Chapter 8

Topics in lectures 15 and 16

– Metabolism

- Chemical foundations
- Catabolism
- Biosynthesis

Metabolism

Chemical Foundations

- -Enzymes
- -REDOX
- Catabolism
 - -Pathways
- Anabolism
 - Principles and pathways

Enzymes

Catalysts for chemical reactions

Reactants <----> Products

- Lower the energy needed for the reaction to occur (activation)
 - Endergonic or exergonic
 ===> Insight 8.1

Redox reactions

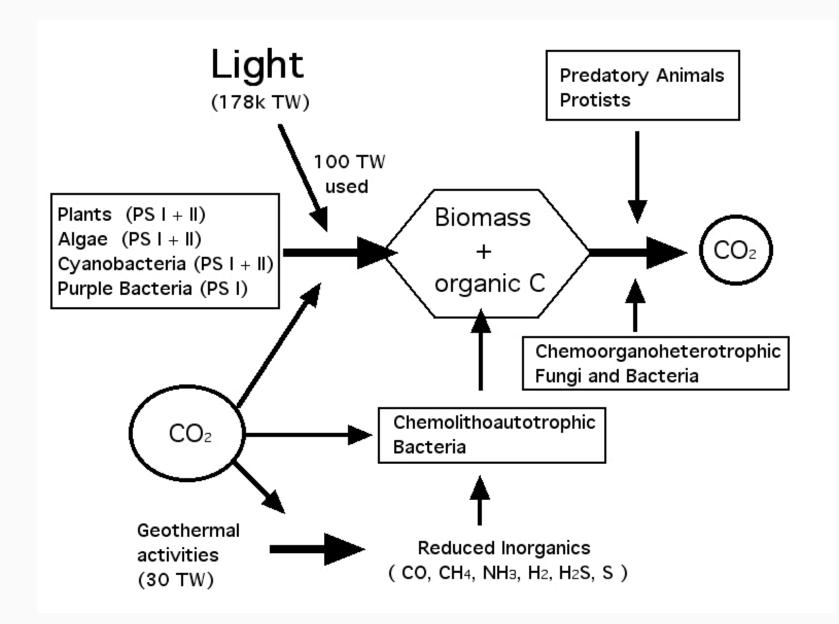
- Reduction and oxidation reaction
- Electron carriers transfer electrons and hydrogens
 - Electron donor
 - Electron acceptor
- Energy is also transferred and captured by the phosphate in form of ATP

REDOXREDOXIDANT + electron <==> REDUCTANT
OXELECTRON ACCEPTOR

OIL --- RIG

Energy

- Cell energetics
 - Exergonic
 - Endergonic
- Redox reaction
- Electron carrier
- Adenosine Triphosphate (ATP)



Integrated model of energy and carbon flow

Metabolism

- Chemical Foundations
 - -Enzymes
 - -REDOX
- Catabolism
 - -Pathways
- Anabolism
 - -Principles and pathways

Catabolism

 Enzymes are involved in the harvest of energy from the environment and their transformation into cell-own, useable energy. Some of this energy needs to be spent in the process on the accession of energy and nutrients (e.g., chemotaxis, transport).

Anabolism

 Enzymes are involved in the use of energy from catabolism in order to synthesize simple and complex compounds, macromolecules and cell structures from simpler compounds).

