

OUTLINE:

Introduction and review
Transport
Glycogenolysis
Glycolysis
Other sugars
Pasteur: Anaerobic vs Aerobic

Exam-1 material

Fermentations

Exam-2 material

Pyruvate

Krebs' Cycle

Oxidative Phosphorylation

Electron transport

Chemiosmotic theory: Phosphorylation

Fat Catabolism

Exam-3 material

Fatty acid Catabolism

Mobilization from tissues (mostly adipose)

Activation of fatty acids

Transport; carnitine

Oxidation: β -oxidation, 4 steps:

Protein Catabolism

Amino-Acid Degradation

Dealing with the nitrogen; Urea Cycle

Dealing with the carbon; Seven Families

Nucleic Acid & Nucleotide Degradation

PHOTOSYNTHESIS:

Overview of Photosynthesis

Key experiments:

Light Reactions

energy in a photon

pigments

HOW

Light absorbing complexes-"red-drop experiment"

Reaction center

Photosystems (PS)

PSII - oxygen from water splitting

PSI - NADPH

Proton Motive Force - ATP

Overview of light reactions

ANABOLISM I: Carbohydrates

Carbon Assimilation - Calvin Cycle

Stage One - Rubisco

Carboxylase

Oxygenase

Glycolate cycle

Stage Two - making sugar

Stage Three - remaking Ru 1,5P₂

Overview and regulation

Calvin cycle connections to biosynthesis

C4 versus C3 plants

Kornberg cycle - glyoxylate

Carbohydrate Biosynthesis in Animals

precursors

Cori cycle

Gluconeogenesis

reversible steps

irreversible steps - four

energetics

2-steps to PEP in mitochondria: Pyr carboxylase-biotin & PEPCK

FBPase

G6Pase

Glycogen Synthesis

UDP-Glc

Glycogen synthase

branching

Pentose-Phosphate Pathway

oxidative-NADPH

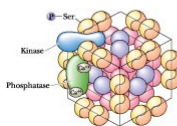
non-oxidative-Ribose 5-P

Regulation of Carbohydrate Metabolism

Anaplerotic reactions

Regulation of Carbohydrate Metabolism

Regulation of Pyruvate Dehydrogenase Complex



- Allosteric regulation by energy charge and substrate/product

-ADP & pyruvate **activates**

-ATP/NADH & acetyl-CoA **inhibit**

- Regulated by reversible phosphorylation of E1

-phosphorylation: **inactive**

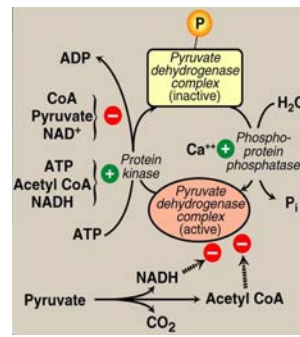
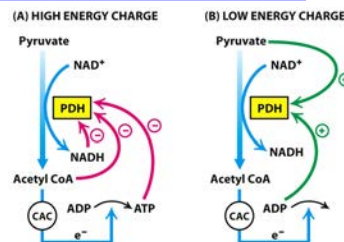
-dephosphorylation: **active**

