

## Review

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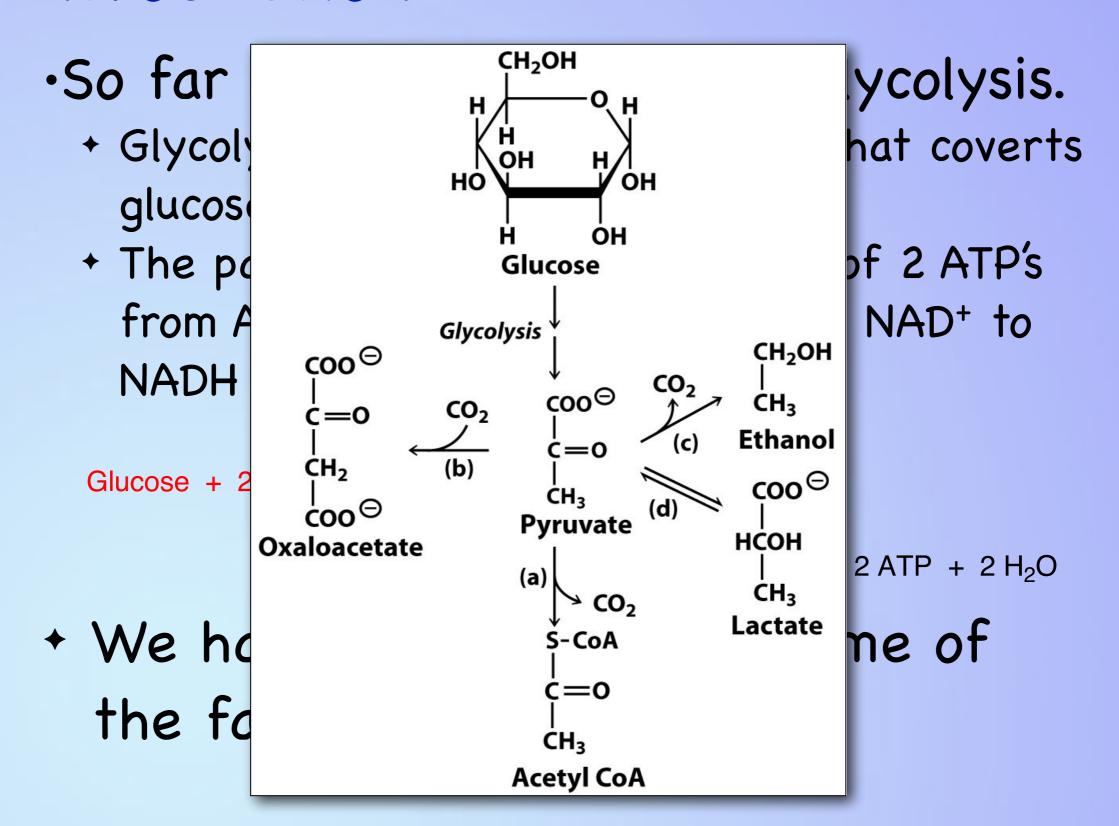
- A. What is the net reaction equation for glycolysis?
- B. What is the metabolic purpose for glycolysis?
- C. Under anaerobic conditions, what are the options for reoxidizing the reduced NADH + H+?

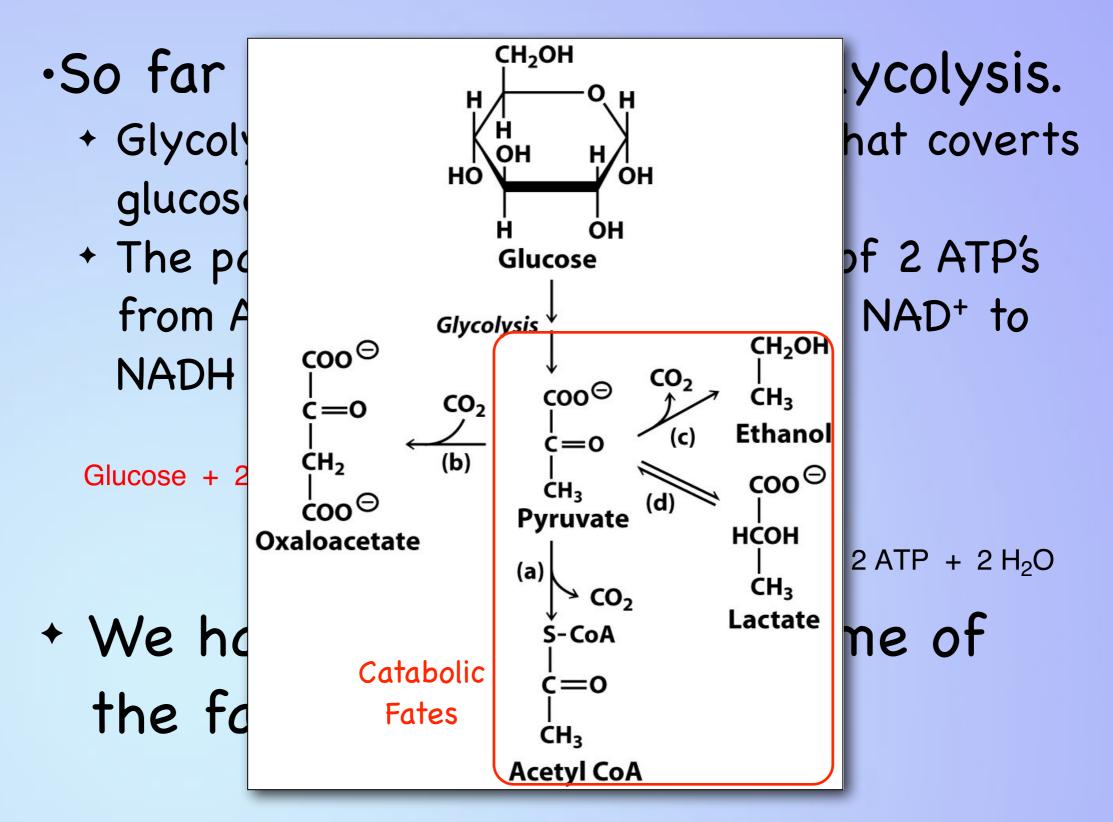
- ·So far we have focused on glycolysis.
  - + Glycolysis is a catabolic pathway that coverts glucose to pyruvate.
  - + The pathway also produces a net of 2 ATP's from ADP and P<sub>i</sub>, plus it reduces 2 NAD+ to NADH + H+.

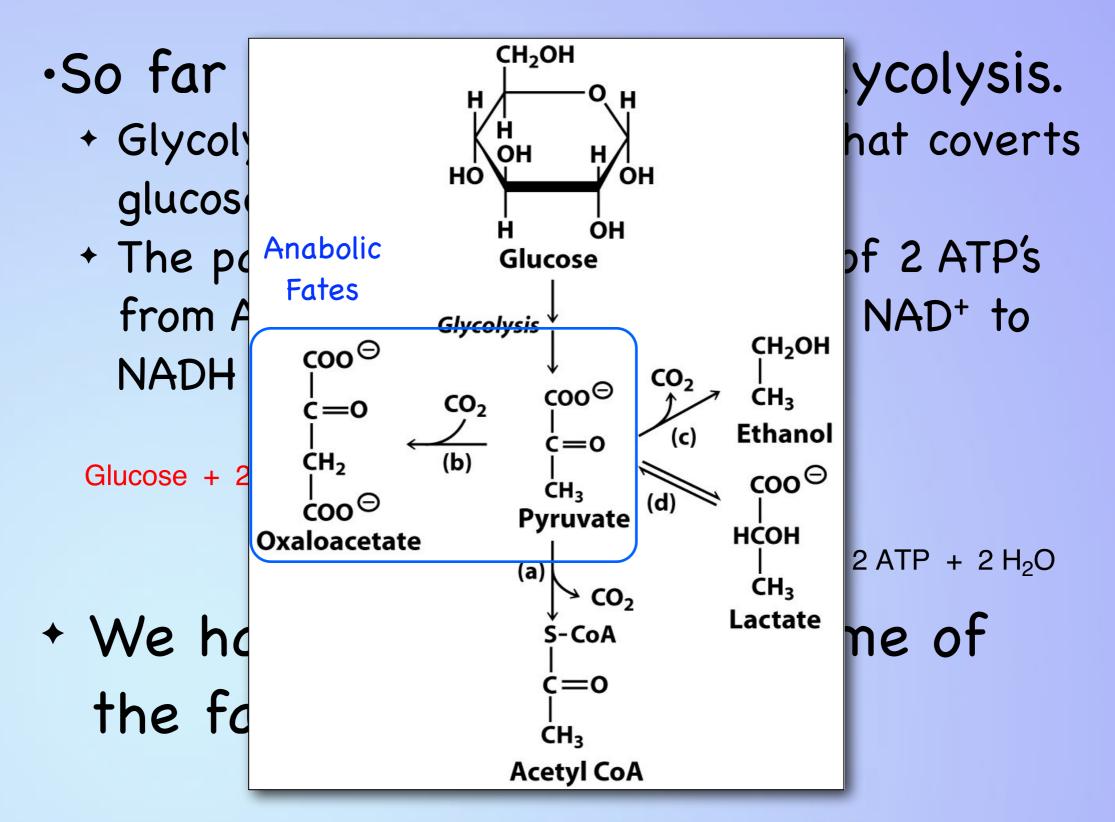
Glucose + 
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 +  $2 \text{ ADP}$  +  $2 \text{ P}_i$ 

2 Pyruvate + 2 NADH + 
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 + 2 ATP + 2  $H_2O$ 

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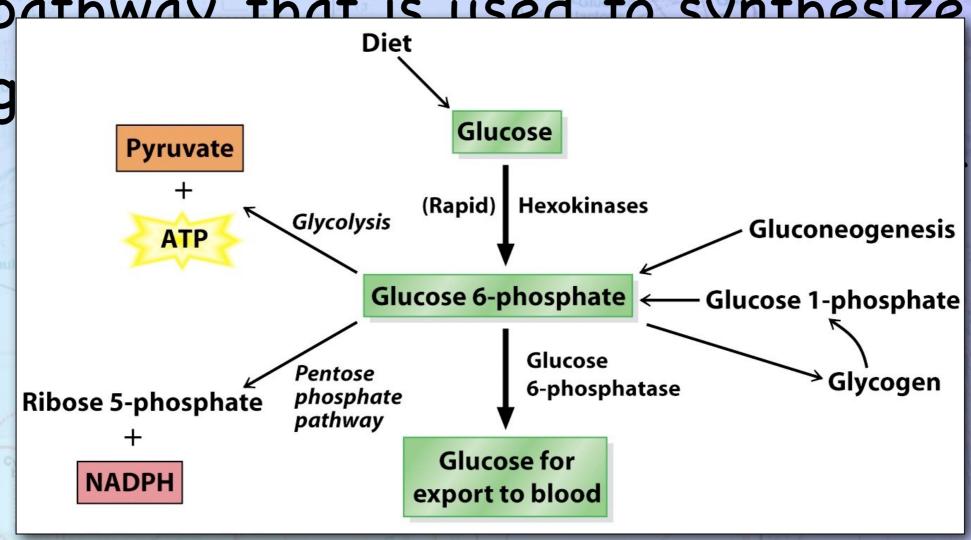
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 We have also considered some of the fates for pyruvate.

- ·Next we will look at an anabolic pathway that is used to synthesize glucose from smaller molecules.
  - + We will also look at the various fates for glucose.
    - Release into the bloodstream
    - Conversion to pentoses along with the production of reduced NADPH + H+ for biosynthetic pathways
    - · Storage as glycogen or starch

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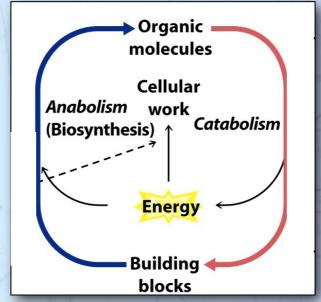


· Storage as glycogen or starch

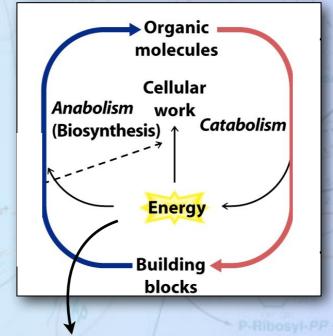
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  - + Catabolism versus Anabolism
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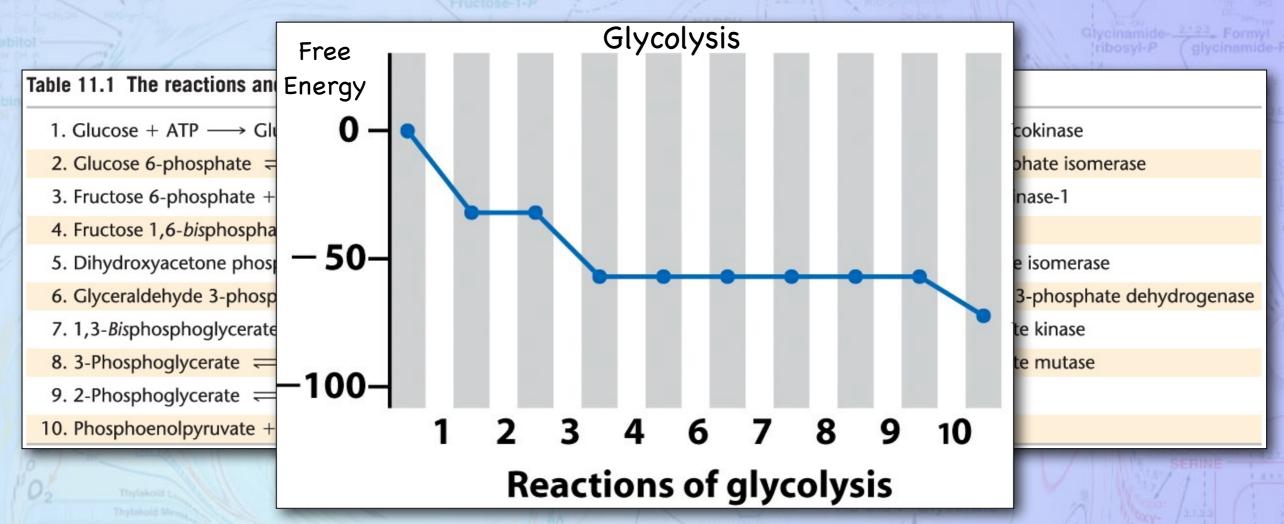
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  - \* Many, but not all of the reactions used in glycolysis are also used in gluconeogenesis.

## ·Gluconeogenesis means "new glucose"

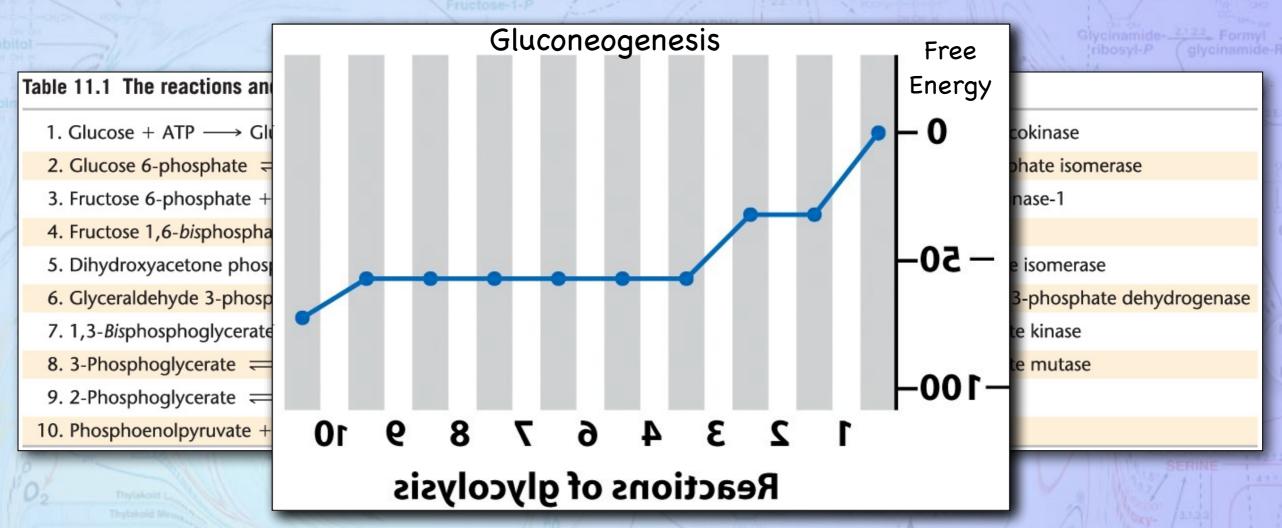
+ Many, but not all of the reactions used in glycolysis are also used in gluconeogenesis.

Table 11.1 The reactions and enzymes of glycolysis			
1. Glucose + ATP $\longrightarrow$ Glucose 6-phosphate + ADP + H $^{\oplus}$	Hexokinase, glucokinase		
2. Glucose 6-phosphate Fructose 6-phosphate	Glucose-6-phosphate isomerase		
3. Fructose 6-phosphate + ATP $\longrightarrow$ Fructose 1,6-bisphosphate + ADP + H $^{\oplus}$	Phosphofructokinase-1		
4. Fructose 1,6-bisphosphate   Dihydroxyacetone phosphate + Glyceraldehyde 3-phosphate	Aldolase		
5. Dihydroxyacetone phosphate Clyceraldehyde 3-phosphate	Triose phosphate isomerase		
6. Glyceraldehyde 3-phosphate + $NAD^{\oplus}$ + $P_i \Longrightarrow 1,3$ -Bisphosphoglycerate + $NADH$ + $H^{\oplus}$	Glyceraldehyde 3-phosphate dehydrogenase		
7. 1,3-Bisphosphoglycerate + ADP === 3-Phosphoglycerate + ATP	Phosphoglycerate kinase		
8. 3-Phosphoglycerate  2-Phosphoglycerate	Phosphoglycerate mutase		
9. 2-Phosphoglycerate   → Phosphoenolpyruvate + H₂O	Enolase		
10. Phosphoenolpyruvate + ADP + $H^{\oplus}$ $\longrightarrow$ Pyruvate + ATP	Pyruvate kinase		

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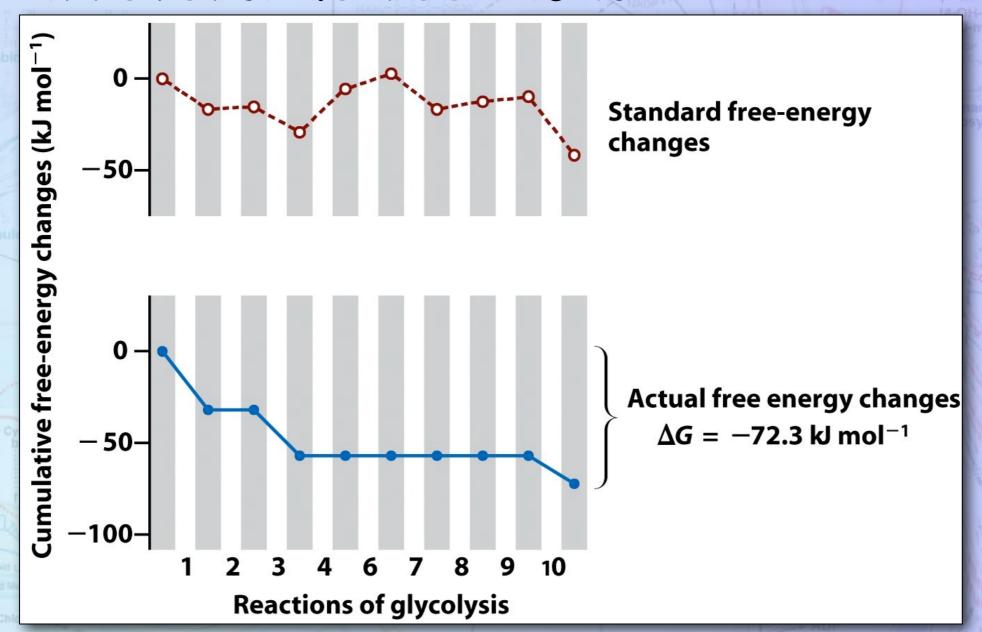
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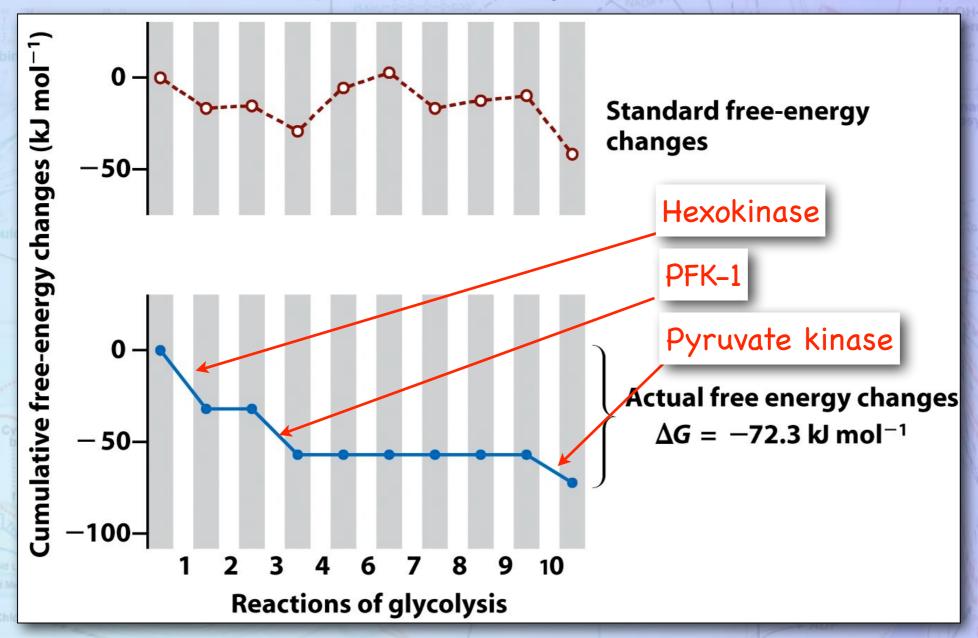
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Step	Reaction	Δ <i>G</i> <sup>01</sup> / (kJ/mol)	ΔG / (kJ/mol)
1	glucose + ATP <sup>4-</sup> → glucose-6-phosphate <sup>2-</sup> + ADP <sup>3-</sup> + H <sup>+</sup>	-16.7	-34
2	glucose-6-phosphate <sup>2-</sup> → fructose-6-phosphate <sup>2-</sup>	1.67	-2.9
3	fructose-6-phosphate2- + ATP4- → fructose-1,6-bisphosphate4- + ADP3- + H+	-14.2	-19
4	fructose-1,6-bisphosphate <sup>4-</sup> → dihydroxyacetone phosphate <sup>2-</sup> + glyceraldehyde-3- phosphate <sup>2-</sup>	23.9	-0.23
5	dihydroxyacetone phosphate <sup>2-</sup> → glyceraldehyde-3-phosphate <sup>2-</sup>	7.56	2.4
6	glyceraldehyde-3-phosphate2- + Pi2- + NAD+ → 1,3-bisphosphoglycerate4- + NADH + H+	6.30	-1.29
7	1,3-bisphosphoglycerate <sup>4-</sup> + ADP <sup>3-</sup> → 3-phosphoglycerate <sup>3-</sup> + ATP <sup>4-</sup>	-18.9	0.09
В	3-phosphoglycerate <sup>3-</sup> → 2-phosphoglycerate <sup>3-</sup>	4.4	0.83
9	2-phosphoglycerate <sup>3-</sup> → phosphoenolpyruvate <sup>3-</sup> + H <sub>2</sub> O	1.8	1.1
10	phosphoenolpyruvate <sup>3-</sup> + ADP <sup>3-</sup> + H <sup>+</sup> → pyruvate <sup>-</sup> + ATP <sup>4-</sup>	-31.7	-23.0

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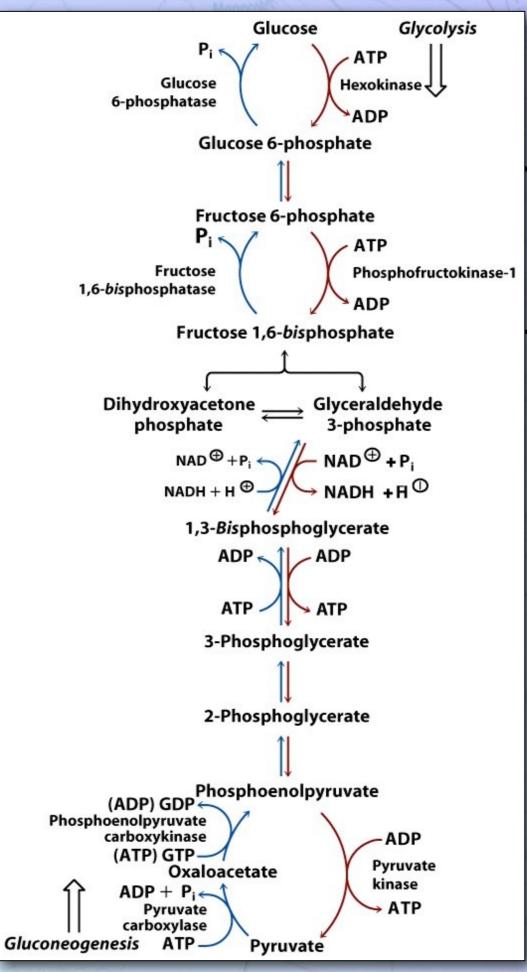
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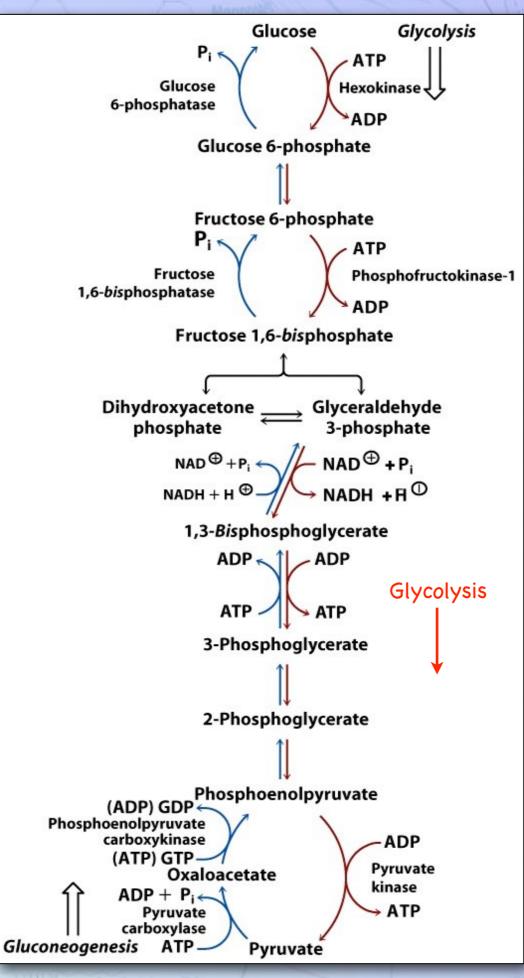


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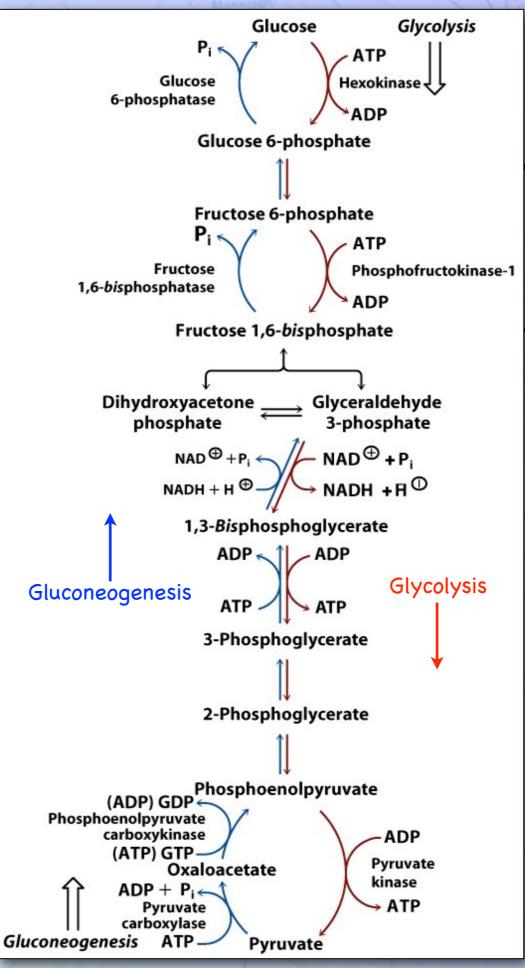


