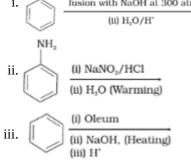
Class XII Session 2023-24 Subject - Chemistry Sample Question Paper - 1

Time Allowed: 3 hours			Maximum Marks: 70	
General Instructions:				
	Read the following instructions carefully.			
	1. There are 33 questions in this question paper with internal choice.			
	2. SECTION A consists of 16 multiple-choice questions carrying 1 mark each.			
	3. SECTION B consists of 5 very short answer questions carrying 2 marks each.			
	4. SECTION C consists of 7 short answer questions carrying 3 marks each.			
	5. SECTION D consists of 2 case-based questions carrying 4 marks each.			
	6. SECTION E consists of 3 long answer questions carrying 5 marks each.			
	7. All questions are compulsory.			
	8. Use of log tables and calculators is not allowed.			
	Section A			
1.	In alkyl halide:		[1]	
	a) All of these	b) the carbon atom of C-halogen bond	bears a	
		partial positive charge		
	c) the halogen atom bears a partial negative	d) the carbon-halogen bond of alkyl ha	lide is	
	charge	polarized		
2.	Maltose is made of:		[1]	
	a) \propto -D-glucose	b) \propto - D-glucose and β - D -glucose		
	c) Glucose and fructose	d) D-fructose		
3.	Which of the following reactions will yield phenol?		[1]	
	L fusion with NaOH at 300 atm			



	iv. (i) NaOH (aq.), 298k/1atm (ii) HCl			
	a) i, iii, iv		b) ii, iii, iv	
	c) i, ii, iii		d) i, ii, iv	
4.	The molecular formula of ethyl acetate	The molecular formula of ethyl acetate is:		[1]
	a) C ₄ H ₈ O		b) C ₄ H ₈ O ₂	
	c) C ₅ H ₁₀ O ₂		d) C ₅ H ₈ O ₂	
5.	Order of the photochemical reaction of	ccurring betwee	en hydrogen and chlorine is	[1]
	a) Second order		b) Third order	
	c) Zero order		d) First order	
6.	Match the column and choose correct	option		[1]
	Vant'Hoff factor	Behaviour of	f compound	
	(a) i = 1	(i) Impossible		
	(b) i > 1	(ii) Associatio	(ii) Association is the solution	
	(c) i < 1	(iii) Dissociation in the solution		
	(d) i = 0	(iv) No disso	(iv) No dissociation or association	
	a) (a) - (iv), (b) - (iii), (c) - (i), (d) -	· (ii)	b) (a) - (iv), (b) - (iii), (c) - (ii), (d) - (i)	
	c) (a) - (iv), (b) - (iv), (c) - (iii), (d)	- (ii)	d) (a) - (iii), (b) - (iv), (c) - (ii), (d) - (i)	
7.			[1]	
	a) Butane < 1-Iodobutane < 1-Bron 1-Chlorobutane	nobutane <	b) Butane < 1-Chlorobutane < 1-Iodobutane < 1-Bromobutane	
	c) Butane < 1-Chlorobutane < 1-Br < 1-Iodobutane	romobutane	d) 1-Iodobutane < 1-Bromobutane < 1- Chlorobutane < Butane	
8.	Red hot steel rods on suddenly immers	sing in water be	ecome:	[1]
	a) Soft and malleable		b) Hard and brittle	
	c) Tough and ductile		d) Fibrous	
9.	The value of decay constant of a comp	oound having a	half life period of 2.95 days is	[1]
	a) $3.0 imes 10^5 ext{s}^{-1}$		b) $2.71 \times 10^{-6} s^{-1}$	
	c) $2.9 \times 10^{-6} s^{-1}$		d) $2.9 \times 10^{6} \text{s}^{-1}$	
10.	Which of the following does not give s	silver mirror tes	st?	[1]

a) CH₃CH₂CHO b) HCOOH c) CH₃CHO d) CH₃COCH₃ One mole of an organic compound 'A' with the formula C₃H₈O reacts completely with two moles of HI to form [1]
 X and Y. When 'Y' is boiled with aqueous alkali forms Z. Z answers the iodoform test. The compound 'A' is

