

Benthic O₂ fluxes measured by Eddy Covariance in a large flume facility

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The Eddy Covariance: a new technique to assess sediment-water O₂ exchanges over various environments

The Eddy Covariance (EC) is a novel technique used to quantify the sedimentary O₂ consumption of marine ecosystems [1]. It measures O₂ exchanges at the sediment-water interface over large spatial scales (> 100 m²) without being intrusive or disturbance of the flow field. This new technique has recently been used in various natural environments as tropical lagoon [2], tidal flat [3], river [4] and deep ocean [5] systems.

In autumn 2011, EC deployments were carried out in a 18 m long flume tank (NIOZ Yerseke, The Netherlands) filled with muddy sediment (Figure 1). Based on simultaneous measurements of near-sediment velocities (ADV Vector, Nortek) and O₂ concentrations (Unisense Clark-type microelectrode) at high frequency (64 Hz), O₂ exchanges at the sediment-water interface were calculated by EC. These EC fluxes were then compared with values based on microsensors profiling and chamber incubation.



Figure 1: The flume Eddy Covariance deployment (Sept. 2011) showing sensor set-up (velocimeter, left and microelectrode, right).

First results and conclusions

EC O₂ fluxes were similar or greater than those obtained by incubation and microsensors profiling, i.e. -23.8~-62.9, -42.2~-56.0 and -15.8~-19.5 mmol m⁻² d⁻¹ respectively. EC fluxes varied with flow velocity, sensor depth and upstream location. The deployment of the EC technique in the flume facility allows to investigate various aspects of boundary layer hydrodynamics under controlled conditions.

[1] Berg et al. (2003) *Mar. Ecol. Prog. Ser.* **261**, 75-83. [2] Hume et al. (2011) *Limnol. Oceanogr.* **56**, 86-96. [3] Kuwae et al. (2006) *Mar. Ecol. Prog. Ser.* **307**, 59-68. [4] McGinnis et al. (2008) *Geophys. Res. Lett.* **35**, doi:10.1029/2007GL032747. [5] Berg et al. (2009) *Limnol. Oceanogr. Meth.* **7**, 576-584.

Theoretical carbon isotope fractionation under deep-earth conditions

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Introduction

In addition to the major component of mantle carbon, which has δ¹³C values of about -5±3‰, some reduced (carbide, hydrocarbons) and neutral (diamond, graphite, dissolved C) carbon compounds from mantle xenoliths and other sources have δ¹³C values in the range of -20 to -35‰. Contamination of organic carbons near the Earth surface and the recycling of sedimentary organic matter via subduction zones have often been invoked to explain the depleted ¹³C signatures of these mantle rocks. However an increasing number of studies do not support this scenario. Two other possibilities exist. First, mantle carbon could have been isotopically heterogeneous since the accretion from the solar nebula and the core-mantle segregation in the first 100 My of Earth history. Chondrites, carbonaceous chondrites particularly, which are widely considered to represent primitive undifferentiated materials from which the Earth accreted, have δ¹³C values ranging widely from -28 to 0‰. Another possibility is that there exist some mechanisms and processes for large carbon isotope fractionation at high temperatures and pressures. For example, Craig [1] and Deines and Wickman [2] reported ca. 12 ‰ differences between graphite and cohenite, (Fe, Ni, Co)₃C, from iron meteorites. A recent experimental study by Satish-Kumar *et al.* [3] reported 2.7 - 4.5 ‰ differences between graphite/diamond and FeC melt at 1350 - 2100°C and 5 - 10 GPa.

Theoretical study

Here, we report our updated results of theoretical calculations on ¹³C/¹²C fractionation among major C-bearing materials (CO₂, calcite, diamond, graphite, and SiC). Our results show that SiC is depleted in ¹³C relative to diamond/graphite and calcite even at very high temperatures: 2.5 - 9 ‰ at 1000 - 2000°C. Our results for SiC is consistent with the experimental results for FeC melt [3]. High pressures (>10's of GPa) under deep-Earth conditions may also affect measurably equilibrium isotope fractionation of C-bearing species [4]. Thus, the recent experimental study and our theoretical calculations clearly demonstrate that carbides (Fe₃C and SiC) are significantly depleted in ¹³C at high temperatures and pressures. We need to revise and improve our understanding of carbon cycles and associated isotope fractionation in the deep-Earth.

[1] Craig (1953) *GCA* **3**, 53-92.

[2] Deines and Wickman (1975) *GCA* **39**, 547-557

[3] Satish-Kumar *et al.* (2011) *EPSL* **310**, 340-348.

[4] Polyakov & Kharlashina (1994) *GCA* **58**, 4739-4750.