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# STUDY MATERIAL CLASS XII (CHEMISTRY) 2023-24

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# Chapter 1 : Solutions

Choose and write the correct option(s) in the following questions.

1. If we place the blood cells in a solution containing less than 0.9% (m/V) sodium chloride. They would swell. This is because:

- (a) the solution is hypotonic
- (b) ) the solution is isotonic with blood
- (c) the solution is hypertonic
- (d) none of these

2. On dissolving sugar in water at room temperature, solution feels cool to touch. Under which of the following cases dissolution of sugar will be most rapid?

- (a) Sugar crystals in cold water.
- (b) Powdered sugar in cold water.
- (c) Sugar crystals in hot water.
- (d) Powdered sugar in hot water.

3. Considering the formation, breaking and strength of hydrogen bond, predict which of the following mixtures will show a positive deviation from Raoult's law?

- (a) Methanol and acetone.
- (b) Chloroform and acetone.
- (c) Nitric acid and water.
- (d) Phenol and aniline.

4. Which of the following aqueous solutions should have the highest boiling point?

- (a) 1.0 M NaOH
- (b) 1.0 M Na<sub>2</sub>SO<sub>4</sub>
- (c) 1.0 M NH<sub>4</sub>NO<sub>3</sub>
- (d) 1.0 M KN<sub>3</sub>

5. Which of the following statements is false?

(a) Two different solutions of sucrose of same molality prepared in different solvents will have the same depression in freezing point.

(b) The osmotic pressure of a solution is given by the equation  $\pi = CRT$  (where

C is the molarity of the solution).

(c) Decreasing order of osmotic pressure for 0.01 M aqueous solutions of barium chloride,

potassium chloride, acetic acid and sucrose is BaCh<sub>2</sub> > KCI > CH<sub>3</sub> COOH > sucrose.

(d) According to Raoult's law, the vapour pressure exerted by a volatile component of a solution is directly proportional to its mole fraction in the solution.

6. At equilibrium the rate of dissolution of a solid solute in a volatile liquid solvent is

- (a) less than the rate of crystallisation
- (b) greater than the rate of crystallisation
- (c) equal to the rate of crystallisation
- (d) zero

7. One kilogram of sea water sample contains 6 mg of dissolved  $O_2$ . The concentration of  $O_2$  in ppm in sample is

- (a) 0.06
- (b) 60
- (c) 6
- (d) 0.6

8. Mole fraction of the solute in a 100 molal aqueous solution is:

- (a) 0.1770
- (b) 0.0177
- (c) 0.0644
- (d) 0.2334

9. Low concentration of oxygen in the blood and tissues of people living at high altitude is due to

- (a) low.temperature
- (b) low atmospheric pressure
- (c) high atmospheric pressure
- (d) both low temperature and high atmospheric pressure

10. A solution of chloroform in diethylether:

- (a) obeysRaoult's law
- (b) shows a positive deviation from Raoult's law
- (c) shows a negative deviation from Raoult's law
- (d) behaves like a near ideal solution

11. Vapour pressure of a pure liquid X is 2 atm at 300 K. It is lowered to 1 atm on dissolving 1 g of Y in 20 g of liquid X. If molar mass of X is 200, what is the molar mass of Y?

- (a) 20
- (b) 50
- (c) 100
- (d) 200

12. In comparison to a 0.01 M solution of glucose, the depression in freezing point of a 0.01 M MgCl<sub>2</sub> solution is\_\_\_\_\_.

- (a) the same
- (b) about twice
- (c) about three times

(d) about six times

13. An unripe mango placed in a concentrated, salt solution to prepare pickle, shrivels because\_\_\_\_\_.

- (a) it gains water due to osmosis.
- (b) it loses water due to reverse osmosis.
- (c) it gains water due to reverse osmosis.
- (d) it loses water due to osmosis.

14. At a given temperature, osmotic pressure of a concentrated solution of a substance\_\_\_\_\_.

- (a) is higher than that of a dilute solution.
- (b) is lower than that of a dilute solution.
- (c) is same as that of a dilute solution.

(d) cannot be compared with osmotic pressure of dilute solution.

15. The values of Van't Hoff factors for KCI, NaCI and K<sub>2</sub>SO<sub>4</sub>, respectively, are

- (a) 2, 2 and 2
- (b) 2, 2 and 3
- (c) 1, 1 and 2
- (d) 2,1 and 2
- 16. Which of the following statements is false?

(a) Units of atmospheric pressure and osmotic pressure are the same.

(b) In reverse osmosis, solvent molecules move through a semipermeable membrane from a region of lower concentration of solute to a region of higher concentration.

(c) The value of molal depression constant depends on nature of solvent.

(d) Relative lowering of vapour pressure, is a dimensionless quantity.

17. The system that forms maximum boiling azeotropes is:

- (a) ethyl alcohol-water
- (b) benzene-toluene
- (c) acetone-chloroform
- (d) carbon disulphide-acetone
- 18. The value of Henry's constant is \_\_\_\_\_.
- (a) greater for gases with higher solubility.
- (b) greater for gases with lower solubility.
- (c) constant for all gases
- (d) not related to the solubility of gases

19. Which of the following factor(s) affect the solvent?

(i) nature and solute (ii) temperature (iii) pressure

(a) (i) and (iii) at constantT
(b) (i) and (ii) at constant P
(c) (ii) and (iii) only
(d) (iii) only

20. Intermolecular forces between two benzene molecules are nearly of same strength as those between two toluene molecules. For a mixture of benzene and toluene, which of the following are not true?

- (a)  $\Delta H_{mixing}$ = zero
- (b)  $\Delta v_{mixing} = zero(c)$
- (c) These will form minimum boiling azeotrope
- (d) These will not form ideal solution

## **Assertion reason type questions**

In the following questions, a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.

(a) Assertion and reason both are correct statements and reason is correct explanation for assertion.

(b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.

(c) Assertion is correct statement but reason is wrong statement.

(d) Assertion is wrong statement but reason is correct statement.

21. Assertion: Molarity of a solution in liquid state changes with temperature. ReasonThe volume of a solution changes with change in temperature.

22Assertion: If more volatile liquid is added to another liquid, vapour pressure of solution will be greater than that of pure solvent.

Reason:Vapour pressure of solution is entirely due to solvent molecules.

23. AssertionWhen NaCl is added to water a depression in freezing point is observed. Reason: The lowering of vapour pressure of a solution causes depression in the freezing point.

24. AssertionWhen a solution is separated from the pure solvent by a semipermeable membrane, the solvent molecules pass through it from pure solvent side to the solution side. Reason: Diffusion of solvent occurs from a region of high concentration solution to a region of low

25.Assertion: The solubility of a gas in a liquid increases with increase of pressure.Reason: The solubility of a gas in a liquid is directly proportional to the pressure of the gas.26. Assertion: The vapour pressure of a liquid decreases if some non-volatile solute is dissolved in it.

