

Tolerance of soil bacterial community to tetracycline antibiotics induced by As, Cd, Zn, Cu, Ni, Cr and Pb pollution

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Abstract. The widespread use of both heavy metals and antibiotics in livestock farming, followed by their subsequent arrival on agricultural soils through manure/slurry spreading, has become a problem of vital importance for human health and the environment. In the current research, a laboratory experiment was carried out for 42 days to study tolerance and co-tolerance of three tetracycline antibiotics (tetracycline, TC; oxytetracycline, OTC; chlortetracycline, CTC) in soils polluted with heavy metals (As, Cd, Zn, Cu, Ni, Cr and Pb) at high concentrations (1000 mg kg⁻¹ of each one, separately). Pollution Induced Community Tolerance (PICT) of the bacterial community was estimated using the leucine incorporation technique. The Log IC₅₀ (logarithm of the concentration causing 50% inhibition in bacterial community growth) values obtained in uncontaminated soil samples for all the heavy metals tested showed the following toxicity sequence: Cu>As>Cr≥Pb≥Cd>Zn>Ni. However, in polluted soil samples the toxicity sequence was Cu>Pb≥As≥Cd≥Cr≥Ni≥Zn. Moreover, at high heavy metal concentrations the bacterial communities showed tolerance to the metal itself, this taking place in the long term for all the metals tested. The bacterial communities of the soil polluted with heavy metals showed also long-term co-tolerance to TC, OTC, and CTC. This kind of studies, focusing on the eventual increases of tolerance and co-tolerance of bacterial communities in agricultural soil, favored by the presence of different kinds of pollutants, is of crucial importance, mostly bearing in mind that the appearance of antibiotic resistance genes in soil bacteria could be transmitted to human pathogens.

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32 **Keywords:** Chlortetracycline; heavy metals; trace elements; oxytetracycline; PICT; tetracycline

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34 1. Introduction

35 The accumulation of heavy metals in soils is a widespread problem all over the world. Some
36 anthropogenic activities such as mining, industrial production, agriculture, or livestock farming, are among the
37 main sources of release of heavy metals to the terrestrial environment (Abdu et al., 2017). The use of heavy
38 metals in livestock production is a common practice, since commercial feeds are enriched with essential
39 elements (such as Cu and Zn) to prevent diseases and obtain an optimal growth rate (Sager, 2007; Wu et al.,
40 2013). In fact, the of presence heavy metals in livestock manure can be seen as generalized, with really high
41 concentrations in swine manure reported by Hölzel et al. (2012) for Zn (up to 8239 mg kg⁻¹), and for Cu (up to
42 3387 mg kg⁻¹), while Wang et al. (2013) found up to 1601.7 mg kg⁻¹ for Cr in layer manure, Zhang et al. (2012)
43 detected up to 300 mg kg⁻¹ for Pb, and up to 100 mg kg⁻¹ for Ni, in poultry manure, and finally Liu et al. (2020)
44 reported up to 59.7 mg kg⁻¹ for Cd, and up to 89.3 mg kg⁻¹ for As, in pig manure. In view of that, these and
45 other heavy metals are considered a serious problem for public health, especially because these toxic elements
46 tend to accumulate in soils, where they may reach very high and progressively increasing concentrations
47 (Kabata-Pendias, 2000; Huang et al., 2007). Moreover, veterinary antibiotics are also widely used for the
48 treatment of bacterial infections (Boxal et al., 2003), and their consumption has been increasing in recent
49 decades, with the estimate of world use for 2030 being 105,596 tons (Van Boeckel et al., 2015). The antibiotics
50 most widely used in veterinary medicine in the European Union are tetracyclines (TCs), and specifically
51 tetracycline (TC), oxytetracycline (OTC) and chlortetracycline (CTC) (European Medicines Agency, 2016).
52 Both heavy metals and antibiotics for veterinary use are poorly absorbed by the intestines of animals, causing
53 that a high percentage of them is excreted in feces and urine (Kornegay et al., 1976; Sarmah et al., 2006). In
54 this sense, it has been reported that tetracycline antibiotics may reach in manures concentrations as high as 746
55 mg kg⁻¹ for CTC (Pan et al., 2011), 211 mg kg⁻¹ for OTC, and 300 mg kg⁻¹ for TC (Widyasari-Mehta et al.,
56 2016). The presence of high levels of heavy metals and veterinary antibiotics in farmland soils is due to the
57 repeated applications of manure and slurries (as well as sewage sludge) as organic fertilizers (Hamscher et al.,
58 2002; Nicholson et al., 2003). Once in the soil, these compounds can interact with soil microbial communities
59 and modify their structure and function (Hattori, 1992; Thiele-Bruhn and Beck, 2005; Chien et al., 2008; Giller
60 et al., 2009; Caban et al., 2018).

61 The increase of the concentration of any pollutant in soils may suppose a selection pressure for soil
62 bacterial communities, causing tolerance to that pollutant (Blanck, 2002). This effect may be useful to quantify
63 the harmful repercussions produced by pollutants on soil bacteria and is called pollution-induced community

64 tolerance (PICT). Agricultural soils, highly influenced by anthropogenic activities, deserve special attention
65 since they are recognized as the largest reservoirs of antibiotic-resistant genes, receiving antibiotics from
66 veterinary use (through repeated applications of manure and slurries, as indicated above), as well as heavy
67 metals (Ji et al., 2012). Therefore, the resistance of soil bacterial communities to antibiotics has become a
68 crucial threat at a world scale, and the study of whether bacterial communities generate co-tolerance to
69 antibiotics in the presence of heavy metals (as well as on how and in which degree is developed) is of vital
70 importance.

