Adsorption of Cadmium and Lead lons on Calcite in Estuarine Waters and Seawater

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According to classification of Whitfield and Turner (1987), toxic cadmium shows deep-sea profiles characteristic of the recycled elements. It is held in the organic matrix via sul-fur bonds. Price and Morel (1990) reported cadmium substitu-tion for zinc in the marine diatom Thalassiosira weissflogii and have attempted to explain cadmium surface depletion rel-ative to deep waters.

Regarding inorganic particles, we have shown that cadmium had very poor affinity for SiO_2 , Al_2O_3 (Bilinski et al., 1976), for MNO₂ (**Bilinski et al**., 1977), for bentonite and kaolinite (**Bilinski et al**., 1990a). Cadmium could be ad-sorbed on synthesized northupite (**Vančina et al**., 1986) and on calcite (**Bilinski et al**., 1990b).

In the present work the primary interest is on the simulta-neous interaction of cadmium and lead ions with calcite, which mineral is present in surface sediments in the Krka River Estuary.

River Estuary. Cadmium and lead ions were added to natural water samples in low concentrations (8 x 10⁻⁸ M) to have solutions undersatu-rated with respect to CdCO₃(s) and PbCO₃(s). Adsorption isotherms of simultaneously and individually adsorbed cad-mium and lead ions show that the two ions do not compete for the same surface sites. This fact can be explained comparing ionic radii and chrystallographic space groups of corre-sponding metal carbonates. So, Ca²⁺ and Cd²⁺ have similar and Pb²⁺ has greater ionic radius (r(Ca²⁺) = 0.99 A; r(Cd²⁺) = 0.97 A; r(Pb²⁺) = 1.20 A). Calcium and cadmium carbonates of the unit cell (ASTM, 8-456 and 5-586) while lead carbon-ate crystallizes in Pmcn group (ASTM, 5-417). Behaviour of cadmium can be directly compared with one observed for zinc by Zachera et al. (1988). Zinc carbonate also shows R3C space group (ASTM, 8-469). Analogously to zinc ion, cadmium can exchange with calcium ion, present at the surface layer of calcite. As Ca²⁺ and CO₃²⁻ species have been identified as the only major surface ions on calcite (Thompson et al., 1989), it can be concluded that lead ion adsorbs via surface CO₃²⁻ group.

LITERATURE:

Whitfield, M., and Turner, D.R. (1987) Ch.17. The role of particles in regulating the composition of semwater. In: Aquatic Sunface Chemistry (ed. W. Stumm), John Wiley & Sons, New York, 457-493.

Price, N.M., and Morel, M.M. (1990), Cadmium and cobalt sub-stitution for zinc in a marine diatom, Nature, 344, 658-660.

Bilinski, H., Kozar, S., and Branica, M. (1976), Adsorption of trace heavy metals on particulate matter in seawater. In: Colloid and Interface Sci. V. (Ed. M. Kerker), Academic Press, New York, 211-231.

Bilinski, H., Kozar, S., Kwokal, Ż., and Branica, M. (1977), Model adsorption studies of Pb(II), Cu(II), Zn(II) and Cd(II) on MnO₂, added to Adriatic seawater samples, Thalassia Jugosl., 13, 101–108.

Bilinski, H., Kozar, S., Plavšić, M., Kwokal, Ż., and Brar ica, M. (1990a), Trace metal adsorption on inorganic solic phases under estuarine conditions, Mar. Chem., in press. solid

Vančina, V., Plavšić, M., Bilinski, H., Branica, M., and Millero, F.J. (1986), Preparation and solublity of northupite from brine and its adsorption properties for Cu(II) and Cd(II) in seawater, Geochim. Cosmochim. Acta 50, 1329-1336.

Bilinski, H., Kwokal, 2., Kozar, S., and Branica, M., (1990b), Scavenging of mercury and cadmium ions in the Krka River Estuary, Book of Abstr., 11th Int. Syme. "Chemistry of the Mediterraneam", "Reactivity of Chemical Species in Aquatic Environments", Primosten, May 9-16, 1990, p. 124.

Zachara, J.M., Kittrick, J.A., and Harsh, J.B. (1988), The mechanism of ${\rm Zn}^{2+}$ adsorption on calcite, Geochim. Cosmochim. Acta, 52, 2281–2291.

