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# Using a Portuguese regosol to derive soil reference values for arsenic

## Utilização de um regossolo português para derivar valores de referência para arsénio

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### ABSTRACT

Soil reference values (SRV) are preliminary screening benchmark values used to evaluate the potential toxicity of specific contaminants in soils. The Portuguese Environmental Agency has proposed SRV for metals and metalloids based on soil uses but without consideration for the soil type. This project aims to derive SRV for arsenic (As) using a natural Portuguese soil since it is known that soil properties can influence the mobility and bioavailability of contaminants. To achieve this aim, a non-contaminated regosol was collected and spiked with a range of concentrations of As. After, a battery of ecotoxicological tests was performed using the spiked soil, including microbial parameters, reproduction with invertebrates, plant growth assays, and aquatic organisms (to evaluate elutriates from the spiked soil) following standard protocols (ISO and OECD) and aimed at deriving concentrations that cause a x% effect (EC<sub>x</sub>) to the tested organism and/or the test endpoints assessed. The endpoints will be used to estimate a SRV (e.g., PNEC- predicted no effect concentration to the ecosystem or HC<sub>5</sub> – hazard concentration for 5% of the species of the ecosystem) using Species Sensitivity Distribution (SSD) models. The results showed that As was very toxic to all the tested organisms, and based on the data we collected, an HC<sub>5</sub> based on 10% effect concentrations HC<sub>5</sub> (EC<sub>10</sub>) of 2.80 mg kg<sup>-1</sup> of soil<sub>d<sub>w</sub></sub> was proposed as SRV for regosol.

**Keywords:** Soil Reference Values, Environmental Risk Assessment, Ecotoxicological Tests, Species Sensitivity Distribution models.

### RESUMO

Os valores de referência do solo (VRS) são usados para avaliações preliminares da potencial toxicidade de contaminantes específicos presentes nos solos. A Agência Ambiental Portuguesa propôs VRS para metais e metalóides com base no uso potencial do solo, mas sem considerar o tipo de solo. Este trabalho tem como objetivo derivar VRS para o arsénio (As) usando um solo natural português, tendo por base o conhecimento de que as propriedades do solo podem influenciar a mobilidade e a biodisponibilidade de contaminantes. Para atingir este objetivo, um regossolo não contaminado foi recolhido e contaminado com uma gama de concentrações de As. Posteriormente, foi realizada uma bateria de testes ecotoxicológicos usando o solo contaminado e incluindo parâmetros microbianos, reprodução com invertebrados, ensaios crescimento e germinação de plantas, assim como ensaios com organismos aquáticos (para avaliação de elutriados do solo contaminado) seguindo protocolos padronizados (ISO e OCDE), de forma a determinar as concentrações que causam um efeito de X% nos organismos ou parâmetros testados (CE ou do Inglês EC<sub>x</sub>). Estes valores serão usados para estimar um VRS (por exemplo, PNEC - concentração para a qual não se prevê a ocorrência de efeitos no ecossistema ou HC<sub>5</sub> - concentração de perigo (do inglês: Hazard Concentration) para 5% das espécies do ecossistema) usando curvas de distribuição da sensibilidade das espécies (do inglês: SSDs). Os resultados mostram que o As é muito tóxico para todos os organismos/parâmetros testados e, com base nos dados recolhidos, um HC<sub>5</sub> baseado em concentrações de efeito de 10% HC<sub>5</sub> (EC<sub>10</sub>) de 2,80 mg kg<sup>-1</sup> de solo<sub>ps</sub> foi proposto como VRS para regossolos.

**Palavras-chave:** Valores de Referência do Solo (VRS), Avaliação de Risco, Testes Ecotoxicológicos, curvas de distribuição de sensibilidade das espécies

## INTRODUCTION

Soil reference values (SRV) are threshold concentrations of pollutants in soil that, when attained, an effect on the terrestrial ecosystem is expected. These values are generic screening standards used majorly to evaluate the potential toxicity of specific contaminants in the soils, providing valid and specific information about the impacts on the terrestrial biota when exposed (Friday, 1999; Pereira *et al.*, 2018).

