# **Class XII Session 2024-25** Subject - Chemistry Sample Question Paper - 4

Time Al	lowed: 3 hours	Maximu	m Marks: 70
General	Instructions:		
	Read the following instructions carefully.		
	1. There are <b>33</b> 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 ques	stions 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 c	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	l.	
	S	ection A	
1.	Chloromethane on treatment with excess of ammonia yields mainly		[1]
	a) Methanamine (CH <sub>3</sub> NH <sub>2</sub> )	b) Mixture containing all these in equal proportion	
	c) N–methylmethanamine (CH <sub>3</sub> —NH—CH <sub>3</sub> )	d) N, N-Dimethylmethanamine	
		$(CH_3 - N < CH_3)$	
2.	Which of the following is a disaccharide?		[1]
	a) Glucose	b) Cellulose	
	c) Lactose	d) Starch	
3.	In the reaction		[1]
	OH ONa		
	$H \to H_2O$		
	Sodium phenoxide	b) They can denote a proton to a stronger base	
	a) Phenois are actuic in nature.	b) They can donate a proton to a stronger base	
4	C) Cleavage of U - H Donu	(1) All of these	[1]
4.			[1]
	a) on its reactivity with other functional	b) size and solubility of the aldehyde and	
	Broups.	actoric morecure.	

5.	Value of Henry's constant K <sub>H</sub> :		[1]
	a) increases with decrease in temperature.	b) remains constant.	
	c) increases with increase in temperature.	d) decreases with increase in temperature.	
6.	Match the item given in Column I with expression g	iven in Column II.	[1]
	Column I	Column II	
	(a) Osmotic Pressure	(i) p = K <sub>H</sub> . $\chi_{\rm B}$	
	(b) Relative lowering of vapour pressure	(ii) $rac{ riangle P}{ extsf{P}^o_A} = \chi_ extsf{B}$	
	(c) Henry Law	(iii) $\triangle T_b = K_b.m$	
	(d) Elevation in boiling point	(iv) p = iCRT	
	a) (a) - (ii), (b) - (iii), (c) - (iv), (d) - (i).	b) (a) - (iv), (b) - (iii), (c) - (ii), (d) - (i).	
	c) (a) - (iii), (b) - (i), (c) - (ii), (d) - (iv).	d) (a) - (iv), (b) - (ii), (c) - (i), (d) - (iii).	
7.	Isomers of the compound $C_4H_9Br$ are:		[1]
	a) 1-Bromo-2-methylbutane and 2-Bromo-2- methylbutane	b) 1 – Bromobutane and 2 – Bromobutane	
	<ul> <li>c) 1 – Bromobutane, 2 – Bromobutane, 1 –</li> <li>bromo- 2 – methylpropane, and 2 – bromo-2</li> <li>– methylpropane</li> </ul>	d) 1 – Bromo - 2 – methylpropane and 2 – Bromo - 2 – methylpropane	
8.	Which one among the following metals of 3d series	has the lowest melting point?	[1]
	a) Cu	b) Fe	
	c) Zn	d) Mn	
9.	The half-life of a reaction is halved as the initial con reaction is:	centration of the reactant is doubled. The order of the	[1]
	a) 1	b) 0	
	c) 2	d) 3	
10.	Which of the following does not give silver mirror to	est?	[1]
	a) CH <sub>3</sub> CH <sub>2</sub> CHO	b) HCOOH	
	c) CH <sub>3</sub> CHO	d) CH <sub>3</sub> COCH <sub>3</sub>	
11.	When diethyl ether is heated with excess of HI, it pr	oduces :	[1]
	a) ethyl iodide	b) ethanol	
	c) methyl iodide	d) iodoform	
12.	Gabriel synthesis is used for the preparation of:		[1]
	a) Quaternary salt	b) Primary amines	
	c) Tertiary amine	d) Secondary amine	

d) moisture of the air.

c) only solubility of aldehydes and ketones.

