

Major Metabolic Pathways of Glucose and Glucose Transport

**Clinical Chemistry Unit Pathology
Department
College of Medicine, KSU**

Objectives

- *By the end of the lecture, students are expected to:*
 - Define a metabolic pathway.
 - Describe the general metabolic pathways for glucose (production and utilization)
 - Briefly describe the HMP
 - Recognize the mechanisms of glucose transport

Metabolic Pathway

Definition

Site:

Cellular (tissue) and Subcellular

Reactions

Rate-limiting enzyme(s)

Regulatory mechanism(s):

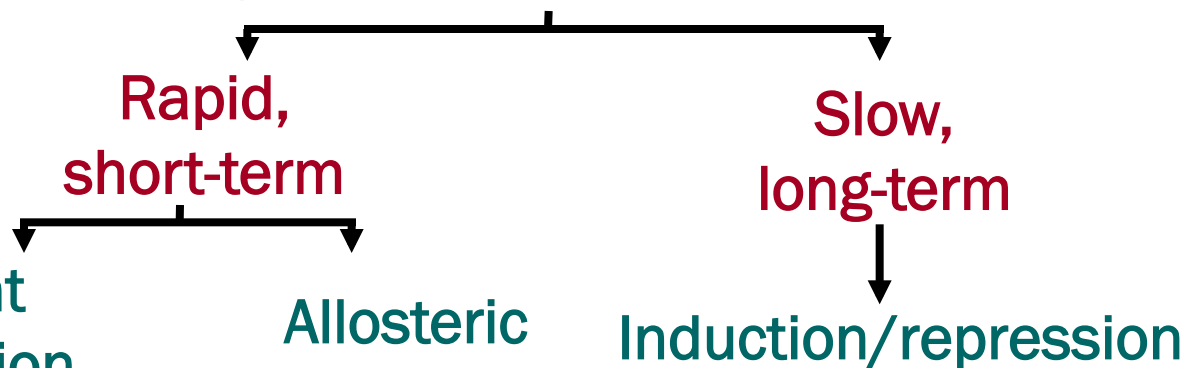
**Rapid,
short-term**

**Slow,
long-term**

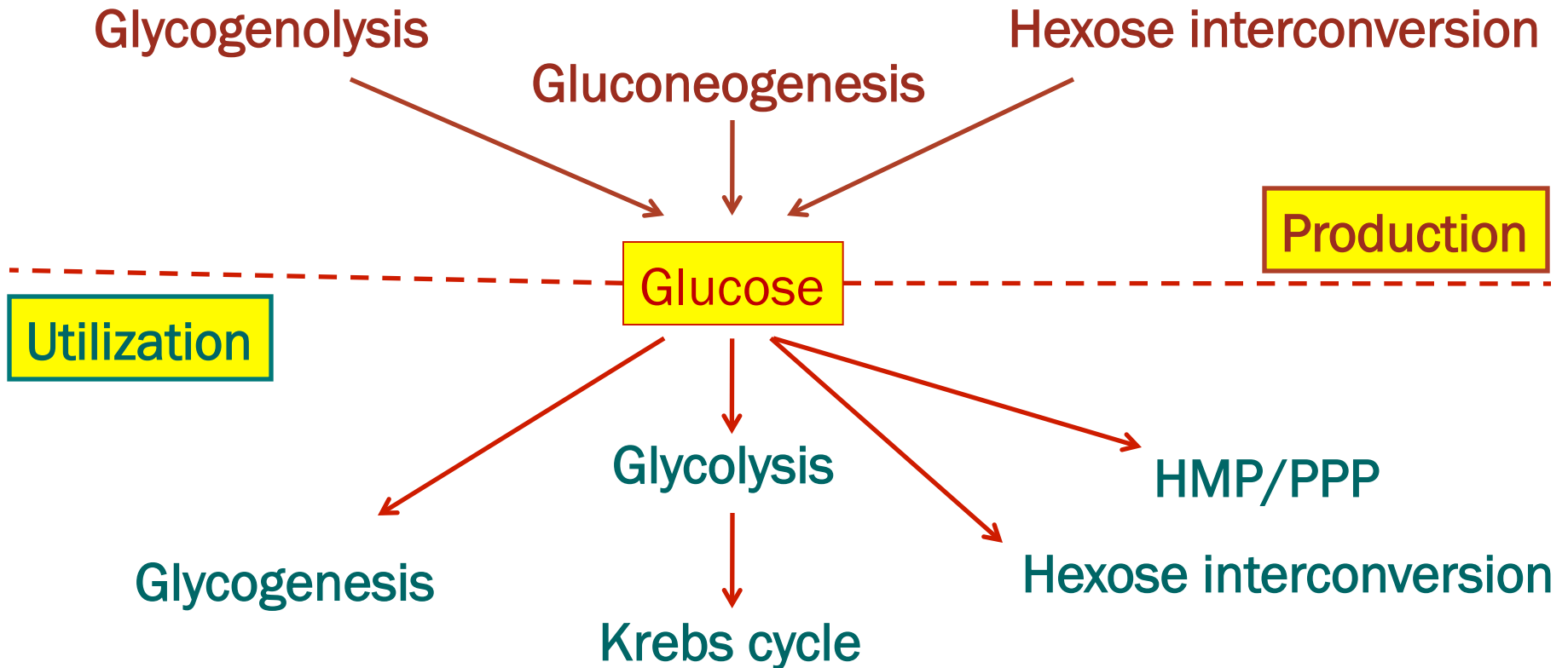
**Covalent
modification**

Allosteric

Induction/repression



Metabolic Pathways of Glucose-production and utilization



Metabolic Pathways of Glucose- catabolic and anabolic

Catabolic cycles

Glycolysis (Mainly)

Krebs (Mainly)

Glycogenolysis

HMP

Anabolic cycles

Gluconeogenesis

Glycogenesis

Glycolysis

- Oxidation of glucose to provide energy.
- Pyruvate is the end product of glycolysis in cells with mitochondria and an adequate supply of oxygen-
aerobic glycolysis
