

Biosynthesis is a huge cost for bacteria.

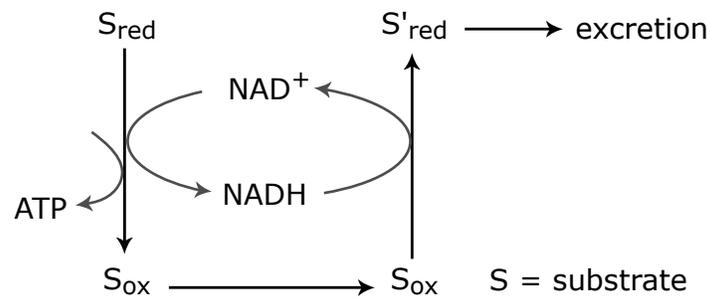
Pathogens are lucky because they use less energy, because their host makes stuff for them. They take up pre-synthesized precursors from host & intermediates.

## 1. Fermentation

No e<sup>-</sup> transport chain

→ Problem: how to reoxidize NADH?

↳ Do internal redox reactions.



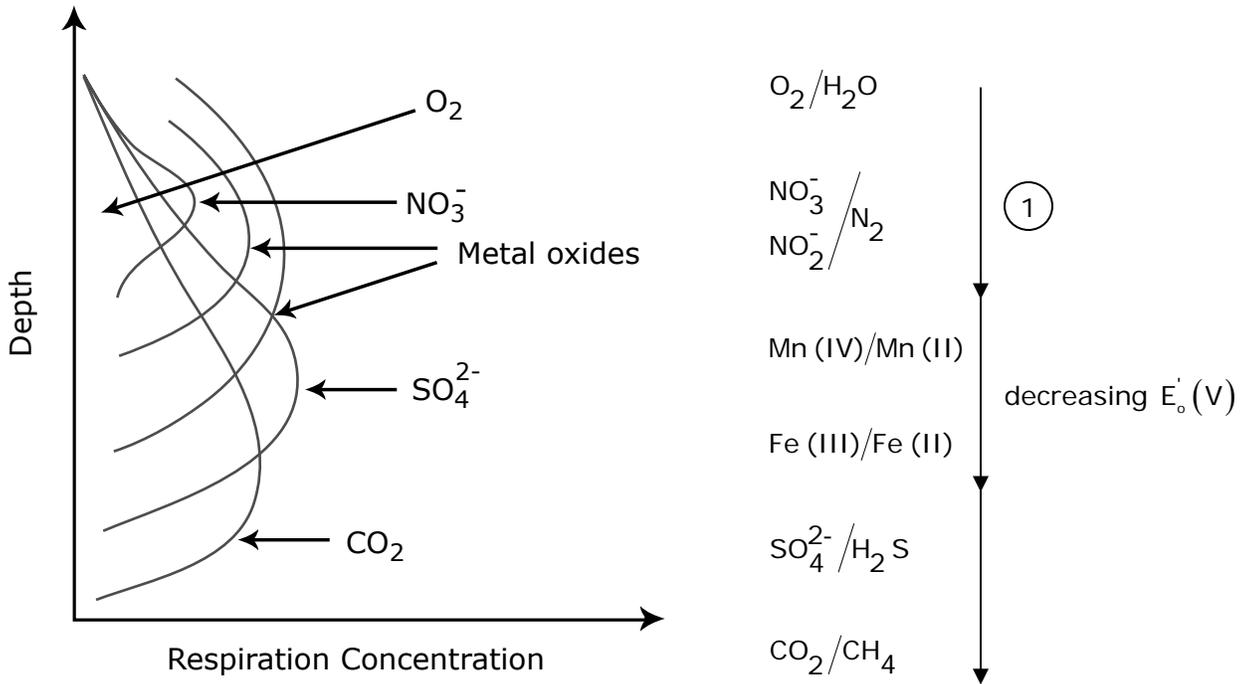
↖ 2-3 ATP/C

### a) Lactic acid fermentation:

- ATP is generated by substrate level phosphorylation
- Strict redox balance: average oxidation state of products is same as substrates  
→ only substrates with intermediate oxidation states can be fermented
- Most involve pyruvate as an intermediate
- Only under strict anaerobic conditions

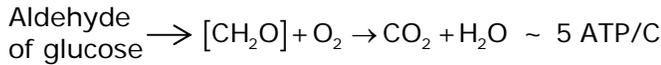
## 2. Respiration

- NADH reoxidation via e<sup>-</sup> transport chain  
↳ ultimate reduction of external e<sup>-</sup> acceptor
- Some of the major e<sup>-</sup> acceptors ⇒



a) Aerobic Respiration:

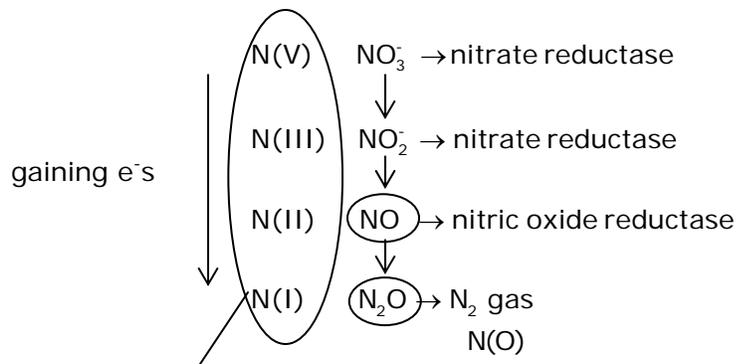
$O_2$ : e- acceptor



- Can generally use a great variety of carbon substrates
- Different species are often specialized in terms of C-substrate key use

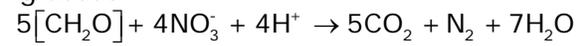
b) Anaerobic Respiration:

- e<sup>-</sup> transport is analogous to aerobes, but use different e<sup>-</sup> carriers with different redox potentials.
- Nitrate – most common form of nitrate respiration = denitrification



What is meant?

"glucose"



Almost as energetically favorable as aerobic respiration (many aerobes can switch to using nitrate for respiration)

- As high as C-substrate diversity of aerobes
- Significance
  - N-removal from systems
  - Production of green house gases –  $\text{N}_2\text{O}$ ,  $\text{NO}$