





Carbohydrates

pentose
hexose

Monosaccharides → 5 or 6 carbons
and are aldehydes and ketones

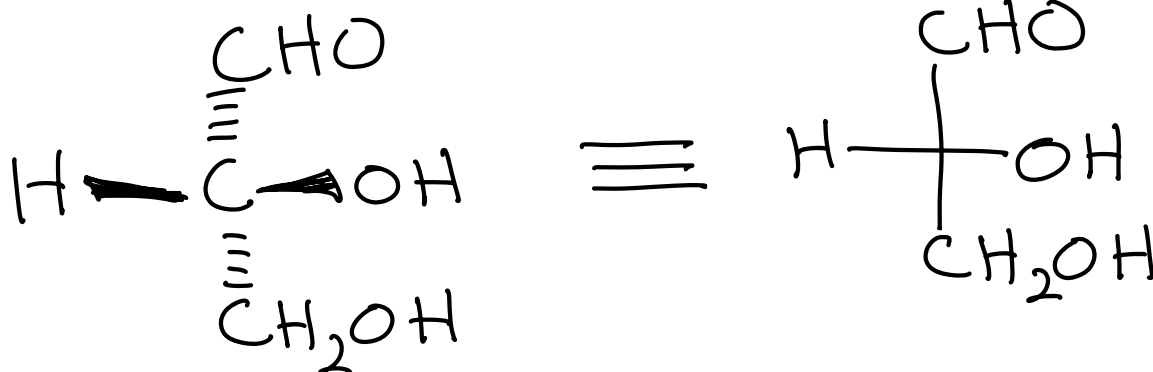
aldose

ketose

Ex.

Glucose is an aldohexose

Stereochemistry



(R)-D-Glyceraldehyde

↗
dextrorotatory
(+)

Called
Fischer
projection

"Like a Teddy Bear
giving you a hug"

Can I have
a hug please?

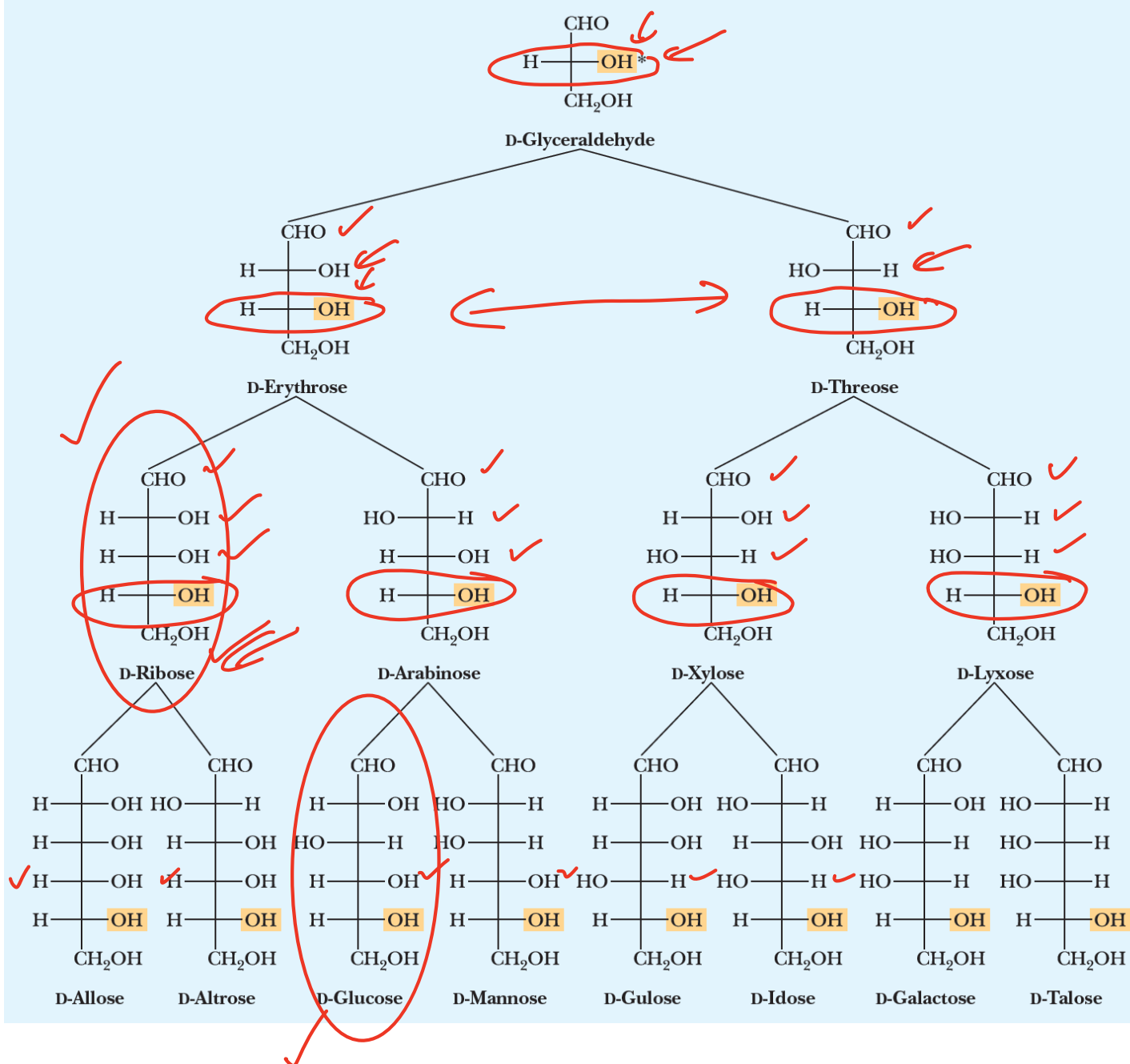


Stereochemistry defines the different carbohydrates

→ Due to the way carbohydrates are synthesized in cells → the common carbohydrates all have the same stereochemistry as D-glyceraldehyde at the carbon farthest from the carbonyl (aldehyde or ketone)