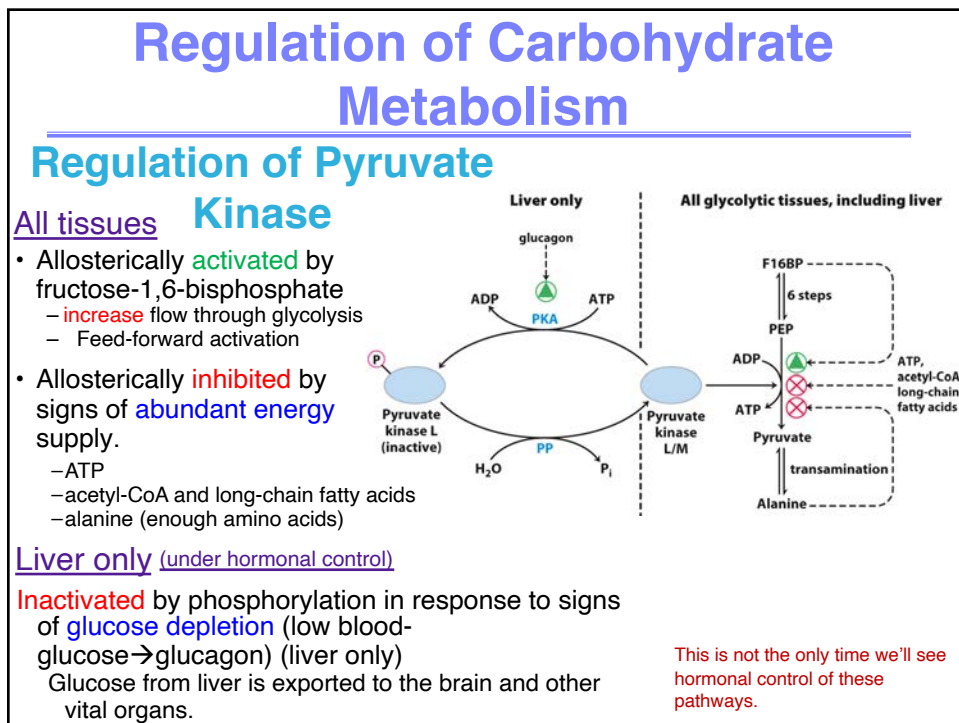
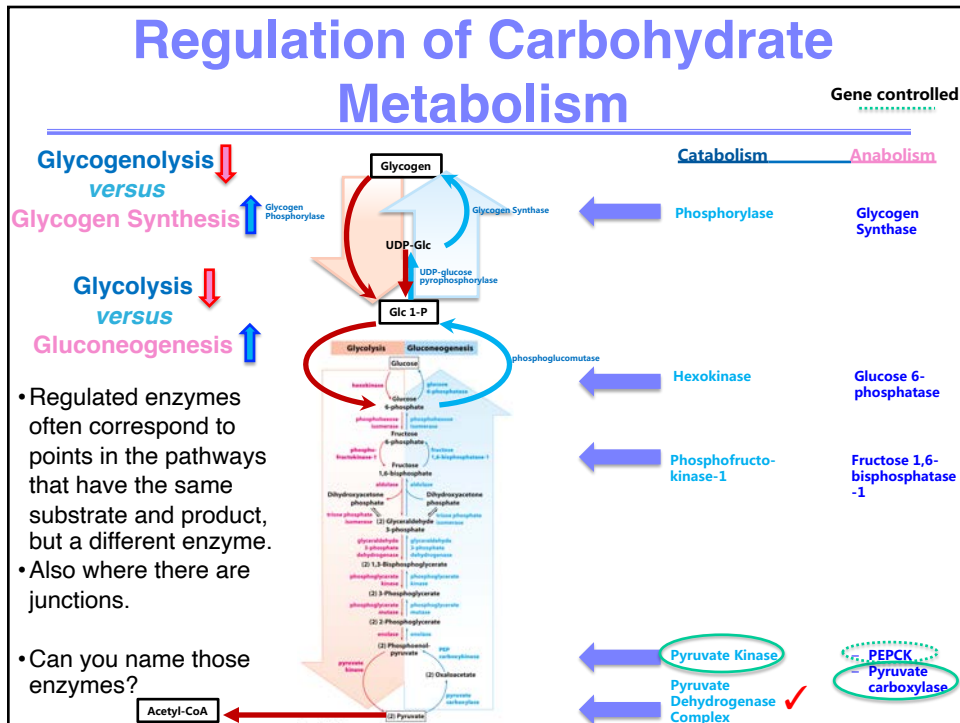
- PDH kinase and PDH phosphatase are part of mammalian PDH complex.