13.	<b>Assertion (A):</b> Glucose and fructose are reducing su <b>Reason (R):</b> Glucose and fructose contain a free ald respectively.	igars. lehydic and ketonic group adjacent to a >CHOH group	[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): Rate of addition of HCN on carbony	l compounds increases in presence of NaCN.	[1]
	<b>Reason (R):</b> Reaction involves the addition of Cn <sup>-</sup> in rate determining step.		
	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 <sub>3</sub> Br in S <sub>N</sub> 2 reactions.	[1]
	<b>Reason (R):</b> $S_N^2$ reactions are always accompanied	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): Aqueous solution of phenol is called	l as carbonic acid.	[1]
	<b>Reason (R):</b> Increasing order of acidity of phenols i	s m nitrophenol >o-nitrophenol >p-nitrophenol.	
	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	
	c) $\Lambda$ is true but $P$ is false	d) Both A and B are false	
	C) A is the but K is faise.	ection B	
17.	Write the formulae for the following coordination co	ompounds:	[2]
	i. Potassium tetrahydroxo sincate (II)	-	
	ii. Potassium trioxalatoaluminate (III)		
	iii. Dichloridobis cobalt III (ethane - 1, 2 diamine)		
18.	Why are $Fe^{2+}$ compounds easily oxidises to $Fe^{3+}$ as	compared to Mn <sup>+2</sup> compounds.	[2]
19.	Answer the following:		[2]
	<ul><li>(a) a. Write the mathematical relation betwee</li><li>b. What is collision frequency?</li></ul>	n rate constant and half-life of a first order reaction.	[1]
	(b) If the rate equation is given below:		[1]
	Rate = $k[A]^2[B]$		
	then what will be the unit of its rate and rat	e constant?	
20.	Calculate the temperature at which the solution cont	aining 54 g of glucose, $C_6 H_{12} O_6$ in 250 g of water will	[2]
	Treeze. ( $K_b$ for water = 1.86 K kg mol <sup>-1</sup> )	OR	
	Define the terms Mass percentage.	OIX	
21.	An organic compound 'A' (molecular formula $C_3H_6$	$_{3}O$ ) is resistant to oxidation but forms a compound 'B' (	[2]

 $C_3H_8O$ ) on reduction. 'B' reacts with HBr to form a bromide'C' which on treatment with alcoholic KOH forms an alkene 'D' ( $C_3H_6$ ). Deduce the structures of A, B, C and D.

#### Section C

22. What are fuel cells? Explain the electrode reactions involved in the working of H<sub>2</sub> - O<sub>2</sub> fuel cell.

23. In a reaction between A and B, the initial rate of reaction was measured for different initial concentrations of A [3] and B as given below:

$A/molL^{-1}$	0.20	0.20	0.40
$B/molL^{-1}$	0.30	0.10	0.05
$r_0/molL^{-1}s^{-1}$	$5.07 imes10^{-5}$	$5.07 imes10^{-5}$	$1.43 imes10^{-4}$

What is the order of the reaction with respect to A and B?

i. Write the mechanism of the following reaction: 24.

$$2CH_{3}CH_{2}OH \xrightarrow[413]{H^{+}} CH_{3}CH_{2}OCH_{2}CH_{3} + H_{2}O$$

ii. Write the preparation of phenol from cumene.

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. How would you bring about the following conversions?

i. Propanal to butanone

ii. Benzaldehyde to benzophenone

iii. Benzoyl chloride to benzonitrile

26. Calculate the emf of the following cell:

$$Mg(s)|Mg^{2+}(0.2M)||Ag^{+}(1 \times 10^{-3}M)|Ag(s)||Mg(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Ag(s)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^{2+}(0.2M)||Mg^$$

$$E^{0}(Ag^{+}/Ag) = 0.80V$$

27. Draw the structures of major monohalo products in each of the following reactions:

i. 
$$OH$$
  
+ SOCl<sub>2</sub>  $\rightarrow$   
ii.  $O_2N$   $CH_2CH_3$   
 $H_2$ , heat or  
UV light  
+ HCl heat

28. State Kohlrausch's law of independent migration of ions. How can the degree of dissociation of acetic acid in a [3] solution be calculated from its molar conductivity data?

