ATÓLICA

## THE UPTAKE OF ZINC IN PLANT SPECIES **INDIGENOUS TO A PORTUGUESE POLLUTED SITE**

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THE USE OF PIONNERS FOR THE PHYTOREMEDIATION OF TOXIC METAL CONTAMINATED SOILS

Pollution of the environment and the non controlled migration of metal contaminants have increased dramatically since the birth of the industrial revolution, causing serious damage on ecosystems and affecting public health. These and other reasons bring up the need for new solutions of remediation. Phytoremediation is an emergent technology that uses plants to remove, degrade or immobilize the contaminants, offering a low cost method for soil remediation.

The use of fast growing pioneer species capable of colonizing poor, contaminated soils is potentially very useful for phytoremediation strategies. Field studies are necessary to obtain information on how these indigenous plants behave under the conditions installed. The region of Estarreja appears as a strong candidate for this kind of research as for many years, several chemical facilities of the region have discharged its solid residues in an improvised park in the surrounding area, and conducted its wastewaters into a stream nearby. In spite of the levels of zinc (amongst other metals as Pb, Hg and As) remaining above the limits established by the european legislation in the sediments of this stream and of the high permeability of the soils in the area, the vegetation remains proliferous on the banks of the stream. Through an observation of the indigenous flora, two high biomass plants appear as the main colonizers: Phragmites australis (common reed) and Rubus ulmifolius (dewberries).

The aim of this study was to determine the distribuition of zinc in different sections - roots, shoots and leaves - of the selected plants and relate that to the zinc content of the soil. The mycorrhizal status of the sampled plants has also been analysed. Arbuscular mycorrhizal fungi (AMF) are a group of soil microorganisms that form symbiotic associations with the plants roots in order to improve growth and reproduction and it has been shown that they are valuable in the adaptation of plants to heavy metals contaminated sites.

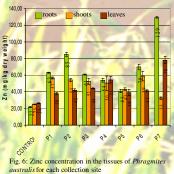
Fig.2: General view of the stream MATERIALS AND METHODS The harvesting occurred in the spring/summer season and was distributed by seven points, separated by 10 m from each other, along the course of the stream, R. ulmifolius was collected from one bank and P. australis from the other. Adjacent soil was also collected. Samples from non metal contaminated sites were taken as controls. The samples were then treated according to the following method . Fig..3: Sample of Rubus ulmifolius Zinc level collected in the area of the stream Incubation with extractant data Fig.4: General view of (EDTA, H<sub>2</sub>0, NH<sub>4</sub>-Ac) Collected analysis AAS Phragmites australis Drying Grinding (ANOVA) soil analysis of present in the area of the stream the digests Hot acid digestion RESULTS Separation AAS Hot acid No AMF were observed in any of the sampled roots: Harvested Washing Drving Grinding of roots. analysis confirmation of the inexistence of AMF should me digestion plants shoots and of the made by a sampling across the climate changes through leaves digests the year. Collection of fresh roots Mycorrhizal - The soil levels of zinc in the soil were variable, samples colonisation analysis ranging from 138 to 993 mg Zn/kg dry weight - For *P.australis* the concentrations of zinc ranged from 38 to 78 ppm in the leaves, 33 to 59 ppm in the 700.00 roots shoots leaves Table 1: Zinc concentration in the different collection sites of Rubus shoots and 41 to 129 ppm in the roots ulmifolius CONCLUSIONS 600,00 -For R. ulmifolius Zn in the tissues ranged from 51 to Zn (mg Zn/kg soil) Collection E 500,00 90 ppm for leaves, from 37 to 108 ppm for shoots and total H,O extract EDTA extract NH4-Ac extract The present field study shows that R1 957 ± 74 ª  $0.8 \pm 0.2$ from 151 to 537 ppm for roots. 87 ± 2 4 ± 0.2 ° 400.00 Rubus ulmifolius and Phragmites 992 ± 30 ª 102 ± 22 R2  $1.1 \pm 0.3$ 12 ± 1 australis colonize a heavy metal, -For P. australis, positive correlations were found 71 ± 2 300,00 715 ± 47 b  $1.1 \pm 0.2$ R3  $12 \pm 1$ specially zinc, contaminated site and between Zn concentrations in the soil with all the 588 ± 23 °  $0.9 \pm 0.2$ 42.3 ± 0.3 R4 3.5 ± 0.3 200,00 uptake and accumulate the metal, with different extractants and the zinc levels in the roots and 853 ± 13<sup>d</sup> R5 0.7 ± 0.3 42 ± 2 <sup>cd</sup> 10.7 ± 0.3 ° L Z no visual toxicity signs. shoots tissues R6 526 ± 12 ° 1.17 ± 0.06 29.0 ± 0.2 d  $3.9 \pm 0.2$ 100.00 In both species, the highest levels of R7 713 ± 55 b 1.0 ± 0.3 b 35 ± 2 d  $2.8 \pm 0.2$  d -For R. ulmifolius, positive correlations were found zinc were always found in the roots 0.00 Rcontro 196 ± 4 f 0.80 ± 0.07 ab 7.77 ± 0.05  $0.90 \pm 0.07^{\circ}$ between Zn concentrations in the soil with different CONTROL 22 80 tissues, which can be interpreted as a 4 20 5 20 Results are expressed as means ± SD (n=3). Means in the same column with extractants (with the exception of water extractable differ are significantly different from each other (p<0.05) acc low metal translocation to the the Tukey test. zinc) and the zinc levels in the roots and leaves tissues with stronger correlations being shown for the roots

