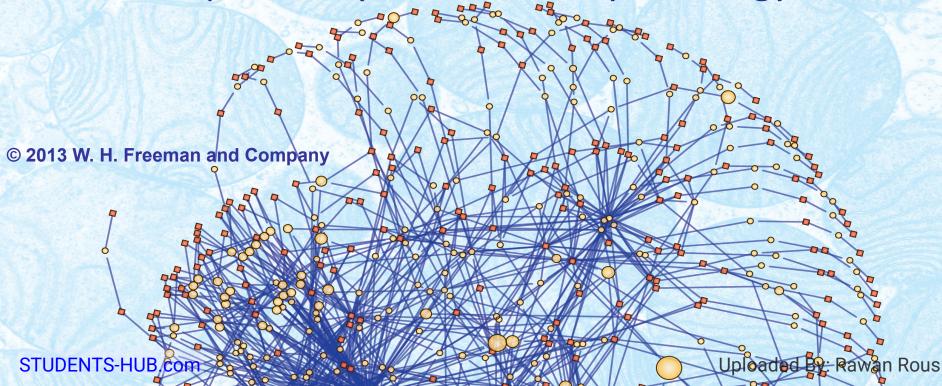
Lehninger

SIXTH EDITION

Principles of Biochemistry

David L. Nelson | Michael M. Cox

7 | Carbohydrates and Glycobiology



Carbohydrates

- Named so because many have formula C_n(H₂O)_n
- Produced from CO₂ and H₂O via photosynthesis in plants
- Range from as small as glyceraldehyde ($M_w = 90 \text{ g/mol}$) to as large as amylopectin ($M_w = 200,000,000 \text{ g/mol}$)
- Fulfill a variety of functions including:
 - energy source and energy storage
 - structural component of cell walls and exoskeletons
 - informational molecules in cell-cell signaling
- Can be covalently linked with proteins to form glycoproteins and proteoglycans

Carbohydrates

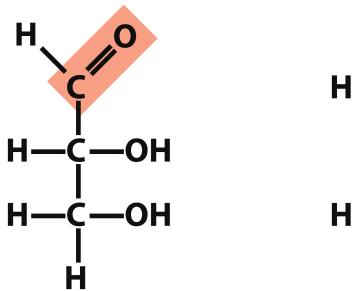
- Monosaccharides (simple sugars) one polyhydroxy aldehyde or ketone unit (glucose)
- Disaccharides two monosaccharide units joined together by a glycosidic linkage (sucrose)
- Oligosaccharides few monosaccharide units joined together (in cells, most oligosaccharides are joined to nonsugar molecule)
- Polysaccharides sugar polymers consisting of >20 monosaccharide units

Monosaccharide Carbon backbone

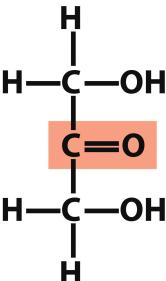
- 3 C → triose
- 4 C → tetrose
- 5 C → pentose
- 6 C → hexose
- 7 C → heptose

Aldoses and Ketoses

- An aldose contains an aldehyde functional group
- A ketose contains a ketone functional group



Glyceraldehyde, an aldotriose



Dihydroxyacetone, a ketotriose

Figure 7-1a
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Stereoisomers

 All monosaccharides (except dihydroxyacetone) are chiral compounds, with <u>at least</u> one chiral carbon

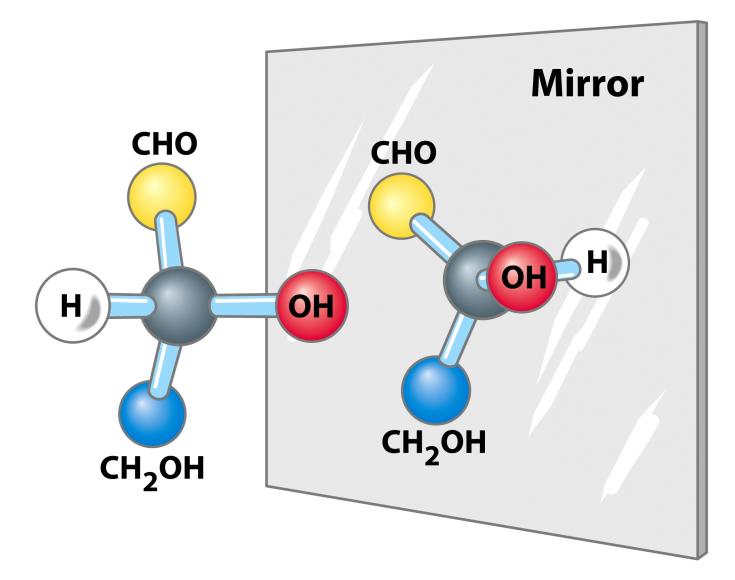
A molecule with n chiral centers can have 2ⁿ stereoisomers

E.g. glyceraldehyde has 2¹ = 2
 aldohexoses have 2⁴ = 16 stereoisomers

Enantiomers

Enantiomers

- Stereoisomers that are nonsuperimposable mirror images
- In sugars that contain many chiral centers, only the one that is most distant from the carbonyl carbon is designated as D (right) or L (left)
- D and L isomers of a sugar are enantiomers
 - For example, L and D glucose have the same water solubility
- Most hexoses in living organisms are D stereoisomers
- Some simple sugars occur in the L-form, such as Larabinose

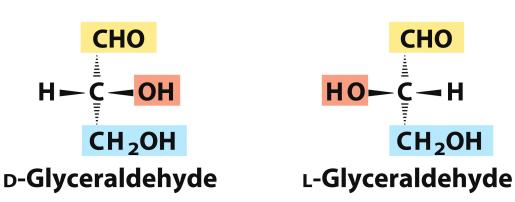


Ball-and-stick models

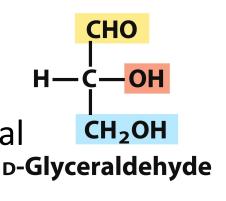
Figure 7-2 part 1 *Lehninger Principles of Biochemistry,* Sixth Edition

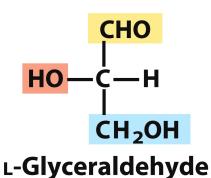
Drawing Monosaccharides

- Chiral compounds can be drawn using perspective formulas
- However, chiral carbohydrates are usually represented by Fischer projections
- Horizontal bonds are pointing toward you; vertical bonds are projecting away



Perspective formulas

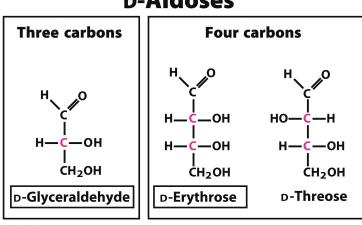




Fischer projection formulas

Diastereomers

- Diastereomers: stereoisomers that are not mirror images
- Diastereomers have different physical properties
 - For example, water solubilities of threose and erythrose are
 different D-Aldoses



Epimers

 Epimers are two sugars that differ only in the configuration around one carbon atom

