



A setup to analyze isotopes of paleoatmospheric CH₄ and N₂O in a single ice core sample

P. Sperlich (1), C. Buizert (1), M. Guillevic (1,2), T. Jenk (1), C.J. Sapart (3), T. Röckmann (3), M. Bock (4), J. Schmitt (4), H. Fischer (4), and T. Blunier (1)

(1) Centre for Ice and Climate, Niels Bohr Institute, University of Copenhagen, Copenhagen, Denmark (sperlich@nbi.ku.dk), (2) Laboratoire des Sciences du Climat et de l'Environnement, Gif sur Yvette, France, (3) Institute for Marine and Atmospheric Research Utrecht, Utrecht University, Utrecht, The Netherlands, (4) Climate and Environmental Physics and Oeschger Centre for Climate Change Research, University of Bern, Bern, Switzerland

The isotopic composition of paleoatmospheric N₂O and CH₄ can be measured in ice core samples to study biogeochemical processes driving their atmospheric budget. We developed a new vacuum-wet-extraction method to measure CH₄ concentrations, $\delta^{15}\text{N}$ and $\delta^{18}\text{O}$ isotope ratios of N₂O, and $\delta^{13}\text{C}$ of CH₄ in a single ice core sample using one mass spectrometer. The ice sample is melted while the air is continuously cryo-pumped onto a liquid nitrogen cooled Haysep trap. N₂ and O₂ are separated and measured with a TCD before CH₄ and N₂O and are measured by GC-IRMS. We determine the CH₄ concentrations with a precision of ± 20 ppbv by relating the IRMS peak size to the TCD peak size.

First measurements of seven pre-industrial ice core samples of 400 – 500 g agree with a standard deviation (1σ) of better than 0.05 ‰ in $\delta^{13}\text{C} - \text{CH}_4$. At this stage, the setup needs improvement for satisfactory measurements of N₂O isotopes and concentrations from ice core samples. Intercomparisons of three standard gases have been shared for N₂O and CH₄ isotopes between three laboratories. The agreement is typically within the standard deviation (1σ) for all isotope measurements.