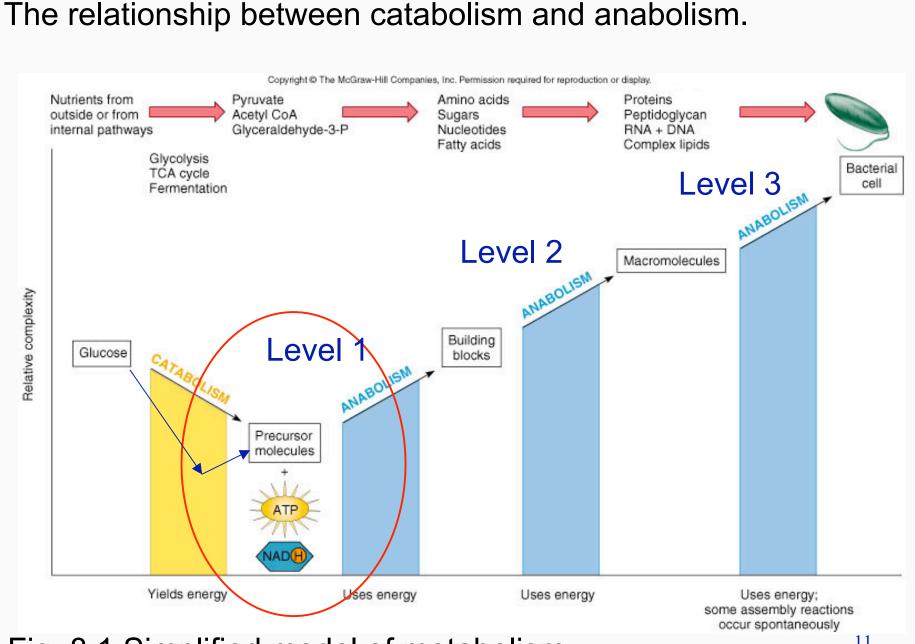


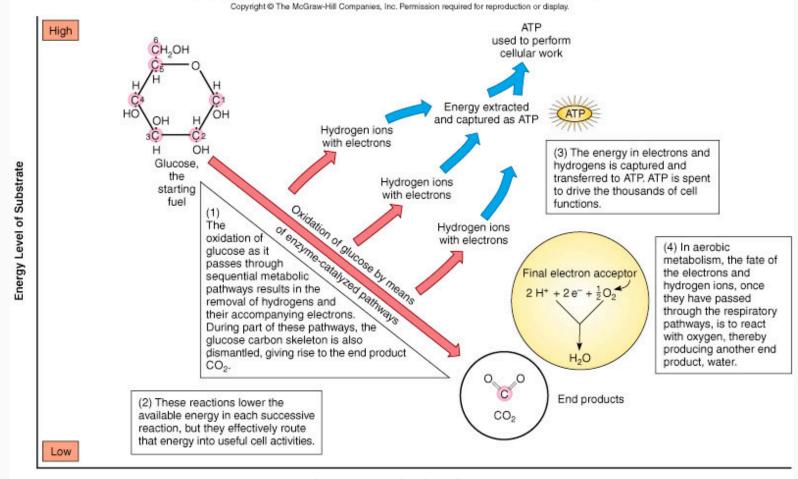
Fig. 8.1 Simplified model of metabolism

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Pathways leading to the 12 precursors

- Central Pathways
 - Mostly catabolism
 - Embden-Meyerhof-Parnas (EMP) pathway [glycolysis]
 - 5 precursors: G6P, 3PG, DHAP, PEP, pyruvate
 - Tricarboxylic acid cycle (TCA)
 - 3 precursors: OAA, alpha-KG, succinyl~CoA
 - Mostly anabolism
 - Pentose phosphate pathway
 - 2 precursors: R5P, E4P
 - Acetyl~CoA,
- Alternate pathways
 - G1P

The general scheme associated with metabolism of organic molecules, the redox reaction, and the capture of energy in the form of ATP.

