

INDIAN SCHOOL AL WADI AL KABIR

UNIT TEST (2024-25)

CLASS: XII Sub: CHEMISTRY (043) MAX.MARKS: 30

DATE: 04/06/2024 Set - I TIME: 1 HOUR

Q.NO.	ANSWER KEY	MARKS
1	С	1
2	D	1
3	С	1
4	A	1
5	В	1
6	A	1
7	С	1
8	A	1
9.	a. b. CH ₃ CH ₂ CH=CH ₂	1+1
10.	a.	1
	By hydroboration-oxidation: Diborane $(BH_3)_2$ reacts with alkenes to give trialkyl boranes as addition product. This is oxidised to alcohol by hydrogen peroxide in the presence of aqueous sodium hydroxide.	
	$\begin{array}{c} \text{CH}_{3}\text{-CH}\text{-CH}_{2} \ + \ (\text{H}\text{-BH}_{2})_{2} \longrightarrow \text{CH}_{3}\text{-CH}\text{-CH}_{2} \\ \text{I} \ \text{BH}_{2} \\ \text{H} \ \text{BH}_{2} \\ \text{CH}_{3}\text{-CH}\text{-CH}_{2} \\ \text{CH}_{3}\text{-CH}_{3} \\ \text{CH}_{3}\text{-CH}_{3} \\ \text{CH}_{3}\text{-CH}_{3} \\ CH$	
	Propan-1-ol b.	
	It is an important laboratory method for the preparation of symmetrical and unsymmetrical ethers. In this method, an alkyl halide is allowed to react with sodium alkoxide.	1
	$R-X+R-\ddot{\ddot{Q}}$ \dot{Na} \longrightarrow $R-\ddot{\ddot{Q}}-R'+Na$ X	
	Ethers containing substituted alkyl groups (secondary or tertiary) may also be prepared by this method. The reaction involves $S_{N}2$ attack of an alkoxide ion on primary alkyl halide.	
11.	a.	
	ÓН	1
	$+ Z_{n} \longrightarrow \bigcirc + Z_{nO}$	

		b.	
			1
		OH OH Br Pr	
		n n	
		+ 3 Br ₂ → + 3H Br	
		Fr.	
		2,4,6-Tribromophenol	1
		c.	1
		011	
		OH ONa OH	
		NaOH (i) CO ₂ COOH	
		(ii) H [*]	
		2-Hydroxybenzoic acid	
F	10	(Salicylic acid)	1
	12.	a. C—Cl bond acquires a partial double bond character due to resonance.b. The carbocation thus formed gets stabilised through resonance	1
			1
F	13.	c. AgCN is mainly covalent, and nitrogen is free to donate electron pairs. a. 5-Ethylheptane-2,4-diol	1 1
	13.	b.	1
		Mechanism	
		Step 1: Formation of protonated alcohol.	
		$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	
		н-С-С- О-н+№ ⇒ н-С-С- О-н н н н н	1
		Ethanol Protonated alcohol (Ethyl oxonium Ion)	
		Step 2: Formation of carbocation: It is the slowest step and hence, the	
		rate determining step of the reaction.	
		H H H H H H H-C-C- O-H Slow H-C-C ⁺ + H₂O H H H H H	1/2
		11 11	
		Step 3: Formation of ethene by elimination of a proton. H H	
		$H - C \stackrel{1}{\longrightarrow} C^{\dagger} \stackrel{H}{\longleftrightarrow} H + H^{\dagger}$	1/
		H H H Ethene	1/2
		The acid used in step 1 is released in step 3 . To drive the equilibrium	
		to the right, ethene is removed as it is formed. \mathbf{OR}	
		i. 4 -Ethoxy-1 -1 – dimethyl cyclohexane	1
		ii.	
		The formation of ether is a nucleophilic bimolecular reaction (S_x 2)	
		involving the attack of alcohol molecule on a protonated alcohol, as indicated below:	
		(i) CH₃-CH₂-O-H + H˙ → CH₃-CH₂-Ö-H	1
		(ii) $CH_3CH_2 - \ddot{O}$: + $CH_3 - CH_2 - \ddot{O}$ $H \rightarrow CH_3CH_2 - \ddot{O} - CH_2CH_3 + H_2O$	1/
		н Н	1/2
		(iii) $CH_3CH_2 = \stackrel{\uparrow}{O} - CH_2CH_3 \longrightarrow CH_3CH_2 - O - CH_2CH_3 + \stackrel{\uparrow}{H}$	
		H H	1/2
1			, ~

14.	a.	1
	Br	
	CH-CH ₃	
	O_2N	
	b. Isopropyl chloride<1-Chloropropane<1-Chlorobutane	1
	c. stereoisomers related to each other as non-superimposable mirror images	1+1
	are called enantiomers. They differ by optical rotation. OR	111
	c. Chiral + Asymmetric carbon	1+1
15.	a.	1/ 1/
	ОН	$\frac{1}{2} + \frac{1}{2}$
	+ CH ₃ CH ₂ Br	
	b.	
	ONa	1
	c.	
	au au	1
	CH - CH ₃	-
	ОН	
	d.	
	он он	
	J. Br J.	1/2 + 1/2
	Br	
	e.	
	ÇH ₃	
		1/2 + 1/2
	$CH_3 - \dot{C} - OCH_3 + NaBr$	
	ĊH ₃	
	OR	
		1