## Table 3: Spearman's correlation coefficients between zind concentration in the plant parts and in the soil

<u> </u>		Zn soil factors				
62	7	total	H <sub>2</sub> O extractable	EDTA extractable	NH <sub>4</sub> -Ac extractable	
Roots	P. australis	0.541 **	0.427 *	0.472 **	0.818 **	
	R. ulmifolius	0.850 **	ns	0.947 *	0.779 **	
Shoots	P. australis	0.720 **	0.795 **	0.737 *	0,489 *	
	R. ulmifolius	ns	ns	ns	ns	
Leaves	P. australis	ns	ns	ns	ns	
	R. ulmifolius	0.557 **	ns	0.510 *	0.685 **	

\*\* correlation is significant at the 0.01 level; \* correlation is significant at the 0.05 level; ns, no significant correlation

Fig. 5: Zinc concentration in the tissues of Rubus folius for each collection site



ſ		Table 2: Zi australis	nc concentrati	on in the differe	ent collection s	ites of Phragm.			
9		Collection	Zn (mg Zn/kg soil)						
ſ		site	24.73		EDTA	NH <sub>4</sub> -Ac			
			total	H <sub>2</sub> O extract	extract	extract			
	Ŧ	P1	452 ± 24 ª	$8.0 \pm 0.4$ <sup>a</sup>	113 ± 3 ª	20 ± 2 ª			
1		P2	328 ± 46 bd	7.8 ± 0.5 ª	91 ± 1 <sup>b</sup>	14.9 ± 0.4 b			
		P3	$257 \pm 6^{d}$	3.2 ± 0.7 b	57 ± 0.8 °	12 ± 1 °			
		P4	360 ± 34 df	4.7 ± 0.4 °	102 ± 9 <sup>d</sup>	12.0 ± 0.4 °			
	Ŧ	P5	138 ± 4 °	$1.0 \pm 0.2$ d	36 ± 0.5 °	11 ± 1 °			

P6

296 ± 9 b P7 1.7 ± 0.06 f 67 ± 3  $21 \pm 1$ Pcontrol 97  $\pm$  2 <sup>g</sup> 1.6  $\pm$  0.1 <sup>f</sup> 15  $\pm$  2 <sup>g</sup> 7.5  $\pm$  0.4 <sup>c</sup> different letters are significantly different from each other (p<0.05) ac the Tukey test. Results are expressed as means ± SD (n=3). Means in the same column with rding to

387 ± 44 <sup>f</sup> 9.2 ± 0.03 <sup>c</sup> 101 ± 7

aboveground tissues. For this, the use of R. ulmifolius and P. australis for phytoextraction purposes does not appear as an effective method of metal removing, but these native metal toleran plant species can be used to reduce the effects of the contamination, avoiding Zn passing into the food chain. The positive correlations between zinc levels in plant tissues and zinc concentrations in the soil indicate that Zn concentrations in plant tissues increase with the increasing concentrations of

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Fig.1: Location of the studied region

total and available metal in the soil.

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f Phragmites

 $27 \pm 2$ 

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