## Net carbon dioxide removal via electrolytic seawater mineralization

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The trapping of carbon dioxide (CO<sub>2</sub>) as an aqueous (bi)carbonate ion (e.g.,  $HCO_3^{-}$ ,  $CO_3^{2-}$ ) or as a mineral solid is attractive because of favorable thermodynamics, and the safety and permanence of storage. We have proposed an approach to rapidly precipitate Ca- and Mg- carbonates and hydroxides from seawater to achieve CO<sub>2</sub> removal. This Equatic<sup>TM</sup> process electrolytically forces mineral carbonate precipitation thereby consuming prevalent CO<sub>2</sub> that is dissolved in seawater by locking it within carbonate minerals, and simultaneously producing alkaline mineral hydroxides that when dissolved in seawater enable the drawdown of atmospheric CO<sub>2</sub> into the seawater, representing net CO2 removal. Specifically, the net reaction for CaCO<sub>3</sub> (aragonite) precipitation is  $Ca^{2+} + CO_2 +$  $2OH^{-} \rightarrow CaCO_3 + H_2O$ . The precipitation of magnesium carbonate is kinetically limited, thus, alkalinity forces the precipitation of Mg<sup>2+</sup> as Mg(OH)<sub>2</sub> (brucite): Mg<sup>2+</sup> + 2OH<sup>-</sup>  $\rightarrow$ Mg(OH)<sub>2</sub>. Therefore, 1 mol of CO<sub>2</sub> is captured by 2 mol of OH<sup>-</sup> through the production of 1 mol CaCO<sub>3</sub>. On the other hand, only 1.2 mol of OH<sup>-</sup> are required per 1 mol CO<sub>2</sub> stored as dissolved  $HCO_3^{-}$  and  $CO_3^{2-}$  ions. This is because when dissolved into seawater, every mol of Mg(OH)2 leads to the absorption of up to ~1.7 mol of CO<sub>2</sub>(actual ratio depends on pH). The formation of magnesium carbonates, e.g., nesquehonite (MgCO<sub>3</sub>·3H<sub>2</sub>O) and hydromagnesite (Mg<sub>5</sub>(CO<sub>3</sub>)<sub>4</sub>(OH)<sub>2</sub>·4H<sub>2</sub>O) can also be achieved by equilibrating alkalinized seawater with air (i.e., ~400 ppm  $CO_2$ ), yielding two limiting cases: (*Case 1*)  $CaCO_3 + Mg(OH)_2$ (i.e., 11% solid + 89% aqueous CO<sub>2</sub>), and (Case 2) CaCO<sub>3</sub> + MgCO<sub>3</sub> production (i.e., 100% solid CO<sub>2</sub> sequestration). Here, we describe the CO<sub>2</sub> mass balances of the process. This analysis offers a quantitative basis for assessing the CO<sub>2</sub> removal potential of the technological approach and developing a robust measurement, reporting, and verification (MRV) strategy. In addition, we describe the pilot project at AltaSea at the Port of Los Angeles in California which will demonstrate an energy intensity of 2.5 MWh per t CO<sub>2</sub> removed, identify the necessary data for carbon accounting and potential design inefficiencies or improvements, and inform design of full-scale operations.