

**GLYCOGEN
METABOLISM, BLOOD
GLUCOSE HOMEOSTASIS,
DIABETES AND INSULIN**

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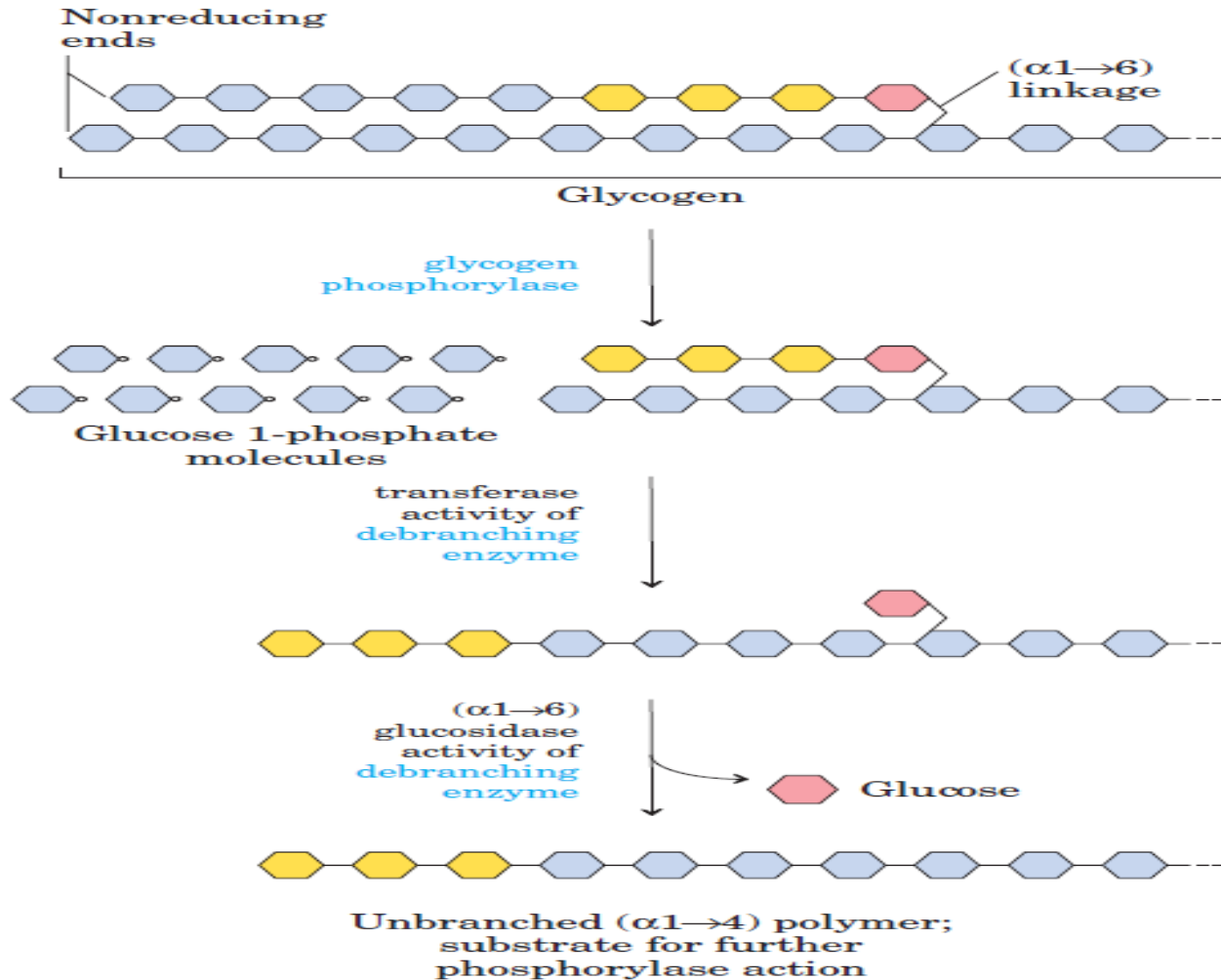
LECTURE CONTENT

- INTRODUCTION
- GLYCOGEN BREAKDOWN (Glycogenolysis)
- GLYCOGEN SYNTHESIS (Glycogenesis)
- REGULATION OF GLYCOGEN METABOLISM
- MAINTENANCE OF BLOOD GLUCOSE
- DIABETES MELLITUS AND INSULIN

