



Effects of elevated atmospheric CO₂, prolonged summer drought and temperature increase on N₂O and CH₄ fluxes in a temperate heathland

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ABSTRACT

In temperate regions, climate change is predicted to increase annual mean temperature and intensify the duration and frequency of summer droughts, which together with elevated atmospheric carbon dioxide (CO₂) concentrations, may affect the exchange of nitrous oxide (N₂O) and methane (CH₄) between terrestrial ecosystems and the atmosphere. We report results from the CLIMAITE experiment, where the effects of these three climate change parameters were investigated solely and in all combinations in a temperate heathland. Field measurements of N₂O and CH₄ fluxes took place 1–2 years after the climate change manipulations were initiated. The soil was generally a net sink for atmospheric CH₄. Elevated temperature (T) increased the CH₄ uptake by on average 10 μg C m⁻² h⁻¹, corresponding to a rise in the uptake rate of about 20%. However, during winter elevated CO₂ (CO₂) reduced the CH₄ uptake, which outweighed the positive effect of warming when analyzed across the study period. Emissions of N₂O were generally low (<10 μg N m⁻² h⁻¹). As single experimental factors, elevated CO₂, temperature and summer drought (D) had no major effect on the N₂O fluxes, but the combination of CO₂ and warming (TCO₂) stimulated N₂O emission, whereas the N₂O emission ceased when CO₂ was combined with drought (DCO₂). We suggest that these N₂O responses are related to increased rhizodeposition under elevated CO₂ combined with increased and reduced nitrogen turnover rates caused by warming and drought, respectively. The N₂O flux in the multifactor treatment TDCO₂ was not different from the ambient control treatment. Overall, our study suggests that in the future, CH₄ uptake may increase slightly, while N₂O emission will remain unchanged in temperate ecosystems on well-aerated soils. However, we propose that continued exposure to altered climate could potentially change the greenhouse gas flux pattern in the investigated heathland.

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1. Introduction

Nitrous oxide (N₂O) and methane (CH₄) are important greenhouse gases (IPCC, 2007), and N₂O is the dominant substance responsible for depletion of the stratospheric ozone layer (Ravishankara et al., 2009). On a global scale, N₂O emissions from unfertilized grassland/steppe, including heathlands, is estimated to be 0.4 Tg N₂O–N y⁻¹ (Stehfest and Bouwman, 2006), which corresponds to about 2% of the total annual N₂O emissions (Fowler et al., 2009). Methane uptake by aerobic soils worldwide is estimated to be 30 Tg CH₄ y⁻¹, counteracting 6% of the total emission of

CH₄ from natural and anthropogenic sources (Wuebbles and Hayhoe, 2002).

Nitrous oxide emitted from soil primarily originates from the two microbial processes nitrification and denitrification, which occur under aerobic and anaerobic soil conditions, respectively (Wrage et al., 2001). Thus, both processes are controlled by soil moisture that regulates oxygen (O₂) availability and by supply of substrates, ammonium (NH₄⁺) or nitrate (NO₃⁻). In addition, denitrifying bacteria need labile carbon (C) compounds as an energy source. Soils may act as a sink for atmospheric N₂O, which has been observed at low mineral nitrogen (N) availability and the responsible organisms could be denitrifying bacteria, but are probably also nitrifying organisms (Chapuis-Lardy et al., 2007).

The CH₄ flux between soil and atmosphere is the net result of CH₄ production by methanogenic archaea and CH₄ oxidation by methanotrophic bacteria (Le Mer and Roger, 2001). In aerobic soils,

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CH₄ oxidation typically proceeds at a greater rate than CH₄ production, resulting in net uptake of atmospheric CH₄. Maximum CH₄ oxidation usually occurs in deeper soil layers. Thus, CH₄ uptake is strongly controlled by the physical diffusion of atmospheric CH₄ through the soil profile, which is mainly regulated by the soil texture and water content (King, 1997), as molecular diffusion in water is four orders of magnitude slower than in air.

The effect of drought on fluxes of CH₄ and N₂O has only been investigated in a limited number of studies in temperate ecosystems on aerobic soils. In a temperate spruce forest, Borken et al. (2000) found that prolonged summer drought increased the annual CH₄ uptake by more than 40%. However, in a deciduous forest on well-drained soil, the CH₄ uptake was only increased by 7% because soil in the control plots already had a low water content due to effective drainage (Borken et al., 2006). Goldberg and Gebauer (2009) reported that summer drought reduced the N₂O emission from a spruce forest.

Elevated atmospheric carbon dioxide (CO₂) concentrations affect soil properties via plant-mediated processes, and thereby potentially the fluxes of N₂O and CH₄. It is well known that plants growing under elevated CO₂ may reduce their transpiration, leading to increased water use efficiency and soil moisture contents (Morgan et al., 2004; Robredo et al., 2007). Furthermore, a common plant response to elevated CO₂ is increased deposition of root-derived C into the rhizosphere (Allard et al., 2006; Pendall et al., 2004).

