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2 **Earthworm assemblages in different intensity of agricultural uses and their**
3 **relation to edaphic variables**

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19 **Abstract**

20 The objective of this study was to relate earthworm assemblage structure with three
21 different soil use intensities, and to indentify the physical, chemical, and
22 microbiological variables that are associated to the observed differences in earthworm
23 assemblage structure between soils. Three soil uses were evaluated: 1- Fifty year old
24 naturalized grasslands; 2- Cattle-grazing fields converted to feedlot within the two years
25 before the start of this work, and 3- Fifty year old intensive agricultural fields. Three
26 different sites for each soil use were evaluated from winter 2008 through summer 2011.
27 Nine earthworm species were identified across all sampling sites. The sites shared five
28 species: the native *Microscolex dubius*, and the introduced *Aporrectodea caliginosa*, *A.*
29 *rosea*, *Octalasion cyaneum*, and *O. lacteum*, but they differed in their relative
30 abundances according to the system. The results show that earthworm community
31 structure is linked to and modulated by soil properties. Both, species abundance and
32 diversity showed significant differences depending on soil use intensity. A PCA
33 analysis showed that species composition is closely related to the environmental
34 variability. The ratio of native to exotic species was significantly lower in the intensive
35 agricultural system when compared to the other two, lower disturbance Systems.
36 *Microscolex dubius* was shown to be related to the naturalized grasslands and it was
37 associated to Ca, pH, Mechanical Resistance, and to respiration. *Aporrectodea*
38 *caliginosa* was related to high K levels, low enzymatic activity, slightly low pH, and
39 low Ca, and appeared related to the highly disturbed environment. *Eukerria stagnalis*
40 and *Aporrectodea rosea*, commonly found un the cattle-grazing system, were related to
41 high soil humidity, low pH, low Ca and low enzymatic activity. These results show that
42 earthworm assamblages can be good descriptors of different soil use intensities. In
43 particular, *Microscolex dubius*, *Aporrectodea caliginosa*, and *Aporrectodea rosea*,
44 showed different temporal patterns and species associations, due to the changes in soil
45 properties attributable tos oil use intensity.

46
47 **Keywords:** soil ecology, soil use intensity, soil biota

48 **1. Introduction**

49 The soil biota plays a crucial role in regulating processes like water infiltration and
50 storage, decomposition and nutrient cycling, humus formation, nutrient transformation
51 and transport; moreover, they stimulate the symbiotic activity in the soil, improve the
52 organic matter storage, and prevent erosion (Lavelle et al., 2006; Coleman and
53 Crossley, 1996).

54 Several of the ecosystem services of the soil depend on the community of soil
55 invertebrates (Lavelle et al., 2006), being earthworms one of the most common
56 component of edaphic communities. Earthworms are considered ecosystem engineers
57 because they improve decomposition processes and nutrient cycling (Lavelle et al,
58 1997; Six et al, 2004) and have a strong effect on the soils' hydraulic properties (Lee,
59 1985; Edwards and Bohlen, 1996; Lavelle and Spain, 2001; Lavelle et al., 2006;
60 Johnson-Maynard, Umiker, and Guy, 2007; Jouquet et al, 2008).

61 The most important factors limiting earthworm populations are food supply, moisture,
62 temperature, and the texture and soil chemical characteristics such as pH, organic matter
63 and macronutrients content (Satchell, 1967; Lee, 1985; Curry, 2004). Earthworm
64 populations are also affected by the direct and indirect effects related to the type and
65 extension of the vegetation cover (Mather and Christensen, 1988; Falco and Momo,
66 1995). The use of soils in agriculture can modify the physical and chemical soil
67 environment thus modulating changes in abundance and composition of earthworm
68 communities (Curry, Byrne and Schmidt, 2002). In this regard, Dale and Polasky (2007)
69 indicate that in agricultural systems, changes in land cover are the direct result of
70 management practices. Therefore, when changes in soil use occur due to different
71 agricultural practices, earthworms' assemblages rapidly respond to them (Lavelle et al,
72 1997; Johnson-Maynard, Umiker, and Guy, 2007).

73 Since earthworm abundance and distribution are strongly influenced by the
74 environmental conditions and the ecological status of the system (Falco and Momo,
75 2010), earthworm community structure can be successfully used as biological indicators
76 of soil conditions (Momo, Falco, Craig, 2003).

77 The use of bioindicators has the advantage of providing historical and functional
78 information about soils. Earthworm community structure integrate this information on
79 soil conditions both in space and time and provide signals of the soil ecological state.

80 In this context, the objectives of this study were: 1) to assess earthworm community
81 composition under three different soil use intensities: intensive agriculture, cattle
82 grazing, and naturalized grasslands. 2) To identify the physical, chemical, and
83 microbiological variables related to the observed community structure. 3) To detect
84 which earthworms species are typical of each set of soil conditions and of each use.

