

CORENZ

Cofactor Regeneration in Enzymatic Systems

Institute for Applied Biotechnology (IAB)

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Project description

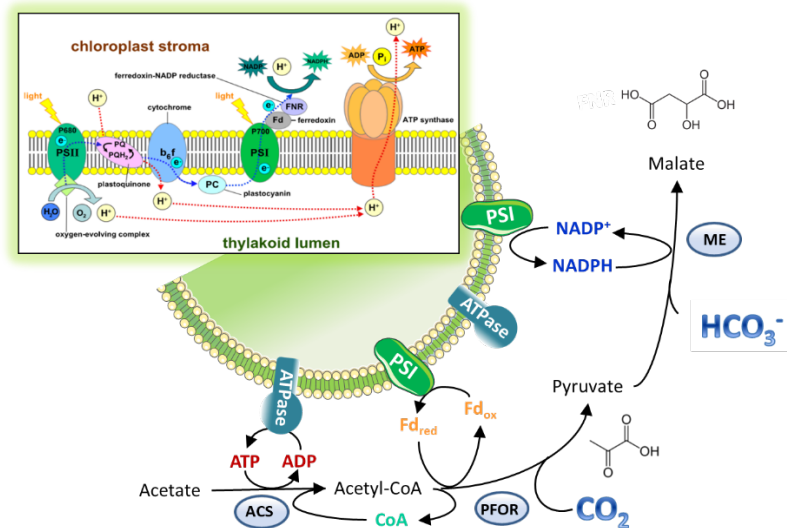
The problem of rising CO₂ levels in the atmosphere as a major cause of global warming, makes it highly attractive to utilize CO₂ as raw material for producing chemicals. Biocatalytic conversion using CO₂-fixing enzymes is particularly interesting since enzymes generally operate with high specificity and selectivity at moderate environmental conditions. We recently demonstrated that natural decarboxylases can be operated in the reverse CO₂-fixing direction in synthetic combinations with low potential redox cofactors *in vitro* (1). We now incorporated further enzymes to establish a synthetic linear pathway for conversion of acetate and 2CO₂ to the C₄-dicarboxylic acid malate. However, enzymatic reactions often depend on cofactors which have to be continuously replenished and lead to high costs in running biocatalytic systems. Furthermore, the accumulation of the used cofactors may cause inhibitory effects and reduce overall efficacy. This project addresses the issue of enzymatic conversion of CO₂ including cofactor recycling within a sustainable closed circular process. As cofactor

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recycling module illuminated chloroplast are incorporated to regenerate the cofactors ATP, NADPH, ferredoxin, and coenzyme A. The pathway potentially provides a biocatalytic route to C4 dicarboxylic acid platform chemicals with CO₂ as a substrate.

(1) Witt A, Pozzi R, Diesch S, Hädicke O, Grammel H. 2019. New light on ancient enzymes – in vitro CO₂ fixation by pyruvate synthase of *Desulfovibrio africanus* and *Sulfolobus acidocaldarius*. FEBS J. 286(22):4494-4508. doi: 10.1111/febs.14981.



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