- In absence of oxygen and in cells that lack mitochondria, the end product is lactate- **anaerobic glycolysis**

Glycogenesis and Glycogenolysis

Glycogenesis:

Synthesis of glycogen from glucose
Mainly liver and muscle, Cytosol

Glycogenolysis

Degradation of glycogen into glucose
Mainly liver and muscle, Cytosol

Gluconeogenesis

- Synthesis of glucose from non-carbohydrate precursors.
- The precursors could be lactate, pyruvate, glycerol and alpha-keto acids.
- It requires both mitochondria and cytosolic enzymes
- Liver and kidney

Hexose Monophosphate shunt(HMP) or Pentose Phosphate Pathway (PPP)

- HMP shunt is an alternative pathway of glucose oxidation
- It is not involved in the generation of energy
- Around 10% of glucose is entered in this pathway
- In liver and kidney, this percentage is upto 30%

Biomedical Importance

- It has two main functions-
 1. Provides NADPH which is required for –
 - synthesis of fattyacids, steroid and some amino acids
 - Detoxification of drugs by cytochrome p450
 - In scavenging the free radicals
 2. Provides Pentoses
 - This pentose and its derivatives are useful in the synthesis of
 - Nucleic acids (DNA and RNA)
 - Nucleotides (ATP, NAD, FAD and CoA)

