

Structural basis of a novel PMF-coupled carbon-concentrating mechanism

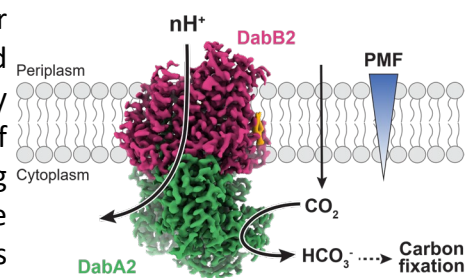
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Microbial carbon dioxide (CO₂) fixation is essential for global primary production and forms the basis of food chains in a variety of ecosystems¹. However, in many natural habitats the process is limited by the scarcity of dissolved CO₂ and the inefficiency of the key CO₂-fixing enzyme

ribulose-1,5-bisphosphate carboxylase/oxygenase (RuBisCO)². To overcome this challenge, many autotrophs evolved specialized adaptations, known as CO₂-concentrating mechanisms (CCMs)^{3,4}.

While cyanobacterial CCMs are relatively well characterized, analogous systems in chemolithoautotrophs, specifically active CO₂ uptake systems have long been overlooked. Here, we present the first cryo-EM structural analysis of the DAB2 complex, an essential membrane-bound CCM for CO₂ uptake in the autotrophic sulfur-oxidizing bacterium *Halothiobacillus neapolitanus*⁵. Our structure reveals a unique β-carbonic anhydrase-like cytoplasmic catalytic subunit (DabA2) and a transmembrane subunit (DabB2) that resembles the proton-conducting subunits of respiratory Complex I. Together with structure-guided mutagenesis and biochemical characterizations, we identified key structural elements possibly couple proton translocation to CO₂ hydration. Our findings establish DAB2 as a prototype of a previously unrecognized family of protonmotive force-coupled vectorial carbonic anhydrases, elucidating a novel strategy for CO₂ capture in non-photosynthetic autotrophs and expanding the mechanistic landscape of bacterial CCMs.



Cryo-EM structure of DAB2 complex

References

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