# Experimental soil warming in a mountain forest ecosystem



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### Background

Global warming will rise forest soil temperatures and thereby accelerate enzyme kinetic driven processes like organic matter decomposition. Already fixed carbon in the humus may become vulnerable to decomposition and leave the soil as CO. This temperature induced increase in soil respiration could, in worst case, have the potential to shift temperate forest ecosystems from a net sink to a net source of

atmospheric CO<sub>o</sub>. The temporal and quantitative degree of the temperature induced feedback on global warming is still under debate.

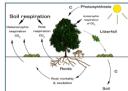


Fig. 1: Carbon cycle

# Methods | Summer

We buried a heating cable into three plots and kept the soil temperature 3°C above the soil temperature of three untreated control plots. Furthermore we created three disturbed control plots where a dummy cable was buried. This enabled us to determine eventual disturbance effects by the cable installation. To separate the root respiration from the heterotrophic respiration we established six trenched plots. All tree roots were cut around the trenched plots one year before starting the measurements. Three of the trenched plots were heated (+3°C). CO\_ fluxes were measured biweekly using a transportable infrared gas analyzer. For high temporal resolution, one plot (5 treatments, 15 chambers) was measured 24 hours, fully automatically throughout the snow free period. Soil temperature and moisture from two soil depths were recorded in half hourly resolution. The automated plot was additionally equipped with ten lysimeter probes.







Fig. 3: (a) Automated plot with automatic chambers, chambers for manual measurements (grange) and the tech box (white) equipped with the infrared gas analyser, the control unit and the data loggers; (b) The heating unit. It contains of 3 transformers and a 50 kA contactor, which is operated by a specially programmed Campbell data logger; (c) Open slot during the trenching procedure

#### Methods | Winter

We measured CO<sub>2</sub> emissions directly from the snow surface using a home made closed dynamic chamber. Additionally, we installed nine Teflon tubes with a small air collector made of a stainless steel mesh at the soil-air interface. The tubes were fixed to tree trunks, so that the other end was accessible even during thick snow cover. CO fluxes were calculated from the concentration gradient (CO at the soil surface; CO at the snow surface) and the snow porosity following Massman et al., 1995. Furthermore we measured concentration gradients through the snow profile using a snow probe in 20 cm depth intervals. This allowed us a better calculation of the flux through snow packs with distinctive layering.

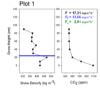






Fig. 5: From left to right: Snow profile with distinct layering chamber for CO. flux measure-Aluminum snow probe for CO. concentration profile measure

- The soil was never frozen under the snow and the CO\_ flux through the snow ranged between 17,5 and 41,0 mg CO\_-C m<sup>-2</sup> h<sup>-1</sup> until the end of March.
- Flux estimates from the chamber measurements on the snow surface were 3-5 times less than flux estimates calculated with the diffusion approach.
- Distinct ice layers can impact the CO flux through snowpacks (Fig. 6).



Results I Winter

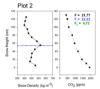


Fig 6: Snow density and CO<sub>2</sub> concentration of two different snow profiles. At plot 1. a distinct ice laver (blue line) acted as barrier for CO. diffusion. The CO. flux calculated from the CO. concentration difference between soil surface and snow surface (F) overestimated the flux compared to the flux calculated from, ice layer surface -> snow surface (Fi). At plot 2, the ice layer was less pronounced and did not act as barrier. Fc is the CO, flux calculated from the linear concentration increase in the chamber on the snow surface.

# Fig. 2: Mean CO, fluxes from the forest soil. The red line shows the date when the heating system was switched on The heated plots were overheated (+8°C) in late August, leading to disproportionately high CO<sub>2</sub> fluxes. In September, the temperature difference was 3°C again. The heating was switched off 10 days past the fist snowfall (dashed red

line). The CO<sub>a</sub> fluxes through snow from December 2005 to March 2006 were calculated with a diffusion approach

# Results | Summer

- Soil respiration from the heated plots was approximately 30% higher than soil respiration from the control plots (Fig. 2).
- Root respiration amounted to 30 40% of the total soil respiration and was generally higher on warming plots.

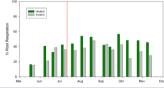


Fig. 4: Percent root respiration as calculated by subtracting the CO. fluxes of the heated and control heated- and control- trenched plots, respectively. The red line shows the start of the heating

### Conclusions

The soil respiration strongly responded (+30%) to a soil temperature increase of 3°C throughout the first observation year. The coming years will show if this response was only an initial flush of easily decomposed organic matter or if higher temperatures will lead to a decomposition of humic substrate, thereby leading to long-term elevated CO<sub>o</sub> emissions from forest

At our site, the winter CO<sub>2</sub> fluxes through snow amounted to 10-15% of the annual soil respiration. This carbon was fixed during summer and was lost to the atmosphere during winter. Therefore we strongly recommend measuring soil respiration in winter, especially if one is calculating annual soil respiration, or carbon budgets, respectively.

### References

Massman W.J., Sommerfeld R.A., Zeller K., Hehn T., Hudnell L, and Rochelle S.G. 1995, CO2 flux through a Wyoming snowpack; diffusional and pressure pumping effects. Biogeochemistry of snowcovered catchments. International Association of Hydrological Sciences, Wallingfort, pp. 71-79.