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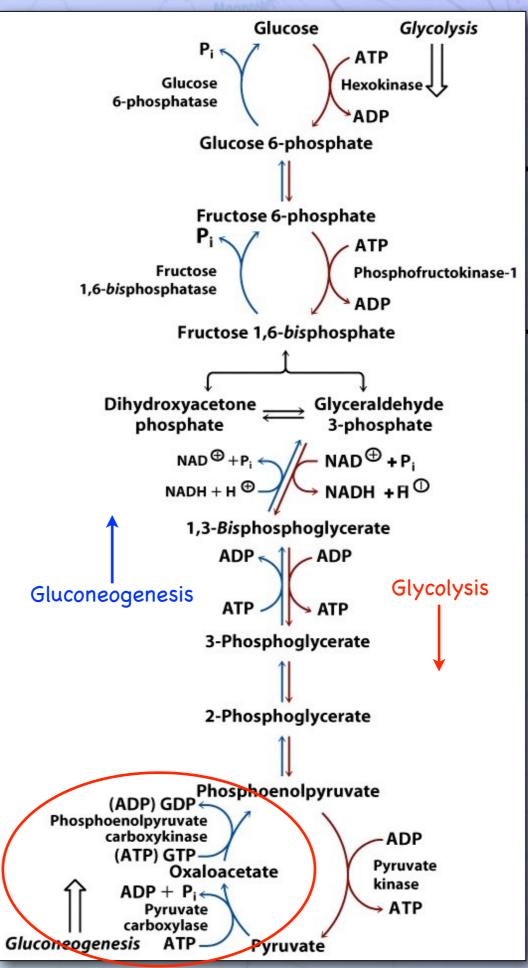


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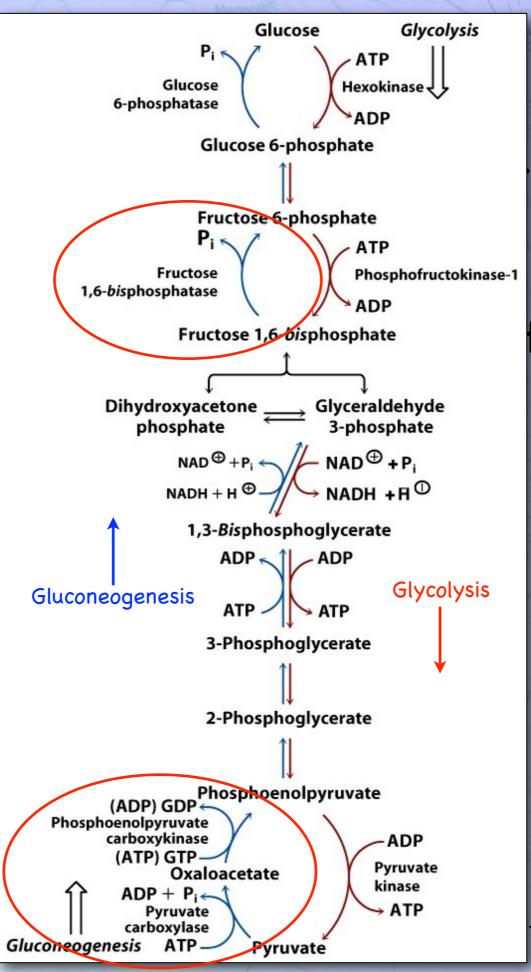


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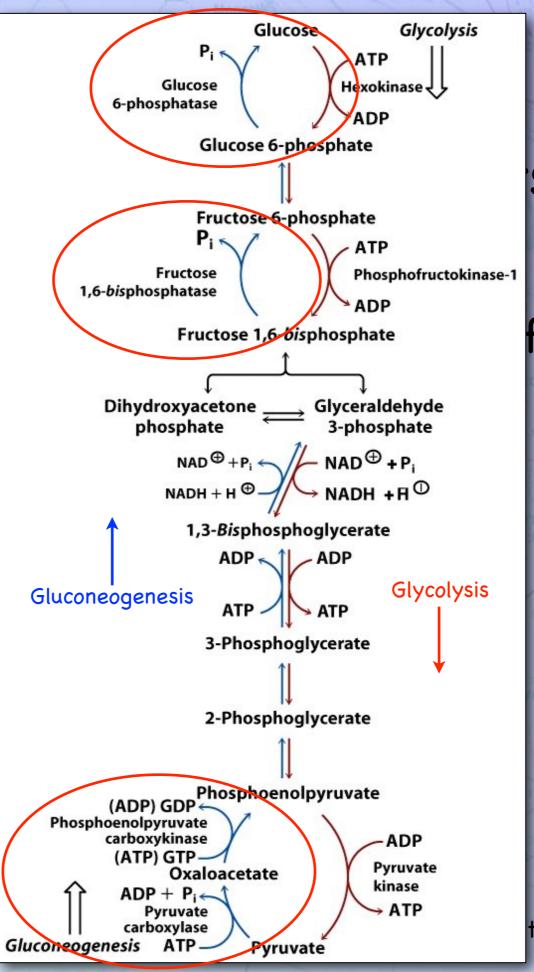


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- ·Starting at pyruvate
  - The pyruvate kinase reaction is replaced with two reactions
    - Pyruvate carboxylase
    - · Phosphoenolpyruvate carboxykinase

### ·Pyruvate carboxylase

COO
$$^{\bigcirc}$$
| C = 0 + HCO $_3^{\bigcirc}$ 
| CH 3 Bicarbonate ATP ADP + P<sub>i</sub>
| CH 2 | COO $^{\bigcirc}$ 
| COO $^{\bigcirc}$ 
| CH 2 | COO $^{\bigcirc}$ 
| CH 2 | COO $^{\bigcirc}$ 
| COO $^{\bigcirc}$ 
| CH 2 | COO $^{\bigcirc}$ 

TABLE 7.2Major coenzymes

Coenzyme	Vitamin source	Major metabolic roles	Mechanistic role
Adenosine triphosphate (ATP)	_	Transfer of phosphoryl or nucleotidyl groups	Cosubstrate
S-Adenosylmethionine	_	Transfer of methyl groups	Cosubstrate
Uridine diphosphate glucose	_	Transfer of glycosyl groups	Cosubstrate
Nicotinamide adenine dinucleotide (NAD $^{\oplus}$ ) and nicotinamide adenine dinucleotide phosphate (NADP $^{\oplus}$ )	Niacin	Oxidation-reduction reactions involving two-electron transfers	Cosubstrate
Flavin mononucleotide (FMN) and flavin adenine dinucleotide (FAD)	Riboflavin (B <sub>2</sub> )	Oxidation-reduction reactions involving one- and two-electron transfers	Prosthetic group
Coenzyme A (CoA)	Pantothenate (B <sub>3</sub> )	Transfer of acyl groups	Cosubstrate
Thiamine pyrophosphate (TPP)	Thiamine $(B_1)$	Transfer of two-carbon fragments containing a carbonyl group	Prosthetic group
Pyridoxal phosphate (PLP)	Pyridoxine (B <sub>6</sub> )	Transfer of groups to and from amino acids	Prosthetic group
Biotin	Biotin	ATP-dependent carboxylation of substrates or carboxyl-group transfer between substrates	Prosthetic group
Tetrahydrofolate	Folate	Transfer of one-carbon substituents, especially formyl and hydroxymethyl groups; provides the methyl group for thymine in DNA	Cosubstrate
Adenosylcobalamin	Cobalamin (B <sub>12</sub> )	Intramolecular rearrangements	Prosthetic group
Methylcobalamin	Cobalamin (B <sub>12</sub> )	Transfer of methyl groups	Prosthetic group
Lipoamide	_	Oxidation of a hydroxyalkyl group from TPP and subsequent transfer as an acyl group	Prosthetic group
Retinal	Vitamin A	Vision	Prosthetic group
Vitamin K	Vitamin K	Carboxylation of some glutamate residues	Prosthetic group
Ubiquinone (Q)	-	Lipid-soluble electron carrier	Cosubstrate

### ·Pyruvate carboxylase

COO
$$^{\bigcirc}$$

C=0 + HCO $_3^{\bigcirc}$ 

Pyruvate carboxylase

CH $_3$ 

Bicarbonate

ATP

ADP +Pi

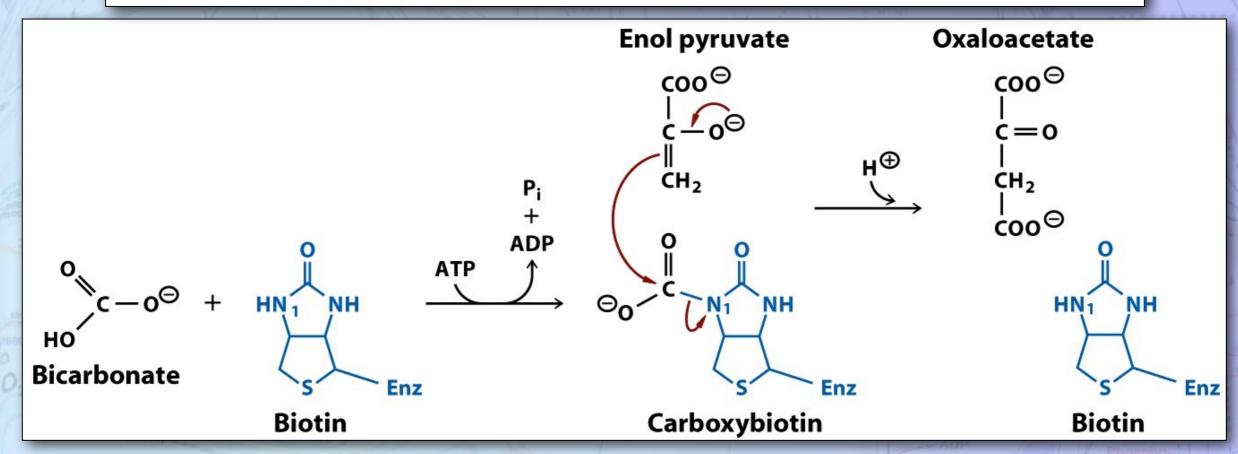
CH $_2$ 

COO $^{\bigcirc}$ 

CH $_2$ 

COO $^{\bigcirc}$ 

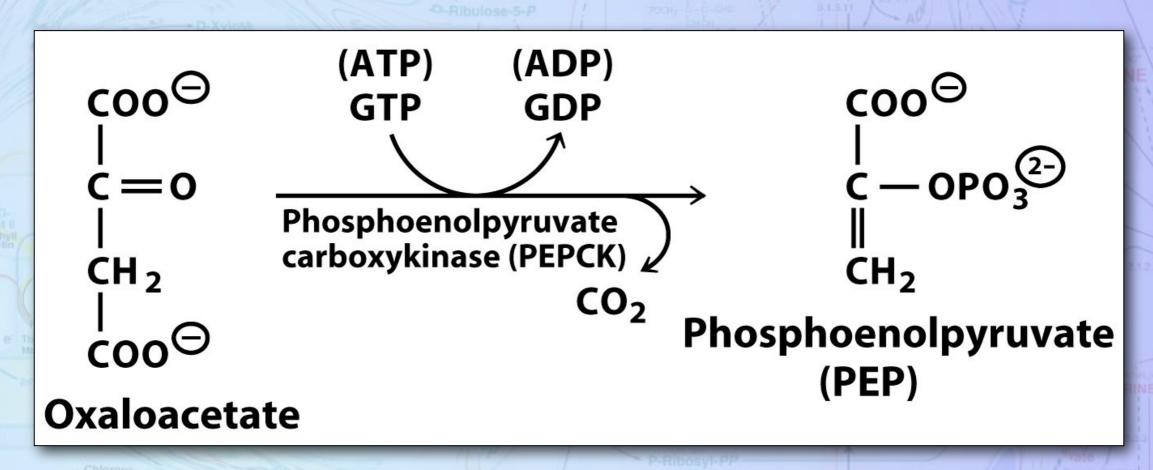
Oxaloacetate



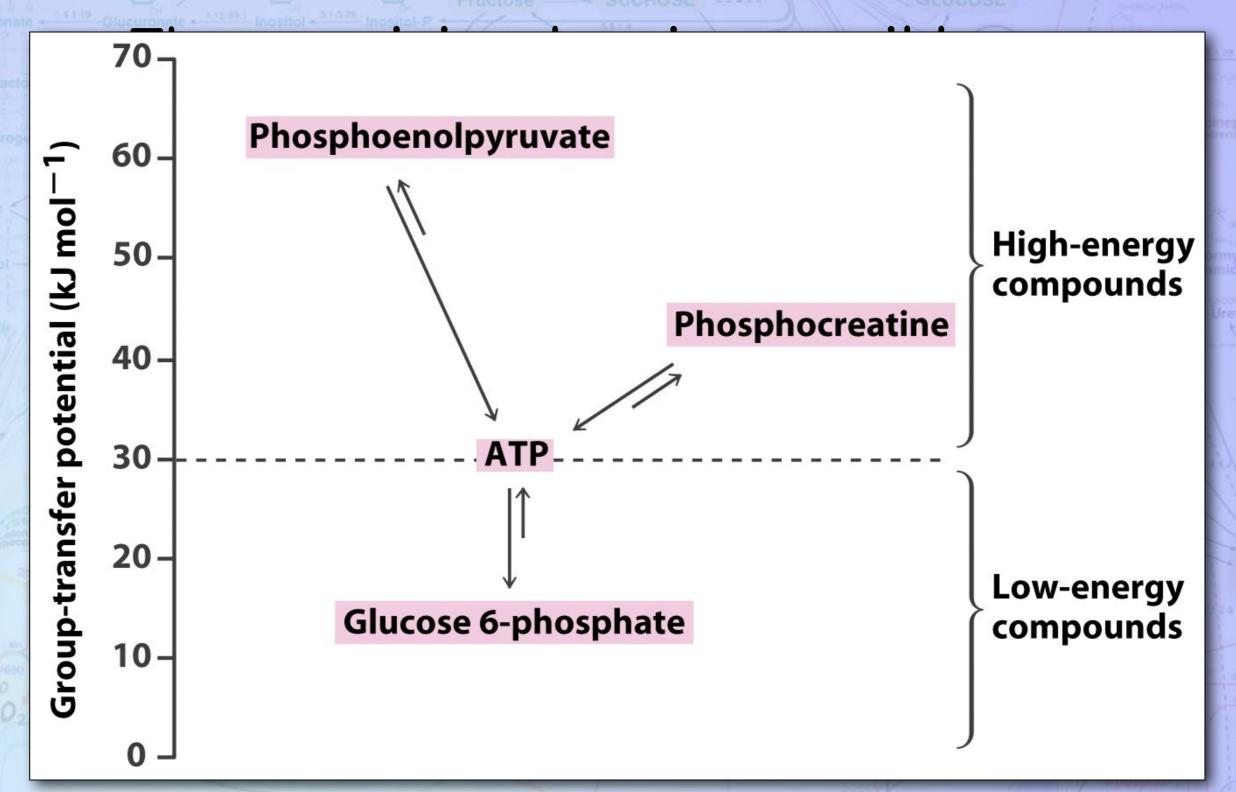
### ·Pyruvate carboxylase

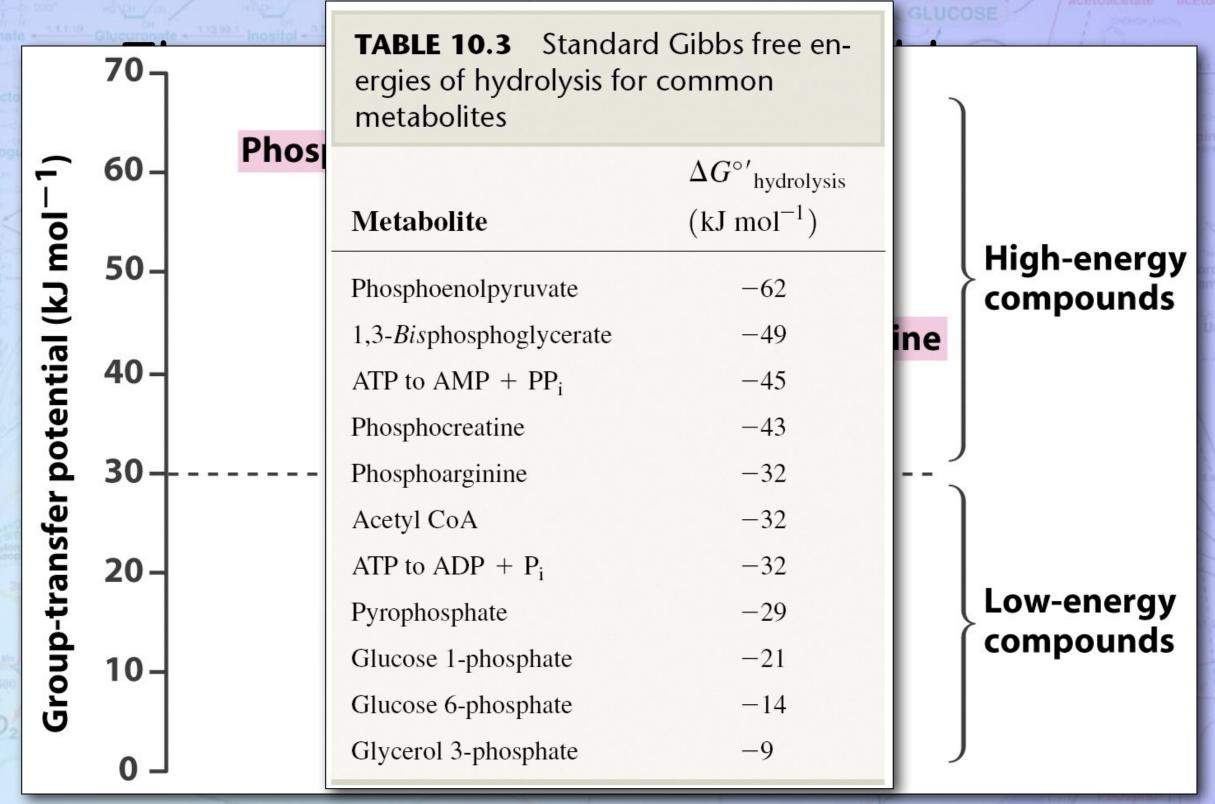
- + The oxaloacetate that is produced in this reaction is a citric acid cycle intermediate.
  - This reaction is also used to increase the quantity of citric acid cycle intermediates
- + Pyruvate carboxylase is stimulated by high Acetyl-CoA levels.
  - High Acetyl-CoA levels signal a need for citric acid cycle intermedates.

- ·Phosphoenolpyruvate carboxykinase
  - + This is a decarboxylation reaction.
    - Decarboxylations have a high negative free energy change (  $\Delta$  G << 0)



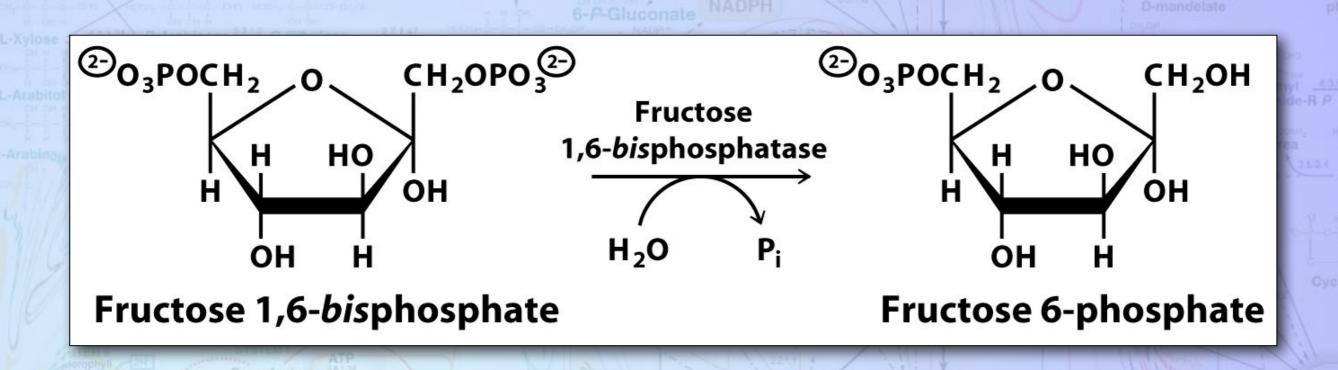
- •The remaining two irreversible reactions from glycolysis which must be circumvented are the two kinase reactions near the beginning of the glycolytic pathway
  - + Both are bypassed using phosphatase reactions, which hydrolyze phosphate esters.



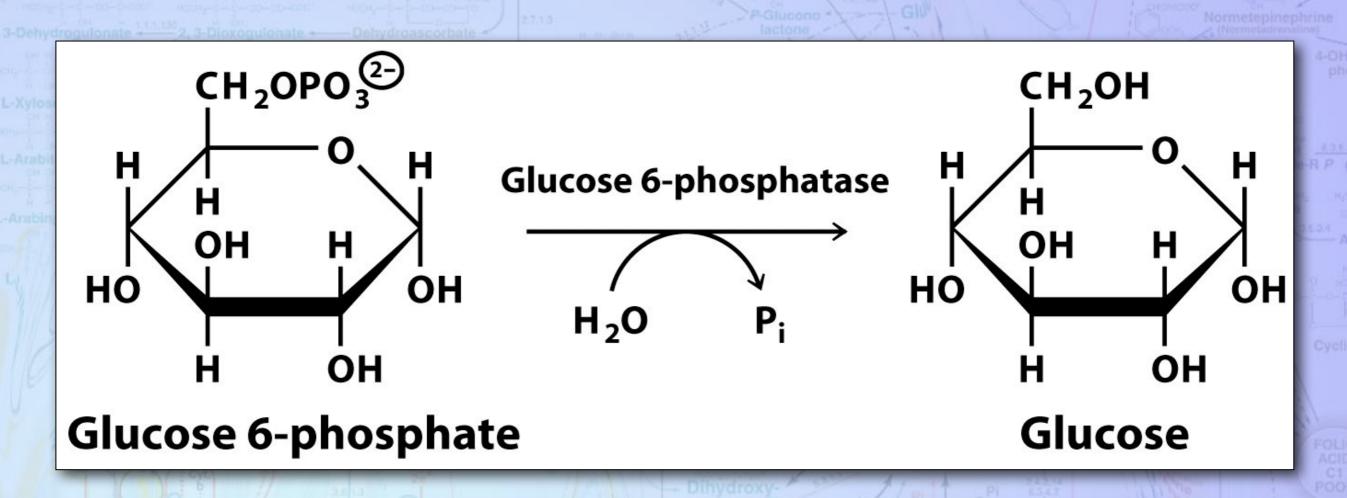


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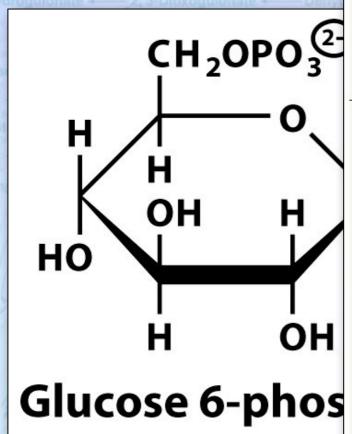
### ·Fructose 1,6-bisphosphatase



### ·Glucose 6-phosphatase

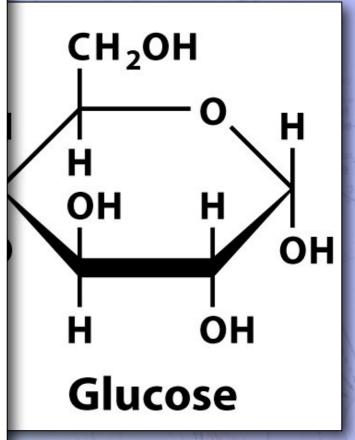


### ·Glucose

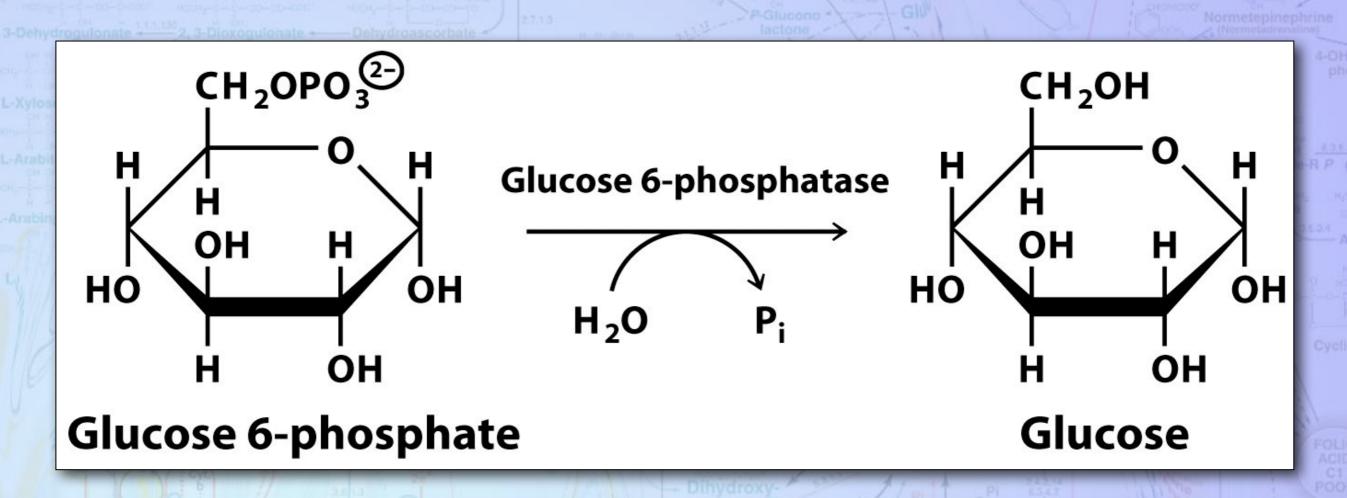


**TABLE 10.3** Standard Gibbs free energies of hydrolysis for common metabolites

	$\Delta G^{\circ}{}'_{ m hydrolysis}$	
Metabolite	$(kJ \text{ mol}^{-1})$	
Phosphoenolpyruvate	-62	
1,3-Bisphosphoglycerate	-49	
ATP to AMP $+$ PP <sub>i</sub>	-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	

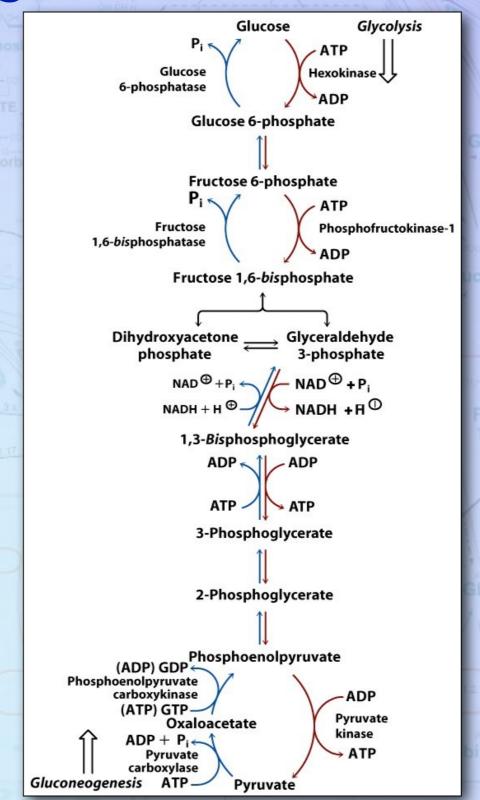


### ·Glucose 6-phosphatase

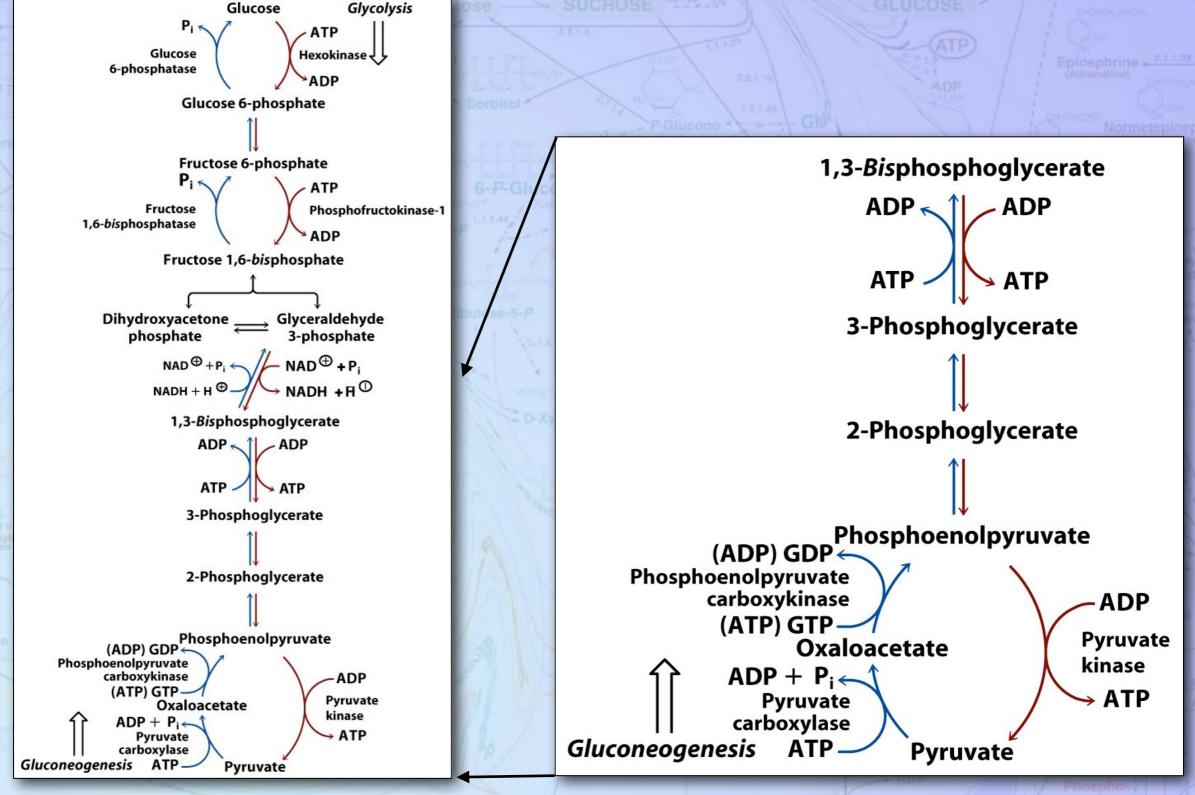


### ·Glucose 6-phosphatase

+ Only the liver, the kidneys and small intestine are capable of carrying out this reaction, for the purpose of increasing the blood glucose levels