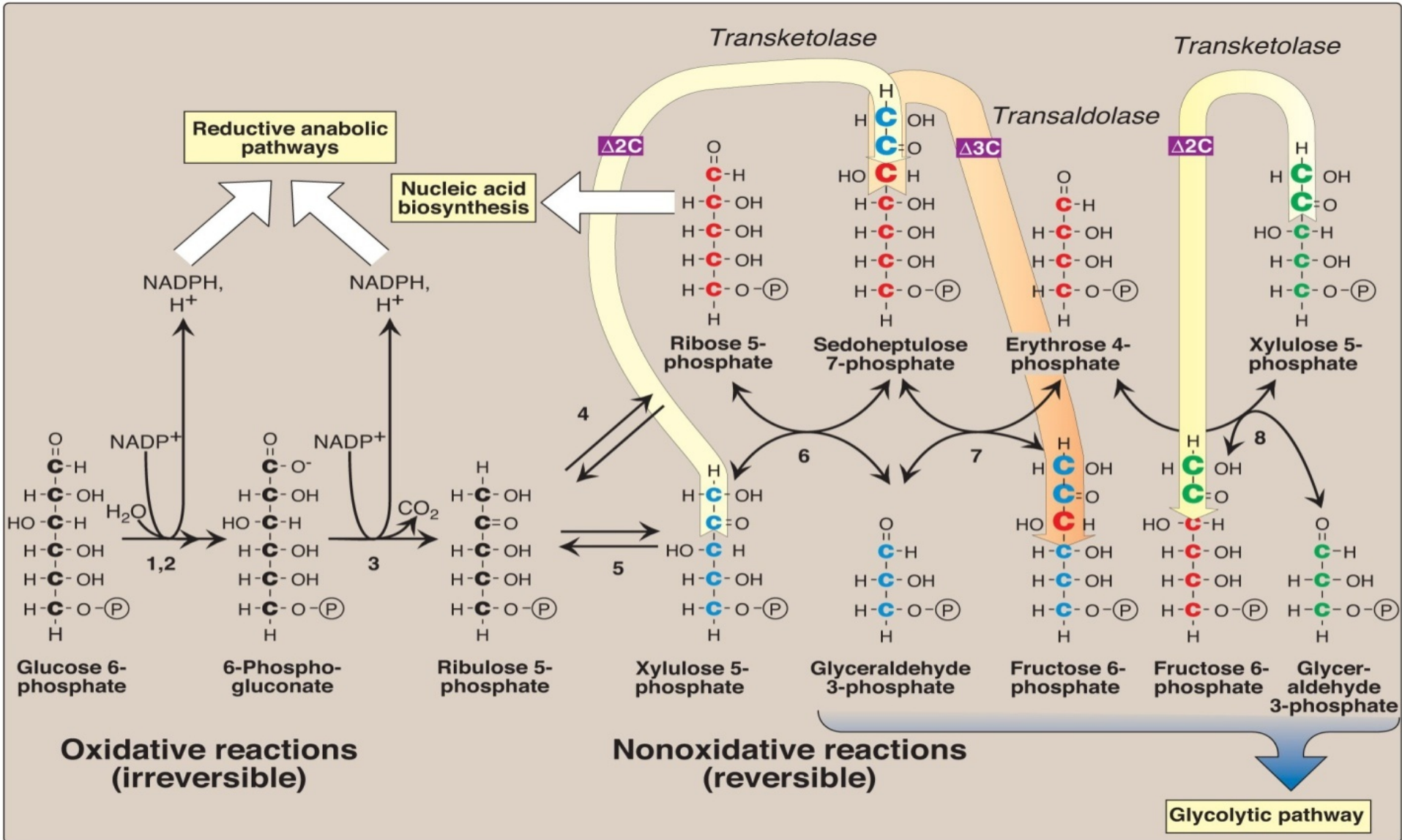
Tissue Distribution

Location- Cytosol

- Liver
- Lactating mammary gland
- Adrenal cortex
- Gonads
- Adipose tissue
- Erythrocytes to reduce glutathione
- Lens and cornea