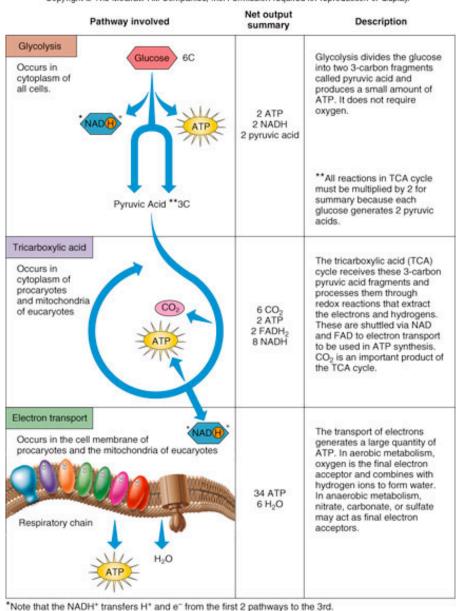


Progress of Energy Extraction Over Time

Fig. 8.12 A simplified model that summarizes the cell's energy machine.13

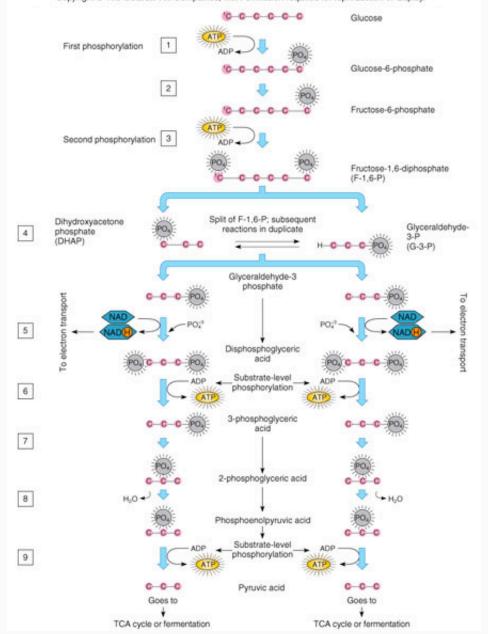
A summary of the metabolism of glucose and the synthesis of energy. Copyright @ The McGraw-Hill Companies, Inc. Permission required for reproduction or display.

Fig. 8.17 Overview of the flow, location, and products of Pathways in aerobic respiration.



The glycolytic steps associated with the metabolism of glucose to pyruvic acid (pyruvate).

Fig. 8.18 Summary of glycolysis



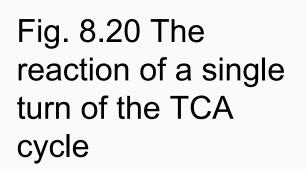
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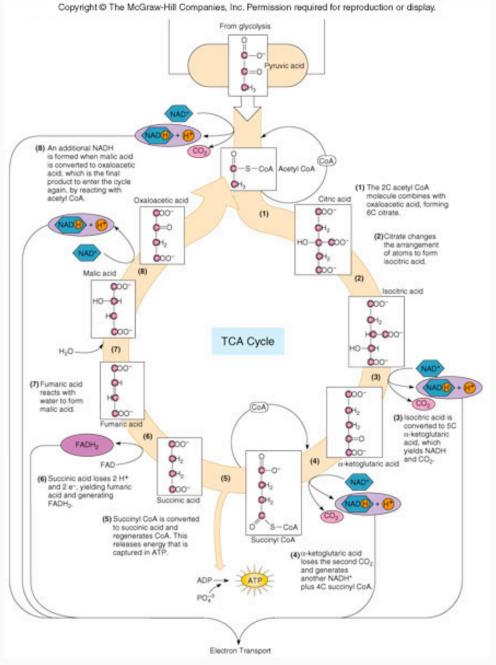
Glycolysis

- Oxidation of glucose to pyruvate
- Splits a 6 carbon sugar into two 3 carbon molecules
- Substrate-level-phosphorylation of some intermediates (Investment of 2 ATPs)
- 2 coenzymes NAD are reduced to 2 NADH
- 2 x substrate-level-phosphorylations (2 x 2 ATPs are synthesized)

Glycolysis continued

- Net yield: 2 ATP & 2 NADH molecules
- Final intermediates are two Pyruvic acid molecules
- Water is generated





The steps associated with TCA cycle.

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TCA cycle

- Each pyruvic acid can be processed to enter the TCA cycle
- CO₂ is generated
- Coenzymes NAD and FAD are reduced to 8 NADH and 2 FADH₂
- Net yield of two ATPs
- Critical intermediates are synthesized

Aerobic respiration

- NADH from Glycolysis
- NADH and FADH₂ from TCA donate electrons into electron transport chain
- Electron transport
- Electron is donated to Oxygen as inorganic, external terminal electron acceptor.

Anaerobic respiration

- NADH from Glycolysis
- NADH and FADH₂ from TCA donate electrons into electron transport chain
- Electron transport
- Electron is donated to nitrate, nitrite, sulfate, sulfite & other oxidized inorganic, external terminal electron acceptors, but not to O₂.

