

## INDIAN SCHOOL AL WADI AL KABIR

## **REHEARSAL EXAMINATION-I**

08/12/2022 CHEMISTRY (043) Maximum Marks: 70

Class: XII Time: 3 Hours

## SET – I – ANSWER KEY

1.	(A) osmotic pressure	1
2.	(C) first order reaction	1
3.	(B) Argon	1
4.	(C) (iii) and (iv)	1
5.	(D) 2	1
6.	(D) +3	1
7.	(A) i, iv	1
8.	(D) Benzyl halides are more reactive than vinyl and aryl halides	1
9.	(B) a dehydrohalogenation reaction	1
10.	(B) Scandium	1
11.	(C) Linkage Isomers	1
12.	(C) Polypeptides	1
13.	(A) Acetone	1
14.	(B) 2-Methylbutan-2-ol	1
15.	(C) Assertion is correct statement but reason is wrong statement.	1
16.	(D) Assertion is wrong statement but reason is correct statement.	1
17.	(A) Assertion and reason both are correct statements and reason is correct explanation for assertion.	1
18.	(C) Assertion is correct statement but reason is wrong statement.	1
19.	a) Oxygen stabilizes Mn more than F due to multiple bonding.	1
	b) This is due to decrease in size and increase in mass from titanium to copper.	1
20.	k = 0.693/t	1/2
	$k = 0.0277 \text{ min}^{-1}$	1/2
	$t_{80\%} = (2.303/0.0277) \log 100/20$	1/2
	= 58.11 min	1/2

21.	<ul> <li>a) In phenol, lone pair of electrons on oxygen are delocalized over benzene ring due to resonance but in alcohol lone pair of electrons on oxygen are localized and hence available for protonation.</li> <li>b) In anisole, O-C<sub>6</sub>H<sub>5</sub> bond is stronger than O-CH<sub>3</sub> bond as O-C<sub>6</sub>H<sub>5</sub> bond has partial</li> </ul>	1
	double bond character due to resonance.	1
22.	a) $OH$ $Na_2Cr_2O_7$ $H_2SO_4$	1
	b) $\rightarrow$	1
	OR	
	a) i) PCC (or any other suitable reagent)	1/2
	ii) Conc. HNO <sub>3</sub>	1/2
	b)	
	CH <sub>2</sub> —CH <sub>-</sub> CH <sub>3</sub>	1
23.	Due to osmosis.	1
	An increase in temperature would accelerate the process of osmosis.	1
24.	a) [Co(NH <sub>3</sub> ) <sub>4</sub> (H <sub>2</sub> O)Cl]Cl <sub>2</sub>	1
	b) Coordination number is 6	1/2
	Oxidation state of chromium is +3	1/2
	OR	

	en Cl Cl en Cl Cl en Cl Cl Cl Trans	1/2 + 1/2
	en 3+  en Co en	1
25.	a) 2,3-Dimethylbutane	1
	b) Isopropyl chloride < 1-Chloropropane < 1-Chlorobutane	1
26.	a) Order of reaction with respect to A is 1	1/2
	Order of reaction with respect to B is 0	1/2
	b) Rate law is, Rate = k[A]	1/2
	Overall order of reaction is 1	1/2
	c) <b>II</b>	1
27.	a) On addition of barium chloride, [Co(NH <sub>3</sub> ) <sub>5</sub> Cl]SO <sub>4</sub> forms white precipitate of	1
41.	BaSO <sub>4</sub> while [Co(NH <sub>3</sub> ) <sub>5</sub> (SO <sub>4</sub> )]Cl does not.	
	b) Diamminechloridonitrito-N-platinum(II)	1
	c) i) Strong field ligand: $t_2g^5 e_g^0$	
	ii) Weak field ligand: $t_2g^3 e_g^2$	1/2
	O.D.	72
	OR	
	a) Type of hybridization $- sp^3d^2$	1
	b) Magnetic property - Paramagnetic	1
	c) Type of complex – Outer orbital complex	1

28.	a) Lead storage battery is a secondary battery.	1/2
	Anode: $Pb(s) + SO_4^{2-}(aq) \rightarrow PbSO_4(s) + 2e^-$	1/2
	Cathode: $PbO_2(s) + SO_4^{2-}(aq) + 4H^+(aq) + 2e^- \rightarrow PbSO_4(s) + 2H_2O(l)$	1/2
	Overall reaction: $Pb(s)+PbO_2(s)+2H_2SO_4(aq) \rightarrow 2PbSO_4(s) + 2H_2O(l)$	1/2
	b) Ions are not involved in the overall cell reaction in the mercury cell.	1
29.	a) i) CH <sub>3</sub> Br	1
	ii) CH <sub>3</sub> CH <sub>2</sub> Cl	1
	b)	
	$CH_{3}-CH_{2}-CH = CH_{2}+ HBr \frac{Peroxide}{(Anti - Markovnitovs reaction)}$ $CH_{3}-CH_{2}-CH_{2}-CH_{2}-I \stackrel{NaI}{\leftarrow} CH_{3}-CH_{2}-CH_{2}-Br$ $1 - Iodobutane acetone 1 - Bromobutane$	1
30.		1
30.	a) Carbon-oxygen bond in phenol has a partial double bond character due to resonance.	1
	b) It is due to symmetry of para-isomers that fits in crystal lattice better as compared	1
	to ortho- and meta-isomers.	1
	c) Grignard reagents are highly reactive and react with any source of proton to give	1
	hydrocarbons.	
	d) Due to -I effect of halogen, ring is deactivated. (Any 3 correct answers)	1
31.	a) E and F	1
	b) A and B	1
	c) Exothermic reaction.	1
	The intermolecular attractive forces between C and D is stronger than those Between C-C and D-D. Therefore, energy is released.	1
	OR	
	c) $P_{\text{total}} = P_1^0 + (P_2^0 - P_1^0) X_2$	1/2
	$400 = 350 + (500-350) X_2$	1/2
	$X_{2} = 1/3 = 0.33$	1/2
	$X_1 = 2/3 = 0.67$	1/2
32.	a) Phosphodiester linkage.	1
	b) The two strands in DNA are complementary to each other because the hydrogen bonds are formed between specific pairs of bases.	1
	c) DNA - Adenine (A), guanine (G), cytosine (C) and thymine (T).	1
	RNA - Adenine (A), guanine (G), cytosine (C) and uracil (U).	1
	OR	