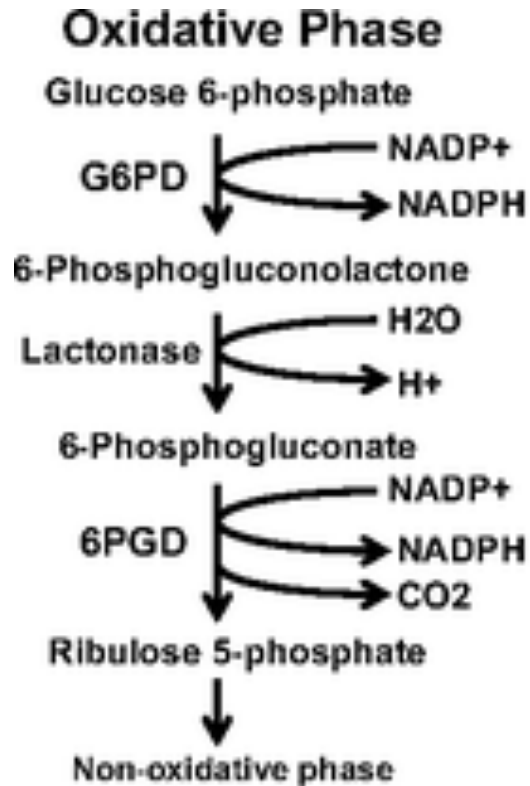
Phases of HMP Shunt

- It has two phases-
 - Oxidative phase
 - Non-oxidative phase



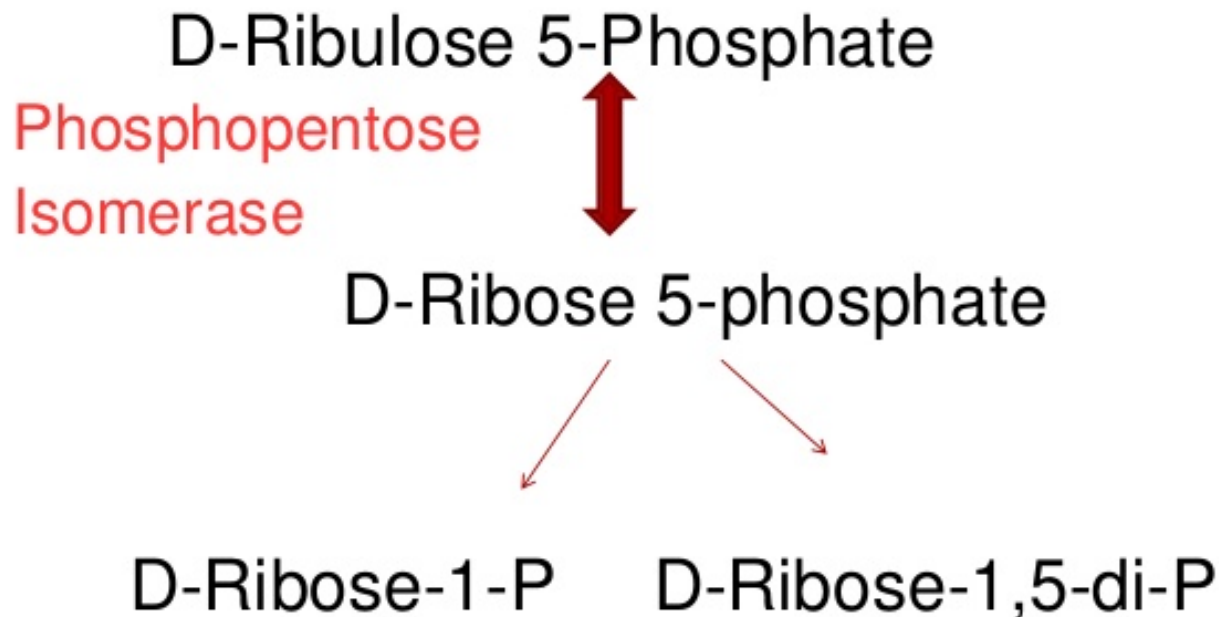
Enzymes numbered above are: 1, 2) *glucose 6-phosphate dehydrogenase* and *6-phosphogluconolactone hydrolase*, 3) *6-phosphogluconate dehydrogenase*, 4) *ribose 5-phosphate isomerase*, 5) *phosphopentose epimerase*, 6 and 8) *transketolase* (coenzyme: thiamine pyrophosphate), and 7) *transaldolase*.

