Implications of initial physiological conditions for bacterial adaptation to changing environments

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Overcoming metabolic transitions is a key determinant of microbial fitness. Some of the most challenging transitions are encountered when microbes are shifted from rapid, often partly fermentative glycolytic to slower, fully respiratory gluconeogenic growth with lag phases that can last many hours without detectable biomass production. Our groups have recently worked extensively to understand the mystery of how such long lag phases emerge and why they cannot be easily overcome by bacteria.

Our independent studies Kotte et al. (2014) and Basan et al. (2020) appear at first glance somewhat contradictory, because we observed very different phenomenologies, regarding population-level heterogeneity and also the extent of observed lag times. While Kotte et al. (2014) found distinct subpopulations of cells, with some cells that grew immediately and others that never resumed growth, Basan et al. (2020) found a unimodal distribution of single cell lag times and the vast majority of cells eventually resumed growth. The resulting lag times from Kotte et al. (2014) were also much longer than those observed in Basan et al. (2020).

These differences are largely the result of different experimental protocols for performing the metabolic shifts. In Basan et al. (2020), bacteria were washed twice on a filter with warmed post-shift medium and then gently resuspended with a pipette in warmed medium, a procedures accomplished in 2–3 min. Thus, cells were instantaneously exposed to the new substrate and lag phases were only a few hours, closely resembling the apparent phases of no growth during the diauxic shifts in batch experiments (Erickson et al., 2017; Basan et al., 2020). In contrast, the protocol of Kotte et al. (2014) included several rounds of centrifugation and washing with ice-cold, substrate-free medium. The entire procedure took about 20 min and likely exposed the bacteria to starvation and cold stress. Perhaps not surprisingly, the protocol by Kotte et al. (2014) resulted in much longer lag times and a different phenomenology. Hence, as illustrated in Fig 1, cells in the two experiments started their adaptation from very different initial physiological conditions. While the “Basan” cells presumably had to mainly reverse their glycolytic flux, the “Kotte” cells likely faced additional challenges that might include a more severe drop in energy supply and additional adaptations of their proteome during the washing phase.

Independent of the physiological starting point, cells in both experiments had to overcome the problem of depletion of key glycolytic metabolites that result from the sudden flux reversal when switching from glycolytic to gluconeogenic growth. This imposes a trade-off between rapid growth and adaptability that was also observed in Kotte et al. (2014), caused by sequential flux limitation in gluconeogenesis (Basan et al., 2020). Imposing additional stress during the transition phase can drive the population into subpopulation heterogeneity, which can be observed as an increased length of the lag phase (Kotte et al., 2014). Thus, rather than contradicting each other, our studies offer a complementary picture of the physiological adaptation from glycolytic to gluconeogenic growth.

Beyond the specific case of understanding the challenges of the diauxic shift, our results highlight the critical importance of the precise environmental conditions prior to introducing any physiological change or starting a measurement. Seemingly small differences such as placing cells on ice, washing in nutrient-free buffer, or length of the procedure can rapidly alter the physiological state of the culture and/or its population structure. Precise reporting of these seemingly trivial procedures is therefore mandatory and must be taken into account when comparing data from different sources.

References

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This piece discusses how the different observations of two independent studies (Kotte et al., 2014; Basan et al., 2020), regarding population-level heterogeneity and lag times during diauxic shift, can be largely explained by different experimental protocols.

**Subject Categories:** Metabolism; Microbiology, Virology & Host Pathogen Interaction