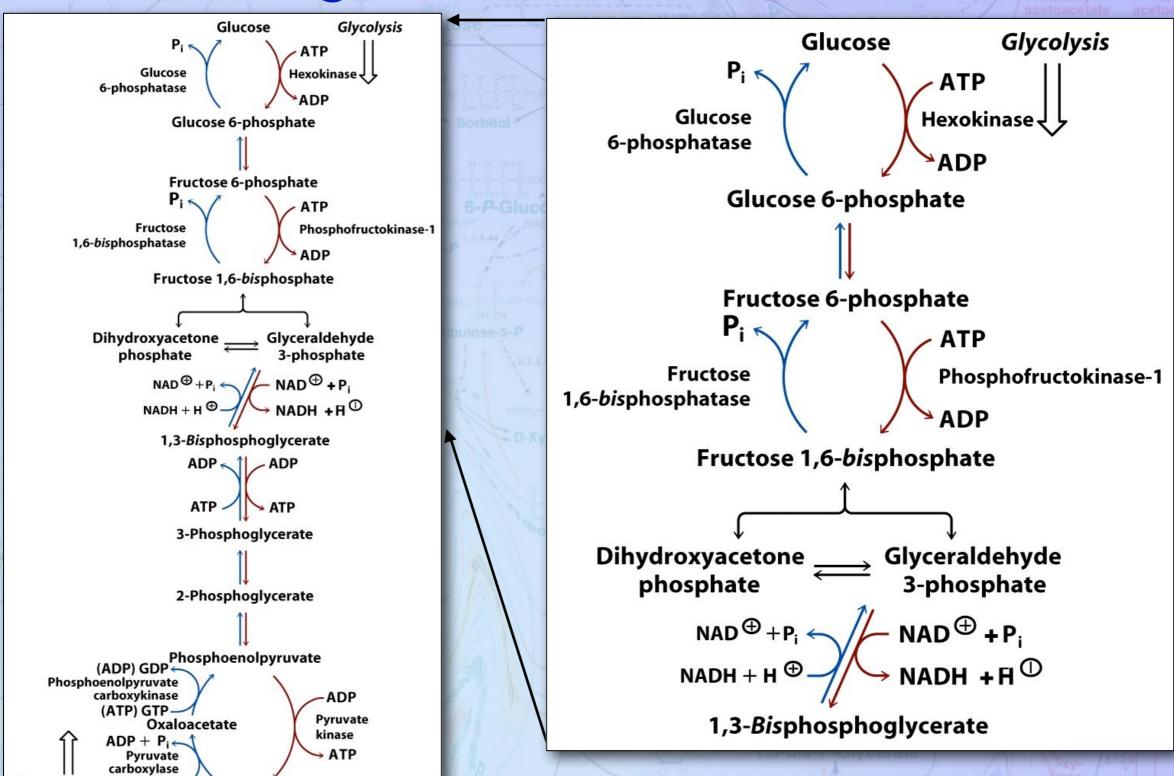
Glucose



Gluconeogenesis

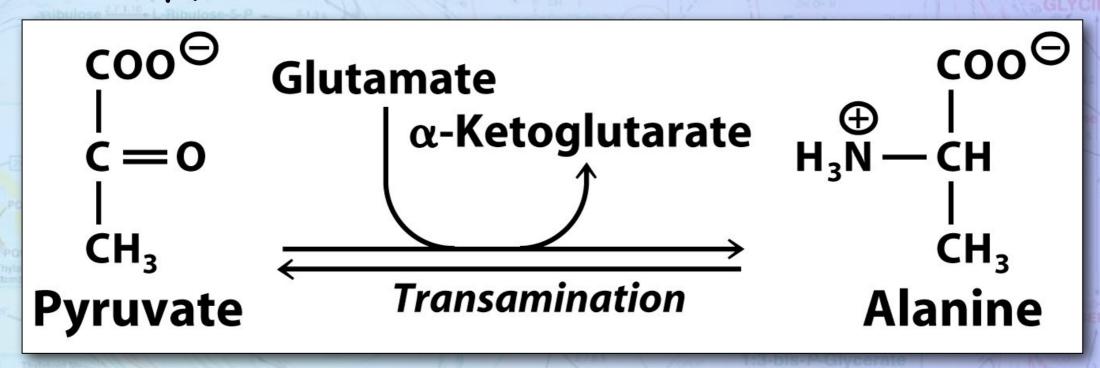
ATP.

Pyruvate

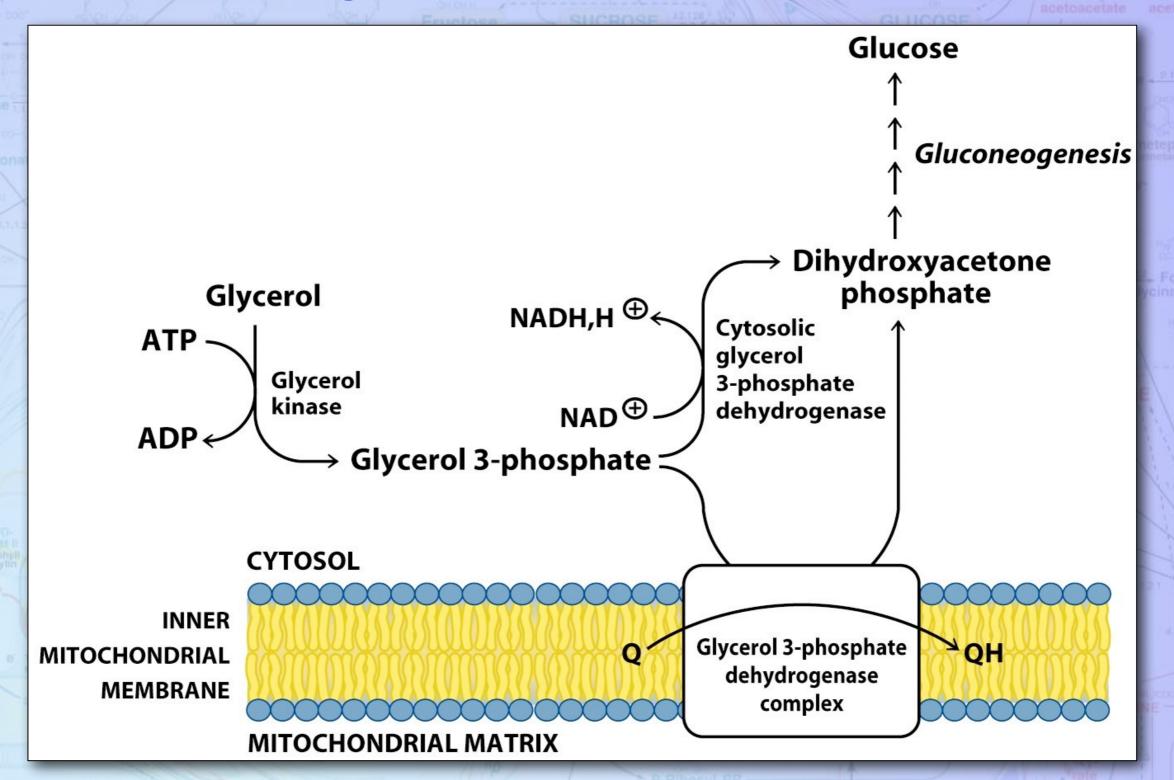


- ·Other molecules can serve as starting material for gluconeogenesis.
  - + Amino Acids
    - Transamination of aspartic acid produces oxaloacetate

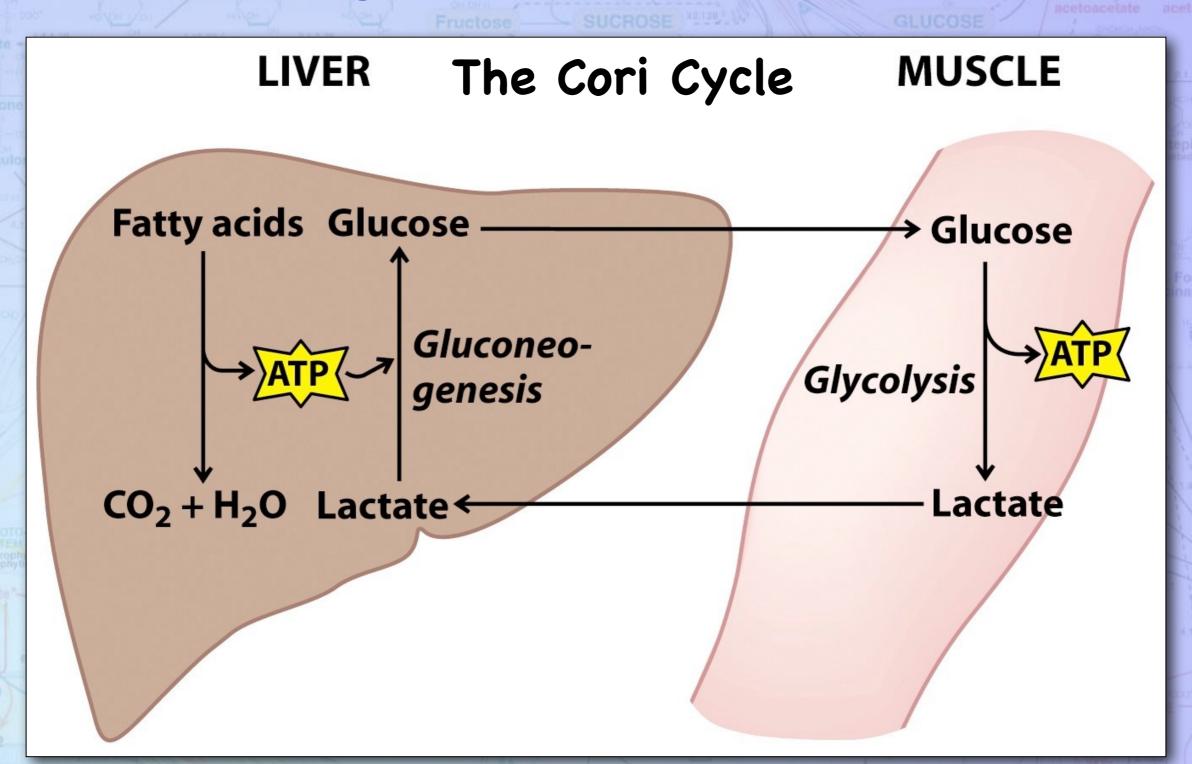
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    - Transamination of alanine produces pyruvate

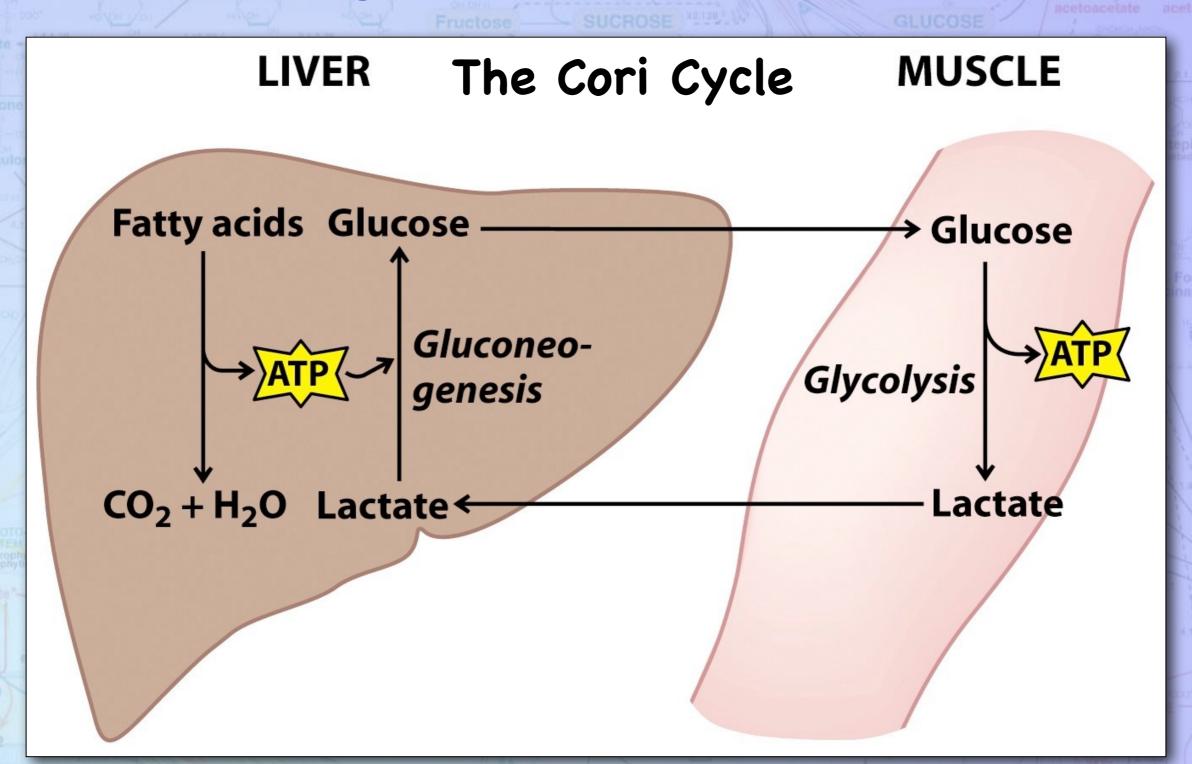


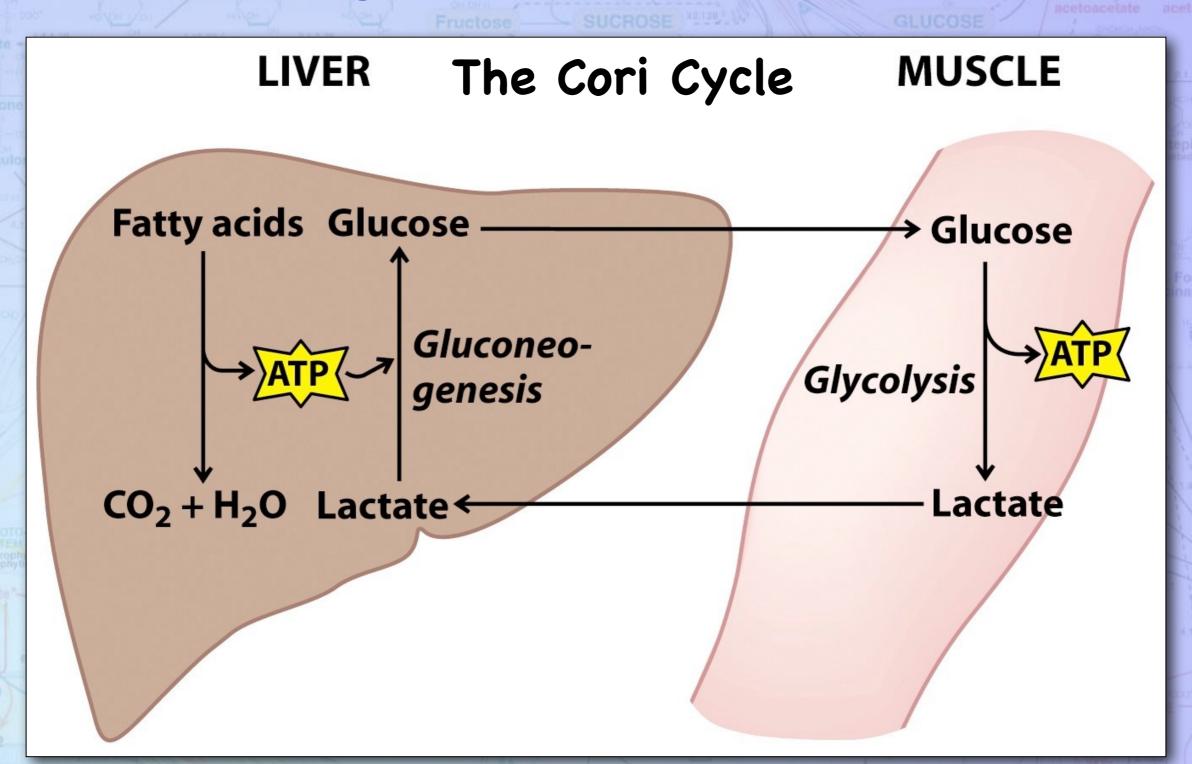
- ·Other molecules can serve as starting material for gluconeogenesis
  - + Fats are broken down to produce Acetyl-CoA and glycerol.
    - Glycerol can be converted to dihydroxyacetone phosphate
    - Acetyl-CoA in some bacteria, plants an fungi (but not humans) can convert acetyl-CoA into oxaloacetate



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  - + Fats are broken down to produce Acetyl-CoA and glycerol.
    - Glycerol can be converted to dihydroxyacetone phosphate
    - Acetyl-CoA in some bacteria, plants an fungi (but not humans) can convert acetyl-CoA into oxaloacetate







### ·Regulation of Gluconeogenesis

#### **Glycolysis**

Glucose + 
$$2 \text{ NAD}^+$$
 +  $2 \text{ ADP}$  +  $2 \text{ P}_i$ 

$$2 \text{ Pyruvate} + 2 \text{ NADH} + \text{H}^+ + 2 \text{ ATP} + 2 \text{ H}_2\text{O}$$

#### Gluconeogenesis

2 Pyruvate + 2 NADH + 
$$H^+$$
 + 6 ATP + 6  $H_2O$ 

Glucose +  $2 \text{ NAD}^+$  + 6 ADP +  $6 \text{ P}_i$ 

#### **Net Reaction**

$$4 \text{ ATP} + 4 \text{ H}_2\text{O} \longrightarrow 4 \text{ ADP} + 4 \text{ P}_i$$

### Gluconeo

# ·Regulation

#### **Glycolysis**

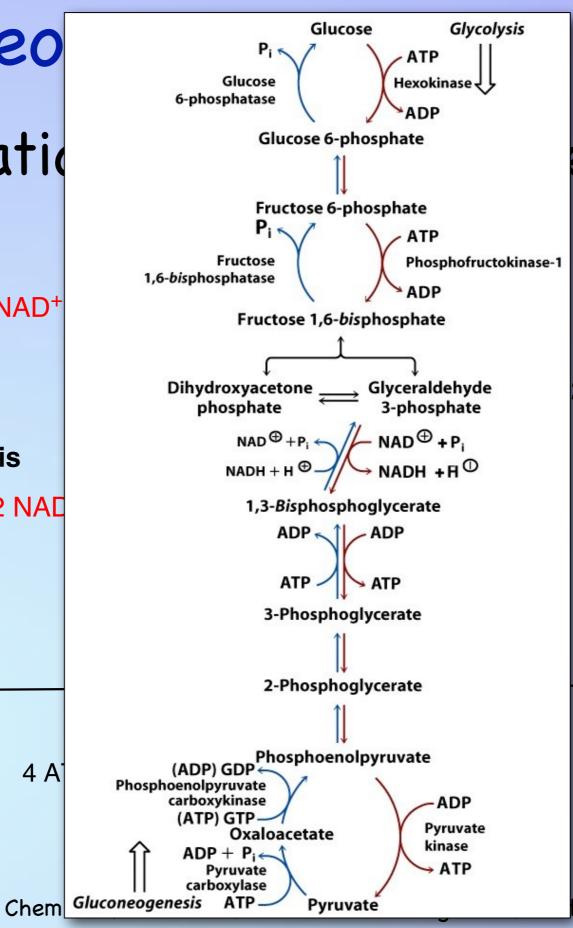
Glucose + 2 NAD+

#### Gluconeogenesis

2 Pyruvate + 2 NAD

#### **Net Reaction**

4 A



Siss

$$2 ATP + 2 H2O$$

 $P_{i}$ 

### ·Regulation of Gluconeogenesis

#### **Glycolysis**

Glucose + 
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 +  $2 \text{ ADP}$  +  $2 \text{ P}_i$ 

$$2 \text{ Pyruvate} + 2 \text{ NADH} + \text{H}^+ + 2 \text{ ATP} + 2 \text{ H}_2\text{O}$$

#### Gluconeogenesis

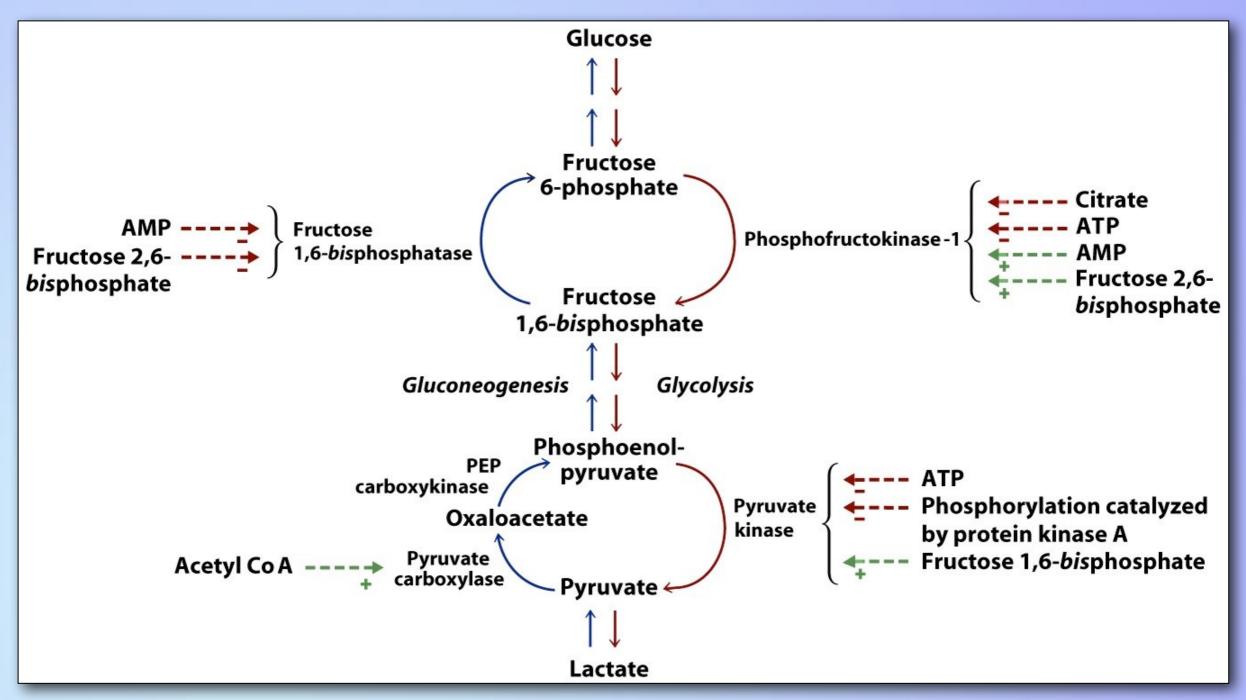
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### ·Regulation of Gluconeogenesis



#### · Deaulation of Gluconeganesis

#### Clicker Question:

Fructose 2,6-bisphosphate coordinately regulates glycolysis and gluconeogenesis. The buildup of this this metabolite is signals which of the following states?

#### Fructose bisphosp

A. There is a buildup of glycolytic intermediates further down stream.

- B. Blood glucose levels are low.
- C. There is a buildup of citric acid cycle intermediates.
- D. ATP levels are low.

E. Blood glucose levels are high.

te

tose 2,6-10sphate

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sphate

Gluconeonesis Glucagon LIVER CELL **EXTRACELLULAR SPACE** Glucagon Click transducer system **CYTOSOL** Fruc and **Fructose** als gluco 6-phosphate whic **ATP** te PFK-2 A. OH activity ADP + Fructose tose 2,6bisphosp **ATP** nosphate **Protein** Fructose **B**. 2,6-bisphosphate kinase A → ADP C. **Fructose** 2,6-bisphosphatase activity D. A talyzed Fructose sphate **Glycolysis** E. 6-phosphate PFK-1

→ Glycolysis

**Fructose** 

1,6-bisphosphate

& Glycogen

#### · Deaulation of Gluconeganesis

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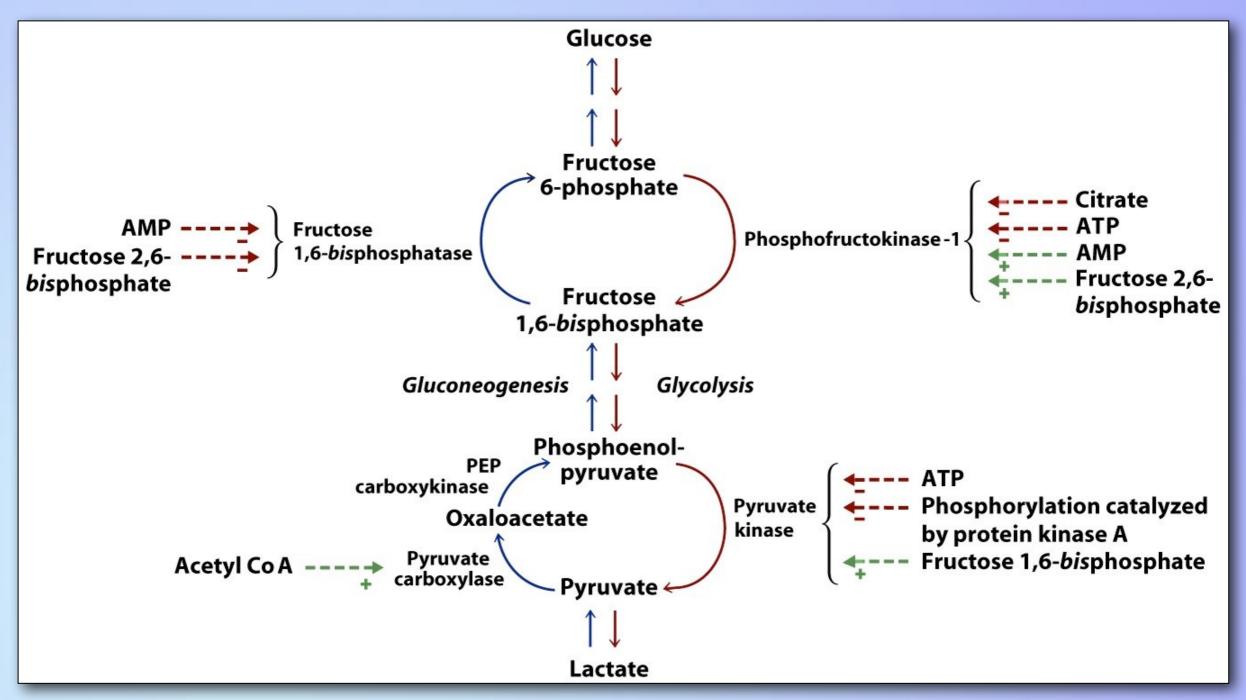
te

