# Glucose Metabolism (Glycolysis)

### **Objectives**

By the end of this lecture, students are expected to:

- □ Recognize glycolysis as the major oxidative pathway of glucose
- ☐ List the main reactions of glycolytic pathway
- □ Discuss the rate-limiting enzymes/Regulation
- ☐ Assess the ATP production (aerobic/anaerobic)
- ☐ Define pyruvate kinase deficiency hemolytic anemia
- □Discuss the unique nature of glycolysis in RBCs.

### **Glycolysis: An Overview**

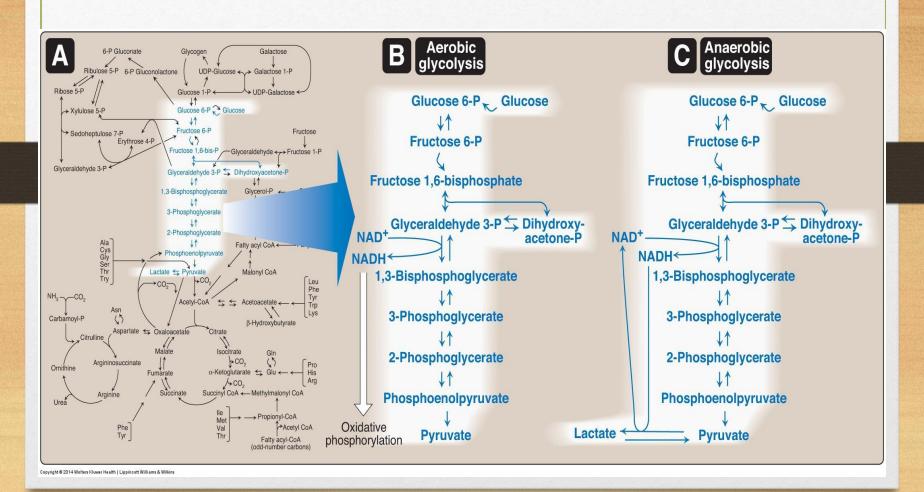
- Glycolysis, the major pathway for glucose oxidation, occurs in the cytosol of all cells.
- ❖ It is unique, in that it can function either aerobically or anaerobically, depending on the availability of oxygen and intact mitochondria.
- ❖ It allows tissues to survive in presence or absence of oxygen, e.g., skeletal muscle.
- \* RBCs, which lack mitochondria, are completely reliant on glucose as their metabolic fuel, and metabolizes it by anaerobic glycolysis.

### **Glycolysis**

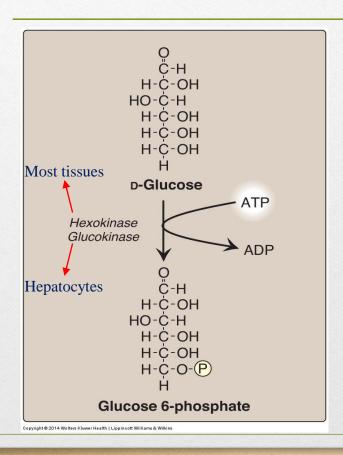
The product of one reaction is the substrate of the subsequent reaction. Glucose 6-P Glucose Fructose 6-P Fructose 1,6-bisphosphate acetone-P 1,3-Bisphosphoglycerate 3-Phosphoglycerate 2-Phosphoglycerate Phosphoenolpyruvate 

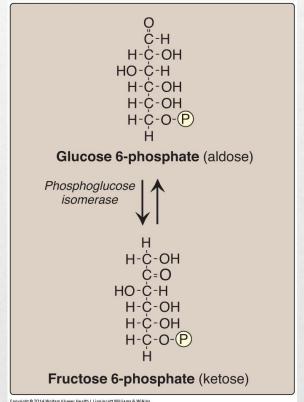
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### **Aerobic Vs Anaerobic Glycolysis**