That is why they are called "D" carbohydrates

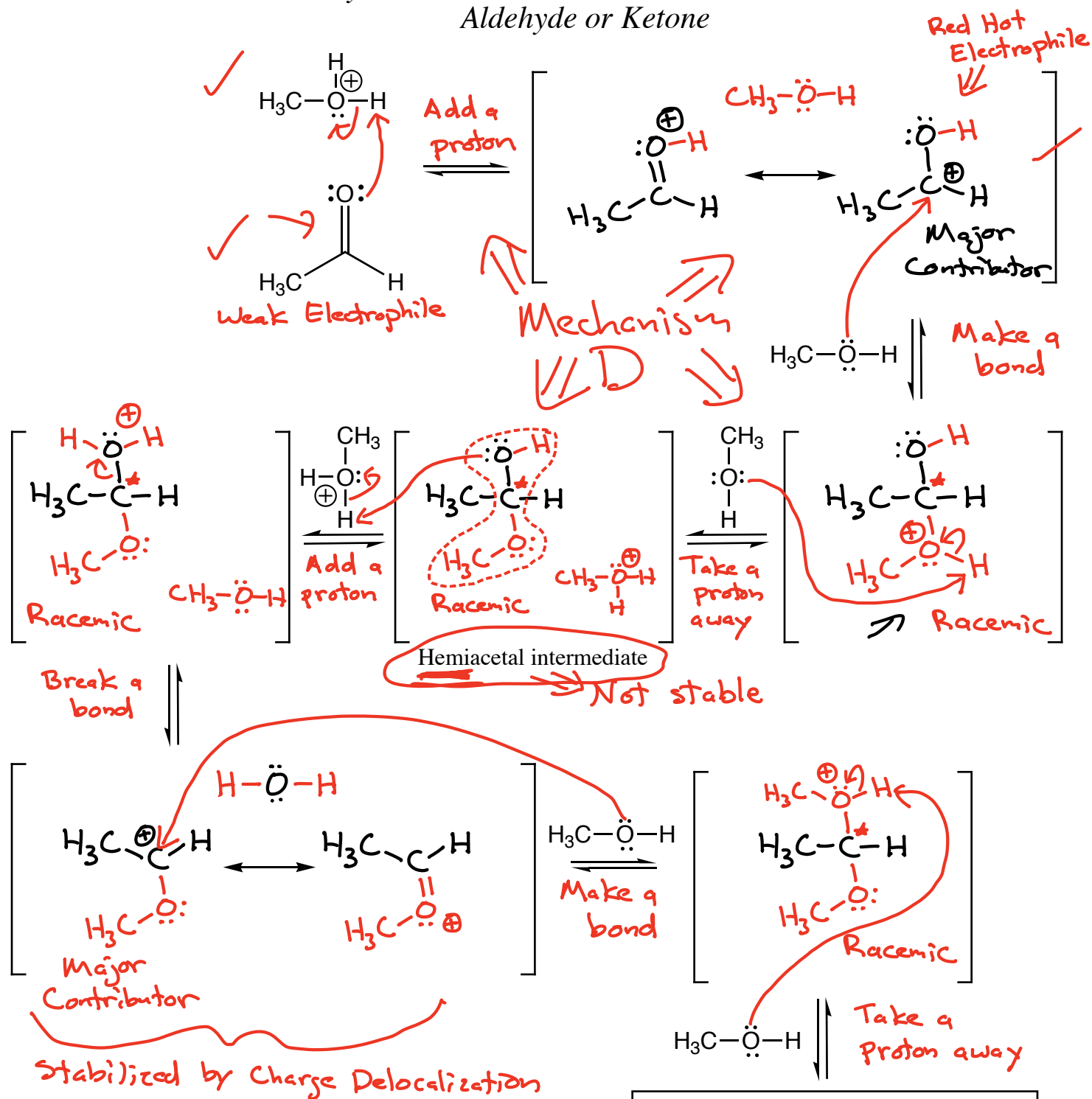
Table 25.1 Configurational Relationships Among the Isomeric D-Aldotetroses, D-Aldopentoses, and D-Aldohexoses





H₂SO₄ "Hex, does that thing have a hemi in it?" "SWEET!"

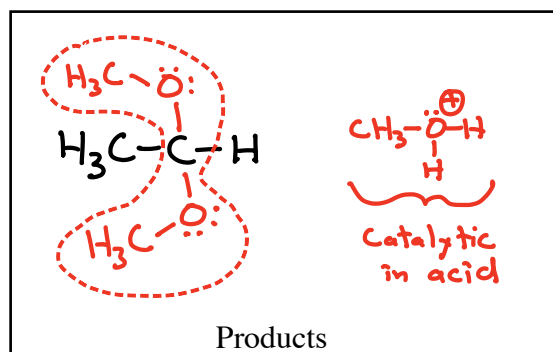
Acid Catalyzed Hemiacetal and Acetal Formation From an Aldehyde or Ketone



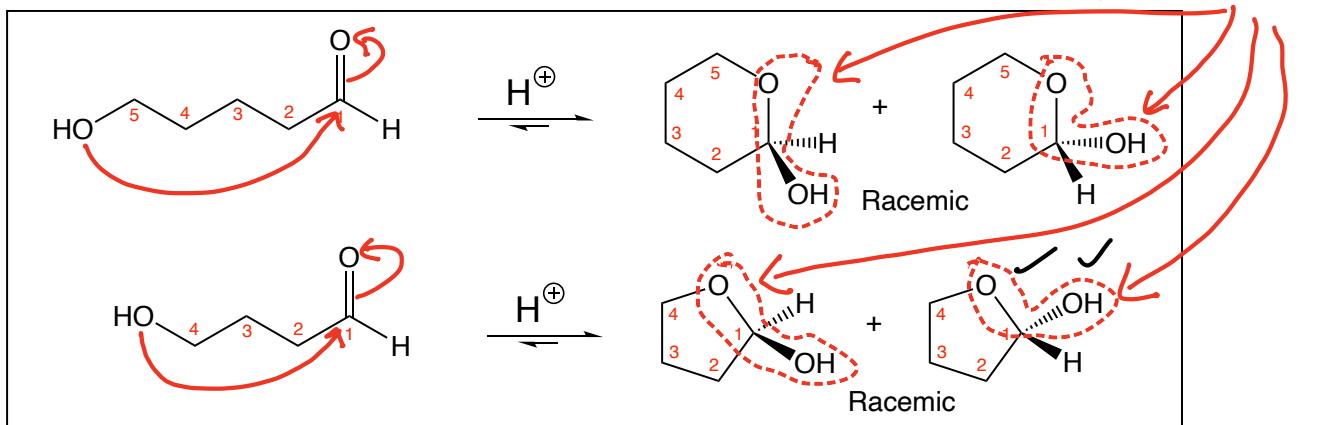
Key Recognition Element (KRE):

Two bonds to O
atoms from an sp³
C atom

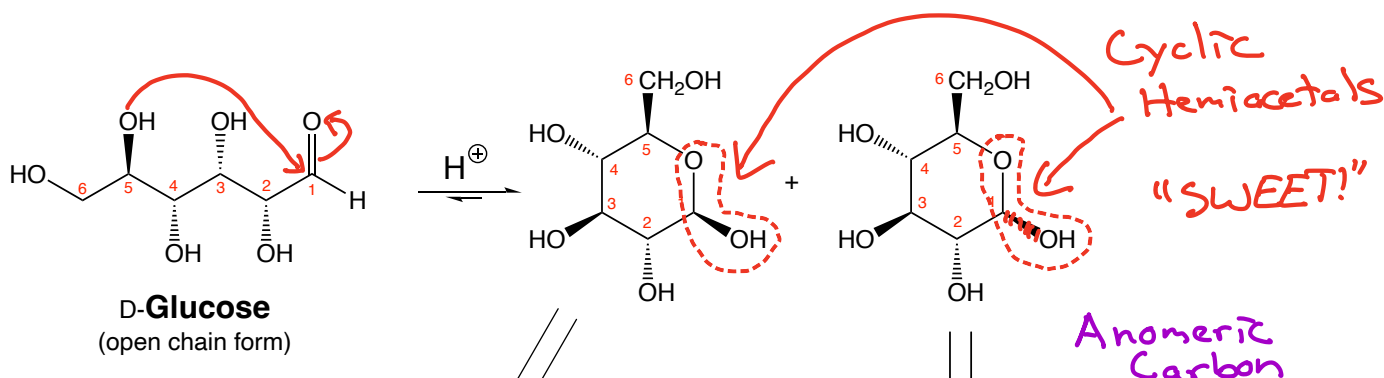
An
acetal



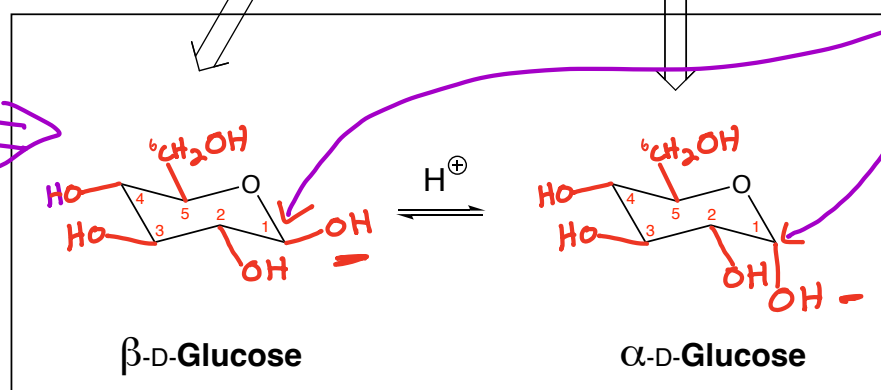
Cyclic Hemiacetals and Carbohydrates



The cyclic form of hemiacetals are stable - "SWEET!"
 → The chelate effect



This interconversion is called "mutarotation"



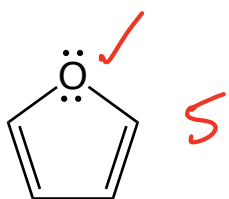
Biochemists call these two forms "anomers"

β -D-Glucopyranose
 means "6-membered ring"

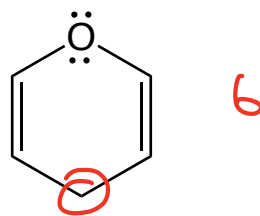
More stable → every group is equatorial!

