

1 **THE EFFECTS OF LEAD SOURCES ON ORAL BIOACCESSIBILITY IN SOIL AND**
2 **IMPLICATIONS FOR CONTAMINATED LAND RISK MANAGEMENT**

3
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14 Abstract

15
16 Lead (Pb) is a non-threshold toxin capable of inducing toxic effects at any blood level but availability
17 of soil screening criteria for assessing potential health risks is limited. The oral bioaccessibility of Pb
18 in 163 soil samples was attributed to sources through solubility estimation and domain identification.
19 Samples were extracted following the Unified BARGE Method. Urban, mineralisation, peat and
20 granite domains accounted for elevated Pb concentrations compared to rural samples. High Pb
21 solubility explained moderate-high gastric (G) bioaccessible fractions throughout the study area.
22 Higher maximum G concentrations were measured in urban (97.6 mg kg⁻¹) and mineralisation (199.8
23 mg kg⁻¹) domains. Higher average G concentrations occurred in mineralisation (36.4 mg kg⁻¹) and
24 granite (36.0 mg kg⁻¹) domains. Findings suggest diffuse anthropogenic and widespread geogenic
25 contamination could be capable of presenting health risks, having implications for land management
26 decisions in jurisdictions where guidance advises these forms of pollution should not be regarded as
27 contaminated land.

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30 Keywords: lead, anthropogenic pollution, geogenic contamination, oral bioaccessibility, human health
31 risk assessment, soil

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38 1. Introduction

39

40 Lead (Pb) is a toxic trace element that has been the subject of extensive human health research. Its
41 neurotoxic effects from the oral exposure pathway, particularly in children, are well documented
42 (EFSA, 2010; ATSDR, 2007; Ryan *et al.*, 2004; CCME, 1999; Rosen, 1995). Some studies also
43 suggest Pb exposure may be associated with increased incidences of violent crime (Meilke & Zahran,
44 2012; Nevin, 2007; Nevin, 2000). Whilst many known toxins have quantifiable threshold exposure
45 levels above which toxic health effects could occur, Pb is currently regarded by the global scientific
46 community as a non-threshold toxin. Non-threshold toxicity indicates that laboratory studies have not
47 identified a minimal risk level (MRL) or a no observed adverse effect level (NOAEL). Adverse
48 health effects could potentially occur at any blood Pb level (EA, 2009; ATSDR, 2007; USEPA, 1988).
49 Therefore, it is arguable that no amount of Pb exposure can be regarded as safe based on available
50 research to date.

51

52 Elevated Pb in the environment is attributed to a number of human activities and industrial processes
53 such as fuel combustion, mining, agricultural slurry spreading, and incineration of municipal wastes
54 (Alloway, 2012; Nriagu & Pacyna, 1989). The reported natural abundance of Pb in the earth's crust
55 ranges from 12 -14 mg kg⁻¹ (Rose *et al.*, 1979; Lee & Yao, 1970; Krauskopf, 1967; Taylor, 1964)
56 although globally reported normal background concentrations (NBC) in soil can substantially vary.
57 The United States Environmental Protection Agency (USEPA) suggests natural Pb concentrations in
58 the United States range from 50 to 400 mg kg⁻¹ (USEPA, 2013). The Canadian Council of Ministers
59 of the Environment (CCME) provides a mean range of 12 – 25 mg kg⁻¹ for Canadian Soils (CCME,
60 1999). The average reported concentration in rural soils in the United Kingdom (UK) is 52.6 mg kg⁻¹,
61 ranging from as low as 2.6 to as high as 713 mg kg⁻¹ (EA, 2007). In Northern Ireland (NI), Jordan *et al.*
62 (2001) reported a mean total Pb soil concentration of 23.2 mg kg⁻¹. More recently, the Tellus
63 Geochemical Survey of NI measured a higher average total Pb concentration of 41.7 mg kg⁻¹, with a
64 maximum extractable Pb concentration exceeding 3,000 mg kg⁻¹ near the Belfast metropolitan area.
65 In rural parts of NI, McIlwaine *et al.* (2014) reported a typical threshold value (TTV) of 63 mg kg⁻¹.

66

67 Due to the global variability in Pb NBCs and also with regards to its non-threshold toxicity,
68 identifying a starting point for contaminated land assessment in a human health context is challenging.
69 The CCME provides a soil quality guideline of 140 mg kg⁻¹ in a residential land use setting (CCME,
70 1999). Following its non-threshold toxicity classification, the generic Pb soil guideline value (SGV)
71 (DEFRA & EA, 2002a) was withdrawn in the UK. Whilst a selection of provisional Category 4
72 screening levels (pC4SL) for Pb were recently published (Harries *et al.*, 2013), final C4SLs have not
73 been issued for any soil contaminant. A C4SL denotes a lower tolerable limit for a contaminant in
74 soil, beneath which human health risk is unlikely to be present.

75

76 Although Pb concentrations in the wider environment have declined since its removal from petrol in
77 the last century, its ubiquitous anthropogenic presence still persists in soils, particularly around urban
78 centres (Harries *et al.*, 2013; Appleton *et al.*, 2012a). Previous research worldwide has highlighted
79 areas of elevated soil Pb concentrations outside of areas where geogenic associations are known to
80 exist, including within the Republic of Ireland (ROI) and NI (Barsby *et al.*, 2012; Bourennane *et al.*,
81 2010; Jordan *et al.*, 2007; Ljung *et al.*, 2006; Zhang, 2006). Such findings demonstrate how
82 anthropogenic pollution sources substantially contribute to elevated soil Pb concentrations. In NI Pb
83 is found in highest total and extractable concentrations around the Belfast urban area and in soils
84 overlying mineral deposits, with high soluble Pb measured in peaty upland areas (Jordan *et al.*, 2001).
85 This latter observation may be accounted for by atmospheric deposition of anthropogenic Pb through
86 rainfall, as precipitation is the primary moisture source in upland peat soils.

87 The large surface area and number of acidic functional groups that are common to peat make it an
88 ideal substrate to bind trace elements either as sedimentary, deposited, particulate matter or as sorbed
89 or complexed metal ions (Brown *et al.*, 2000). The major sources of metals in the peat mass of
90 ombrotrophic peat bogs has been shown to come from atmospheric precipitation (Steinnes and
91 Friedland, 2006) which has been specifically illustrated in Ireland (Coggins *et al.*, 2006). Whilst the
92 ability of peat to accumulated trace metals has been well documented there is little data on the
93 bioaccessibility of the trace elements in this peat rich soils.

94
95 Four Pb source domains were previously identified in NI accounting for elevated soil Pb
96 concentrations (McIlwaine *et al.*, 2014). A domain is an area where a distinguishable factor is
97 recognised as controlling the concentration of an element. Urban, peat, granite and mineralisation Pb
98 source domains were related to elevated concentrations of Pb, with typical threshold values (TTV)
99 higher than the TTV calculated for the remaining rural domain. TTVs aim to identify the threshold
100 between diffuse and point source anthropogenic contamination, thereby giving an indication of typical
101 concentrations within defined geographical areas. Urban Pb source domains are likely to be directly
102 attributable to anthropogenic activity. Some anthropogenic pollution sources are potentially more
103 soluble in the environment and resultantly more bioavailable (Ljung *et al.*, 2007; Appleton *et al.*,
104 2012b), in turn posing a greater risk to human health.