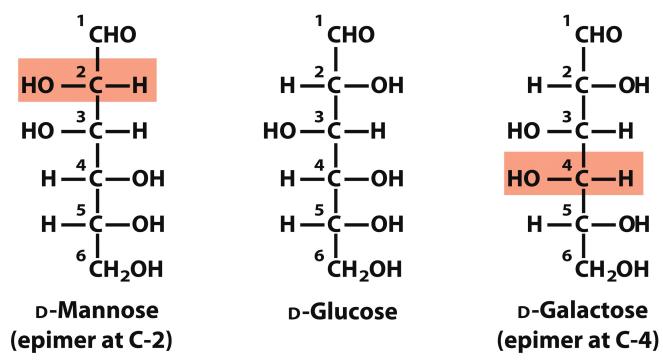


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Structures to Know

- Glyceraldehyde & dihydroxyacetone are the standard three-carbon sugars
- Ribose is the standard five-carbon sugar
- Glucose is the standard six-carbon sugar
- Galactose is an epimer of glucose
- Mannose is an epimer of glucose
- Fructose is the ketose form of glucose

D-Aldoses

Three carbons H-C-OH CH₂OH D-Glyceraldehyde

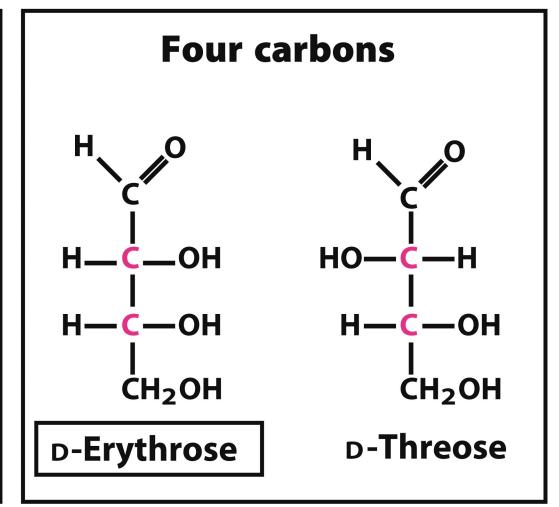


Figure 7-3a part 1

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D-Aldoses

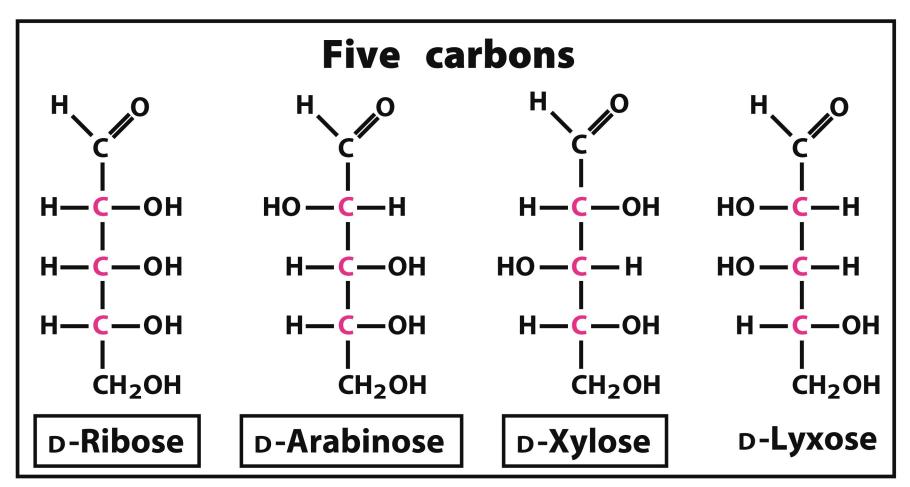


Figure 7-3a part 2
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D-Aldoses

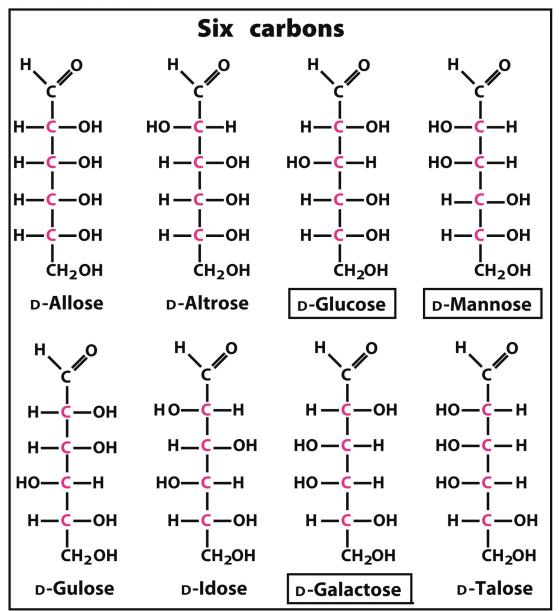


Figure 7-3a part 3

D-Ketoses

Three carbons

Dihydroxyacetone

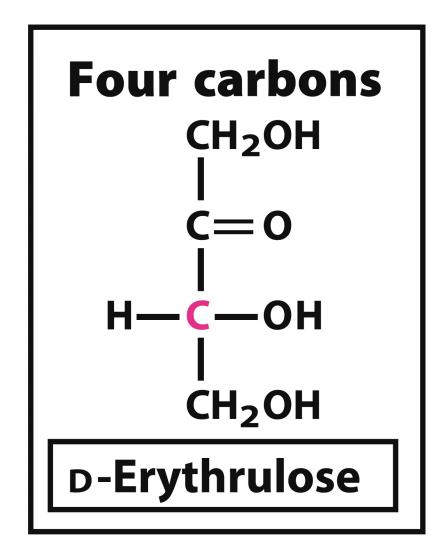
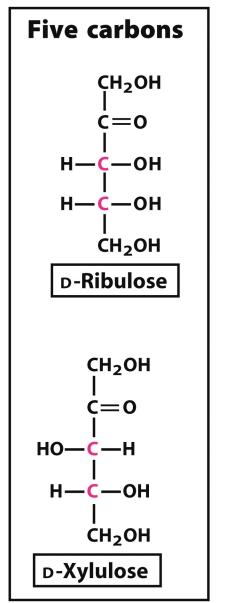


Figure 7-3b part 1

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D-Ketoses



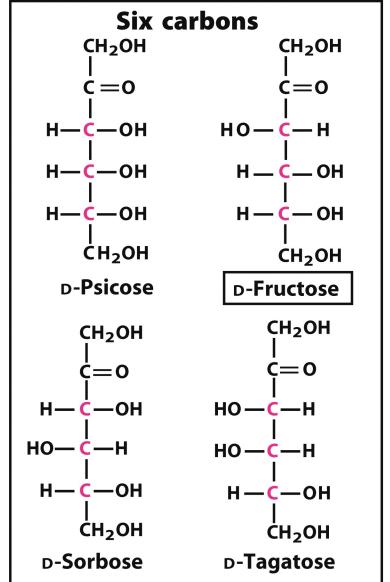


Figure 7-3b part 2

Common Monosaccharides Have Cyclic Structures

 In solution, monosaccharides with 4 or more carbons form cyclic compounds (ring structures)

