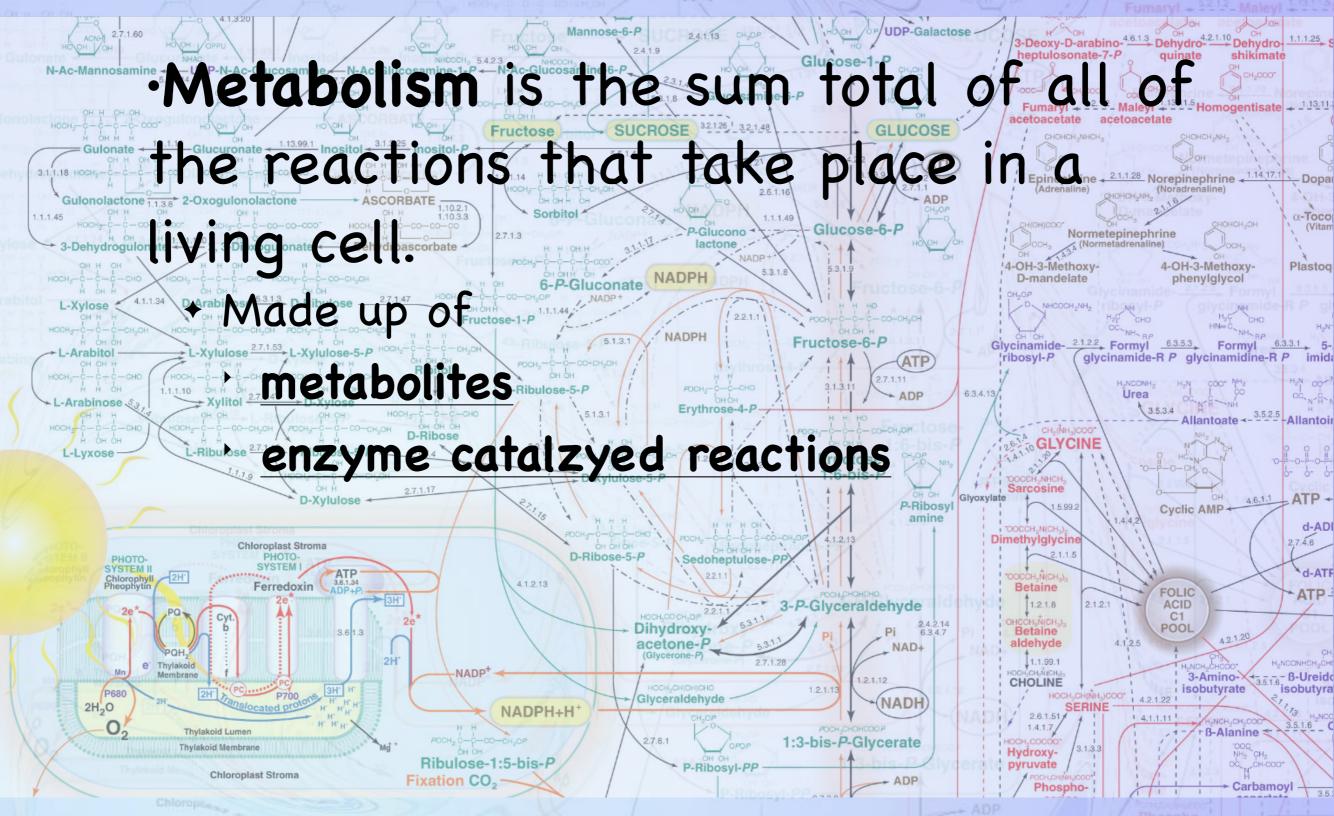
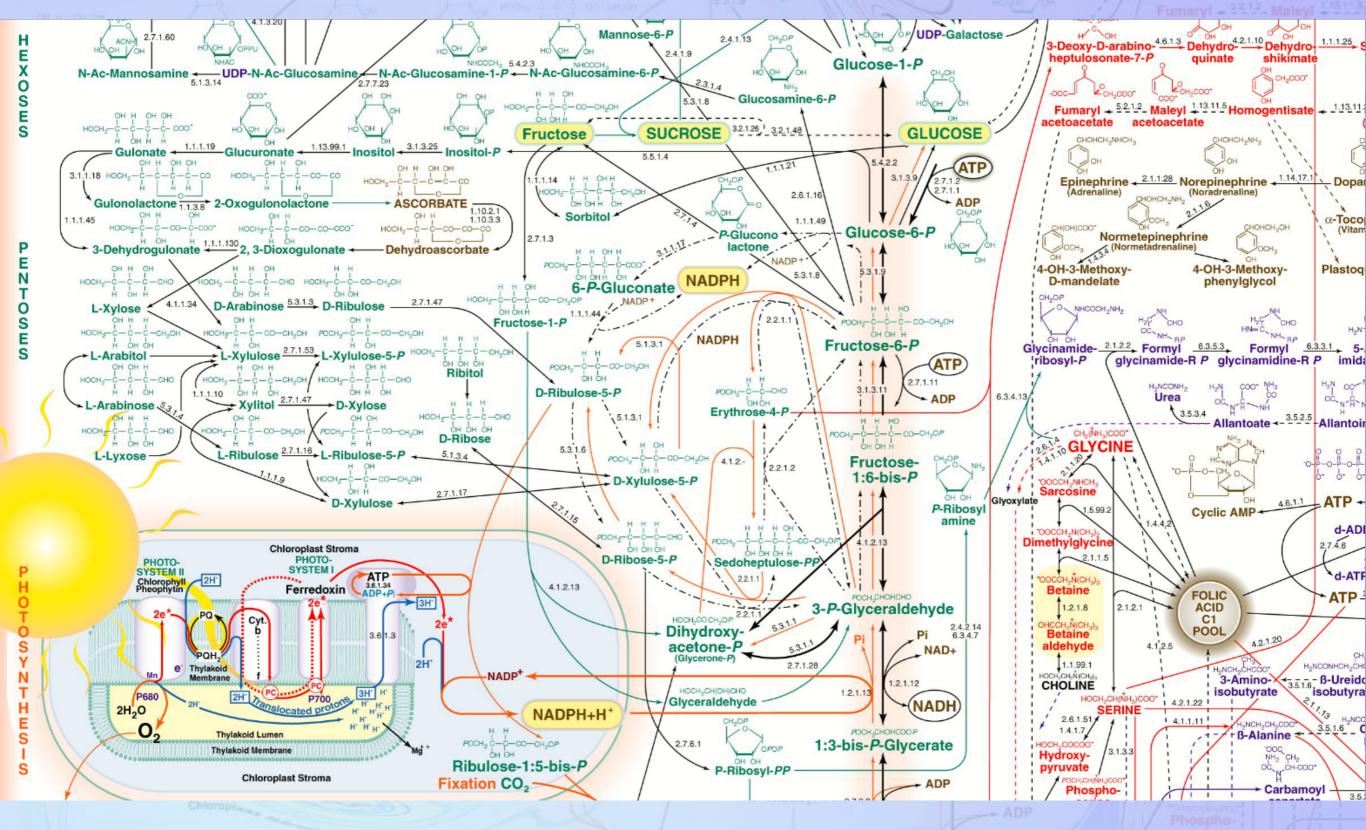


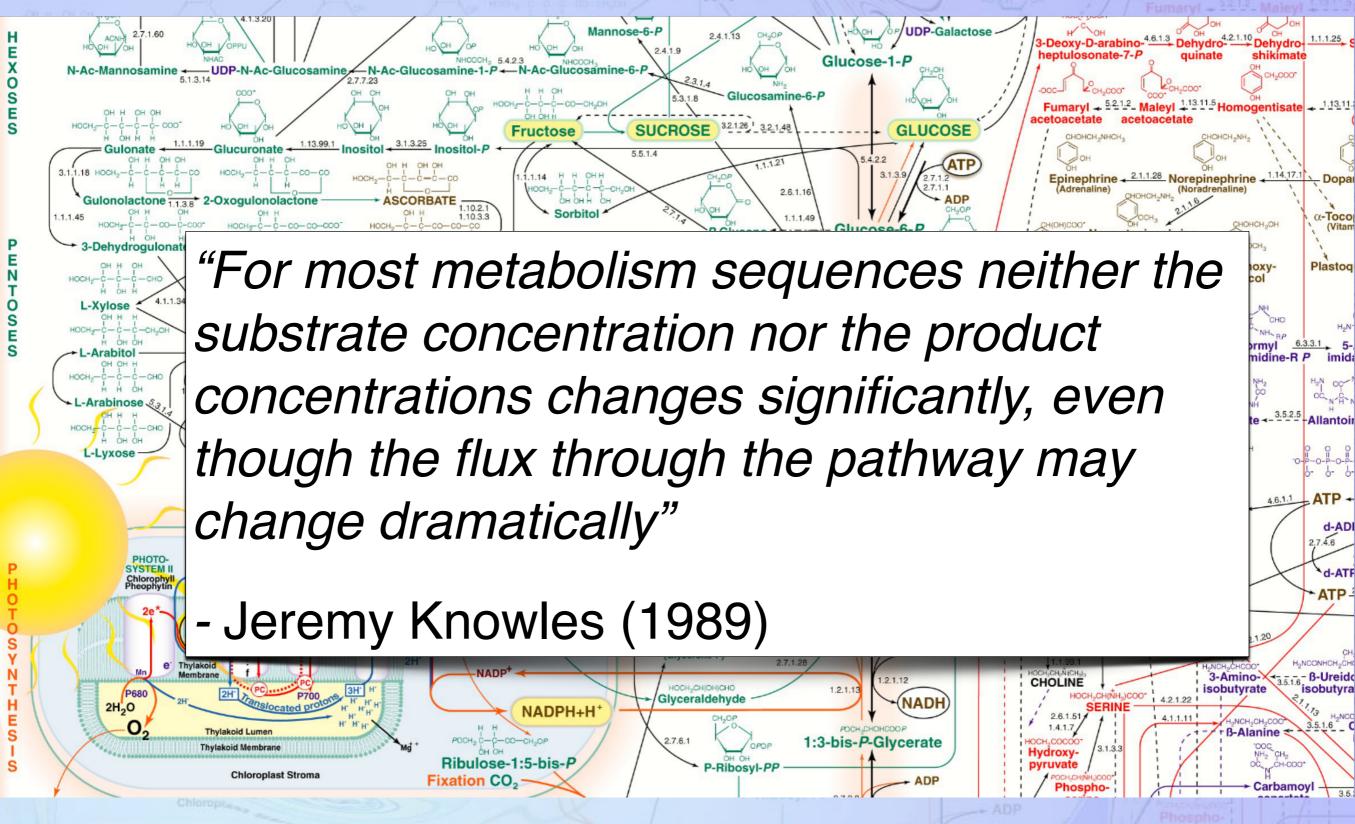
- ·Metabolism is the sum total of all of the reactions that take place in a living cell.
 - + Made up of
 - · metabolites
 - enzyme catalzyed reactions