Reason: The relative lowering of vapour pressure of a solution containing a non-volatile solute is equal to the mole fraction of the solute in the solution.

27.Assertion: Molecular mass of polymers cannot be calculated using boiling point or freezing point

Reason: Polymers solutions do not possess a constant boiling point or freezing point.

28.Assertion: The boiling point of 0.1 M urea solution is less than that of 0.1 M KCl solution. ReasonElevation of boiling point is directly proportional to the number of species present

29. Assertion: Non-ideal solutions always form azeotropes.

Reason: Boiling point of an azeotrope may be higher or lower than boiling points of both components.

30.Assertion: Lowering of vapour pressure is directly proportional to osmotic pressure of the solution.

Reasonn: Osmotic pressure is a colligative property.

### Answer kev

1. a
2. d
3. a
4. b
5. a
6. c
7. c
8. b
9. b
10. c
11. a
12. c
13. d
14. a
15. b
16. b

17. c 18. b 19. (a),(b) 20. (c) ,(d) 21. a 22. c 23. a 24. c 25. a 26. b 27. c

28. a

29. d

30. b

## **CHAPT2 - ELECTROCHEMISTRY**

## **Multiple Choice Questions**

Choose and write the correct option(s) in the following questions. 1. Which cell will measure standard electrode potential of copper electrode? (a)  $Pt(s)|H_2(g, 0.1 bar)|H^+(aq., 1 M)||Cu^{2+}(aq., 1M)|Cu$ (b)  $Pt(s)|H_2(g, 1 bar) |H^+(aq., 1 M) ||Cu^{2+}(aq., 2M)|Cu$ (c)  $Pt(s)|H_2(g, 1 bar) |H^+(aq., 1 M)||Cu^{2+}(aq., 1M)|Cu$ (d)  $Pt(s)|H_2(g, 1 bar) |H^+(aq., 0.1 M)| |Cu^{2+}(aq., 1M)|Cu$ 2. A hypothetical electrochemical cell is shown below:  $A | A^{+}(xM) || B^{+}(yM) | B$ The emf measured is +0.20 V. The cell reaction is (a)  $A + B^+ \longrightarrow A^+ + B$ (b)The cell reaction cannot be predicted. (c)  $A^+ + e^- \longrightarrow A$ ,  $B^+ + e^- \longrightarrow B$  $(d) \mathbf{A}^+ + \mathbf{B} \longrightarrow \mathbf{A} + \mathbf{B}^+$ 3. A device that converts energy of combustion of fuels like hydrogen and methane directly into electrical energy is known as: (a) dynamo (b) Ni-Cd cell (c) fuel cell (d) electrolytic cell 4. The difference between the electrode potentials of two electrodes when no current is drawn through the cell is called (a) Cell potential (b) Cell emf (c) Potential difference (d) Cell voltage 5. Which of the following statement is not correct about an inert electrode in a cell? (a) It does not participate in the cell reaction. (b) It provides surface either for oxidation or for reduction reaction. (c) It provides surface for conduction of electrons. , (d) It provides surface for redox reaction. 6. Charge carried by 1 mole of electrons is (a)  $6.023 \times 10^{23}$  coulomb (b)  $9.65 \times 10^4$  coulomb (d)  $6.28 \times 10^{19}$  coulomb (c)  $1.6 \times 10^{-19}$  coulomb 7. Standard electrode potential for  $Sn^{4+}/Sn^{2+}$  couple is +0.15 V and that for the  $Cr^{3+}/Cr$  couple is -0.74 V. The two couple in their standard states are connected to make cell. The cell potential will be: (b) 0.89 V (a) +1.19 V (d) +1.83 V · (c) +0.18 V 8. Which one of the following is always true about the spontaneous cell reaction in a galvanic cell? (b)  $E_{cell}^{o} < 0, \Delta G^{o} < 0, Q < K_{c}$ (a)  $E_{cell}^{o} > 0, \Delta G^{o} < 0, Q > K_{c}$ (d)  $E_{\text{cell}}^{\circ} > 0, \Delta G^{\circ} < 0, Q < K_{c}$ (c)  $E_{\text{cell}}^{o} > 0, \Delta G^{o} > 0, Q > K_{c}$ 9. The electrode potential data are given below  $\mathrm{Fe}^{3^+} + e^- \longrightarrow \mathrm{Fe}^{2^+}.$  $E^{0} = +0.77 \text{ V}$  $Al^{3+} + 3e^- \longrightarrow Al, \qquad E^0 = -1.66 V$  $Br_2 + 2e^- \longrightarrow 2Br^ E^{0} = +1.80 \text{ V}$ Based on the data, the reducing power of Fe2+, Al, Br- will increase in the order (a)  $Br^{-} < Fe^{2+} < Al$ (b)  $Fe^{2+} < Al < Br^{-}$ (c) Al  $< Br^{-} < Fe^{2+}$ (d)  $Al < Fe^{2+} < Br^{-1}$ 