71 Several previous studies have focused on co-tolerance among different heavy metals and antibiotics (Berg et
72 al., 2010; Sarma et al., 2010; Fernández-Calviño and Bååth, 2013; Song et al., 2017; Santás-Miguel et al.,
73 2020a; Zhong et al., 2021). Specifically, Fernández-Calviño and Bååth (2013) evaluated the tolerance of soil
74 bacterial community in one soil polluted with different Cu concentrations (2-32 mmol Cu kg⁻¹) to three
75 antibiotics (vancomycin, tetracycline and tylosin) and observed an increase in the co-tolerance of soil bacterial
76 communities to tetracycline (at ≥ 8 mmol Cu kg⁻¹), tylosin (at ≥ 16 mmol Cu kg⁻¹) and vancomycine (at ≥ 16
77 mmol Cu kg⁻¹). Song et al. (2017) assessed the effects of adding different concentrations of Cu (33.3- 1000 mg
78 kg⁻¹) and Zn (165-5000 mg kg⁻¹) on the tolerance of soil bacterial communities to tetracycline, finding that the
79 co-tolerance to the antibiotic significantly increased in soils where Cu concentration were ≥ 365 mg kg⁻¹, and
80 Zn concentrations were ≥ 264 mg kg⁻¹. Zhong et al. (2021) studied the tolerance to tetracycline and vancomycin
81 in 10 mine soil samples with the presence of Cu (361-4399 mg k⁻¹), Zn (33-3811 mg kg⁻¹) and Pb (195-20,239
82 mg k⁻¹). These authors found that, in general, there were increases in tolerance of soil bacterial communities to
83 antibiotics, although they did not detect a systematic pattern in the co-tolerance to tetracycline associated to
84 heavy metals concentrations, whereas the induced levels of heavy metal tolerance coincided with elevated
85 levels of tolerance to vancomycin. However, until now, there were not studies evaluating the effect of a wide
86 range of different heavy metals present in livestock manure on the tolerance shown by soil bacterial
87 communities against each of the three tetracycline antibiotics most used in animal husbandry. Therefore, the
88 objective of this study is to determine the eventual development of tolerance in soil bacterial communities to
89 heavy metals in agricultural soils contaminated individually with 1000 mg kg⁻¹ of As, Cd, Zn, Cu, Ni, Cr and
90 Pb, and also the eventual generation of co-tolerance to the antibiotics tetracycline, oxytetracycline and
91 chlortetracycline, using the leucine incorporation technique as the endpoint. Previous works using Cu and/or
92 Zn showed that the heavy metal concentrations needed to induce bacterial community tolerance to antibiotics
93 in soils is high, generally ≥ 1000 mg kg⁻¹ (Fernández-Calviño and Bååth, 2013; Song et al., 2017). Although

94 these values are nor very common, they may be reached after accumulation with time, since heavy metals are
95 not degraded in soils. In this regard, Mirlean et al. (2007) found Cu concentrations higher than 1000 mg kg⁻¹
96 in Brazilian vineyard soils. On the other hand, soils contaminated with Zn can reach values much higher than
97 1000 mg kg⁻¹, as is the case of soils dedicated to orchards in England, with concentrations of 1800 mg kg⁻¹
98 (Brümmer and Herms, 1983), while amended agricultural soils may reach scores around 7000 mg kg⁻¹ (Kabata-
99 Pendias, 2000). Soils dedicated to gardens and orchards located in England also reached As concentrations of
100 around 900 mg kg⁻¹ (Xu and Thornton, 1985). The same occurs with soils contaminated with Pb, achieving up
101 to 12000 mg kg⁻¹ (Godzik et al., 1995) in soils of orchards and gardens in Poland. Therefore, the results of the
102 current research could be relevant in order to define appropriate management practices for wastes and fertilizers
103 in agricultural soils.

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107 **2. Material and methods**

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109 **2.1. Chemicals**

110 Tetracycline hydrochloride (CAS 64-75-5; ≥95% in purity), oxytetracycline hydrochloride (CAS
111 2058-46-0; ≥95% in purity) and chlortetracycline hydrochloride (CAS 64-72-2; ≥97% in purity), all three
112 supplied by Sigma–Aldrich (Steinheim, Germany), were used for soil spiking.

113 The heavy metals added to the soil samples for determining bacterial community tolerance were
114 arsenic (as Na₂HAsO₄ 7H₂O, CAS; 10048-95-0), cadmium (as Cd(NO₃)₂ 4H₂O, CAS 10022-68-1), zinc (as
115 Zn(NO₃)₂ 6H₂O, CAS; 10196-18-6), copper (as Cu(NO₃)₂ 2.5H₂O, CAS 19004-19-4), nickel (as Ni(NO₃)₂
116 6H₂O, CAS 13478-00-7), chromium (as K₂Cr₂O₇, CAS 7778-60-9), and lead (as Pb(NO₃)₂, CAS 10099-74-8),
117 all of them supplied by Panreac (Barcelona, Spain).

118

119 **2.2. Soil samples**

120 An agricultural soil from Sarria (NW of Spain) was selected form a set of soils previously analyzed
121 by Conde-Cid et al. (2018). Total concentrations of Na, K, Ca, Mg, Al, Fe, Mn, as well as of As, Cd, Cr, Cu,
122 Ni, Pb, and Zn, were determined using ICP-mass spectrometry (820-NS, Varian, Palo Alto, CA, USA), after
123 nitric acid (65%) microwave assisted digestion. As, Cd, Cr, Cu, Ni, Pb, and Zn bioavailability at 0 and 42 days

124 of incubation was assessed performing extractions with 0.01 M CaCl₂ (Novozamsky et al., 1993), and with
125 EDTA (ethylenediamine tetra-acetic acid) (Lakanen and Erviö, 1971). The electrical conductivity (EC) was
126 analyzed using distilled water as extraction solution ratio 1:5 (soil:water ratio 1:5) and measuring it in a
127 conductivity-meter. Moreover, this soil was previously used for measurement of the bacterial growth after being
128 polluted with antibiotics (tetracycline, oxytetracycline and chlortetracycline), with results shown in Santás-
129 Miguel et al. (2020b) and Santás-Miguel et al. (2020c).

130 The main characteristic of the soil studied are shown in Table S1 (Supplementary Material). Briefly, its texture
131 was silt loam, with pH in water of 6.0 and pH in KCl (0.1 M) of 5.2. The organic carbon (C) and total nitrogen
132 (N) contents were 1.8% and 0.2%, respectively and the effective cation exchange capacity (eCEC) was 13.16
133 cmol_c kg⁻¹. The electrical conductivity measured was 298 μS cm⁻¹. The total contents (in mg kg⁻¹) for the
134 measured heavy metals were the following: Cr_T 43.4, Ni_T 25.4, Cu_T 43.4, As_T 27.5, Cd_T <detection limit, Pb_T
135 17.1, and Zn_T 135.5. These values were similar to those found in non-polluted soils in the study area (Macías
136 and Calvo, 2008).