Electron transport

- Membrane bound carriers transfer electrons (redox reactions)
- Proton motive force (PMF)
- Chemiosmosis

Chemiosmosis entails the electron transport and formation of a proton gradient (proton motive force).

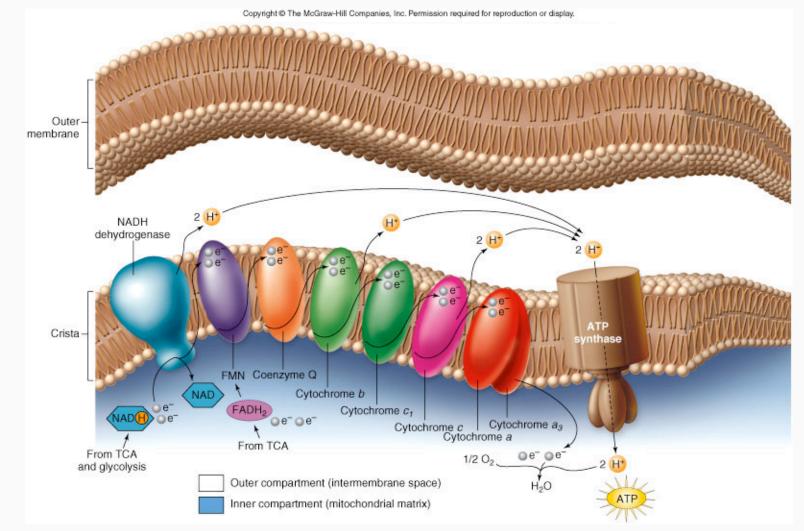


Fig. 8.22 The electron transport system and oxidative phosphorylation

Electron transport chain

- Cytoplasmic membrane
 Prokaryotes
- Mitochondria
 - eukaryotes

Total maximum yield of ATP for one glucose molecule from aerobic respiration.

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TABLE 8.4 Summary of Aerobic Respiration for One Glucose Molecule

	Glycolysis*	Net Output	TCA Cycle*	Net Output	Respiratory Chain	Net Output	Total Net Output per Glucose
ATP produced ATP used	$2 \times 2 = 2$	2	$1 \times 2 = 0$	2	$17 \times 2 = 0$	34	$40 - 2$ (used) = 38^{**}
NADH produced	$1 \times 2 =$	2	$4 \times 2 =$	8	0		10
FADH produced	0		$1 \times 2 =$	2	0		2
CO2 produced	0		$3 \times 2 =$	6	0		6
O2 used	0		0		$3 \times 2 =$	6	
H ₂ O produced	2		0		$3 \times 2 =$	6	8 - 2 (used) = 6
H ₂ O used	0		2		0		

*Products are multiplied by 2 because the first figure represents the amount for only one trip through the pathway, and two molecules make this trip for each glucose. **This amount can vary among microbes.

Table 8.4 Summary of aerobic respiration for one glucose molecule

Fermentation

- NADH from only glycolysis is used to reduce the organic products
- NADH produced by TCA (if present) is not used
- Internal organic compounds serve as the final electron acceptors
- ATP yields are small (per glucose molecule), compared to respiration
- Must metabolize large amounts of glucose to produce equivalent of ATP from respiration

The fermentation of ethyl alcohol and lactic acid.

Copyright @ The McGraw-Hill Companies, Inc. Permission required for reproduction or display. System: Yeasts System: Homolactic bacteria; Human muscle Glucose NAE NAD(Glycolysis н-С —он Pyruvic acid CO н-с -с-н 0 Acetaldehyde NAD NAD н н NAD H-C-C-0 OH Ethyl alcohol Lactic acid

Fig. 8.24 The chemistry of fermentation systems

Types of fermenters

- Facultative anaerobes
 - Fermentation in the absence of oxygen
 - Respiration in the presence of oxygen
 - Ex. Escherichia coli, yeast
- Strict anaerobic fermenters
 - No respiration
 - Ex. Clostridium botulinum, Streptococcus mutans

Products of fermentation

- Alcoholic fermentation
- Acidic fermentation
- Mixed acid fermentation

An example of mixed acid fermentation and the diverse products synthesized.

