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Ecological drivers influence the distributions of two cryptic lineages in an earthworm morphospecies

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26 ABSTRACT

27 Substantial genetic diversity exists within earthworm morphotypes, such that traditional

28 species designations may be incomplete. It is, however, currently not known whether these

29 different genetic variants show ubiquity or specialty in their distribution across separated sites

30 subject to different climatic, biotic or soil physicochemical factors. Here we report on the results

31 of a survey in which individuals of the *Lumbricus rubellus* morphotype, a species known to

32 comprise two deeply divergent genetic lineages in England and Wales, were sampled from 26

33 plots. Sequences from the mitochondrial cytochrome oxidase I gene were used to distinguish

34 lineages for 787 individuals. In conjunction, a range of geographic, climatic, biotic and soil

35 physiochemical variables were also collected for each locality.

36

37 Genotyping indicated that Lineage A was more common than Lineage B, comprising 58% of

38 the collected *L. rubellus*. Six site populations comprised only Lineage A, while only a single

39 site comprised entirely Lineage B. The remaining 20 sites containing both lineages. A

40 multivariate ordination of site variables identified major difference between sites were

41 associated with low pH, organic-rich soils in Western wet upland areas and pollutant levels

42 associated with sites in the South. Earthworm genotype (as proportion of Lineage A) was not

43 correlated with either of these major environmental axes. When individual variables of soil pH

44 and the percentage of soil organic matter, which are known to be key driver of soil species

45 distributions, were investigated as single variables significant relationship with lineage

46 frequency were found. Soil organic matter content was significantly negatively correlated with

47 Lineage A proportion, while pH was significantly positively correlated. This lineage preference

48 may be related to lineage metabolism and/or behavioral differences.

49

50 Measurement of tissue metal concentrations in worms from 17 sites identified a significant site

51 effect in all cases, but a lineage effect only for arsenic (higher Lineage B). Tissue arsenic

52 concentrations varied between lineages, supporting previous observations that there are

53 differences in the way the two lineages have adapted to manage exposure to this metalloid.

54

55 Keywords: Biogeography, Earthworm, Cryptic species, pH, Soil organic matter

1. INTRODUCTION

57 Soils contain a wealth of invertebrate biodiversity recognised for their important contributions

58 to ecological processes (Bardgett and van der Putten, 2014; Fitter et al., 2005; Giller, 1996).

59 One key group of species are the "ecosystem engineers": those organisms that modify the

60 physical state of the soil and resource availability for other species. Earthworms are known as

61 a key group of ecosystem engineers in many habitats. They perform a range of physical

62 (aeration, bioturbation, litter fragmentation) and biological (microbial interactions, exudate

63 production) roles in soil (Blouin et al., 2013; Lavelle et al., 1997; Sackett et al., 2013; Umarov

64 et al., 2008). Because of their functional importance, earthworms have emerged as a major

65 taxon for biomonitoring and biomarker assessments of human induced pressures on soil

66 communities (Cluzeau et al., 2012; Rutgers et al., 2009).

67

68 As soil invertebrate species, including earthworms, have been shown to be sensitive to a

69 range of land use change and pollution impacts (Bundy et al., 2007; Cluzeau et al., 2012),

70 different soil taxa have become a natural focus for research on the relationships between

71 environmental pressures, biodiversity and soil functioning (Bartlett et al., 2010; Leveque et al.,

72 2015; Rutgers et al., 2016). For community studies, a major constraint relates to current

73 uncertainties in earthworm taxonomy. Traditionally earthworm identification has relied on

74 morphology, but the paucity of suitable local keys and problems with application to juveniles

75 has also recently encouraged the use of molecular methods (Dominguez et al., 2015;

76 Emerson et al., 2011; Klarica et al., 2012). These genotyping studies have begun to challenge

77 current understanding of diversity through the identification of genetically distinct cryptic

78 lineages within previously established morphospecies.

79

80 Earthworm species in which cryptic lineage diversity has to date been identified include

81 *Eisenia fetida/andrei* (Römbke et al., 2016), *Lumbricus terrestris* (James et al., 2010),

82 *Aporrectodea caliginosa* (PerezLosada et al., 2009), *Allolobophora chlorotica* (King et al.,

83 2008), *Amynthas gracilis / Amynthas cortici* (Novo et al., 2015) and *Lumbricus rubellus*. For

84   *L. rubellus*, genotyping studies based on mitochondrial cytochrome oxidase I and II markers

85   have identified as many as 6 cryptic lineages across Europe (Giska et al., 2015), two of which

86   are found in the UK (Andre et al., 2010; Kille et al., 2013). The two UK lineages have 10-15%

87   divergence for the mitochondrial COI and COII sequences. While this implies they may

88   actually be cryptic species, recent analysis of multiple nuclear markers using RADseq has not

89   supported this interpretation, instead suggesting that different *L. rubellus* lineages may

90   actually correspond to a single highly polymorphic species (Giska et al., 2015). Comparative

91   studies of the two lineages in the UK have, nonetheless, identified physiological differences

92   between them, including variation in pheromone production (Jones et al., 2016), maturation

93   time (Anderson et al., 2013), metabolic profiles (Liebeke et al., 2014), mechanism of arsenic

94   adaptation (Kille et al., 2013), trace element metabolism (Andre et al., 2010), and microbiome

95   complement (Pass et al., 2015).

96

97   Despite known biological differences, the extent to which differences in distribution and

98   physiology are related to different geographical, climate and soil physicochemical preferences

99   between the two known UK lineages of *L. rubellus* is not established. The two lineages found

100   co-occur at some, but not all, sites meaning that they have some likely niche divergence that

101   facilitates coexistence (Andre et al., 2010; Giska et al., 2015; Kille et al., 2013). We aim to

102   better understand the nature of the spatial and geochemical drivers of lineage relative

103   abundance, and so here we test the hypothesis that the site distribution of the two cryptic *L.*

104   *rubellus* lineages is based on one or more geographical, climatic, physiochemical or biotic

105   drivers. We collected and genotyped morphotype *L. rubellus* at multiple well-characterized

106   sites that differed in their properties to investigate the relationships that determine lineage

107   distributions. Tissue metal concentrations were also measured to assess if trace metal levels

108   could also influence distributions, as could be the case if the two lineages had different

109   sensitivity to specific contaminants.

2. METHODS

2.1 Site selection

Twenty six sites located across England and Wales (Fig. 1) were visited between four times (for Devon Great Consols Mine and Control, Shipham Mine and Control, Cwmystwyth Mine and Control) and a single visit (for Porton Down, Parys Mountain, Castell, Clydach, Roman Gravel, Didcot) over four separate sampling events from Spring 2011 to Spring 2014. The chosen sites were selected to capture a range of the habitats and soil conditions under which morphotype *L. rubellus* can be collected. Land-uses covered included arable systems, broadleaf woodland, rough grassland and improved pasture habitats. Sites included both mineral and organic soils, although not true peats.

To allow the role of soil geochemistry and pollution status on lineage distribution to be addressed, sites of different known pollution history were sampled. Sites corresponded to three groups with respect to past land-use and associated expected contamination level. These were: 1) sites with no known pollution source (Unpolluted); 2) sites near to industrial facilities expected to be characterised by moderate pollution (Industrial polluted); and 3) sites at abandoned mining sites that can be expected to have high pollution (Mine polluted). For expected polluted sites from categories 2 and 3, a local control site was also sampled. This reference site was located outside of the area that was expected to be strongly influenced by the main pollution source and so was on soil expected to contain regional background pollutant concentrations.