### **Aerobic Glycolysis** (1st and 2nd reactions)

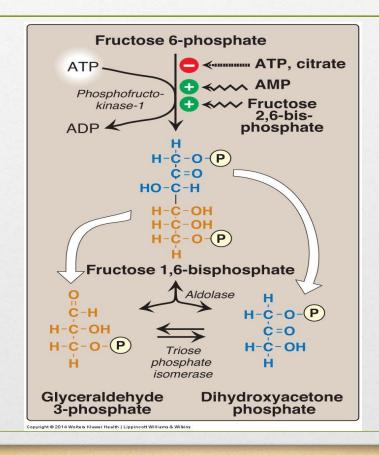




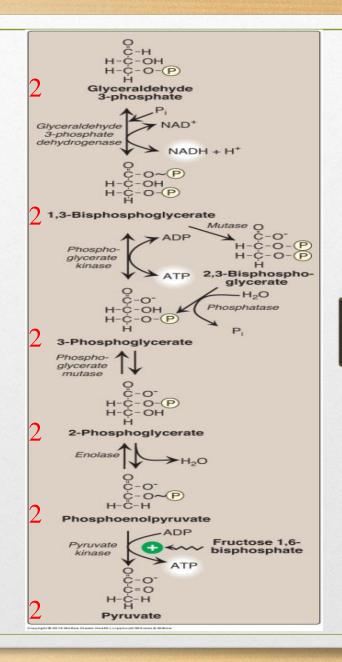
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## **Aerobic Glycolysis**

(Reactions: 3<sup>rd</sup> - 5<sup>th</sup>)



## Aerobic Glycolysis (Reactions: 6<sup>th</sup> – 10<sup>th</sup>)



## Regulation: Glucokinase/Hexokinase

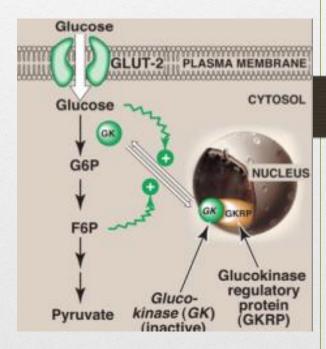
• Hexokinase – it is inhibited by the reaction product, glucose-6-P which accumulates when further metabolism of this hexose is reduced

 Glucokinase – It is inhibited indirectly by Fructose-6-P and is indirectly stimulated by glucose

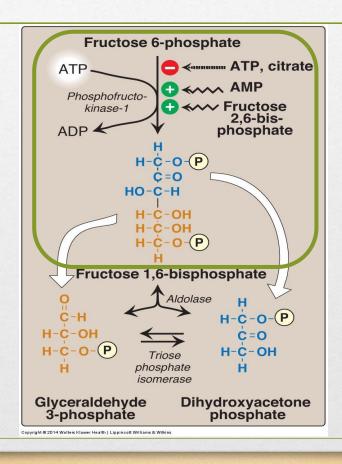
## Glucokinase (GK) Regulation

• In the presence of high fructose-6-phosphate, GK translocates and binds tightly to GKRP (glucokinase regulatory protein) in the nucleus, making it inactive

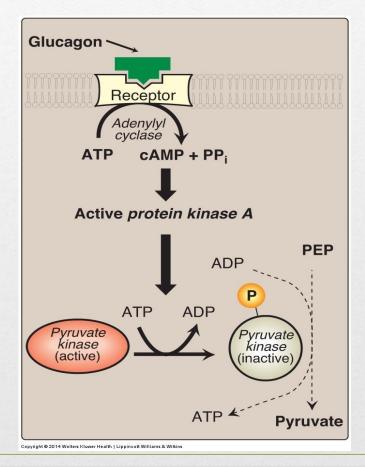
• When glucose levels are high in blood and hepatocytes (GLUT-2), GK is released from GKRP and enters the cytosol



## **Regulation: PFK-1**



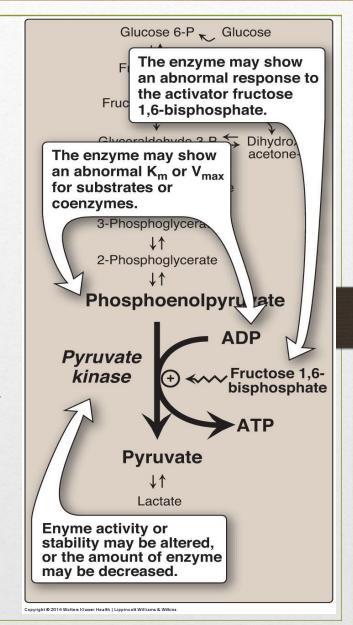
## Pyruvate Kinase Covalent Modification