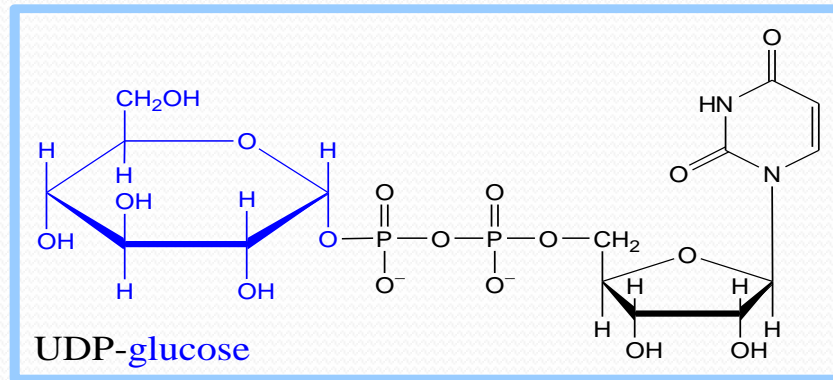
INTRODUCTION

- In many organisms, excess glucose is converted to polymeric forms for storage – glycogen
- In vertebrates, glycogen is found primarily in the liver and skeletal muscle
- Liver glycogen serves as a reservoir of glucose for other tissues when dietary glucose is not available (between meals or during a fast)
- In humans, the total amount of energy stored as glycogen is far less than the amount stored as fat (triacylglycerol)
- Glycogen granules are complex aggregates of glycogen and the enzymes that synthesize and degrade it as well as the machinery for regulating these enzymes

