



## **Contrasting isotope patterns of NO<sub>3</sub>- and N<sub>2</sub>O in polluted ombrotrophic peat bogs help to explain negligible N<sub>2</sub>O emissions**

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A quantitative assessment of N<sub>2</sub>O production and consumption in soils under various environmental conditions is needed for the formulation of efficient mitigation strategies. In an era of climatic change, wetlands may serve as a major source of N<sub>2</sub>O for the atmosphere, and thus contribute to further warming. We studied N cycling in three high-elevation, rain-fed, Sphagnum-dominated peat bogs in the Czech Republic (Central Europe). Despite a history of medium-to-high N deposition rates (10-40 kg N/ha/yr), we found negligible N<sub>2</sub>O emission rates in the Eagle Mts. and Sumava Mts. At a depth of 60 cm, the least N-polluted site, exhibited nearly 7 times higher porewater N<sub>2</sub>O concentrations, compared to the most N-polluted site. This was probably related to the higher denitrification potential, expressed in the abundance of nirK and nirS genes. Upcore, N<sub>2</sub>O concentrations dramatically decreased, while delta15N-N<sub>2</sub>O values systematically increased. At one mountain-top peat bog in the Ore Mts., we simultaneously determined vertical delta15N trends in peat porewater NO<sub>3</sub>- and N<sub>2</sub>O. Our study, the first in its kind, revealed a strong negative correlation between these two variables at depths where most N<sub>2</sub>O is formed (30-60 cm below peat surface). N<sub>2</sub>O concentrations in the peat pore water were 20 % lower than in the lowermost atmosphere, indicating that the peat serves, at least intermittently, as a sink for atmospheric N<sub>2</sub>O, not as a N<sub>2</sub>O source. Upcore, higher delta15N values of N<sub>2</sub>O were accompanied by lower delta15N values of NO<sub>3</sub>-. We suggest that such N isotope systematics can help to distinguish between N<sub>2</sub>O diffusion in peat pore water, N<sub>2</sub>O production from NO<sub>3</sub>- via denitrification, and N<sub>2</sub>O consumption via further reduction to harmless N<sub>2</sub>. Heavier N<sub>2</sub>O-N upcore corresponded to residual N<sub>2</sub>O following partial reduction to N<sub>2</sub> and a loss of warming potential. Upcore, increasing proportion of atmospheric N<sub>2</sub>O was also isotopically detected.