Atmospheric Methane over the past 400 million years

ROBERT A. BERNER¹, DAVID BEERLING², FRED T. MACKENZIE³, MICHAEL B. HARFOOT⁴ AND JOHN A. PYLE⁴

¹Yale University, New Haven, CT

(*correspondence: robert.berner@yale.edu)

²University of Sheffield, Sheffield, UK

³University of Hawaii, Honolulu, HI

⁴Centre for Atmospheric Science, University of Cambridge, UK

Long term variations in atmospheric CH₄ concentration, excluding short bursts from hydrate reservoirs, have been estimated for the past 400 Ma. We constructed a steady state model that combines estimates of wetland CH₄ emission, based on the percentage abundance of coal basin sediments and original global rates of clastic sedimentation, with a Cambridge 2-D chemistry-transport model to determine loss by reaction with OH radical. The results [1] indicate that tropospheric CH₄ concentrations reached exceptionally high values of ~12,000 ppb during the Permo-Carboniferous, when tropical swamplands were widespread, fell to minimum levels (~100 ppb) during the Triassic 'coal gap', averaged around 2000-4000ppb during the Mesozoic and < 1000 ppb in the Cenozoic. The high Permo-Carboniferous levels made an appreciable methane contribution to mean earth surface temperature (see Figure) that allowed for lack of glaciation in paleo-Siberia while still permitting continent-wide glaciation in the southern hemisphere. The moderately high Mesozoic CH₄ levels were due partly to CO₂-driven warming resulting in increased CH4 release from wetlands. Other than during the Permo-Carboniferous, over the rest of Phanerozoic history methane appears to have played a secondary role to atmospheric CO₂ and the evolution of the Sun in driving climate change.

[1] Beerling et al. (2009) Am J. Sci. 309, 97-113.

Microbial metal reduction: Insights from genomics and transcriptomics

RIZLAN BERNIER-LATMANI* AND PILAR JUNIER

Environmental Microbiology Laboratory, Ecole Polytechnique Federale de Lausanne, Switzerland (*correspondence: rizlan.bernier-latmani@epfl.ch)

Microbial metal reduction can be catalyzed by metal- and sulfate-reducing bacteria. In particular, Shewanella oneidensis MR-1, a metal-reducing bacterium (MRB), and Desulfotomaculum reducens MI-1, a Gram positive sulfatereducing bacterium (SRB), have been shown to couple growth to Fe(III) reduction and to reduce U(VI). The mechanisms of Fe(III) and U(VI) reduction by S. oneidensis have been studied extensively using biochemical, transcriptomic and mutagenic approaches and have been shown to involve periplasmic and outer-membrane *c*-type cytochromes for both metals. The genome of S. oneidensis contains 42 c-type cytochromes which account for the metabolic versatility of this organism. In contrast, the mechanism of metal reduction by D. reducens remains unknown. Recent results have shown that U(VI) and Fe(III) reduction in D. reducens proceed through different pathways as vegetative cells grow with Fe(III) and spores reduce U(VI) with no apparent metabolic benefit. Reduction by spores requires the presence of spent growth medium suggesting the requirement for an endogenous factor to reduce U(VI). Furthermore, Fe(III) reduction by D. reducens differs significantly from that in S. oneidensis as the former lacks an outer membrane. In addition, the D. reducens genome only encodes for a single triheme c-type cytochrome annotated as a nitrite reductase. A transcriptomic study targeting Fe(III) reduction in D. reducens was conducted in order to investigate the role of the single *c*-type cytochrome and that of the nine hydrogenases in the genome. In addition, a second transcriptomic study was conducted to evaluate the effect of U(VI) on the metabolism of fermentatively-grown D. reducens cells.We report the results of these transcriptomic studies and compare them to previous results for S. oneidensis.