

Board Paper of Class 12-Science 2023 Chemistry Delhi(Set 1)

Total Time: 180

Total Marks: 70.0

Section A

Q.No.1: Which of the following molecules has a chiral centre correctly labelled with an asterisk (*)?

- (a) CH₃C*HBrCH₃
- (b) CH₃C*HC/CH₂Br
- (c) HOCH₂C*H(OH)CH₂OH
- (d) CH₃C*Br₂CH₃

Q.No.2: Which of the following alcohols will not undergo oxidation?

- (a) Butanol
- (b) Butan-2-ol
- (c) 2-Methylbutan-2-ol
- (d) 3-Methylbutan-2-ol

Q.No.3: A voltaic cell is made by connecting two half cells represented by half equations below:

$$\mathrm{Sn}^{2+}{}_{\mathrm{(aq)}} + 2\mathrm{e}^{-}
ightarrow \mathrm{Sn}_{\mathrm{(s)}} \,\, \mathrm{E}^{\,\circ} = -0.14 \,\, \mathrm{V}$$

$${
m Fe}^{3+}{}_{
m (aq)} + {
m e}^{-}
ightarrow {
m Fe}^{2+}{}_{
m (aq)} {
m E}^{\,\circ} = +0.77 {
m V}$$

Which statement is correct about this voltaic cell?

- (a) Fe^{2+} is oxidised and the voltage of the cell is -0.91 V
- (b) Sn is oxidised and the voltage of the cell is 0.91 V
- (c) Fe^{2+} is oxidised and the voltage of the cell is 0.91 V
- (d) Sn is oxidised and the voltage of the cell is 0.63 V Marks:[1.00]

Marks:[1.00]

Marks:[1.00]

Q.No.4: Four half reactions I to IV are shown below:

I.
$$2Cl^- o Cl_2+2e^-$$

II.
$$4\,\mathrm{OH}^-
ightarrow \mathrm{O}_2 + 2\mathrm{H}_2\mathrm{O} + 2\mathrm{e}^-$$

III.
$$Na^+ + e^- \rightarrow Na$$

IV.
$$2H^+ + 2e^- \rightarrow H_2$$

Which two of these reactions are most likely to occur when concentrated brine is electrolysed?

- (a) I and III
- (b) I and IV
- (c) II and III
- (d) II and IV Marks:[1.00]

Q.No.5: Which property of transition metals enables them to behave as catalysts?

- (a) High melting point
- (b) High ionisation enthalpy
- (c) Alloy formation
- (d) Variable oxidation states

Marks:[1.00]

Q.No.6: In the two tetrahedral structures of dichromate

- (a) 4 Cr O bonds are equivalent in length.
- (b) 6 Cr O bonds are equivalent in length.
- (c) All Cr O bonds are equivalent in length.
- (d) All Cr O bonds are non-equivalent.

Marks:[1.00]

Q.No.7: 1 mole of liquid A and 2 moles of liquid B make a solution having a total vapour pressure 40 torr. The vapour pressure of pure A and pure B are 45 torr and 30 torr respectively. The above solution (a) is an ideal solution.

- (b) shows positive deviation.
- (c) shows negative deviation.
- (d) is a maximum boiling azeotrope.

Marks:[1.00]

Q.No.8: Which of the following would not be a good choice for reducing nitrobenzene to aniline? (a) LiA/H₄ (b) H_2/Ni (c) Fe and HC/ (d) Sn and HCI Marks:[1.00] Q.No.9: If molality of a dilute solution is doubled, the value of the molal elevation constant (K_b) will be (a) halved (b) doubled (c) tripled Marks:[1.00] (d) unchanged Q.No.10: Hydrolysis of sucrose is called (a) inversion (b) hydration (c) esterification (d) saponification Marks:[1.00] **Q.No.11:** Which one of the following has lowest pK_a value? СН, - СООН (a) $O_2N - CH_2 - COOH$ (b) (c) (d) HCOOH Marks:[1.00] **Q.No.12:** Which of the following cell was used in Apollo space programme? (a) Mercury cell (b) Daniel cell $\mathrm{H_2}\mathbf{-}\mathrm{O_2}$ Fuel cell (d) Dry cell Marks:[1.00]

Q.No.13: The following experimental rate data were obtained for a reaction carried out at 25°C:

 $A_{(g)} + B_{(g)} + C_{(g)} + D_{(g)}$

(-)	X=7	
Initial [A _(g)]/mo/ dm ⁻³	Initial $[B_{(g)}]/mol dm^{-3}$	Initial rate/mo/ dm ⁻³ s ⁻¹
3.0×10^{-2}	2.0×10^{-2}	1.89×10^{-4}
3.0×10^{-2}	4.0×10^{-2}	1.89×10^{-4}
6.0×10^{-2}	4.0×10^{-2}	7.56×10^{-4}

What are the orders with respect to $A_{(q)}$ and $B_{(q)}$?

