



## Isotopic signatures of H<sub>2</sub> and CO uptake and emissions by soil

Maria Elena Popa (1), Qianjie Chen (2), and Thomas Röckmann (1)

(1) Institute for Marine and Atmospheric research Utrecht, Utrecht University, Utrecht, the Netherlands (epopa2@yahoo.com), (2) Department of Atmospheric Sciences, University of Washington, Seattle, Washington, USA

We performed soil chamber experiments in order to determine the isotopic signatures of the exchange of H<sub>2</sub> and CO between soil and atmosphere. The experiments took place at two sites in the Netherlands, a forest (Speuld) and a grass field (Cabauw). Flask samples were filled from the soil chamber, and then analyzed in the laboratory for  $\delta D$  in H<sub>2</sub> and  $\delta^{13}C$  and  $\delta^{18}O$  in CO.

The isotope results prove that, for both species, uptake and emission occur simultaneously regardless of the direction of the net flux. We were able to determine separately the isotopic effects of the two fluxes. For both H<sub>2</sub> and CO, soil uptake is associated with a small positive fractionation (the lighter molecule is taken up faster). The soil uptake fractionation ( $\alpha = k_{heavy}/k_{light}$ ) was  $0.945 \pm 0.004$  for H<sub>2</sub>; for CO, the fractionation was 0.992 for <sup>13</sup>C and 0.985 for <sup>18</sup>O.

The isotopic composition of the H<sub>2</sub> emitted from the grassland was  $-530 \pm 40$  ‰ less depleted than what is expected from the isotopic equilibrium of H<sub>2</sub> with water. For CO, the isotopic composition of the soil emission is depleted in <sup>13</sup>C compared to atmospheric CO, and lower than the average isotopic composition of plant or soil organic matter.