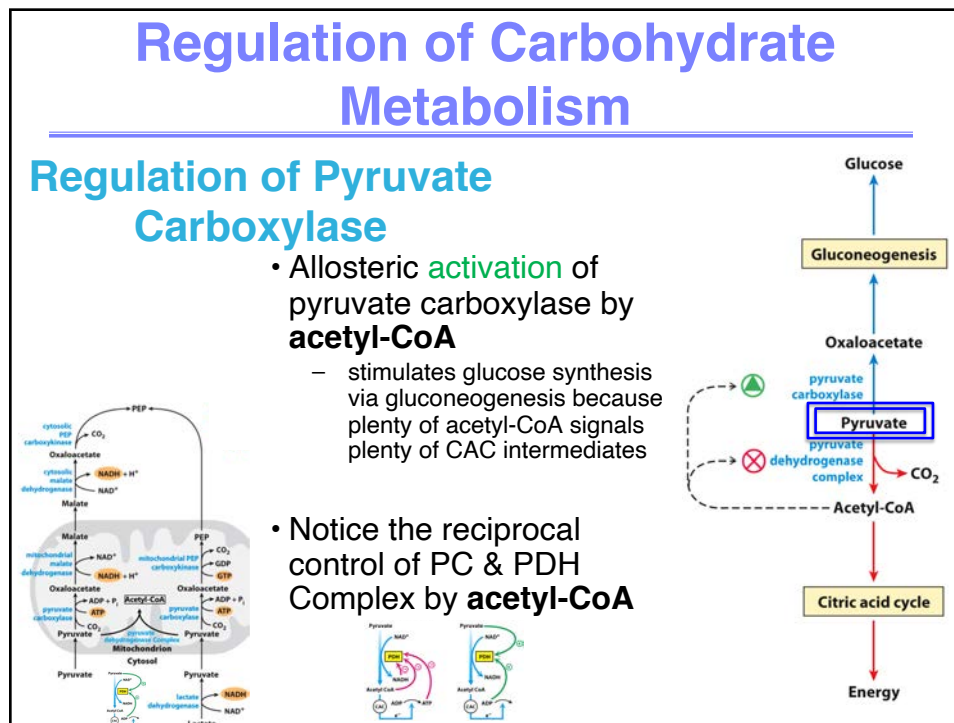
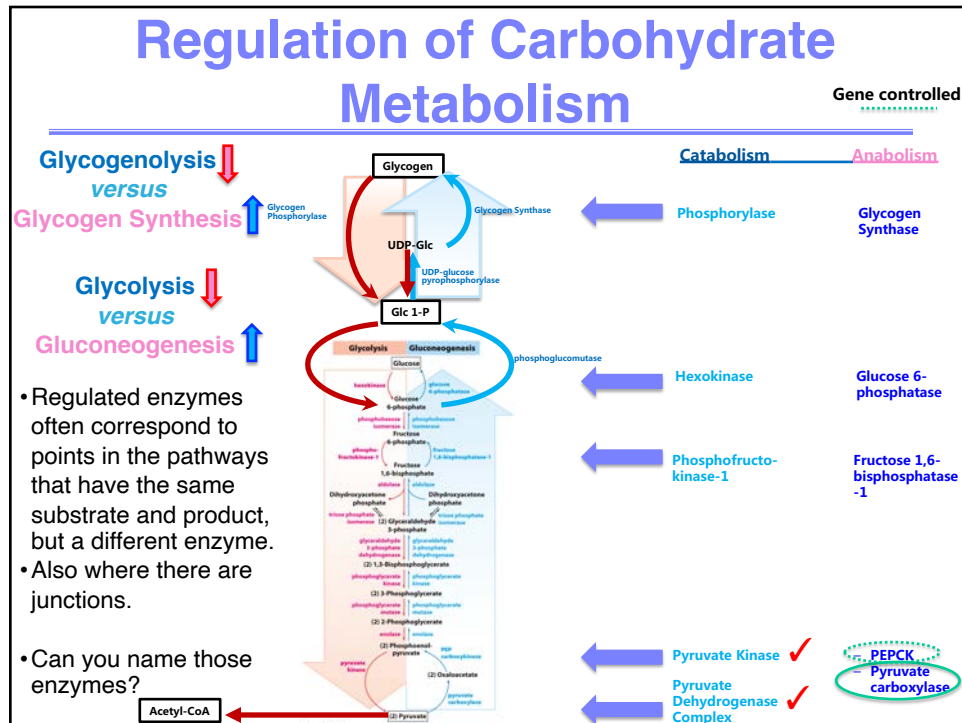
-Kinase is activated by ATP.