	,		
	a) methoxyethane	b) ethoxyethane	
	c) Propan – 2 – o1	d) Propan – 1 – o1	
12.	Hinsberg's reagent is:		[1]
	a) Benzene sulphonic acid	b) Benzene sulphonamide	
	c) Phenyl isocyanide	d) Benzene sulphonyl chloride	
13.	Assertion (A): Insulin is water-soluble. Reason (R): Insulin is a globular protein.		[1]
	a) Both A and R are true and R is the correct explanation of A.	b) Both A and R are true but R is not the correct explanation of A.	
	c) A is true but R is false.	d) A is false but R is true.	
14.	Assertion (A): Lower aldehydes and ketones are so masses increase.Reason (R): Distinction between aldehydes and ket	luble in water but the solubility decreases as the molecular cones can be made by Tollen's test.	[1]
	a) Both A and R are true and R is the correct explanation of A.	b) Both A and R are true but R is not the correct explanation of A.	
	c) A is true but R is false.	d) A is false but R is true.	
15.	Assertion (A): Isopropyl chloride is less reactive th	an CH_3Br in S_N^2 reactions.	[1]
	Reason (R): S_N^2 reactions are always accompanied	l by inversion of configuration.	
	a) Both A and R are true and R is the correct explanation of A.	b) Both A and R are true but R is not the correct explanation of A.	
	c) A is true but R is false.	d) A is false but R is true.	
16.	Assertion (A): \rightarrow is more acidic than	OH CH ₃	[1]
	1.02	ase acidity while electron-withdrawing groups (EWG)	
	a) Both A and R are true and R is the correct explanation of A.	b) Both A and R are true but R is not the correct explanation of A.	
	c) A is true but R is false.	d) A is false but R is true.	
	S	ection B	
17.	For the complex $[NiCl_4]^{2-}$, write,		[2]
	i. the IUPAC name ii. the hybridisation type		

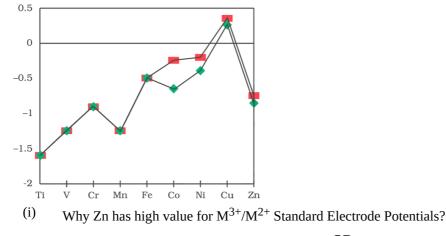
	iii. the shape of the complex (Atomic no. of Ni = 28)		
18.	Write the electronic configuration of the elements with the atomic numbers 61, 91, 101 and 109. [2]		
19.			
	(i) Consider the following equation:	[1]	
	$2NO(g)+2H_2(g) ightarrow N_2(g)+2H_2O(g)$		
	The rate law for this reaction is first order with respect to H_2 and second order with respect to NO.		
	Write the rate law for this reaction.		
	(ii) Oxygen is available in plenty in air yet fuels do not burn by themselves at room temperature. Explain.	[1]	
20.	Define the terms: Van't Hoff factor	[2]	
	OR		
	Under what condition do non-ideal solutions show negative deviations?		
21.	Arrange the following compounds in increasing order of their boiling points :	[2]	
	CH ₃ CHO, CH ₃ CH ₂ OH, CH ₃ OCH ₃ , CH ₃ CH ₂ CH ₃		
	Section C		
22.	What are fuel cells? Why we need these types of fuel cells? Explain with an example.	[3]	
23.	For a reaction: $\mathrm{H}_2 + \mathrm{Cl}_2 \stackrel{\mathrm{hv}}{\longrightarrow} 2\mathrm{HCl}$ Rate = k	[3]	
	i. Write the order and molecularity of this reaction.		
	ii. Write the unit of k.		
24.	An aromatic compound (A) having molecular formula $ m C_6H_6O$ on treatment with $ m CHCl_3$ and KOH gives a	[3]	
	mixture two isomers B and C both of B & C give same product D when distilled with Zn dust. Oxidation of D		
	gives E of formula C ₇ H ₆ O ₂ The sodium salt of E on heating with soda lime gives F which may also be obtained		
	by distilling A with zinc dust. Identify compounds A to F giving sequence of reactions.?		
	OR		
	Write the reactions and conditions for the following conversions:		
	i. 2-Propanone into 2-methyl-2-Proponal		
	ii. n-Propyl alcohol into hexane		
25.	Complete each synthesis by giving missing starting material, reagent or products.	[3]	
	+ H ₃ C ^{-C} CHO dilute NaOH		
26.	Calculate the maximum work and log Kc for the given reaction at 298 K :	[3]	
	Ni (s) + 2 Ag ⁺ (aq) \rightleftharpoons Ni ²⁺ (aq) + 2Ag (s)		
	Given:		
	$E_{Ni^{2+}/Ni}^{o}$ = -0.25 V, $E_{Ag^{+}/Ag}^{O}$ = +0.80 V		
	$1F = 96500 \text{ C mol}^{-1}$		
27.	What are haloarenes? How are they classified? Give one method each for the preparation of nuclear and side	[3]	
	chain substituted halorenes.		
28.	The E^0 values at 298 K corresponding to the following two reduction electrodes processes are:	[3]	
	i. $Cu^+/Cu = +0.52V$		
	ii. $Cu^{2+}/Cu^{+} = +0.16V$		