85

86 **2. Materials and Methods**

87 **2.1 Sampling sites**

88 This study was performed in the rolling pampas within the Argentine pampas, a wide
89 plain with more than 52 million hectares (520.000 km²) of land suitable for cattle rising
90 and cropping (Viglizzo et al, 2004). It is one of the largest and most productive
91 agricultural regions in the world.

92 Agricultural systems based on crop–crop and crop–pasture rotations under grazing
93 conditions have been very common in the region for over a century until the 1980s. The
94 adoption of conservative tillage and no-till practices has significantly increased during
95 the 1980s and 1990s. Although pesticides were extensively used since the 1960s, crops
96 and pasture fertilization increased noticeably only during the 1990s (Viglizzo et al,
97 2003). The expansion of the land area used for annual crops on different environments
98 means that the pampean ecosystem is currently under an intense disturbance regime
99 (Navarrete et al, 2009).

100 The selected study sites are located in central Argentina, on argiudol soils, (Mollisols,
101 Typical argiudols (USDA, 2010)). The study sites were privately owned fields located
102 in Navarro, Buenos Aires Province (34° 49' 35" S, 59° 10' 38" W), and Chivilcoy (35°
103 03' 10" S; 59° 41' 08" W) approximately 75 and 150 km west of Buenos Aires City,
104 respectively.

105 Weather regime in this region is temperate humid, with an average annual rainfall
106 around 1000 mm. The mean annual temperature is 17 °C. Phytogeographically, it is
107 within the neotropical region, oriental district of the Pampean Province, and the
108 dominant vegetation is a gramineous steppe (Cabrera and Willink, 1973).

109

110 **2.2 Land use intensity in the selected sites**

111 The systems analyzed differed only in their use intensity. Samplings were carried out on
112 three different type of soil uses (agroecosystems) which represent three different levels
113 of disturbance of the same argiudol soil:

114 Agroecosystem 1: agricultural systems, sites with 50 years of continuous intensive
115 agricultural practices, and under no-tillage during the last 16 years. Under a regular
116 corn-wheat-soybean rotation, currently under no-tillage, chemical weed control is used.
117 During the cropping season, heavy machinery is used and insecticides, herbicides, and
118 fertilizers are applied several times a year.

119 Agroecosystem 2: Cattle-grazing systems, sites with 40 years under direct grazing,
120 turned to a feedlot system within the two years before the start of this work. Originally
121 managed under grazing with high instantaneous animal load per hectare, it moved to
122 bale production (oat, maize, and sorghum) two years prior to the start of this study.

123 Agroecosystem 3: Naturalized grasslands, sites with no significant anthropic impact
124 during the last 50 years.

125 Nine sampling fields (three replicates for each one of the three treatments) were
126 evaluated. At each site, five samples were taken every three months covering a two year
127 period.

128 Each sampling date, five random samples were taken 25 meters apart from each other
129 per each replicate (3) and treatment (3). Thus, a total of 45 samples were taken per
130 sampling date. The size of each sample was of 25 x 25 x 25 cm.

131 The measured environmental variables were bulk density (BD), mechanical resistance
132 (MR), humidity (RH), electrical conductivity (EC), organic mater (OM), pH, N, P, Ca,
133 Mg, K, and Na. To characterize the sites, microbiological activity was assessed through
134 soil respiration and free nitrogen-fixing bacteria activity (Nitrogenase Acetylene
135 Reduction Activity, ARA). Methods used for chemical and physical analyses are shown
136 in Table 1.

137 Earthworm extraction from the soil samples was performed by handsorting. Earthworms
138 were preserved with soil until identification in the laboratory, and later fixed and
139 preserved in alcohol- formalin - glycerin following Righi (1979) and identified by
140 external morphology using keys from Righi (1979) and Reynolds (Reynolds, 1996).
141 Clitelated individuals were identified to species level and pre-clitelated ones to genus.

142 At each site, earthworm taxonomic composition and population density were measured.
143 Earthworm communities were characterized at each soil use intensity by population
144 density, species richness, both observed and estimated (Chao index), and diversity
145 (Shannon).

146

147 **2.3 Statistical analyses**

148 Due to non-normal distributions of the physical and chemical data, Kruskal-Wallis
149 ANOVA tests were carried out to compare variables between treatments.

150 The relationship between environmental variables and earthworm species abundances
151 was further analyzed at the genus level to include non-clitellated individuals, by means
152 of a principal component analysis (PCA) using abundances. Prior to analysis, the
153 species abundances data were transformed using the Hellinger method of Legendre &
154 Gallagher (2001) such that the resulting PCA represents Hellinger distance between
155 samples rather than Euclidean distance. Physical and chemical variables were then fitted
156 into the ordination space described by the first two principal components of the
157 earthworm data by projecting biplot vectors. The statistical significance of the
158 environmental variables is based on random permutations of the data and P-values were
159 adjusted by a sequential multiple test procedure Hommel (1988). The ordination
160 analysis and vector fitting were produced using the R statistical language (R Core team,
161 2012) and the Vegan package (Oksanen et al, 2011).