	(9)				
	Order with respect to $A_{(g)}$	Order with respect to $B_{(g)}$			
(a)	Zero	Second			
(b)	First	Zero			
(c)	Second	Zero			
(d)	Second	First			

Marks:[1.00]

Q.No.14: The magnetic moment of $[NiCl_4]^{2-}$

- (a) 1.82 BM
- (b) 2.82 BM
- (c) 4.42 BM
- (d) 5.46 BM

[Atomic number: Ni = 28]

Marks:[1.00]

Q.No.15: Assertion (A): Proteins are polymers of α-amino acids connected by a peptide bond.

Reason (R): A tetrapeptide contains 4 amino acids linked by 4 peptide bonds.

- (a) Both Assertion (A) and Reason (R) are true and Reason (R) is the correct explanation of the Assertion (A).
- (b) Both Assertion (A) and Reason (R) are true, but Reason (R) is not the correct explanation of the Assertion (A).
- (c) Assertion (A) is true, but Reason (R) is false.
- (d) Assertion (A) is false, but Reason (R) is true.

Marks:[1.00]

Q.No.16: Assertion (A): For a zero order reaction the unit of rate constant and rate of reaction are same.

Reason (R): Rate of reaction for zero order reaction is independent of concentration of reactant.

- (a) Both Assertion (A) and Reason (R) are true and Reason (R) is the correct explanation of the Assertion (A).
- (b) Both Assertion (A) and Reason (R) are true, but Reason (R) is not the correct explanation of the Assertion (A).
- (c) Assertion (A) is true, but Reason (R) is false.

Marks:[1.00]

Q.No.17: Assertion (A): Acetic acid but not formic acid can be halogenated in presence of red P and Cl_2 .

Reason (R): Acetic acid is a weaker acid than formic acid.

- (a) Both Assertion (A) and Reason (R) are true and Reason (R) is the correct explanation of the Assertion (A).
- (b) Both Assertion (A) and Reason (R) are true, but Reason (R) is not the correct explanation of the Assertion (A).
- (c) Assertion (A) is true, but Reason (R) is false.
- (d) Assertion (A) is false, but Reason (R) is true.

Marks:[1.00]

Q.No.18: Assertion (A): Trans $\left[\mathrm{CrCl}_2\left(\mathrm{ox}\right)_2\right]^{3-}$ shows optical isomerism.

Reason (R): Optical isomerism is common in octahedral complexes involving didentate ligands.

- (a) Both Assertion (A) and Reason (R) are true and Reason (R) is the correct explanation of the Assertion (A).
- (b) Both Assertion (A) and Reason (R) are true, but Reason (R) is not the correct explanation of the Assertion (A).
- (c) Assertion (A) is true, but Reason (R) is false.
- (d) Assertion (A) is false, but Reason (R) is true.

Marks:[1.00]

Section B

Q.No.19: (i) What should be the signs (positive/negative) for $E_{\rm cell}^{\rm o}$ and ΔG° for a spontaneous redox reaction occurring under standard conditions? (ii) State Faraday's first law of electrolysis.

Calculate the emf of the following cell at 298 K:

$$\operatorname{Fe}_{(\mathrm{s})} \mid \operatorname{Fe}^{2+} \left(0.01\mathrm{M}\right) \mid \mid \operatorname{H}^{+}_{(1\mathrm{M})} \mid \operatorname{H}_{2(\mathrm{g})} \left(1 \text{ bar}\right), \operatorname{Pt}_{(\mathrm{s})}$$
Given $\operatorname{E}^{\mathrm{o}} = 0.44 \, \mathrm{V}$

Given $E_{cell}^{o} = 0.44 \text{ V}.$

Marks:[2.00]

Q.No.20: What happens to the rate constant k and activation energy E_a as the temperature of a chemical reaction is increased? Justify. Marks:[2.00]

Q.No.21: (a) Which of the following species cannot act as a ligand? Give

reason.

 $OH^{-}, NH_{4}^{+}, CH_{3}NH_{2}, H_{2}O$

- (b) The complex $\left[\mathrm{Co}\left(\mathrm{NH_3}\right)_5\left(\mathrm{NO_2}\right)\right]\mathrm{Cl_2}$ is red in colour. Give IUPAC name of its linkage isomer. **Marks:[2.00]**
- **Q.No.22:** Why is boiling point of o-dichlorobenzene higher than p-dichlorobenzene but melting point of para isomer is higher than ortho isomer?