2.2 Site geographical, biological and soil physiochemical characterisation

To allow the assessment of environmental drivers relating to lineage distribution, we used both publically available resources as well as our own analyses to gather data on each sites. Site geographical locations were collected as Easting and Northings from www.gridreferencefinder.com and site altitudes from www.freemaptools.com/elevation-

137 [finder.htm](). A series of site climate conditions were also assembled from

138 [www.metoffice.gov.uk/](). These were: annual average maximum temperature, annual average

139 minimum temperature, average January minimum temperature, average July minimum

140 temperature, average annual rainfall, average annual rain days and average annual frost

141 days. Initial visits to each site recorded main land-use (arable, broadleaf woodland, rough

142 grassland and improved pasture) and where present the average sward height of vegetation

143 at collection locations. The site was identified according to the level of shade (open, part

144 shaded, shaded) and the presence of livestock was noted.

145

146 An initial site survey identified points on the site where morphospecies *L. rubellus* could be

147 found. Thereafter all collections were focussed on these locations. For any one sampling event

148 at each site, between 6 and 25 fully clitellate *L. rubellus* were collected by digging and hand-

149 sorting from the soil to 20 cm depth. Generally the required number of worms could be

150 collected within a reasonable search period (approximately 2 h duration). There were,

151 however, some locations where this was not possible for particular sampling events. Climate

152 factors (notably dry soils), low frequency of adults in the population or the requirement to limit

153 site damage caused by digging were the major constraints. During collection, the presence of

154 other earthworm morphospecies was noted. Only common species were recorded (>5

155 individuals observed). In total 10 other species were found: *Aporrectodea caliginosa*,

156 *Aporrectodea rosea*, *Aporrectodea longa*, *Allolobophora chlorotica*, *Lumbricus castaneus*,

157 *Dendrobaena rubida*, *Lumbricus terrestris*, *Lumbricus festivus*, *Octolasion cyaneum*, and

158 *Octolasion tyrtaeum tyrtaeum*. At the end of sampling, the *L. rubellus* collected were washed

159 and blotted dry on-site and then snap frozen in liquid nitrogen before being transfered to the

160 laboratory under dry ice storage.

161

162 Triplicate soil samples from surface to 5 cm depth were collected from each site collection

163 location. All soil samples were oven dried at 80°C to constant weight and then sieved through

164 a 2 mm mesh to remove large roots and stones. Total concentrations of aluminium, arsenic,

165 barium, cadmium, cobalt, chromium, copper, iron, lead, manganese, mercury, molybdenum,

166 nickel, selenium, titanium, vanadium, zinc, calcium and total phosphorous were determined in

167 a 1 g sample of this processed soil following an aqua regia digestion protocol (Arnold et al.,

168 2008; Emmett et al., 2010; Spurgeon et al., 2008). Digests were subsequently analysed on a

169 Perkin Elmer Optima 7300 DV inductively coupled plasma optical emission spectrometry

170 instrument. For quality control, an in house reference traceable to BCR-143R (Commission of

171 the European Communities, Community Bureau of Reference) was included with each batch

172 of digestions. Measured concentrations were within 10% of certified values for all measured

173 elements with the exception of Al where the value was 55%. Organic matter content of each

174 soil sample was measured by proxy using loss on ignition following combustion at $500^{\circ}C$

175 (Rowell, 1994) and soil pH was quantified by electrode from a 1:2.5 volume soil:water mix (i.e.

176 1 volume soil with 2.5 volumes water added)(International Organisation for Standards, 2005).

177

178 *2.3 Lineage assignment by mitochondrial cytochrome oxidase I (COI) sequencing*

179 DNA was extracted from ~10 mg of frozen tissue (taken from the tail of each individual using

180 a scalpel) by automated DNA extraction using a Nucleoplex Plants Tissues DNA Extraction

181 Kit (Nucleoplex, Manchester, UK). After DNA quantification using Nanodrop (Thermo

182 Scientific, Willmington, DE), polymerase chain reaction amplification of the COI gene was

183 conducted using a set of established forward (GGTCAACAAATCATAAAGATATTGG) and

184 reverse (TAAACTTCAGGGTGACCAAAAAATCA) primers (Folmer et al., 1994) amplified after

185 5 minutes at $95^{\circ}C$ over 40 cycles of 30 sec $95^{\circ}C$, 30 seconds $48^{\circ}C$ and 60 seconds $48^{\circ}C$. A

186 sub-set of all PCR products were checked by gel electrophoresis to ensure successful

187 amplification and purified for sequencing using 0.25 U each of Exonuclease I and Shrimp

188 Alkaline Phosphatase (NEB, Hitchin, UK), incubated at $37^{\circ}C$ for 45 minutes and 80 $^{\circ}C$ for 15

189 minutes. Purified PCR products were then sequenced as in Andre et al. (2010), using ABI

190 PRISM® BigDye v3.1 Terminator Sequencing technology (Applied Biosystems, USA).

191

192 Sequences were aligned and trimmed for tree construction using the Maximum Likelihood

193 method and General Time Reversible substitution model with a gamma distribution in Mega

194 v5.01. Sequences for *L. rubellus* associated with specific mitochondrial lineages already

195 documented in the UK were incorporated into the analysis as anchor sequences (Anderson

196 et al., 2013), with sequences for *L. terrestris*, *L, festivus* and *L. castaneus* included as an out-

197 group. Tree topology was supported by bootstrap analyses over 1000 iterations. Individuals

198 that showed a close relationship with one of the two previously identified UK *L. rubellus*

199 lineages were identified from the analysis. Any individuals showing intermediate status

200 resulting from probable sequencing errors were excluded from further analysis.

201

202 *2.4 Earthworm tissue trace element concentrations*

203 Earthworm tissues from 494 individuals taken from a sub-set of 17 sites (Alice Holt ECN

204 Control, Avonmouth Control, Avonmouth Incinerator, Avonmouth Savalco, Cwmystwyth

205 control, Cwmystwyth mine, Devon Great Consols Control, Devon Great Consols Mine,

206 Drayton ECN Control, Port Talbot Control, Port Talbot blast furnace, Porton Down ECN,

207 Scunthorpe blast furnace, Scunthorpe Control, Shipham control, Shipham mine, Snowdown

208 ECN control) were prepared for analysis (nb samples from remaining sites were lost due to

209 storage issues). These samples were analysed for tissue Al, As, Cd, Co, Cr, Cu, Fe, Mn, Mo,

210 Ni, Pb, Se, Sr and Zn concentrations. Whole earthworms, after tail removal for DNA extraction,

211 were initially ground to powder under liquid nitrogen in a cryogenic mill. The powder was

212 freeze-dried and a 100 mg sample digested with 10 ml of 70% HNO_3 (Ultrapure) at 200°C for

213 15 minutes within a microwave vessel. Samples were run as two batches on a Perkin Elmer

214 DRCII ICP-MS. Each batch included multiple certified reference material samples for TORT-

215 2 and DOLT-4 (National Research Council, Canada). Certified values for reference materials

216 corroborated well with measured values. Average recovery was 91% (range 85% for Se to

217 110% for Pb) in the first batch of samples and 94.8% (range 53.9 for Al to 129% for Se) in the

218 second batch. Recoveries of only two metals, Al and Se, were outside 80% of certified values

8

219   for any run, with 19 of 27 determinations within 10%. With systematic bias absent, acquired

220   data can be used for statistical processing without requirement for recovery correction.