162

163 **3. Results**

164

165 **3.1 Physical and chemical soil parameters**

166 Of all the physical - chemical and microbiological parameters evaluated, only four
167 variables (Na, EC, MR, and respiration) showed significant statistical differences
168 between each of the three systems and only OM presented no differences (Table 1).
169 From the four variables that separate the three systems, the naturalized grasslands
170 showed the highest Na and EC values.

171 Microbiological activity and soil microfauna were assessed through soil respiration and
172 nitrogen fixing bacteria activity, which separated the naturalized grasslands for their
173 high value when compared to the other two agroecosystems.

174 3.2 Earthworm assemblage response to soil use intensity

175 Results show that each soil use presents a different species composition and abundance
176 (Fig 1). The relative abundances of the earthworm species found in each system is
177 shown in figure 2.

178 A total of 9 earthworm species were identified across all systems. Five species were
179 common to all of them: the native *Microscoclex dubius* and the exotic *Aporrectodea*
180 *caliginosa*, *A. rosea*; *Octolasion cyaneum*, *O. lacteum*, but differed in their abundances.
181 The differences in abundance explain the significant differences found for the Shannon
182 index values (ANOVA test $p < 0.05$). The richness estimate (Chao) and the observed
183 richness only differed in the cattle grazing system (Table 2).

184 In the naturalized grasslands the species identified as being the dominant (44% of all
185 the individuals collected) was the epigeic native *Microscoclex dubius*, followed by the
186 endogeic exotics *Aporrectodea caliginosa*, *A. rosea*, *Octolasion cyaneum*, and *O.*
187 *lacteum*. The other endogeic species, *A. trapezoides*, and the native *M. phosphoreus*
188 were less frequent (Fig. 2a). Forty seven percent of all the individuals collected
189 belonged to native species, and the ratio natives / exotics was 1:2.5.

190 In the cattle-grazing system the endogeic native *Eukerria satgnalis* was dominant and
191 the exotic *A. rosea* was also common. Other species that were present albeit with a low
192 frequency were *A. caliginosa*, and *Microscoclex dubius*. *M. phosphoreus*; *Octolasion*
193 *cyaneum*, and *O. lacteum* appeared on either one or two sampling dates only. In this
194 system, *E. stagnalis* represents 68% of all the individuals collected and *A. rosea*
195 represents 22% (Fig. 2b). The ratio of native species / exotic was 1: 1.3.

196 In the agricultural system, the most common species were the endogeic exotics
197 *Aporrectodea caliginosa*, *A. rosea*, and *A. trapezoides*. The other endogeic species
198 *Octolasion cyaneum*, and the epigeic native *Microscoclex dubius* were less frequent.
199 *Octolasion tyrtaeum* was only detected in the first sampling date, and *O. lacteum*
200 appears in two sampling dates with a single individual each. Here, the exotic species
201 represent 95% of the individuals (Fig. 2c). The agricultural system also had the lowest
202 ratio of native species /exotic (1:6).

203 The differences in the chemical and physical soil parameters, as well as the different
204 temporal distribution and species requirements determined the species' co-occurrences
205 found in each system. We observed these associations involving both native and

206 introduced species, and combining different ecological categories. In this way the
207 associations most frequently found in naturalized grasslands were: *A. rosea* –
208 *Microscoclex dubius* (appearing together in 33% of the samples), *Octolasion cyaneum* –
209 *O. lacteum* (10%), and *A. rosea* –*Octolasion cyaneum* (10%). In the cattle grazing sites
210 *A. rosea* –*Eukerria stagnalis* (67%) and in the agricultural system the most common
211 associations were *A. caliginosa* –*A. rosea* (12.5%), and *A. rosea* – *Microscoclex dubius*
212 (12,5%).

213 The relationship between the characteristics of the environment and earthworm
214 presence was further analyzed at the genus level, assessing the sensitivity of the groups
215 with the soil parameter values through a Mann-Whitney U-test (Table 3).

216 *Aporrectodea*, *Octolasion* and *Microscoclex* were present in samplings with the same
217 levels of Mg, K, and BD. *Octolasion* separated from *Aporrectodea* only for Ca levels,
218 and its response to soil humidity, MR, and Respiration put it close to *Microscoclex*. In
219 turn, *Microscoclex* differed from the other groups due to Na, pH, ARA, and high MR
220 (RM 10 cm). On the other hand, *Eukerria* was related to places with low levels of Ca,
221 K, pH, EC, ARA, BD, MR and high humidity.