GLYCOGEN BREAKDOWN



GLYCOGEN SYNTHESIS

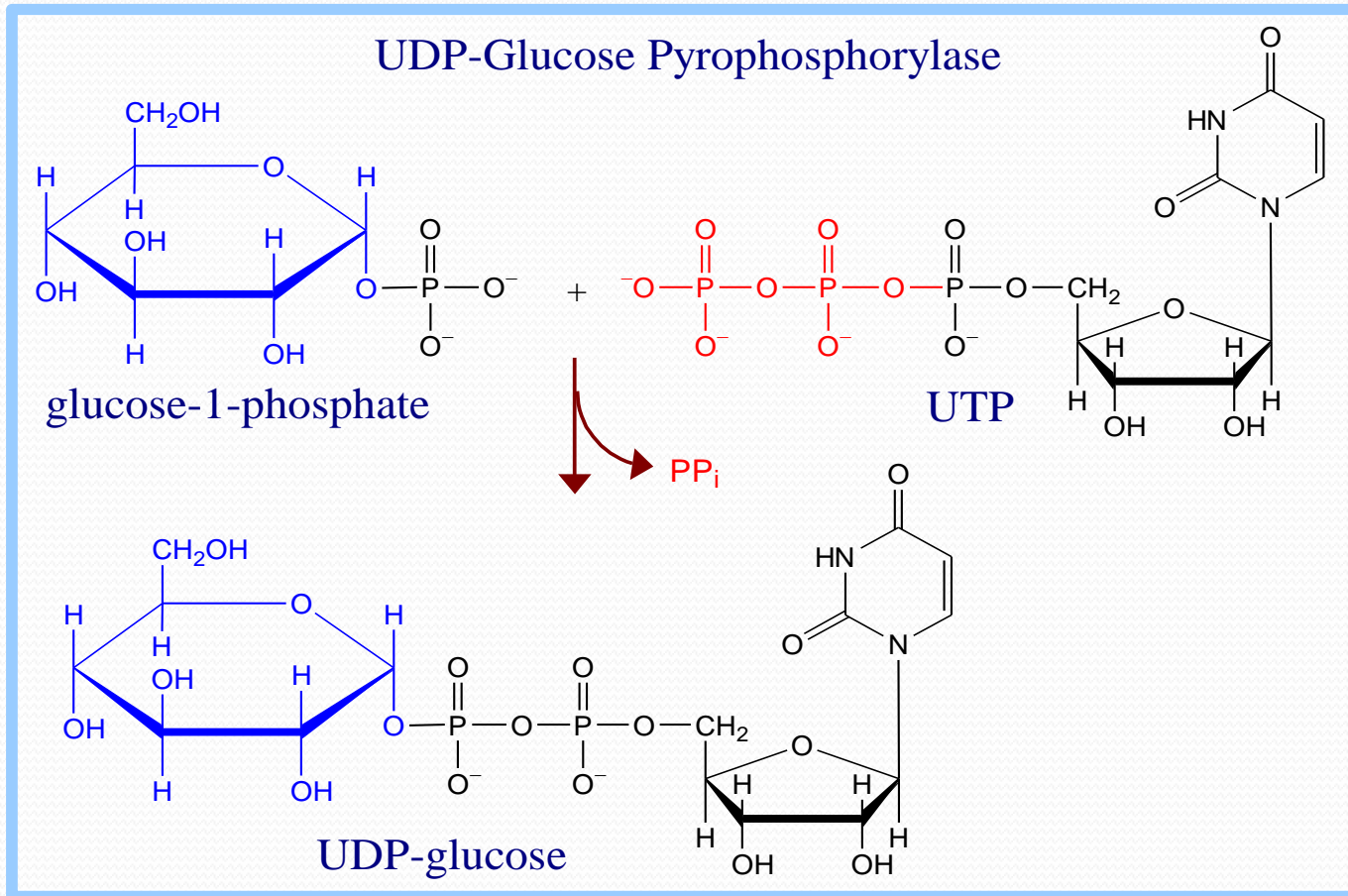


Uridine diphosphate glucose (UDP-glucose) is the immediate precursor for glycogen synthesis.

As glucose residues are added to glycogen, UDP-glucose is the substrate and UDP is released as a reaction product.

Nucleotide diphosphate sugars are precursors also for synthesis of other complex carbohydrates, including oligosaccharide chains of glycoproteins, etc.

GLYCOGEN SYNTHESIS



UDP-glucose is formed from glucose-1-phosphate:

- ◆ **glucose-1-phosphate + UTP \rightarrow UDP-glucose + PP_i**
- ◆ **PP_i + H₂O \rightarrow 2 P_i**

Overall:

- ◆ **glucose-1-phosphate + UTP \rightarrow UDP-glucose + 2 P_i**

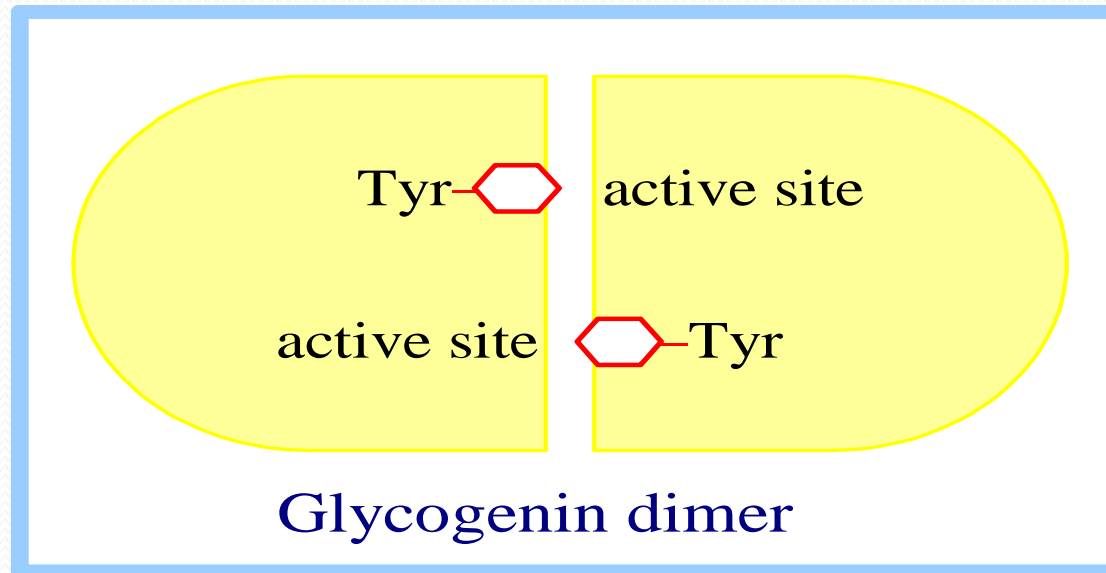
Spontaneous hydrolysis of the ~P bond in PP_i (P~P) drives the overall reaction.