 Carbonyl C is attacked by the hydroxyl O forming a covalent bond

Hemiacetals and Hemiketals

- Aldehyde and ketone carbons are electrophilic
- Alcohol oxygen atom is nucleophilic
- When aldehydes are attacked by alcohols, hemiacetals form
- When ketones are attacked by alcohols, hemiketals form

$$R \xrightarrow{1} C \xrightarrow{O} + HO - R^{2} \Longrightarrow R \xrightarrow{1} C - OR^{2} \xrightarrow{HO - R^{3}} R \xrightarrow{OR^{3}} R \xrightarrow{OR^{2} + HOH}$$
Aldehyde Alcohol Hemiacetal Acetal

$$R \xrightarrow{1} C = O + HO - R^{3} \Longrightarrow R \xrightarrow{1} C - OR^{3} \Longrightarrow HO - R^{4} \qquad R \xrightarrow{1} C - OR^{3} + HOH$$

$$R \xrightarrow{1} R \xrightarrow{$$

Figure 7-5

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When the second alcohol is part of another sugar molecule, the bond produced in a law bond us

Cyclization of Monosaccharides

- Pentoses and hexoses readily undergo intramolecular ring formation
- The former carbonyl carbon becomes a <u>new chiral center</u>, called the <u>anomeric carbon</u>
- The former carbonyl oxygen becomes a hydroxyl group; the position of this group determines if the anomer is α or β
- If the hydroxyl group is on the opposite side (trans) of the ring as the CH₂OH moiety the configuration is α
- In the hydroxyl group is on the same side (cis) of the ring as the CH₂OH moiety, the configuration is β

Mutarotation – the interconversion of α and β anomers.

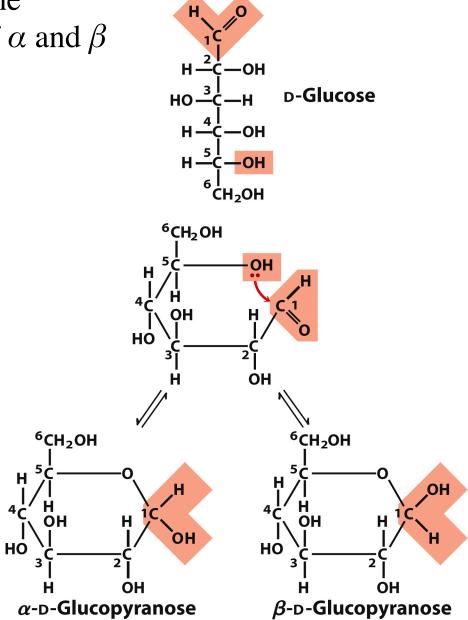
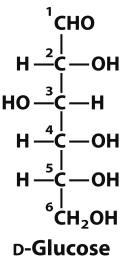


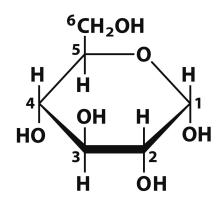
Figure 7-6
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Pyranoses and Furanoses

- Six-membered oxygen-containing rings are called pyranoses
- Five-membered oxygen-containing rings are called furanoses
- The anomeric carbon is usually drawn on the right side
- Cyclic sugar structures are more accurately represented in Haworth perspective formulas



D-Glucose Fischer projection



α-D-Glucopyranose
Haworth perspective
Uploaded By: Rawan Rous

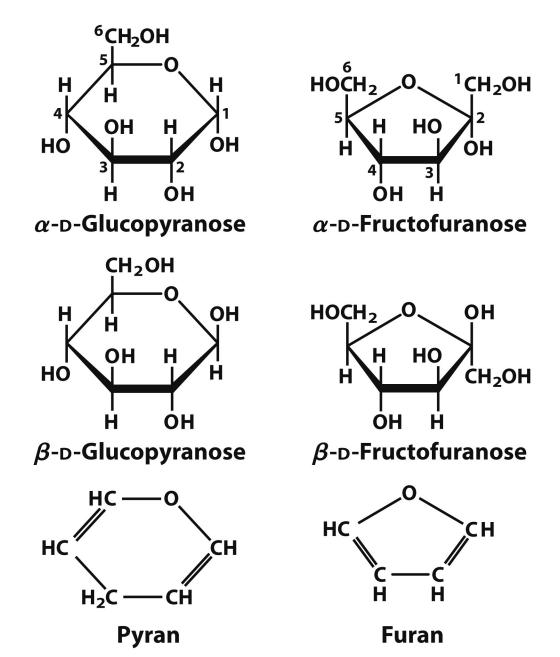


Figure 7-7
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Converting linear to cyclic structures

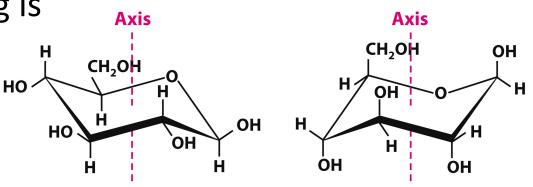
- For any linear hexose:
- 1. Draw 6-membered ring (O at the upper right)
- 2. Number C's clockwise (start with anomeric)
- Insert the –OH
 a. if –OH is on the right, draw it pointing down
 b. if –OH is on the left, draw it pointing up
- Insert the terminal –CH₂OH
 a. if the sugar is the p-isomer, draw it upwards
 b. if it is the μ-isomer, draw it downwards
- Draw the –OH on the anomeric C
 a. if it faces the same side as the –CH₂OH, it's β

The ring is not planar

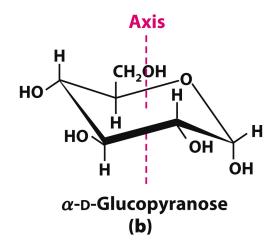
- In solution pyranose ring is not planar but it can assume two chair
- Boat conformation is possible but very rare

conformations

 The two chair conformations are not readily interconvertible (one conformer is more stable than the other)



Two possible chair forms of β -D-glucopyranose (a)



Chain-Ring Equilibrium and Reducing Sugars

- The ring forms exist in equilibrium with the open-chain forms
- Aldehyde can reduce Cu²⁺ to Cu⁺ (Fehling's test)
- Aldehyde can reduce Ag⁺ to Ag⁰ (Tollens' test)
- Allows detection of reducing sugars, such as glucose (by measuring the amount of oxidizing agent reduced by a sugar solution, the sugar concentration can be estimated)

The cuprous ion (Cu⁺) produced forms a red cuprous oxide precipitate.

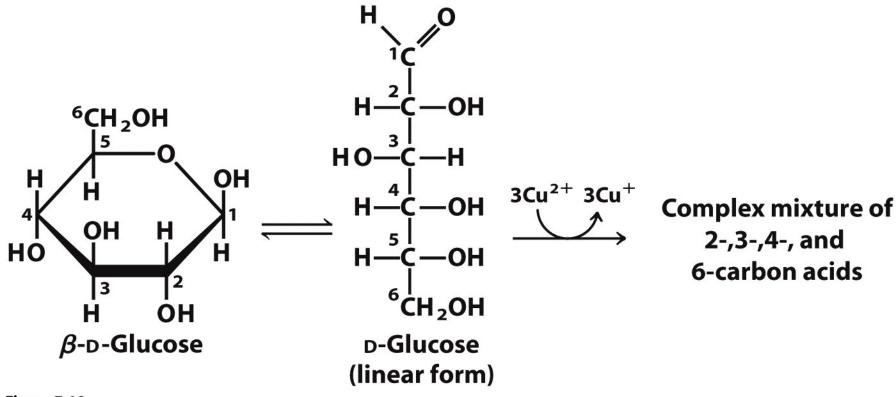
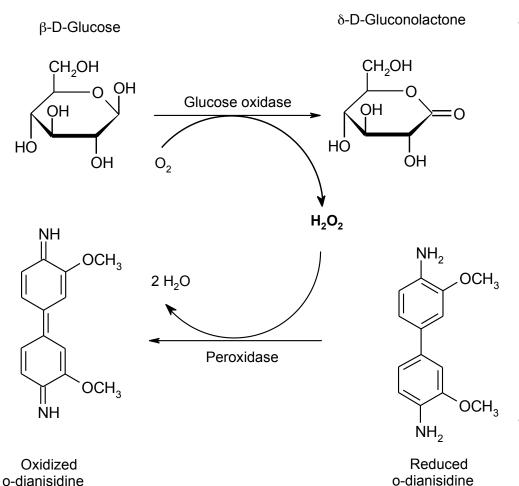