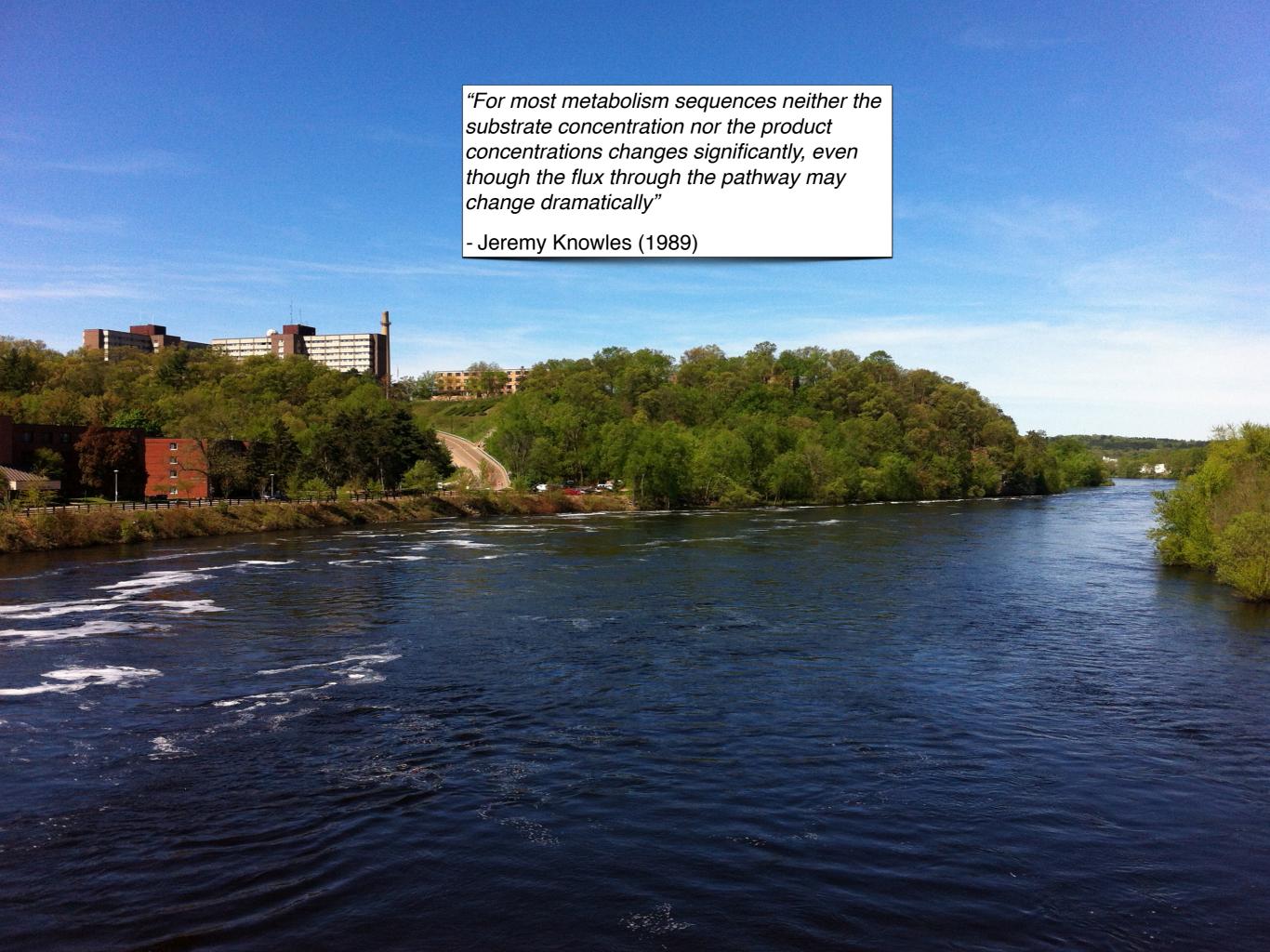


Chem 352, Lecture 7, Introduction to Metabolism

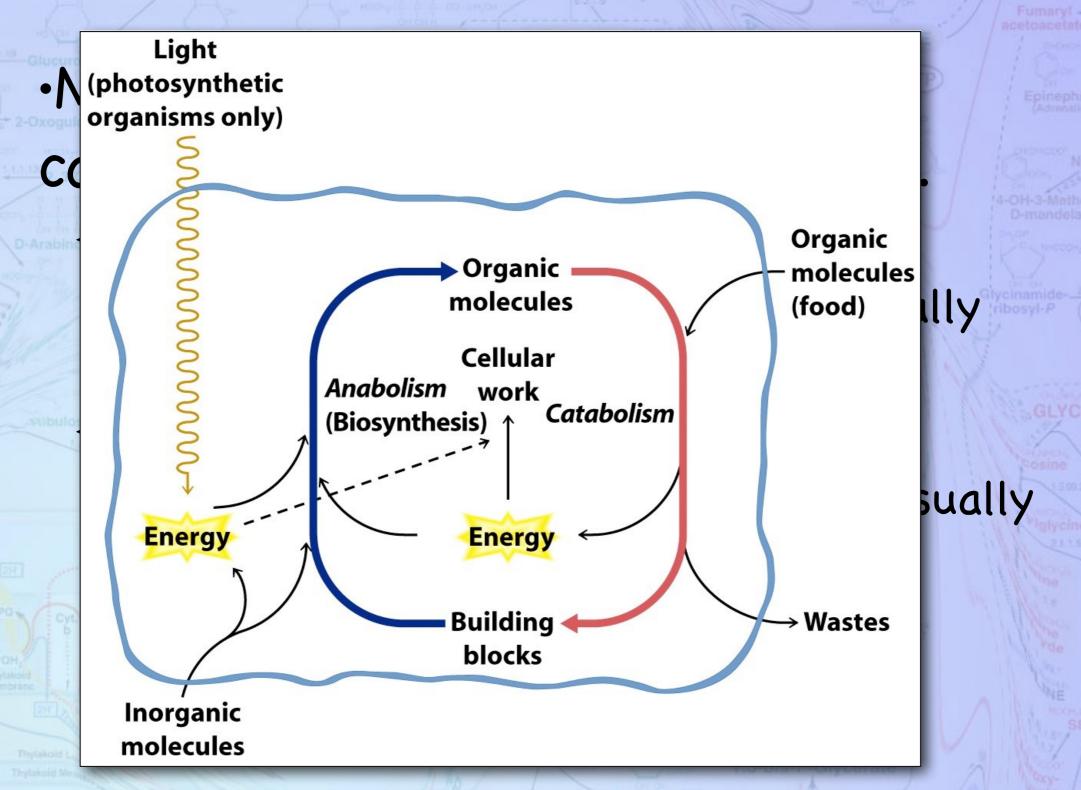
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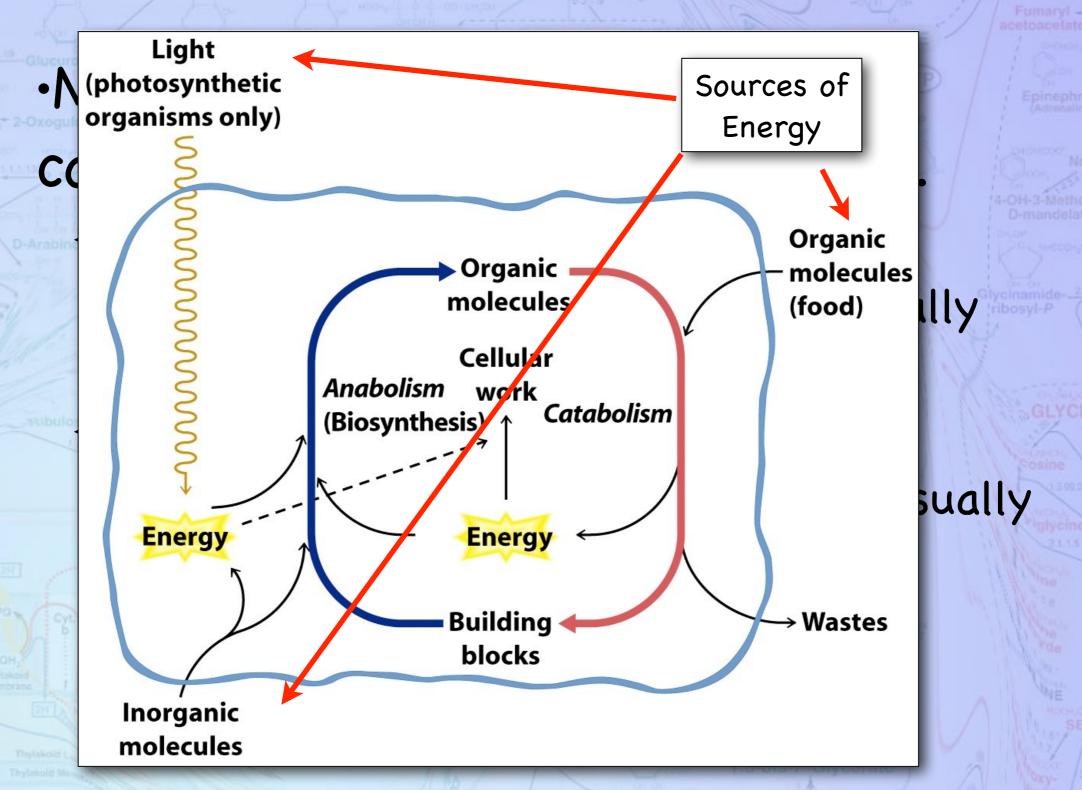


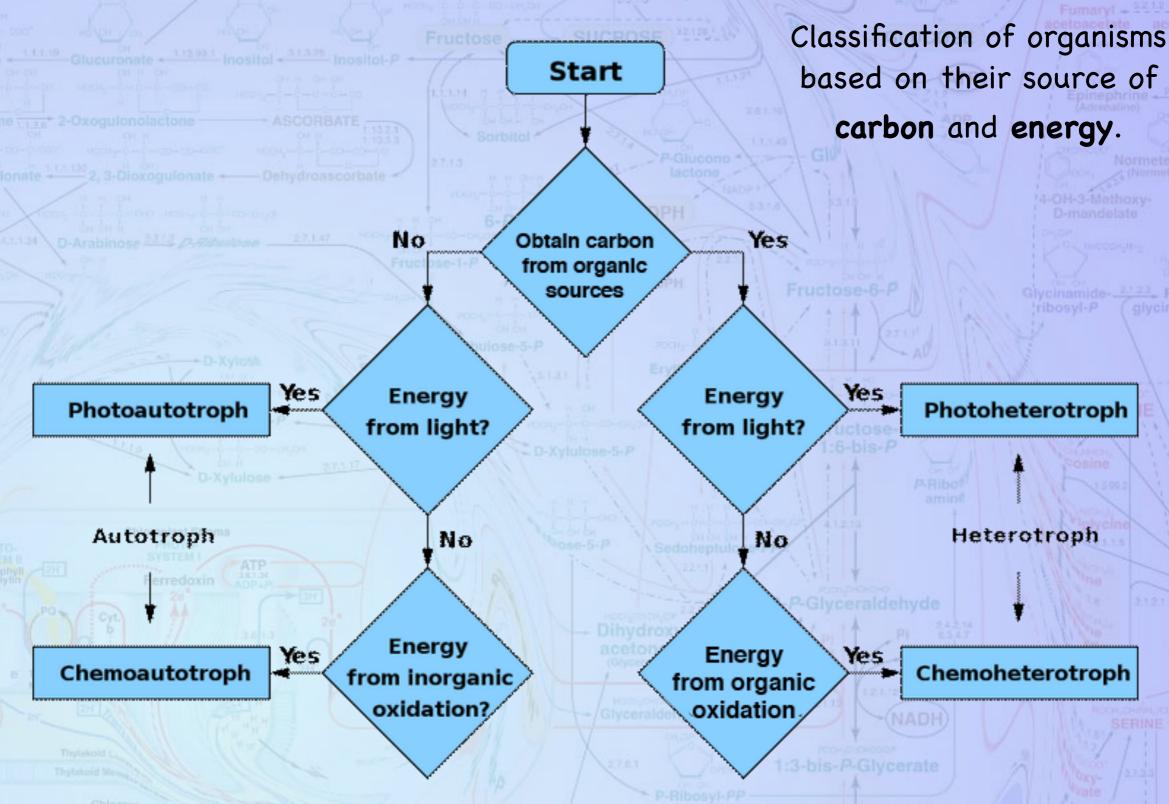




- ·Metabolism is divided into two complimentary sets of reactions.
 - * Anabolic reactions (anabolism)
 - The synthetic reactions, which usually require an input in free energy.
 - + Catabolic reaction (catabolism)
 - The degradative reactions, which usually lead to a release of free energy.







Common themes in metabolism:

Chem 352, Lecture 7, Introduction to Metabolism

Common themes in metabolism:

 Organisms or cells maintain specific internal concentrations of inorganic ions, metabolites, and enzymes. Cell membranes provide the physical barrier that segregates cell components from the environment.