## Pyruvate Kinase Deficiency Hemolytic Anemia

#### PK Mutation may lead to:

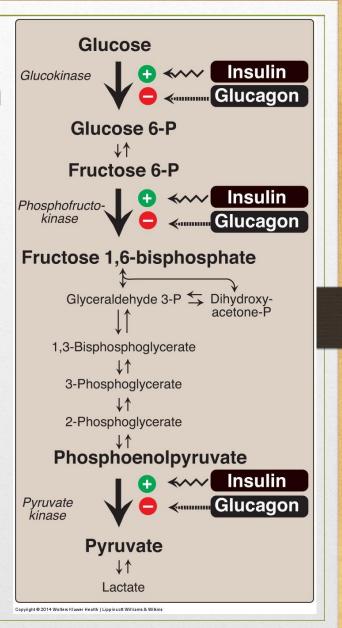
- 1. Altered Enz. Kinetics.
- 2. Altered response to activator.
- 3. Decreased the amount of the Enz. or its stability



## Long-Term Regulation of Glycolysis

**Insulin:** Induction

Glucagon: Repression



## **Summary** (Regulation of Glycolysis)

#### Regulatory Enzymes (Irreversible reactions):

Glucokinase/hexokinase

PFK-1

Pyruvate kinase

#### **Regulatory Mechanisms:**

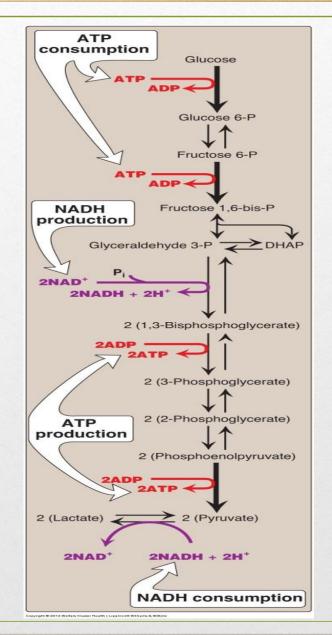
Rapid, short-term: Allosteric, Covalent modifications

Slow, long-term: Induction/repression

Apply the above mechanisms for each enzyme where applicable

## **Glycolysis**

For each NADH, 3 ATP will be produced by ETC in the mitochondria



## Substrate-level phosphorylation vs. Oxidative phosphorylation

- Phosphorylation is the metabolic reaction of introducing a phosphate group into an organic molecule.
- Oxidative phosphorylation: The formation of high-energy phosphate bonds by phosphorylation of ADP to ATP <u>coupled to</u> the transfer of electrons from reduced coenzymes to molecular oxygen via the electron transport chain (ETC); it occurs in the mitochondria.
- Substrate-level phosphorylation: The formation of high-energy phosphate bonds by phosphorylation of ADP to ATP (or GDP to GTP) coupled to cleavage of a high-energy metabolic intermediate (substrate). It may occur in cytosol or mitochondria

## Aerobic Glycolysis (Net ATP produced)

#### **ATP Consumed:**

2 ATP

#### **ATP Produced:**

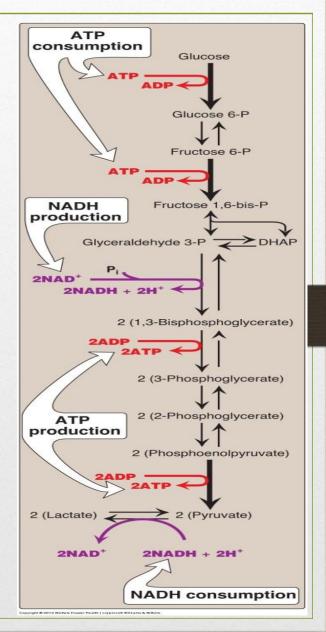
Substrate-level  $2 \times 2 = 4$  ATP Oxidative-level  $2 \times 3 = 6$  ATP Total 10 ATP

**Net:** 10 - 2 = 8 ATP

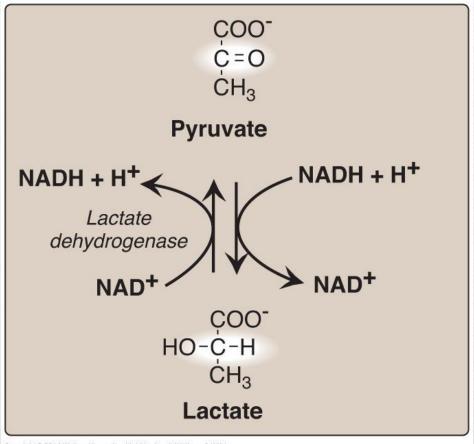
### **Anaerobic Glycolysis**

- NADH produced cannot be used by ETC for ATP production.
  (No O<sub>2</sub> and/or No mitochondria)
- Less ATP production, as compared to aerobic glycolysis.
- Lactate is an obligatory end product, Why?