• high ATP \rightarrow phosphorylated PDH \rightarrow less acetyl-CoA made

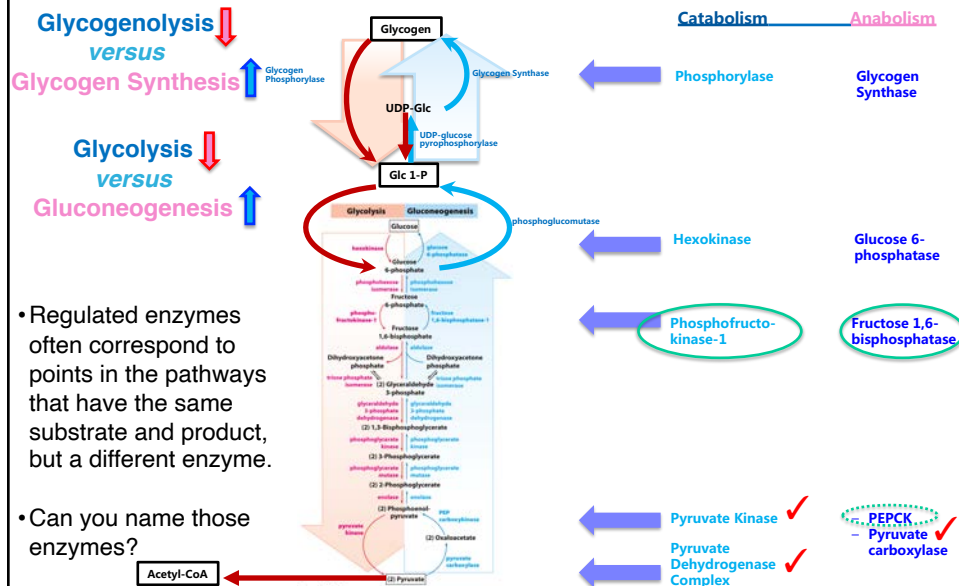
• low ATP \rightarrow kinase is less active and phosphoprotein phosphatase removes phosphate from PDH \rightarrow more acetyl-CoA made



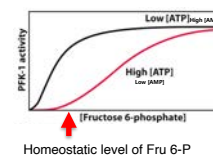




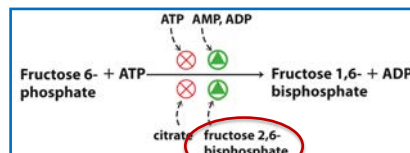
Gene controlled



Regulation of Phosphofructokinase-1 *versus*

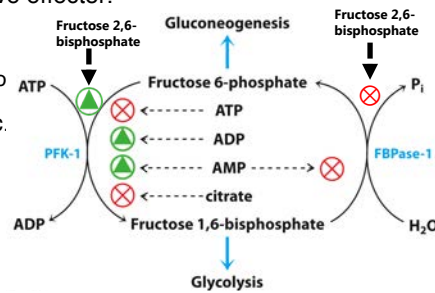


- Fructose-6-phosphate \rightarrow fructose 1,6-bisphosphate is the **commitment step in glycolysis**
 - While ATP is a substrate, ATP is also a negative effector.
 - Do not spend glucose in glycolysis if there is plenty of ATP.
 - Same for citrate, if there is plenty of citrate, do not waste glucose
 - Low energy charge inhibits biosynthesis of Glc.
-
- The diagram illustrates the regulation of glycolysis and gluconeogenesis. It shows the interconversion of Fructose 6-phosphate and Fructose 2,6-bisphosphate. Glycolysis (downward arrow) is activated by ATP (green triangle) and inhibited by ATP (red X). Gluconeogenesis (upward arrow) is inhibited by Fructose 2,6-bisphosphate (red X) and activated by ATP (green triangle). The conversion of Fructose 2,6-bisphosphate to Fructose 6-phosphate is activated by P_i (green triangle) and inhibited by ATP (red X).