137

138 **2.3. Experimental design**

139 The agricultural soil was polluted, separately, with 1000 mg kg⁻¹ of As, Cd, Zn, Cu, Ni, Cr, and Pb.
140 The procedure was: an amount of 90 g of the air-dried soil was placed in a polypropylene tube (500 mL) and
141 then moistened up to 60-80% of the water holding capacity and incubated for 1 week at 22 °C in the dark, this
142 time being adequate for the recovery and stabilization of the growth of the bacterial communities after moisture
143 adjustment (Meisner et al., 2013). After this time, the soil sample was distributed in 8 polypropylene tubes
144 (100 mL) (putting 12 g of soil, dry weight, in each), with 7 of them being individually polluted with 1000 mg
145 kg⁻¹ of one of the 7 heavy metals (As, Cd, Zn, Cu, Ni, Cr, or Pb), and the 8th tube acting as control (containing
146 soil without metal). Then, the 8 microcosms were distributed in 24 polypropylene tubes (50 mL) (8 microcosms
147 x 3 replicates), placing 3.75 g (dry weight) in each tube, and they were incubated for 42 days in the dark. After
148 1 and 42 days of incubation, 1 g of each tube was used to measure the growth of bacterial communities, and
149 after 42 days, 1.75 g samples were used to estimate the bacterial community tolerance to heavy metals. A
150 schematic description of the experimental design is shown in Figure S1 (Supplementary Material).

151 The concentration of bioavailable heavy metals at days 0 and 42 is shown in Table S2. The
152 concentrations of bioavailable heavy metals at day 0, determined in polluted soil samples extracted with CaCl₂,
153 were 617.4, 333.4, 834.8, 71.1, 429.0, 220.3, and 302.2 mg kg⁻¹ (for As, Cd, Cr, Cu, Ni, Pb, and Zn

154 respectively), whereas, for those extracted with EDTA, they were 917.4, 1016.9, 904.4, 982.1, 802.6, 1080.7,
155 and 953.2 mg kg⁻¹ for As, Cd, Cr, Cu, Ni, Pb, and Zn respectively (Table S2, Supplementary Material). The
156 bioavailable concentrations determined in polluted soil samples at day 42 of incubation were 95.7, 140.9, 191.3,
157 17.4, 252.2, 3.5 and 146.1 mg kg⁻¹, when extracted with CaCl₂, while they were 426.1, 852.2, 252.2, 687.0,
158 800.0, 765.2 and 713.0 mg kg⁻¹ when extracted with EDTA, corresponding to As, Cd, Cr, Cu, Ni, Pb, and Zn,
159 respectively, in both extractants (Table S2, Supplementary Material).

160 The toxicity exerted individually by the heavy metals on the bacterial communities was measured by
161 estimating the bacterial growth on days 1 and 42 of incubation, following the protocol established by Bååth
162 (1994) and Bååth et al. (2001). The bacterial community tolerance was determined according to Bååth (1992),
163 and to Díaz-Raviña et al. (1994), with certain modifications indicated below, and using the leucine
164 incorporation method (Bååth, 1994; Bååth et al., 2001). Briefly, soil samples obtained from each microcosm
165 were mixed with distilled water (rate 1:20 (w/v)), using a multi-vortex shaker for 3 min at maximum intensity.
166 Then, the soil/water mixture was centrifuged at 1000 x g for 10 min to obtain the soil bacterial suspension
167 (supernatant). Aliquots of 1.50 mL of the supernatant were transferred to 2 mL centrifugation tubes, in which
168 150 µL of pollutant solutions (containing heavy metals or antibiotics) were subsequently added. As a result, a
169 total of 7 heavy metals (As, Cd, Zn, Cu, Ni, Cr and Pb) and 3 antibiotics (TC, OTC and CTC) were added to
170 the bacterial suspensions (thus, a total of 10 pollutants were studied), and these plus a control (only water
171 added), were used for each sample. The final concentrations of heavy metals added ranged between 10⁻² and
172 10⁻⁶ mol L⁻¹, while the range for antibiotics went from 400 to 6x10⁻³ mg L⁻¹. The bacterial community growth
173 was estimated after a pre-incubation step of 24 h, where bacterial suspensions were added with the different
174 concentrations of antibiotics before the leucine incorporation assay (Berg et al., 2010; Fernández-Calviño and
175 Bååth, 2013). The [³H] leucine incorporation was then measured on each micro-centrifugation tube, as follows:
176 a volume of 0.2 µL [³H] Leu (3.7 MBq mL⁻¹ and 0.574 TBq mmol⁻¹; Amersham) was added with non-labeled
177 Leu to each tube, resulting in 275 nM Leu in the bacterial suspensions. After 3 h of incubation, the bacterial
178 growth was terminated by adding 75 mL of 100% trichloroacetic acid. Washing was performed as described
179 by Bååth et al. (2001), and a subsequent measurement of radioactivity was carried out using a liquid
180 scintillation counter (Tri-Carb 2810 TR, PerkinElmer, USA).

181

182 **2.4. Data analyses**

183 The tolerance of the bacterial community to the seven heavy metals (As, Cd, Zn, Cu, Ni, Cr and Pb)
184 and the three tetracycline antibiotics (TC, OTC and CTC) was estimated as Log IC₅₀, the logarithm of the
185 concentration that resulted in 50% inhibition of bacterial community growth. Log IC₅₀ was calculated using a
186 logistic model, as follows:

$$187 \quad Y = c/[1 + e^{b(X-a)}]$$

188 where Y is the measured level of Leu incorporation, X is the logarithm of the contaminant substance (heavy
189 metals (As, Cd, Zn, Cu, Ni, Cr, Pb) or antibiotics (TC, OTC or CTC)) added to the bacterial suspension, a is
190 the log IC₅₀, c the bacterial growth rate in absence of the toxic substance, and b is a slope parameter indicating
191 the inhibition rate. A higher value of log IC₅₀ indicates a higher community tolerance, while a lower value
192 indicates that the pollutants (heavy metals or antibiotics) are more toxic to the bacterial community.

193

194 3. Results and Discussion

195 3.1. Toxicity of heavy metals on soil bacteria

196 The toxicity of the heavy metals on soil bacterial communities was tested in the agricultural soil in the
197 short and long terms (1 and 42 days of incubation). The bacterial growth results obtained for the soil samples
198 polluted with 1000 mg kg⁻¹ of As, Cd, Zn, Cu, Ni, Cr, or Pb (separately), and for the unpolluted soil (0 mg kg⁻¹),
199 allowed to establish a toxicity index for these heavy metals (Table 1). On day 1, the toxicity sequence was
200 the following: Cr>Cu>Ni>Cd>Zn>As>Pb. After 42 days of incubation the sequence was similar, with changes
201 mainly affecting to Ni and Cd: Cr>Cu>Cd>Ni>Zn>As>Pb.