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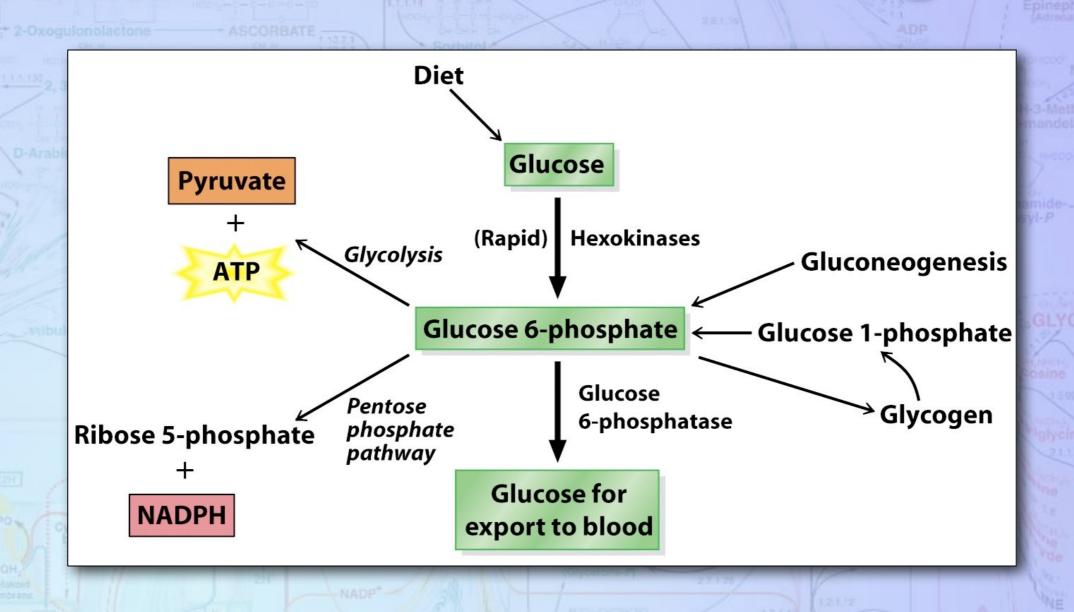
talyzed

sphate

### ·Regulation of Gluconeogenesis



# Fates for Glucose 6-Phosphate

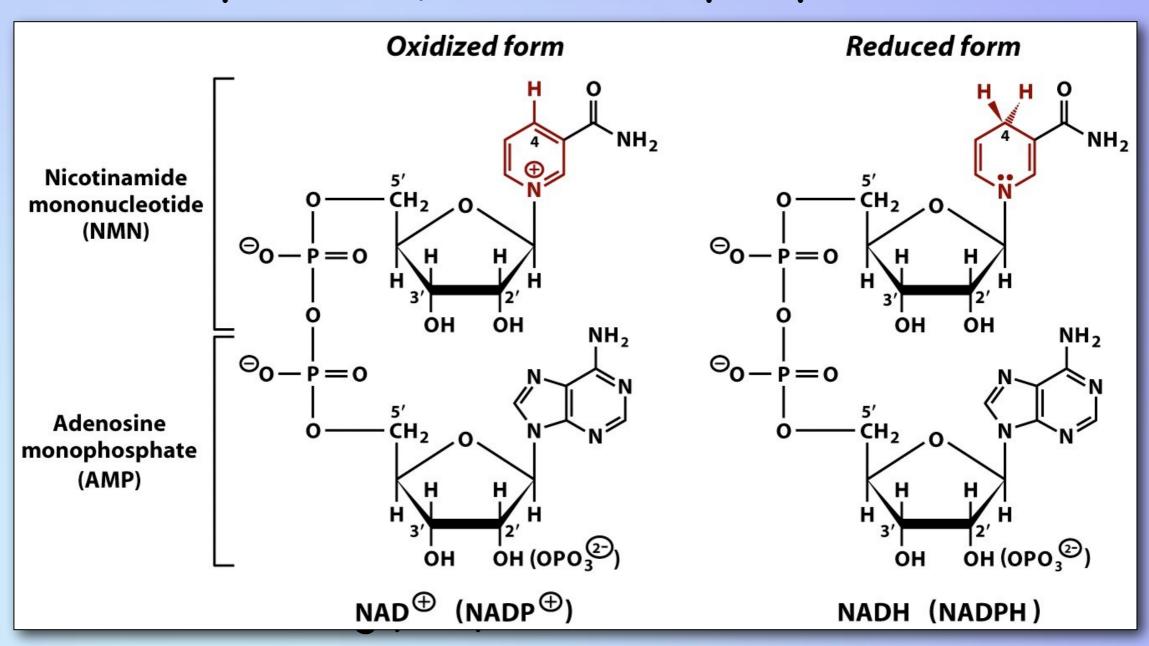


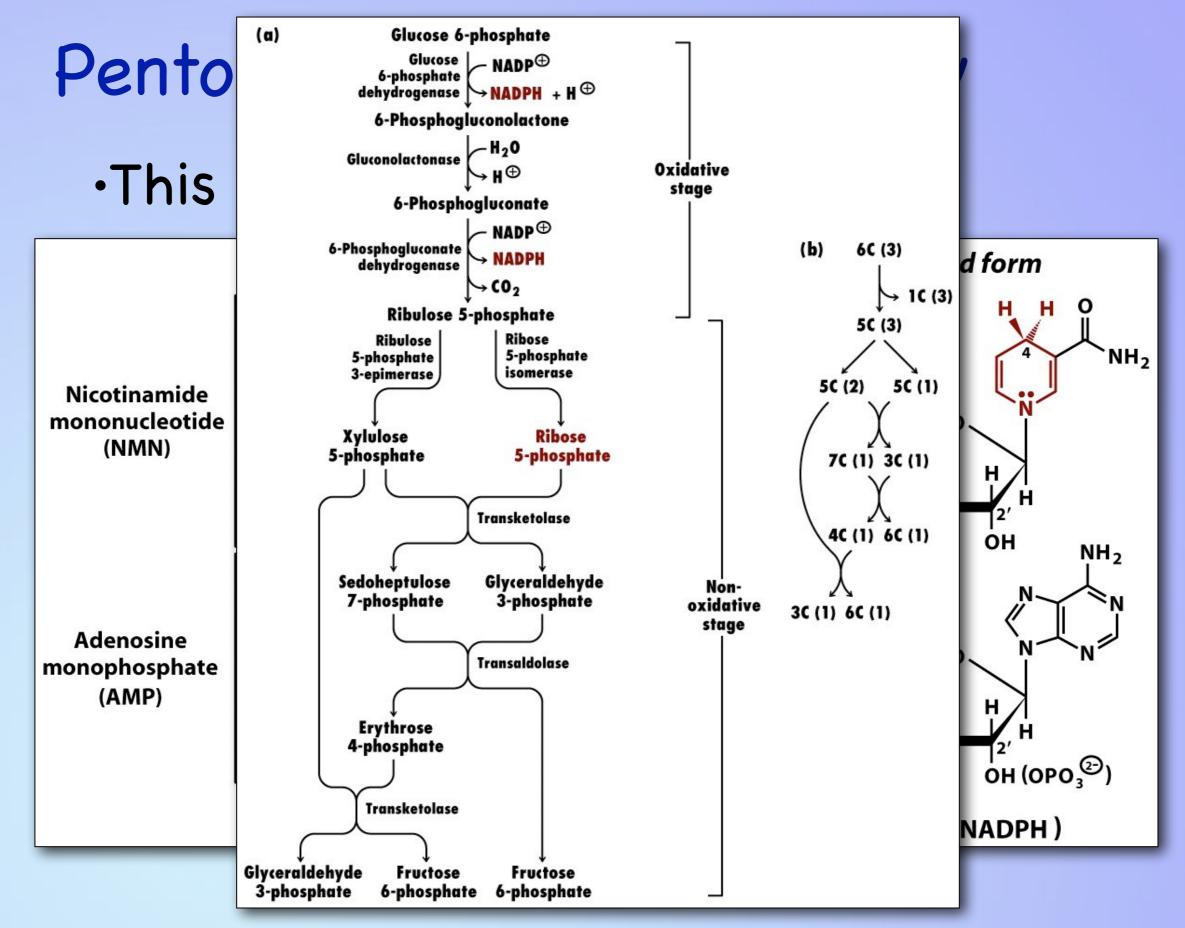
### Pentose Phosphate Pathway

- ·This pathway has two purposes
  - + Produce pentoses for nucleotide biosynthesis
  - \* **Produce reducing power**, in the form of reduced NADPH + H<sup>+</sup>, for biosynthetic reactions.
- ·There are two stages to this pathway
  - \* Oxidative, which produces the pentose and the reduced NADPH + H+.
  - \* Non-oxidative, which converts the pentoses back into glycolytic intermediates.

### Pentose Phosphate Pathway

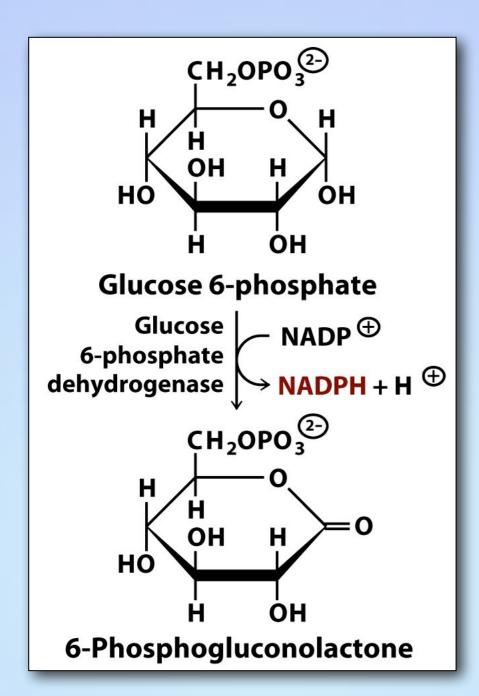
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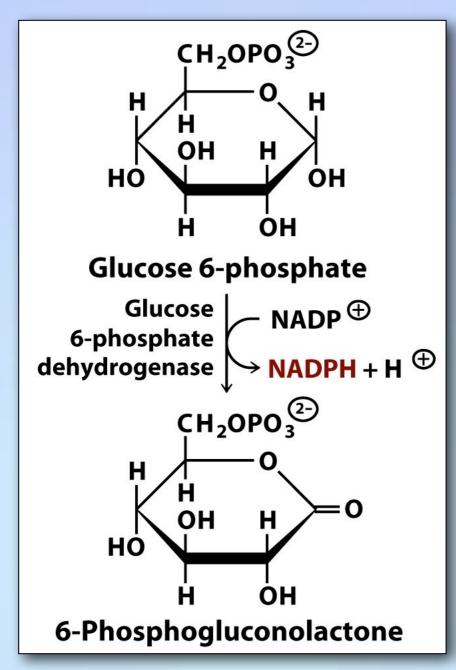


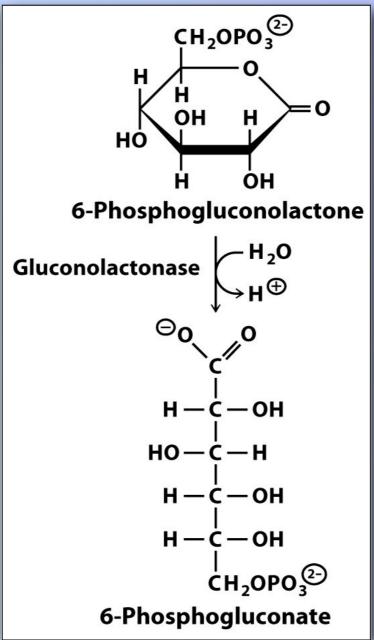


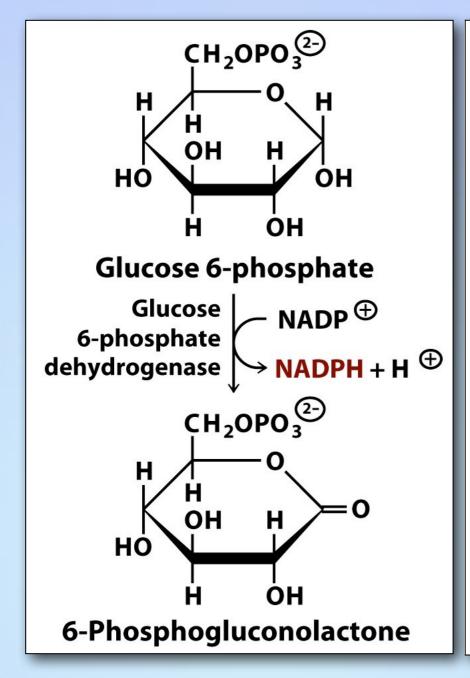
Chem 352, Lecture 8, Part II: Gluconeogenesis, Pentose Phosphate & Glycogen

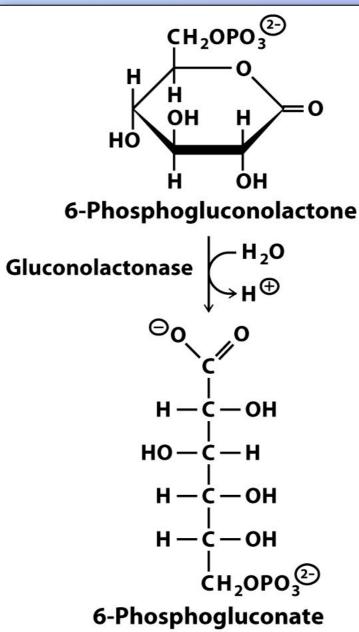
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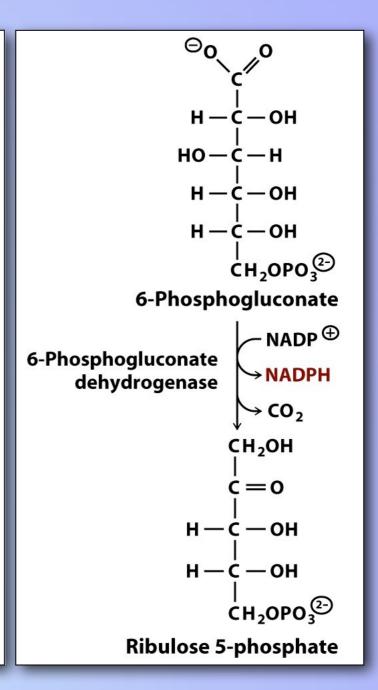






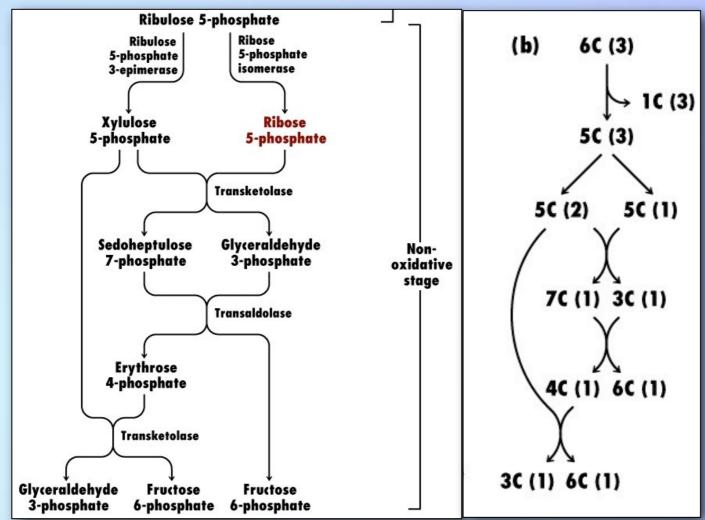






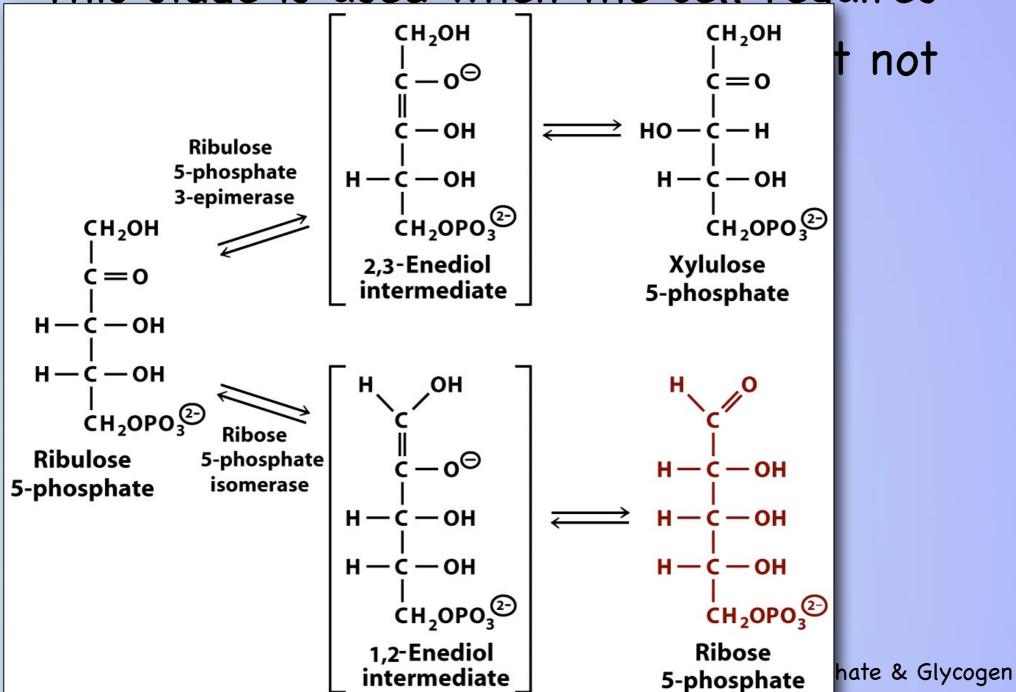
- + This is the only stage that is used if the cell requires **both** pentoses and biosynthetic reducing power.
  - Such is the cases when cells are rapidly growing and dividing.

- ·Nonoxidative Stage:
  - + This stage is used when the cell requires only pentoses or reducing power, but not both.

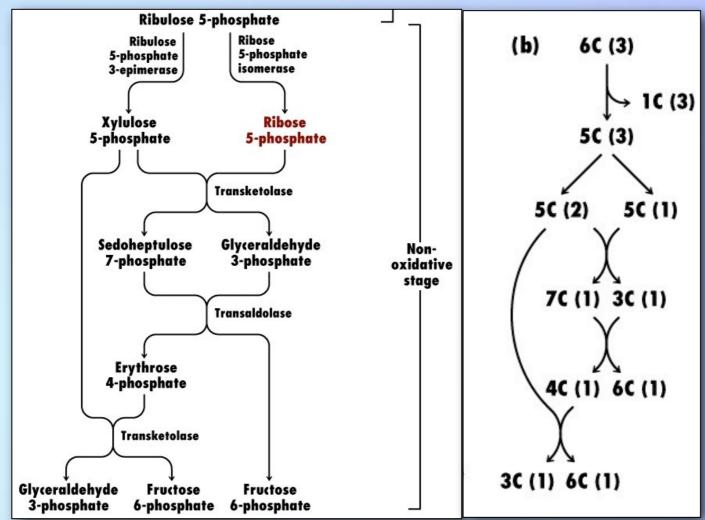


### ·Nonoxidative Stage:

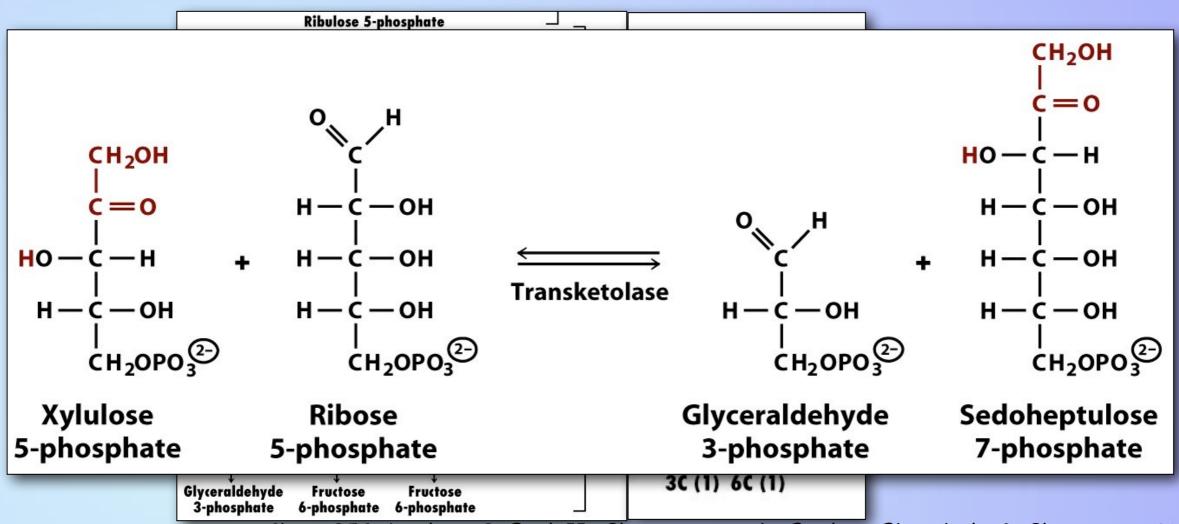
+ This stage is used when the cell requires



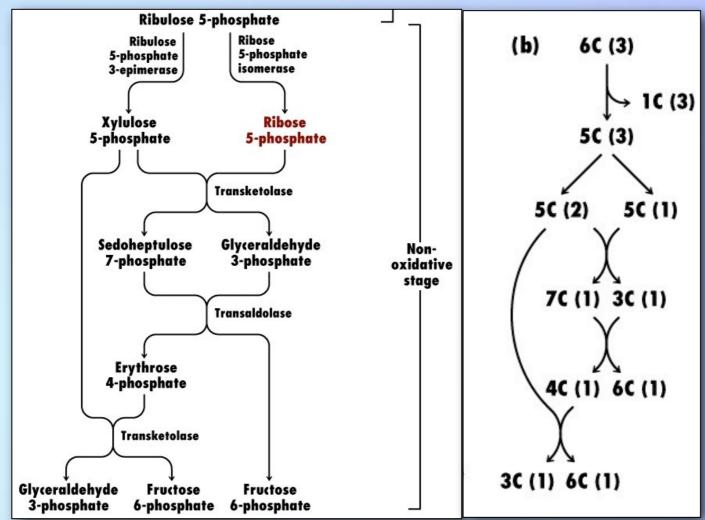
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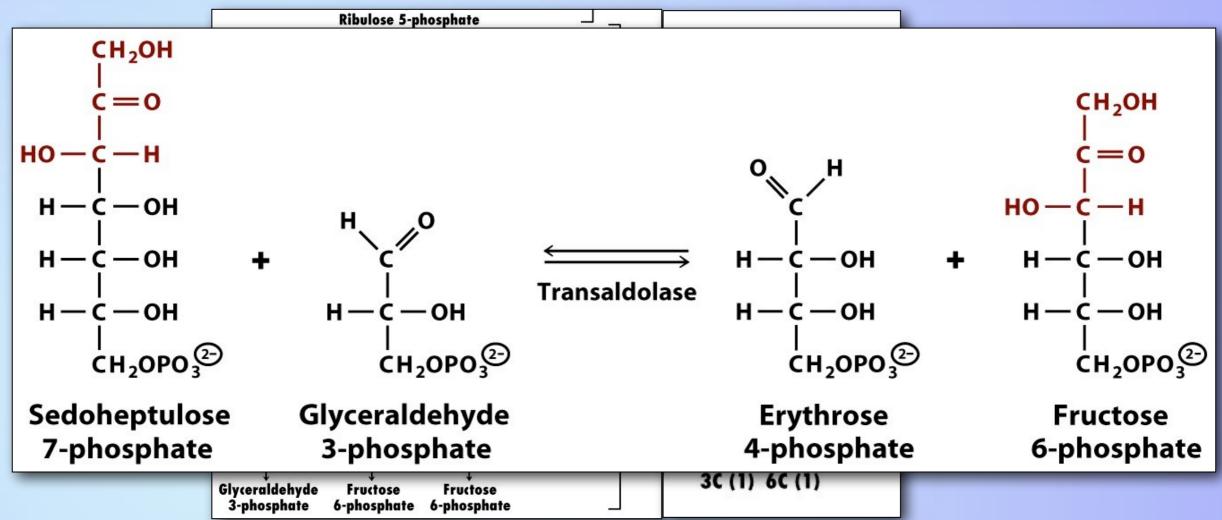
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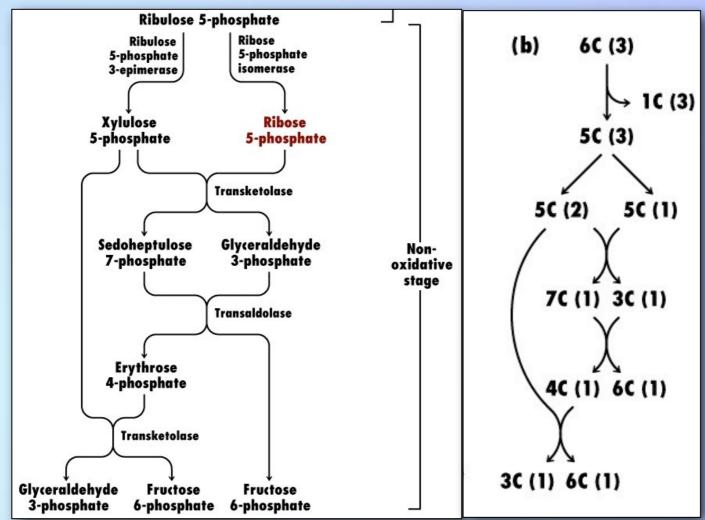
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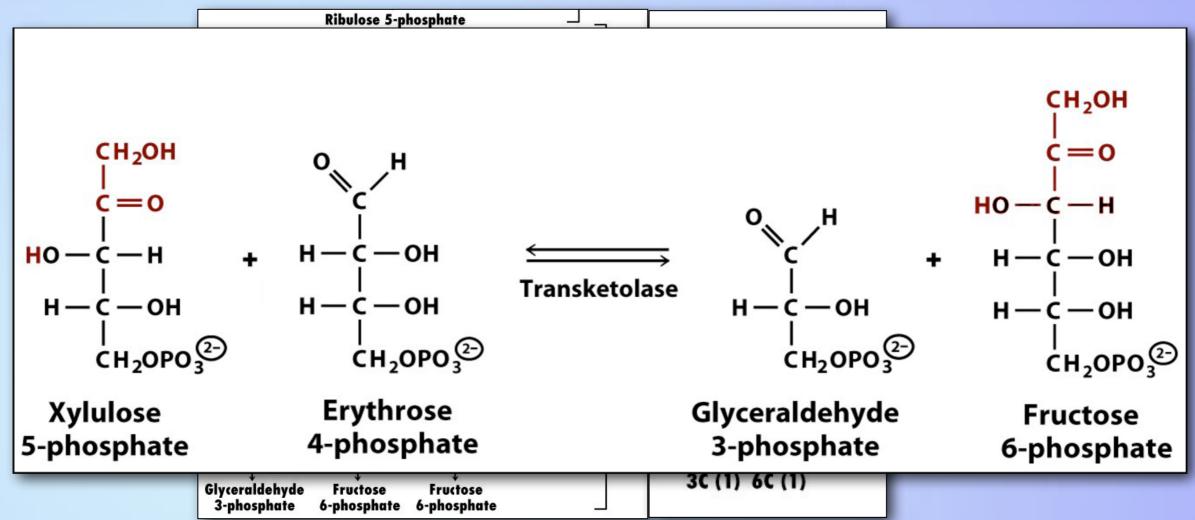
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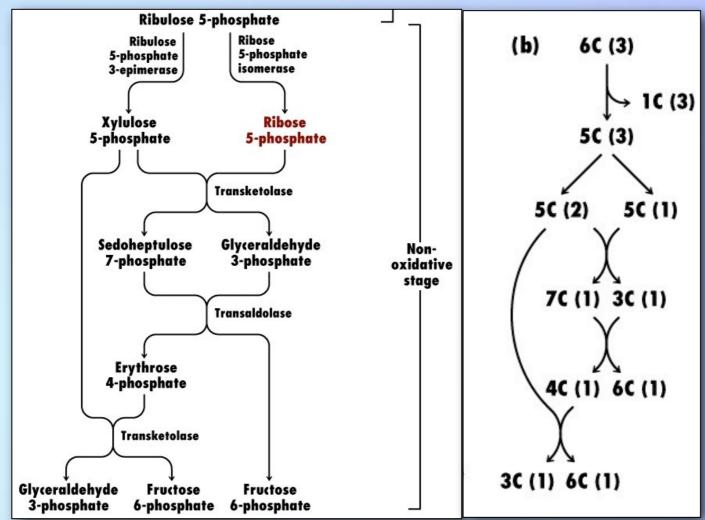
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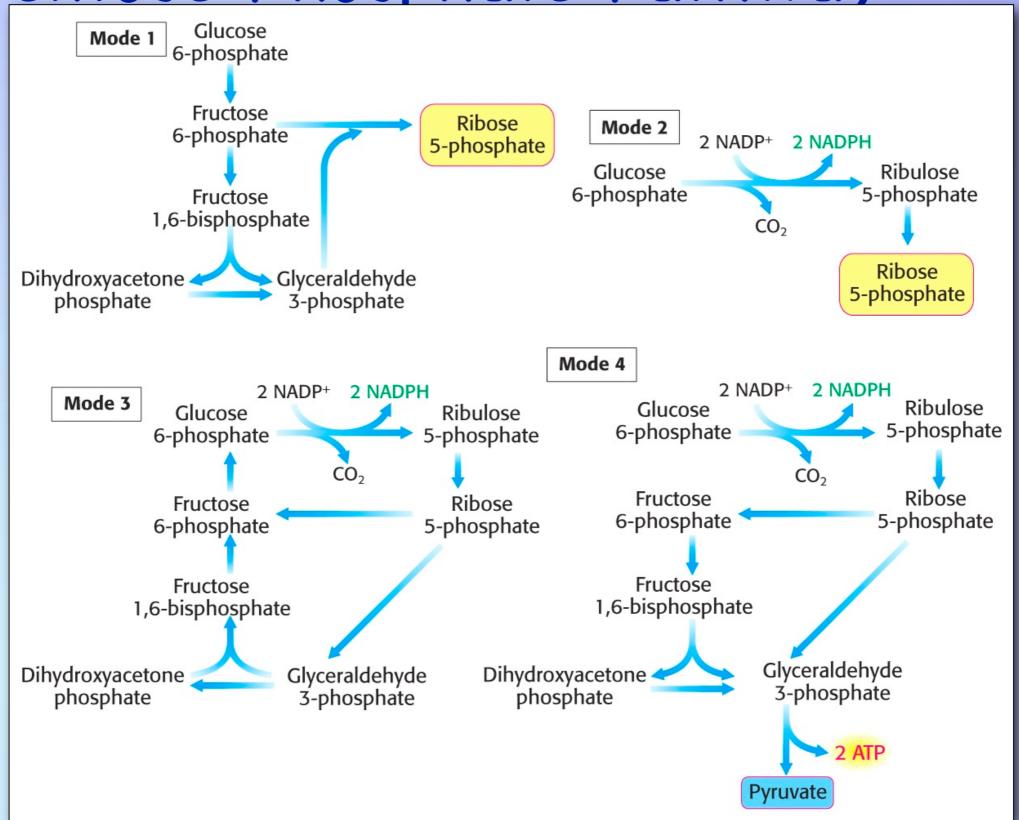