### Section D

#### Read the following text carefully and answer the questions that follow: 29. [4]

Observe the graph of transition metal and their melting points

[3]

[3]

[3]

[3]



- i. Why does W (tungsten) has highest melting point? (1)
- ii. Which element in 3d series has lowest enthalpy of atomisation and why? (1)
- iii. Why is mercury liquid? (2)

## OR

Why are transition metals less electropositive than 's'-block elements? (2)

#### 30. Read the following text carefully and answer the questions that follow:

A raw mango placed in concentrated salt solution loses water via osmosis and shrivel into pickle. Wilted flowers revive when placed in fresh water. A carrot that has become limp because of water loss into the atmosphere can be placed into the water making it firm once again. Water will move into its cells through osmosis. When placed in water containing less than 0.9% (mass/volume) salt, blood cells swell due to flow of water in them by osmosis.

[4]

- i. People taking a lot of salt or salty food suffer from puffiness or edema. What is the reason behind this?
- ii. The preservation of meat by salting and of fruits by adding sugar protects against bacterial action. How?
- iii. Why the direction of osmosis gets reversed if a pressure larger than the osmotic pressure is applied to the solution side? Write its one application.

#### OR

What care is generally taken during intravenous injections and why?

#### Section E

31.	Attem	pt any five of the following:	[5]
	(a)	Define native state in reference to proteins.	[1]
	(b)	Define the following terms:	[1]
		a. Polysaccharides	
		b. Nucleotides	
	(c)	Deficiency of which vitamin causes scurvy?	[1]
	(d)	What happens when D-glucose is treated with the following? Give equations to support your answer.	[1]
		a. HI	
		b. HNO <sub>3</sub>	
	(e)	Name the disaccharide which on hydrolysis gives glucose and galactose.	[1]
	(f)	Write the products obtained after hydrolysis of lactose.	[1]
	(g)	Give the reaction of glucose with hydrogen cyanide. Presence of which group is confirmed by this	[1]
		reaction?	
32.	Draw a	Ill the isomers of: (geometrical and optical)	[5]

- i. [CoCl<sub>2</sub>(en)<sub>2</sub>]<sup>+</sup>
- ii. [Co(NH<sub>3</sub>)Cl(en)<sup>2</sup>]<sup>2+</sup>
- iii. [Co(NH<sub>3</sub>)<sub>2</sub>Cl<sub>2</sub>(en)]<sup>+</sup>

OR

Using crystal field theory, draw energy level diagram, write the electronic configuration of the central metal atom/ion and determine the magnetic moment value in the following:

i. [CoF<sub>6</sub>]<sup>3-</sup>, [Co(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup>, [Co(CN)<sub>6</sub>]<sup>3-</sup>

ii. [FeF<sub>6</sub>]<sup>3-</sup>, [Fe(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup>, [Fe(CN)<sub>6</sub>]<sup>4-</sup>

33. a. Give plausible explanation for each of the following:

i. Why are amines less acidic than alcohols of comparable molecular masses?

ii. Why are primary amines highest boiling than tertiary amines?

iii. Why are aliphatic amines stronger bases than aromatic amines?

b. Complete the following reactions:

- i.  $C_6H_5N_2Cl + C_2H_5OH \rightarrow$
- ii. C<sub>6</sub>H<sub>5</sub>NH<sub>2</sub> + (CH<sub>3</sub>CO)<sub>2</sub>O  $\rightarrow$

#### OR

Write the structure of A, B, C, D and E in the following reactions:

 $C_6H_5NO_2 \xrightarrow{Sn/HCl} A \xrightarrow{(CH_3CO)_2O} B \xrightarrow{HNO_3 + H_2SO_4} C \xrightarrow{OH^- \text{ or } H^+} D$  $H_2SO_4$ 

## Solution

#### Section A

1. **(a)** Methanamine (CH<sub>3</sub>NH<sub>2</sub>)

## **Explanation:** $CH_3Cl + NH_3 \rightarrow CH_3NH_2 + HCl$

Ammonia molecule is a nucleophile in nature as it has unpaired electrons. This nucleophile attacks the chloromethane  $CH_3Cl$  molecule and forms methylamine or methenamine by a nucleophilic substitution reaction mechanism. The carbon atom is partially positive in the molecule, due to the electronegativity of the halide attached which is partially negative. The electron-rich nucleophile attacks the positive ion, causing the halide ion to be separated from the molecule.