202 The data obtained in this study differ from those reported by other authors, which was expected, since
203 the toxicity of heavy metals on soil microorganisms depends on the physicochemical characteristics of the soils
204 (such as pH and carbon content), the concentrations of heavy metals added, and the incubation period (Hattori,
205 1992; Diaz-Raviña and Bååth, 1996; Nannipieri et al., 1997; Ahmad et al., 2005). Hattori (1992) evaluated the
206 influence of Cd, Cr, Cu, Ni, Pb, and Zn on the microorganisms present in two soils after four weeks, and
207 obtained the toxicity sequence Cd> Cu> Ni> Cr> Zn for one of them (a Gleysol, with pH = 5.8 and C = 0.5%),
208 and the sequence Cu> Cd> Ni> Zn> Cr for the other soil (an Andosol, with pH = 6.4 and C = 3.2%). Ahmad
209 et al. (2005) measured the response of the microbial populations of one soil (pH = 7.6 and C = 0.39%) to Cu,
210 Cd, Cr, Hg, Mn, Ni, Pb and Zn at five incubation times, finding that the sensitivity of the bacterial populations
211 depended on the functional group to which they belonged, being the aerobic-heterotrophic bacteria more
212 sensitive to Ni, Pb and Cu, while nitrogen-fixing bacteria were more sensitive to Cd and Pb. Bååth (1989)
213 observed that the relative toxicity of metals decreased in the order Cd> Cu> Zn> Pb, although some differences
214 were found among different investigations. Analyzing data obtained in different studies, it can be observed
215 that, in general, the heavy metals that exert a higher effect on soil microorganisms are Cd and Cu. These results
216 agree with those obtained using as endpoints other organisms, such as *Daphnia magna*, *Vibrio fischeri*, *Lemma*
217 *minor* and *Eisenia fetida*, which show that there is great variability in the toxicities caused by heavy metals,
218 but also that Cd and Cu are the most toxic (Neuhauser et al., 1985; Arambašić et al., 1995; Huynh and Bulich,
219 1995; Dirilgen, 2001; Hsieh et al., 2004; Teodorovic et al., 2009).

220

221 3.2. Soil bacterial communities' tolerance to heavy metals in the long-term

222 Figure 1 shows the dose-response curves obtained for each heavy metal after 42 days of incubation,
223 including data for each microcosm. These dose-response curves are sigmoidal, such as those reported by other

224 authors (Díaz-Raviña et al., 1994; Cruz-Paredes et al., 2017; Song et al., 2017; Santás-Miguel et al., 2020a).
225 The inhibition curves show the absence of bacterial growth inhibition at low heavy metals' concentrations, but
226 inhibition takes place at high doses, and it increased with dose. As general trend, it is observed that the
227 inhibition curves obtained for all heavy metals tested shift to the right with respect to the control (unpolluted
228 soil). This shift to the right in the dose-response curves suggests the existence of soil bacterial community
229 tolerance to the high heavy metals' concentrations added on the soil used in this research (1000 mg kg⁻¹).

230 In this study, the dose-response curves fitted well to the logistic model, with R² ranging between 0.966
231 and 0.991 (mean R²=0.983) for the control sample (0 mg kg⁻¹), and R² between 0.861 and 0.987 (mean R²=
232 0.950) for samples polluted with heavy metals (1000 mg kg⁻¹ of each one, individually).

233 Table 2 shows the Log IC₅₀ values obtained from each dose-response curve. The unpolluted soil
234 showed the lowest Log IC₅₀ value for Cu (-4.17±0.05), and the highest value for Ni (2.56±0.08), i.e, the bacteria
235 found in the bacterial suspension of unpolluted soils show high toxicity effects due to Cu, and lower toxicity
236 due to Ni. As regards the Log IC₅₀ values obtained for the whole set of heavy metals tested, the following
237 toxicity sequence was observed for the control (unpolluted soil): Cu>As>Cr≥Pb≥Cd>Zn>Ni. This toxicity
238 sequence is similar to that reported by Díaz-Raviña et al. (1994), who studied the bacterial community tolerance
239 to Cu, Cd, Zn, Ni and Pb using the thymidine incorporation technique, and found that Cu was the most toxic,
240 while Ni showed the lowest toxicity on bacterial communities. However, the order of the rest of heavy metals
241 differed from that observed in the current study, with the sequence being: Cu> Cd> Zn> Pb> Ni.

242 Regarding the samples polluted with 1000 mg kg⁻¹ of each of the heavy metals separately, the lowest
243 Log IC₅₀ score corresponded to Cu (-2.61±0.12), and the highest to Zn (-1.63±0.09) (Table 2), i.e., the bacterial
244 communities present in these microcosms are more sensitive to Cu than to Zn. The toxicity sequence, in
245 function of the Log IC₅₀ values, was: Cu>Pb≥As≥Cd≥Cr≥Ni≥Zn.

246 The LogIC₅₀ values obtained for each soil sample polluted with the highest heavy metal concentrations
247 increased with respect to the control (unpolluted soil), indicating that the bacteria exposed to high levels of
248 heavy metals developed tolerance to these pollutants. The highest increases in Log IC₅₀ (ΔLog IC₅₀, Table 2)
249 corresponded to Cr and Cu (1.6 units), followed by As, Zn, Cd and Pb (1.4, 1.3, 1.2 and 1.1 units, respectively),
250 whereas the increase was of 0.7 units for Ni. To note that the presence of high amounts of heavy metals in the
251 soil causes that bacterial communities increase their tolerance to these pollutants (Díaz-Raviña et al. 1994;
252 Díaz-Raviña and Bååth, 1996; Díaz-Raviña and Bååth, 2001; Fernández-Calviño and Bååth, 2013; Santás-

253 Miguel et al., 2020a; Zhong et al., 2021), although the magnitude of these increases seems to depend on the
254 toxicity or initial pressure exerted by the metal on the soil bacterial communities.

255 Figure S2 (Supplementary Material) shows the relation between the increase in tolerance to heavy
256 metals and the toxicity exerted by each metal on the bacterial growth in the short term (1 day) and in the long
257 term (42 days). It can be observed that the higher is the bacterial growth inhibition on days 1 and 42, the greater
258 is the increase in tolerance generated by bacterial communities to heavy metals. This behavior was observed
259 for all the heavy metals studied except Ni. In fact, eliminating Ni the relation becomes significant at day 1
260 ($r=0.789$, $P < 0.05$) and at day 42 ($r=0.730$; $P < 0.05$). The toxicity of heavy metals on soil bacterial communities
261 is related to the bioavailability of these heavy metals in the soil. The time-course evolution of heavy metals'
262 bioavailability is shown in Table S2 (Supplementary Material), indicating that, in general, they have a high
263 bioavailability in the short term (Ni being the least bioavailable), while, in the long term (concentrations in
264 EDTA), the bioavailability of heavy metals decreases in a percentage of between 20 and 70%, being less in the
265 case of Ni, which remains constant over time.