Formulae the galvanic cell for their combination. What will be the cell potential? Calculate the $\Delta_r G^0$ for the cell reaction. (1 F = 96500 C mol⁻¹)

Section D

29. Read the text carefully and answer the questions:

The unique behaviour of Cu, having a positive E° accounts for its inability to liberate H₂ from acids. Only oxidising acids (nitric and hot concentrated sulphuric) react with Cu, the acids being reduced. The stability of the half-filled d sub-shell in Mn²⁺ and the completely filled d¹⁰ configuration in Zn²⁺ are related to their E° values, whereas E° for Ni is related to the highest negative Δ_{hyd} H°. An examination of the $E^o_{(M^{3+}/M^{2+})}$ values the low value for Sc reflects the stability of Sc³⁺ which has a noble gas configuration. The comparatively high value for Mn shows that Mn²⁺(d⁵) is particularly stable, whereas a comparatively low value for Fe shows the extra stability of Fe³⁺ (d⁵). The comparatively low value for V is related to the stability of V²⁺ (half-filled t_{2g} level).



OR

Why Cu^{2+} is more stable than Cu^+ ?

- (ii) Transition metals, despite high E° oxidation, are poor reducing agents. Justify.
- (iii) Why is Cr^{2+} reducing and Mn^{3+} oxidising when both Cr and Mn have d⁴ configuration?

30. **Read the text carefully and answer the questions:**

In order to overcome the scarcity of drinking water in a remote village in Gujarat, Arnav and Aariv two young entrepreneurs still in their high school, have developed a unique water purifier that is capable of converting sea water into drinking water. It works on the principle of concentration difference between two solutions.

- (i) Name the phenomenon/process based on which this product is made?
- (ii) How difference in concentration of solutions help in converting sea water into drinking water?
- (iii) What arrangement they must have created in their product to covert sea water into drinking water?

OR

Equimolar solutions of NaCl and glucose are not isotonic. Why?

Section E

31.	Attem	pt any five of the following:	[5]
	(i)	Write the full forms of DNA and RNA.	[1]
	(ii)	How do enzymes help a substrate to be attacked by the reagent effectively?	[1]
	(iii)	Give two examples of reducing sugars.	[1]
	(iv)	What type of linkage holds together the monomers of DNA?	[1]
	(v)	Name the enzyme which convert surcrose into glucose and fructose.	[1]
	(vi)	What are polypeptides?	[1]

Page 5 of 14

[4]

(vii) Define the following term: Anomers.

32. i. Ratio of $[Ar(NH_3)_2]^+$ and $[Ar^+]$ in 0.1 M NH_3 solution.

ii. Ratio of $[Ag(S_2O_3)_2]^{3-}$ and [Ag+] in 0.1 M $S_2O_3^{2-}$ solution Given that the stability/ formation constant (k_f) for $[Ag(NH_3)_2]^+a$ and $[Ag(S_2O_3)_2]^{3-}$ are 1.7×10^7 and 1.0×10^{13} respectively.

OR

[1]

[5]

[5]

Give the oxidation state, d-orbital occupation and coordination number of the central metal ion in the following complexes:

i. $K_3 [Co(C_2O_4)_3]$ ii. cis $- [Cr(en)_2Cl_2]Cl$ iii. $(NH_4)_2 [CoF_4]$ iv. $[Mn(H_2O)_6]SO_4$

- 33. i. Write one chemical reaction for each
 - a. Carbylamine reaction
 - b. Acetylation reaction
 - ii. Write structure of N,N-ethylmethylethanamide

OR

- i. Write the structures of A and B in the following reactions:
 - a. $C_6H_5N_2^+Cl^- \xrightarrow{CuCN} A \xrightarrow{H_2O/H^+} B$ b. $CH_3COOH \xrightarrow{NH_3} A \xrightarrow{NaOBr} B$
- ii. Write the chemical reaction of methyl amine with benzoyl chloride and write the IUPAC name of the product obtained.
- iii. Arrange the following in the increasing order of their pK_b values: C₆H₅NH₂, NH₃, C₂H₅NH₂, (C₂H₅)₂ NH

Solution

Section A

1. (a) All of these

Explanation: Since halogen atoms are more electronegative than carbon, the carbon-halogen bond of alkyl halide is polarized; the carbon atom bears a partial positive charge whereas the halogen atom bears a partial negative charge.