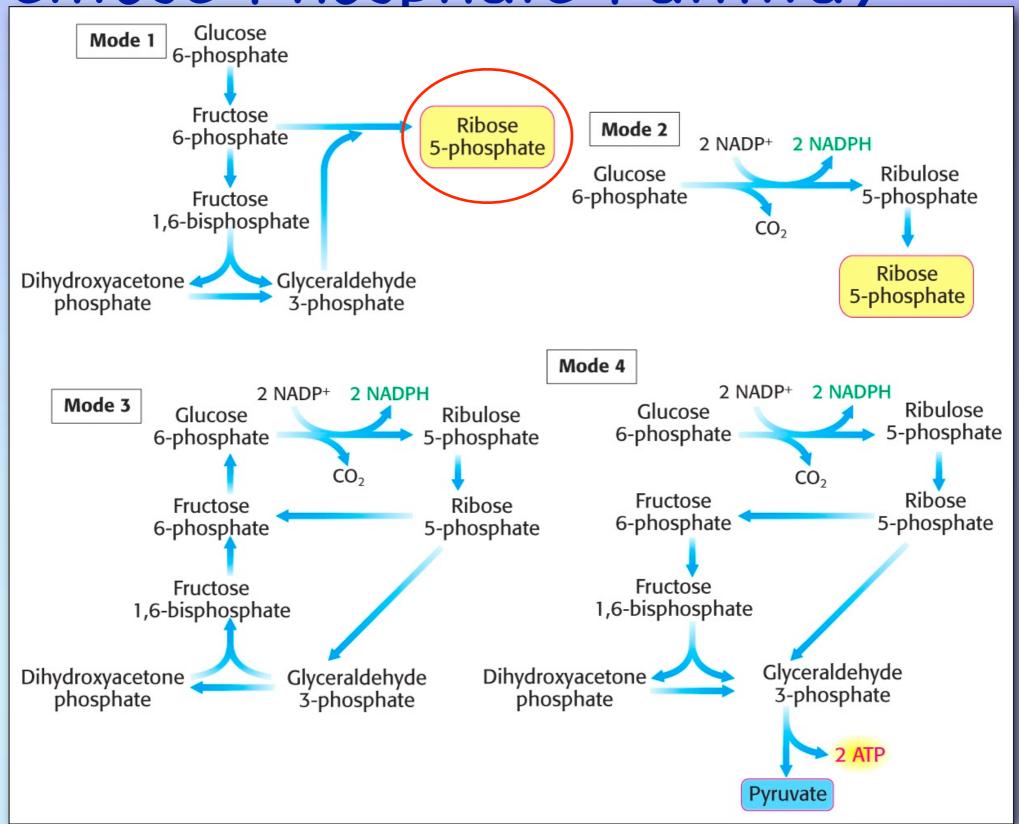
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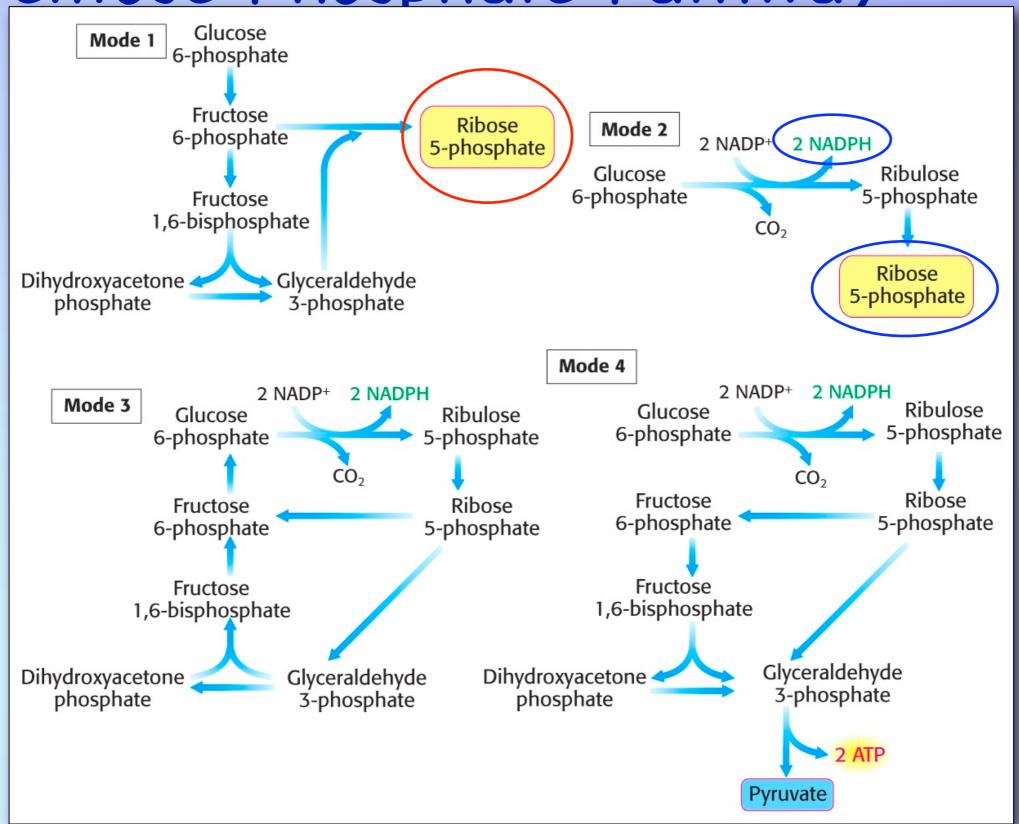


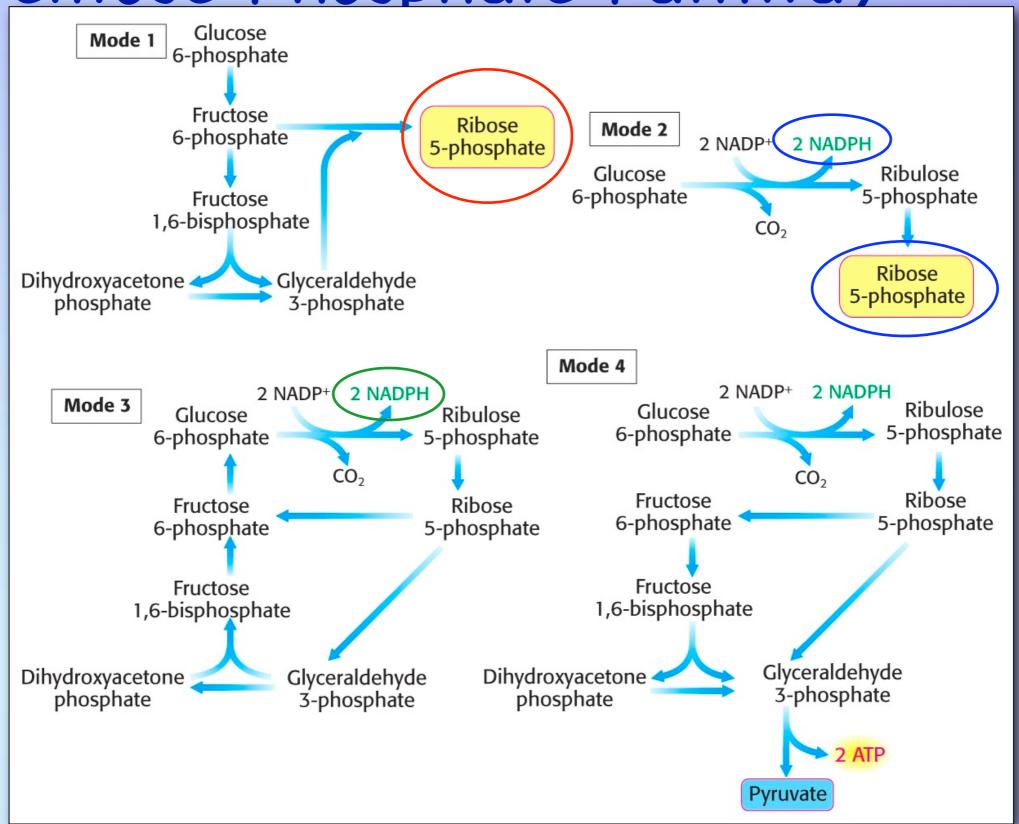
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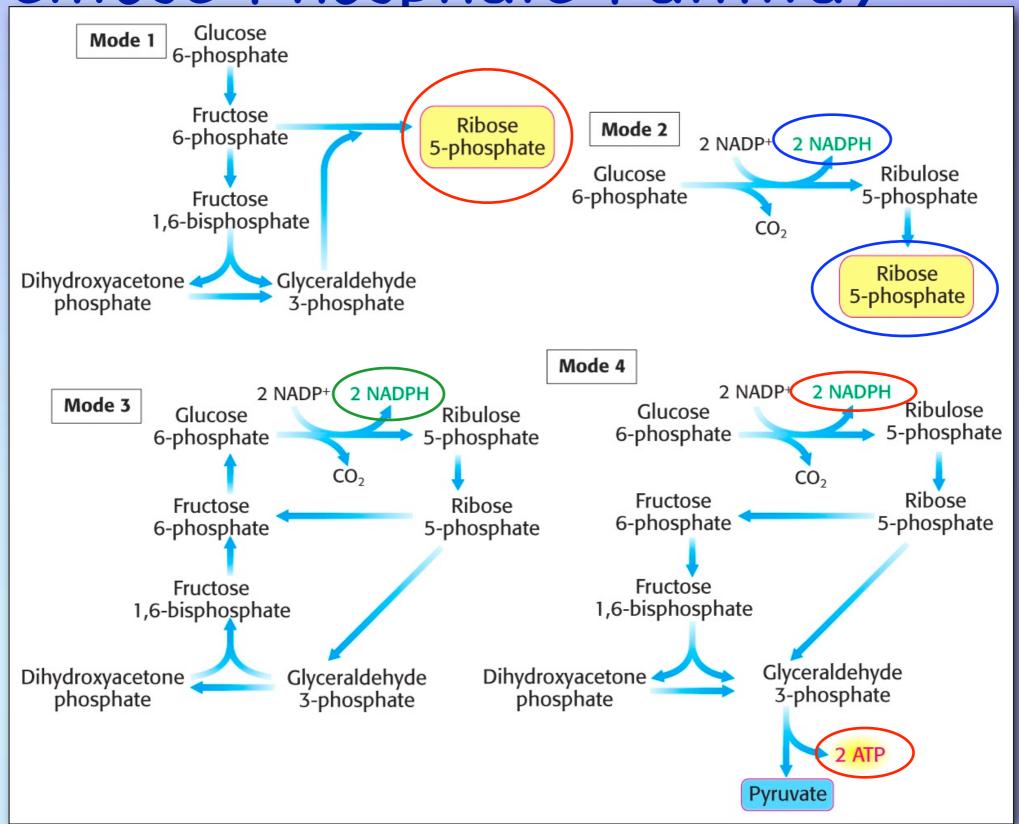








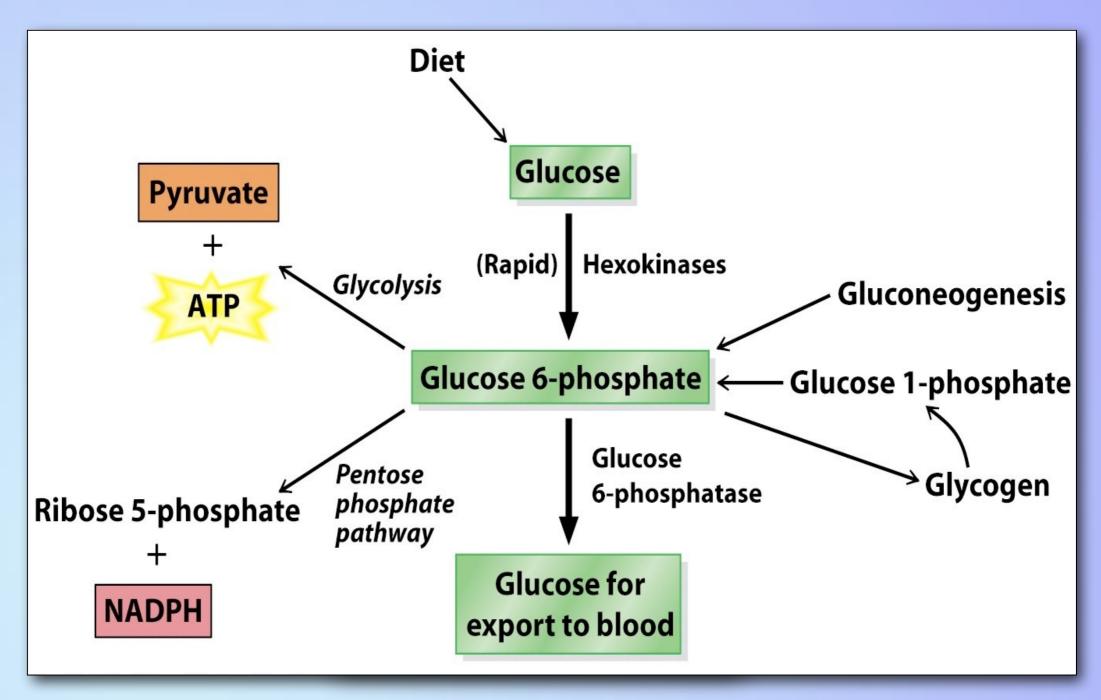




Chem 352 - Lecture 8
Carbohydrate Metabolism
Part II: Gluconeogenesis, Pentose Phosphate
Pathway & Glycogen Metabolism
9. April, 2014

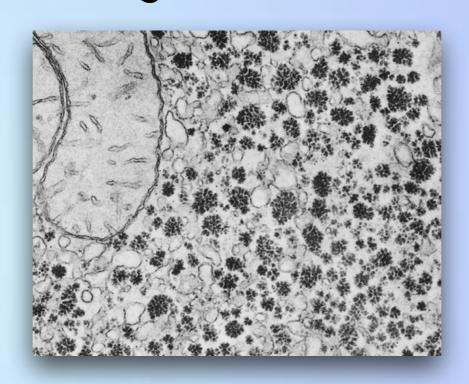
Question for the Day: How is the storage of glucose as glycogen in the liver tied to the blood glucose levels?

Glycogen is a storage form of glucose.



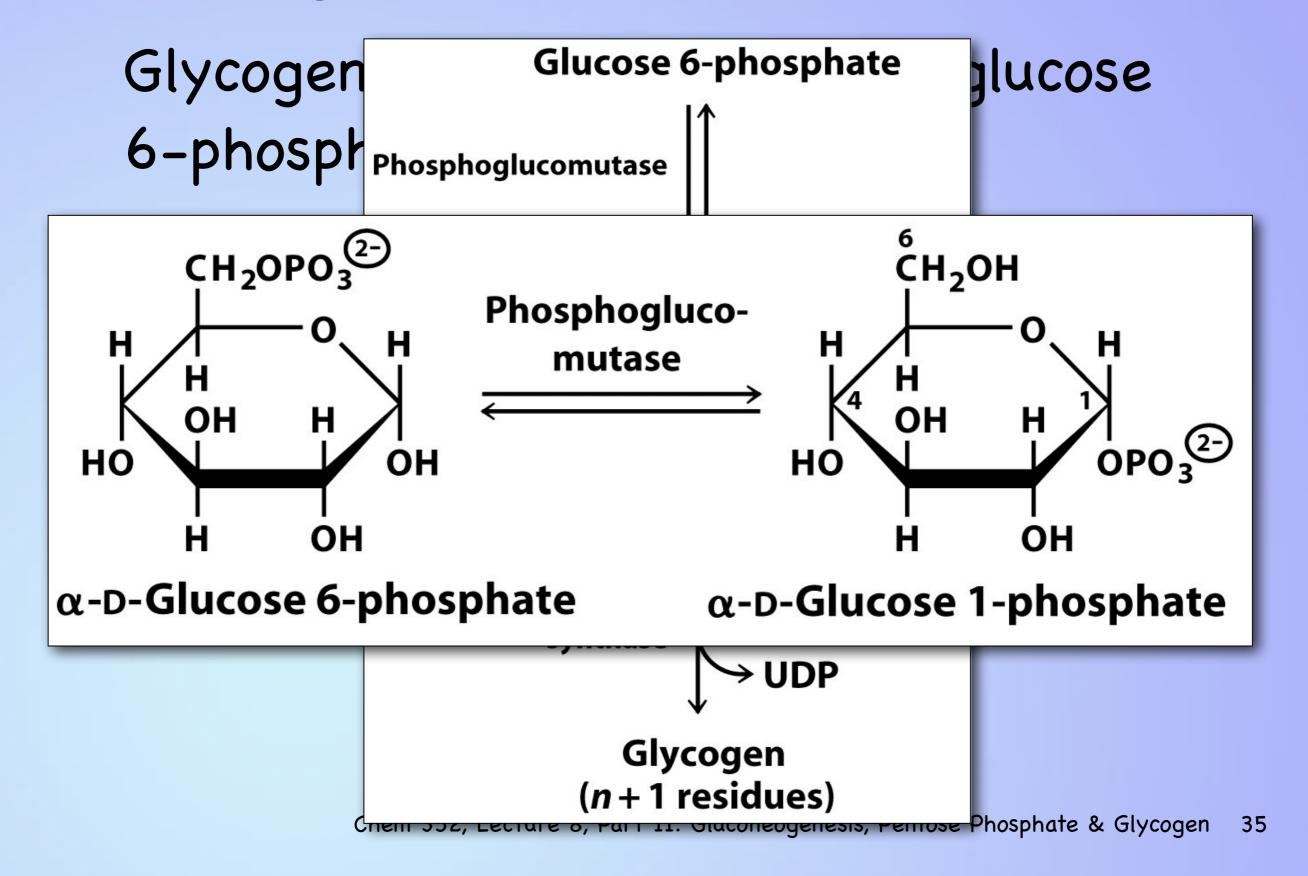
### Glycogen is a storage form of glucose.

- + It is stored in muscles as a readily available energy resource for future activity.
- + It is stored in the liver as a resource for regulating blood glucose levels.

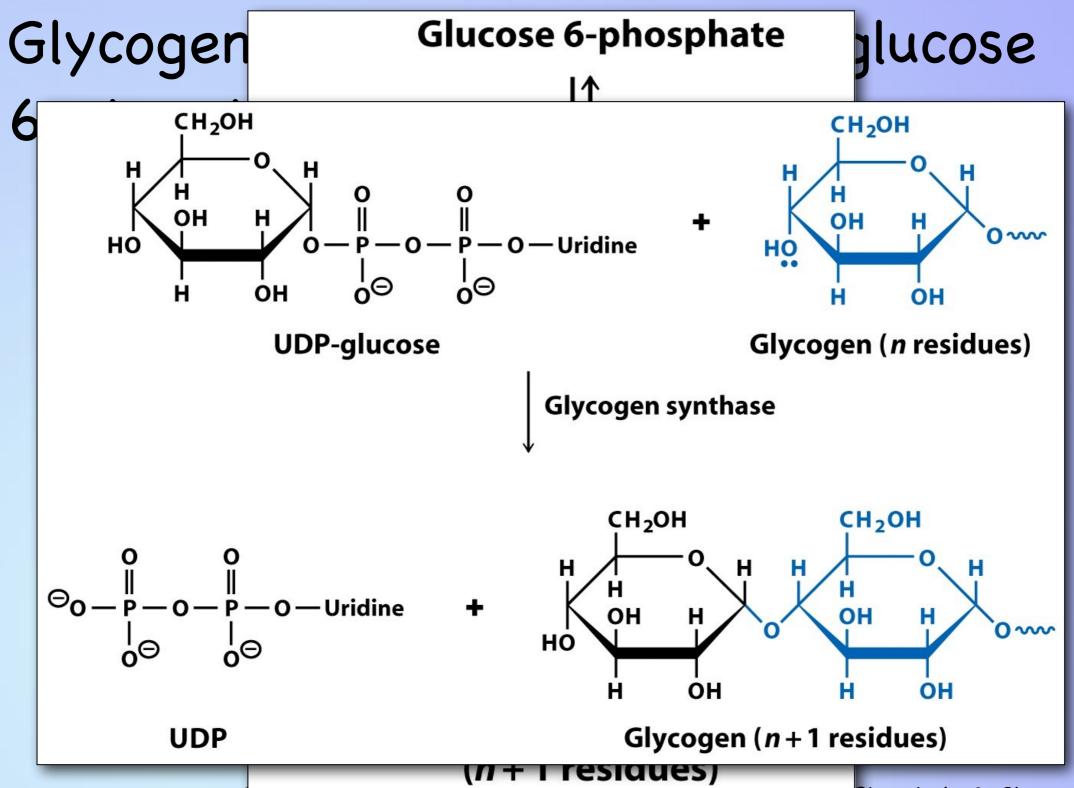


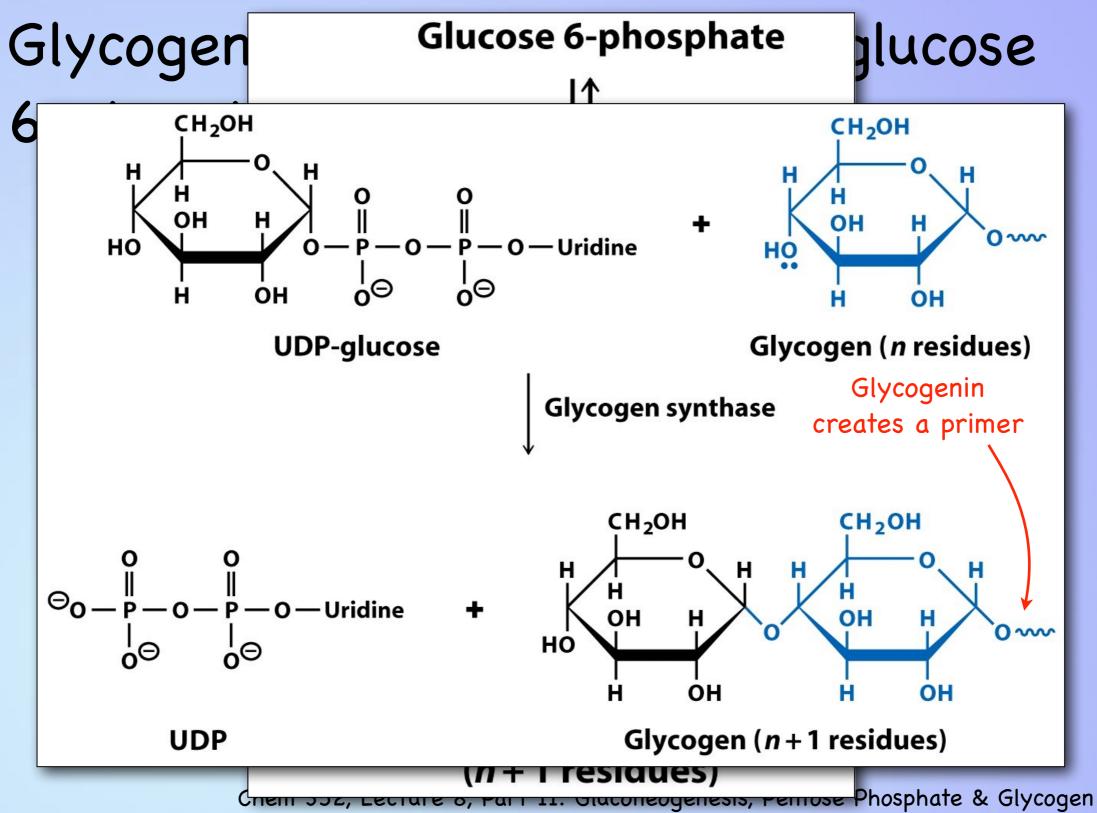
Glycogen is synthesized from glucose 6-phosphate.

Glycogen **Glucose 6-phosphate** glucose 6-phosphoglucomutase **Glucose 1-phosphate** UDP-glucose pyrophosphorylase PD **UDP-glucose** Glycogen (n residues) Synthase UDP Glycogen (n+1 residues)



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35

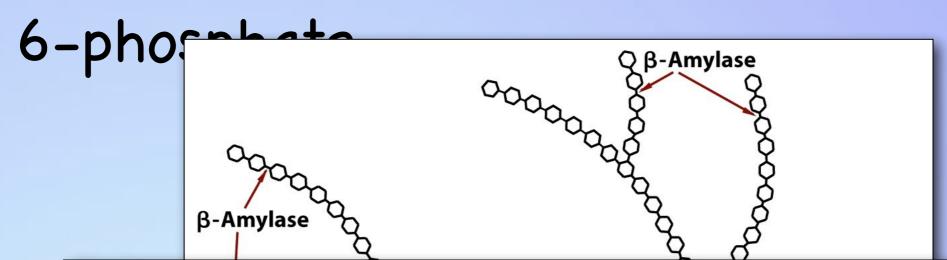
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Glycogen is synthesized from glucose

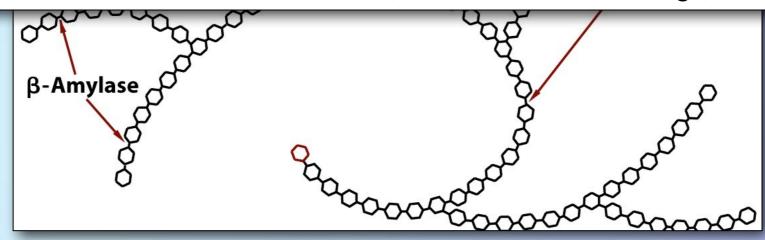
Q β-Amylase α-Amylase  $\alpha$ -Amylase

Glycogen is synthesized from glucose



The  $\alpha(1\rightarrow 6)$  glycosidic links are made transferring an oligosaccharide unit from a non reducing end to a point further in along the polymer.

• Reaction catalyzed by amylo-(1,4  $\rightarrow$ 1,6)-transglycosylase.