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Colorimetric Glucose Analysis

(faint orange)



- Nowadays, enzymatic methods are used to quantify reducing sugars such as glucose
 - The enzyme glucose oxidase catalyzes the conversion of glucose to glucono- δ -lactone and hydrogen peroxide
 - Hydrogen peroxide oxidizes organic molecules into highly colored compounds
 - Concentrations of such compounds is measured colorimetrically
- Electrochemical detection is used in portable glucose sensors

(bright orange)

Blood glucose measurements in the diagnosis and treatment of diabetes

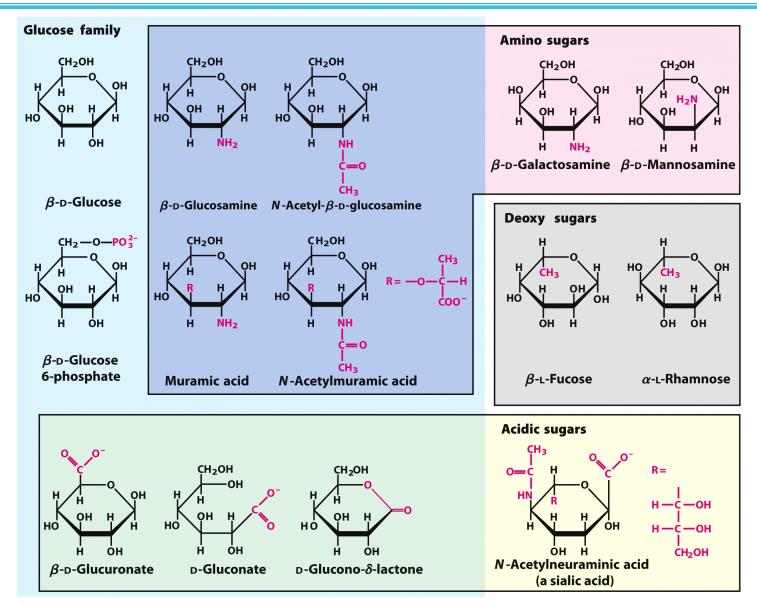
- Untreated diabetes has several long-term consequences: kidney failure, cardiovascular disease, blindness, impaired wound healing, etc.
- Average [glc]_{blood} over days can be measured because of a nonenzymatic reaction between glc and primary amino groups in Hb
- The amount of glycated Hb (GHB) reflects the average [glc]_{blood} over the circulating lifetime of RBC (~120 days)
- Normal levels ~5% of Hb is GHB
- Diabetic people may have it as high as 13%
- Advanced glycation endproducts (AGEs) contribute to the

Damage to kidneys, retinas, cardiovascular system

↓?

↓?

Organisms contain many hexose derivatives



The Glycosidic Bond

- Two sugar molecules can be joined via a glycosidic bond between an anomeric carbon and a hydroxyl carbon
- The resulting compound is a glycoside
- The glycosidic bond (an acetal) between monomers is less reactive than the hemiacetal at the second monomer
 - Second monomer, with the hemiacetal, is reducing
 - Anomeric carbon involved in the glycosidic linkage is nonreducing
- The disaccharide formed upon condensation of two glucose molecules via $1 \rightarrow 4$ bond is called maltose

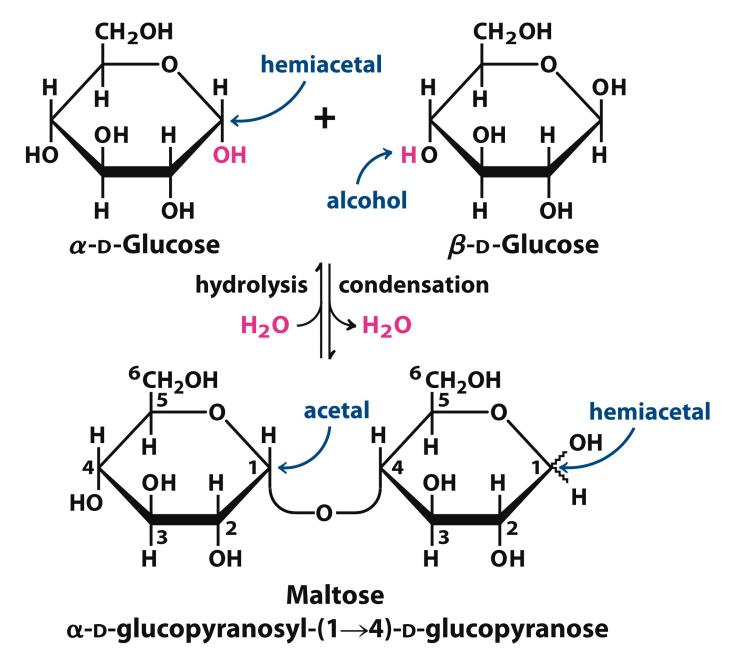


Figure 7-10

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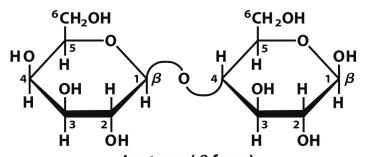
Nonreducing Disaccharides

- Two sugar molecules can be also joined via a glycosidic bond between two anomeric carbons
- The product has two acetal groups and no hemiacetals
- There are no reducing ends, this is a nonreducing sugar
- Trehalose is a constituent of hemolymph of insects
 - Provides protection from drying

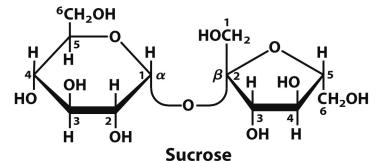
Naming disaccharides

- 1. Find the nonreducing end
- 2. Give the configuration (α or β) at the anomeric C joining the first monosaccharide to the other
- 3. Name the nonreducing residue (use "furano" for 5-membered rings or "pyrano" for 6-membered rings)
- Add the glycosidic bond in parenthesis (from which C to which C)
- 5. Name the second residue
- 6. If there are more residues, repeat step 2

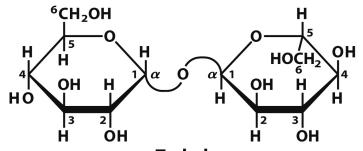
^{*} Abbreviations can be used (Glc, Fru, Gal, Man, GlcN, etc.)