#### 2.

(c) Lactose

Explanation: Lactose

#### 3.

### (d) All of these

**Explanation:** Phenol reacts with sodium hydroxide solution to give a colourless solution containing sodium phenoxide. In this reaction, the hydrogen ion has been removed by the strongly basic hydroxide ion in the sodium hydroxide solution.

#### 4.

(b) size and solubility of the aldehyde and ketone molecule.

**Explanation:** Size and solubility of aldehyde and ketone determine fragrance. For example, aldehyde C-10 is used in floral blends like rose, jasmine, etc. While aldehyde C-11 has a strong citrus smell and aldehyde C-16 has a strong strawberry smell.

#### 5.

(c) increases with increase in temperature.

Explanation: Value of henry constant increases with increase in temperature.

#### 6.

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

#### 7.

(c) 1 – Bromobutane, 2 – Bromobutane, 1 – bromo- 2 – methylpropane, and 2 – bromo-2 – methylpropane

**Explanation:** Isomers are compounds having the same molecular formula but different structures. In the given straight-chain 4-C compounds, Br is placed at different positions but the molecular formula is same.

 $CH_3CH_2CH(Br)CH_3$ ,  $CH_3CH_2CH_2CH_2Br_{2-Bromobu \tan e}$  1-Bromobu  $\tan e$ 

In the given branched 4-C compounds, there is a branching of -CH<sub>3</sub> at carbon 2 but Br is placed at positions 1 and 2

#### respectively.

 $CH_3CH(CH_3)CH_2Br,\ CH_3C(Br)(CH_3)CH_3$ 

 $1-Bromo-2-methyl propane \qquad 2-Bromo-2-methyl propane$ 

So, all of these are isomers since they have the same molecular formula but different structures.

## 8.

**(c)** Zn

Explanation: Zn metal of 3d series has the lowest melting point.

## 9.

**(c)** 2

**Explanation:** For 2<sup>nd</sup> order reaction, the half-life is inversely related to the concentration of the reactant.  $t_{1/2}$  for second order reaction  $\alpha \frac{1}{|D|}$ 

## 10.

(d) CH<sub>3</sub>COCH<sub>3</sub>

**Explanation:** CH<sub>3</sub>COCH<sub>3</sub> 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) ethyl iodide

Explanation: ethyl iodide

## 12.

(b) Primary amines

**Explanation:** In Gabriel Phthalimide reaction, the sodium or potassium salt of phthalimide is N-alkylated with a primary alkyl halide to give the corresponding *N*-alkylphthalimide for producing primary amines. This is because of the reaction of sodium or potassium salt of phthalimide with alkyl halide impure SN<sub>2</sub> reaction.

- 13. (a) Both A and R are true and R is the correct explanation of A.
   Explanation: Reducing sugars contain a free aldehydic or ketonic groupadjacent to a ><sub>CHOH</sub> group and reduce Tollen's reagent, Schiff's reagent or Benedict's solution.
- 14. **(a)** Both A and R are true and R is the correct explanation of A.

**Explanation:** The addition of HCN to carbonyl compounds involves the addition of CN<sup>-</sup> in rate determining step which are supplied easily by NaCN and thus addition becomes fast.

## 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.

(d) Both A and R are false.

## **Explanation:**

Aqueous solution of phenol is called carbolic acid. Electron withdrawing groups such as -NO<sub>2</sub>, -CN, -X- increase the acidity. The increase is more at o - and p - positions than at m-position.

i.e. p-nitrophenol >o-nitrophenol >m-nitrophenol.

