamento de Biología, Universidad Nacional Autónoma de México, Mexico City. 23 p.
- Palacios-Vargas, J. G. and B. E. Mejía. 2007. Técnicas de colecta, montaje y preservación de microartrópodos edáficos. Facultad de Ciencias, Universidad Nacional Autónoma de México, Mexico City. 74 p.
- Palacios-Vargas, J. G. and J. Villarreal-Rosas. 2013. A new Mexican species of *Folsomides* (Collembola: Isotomidae). Revista Mexicana de Biodiversidad 84:480-484.
- Rivera-Aguilar, V., I. M. Cacheux and H. Godínez-Álvarez. 2004. Las costras biológicas del suelo y las zonas áridas. Ciencias 75:24-27.

- Sandor, M. and A. Maxim. 2008. Considerations concerning the role and functions of soil biodiversity in ecosystems. ProEnvironment 1:44-49.
- Santos, P. F. and W. G. Whitford. 1983. Seasonal and spatial variation in the soil microarthropod fauna of the white sands national monument. The Southwestern Naturalist. 28:417-421.
- Schenürer, J., M. Clarholm and T. Rosswall. 1985. Microbial biomass and activity in an agricultural soil with different organic matter contents. Soil Biology and Biochemestry 17:611-618.
- StatSoft. 1995. STATISTICA data analysis software system and computer manual. Version 6.0. StatSoft, Tulsa.
- Steinberger, Y. 1991. Biology of arid region soils: faunal components. *In* Semiarid lands and deserts: soil resource and reclamation, J. Skujins (ed.). Marcel Dekker, New York. p. 173-192.
- Steinberger, Y. and W. G. Whitford. 1985. Microarthropods of a desert tabosa grass (*Hilaria mutica*) swale. American

Midland Naturalist 114:225-234.

- Suhardjono, Y. R. and P. Greenslade. 1994. Folsomides arnoldi n. sp. (Isotomidae): a new collembolan abundant in arid Australia, with a redescription of Folsomides denisi (Womersley). Proceedings of the Linnean Society of New South Wales 114:21-27.
- Wallwork, J. A., B. W. Kamill and W. G. Whitford. 1984. Life styles of desert litter-dwelling microarthropods: a reappraisal based on the reproductive behavior of Cryptostigmatid mites. South African Journal of Science 80:163-169.
- Wallwork, J. A., M. MacQuitty, S. Silva and W. G. Whitford. 1986. Seasonality of some Chihuahuan Desert soil oribatid mites (Acari: Cryptostigmata). Journal of Zoology (London) 208:403-416.
- Walter, D. E., E. E. Lindquist, I. M. Smith, D. R. Cook and G. W. Krantz. 2009. Order rombidiformes. *In* A manual of acarology, third edition, G. W. Krantz and D. E. Walter (eds.). Texas Tech University Press, Lubbock. p. 233-420.