10. An electrochemical cell can behave like an electrolytic cell when \_ (d)  $E_{\text{cell}} = E_{\text{ext}}$ (c)  $E_{\text{ext}} > E_{\text{cell}}$ (b)  $E_{\text{cell}} > E_{\text{ext}}$ (a)  $E_{\text{cell}} = 0$ 11. Which of the statements about solutions of electrolytes is not correct? (a) Conductivity of solution depends upon size of ions. (b) Conductivity depends upon viscosity of solution. (c) Conductivity does not depend upon solvation of ions present in solution. (d) Conductivity of solution increases with temperature. 12. Using the data given below find out the strongest reducing agent.  $E_{\rm Cl_2/Cl^-}^{\rm o} = 1.36 \, {\rm V}$  $E_{\rm Cr_2O_2^{2-}/Cr^{3+}}^{\rm o} = 1.33 \, \rm V$  $E_{MnO_4^-/Mn^{2+}}^{o} = 1.51 V$   $E_{Cr^{3+}/Cr}^{o} = -0.74 V$ (c) Cr<sup>3+</sup>  $(d) \text{ Mn}^{2+}$ (a) Cl<sup>-</sup> (b) Cr 13. Using the data given in Q. 12 find out in which option the order of reducing power is corr. (b)  $Mn^{2+} < Cl^{-} < Cr^{3+} < Cr$ (a)  $Cr^{3+} < Cl^{-} < Mn^{2+} < Cr$ (c)  $Cr^{3+} < Cl^{-} < Cr_2 O_7^{2-} < MnO_4^{-}$ (d)  $Mn^{2+} < Cr^{3+} < Cl^{-} < Cr$ 14. Use the data given in Q. 12 find out the most stable oxidised species. (*d*)  $Mn^{2+}$ (a)  $Cr^{3+}$ (b)  $MnO_4^-$ (c)  $Cr_2O_7^{2-}$ 15. The quantity of charge required to obtain one mole of aluminium from Al<sub>2</sub>O<sub>3</sub> is (a) 1F (b) 6F (c) 3F (d) 2F 16. The standard emf of a galvanic cell involving 3 moles of electrons in a redox reaction is 0.59 V. The equilibrium constant for the reaction of the cell is: (a)  $10^{10}$  $(b) 10^{20}$ (c)  $10^{30}$  $(d) 10^{15}$ 17.  $\Lambda^{\circ}_{m(NH_4OH)}$  is equal to \_\_\_\_\_ (a)  $\Lambda^{\circ}_{m(\mathrm{NH}_4\mathrm{OH})} + \Lambda^{\circ}_{m(\mathrm{NH}_4\mathrm{Cl})} - \Lambda^{\circ}_{(\mathrm{HCl})}$ (b)  $\Lambda^{\circ}_{m(\mathrm{NH},\mathrm{C})} + \Lambda^{\circ}_{m(\mathrm{NaOH})} - \Lambda^{\circ}_{(\mathrm{NaCh})}$ (c)  $\Lambda^{\circ}_{m(\mathrm{NH}_{4}\mathrm{Cl})} + \Lambda^{\circ}_{m(\mathrm{NaCl})} - \Lambda^{\circ}_{(\mathrm{NaOH})}$ (d)  $\Lambda^{\circ}_{m(\text{NaOH})} + \Lambda^{\circ}_{m(\text{NaCl})} - \Lambda^{\circ}_{(\text{NH, Cl})}$ 18. In the electrolysis of aqueous sodium chloride solution which of the half cell reaction will occur at anode? (a)  $\operatorname{Na}^+(aq) + e^- \longrightarrow \operatorname{Na}(s);$  $E_{cell}^{o} = -2.71 \text{ V}$ (b)  $2H_2O(l) \longrightarrow O_2(g) + 4H^+(aq) + 4e^-;$  $E_{cell}^{o} = 1.23 V$ (c)  $H^+(aq) + e^- \longrightarrow \frac{1}{2}H_2(g);$  $E_{\rm cell}^{\rm o} = 0.00 \, {\rm V}$ (d)  $\operatorname{Cl}^{-}(aq) \longrightarrow \frac{1}{2} \operatorname{Cl}_{2}(g) + e^{-};$  $E_{cell}^{o} = 1.36 V$ 19. The positive value of the standard electrode potential of Cu<sup>2+</sup>/Cu indicates that (a) this redox couple is a stronger reducing agent than the  $H^+/H_2$  couple. (b) this redox couple is a stronger oxidising agent than  $H^+/H_2$ . (c) Cu can displace  $H_2$  from acid. (d) Cu cannot displace  $H_2$  from acid. 20.  $E_{cell}^{o}$  for some half cell reactions are given below. On the basis of these mark the correct answer. (i)  $H^+(aq) + e^- \longrightarrow \frac{1}{2}H_2(g)$  $E_{cell}^{0} = 0.00 V$ (*ii*)  $2H_2O(l) \longrightarrow O_2(g) + 4H^+(aq) + 4e^-$ (*iii*)  $2SO_4^{2-}(aq) \longrightarrow S_2O_8^{2-}(aq) + 2e^-;$   $E_{cell}^0 = 1.23 V$   $E_{cell}^0 = 1.96 V$ 

- (a) In dilute sulphuric acid solution, hydrogen will be reduced at cathode.
- (b) In concentrated sulphuric acid solution, water will be oxidised at anode.
- (c) In dilute sulphuric acid solution, water will be oxidised at anode.
- (d) In dilute sulphuric acid solution,  $SO_4^{2-}$  ion will be oxidised to tetrathionate at anode.

## **Assertion-Reason Questions**

In the following questions a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.

- (a) Assertion and reason both are correct statements and reason is correct explanation for assertion.
- (b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
- (c) Assertion is correct statement but reason is wrong statement.
- (d) Assertion is wrong statement but reason is correct statement.

21. Assertion	1 (A)	):	Cu is less reactive than hydrogen.
Reason	( <b>R</b> )	):	$E_{Cu^{2+}/Cu}^{o}$ is negative.
22.Assertion	<b>(A</b> )	):	$E_{\text{cell}}$ should have a positive value for the cell to function.
Reason	( <b>R</b> )	):	$E_{\rm cathode} < E_{\rm anode}$
23.Assertion	(A)	:	Conductivity of all electrolytes decreases on dilution.
Reason	(R)	:	On dilution number of ions per unit volume decreases.
24 Assertion	(A)	:	$\Lambda_m$ for weak electrolytes shows a sharp increase when the electrolytic solution is diluted.
Reason	(R)	:	For weak electrolytes degree of dissociation increases with dilution of solution.
25Assertion	(A)	:	Mercury cell does not give steady potential.
Reason	( <b>R</b> )	:	In the cell reaction, ions are not involved in solution.
26Assertion	(A)	:	Current stops flowing when $E_{cell} = 0$ .
Reason	(R)	:	Equilibrium of the cell reaction is attained.
27Assertion	(A)	:	$E_{Ag^{+}/Ag}$ increases with increase in concentration of Ag <sup>+</sup> ions.
Reason	( <b>R</b> )	:	$E_{Ag^{+}/Ag}$ has a positive value.
28.Assertion	(A)	:	Copper sulphate cannot be stored in zinc vessel.
Reason	(R)	:	Zinc is less reactive than copper.
29.Assertion	(A)	:	Electrolysis of NaCl solution gives chlorine at anode instead of O2.
Reason	( <b>R</b> )	:	Formation of oxygen at anode requires overvoltage.
	(A)		For measuring resistance of an ionic solution an AC source is used.
Reason	(R)		Concentration of ionic solution will change if DC source is used.

## Answers

<b>1.</b> (c)	<b>2.</b> (a)	<b>3.</b> (c)	<b>4.</b> (b)	5. (d)	<b>6.</b> (b)	7. (b)	<b>8.</b> ( <i>d</i> )
<b>9.</b> (a)	<b>10.</b> (c)	<b>11.</b> (c)	12. (b)	13. (b)	14. (a)	15. (c)	<b>16.</b> (c)
<b>17.</b> (b)	<b>18.</b> ( <i>d</i> )	<b>19.</b> (b), (d)	<b>20.</b> (a), (c)			-200 Y	

21.(C)22.(C)23.(a)24.(a)25.(d)26.(a)27.(b)28.(C)29.(a)30.(a)