 Marks:[2.00]

Q.No.23: For the pair phenol and cyclohexanol, answer the following:

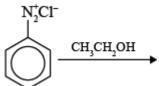
- (a) Why is phenol more acidic than cyclohexanol?
- (b) Give one chemical test to distinguish between the two. Marks:[2.00]

Q.No.24: (i) Draw the zwitter ion structure for sulphanilic acid.

(ii) How can the activating effect of $-NH_2$ group in aniline be controlled?

OR

(i) Complete the reaction with the main product formed:



(ii) Convert Bromoethane to Propanamine.

Marks:[2.00]

Q.No.25: Give the reaction of glucose with hydrogen cyanide. Presence of which group is confirmed by this reaction?

Marks:[2.00]

Section C

Q.No.26: (a) For the reaction $2N_2O_{5(q)} \rightarrow 4NO_{2(q)} + O_{2(q)}$ at 318 K

calculate the rate of reaction if rate of disappearance of $N_2O_{5(g)}$ is 1.4×10^{-3} m s⁻¹.

(b) For a first order reaction derive the relationship $t_{99\%} = 2t_{90\%}$ Marks:[3.00]

Q.No.27: (a) On the basis of crystal field theory write the electronic configuration for d^5 ion with a strong field ligand for which $\Delta_0 > P$.

(b) $[Ni(CO)_4]$ has tetrahedral geometry while $[Ni(CN)_4]^{2-}$ has square planar yet both exhibit dimagnetism. Explain.

[Atomic number: Ni = 28] **Marks:[3.00]**

Q.No.28: (a) Illustrate Sandmeyer's reaction with an equation.

(b) Explain, why $(CH_3)_2NH$ is more basic than $(CH_3)_3N$ in aqueous solution.

Marks:[3.00]

Q.No.29: Give reasons for any 3 of the following observations:

- (a) Penta-acetate of glucose does not react with hydroxylamine.
- (b) Amino acids behave like salts.
- (c) Water soluble vitamins must be taken regularly in diet.
- (d) The two strands in DNA are complimentary to each other. Marks:[3.00]

Q.No.30: (i) Why is the C–O bond length in phenols less than that in methanol?

(ii) Arrange the following in order of increasing boiling point:

Ethoxyethane, Butanal, Butanol, n-butane

(iii) How can phenol be prepared from anisole? Give reaction.

OR

(i) Give mechanism of the following reaction:

$$CH_3CH_2OH \xrightarrow{H_2SO_4} CH_3CH_2 - O - CH_2CH_3 + H_2O$$

(ii) Illustrate hydroboration-oxidation reaction with an example. Marks:[3.00]

Section D

Q.No.31: The following question is case - based question. Read the passage carefully and answer the question that follow:

Nucleophilic Substitution

Nucleophilic Substitution reaction of haloalkane can be conducted according to both S_NI and S_N2 mechanisms. S_N1 is a two step reaction while S_N2 is a single step reaction. For any haloalkane which mechanism is followed depends on factors such as structure of haloalkane, properties of leaving group, nucleophilic reagent and solvent.

Influences of solvent polarity: In S_NI reaction, the polarity of the system

increases from the reactant to the transition state, because a polar solvent has a greater effect on the transition state than the reactant, thereby reducing activation energy and accelerating the reaction. In S_N2 reaction, the polarity of the system generally does not change from the reactant to the transition state and only charge dispersion occurs. At this time, polar solvent has a great stabilizing effect on Nu than the transition state, thereby increasing activation energy and slow down the reaction rate. For example, the decomposition rate (S_N1) of tertiary chlorobutane at 25°C in water (dielectric constant 79) is 300000 times faster than in ethanol (dielectric constant 24). The reaction rate (S_N2) of 2-Bromopropane and NaOH in ethanol containing 40% water is twice slower than in absolute ethanol. Hence the level of solvent polarity has influence on both S_NI and S_N2 reaction, but with different results. Generally speaking weak polar solvent is favourable for S_N2 reaction, while strong polar solvent is favourable for S_NI. Generally speaking the substitution reaction of tertiary haloalkane is based on S_NI mechanism in solvents with a strong polarity (for example ethanol containing water).

Answer the following questions

- (a) Why racemisation occurs in S_NI ?
- (b) Why is ethanol less polar than water?
- (c) Which one of the following in each pair is more reactive towards S_N2 reaction ?

(i)
$$CH_3 - CH_2 - I$$
 or $CH_3CH_2 - Cl$
(ii) $CH_3 - CH_2 - Cl$

- (c) Arrange the following in the increasing order of their reactivity towards $\mathsf{S}_N\mathsf{I}$ reactions :
 - (i) 2-Bromo-2-methylbutane, 1-Bromopentane, 2-Bromopentane
 - (ii) 1-Bromo-3-methylbutane, 2-Bromo-2-methylbutane, 2-Bromo-3-methylbutane

Marks:[4.00]

Q.No.32: Rahul set-up an experiment to find resistance of aqueous KCl solution for different concentrations at 298 K using a conductivity cell connected to a Wheatstone bridge. He fed the Wheatstone bridge with a.c. power in the audio frequency range 550 to 5000 cycles per second. Once the resistance was calculated from null point, he also calculated the conductivity, k and molar conductivity k and recorded his readings in tabular form.

