

The origin and fate of trace metals during CO₂-charged water injection into basaltic geothermal reservoir, Nesjavellir SW Iceland

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The Nesjavellir geothermal power plant is located within Hengill volcanic system (SW Iceland) where bedrock consists of successions of hyaloclastites and lava sequences formed during glacial and interglacial periods, respectively. The common alteration minerals include smectites, zeolites, quartz, calcite, prehnite, sulfides, feldspars, epidote, chlorite, wairakite and wollastonite (e.g., Franzson, 1998). The hydrothermal fluids in the geothermal system are meteoric in origin and dilute with chloride concentrations <250 ppm. Among the major solutes are CO₂ and H₂S which are emitted during geothermal utilization in quantities of 15 and 8 kt/yr, respectively. Based on carbon and sulfur isotope systematics, the main sources of CO₂ and H₂S in these hydrothermal fluids are magmatic gas input and host rock leaching (Stefánsson et al., 2017). To limit the power plant emissions, a pilot project using Carbfix technology has been conducted in Nesjavellir, where CO₂ and H₂S are separated from the power plant exhaust stream, dissolved in condensed steam, and injected together with separated geothermal water into the injection reservoir (Gunnarsson et al., 2018). Here this mixture reacts with basaltic host rock resulting in mineralization of these gases and therefore providing their safe and permanent underground storage. One of the main concerns related to a subsurface injection of acidic CO₂-H₂S fluids is enhanced mobilization of trace metals which constitute the reservoir rocks. This increased metal mobility has been observed during laboratory experiments mimicking water-CO₂-basalt interaction (e.g., Galeczka et al., 2013). To assess the possibility of toxic metals plume forming during the injection of CO₂-H₂S charged water, we investigate sources and sinks of trace metals in the storage reservoir. We take into account the chemical composition of Nesjavellir boreholes' drill cuttings and the geothermal fluid at depth and on the surface after depressurization. We demonstrate that quantification of metal fluxes in such conditions is crucial for predicting long-term environmental impact of such gas injections especially when considering scaling up of the operations.

References

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