α -D-Glucopyranose
 Less stable → one -OH is axial

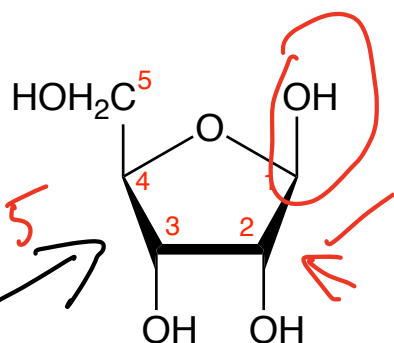
For Glucose:
 " α is axial!"



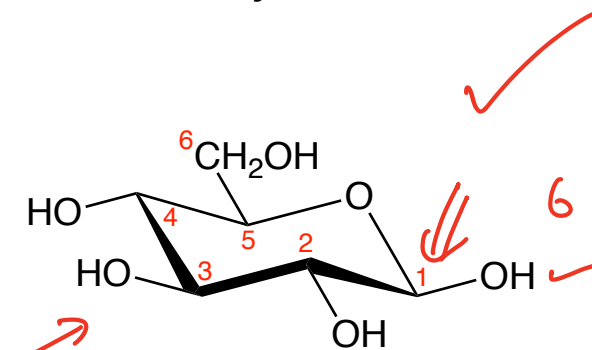
Furan ✓



Pyran ✓

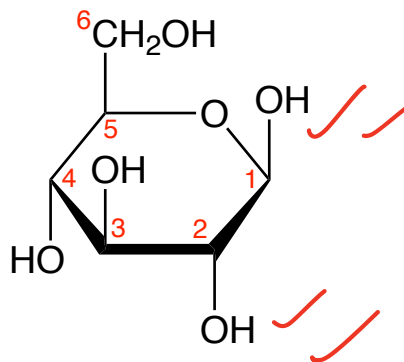


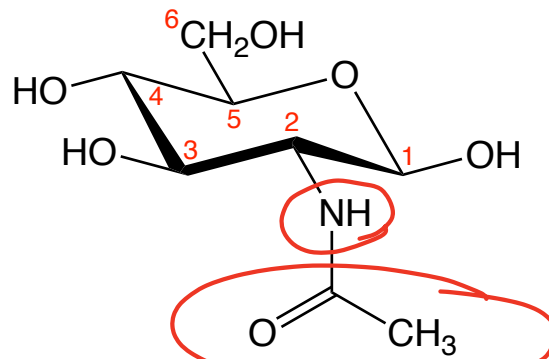
A furanose
 β -D-Ribose
 or β -D-Ribofuranose



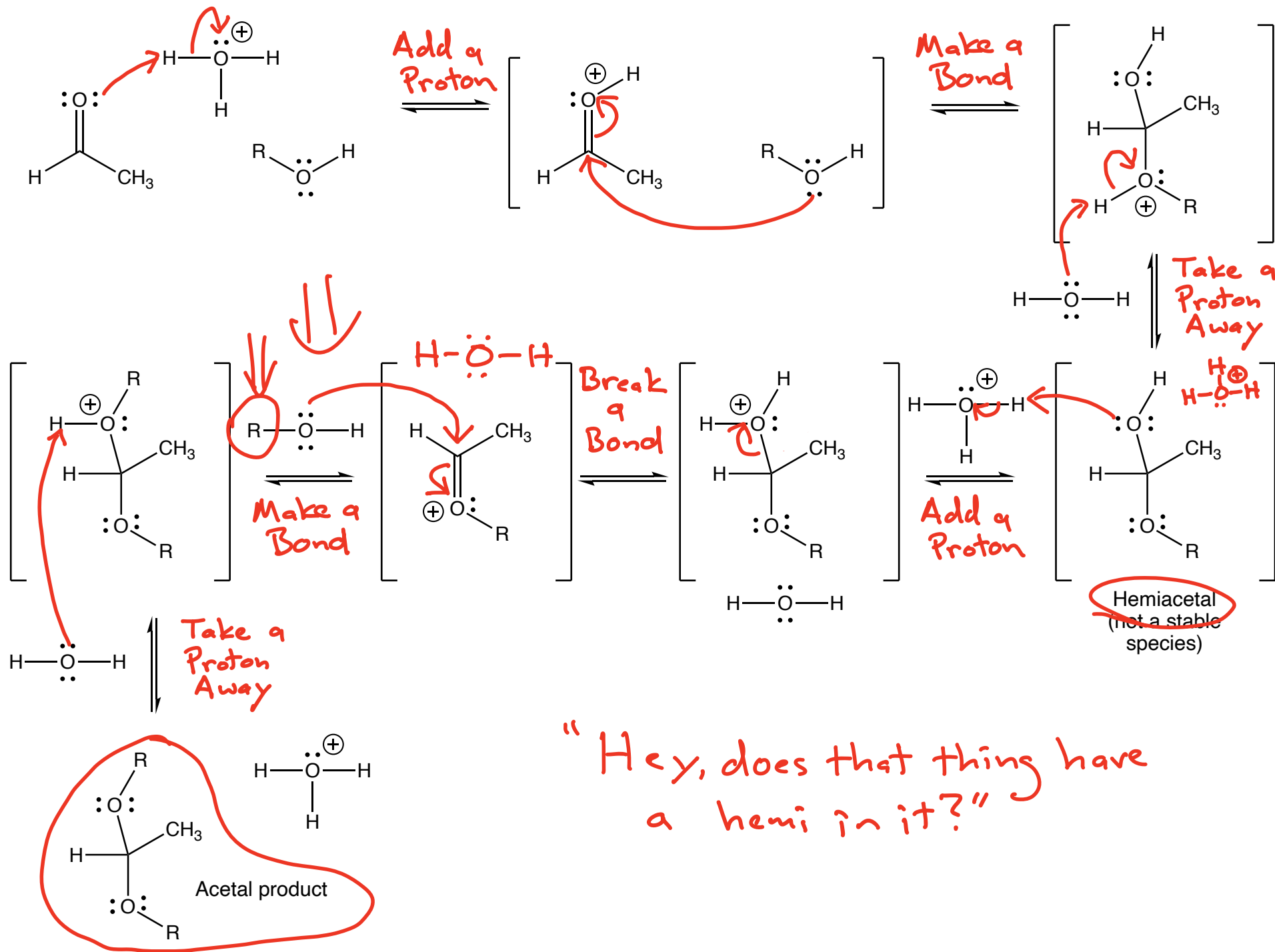
A pyranose
 β -D-Glucose
 or β -D-Glucopyranose