Common themes in metabolism:

 Organisms extract energy from external sources to drive energy-consuming reactions. Photosynthetic organisms derive energy from the conversion of solar energy to chemical energy. Other organisms obtain energy from the ingestion and catabolism of energyyielding compounds

Common themes in metabolism:

• The metabolic pathways in each organism are specified by the genes it contains in its genome.

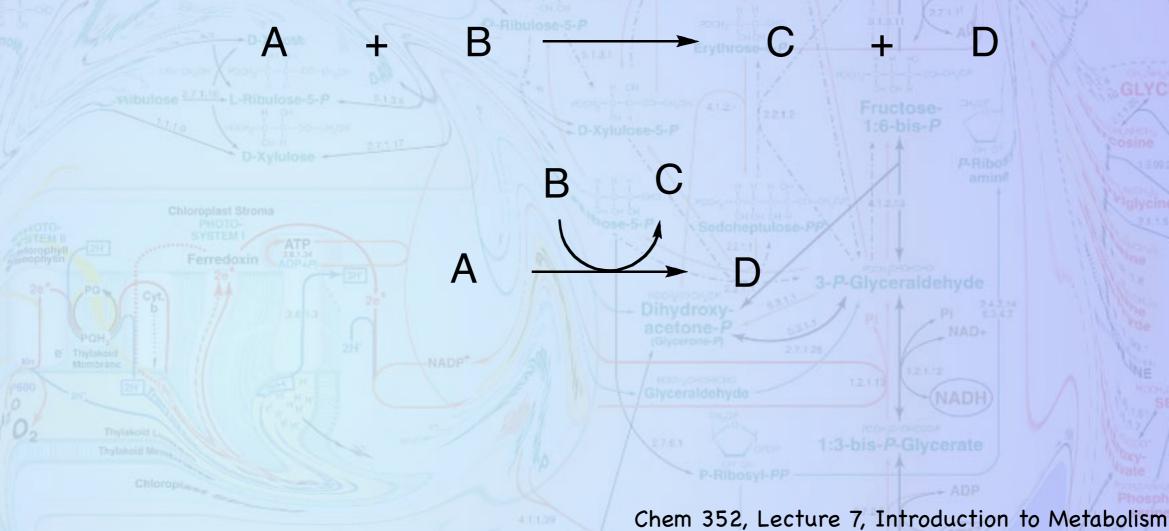
Common themes in metabolism:

• Organisms and cells interact with their environment. The activities of cells must be geared to the availability of energy. Organisms grow and reproduce when the supply of energy from the environment is plentiful. When the supply of energy from the environment is limited, energy demands can be temporarily met by using internal stores or by slowing metabolic rates as in hibernation, sporulation, or seed formation. If the shortage is prolonged, organisms die.

Common themes in metabolism:

• The cells of organisms are not static assemblies of molecules. Many cell components are continually synthesized and degraded, that is, they undergo turnover, even though their concentrations may remain virtually constant. The concentrations of other compounds change in response to changes in external or internal conditions.

The enzymes arrange the metabolites into pathways.



The enzymes arrange the metabolites into pat 3-Phosphoglycerate

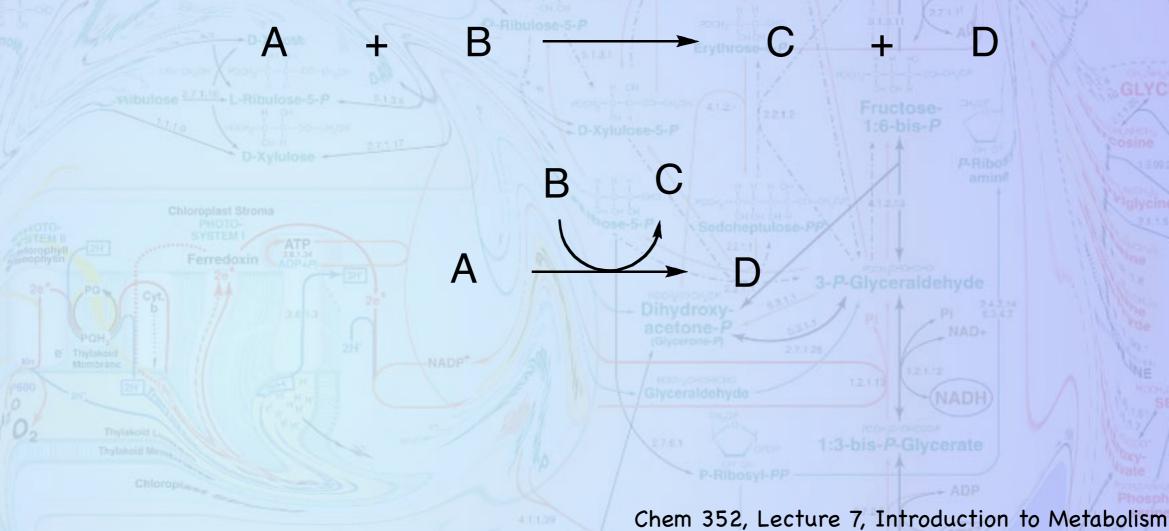


3-Phosphoserine

Linear Pathway

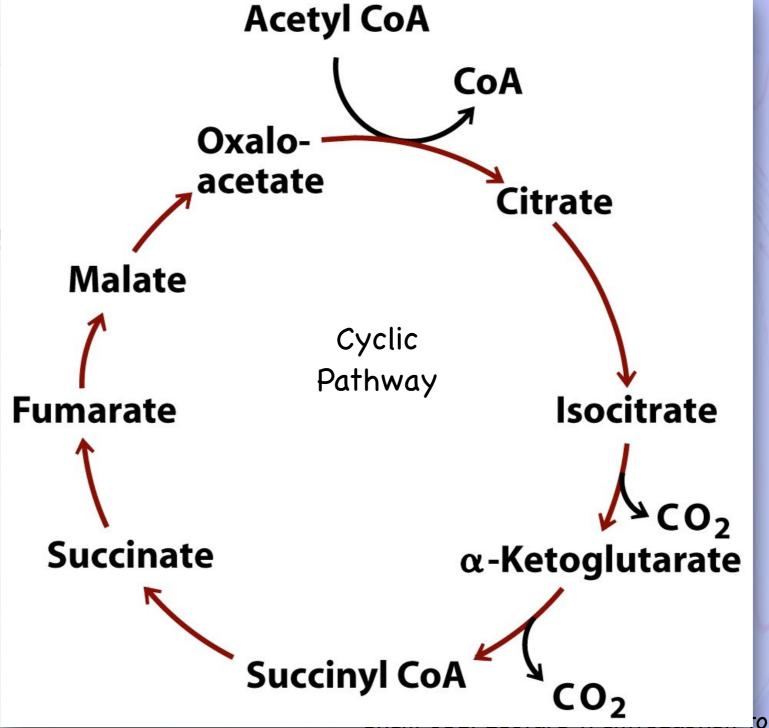
Serine

The enzymes arrange the metabolites into pathways.

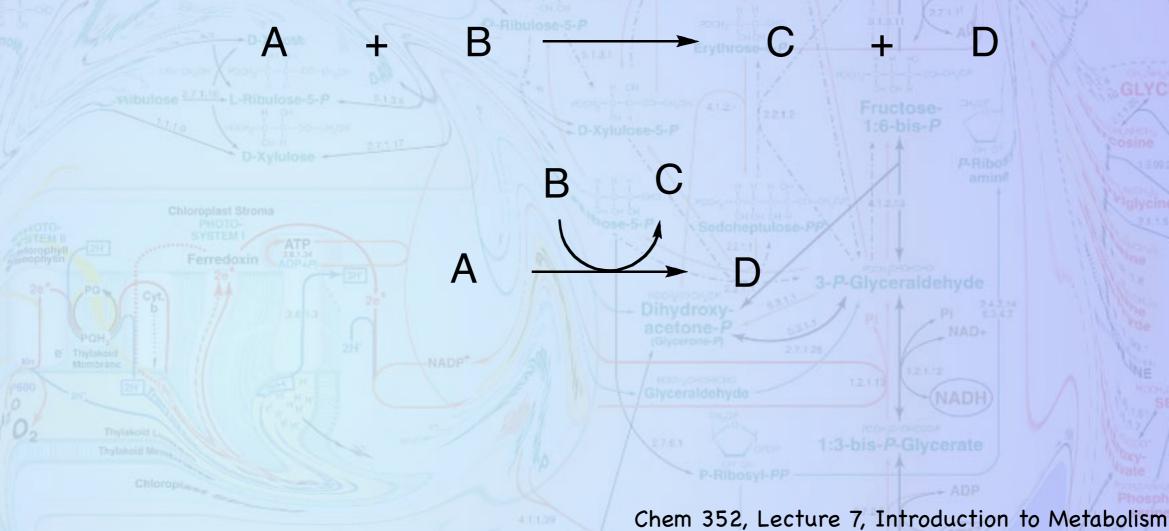


The enzymes arrange the metabolites

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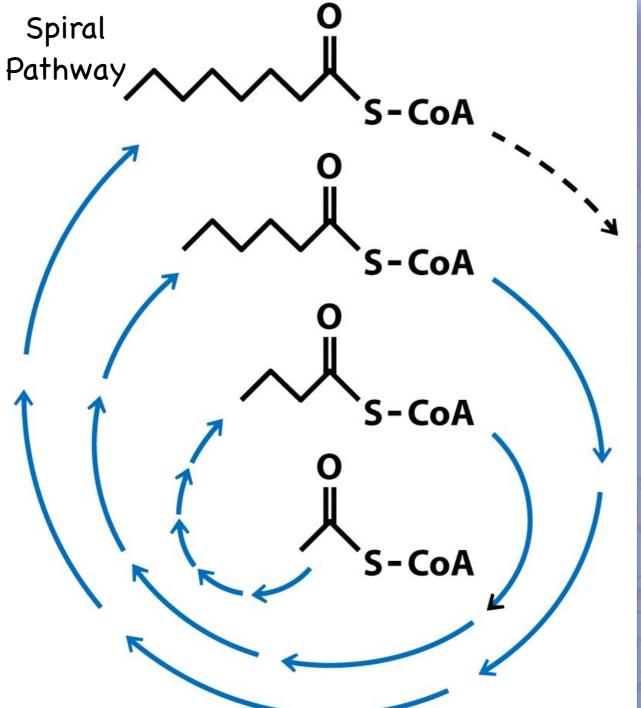


The enzymes arrange the metabolites into pathways.