## <u>CHAPTER 3</u> CHEMICAL KINETICS

### MCQ

- 1. The reaction rate constant can be defined as the rate of reaction when each reactant's concentration is
- a) Zero
- b) Unity
- c) Doubled the initial concentration
- d) Infinite
- 2. What effect does temperature have on the half-life of a first-order reaction?
- a) It increases
- b) It decreases
- c) It remains the same
- d) Both increases as well as decrease
- 3. The role of catalyst is to change
- a) Gibb's energy of the reaction
- b) Enthalpy of the reaction
- c) Activation energy of the reaction
- d) Equilibrium constant
- 4. Activation Energy of a chemical reaction can be determined by
- a) Determining the rate constant at standard temperature
- b) Determining the rate constant at two temperatures
- c) Using catalyst
- d) Determining probability of collision

5. If half - life period of a reaction is inversely proportional to initial concentration of reactant, What is the order of reaction is

- a) first order
- b) second order
- c) Third order
- d) Zero order
- 6. Which of the following statement is not correct
- a) Order of a reaction can be fractional number
- b) Order of a reaction is experimentally determined quantity

c) Order of a reaction is always equal to the sum of the stoichiometric coefficient of reactant in the balanced chemical reaction

d) The order of the reaction is the sum of the powers of molar concentration of the reactants in the rate law expression

7. Hydrolysis of ester follows the kinetics of:

a)first order b) second order c) zero order d) pseudo first

8. Rate Law for the reactionA + 2B -----  $\rightarrow$  C is found to be Rate = K [A][B]. Concentration of B is doubled keeping the concentration of A constant, the value of rate constant will be:

a) Remain same

b) Doubled

c) Quadrupled

d) Halved

9. If 75% of the first order reaction was completed in 32minutes , 50% of the same reaction would be completed in

a) 8min

b) 4 minutes

c) 16 minutes

d) 24minutes

10. For Zero Order reaction a)  $t_{1/2}\alpha$  a b)  $t_{1/2}\alpha$  1/a

c)  $t_{1/2} \alpha a^2$ 

d)  $t_{1/2} \alpha 1/a^2$ 

11. For the reaction :  $2NH_3$  (Pt)-- $\rightarrow$  N<sub>2</sub> +  $3H_2(g)$  Rate = K[NH<sub>3</sub>]<sup>0</sup>. Order of the reaction is

a) First order

b) Second order

c) Zero order

d)Third order

12. The half -life of first order reaction is 69.3 sec. The value of rate constant of the reaction is: a)  $1.0 \text{ s}^{-1}$  b)  $0.1 \text{ s}^{-1}$  c)  $0.01 \text{ s}^{-1}$  d)  $0.001 \text{ s}^{-1}$ 

13. In a reaction  $2A - --- \rightarrow product$ . Concentration of A decreases from 0.5 mol L<sup>-1</sup> to 0.4molL<sup>-1</sup> in10 minutes. The rate of the reaction during the interval is a) 5 mol L<sup>-1</sup> min b) 0.5 mol L<sup>-1</sup> min c) 0.05 mol L<sup>-1</sup> min d) 0.005 mol L<sup>-1</sup> min

14. For a reaction  $A + B - --- \rightarrow$  Product. The rate Law is given as rate = k [A]<sup>1/2</sup>[B]<sup>2</sup>. What is the order of reaction ?

a) 2 b) 2/5 c) 5/2 d) 3

15. From the rate expressions for the following reaction determine the dimensions of the rate constant .3NO (g) ---→ N<sub>2</sub>O (g) NO<sub>2</sub>(g) rate = K[NO]<sup>2</sup>
a) mol<sup>-1</sup> L s<sup>-1</sup>
b) s<sup>-1</sup>
c) mol<sup>-2</sup> L<sup>2</sup> s<sup>-1</sup>
d) mol<sup>-2</sup> L s<sup>-1</sup>
16. For the reaction A+B→C molecularity is?

a) Molecularity is more than Zero

b) Molecularity is less than Zero

c) Molecularity is equal Zero

d) Molecularity is infinite

17. A reaction is second order with respect to reactant. How is the rate affected If the concentration of reactant is doubled

a) increase by 2 times of initial rate

b) decrease by 2 times of initial rate

c) increase by 4 times of initial rate

d) decrease by 4 times of initial rate

18. Calculate the half – life of first order reaction if rate constant is given as 4 years a)  $1.73 \times 10^{-1}$  year b)  $1.73 \times 10^{-1}$  min c)  $2.37 \times 10^{-2}$  year d)  $17.3 \times 10^{-1}$  year

19. The decomposition of NH<sub>3</sub> on platinum surface is zero order. What is the rate of production of H<sub>2</sub> if  $k = 2.5 \times 10^{-4} \text{mol}^{-1} \text{ L s}^{-1}$ a) 2.5 x 10<sup>-4</sup>mol<sup>-1</sup> L s<sup>-1</sup> b) 7.5 x 10<sup>-4</sup>mol<sup>-1</sup> L s<sup>-1</sup> c) 25.5 x 10<sup>-4</sup>mol<sup>-1</sup> L s<sup>-1</sup> d) 20.5 x 10<sup>-4</sup>mol<sup>-1</sup> L s<sup>-1</sup>

20. Half-Life of zero order reaction is

a) 0.693 / k

b)  $[R]_0/2k$ 

c) [R]<sub>0</sub>/k

d)  $t_{1/2}$  is independent of  $[R]_0$ 

21. Elementary reactions are those reactions which takes place in

a) multiple steps

b) two steps

c) single step

d) none of the above

22. For a reaction,  $A + H_2O - B$ . Rate  $\alpha$  [A], the order and molecularity of reaction is

a) order = 1 & molecularity = 1

b) order = 2 & molecularity = 2

c) order = 3 &molecularity = 2

d) order = pseudo order &molecularity = 2

23. In a reaction, when the concentration of reactant is increased two times, the increase in rate of reaction was four times. Order of reaction is  $(a) \operatorname{zero}(b) 1 (c) 2 (d) 3$ 