105
106 Not all toxins that are rendered bioaccessible will be bioavailable, but *in vitro* bioaccessibility tests
107 can better refine the contaminated land risk assessment process by reducing reliance on total soil
108 contaminant concentrations. Such reliance may overestimate health risks (CIEH, 2009; Scheckel *et*
109 *al.*, 2009; Nathanail & Smith, 2007; Nathanail, 2006; Ruby *et al.*, 1999). The Unified BARGE
110 (Bioaccessibility Research Group of Europe) Method (UBM) is therefore a useful extraction method
111 to employ in risk assessment scenarios where oral contaminant exposure is expected to contribute to
112 toxic health effects. The UBM is a robust soil extraction technique that measures *in vitro* the oral

113 bioaccessibility of contaminants by mimicking the conditions of the human stomach and upper
114 intestine (BARGE/INERIS, 2010). The method has been validated for Pb, arsenic (As) and cadmium
115 (Cd) using *in vivo* swine data (Denys *et al.*, 2012; Caboche, 2009) and has also been subjected to
116 global inter-laboratory trials (Wragg *et al.*, 2011). Data obtained from UBM extractions provide an
117 indication of what fraction of a contaminant may be solubilised in the gastro-intestinal (GI) tract (the
118 bioaccessible portion) and therefore potentially available for absorption resulting in toxic health
119 effects (the bioavailable portion).

120

121 The aim of this research was to measure the oral bioaccessibility of Pb in soil and attribute findings to
122 different Pb sources through solubility estimation and source domain identification. This aim was met
123 through 1) exploratory geochemistry data analysis (EDA) to identify areas of elevated soil Pb
124 concentrations and examine associated spatial structures, 2) comparison of total and extractable Pb
125 concentrations to estimate Pb solubility and 3) source domain identification to determine whether
126 elevated Pb concentrations are the result of geogenic or anthropogenic processes. Lastly, measured
127 oral bioaccessibility was compared across the identified Pb source domains.

128 2. Methodology and Study Area

129 2.1 Study Area

130

131 The study area is located in the UK and Ireland, consisting of NI and neighbouring County (Co.)
132 Monaghan in ROI (Fig. 1). The estimated cumulative population of NI and Co. Monaghan is 1.9
133 million, with a low average population density of 130 per km² (ONS, 2013; CSO, 2011). Current and
134 historical industrial activities concentrated around the Belfast metropolitan area include textiles
135 manufacturing, shipbuilding and aerospace engineering. In addition, quarrying activities are
136 widespread throughout the region with active mines also present, particularly near the Antrim Glens
137 in the northeast (GSNI, 2014). Outside of the larger urban areas of Belfast and Londonderry, land is
138 largely rural and used for agricultural purposes, with metropolitan areas accounting for less than 4%
139 of land use across the study area (European Environment Agency, 2012). As a result, the study area is
140 perceived to be relatively unspoiled from an anthropogenic pollution perspective (Zhang, 2006).

141

142 Soil types present in the study area include peats, humic and sand rankers, brown earths, podzols,
143 mineral gleys and alluviums. Soil pH falls within a narrow acidic range of approximately 5.0 to 6.0
144 (Jordan *et al.*, 2001), with the NI Tellus geochemical survey more recently recording an average pH
145 of 4.7. This decrease in pH over time suggests acidification of soils may be increasing in the study
146 area. The climate is temperate and average annual rainfall ranges from a low of 800 mm in the
147 eastern region to a high of over 1900 mm in the west. The Antrim Glens in the northeast, the Sperrin

148 Mts. in the west and the granitic Mourne Mts. in the southeast intercept much of the precipitation
149 borne by air currents which have travelled over the Atlantic Ocean, although the western half of the
150 study area is most significantly affected by these Atlantic weather patterns (Met Office, 2012).

151

152 2.2 Geochemistry Data Analysis

153

154 Total and extractable Pb concentration data from the NI Tellus and Tellus Border geochemical
155 surveys were provided by the Geological Survey of Northern Ireland (GSNI) and by the Geological
156 Survey of Ireland (GSI), respectively. Rural NI Tellus Survey soil samples were collected on a 2 km²
157 grid at depths of 5 – 20 cm ('A') and 35 – 50 cm ('S'). 'A' samples were analysed for total Pb
158 concentrations by x-ray fluorescence spectrometry (XRFS) and for extractable concentrations by
159 inductively coupled plasma mass spectrometry (ICP-MS) following an *aqua regia* digest. 'S' NI
160 Tellus Survey soil samples were also digested by *aqua regia* and analysed by ICP-MS. Tellus Border
161 'A' samples were collected on a 4 km² grid and analysed only by ICP-MS following an *aqua regia*
162 digest to yield extractable concentration data. Full analytical and field methods employed by these
163 comprehensive regional geochemical surveys can be found in Smyth (2007) and Knights and Glennon
164 (2013).

165

166 As part of this research, additional XRFS analysis was conducted at the British Geological Survey
167 (BGS) Analytical Geochemistry Facility on a sub-set of 18 Tellus Border 'A' samples in Co.
168 Monaghan according to the same methods described in the NI Tellus Survey methodology (Smyth,
169 2007). The additional XRFS data was required for solubility estimation and for calculation of UBM
170 bioaccessible fractions (BAF) in Co. Monaghan. Geochemistry data were handled in SPSS v.19.0, R
171 (R Core Team, 2013) and MS Excel 2010.

172

173 2.3 Geostatistical Analysis and Interpolation

174

175 Interpolation by ordinary kriging (OK) (Matheron, 1965) was conducted in ArcMap 10.0 (ESRI,
176 2010). The OK model yielding a mean prediction error closest to zero was selected as the final model
177 for generating interpolated surfaces and geostatistics (Lloyd, 2010). Geostatistical outputs can be
178 influenced by a nonparametric data distribution (Lloyd, 2010; Clarke, 2001; Einax & Soldt, 1999).
179 Total and extractable Pb concentration data were therefore log-transformed prior to interpolation. OK
180 models were checked for robustness using cross validation statistics and a visual assessment of the
181 best fit semi-variogram using a maximum search neighbourhood of 12 nearest sample locations.

