



Biomineralisation and Gene Expression in *Emiliana huxleyi*

Luke Mackinder (1,2), Glen Wheeler (2,3), Declan Schroeder (2), Ulf Riebesell (1), and Colin Brownlee (2)

(1) IFM-GEOMAR, Kiel, Germany (lmackinder@ifm-geomar.de), (2) Marine Biological Association, The Laboratory, Citadel Hill, Plymouth, UK, (3) Plymouth Marine Laboratory, Prospect Place, Plymouth, UK

Biomineralisation in the marine phytoplankton *Emiliana huxleyi* is a stringently controlled intracellular process. The molecular basis of coccolith production is still relatively unknown although its importance in global biogeochemical cycles and sensitivity to increased pCO₂ levels has been well documented. Unravelling the cellular processes of biomineralisation including ion transport pathways and mineral nucleation will aid in our understanding of the effects of a changing ocean on this group of phytoplankton and could also shed light on isotope fractionation of elements used for proxies. This study uses three calcifying vs. non-calcifying comparisons to investigate the role of putative calcification genes in *E. huxleyi*. It analyses relative transcript levels using quantitative reverse transcription PCR. Calcification consistently induces the up-regulation of genes encoding a putative bicarbonate transporter belonging to the solute carrier 4 family, a calcium proton exchanger and a vacuolar H⁺-ATPase in comparison to genetically identical non-calcifying cells. In contrast up to a 26 fold down-regulation of GPA, a calcium binding protein, is seen when cells are in their calcifying form. This data provides strong evidence that these genes play key roles in *E. huxleyi* biomineralisation.