(a) zero(b) 1 (c) 2 (d) 3

24. If the rate of a gaseous reaction is independent of pressure, the order of reaction is:

(a) 0 (b) 1 (c) 2 (d) 3

25. Order of reaction can be

(a) Zero(b) fraction (c) whole number (d) integer, fraction, Zero

26. Instantaneous rate of a chemical reaction is

(a) rate of reaction in the beginning (b) rate of reaction at the end

- (c) rate of reaction at a given instant (d) rate of reaction between two specific time intervals
- 27. In the rate equation, when the concentration of reactants is unity then rate is equal to
- (a) specific rate constant
- (b)average rate constant
- (c) instantaneous rate constant
- (d) None of the above
- 28. The activation energy in a chemical reaction is defined as
- (a) the difference in energies of reactants and products
- (b) the sum of energies of reactants and products
- (c) the difference in energy of intermediate complex with the average energy of reactants and products
- $\left(d\right)$  the difference in energy of intermediate complex and the average energy of reactant
- 29. A plot of log (a x) against time t is a straight line. This indicates that the reaction is of (a) zero order (b) first order (c) second order (d) third order
- 30. Rate of which reaction increases with temperature?
- (a) Exothermic reaction (b) Endothermic reaction (c) (a) and (b) both (d) None of the above
- 31. Radioactive disintegration is an example of
- (a) zero order reaction (b) first order reaction (c) second order reaction (d) third order reaction
- 32. The overall rate of a reaction is governed by
- (a) the rate of fastest intermediate step
- (b) the sum total of the rates of all intermediate steps
- $(\mathbf{c})$  the average of the rates of all the intermediate steps
- (d) the rate of slowest intermediate step
- 33. For a unimolecular reaction
- (a) the order and molecularity of the slowest step are equal to one
- (b) molecularity of the reaction can be zero, one or two
- (c) molecularity of the reaction can be determined only experimentally
- (d) more than one reacting species are involved in one step
- 34. The unit of rate and rate constant are same for a
- (a) zero order reaction (b) first order reaction (c) second order reaction (d) third order reaction
- 35. The rate constant of a reaction depends upon
- (a) temperature of the reaction (b) extent of the reaction
- (c) initial concentration of the reactants (d) the time of completion of reaction
- 36. In chemical equation H<sub>2</sub> (g) + I<sub>2</sub> (g)  $\rightleftharpoons$  2HI (g) the equilibrium constant Kp depends on
- (a) total pressure
- $(b) \ catalyst \ used$
- $(c) \ amount \ of \ H_2 \ and \ I_2$
- (d) temperature

#### ANSWER KEY

(1) b	(2) b	(3) c	(4) b	(5) b	(6) c
(7) d	(8) a	(9) c	(10) a	(11) c	(12) c
(13) d	(14) c	(15) a	(16) a	(17) c	(18) a
(19) b	(20) b	(21) c	(22) d	(23) c	(24) a
(25) d	(26) c	(27) a	(28) d	(29) b	(30) b
(31) b	(32) d	(33) a	(34) a	(35) a	(36) d

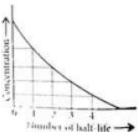
## CASE BASED STUDY

## CHEMICAL KINETICS

#### **1. Read the passage given below and answer the following questions:**

The half-life of a reaction is the time required for the concentration of reactant to decrease by half

i.e., $[A]_t = [A]/2$ For first order reaction, $t_{1/2} = 0.693$ /kthis means  $t_{1/2}$  is independent of initial concentration. Figure shows that typical variation of concentration of reactant exhibiting first order kinetics. It may be noted that though the major portion of the first order kinetics may be over in a finite time, but the reaction will never cease as the concentration of reactant will be zero only at infinite time.



The following questions are multiple choice question. Choose the most appropriate answer: (i) A first order reaction has a rate constant  $k=3.01 \times 10^{-3}$  /s. How long it will take to decompose half of the reactant?

(a) 2.303 s (b) 23.03 s (c) 230.3 s (d) 2303 s

# (ii) The rate constant for a first order reaction is 7.0 x $10^{-4}$ s<sup>-1</sup>. If initial concentration of reactant is 0.080 M, what is the half-life of reaction?

a) 990 s b) 79.2 s c) 12375 s d) 10.10 x 10<sup>-4</sup> s

# (iii) For the half-life period of a first order reaction, which one of the following statements is generally false?

a) It is independent of initial concentration. b) It is independent of temperature.

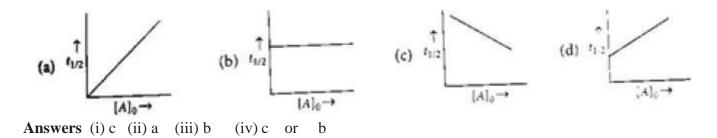
c) It decreases with the introduction of a catalyst. d) None of these.

(iv) The rate of a first order reaction is 0.04 mol  $L^{-1}$  s<sup>-1</sup> at 10 minutes and 0.03 mol  $L^{-1}$  s<sup>-1</sup> at 20 minutes after initiation. The half life of the reaction is

a) 4.408 min b) 44.086 min c) 24.086 min d)2.408 min

#### **OR**

The plot of  $t_{1/2}$  vs initial concentration [A]<sub>0</sub> for a first order reaction is given by



## Passage -2Read the passage given below and answer the following questions:

The branch of physical chemistry that is concerned with understanding the rates of chemical reactions. It is to be contrasted with thermodynamics which deals with the direction in which a process occurs but in itself tells nothing about its rate. Thermodynamics is time's arrow while chemical kinetics is time's clock. Chemical kinetics relates to many aspects of cosmology, geology, biology, engineering and even psychology and thus has far-reaching implications. The principles of chemical kinetics applied to purely physical processes as well as to chemical reactions. The rate of a chemical reaction is defined in terms of the rates with which the products are formed and the reactants (the reacting substances) are consumed. For chemical systems, it is usual to deal with the concentrations of substances, which is defined as the amount of substances per unit volume. The rate can then be defined as the concentration of a substance that is consumed or produced in unit time. Sometimes it is more convenient to express rates as number of molecules formed are consumed in unit time.

READ THE STATEMENTS GIVEN AS ASSERTION & REASON BOTH AND CHOOSE THE CORRECT OPTION AS (A)/(B)/(C)/(D) AS PER THE FOLLOWING INSTRUCTIONS

(A) – if both assertion & reason are correct statements and reason is the correct explanation of assertion

(B)– if both assertion & reason are correct statements and reason is not the correct explanation of assertion

(C) if assertion is the correct statement & reason is incorrect statement.

(D) – if assertion is incorrect statement and reason is the correct statement.

1.Assertion: All collisions of reactant molecules do not lead to product formation.

Reason: Only those collisions in which molecules have correct orientation and sufficient kinetic energy lead to compound formation.

2.Assertion: The enthalpy of reaction remains constant in the presence of a catalyst

Reason: A catalyst lowers down the activation energy but the difference in energy of reactant and product remains same.

3Assertion: The rate of reaction is the rate of change of concentration of a reactant or product. Reason: The rate of reaction remains constant during the course of a reaction.

4.Assertion: Instantaneous rate is used to predict the rate of reaction at a particular moment.Reason: Average rate is constant for the time interval for which it is calculated.

ANSWERS 1.A 2. A 3. C 4. B

## Passage – 3

**Read the passage given below and answer the following questions:** The principles of chemical kinetics applied to purely physical processes as well as to chemical reactions. The rate of a chemical reaction is defined in terms of the rates with which the products are formed and the reactants (the reacting substances) are consumed. For chemical systems, it is usual to deal with the concentrations of substances, which is defined as the amount of substances per unit volume. The rate can then be defined as the concentration of a substance that is consumed or produced in unit time. Sometimes it is more convenient to express rates as number of molecules formed are consumed in unit time.

### II. MULTIPLE CHOICE QUESTIONS

1. A reaction is first order with respect to reactant A. What will be the initial rate, if concentration of reactant is reduced by half?