	c)			
		DNA	RNA	2
	It h	nas a double stranded α-helix structure	It has a single stranded structure	
		The sugar present is 2-deoxyribose	Sugar is ribose	
			(Any 2 points)	
33.	a)	$E_{cell} = (E_c^o - E_a^o) - \underline{0.059}$ log [	Mg <sup>2+</sup> ]	1/2
		E <sub>cell</sub> = [0.80 – (-2.37)] - <u>0.059</u>	$\log \frac{10^{-2}}{(10^{-4})^2}$	1/2
		Ecell = $3.17 - 0.0295 \log 10^6$	= 2.993 V	2
	b)	Cells that convert the energy of comb methane, methanol, etc.) directly into	oustion of fuels (like hydrogen, electrical energy are called fuel cells.	1
	c)	Due to overpotential/ Overvoltage of	$O_2$	1
		OR		
	a)	$\Lambda^{c}_{m} = \kappa \times 1000/M$		1/2
		$= 4.95 \times 10^{-5} \times 1000/$	0.001	1/2
		= $49.5 \text{ S cm}^2/\text{mol}$ $CH_3 \text{ COOH} \rightarrow CH_3\text{COO}^2$	. U <sup>+</sup>	1/2
		$\Lambda^0$ CH <sub>3</sub> COOH $\rightarrow$ CH <sub>3</sub> COO		
		= 40.	9 +349.6	
		$\Lambda^0$ CH <sub>3</sub> COOH = 390.5	S cm <sup>2</sup> /mol	1/2
		$\alpha = \frac{\Lambda_{\rm m}}{\alpha}$		17
		$\mathcal{L} = \mathcal{A}_{\mathrm{m}}^{0}$		1/2
		= 49.5/ 390.5		
		= 0.127		1/2
	b)	$E^{\circ}_{cell} = E^{\circ}_{(Ag+/Ag)} - E^{\circ}_{(Zn2+}$	-/Zn)	
		= 0.80 - (-0.76)		1/2
		= 1.56V		1/2
		$\Delta G^{\circ} = - nFE^{\circ}_{cell}$		1/2
		= - 2 × 96500 × 1.56 = - 301080 joules/mol		
		= -301.080 kJ/mol		1/2

34.	a)	Step 1: Protonation of alkene to form carbocation by electrophilic attack of $H_3O^+$ .	1
		$H_2O + H^+ \rightarrow H_3O^+$	
		Н н	
		$>C = C < + H - \ddot{O} + H \Longrightarrow - \ddot{C} - \dot{C} < + H_2 \ddot{O}$	
		Step 2: Nucleophilic attack of water on carbocation.	1
		$-\overset{H}{_{}{_{}{}{}}}-\overset{}{_{}{}$	
		Step 3: Deprotonation to form an alcohol.	1
		$-\overset{H}{_{}{_{}{}{}}}-\overset{}{}$	_
	b)	$A=$ $OH$ $COONa$ $OH$ $COOH$ $OCOCH_3$ $OH$ $OH$ $OH$ $OH$ $OCOCCH_3$	2
		OR	
	a)	Step 1: Formation of protonated alcohol.	1
		$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	
		H H H H H 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.	1
		$ \begin{array}{cccc} H & H & H & H & H \\ H - C - C & O - H & \longrightarrow & H - C - C + H_2O \\ H & H & H & H \end{array} $	
		Step 3: Formation of ethene by elimination of a proton.	1
		$H - \stackrel{H}{C} \stackrel{H}{\longrightarrow} \stackrel{H}{C} = \stackrel{H}{C} + \stackrel{H}{H}$	-
		Ethene	
	b) i)	$ \begin{array}{c} OCH_3 \\ \hline H_2SO_4 \\ HNO_3 \end{array} + \begin{array}{c} OCH_3 \\ \hline \end{array} $	1
		NO <sub>2</sub>	
	ii)	$R-X + R'- $ $\stackrel{\bullet}{\bigcirc}$ $Na \longrightarrow R- \stackrel{\bullet}{\bigcirc} -R' + Na X$ (or any specific example)	1

35.	a) i) Oxygen and fluorine have small size and high electronegativity. Hence, they	1
	can oxidize the metal to highest oxidation states.	
	ii) This is because small atoms like B, C, H, N etc. can occupy interstitial sites	1
	in the lattice of transition elements.	
ļ	iii) This is because 5f electrons in actinoids have poorer shielding effect than 4f	1
ļ	electrons in lanthanoids.	
	b) $5NO_2^- + 2MnO_4^- + 6H^+ \rightarrow 2Mn^{2+} + 5NO_3^- + 3H_2O$	1
	c) $2KMnO_4 \rightarrow K_2MnO_4 + MnO_2 + O_2$	1