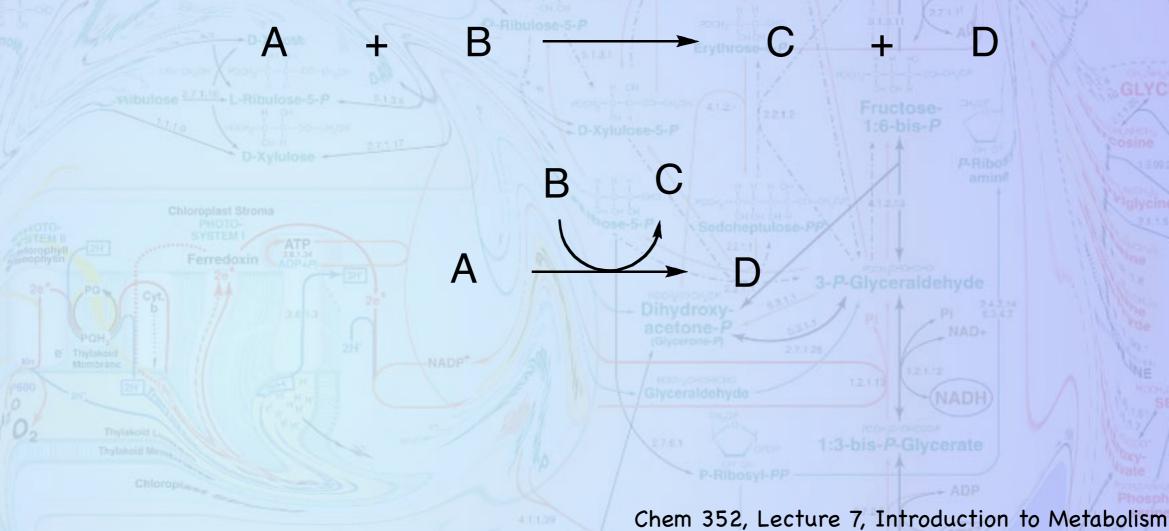
The enzymes arrange the metabolites

into path Spiral Pathway



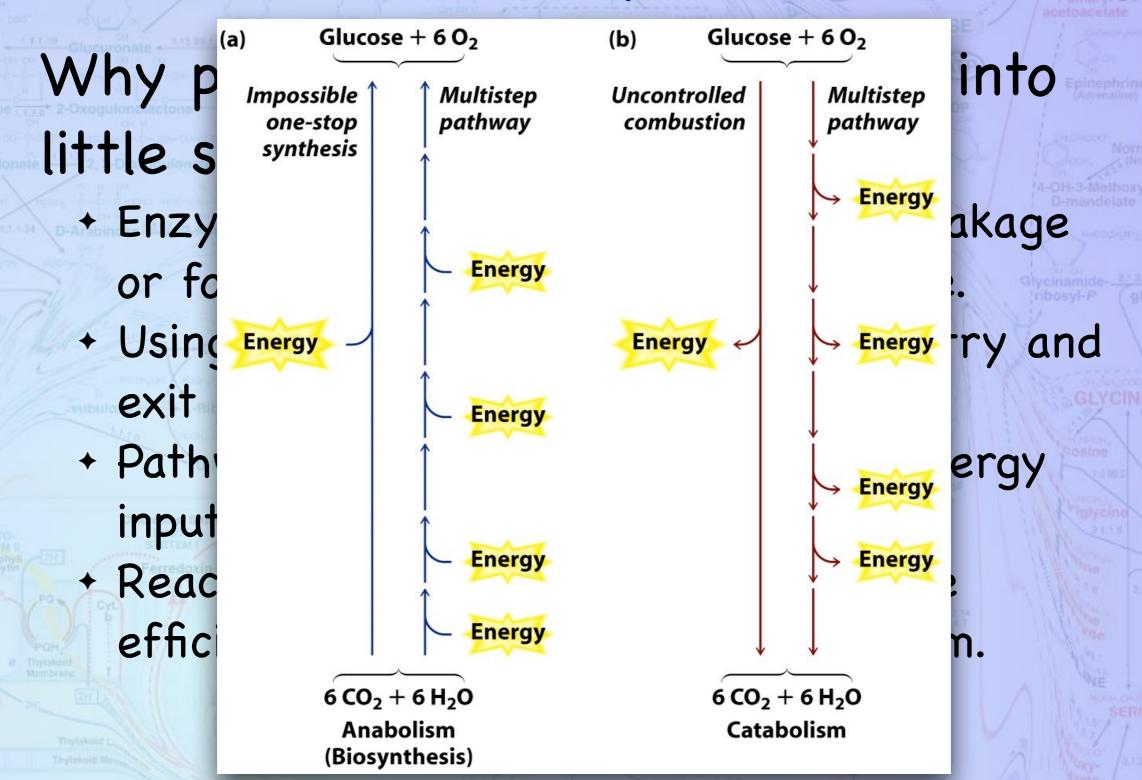
iction to Metabolism

The enzymes arrange the metabolites into pathways.



Why pathways are broken down into little steps.

- + Enzyme specificity allows only for breakage or formation of a few bonds at a time.
- + Using pathways allows for multiple entry and exit points for metabolites.
- + Pathways allow for finer control of energy input and output.
- * Reactions are thermodynamically more efficient if carried out near equilibrium.



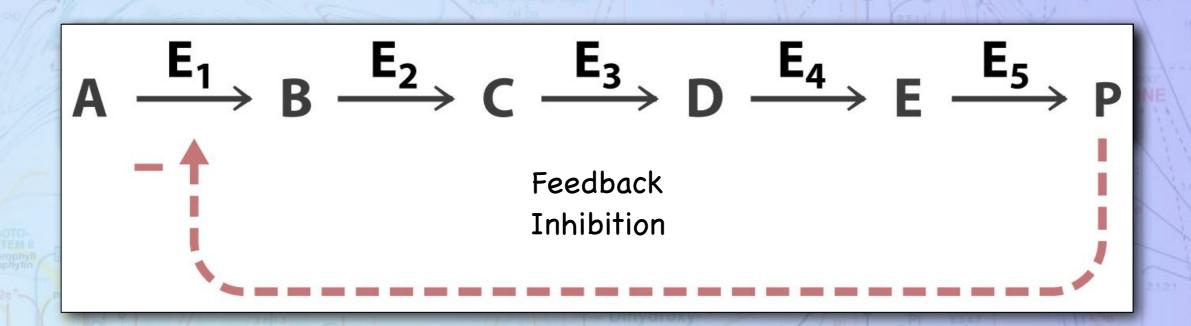
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- ·Pathways are regulated
 - + To control the flow of metabolites through a pathway

$$A \xrightarrow{E_1} B \xrightarrow{E_2} C \xrightarrow{E_3} D \xrightarrow{E_4} E \xrightarrow{E_5} P$$

- ·Pathways are regulated
 - + To control the flow of metabolites through a pathway



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$$A \xrightarrow{E_1} B \xrightarrow{E_2} C \xrightarrow{E_3} D \xrightarrow{E_4} E \xrightarrow{E_5} P$$
Feed-forward
Activation

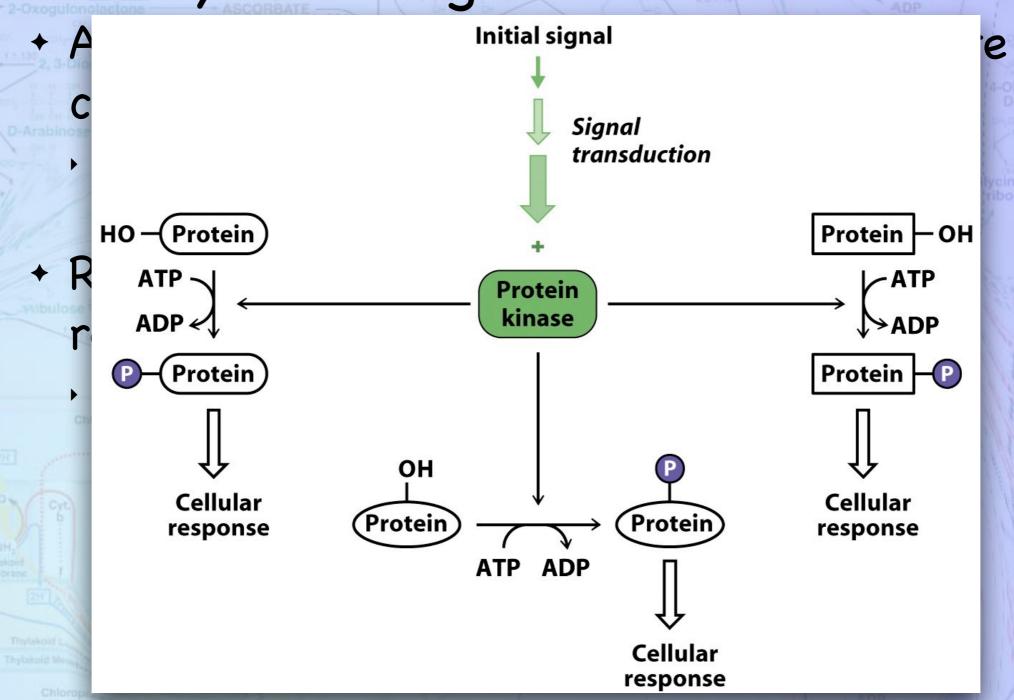
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Pathways are regulated

- + Allosteric regulation responds to immediate conditions within the cell,
 - · And have short term response times.
- * Reversible covalent modifications typically respond to extracellular signals,
 - · And have longer term response times.

Pathways are regulated



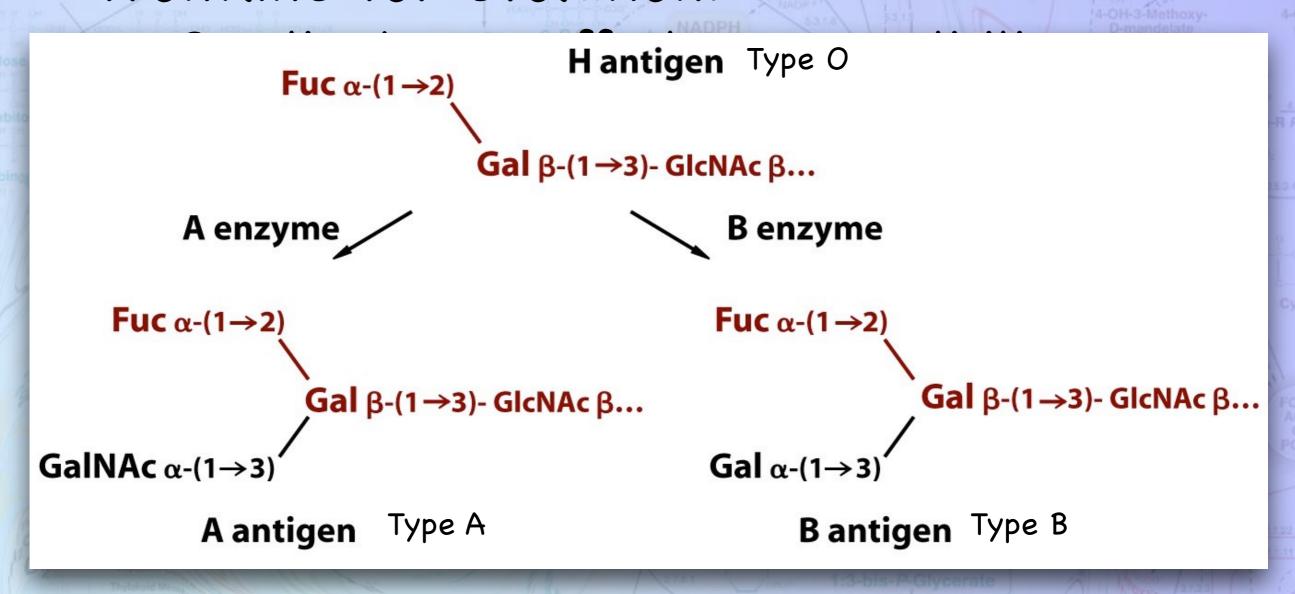
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Metabolic pathways represent the frontline for evolution.

 Genetic changes affect enzyme activities, which in turn, affect the flow of material through metabolic pathways.

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Metabolic Pathways

Metabolic pathways represent the frontline for evolution.

+ Genetic chal which in tur through met



Andi Stempniak

Jennifer Czubakowski holds a picture of her brother, Tommy along with her parents Theresa and Ron in the living room of their Eau Claire home. Jenny Czubakowski held a picture of her late brother, Tommy, at the rural Eau Claire home she shares with their parents, Theresa and Ron. The family is trying to help people be aware of ornithine transcarbamylase deficiency, or OTC, which Tommy had.

