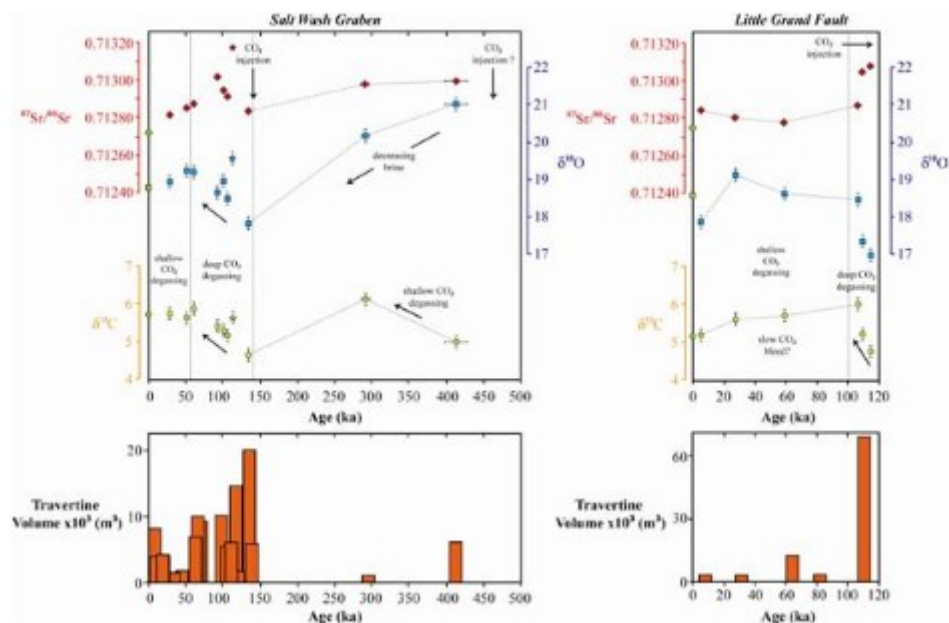


Coupled CO₂-leakage and in situ fluid-mineral reactions in a natural CO₂ reservoir, Green River, Utah

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Surface travertine deposits and carbonate veining within the footwall of Little Grand Fault and Salt Wash Graben, Utah, record a 413 ka history [1] of CO₂ leakage [2]. Isotopic analysis of U-series dated carbonate veins reveals a coupling between CO₂ injection into the host aquifer and the rates of surface leakage, CO₂-promoted silicate mineral hydrolysis reactions within the reservoir [3] and carbonate deposition within fracture conduits through which CO₂-charge fluids ascend to the surface (Fig. 1). Rapid carbonate precipitation rates, recorded in the kinetic fractionation of $\delta^{13}\text{C}$ HCO₃ and $\delta^{18}\text{O}$ HCO₃, reflect an increase in in situ p CO₂ which elevates concentrations of Ca²⁺ and HCO₃⁻, lowering the point at which the ascending fluid reaches carbonate supersaturation due to CO₂ degassing. The spatial and temporal relationship of travertine deposition to CO₂ injection, suggests that rapid rates of carbonate deposition initially plug easily exploited leakage pathways causing leakage sites to propagate laterally, but that this blocking rate decreases with dissipation of the CO₂ charge. This has important implications for the prediction of leakage behaviour in storage sites and for modeling the coupling of subsurface geochemical processes to the evolution of surface leakage.



[1] Burnside et al. (2009) in *Faults & Top seals*, EAGE. Montpellier, France.

[2] Dockrill et al. (2010) *J. Struc. Geol.* in press

[3] Kampman et al. (2009) *Earth Planet. Sci. Lett.* 284, 473–488.

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