222 In order to know how the species' composition explain the environmental variability, an
223 indirect ordination PCA analysis was used, followed by a vector fitting (Fig. 3).

224 Interestingly, the analysis showed no relationship between species with fertility levels
225 (N, P and OM), but it did with elements of low soil mobility.

226 The first two axes explain 57% of the variance. The environmental variables that were
227 significantly related to the species ordination were: RH, K, ARA, Respiration, MR, Ca,
228 and pH (adjusted $P < 0.05$).

229 As it can be seen in Fig. 3, the ordination method shows that *Microscoclex dubius*
230 appeared related to the levels of Ca, pH, MR and respiration. This species is well
231 adapted to environments rich in Ca, neutral pH, high microbiological activity, and high
232 mechanical resistance. The environment defined by *Microscoclex dubius* was related to
233 the characteristics of the Naturalized Grassland system, and this species can be
234 considered as indicative of the conditions prevailing in this system. In the same way
235 *Aporrectodea caliginosa* (Fig. 3) is related to high K levels, low enzymatic activity, low
236 pH, and low Ca. These are characteristic of the Agricultural system, being this
237 cosmopolite, invasive species a good indicator of high perturbation sites. Finally,

238 *Eukerria stagnalis* and *Aporrectodea rosea*, were related to the second ordination
239 factor, and they describe an environment with high soil humidity, low pH, low Ca
240 levels, and low ARA. These characteristics describe the Naturalized Grassland system.
241 In this way, these species are clearly good descriptors of the three studied use intensity
242 regimes of the same soil.

243

244 **4. Discussion**

245 These results show that the structure of the earthworm assemblage changes in relation to
246 differences in soil use intensity in terms of its composition, abundance, seasonal
247 dynamics and species associations. The data presented here show that, on the same soil
248 and the same regimen of temperature and precipitations, the earthworm assemblage
249 composition and abundance varied across the different systems studied, thus reflecting
250 the differences due to land use intensities and their associated management practices.
251 Tillage, weed control, fertilization and soil cover are parameters that best characterize
252 the different land use intensities (Decaëns et al, 2008; Viglizzo et al, 2004; Curry,
253 2004), modifying the physical (water and air movement) and chemical environment,
254 thus changing habitat suitability.

255 In the AG system under highest use intensity, earthworm communities were affected
256 directly by the changes caused by tillage practices or indirectly through changes in food
257 supply. Several studies indicate that earthworm communities are more abundant and
258 rich in species in undisturbed soils when compared to cropland (Feijoo et al, 2011;
259 Felten and Emmerling, 2011; Emmerling 2001; Curry et al, 2008; Decaëns et al, 2008).
260 In this study, however, this pattern was not observed. All three systems have the same
261 richness value and the abundances are consistently higher in the AG system with
262 highest use intensity. This system also showed the highest native species replacement
263 by exotic ones (ratio 1:1.6). These results agree with those of Lee (1985), Paoletti
264 (1999), and Smith et al. (2008), who found that annual croplands have higher
265 earthworm abundance than older fields. The dominance of introduced species is another
266 characteristic of highly disturbed sites, as pointed out by Fragoso et al. (1999),
267 Winsome (2006), and Chan and Barchia (2007).

268 The results presented in this work indicate that earthworm assemblage response to the
269 same soil subjected to different use intensities can be used as indicator of

270 agroecosystem soil use intensity. This response can be explained in terms of quality and
271 quantity of food (Bohlen et al, 1997), the long term use of inorganic fertilizers which
272 have a positive effect on the total number of worms (Edwards and Bohlen, 1996; Curry,
273 2004), pH changes and the level of Ca in the soil (Lee, 1985; Paoletti, 1999; Smith et al,
274 2008).

275 In the agricultural and cattle grazing systems, microbiological activity was low (as
276 assessed through respiration and ARA) when compared to the naturalized grasslands.
277 This can be explained, as the result of a reduction in pH and Ca, as well as to the
278 ecological categories of species present (Scheu et al, 2002). Indeed, Scheu (2003)
279 indicates that the presence of endogean species significantly reduces bacterial biomass
280 and the functioning of the microbial assemblage. In AG, 95% of the species present are
281 exotic endogean, while in the CG system the 97% are endogean (70 % native, 30%
282 exotic).

283 These results show that the ecological categories of the earthworm assemblages are also
284 related to the microbiological activity of each studied system, being another indication
285 that earthworms are good descriptors of the functioning of the edaphic environment.

286 Soil use intensity is also indicated by the presence of a few species that closely related
287 to environmental variability. The intensification of the agricultural activities in the
288 Pampas determine up to a 50% reduction in the calcium level (Casas, 2005). The
289 ordination analysis related *Microscolex dubius* with high Ca levels and thus, to less
290 disturbed environments. On the other hand, *A. caliginosa* and *Eukerria stagnalis* are
291 present in low Ca soils.