182

183 Semi-variogram parameters give an indication of the spatial structure that exists within a data set.
184 This in turn can help explain geochemical or environmental processes that affect the spatial
185 distributions of elements (Goulard & Voltz, 1992; McBratney *et al.*, 1982). The semi-variogram sill
186 (C_1) is synonymous with the sample variance and represents the maximum variance that exists
187 between measured sample values within the range of spatial correlation (a). Beyond the distance a ,
188 samples are no longer spatially correlated (Clarke, 2001; Gringarten & Deutsch, 2001). The nugget
189 variance (C_0) is attributed to micro-scale variance outside of sampling resolutions. Although the
190 nugget effect is commonly regarded as an indication of measurement error or random semi-variogram
191 behaviour, micro-scale processes which control element distributions may also be accounted for by
192 the nugget variance. For example, Imrie *et al.* (2008) found that factors attributed to anthropogenic
193 land use patterns were accounted for by a nugget effect. Dobermann *et al.* (1995) concluded buffalo
194 excrement influenced soil chemistry over a range that occurred within the nugget variance. Functions
195 with a high proportion of total variance ($C_0 + C_1$) accounted for by the nugget variance may therefore
196 be indicative of anthropogenic processes or land use behaviours which are significantly affecting
197 element distributions but occurring over short spatial scales not detected by the primary range (a) of
198 the function.

199

200 OK yields results which increase in accuracy in line with increasing sample numbers (Einax & Soldt,
201 1999). Due to lower sample numbers than were available for total and extractable Pb concentration
202 data, bioaccessible Pb concentrations were interpolated using inverse distance weighting (IDW) with
203 a maximum search neighbourhood of five neighbouring sample locations. IDW is an exact
204 interpolator (Lloyd, 2010) and this method therefore yielded a more accurate range of Pb
205 bioaccessible concentration values across the interpolated surface.

206

207 2.4 Pb Solubility Estimation

208

209 A method for estimating element solubility in soil at a regional scale was applied to the NI Tellus and
210 Tellus Border XRFS and ICP-MS data, similar to approaches used previously in Finland (Jarva *et al.*,
211 2009) and Cyprus (Cohen *et al.*, 2012). XRFS measures total element concentrations in soils whereas
212 ICP concentrations rely on the antecedent *aqua regia* acid extraction. Although *aqua regia* acid is
213 said to effectively leach many metals (Gill, 1997), the solubility of elements will affect how easily
214 they are leached from the soil (Delgado *et al.*, 2011). Therefore, by comparing the concentrations
215 measured by the two methods, element solubility at a regional scale can be estimated. Elements
216 which are more soluble in the environment generally exhibit higher oral bioaccessibility (Finžgar *et*
217 *al.*, 2007).

218

219 ICP extractable concentrations were plotted against XRFS total concentrations using the R statistical
220 software package (R Core Team, 2013) to explore the relationship between the two analytical
221 methods. The ratio of XRFS/ICP Pb concentrations was mapped by OK to illustrate geographical
222 trends in Pb solubility. The classes on the map were defined by the boxplot classes method
223 (McIlwaine *et al.*, 2014) with an additional class added where the ratio was equal to one, i.e. where
224 the two analytical methods are equal. Boxplot classifications retained the appropriate amount of
225 detail to allow a direct comparison with the mapped bioaccessibility results.

226

227 2.5 Pb Domain Identification

228

229 Domains were previously identified for Pb in NI as described in McIlwaine *et al.* (2014). Total XRFS
230 concentrations in shallow soils were mapped using empirical cumulative distribution function (ECDF)
231 classes and compared to the main factors identified as controlling element concentrations—bedrock
232 geology, superficial geology, land use classification and mineralisation. Elevated concentrations of
233 Pb were attributed to urban, granite, mineralisation and peat source domains in NI with the remaining
234 rural domain hosting lower Pb concentrations.

235

236 Additional Co. Monaghan data were obtained to identify Pb domains across the extent of the study
237 area for this research. Bedrock geology data were obtained from the GSI 1:500000 Bedrock Geology
238 map (GSI, 2005). Superficial peat cover was identified using the Irish Environmental Protection
239 Agency Soils and Subsoils Mapping Project data completed by Teagasc (Fealy & Green, 2009). The
240 Corine land cover data (European Environment Agency, 2012) was used to identify urban and rural
241 land use within the study area. Areas of known or suspected mineralisation in Co. Monaghan are
242 identified in the Tellus Border prospectivity map (Coulter and Stinson, 2013).

243

244 Data used for identifying the mineralisation source domain in NI and Co. Monaghan relied upon
245 prospectivity maps (Coulter & Stinson, 2013; Lusty *et al.*, 2012) and not the locations of working or
246 historic mines. The mineralisation domain and associated soil Pb is therefore regarded as geogenic
247 and naturally occurring for the purposes of this research.

248

249 2.6 Oral In Vitro Bioaccessibility Testing

250

251 UBM extractions were carried out in 2009 (Barsby *et al.*, 2012) and 2013 at the BGS Analytical
252 Geochemistry Facility following the published method (BARGE/INERIS, 2010). The 2009 and 2013
253 data sets were joined to create a UBM data set of 163 samples for this research. Samples (< 2 mm

254 fraction) were selected from the NI Tellus Survey and Tellus Border soil archives to cover a wide
255 range of soil and underlying bedrock types present in the study area.

256

257 Standard BGS internal laboratory procedures were followed during UBM extraction and analysis
258 according to UK Accreditation Service national laboratory requirements. Reagents were sourced by
259 BGS from Merck, Sigma, Baker and Carl Roth. Saliva, gastric, duodenal and bile solutions were
260 prepared one day prior to soil extractions to permit stabilisation. Solution pH was adjusted as
261 required according to UBM specifications using either 37% HCl or 1M HNO₃ (Table 1). Soils not
262 adhering to pH specifications (pH < 1.5) after one hour of gastric extraction were discarded and re-
263 extracted at a later date.

264

265 Extracts were analysed by an Agilent 7500cx series ICP-MS employing an octopole reaction system
266 in combination with a CETAC autosampler. The instrument was calibrated at the beginning of every
267 analytical run using a minimum of three standards and one blank for each trace element. Multi-
268 element quality control check standards were analysed at the start and end of each run and after every
269 25 samples at minimum.

270

271 One blank, one duplicate and one certified BGS102 reference soil (Wragg, 2009) were included in the
272 extraction run for each of seven soils extracted in 2013. The BGS102 certificate of analysis provides
273 certified UBM values for acceptable ranges of gastric (G) Pb concentrations. Average measured G Pb
274 in reference soils was within one standard deviation of the certified BGS102 value. The mean relative
275 per cent difference (RPD) for gastric Pb in study area soil samples was 8%. In line with the available
276 BGS102 certified reference value for G Pb, G data are presented in the following results as it is
277 common practice to report the results yielding the highest bioaccessibility to ensure health risks are
278 not underestimated. This approach also adheres to the precautionary principle advocated by UK
279 contaminated land legislation and guidance (DEFRA, 2012). Details of quality control for 2009
280 extractions are similar to the above and are described in detail in Barsby et al. (2012).