266 The results suggest that the increase in soil bacterial community tolerance after the addition of heavy
267 metals can be attributed to the immediate effect caused by the decay of sensitive species, and to a posterior
268 effect caused by different competitive capacities and ulterior adaptation in the bacteria remaining. This effect
269 of heavy metals in soil has been observed previously by Diaz-Raviña and Bååth (1996) for Zn, Cd, Ni, Cu and
270 Pb, and by Fernandez-Calviño and Bååth (2016) for Cu.

271

272 **3.3. Soil bacterial communities' tolerance to tetracycline antibiotics in the long-term**

273 Figures 2, 3, and 4 show dose-response curves obtained at 42 days of incubation for each heavy metal
274 added to the soil at 1000 mg kg^{-1} , after the exposure of bacterial suspension from each microcosm to
275 concentrations of tetracyclines antibiotics (TC, OTC and CTC) ranging from 400 to $6 \times 10^{-3} \text{ mg L}^{-1}$. The
276 inhibition curves obtained for these tetracycline antibiotics are sigmoidal. As general trend, the dose-response
277 curves are shifted to the right respect to the control, and in some cases the tetracycline antibiotics did not show
278 an inhibitory effect on bacterial communities.

279 These dose-response curves fitted well to the logistic model, with R^2 ranging between 0.925 and 0.982
280 (mean $R^2=0.956$) for TC samples, between 0.771 and 0.997 (mean $R^2= 0.952$) for OTC samples, and from
281 0.959 to 0.992 (mean $R^2=0.976$) for CTC samples.

282 Table 3 shows the Log IC₅₀ values obtained from each dose-response curve, being 1.83±0.06 for the
283 control sample after exposure of the bacterial suspension to different concentrations of TC. In general, the Log
284 IC₅₀ values obtained for each microcosm polluted with 1000 mg kg⁻¹ of each heavy metal, and subsequently
285 exposed to TC, increased with respect to the control. The magnitude of these increases varies in function of the
286 heavy metal, being around 0.4 units for Pb, Zn and As, and 0.5 units to Ni. However, the bacterial communities
287 of soils contaminated with Cd, Cu and Cr do not show inhibition in their growth when exposed to TC, and,
288 therefore, their LogIC₅₀ cannot be determined due to the high community tolerance to TC. Consequently, heavy
289 metals that do not show inhibition have been assigned a Log IC₅₀ value of >2.6, since it is the maximum
290 concentration tested in the current study.

291 The Log IC₅₀ value obtained for the control soil in presence of OTC was 1.11±0.08. Regarding the
292 Log IC₅₀ values obtained for the samples polluted with heavy metals, it can be observed that, after the exposure
293 of the bacterial suspension to OTC, there was an increase in these values with respect to the control, going
294 from 0.2 to 0.6 for Pb, Cd, Zn, As and Ni, and being 1.6 for Cu. Regarding Cr, the exposure of the soil bacterial
295 suspension to OTC did not cause inhibition of its growth, and, therefore, as in the case of Cd, Cu and Cu for
296 TC, the value of >2.6 (maximum concentration tested) was assigned.

297 Finally, the Log IC₅₀ value obtained for the control soil added with CTC was 1.52±0.10. After
298 exposure of the bacterial suspensions to CTC, the Log IC₅₀ values obtained for the samples polluted with heavy
299 metals showed an increase with respect to the control, going from 1.7 to 2.0 for all the heavy metals except for
300 Cr, which did not show inhibition of bacterial growth when exposed to different concentrations of CTC.
301 Therefore, it can be indicated that bacterial communities exposed to high concentrations of heavy metals show
302 great co-tolerance to CTC. In general, the increase in tolerance obtained for TC, OTC and CTC cannot be
303 related to the toxicity of heavy metals in the soil, at 1 and 42 days of incubation (data not shown).

304 Co-tolerance to TC generated by soil bacterial communities exposed to heavy metals has been
305 previously demonstrated (Berg et al., 2010; Fernández-Calviño and Bååth, 2013; Song et al. 2017). Song et al.
306 (2017) observed the existence of co-tolerance to TC in a microcosm exposed to concentrations of Cu ≥333 mg
307 kg⁻¹ and Zn ≥500 mg kg⁻¹. Fernández-Calviño and Bååth (2013) found co-tolerance to TC of soil bacterial
308 communities in contaminated soils with concentrations of Cu ≥500 mg kg⁻¹. The results shown in these studies
309 agree with what was observed in the present work, indicating that soil bacterial communities exposed to high
310 concentrations of heavy metals (1000 mg kg⁻¹) generate co-tolerance to tetracycline. However, Zhong et al.

311 (2021) did not observe co-tolerance to TC in soils contaminated with Cu, Zn and Pb, while they found co-
312 tolerance of soil bacterial communities to vancomycin.

313 This co-tolerance of soil bacterial communities to TC at high concentrations of heavy metals is also
314 observed for OTC and CTC. The results obtained in this study agree with the data reported in studies previously
315 carried out by Santás-Miguel et al. (2020a), where co-tolerance of soil bacterial communities to tetracycline
316 antibiotics (TC, OTC and CTC) was observed in soils with concentrations of Cu ≥ 1000 mg kg⁻¹ after 42 days
317 of incubation.

318 It is relevant that co-tolerance studies, in general, focus on a reduced number of heavy metals
319 (generally Cu) and tetracycline antibiotics, while the present work shows for the first time the effect of 7
320 different heavy metals as regards the co-tolerance to the 3 most used tetracycline antibiotics (TC, OTC and
321 CTC).

322 It is also remarkable that the role of heavy metals as a co-selection factor to generate antibiotic
323 resistance genes has been studied for a wide variety of heavy metals and antibiotics (Berg et al., 2010;
324 Dickinson et al., 2019; Cao et al., 2020; Mazhar et al., 2021), and, although in the present study antibiotic
325 resistance genes (ARGs) have not been studied directly, the resistance of soil bacterial communities to heavy
326 metals and tetracycline antibiotics (TC, OTC, CTC) could be detected using the Leucine incorporation method.
327 The heavy metal levels of the tested microcosms (1000 mg kg⁻¹) are really very high, but they should not be
328 considered unrealistic, since heavy metals are not easily biodegradable, and therefore their concentrations can
329 increase over time (Li et al., 2015), reaching very high levels in the soil (Smolders et al., 2004; Spiteri et al.,
330 2005; Zhuang et al., 2009; Komárek et al., 2010). Therefore, soils contaminated with high concentrations of
331 heavy metals could become hotspots of ARGs, which would pose a potential danger to public health, since
332 they could be transmitted to humans (Forsberg et al., 2012).