Lactose (β form) β -D-galactopyranosyl-(1 \rightarrow 4)- β -D-glucopyranose Gal(β 1 \rightarrow 4)Glc



 β -D-fructofuranosyl α -D-glucopyranoside Fru($2\beta \leftrightarrow \alpha 1$)Glc \equiv Glc($\alpha 1 \leftrightarrow 2\beta$)Fru



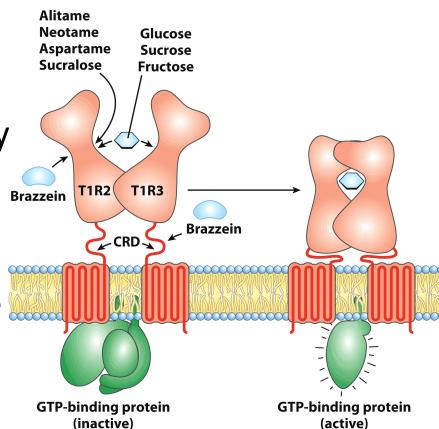
Trehalose α -D-glucopyranosyl α -D-glucopyranoside $Glc(\alpha 1 \leftrightarrow 1\alpha)Glc$

Figure 7-11
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Sweet tooth? Anyone?

Res 73 Special Management of M

- Most people like sweets
- Sweetness is one of 5 tastes humans can taste
- Due to receptors on gustatory cells on the surface of tongues... T1R2 and T1R3
- Binding of molecules to these molecules to the second to the molecules to t



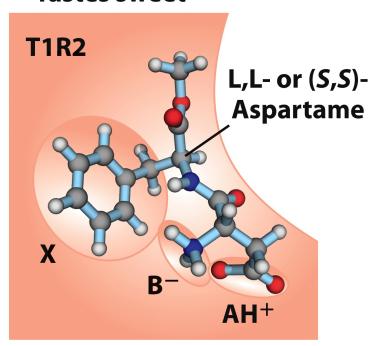


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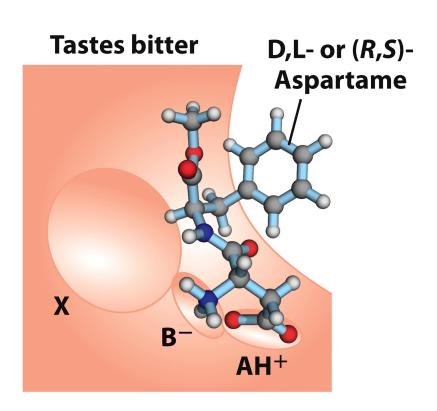


L,L-Aspartame is sweet; D,L-Aspartame is bitter

Tastes sweet

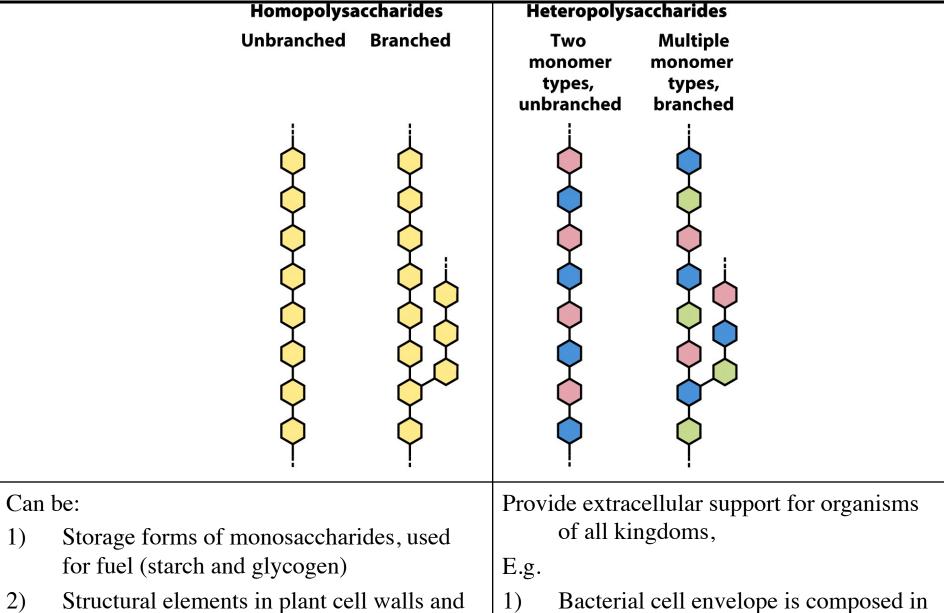


Box 7-2 figure 3 *Lehninger Principles of Biochemistry,* Sixth Edition © 2013 W. H. Freeman and Company



Polysaccharides

- Natural carbohydrates are usually found as polymers
- These polysaccharides can be
 - homopolysaccharides
 - heteropolysaccharides
 - linear
 - branched
- Polysaccharides do not have a defined molecular weight.
 - This is in contrast to proteins because unlike proteins, no template is used to make polysaccharides



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animal exoskeletons (cellulose and chitin)

1)

part of a heteropolysaccharide

Extracellular matrix in animal cells 2) <u> Unloaded Ry: Rawan Rous</u>

Glycogen

- Glycogen is a branched homopolysaccharide of glucose
 - Glucose monomers form ($\alpha 1 \rightarrow 4$) linked chains
 - Branch-points with ($\alpha 1 \rightarrow 6$) linkers every 8–12 residues
 - Molecular weight reaches several millions
 - Functions as the main storage polysaccharide in animals

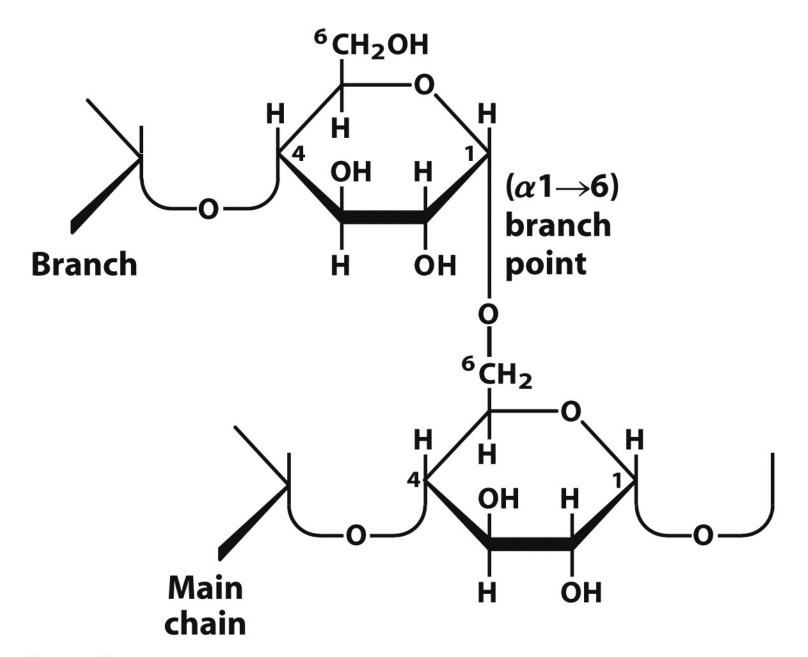


Figure 7-14b

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Starch

- Starch is a mixture of two homopolysaccharides of glucose
 - Amylose is an unbranched polymer of ($\alpha 1 \rightarrow 4$) linked residues
 - Amylopectin is branched like glycogen but the branchpoints with ($\alpha 1 \rightarrow 6$) linkers occur every 24–30 residues
 - Molecular weight of amylopectin is up to 200 million
- Starch is the main storage polysaccharide in plants

