

RHIZOREMEDIATION OF CO-CONTAMINANTS USING AUSTRALIAN NATIVE VEGETATION

Nanthi Bolan^{1, 2}, A. Kunhikrishnan³, B. Seshadri^{1, 2}, R. Naidu^{1, 2}

¹Global Centre for Environmental Remediation, The University of Newcastle (UON), Callaghan, New South Wales, Australia ²Cooperative Research Centre for Contamination Assessment and Remediation in the Environment, Salisbury, South Australia, Australia ³Chemical Safety Division, Department of Agro-Food Safety, National Academy of Agricultural Science, Wanju-gun, Jeollabuk-do, Republic of Korea <u>Nanthi.Bolan@newcastle.edu.au</u>

Keywords: Chromate; Trichloroethylene; Rhizosphere; Bioavailability

Introduction

Rhizosphere influences the dynamics of nutrients and contaminants through increased microbial activity, release of root exudates, and alteration of pH (Hinsinger et al., 2009). Chromate [Cr(VI)] and trichloroethylene (TCE) occur together in many contaminated sites (Lo et al., 2005). The bioavailability and ecotoxicity of Cr(VI) and TCE can be mitigated by enhancing their reduction reactions (Bolan et al., 2013a; Lo et al., 2005). The objective of this study was to evaluate the rhizosphere-induced reduction (i.e., rhizoreduction) and redistribution of Cr(VI) and TCE in Australian native vegetation in relation to their bioavailability.

Methods

The reduction of Cr(VI) and microbial activity by basal respiration were examined using rhizosphere and bulk soil samples from a number of Australian native vegetation (*Acacia pubescens, Eucalyptus camaldulensis, Enchylaena tomentosa, Templetonia retusa, Dichantheum sericeum,* and *Austrodanthonia richardsonii*). Naturally Cr(VI) contaminated and TCE spiked soils were used to examine the effect of *D. sericeum* on the redistribution and bioavailability of these contaminants following the modified plant growth experiment of Stanford and Dement (1957).

Results and Discussion

The rhizosphere soil contained higher levels of microbial activity, dissolved organic carbon (DOC), and organic acid content than the non-rhizosphere soil (Table 1). The batch experiment indicated that the rhizosphere soil caused up to 5.1 fold increases in the reduction rate of Cr(VI) (Table 1). There was a significant relationship between rhizosphere-induced increases in microbial activity and Cr(VI) reduction, indicating the role of increased microbial activity in rhizosphere soil on Cr(VI) reduction. In the plant growth experiment, *D. sericeum* decreased the concentrations of Cr(VI) and TCE in soil (Table 2). The decrease in Cr(VI) concentration was attributed mainly to the reduction of Cr(VI) to Cr(III), thereby reducing its bioavailability and mobility (Bolan et al., 2013b). The decrease in TCE concentrations may be

Proceedings of the 18th International Conference on Heavy Metals in the Environment, 12 to 15 September 2016, Ghent, Belgium *This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License.*

attributed to a number of processes including plant uptake, volatilization, and reduction (Graber et al., 2007). Reduction of TCE to ethylene has been shown to reduce its ecotoxicity.

 Table 1. Chemical and biological properties of rhizosphere and non-rhizosphere soils, and rhizosphere effect of Cr

 reduction

Plant species	Rhizosphere/non- rhizosphere	DOC (mg/kg)	Basal respiration (mg CO ₂ /kg)	Organic acid (mmol/kg)*	Rhizosphere effect of Cr reduction**
Acacia pubescens	Rhizosphere	2280	78.5	79.3	2.45
	Non-rhizosphere	674	22.2	15.3	
Eucalyptus	Rhizosphere	1154	39.2	45.1	2.56
camaldulensis	Non-rhizosphere	587	23.7	9.24	
Enchylaena	Rhizosphere	1212	44.3	34.5	3.89
tomentosa	Non-rhizosphere	347	18.7	7.12	
Templetonia retusa	Rhizosphere	1624	78.5	65.4	3.80
1	Non-rhizosphere	456	23.3	3.45	
Dichantheum	Rhizosphere	1897	95.3	54.1	5.07
sericeum	Non-rhizosphere	345	25.2	3.45	
Austrodanthonia	Rhizosphere	1434	36.5	31.3	2.66
richardsonii	Non-rhizosphere	768	24.3	6.45	

*mainly acetic and lactic acids in the non-rhizosphere soils; a range of acids including citric, malic, oxalic, malonic, fumaric, pyruvic, and trans-aconitic in the rhizosphere soil

**Rhizosphere effect = Ratio of rate of Cr reduction between rhizosphere and non-rhizosphere soils

Table 2. Cr(VI) and TCE distribution in soil in plant growth experiment

Contaminants*	Soil concentration	on (mg/kg)			
	With plants	Without plants			
Cr(VI)	43.7±7.25	124±18.2			
TCE	37.5±19.2	89.3±36.7			
*Original soil concentration (mg/kg): Cr(VI) = 257; TCE = 100					

Conclusion

Depending on the nature of contaminants present in soil, the rhizosphere-induced reduction by plant

species such as D. sericeum has implications to both their bioavailability to higher plants and

microorganisms, and remediation of contaminated soils. Further research is required to examine the plant-

induced decrease in TCE concentration in rhizosphere soil.

References

Bolan, N.S.; Kunhikrishnan, A.; Gibbs, J. (2013a). Rhizoreduction of Arsenate and Chromate in Australian Native Grass, Shrub and Tree Vegetation. *Plant Soil*, 367, 615–625.

Bolan, N.S.; Choppala, G.; Kunhikrishnan, A.; Park, J.H.; Naidu, R. (2013b). Microbial Transformation of Trace Elements in Soils in Relation to Bioavailability and Remediation. *Rev. Environ. Contam. Toxicol.*, 225, 1-56.

Graber, E.R.; Sorek, A.; Tsechansky, L.; Atzmon, N. (2007). Competitive Uptake of Trichloroethene and 1,1,1-Trichloroethane by *Eucalyptus camaldulensis* Seedlings and Wood. *Environ. Sci. Technol.*, *41*, 6704-10.

Hinsinger, P.; Bengough, G.; Vetterlein, D.; Young, I.M. (2009). Rhizosphere: Biophysics, Biochemistry and Ecological Relevance. *Plant Soil*, 321, 117-152.

Lo, I.; Lam, C.; Lai, K. (2005). Competitive Effects of Trichloroethylene on Cr(VI) Removal by Zero-valent Iron. J. Environ. Eng., 131, 1598–1606.

Stanford, G.; Dement, J.D. (1957). A Method for Measuring Short Term Nutrient Absorption by Plants: I. Phosphorus. Soil Sci. Soc. Am. J., 21, 612–617.

Proceedings of the 18th International Conference on Heavy Metals in the Environment, 12 to 15 September 2016, Ghent, Belgium *This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License.*