(a) Halved (b) doubled (c) tripled (d) not changed

2. What is the ratio of the rate of decomposition of  $N_2O_5$  to the rate of formation of  $NO_2$ ?

 $2N_2O_5(g) - --- > 4NO_2(g) + O_2(g)$ 

(a) 1:4 (b) 2;1 (c) 4:1 (d) 1:2

3. In the first order reaction, the concentration of the reactant is reduced to 1/4th in 60 minutes. What will be its half-life?

(a) 120 min (b) 40 min (c) 30 min (d) 25 min

- 4. Which of the following reaction ends in infinite time?
- (a) Zero order (b) First order (c) Second order (d) Third order
- 5. The rate constant of a zero order reaction is-
- (a) Directly proportional to initial concentration (b) Doesn't depend on concentration
- (c) Inverselyproportional to initial concentration (d) Can't say
- 6. The first order reaction takes 80 minutes to complete 99.9%. What will be its half-life?

a) 8 min (b) 16 min (c) 24 min (d) 32 min

7. A catalyst increases the reaction rate by:

- (a) decreasing enthalpy (b) increasing internal energy
- (c) decreasing activation enthalpy (d) increasing activation enthalpy
- 8. Chemical kinetics is the study to find out:
- (a) The feasibility of a chemical reaction(c) Extent to which a reaction will proceed
- (b) Speed of a reaction (d) All of the above

9. The rate of a reaction:

(a) Increases as the reaction proceeds (b) Decreases as the reaction proceeds (c) Remains the same as the reaction proceeds (d) May increase or decrease as the reaction proceeds

10. Which of the following influences the reaction rate performed in a solution?

(a) Temperature (b) Activation energy (c) Catalyst (d) All of the above

### passage 4:

## Read the passage given below and answer the following questions:

The rate of a chemical reaction is expressed either in terms of decrease in the concentration of a reactant per unit time or increase in the concentration of a product per unit time. Rate of the reaction depends upon the nature of reactants, concentration of reactants, temperature, presence of catalyst, surface area of the reactants and presence of light. Rate of reaction is directly related to the concentration of reactant. Rate law states that the rate of reaction depends upon the concentration terms on which the rate of the reactants in the rate as observed experimentally. The sum of powers of the concentration of the reactants in the rate law expression is called order of reaction while the number of reacting species taking part in an elementary reaction which must collide simultaneously in order to bring about a chemical reaction is called molecularity of the reaction.

- 1. Why do pieces of wood burn faster than log of wood of same mass?
  - a) Pieces of wood has less surface area
  - b) Pieces of wood has large surface area
  - c) Log of wood more mass
  - d) Pieces of wood has less mass
- 2. What is a complex reaction
  - a) Reaction which takes place in single step
  - b) Reaction takes place in two or more steps
  - c) Reaction which form complex
  - d) None of the above
- 3. When initial concentration of reactact is doubled the half life period of zero order reaction
  - a) Remain unchanged b) is halved c) is tripled d) is Doubled

4. Addition of catalyst during a chemical reaction alters which of the following

a) Entropy b) Activation energy c) internal enery d) Enthalpy

OR If the temperature of a reaction is increased by 20°C the rate of reacrion increase by a) One time b) two times c) three times d) four times Answer :

1. B 2. B 3. D 4. B OR D

#### Passage 5:

## Read the passage given below and answer the following question

Temperature has a marked effect on the rate of reaction. For most of the reactions, the rate of reaction becomes nearly double for every  $10^{\circ}$  degree rise in temperature. The effect of temperature is usually expressed in terms of temperature coefficient. The quantitative dependence of reaction rate on temperature was first explained by Swante Arrhenius. Arrhenius proposed a simple equation called as Arrhenius equation :  $k = A e^{-Ea/RT}$ . The equation provides the relationship between the rate constant (k) of a reaction and the Temperature of the system (T). (A) is the Arrhenius factor or pre-exponential factor. (E<sub>a</sub>) is the activation energy and (R) is gas constant.

- Activation energy of a reaction from the slope of which of the following graph?
   a) In k vs T
   b) In k / T vs T
   c) In k vs 1/T
   d) T / InK vs 1/T
- 2. According to Arrhenius equation ,rate constant of a chemical reaction isequal to a)  $K = A e^{-Ea/RT}$  b)  $k = A e^{Ea/RT}$  c)  $k = A e^{-RT/Ea}$  d)  $k = A e^{RT/Ea}$
- Rate of a reaction is doubled for every 10°C raise in temperature. If the temperature is raised by 20°C the rate of reaction increases by

a) 2 times b) 4 times c) 3 times d) no change

- 4. Activation energy of a chemical reaction can be determined by
  - a) Determining the rate constant at standard temperature
  - b) Determining the rate constant at two temperatures
  - c) Determining probability of reaction

d) Using catalyst

**ANSWER**: 1 = a, 2 = a, 3 = b, 4 = b

## Passage 6:

## Read the passage given below and answer the following question

Rate = k [A]x [B]y x and y indicate how sensitive the rate is to the change in concentration of A and B. Sum of these exponents, i.e., x + y in (3.4) gives the overall order of a reaction whereas x and y represent the order with respect to the reactants A and B respectively. Hence, the sum of powers of the concentration of the reactants in the rate law expression is called the order of that chemical reaction. Order of a reaction can be 0, 1, 2, 3 and even a fraction. A zero order reaction means that the rate of reaction is independent of the concentration of reactants. A balanced chemical equation never gives us a true picture of how a reaction takes place since rarely a reaction gets completed in one step. The reactions taking place in one step are called elementary reactions. When a sequence of elementary reactions (called mechanism) gives us the products, the reactions are called complex reactions.

- Calculate the overall order of a reaction which has the rate expression Rate = k [A]1/2 [B]3/2
  - a) Second order b) first order c) Zero order d) Fraction order
- 2. For first order reaction which of the following statement is correct
  - a) T1/2 independent of [R]o
  - b) t1/2 dependent of [R]o
  - c) t1/2 is equal to [R]o
  - d) None of above
- 3. For zero order reaction rate of reaction is
  - a) independent of the concentration of reactants.
  - b) dependent of the concentration of reactants.
  - c) independent of the concentration of products
  - d) dependent of the concentration of reactants.
- 4. In first order reaction Which of the following statement is correct
  - a) Time taken for completion of 99.9% reaction is 10 times of t1/2 of reaction
  - b) Time taken for completion of 99.9% reaction is 20 times of t1/2 of reaction
  - c) Time taken for completion of 99.9% reaction is 99.9 times of t1/2 of reaction

d) Time taken for completion of 99.9% reaction is equal to times of t1/2 of reaction ANSWERS 1- A, 2 - A, 3 - A + A