S. No.	Conc. (M)	k S cm ⁻¹	∧ _m S cm ² mol ⁻¹
1.	1.00	111.3×10^{-3}	111.3
2.	0.10	12.9×10^{-3}	129.0
3.	0.01	1.41×10^{-3}	141.0

Answer the following questions:

- (a) Why does conductivity decrease with dilution?
- (b) If \wedge_m^o of KC/ is 150.0 S cm 2 mol $^{-1}$, calculate the degree of dissociation of 0.01 M KCl.
- (c) If Rahul had used HCl instead to KCl then would you expect the \land_m values to be more or less than those per KCl for a given concentration. Justify.

OR

(c) Amit, a classmate of Rahul repeated the same experiment with CH₃COOH solution instead of KCl solution. Give one point that would be similar and one that would be different in his observations as compared to Rahul. **Marks:[4.00]**

Q.No.33: (i) Why is boiling point of 1M NaCl solution more than that of 1M glucose solution?

- (ii) A non-volatile solute 'X' (molar mass = 50 g mol^{-1}) when dissolved in 78g of benzene reduced its vapour pressure to 90%. Calculate the mass of X dissolved in the solution.
- (iii) Calculate the boiling point elevation for a solution prepared by adding 10g of $MgCl_2$ to 200g of water assuming $MgCl_2$ is completely dissociated.

 $(K_b \text{ for Water} = 0.512 \text{ K kg mol}^{-1}, \text{ Molar mass MgCl}_2, = 95 \text{g mol}^{-1})$

OR

- (i) Why is the value of Van't Hoff factor for ethanoic acid in benzene close to 0.5?
- (ii) Determine the osmotic pressure of a solution prepared by dissolving 2.32 \times 10^{-2} g of K₂SO₄ in 2L of solution at 25 °C, assuming that K₂SO₄ is completely dissociated.
- $(R = 0.082 \text{ L atm } K^{-1} \text{ mol}^{-1}, \text{ Molar mass } K_2SO_4 = 174g \text{ mol}^{-1})$
- (iii) When 25.6g of Sulphur was dissolved in 1000g of benzene, the freezing point lowered by 0.512 K. Calculate the formula of Sulphur (S_x) .

 $(K_f \text{ for benzene} = 5.12 \text{ K kg mol}^{-1}, \text{ Atomic mass of Sulphur} = 32g \text{ mol}^{-1})$

Marks:[5.00]

Q.No.34: (i) Write the reaction involved in Cannizaro's reaction.

- (ii) Why are the boiling point of aldehydes and ketones lower than that of corresponding carboxylic acids ?
- (iii) An organic compound 'A' with molecular formula $C_5H_8O_2$ is reduced to n-pentane with hydrazine followed by heating with NaOH and Glycol.

'A' forms a dioxime with hydroxylamine and gives a positive Iodoform and Tollen's test. Identify 'A' and give its reaction for Iodoform and Tollen's test.

OK

- (i) Give a chemical test to distinguish between ethanal and ethanoic acid.
- (ii) Why is the a-hydrogens of aldehydes and ketones are acidic in nature?
- (iii) An organic compound 'A' with molecular formula $C_4H_8O_2$ undergoes acid hydrolysis to form two compounds 'B' and 'C'.

Oxidation of 'C' with acidified potassium permanganate also produces 'B'.

Sodium salt of 'B' on heating with soda lime gives methane.

- (1) Identify 'A', 'B' and C'.
- (2) Out of 'B' and 'C', which will have higher boiling point? Give reason.

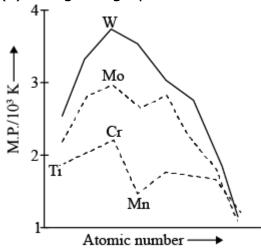
Marks:[5.00]

Q.No.35: (a) Why is chemistry of actinoids complicated as compared to lanthanoids?

(b) Complete the following reaction and justify that it is a disproportionation reaction:

 $3\,{\rm MnO_4^{2-}} + \,4{\rm H^+} \rightarrow \underline{\hspace{1cm}} + \,2{\rm H_2O}.$

(c) The given graph shows the trends in melting points of transition metals:



Explain the reason why Cr has highest melting point and manganese (Mn) a lower melting point.

Marks:[5.00]