

Low-temperature reactivity of Fe-brucite in serpentinites: Experiments and natural sample characterization

PAULA DÖRFLER¹, BENJAMIN MALVOISIN², BRUNO LANSO³, FABRICE BRUNET⁴, ANGELIKA D. ROSA⁵, ANNE LINE AUZENDE⁶ AND STANISLAV JELAVIC⁷

¹Université Grenoble Alpes

²ISTerre, Univ. Grenoble Alpes

³ISTerre

⁴Univ. Grenoble Alpes

⁵ESRF

⁶ISTerre-UGA

⁷ISTerre, UGA

The low-T alteration (< 150 °C) of serpentinites is of primary importance for natural CO₂ storage and H₂ generation under conditions where life may exist. Reactions happening in this temperature range are difficult to study in the laboratory for kinetic reasons.

While missing in most hand-collected or dredged oceanic samples, (Fe-bearing) brucite is abundant in cores of oceanic serpentinites retrieved from depths > 200 m [1]. This ferroan brucite, (Mg,Fe)(OH)₂, is one of the most reactive phases in changing redox and temperature conditions, owing also to its small grain size often ranging between 20-100 nm. For example, rapid oxidation of ferroan brucite into ferrian (Fe³⁺) brucite has been observed at T < 150°C in the laboratory. An isostructural Fe²⁺ - Fe³⁺ oxidation was inferred, involving charge-balance by deprotonation [2].

In this study, reactivity of synthetic ferroan brucite was investigated through time-resolved oxidation experiments on synthetic and lab air at ambient temperature. Based on XRD and the UV-Vis spectrophotometric analytical technique, the Fe²⁺/Fe_{tot} was analyzed both in initial and reacted brucite. Synthetic ferroan brucite oxidation reached a close-to-maximum reaction progress already after 5 days in lab experiments. Progressive oxidation was also observed in experiments using natural sample powders as starting material.

Additionally, in-situ synchrotron XAS experiments in a diamond-anvil cell showed a decreasing Fe-O bond length during the isostructural oxidation, consistent with the hypothesized charge-balance mechanism.

Further, phase relationships between brucite and serpentine in a natural sample studied by TEM, coupled to EDX chemical analysis, indicate that ferroan brucite often occurs together with lizardite, in reaction zones where Mg-brucite is abundant.

[1] Klein et al. (2020). Brucite formation and dissolution in oceanic serpentinite. *Geochemical Perspectives Letters*, 16, 1-5.

[2] Carlin et al. (2023). Fe(III)-substituted brucite: Hydrothermal synthesis from (Mg_{0.8}Fe_{0.2})²⁺-brucite, crystal chemistry and relevance to the alteration of ultramafic rocks. *Applied Clay Science*, 234, 106845.