## **Tunable biopolymer production from renewable methanol**

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Methylotrophic bacteria are microorganisms able to use reduced C1 compounds such as methanol as carbon and energy source and convert them into various valuable multicarbon chemicals and materials. As the production capacity of renewable methanol is increasing worldwide, hybrid concepts of abiotic methanol production and biotic methanol upgrading are gaining interest because of high overall energy efficiencies. This is particularly relevant for the production of complex molecules which can only be obtained through microbial routes, such as biopolymers.

We will discuss biopolymer production by the microorganism *Methylorubrum extorquens*. We developed and intensified a fermentative process for the production of microbial proteins and polyhydroxyalkanoates (PHA) and report for the first time on polyhydroxybutyrate/ valerate production on methanol-pentanol mixtures using an online methanol sensor to control methanol addition at bioreactor scale. Biomass production and biopolymer production phases are optimized. The impact of increasing pentanol levels on biopolymer production and properties is studied and the PHB(V) polymers benchmarked against literature data on methylotrophic bacteria.

The interest of producing microbial proteins (in combination with PHA) goes well beyond their application in feed (and food). While it is known that certain proteins can be used for food packaging, the potential of microbial proteins is largely unexplored. The ongoing PROMIPOL project therefore aims to finetune the combined production of microbial proteins and PHA, study their joint or separate extraction and investigate their properties in view of processing for food packaging or other applications.

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## Technological advances in Bio/Electrochemical carbon dioxide capture & utilization (CCU)

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Recent years have seen a sharp rise in the share of renewable energy production. This renewable power usually often leads to energy surplus. Storage (batteries, water pumping or hydrogen production) have been proposed to exploit this surplus. One of the novel alternatives is to use excess electricity to convert CO<sub>2</sub> into organic chemicals and fuels. Growing rapidly, now comprise approximately 25% of the global electricity capacity is now comprised of renewable energy sources, thus providing great opportunity for CO<sub>2</sub> reduction (CO2R). The International Renewable Energy Agency (IRENA) estimated at the end of 2018 the installed global renewable energy generation across wind, solar, hydroelectric, and other sources to be 2351 GW. CO2R pathways can utilize electricity directly in the conversion step or indirectly via other energy carriers (e.g., H<sub>2</sub>) in the so called Powerto-X approach. Most common conversion pathways include electrochemical, bioelectrochemical, plasma and thermochemical conversion. Non-reductive routes which are commercially mature such as enhanced oil recovery, food and beverage, and concrete curing also exist. While the latter have a higher relative level of maturity, they provide fewer opportunities for electricity utilization compared to reductive routes. The electrochemical CO<sub>2</sub> reduction provides a viable option for reducing anthropogenic CO<sub>2</sub> emissions, while at the same time closing the carbon cycle, by selectively converting CO<sub>2</sub> to fuels/chemicals. While reduction to CO or formic acid only requires two protoncoupled electron transfers, the reduction of CO<sub>2</sub> to ethylene or ethanol consumes 12 electrons. Currently as best possible option as each has been synthesized electrochemically with partial current densities for the indirect route, methanol and methane are at high current TRL (i.e., low technical barriers to, CO, ethylene, and formate are considered >100 mA cm2, often considered as commercially relevant current density, at a faradaic efficiency >60%. formation) and high achievable rates of formation. In this presentation, along with a general overview of CCU, VITO's research on CO<sub>2</sub> conversion using bioelectrochemical and electrochemical approach will be presented.