Glycosidic Linkages in Glycogen and Starch

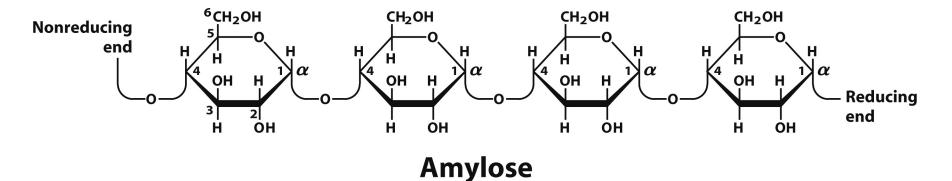
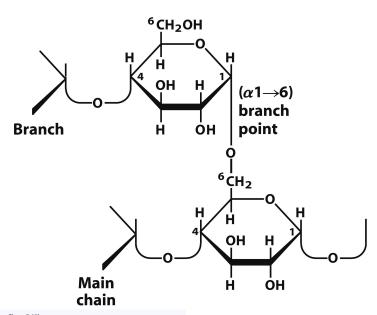


Figure 7-13a
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Mixture of Amylose and Amylopectin in Starch

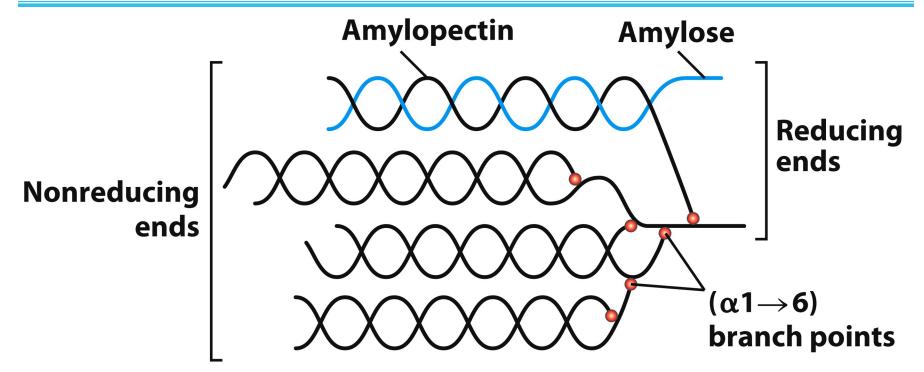


Figure 7-13c
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In starch granules, strands of amylopectin (blue) form double-helical structures with each other or with amylose strands (blue). Glucose residues at the nonreducing ends of the outer branches are removed enzymatically during the mobilization of starch for energy production.

Glycogen has a similar structure but is more highly branched and more Uploaded By: Rawan Rous

Metabolism of Glycogen and Starch

- Glycogen and starch often form granules in cells
- Granules contain enzymes that synthesize and degrade these polymers
- Glycogen and amylopectin have one reducing end but many nonreducing ends (a polymer with n branches has n + 1 nonreducing ends and only 1 reducing end)
- Enzymatic processing occurs simultaneously in many nonreducing ends

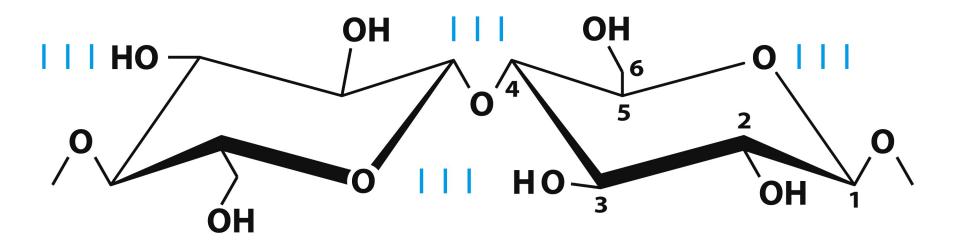
Dextrans

- Bacterial and yeast polysaccharides
- $(\alpha 1 \rightarrow 6)$ -linked poly-D-glucose
- All units have $(\alpha 1 \rightarrow 3)$ branches
- Some have also (α1 → 2) or (α1 → 4) branches
- Dental plaque (formed by bacteria on the surface of teeth) is rich in dextrans

Cellulose

- Cellulose is a branched homopolysaccharide of glucose
 - Glucose monomers form ($\beta 1 \rightarrow 4$) linked chains
 - Hydrogen bonds form between adjacent monomers
 - Additional H-bonds between chains
 - Structure is now tough and water-insoluble
 - Most abundant polysaccharide in nature
 - Cotton is nearly pure fibrous cellulose

Hydrogen Bonding in Cellulose



 $(\beta 1 \rightarrow 4)$ -linked D-glucose units

Figure 7-14

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Cellulose Metabolism

- The fibrous structure and water-insolubility make cellulose a difficult substrate to act on
- Fungi, bacteria, and protozoa secrete cellulase, which allows them to use wood as source of glucose
- Most animals cannot use cellulose as a fuel source because they lack the enzyme to hydrolyze ($\beta 1 \rightarrow 4$) linkages
- Ruminants and termites live symbiotically with microorganisms that produces cellulase
- Cellulases hold promise in the fermentation of biomass into biofuels



Figure 7-16
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Chitin

- Chitin is a linear homopolysaccharide of N-acetylglucosamine
 - N-acetylglucosamine monomers form ($\beta 1 \rightarrow 4$)-linked chains
 - Forms extended fibers that are similar to those of cellulose
 - Hard, insoluble, cannot be digested by vertebrates
 - Structure is tough but flexible, and water-insoluble
 - Found in cell walls in mushrooms, and in exoskeletons of insects, spiders, crabs, and other arthropods
 - 1 billion tons of chitin is produced in the biosphere per year

Chitin

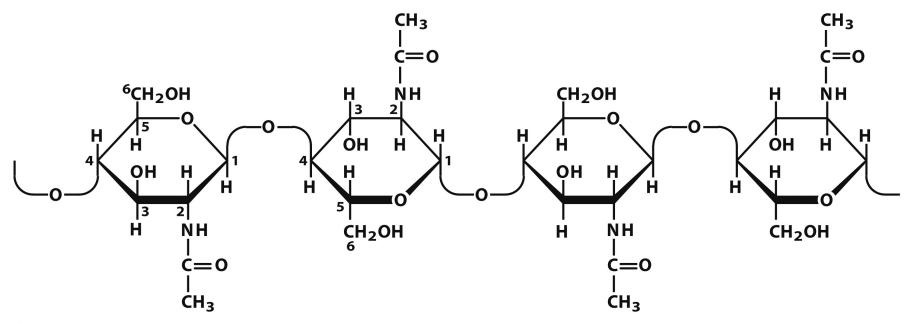


Figure 7-16a

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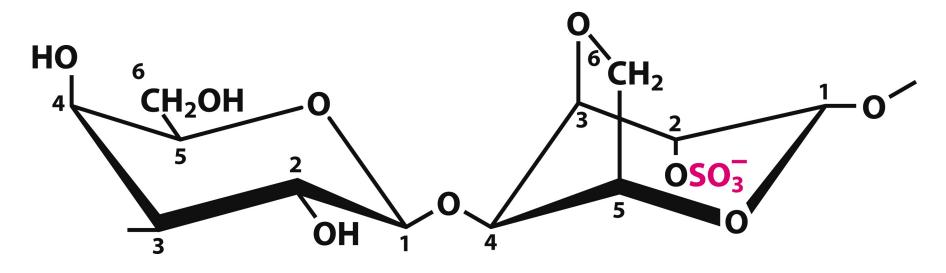