2. (a) \propto -D-glucose

Explanation: A disaccharide, maltose is composed of two α -D-glucose units in which C₁ of one glucose (I) is linked to C₄ of another glucose unit (II).

3.

(c) i, ii, iii

Explanation:

- Preparation of phenols from haloarenes: Chlorobenzene is an example of haloarenes which is formed by
 monosubstitution of the benzene ring. When chlorobenzene is fused with sodium hydroxide at 623K and 320 atm
 sodium phenoxide is produced. Finally, sodium phenoxide on acidification gives phenols.
- Preparation of phenols from diazonium salts: When an aromatic primary amine is treated with nitrous (NaNO₂ + HCl) acid at 273 278 K, diazonium salts are obtained. These diazonium salts are highly reactive in nature. Upon warming with water, these diazonium salts finally hydrolyse to phenols. Phenols can also be obtained from diazonium salts by treating it with dilute acids.
- Preparation of phenols from benzene sulphonic acid: Benzenesulphonic acid can be obtained from benzene by
 reacting it with oleum. Benzenesulphonic acid thus formed is treated with molten sodium hydroxide at high temperature
 which leads to the formation of sodium phenoxide. Finally, sodium phenoxide on acidification gives phenols.

4.

(b) C₄H₈O₂

Explanation: Its molecular formula is C₄H₈O₂ and its chemical formula is CH₃COOC₂H₅.

5.

(c) Zero order

Explanation: light-dependent Photochemical reaction between H_2 and $Cl_2 [H_2(g) + Cl_2(g) \rightarrow 2HCl(g)]$ is 0 (zero) w.r.t. to reactants.

6.

(b) (a) - (iv), (b) - (iii), (c) - (ii), (d) - (i) **Explanation:** (a) - (iv), (b) - (iii), (c) - (ii), (d) - (i)

7.

(c) Butane < 1-Chlorobutane < 1-Bromobutane < 1-Iodobutane

Explanation: Due to the polar nature of alkyl halides and the increase in molecular weight compared to their parent alkanes, the boiling points of alkyl halides are higher than that of their parent alkanes. The boiling points of alkyl halides depend on the molecular mass and the size of the halogen atom (decrease from I to F). With the increase in size, mass, and the number of electrons in halogen atoms, the magnitude of Van Der Waals forces increase and the boiling point also increases. The boiling point of alkyl halides reduces in the order RI > REr > RCl > RF.

Therefore, the order of increasing order of boiling points should be Butane < 1-Chlorobutane < 1-Iodobutane < 1-Iodobutane.

8.

(b) Hard and brittle

Explanation: Red hot steel rods on suddenly immersing in water become hard and brittle.

9.

(b) $2.71 \times 10^{-6} \text{s}^{-1}$

Explanation: Decay constant(k) = $\frac{0.693}{t_{1/2}}$

$$k = \; rac{0.693}{2.95 imes 24 imes 60 imes 60} \, = 2.71 imes 10^{-6} \; s^{-1}$$

10.

(d) CH₃COCH₃

Explanation: CH₃COCH₃ will not give a silver mirror test (Tollens Test). Tollens test is given by aldehydes only and HCOOH is the only acid that gives tollen's test. Ketones do not give tollen's test.

11. (a) methoxyethane

Explanation: Ether reacts with HI to form alcohol and alkyl iodide. Alcohol on oxidation will give the iodoform test.

12.

(d) Benzene sulphonyl chloride

Explanation: Benzene sulphonyl chloride, $C_6H_5SO_2Cl_2$, is called Hinsberg reagent. It is used to distinguish between primary, secondary and tertiary amines.

13.

(b) Both A and R are true but R is not the correct explanation of A. **Explanation:** Insulin is a globular protein. This protein has a three-dimensional folded structure. These are stabilised by internal hydrogen bonding, hence, they are water-soluble.

14.

(b) Both A and R are true but R is not the correct explanation of A.

Explanation: The assertion is true that the aldehydes have an oxygen atom that can form a hydrogen bond with the hydrogen atoms of water. So lower aldehydes are soluble in water. But as the chain length increases, the hydrophobic part (the alkyl chain) dominates over the hydrophilic part (the polar -CHO group). So higher molecular weight aldehydes are insoluble in water but lower molecular weight aldehydes are easily soluble in water.

The reason is also correct because aldehydes form a silver mirror with Tollen reagent but ketones do not.

15.

(b) Both A and R are true but R is not the correct explanation of A.