292 In this sense, Mele and Carter (Mele and Carter, 1999) point out that the distribution
293 and number of native species are negatively correlated with P, K, and Mg levels, these
294 species being adapted to lower nutrient levels. In our study the only species that is
295 related to higher K levels is *A. caliginosa*, which is the most abundant in the agricultural
296 system.

297

298 **5. Conclusions**

299

300 The richness, composition and abundance as well as the species associations found,
301 reflected the physical, chemical, and biological changes, brought about as a result of the

302 different intensities of the agricultural practices used on each tested system. The data
303 gathered indicate that the different environments are well characterized by the levels of
304 cations (Ca, K), pH, microbiological activity, and physical variables such as mechanical
305 resistance and moisture. Earthworm species assemblage reflected the changes in these
306 variables and are therefore good descriptors of the studied systems.

307 *Microscolex dubius* was associated to sites with high levels of calcium, microbiological
308 activity and high mechanical resistance and describes the naturalized grassland.

309 *Eukerria stagnalis* is primarily associated with high humidity as seen in the cattle
310 grazing system in which it is the dominant species. *Aporrectodea caliginosa* is
311 associated to highly disturbed environments, with high K levels, low CE and NA, and
312 low microbiological activity, all typical of the Agricultural system. It is interesting to
313 note that the earthworm species most related to the different systems, are not related to
314 the variables most usually measured: OM, N, and P. Therefore, monitoring these
315 species would provide indirect estimations of those scarcely measured variables, thus
316 complementing the information provided by other more common soil analyses in
317 agroecosystems.

318 *Eukerria stagnalis* is indicative the high humidity, increased soil acidity, and a
319 reduction in the levels of calcium and potassium, which are conditions prevalent in the
320 intermediate use intensity system.

321 *Aporrectodea caliginosa* is the species best adapted to the most disturbed environment.
322 This implies that the population recovers quickly after a disturbance (Curry, 2004,
323 Felten and Emmerling, 2011; Decaëns, 2011), it is not significantly affected by changes
324 in litter quality (Curry and Schmidt, 2007).

325 The spatial and temporal patterns in the distribution and abundance of earthworm
326 species observed in this work followed the differences in the physical and chemical
327 variables measured on the different systems studied. These differences are, in turn, a
328 reflection of the different management practices applied to the same argiudol soil.
329 Therefore, these results show that the structure of the earthworm assemblages can be
330 reliably used for monitoring different soil use intensity management practices.

331

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333

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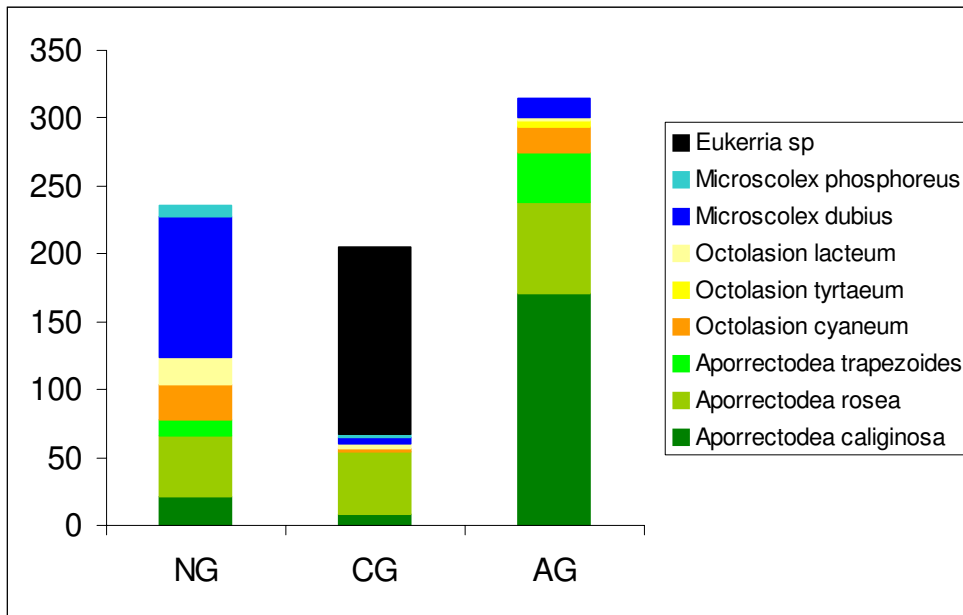


Figure 1: Abundance (N) of each earthworm species throughout the total sampling period for each system.