281

282 **3. Results**

283 3.1 Lead Soil Concentrations

284

285 The highest extractable Pb concentrations are found in soils along the northeast coast near the Antrim
286 Glens, extending south into the Belfast metropolitan area and Ards Peninsula, with the occurrence of
287 elevated Pb concentrations continuing along the southeast NI-Co. Monaghan border (Fig. 1, Fig. 2A).
288 Peat soils overlying the Sperrin Mts. in the northwest also host elevated concentrations of Pb. It is
289 this part of the study area that receives the most precipitation borne from Atlantic Ocean air currents

290 (Met Office, 2012). The maximum measured extractable Pb concentration occurs near the greater
291 Belfast metropolitan area ($> 3000 \text{ mg kg}^{-1}$).

292

293 Although no SGV is currently available for Pb in the UK, Table 2 provides an overview of how Pb
294 concentrations compare against historic withdrawn and current provisional soil screening criteria.
295 Fig. 2B illustrates where extractable Pb concentrations exceed the lowest published pC4SL of 30 mg
296 kg^{-1} by at least 10%. This criterion is applicable for a female child in an allotment setting (Harries *et*
297 *al.*, 2013). Pb concentrations were flagged as exceeding the pC4SL only where the concentration met
298 or exceeded 33 mg kg^{-1} to ensure the measured Pb concentration was sufficiently above the pC4SL.
299 Extractable Pb concentrations exceed the pC4SL at over 2,208 of 7,234 NI Tellus and Tellus Border
300 soil sample locations in the study area (Fig. 2B). When total XRFS Pb concentrations are compared
301 with the pC4SL, the number of occurrences where the screening criterion is exceeded increases to
302 2,629 (not illustrated).

303

304 Given the withdrawn and provisional nature of the Pb SGV and pC4SL, respectively, it would not be
305 possible to assess the potential level of health risk from the values presented in Table 2 and Figs. 2A-
306 B alone. Although Fig. 2B shows a geographically widespread occurrence of extractable Pb
307 concentrations exceeding the lowest published pC4SL, it is important to note that individual sites
308 must be assessed on a case by case basis taking relevant land use scenarios and all likely risk exposure
309 pathways into account.

310

311 XRFS Pb concentrations were mapped previously by Barsby *et al.* (2012) and exhibit similar spatial
312 patterns to extractable Pb concentrations. Extractable Pb concentrations are controlled by a spatial
313 function with a moderate range (a) of 22.8 km. Short to medium range spatial functions are sometimes
314 associated with processes that have a higher frequency of variation over short distances. Such
315 functions can be the result of smaller scale processes such as anthropogenic interactions with the
316 environment,

317 while long range functions capture the effects of larger scale geologic forming processes (Imrie *et al.*,
318 2008; Dobermann *et al.*, 1995). Pb exhibits a spatial structure in the study area that varies over a short
319 scale in terms of its range relative to trace elements of known geogenic origin such as nickel or
320 chromium which are controlled by longer range functions ($>70 \text{ km}$; McIlwaine *et al.*, 2014; Barsby *et*
321 *al.*, 2012). The high proportion of nugget variance (63%) for Pb spatial distributions (Table 3) also
322 suggests a high degree of micro-scale variation or spatial variability not detected by the primary range
323 of the Pb function.

324

325 Fig. 3 illustrates the difference between Pb extractable concentrations in NI Tellus Survey 'A' soils
326 and 'S' soils as measured by ICP-MS following an *aqua regia* digest. Pb is present at higher average

327 and maximum concentrations in 'A' soils than 'S' soils. Anthropogenic and atmospheric Pb
328 deposition to soil is expected to be most pronounced at surface level ('A').

329

330 3.2 Lead Solubility and Domain Identification

331

332 Figs. 4 and 5 illustrate comparative differences in Pb extractable and total concentrations. Such
333 information can provide insight into contaminant sources. For example, geogenic metals are often
334 highly insoluble and exhibit lower bioaccessibility (Cox *et al.*, 2014) whilst other forms of
335 anthropogenic pollution tend to be more soluble and more bioaccessible (Ljung *et al.*, 2006).

336

337 Fig. 4 plots the relationship between XRFS and ICP Pb concentrations, with a 1-1 ratio represented by
338 the dashed line shown on the scatterplot. Although soil analysis by XRFS detects an additional
339 insoluble portion of Pb, the cluster of most points around the 1-1 line shows that a significant
340 proportion of total Pb soil concentrations was detectable by ICP, with XRFS concentrations exceeding
341 ICP concentrations by no more than 15%. This suggests the majority of Pb in soil is soluble and not
342 encapsulated by an insoluble mineral matrix. Pb encapsulated by insoluble minerals generally
343 displays decreased bioavailability and bioaccessibility (Ruby *et al.*, 1999).

344

345 Fig. 5 illustrates the geographic variability in XRFS/ICP concentrations ratios. Higher levels of Pb
346 solubility occur in the darker areas of the map, where the ratio is less than one. Where the map
347 becomes lighter Pb is less soluble. Higher levels of solubility are observed along the central and
348 western NI-ROI border and throughout the eastern coast. One area of higher solubility strongly aligns
349 with an identified mineralisation source domain (Fig. 6A). Higher proportions of insoluble Pb occur
350 in the southeast and northwest near the Mourne and Sperrin Mts., respectively, with the Mourne Mts.
351 comprising the granite source domain and the Sperrin Mts. and associated geology aligning with the
352 peat source domains. Rural, peat and urban domains host moderately soluble portions of Pb.
353 Although an elevated peat source domain was also identified in Fig. 6A, Pb solubility trends in Fig. 5
354 do not clearly align spatially with patterns illustrated for the peat source domain. Instead Pb solubility
355 in peat is comparable to the intermediate solubility observed within urban source domains.

356

357 3.3 Lead Bioaccessibility

358

359 The results of UBM extractions are summarised in Table 4. G bioaccessibility was higher than
360 gastro-intestinal (GI) bioaccessibility due to the lower pH of the G digestion which increases Pb
361 mobility in solution (Denys *et al.*, 2012; Farmer *et al.*, 2011; Denys *et al.*, 2007). The maximum G
362 bioaccessible concentration was 199.8 mg kg⁻¹, accounting for 68.6% of total Pb. The median G ICP-

363 BAF was 40.3%, decreasing to 15.6% of extractable Pb concentrations in the GI phase. XRFs-BAF
364 values Pb did not differ greatly from ICP-BAFs as a result of most Pb in soils in the study area being
365 detectable by ICP-MS (Fig. 4). Pb gastric bioaccessibility exceeded 50% of total concentrations at 13
366 different soil locations across the study area (Fig. 6A).

367

368 Fig. 6A and Table 5 compare elevated Pb source domains with measured gastric oral bioaccessibility.
369 Across the five source domains (inclusive of the rural domain), the mean ICP-BAF range was 35.6% -
370 46.4%. The highest maximum BAFs and the highest mean and maximum bioaccessible
371 concentrations occurred in the mineralisation domain. Despite insoluble portions of Pb observed near
372 the granite domain (Fig. 5), the highest average BAFs were measured in soils overlying this domain.
373 Rural areas had the lowest mean and minimum bioaccessible Pb concentrations and the lowest
374 average BAFs (Table 5). Urban domains accounted for the second highest maximum bioaccessible
375 Pb concentrations, although peat and urban domains each hosted intermediate levels of bioaccessible
376 Pb in general when compared to the other source domains (Table 5). Where small urban domains
377 overlapped with the extent of the mineralisation domain, it was assumed mineralisation acted as the
378 primary Pb source and samples were assigned to the mineralisation domain.