Is this a typo?

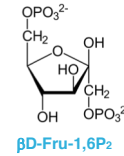
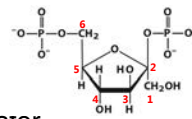
Bumble bees are missing an FBPase that responds to AMP



- **Go glycolysis** if AMP is high and ATP is low.
- **Go gluconeogenesis** if AMP is low.

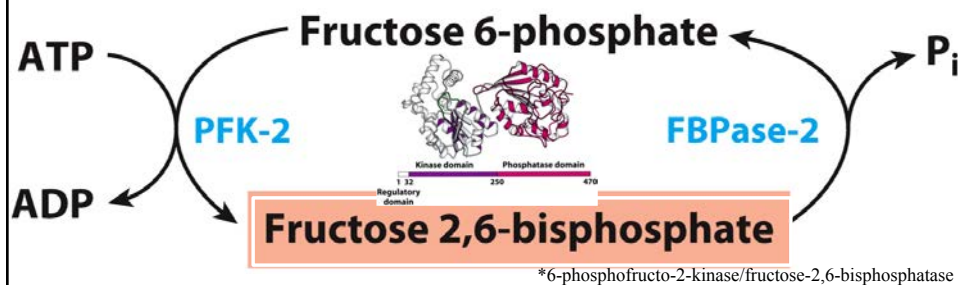
Regulation of Carbohydrate Metabolism

Fructose 2,6-(bis)phosphate (β D-Fru-2,6P₂)



- **NOT** a glycolytic intermediate, only a regulator
- Produced specifically to regulate glycolysis and gluconeogenesis
 - **activates** phosphofructokinase-1 (PFK-1) (glycolysis)
 - **inhibits** fructose 1,6-bisphosphatase (FBPase-1) (gluconeogenesis)

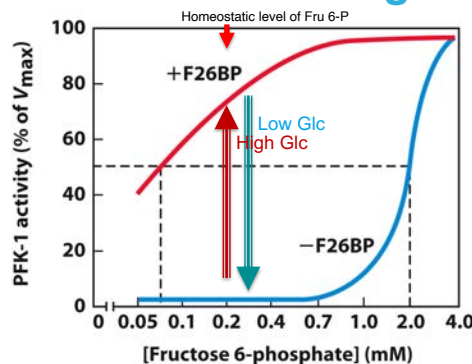
Enzyme for synthesis and degradation of Fru 2,6P₂ done with a dual-function enzyme: **PFK-2/FBPase-2***



*6-phosphofructo-2-kinase/fructose-2,6-bisphosphatase

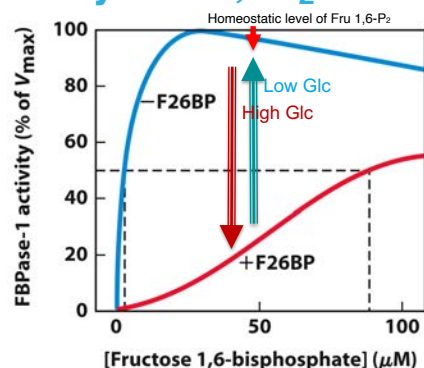
Regulation of Carbohydrate Metabolism

Regulation of Glycolysis and Gluconeogenesis by Fru-2,6P₂



- Without Fru 2,6P₂, **STOP** glycolysis
- **GO** gluconeogenesis.

