

Natural carbonate precipitation rates from bomb-pulse radiocarbon dating

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Precipitation of carbonate minerals is one of the most stable methods to remove carbon dioxide (CO₂) from the atmosphere and sequester it within a solid crystal structure, but the process is slow relative to rates of global CO₂ production. Mineralization of atmospheric CO₂ occurs naturally during alteration of Mg-rich minerals (e.g., mining waste piles, fractured ultramafic systems). Previous studies have confirmed that actively precipitating systems incorporate modern atmospheric CO₂ by measuring F¹⁴C > 1. We report here on our efforts to develop a ¹⁴C bomb-curve dating method for comparing mineralization rates of secondary carbonate minerals from both natural and anthropogenically-influenced systems.

Carbon mineralization occurs through reaction of aqueous cations (Mg²⁺, Ca²⁺) and dissolved bicarbonate (HCO₃⁻). If atmospheric CO₂ is the sole carbon source, radiocarbon dating utilizing the “bomb curve,” or rise in atmospheric ¹⁴C content due to weapons testing, should provide the year (± 1 year) of carbonate precipitation. We first validate this hypothesis by analyzing fresh (“zero-age”) carbonates collected from both an ultramafic-hosted landslide in northwestern Washington (USA) and an alkaline spring in coastal California (USA). Method accuracy is tested on carbonate samples with known ages of precipitation at three time-points along the ¹⁴C bomb curve, and is then applied to mine tailings in Australia and Canada. Future application of the method along transects of a single sample may provide information on the natural rate of precipitation.

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