333

334 **4. Conclusions**

335 In the current study it was evidenced that heavy metals present in soils show wide variability as regards
336 the toxicity they exert on bacterial communities, with Cd and Cu being the most toxic. Specifically, soil samples
337 contaminated with a high concentration (1000 mg kg⁻¹) of each of 7 different heavy metals (As, Cd, Zn, Cu,
338 Ni, Cr and Pb), added separately, showed tolerance of the soil bacterial communities to the heavy metals
339 themselves, compared to the control soil. Besides, it was shown that the higher the effect of heavy metals in
340 the short and long terms, the greater the increase in tolerance to metals generated in the soil bacterial

341 communities (except for Ni). In addition, soil samples contaminated with high concentrations of heavy metals
342 also showed co-tolerance of soil bacterial communities to tetracycline, oxytetracycline and chlortetracycline
343 for all the metals tested, compared to the control soil. However, in general, there was no correlation between
344 the initial damage exerted by heavy metals in the short and long terms and the co-tolerance shown by soil
345 bacterial communities to tetracycline antibiotics. Future studies could give further details about the potential
346 interactions of different pollutants present in agricultural soils, which can be considered relevant as regards
347 environmental preservation and public health.

348

349 **Acknowledgment**

350 This study has been funded by the Spanish Ministry of Economy and Competitiveness through the projects
351 CGL2015-67333-C2-1-R and -2-R (FEDER Funds). David Fernández Calviño holds a Ramón y Cajal contract
352 (RYC-2016-20411), financed by the Spanish Ministry of Economy, Industry and Competitiveness. Vanesa Santás
353 Miguel holds a pre-doctoral fellowship (ED481A-2020/089) financed by Xunta de Galicia.

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552 **Tables and figures**

553

554 **Table 1.** Toxicity index values \pm error of soil samples containing 0 mg kg⁻¹ (control soil) and 1000 mg kg⁻¹ of

555

As, Cd, Zn, Cu, Ni, Cr or Pb (individually), after 1 and 42 days of incubation.

556

Heavy metal	Day 1	Day 42
	Toxicity Index \pm error	Toxicity Index \pm error
As	0.69 \pm 0.04	2.39 \pm 0.18
Cd	0.12 \pm 0.01	0.15 \pm 0.00
Zn	0.15 \pm 0.04	2.36 \pm 0.04
Cu	0.03 \pm 0.01	0.06 \pm 0.01
Ni	0.05 \pm 0.02	2.18 \pm 0.39
Cr	0.01 \pm 0.01	0.01 \pm 0.00
Pb	0.72 \pm 0.04	2.76 \pm 0.06

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558

559 **Table 2.** Estimated values of $\text{LogIC}_{50 \pm \text{error}}$ and R^2 from fits to a logistic model of soil samples containing 0 mg
 560 kg^{-1} (control soil) and 1000 mg kg^{-1} of As, Cd, Zn, Cu, Ni, Cr or Pb (individually). The increases in tolerance (Δlog
 561 $\text{IC}_{50} = \text{LogIC}_{50 \text{ polluted}} - \text{LogIC}_{50 \text{ control}}$) were calculated for each heavy metal

Heavy metal	Control soil		Heavy metal-polluted soil (1000 mg kg^{-1})		$\Delta \text{log IC}_{50}$
	$\text{Log IC}_{50 \pm \text{error}}$	R^2	$\text{Log IC}_{50 \pm \text{error}}$	R^2	
As	-3.63±0.11	0.970	-2.28±0.04	0.987	1.4
Cd	-3.37±0.07	0.988	-2.21±0.06	0.966	1.2
Zn	-2.89±0.06	0.988	-1.63±0.09	0.966	1.3
Cu	-4.17±0.05	0.991	-2.61±0.12	0.973	1.6
Ni	-2.56±0.08	0.966	-1.82±0.11	0.861	0.7
Cr	-3.44±0.07	0.987	-1.86±0.19	0.938	1.6
Pb	-3.42±0.07	0.989	-2.34±0.06	0.977	1.1

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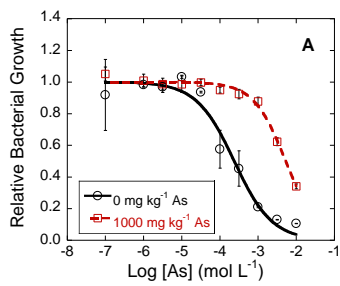
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564 **Table 3.** Estimated values of $\text{LogIC}_{50} \pm \text{error}$ and R^2 from fits to a logistic model of soil samples polluted with 1000
565 mg kg^{-1} of As, Cd, Zn, Cu, Ni, Cr and Pb. The increases in tolerance ($\Delta \log \text{IC}_{50} = \text{LogIC}_{50} \text{ polluted} - \text{LogIC}_{50} \text{ control}$)
566 were calculated for tetracycline (TC), oxytetracycline (OTC), and chlortetracycline (CTC)

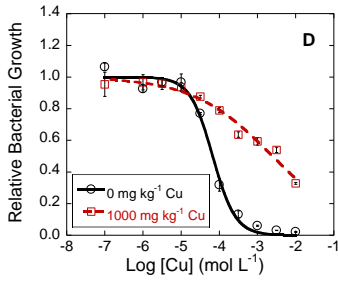
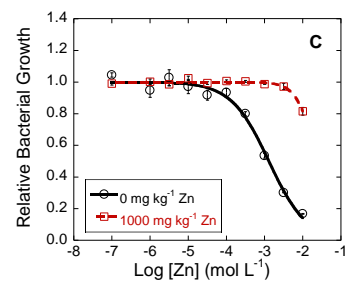
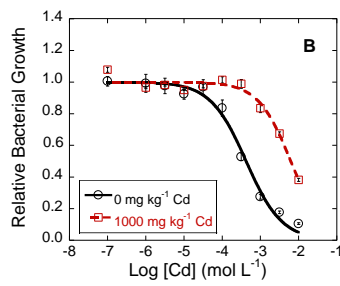
	TC			OTC			CTC		
	$\text{Log IC}_{50} \pm \text{error}$	R^2	$\Delta \log \text{IC}_{50}$	$\text{Log IC}_{50} \pm \text{error}$	R^2	$\Delta \log \text{IC}_{50}$	$\text{Log IC}_{50} \pm \text{error}$	R^2	$\Delta \log \text{IC}_{50}$
Control	1.83±0.06	0.975	--	1.11±0.08	0.982	--	1.52±0.10	0.959	--
As	2.26±0.09	0.925	0.4	1.65±0.02	0.997	0.5	1.91±0.04	0.985	0.4
Cd	≥2.6*	--	0.8	1.47±0.09	0.973	0.4	1.91±0.01	0.982	0.4
Zn	2.28±0.05	0.971	0.4	1.64±0.04	0.991	0.5	2.19±0.11	0.977	0.7
Cu	≥2.6*	--	0.8	2.67±0.37	0.771	1.6	1.82±0.08	0.959	0.3
Ni	2.33±0.04	0.982	0.5	1.68±0.07	0.968	0.6	1.75±0.04	0.992	0.2
Cr	≥2.6*	--	0.8	≥2.6*	--	1.5	≥2.6*	--	1.1
Pb	2.23±0.08	0.926	0.4	1.34±0.08	0.981	0.2	1.99±0.03	0.976	0.5