Phase 1- Oxidative pathway

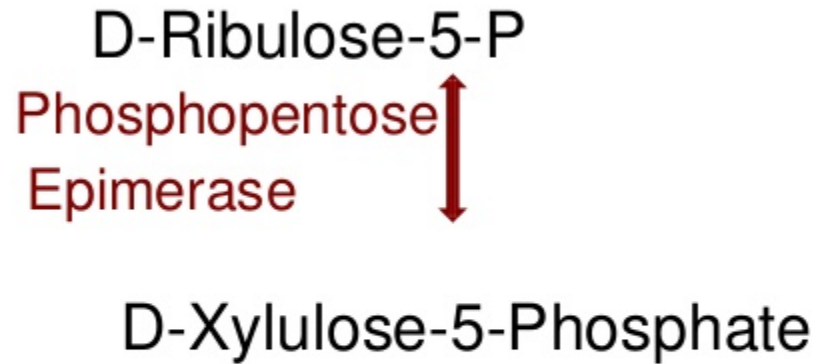


Phase 2- Non-oxidative

a) Interconversion of pentoses




Phase 2- Non-oxidative



Phase 2- Non-oxidative

a) Conversion of pentose phosphate to hexose phosphates


 2 Particular Enzymes are required:

1) TRANSKETOLASE

2) TRANSALDOLASE

Transketolation

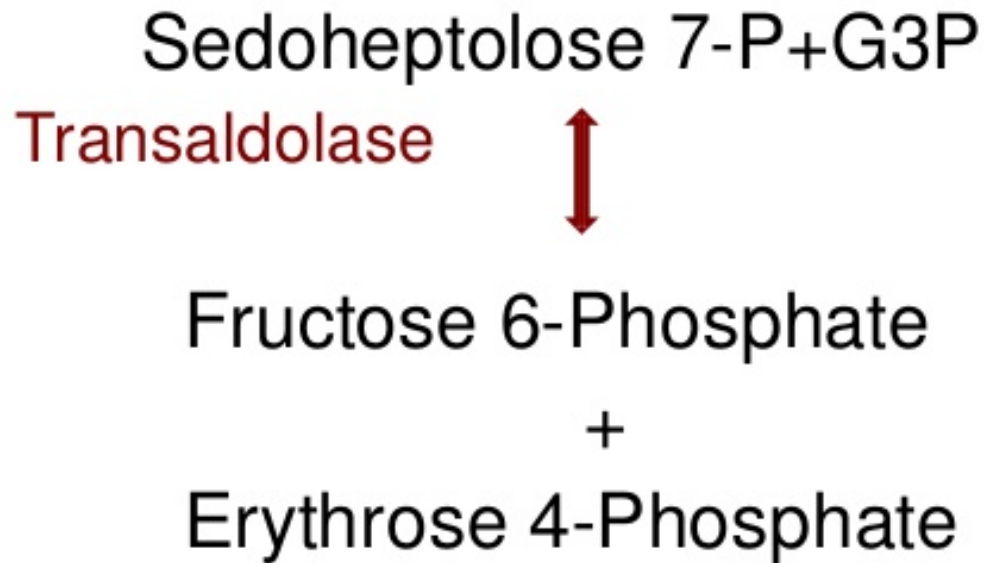
1) Xylulose-5-P + Ribose-5-P

Transketolase  TPP

Sedoheptolose 7-Phosphate +

Glyceraldehyde-3-Phosphate

Transaldolation



