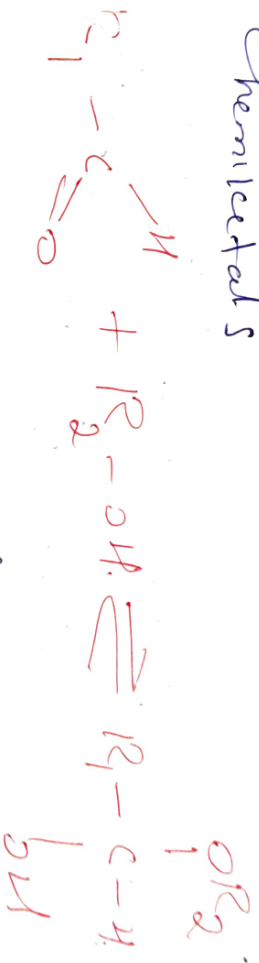


STRUCTURE OF ~~Glucose~~ Glucose

→ For a better understanding of glucose structure, let us consider the formation of hemiacetals and

Chemiacetals



Aldehyde

Alcohol

Hemiacetal

Mechanism:

→ When alcohol oxygen adds to the

Carbonyl Carbon of an aldehyde or ketone

→ There will be a nucleophilic attack of hydroxyl group at the electrophilic Carbonyl group

→ Since the alcohols are weak nucleophiles, the attack on the carbonyl carbon is usually promoted by protonation at the carbonyl oxygen

→ When this reaction takes place with an aldehyde, the product is called a hemiacetal and when the reaction takes place with a ketone, the product is referred to as a hemiketal

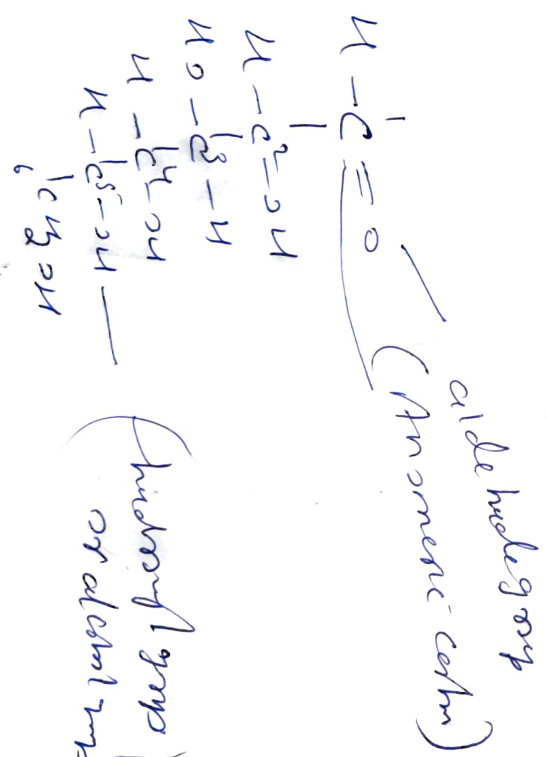


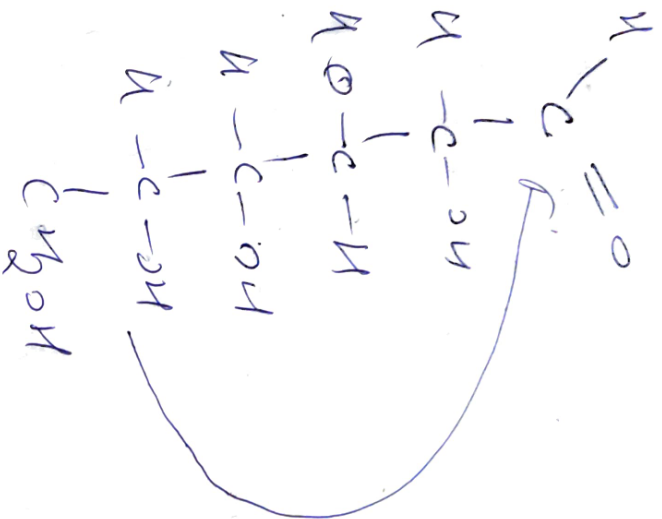
(2)

→ In the case of monosaccharides, the hydroxyl group ~~of~~ can react with its own aldehyde or keto functional group to form hemiacetal and hemiacetal.

→ Thus, the aldehyde group of glucose at C_1 reacts with alcohol group at C_5 to form two types of cyclic hemiacetals namely α and β

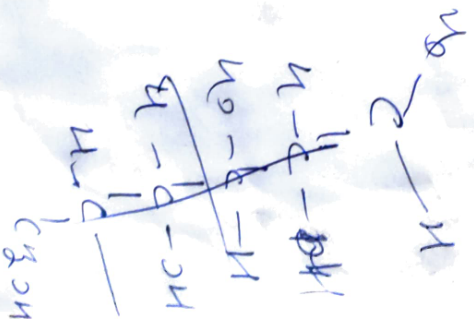
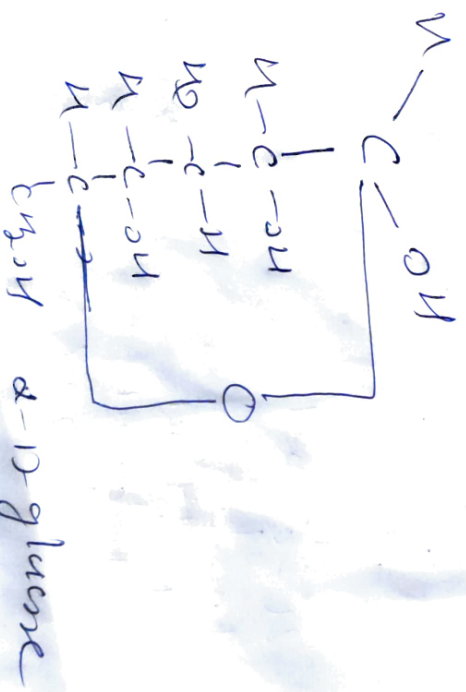
D-glucose





