Impact of cyanobacteria *Gloeocapsa* sp. on calcium carbonate precipitation rates

L.S. SHIROKOVA¹, O.S. POKROVSKY²*, J. SCHOTT², E.H. OELKERS², R. MARTINEZ² AND P. BENEZETH²

¹Institute of Ecological Problems of the North, Russian Academy of Science, Arkhangelsk, Russia (lshirocova@yandex.ru)

²Géochimie et Biogéochimie Experimentale, LMTG, CNRS, Universite de Toulouse, 14 Avenue Edouard Belin 31400, Toulouse, France (oleg@lmtg.obs-mip.fr)

The present study is aimed at determining the physicochemical factors governing calcium carbonate precipitation rates and mechanisms in the presence of cyanobacteria metabolic activity. The cyanobacteria Gloeocapsa sp. f-6gl were cultured for 2-14 days in calcite supersaturated, phosphate-free, Castenholz media at 25°C with and without air bubbling at pH from 7.8 to 10.8. Initial concentrations of Ca and HCO₃ ranged from 0.01 to 0.001 M. pH, optical density (cell biomass), [Ca], and [Alk] were measured as a function of time during the experiments. Adsorption of Ca on live and dead cyanobacteria was studied as a function of pH (2 to 12), Ca concentration in solution (0.1 µM to 1 mM), duration of exposure (1 hr to 2 days), and the metabolizing state of the culture (e.g. actively growing photosynthesizing cells under the light, non-photosynthesizing cells in the darkness, poisoned by contact with NaN3/formaline and sterilized biomass). These measurements allowed quantification of the number of surface sites able to participate in CaCO₃ precipitation via Ca²⁺ binding at the cell surface. The number of Ca²⁺-active surface sites is higher for inactivated cells and it does not depend on pH from 8.2 to 11.5. CaCO₃ precipitation rates appear be controlled by bulk solution pH and supersaturation state rather than live bacterial surfaces. SEM and XRD observations demonstrate that the cells are embedded in the calcite matrix stemming from non-metabolic mediated precipitation.

Selenium isotopes: a new oceanic paleoredox proxy?

ANDREW SHORE¹*, GAWEN JENKIN¹ AND THOMAS JOHNSON²

 ¹Department of Geology, University of Leicester, Leicester, LE1 7RH, UK (*correspondence: ajts1@le.ac.uk)
²Department of Geology, University of Illinois, Urbana-

Champaign, Illinois, 61801, USA

Previous work on selenium has tended to focus on understanding the biogeochemical cycle of selenium, driven by the need to understand and manage its environmental behaviour due to its toxicity in moderate concentrations. Due to their redox-sensitive nature, and the tendency for massdependent fractionation to be induced by redox reactions, some authors have begun investigating whether the isotopes of elements such as Mo, Fe and Se can be used in palaeoredox studies (e.g. Mo [1]).

A record of sedimentary selenium isotope ratios (⁸²⁷⁶Se), spanning the last 500 ka, has been produced. We propose differences in the cycling of Se due to variations in water column oxygen content will affect seawater selenium isotope ratios and sediment components deposited from the seawater. Sediments deposited under an anoxic water column are dominated by organically - bound Se which has little fractionation with the water. In contrast, samples deposited under an oxic water column of authigenic Se which should be enriched in light isotopes relative to the water.

Oxygenation of the oceans is difficult to constrain. Determination of past ocean oxygenation provides important constraints on atmospheric O_2 and the overall "health" of the oceans, information crucial for understanding the development of life.

[1] Siebert *et al.* (2003) *Earth & Planetary Science Letters* **211**, 159 – 171.