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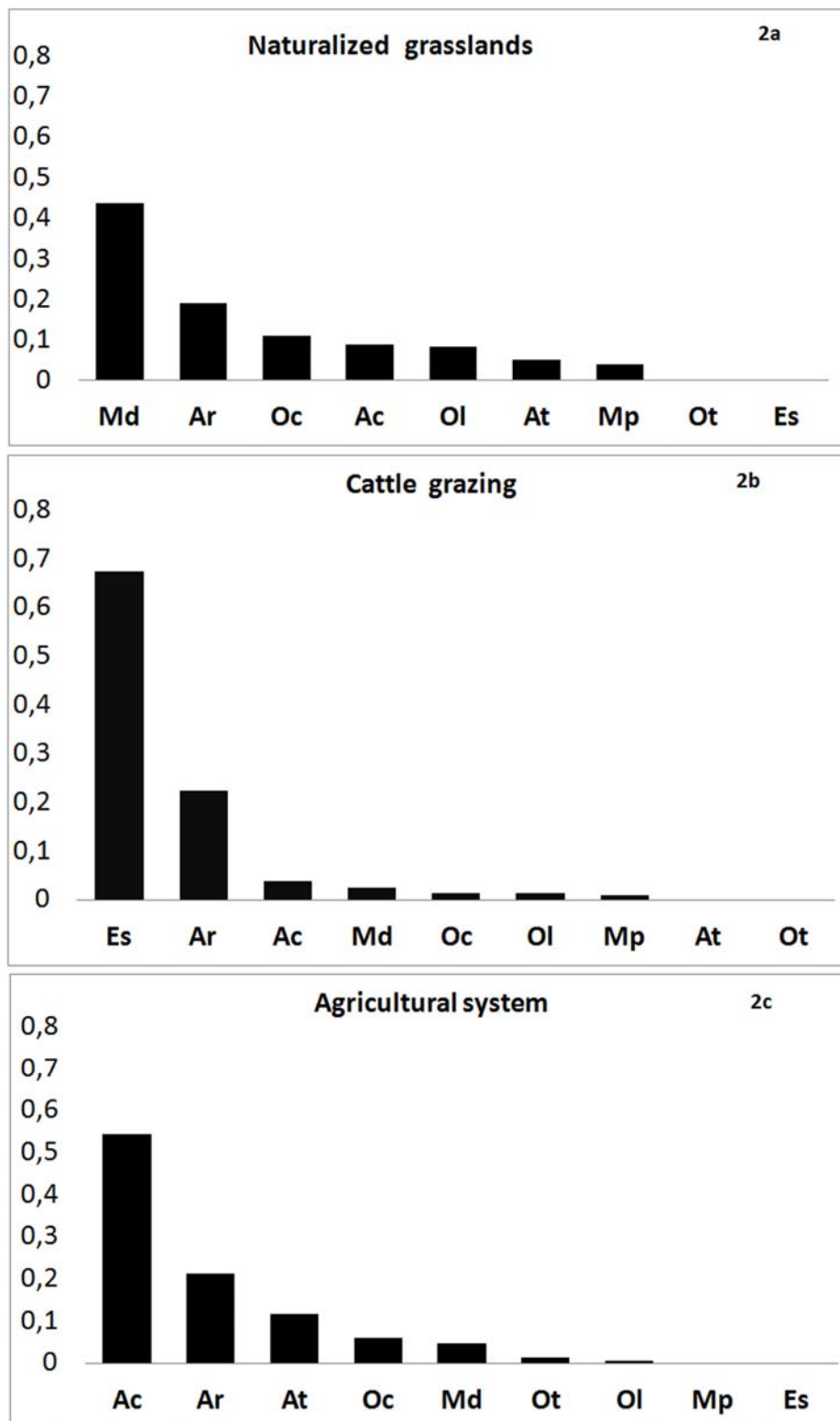
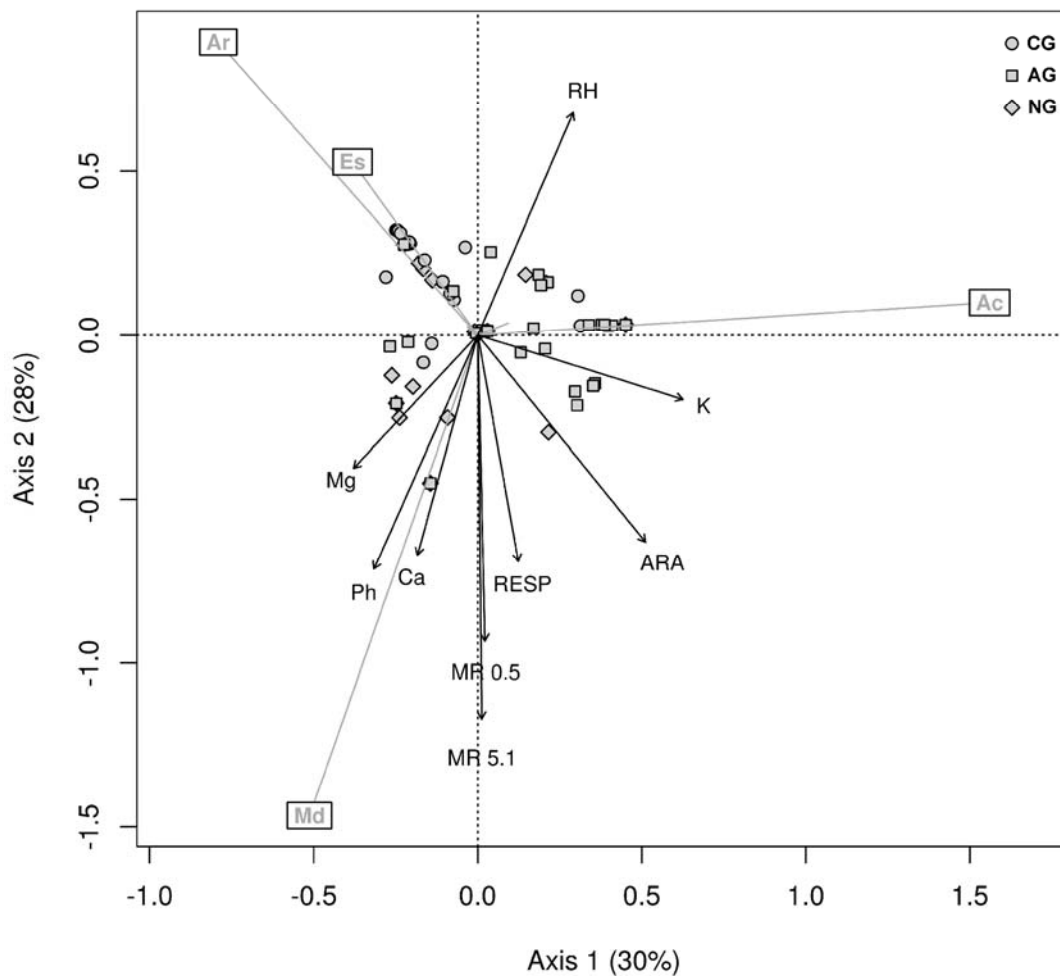