Thompson, D.W., and Pownall, P.M. (1989). Surface electrical properties of calcite, J. Colloid. Int. Sci., 131, 74-81.

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Fate of Trace Elements (Co, Sb, Zn) Entering Saronikos Gulf, Greece

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INTRODUCTION

Saronikos Gulf receiving the industrial and domestic wastes of the greater Athens area is among the most polluted gulfs of Greece. In order to study the fates and pathways of trace elements in Saronikos Gulf, several sampling cruises were organised and a big number of seawater and sediment core samples were collected from several stations (See Fig. 1). The trace element content of the samples was determined by Instrumental Neutron Activation Analysis. The results obtained for three selected trace metals namely cobalt, antimony and zinc, are presented and discussed.



Figure 1. Sampling Stations

DISCUSSION

DISCUSSION
In the main pollution sources of the trace elements studied are the hears Sewage Outfall (ASO) and a big Fertiliser Plant (FP) at the sources of Piraeus Harbour where the highest concentrations of Co. So and Zo in seawater and sediments were observed (See Tab. 1 and 2).
In settling of particles and flocs in the area of the outfall water column, since their concentrations decrease with distance of the sources to reach those of unpolluted coastal regions at a distance of 5-7 km. The rapid sedimentation was confirmed by the analysis of particulate matter which revealed that with the exception of the area close to the point sources where 15-25% of total Co. So and Zn were found to be associated with particulate matter, the buk of the area close to the point sources where 15-25% of total Co. So and Zn were found to be associated with particulate matter, the source stores of their total concentrations in the rest of the stations.
The concentrations of Co and Sb in sewater and sediments of Maronikos were almost equal to those of the Open Saronikos, were found to hear considered as typical concentrations for the sources was possible by means of some sources of 9.210.41 µg/g, Sb: 0.4340.13 µg/g and Zn: 4.813.0 µg/l. For the open Saronikos sediments the "respective of a disting the restment with 0.55 MCl, which extracts the "anthropogenic" fraction of trace elements (Aremia and Chau, 1976). It was found that outfle cores, while in sediments affected by the fertiliser plant sources was possible by means of from on the "anthropogenic" fraction of the sediments and examples, it was found that for any first the distribution of Co. Sb and Zn in the samples of trace elements of on Saronikos distribution in sediments affected by the fertiliser plant with other "anthropogenic" fraction of the sediments and examples, it was found that force samples, it was found that of the sediments fraction in a more for trace elements of on Saronikos Gult behave in a similar manner with other trace elements o

TABLE 1. Cobalt, antimony and zinc TABLE 2. Cobalt antimony and zinc in seawater $(\mu g/1)$. Average values are given silt-clay fraction of sediments $(\mu g/g)$. Average values are given in parentheses.

AREA	COBALT	ANTIMONY	ZINC	AREA	COBALT	ANTIMONY	ZINC
ASO, FP	0.12-0.60	0.60-1.40 (1.0)	9.0-41 (19)	Fertiliser Plant	35-87 (54)	14-100 (60)	840-2100 (1520)
Inner Saronikos	0.03-0.18 (0.10)	0.16-0.55 (0.29)	0.5-19 (10)	Sewage Outfall	13-24 (18)	6.6-32 (16)	220-620 (430)
Outer Saronikos	0.03-0.16 (0.09)	0.11~0.31 (0.23)	0.5-14 (4.8)	Inner Saronikos	8.6-9.6 (8.0)	0.48~0.72 (0.57)	81-110 (91)
				⁻ Outer Saronikos	7.8-10 (9.2)	0.31 - 0.60 (0.43)	68~100 (83)

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REFERENCES

Rapp. Comm. int. Mer Médit., 32, 1 (1990).

AGEMIAN, F. and A.S.Y. CHAU, 1976. Analyst, 101, 761-767. GRIMANIS A.P., D. ZAFIROPOULOS, N. KALOGEROPOULOS and M. VASSILAKI GRIMANIJ 1984. VIIes Journ. Etud. Pollution. Lucerne. CIESM, 391-397. GRIMANIS A.P., N. KALOGEROPOULOS and M. VASSILAKI-GRIMANI, 1988. Rapp. Comm. Int. Mer. Medit., 31, 2, p. 160. KALOGEROPOULOS N., M. SCOULLOS, M. VASSILAKI-GRIMANI and A.P. GRIMANIS, 1989. Sci. Total Environ., 79, 241-252.