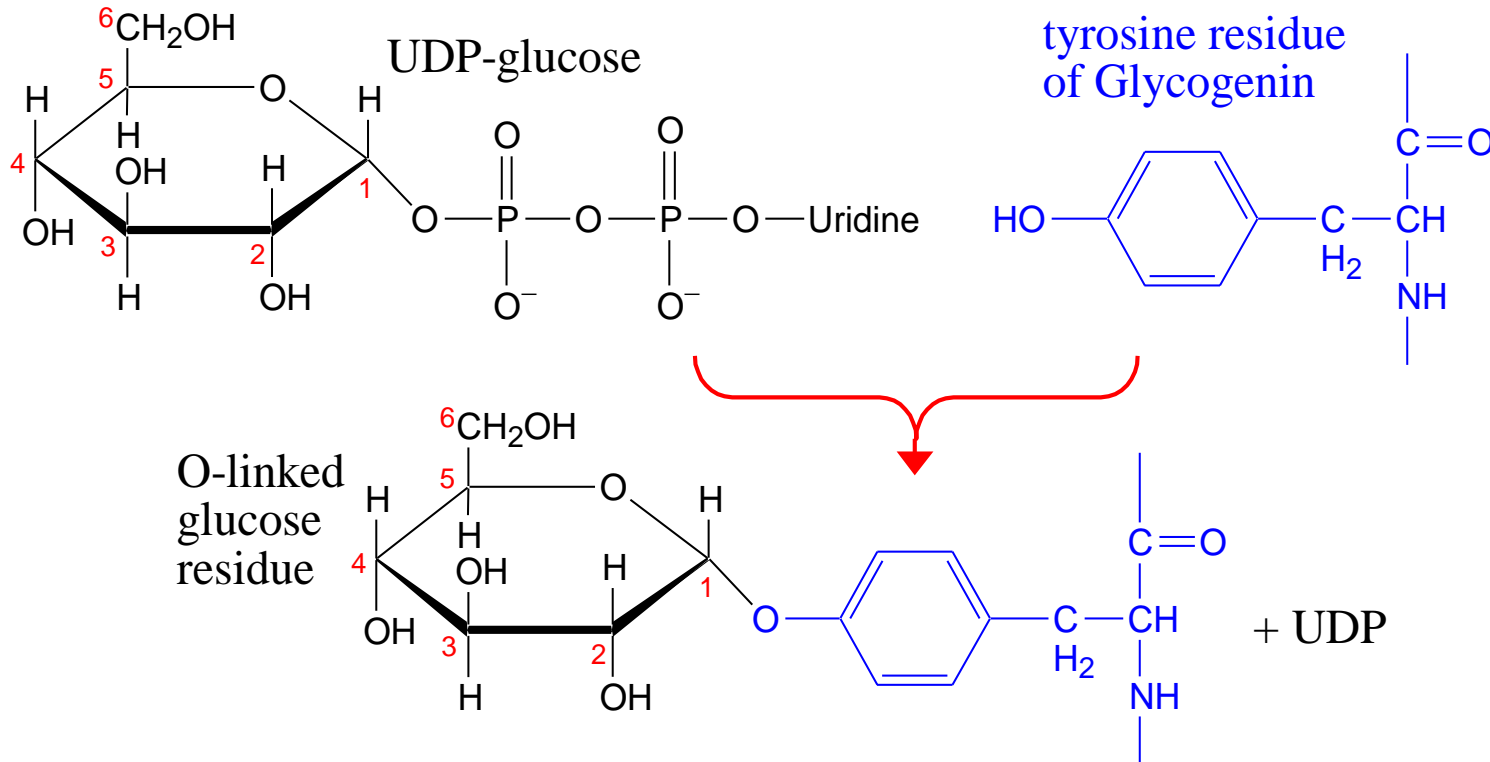
Cleavage of PP_i is the only energy cost for glycogen synthesis (one ~P bond per glucose residue).

Glycogenin initiates glycogen synthesis.