Toxicity data for plants, invertebrates, soil microbial activity, and sometimes mammals and birds, or even background concentrations obtained from natural soils, are frequently used to derive SRV for environmental risk assessment (Caetano *et al.*, 2016; Pereira *et al.*, 2018). The effects of the contaminant on the soil biota are tested at different endpoints and organisms. Then probabilistic or deterministic methods are applied for the estimations of the reference values for the pollutant, following different standard guidance documents, such as the European Commission Technical Guidance Document on Risk Assessment (European Commission, 2003), United States Environmental Protection Agency or the United States Environmental Protection Agency (USEPA), and the methods proposed by The Netherlands (Swartjes *et al.*, 2012).

In Portugal, derivation of SRV started with a cambisol, which is the most common soil category in the country's north and center. For this kind of soil, SRV for copper, cadmium, and uranium have already been suggested (Caetano *et al.*, 2016). The major objective of this work is to generate an ecotoxicological dataset using a specific type of Portuguese natural soil (regosol) for the derivation of soil reference values for arsenic (As). The regosol is the dominant type of soil at Estarreja region (center of Portugal), where historical problems of soil contamination were caused by the Estarreja Chemical Complex. Obtaining regional SRVs of the most concerning contaminants in the region, as for example As, to be used in for first-tier soil risk assessment purposes is of utmost importance for a site-specific evaluation.

## MATERIAL AND METHODS

Based on the area's geological maps and historical soil uses, a representative regosol soil from Estarreja was collected from three different sites (40°45'47.5"N 8°35'49.6"W) randomly and mixed thoroughly to obtain a composite sample, as a true representative of a regosol from that area. Soil samples were collected (0-10 cm depth), labelled, transported, processed adequately (2 mm sieve was used for the enzyme and aquatic toxicity assays and 4 mm for the assays with terrestrial invertebrates and plant assays), and stored depending on the parameters to be analysed (e.g., oven/air-dried for physical-chemical analysis).

The soil was spiked with a range of concentrations, defined based on the Effect concentration (EC<sub>x</sub>) sampling design (12 concentrations with minimum of 3 replicates per concentrations) and left for 48 hours. The concentrations tested were 0, 1.86, 3.25, 5.68, 9.95, 17.41, 30.46, 54.31, 93.29, 163.27, 285.71 and 500 mg of As kg<sup>-1</sup> of soil<sub>dw</sub> respectively. After 48 hours, the spiked soil samples were used to perform a battery of terrestrial ecotoxicological tests with *Eisenia fetida* and *Folsomia candida* to assess the effects on the reproduction of terrestrial invertebrates following the OECD 232 and 222 criteria, the seedling emergence and growth of terrestrial plants (at least two monocotyledonous and two dicotyledonous species) following the OECD 208 protocol. In our case, we used *Avena sativa*, *Triticum aestivum*, *Lactuca sativa* and *Solanum lycopersicum* respectively. The contaminated soils were also brought in contact with deionized water containing 0.001 mol/l of CaCl<sub>2</sub> solution (in a liquid to solid ratio of 4:1 l/kg) and agitated for 24 hours until near equilibrium between liquid and solid phase was achieved. After 24 hours, the samples were taken to the centrifuged for 20 min and then the supernatant was collected, sieved with a 0.45 μm syringe filters and then used for the aquatic toxicity assays with *Daphnia magna* (immobilization test following OECD 202), *Raphidocelis subcapitata* and *Lemna minor* (Growth inhibition tests following OECD 201 and 221) and *Allivibrio fischeri* (following the Microtox® model 500 Toxicity Analyzer (Modern Water, New Castle, DE) with the 81.9% basic test protocol.

A sample of fresh soil was also contaminated with the range of concentrations of As as stated previously and incubated for 30 days under a well-controlled photoperiod and temperature conditions, for microbial parameters. After 30 days, the contaminated soil was used to assess the activity of several soil microbial enzymes (dehydrogenases, cellulase (Schinner & von Mersi, 1990; Schinner *et al.*, 1996), arylsulfatase (Tabatabai & Bremner, 1970; Schinner *et al.*, 1996), urease (Schinner *et al.*, 1996; Kandeler & Gerber, 1998), phosphatases, potential nitrification and nitrogen mineralisation (Keeney, 1983; Schinner *et al.*, 1996).