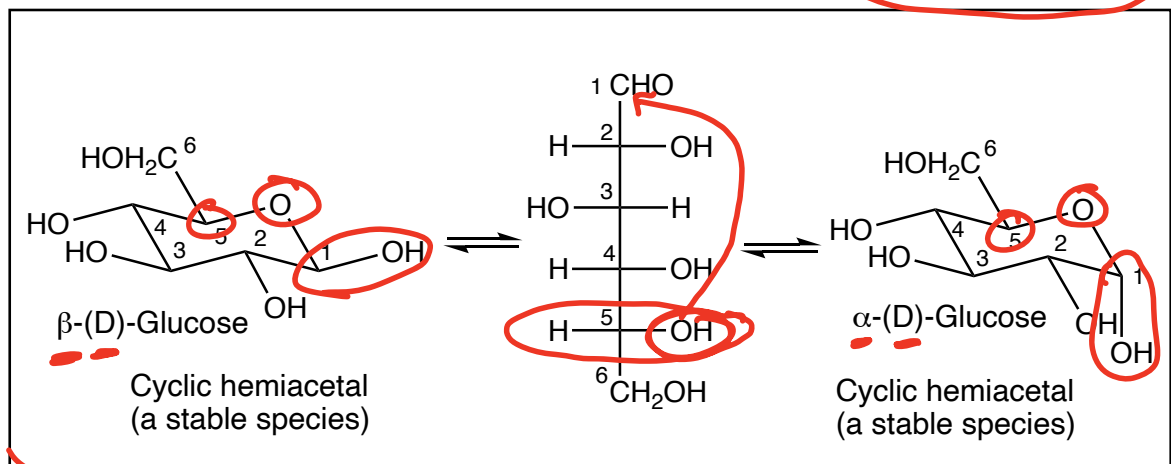
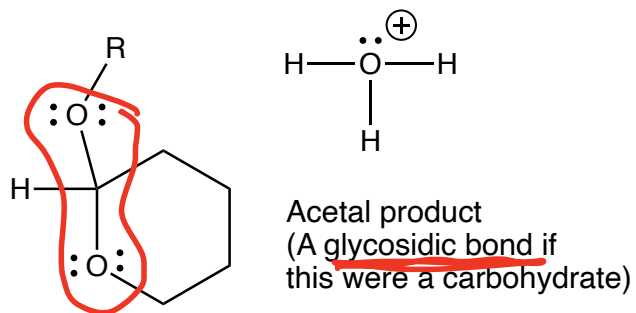
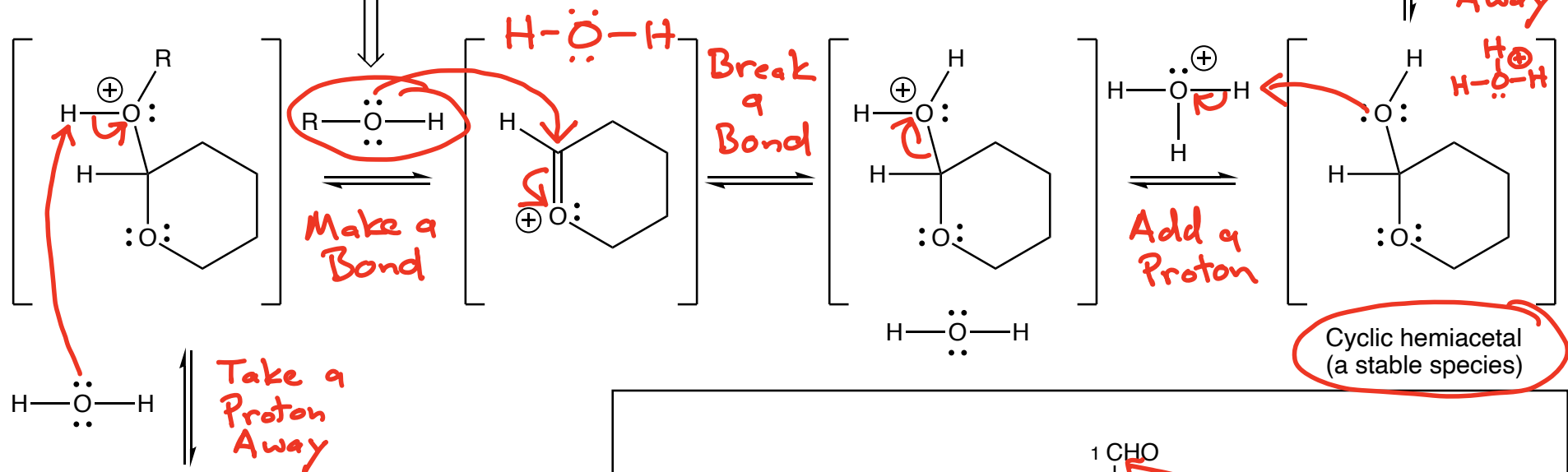
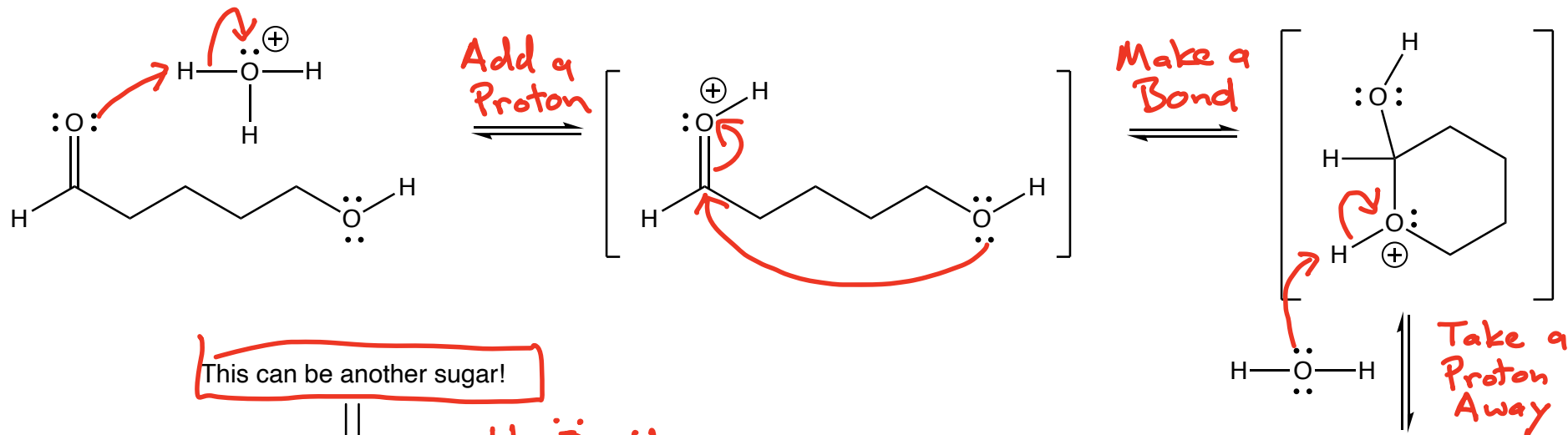
This is called a
 Haworth projection