## Section B

- 17. i. K<sub>2</sub>[Zn(OH)<sub>4</sub>]
  - ii. K<sub>3</sub>[Al(C<sub>2</sub>O<sub>4</sub>)<sub>3</sub>]
  - iii.  $[CoCl_2(en)_2]^+$
- 18. Mn<sup>2+</sup> compounds are less readily converted to its +3 oxidation state (Mn<sup>3+</sup>) as compared to Fe<sup>2+</sup> because Mn<sup>2+</sup> (3d<sup>5</sup>) is more stable than Mn<sup>3+</sup> (3d<sup>4</sup>) due to stable half filled d-orbital electronic configuration. Whereas Fe<sup>3+</sup> (3d<sup>5</sup>) is more stable than Fe<sup>2+</sup> (3d<sup>6</sup>) due to half filled d-orbitals therefore Fe<sup>2+</sup> gets easily converted to Fe<sup>3+</sup>

## 19. Answer the following:

(i) a. 
$$t_{1/2} = \frac{0.693}{k}$$

b. The number of collisions per second per unit volume of the reaction mixture.

(ii) Unit of rate = mol  $L^{-1}s^{-1}$ 

Unit of rate constant (k) = 
$$\frac{\text{Unit of rate}}{\text{Unit of } [A^2] \times \text{Unit of } [B]}$$

$${{{\left( {{mol}{L^{ - 1}}} 
ight)}^2}{\left( {{mol}{L^{ - 1}}} 
ight)}^2}}{{mol}^{ - 2}}L^2{s^{ - 1}}}$$

20. Molecular mass of glucose

$$egin{aligned} M_B &= 72 + 12 + 96 = 180\,g\,mol^{-1} \ \Delta T_f &= rac{K_f imes w_B imes 1000}{M_B imes w_A} \ &= rac{1.86 imes 54 imes 1000}{180 imes 250} = 2.23\,\,\,\mathrm{K} \end{aligned}$$

Freezing point of solution =  $T_{f}^{0}$  -  $\triangle Tf$  = 273 - 2.23 = 270.77*K* 

**Mass percentage:** The mass percentage of a component in a given solution is defined as the mass of the component per 100g of the solution.

 $Mass~\%~of~a~component = \frac{Mass~of~the~component~in~the~solution}{Total~mass~of~the~solution} \times 100$ 

21. Structure of A, B, C and D are deduced in the following manner.

$$\begin{array}{c} CH_3 - \underset{(A)}{CO} - CH_3 \xrightarrow[\text{Reduction}]{LrAH_4} CH_3CH(OH)CH_3\\ \hline CH_3CH(OH)CH_3 + HBr \rightarrow CH_3CH(Br)CH_3\\ \hline (B) & (C) \\ CH_3CH(Br)CH_3 + KOH(alc.) \rightarrow CH_3 - CH = CH_2 + KBr + H_2O\\ \hline (C) & (D) \end{array}$$
Section C

22. **Fuel cells :** Those galvanic cells in which chemical energy of combustion of fuels like hydrogen, methane, etc. is converted into electrical energy are called fuel cells.

H<sub>2</sub>-O<sub>2</sub> fuel cell: The cell consists of three compartments separated by a porous electrode. Hydrogen gas is introduced into one

compartment and oxygen into another compartment. These gases then diffuse slowly through the electrode and react with an electrolyte that is in the central part of the cell. The electrodes are made of porous carbon and electrolyte is a resin containing concentrated aqueous sodium hydroxide solution. Hydrogen is oxidized at anode and oxygen is reduced at the cathode.



The electrode reactions involved in the working of the H<sub>2</sub>-O<sub>2</sub> fuel cell are as:

At cathode,

 $O_2(g) + 2H_2O(l) + 4e^- \longrightarrow 4OH^-(aq)$ At anode,  $2H_2(g) + 4OH^-(aq) \longrightarrow 4H_2O(l) + 4e^-$ Overall cell reaction:  $2H_2(g) + O_2(g) \longrightarrow 2H_2O(l)$ 