221

222   *2.5 Data handling and statistical analysis*

223   The number of *L. rubellus* returning COI sequences that were closely related to reference

224   sequences from previously collected lineage A and Lineage B individuals were counted for

225   each study site. These were calculated as proportions before being logit transformed as the

226   most appropriate transformation for biological proportion data (Warton and Hui, 2011), with

227   value of zero and one modified by addition and subtraction of half of the lowest proportion

228   respectively. Environmental drivers were established as either categorical (e.g. site type, site

229   shading, livestock presence/absence, earthworm species presence/absence) or as

230   continuous measured variables. The values for soil metal concentrations were log transformed

231   to obtain a Gaussian distribution in accordance with established practice (Davies, 1989).

232

233   Relationships amongst site geographical, climate and soil variables (after appropriate

234   transformation) were initially investigated using principal component analysis in Minitab 14

235   (Minitab, PA, USA). This reduces the dimensionality of the original complex dataset in an

236   unsupervised fashion, by successively generating new axes (principal components), that are

237   the linear combinations of the original data that explain greatest overall variance. The principal

238   components (PCs) arising can be interpreted as ordinations representing a summary of

239   environmental factors. Pearson correlations between (logit transformed) proportions of

240   Lineage A individuals at each site and the individual principal component scores were then

241   calculated to investigate the relationships between ordinated site characteristics and

242   genotype. Based on this analysis and prior knowledge, possible individual primary driver

243   variables were identified that were in turn assessed for Pearson correlation with genotype.

244   Because of this key role, percentage organic matter content and soil pH were selected as

245   focus variables. Site and lineage effects on earthworm tissue log transformed trace element

246   concentrations were analysed using a mixed model based general linear model in Minitab 14.

247   Within the model, site and lineage were included as fixed variables, with sampling campaign

248   (1-4) as a random factor.

249   *3. RESULTS*

250   High quality *L. rubellus* COI sequences were obtained from DNA samples taken from 787

251   earthworms for assignment as either Lineage A or Lineage B individuals. The maximum

252   number of sequences from any one site was 73, from Cwmystwyth Mine, and the minimum 3,

253   from Avonmouth Control (Fig. 1). In total, 457 individuals were assigned as Lineage A, 58%

254   of the number collected. The remaining 330 (42%) were assigned as Lineage B. Eight sites

255   (Avonmouth Savalco, Avonmouth Incinerator, Clydach Smelter, Didcot Power Station, Dinas

256   Powys, Parys Mountain, Scunthorpe Control and Scunthorpe) had populations comprising

257   only Lineage A individuals. Seven sites (Avonmouth Control, Castell Mine, Cwmystwyth

258   Control, Cwmystwyth Mine, Drayton ECN Control, Shipham Control and Shipham Mine)

259   contained populations comprised largely Lineage B, although only Castell Mine was

260   exclusively B. All remaining sites had mixed lineage populations, although with more Lineage

261   A than B individuals.

262

263   The site geographical, physical and soil characteristics were analysed using principal

264   component analysis. The first PC explained 21.4% of total variance. A number of parameters

265   were positively correlated with this axis, including soil % loss on ignition (LOI); soil log Fe, log

266   Co and log Al concentrations; some earthworm species; and site altitude and climate variables

267   including average rainfall and number of rain days. Negatively correlated variables included

268   soil pH; average July max temperature and average temperature; log Ca and log P

269   concentrations; and Easting (Fig 2). This first PC axis could therefore be interpreted as

270   representing a set of variables characterised by the presence of high organic matter, low pH

271   soils, associated with wetter and colder upland regions located mainly in the West of England

272   and Wales. The second PC axis explained a further 17% of variation, and was positively

273   associated with Northing and the weather variable of average frost days and average rain

274   days. Variables negatively associated with this axis included pollutant metal concentrations

275   such as log Pb, log Zn and log Cd concentrations (Fig 2). This axis can be interpreted as

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276   representing a gradient of metal contamination of sites located primarily in the South of

277   England and Wales.

278

279   To assess if the site characteristics summarised by the two first PCs potentially act as drivers

280   of lineage distribution, the PC1 and PC2 scores were correlated with the (logit transformed)

281   Lineage A proportion at each location. This did not identify significantly relationships between

282   Lineage A proportion with site PC1 or PC2 score (p=0.25 and p=0.074 respectively). As sites

283   PC scores were not significant, we next went on to investigate if individual variables measured

284   are related to the relative frequency of *L. rubellus* lineages. Specifically we selected soil pH

285   and LOI for initial assessment, as these are established as drivers of patterns of diversity

286   (Griffiths et al., 2011; Raty and Huhta, 2003). Both variables were significantly correlated with

287   logit Lineage A proportion (soil % OM -0.529, p=0.005; pH -0.392, p=0.048). The nature of

288   these two relationships were summarised by locally weighted scatterplot smoother model fits.

289   These indicate a decline in proportion of Lineage A (higher logit transformed values) as site

290   soil % LOI increases from 0-20%, thereafter remaining constant. The model fits for pH

291   indicated an initial decline in the proportion of Lineage A individuals (higher logit transformed

292   values) as pH increases from 4.5 to 5, with, thereafter, an increase in frequency (lower logit

293   transformed values) where site pH increases from 5 to 7.5 (Fig. 4 a,b). Amongst other

294   measured variables, only log soil Ca concentrations (-0.487, P=0.012) and the average annual

295   number of rain days (0.433, p=0.027) were also significantly correlated with Linage A

296   proportion. Both of these variables are, however, also significantly correlated with soil pH

297   (Annual rain days: -0.584, p=0.002; log soil Ca concentration:-0.751, p<0.001) making precise

298   attribution of cause challenging.

299

300   Separate univariate models were generated to analyse tissue metal concentrations in relation

301   to collection site and lineage. The collection site had a highly significant (p<0.001) influence

302   on tissue concentrations for all analysed trace elements. This is only to be expected, given

303   that the sites include locations with no history of local pollution, to highly contaminated

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304   industrial and mine sites. Lineage was also a significant factor in the model for As (p<0.02).

305   This difference is, however, based on only a relatively small difference in average tissue

306   concentrations between lineages across all sites. Thus, average concentrations in Linage A

307   of 10.69 (n=300) was slightly lower than the average tissue arsenic concentrations of 11.7

308   mg/kg (n=195) for Lineage B, Hence although statistically significant, the absolute magnitude

309   of difference in tissue As concentrations between lineages is small. For all other analysed

310   metals, there was no significant effect of lineage on tissue concentration (p>0.05).

4. DISCUSSION

Species distributions can be affected by a range of environmental drivers, including physiological tolerances, dispersal constraints, biotic interactions and anthropogenic influences (Dennis and Hellberg, 2010; Gaston, 2003). Among earthworms, species show preference for certain habitats, for example common compost earthworm species such as *Eisenia fetida*, *Perionyx excavatus* and *Eudrilus eugeniae* preferentially occupy organic matter rich habitats associated with animal manure or composting vegetation (Edwards, 2004). Further, some species also have preference for different soil physiochemical properties. For example, Jaensch et al (2013) found differences in morphospecies preference across different soil pH classes, with species such as *Allolobophora chlorotica*, *Aporrectodea rosea*, *Aporrectodea longa* and *Lumbricus terrestris* preferring soils with pH >5.6, and *Dendrodrilus rubida*, *Dendrobaena octaedra* and *L. rubellus* soils with pH <5.6.