Figure 2: Relative abundance (ni/N) for each earthworm species identified in the three systems. 2a Naturalized grassland (NG), 2b Cattle grazing (CG), 2c Agricultural system (AG). Ac :*Aporrectodea caliginosa*, Ar: *Aporrectodea rosea*; At: *Aporrectodea trapezoides*; Ot: *Octolasion tyrtaeum*; Md: *Microscolex dubius*; Oc: *Octolasion cyaneum*; OI: *Octolasion lactuem*; Es: *Eukerria stagnalis*; Mp: *Microscolex phosphoreus*.

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Figure 3: PCA biplot of Hellinger transformed earthworm species, only the four most abundant ones are shown. The arrows are significant environmental variables fitted into the ordination space. The percentage of explained variance is shown in each axis. Ac: *Aporrectodea caliginosa*; Ar: *Aporrectodea rosea*; Es: *Eukerria stagnalis*; Md: *Microscolex dubius*.

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Table 1: Physicochemical and microbiological parameters measured (n=150 per system)

Parameters	Method	System					
		NG		CG		AG	
P (ppm)	Kurtz y Bray	11+/- 8,5	b	15 +/- 12	a	14+/- 12	a
OM (%)	Walkey-Black	4 +/- 1,5	a	4 +/- 1,5	a	4 +/- 1,4	a
Ec (dS*m ⁻¹)	conductivimeter	1,5+/- 1,3	a	0,8 +/- 0,5	b	0,7+/- 0,5	c
pH		7,5+/- 1	a	6 +/- 0,6	b	6 +/- 0,5	b
Bulk density (gr*cm ⁻³)	Porta	1,2+/-0,2	a	1,1 +/- 0,1	b	1,2+/- 0,1	a
Rh (%)	calculation	0,2+/- 0,1	a	0,3 +/- 0,1	b	0,2+/- 0,1	a
Ca (cmol*Kg soil ⁻¹)	titration with EDTA	6,7 +/-1,3	a	5 +/- 0,5	b	6 +/- 0,7	a
Mg (cmol*Kg soil ⁻¹)	titration with EDTA	1,8+/- 0,4	a	1,5+/- 0,7	b	1,6+/- 0,5	b
Na (cmol*Kg soil ⁻¹)	flame photometry	1,3+/- 0,5	a	0,8 +/- 0,2	b	0,7 +/- 0,2	c
K (cmol*Kg soil ⁻¹)	flame photometry	1,6 +/- 0,5	a	1,3+/-0,3	b	1,6+/- 0,5	a
N (%)	Kjeldahl	0,28 +/- 0,1	a	0,32+/- 0,1	b	0,29+/-0,05	b
Nitrogenase activity (nanolitres of ethylene* gr dry soil*incubation hour ⁻¹)	ARA	0,3+/- 0,3	a	0,2+/-0,2	b	0,2+/-0,3	b
Respiration (mg de CO ₂ *gr dry soil day ⁻¹)	alkaline incubation	0,09+/-0,06	a	0,07+/-0,05	b	0,05+/-0,05	c
MR 0-5 (Kg*cm ⁻²)	cone	10+/- 6	a	2,5+/- 3	b	5,5 +/- 4	c
MR 5=10 (Kg*cm ⁻²)	cone	13+/-7	a	5 +/-5	b	8+/- 5	c