Explanation: As the size of the alkyl groups increases, the S_N^2 reactivity decreases, further C - Cl bond is stronger and more difficult to cleave than C - Br bond. So CH_3Br is more reactive than $(CH_3)_2CHCl$.

16.

(c) A is true but R is false.Explanation: A is true but R is false.

Section B

17. i. The IUPAC name of [NiCl₄]²⁻ is tetra chloridonickelate (II) ion.

ii. sp³-hybridisation.

- iii. Due to sp³-hybridisation, its shape is tetrahedral.
- 18. Z = 61 has electronic configuration as $[Xe]^{54} 4f^5 5d^0 6s^2$
 - Z = 91 has electronic configuration as $[Rn]^{86}$ 5f² 6d¹7s²
 - Z = 101 has electronic configuration as $[Rn]^{86} 5f^{13} 6d^0 7s^2$
 - Z = 109 has electronic configuration as $[Rn]^{86} 5f^{14} 6d^7 7s^2$

19. Answer the following:

- (i) Rate = k $[NO]^2 [H_2]^1$
- (ii) The activation energy for combustion reactions of fuels is very high at room temperature, therefore, they do not burn by themselves at room temperature.
- 20. Van't Hoff factor: Van't hoff factor 'i' is a correction factor defined as the ratio between the actual concentration of particles produced when the substance is dissolved and the concentration of a substance as calculated from its mass.

 $= \frac{\text{Observed value of colligative property}}{\text{Normal value of colligative property}}$

OR

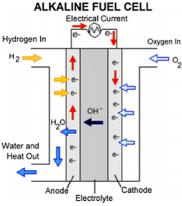
When the new forces of attraction between components are greater than those in the pure components. That is when two components A and B are mixed, the interactions between A....B is greater than A...A and B...B interaction then the binary non-ideal solution would show negative deviation from Raoult's law.

21. Boiling point depends upon the size of the molecules, branching in the molecules and intermolecular forces of attraction between the molecules.

On the basis of this, the increasing order of boiling point are: CH₃CH₂CH₃ < CH₃OCH₃ < CH₃CHO < CH₃CH₂OH

Section C

22. Galvanic cells (or Voltaic cells) that are designed to convert the energy of combustion of fuels like hydrogen, methane, methanol, etc. directly into electrical energy are called fuel cells. In these cells, the reactants are continuously supplied to electrodes. One of the most successful fuel cells uses the reaction of hydrogen with oxygen to form water as shown in the figure.



The cell was used for providing electrical power in the Apollo space programme. The water vapours produced during the reaction were condensed and added to the drinking water supply for the astronauts. In the cell, hydrogen and oxygen are bubbled through porous carbon electrodes into concentrated aqueous sodium hydroxide solution. Catalysts like finely divided platinum or palladium metal are incorporated into the electrodes for increasing the rate of electrode reactions.

The electrode reactions are given below:

 $Cathode: O_2(g) + 2H_2O(l) + 4e^- \rightarrow 4OH^-(aq)$

Anode : $2H_2(g) + 4OH^-(aq) \rightarrow 4H_2O(l)$

Overall reaction being:

 $2\mathrm{H}_2(g) + \mathrm{O}_2(g) \to 2\mathrm{H}_2(g)$

The cell runs continuously as long as the reactants are supplied. Fuel cells produce electricity with an efficiency of about 70% compared to thermal plants whose efficiency is about 40%. There has been tremendous progress in the development of new electrode materials, better catalysts and electrolytes for increasing the efficiency of fuel cells. These have been used in automobiles on an experimental basis. Fuel cells are pollution free and in view of their future importance, a variety of fuel cells have been fabricated and tried.

23. i. For a reaction, $H_2 + Cl_2 \xrightarrow{hv} 2HCl$

Rate= k, suggests that the reaction is of zero order. Further, the molecularity of a given reaction is 2 as two molecules are participating in the reaction.

Hence, order = zero and molecularity = two.

ii. The unit of k for zero order reaction is equal to the rate of a reaction which is mol $L^{-1}s^{-1}$.

Hence, the unit of k for the given reaction is mol $L^{-1}s^{-1}$.