The UK earthworm fauna is notably denuded, comprising only around 20-25 native species, compared to about 180 species that are found in neighboring France (Bouché, 1972; Sims and Gerard, 1985). The reduced earthworm fauna of the UK can be linked to its recent history of glaciation and the severing of the land bridge to Europe that restricted earthworm colonization after glacial retreat. This influence of quaternary glaciation is consistent with what is known about the current distribution and genetic structure of a range of species across Europe and the UK (Hewitt, 2000). Among UK earthworm species, the majority show a widespread and cosmopolitan distribution (Boag et al., 1997; Carpenter et al., 2012; Rutgers et al., 2016; Sims and Gerard, 1985). Habitat preferences are known, such as those for pH and for organic rich habitats as discussed previously, however the spatial heterogeneity of terrestrial habitats means that at coarse recording scales (e.g. 10 km^2 or even 1 km^2), a significant proportion of UK earthworm species may be present in any given sampling area (e.g. a mixed land-use area subject to comprehensive earthworm sampling within different vegetation stands and habitats).

339 Genetic marker studies have identified deeply divergent cryptic lineages within many common

340 UK earthworm morphospecies based on mitochondrial or nuclear genetic marker analysis. An

341 active debate currently surrounds the question of whether these cryptic lineages correspond

342 to cryptic species or highly polymorphic species variants (Blakemore et al., 2010; Giska et al.,

343 2015; King et al., 2008). In the specific case of *L. rubellus*, the presence of cryptic lineages is

344 established from studies conducted from measurement of highly divergent (13-15%)

345 sequences for both of the cytochrome oxidase I and II mitochondrial genes (Andre et al., 2010;

346 Donnelly et al., 2014; Kille et al., 2013). Pan-European studies have shown that at continental

347 scale, morphotype *L. rubellus* may comprise of 5 or more such deeply divergent lineages

348 (Giska et al., 2015), two of which were here found across sites in England and Wales (Fig 1).

349 Recently RADseq analysis suggests that cryptic *L. rubellus* lineages may represent a case of

350 a highly polymorphic single species rather than true cryptic species (Giska et al., 2015).

351 Nonetheless, previous studies of the lineage physiology have identified a number of

352 differential responses between lineages (as previously outlined notably for the two UK

353 lineages). For example, Jones et al. (2016) found that the two lineage were favorably attracted

354 to soils that had previously been worked by earthworm of their own rather than the alternative

355 lineage. These results suggests that pheromone attractants may allow mate selection in mix

356 populations,. such as those that are found at the majority of our sampled site. Such selection

357 has the potential to underpin lineage differences in habitat preference and, as a consequence,

358 different spatial distributions at local scale.

359

360 Earthworms are key ecosystem engineers for the role that play an important role in the

361 creating of the spatial structure and chemistry of the soil habitat through bioturbation, litter

362 degradation and nutrient cycling (Edwards, 2004; Lavelle et al., 1997; Liebeke et al., 2015).

363 The extent to which the divergent lineages of common earthworm species overlap in respect

364 of habitat preference will be an important determinant of morphospecies contributions to

365 different ecosystem processes across space and time. The analysis here suggests that, in the

366 case of the two UK lineages of *L. rubellus*, there are ecological drivers of distribution.

367   Individually, soil pH and % OM were both significant correlated with the proportions of *L.*

368   *rubellus* Lineage A (and conversely Lineage B) collected across the 26 sample sites. These

369   two measurement parameters were selected for particular focus because they are recognized

370   as important environmental drivers of the distribution of a number of soil taxa (Cassagne et

371   al., 2003; Griffiths et al., 2011; Raty and Huhta, 2003). Additionally, there are also correlations

372   with other climate and soil variable that are themselves know to influence soil pH through soil

373   geochemistry and leaching.

374

375   Different soil pH preferences have direct effects on earthworm traits including reproduction,

376   growth and survival (Baker and Whitby, 2003; Spurgeon et al., 2006; Van Gestel et al., 1992).

377   *L. rubellus* is tolerant of relatively low pH, being commonly (and even preferentially) found in

378   moderately acidic soils (Jaensch et al., 2013). Results here suggest that this cosmopolitan

379   nature could partly arise from different lineage pH preferences, with Lineage B found in more

380   acid habitats from pH 4.5 to 5.5 and Lineage A preferentially in nearer neutral pHs of 5.5 and

381   above. Thus, within the current study, Lineage B was absent from 6 of 26 sampled sites, while

382   Lineage A was found at all except one of the sampled sites. The detailed genetics of the two

383   cryptic lineages may provide some clues to the basis of such differences. Studies of

384   mitochondrial and genetic marker genes have established that Lineage B has lower genetic

385   diversity of measured traits than Lineage A (Donnelly et al., 2014; Kille et al., 2013). This

386   suggests that Lineage B may have undergone a population bottleneck that restricted the

387   genetic diversity, and possibly, the colonization capacity of this lineage.

388

389   The strongest correlate of lineage frequencies was soil organic matter (% loss on ignition).

390   The fresh and partially degraded soil organic component provides earthworms with food. It is,

391   therefore, possible that this association is driven by different dietary requirements of the two

392   lineages, as has been recognized for different earthworm species (Piearce, 1978). However,

393   in addition to acting as food, soil organic matter also contributes to soil structure and moisture

394   retention. Earthworms are known to be sensitive to soil texture, with regional studies linking

395 species distributions to soil sand, clay and organic matter content (Joschko et al., 2006;

396 Salome et al., 2011). Soils lacking in organic matter are also vulnerable to prolonged periods

397 of high soil moisture deficit. This can be challenging for earthworms given their critical need

398 to retain water balance. The significant correlation with site average rain days also points to a

399 possible influence of soil hydrology on distributions. Metabolomic analyses have identified that

400 many earthworm species contain a high number of betaines which likely act as osmolytes that

401 help to retain soil water balance (Liebeke and Bundy, 2013). Any differences in the extent of

402 such protection between lineages may influence colonization ability for more drought

403 susceptible soils.

404

405 Although there is correlation of lineage frequency with both soil pH and soil organic matter,

406 the fact that these two soil variables are co-correlated to other environmental variables makes

407 it hard to unequivocally assign them as the major drivers of lineage distribution. For example,

408 high organic matter/low pH soils are more common in the West of England and Wales than in

409 the East. This geographic relationship could potentially be associated with different

410 recolonization histories for the two lineages, e.g. perhaps recolonization from different glacial

411 refugia (Hewitt, 2000). However, as there is no significant correlation of Easting to lineage

412 proportion, this seems less likely than direct effects of soil organic matter/pH. Ultimately, to

413 tease apart the drivers of lineage preference, higher resolution collection and mapping and

414 experimental manipulation of habitats would be required.

415

416 Differences in physiology that separate species in relation to habitat preference could also

417 affect the way that the two lineages handle and accumulate different trace elements. For the

418 site-level analysis, the soil concentrations of major pollutant metals were correlated with PC2,

419 which was not associated with lineage. For the individual analysis of tissue metals, arsenic

420 was the only one found to vary with lineage (significantly higher in Lineage B individuals).

421 Previous work has indicated that the two lineages differ in the genetic mechanisms underlying

422 the development of arsenic tolerance. Analysis of amplified fragment length polymorphisms

423 indicated that Lineage A showed differences in patterns of nuclear markers indicating genetic

424 tolerance, while Lineage B showed a difference in DNA methylation patterning, but not genetic

425 differences (Kille et al., 2013). The observed difference here in As accumulation between

426 lineages across sites suggests that these genetic differences lead to phenotypic differences

427 in the handling of As.

428

429 *5. CONCLUSIONS*

430 Earthworms represent 'super-sentinels' exploited for environmental monitoring and

431 ecotoxicology, as well as being keystone soil engineers essential for soil quality. The

432 identification of possible drivers of species and lineage distributions has potential implications

433 for their use in environmental assessment as well as in studies of ecosystem service delivery.

