Mercury reduction by vivianite

M. ETIQUE¹, S. BOUCHET¹, J. BYRNE², L. THOMAS ARRIGO¹, R. KAEGI³, R. KRETZSCHMAR¹

¹Soil Chemistry Group, IBP, ETH Zürich, Universitaetstrasse 16, CH-8092 Zürich (marjorie.etique@usys.ethz.ch)

²Geomicrobiology, Center for Applied Geosciences, University of Tuebingen, Sigwartsrasse 10, D-72076 Tuebingen

³Eawag, Swiss Federal Institute of Aquatic Science and Technology, Ueberlandstrasse 133, CH-8600 Duebendorf

Mercury (Hg) is a toxic element of global environmental concern, released into the environment by human activities since the late 1700s. In aquatic/terrestrial systems, Hg can be converted by anaerobic microorganisms to monomethylmercury (MeHg), a potent neurotoxin bioaccumulated and bioamplified in food webs. In iron-rich sediments of eutrophicated lakes, where Hg is methylated, vivianite, a ferrous phosphate mineral, commonly occurs in immediate vicinity of organic remains. However, Fe^{II}-bearing minerals such as magnetite [1] or green rusts [2] are recognized as Hg^{II} reducers. Here, the reduction of HgII to Hg0 by vivianite (Fe^{II}₃(PO₄)_{2.8H₂O) was investigated by fluctuating Hg/Fe} ratios (0.1, 1, 100) at circumneutral pH under anoxic conditions to illuminate kinetic parameters and the nature of the Fe^{III}-bearing minerals formed. The ability of vivianite to reduce inorganic divalent Hg is of high importance to better understand (i) the interplay between the Hg and Fe biogeochemical cycles, and to (ii) its potential role in limiting the production of MeHg in suboxic/anoxic sediments by the formation of elemental Hg, less available for methylating bacteria.



Figure 1. Percentage of inorganic divalent mercury (Hg2+) reduced by vivanite with various Fe/Hg ratios (100:1, 1:1, 1:10); (A). Elemental distribution map of vivianite nanoparticles after mercury reduction (B).

REFERENCES

[1] Wiatrowski H.A., et al. (2009) *Environ. Sci. Technol.*, 43:5307-5313. [2] Remy P.-Ph., et al. (2015) *Water Res.*, 70:266-278.