

Extracellular pathways and fluxes in bioelectrochemical CO₂ reduction to organics

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Description

Global climate change due to anthropogenic fossil fuel consumption, resource scarcity and dependency on foreign influences are primary triggers to transition to a biobased and circular economy. This means that not only energy has to be produced from renewable sources but also fuels, chemicals, feed, food and pharmaceuticals. Electrochemical technologies such as water splitting for hydrogen production and CO₂ reduction are being researched to answer this challenge.

Microbial electrosynthesis (MES) is an emerging novel technology in which microorganisms that are able to use (renewable) electricity, via an electrode (cathode), act as biocatalyst to convert CO₂ to valuable chemicals (see figure). This technology potentially has many advantages such as no net CO₂ emissions (when renewable electricity is used), low water footprint, large product variety, high electron recovery and long life time of the catalyst (years). Most research focuses on producing acetic acid from CO₂, but recently CO₂ elongation to longer chained carboxylic acids (butanoic and hexanoic acid) has been demonstrated.

Since MES is an emerging technology, many fundamental mechanisms (such as electron and carbon fluxes, see figure 2) are still unknown. Unravelling these mechanisms will allow to overcome process limitations on microscale as shown in figure 3.

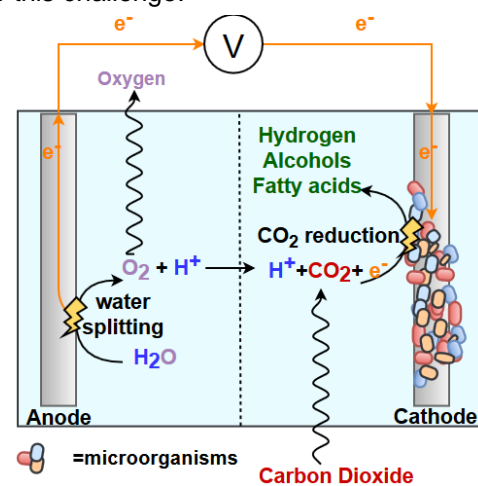


Figure 1: schematic overview of MES

In my PhD research I aim to give insights on what and how (bio-, electro-, and/or bioelectro)-catalytic processes occur and what limits performance of CO₂ reduction to organics by MES at the μm to cm -scale (i.e. electrode-biocatalyst scale). Both experimental and mathematical approaches will be exploited to provide answers to the research questions.

electron flux & transfer mechanism
carbon flux & catabolic processes

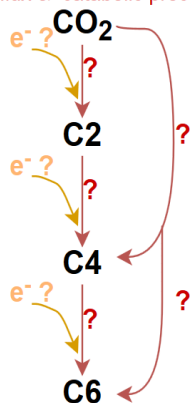


Figure 2: unknown fluxes processes in bioelectrochemical CO₂ reduction to longer chained organics

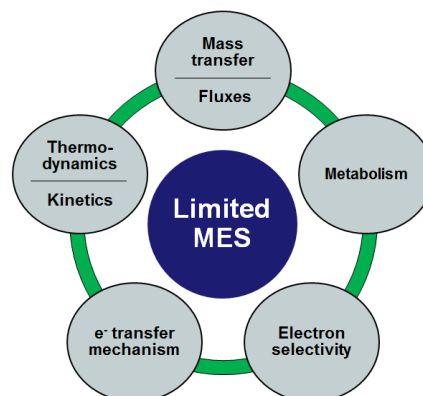


Figure 3: potential limitations in MES on microscale