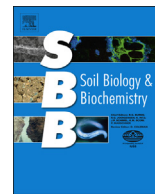




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Fire increases the risk of higher soil N₂O emissions from Mediterranean Macchia ecosystems



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ABSTRACT

Intensification of droughts under climate change is projected to increase fire frequency in the Mediterranean region. Fires cause direct emission of greenhouse gases (GHG) such as carbon dioxide (CO₂) and nitrous oxide (N₂O), due to the combustion of organic matter, creating a positive feedback on climate change. However, the potential importance of indirect GHG emissions due to changes in soil biological and chemical properties after fire is less well known. Increased soil mineral nitrogen (N) concentrations after fire pose a risk for increased emissions of gaseous N, but studies on the post-fire N₂O production and soil N turnover rates (mineralization, nitrification, microbial immobilization, denitrification) are still rare. We determined N₂O production, rates of N turnover and pathways for N₂O production from the soil of burned and unburned plots of a Macchia shrubland in central Spain using a ¹⁵N labelling approach. Measurements were initiated before the controlled burning and continued for up to half a year after fire. Fire markedly increased the risk of N₂O emissions from soil through denitrification (N₂O production rate was 3 to ≈ 30 times higher in burned soils compared to control, with N₂O being produced solely from soil nitrate). In contrast, soil gross N cycling rates were not accelerated after fire. Thus, the increased N₂O production was not closely linked with N mineralization, but may be explained by increased mineral N availability from ash, increased pH in burned plots, and less competition for available N and C sources due to absence of plants.

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1. Introduction

The Mediterranean area is considered to be especially vulnerable to climate change, because it lies at the transition between arid and humid regions of the world (Scarascia-Mugnozza et al., 2000). Climate change projections for the Mediterranean region include warmer temperatures and decreased precipitation, leading to intensification of water scarcity (droughts) (IPCC, 2013). Mediterranean Macchia shrubland ecosystems are naturally affected by fires, but models project increased occurrence of fires for the Mediterranean region with global warming (e.g. Carvalho et al., 2011). This is because reduced water availability in combination with higher temperatures increases fire risk and the length and

severity of the fire season (Moreno et al., 2010). Severe fire events during recent years in Spain, Italy and Greece have been linked to temperature anomalies (Amraoui et al., 2013), and heat waves like this are projected to increase with global warming (IPCC, 2013).

Direct CO₂ production during fires significantly contributes to GHG emissions in Mediterranean countries, and thus further accelerates climate change, if the area burned increases due to global warming (Miranda et al., 1994). Burning biomass is also a source of other GHGs such as N₂O and other air pollutants (Crutzen et al., 1979; Laursen et al., 1992). However, the post-fire effects on N₂O emissions due to mid-to long term changes in soil biological and chemical properties are less well known. Only a few studies from Mediterranean ecosystems have concentrated on changes in N₂O production and N cycling after fire (Dannenmann et al., 2011; Fierro and Castaldi, 2011; Inclán et al., 2012). Therefore, the risk for increased post-fire N₂O emissions (Skiba and Smith, 2000) from these ecosystems, and the magnitude of the potential feedback to climate change remains uncertain. Emissions of N₂O could increase as a consequence of changes in C and N cycling after fire (Brown et al., 2012). Fire disturbance combined with other climate change drivers (elevated CO₂, heat, altered precipitation and enhanced N deposition) has been suggested to cause strongly positive interactions between these factors, leading to higher N₂O fluxes than would be expected based on single-factor studies (Niboyet et al., 2011; Brown et al., 2012). This highlights the importance of studying the post-fire effects on soil N₂O fluxes under a global change context.

Our understanding of how fire may affect soil microbial N cycling in Mediterranean ecosystems is still limited. Fire consumes a large proportion of total aboveground N (up to 80% depending on fire severity, Boby et al., 2010), but it also enhances mineral N availability, as N uptake is drastically diminished, and N remains available in the ash mainly as ammonium (Levine et al., 1988; Marion et al., 1991; Prieto-Fernández et al., 2004). However, the extent to which the soil mineral N concentrations could be affected indirectly by altered inorganic N production (mineralization of organic N to ammonium (NH₄⁺), and nitrification of NH₄⁺ to nitrate (NO₃⁻) and uptake rates by soil micro-organisms (immobilization of NH₄⁺ or NO₃⁻) is largely unknown (Dannenmann et al., 2011). Fire directly produces NH₄⁺, but NO₃⁻ is not formed in the fire, and requires nitrification to form (Knicker, 2007). Soil pH increases after fire, due to the high cation content of the ash (Jensen et al., 2001), and the increase in pH could stimulate nitrification (Ste-Marie and Paré, 1999) and denitrification (Skiba and Smith, 2000).

