SHIKSHA CLASSES

Subject : Chemistry BOARD ANSWEF		SWER PAPER Total Marks	: 20
Class : XII Topic: 10. Halogen Derivatives			
Section (A) Q. 1. (a): Select and write the most appropriate answer from given alternatives in each sub-question [5]		Ans. : Enantiomer : The optical isomer whic non superimposable mirror images of other are called enantiomer.	h are each
i)	The strongest C-Cl bond present in	mixture of enantiomer (dextrorotatary laevorotatory) is called racemic mixture.	v and
Ans: (a)		ii) What is primary Alkyl Halide?	
	Aromatic C - Cl bond length is less than aliphatic C - Cl. Thus less bond distance stronger the bond.	Ans.: In this type of alkyl halide, the carbon w is bonded to the halogen family will be attached to one other alkyl group. It do matter how much bulky group is attached	which only esn't to it.
ii)	Which of the following structure is Vinyl	Section (B)	
	Chloride?	Q.2 Answer the following question.	
		(Any three)	[6]
Ans.: (a) 0		i) What are the halogen derivative alkane, classified with examples?	es of
	CI	Ans:Halogen deriratives of alkane :	
	Vinyl	The hydrocarbon in which, replacement o or more than one hydrogen atom by one r than one halogen atom is known as Hal- derivates of hydrocarbon or alkane'.	f one more ogen
)	SP-Hybridisation carbon atom	The general molecular formula of Hale	ogen
Ш) Алс	i ne naiogenating reagent is	derivatie of alkane is	
iv)	Which of the following is better reactant for SN ¹ Reaction?	$\begin{bmatrix} C_n H_{2n+2} & \xrightarrow{-H} \\ C_n H_{2n+2} & \xrightarrow{-2H} \\ C_n H_{2n+2} & \xrightarrow{-2H} \\ C_n H_{2n+2} & \xrightarrow{-3H} \\ C_n H_{2n+2} & \xrightarrow{-3H} \\ \hline \end{bmatrix} $ "Halogen derivatives of Allowed environments of Allowed environments of the set o	lkane"
Ans: (d)		The halogen derivatives of alkanes classiffied on the basis of number of hal atom present in alkane.	s are ogen
.,	Ans: a) 3-chloroprop-1-ene	1) Mono halogen derivative of Alkane.	
Q.1	(b) : Very short answer type Question [2]	2) Poly halogen derivative of Alkane.	
i)	Write defination of enantiomer and racemic mixture?	1) Mono halogen derivative of Alkan	ne :





y) What are Optical Activity? Explain with examples.

Ans. : Optical Activity : When plane polarized light is pass through glucose, lactic acid(organic solution), the plane polarized light gets rotated through certain angle either clockwise direction or anti-clockwise direction (right hand side & left hand side) the phonomenon is known as "optical activity".

Example : When an aqueous solution of certain organic compounds like sugar, lactic acid is placed in the path of plane polarized light, the transmitted light has oscillations in a different plane than the original. In other words, the incident light undergoes rotation of its plane of polarization. The plane of polarization rotates either to the right (clockwise) or to the left (anticlockwise).

Thus, this process / phenomenon is known as "optical activity."

Q.3 Answer the following question (Any one) [3]

Explain the nature of C-X bond in aliphatic i) and aromatic halogen compounds.

Ans. : Nature of C-X bond :

(A)
$$\begin{array}{c} H_3C - C1 \\ \uparrow \\ 1.77A^0 \end{array}$$
 (B)
$$\begin{array}{c} Ar - C1 \\ \uparrow \\ 1.70A^0 \end{array}$$

Aliphatic compound

(b)

Aromatic compound

i)

The lone pair ele. present on chlorine atom going to delocalization and thus create a π -bond between carbon of aromatic nuclears and chlorine atom thus bond distance automatically decreases than C-Cl aliphatic or single bond between C-Cl/C-Xspeices.

5% is less

5% is more Thus, less bond difference between aromatic carbon & chlorine atom.

Explain Friedel Craft Alkylation and ii) Acylation reaction with its examples.

Ans: Alkylation : Aryl halide react with methyl chloride in presence of anhydrous AlCl, it gives methyl arylhalide

Acvlation: When Chlorobenzence react with acyl chloride in presence of unhydrous AlCl, it gives 1-chloro acetophenone and 4-chloro acetophenone.

to form 2-methyl chlrobenzene (minor) & 4methyl chlorobenzene (major) products. Such types of reaction is "Fredel craft Alkylation Reaction of chlorobenzene.

2) Friedal Craft Acylation Reaction of Chlorobenzene

When chlorobenzene is heated with acetyl chloride or acetic anhyldride in presence of aluminium trichloride (lewis acid) to form 2chloro acetophenone (minor) and 4-chloro acetophenone is major and AlCl₂ remain it (as -catalyst).

The 4-chloro acetophenone is major this is due

to absence of steric crowding between chlorine atom and acetyl group.

OR

ii) What are Finkelstein and Swartz reaction?

- **Ans. : Finkelstein reaction :** Alkyl iodides are prepared conveniently by treating alkyl chlorides or bromides with sodium iodide in methanol or acetone. This reaction is called finkelstein reaction.
- $R Cl + NaI \xrightarrow{Acetone} R I + NaCl \downarrow$

Swartz Reaction : Alkyl fluorides are prepared by heating alkyl chlorides or bromides with metal fluorides such as AgF, Hg_2F_2 , AsF_3 , SbF_3 etc. is called Swartz reaction.

 $R-Cl+AgF \longrightarrow R-F+AgCl \downarrow$

- iii) What are the SN² reaction, explain with an example?
- Ans.: Bimolecular Nucleophilic Substitution Reaction (BNSR)- is a second order one step reaction.

When methyl bromide is treated with sodium hydroxide formed methyl alcohol and NaBr-Sodium bromide as side product.

Reaction :

The Nucleophilic Hydroxy group from sodium hydroxide i) attack on carbon atom of methyl bromide from backside and generate transition state and fast generate product.

Thus, this is an first step of reaction mechanism (Simultaniously nucleophile attack and Leaving of nucleophile).

Kinetic :

Rate \propto [CH₃Br] [NaOH] \ominus Rate \propto [CH₃Br] [OH] \therefore Rate = K [CH₃Br] [OH] K = Rate constant

The rate of reaction is depends upon both reactants i.e. $CH_3Br + NaOH$.

 \therefore SN² reaction is second order one step Reaction.

EPD For SN² Reaction :

The graph is ploted potential energy verses reaction co-ordinate is energy profile diagram. The energy difference between reactant and product is heat of enthalpy (Δ H).

 $\therefore \Delta H = R - P.$

The energy difference between reactant and transition state is known as "change in energy ΔE "

 $\therefore \Delta E = R - T$. State.

On the basis of above representation.

In case heat is absorbed then $\Delta H = \bigoplus$ ve (endothermic), then product formed is less stable, while heat is released then ΔH is \bigoplus ve (exothermic) & product formed is stable.

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