23. Consider the order of the reaction with respect to A is x and with respect to B is y.

Therefore,  $r_0 = k[A]^x [B]^y$  $5.07 \times 10^{-5} = k[0.20]^x [0.30]^y$  ..... (i)  $5.07 \times 10^{-5} = k[0.20]^x [0.10]^y$  ..... (ii)  $1.43 \times 10^{-4} = k[0.40]^x [0.05]^y$  ..... (iii) Dividing equation (i) by (ii), we obtain  $k[0.20]^x[0.30]^y$  $5.07{ imes}10^{-5}$  $\frac{10^{-5}}{5.07 \times 10^{-5}} =$  $k[0.20]^x[0.10]^y$  $\frac{\left[0.30\right]^{y}}{\left[0.10\right]^{y}} \left(\frac{0.30}{0.10}\right)^{y}$  $\mathbf{v} = \left(\frac{0.30}{0.10}\right)^y$ 1 =y = 0Dividing equation (iii) by (ii), we obtain  $k[0.40]^x[0.05]^y$  $1.43{ imes}10^{-4}$ = $5.07{\times}\overline{10^{-5}}$  $k[0.20]^x[0.30]^y$  $\frac{[0.40]^y}{[0.20]^y}$  $\underline{1.43 \times 10^{-4}}$  $[Since y = 0, [0.05]^y = [0.30]^y = 1]$  $\overline{5.07 \times 10^{-5}}$  $2.821 = 2^x$  $\log 2.821 = x \log 2 \ (Taking \ \log \ on \ both \ sides) x = rac{\log 2.821}{\log 2}$ = 1.496

= 1.5 (approximately)

Hence, the order of the reaction with respect to A is 1.5 and with respect to B is 0.



ii. To prepare phenol, cumene is first oxidized in the presence of air of cumene hydro-peroxide.



Then, cumene hydroxide is treated with dilute acid to prepare phenol and acetone as by-products.

$$CH_{3} \longrightarrow C \longrightarrow O \longrightarrow H \xrightarrow{H^{+}} OH \xrightarrow{OH} H^{+} \xrightarrow{H^{+}} H^{+} \xrightarrow{OH} H^{+} \xrightarrow{H^{+}} H^{+} H^{+} \xrightarrow{H^{+}} H^{+} H^{+} \xrightarrow{H^{+}} H^{+} \xrightarrow{H^{+}} H^{+} H^{+}$$

$$\text{i. } CH_3 - \underbrace{C}_{\substack{||\\ 2-propanone}} - CH_3 + CH_3MgBr \xrightarrow{dry \ ether}_{addition} \begin{bmatrix} CH_3 \\ | \\ CH_3 - \underbrace{C}_{l} - CH_3 \\ | \\ OMgBr \end{bmatrix} \xrightarrow{H_2O/H^+} CH_3 - \underbrace{CH_3 \\ | \\ CH_3 - CH_3 \\ | \\ OH \\ 2-methyl-2-propanole \end{bmatrix} \xrightarrow{H_2O/H^+}_{addition} CH_3 = \underbrace{CH_3 \\ | \\ CH_3 - CH_3 \\ | \\ OH \\ 2-methyl-2-propanole \end{bmatrix} \xrightarrow{H_2O/H^+}_{addition} CH_3 = \underbrace{CH_3 \\ | \\ CH_3 - CH_3 \\ | \\ OH \\ 2-methyl-2-propanole \end{bmatrix} \xrightarrow{H_2O/H^+}_{addition} CH_3 = \underbrace{CH_3 \\ | \\ CH_3 - CH_3 \\ | \\ OH \\ 2-methyl-2-propanole \end{bmatrix} \xrightarrow{H_2O/H^+}_{addition} CH_3 = \underbrace{CH_3 \\ | \\ CH_3 - CH_3 \\ | \\ CH_3 - CH_$$

ii. n-Propyl alcohol to hexane

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

25. The following steps are involved in the conversions:

i. Propanal to butanone

$$\begin{array}{c} \begin{array}{c} O\\ H\\ CH_{3}-CH_{2}-C-H+CH_{3}MgBr \longrightarrow CH_{3}-CH_{2}-CH-CH_{3} & \underbrace{H_{2}O/H^{+}}_{I} \\ \end{array} \\ O\\ CH_{3}-CH_{2}-C-CH_{3} & \underbrace{Cu}_{573 \ K} & CH_{3}-CH_{2}-CH-CH_{3} \\ \end{array} \\ \begin{array}{c} O\\ H\\ H\\ Butanone \end{array}$$

