## Iron Dynamics in Basalt-rich Andosols – Implications for Enhanced Rock Weathering

**TOBIAS LINKE**<sup>1,2</sup>, KNUD DIDERIKSEN<sup>3,4</sup>, KAREN MARIA DIETMANN<sup>5,6</sup>, ERIC H. OELKERS<sup>1,7</sup>, JENS HARTMANN<sup>2</sup> AND SIGURDUR R. GISLASON<sup>1,7</sup>

<sup>1</sup>Institute of Earth Sciences, University of Iceland

<sup>6</sup>Grupo de Investigación Reconocido-Química del Estado Sólido, Materiales y Catálisis Heterogénea (GIR-QUESCAT), University of Salamanca

<sup>7</sup>Ali I. Al-Naimi Petroleum Engineering Research Center, KAUST

Various carbon dioxide removal (CDR) techniques are currently under consideration to mitigate global warming and achieve proposed climate targets. Among these, enhanced rock weathering (ERW) aims to drawdown CO<sub>2</sub> by accelerating natural weathering processes utilizing abundant, low-cost materials, such as basalt. The chemical composition of basalt, particularly the role of iron (Fe), is critical for the success of this approach, as the Fe<sup>2+</sup> content in basalt can vary depending on its

This study presents findings from a field investigation of an Icelandic Andosol. The site has received a high dust flux of basaltic material for the past 3,000 years, making this an ideal analogue to study long-term ERW applications. Our results reveal that the rapid oxidation of  $Fe^{2+}$  in the oxic zones of the system leads to the rapid precipitation of iron-(oxy)-hydroxides. In contrast, reduced conditions in deeper soil horizons favor the preservation of  $Fe^{2+}$  in both the soil waters and in  $Fe^{2+}$ -carbonate and -sulfide minerals. Notably, the formation of siderite provides an additional carbon sink, but its potential re-dissolution and oxidation of  $Fe^{2+}$ -under fluctuating soil conditions can lead to the loss of this stored carbon.

The weathering of soil minerals and basaltic material increases substantially the alkalinity of soil waters. In iron-rich systems, a significant portion of this alkalinity is attributed to Fe<sup>2+</sup>. However, transitioning to an oxic environment causes the redissolution of iron minerals, oxidation of Fe<sup>2+</sup>, and the precipitation of Fe<sup>3+</sup>-minerals, resulting in a loss of the alkalinity previously associated with Fe<sup>2+</sup>. Field evidence demonstrates the large impact of this process on the CO<sub>2</sub> drawdown potential of ERW efforts. Simultaneously, the formation of new Fe<sup>3+</sup>-minerals, such as ferrihydrite, provides ample surfaces to immobilize heavy metals. The formation of these Fe<sup>3+</sup>-minerals may also affect nutrient cycling, including phosphate availability.

<sup>&</sup>lt;sup>2</sup>Institute for Geology, University of Hamburg

<sup>&</sup>lt;sup>3</sup>Geological Survey of Denmark & Greenland (GEUS)

<sup>&</sup>lt;sup>4</sup>Nano-Science Center, University of Copenhagen

<sup>&</sup>lt;sup>5</sup>Landesamt für Umwelt, Naturschutz und Geologie Mecklenburg-Vorpommern