



Continuous analysis of stable carbon isotopes in CO₂ and CH₄ fluxes with an automatic chamber flux system at Mer Bleue bog

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Northern peatlands are one of the largest natural sources for methane (CH₄) to the atmosphere and thus patterns and controls of CH₄ fluxes from these systems are intensively studied. To understand the atmospheric CH₄ budget, different isotopic signatures are typically assigned to specific sources. However, isotopic signatures of CH₄ from wetlands may not be stable and instead vary with methanogenic and emission pathways which influence sources of carbon and fractionation. The aim of this study was to assess the temporal and spatial variability in the isotopic composition of CH₄ and carbon dioxide (CO₂) emitted throughout the growing season at Mer Bleue, a temperate ombrotrophic bog in Canada. To this end, laser based stable isotope analyzers for ¹³C in CO₂ and CH₄ were attached to an automated flux chamber system operated at the site. Using 12 different transparent chambers, we measured differences in isotopic composition of CO₂ and CH₄ and compared the observed patterns to vegetation characteristics and air and soil temperatures. We specifically compared sedge dominated chambers (affected by aerenchyma gas transport), shrub (mostly *Chamaedaphne calyculata*), and *Sphagnum* moss dominated chambers.

The CH₄ fluxes in the sedge dominated chambers were much higher than the other vegetation types, as expected, but also had a different isotopic composition, presumably due to the different gas transport mechanisms. In the presence of aerenchyma, the methanotrophic layer is effectively bypassed and aerobic CH₄ oxidation is significantly reduced. This lead to lower δ¹³C values (in the range of 5-10 ‰ in CH₄ emitted on sedge dominated plots, compared to less negative δ¹³C values observed at plots dominated by shrubs or *Sphagnum*. As the chamber system uses transparent chambers, δ¹³C values in CO₂ were predominantly driven by photosynthetic activity, heterotrophic and autotrophic respiration. Data analysis is still ongoing to attempt to identify the effects of CH₄ oxidation δ¹³C values in CO₂. Overall our study demonstrates the value of online, laser-based stable isotope analysis in CH₄ and CO₂, although the accuracy and calibration of such instruments at low atmospheric concentrations remains challenging.