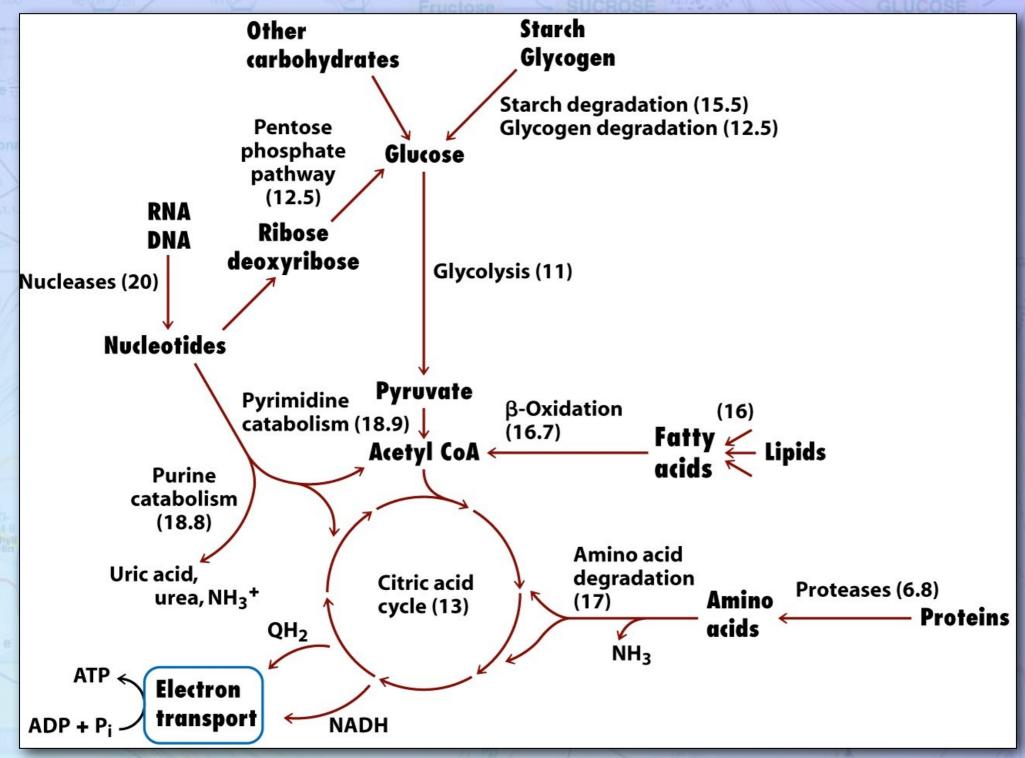
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Metabolic Pathways

Metabolic pathways represent the frontline for evolution.

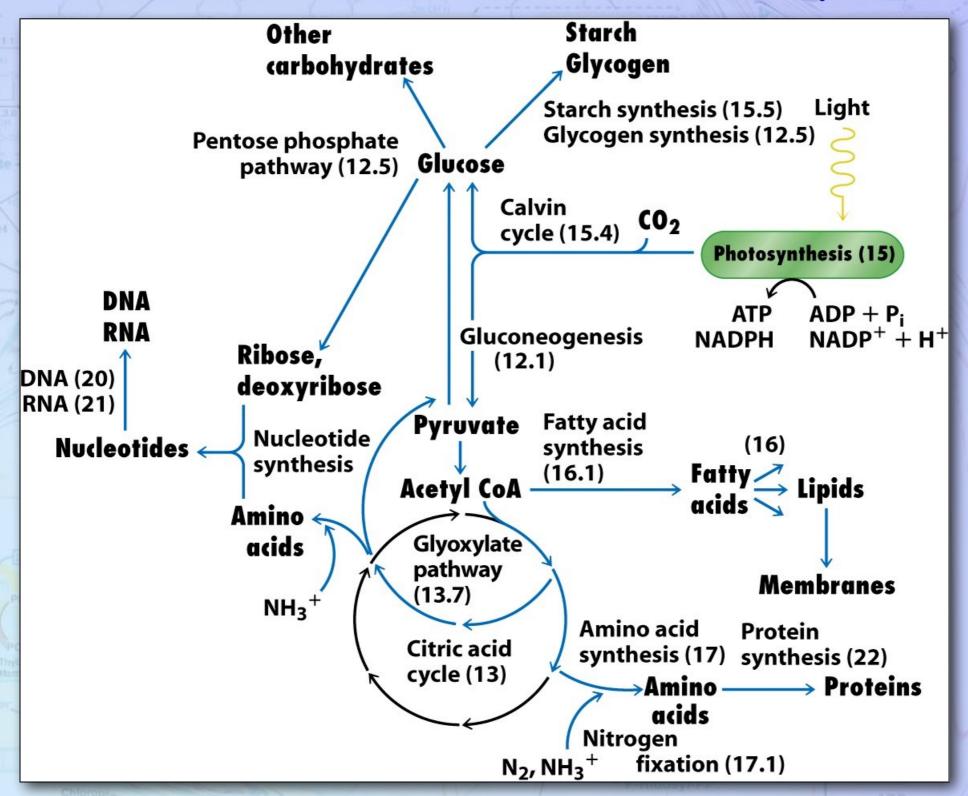
 Genetic changes affect enzyme activities, which in turn, affect the flow of material through metabolic pathways.

Major Metabolic Pathways



Catabolic Pathways

Major Metabolic Pathways



Anabolic Pathways

Major Metabolic Pathways

+ In many organisms, the various pathways are regulated through compartmentalization.

Golgi apparatus P (end-on view) sorting and secretion of some proteins

Mitochondria: citric acid cycle, electron transport + ATP synthesis, fatty acid degradation

Lysosome: degradation of proteins, lipids, etc.

Plasma membrane

Cytosol: fatty acid synthesis, glycolysis, most gluconeogme:s reaction pentose phosphase pathwwary

Nucleus: nucleic acid synthesis

Endoplasmic reticulum: delivery of proteins and synthesis of lipids for membranes

Nuclear membranes

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•The spontaneity (favorability) of a chemical reaction can be determined from its Gibbs Free Energy (Δ G)

$$\Delta G = \Delta G^{0'} + RT \ln \left(\frac{[products]}{[reactants]} \right)$$

Under conditions of constant temperature and pressure there are two contributions to the free energy change

- + Enthalpy, H
- + Entropy, S

$$\Delta G = \Delta H - T \Delta S$$

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Change in heat content

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- + Enthalpy, H
- + Entropy, S

$$\Delta G = \Delta H - T \Delta S$$
Change in heat content Change in disorder

Change in hear contem

•The actual conditions within the cell must be considered when determining a ΔG value.

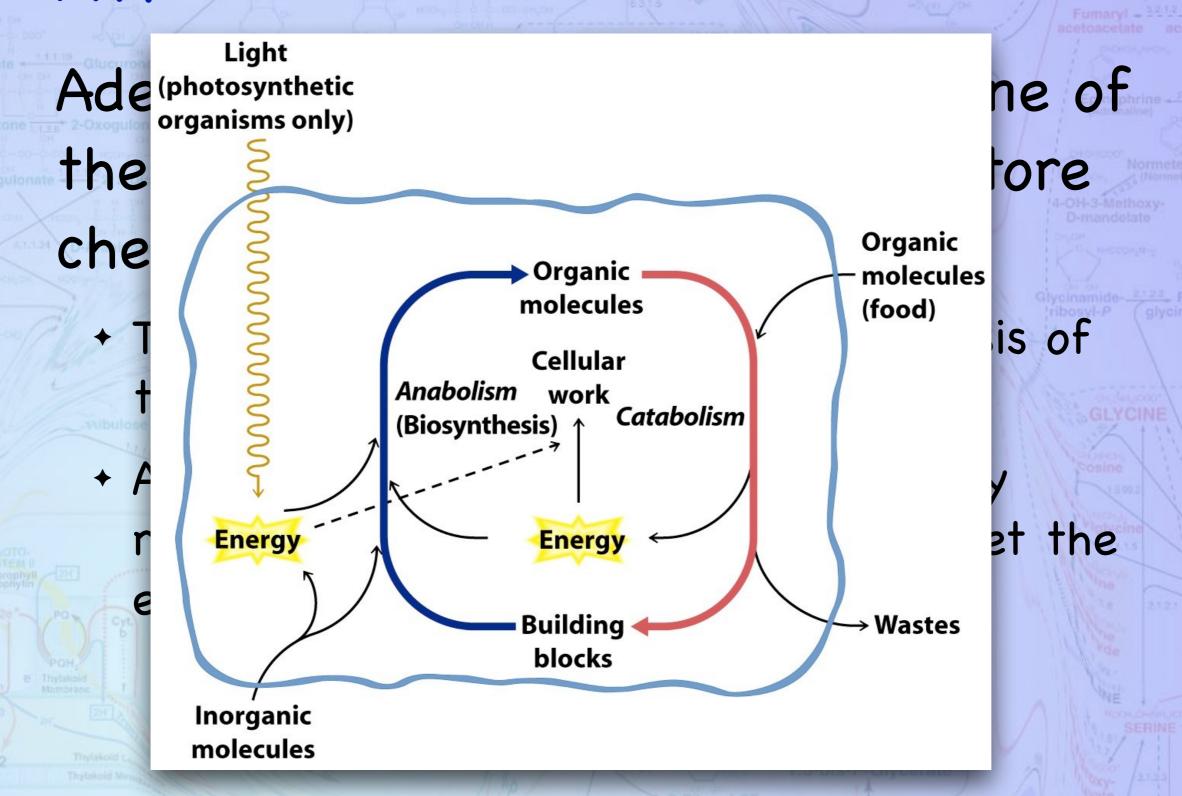
$$\Delta G = \Delta G^{\circ'} + RT \ln \left(\frac{[C][D]}{[A][B]} \right)$$

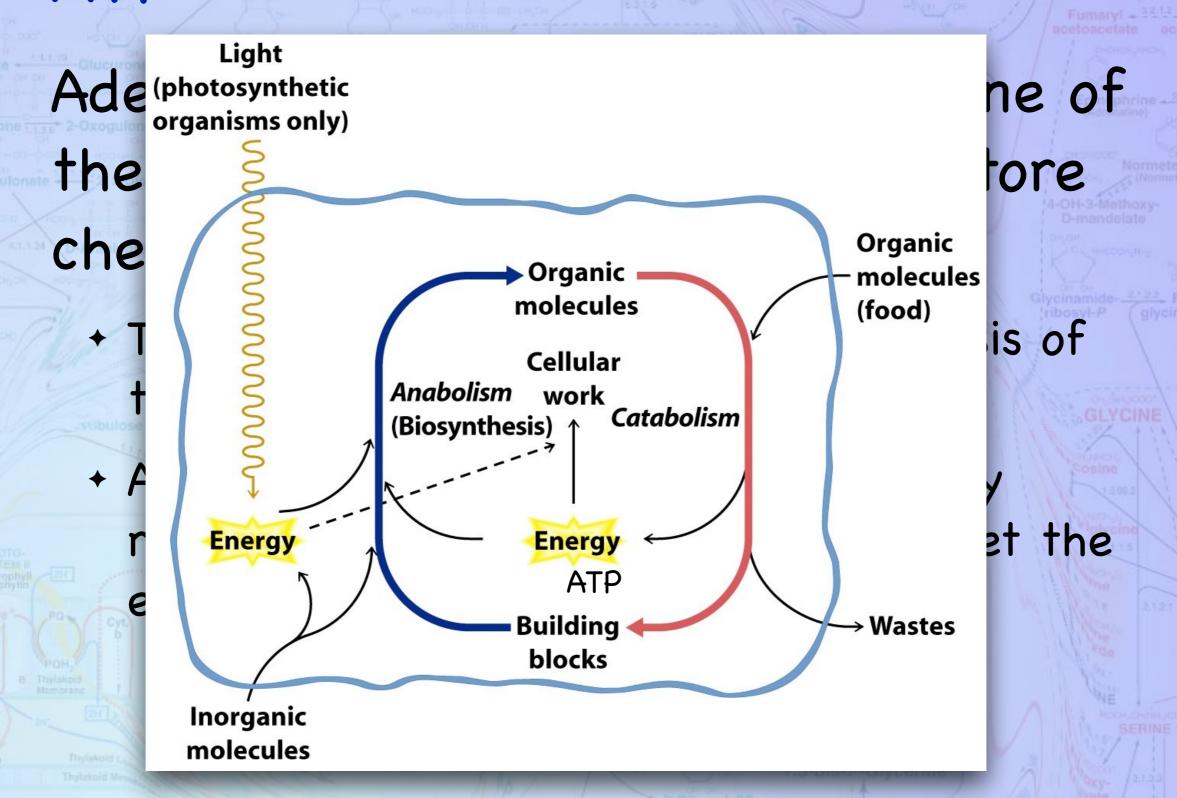
$$Q = \begin{pmatrix} [C][D] \\ [A][B] \end{pmatrix}$$
 is the mass action ratio

- •When $Q \approx K_{eq}$ a reaction is reversible.
- •When $Q < K_{eq}$ a reaction is spontaneous and irreversible.
- •When Q > K_{eq} a reaction is nonspontaneous and irreversible

Adenosine Triphosphate (ATP) is one of the molecules used by a cell to store chemical energy.