379

380 Despite the lower solubility of Pb in soils overlying the Sperrin Mts. (Fig. 5), Pb from the peat source
381 domain present in this area is still moderately bioaccessible (Fig. 6A, Table 5). Similarities in Pb
382 bioaccessibility between the peat and urban domains may suggest that bioaccessible Pb in these
383 domains arises from similar sources, such as atmospheric deposition from urban or industrial
384 emissions. Alternatively, this observation in peat may be coincidental and instead governed by the
385 presence of dissolved organic matter, low pH and reducing conditions in peat soils that are conducive
386 to higher levels of trace element mobility and bioaccessibility (Appleton *et al.*, 2013; Palmer *et al.*,
387 2013; Yang *et al.*, 2003).

388

389 Regional trends in measured gastric bioaccessible Pb concentrations are illustrated by Fig. 6B.
390 Higher levels of gastric bioaccessibility are present around the Belfast metropolitan area, along the
391 extent of the NI-ROI border, and also along the northeast coast. In addition to a peat source domain
392 immediately north of this latter coastal location, mining activity occurs in this area (GSNI, 2014),
393 although a Pb mineralisation domain is not present. Another area where measured bioaccessible Pb
394 concentrations are high is south of Lough Neagh in proximity to an urban source domain. In general,
395 observed spatial patterns in Pb bioaccessibility closely align with those observed for elevated Pb soil
396 concentrations (Fig. 2A), areas of higher Pb solubility (Fig. 5) and also with elevated mineralisation,
397 urban, and peat Pb source domains (Fig. 6A). These findings may suggest that both diffuse
398 anthropogenic and widespread geogenic Pb sources are capable of presenting health risks from the
399 oral exposure pathway.

400 4. Discussion

401

402 Part IIA of the 1990 Environmental Protection Act (EPA 1990) outlines the statutory obligations in
403 England, Wales and Scotland for assessing potentially contaminated areas of land to determine if sites
404 are fit for proposed land use. At the time of writing no cohesive contaminated land legislative
405 framework has been officially adopted in NI or ROI for assessing potential risks to human health.
406 Guidance on the NI Environment Agency (NIEA) web site directs users to English Environment
407 Agency (EA) publications as official adoption and enforcement of Part 3 of the enacted Waste and
408 Contaminated Land Order (NI) 1997 has yet to occur (NIEA, 2010). The Irish Environmental
409 Protection Agency is currently in the process of developing its own framework.

410

411 Toxic elements from some types of anthropogenic pollution may be more bioaccessible than those
412 associated with geogenic sources due to the more soluble phases in which they exist in soil (Appleton
413 *et al.*, 2012b; Cave *et al.* 2007; Ljung *et al.*, 2007; Cave *et al.* 2003), although this study also found
414 that Pb attributed to geogenic sources displayed higher average BAFs than Pb from other source
415 domains. Despite the knowledge that soluble and anthropogenic forms of pollution in the
416 environment may be more likely to cause harm due to their increased bioavailability and
417 bioaccessibility, sections 3.21 - 3.26 of the 2012 DEFRA guidance for Part IIA of EPA 1990 state that
418 soils hosting widespread geogenic contamination or diffuse anthropogenic pollution should not be
419 regarded as contaminated land. The exception is where strong scientific evidence concludes that
420 significant health risks are being caused or are likely to occur (DEFRA, 2012). This approach is not
421 unique to the UK. For example, a similar regime is in place in Finland, where a Government Decree
422 on the Assessment of Soil Contamination and Remediation Needs (214/2007) (Ministry of the
423 Environment Finland, 2007) states that the assessment process shall regard natural geological
424 concentrations and diffuse anthropogenic pollution as contributing to background concentrations
425 (Jarva *et al.*, 2010). However, such guidance may be misaligned with our knowledge concerning the
426 health effects from oral Pb exposure in soil, particularly with regard to its non-threshold toxicity
427 (ASTDR, 2007; USEPA, 1988). Gathering more evidence on other risk pathways for Pb exposure
428 such as inhalation would help underpin with more certainty the potential health effects from exposure
429 to low level diffuse anthropogenic pollution or widespread geogenic contamination.

430

431 The high solubility of Pb in surface soils and reduced Pb concentrations in deep soils in the study area
432 suggests that a portion of elevated Pb concentrations is from diffuse anthropogenic pollution sources.
433 This finding is supported by the observed spatial trends in Pb soil distributions where elevated
434 concentrations align with urban and peat source domains. Upland peat soils may be intercepting
435 anthropogenic Pb carried in rainfall. The medium range spatial structure observed for extractable Pb
436 concentrations also supports the conclusion that anthropogenic processes may be influencing or have

437 historically influenced Pb soil concentrations. Bioaccessibility in the urban domain was higher than
438 that observed in the remaining rural domain, demonstrating the anthropogenic effects of industrial
439 activity and higher population densities over Pb distributions and associated possible health effects.

440

441 From an oral risk exposure standpoint, the region identified as a mineralisation domain hosted the
442 highest concentrations of bioaccessible Pb. Although the granite domain accounted for lower
443 maximum levels of bioaccessibility compared to the other domains, average BAFs were highest in
444 soils overlying the granite domain. These findings suggesting that risk associated with geogenic
445 sources of Pb should also be taken into consideration.

446 5. Conclusion

447

448 Combining existing knowledge surrounding non-threshold toxicity with the findings that Pb in the
449 study area displays moderate to high solubility and oral bioaccessibility and warrants more detailed
450 risk evaluation for Pb in soil. The findings of this study should be taken into account during the
451 development of final Pb soil screening levels and the adoption of an official Irish or Northern Irish
452 contaminated land regime, if or when such measures take place.

453

454 Conclusions regarding toxicity risks from oral soil Pb exposure can only be made on a site specific
455 basis taking all exposure pathways and relevant land use scenarios into account. However, the
456 findings of this research suggest that diffuse anthropogenic forms of pollution and the presence of
457 natural geogenic contaminants should be considered more carefully in a health risk context,
458 particularly in the case of a non-threshold toxin such as Pb.