The effect of elevated atmospheric CO₂ on N₂O fluxes has been investigated intensively through the last decade in a variety of ecosystems. Long-term studies (≥ 1 year) in natural or semi-natural ecosystems on well-aerated soils with low N availability show either no response in annual N₂O emissions rates to atmospheric CO₂ levels (Baggs et al., 2003; Mosier et al., 2002; Phillips et al., 2001b) or increased N₂O emissions from soil under elevated CO₂ (Kammann et al., 2008). The positive effect of elevated CO₂ on the N₂O emission was explained by increased rhizodeposition of labile C substrates stimulating denitrification. In an N-limited pine forest, Phillips et al. (2001b) observed seasonal variability in the N₂O response to elevated atmospheric CO₂ compared to an ambient control. This included reduced N₂O emissions during the growing season due to high plant-microbial competition for N, and enhanced N₂O emissions during winter possibly because denitrification was stimulated by greater soil moisture and labile C sources under elevated CO₂. In a short-term study during autumn, Arnone and Bohlen (1998) also found that elevated atmospheric CO₂ increased N₂O emission, which they ascribed to improved soil moisture conditions in a relatively dry grassland favouring the microbial transformation of N.

Reduced CH₄ consumption under elevated atmospheric CO₂ has been observed in several ecosystems on undisturbed aerobic soils (Ambus and Robertson, 1999; Baggs and Blum, 2004; Dubbs and Whalen, 2010; Ineson et al., 1998; Phillips et al., 2001a), but the exact mechanism is not well known and may vary between ecosystems. Two possible mechanisms have been suggested for the reducing effect of elevated CO₂ on net CH₄ consumption: i) decreased CH₄ oxidation due to higher soil water content and thereby reduced diffusion of CH₄ (Ambus and Robertson, 1999; Baggs and Blum, 2004) and ii) increased CH₄ production due to anaerobic microsites caused by reduced O₂ diffusion and increased respiration (McLain and Ahmann, 2008).

Effects of elevated temperature on N₂O and CH₄ fluxes have only been investigated in a few field-scale warming experiments located in temperate ecosystems on well-aerated soils (McHale et al., 1998; Peterjohn et al., 1994; Rustad and Fernandez, 1998). The artificial warming had either no effect on CH₄ flux or it increased CH₄ uptake, which could be related to the observed warming-induced

declines in soil moisture in certain soil layers. Warming may reduce soil moisture by increasing the evapotranspiration (Dermody et al., 2007). No effect of warming on N₂O flux was observed (McHale et al., 1998; Peterjohn et al., 1994), although elevated temperature is known to increase net N mineralization (Rustad et al., 2001).

In the current study, the effects of future climatic and atmospheric conditions on the biosphere-atmosphere exchange of N₂O and CH₄ were investigated in a temperate heathland at the CLIMAITE experimental site (www.climaite.dk). CLIMAITE was initiated in 2005 to improve our understanding of how biological processes in natural terrestrial ecosystems may be affected under future environmental conditions involving elevated temperature, elevated concentration of atmospheric CO₂ and prolonged summer drought, simulating *in situ* the climatic scenario as predicted for Denmark in year 2075 (Mikkelsen et al., 2008). Previous studies have examined the effects on greenhouse gas fluxes of warming, elevated CO₂ and summer drought, but to our knowledge field studies combining all three factors in a full-factorial design have not been reported. We formulated four hypotheses for the responses in N₂O and CH₄ fluxes to the climate change parameters investigated in the experiment.

- 1) Prolonged summer drought will stimulate CH₄ uptake and reduce N₂O emissions.
- 2) Elevated CO₂ will reduce CH₄ uptake due to higher soil moisture caused by plant water saving mechanisms. Nitrous oxide emissions will remain unchanged under elevated CO₂ because the stimulating effects of increased soil moisture and availability of labile carbon compounds will be offset by increased plant-microbial competition for N.
- 3) Warming will increase CH₄ uptake because of reduced soil moisture. On an annual basis N₂O emission will be unaffected by warming, but this may include a reduction during spring and summer due to reduced soil moisture and an increased emission in autumn due to higher N turnover rates.
- 4) In the combinations of two or three treatments, the treatment effects will either counteract each other, if in opposite directions, or intensify each other, if in the same direction.

To address these hypotheses, we conducted a full-factorial study including all treatments. In addition, a more intense study involving a subset of five treatments was carried out to focus on treatment effects during the experimental summer drought and the subsequent rewetting period.

2. Materials and methods

2.1. Field site

The study took place at the CLIMAITE experimental site (Mikkelsen et al., 2008) situated at Brandbjerg (55°53' N, 11°58' E) about 50 km NW of Copenhagen, Denmark. The site is a dry, temperate heathland on a hilly nutrient-poor sandy podzol (FAO classification). The mineral soil consists of 20.5% coarse sand, 71.6% fine sand, 5.8% silt and 2.2% clay, and has a pH_{H₂O} of 4–5. Mean pore volume and field capacity of the upper 15 cm of the mineral soil is 42 and 17 vol%, respectively. The content of carbon and nitrogen decline sharply from 6.4% and 0.34% in the upper 2 cm of the mineral soil to 0.39% and 0.02% in the 10–30 cm layer, respectively. Above the mineral soil is an organic top layer of about 5 cm in depth containing approximately 23% carbon and 1.2% nitrogen. The vegetation is dominated by *Calluna vulgaris* (L.), *Deschampsia flexuosa* (L.) and various mosses. Annual mean temperature is 8.0 °C, annual mean precipitation is 613 mm (www.DMI.dk) and the N

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