434 For example, when assessing biodiversity effects of pollution and land-use change it may be

435 valuable to consider the occurrence of different lineages to understand how populations may

436 adapt to change through changes in lineage frequency. This analysis may be required

437 because the two widespread cryptic lineages of *L. rubellus* differ in their habitat preferences

438 with frequencies changing as conditions change. Given that bacterial communities are also

439 known to differ in relation to soil pH, then difference in the nature and strengths of earthworm

440 and microbial interactions can be expected between lineages. These relationships between

441 soil macrofauna and microbes are key to soil carbon turnover, nutrient cycling and soil

442 structural characteristics and this aspect warrants further investigation. Earthworms are also

443 valuable for metal biomonitoring. Our results suggest that the lineages behave identically with

444 respect to metal bioaccumulation, with the exception of As. Thus, selection of morphotype *L.*

445 *rubellus* will provide a coherent picture of metal accumulation independent of lineage, unless

446 As is a specific focus of any assessment.

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615

616   LEGENDS TO FIGURES

617

618   Figure 1. Location of collection sites and the proportion of Lineage A (dark blue shading) and

619   Lineage B (light yellow shading) *L. rubellus* based on the total number of collected and

620   assigned genotyped individual (given in brackets) for the 26 sites visit over four separate

621   collection campaigns

622

623   Figure 2. Principal component analysis results show the ordination of site geographical,

624   climatic, biotic and soil chemical variables of sample sites showing the major related site

625   characteristic variables.

626

627   Figure 3. Boxplots showing median (centre line), upper and lower quartile (box limits) and

628   upper and 95% confidence intervals (whiskers) of trace metal concentrations measured

629   across 17 samples site for assigned Lineage A and Lineage B *L. rubellus.*
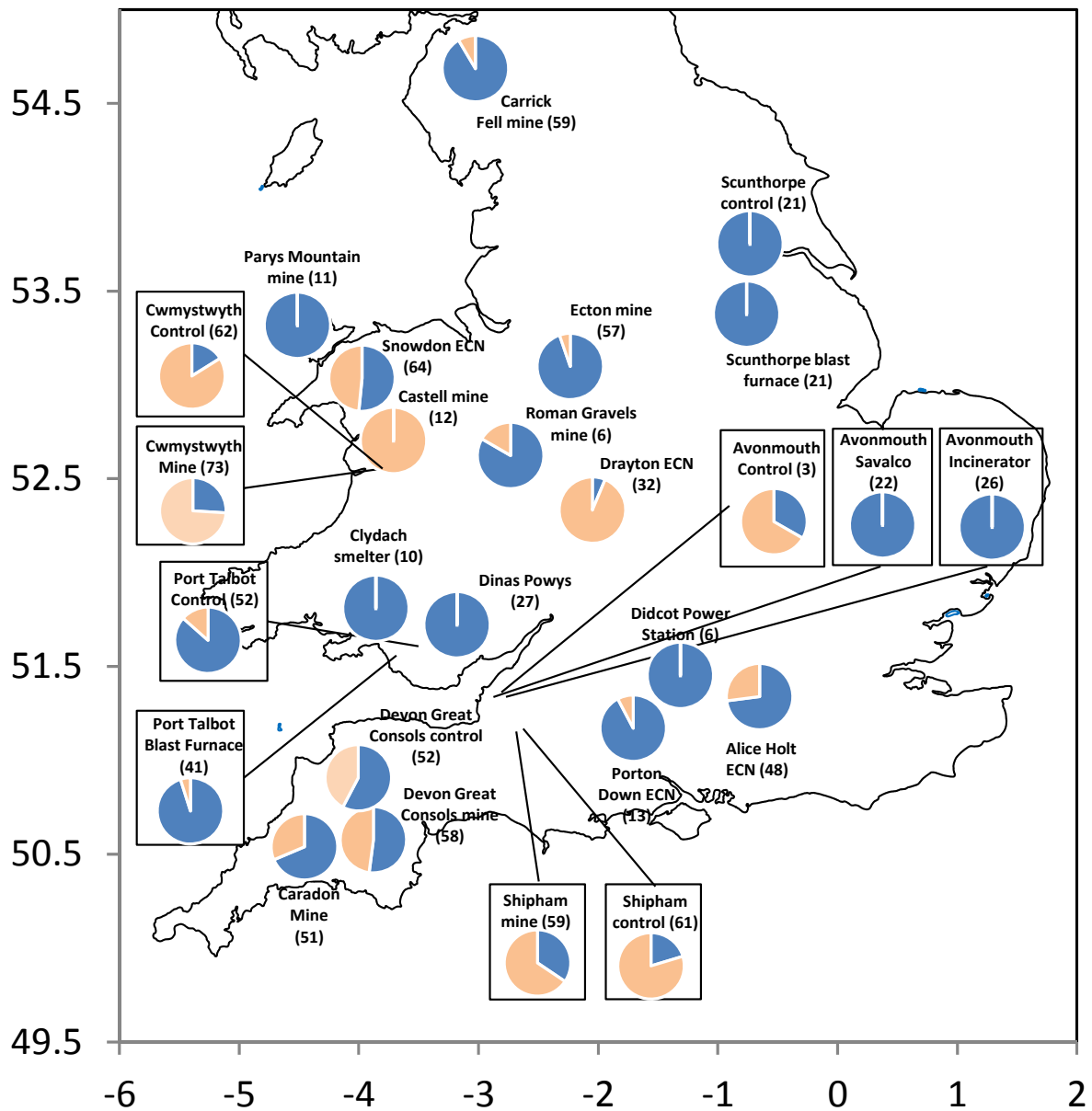
630