Figure 7-16b
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Agar and Agarose

- Agar is a complex mixture of hetereopolysaccharides containing modified galactose units
- Agar serves as a component of cell wall in some seaweeds
- Agarose is one component of agar
- Agar solutions form gels that are commonly used in the laboratory as a surface for growing bacteria
- Agar is also used for capsules in which some drugs and vitamins are packaged
- Agarose solutions form gels that are commonly used in the laboratory for separation DNA by electrophoresis

Agar and Agarose



Agarose 3)D-Gal(β 1 \rightarrow 4)3,6-anhydro-L-Gal25(α 1 repeating units)

Figure 7-21
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Glycosaminoglycans

- Linear polymers of repeating disaccharide units
- One monomer is either
 - N-acetyl-glucosamine or
 - N-acetyl-galactosamine
- Negatively charged
 - Uronic acids (C6 oxidation)
 - Sulfate esters
- Extended hydrated molecule
 - Minimizes charge repulsion
- Forms meshwork with fibrous proteins to form extracellular matrix
 - Connective tissue
 - Lubrication of joints

Glycosaminoglycan Repeating disaccharide Number of CH₂OH disaccharides per chain Н COO HO 0 **Hyaluronate** $(\beta 1\rightarrow 4)$ Н NH~50,000 OH C = 0*(β*1→3) CH₃ OH **GIcA GIcNAc** CH₂OH ⁻0₃SO COO Chondroitin $(\beta 1\rightarrow 4)$ Н 4-sulfate NH OH 20-60 C = 0*(β*1→3) OH CH₃ **GIcA** GalNAc4S

Figure 7-22 part 1a

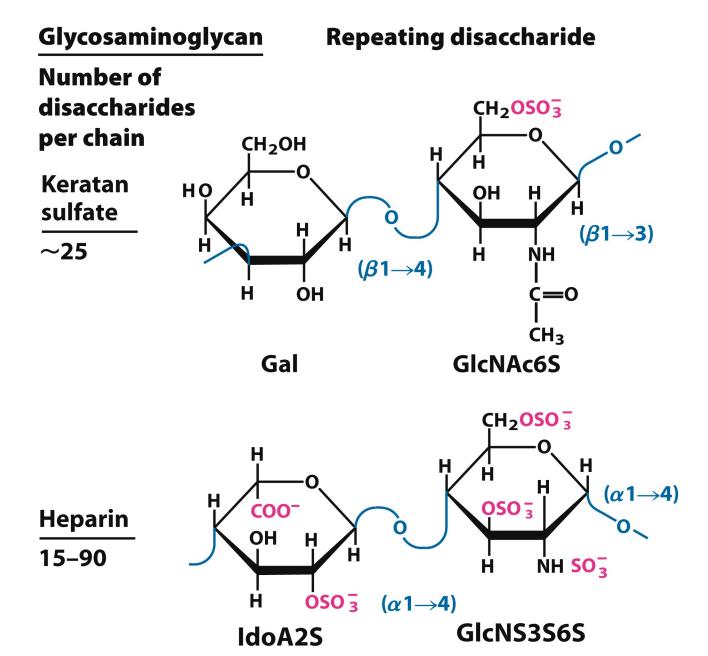
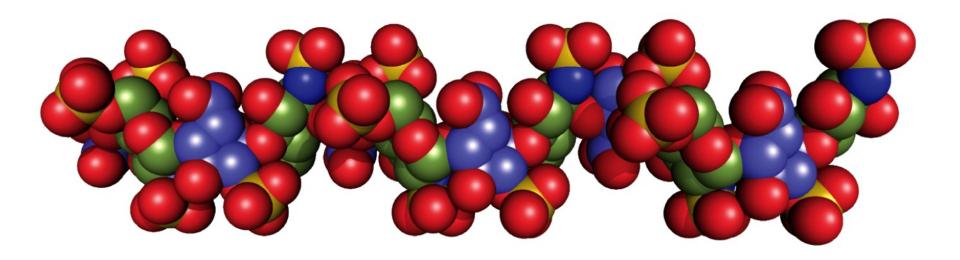


Figure 7-22 part 1b

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Heparin and Heparan Sulfate

- Heparin is linear polymer, 3–40 kDa
- Heparan sulfate is heparin-like polysaccharide but attached to proteins
- Highest negative charge density biomolecules
- Prevent blood clotting by activating protease inhibitor antithrombin
- Binding to various cells regulates development and formation of blood vessels
- Can also bind to viruses and bacteria and decrease their virulence

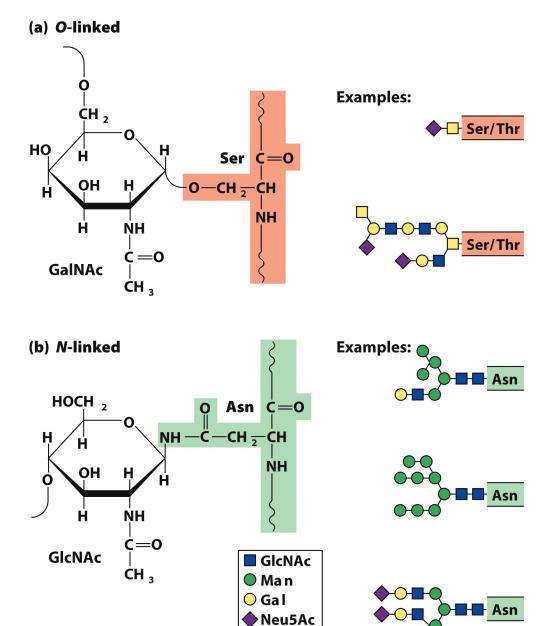


Heparin segment

Figure 7-22 part 2 Lehninger Principles of Biochemistry, Sixth Edition © 2013 W. H. Freeman and Company

Glycoconjugates: Glycoprotein

- A protein with small oligosaccharides attached
 - Carbohydrate attached via its anomeric carbon
 - About half of mammalian proteins are glycoproteins
 - Carbohydrates play role in protein-protein recognition
 - Only some bacteria glycosylate few of their proteins
 - Viral proteins heavily glycosylated; helps evade the immune system



☐ GalNAc

Figure 7-30
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Glycoconjugates: Glycolipids

- A lipid with covalently bound oligosaccharide
 - Parts of plant and animal cell membranes
 - In vertebrates, ganglioside carbohydrate composition determines blood groups
 - In gram-negative bacteria, lipopolysaccharides cover the peptidoglycan layer

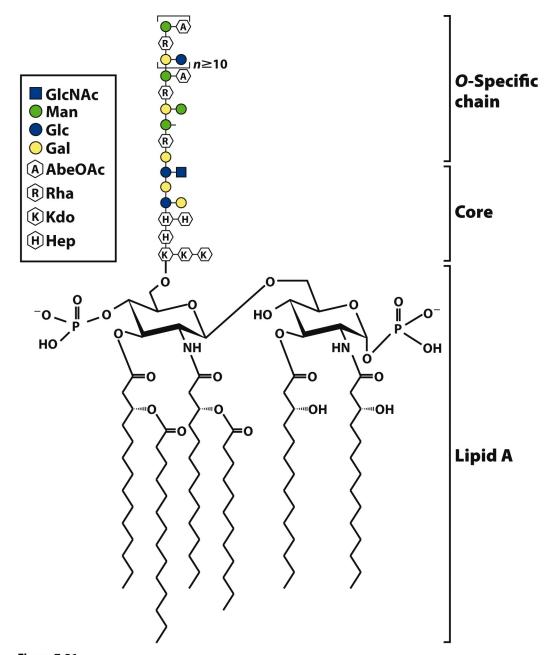


Figure 7-31
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Glycoconjugates: Proteoglycans