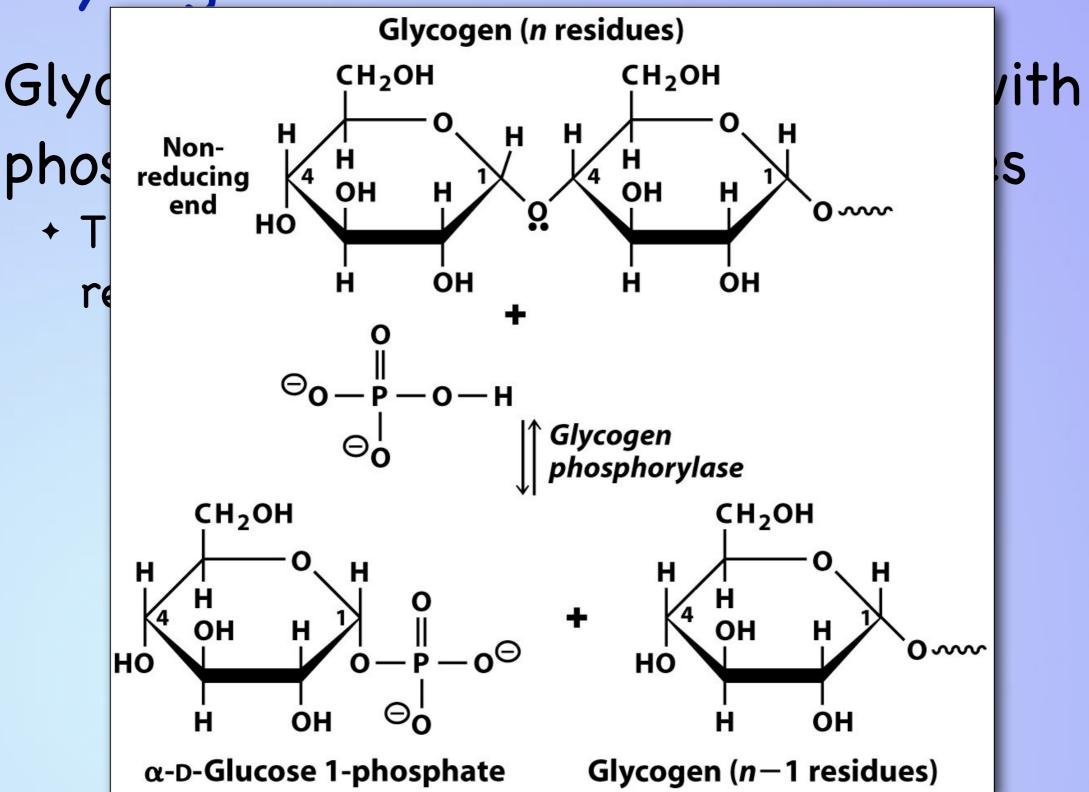


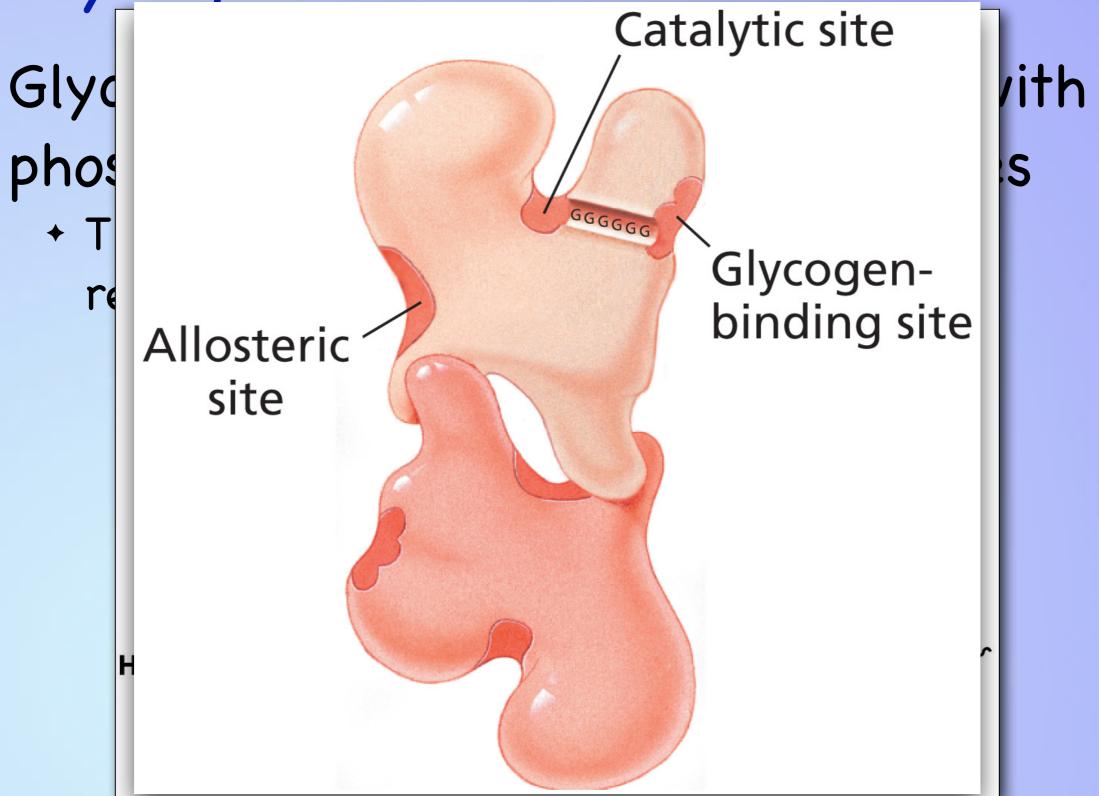
Glycogen is synthesized from glucose

Q β-Amylase α-Amylase  $\alpha$ -Amylase

Glycogen degradation is catalyzed with phosphorylases instead of hydrolases

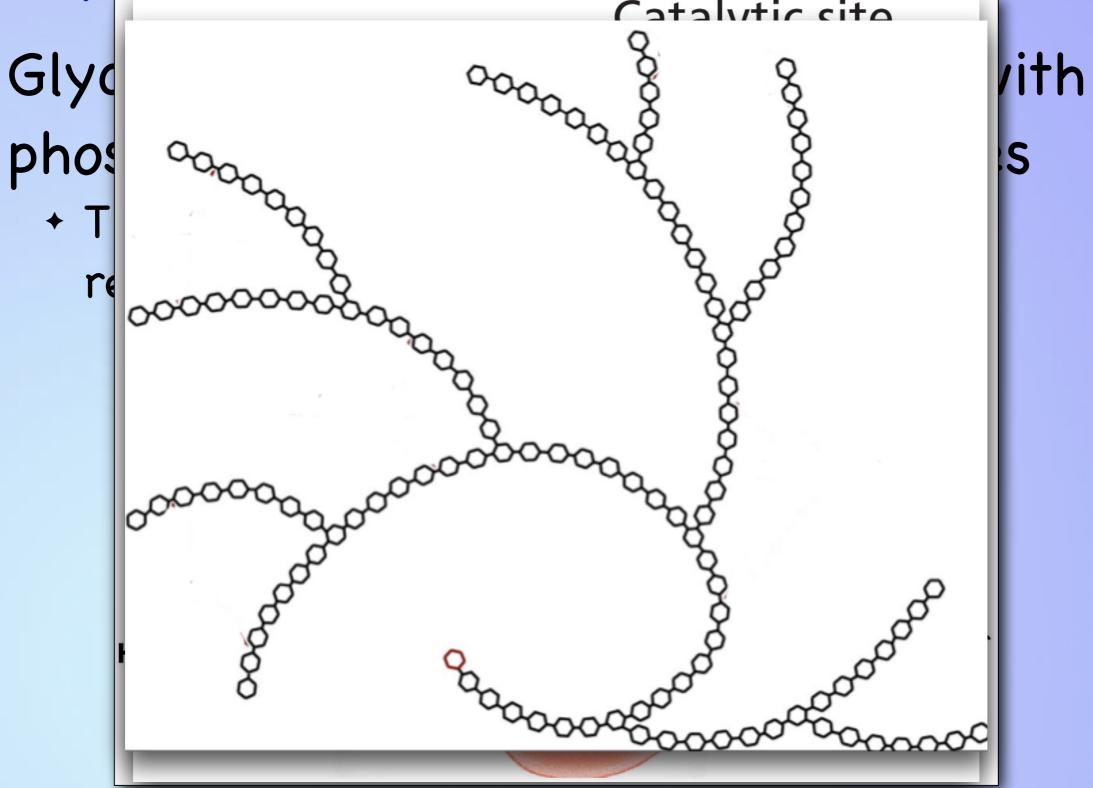
+ This leaves the glucose units that are removed with a phosphate attached.

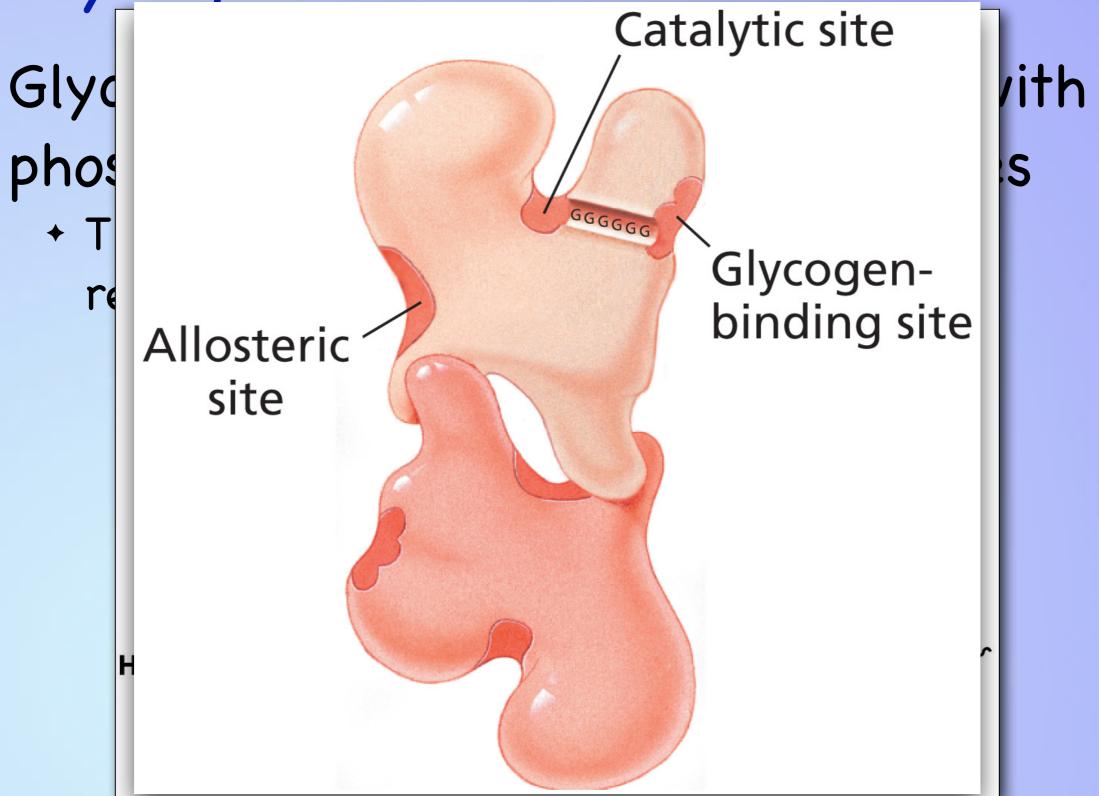


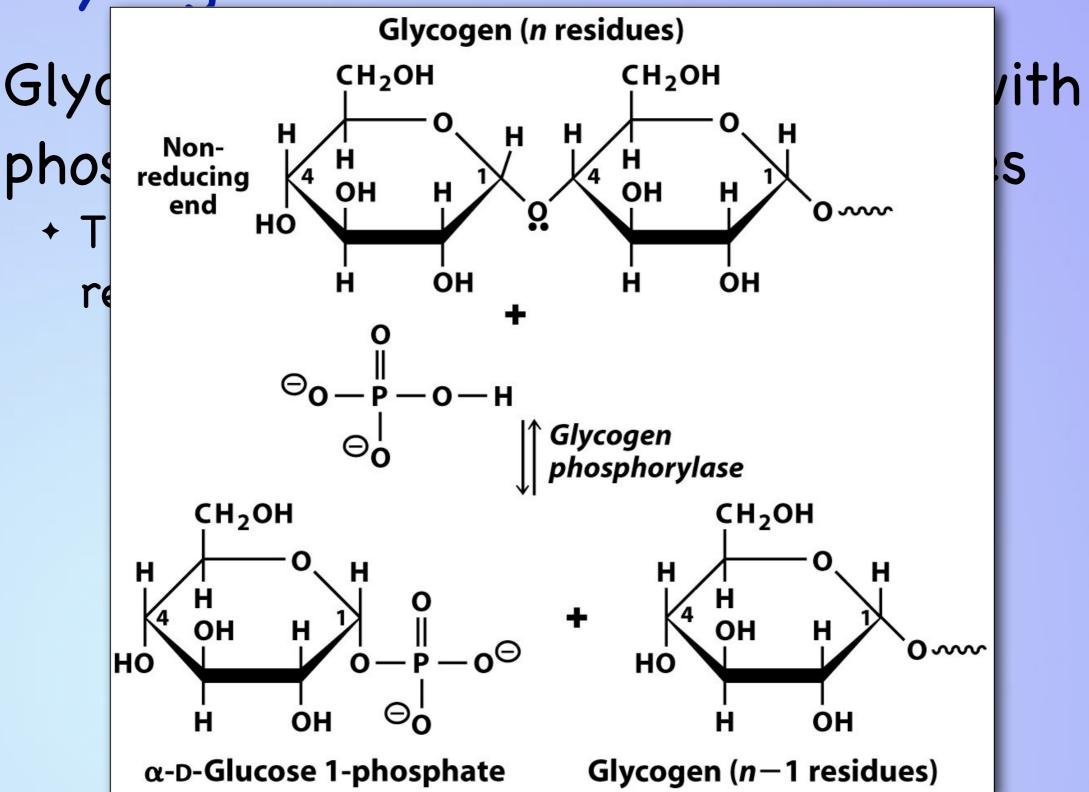


Glycogen Metabolism

Catalytic site







Glycogen degradation is catalyzed with phosphorylases instead of hydrolases

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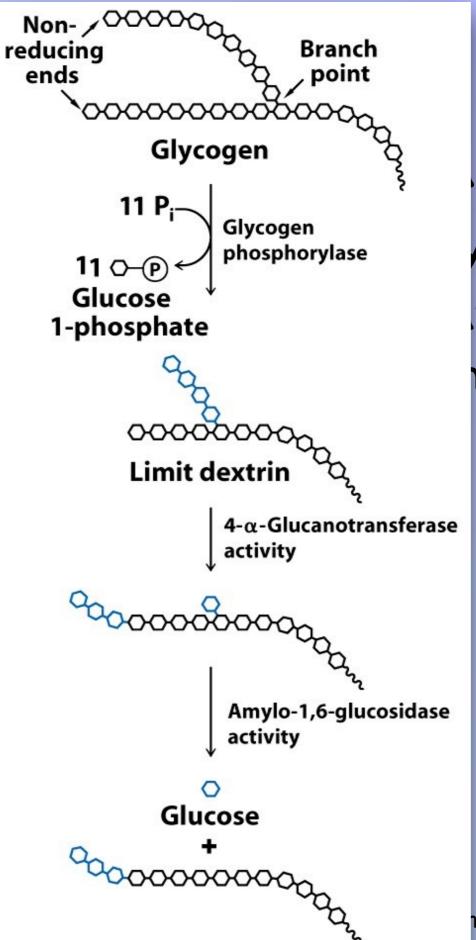
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# Glycogen

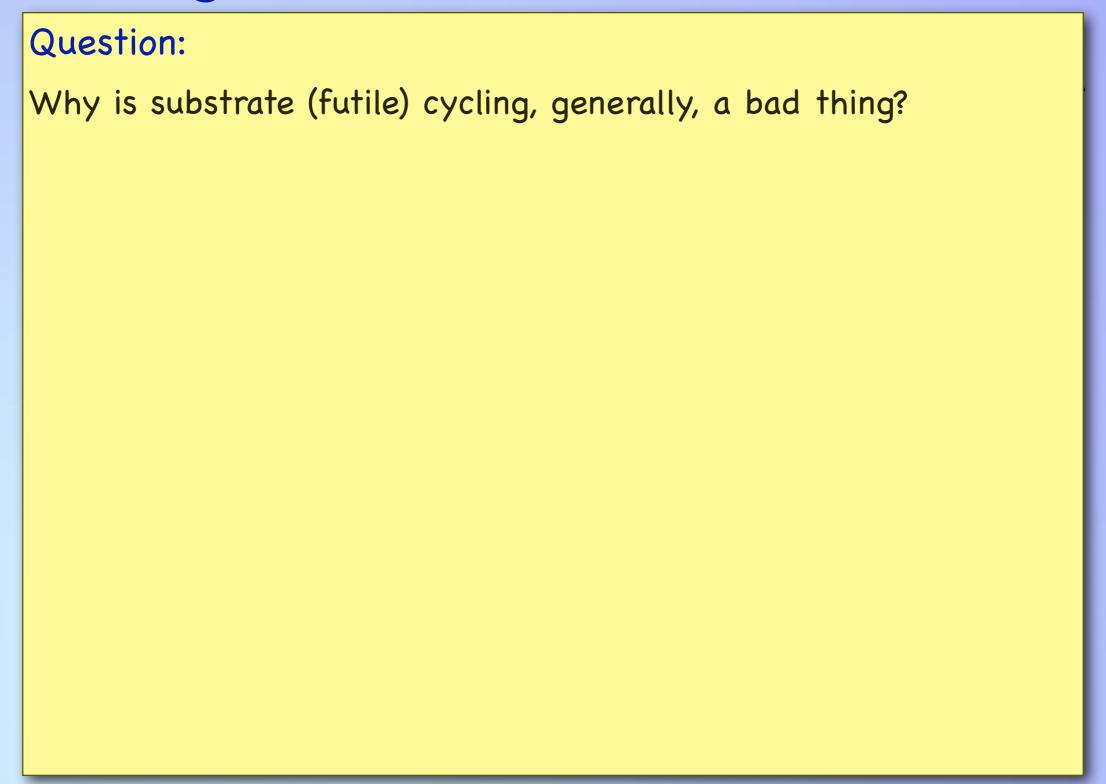
Glycogen de phosphoryle

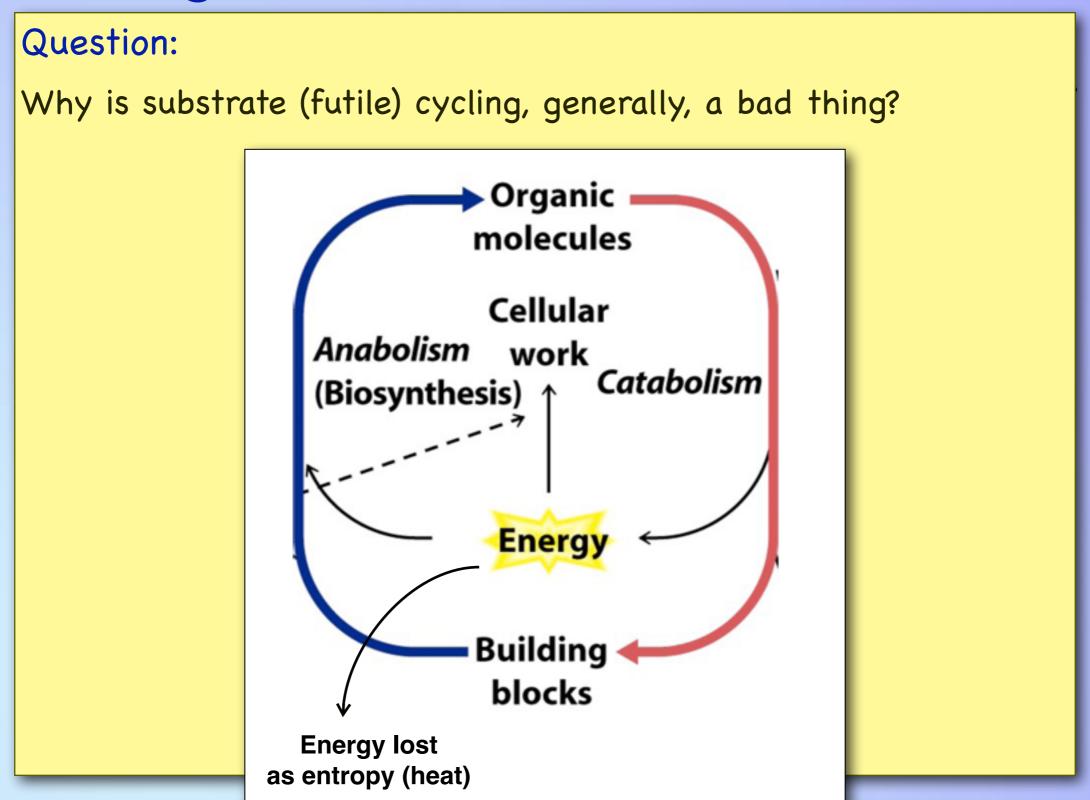
+ This leaves removed w



lyzed with drolases t are red.

- ·Regulation of glycogen synthesis and degradation are coordinated.
  - \* As we saw with glycolysis and gluconeogenesis, this is done to prevent substrate (futile) cycling.
- ·Insulin, glucagon and epinephrine (adrenaline) are the hormones that regulate glycogen metabolism.



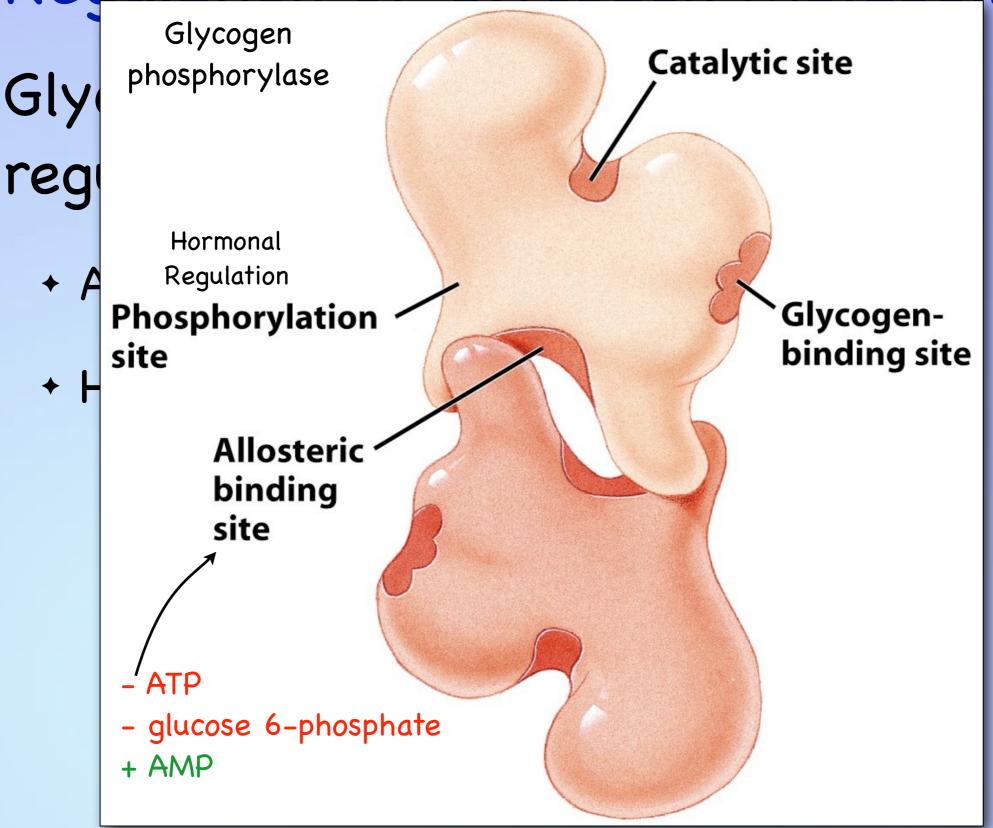


Chem 352, Lecture 8, Part II: Gluconeogenesis, Pentose Phosphate & Glycogen

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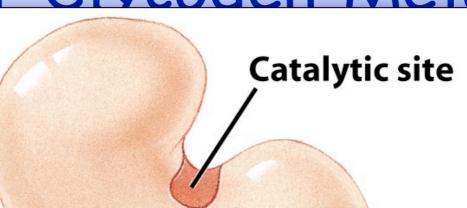
Glycogen metabolism is strictly regulated.

- + Allosterically
- + Hormonally



req

Glycogen phosphorylase



Hormonal

**Glycogen synthase** 

Regulation

Active form (a)

Inactive form (b)