631   Figure 4. Scatterplots with fitted locally weighted scatterplot smoother line of proportion of

632   Lineage A *L. rubellus* in relation to (a) Soil % OM and (b) soil pH.
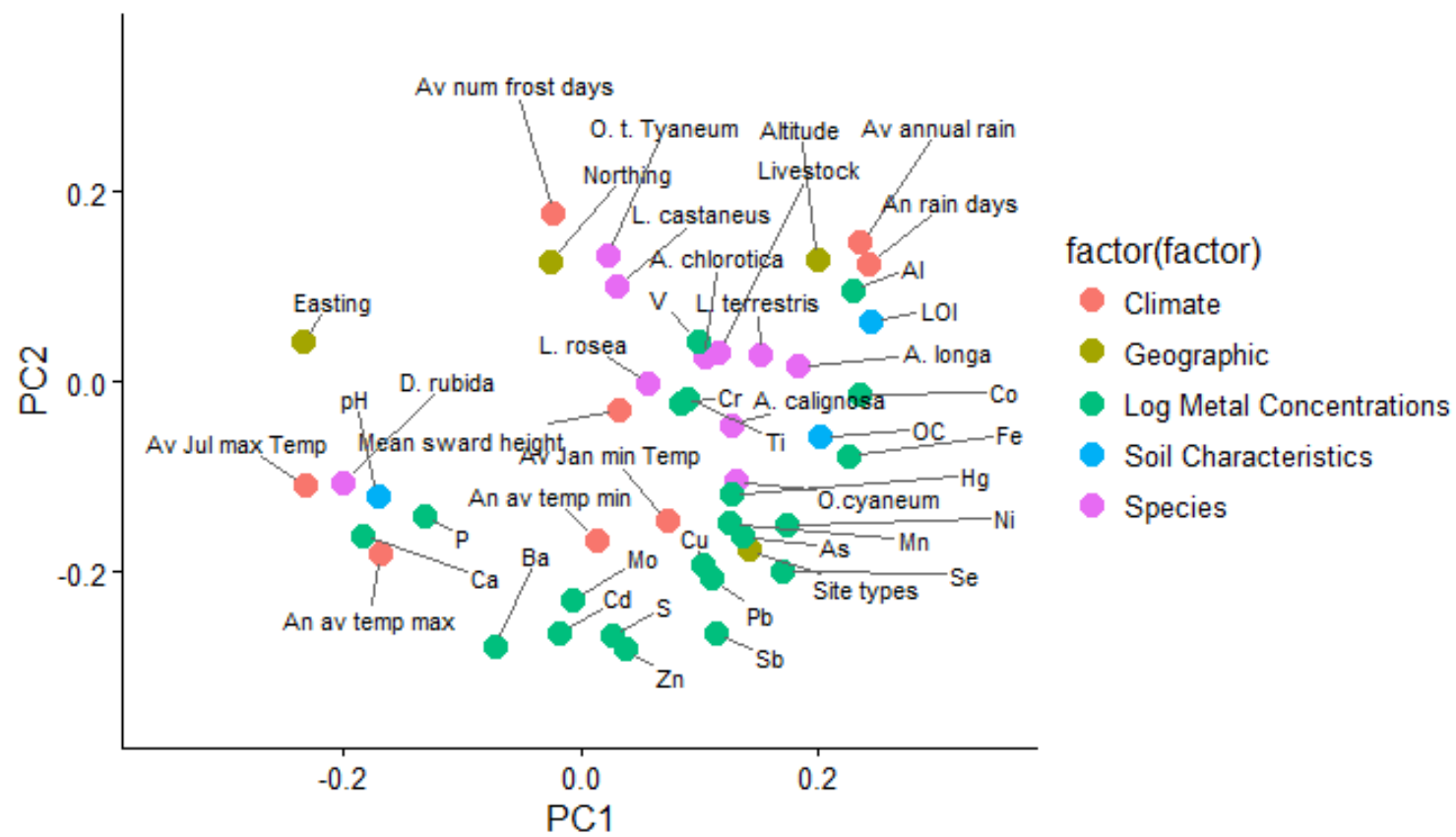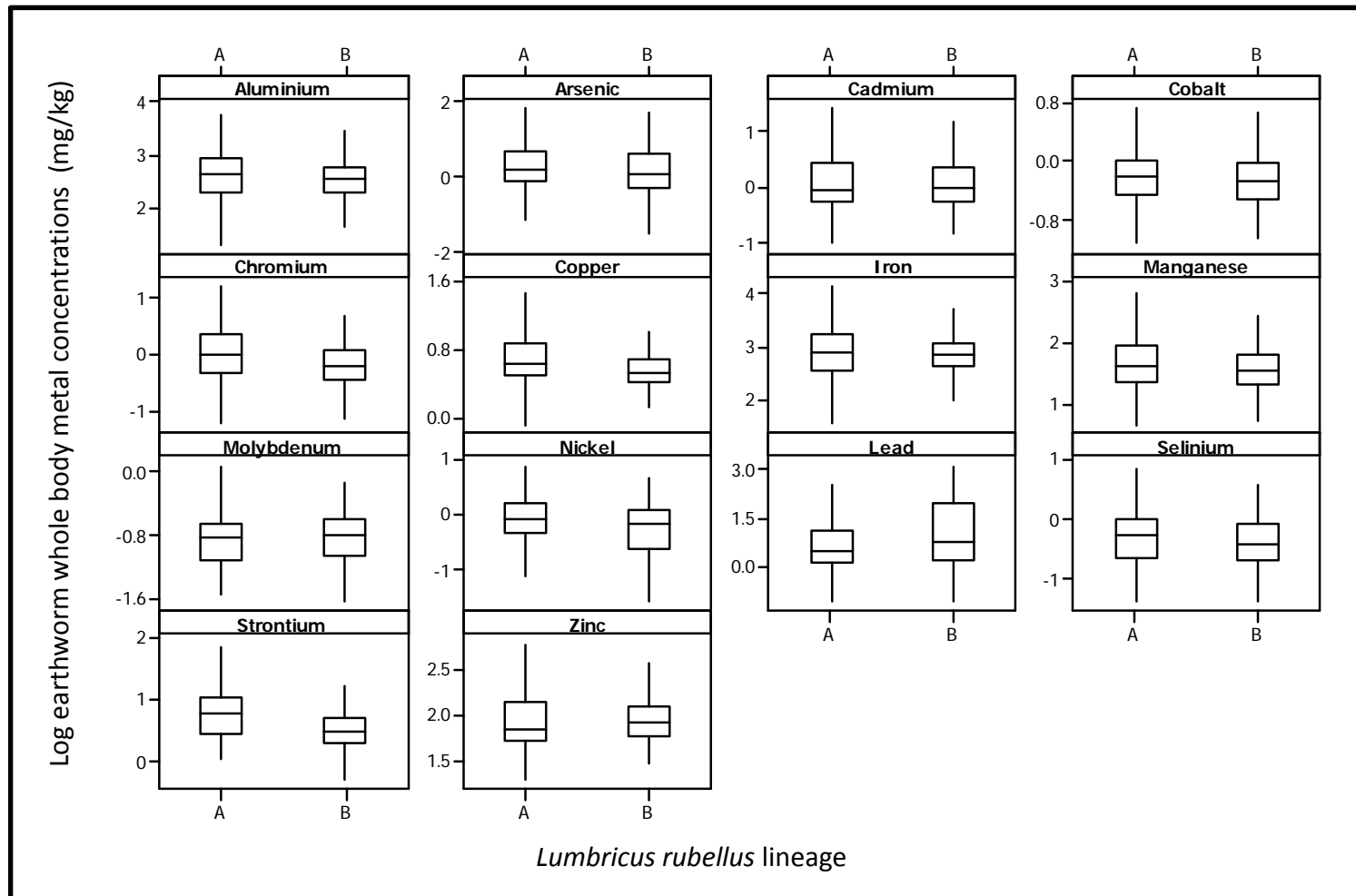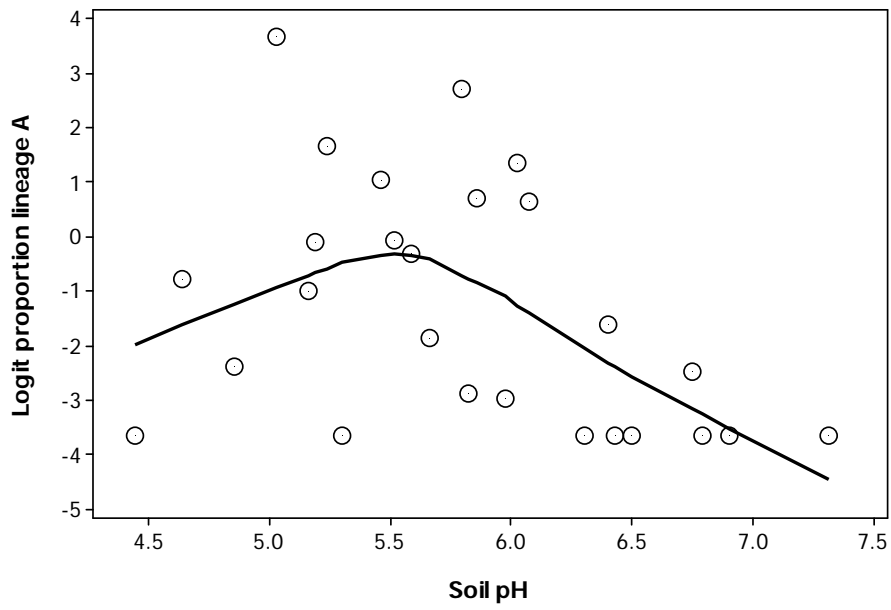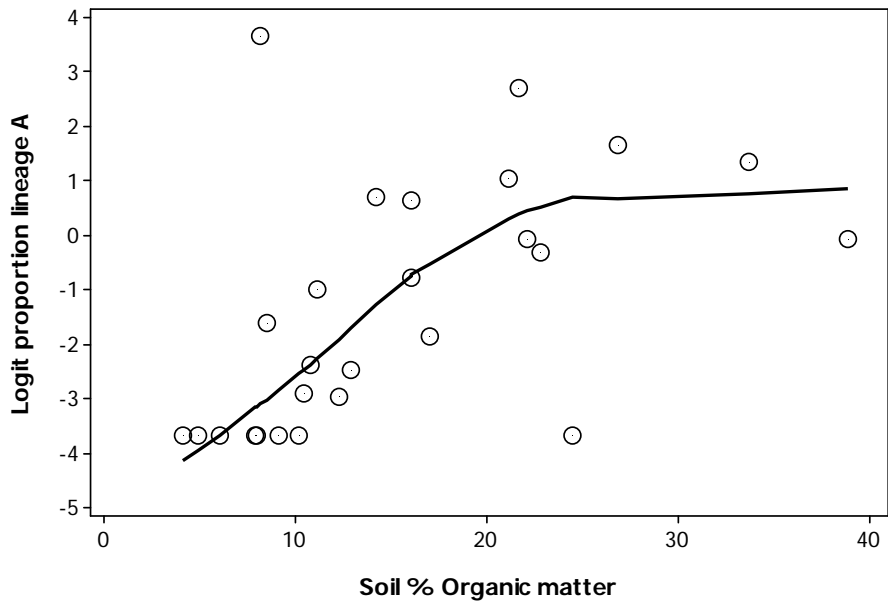
633

