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Burmeister et al.,

## **1 Supplementary Videos**

## 2 Title: A microfluidic co-cultivation platform to investigate microbial

## 3 interactions on single-cell level

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## 18 V1:

- 19 Time-lapse video of one microfluidic co-cultivation chamber with growing *C. glutamicum* DM1800 and
- 20 *C. glutamicum* Δ*lysA* pEKEX2-eYFP cells. Cells are supplied with CGXII medium and 10 mM lysine.
- 21 Images were taken in an interval of 10 min.
- 22
- 23 V2:
- 24 Microfluidic control experiments with *C. glutamicum* Δ*lysA* pEKEX2-eYFP. Left frame shows time-lapse
- 25 images of the growing lysine auxotrophic strain supplied with CGXII and 10 mM lysine. Right frame
- 26 shows time-lapse images of the lysine auxotrophic strain in CGXII medium without additional lysine.
- 27 Images were taken in an interval of 10 min.

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- 29 V3:
- 30 Contact-dependent interaction experiment with *E. coli* S17-1 pRhokHi-2-eYFP and *P. putida* KT2440
- 31 pJT'Tmcs-mCherry in microfluidic co-cultivation chambers. Cells are supplied with LB medium.
- 32
- 33 V4:
- Contact-dependent interaction experiment with *E. coli* S17-1 pRhokHi-2-eYFP and *P. putida* KT2440 pJT'Tmcs-mCherry in microfluidic monolayer growth chambers. Direct cell contact is allowed and
- 36 plasmid transfer occurs, which leads to a colour shift in single cells. Cells are supplied with LB medium.
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