- + This energy is released by the hydrolysis of the two phosphate anhydride bonds.
- + ATP is one of the ways that the energy released from catabolism is used to meet the energy requirements of anabolism





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A th

Adenosine 5'-triphosphate (ATP (4))

$$\Theta_{O} - P - O - P - O - Adenosine$$

$$(2) \begin{array}{c} H_2O \\ H^{\oplus} \end{array}$$

$$\bigcirc O - P \stackrel{\square}{=} O - Adenosine$$

Adenosine 5'-diphosphate (ADP (3-)) Adenosine 5'-monophosphate (AMP (2-))

Inorganic pyrophosphate (PP_i)

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of

the



of

TABLE 10.1 Standard Gibbs free energies of hydrolysis for ATP, AMP, and pyrophosphate.

Reactants and products	$\Delta G^{\circ}{}'_{ m hydrolysis}$ l $({ m kJ~mol}^{-1})$
$ATP + H_2O \rightarrow ADP + P_i + H^{\oplus}$	-32
$ATP + H_2O \rightarrow AMP + PP_i + H^{\oplus}$	-45
$AMP + H_2O \rightarrow Adenosine + P_i$	-13
$PP_i + H_2O \rightarrow 2 P_i$	-29

 P_i (inorganic phosphate) = HPO_4^{2-}

 PP_i (pyrophosphate) = $HP_2O_7^{3-}$

$$Ho - \ddot{\beta} - o^{\Theta}$$
 $Ho - \ddot{\beta} - o - \ddot{\beta} - o^{\Theta}$ O^{Θ} O^{Θ} O^{Θ} Inorganic pyrophosphate (PP_i)

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- + ATP is one of the ways that the energy released from catabolism is used to meet the energy requirements of anabolism

- ·Adenosine Triphosphate (ATP) is just one of the molecules used by a cell to store chemical energy.
 - + The other ribonucleotide triphosphates are also used for this same purpose.
 - · Guanosine triphosphate (GTP)
 - · Cytidine triphosphate (CTP)
 - · Uridine triphosphate (UTP



Question:

In a rat hapatocyte, the concentrations ATP, ADP and Pi are 3.4 mM, 1.3 mM and 4.8 mM, respectively. Calculate the Gibbs free energy for the hydrolysis of ATP in this cell. How does this compare to the standard free energy change?

Question:

 PP_i (pyrophosphate) = $HP_2O_7^{3-}$

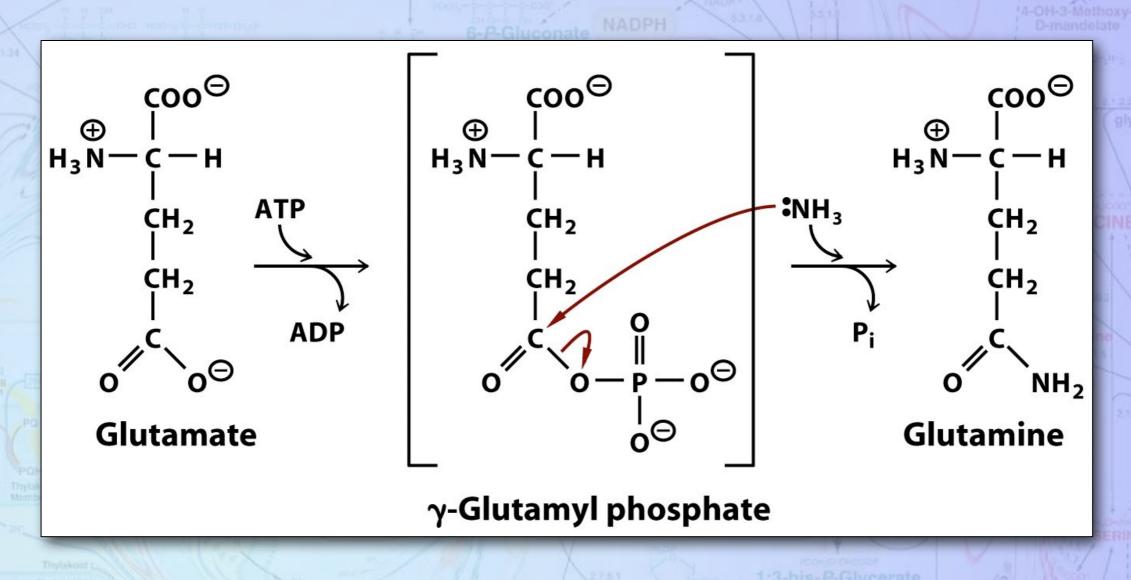
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·The hydrolysis of ATP can be used to drive unfavorable reactions





·Phosphoryl-group-transfer potential

TABLE 10.3	Standard Gibbs free en-
ergies of hydi	olysis for common
metabolites	

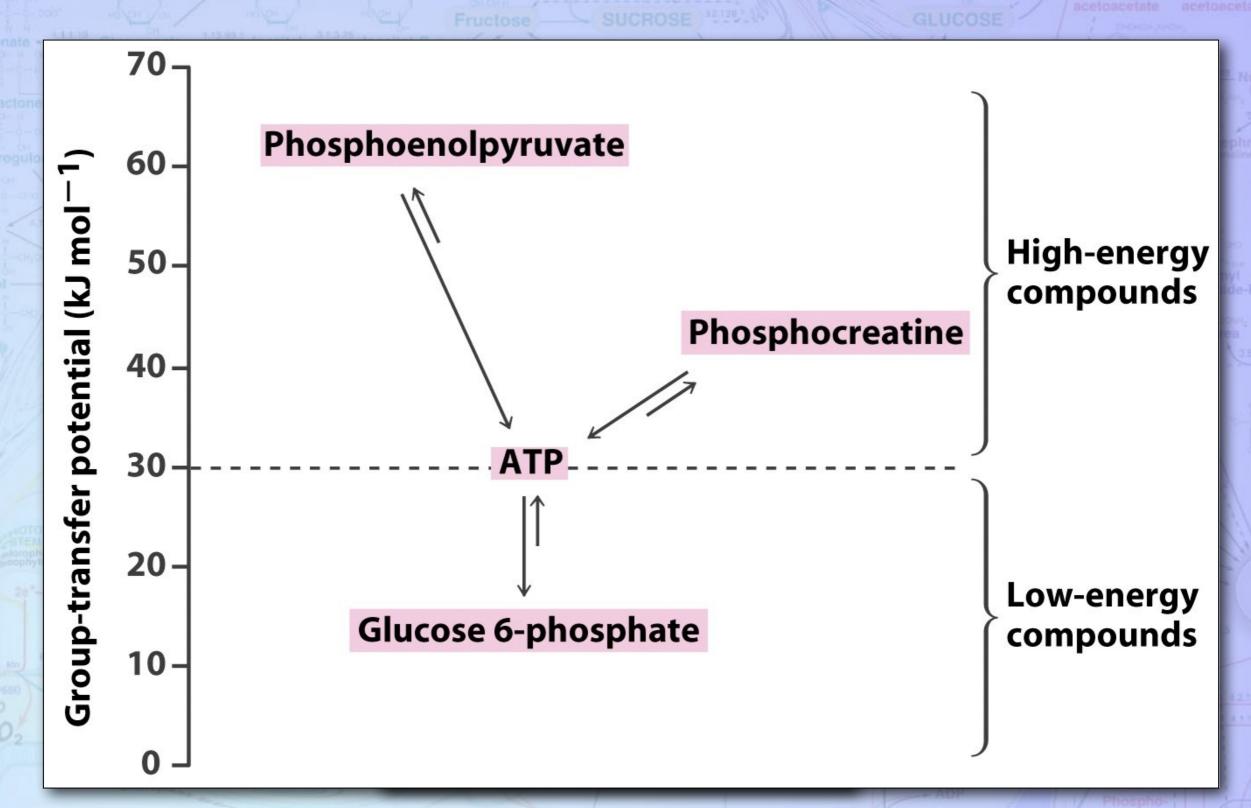
Metabolite	$\Delta G^{\circ}{}'_{ m hydrolysis}$ (kJ mol $^{-1}$)
Phosphoenolpyruvate	-62
1,3-Bisphosphoglycerate	-49
ATP to AMP $+$ PP _i	-45
Phosphocreatine	-43
Phosphoarginine	-32
Acetyl CoA	-32
ATP to ADP $+ P_i$	-32
Pyrophosphate	-29
Glucose 1-phosphate	-21
Glucose 6-phosphate	-14
Glycerol 3-phosphate	-9



·Phosphoryl-group-transfer potential

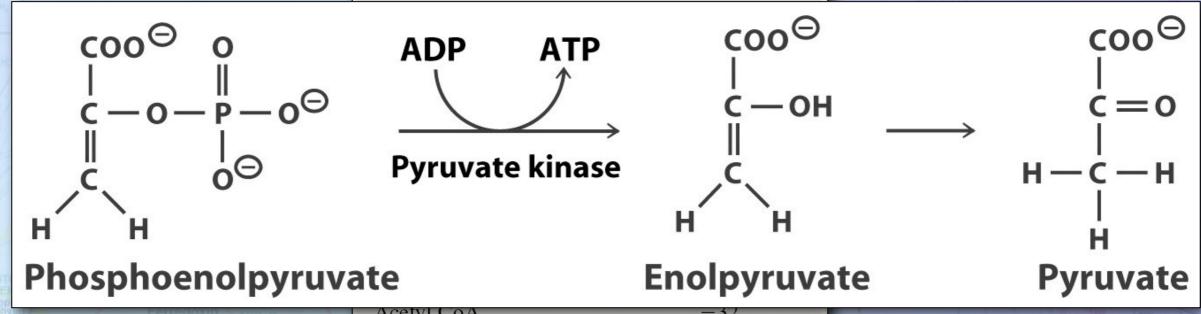
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TABLE 10.3 Standard Gibbs free energies of hydrolysis for common metabolites



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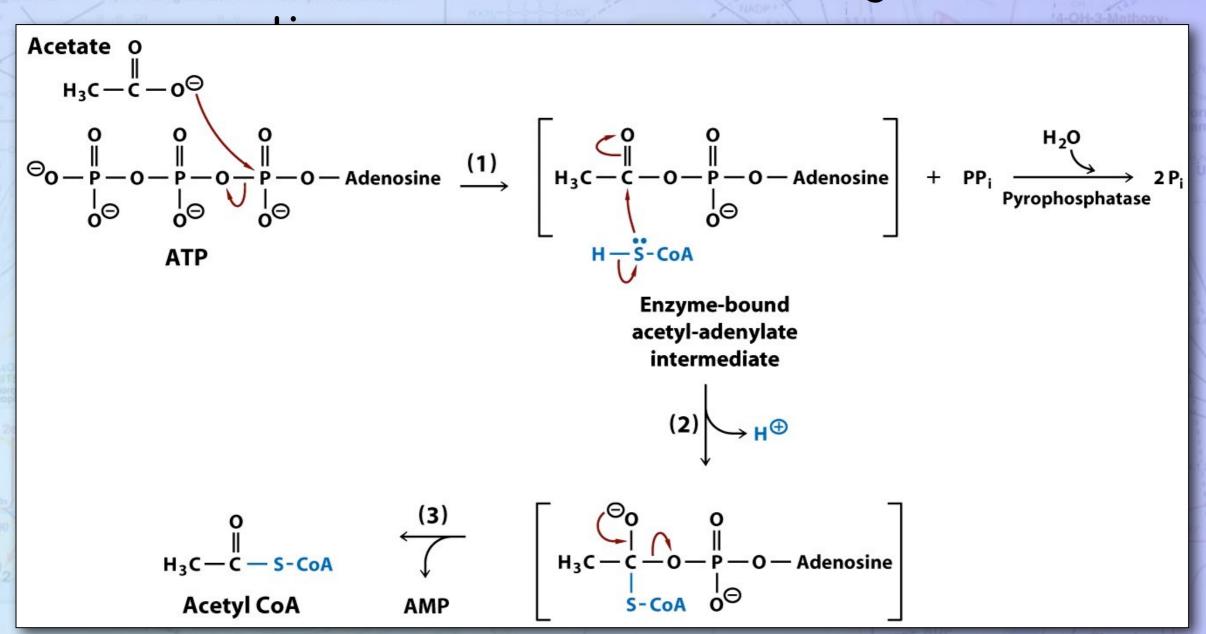