634   FIG. 1

635

636

FIG. 2

FIG. 3

FIG. 4

Supplementary Table 1. Geographical locations and reported climatic conditions of the 23 sites used for the collection of morphotype *L. rubellus*.

Site name	Site types	Land use	Ordnance Survey Grid reference	Easting	Northing	Altitude	Annual Average temp Max	Annual average temp min	Average Jan Min Temp	Average Jul Max Temp	Average Annual Rain	Annual rain days	Average No Forst Days
Alice Holt ECN Control	Unpolluted	Broadleaf woodland	SU 80060 39821	480060	139821	88.6	14.1	6.4	1.6	21.9	755	121	45.9
Avonmouth Control	Unpolluted	Improved pasture	ST 57006 82149	357006	182149	7	14.2	7	2.2	21.5	802	126	34.9
Avonmouth Incinerator	Industrial polluted	Rough grassland	ST 54099 81659	354099	181652	6	14.2	7	2.2	21.5	802	126	34.9
Avonmouth Savalco	Industrial polluted	Rough grassland	ST 53859 79411	353859	179411	6	14.2	7	2.2	21.5	802	126	34.9
Caradon Mine	Mining polluted	Rough grassland	SX 25624 69792	225624	69792	226	13.2	7	3	19.1	1385	172	30.6
Carrick Fell Mine	Mining polluted	Rough grassland	NY 32211 32982	332211	532982	661	13	5.8	1.6	19.7	1521	176	56.5
Castell Mine	Mining polluted	Rough grassland	SN 77415 81254	277415	281254	297	11.9	5.2	1	18.2	186	191	58.4
Clydach Smelter	Industrial polluted	Broadleaf woodland	SN 69587 01409	269587	201409	25	13.5	8.5	4	19.6	999	148	9.7
Cwmystwyth control	Unpolluted	Rough grassland	SN 79598 74222	279598	274222	198	11.9	5.2	1	18.2	1856	191	58.4
Cwmystwyth mine	Mining polluted	Rough grassland	SN 80852 75166	280852	275166	177	11.9	5.2	1	18.2	1856	191	58.4
Devon Great Consouls Control	Unpolluted	Improved pasture	SX 42560 74019	242560	74019	133	14	8.1	4	19.9	1007.4	142	16.3
Devon Great Consouls Mine	Mining polluted	Broadleaf woodland	SX 42385 73152	242385	73152	133	14	8.1	4	19.9	1007.4	142	16.3
Didcot Power Station	Industrial polluted	Broadleaf woodland	SU 51645 91402	451645	191402	53	14.4	5.9	1.2	22.6	661	112	57.7
Drayton ECN Control	Unpolluted	Improved pasture	SP 16391 55061	416391	255061	66	14.5	5.9	1.3	22.8	614	114	52.2
Dinas Powys	Unpolluted	Broadleaf woodland	ST 15868 70431	315868	170431	57	14.7	7	2.3	21.7	1151.9	149	35.7
Ecton Mine	Mining polluted	Broadleaf woodland	SK 09698 58263	409698	358263	103	13.9	6	1.2	22.1	598	112	49.1
Parys Mountain	Mining polluted	Rough grassland	SH 43829 89971	243829	389971	117	13.2	7.7	3.6	18.8	841	143	20.3
Port Talbot Control	Unpolluted	Rough grassland	SS 83690 84574	283690	184574	150	13.5	8.5	4	19.6	999	148	9.7
Port Talbot Blast Furnace	Industrial polluted	Rough grassland	SS 79001 85463	279001	185463	6.1	13.5	8.5	4	19.6	999	148	9.7
Porton Down ECN	Unpolluted	Improved pasture	SU 19575 37692	419575	137692	102	14.1	6.2	1.4	21.9	749	122	47.6
Roman Gravels Mine	Mining polluted	Rough grassland	SJ 33592 00339	333592	300339	368	14.1	5.6	1.3	21.6	668	126	51.8
Scunthorpe blast furnace	Industrial polluted	Arable	SE 94800 15840	494800	415835	19.7	13.4	5.7	0.9	21.3	613	115	49.8
Scunthorpe Control	Unpolluted	Arable	SE 93156 12000	493156	412000	10.7	13.4	5.7	0.9	21.3	613	115	49.8
Shipham control	Unpolluted	Improved pasture	ST 46312 59409	346312	159409	54	14.6	7	2.6	21.7	899	134	28.9
Shipham mine	Mining polluted	Improved pasture	ST 44799 57273	344799	157273	169	14.6	7	2.6	21.7	899	134	28.9
Snowdown ECN control	Unpolluted	Rough grassland	SH 63674 55116	263674	355116	748	12	5.9	1.8	18.1	2612	199	50.1

Supplementary Table 2. Vegetation and presence (shaded) or absence (unshaded) for earthworm species at the 23 sites used for the collection of morphotype *L. rubellus.*