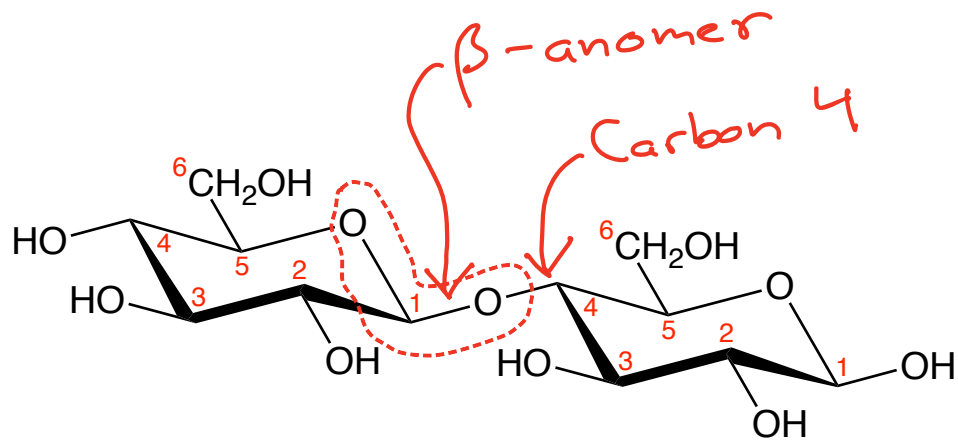


N-Acetyl-D-Glucosamine
(GlcNAc) ✓

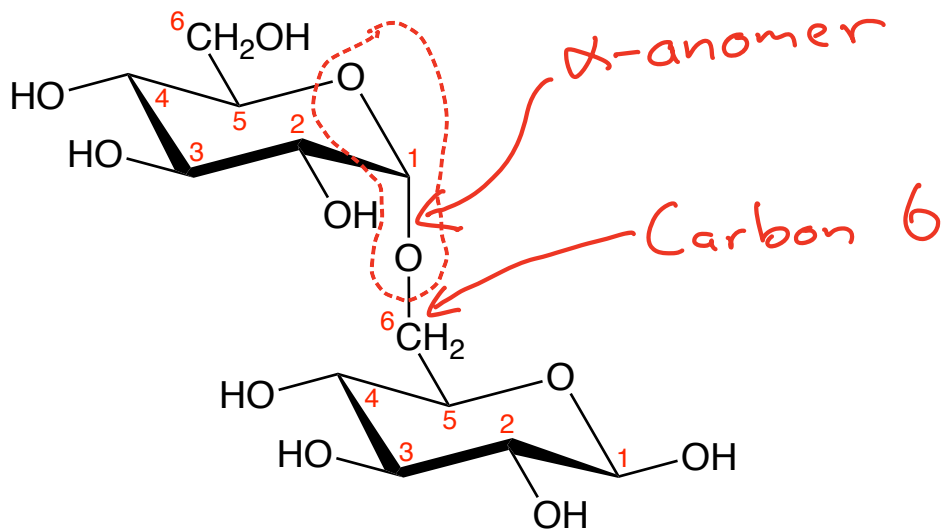




This process is called "Mutarotation"