459

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461

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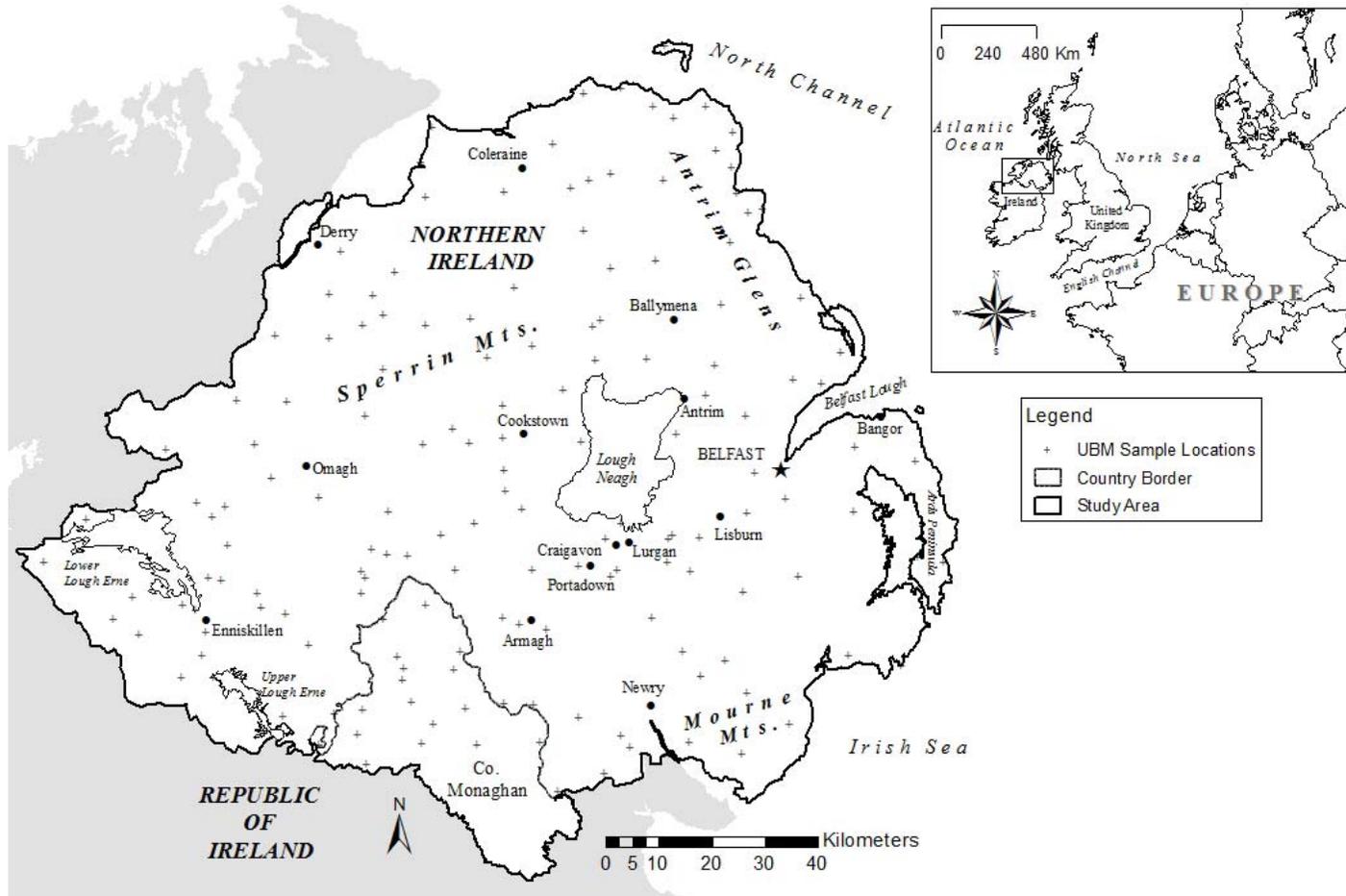
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760 Figures

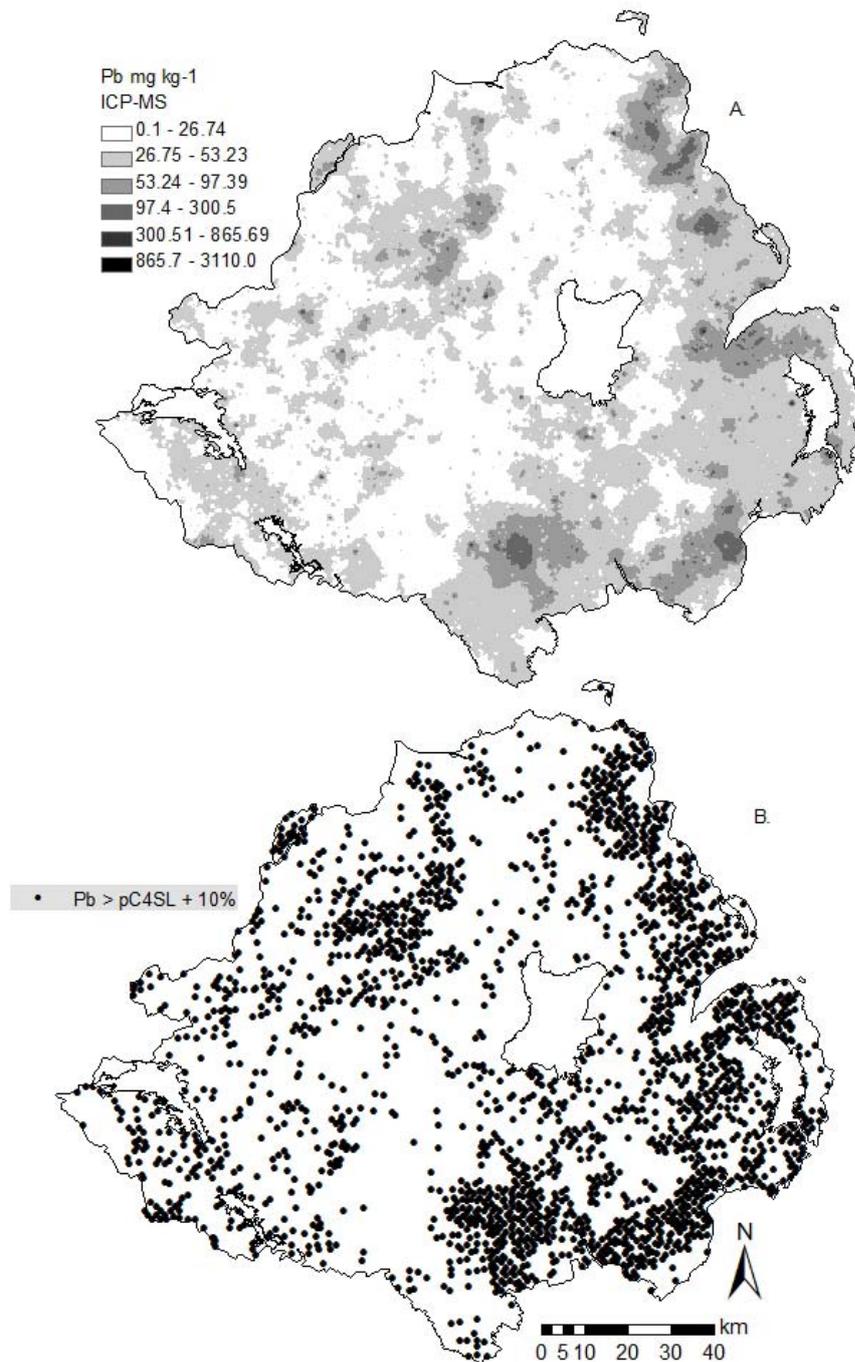
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763 Fig. 1 Study area map showing location in Europe and 163 UBM sample locations across Northern Ireland and Co. Monaghan in the Republic of Ireland