Site name	Mean sward height	Livestock	Exposure	*Aporrectodea calignosa*	*Aporrectodea rosea*	*Lumbricus castaneus*	*Dendrobaena rubida*	*Aporrectoda longa*	*Lumbricus terrestris*	*Lumbricus festivus*	*Octolasion cyaneum,*	*Allolobophora chlorotica*	*Octolasion tyrtaeum tyaneum*
Alice Holt ECN Control	-	Deer	shaded										
Avonmouth Control	20 cm	None	part-shaded	■				■			■		
Avonmouth Incinerator	10 cm	None	open			■			■			■	
Avonmouth Savalco	35 cm	None	open						■				■
Caradon Mine	5 cm	horses	open										
Carrick Fell Mine	10 cm	sheep	open						■				
Castell Mine	5 cm	None	open				■			■			
Clydach Smelter	5 cm	None	shaded	■	■		■			■			
Cwmystwyth control	10 cm	None	open					■					
Cwmystwyth mine	10 cm	sheep	open										
Devon Great Consouls Control	10 cm	None	open	■					■			■	
Devon Great Consouls Mine	10 cm	None	part-shaded				■						
Didcot Power Station	-	None	Shaded	■		■		■			■		
Drayton ECN Control	15 cm	None	open	■	■	■		■	■			■	
Dinas Powys	-	None	shaded	■	■	■			■			■	
Ecton Mine	-	None	part-shaded						■				
Parys Mountain	5 cm	None	part-shaded	■									
Port Talbot Control	10 cm	None	part-shaded	■	■	■			■			■	
Port Talbot Blast Furnace	10 cm	horses	open										
Porton Down ECN	8 cm	None	part_shaded										
Roman Gravels Mine	3 cm	horses	open										
Scunthorpe blast furnace	5 cm	None	open	■		■		■	■			■	
Scunthorpe Control	6 cm	None	part-shaded	■								■	
Shipham control	10 cm	None	open					■	■				
Shipham mine	10 cm	cattle	part-shaded			■		■					
Snowdown ECN control	15 cm	sheep	open	■		■	■						

Supplementary Table 3. Arithmetic mean of measured soil chemical properties for pH, loss on ignition and concentrations of a suite of trace lements based on analysis of three samples collected from sites the 23 sites used for the collection of morphotype *L. rubellus*.

Site name	Soil pH	% Soil loss on ignition	Al (mg/kg)	As (mg/kg)	Ba (mg/kg)	Cd (mg/kg)	Co (mg/kg)	Cr (mg/kg)	Cu (mg/kg)	Fe (mg/kg)	Hg (mg/kg)	Mn (mg/kg)	Mo (mg/kg)	Ni (mg/kg)	Pb (mg/kg)	Sb (mg/kg)	Se (mg/kg)	Ti (mg/kg)	V (mg/kg)	Zn (mg/kg)	Ca (mg/kg)	P (mg/kg)	S (mg/kg)
Alice Holt ECN Control	5.16	11.15	8737	13.2	19.3	0.1	6	12.6	12.4	16500	0.17	78	0.3	9.5	27.1	0.4	0.5	7	24.6	43	1640	424	202
Avonmouth Control	5.86	14.23	11130	20.9	356	2.3	10	46.3	75.1	26133	0.76	539	1.2	25.1	207	1.3	1.7	32.9	26.9	697	68633	524	1029
Avonmouth Incinerator	6.5	7.94	11000	36.9	439	50.6	12.3	23.6	204.3	26133	1.07	1010	2	25.7	1943	14.9	5.6	34.4	31.8	4640	39267	1600	3190
Avonmouth Savalco	6.3	24.51	14733	7.8	75.1	2.6	8.1	25.1	29.8	19233	0.25	663	0.6	18.4	99.6	1.2	0.6	95.5	29.6	299	4777	1233	734
Caradon Mine	4.64	16	5147	407	33.7	1.17	2.03	3.9	609	20633	0.33	221	2.57	4.27	69.1	3.17	1.49	27.4	13.8	43.5	342	776	884
Carrick Fell Mine	4.85	10.8	16300	737	33.5	3.07	8.18	16.4	59	33300	2.06	837	18.10	11.4	173	7.01	3.85	165	92.6	282	1607	961	1457
Castell Mine	5.03	8.15	14333	14.7	15.7	9.03	12.5	16.4	60	46267	0.5	844	1.16	16.5	210	2.28	1.35	8.1	26.8	1792	321	506	312
Clydach Smelter	5.3	10.2	6780	44.1	142	1.61	90.1	32.3	465	25067	0.34	1369	2.68	1799	278	3.57	14.43	70.9	47.2	370	39277	945	1230
Cwmystwyth control	5.23	26.89	16167	19.8	14.6	0.1	6.7	18.7	19.7	38033	0.12	467	0.9	14.3	626	1.6	1.2	6.6	24.6	116	209	688	709
Cwmystwyth mine	5.46	21.13	20033	49	29.8	0.2	41.2	24.1	28.8	50433	112	2597	1.5	23.3	657	2.5	1.4	22	33.3	127	551	689	534
Devon Great Consols Control	5.58	22.79	21533	310	45.5	0.4	14.8	31.7	107.3	45800	0.05	585	1	27.5	68	1.4	2.3	41.8	38.7	140	2213	1177	919
Devon Great Consols Mine	5.19	38.9	17300	6270	45.9	0.2	25.7	17.8	2647	79600	0.64	630	1.3	22.3	225	13.9	2.6	38.4	38.5	277	3340	286	1503
Didcot Power Station	7.31	4.17	10570	14.5	139	0.53	8.89	14.7	41	22900	0.34	309	0.93	24.1	102	2.42	1.40	15.1	26.9	181.00	37600	901	1250
Drayton ECN Control	5.79	21.65	12833	25.1	48.4	0.4	4.9	15.8	46	28533	0.1	805	2.3	10.7	30.2	0.6	0.5	34.3	32	176	35000	2000	1183
Dinas Powys	6.9	8	17166	23	85.4	1.2	16.1	39	42.6	24933	0.1	1120	1.7	45.1	109	*	1.2	41.5	45.7	770	12766	2070	*
Ecton Mine	5.82	10.4	1403	136	537	61.4	33.2	5.4	5787	12267	0.35	965	101	72.4	1553	92.1	5.42	20.1	20.2	6047	116200	203	9280
Parys Mountain	4.44	4.95	681	1480	96	8.77	7.78	5	3673	175667	3.34	35.7	37.5	3.17	29033	210	42	54.4	19	2333	1749	26.7	22467
Port Talbot Control	5.66	16.99	15300	14.5	127	0.6	11.3	37.7	33.8	31467	0.27	1287	0.9	17.4	39.1	0.8	1.2	15.7	62.9	480	13903	1427	1083
Port Talbot Blast Furnace	5.98	12.3	10323	14.5	287	0.8	4.7	129	35.5	34967	0.33	3550	1.7	16.8	117	2.2	1.1	230.7	140	341	89133	1227	1680
Porton Down ECN	6.75	12.93	6153	21	88.3	0.7	8	20	39.2	13567	0.37	611	0.9	17.3	109	1.4	0.5	46.3	17.2	184	148333	3100	1423
Roman Gravels Mine	6.4	8.48	10633	13.7	123	14.2	14.3	13.3	99	30133	0.33	582	1.17	24.8	1125	2.48	1.33	15.7	18.4	1788	6367	455	878
Scunthorpe blast furnace	6.43	9.14	10803	40.6	90.3	0.3	13.1	42	25.4	59900	0.33	1227	1.5	29.5	124	1.8	1	68.3	128.7	183	13500	1497	453
Scunthorpe Control	6.79	6.04	7197	24.6	47.4	0.4	4.9	15.4	20.2	21767	0.12	535	2.3	11.2	30.0	0.6	0.5	33.6	30.7	69.1	130667	1253	1167
Shipham control	6.07	16.1	12967	37.1	302	2.1	7.5	23.1	19.1	26500	0.03	465	0.7	16.4	163	0.9	0.4	53.9	34.8	328	3520	972	584
Shipham mine	6.03	33.7	9907	867	1526	404	12.5	21	85.1	85100	12.3	2277	5.1	31.6	7260	48.8	3.8	51.4	28.5	31833	16117	2370	3547
Snowdown ECN control	5.52	22.07	38600	17.5	10.1	0.3	30.6	78.2	19.5	70733	0.19	1350	0.5	30.4	37	0.5	1.4	1530	239	114	1111	524	658