## <u>CHAPTER 4</u> <u>d& f block elements</u>

1. The number of unpaired electrons in gaseous species of  $Mn^{3+}$ ,  $Cr^{3+}$  and  $V^{3+}$  respectively are:

- a) 4, 4 and 2
- b) 3, 3 and 2
- c) 4, 3 and 2
- d) 3, 3 and 3
- 2. Gun metal is an alloy of:
- a) Cu and Al
- b) Cu and Sn
- c) Cu, Zn and Sn
- d) Cu, Zn and Ni

**3.** Which one of the following elements shows the maximum number of different oxidation states in its compounds?

- a) Eu
- b) La
- c) Cd
- d) Am

4. Zinc and mercury do not show variable valency like d-block elements because

a) they are soft

- b) their d-shells are complete
- c) they have only two electrons in the outermost subshell.
- d) their d-shells are incomplete
- 5. The pair of elements that has similar atomic radii is
- a) Mn and Re
- b) Ti and Hf
- c) Sc and Ni
- d) Hf and Zr

6. Silver nitrate produces a black stain on the skin due to

- a) being a strong reducing agent
- b) its corrosive action
- c) formation of complex compound
- d) its reduction to metallic silver

7. The transition metal that exhibit largest number of oxidation state is

- a) Cr
- b) Mn
- c)Zn
- c) Sc

8.  $Mn^{2+}$  compounds are more stable than  $Fe^{2+}$  compounds towards oxidation in +3 statebecause

- a) 3d<sup>5</sup> configuration is more stable than 3d<sup>6</sup> configuration
- b) 3d<sup>6</sup> configuration is more stable than 3d<sup>5</sup> configuration
- c) 3d<sup>5</sup> configuration is less stable than 3d<sup>6</sup> configuration
- d)  $3d^6$  configuration is less stable than  $3d^5$  configuration
- 9. Transition metals show paramagnetic behaviour due to:
- a) Paired electrons in the (n-1)d orbital
- b) Unpaired electrons in the (n-1)d orbital
- c) Due to  $d^{10}$  configuration
- d) None of the above
- 10. The correct statement(s) about interstitial compounds is/are:
- a) Transition elements are large in size and contain multiple interstitial sites
- b) Transition elements can trap atoms of other elements
- c) Interstitial elements are chemically innert

d) All the above

11. The d block elements that may not be regarded as transition elements are

- a) Mn, Fe, Ni due to partially filled d orbital
- b) Mn, Fe, Ni due to completely filled d orbital
- c) Zn, Cd, & Hg due to completely filled d orbital

d) ZN, Cd, & Hg due to partially filled d orbital

12. Element that does not show variable oxidation state is

a) Fe b) Mn c) Cu d) Zn

13. The magnetic momentum of  $Mn^{2+}$  is

a) 5.92BM

b) 2.80Bm

c) 8.95Bm

d) 3.9BM

14. The enthalpy of atomisation of transition metals is high due to

- a) High effective nuclear charge
- b) A large number of valence electrons
- c) very strong metallic bond
- d) All the above
- 15. Which of the following is colourless ion

a) Cr<sup>4+</sup>

b)  $Sc^{3+}$ 

c) Ti<sup>3+</sup>

- d)  $V^{3+}$
- 16. The number of 3d electrns in Cr3+ ion is

a) 6 b) 3 c)5 d) 4

17. Transition metals and their compounds act as good catalyst due to

a) Variable oxidation state

b) unstable intermediate

c)provide a suitable surface

d) All the above

18. Which of the following is in the correct order of increasing oxidising power? a)  $VO_2^+ < Cr_2O_7^{2-} < MnO_4^{-1}$ b)  $VO_2^+ < MnO_4^{-1} < Cr_2O_7^{2-}$ c)  $Cr_2O_7^{2-} < VO^+ < MnO_4^{-1}$ d)  $Cr_2O_7^{2-} < MnO_4^{-1} > VO_2^+$ 

19. The electronic configuration of gadolinium (atomic no 64 )is a) [Xe]  $4f^35d^56s^2$ b) [Xe]  $4f^75d^26s^1$ c) [Xe]  $4f^75d^16s^2$ d) [Xe]  $4f^85d^6s^2$ 

20. Which of the following statement is correct
a) Cu(II)aq is more stable
b) Cu(II)aq is less stable
c)Cu(I)aq& Cu(II)a are equally stable
d) stability of Cu(I) & Cu(II)depends upon nature of copper salt

21. The element that has maximum composition in Mischmetal and used in gas lighter is a) Ce b) Fe c) Sm d) La 22. When neutral alkaline KMnO<sub>4</sub> is treated with potassium iodide, the iodide ion is converted into X. X is a)  $I_2$  b)  $IO_4^-$  c)  $IO_3^-$  d)  $IO^-$ 

23. Different oxidation states exhibited by the lanthanoids are

a) +2 , +3 ,+4 b) +2 , +3 , -4 c) -2 , -3 -4 d) +2 , +4 , +5

24. The most stable oxidation state of lanthanoids and actinoids is a) +5 b) +2 c) +3 d) +1

25. On addition of small amount of KMnO4 to concentrated  $H_2SO_4$ , a green oily compound sobtained . which is highly explosive in nature . Identify the compound a)  $Mn_2O_7$  b)  $MnO_2$  c)  $MnSO_4$  d)  $Mn_2O_3$ 

26. Actinoids exhibit greater number of oxidation state than lanthanoid .The reason is

a) more energy difference between 5f, 6d than 4f and 5f

b) 4f orbitals are more diffuse than 5f orbital

c) Lesser energy difference between 5f and 6d than 4f and 5dorbtals

d) More reactive nature of actinoids than lanthanoids

27. Transition metals and their compounds show coloured salts because of

- a) presence of paired electrons in d orbital
- b) absence of unpaired electrons in d orbital
- c) presence of unpaired electron in d orbital
- d) Presence of paired electron in f orbital

28. Highest oxidation state of manganese in fluorides is +4 (MnF<sub>4</sub>) but with oxygen is +7 (Mn<sub>2</sub>O<sub>7</sub>)because

a) Fluorine is more electronegative than oxygen

b) Fluorine does not contain d orbital

c) Fluorine stabilises lower oxidation state

d) In covalent compounds fluorine can form single bond while oxygen can form double bond

29. Why is HCl not used to make the medium acidic in oxidation reaction of KMnO<sub>4</sub>?

a) Both HCl& KMnO<sub>4</sub> act as oxidising agents

- b) KMnO<sub>4</sub> oxidises HCl to Cl<sub>2</sub>
- c) KMnO<sub>4</sub> is a weaker oxidising agent than HCl
- d) KMnO<sub>4</sub> act as reducing agent in presence of HCl

30. Although Zirconium belongs to 4d transition series and Hoffnium to 5d series they show similar physical and chemical properties because:

a) Both belong to d block

- b) Both have same number of electrons
- c) Both have similar atomic radius
- d) Both belong to same group of the periodic table

