

Surface-induced redox processes: Oxidative transformation of oxytetracycline by Fe- and Mn- containing minerals

MARINA KARPOV¹, BETTINA SEIWERT², THORSTEN REEMTSMA², TAMARA POLUBESOVA¹ AND BENNY CHEFETZ^{1*}

¹ Department of Soil and Water Sciences, Faculty of Agriculture, Food and Environment, The Hebrew University of Jerusalem, P.O. Box 12, Rehovot, 7610001, Israel

(*correspondence: benny.chefetz@mail.huji.ac.il)

² Department of Analytical Chemistry, Helmholtz Centre for Environmental Research GmbH - UFZ, 04318 Leipzig, Germany

Pharmaceuticle degradation by Fe and Mn minerals

Transformation of pharmaceuticals by mineral surfaces significantly affects their environmental behaviour [1]. Manganese oxides and iron-enriched minerals are common redox-reactive species, participating in a variety of redox reactions resulting in the transformation of organic and inorganic pollutants in the environment [2].

In this work, we compared reaction kinetics and mechanisms of adsorption and oxidation of oxytetracycline (OTC), an antibacterial agent and an environmental pollutant [3], by Fe(III)-enriched montmorillonite and birnessite (δ -MnO₂).

This is the first work which investigated the oxidative transformation of OTC by Fe(III)-enriched mineral surface.

Results and discussion

OTC sorption (including inter-layer adsorption in case of Fe(III)-montmorillonite) and oxidation by both minerals were accompanied by their reductive dissolution. Participation of various radicals in OTC oxidative transformation, forming multiple detected transformation products, was suggested for both OTC-mineral systems. However, OTC transformation kinetics, its oxidation intensity and pathways, and transformation products, were distinct for Fe- and Mn-containing minerals due to different reduction potential and structure of the mineral surface.

[1] Qin, Chen & Zhuang (2015), *Crit. Rev. Environ. Sci. Technol.* **45**, 13, 1379–1408. [2] Borch, Kretzschmar, Kappler, Cappellen, Ginder-Vogel, Voegelin & Campbell (2010), *Environ. Sci. Technol.* **44**, 15-23. [3] Wang, Yao, Sun, Li & Huang (2016), *Environ. Sci. Technol.* **50**, 1, 145–153.