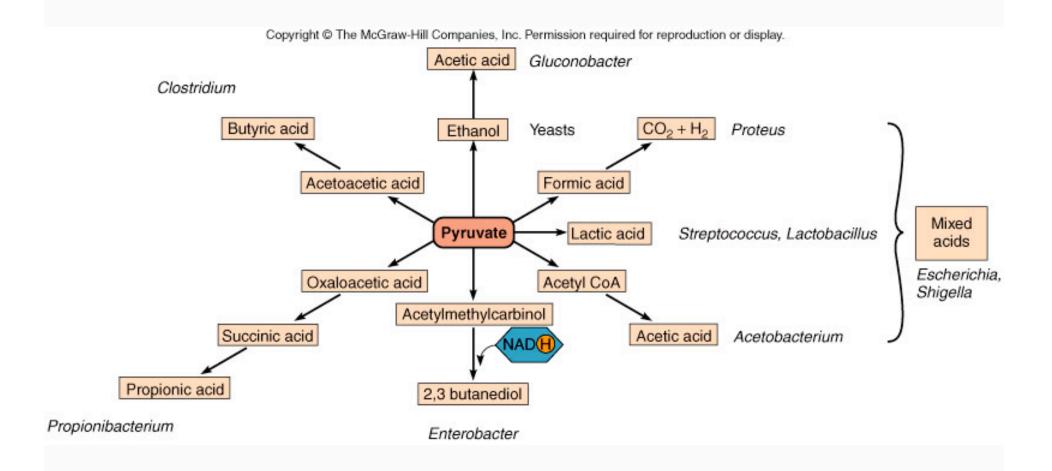


Fig. 8.25 Miscellaneous products of pyruvate

Biosynthesis anabolism

Enzymes are involved in the **use of energy** from catabolism in order to synthesize simple and complex compounds, macromolecules and cell structures from simpler compounds).

The three levels of Anabolism

- Synthesis of Building Blocks, namely complex compounds including the four monomers
- Synthesis of the four Macromolecules
- Assembly of Cellular Structures

Monomer Intermediates

- Cellular building blocks include the four monomers needed for the synthesis of macromolecules:
 - Amino acids
 - Nucleotides
 - Monosaccharides
 - Glycerol and Fatty acids

Macromolecules

- The second level of anabolism is the synthesis of macromolecules:
 - **Proteins** (from amino acids)
 - Nucleic acids (from nucleotides)
 Polysaccharides (from monosaccharides)
 - Lipids (from glycerol and fatty acids)

Cellular Structures

- The third level of anabolism is the assembly of cellular structures from macromolecules:
 - Membranes (from lipids, proteins & polysaccharides)
 - Cell walls (from polysaccharides, proteins & lipids)
 - Ribosome (from nucleic acids & proteins)
 - Flagellum (from proteins)
 - Endospores (from lipids, proteins, nucleic acids & Polysaccharides
 - Inclusion bodies
 - Etc.

Examples for Biosynthesis

- Macromolecule synthesis
 - transcription and translation
- Gluconeogenesis
- Amination
- Transamination

Gluconeogenesis

- When the glucose supply is low, Pyruvate (intermediate) can be converted to glucose.
- Autotrophic organisms, which fix carbon solely form CO₂, funnel the resulting intermediate into reverse glycolysis.