Transketolation

2) Xylulose 5-P + Erythrose 4-P

Transketolase \updownarrow TPP

Fructose 6-Phosphate + G3P

\updownarrow
Dihydroxy-acetone-P + G3P

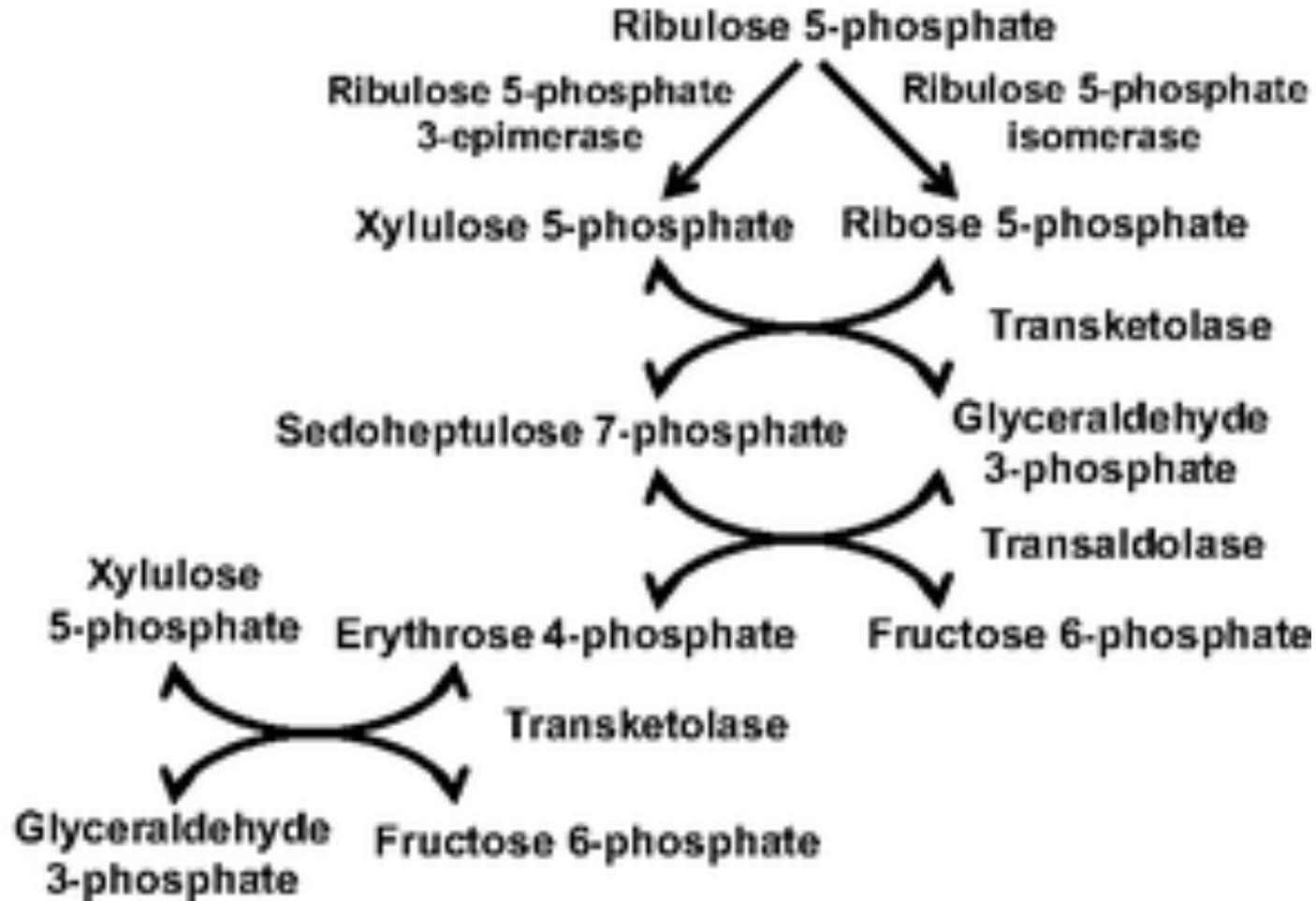
\updownarrow
Fructose-1,6-bi-P

Recycles
the Pathway

\updownarrow
Glucose-6-P \rightleftharpoons Fructose-6-P



Non-Oxidative Phase



Clinical Correlations

G-6-PD deficiency results in:

➤ Hemolytic Anemia

➤ Neonatal Jaundice

➤ Kidney failure

Glucose Transport

Na⁺-Monosaccharide Cotransporter:

Against concentration gradient

Energy dependent

Carrier-mediated

Coupled to Na⁺ transport

Small intestine, renal tubules & choroid plexus

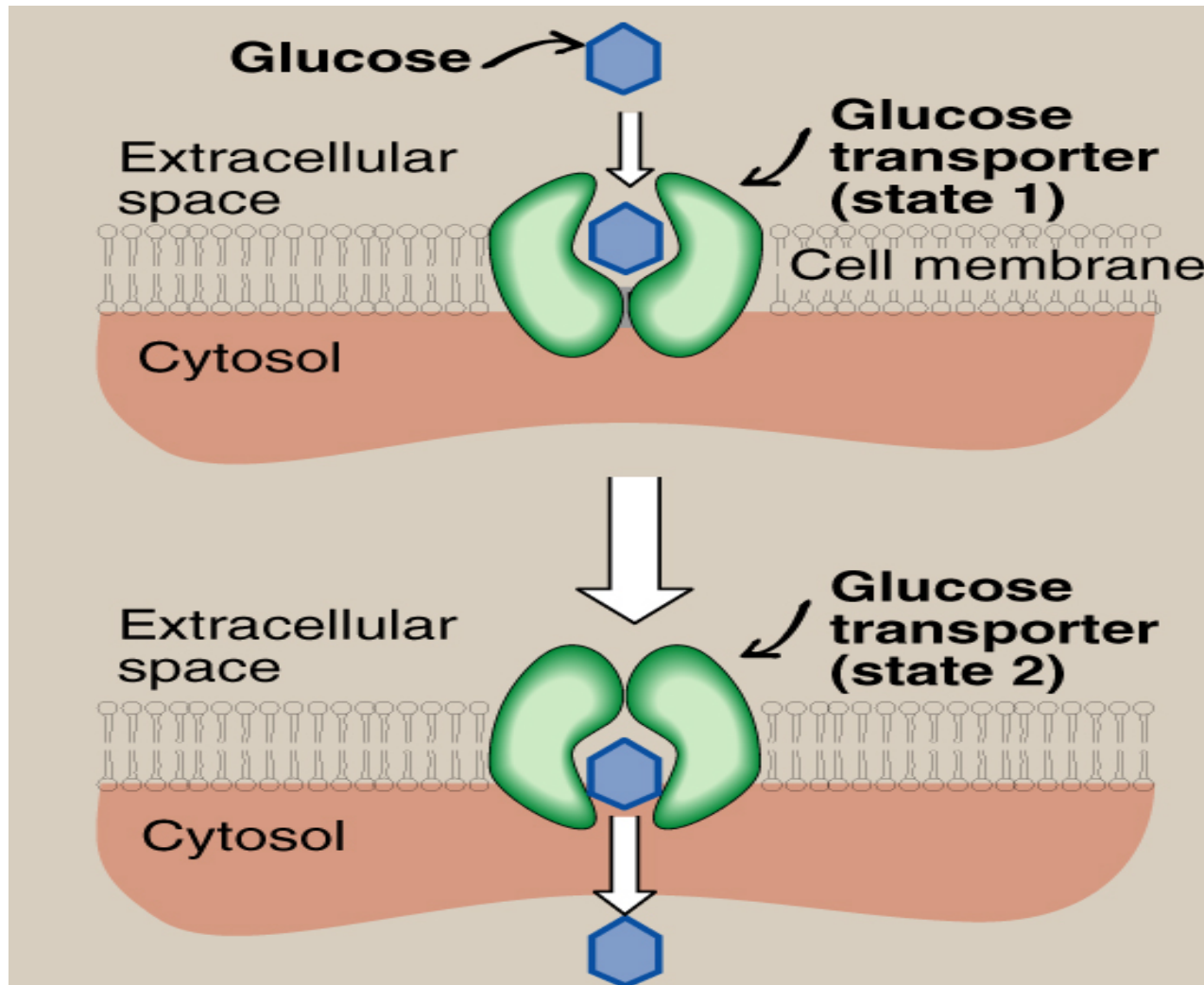
Na⁺-Independent Facilitated Diffusion:

Down the concentration gradient

Energy Independent

Glucose Transporters (GLUT 1-14)

Glucose Transport: Facilitated Diffusion



Glucose Transporters

- **Tissue-specific expression pattern**

GLUT-1

RBCs and brain

GLUT-2

Liver, kidney & pancreas

GLUT-3

Neurons

GLUT-4

Adipose tissue & skeletal muscle

GLUT-5

Small intestine & testes

GLUT-7

Liver (ER-membrane)

- **Functions:**

GLUT-1, 3 & 4 Glucose uptake from blood

GLUT-2

Blood & cells (either direction)

GLUT-5

Fructose transport

Take Home Message

- There are multiple pathways for glucose that can be grouped in to catabolic (utilizing glucose) or anabolic (producing glucose)
- Glycolysis is the major metabolic pathway of glucose breakdown to provide energy

Take Home Message - HMP

- Alternative pathway for glucose oxidation but not meant for producing energy
- Has two phases- oxidative and non-oxidative
- During oxidative phase, glucose-6-P is oxidized with generation of 2 moles of NADPH, and one mole of pentose phosphate, with liberation of CO₂
- During non-oxidative phase, pentose phosphate is converted to intermediates of glycolysis

References

- Lippincott's Illustrated Reviews- Biochemistry 6th Edition- pages: 96-97, 117, 126, 128, 145-147
- <http://www.biochemden.com/the-hexose-monophosphate-shunt/>