567 * The value of 2.6 is the maximum value tested

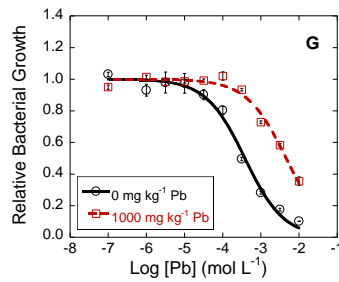
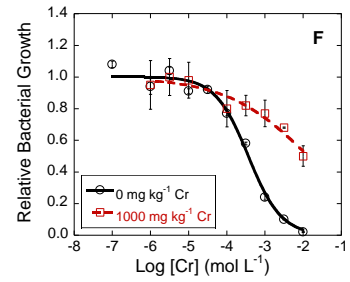
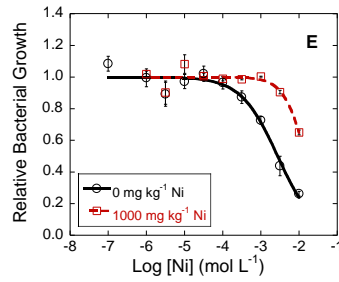
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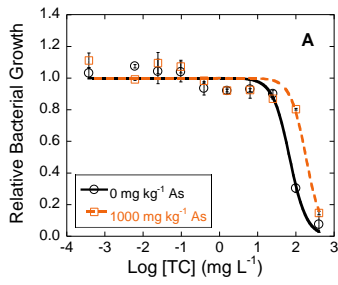


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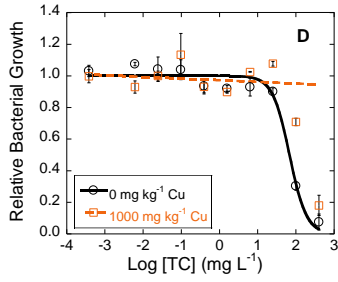
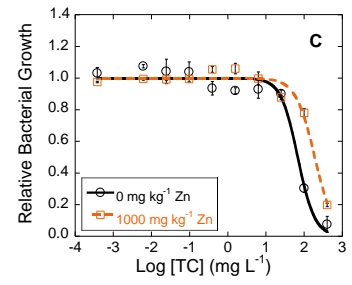
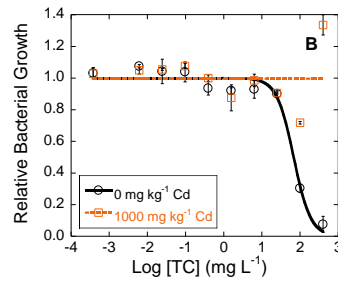
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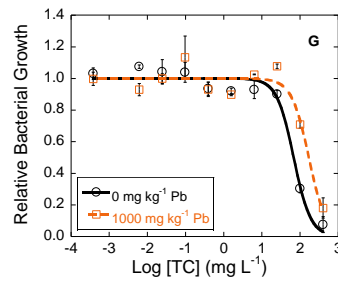
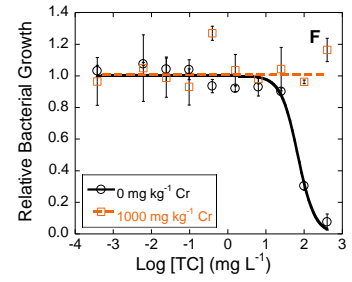
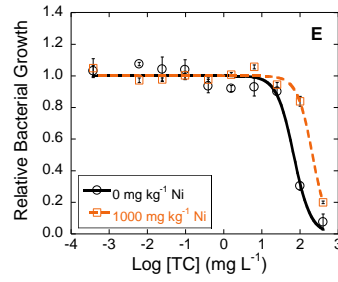
574 **Figure 1.** Dose-response curves obtained after the addition of 10 different concentrations of the heavy metals As (A), Cd
 575 (B), Zn (C), Cu (D), Ni (E), Cr (F) and Pb (G) to bacterial suspensions extracted from soils polluted with 1000 mg kg⁻¹
 576 of each of the same heavy metals, separately, after 42 days of incubation. Black lines represent the dose-response curves
 577 corresponding to the unpolluted soil (control) and discontinued red lines are dose-response curves corresponding to the
 578 soil polluted with 1000 mg kg⁻¹ of heavy metal. Average values (n=3) with coefficients of variation always <5%



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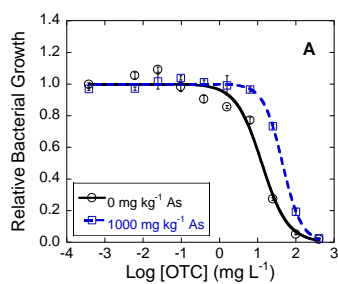
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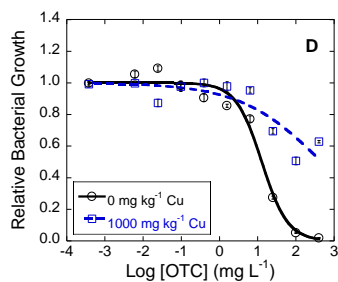
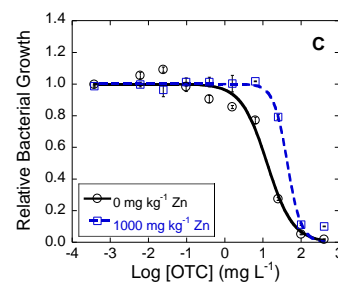
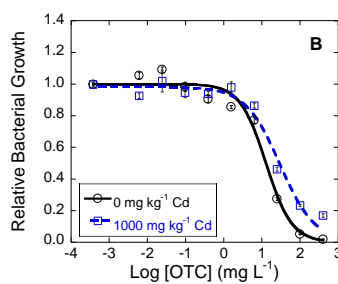
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584 **Figure 2.** Dose-response curves obtained after the addition of 10 different concentrations of tetracycline (TC) to
585 bacterial suspension extracted from soil samples polluted with 1000 mg kg⁻¹ of As (A), Cd (B), Zn (C), Cu (D), Ni
586 (E), Cr (F) or Pb (G), each of them added separately, after 42 days of incubation. Black lines represent the dose-
587 response curves obtained for the unpolluted soil (control) and discontinued orange lines are dose-response curves
588 for soils polluted with 1000 mg kg⁻¹ of one of the heavy metals. Average values (n=3) with coefficients of variation
589 always <5%.