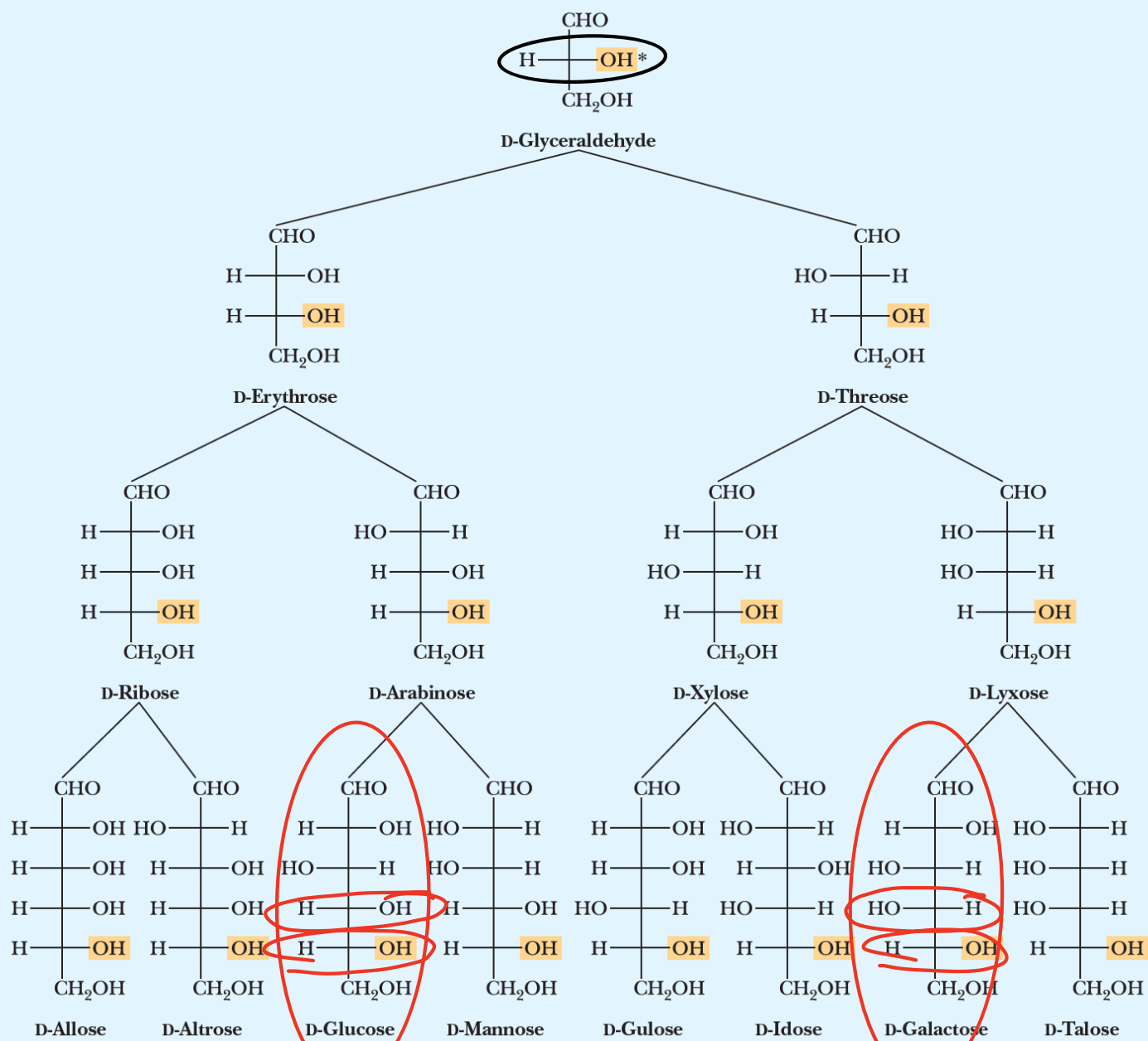


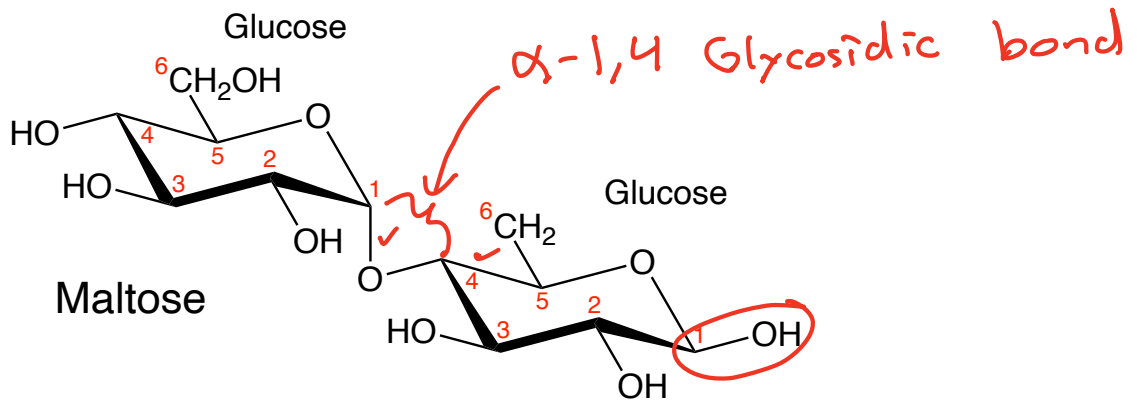
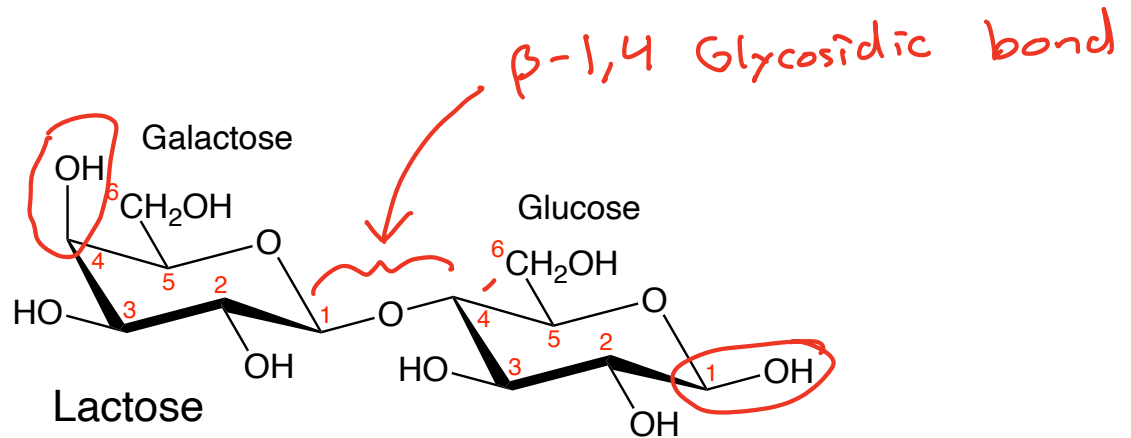
This is a β -1,4-Glycosidic Bond



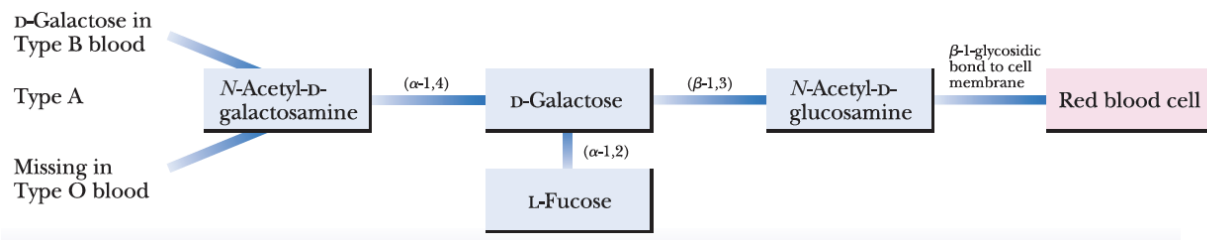
This is an α -1,6-Glycosidic Bond

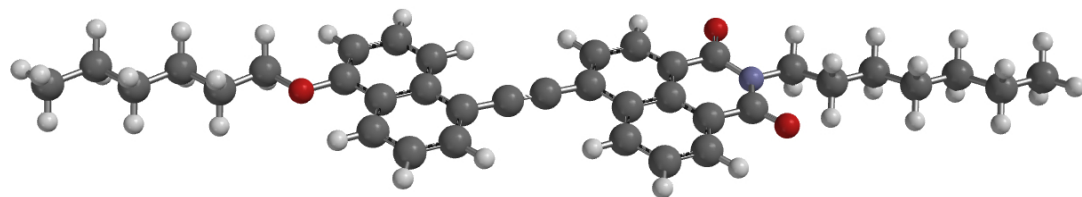
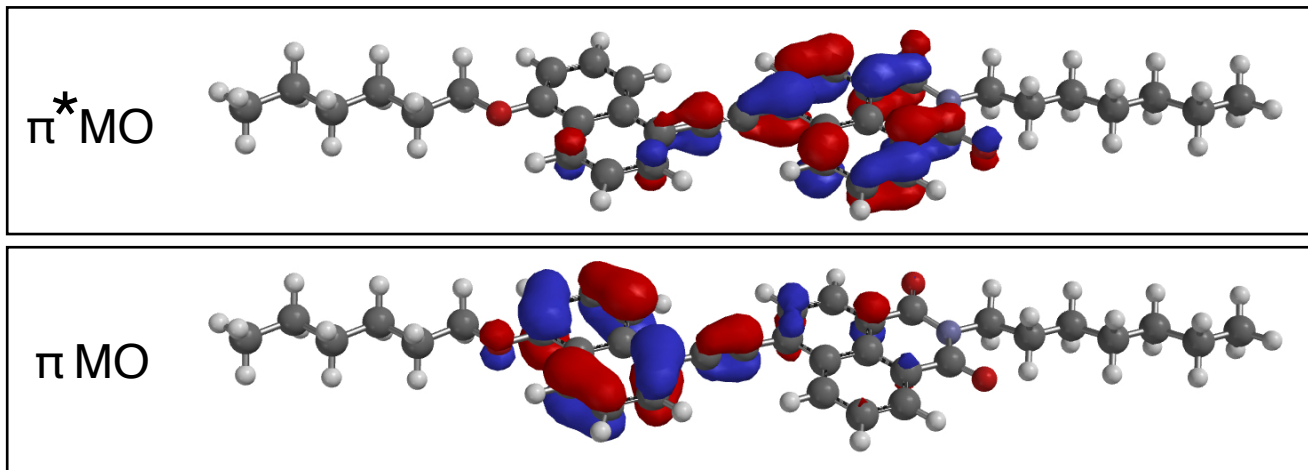
Table 25.1 Configurational Relationships Among the Isomeric D-Aldotetroses, D-Aldopentoses, and D-Aldohexoses





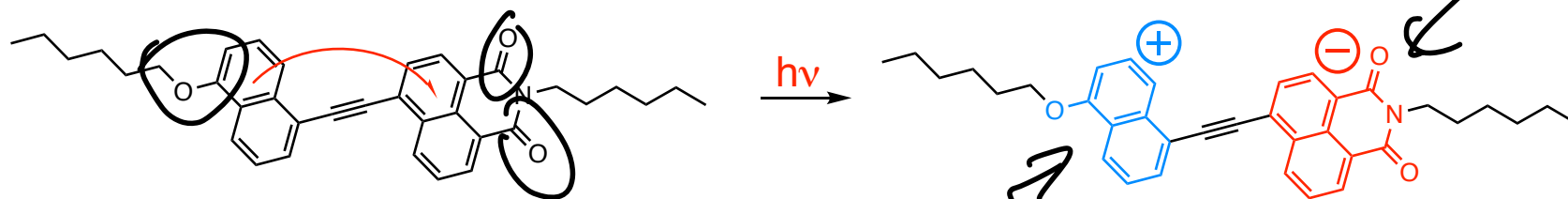
We can link more carbohydrates together, always at Carbon 1, with α or β linkages at carbons 2, 3, 4 or 6



