

Source portioning of N₂O emissions after long term elevation of soil temperature in a permanent grassland soil

Anne Jansen-Willems (1,2), Gary Lanigan (1), Timothy Clough (3), Louise Andresen (2), Christoph Müller (2,4)
(1) Teagasc Johnstown Castle, Wexford, Co. Wexford, Ireland (Anne.jansen@teagasc.ie), (2) Department of Plant Ecology, Justus Liebig University Giessen, Heinrich-Buff-Ring 26, 35392 Giessen, Germany (pflanzenoekologie@bio.uni-giessen.de), (3) Department of Soil and Physical Sciences, Faculty of Agriculture and Life Sciences, New Zealand (Timothy.Clough@lincoln.ac.nz), (4) School of Biology and Environmental Science, University College Dublin, Dublin, Ireland (Christoph.mueller@ucd.ie)

Several methods, such as source portioning, have been used to quantify the contributions of individual N pools to N₂O emissions. These methods however, assume the absence of hybrid reactions such as co-denitrification, which were previously identified as important. A straight forward method portioning N₂O fluxes into four different production processes, including a hybrid reaction, was therefore developed. This method portioned the N₂O fluxes in nitrification, denitrification, oxidation of organic matter and co-denitrification, using data on ⁴⁵R and ⁴⁶R of the N₂O flux and the ¹⁵N content of the NO₃⁻ and NH₄⁺ in the soil. This newly developed method was used to analyse the N₂O emissions from incubated soil, which was previously subjected to 6 years of elevated soil temperature of +0, +1, +2 or +3 °C. N₂O emissions were measured and analysed at four time points in the six days following, NO₃¹⁵NH₄ Gly or ¹⁵NO₃NH₄ Gly, label addition. The oxidation of organic N was found to be the main source of N₂O fluxes at all sampling dates, comprising between 63 and 85% of the total N₂O flux. The percentage contribution made by organic N to N₂O fluxes increased over the sampling period, rising from a minimum of 40% in the control treatment, to virtually 100% across all treatments by Day 6. Compared to the control treatment, denitrification contributed less to N₂O from soil subjected to +2 and +3 °C warming (p < 0.0001 and p = 0.002, respectively). Co-denitrification only contributed to the N₂O flux during the first day after substrate addition. The highest amount of N₂O produced via co-denitrification was found under the control treatment. From soil subjected to +2 and +3 °C treatments, the contribution of co-denitrification was minor. However, these differences in co-denitrification were not significant. This research showed the importance of the oxidation of organic N in N₂O emissions. It should therefore not be omitted as a potential source in source portioning. Emissions of N₂O in the first six days after fertilisation decreased for soils previously subjected to higher temperatures as a consequence of a reduction in the rates of denitrification and the oxidation of organic N.