A one-way analysis of variance test was used to test for a significant effect of the pollutant concentrations on each of the tested species/parameters. After all the ANOVA assumptions had been met, Dunnett's post-hoc test was employed to determine which concentrations differed significantly from the control and the LOEC (low observed effect concentration) and NOEC (no observed effect concentration) values obtained. The  $EC_{50}$ ,  $EC_{20}$  and  $EC_{10}$  values were calculated using the DRC package in R and compared with the STATISTICA® 7.0 software (StatSoft, Inc., Tulsa, OK, USA). All the estimations were recorded at a 95% confidence interval and alpha 0.05.

The USEPA species sensitivity distribution software and the ssdtool package in R were used to estimate the percentage hazard concentration ( $HC_p$ ), and the protection level was determined and proposed as the SRV for As in regosol.

## RESULTS AND DISCUSSION

Arsenic was very toxic for all the plant species tested, with total inhibition noted at  $\geq 281 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$ . However, *L. sativa* had a better germination percentage than *H. vulgare*, but the dry mass of *H. vulgare* proved to be a more sensitive endpoint amongst the parameters tested for plants. Figure 1 shows the effect of As on the growth of one of the terrestrial plants (*L. sativa*).

For the invertebrates, As significantly inhibited the reproduction of *E. fetida* ( $p < 0.01$ ) and *F. candida* ( $p < 0.01$ ) as the concentration increases, with a total inhibition noted at  $\geq 93.29 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  as

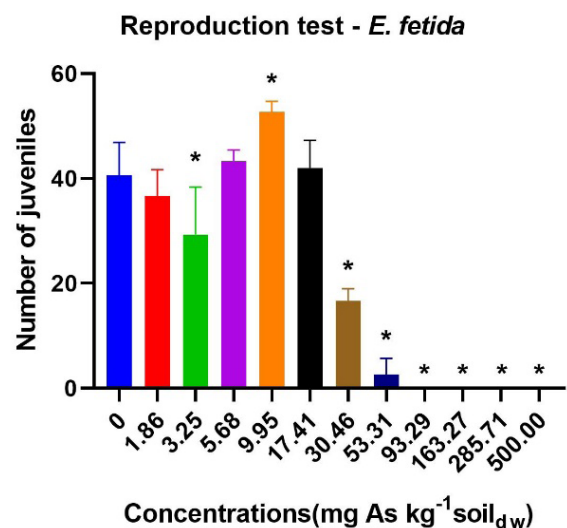


**Figure 1** - Phytotoxicity of As to *L. sativa*.

shown in Figure 2. The NOEC and LOEC values recorded were  $1.86 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  and  $3.35 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  and  $5.68 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  and  $9.95 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  respectively, for these organisms.

For the microbial parameters, As inhibited the activity of acid phosphatase as the concentration increases with concentrations  $\geq 93.95 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  being significantly different from the control. Furthermore, As inhibited dehydrogenase activity with the activity at concentrations  $\geq 30.46 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$  being significantly different from the control.

For the aquatic organisms, As was extremely toxic to *D. magna*, with total immobilization being noted in the elutriate prepared from soils with concentrations above  $17.41 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$ .



**Figure 2** - Average number of juveniles produced by *E. fetida* exposed to several concentrations of As in the regosol. The asterisks sign out for significant differences relatively to the control ( $0.0 \text{ mg As kg}^{-1} \text{ soil}_{\text{dw}}$ ) (Dunnett test,  $p < 0.05$ ).

## CONCLUSIONS

The ecotoxicological data gathered in this study showed that As has an impact on the soil and aquatic community, even at the lowest concentrations tested, when the organisms are exposed through a regosol.

The estimated Effect Concentrations (EC<sub>x</sub>) that were calculated from each of the experiments were used to derive the reference values.

This was achieved by fitting the most sensitive endpoints from each of the assays to a mathematical model using the USEPA SSD software to derive a HC<sub>5</sub> for EC<sub>10</sub>, EC<sub>20</sub> and EC<sub>50</sub>. Based on the data

collected, an SRV of 2.80 mg As kg<sup>-1</sup> soil<sub>dw</sub> with R<sup>2</sup> of 0.96 was proposed.

Other assays are currently running, and the data would be used to generate an SRV that genuinely represents the ecological reality of the study location, which is related to agricultural soils.

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