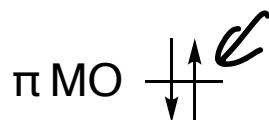


Blue Light = higher energy

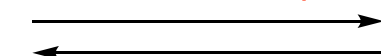
Orange light = lower energy




$\pi^* \text{MO}$ —




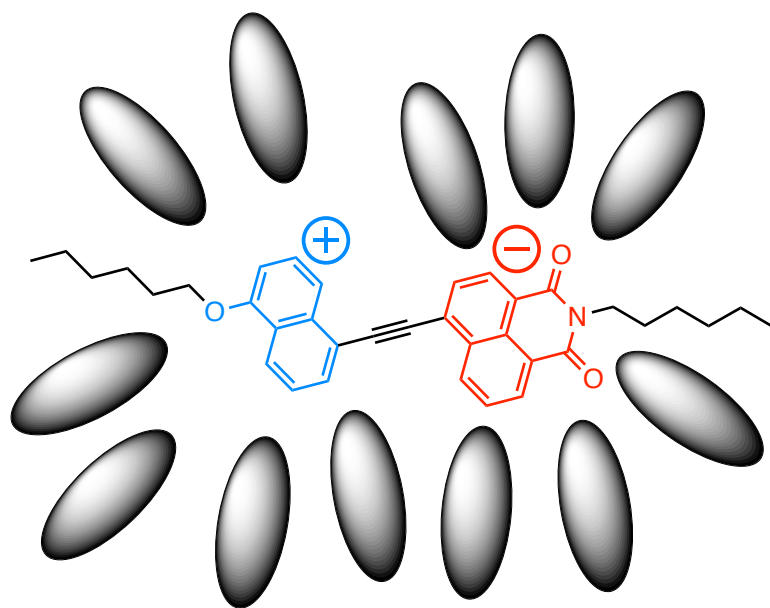
$h\nu$ absorb (UV)



$h\nu$ fluorescence (visible)

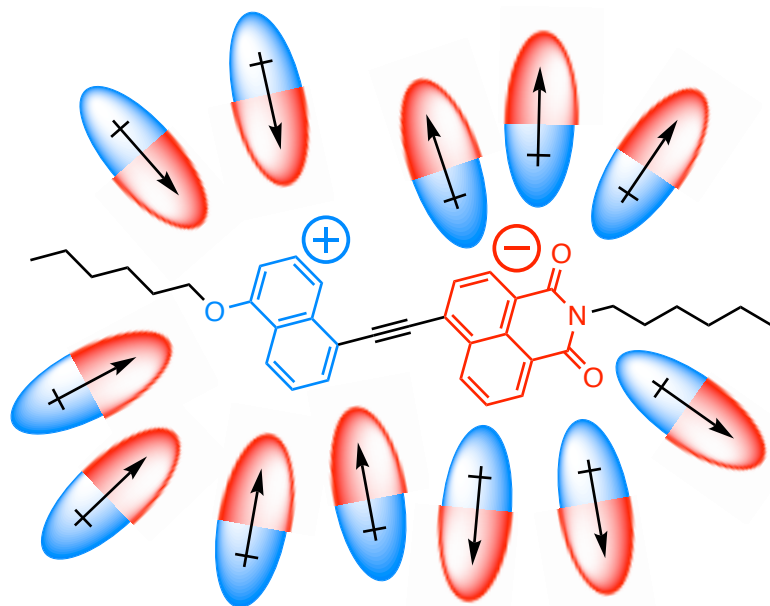
$\pi^* \text{MO}$ 

$\pi \text{ MO}$ 



Non-polar Solvent Molecule

No stabilization of the charged excited state



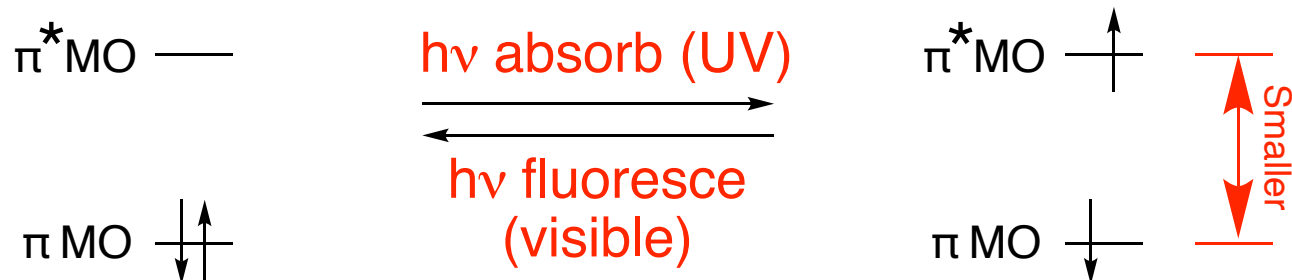
Polar Solvent Molecule

Stabilization of the charged excited state

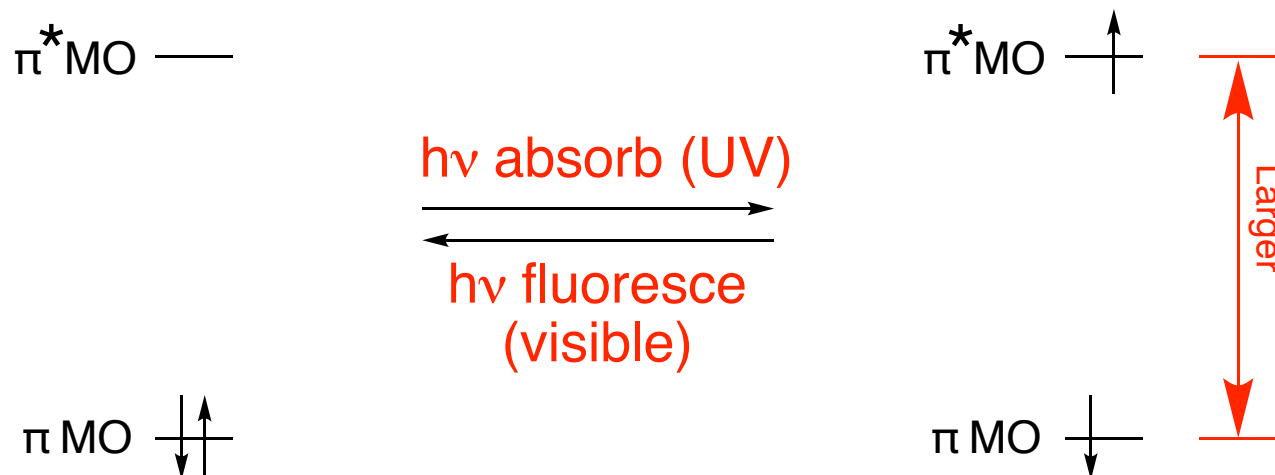
Which is blue?
Polar or
Non-polar
solvent?



← Less Polar Solvents
More Polar Solvents →



Polar Solvent - the charged excited state is well solvated because the solvent molecules have a relatively large molecular dipole that interacts favorably with the charges. There is a smaller energy difference between the ground state and excited state, corresponding to **orange light**



Non-polar Solvent - the charged excited state is disfavored because it is not well-solvated. There is no molecular dipole in the solvent to interact favorably with the charges. This makes the charged excited state higher in energy. There is a larger energy difference between the ground state and excited state, corresponding to **blue light**

