

Expression of *Duneliata* sp. Carbonic Anhydrase and Its Application to CO₂ Capture/Conversion

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Carbonic anhydrases (CAs) has been focused as biological catalysis for CO₂ sequestration process because the enzyme is known to have ability to convert CO₂ to bicarbonate. In this report, we made codon-optimized sequence of a type-CA cloned from *Duneliata tertiolecta*. (Dt-aCAopt) and subsequently characterized its catalyzing properties to apply for CO₂ capture technology. The expression level in *E. coli* BL21 (DE3) was better for codon-optimized Dt-aCAopt than intact sequence of Dt-aCA. The expressed amount of Dt-aCAopt is 27.79 mg/L at 1.0 mM of IPTG induction and 20°C of growth temperature (for the case of intact Dt-aCA, negligible). Dt-aCAopt enzyme shows half-denaturation temperature at 45°C and show high-stability at pH 7.6/10.0. Apparent values of K_m and V_{max} for p-nitrophenylacetate substrate are 0.9095 mM and 3.303×10^{-8} mM min⁻¹. The effects of metal ions and inhibitors were investigated to find out adequate reaction conditions for Dt-aCAopt application. In final, we showed that in the Ca²⁺ solution, Dt-aCAopt enzyme can catalyze well the conversion of CO₂ to CaCO₃, as the calcite form.