 $P_{-} O_{-}$ 

ii. Benzaldehyde to Benzophenone



iii. Benzoyl Chloride to Benzonitrile

$$C_6H_5COCl \xrightarrow{H_{13}} C_6H_5CONH_2 \xrightarrow{H_{205}} C_6H_5C \equiv N$$
  
Benzoyl chloride Heat Benzamide heat Benzonitrile

26. 
$$\frac{Mg(s) \longrightarrow Mg^{2^+}(aq) + 2e^-}{Mg(s) + 2Ag^+(aq) + 2e^- \longrightarrow 2 Ag(s)}$$
$$\frac{Mg(s) + 2Ag^+(aq) \longrightarrow Mg^{2^+}(aq) + 2Ag(s)}{Mg^{2^+}(aq) + 2Ag(s)}$$

applying nernst equation

$$egin{aligned} &E_{cell} = E_{cell}^0 - rac{0.0591}{2} \log rac{[Mg^{2+}]}{[Ag^{2+}]} \ &E^0(Ag^+/Ag) - E^0(Mg^{2+}/Mg) - rac{0.0591}{2} \log rac{0.2}{(10^{-3})^2} \ &= +0.80V - (-2.37V) - rac{0.0591}{2} \log (2 imes 10^5) \ &= +3.17V - rac{0.0591}{2} [\log 2 + \log 10^5] \ &= +3.17V - rac{0.0591}{2} imes 5.3010 \ &= +3.17V - 0.1566V \ &= 3.0134 \ \mathrm{V} \end{aligned}$$

27. Major monohalo product are as follows:



28. **Kohlrausch law of independent migration of ions:** It states that limiting molar conductivity of an electrolyte can be represented as the sum of the individual contributions of the anion and cation of the electrolyte. If  $\lambda^o Na^+$  and  $\lambda^o Cl^-$  are limiting molar conductivity for sodium chloride is given by

 $\lambda^0_m\left(NaCl
ight)=\lambda^oNa+\lambda^0Cl^-$ 

Calculation of degree of dissociation of weak electrolyte like acetic acid. The degree of dissociation  $\alpha$  is given by:  $\alpha = \frac{\lambda_m}{\lambda_m^0}$ 

where 
$$\lambda_m$$
 be molar conductivity and  $\lambda_m^0$  be the limiting molar conductivity.

#### Section D

- 29. i. It is due to presence of most number of unpaired electrons and there is more frequent metal-metal bonding in 5d series than 3d and 4d series.
  - ii. Zinc has lowest enthalpy of atomisation due to weak metallic bond which is due to absence of unpaired electrons.
  - iii. It is due to larger size, absence of unpaired electron and weak interatomic attraction and weaker metallic bond. **OR**

It is due to smaller atomic size and higher ionisation enthalpies.

- 30. i. People experience water retention in tissue cells and intercellular spaces due to osmosis.
  - ii. Through the process of osmosis, a bacterium on salted meat or candid fruit loses water, shrivels and dies.
  - iii. The pure solvent flows out of the solution through the semi permeable membrane due to reverse osmosis. It is used in desalination of sea water.

OR

During intravenous injection, the concentration of the solution should be same as that of blood so that they are isotonic. Because if the solution concentration is hypertonic than blood cell will shrink and if it is hypotonic than blood cell will swells/burst.

#### Section E

- 31. Attempt any five of the following:
  - (i) Native state of protein is the sequence in which the amino acids are linked together with the help of peptide bond.
  - (ii) a. Polysaccharides contain a large number of monosaccharide units joined together by glycosidic linkages.
    b. Sugar + Phosphate + base / Nucleoside linked to a phosphate group.