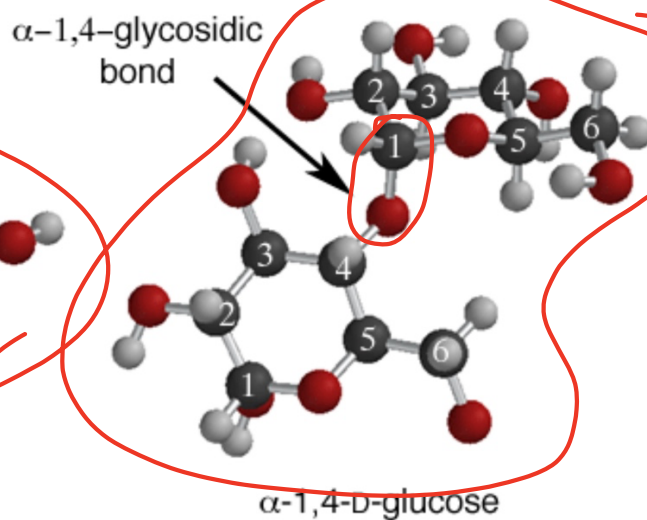
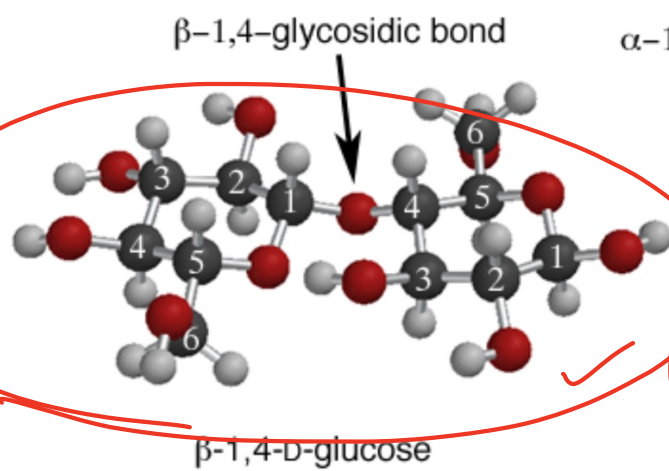
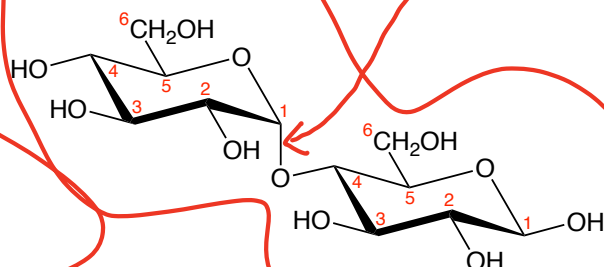
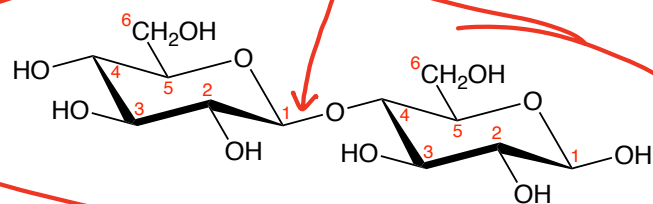
Answer to pop-quiz question

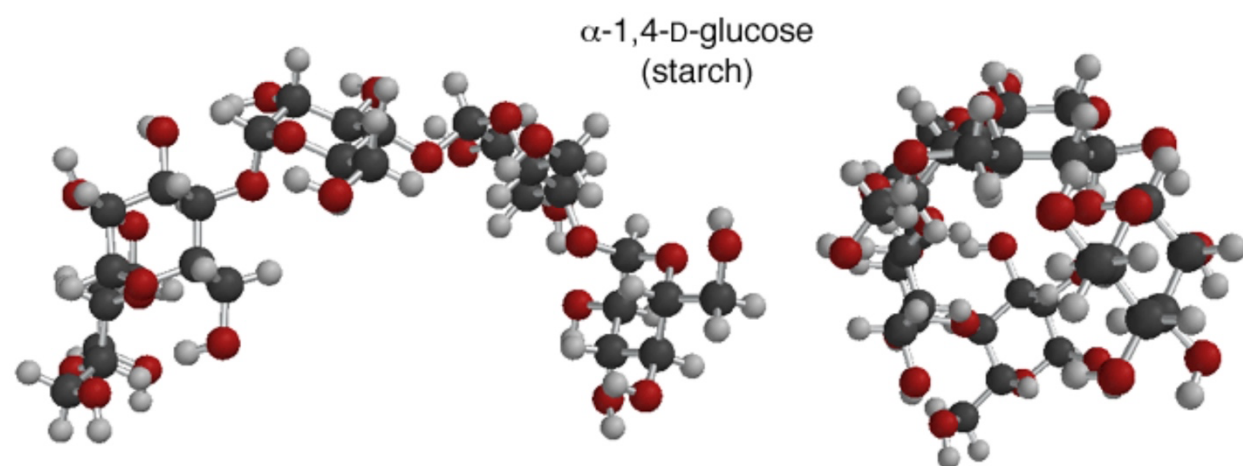
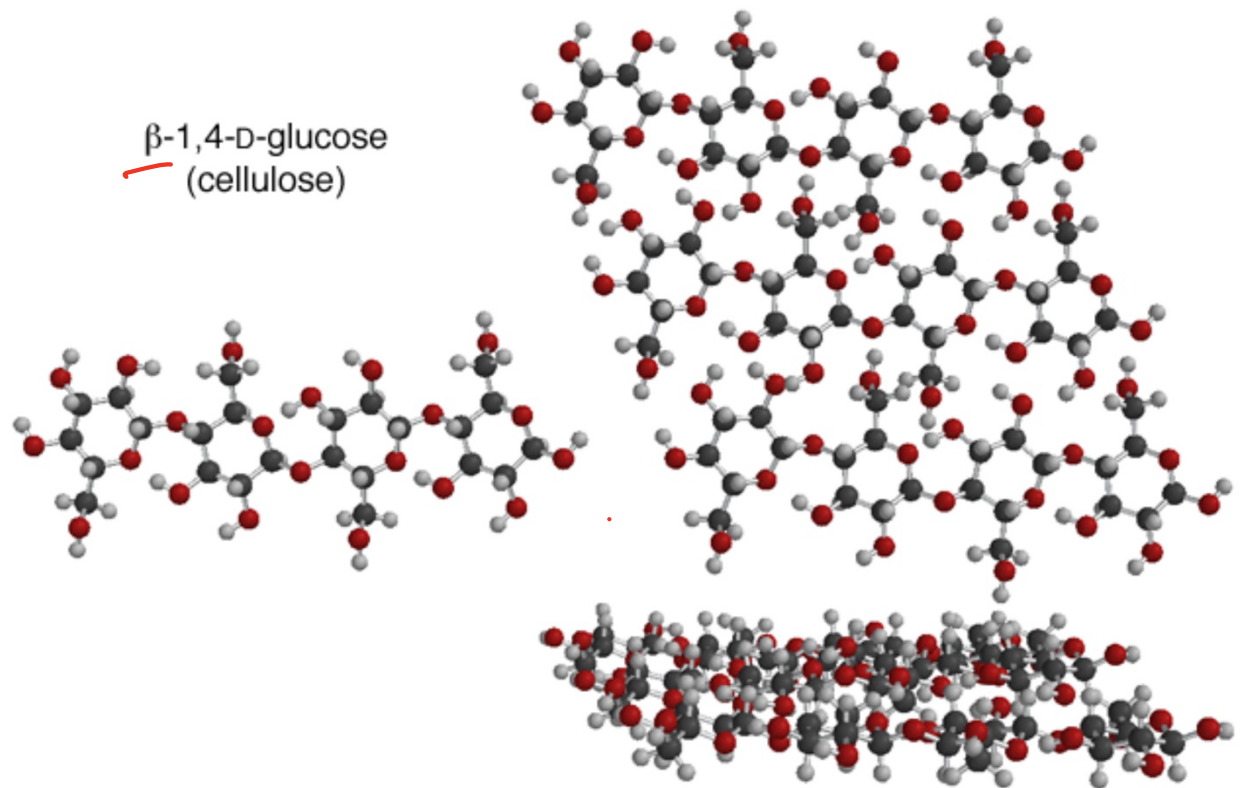


VS.



What is the difference —
They are both polymers
of D-Glucose







vs.