A (
$$C_6H_6O$$
)
CHCl₃ B + C
24.
CHCl₃ B + C
Zn
D (0) E ($C_7H_6O_2$) NaOH + CaO
Sodium salt of E F $\stackrel{Zn}{\checkmark}$ A

The aromatic compound having molecular formula C₆H₆O and which gives a mixture of two isomers on reacting with CHCl₃ and

KOH is phenol i.e. The reactions are (A) Therefore OH сно снсі_з + кон ОН O сно \bigcirc (B) 0 (A) сно (A) (B) (C) ОН сно Zn Zn (C) (O) сно соон (D) (0) 0 сно (D) (E) он соон COONa NaOH / CaO Zn (F) [O] õ (E) 0 C decarboxylation (A) OR $\text{i. } CH_3 - \underbrace{C}_{\substack{\parallel\\ 2-propanone}} - CH_3 + CH_3MgBr \xrightarrow{dry \ ether}_{addition} \begin{bmatrix} CH_3 \\ \downarrow \\ CH_3 - \underbrace{C}_{l} \\ \downarrow \\ OMgBr \end{bmatrix} \xrightarrow{H_2O/H^+} CH_3 - \underbrace{C}_{l} \\ \downarrow \\ OH \\ OH \end{bmatrix}$ 2-methyl-2-propanol

ii. n-Propyl alcohol to hexane

$$\begin{array}{c} \mathrm{CH_3CH_2CH_2OH}+\mathrm{PCl_5}\rightarrow\mathrm{CH_3CH_2CH_2Cl}+\mathrm{POCl_3}+\mathrm{HCl}\\ \\ 2CH_3CH_2CH_2Cl+2Na \xrightarrow{Dry} CH_3CH_2CH_2C_2CH_3+2NaCl\\ 1-Chloropropane \xrightarrow{Dry} either \xrightarrow{n-Hexane} \end{array}$$

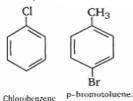
25. The reaction between an aldehyde/ketone and an aromatic carbonyl compound lacking an α -hydrogen is called Claisen–Schmidt condensation.

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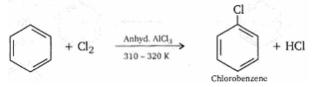
$$\begin{array}{c} \begin{array}{c} \mathsf{CH0} \\ \begin{array}{c} \mathsf{H}_{3} \\ \\ \mathsf{CH0} \\ \end{array} \\ \begin{array}{c} \mathsf{H}_{3} \\ \\ \mathsf{H}_{3} \\ \\ \mathsf{C}_{4} \\ \\ \mathsf{Ch0} \\ \end{array} \\ \begin{array}{c} \mathsf{H}_{4} \\ \\ \mathsf{C}_{4} \\ \\ \mathsf{C}_{4}$$

27. **Haloarenes:** The replacement of hydrogen atoms in a aromatic hydrocarbon by halogen atoms results in the formation of aryl halide (haloarene). Haloarenes contain halogen atoms attached to sp² hybridized carbon atoms of an aryl group. They are classified as:

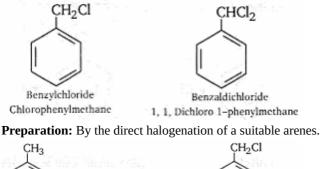
i. **Nuclear halogen derivatives:** Halogen derivatives of aromatic hydrocarbons in which the halogen atom (F, Cl, Br, or I) is directly attached to an aromatic ring are called nuclear halogen derivatives. Some examples are:

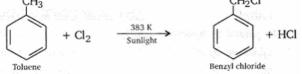


It is prepared by the direct chlorination of aromatic hydrocarbon.



ii. **Side chain halogen derivatives:** Halogen derivatives of aromatic hydrocarbons in which the halogen atom is linked to one of the carbon atoms of the side chain carrying the aryl group are called side chain halogen derivatives. For example,





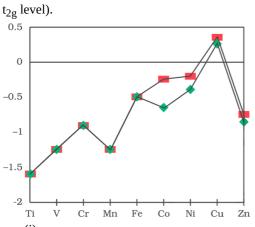
28. cell representation is

$$\begin{split} & \text{Cu}^{+}|\text{Cu}^{2+}||\text{Cu}^{+}|\text{Cu}\\ & E_{cell}^{0}=E_{reduction(cathode)}^{0}-E_{oxidation(anode)}^{0}\\ & =\text{E}^{0}(\text{Cu}^{+}/\text{Cu})-\text{E}^{0}(\text{Cu}^{2+}/\text{Cu}^{+})\\ & =+0.52\text{V}-0.16\text{V}\\ & =0.36\text{ V}\\ & Cu^{+}\rightarrow Cu^{2+}+e^{-}\\ & Cu^{+}\rightarrow e^{-}\rightarrow Cu\\ & \text{n}=1\\ & \Delta_{r}G^{0}=-nE^{0}F\\ & =-1\times0.36V\times96500\\ & =-34740\text{ Jmol}^{-1}\\ & =-34.74\text{ KJ mol}^{-1} \end{split}$$