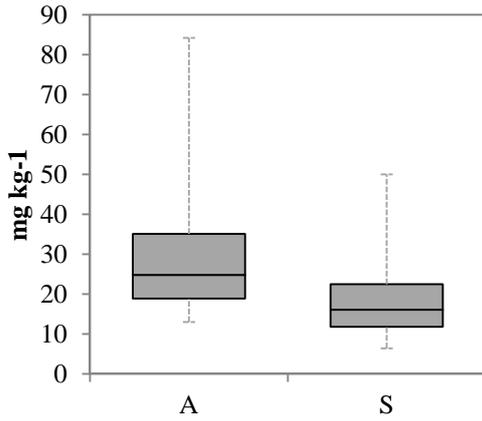
764 (ROI); Northern Ireland images are GSNI Crown Copyright



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767 Fig. 2 (A) ICP-MS Pb concentrations (mg kg^{-1}) in 'A' soils across study area and (B) soil sample
 768 locations from NI Tellus and Tellus Border geochemical surveys where 'A' soil concentrations
 769 exceed the lowest published pC4SL of 30 mg kg^{-1} by 10% or greater ($n = 2,208$) for a female child
 770 receptor in an allotment setting (Harries *et al.*, 2013)

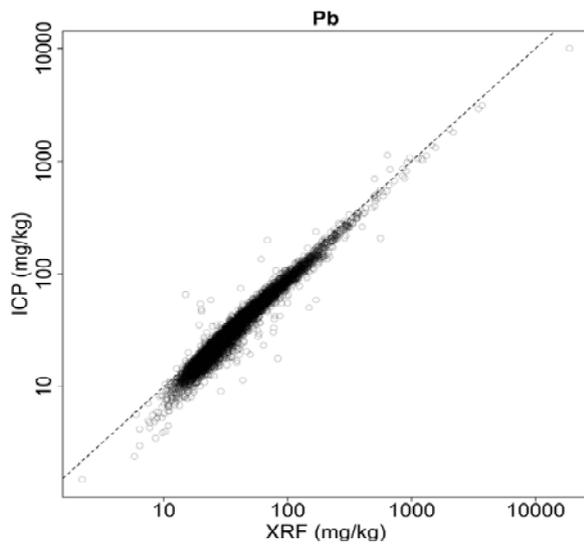
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773 Fig. 3 Surface ('A') Pb extractable concentrations versus beneath-surface ('S') Pb extractable
 774 concentrations as measured by ICP-MS following an *aqua regia* digest in Northern Ireland. Lower
 775 and upper error bars represent 5th and 95th percentile ranges, respectively.

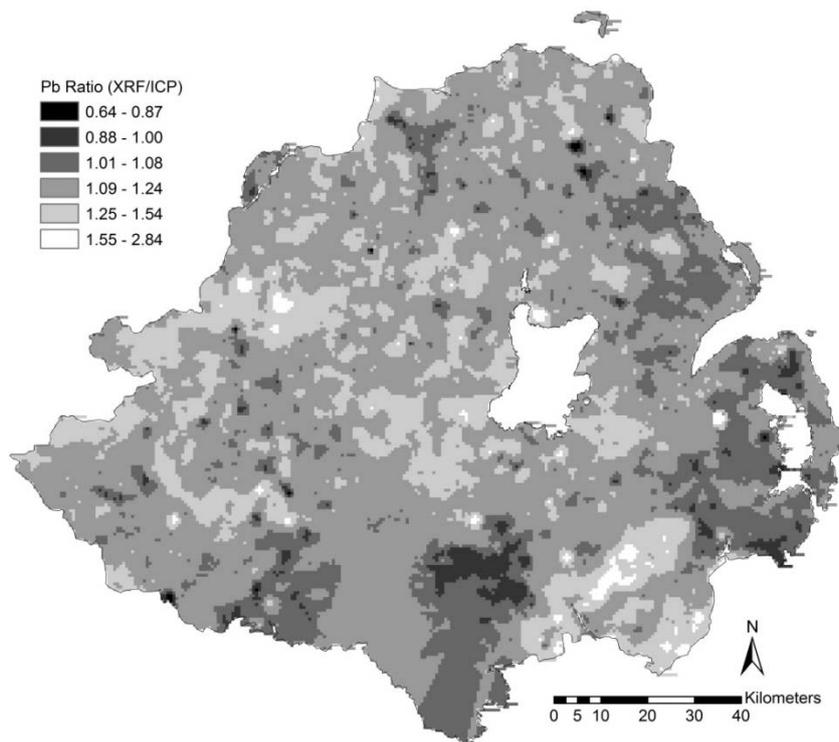
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778 Fig. 4 Pb extractable concentrations in 'A' soils measured by ICP-MS plotted against total Pb XRFs
 779 concentrations in NI and Co. Monaghan; dashed line indicates a 1-1 ratio between the two
 780 concentrations