Because if not formed, All cellular NAD<sup>+</sup> will be converted to NADH, with no means to replenish the cellular NAD  $\rightarrow$  Glycolysis stops  $\rightarrow$  death of the cell



### Lactate Dehydrogenase



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## Anaerobic Glycolysis (Net ATP produced)

#### **ATP Consumed:**

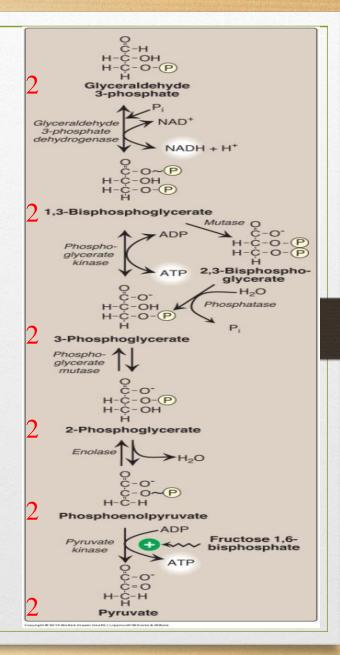
ATP

#### **ATP Produced:**

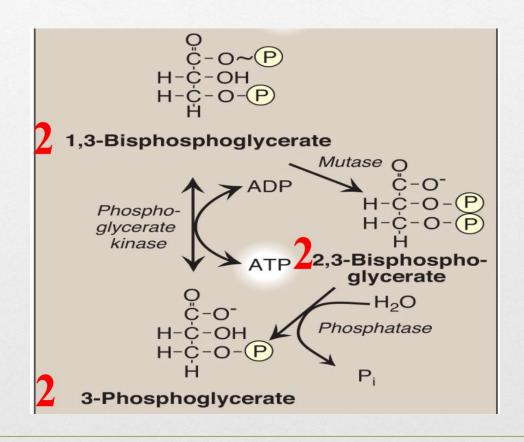
Substrate-level  $2 \times 2 = 4$  ATP Oxidative-level  $2 \times 3 = 6$  ATP Total 4 ATP

**Net:** 4-2=2 ATP

### Anaerobic Glycolysis in RBCs (2,3-BPG Shunt)



## Anaerobic Glycolysis in RBCs (2,3-BPG Shunt)



### Glycolysis in RBCs (Net ATP produced)

#### **ATP Consumed:**

**ATP** 

#### **ATP Produced:**

Substrate-level

or  $2 \times 2 = 4$ 

ATP

 $1 \times 2 = 2$ 

Oxidative-level

Total

 $2 \times 3 = 6$ 

ATP

4 or 2 ATP

Net:

4 - 2 = 2

ATP

2-2=0

ATP

## Glycolysis in RBCs (Summary)

#### **End product:**

Lactate

No net production or consumption of NADH

#### **Energy yield:**

If no 2,3-BPG is formed: 2 ATP

If 2,3-BPG shunt occurs: 0 ATP

#### PK Deficiency hemolytic anemia depends on:

Degree of PK Deficiency

Compensation by 2,3-BPG

## **Take Home Messages**

- ☐ Glycolysis is the major oxidative pathway for glucose
- ☐ Glycolysis is employed by all tissues
- ☐ Glycolysis is a tightly-regulated pathway
- □ PFK-1 is the rate-limiting regulatory enzyme
- ☐ Glycolysis is mainly a catabolic pathway for ATP production, But it has some anabolic features (amphibolic)
- ☐ Pyruvate kinase deficiency in RBCs results in hemolytic anemia

## **Take Home Messages**

**□** Net energy produced in:

Aerobic glycolysis: 8 ATP

Anaerobic glycolysis: 2 ATP

**☐** Net energy produced in glycolysis in RBCs:

Without 2,3 BPG synthesis: 2 ATP

With 2,3 BPG synthesis: 0 ATP

### Reference

Lippincott Illustrated Review of Biochemistry, 6<sup>th</sup> edition, 2014, Unit 2, Chapter 8, Pages 91-108.