(iii)Vitamin C

(iv) a. n-hexane is formed

$$\begin{array}{c} CHO \\ (CHOH)_4 & \xrightarrow{HI, \ \bigtriangleup} & CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3 \\ \downarrow \\ CH_2OH \\ \end{array}$$
Saccharic acid is formed  

$$\begin{array}{c} CHO \\ CHO \\ CHO \\ \end{array}$$

b.

$$( \overset{|}{C} HOH)_4 \xrightarrow{Oxidation} ( \overset{|}{C} HOH)_4 \xrightarrow{Oxidation} ( \overset{|}{C} HOH)_4 \xrightarrow{Oxidation} ( \overset{|}{C} HOH)_4$$

(v) Lactose

(C)

(vi)Hydrolysis of Lactose gives D-galactose and D-glucose.



(vii)<sub>CHO</sub> CN CH< OH HCN (CHOH), (CHOH), -> ĊH₂OH CH<sub>2</sub>OH Confirms the presence of aldehydic/carbonyl group.

32. i. [CoCl<sub>2</sub>(en)<sub>2</sub>]<sup>+</sup>



Trans [CoCl2(en)2] tisomer-optically inactive (Superimposable mirror images)



Mirror

Cis [CoCl2(en)2]<sup>+</sup>isomer-optically active (Non-superimposable mirror images)



Trans [CoCl<sub>2</sub>(en)<sub>2</sub>]<sup>+</sup> isomer-optically inactive

(Superimposable mirror images)



Cis  $[CoCl_2(en)_2]^+$  isomer-optically active

(Non-superimposable mirror images)

In total, three isomers are possible. ii. [Co(NH<sub>3</sub>)Cl(en)<sup>2</sup>]<sup>2+</sup>

Trans-isomers are optically inactive. Cis-isomers are optically active.

iii. [Co(NH<sub>3</sub>)<sub>2</sub>Cl<sub>2</sub>(en)]<sup>+</sup>



Synergic bonding

OR



 $Co^{2+} = 3d^7$ 

Number of unpaired electrons = 4

Magnetic moment =  $\sqrt{n(n+2)} = \sqrt{4(4+2)} = 4.9$  B.M

[Co(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup>:



 $Co^{2+} = 3d^7$ Number of unpaired electrons = 3 Magnetic moment =  $\sqrt{3(3+2)}$  = 3.87 B.M

[Co(CN)<sub>6</sub>]<sup>3-</sup> :



 $Co^{3+} = 3d^6$ Number of unpaired electrons = 0 Diamagnetic in nature.

ii.  $FeF_6^{3-}$ :



 $Fe^{2+} = 3d^6$ 

 $[Fe(H_2O)_6]^{2+}: t^4_{2g} e_g^2$ Number of unpaired electrons = 4 Magnetic moment =  $\sqrt{4(4+2)}$  = 4.9 B.M [Fe(CN)\_6]^4- :



 $Fe^{2+} = 3d^6$ 

Diamagnetic in nature.

33. a. i. Loss of proton from amines give ion whereas loss of a proton from alcohol gives an alkoxide ion.

Since O is more electronegative than N, therefore, RO<sup>-</sup> can accommodate the -ve charge more easily than RNH<sup>-</sup>.

Consequently, RO<sup>-</sup> is more stable than RNH<sup>-</sup>. Thus, alcohols are more acidic than amines.

- ii. Primary amines (RNH<sub>2</sub>) have two hydrogen atoms on the N atom and therefore, form intermolecular hydrogen bonding.
   Tertiary amines (R<sub>3</sub>N) donot have hydrogen atoms on the N atom and therefore, these donot form hydrogen bonds. As a result of hydrogen bonding in primary amines, they have higher boiling points than tertiary amines of comparable molecular mass.
- iii. Both arylamines and alkylamines are basic in nature due to the presence of lone pair on N-atom. But arylamines are less basic than alkylamines.
- b. i.  $C_6H_5N_2Cl + C_2H_5OH \xrightarrow{Reduction} C_6H_6 + CH_3CHO + N_2 + HCl$   $\xrightarrow{Benzene} Ethanal$ ii.  $C_6H_5NH_2 + (CH_3CO)_2O \xrightarrow{CH_3COOH} C_6H_5CONHCH_3 + CH_3COOH$  $Ace \tan ilide$