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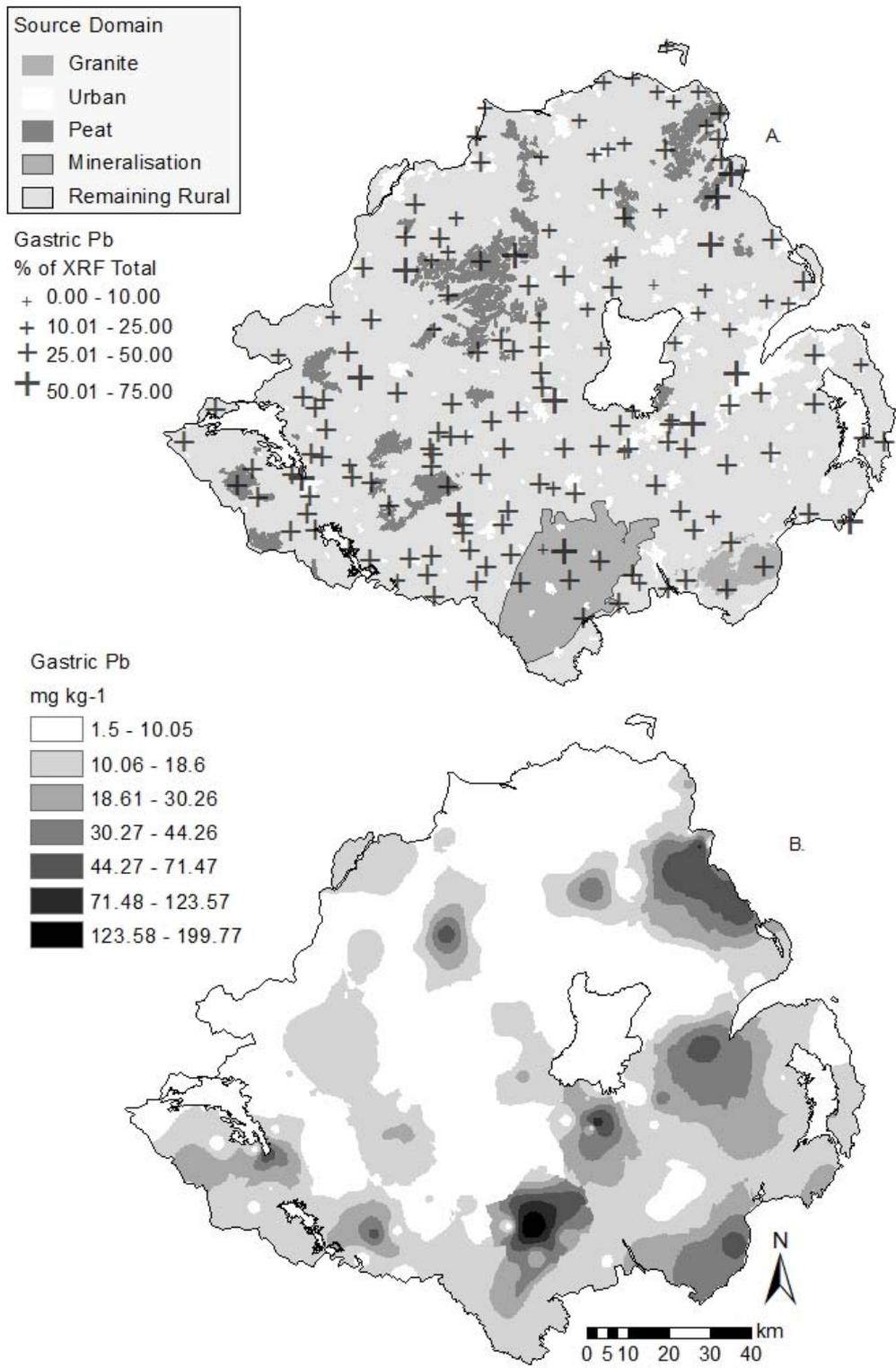


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783 Fig. 5 Map of XRF/ICP extractable Pb concentrations; high ratios illustrate areas of lower Pb
 784 solubility and low ratios illustrate where high Pb solubility exists

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Fig. 6 (A) Elevated Pb source domains within study area identified by ECDF method overlain with measured Pb G BAFs expressed as percentage of total XRFs concentrations and (B) Pb oral bioaccessibility (mg kg⁻¹) interpolated by IDW, 5 nearest neighbours; *n* = 163

794 Tables

795

796 Table 1 pH tolerances for stabilised UBM digestive fluids

Solution	pH Tolerance
Saliva	6.5 +/- 0.5
Gastric	1.0 +/- 0.1
Duodenal	7.4 +/- 0.2
Bile	8.0 +/- 0.2

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798

799 Table 2 Summary of study area and sample set 'A' soil Pb concentrations compared against historic
800 and provisional generic UK soil assessment criteria and study area typical threshold values (TTVs);
801 all values in mg kg⁻¹

Historic UK SGV ^a	pC4SL ^b	Study Area Source Domain TTVs ^c		Pb 'A' Soil Concentrations						
				Method	Study Area <i>n</i> = 7,234 ^d			Study Sample Set <i>n</i> = 163 ^e		
					Mean	Med	Max	Mean	Med	Max
Res/Allot		Mineral	Granite	XRFS	41.6	28.8	18,756	39.8	28.7	291.0
450	30-330	110	170							
Commercial		Peat	Urban	ICP-MS	31.5	23.2	3,110	35.4	23.5	268.0
750	1100-6000	160	220							
		Remaining Rural								
63										

802 ^aDEFRA & EA, 2002a & 2002b

803 ^bRange of values dependent on different modelled exposure and land use scenarios as presented in Harries *et al.* (2013). Res = residential,
804 allot = allotment.

805 ^cThe unique values calculated for this study incorporating 18 sample locations in Co. Monaghan did not yield different TTV results from
806 McIlwaine *et al.*, 2014. Mineral = mineralisation domain.

807 ^dBased on NI Tellus and Tellus Border geochemical survey data where 7,234 samples were analysed by ICP-MS in NI and Co. Monaghan.
808 6,862 were analysed by XRFS in NI with 18 additional samples in Co. Monaghan analysed by XRFS outside of routine Tellus Border
809 survey analyses.

810 ^eInclusive of 90 samples from Barsby *et al.*, 2012

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813 Table 3 Pb geostatistical summary showing 98% of total variance in Pb extractable soil distributions
814 is accounted for by a short-range function as modelled in Fig. 2 while total concentrations show Pb
815 concentrations are controlled by a longer range function suggestive of geogenic processes

	Nugget Variance C_0	Function Variance C_1	Range (km) a	Total Variance $C = C_0 + C_1$	Unexplained Variance $\% C_0 = (C_0/C_1)*100$
Extractable Pb (ICP)	0.004	0.197	1.2	0.201	2%
Total Pb (XRFS)	0.193	0.100	32.9	0.293	66%
XRFS/ICP Ratio	0.004	0.014	3.5	0.018	21%

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818 Table 4 Summary of gastric (G) and gastro-intestinal (GI) Pb bioaccessible concentrations (mg kg⁻¹)
 819 and bioaccessible fractions (BAF, %) in study area (*n* = 163)

	mg kg ⁻¹			XRFS-BAF			ICP-BAF		
	Med	Max	Min	Med	Max	Min	Med	Max	Min
G Pb	8.6	199.8	1.5	33.9	68.6	8.2	40.3	74.6	9.7
GI Pb	3.8	85.9	0.0	12.8	35.1	0.2	15.6	38.1	0.3

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822 Table 5 Comparison of Pb G bioaccessible concentrations (mg kg⁻¹) and BAF (%) against identified
 823 Pb concentrations in 'A' soils overlying five Pb source domains in study area

Domain	Mean	G Pb			ICP-BAF			XRFS-BAF		
		Mean	Max	Min	Mean	Max	Min	Mean	Max	Min
Mineralisation <i>n</i> = 9	36.4	199.8	6.9	42.7	74.6	15.8	37.3	68.6	8.2	
Granite <i>n</i> = 3	36.0	49.0	16.3	46.4	52.2	37.5	40.2	46.8	34.4	
Peat Soil <i>n</i> = 18	20.6	74.9	2.6	42.6	66.2	13.4	37.3	64.4	12.3	
Urban <i>n</i> = 31	19.9	97.6	2.3	41.7	67.1	14.3	36.4	64.0	13.0	
Remaining Rural <i>n</i> = 102	9.7	51.3	1.5	35.6	65.8	9.7	30.5	58.9	9.8	

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