- Sulfated glucoseaminoglycans attached to a large rodshaped protein in cell membrane
 - Syndecans: protein has a single transmembrane domain
 - Glypicans: protein is anchored to a lipid membrane
 - Interact with a variety of receptors from neighboring cells and regulate cell growth

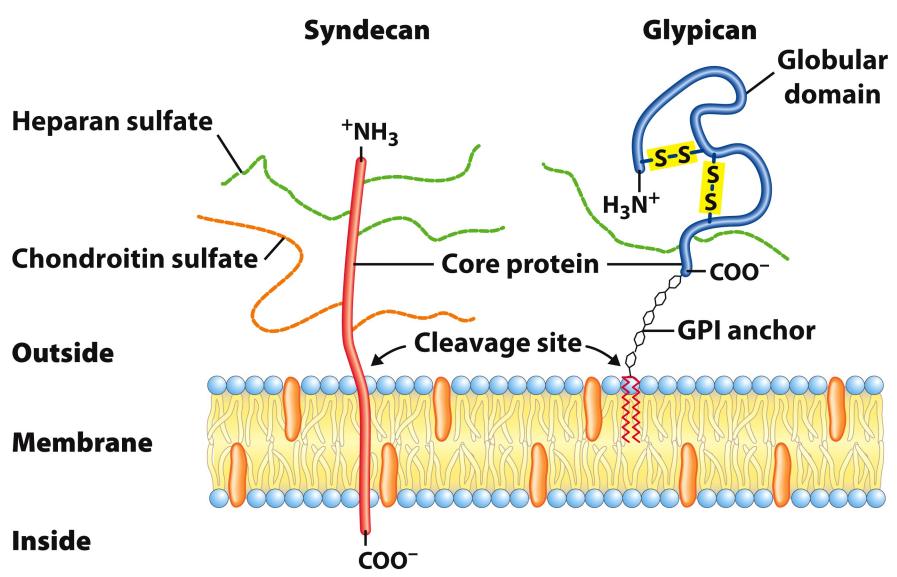


Figure 7-26a
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Heparan sulfate

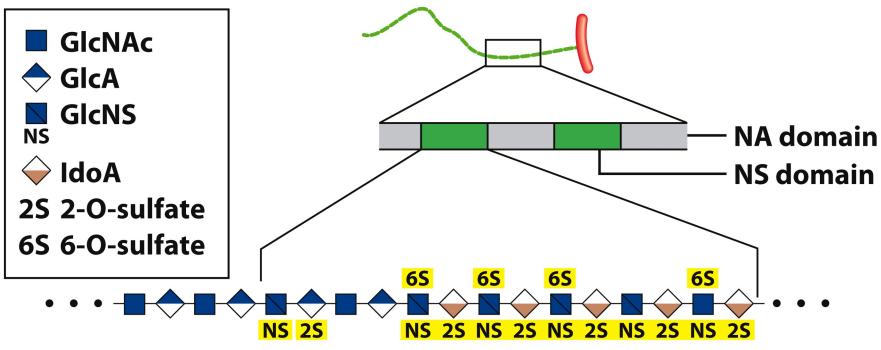


Figure 7-26b

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Proteoglycans

- Different glycosaminoglycans are linked to the core protein
- Linkage from anomeric carbon of xylose to serine hydroxyl
- Our tissues have many different core proteins; aggrecan is the best studied

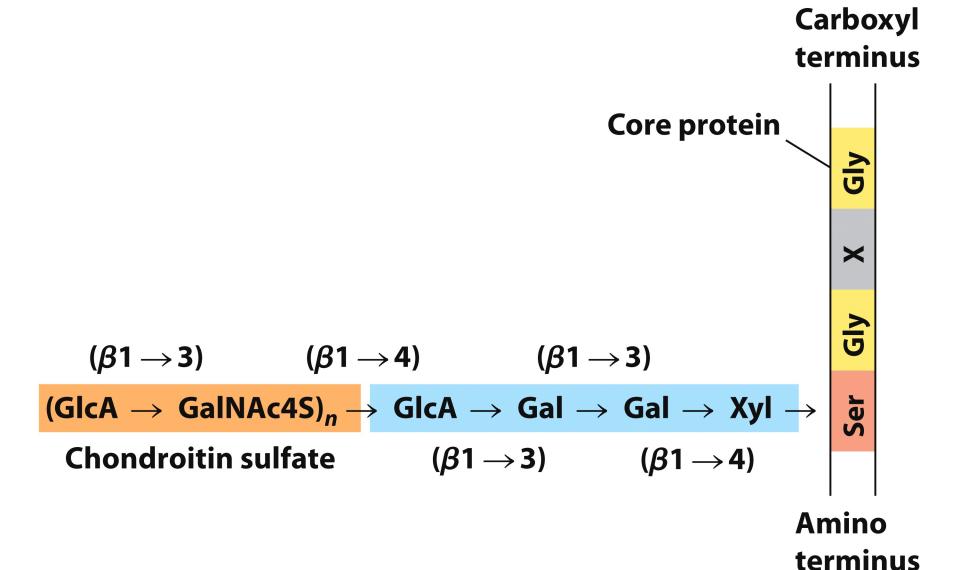


Figure 7-25

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Proteoglycan Aggregates

- Hyaluronan and aggrecan form huge (M_r > 2•10⁸) noncovalent aggregates
- Hold lots of water (1000× its weight); provides lubrication
- Very low friction material
- Covers joint surfaces: articular cartilage
 - Reduced friction
 - Load balancing

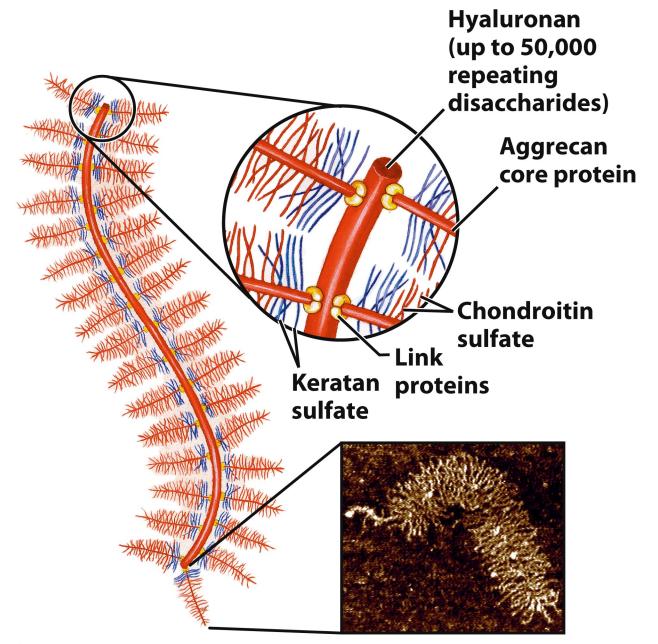


Figure 7-28
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