OH

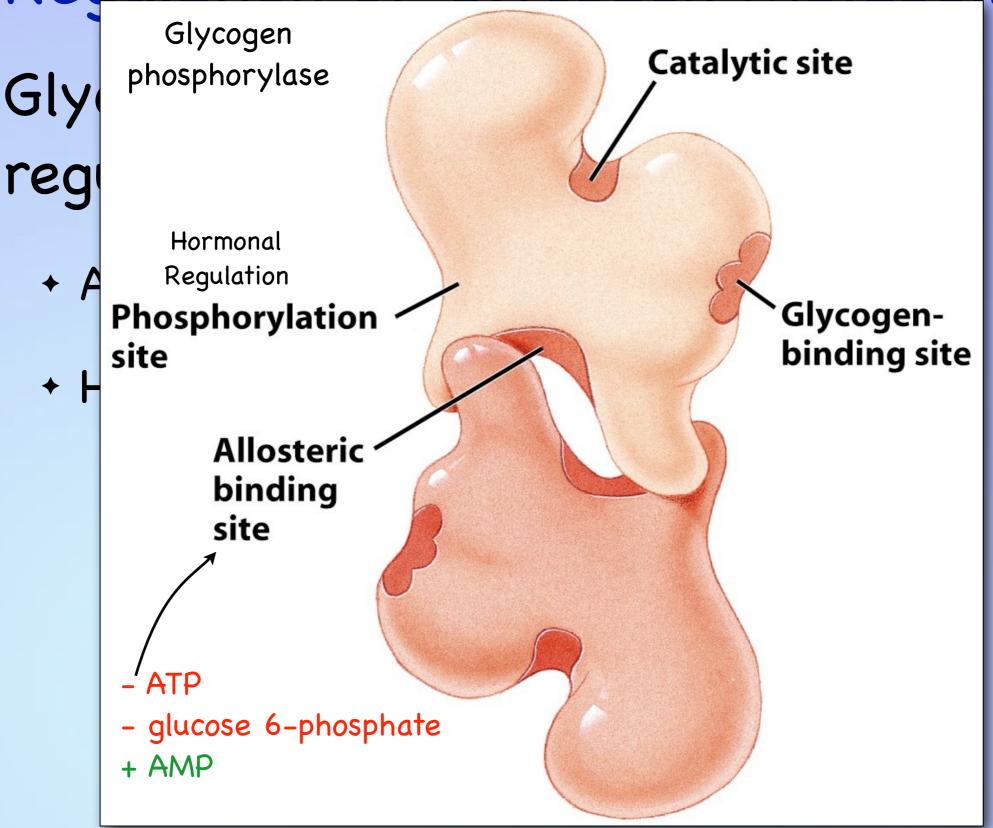
Glycogen phosphorylase

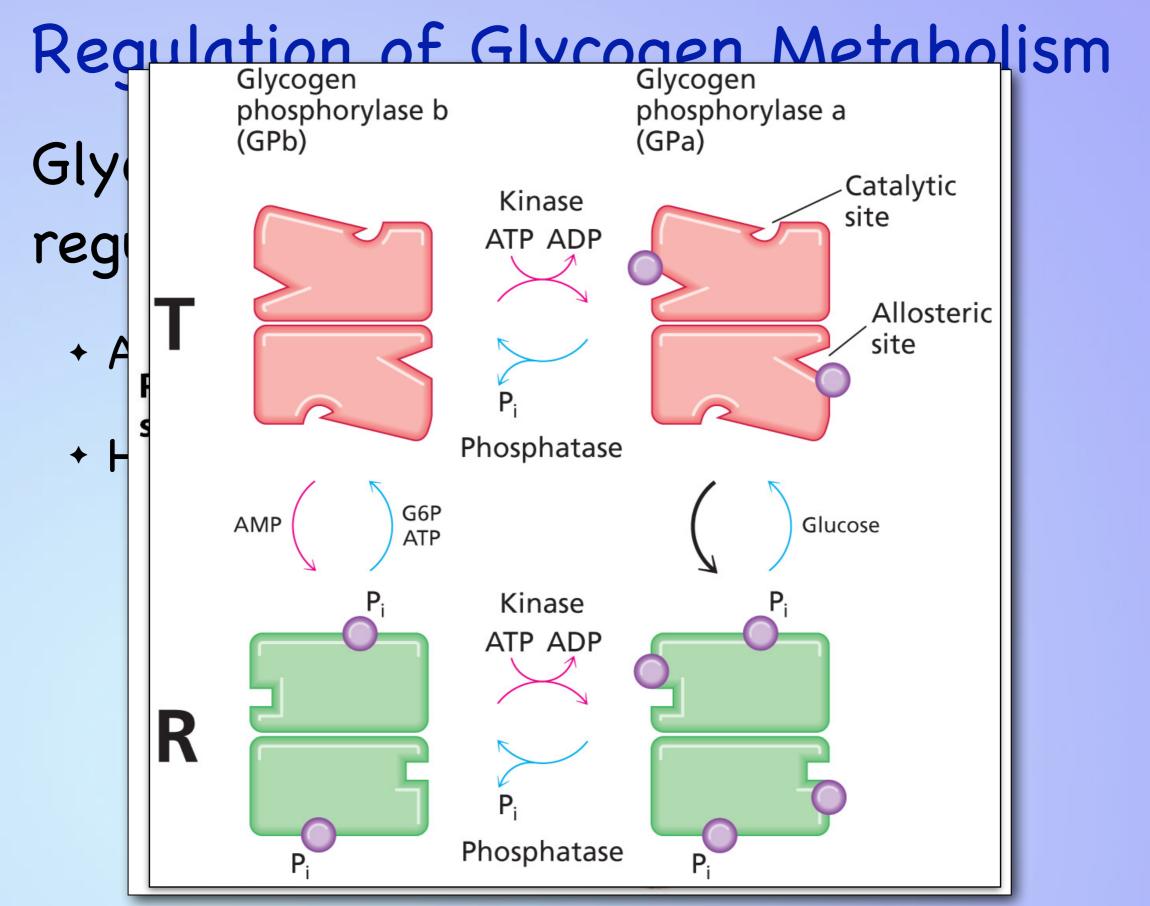


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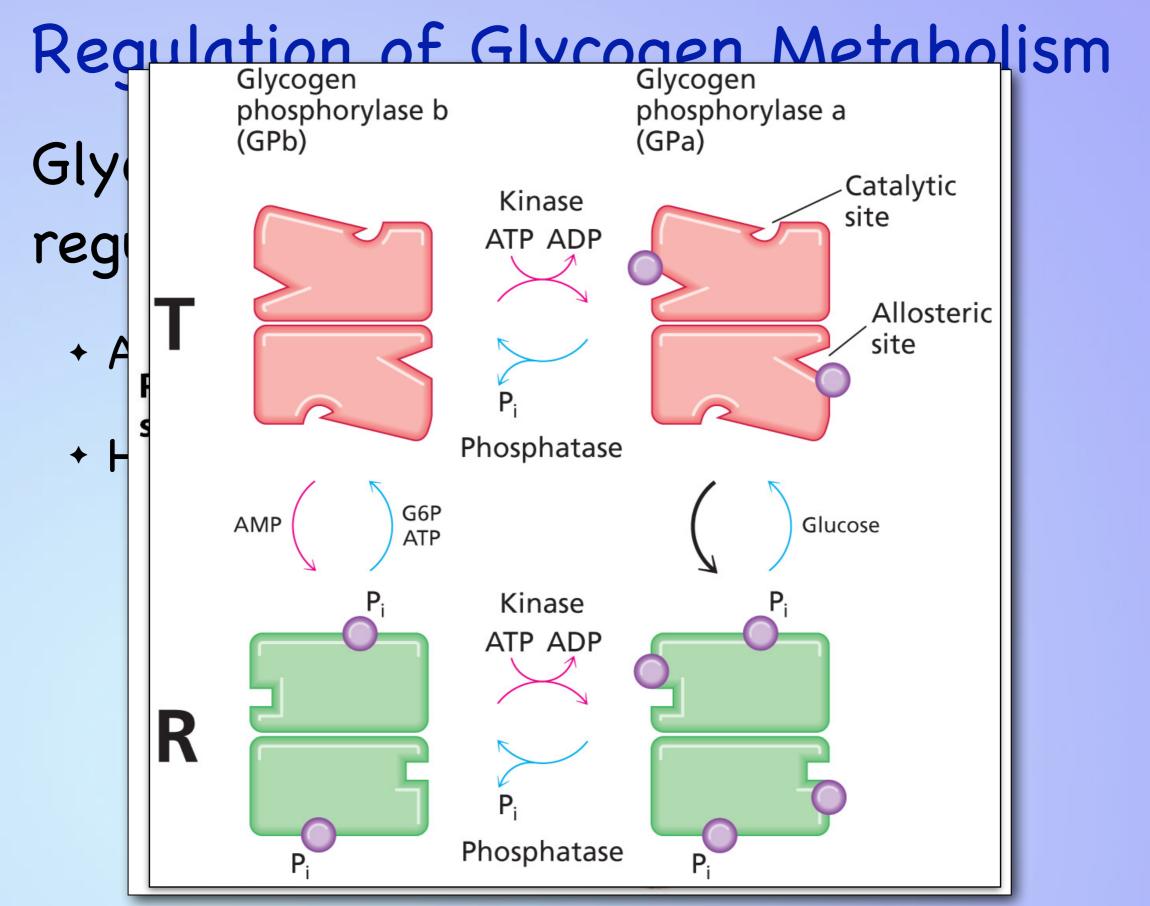
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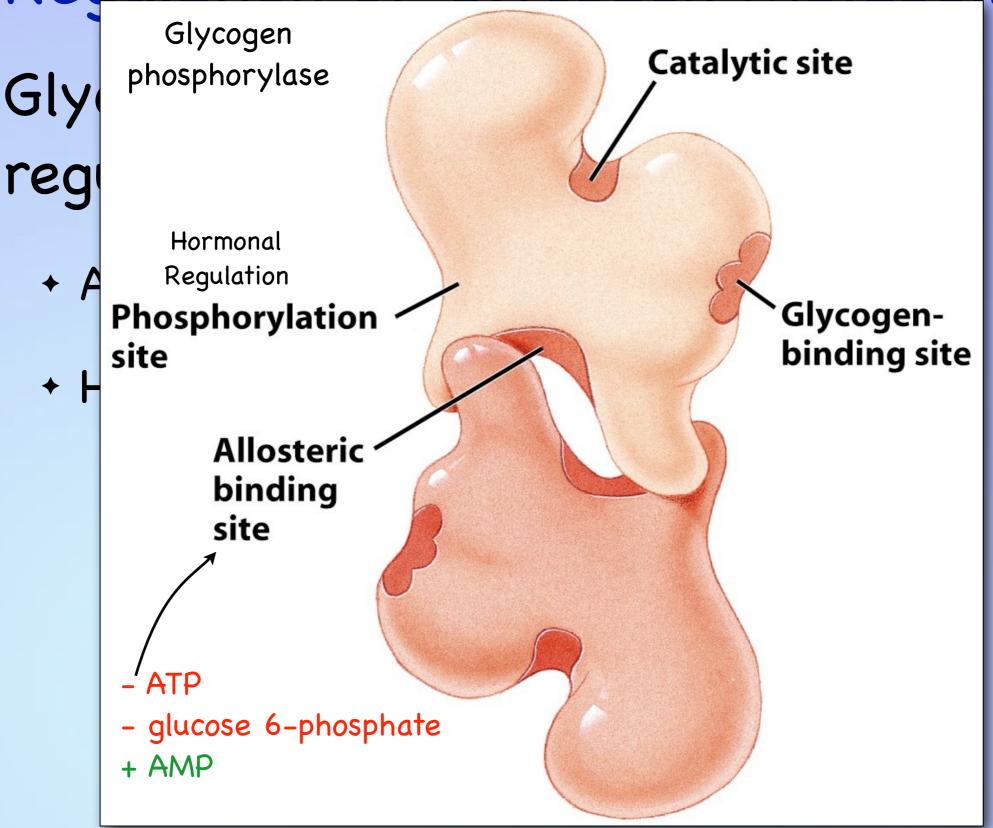
+ AMP





#### Regulation of Glycogen Metabolism Glycogen nhosnhorvlase h nhosphorylase a T state R state Glycogen Phosphorylase b Glycogen Phosphorylase a Catalytic site Catalytic site PLP' PLP' Ser-14' shift Ser-14' subunit 34Å rotation 10° Arg-69' Ser14-P' Tower Tower helices helices Arg-14 Ser-14 Catalytic site Catalytic site PLP PLP Phosphatase





Glycogen metabolism is strictly regulated.

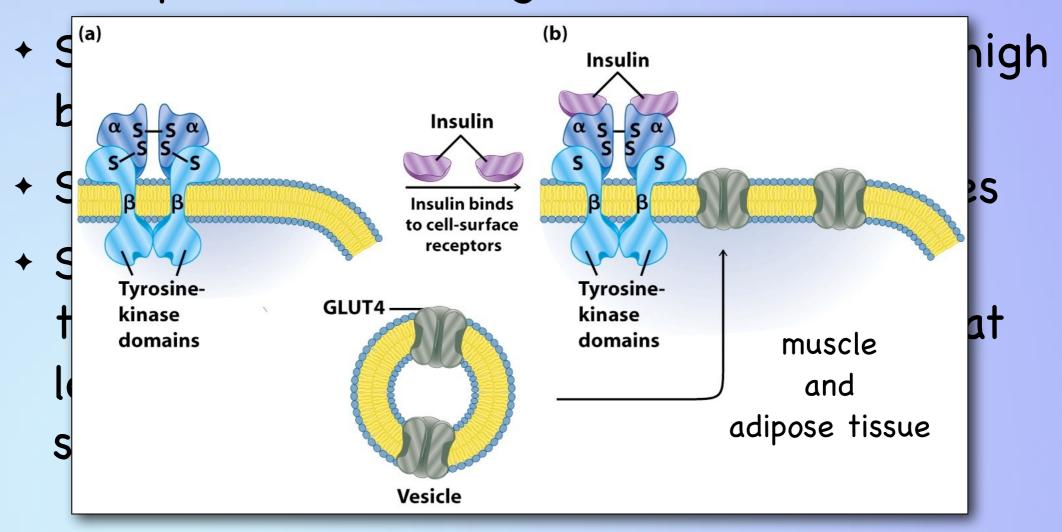
- + Allosterically
- + Hormonally

#### Insulin

- + Is a protein containing 51 amino acids
- + Secreted by the pancreas in response to high blood glucose levels
- + Stimulates the uptake of glucose by tissues
- \* Stimulates glycogen synthesis in the liver through a signal transduction pathway that leads to dephosphorylation of glycogen synthetase and glycogen phosphorylase

### Insulin

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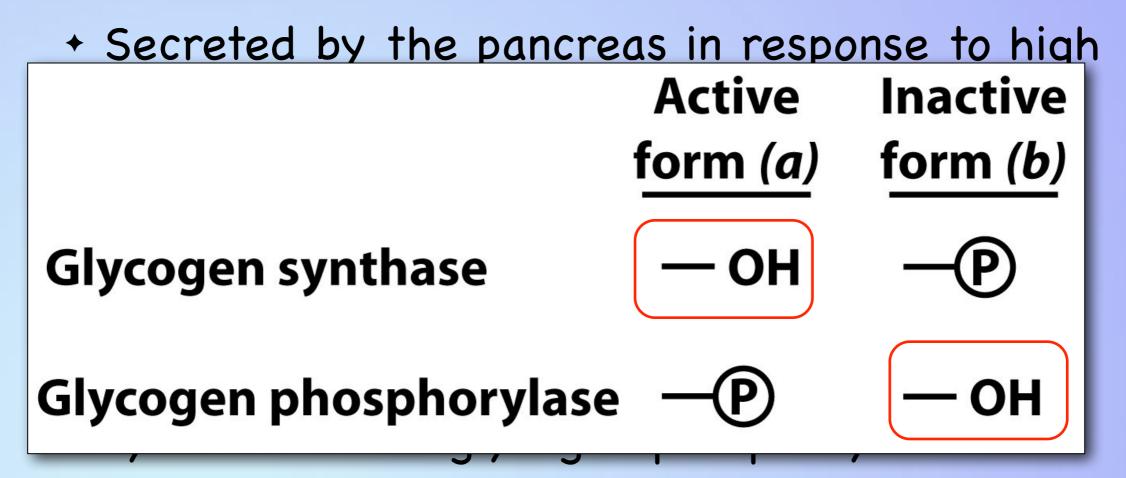


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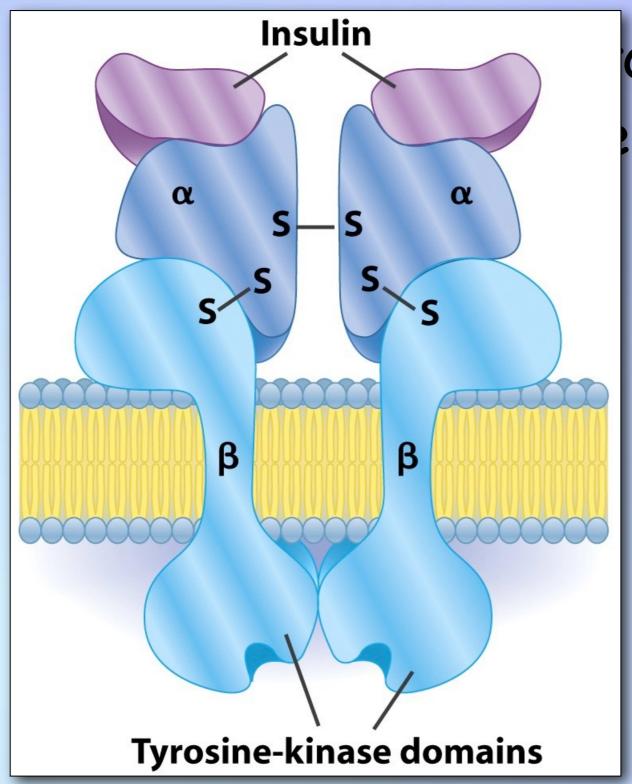
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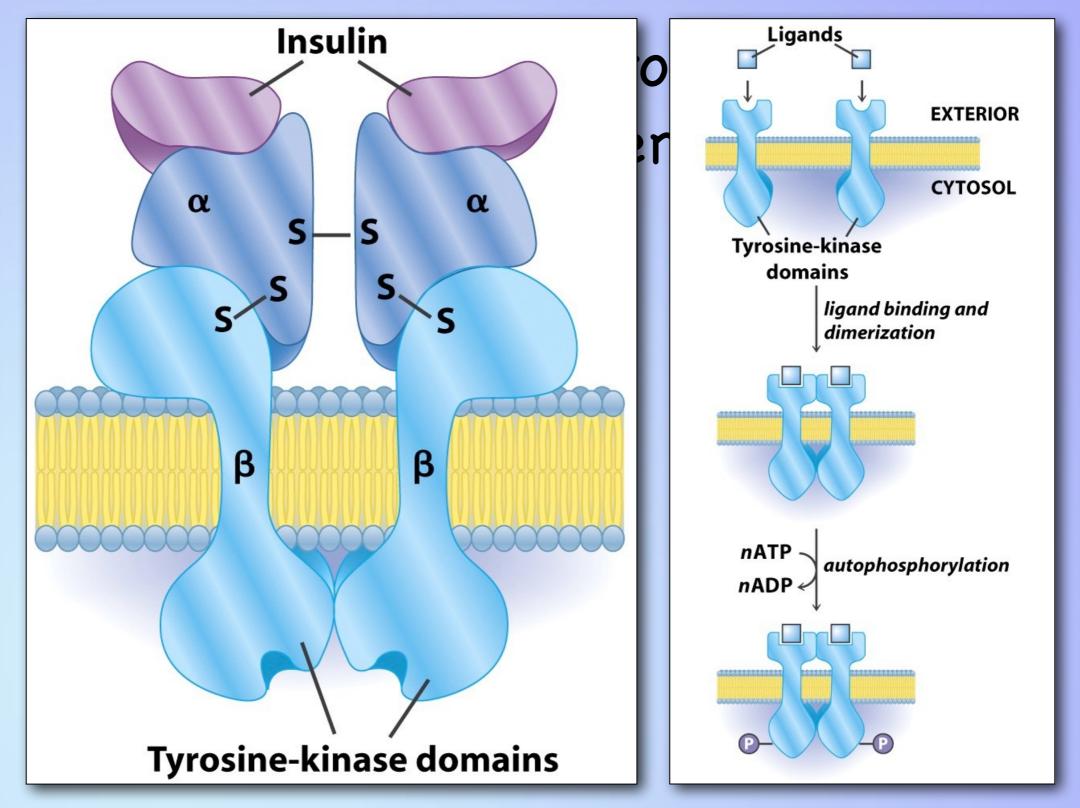
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•Insulin binds to a tyrosine kinase type receptor. (Chapter 9.12, Section D)



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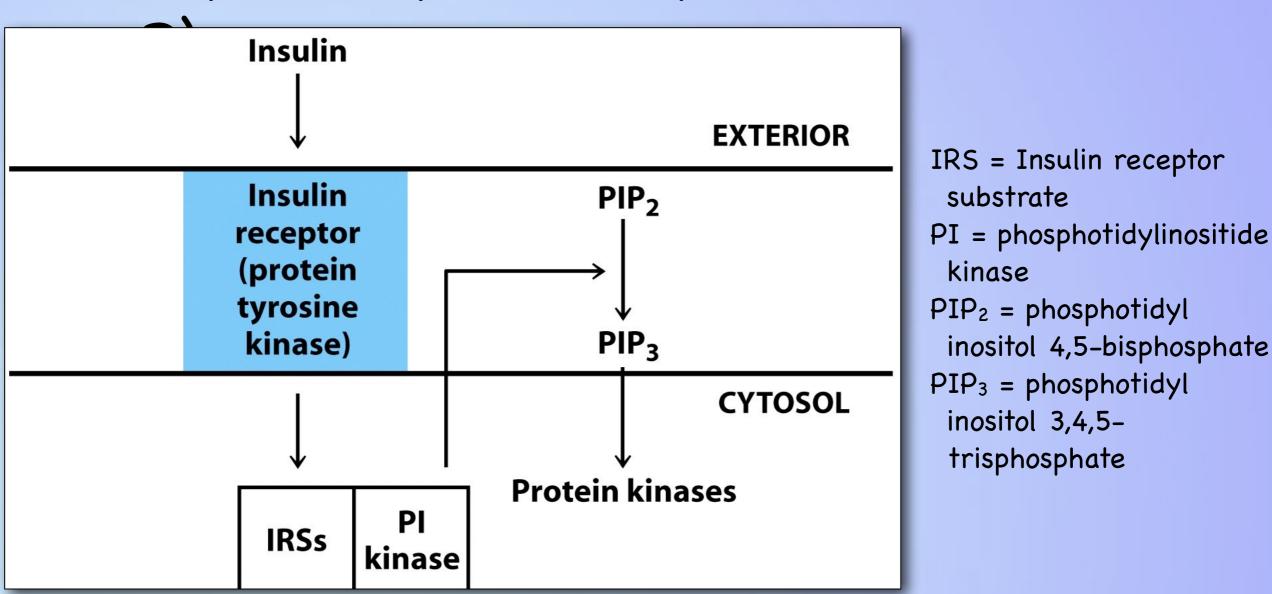
Chem 352, Lecture 8, Part II: Gluconeogenesis, Pentose Phosphate & Glycogen



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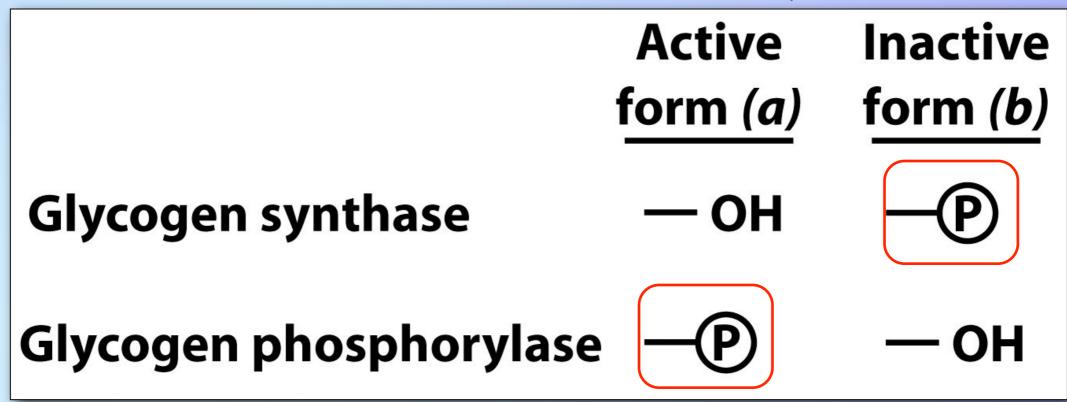
·Insulin binds to a tyrosine kinase type receptor. (Chapter 9.12, Section



### ·Glucagon

- + Is a peptide hormone containing 29 amino acids
- + Secreted by the  $\alpha$  cells in the pancreas in response to low blood glucose levels
- \* Stimulates glycogen degradation in the liver through a signal transduction pathway that leads to phosphorylation of glycogen synthetase and glycogen phosphorylase

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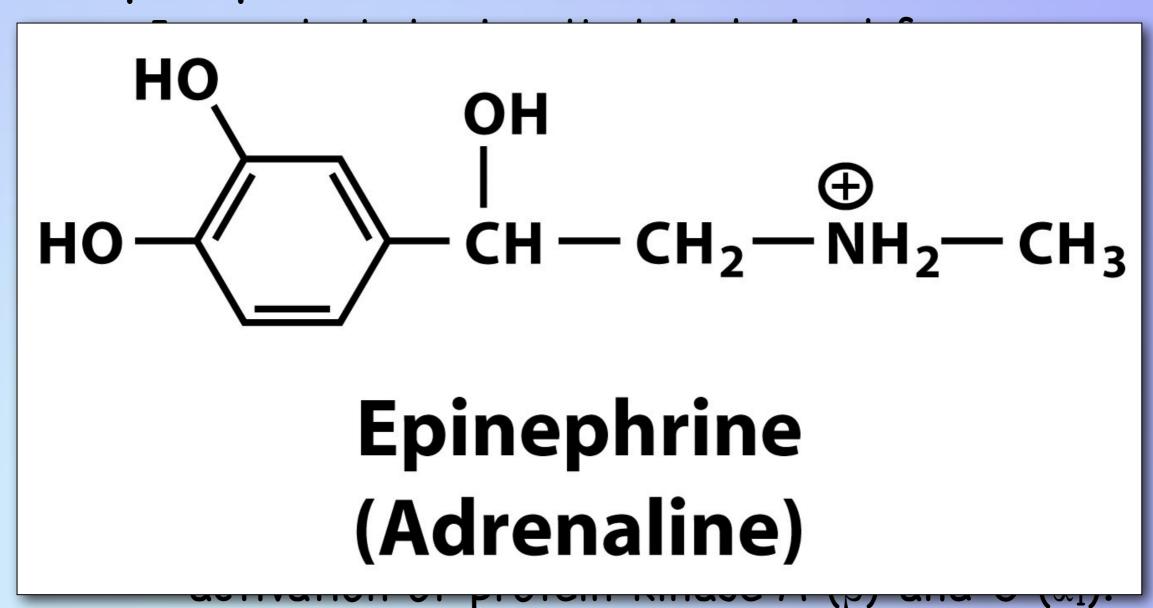
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### ·Epinephrine

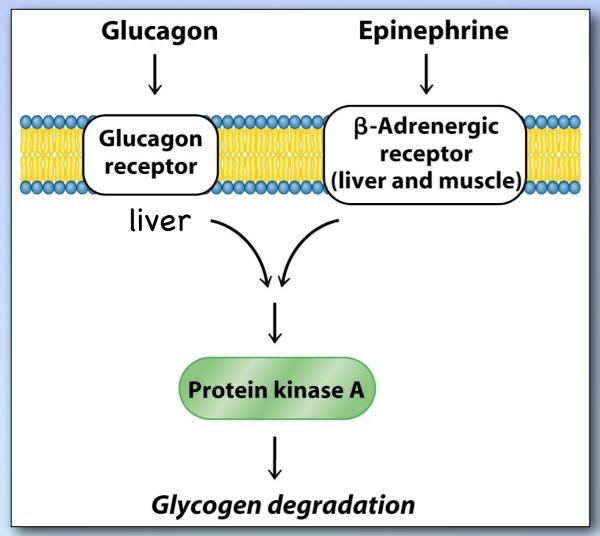
- + Is a catacholamine that is derived from tyrosine.
- \* Released by the adrenal glands by a neural signal that is triggered by the "Fight-or-Flight" response.
- + Stimulates the breakdown of glycogen
- \* Binds to adrenergic receptors in the muscle  $(\beta)$  and the liver  $(\alpha_1)$ .
  - Signal transduction pathway leads to activation of protein kinase A ( $\beta$ ) and C ( $\alpha_1$ ).

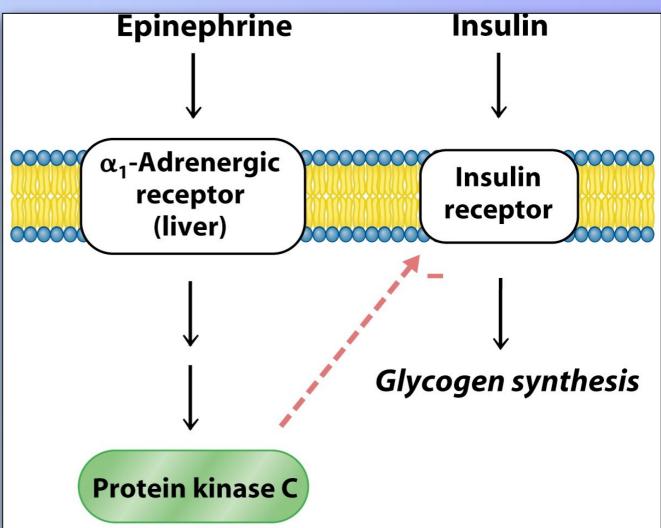
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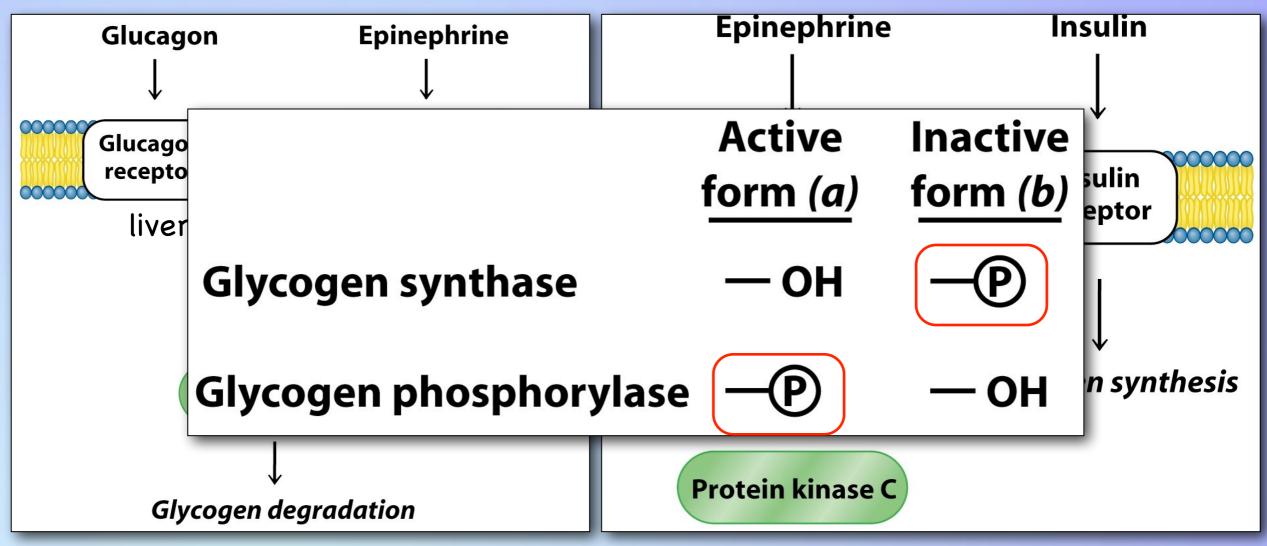




Both activate glycogen degradation by phosphorylating Glycogen synthase (inactive) and Glycogen phosphorylase (active)

Both inhibit the action of insulin by phosphorylating the insulin receptor

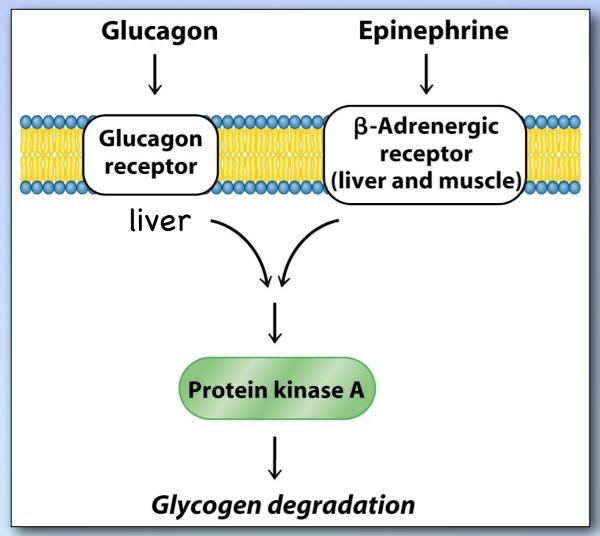
Chem 352, Lecture 8, Part II: Gluconeogenesis, Pentose Phosphate & Glycogen

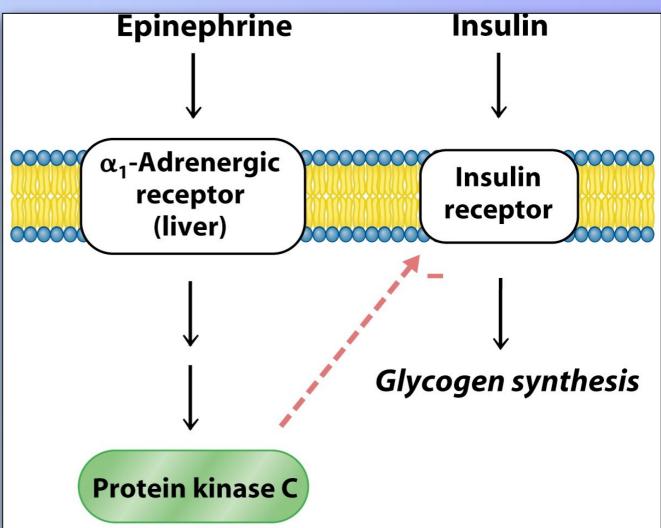


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Chem 352, Lecture 8, Part II: Gluconeogenesis, Pentose Phosphate & Glycogen

