

## Organic phosphorus mineralisation in a temperate grassland soil under elevated atmospheric carbon dioxide concentrations

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**Background:** Phosphorus (P) is an essential nutrient for all biota and significant proportions of P in soil are present in organic form. Increased atmospheric concentrations of carbon dioxide ( $[\text{CO}_2]$ ) have been shown to influence plant P uptake traits, resulting in plant-mediated changes in soil P pools. However, little is known on the effect of elevated  $[\text{CO}_2]$  on organic P mineralisation rates in soil.

**Study design & hypotheses:** A  $^{33}\text{P}$  isotopic dilution experiment was performed with soils of the 17-year-old Giessen free air carbon dioxide enrichment (GiFACE) – trial. At the GiFACE, three plots are treated with 20 % elevated  $[\text{CO}_2]$  while three control plots receive ambient air. We hypothesised that i) the observed positive effect of elevated  $[\text{CO}_2]$  on plant growth translates into differences in soil organic P mineralisation rates between treated and untreated plots, resulting in ii) differences in soil organic P pools.

**Methods:** Fresh soil (0-8 cm) was sampled from each plot, labelled with a carrier free  $^{33}\text{P}$  solution and incubated for 36 days at 19°C in the dark. On six time points, inorganic P and  $^{33}\text{P}$  in soil filtrates, soil microorganisms (by liquid fumigation) and resin extractable P were quantified. The baseline of  $^{33}\text{P}$  isotopic dilution was assessed from a short term batch experiment and extrapolated for 36 days. Gross organic P mineralisation rates were determined as the difference between isotopic dilution in the incubated soils (physicochemical + biological processes) minus extrapolated values (physicochemical processes only). Additionally, enzyme addition assays on alkaline soil extracts were performed to quantify different soil organic P classes, using enzymes with a known substrate specificity.

**Results & Discussion:** Gross organic P mineralisation rates were high during the first three days ( $5.5 - 34.3 \text{ mg P kg}^{-1} \text{ d}^{-1}$ ), possibly due to the soil disturbance at labelling soils with  $^{33}\text{P}$ . However, gross organic P mineralisation decreased rapidly to rates between 0.7 and  $4.2 \text{ mg P kg}^{-1} \text{ d}^{-1}$ . Net organic P mineralisation during this phase was quantified by subtracting microbial P immobilisation and ranged between 0.3 and  $3.4 \text{ mg P kg}^{-1} \text{ d}^{-1}$ . We did not detect a significant effect of elevated  $[\text{CO}_2]$  on soil organic P transformation rates, however, relatively high P fertilisation since the start of the GiFACE-trial (equal to  $5.6 \text{ g P m}^{-2}$ ) may have masked subtle changes in mineralisation rates. Significantly higher concentrations of phytate-like P were found in plots treated with elevated  $[\text{CO}_2]$ . We assume that increased plant biomass production under elevated  $[\text{CO}_2]$  caused the formation of more inositol-hexakisphosphate (phytate), a P storage compound in plants.

**Conclusions & Outlook:** Elevated  $[\text{CO}_2]$  had no clear effect on soil organic P mineralisation rates, but soil organic P classes were clearly affected. Organic P transformation rates will also be verified with a recently developed numerical model.