This would be the state of **low Glc**



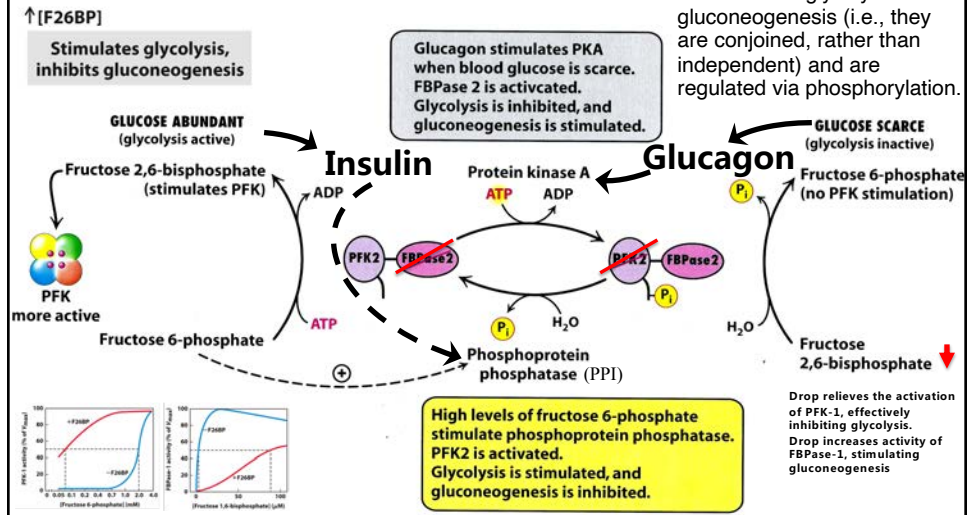
- With Fru 2,6P₂ (130 nM), **GO** glycolysis.
- With Fru 2,6P₂ (1300 nM), **STOP** gluconeogenesis.

This would be the state of **high Glc**

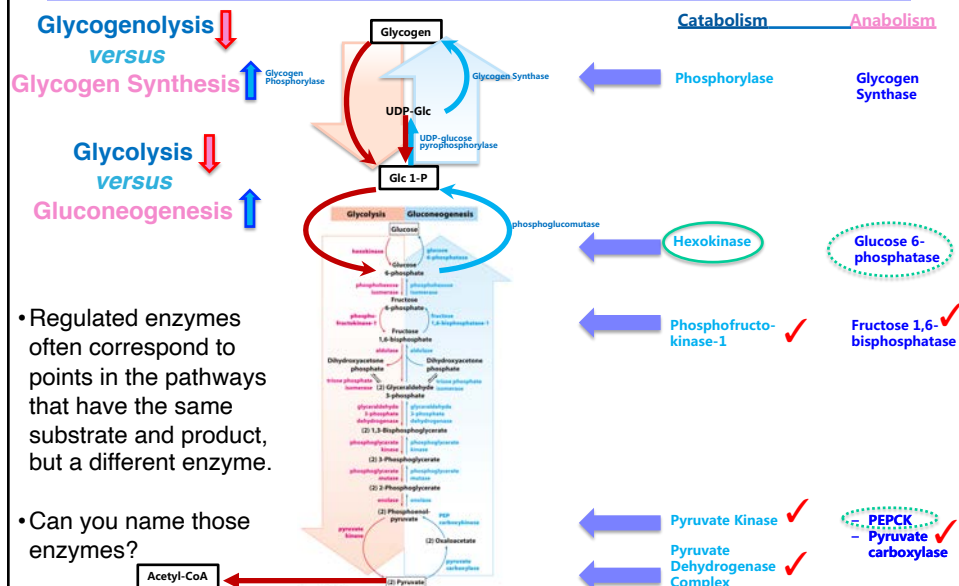
What controls PFK-2/FBPase-2?

Regulation of Carbohydrate Metabolism

Regulation of Fru-2,6-P₂ Levels



Regulation of Carbohydrate Metabolism

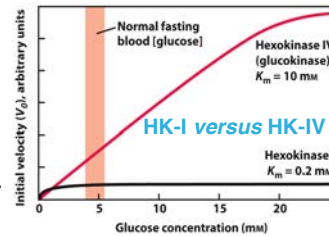


Regulation of Carbohydrate Metabolism

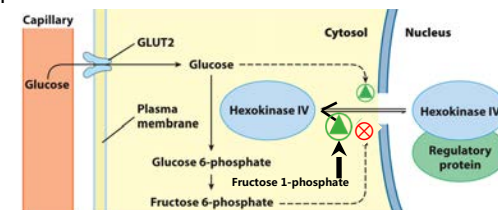
Regulation of Hexokinase

There Are Four Isozymes of Hexokinase (I-IV)