·Nucleotidyl group transfer

+ Used to activate substrates in ligase reactions

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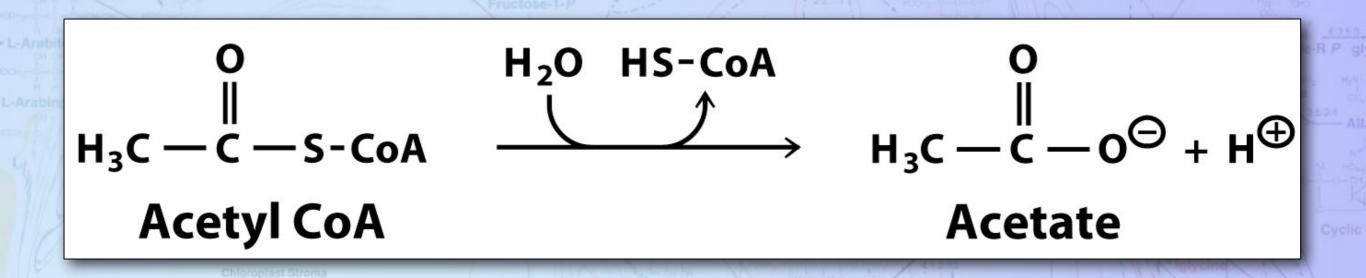




·Nucleotidyl group transfer

+ Used to activate substrates in ligase reactions

·The thioester group also has a high energy for hydrolysis



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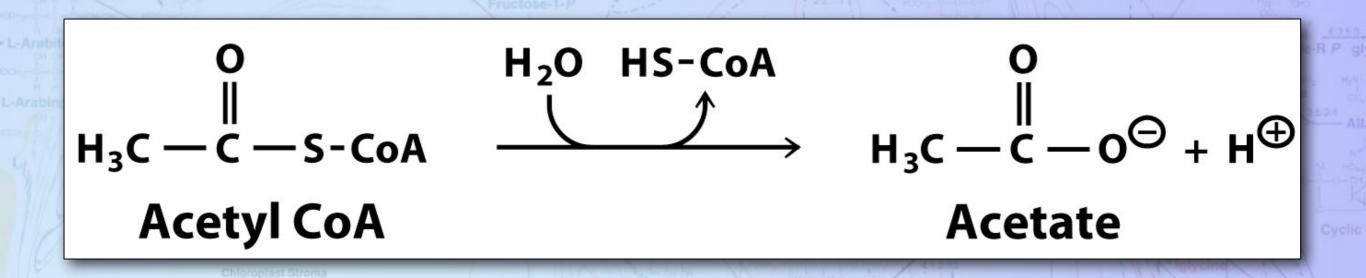
TABLE 10.3 Standard Gibbs free enenergy for ergies of hydrolysis for common metabolites

0
$H_3C - \ddot{C} - S - CoA$
A cotud CoA
Acetyl CoA

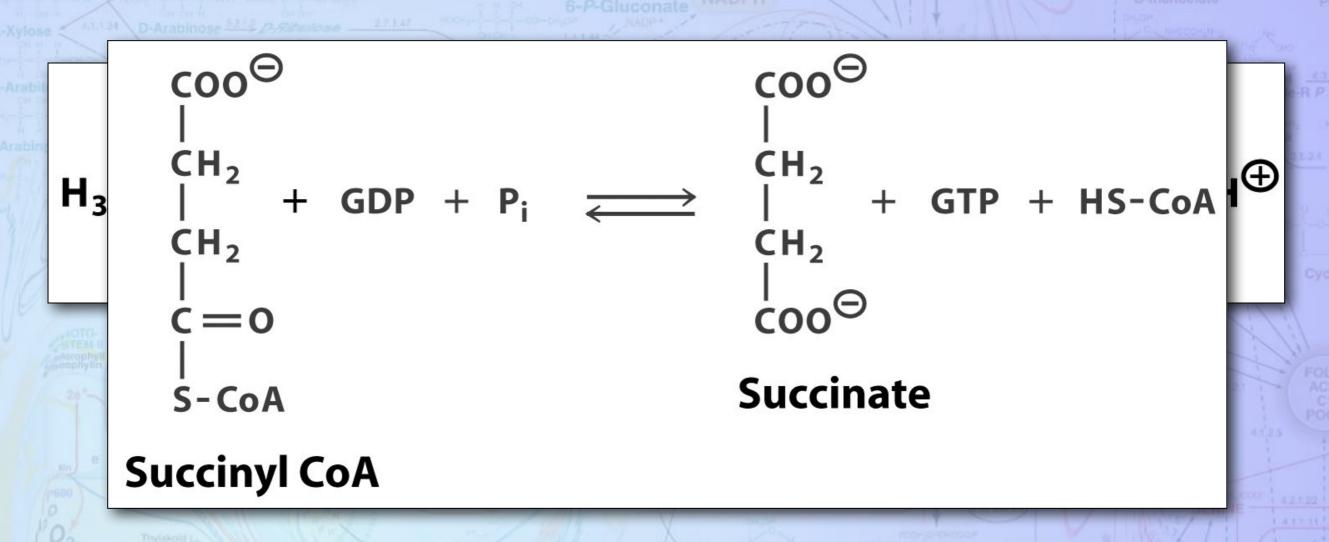
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Glycerol 3-phosphate	-9

· c" — o[⊜] + H[⊕] **Acetate**

·The thioester group also has a high energy for hydrolysis

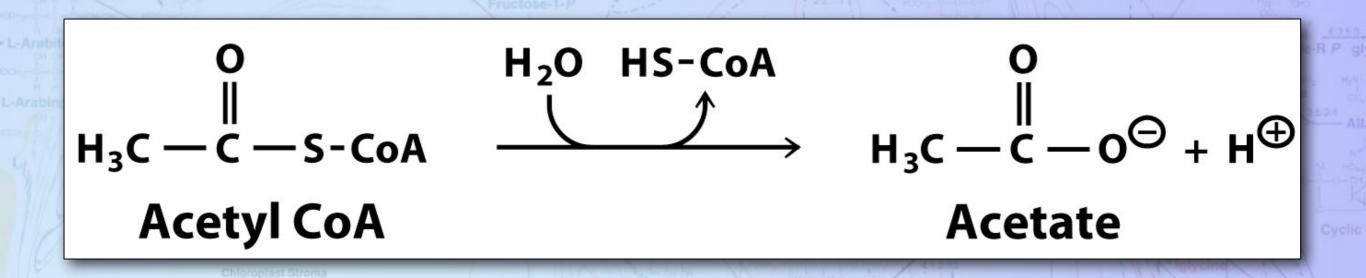


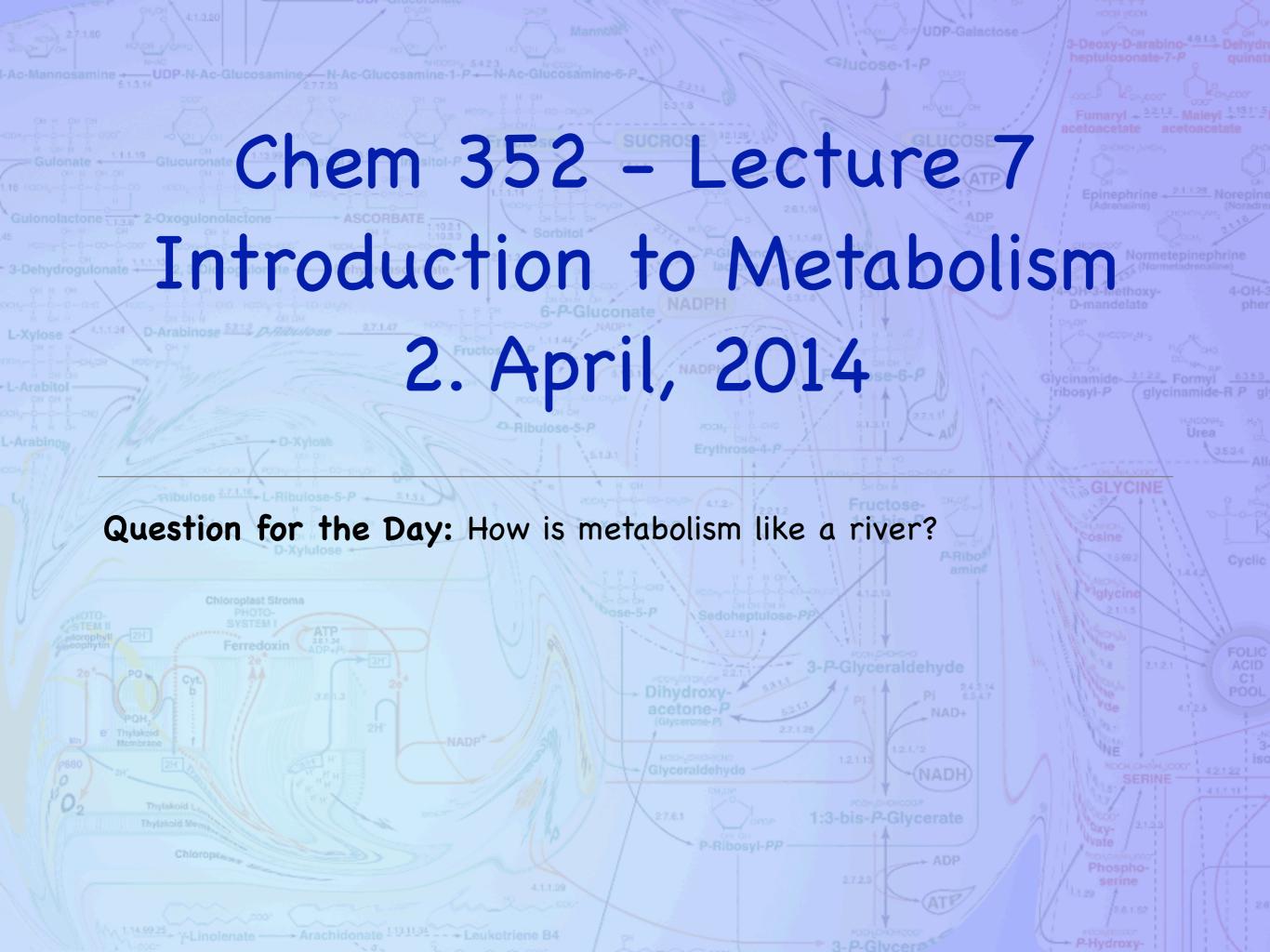
·The thioester group also has a high energy for hydrolysis



Thioesters as High Energy Compounds

·The thioester group also has a high energy for hydrolysis





- •Reduced coenzymes (NAD, NADP, FAD, FMN, ubiquinone) provide another way to store chemical energy.
 - + They can be used to store the free energy that is released in oxidation reactions.
 - + The electrons released in these reactions are transferred to the coenzyme, usually in the form of a hydride (H:-) ion.

Reduction potentials can be used to measure the ability of a molecule to serve as a reducing agent in an oxidation/reduction reaction

Aoxidized + Breduced -> Areduced + Boxidized

(B is the reducing agent in this reaction)

Reduction potentials can be measured with an electrochemical cell.

