

Biomarker and stable isotope analysis linked to methane cycling in Lake Untersee, Antarctica

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Lake Untersee, in East Antarctica is a perennially ice-covered lake (2 – 4 m) consisting of two basins; a well mixed oxic basin ca. 160 m deep and a smaller, anoxic basin ca. 100 m separated by a sill at 50 m depth. Analysis of microbial phospholipid fatty acids (PLFA) and ether lipids (archaeol and GDGT-0) and their isotopic compositions was used to constrain methane cycling and the recycling of carbon within the anoxic basin water column and sediments.

High concentrations of methane (8.5 mmol/L) were present within the bottom waters of the anoxic basin, derived from sedimentary methanogens as shown by archaeol concentrations of 25 µg/g. Methane concentrations decreased slightly towards the anoxic/oxic interface. Methane is derived from CO₂ reduction based on its isotopic composition (δ¹³C: -50‰; δD: -438‰ at 95m) and that of the corresponding DIC (δ¹³C ca. +25‰ at 95m). Methane oxidation occurred within a suboxic transition zone present at ca. 70 – 80m. Within this zone [CH₄] decreased from 7.3 mmol/L at 85 m to 0.02 mmol/L by 72 m. Concurrently, δ¹³C-CH₄ values increased, from -51‰ at 85m to -40‰ at 72m, consistent with CH₄ oxidation.

PLFA concentrations varied throughout the water column, but generally increased with depth. Between 20 and 72m, values ranged from 0.2 – 3.8 µg/L (72m: 3.5 x 10⁵ cells/L). Higher biomass was observed at the bottom of the suboxic zone: 78 and 80 m were 8.5 and 12.9 µg/L respectively (80m: 1.3 x 10⁶ cells/L). The increase in PLFA concentrations at the bottom of the suboxic zone, combined with depleted δ¹³C_{PLFA} values (δ¹³C-16:1: -36.8‰ at 76m) was consistent with a methanotrophic community supported by methane derived from the sediments. PLFA concentrations of 57 µg/g (5.4 x 10⁹ cells/g) and δ¹³C_{PLFA} values of ca. -20 to -25‰ within the sediments suggests an active heterotroph population producing CO₂ via organic degradation supporting the methanogenic community.

Identification of biomarkers created by the modern, viable bacterial and archaeal community is fundamental to understanding the preservation of biomarkers within palaeo-lacustrine environments that are relevant analogues to early Earth and Mars.