- Isozymes are different enzymes that catalyze the **same reaction**.
 - typically share **similar sequences**
 - **may have different kinetic properties**
 - can be regulated differently
- HK I is expressed in all tissues, to different levels.
- HK IV (glucokinase) is only expressed in the liver and pancreas.
 - has higher K_m , so responsive to higher [glucose]
 - not inhibited by glucose-6-phosphate, so can function at higher [glucose]
 - functions to clear blood glucose at higher [glucose] for storage as glycogen **also Fru1P**
 - Glc activates release/Fru6P inhibits

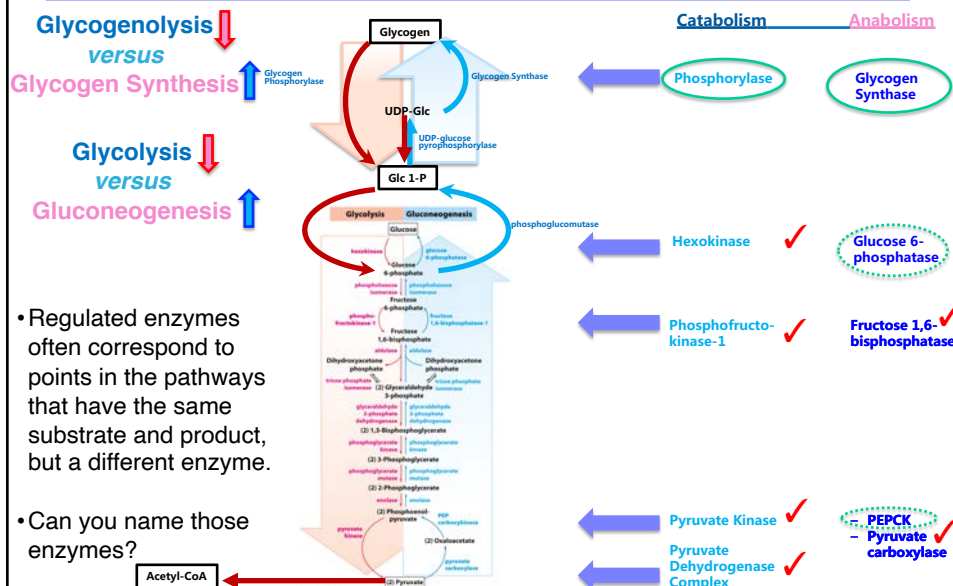


Glucokinase Is Regulated by Sequestration



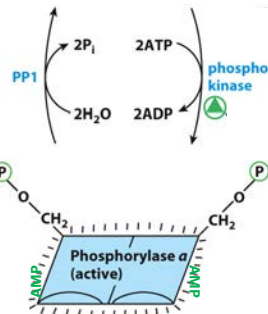
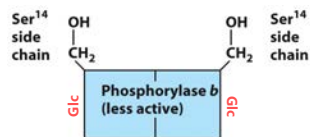
Regulation of Carbohydrate Metabolism

Gene controlled

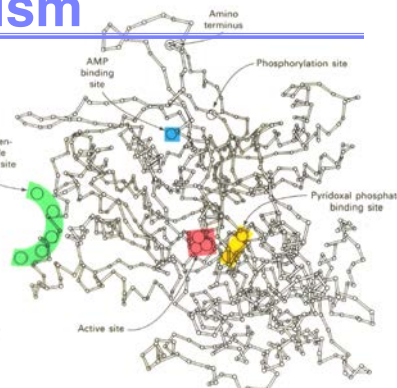


Regulation of Carbohydrate Metabolism

Regulation of Glycogen Phosphorylase



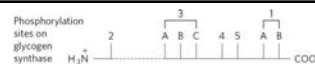
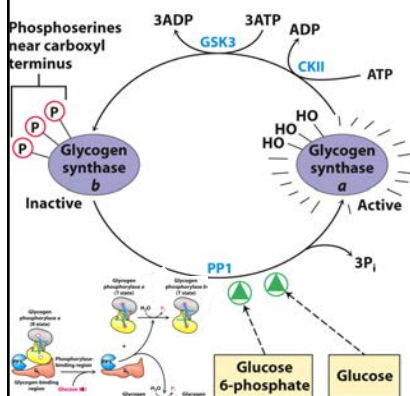
Nobel Prize 1972
Earl Sutherland
1915-1974