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NG: naturalized grassland; CG: cattle grazing, AG; agricultural system. Different letters within a row indicate significant differences between systems, P < 0.05 Kruskal-Wallis ANOVA tests.

499 **Table 2:** Observed and estimated species richness, mean density, and Shannon
500 diversity index.
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	Richness observed (Sob)	Richness estimate (Chao)	Density (ind/m ²)	Shannon index
Naturalized grassland	7	7 +/- 0	46 +/- 19	0,53 a
Cattle - grazing	7	8,5 +/- 1,5	40 +/- 55	0,37 b
Agricultural system	7	7,25 +/- 0,4	76 +/- 56	0,57 a

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504 **Table 3:** Mean (range) values of each measured variable as they relate to earthworm
 505 genus presence (Non-clitellated specimens included)

Parameters	Genus			
	<i>Aporrectodea</i>	<i>Octolasion</i>	<i>Microscolex</i>	<i>Eukerria</i>
OM (%)	4,4 (3,7-5,3)	4,8 (4,4-6,1)	4,9 (3,2-5,9)	4,7 (4-5,9)
N (%)	0,29 (0,26-0,33)	0,29 (0,26-,33)	0,29 (0,27-0,34)	0,29 (0,25-0,34)
P (ppm)	8,7 (4,4-17,6)	7,7 (3,6-15,2)	9,3 (4,8-15)	6,8 (4,4-14,1)
Ca (cmol*Kg soil ⁻¹)	6,0 (5,5-6,4) a	6,6 (6,1-9) b	6,1 (5,8-7) c	5 (4,6-5,4) d
Mg (cmol*Kg soil ⁻¹)	1,7 (1,1-2) a	1,7 (1,1-1,9) a	1,6 (1,5-1,9) a	1,1 (1-1,6) b
Na (cmol*Kg soil ⁻¹)	0,8 (0,7-1) a	0,74 (0,4-0,9) a	0,9 (0,7-1,1) b	0,8 (0,7-1,1) a
K (cmol*Kg soil ⁻¹)	1,3 (1,1-1,7) a	1,5 (1,2-1,8) a	1,3 (1,1-1,8) a	1,1 (1-1,4) b
pH	6,2 (5,8-7) a	6,3 (6-6,8) a	6,8 (6,2-7,2) b	6 (5,6-6,5) c
Ec (dS*m ⁻¹)	0,6 (0,3-0,9) a	0,6 (0,3-0,9) a	0,7 (0,3-1,2) a	0,4 (0,2-0,7) b
Nitrogenase activity (nanolitres of ethylene* gr dry soil*incubation hour ⁻¹)	0,15 (0,07-0,3) a	0,26 (0,11-0,35) a	0,27 (0,14-0,37) b	0,15 (0,12-0,18) a
Respiration (mg de CO ₂ *gr dry soil day ⁻¹)	0,04 (0,03-0,09) a	0,04 (0,03-0,09) ab	0,07 (0,04-0,1) b	0,05 (0,02-0,07) a
Rh (%)	0,3 (0,2-0,3) a	0,25 (0,17-0,29) ab	0,2 (0,2-0,3) b	0,3 (0,3-0,4) c
Bulk density (gr*cm ⁻³)	1,2 (1,1-1,3) a	1,21 (1,1-1,3) a	1,2 (1,1-1,3) a	1,1 (1-1,2) b
MR 0-5 (Kg/cm ²)	4,6 (2,25-8,2) a	4,9 (3-8) ab	7,8 (4,3-12,5) b	0,78 (0-3) c
MR 5=10 (Kg/cm ²)	6,5 (3,5-10,8) a	7,6 (4-11,5) a	10 (7-17) b	2,6 (0,8-5,5) c

506 Variables measured at the sampling points were each earthworm genus was recorded.
 507 Different letters within each row indicate significant differences between earthworm
 508 genus, P < 0,05, Mann–Whitney U-test pairwise comparisons.
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