Section D

29. Read the text carefully and answer the questions:

The unique behaviour of Cu, having a positive E° accounts for its inability to liberate H₂ from acids. Only oxidising acids (nitric and hot concentrated sulphuric) react with Cu, the acids being reduced. The stability of the half-filled d sub-shell in Mn²⁺ and the completely filled d¹⁰ configuration in Zn²⁺ are related to their E° values, whereas E° for Ni is related to the highest negative Δ_{hyd} H°. An examination of the $E^o_{(M^{3+}/M^{2+})}$ values the low value for Sc reflects the stability of Sc³⁺ which has a noble gas configuration. The comparatively high value for Mn shows that Mn²⁺(d⁵) is particularly stable, whereas a comparatively low value for Fe shows the extra stability of Fe³⁺ (d⁵). The comparatively low value for V is related to the stability of V²⁺ (half-filled



(i) Due to the removal of an electron from the stable d^{10} configuration of Zn^{2+} .

OR

The Stability of Cu^{2+} is more than Cu^+ as stability depends on the hydration energy of the ions when they bond to the water molecules. The Cu^{2+} ion has a greater charge density than Cu^+ ion and thus forms much stronger bonds releasing more energy.

- (ii) Transition metals despite having high E° oxidation, are poor reducing agents because of their high heat of vaporization, high ionisation energies and low heats of hydration.
- (iii) Cr^{2+} is reducing as its configuration changes from d₄ to d₃, the having a half-filled t_{2g} level. On the other hand, the

change from Mn^{3+} to Mn^{2+} results in the half-filled (d₅) configuration which has extra stability.

30. Read the text carefully and answer the questions:

In order to overcome the scarcity of drinking water in a remote village in Gujarat, Arnav and Aariv two young entrepreneurs still in their high school, have developed a unique water purifier that is capable of converting sea water into drinking water. It works on the principle of concentration difference between two solutions.

- (i) The product is based on the phenomenon of Reverse Osmosis between solutions of two different concentration.
- (ii) When solutions of two different concentration are separated by a semipermeable membrane and excess pressure is applied on the solution of higher concentration, solvent flow from higher concentration to lower concentration. This is called Reverse Osmosis and same can be used to treat seawater and convert into drinking water.

(iii)Following arrangement must have been made:

- i. Use of a semipermeable membrane.
- ii. Separate Compartment having seawater and Drinking Water separated by semipermeable membrane.
- iii. Excess pressure applied in compartment having sea water.

OR

NaCl is an electrolyte and gets dissociated to two ions (Na⁺ and Cl⁻) and exerts almost double osmotic pressure than glucose which is a non-electrolyte and does not dissociate.

Section E

- 31. Attempt any five of the following:
 - (i) DNA \longrightarrow Deoxyribonucleic acid
 - $\mathsf{RNA} \longrightarrow \mathsf{Ribonucleic} \ \mathsf{acid}$
 - (ii) Active site of enzymes holds the substrate molecule in a suitable position so that the enzyme help substrate can be attacked by the reagent effectively.
 - (iii)Examples of reducing sugars Maltose and Lactose.
 - (iv)The 2 strands of DNA are attached to each other by hydrogen bonds that connect the nitrogenous bases of one strand to the bases of the other strand (this is called complementary base pairing) where Adenine pairs with Thymine by 2 hydrogen bonds and Guanine pairs with Cytosine using 3 hydrogen bonds.

While nucleotides of DNA are held together by Phosphodiester linkages. The structure of DNA holds the nucleotides in place using phosphodiester bonds. A phosphodiester bond occurs when exactly two of the hydroxyl group in phosphoric acid react with hydroxyl groups on other molecules to form two ester bonds. The phosphodiester bond is the linkage between the 3' carbon atom of one sugar molecule and the 5 ' carbon atom of another, deoxyribose in DNA.

(v) Invertase

(vi)Polypeptides are formed when several molecules generally more than ten of α -amino acids are joined together by a peptide bond.

(vii) somers which differ only in the configuration of hydroxyl group at C_1 are called anomers. e.g. α and β form of glucose.