D-Glucose

(Fischer Projection)

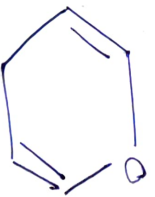


Pyranose and furanose Structures ⁽³⁾

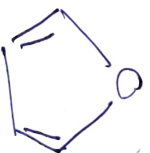
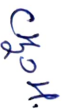
(Haworth Projection formulae)

→ Haworth projection formulae are depicted by a six-membered ring pyranose (based on pyran) or a five membered ring ~~&~~ furanose based on furan.

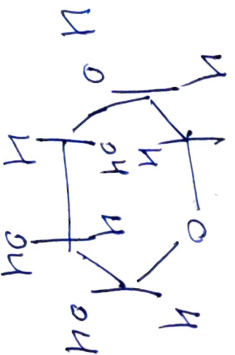
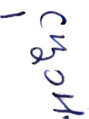
→ The cyclic forms of glucose are known as α-D-glucopyranose and α-D-glucofuranose.



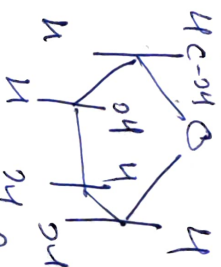
Pyran.



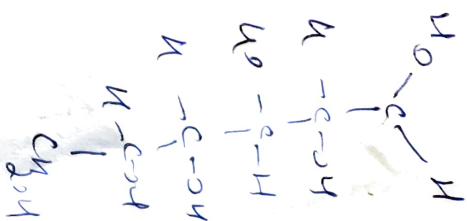
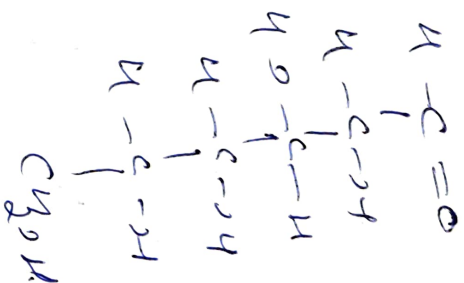
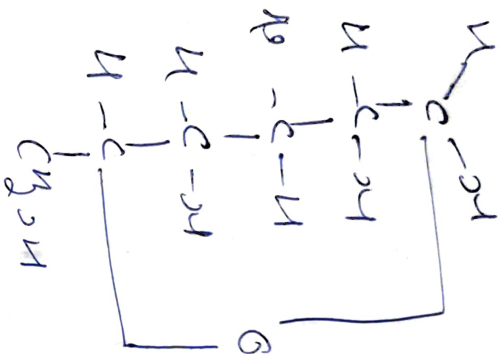
Furan



α-D-glucopyranose



α-D-glucofuranose.

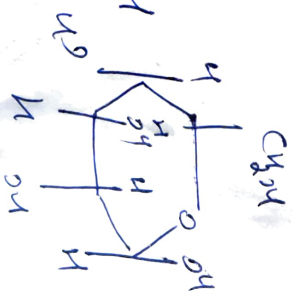
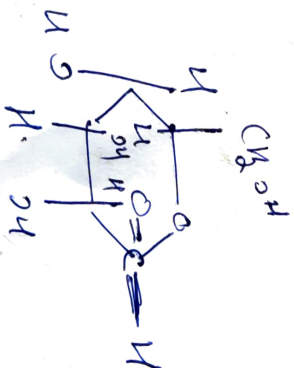
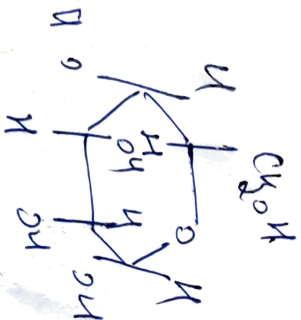


α-D-Glucose

D-glucose

Fischer projection formulae.

β-D-glucose



α-D-Glucopyranose.

D-glucose.

β-D-glucopyranose.

(Newman Projection formulae)

Anomers Mutarotation (2)

- The α and β cyclic forms of D-glucose are known as anomers
- They differ from each other in the configuration only around C₁ known as anomeric carbon (hemiacetal carbon).
- In case of anomer, the -OH group held by anomeric carbon is on the opposite side of the group -CH₂OH of sugar molecule
- The reverse is true for β -anomer
- The anomers differ in certain physical and chemical properties
- Mutarotation: The α and β anomers of glucose have different optical rotations
- The specific optical rotation of a freshly prepared glucose (α anomer) solution in water is $+112.2^\circ$ which gradually changes and attains an equilibrium with a constant value of $+52.7^\circ$.

→ In the presence of alkali, the decrease in optical rotation is rapid

→ The optical rotation of β -glucose is $+18.7^\circ$.

→ Mutarotation is defined as the change in the specific optical rotation representing the interconversions of α and β forms of D-glucose to an equilibrium mixture

α -D-glucose $\rightleftharpoons$ Equilibrium mixture $\rightleftharpoons$ β -D-glucose
 $+112.2^\circ$ $+52.7^\circ\text{C}$ $+18.7^\circ\text{C}$

→ The equilibrium mixture contains 63% β anomer and 36% α anomer of glucose with 1% open chain form

→ β -anomer form is more predominant due to its stable conformation