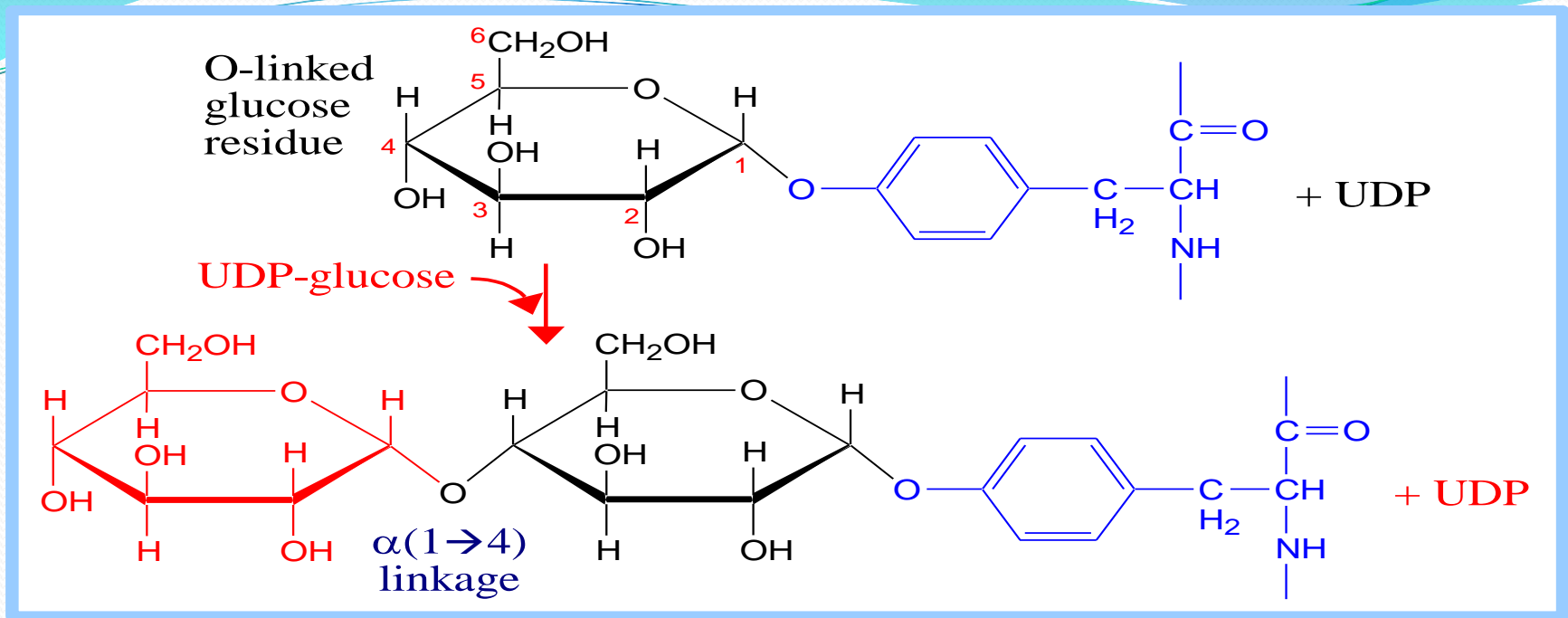
Glycogenin is an enzyme that catalyzes attachment of a **glucose** molecule to one of its own **tyrosine** residues.

Glycogenin is a **dimer**, and evidence indicates that the 2 copies of the enzyme glucosylate one another.





A **glycosidic bond** is formed between the anomeric C₁ of the glucose moiety derived from UDP-glucose and the hydroxyl oxygen of a **tyrosine** side-chain of **Glycogenin**.
 UDP is released as a product.



Glycogenin then catalyzes glucosylation at C₄ of the attached glucose (UDP-glucose again the donor), to yield an O-linked disaccharide with a(1-4) glycosidic linkage.

This is repeated until a short linear glucose polymer with a(1-4) glycosidic linkages is built up on Glycogenin

Glycogen Synthase catalyzes transfer of the glucose moiety of UDP-glucose to the hydroxyl at C₄ of the terminal residue of a glycogen chain to form an $\alpha(1 \rightarrow 4)$ glycosidic linkage:



A branching enzyme transfers a segment from the end of a glycogen chain to the C₆ hydroxyl of a glucose residue of glycogen to yield a branch with an $\alpha(1-6)$ linkage.

REGULATION OF GLYCOGEN METABOLISM

- Both synthesis & breakdown of glycogen are spontaneous.

If both pathways were active simultaneously in a cell, there would be a "**futile cycle**" with cleavage of **one ~P bond per cycle** (in forming UDP-glucose).

To prevent such a futile cycle, Glycogen Synthase and Glycogen Phosphorylase are **reciprocally regulated**

- Regulation of glycogen metabolism is achieved through (1) Covalent modification of key enzymes (2) Allosteric mechanisms and (3) Regulations by hormones (glucagon, epinephrine and insulin)

