Geophysical Research Abstracts, Vol. 8, 05410, 2006 SRef-ID: 1607-7962/gra/EGU06-A-05410 © European Geosciences Union 2006



## The unexpected impact of ocean acidification on the production of dimethyl sulfide and other important climate relevant organic gases

**O. W. Wingenter** (1), K. B. Haase (1), M. Zeigler (1), D. R. Blake (2), K. Schulz (3), U. Riebesell (3)

(1) Department of Chemistry and the Geophysical Research Center, New Mexico Institute of Mining and Technology, Socorro, NM, 87801 USA, (2) Department of Chemistry, University of California, Irvine, CA, 92697 USA, (3) Leibniz Institute for Marine Sciences IFM-GEOMAR, Kiel, Germany

Nearly 60 hydrocarbons, halocarbons, alkyl nitrates and organic sulfur trace gases were quantified from sea water and air samples taken from the nine mesocosms over the 24 day PeECE III experiment. Three replicate mesocosms of three scenarios representing the present day  $CO_2$  atmospheric concentrations (equilibrated to 375 ppmv of atmospheric  $CO_2$ ), future  $CO_2$  concentrations anticipated by the end of this century (when considering the IPCC's business as usual scenario, 750 ppmv of  $CO_2$ ), and a triple present day  $CO_2$  case (1150 ppmv  $CO_2$ ) were sampled each day. Peak dimethyl sulfide (DMS) concentrations corresponded with the bloom in *Emiliania huxleyi*. The peak in DMS came later in the double and triple  $CO_2$  mesocosms and the time integrated average amount of DMS was 20 and 15 percent higher in the double and triple CO2 "Worlds", respectively. Results of DMS measurements made by other researchers showed similar results. Chloroiodomethane had its peak concentration much later than the maximum in chlorophyll corresponding to the decline of the phytoplankton bloom. (At this time DMS experienced a secondary maximum.) When integrating the  $CH_2CII$ concentration with respect to time, we estimate that production of  $CH_2CII$  is about 40 percent higher in the 2xCO<sub>2</sub> and about 100 percent higher in the 3xCO<sub>2</sub> mesocosms.

Marine production of DMS critically impacts the radiative properties of the Earth because it is a major source of cloud condensation nuclei in the clean marine environment. A significant perturbation to its production and subsequent sea-to-air flux will alter the Earth's albedo. Marine production of  $CH_2CII$  is a major source of atmospheric iodine (I). Once released to the atmosphere, it participates in catalytic ozone (O<sub>3</sub>) destruction. The concentration of O<sub>3</sub> is pivotal to the oxidative capacity of the atmosphere because it is a primary hydroxyl (HO) precursor. Significant future  $CH_2CII$ emissions would result in lower O<sub>3</sub> mixing ratios and ultimately lead to longer lifetimes of some greenhouse gases such as methane. Results and climatic implications of other gases will also be presented. Perturbations of these gases, by the few marine microorganisms studied during PeECE III, suggests the potential impact on other marine climate relevant gases as a result of projected ocean acidification.