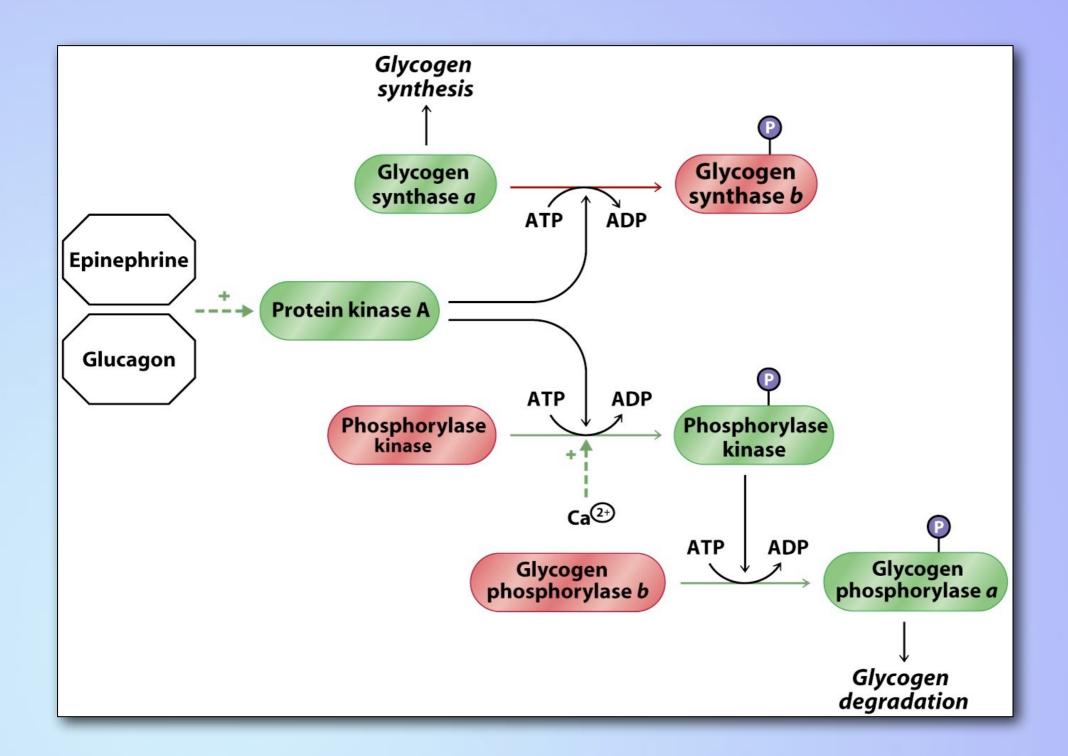


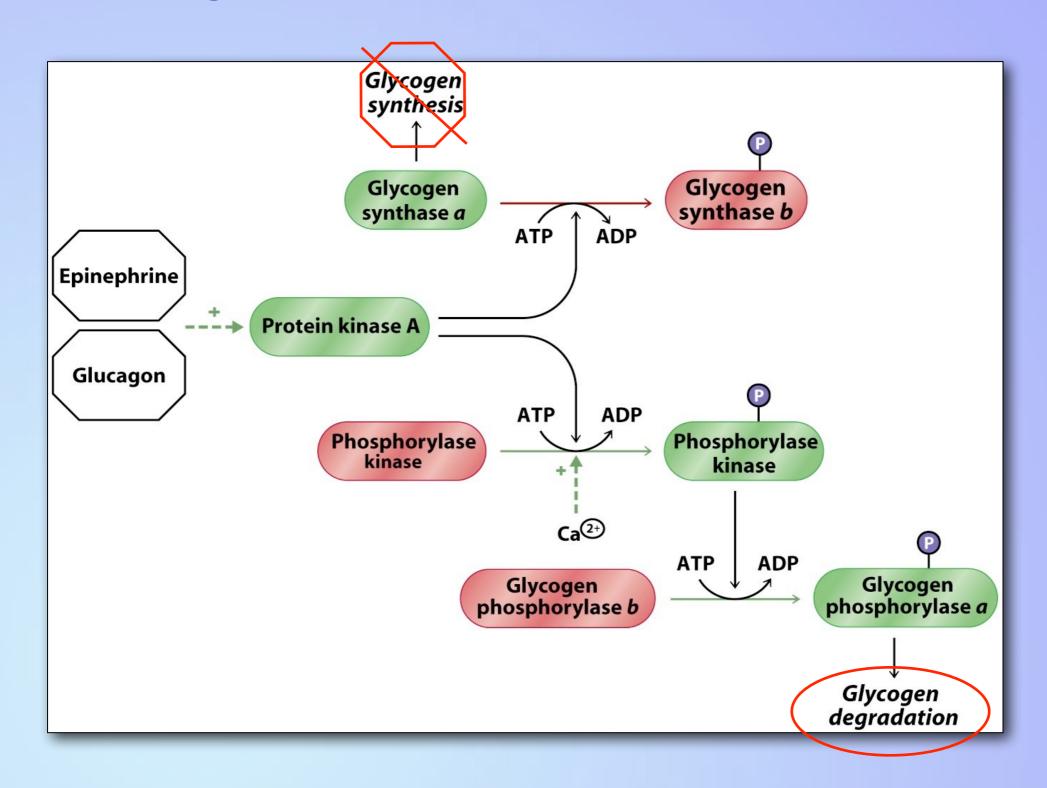


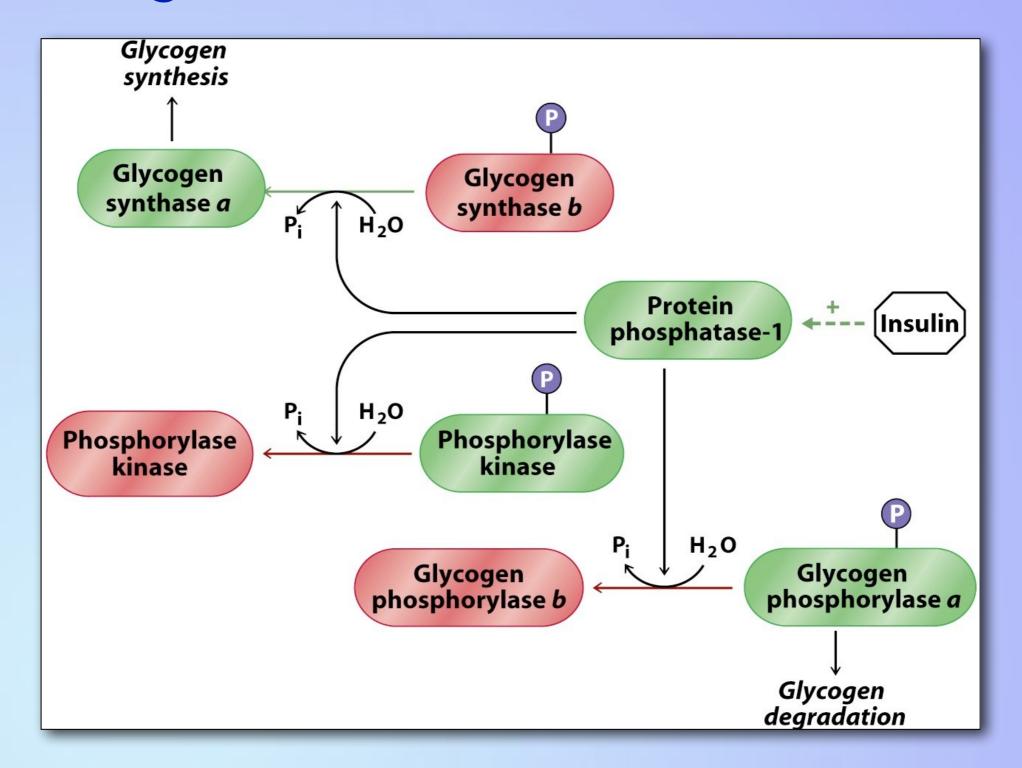
Both activate glycogen degradation by phosphorylating Glycogen synthase (inactive) and Glycogen phosphorylase (active)

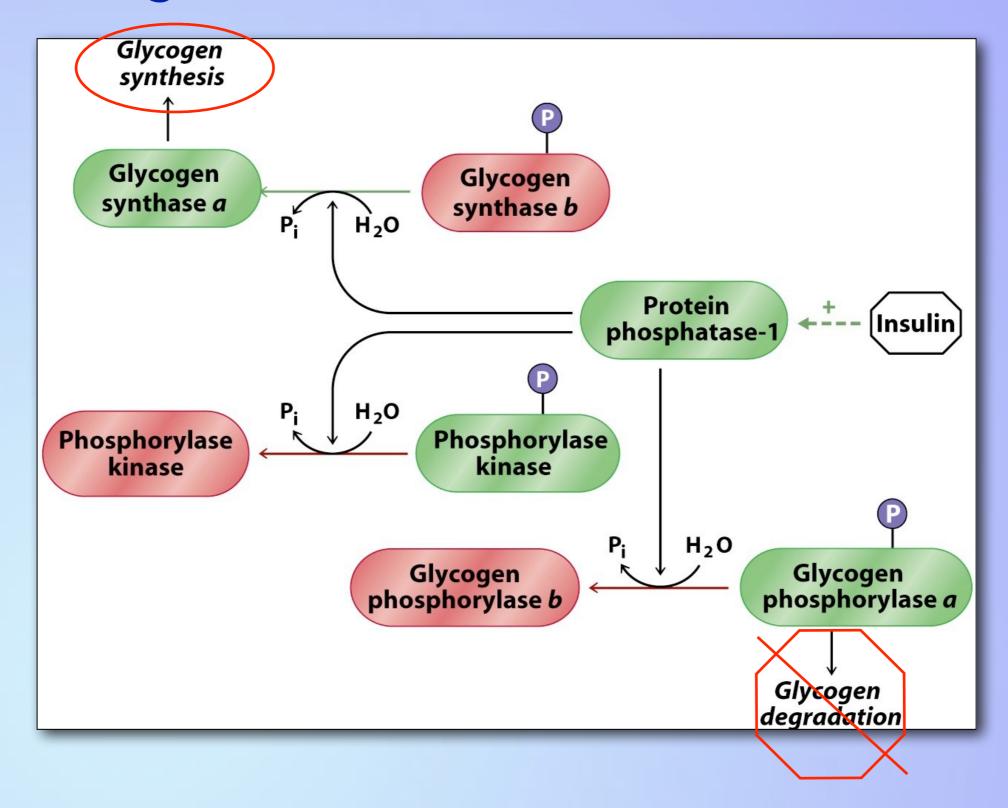
Both inhibit the action of insulin by phosphorylating the insulin receptor

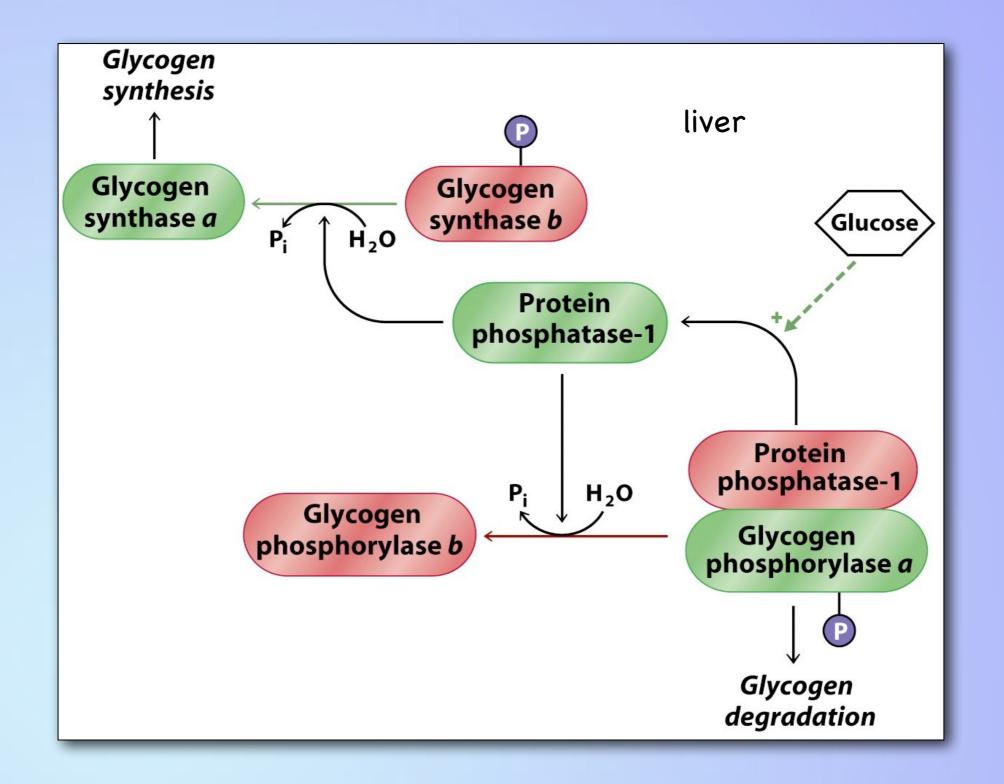
Chem 352, Lecture 8, Part II: Gluconeogenesis, Pentose Phosphate & Glycogen

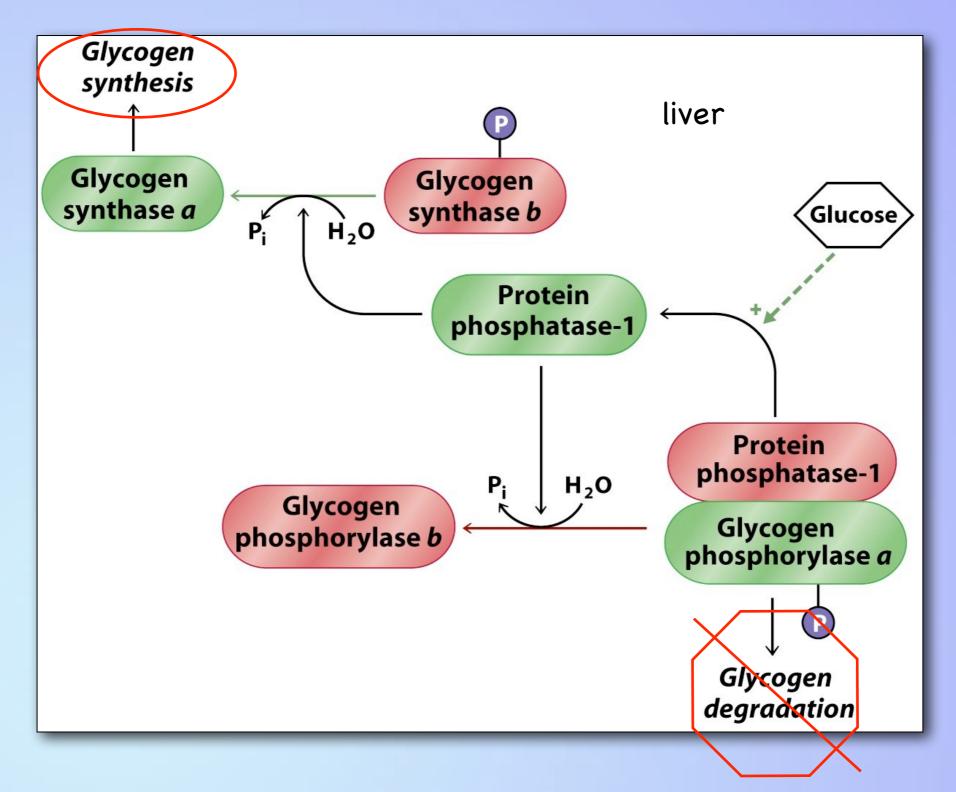




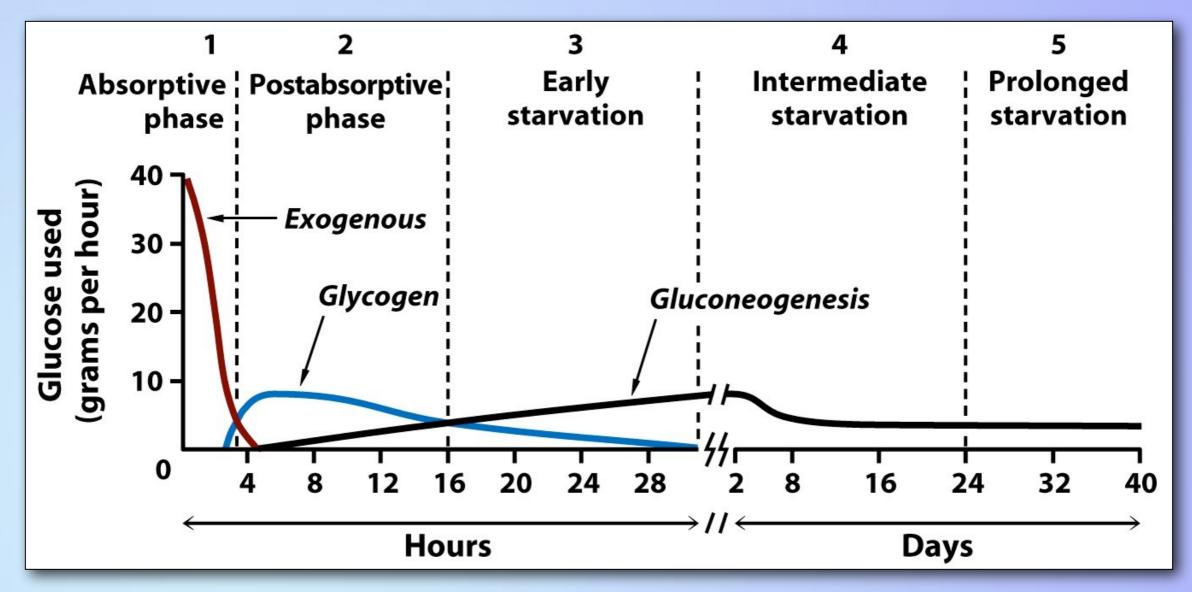








- ·Time since last meal
  - + 40 days in the desert after eating a sugary dessert.



#### Question:

Tumor cells often lack an extensive capillary network and must function under conditions of limited oxygen supply. Explain why these cancer cells take up far more glucose than other tissues and may overproduce some of the glycolytic enzymes?

#### Question:

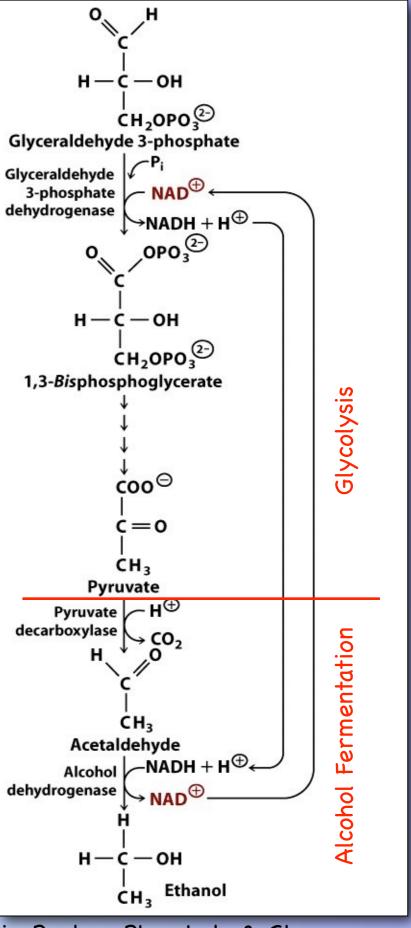
Rapid glycolysis during strenuous exercise provides the ATP needed for muscle contraction.

- A. Since the lactate dehydrogenase reaction does not produce any ATP, would glycolysis be more efficient if pyruvate rather than lactate were the end product?
- B. What is the fate of the lactate that is produced in the muscles during strenuous exercise?

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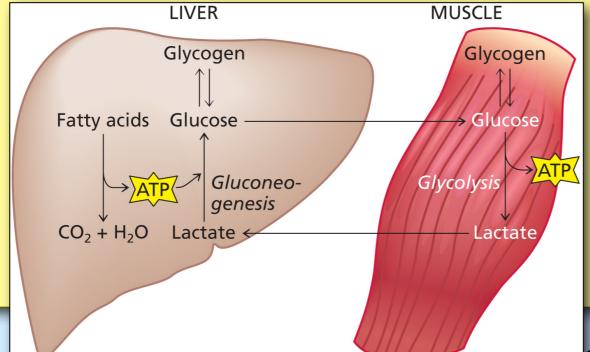
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Jeogenesis, Pentose Phosphate & Glycogen

#### Question:

Epinephrine promotes the utilization of stored glycogen for glycolysis and ATP production in muscles. How does epinephrine promote the use of liver glycogen stores for the generating the energy needed by contracting muscles?

Question: Epinephrine promotes the utilization of stored alycogen for Epinephrine Epinephrine Insulin **Epinephrine** Glucagon 000000 **β-Adrenergic** Glucagon receptor  $\alpha_1$ -Adrenergic receptor Insulin (liver and muscle) receptor receptor (liver) 00000 Glycogen synthesis **Protein kinase A Protein kinase C** Glycogen degradation

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#### Question:

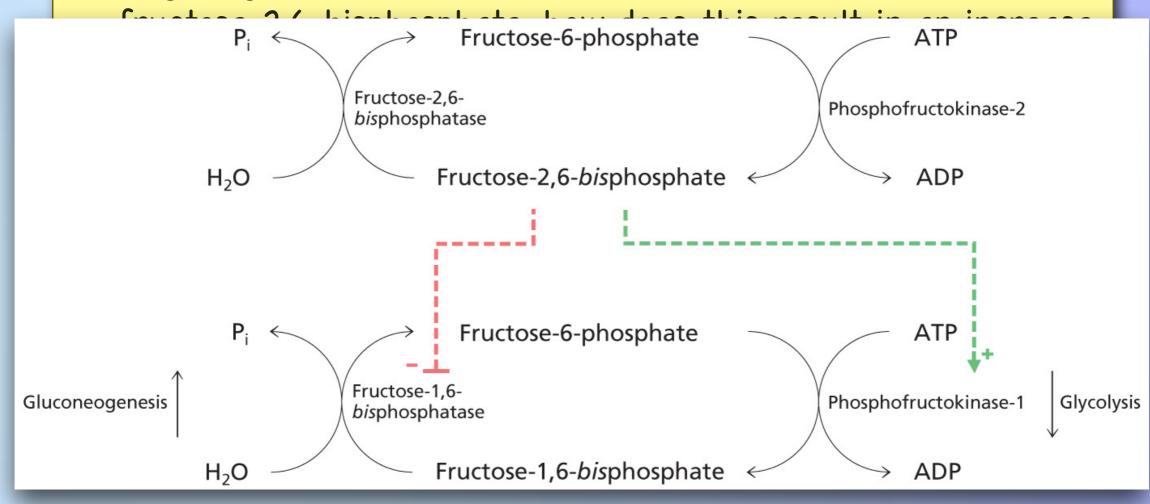
The polypeptide hormone glucagon is released from the pancreas in response to low blood glucose levels. In liver cells, glucagon plays a major role in regulating the rates of the opposing glycolysis and gluconeogenesis pathways by influencing the concentrations of fructose 2,6-bisphosphate.

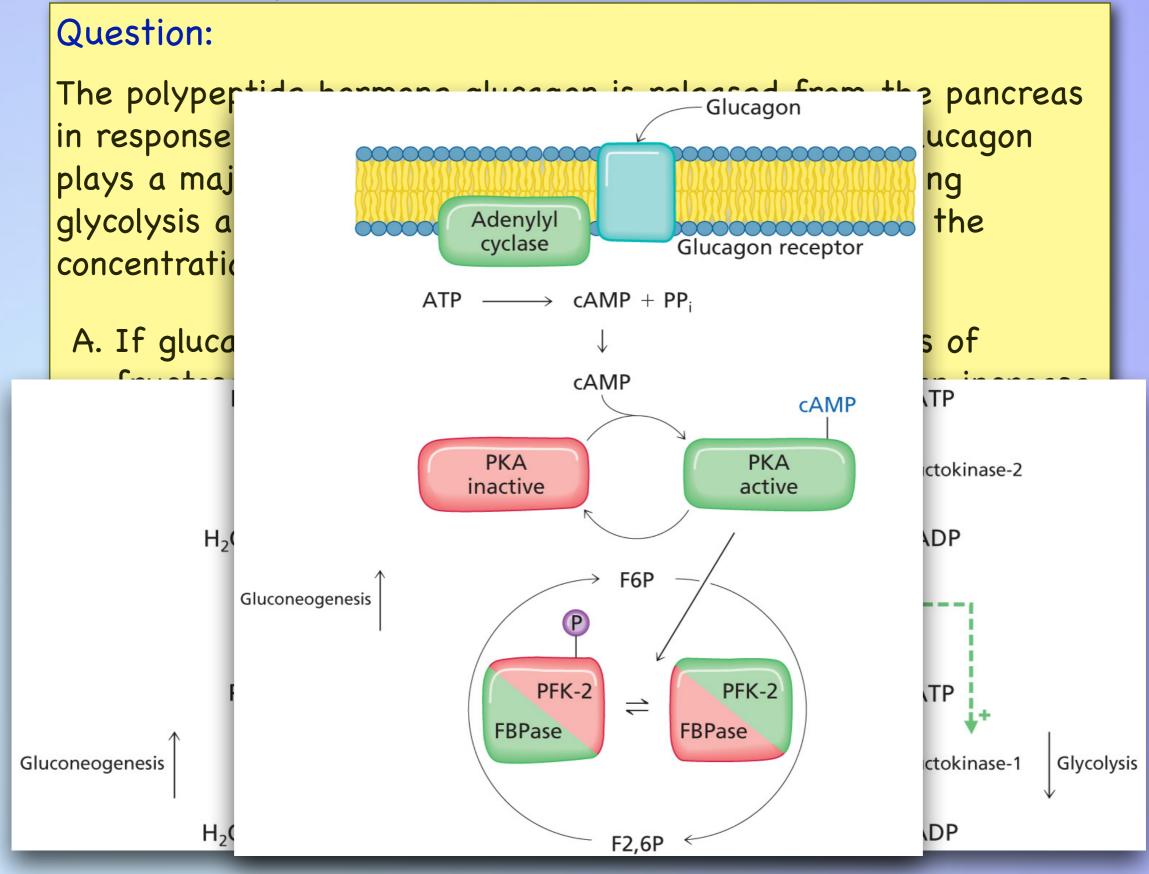
A. If glucagon causes a decrease in the concentrations of fructose 2,6-bisphosphate, how does this result in an increase in blood glucose levels?

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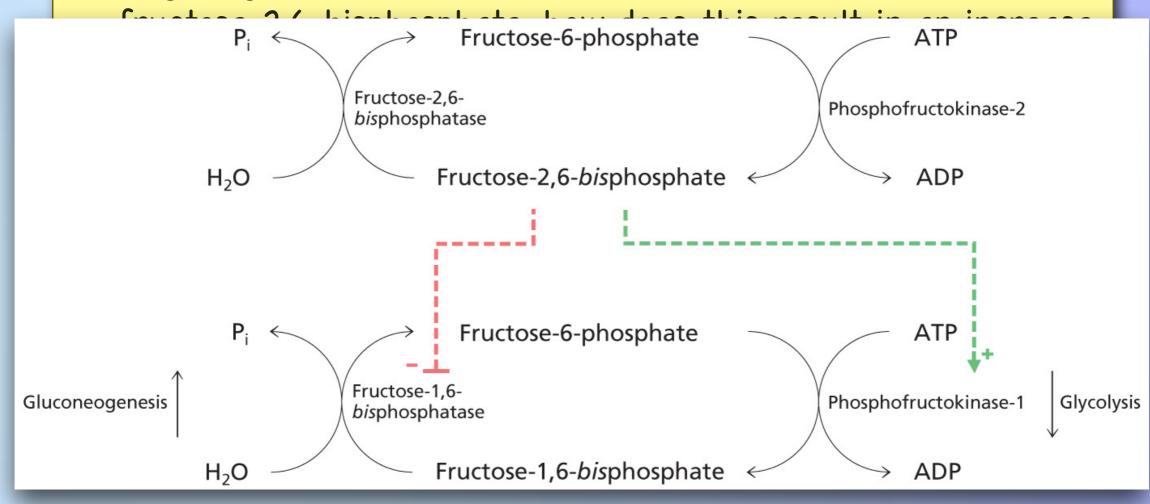




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#### Question:

The pentose phosphate pathway and the glycolytic pathway are interdependent, since they have in common several metabolites whose concentrations affect the rates of enzymes in both pathways.

A. Which metabolites are common to both pathways?

## Next Up

### ·Lecture 8 - Carbohydrate Metabolism

+ Part III: Citric Acid Cycle (Chapter 13)