It is uncertain whether organic N and C availability to soil microbes increases or decreases after fire (Prieto-Fernández et al., 2004), as there are factors affecting into opposite directions. Microbial biomass in the surface soils decreases immediately after fire (Dumontet et al., 1996; Rutigliano et al., 2007; Fontúrbel et al., 2012), and the released organic C and N from the micro-organisms killed by the heat could increase substrate availability for the surviving microbes (Andersson et al., 2004), and thus increase mineralization. Organic N content has been found to increase, decrease or stay the same after fire, but in each case with decreased average lability (Prieto-Fernández et al., 2004). Therefore, it is also possible that N mineralization decreases after fire, as the remaining organic N is mainly stored in recalcitrant hetero-aromatic N structures formed in the fire (Knicker, 2007). Soil micro-organisms are important for determining the fate of the mineral N released in the fire: it could be either retained in the biomass (Bell and Binkley, 1989) or lost from the ecosystem through the microbial nitrification and denitrification (a process converting NO₃⁻ to N₂O and N₂), both of which are known to be stimulated by N additions (e.g. Levine et al., 1988; Barnard et al., 2005). The fate of the N has also consequences for the plant community, as losses of N through

leaching and denitrification could potentially reduce the labile pool of N available for plant growth slowing recovery (Knicker, 2007).

The aim of this study was to measure N₂O emissions, pathways of N₂O production (nitrification and denitrification) and rates of soil N turnover after a fire in a Mediterranean Macchia ecosystem. We hypothesised that N₂O emissions would increase after the fire, due to increased mineral N availability in the soil. We also hypothesised that the effect would decline with time since fire, as recovery of vegetation would increase competition for available N.

2. Materials and methods

2.1. Site description and soil sampling

The site was established on a Macchia shrubland 160 km south-west from Madrid at the Coto Nacional de los Quintos de Mora (39°25'27" N 4°04'17" W), as part of the SECCIA (*Simulation of Effects of Climate Change in a Shrubland Affected by Fire*) project. The field work was initiated in September 2009 and continued until April 2010 to follow the mid-to long-term changes in the N cycling rates. Yearly precipitation at the site is 622 mm and mean annual temperature (MAT) is 14.9 °C (1948–2006 “Los Cortijos” meteorological station; 39°19' N, 4°04' W; AEMET, Ministerio de Medio Ambiente, Medio Rural y Marino, Spain), but in 2009 the precipitation was 427 mm and thus lower than the long-term average (Parra et al., 2012). Warmest month of the campaign period (September 2009–April 2010), was September (19 °C), October and November were the driest months, while most rain (60% of the period) occurred between December and February (Fig. 1). Altitude of the site is 900–925 m and slope is 15–25%. The soil is a Dystric Cambisol, formed on Quartzite bedrock. Average soil properties at 0–5 cm depth are given in Table 1. The dominating plant species are *Cistus ladanifer* L., *Erica arborea* L., *Phillyrea angustifolia* L., *Erica scoparia* L. and *Rosmarinus officinalis* L. A more detailed site description has been published by Parra et al. (2012). Other aspects of plant and ecosystem responses to drought and fire have already been investigated on the site (Hijonosa et al., 2012; Parra et al., 2012; Ramírez et al., 2012).

For the purpose of the present study, six plots of 25 m² were randomly assigned either into burned or control block (n = 3, see Fig. 2). The plots were burned on September 23rd 2009. The fire intensity was high (mean maximum temperature of 710.0 ± 25.4 °C), and similar among all burned plots (Parra et al., 2012). Soil samples (0–5 cm top-most soil, excluding litter layer) were collected from a regular (50 × 50 cm²) grid (80–100 subsamples per plot) with a small soil corer, pooled and immediately sieved (5 mm). Soils were first sampled 1 day before the burning, and sampling was repeated 3, 13, 35 and 204 days after the controlled burning. Soils were transported to a nearby field-laboratory within the day of sampling on –1, 3 and 13 days after the fire, and gravimetric water contents were determined by oven drying at 105 °C to constant weight. Incubations started the following day. Soils collected 35 and 204 days after the fire were placed in insulated cool-boxes, and shipped to a laboratory in Denmark for processing within less than one week.

2.2. ¹⁵N labelling and incubation experiment

Soil samples (20 g) were incubated in 2 l glass jars for 48 h, after labelling with ammonium nitrate (NH₄NO₃) (either double labelled ¹⁵NH₄¹⁵NO₃, or single labelled ¹⁵NH₄NO₃, 5 atom% excess, 10 µg N/g moist soil). Soils were adjusted to the same water content as on the 1st incubation time before burning (35% gravimetric water content after adding the labelling solutions) to keep the incubations comparable. All incubations were conducted at a constant room

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