+ The oxidation and reduction are separated by a wire.

The reduction of Cu2+ by Zn

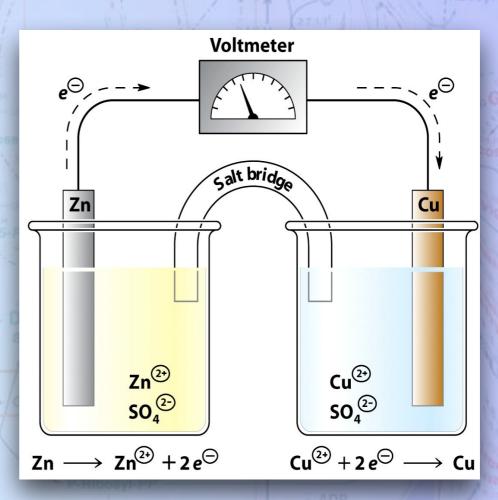
$$Zn + Cu^{2+} \leftrightarrow Zn^{2+} + Cu$$

$$\Delta G' = -n\mathcal{F}\Delta E'$$
 (Nernst Equation)

 $\Delta E'$ = potential measured by the voltmeter

n = number of electrons transferred

 $F = Faraday's constant (96,586 JV^{-1}mol^{-1})$



The change in the reduction potential for an oxidation/reduction reaction $(\Delta E^{o'})$ can be used to determine the change in Free energy for the reaction.

$$\Delta E^{o'} = E^{o'}_{\text{electron acceptor}} - E^{o'}_{\text{electron donor}}$$

$$\Delta G^{o'} = -n\mathscr{F} \Delta E^{o'}$$

n = number of electrons transferred

 $F = Faraday's constant (96,586 JV^{-1}mol^{-1})$

Standard reduction potentials, Eo', are usually measured with respect to the reduction potential for of 2 H⁺(aq) \rightarrow H₂(g)

TABLE 10.4 Standard reduction potentials of some important biological half-reactions

Reduction half-reaction	$E^{\circ \prime}$ (V)
Acetyl CoA + CO ₂ + H $^{\oplus}$ + 2 e^{\ominus} \rightarrow Pyruvate + CoA	-0.48
Ferredoxin (spinach), $Fe^{\bigcirc} + e^{\bigcirc} \rightarrow Fe^{\bigcirc}$	-0.43
$2 \text{ H}^{\oplus} + 2e^{\bigcirc} \rightarrow \text{H}_2 \text{ (at pH 7.0)}$	-0.42
α -Ketoglutarate + CO ₂ + 2 H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow Isocitrate	-0.38
Lipoyl dehydrogenase (FAD) + 2 H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow Lipoyl dehydrogenase (FADH ₂)	-0.34
$NADP^{\oplus} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow NADPH + H^{\oplus}$	-0.32
$NAD^{\oplus} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow NADH + H^{\oplus}$	-0.32
Lipoic acid $+ 2 H^{\oplus} + 2e^{\ominus} \rightarrow$ Dihydrolipoic acid	-0.29
Glutathione (oxidized) + 2 H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow 2 Glutathione (reduced)	-0.23
$FAD + 2 H^{\oplus} + 2e^{\ominus} \rightarrow FADH_2$	-0.22
$FMN + 2 H^{\oplus} + 2e^{\ominus} \rightarrow FMNH_2$	-0.22
Acetaldehyde + 2 H $^{\oplus}$ + 2 e^{\ominus} \rightarrow Ethanol	-0.20
Pyruvate $+ 2 H^{\oplus} + 2e^{\ominus} \rightarrow Lactate$	-0.18
Oxaloacetate + $2 H^{\oplus} + 2e^{\ominus} \rightarrow Malate$	-0.17

TABLE 10.4 Standard reduction potentials of some important biological half-reactions

Reduction half-reaction	$E^{\circ}{}'$ (V)
Cytochrome b_5 (microsomal), $F_e^{(3-)} + e^{\bigcirc} \rightarrow F_e^{(2-)}$	0.02
Fumarate $+ 2 H^{\oplus} + 2e^{\ominus} \rightarrow Succinate$	0.03
Ubiquinone (Q) + 2 H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow QH ₂	0.04
Cytochrome b (mitochondrial), $F_e^{\stackrel{\text{(3)}}{\ominus}} + e^{\stackrel{\text{(2)}}{\ominus}} \rightarrow F_e^{\stackrel{\text{(2)}}{\ominus}}$	0.08
Cytochrome c_1 , $Fe^{\bigoplus} + e^{\bigoplus} \to Fe^{\bigoplus}$	0.22
Cytochrome c , Fe ³⁺ + $e^{\ominus} \rightarrow$ Fe ²⁺	0.23
Cytochrome a , Fe ^{3\rightarrow} + e ^{2\rightarrow} Fe ^{2\rightarrow}	0.29
Cytochrome f , $Fe^{\oplus} + e^{\ominus} \rightarrow Fe^{\ominus}$	0.36
Plastocyanin, $Cu^{2+} + e^{\bigcirc} \rightarrow Cu^{+}$	0.37
$NO_3^{\ominus} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow NO_2^{\ominus} + H_2O$	0.42
Photosystem I (P700)	0.43
$Fe^{\stackrel{\text{(3)}}{\rightarrow}} + e^{\stackrel{\text{(2)}}{\rightarrow}} Fe^{\stackrel{\text{(2)}}{\rightarrow}}$	0.77
$^{1}/_{2}O_{2} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow H_{2}O$	0.82
Photosystem II (P680)	1.1

·Like Δ G, the observed change in the reduction potential for a reaction, (Δ E), can be determined relative to the change in the standard reduction potential, (Δ E°):

$$\Delta E = \Delta E^{o'} - \frac{RT}{n\mathcal{F}} \ln \left(\frac{[A_{ox}][B_{red}]}{[A_{red}][B_{ox}]} \right)$$

Problem:

Determine the standard free energy change for the oxidation of NADH + H^+ by O_2 .

$$1/2 O_2 + NADH + H^+ \rightarrow H_2O + NAD^+$$

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$$1/2 O_2 + NADH + H^+ \rightarrow H_2O + NAD^+$$

$$\Delta G^{\circ} = -n \mathcal{F} \Delta E^{\circ}$$

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Fumarate + $2 \text{ H}^{\oplus} + 2e^{\bigcirc} \rightarrow \text{Succinate}$	0.03		
Ubiquinone (Q) + 2 H $^{\oplus}$ + 2 e^{\ominus} \rightarrow QH ₂	0.04		
Cytochrome b (mitochondrial), $F_e^{\stackrel{\textcircled{3}}{\ominus}} + e^{\bigcirc} \rightarrow F_e^{\stackrel{\textcircled{2}}{\ominus}}$	0.08		
Cytochrome c_1 , Fe ^{3\oplus} + $e^{\ominus} \rightarrow$ Fe ^{2\oplus}	0.22		
Cytochrome c , $Fe^{\bigcirc} + e^{\bigcirc} \rightarrow Fe^{\bigcirc}$	0.23		
Cytochrome a , $F_e^{\bigoplus} + e^{\bigoplus} \rightarrow F_e^{\bigoplus}$	0.29		
Cytochrome f , Fe ³⁺ + $e^{\bigcirc} \rightarrow$ Fe ²⁺	0.36		
Plastocyanin, $Cu^{2+} + e^{\Theta} \rightarrow Cu^{+}$	0.37		
$NO_3^{\ominus} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow NO_2^{\ominus} + H_2O$	0.42		
Photosystem I (P700)	0.43		
$F_e^{\oplus} + e^{\ominus} \rightarrow F_e^{\oplus}$	0.77		
$^{1}/_{2}O_{2} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow H_{2}O$	0.82		
Photosystem II (P680)	1.1		

Pr De	TABLE 10.4 Standard reduction potentials of some important biological half-reactions		
of.	Reduction half-reaction	$E^{\circ \prime}$ (V)	
	Acetyl CoA + CO ₂ + H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow Pyruvate + CoA	-0.48	
	Ferredoxin (spinach), $Fe^{\bigcirc} + e^{\bigcirc} \rightarrow Fe^{\bigcirc}$	-0.43	
	$2 \text{ H}^{\oplus} + 2e^{\ominus} \rightarrow \text{H}_2 \text{ (at pH 7.0)}$	-0.42	
	α -Ketoglutarate + CO ₂ + 2 H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow Isocitrate	-0.38	
	Lipoyl dehydrogenase (FAD) + 2 H $^{\oplus}$ + 2 e^{\ominus} \rightarrow Lipoyl dehydrogenase (FADH ₂)	-0.34	
	$NADP^{\oplus} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow NADPH + H^{\oplus}$	-0.32	
	$NAD^{\oplus} + 2 H^{\oplus} + 2e^{\ominus} \rightarrow NADH + H^{\oplus}$	-0.32	
	Lipoic acid $+ 2 H^{\oplus} + 2e^{\ominus} \rightarrow$ Dihydrolipoic acid	-0.29	
	Glutathione (oxidized) + 2 H $^{\oplus}$ + 2 e^{\ominus} \rightarrow 2 Glutathione (reduced)	-0.23	
	$FAD + 2 H^{\oplus} + 2e^{\ominus} \rightarrow FADH_2$	-0.22	
	$FMN + 2 H^{\oplus} + 2e^{\ominus} \rightarrow FMNH_2$	-0.22	
	Acetaldehyde + 2 H $^{\oplus}$ + 2 e^{\bigcirc} \rightarrow Ethanol	-0.20	
	Pyruvate $+ 2 H^{\oplus} + 2e^{\ominus} \rightarrow \text{Lactate}$	-0.18	
42	Oxaloacetate + $2 \text{ H}^{\oplus} + 2e^{\bigcirc} \rightarrow \text{Malate}$	-0.17	

Problem:

Determine the standard free energy change for the oxidation of NADH + H^+ by O_2 .

$$1/2 O_2 + NADH + H^+ \rightarrow H_2O + NAD^+$$

$$\Delta G^{\circ} = -n \mathcal{F} \Delta E^{\circ}$$

·Like Δ G, the observed change in the reduction potential for a reaction, (Δ E), can be determined relative to the change in the standard reduction potential, (Δ E°):

$$\Delta E = \Delta E^{o'} - \frac{RT}{n\mathcal{F}} \ln \left(\frac{[A_{ox}][B_{red}]}{[A_{red}][B_{ox}]} \right)$$

Problem:

Determine the maximum number of ATP's that could be synthesized from ADP and P_i if coupled to the oxidation of NADH + H⁺ by O_2 .

$$1/2~O_2~+~NADH~+~H^+~\rightarrow~H_2O~+~NAD^+$$

$$ADP~+~P_i~\rightarrow~ATP~+~H_2O$$

Problem:

Determine the max synthesized from A NADH + H⁺ by O₂.

 $1/2 O_2 + N$

TABLE 10.3 Standard Gibbs free energies of hydrolysis for common metabolites

	$\Delta G^{\circ}{}'_{ m hydrolysis}$
Metabolite	(kJ mol ⁻¹)
Phosphoenolpyruvate	-62
1,3-Bisphosphoglycerate	-49
ATP to AMP $+$ PP _i	-45
Phosphocreatine	-43
Phosphoarginine	-32
Acetyl CoA	-32
ATP to ADP $+ P_i$	-32
Pyrophosphate	-29
Glucose 1-phosphate	-21
Glucose 6-phosphate	-14
Glycerol 3-phosphate	-9

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Problem:

Determine the maximum number of ATP's that could be synthesized from ADP and P_i if coupled to the oxidation of NADH + H⁺ by O_2 .

$$1/2~O_2~+~NADH~+~H^+~\rightarrow~H_2O~+~NAD^+$$

$$ADP~+~P_i~\rightarrow~ATP~+~H_2O$$

Next Up

·Lecture 8 - Carbohydrate Metabolism

+ Part I: Glycolysis (Moran et al., Chapter 11)