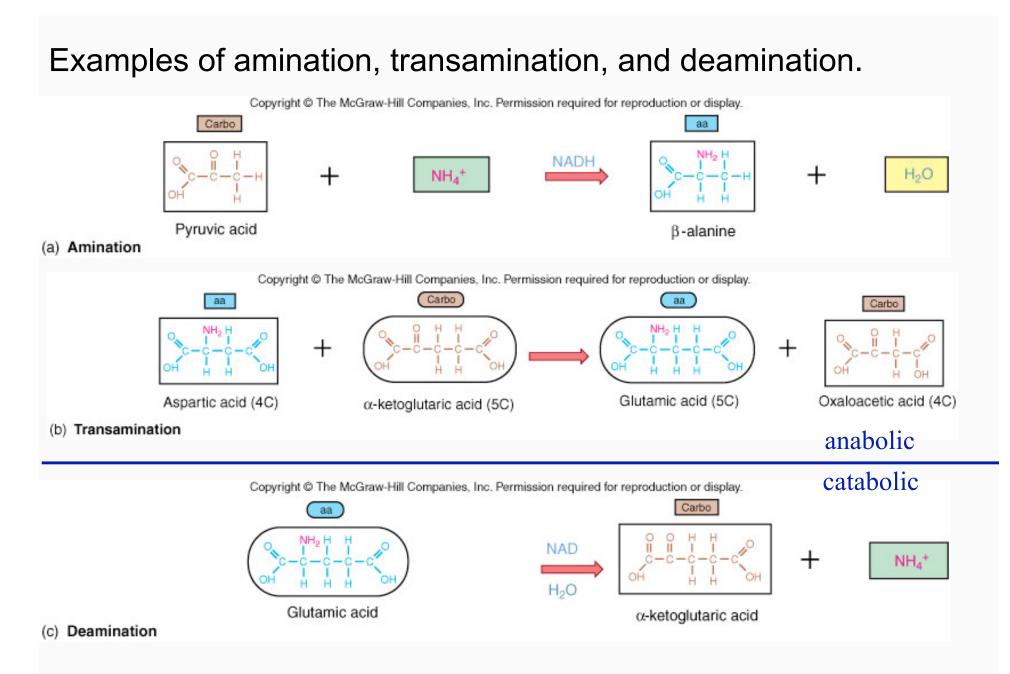


Fig. 8.27 Reactions that produce and convert amino acids ³⁸

Amphibolic

- Integration of the catabolic and anabolic pathways
- Intermediates serve multiple purposes

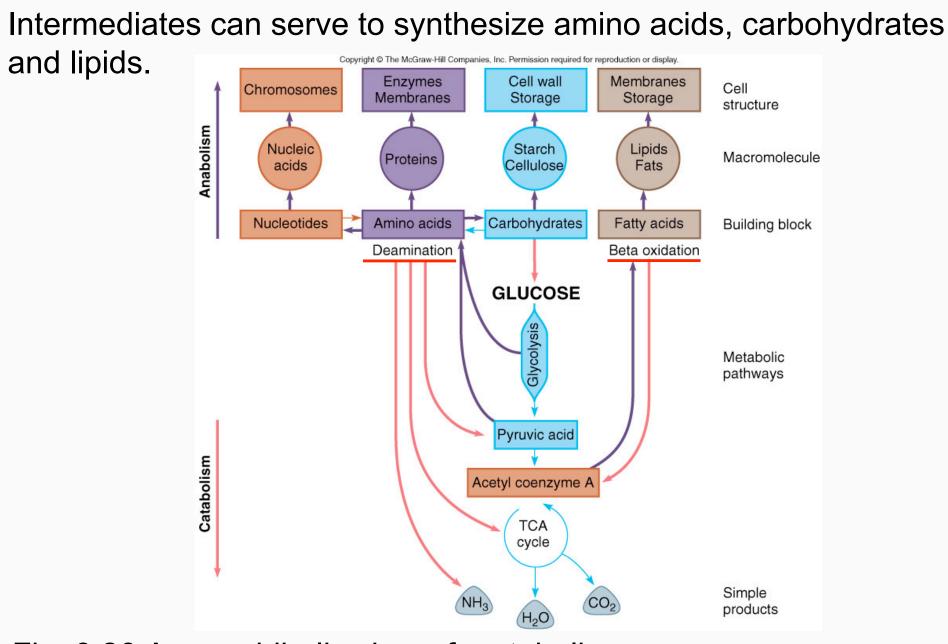


Fig. 8.26 An amphibolic view of metabolism

Beta oxidation

 Alternate catabolism of fatty acids into acetyl, which can then enter the TCA cycle as acetyl CoA.