32. i. The equilibrium is:

 $egin{aligned} &Ag^+ + 2NH_3[Ag(NH_3)_2]^+\ arphi & \mathrm{Stability\ constant}, \ &\mathrm{K}_f = rac{[\mathrm{Ag}(\mathrm{NH}_3)_2]^+}{[\mathrm{Ag}^+][\mathrm{NH}_3]^2}\ &= 1.7 imes10^7\ (\mathrm{Given})\ &\mathrm{or}\ rac{[Ag(NH_3)_2]^+}{[Ag^+]}\ &= 1.7 imes10^7 imes[NH_3]^2\ &= 1.7 imes10^7 imes[NH_3]^2\ &= 1.7 imes10^7 imes(0.1)^2\ &= 1.7 imes10^5 \end{aligned}$

ii. The equilibrium is

$$egin{aligned} &\mathrm{Ag}^+ + 2\mathrm{S}_2\mathrm{O}_3^{2^-} \stackrel{\mathrm{k}_f}{\rightleftharpoons} \left[\mathrm{Ag}(\mathrm{S}_2\mathrm{O}_3)_2
ight]^{3^-} \ &\therefore \ &\mathrm{Stability\ constant}, \ &\therefore \ & rac{\left[Ag(S_2O_3)_2
ight]^{3^-}}{\left[Ag^+
ight]} = 1.0 imes 10^{13} imes [S_2O_3^{2^-}]^2 = 1.0 imes 10^{13} (0.1)^2 = 1 imes 10^{11} \ &\therefore \ & rac{\left[Ag(S_2O_3)_2
ight]^{3^-}}{\left[Ag^+
ight]} = 1.0 imes 10^{13} imes [S_2O_3^{2^-}]^2 = 1.0 imes 10^{13} (0.1)^2 = 1 imes 10^{11} \end{aligned}$$

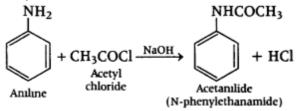
i. $K_3 [Co(C_2O_4)_3]$ The central metal ion is Co. Its coordination number is 6. The oxidation state can be given as: x - 6 = -3x = +3The d orbital occupation for $Co^{3+}is$, $t_{2g}{}^6e_g{}^0$ ii. cis- $[Cr(en)_2 Cl_2] Cl$ The central metal ion is Cr. The coordination number is 6. The oxidation state can be given as: x + 2(0) + 2(-1) = +1x - 2 = +1x = +3 The d orbital occupation for Cr^{3+} is t_{2q}^{3} . iii. $(NH_4)_2 [CoF_4]$ The central metal ion is Co. The coordination number is 4. The oxidation state can be given as: x - 4 = -2x = + 2 The d orbital occupation for $Co^{2+}is, e_g{}^4t_{2g}{}^3.$ iv. $[Mn(H_2O)_6] SO_4$ The central metal ion is Mn. The coordination number is 6. The oxidation state can be given as: x + 0 = +2x = +2The d orbital occupation for Mn is $t_{2g}{}^3e_g{}^2.$

33. i. **Carbylamine reaction:** Aliphatic or aromatic primary amines on heating with chloroform and ethanolic potassium hydroxide form isocyanides or carbylamine which are foul smelling substances. This reaction is known as carbylamine reaction. $C_6H_5NH_2$ Δ

$$\begin{array}{c} \overset{\sim}{\rightarrow} C_{6}H_{5}NC + 3KCl + 3H_{2}O \\ \text{Aniline} \end{array}$$

This reaction is used as a test for primary aliphatic and aromatic amine.

ii. Acetylation:

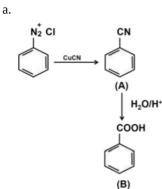


iii. The structure of N,N-ethylmethylethanamide is:

$$C_2H_5- egin{array}{ccc} N & -C & -CH_3 \ & & \parallel \ & CH_3 & O \end{array}$$

OR

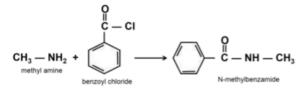
i. The structure of A and B on following reaction is:



b.

$$\begin{array}{ccc} CH_{3}COOH \xrightarrow{NH_{3}}{\Delta} & CH_{3} \xrightarrow{O}{\overset{\parallel}{\leftarrow}} & NH_{2} + H_{2}O \\ & (A) & & \\ & & \downarrow & NaOBr \\ & & CH_{3} - NH_{2} + CO_{2} \\ & (B) & (1^{\circ} \text{ amine}) \end{array}$$

ii. The chemical reaction of methyl amine with benzoyl chloride and IUPAC name of the product form is as follows:



iii. Increasing order of pKb values

 $(C_2H_5)_2 \, \overset{\,\,{}_\circ}{N}\, H < C_2H_5NH_2 < \overset{\,\,{}_\circ}{N}\, H_3 < C_6H_5 \, \overset{\,\,{}_\circ}{N}\, H_2$