Engineering Conferences International ECI Digital Archives

Microbial Engineering II

Proceedings

4-3-2022

Lactic acid production in the synthetic autotroph Komagataella phaffii

Michael Baumschabl

Thomas Gaßler

Özge Ata

Diethard Mattanovich

Follow this and additional works at: https://dc.engconfintl.org/microbial_ii

LACTIC ACID PRODUCTION IN THE SYNTHETIC AUTOTROPH KOMAGATAELLA PHAFFII

Michael Baumschabl, Austrian Centre of Industrial Biotechnology (ACIB GmbH); University of Natural Resources and Life Sciences (BOKU), Austria michael.baumschabl@boku.ac.at Thomas Gaßler, University of Natural Resources and Life Sciences (BOKU), Austria; Institute of Microbiology, ETH Zurich, Switzerland Özge Ata, Austrian Centre of Industrial Biotechnology (ACIB); University of Natural Resources and Life Sciences (BOKU), Austria Diethard Mattanovich, Austrian Centre of Industrial Biotechnology (ACIB); University of Natural Resources and Life Sciences (BOKU), Austria

Key Words: Komagataella Phaffii, Lactic acid, CO2 assimilation, Calvin cycle

These guidelines have been prepared in the format that should be used for the abstract submission. Authors The methylotrophic yeast Komagataella phaffii was recently converted to a synthetic autotroph by the integration of the Calvin Benson Bassham cycle. The key step here is the fixation of one molecule CO2 by the enzyme RuBisCO. We could already prove that this strain could grow using CO2 as a carbon source. The next step was to test the ability of this strain to produce organic acids.

We chose lactic acid as a product of choice. It is a hydroxycarboxylic acid used in food, pharmaceutical and chemical industry. Furthermore, it is the precursor of the biodegradable polymer poly-lactic acid (PLA). Lactic acid can easily be produced in Komagataella phaffii by integration of a lactate dehydrogenase gene.

In this work we assessed the lactic acid production in the autotrophic strains using CO2 as carbon source and the LDH gene under the control of the AOX1 promoter. This strain was able to produce up to 150 mg L-1 in approximately 200 hours of cultivation time. Titers were further improved up to 300 mg L-1 by the deletion of the CYB2 gene. Additionally, we compared the lactic acid consumption kinetics of the CYB2 knock-out strain compared to its parental strain.

With this work we were able to show that lactic acid can be produced under autotrophic conditions. We were able to further improve the titers by the knock-out of CYB2 which reduced the ability to consume the produced lactic acid.