

# *In situ* spectral kinetics of Cr(VI) reduction by *c*-type cytochrome in a suspension of living *Shewanella putrefaciens* 200

XIAOMIN LI<sup>1,2</sup>, TONGXU LIU<sup>1\*</sup> AND FANGBAI LI<sup>1\*</sup>

<sup>1</sup>Guangdong Institute of Eco-Environmental and Soil Sciences, Guangzhou, P. R. China 510650

<sup>2</sup>School of Civil and Environmental Engineering, University of New South Wales, Sydney, NSW, Australia 2052

(\*E-mail: txliu@soil.gd.cn; cefbli@soil.gd.cn)

Although *c*-type cytochromes (*c*-Cyts) mediating metal reduction have been mainly investigated with *in vitro* purified proteins of dissimilatory metal reducing bacteria, the *in vivo* behavior of *c*-Cyts is still unclear. Here, *c*-Cyts in living *Shewanella putrefaciens* 200 (SP200) were successfully quantified using diffuse-transmission UV/Vis spectroscopy, and the *in situ* spectral kinetics of Cr(VI) reduction by *c*-Cyts were examined. A brief kinetic model was established with two predominant reactions, redox transformation of *c*-Cyts and Cr(VI) reduction by reduced *c*-Cyts, but the fitting curves were not well-matched with the experimental data from *c*-Cyts. The Cr(III)-induced toxic effect was then added to the model, resulting in substantially improved fitting curves to the experimental data. The Cr-induced toxic effect to the cellular function of redox transformation of *c*-Cyts was further confirmed by 16S rRNA analysis. This study provides a molecular-level insight into *in situ* microbial metal reduction processes and toxic effects of heavy metals under physiological conditions.