- Glycogen phosphorylase cleaves glucose residues off glycogen, generating glucose-1-phosphate (Glc 1P).
- Phosphorylation **activates** glycogen phosphorylase-b
 - Phosphorylase-b Kinase
 - Accentuated by allosteric binding of AMP (muscle only)
- Dephosphorylation **inhibits** glycogen phosphorylase-a
 - Phosphoprotein phosphatase-1 (PP1)
 - Accentuated by allosteric binding of Glc (in liver only)

Regulation of Carbohydrate Metabolism

Regulation of Glycogen Synthase



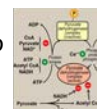
Kinase	Phosphorylation sites	Degree of synthase inactivation
Protein kinase A	1A, 1B, 2, 4	+
Protein kinase G	1A, 1B, 2	+
Protein kinase C	1A	+
Ca ²⁺ /calmodulin kinase	1B, 2	+
Phosphorylase b kinase	2	+
Casein kinase I	At least nine	+++
Casein kinase II	5	0
Glycogen synthase kinase 3	3A, 3B, 3C	+++
Glycogen synthase kinase 4	2	+

Need energy: stop storing & release

- Glycogen synthase adds glucose residues to glycogen using UDP-Glc.
- Phosphorylation **inhibits** glycogen synthase-a
 - Its complicated, responding to multiple signals
 - Example: First Casein Kinase-2 (CKII), then Glycogen Synthase Kinase-3 (GSK3) [inactivated by insulin]

- Dephosphorylation **activates** glycogen synthase-b
 - Phosphoprotein phosphatase-1 (PP1) (in liver it's a different PP)
 - PP1 is bound to GS-b
- Also, feedforward control by glucose and Glc-6P
 - Binding causes a conformation favorable for PP-1 binding
 - Binding does not allow GSK-3 access to phosphorylation sites

Regulation of glycogen synthase/glycogen phosphorylase is somewhat similar to regulation of PDH complex



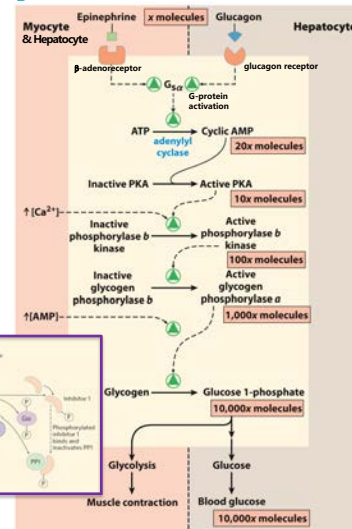
Regulation of Carbohydrate Metabolism

Glycogen Phosphorylase Cascade



Nobel Prize 1992
Edwin Krebs
1918-2009

- **Glucagon/epinephrine** signaling pathway **activated** when there is a **NEED** for energy
 - starts phosphorylation cascade via cAMP
 - cAMP **activates** PKA
 - PKA **activates** phosphorylase-b kinase
 - this kinase **activates** glycogen phosphorylase
 - Massive degradation of glycogen
 - In muscle: Glc1P → Glc6P → glycolysis
 - In liver: Glc1P → Glc6P → Blood glucose
- There is reciprocal **inhibition** of GS
 - PKA phosphorylates G_M , which is bound to PP1 on GS, thus dissociating it.
 - PKA also phosphorylates **PP1-inhibitor protein**, which binds and **inactivates** the free PP1, thus leaving $GS-\textcircled{C}$ and inactive



Regulation of Carbohydrate Metabolism

Gene controlled