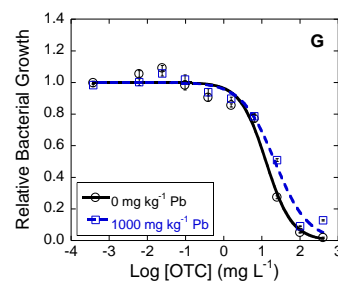
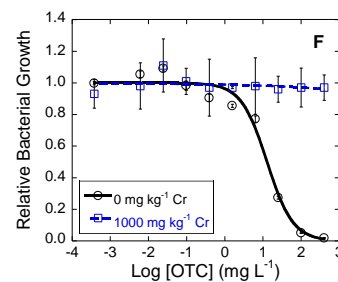
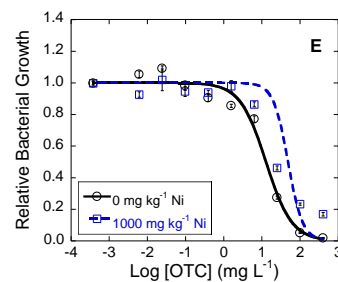
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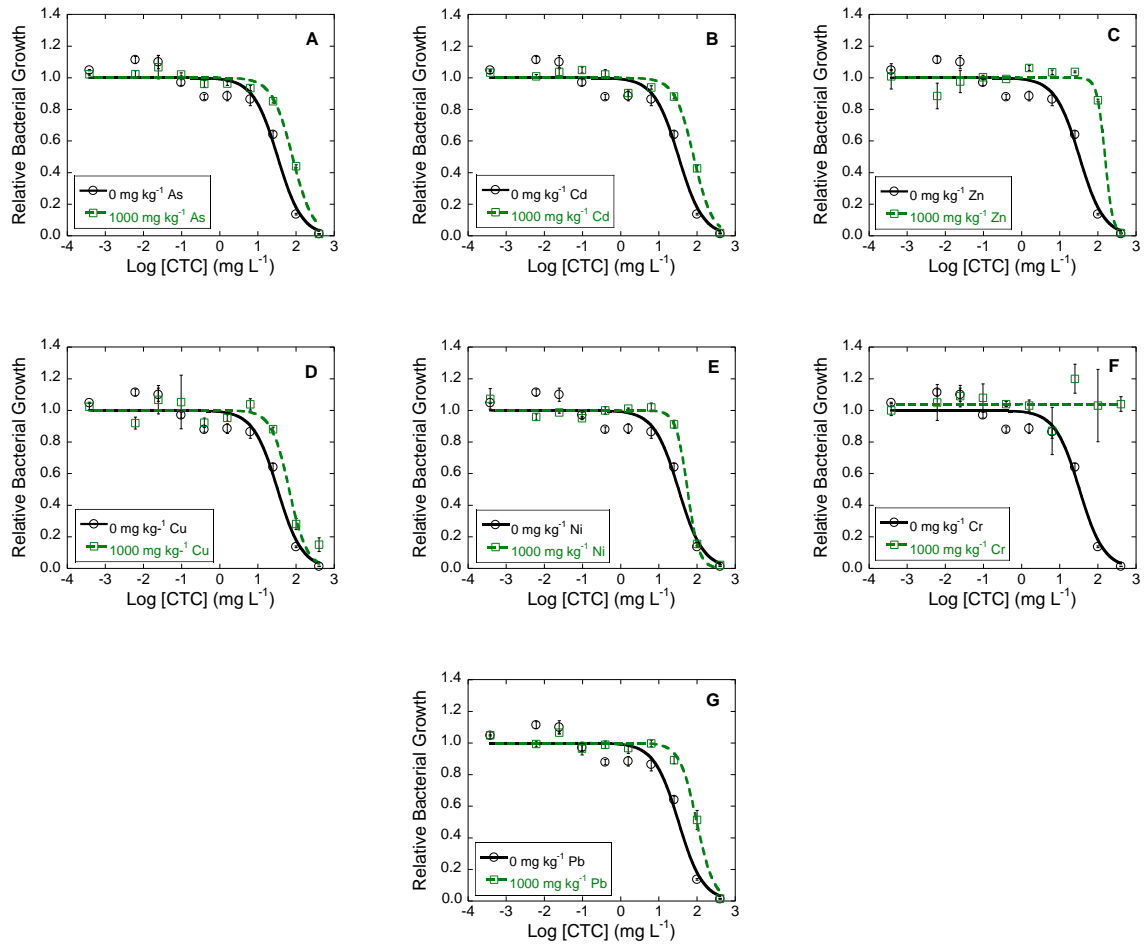
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596 **Figure 3.** Dose-response curves obtained after the addition of 10 different concentration of oxytetracycline (OTC)
 597 to bacterial suspension extracted from soil samples polluted with 1000 mg kg⁻¹ of As (A), Cd (B), Zn (C), Cu (D),
 598 Ni (E), Cr (F) or Pb (G), each of them added separately, after 42 days of incubation. Black lines represent the dose-
 599 response curves obtained for the unpolluted soil (control) and discontinued blue lines are dose-response curves for
 600 soils polluted with 1000 mg kg⁻¹ of one of the heavy metals. Average values (n=3) with coefficients of variation
 601 always <5%.

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611 **Figure 4.** Dose-response curves obtained after the addition of 10 different concentration of chlortetracycline (CTC)
612 to bacterial suspension extracted from soil samples polluted with 1000 mg kg⁻¹ of As (A), Cd (B), Zn (C), Cu (D),
613 Ni (E), Cr (F) or Pb (G), each of them added separately, after 42 days of incubation. Black lines represent the dose-
614 response curves obtained for the unpolluted soil (control) and discontinued green lines are dose-response curves
615 for soils polluted with 1000 mg kg⁻¹ of one of the heavy metals. Average values (n=3) with coefficients of variation
616 always <5%.

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