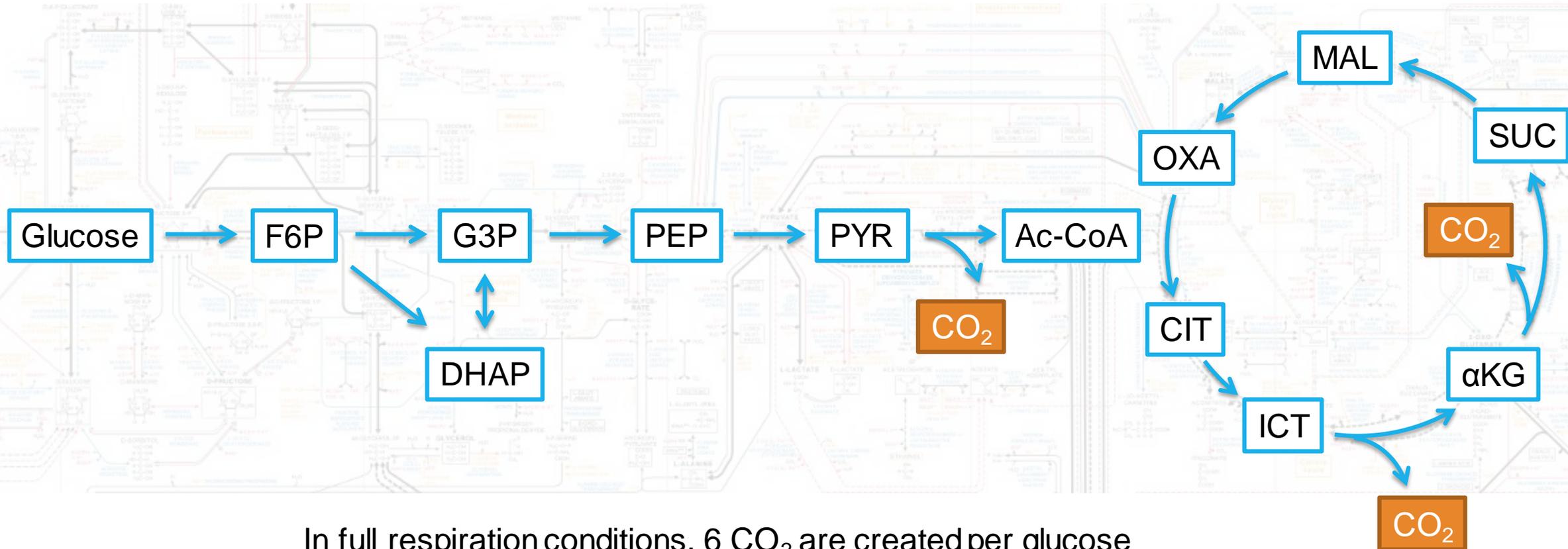




# Carbon-Optimized Bioconversion

Breakout Questions – Day 1

# Glycolysis and the TCA cycle waste CO<sub>2</sub>



In full respiration conditions, 6 CO<sub>2</sub> are created per glucose  
1 per 3-C pyruvate to 2-C acetyl-CoA (x2)  
1 from 6-C isocitrate to 5-C a-ketoglutarate (x2)  
1 from 5-C α-ketoglutarate to 4-C succinate (x2)

# Breakout Session 1 – SynBio/Fermentation

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- ▶ What metabolic pathways could be optimized to avoid CO<sub>2</sub> evolution, or allow for maximal internal recycling of carbon?
- ▶ What does a large scale carbon optimized fermentation process look like?
  - Microbe, Fermentation process, External energy accommodation, Downstream processing
- ▶ What products could be made that are attractive for translation?
  - Aim for high volume, low cost, displacing/replacing petrochemicals

# Breakout Session 1 – Cell-Free

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- ▶ What advantages could cell-free systems offer as a bioconversion platform?
  - What bioproducts are best suited for cell-free systems at scale?
- ▶ What advantages could cell-free systems offer with respect to optimization of carbon efficiency during bioconversion?
- ▶ To what extent can cell-free prototyping be used to develop/test carbon optimization tools for traditional whole-cell fermentation systems?
- ▶ What are the barriers to entry to large-scale cell-free bioprocessing?
  - Enzyme stability, Enzyme production, Co-factor generation/regeneration