31. Cu+ is not stable in aqueous solutions because

- a) In aqueous solution it undergo disproportionation reaction to form stable Cu<sup>2+</sup> ion
- b) It contain d<sup>10</sup> configuration
- c) it is colourless in cu<sup>+1</sup> state
- d) none of the above

- 32. Transition metals processes high melting point because
- a) presence of paired electrons
- b) of weak metallic bond
- c) due to strong metallic bond
- d) of covalent bond

### 33. Name a member of the lanthanide series which is well known to exhibit +4 oxidation state.

- (a) Cerium (X=58)
- (b) Europium (Z=63)
- (c) Lanthanum (Z=57)
- (d) Gadolinium (Z=64)

### 34. Identify the incorrect statement among the following.

- (a) Lanthanide contraction is the accumulation of successive shrinkages.
- (b) the different radii of Zr and Hf due to consequences of the lanthanide contraction.
- (c) Shielding power of 4f electrons is quite weak.
- (d) None of the above

### 35. Though Silver contain completely filled d orbital (d<sup>10</sup>) still called transition element

- a) It shows +1 oxidation state
- b) It shows +2 oxidation state with d<sup>9</sup> configuration
- c) It is a metal
- d) It is a soft metal

### Answer key

- 1. c
- 2. c
- 3. d
- 4. b
- 5. d
- 6. d
- 7. b
- 8. a
- 9. b
- 10. d
- 11. c
- 12. d
- 13. a
- 14. d 15. b
- 15.0
- 16. b 17. d
- 17. u 18. a

19. c 20. a 21. a 22. c 23. a 24. c 25. a 26. c 27. c 28. d 29. b 30. c 31. a 32. c 33. a

34. a

35. b

## d& f Block elements CASEBASE STUDY

## Passage -1

## Read the passage given below and answer the following questions:

The f-block elements are those in which the differentiating electrons enters the (n-2)f orbitals. There are two series of f-Block elements corresponding to filling of 4f and 5f-orbitals. The series of 4f-orbitals is called lanthanides. Lanthanides show different oxidation states depending upon stability of  $f^0$ ,  $f^7$  and  $f^{14}$  configurations, though the most common oxidation states is +3. There is a regular decrease in the size of lanthanides ions with increase in atomic number which is known as lanthanides contraction.

The following questions are multiple choice question. Choose the most appropriate answer:

(i) The atomic number of three lanthanides elements X, Y and Z are 65, 68 and 70 respectively, their  $Ln^{3+}$  electronic configuration is

(a)  $4f^8$ ,  $4f^{11}$ ,  $4f^{13}$ (b)  $4f^{11}$ ,  $4f^8$ ,  $4f^{13}$ (c)  $4f^0$ ,  $4f^2$ ,  $4f^{11}$ (d)  $4f^3$ ,  $4f^7$ ,  $4f^9$ 

- (ii) lanthanide contraction is observed in
- (a) Gd
- (b) At
- (c) Xe
- (d) Te

#### (iii) Which of the following is not the configuration of lanthanide?

(a)  $[Xe]4f^{10}6s^2$ (b)  $[Xe]4f^{15}d^{1}6s^2$ (c)  $[Xe]4d^{14}5d^{10}6s^2$ (d)  $[Xe]4f^75d^{1}6s^2$ 

Or

#### Name a member of the lanthanide series which is well known to exhibit +4 oxidation state.

- (a) Cerium (X=58)
- (b) Europium (Z=63)
- (c) Lanthanum (Z=57)
- (d) Gadolinium (Z=64)

#### (iv) Identify the incorrect statement among the following.

(a) Lanthanide contraction is the accumulation of successive shrinkages.

(b) the different radii of Zr and Hf due to consequences of the lanthanide contraction.

(c) Shielding power of 4f electrons is quite weak.

(d) There is a decrease in the radii of the atoms or ions proceeds from La to Lu

Answer(i) a (ii) a (iii) c or a (iv) b

## Passage 2

## Read the passage given below and answer the following questions:

The D-Block elements are those elements in which the last electron enters the d-subshell of the penultimate shell. The general electronic configuration of these elements is (n-1) d <sup>1-10</sup>ns<sup>1-2</sup>, where n is the outermost shell. The d-block consisting of groups 3-12 occupies the large middle section of the periodic table. The elements of the d-block are also known as transition elements as they possess properties that are transitional between s&p block elements. A transitional element is defined as the elements which has incompletely filled d-orbitals in its ground state or any other of its oxidation states. There are four series of transition elements spread between group 3&12. First transition series are 3d series: Scandium to Zinc. Second transition series are 4d series: Yttrium to Cadmium. Third transition series are 5d series: Lanthanum to Mercury. Fourth transition series are 6d series: Begins with Actinium and is still incomplete.

- 1. Which block elements are called transitional elements
  - a) S-block b) p-block c) d-block d) f-block
- 2. Why is the Zinc compounds are colourless
  - a. Absence of unpaired electrons in p orbital
  - b. Presence of unpaired electrons in p orbital
  - c. Presence of unpaired electrons in d- orbital
  - d. Absence of unpaired electrons in d- orbital
- 3. Though Silver contain completely filled d orbital (d<sup>10</sup>) still called transition element
  - a. It shows +1 oxidation state
  - b. It shows +2 oxidation state with d<sup>9</sup> configuration
  - c. It is a metal
  - d.It is a soft metal
- 4. Which of the following d block element exist in liquid state
  - a. Zn b) Mn c) Hg d) Fe
    - Or

Name the transition element used as filament in bulb a) Copper b) Iron c) Alluminium d) Tungsten ANSWERS : 1 - C, 2 - c, 3 - b, 4 - c OR d

## Passage 3:

## Read the passage given below and answer the following questions:

Potassium permanganate is prepared by fusion of pyrolusite MnO<sub>2</sub> with KOH in the presence of an oxidizing agent like KNO<sub>3</sub>. This produces the dark green Potassium manganate K<sub>2</sub>MnO<sub>4</sub> which disproportionates in a neutral or acidic solution to give purple permanganate ion. Potassium permanganate is an important oxidizing agent in acidic, alkaline, as well as neutral medium.

- 1. What is the state of hybridisation of Mn in MnO<sub>4</sub>
- a) SP2 b) SP3 c) SP d) dSP3
- 2. The colour of  $KMnO_4$  is due to
  - a) Potassium ion
  - b) Charge transfer
  - c) d d transition
  - d) K and O double bond
- The products obtained by oxidation of KI in acid medium by KMnO<sub>4</sub> is

   a) Iodine b) iodate ion c) water d) MnI<sub>2</sub>
- The number of electrons transferred when KMnO₄ act as oxidising agent in acid medium is To give Mn<sup>2+</sup>
  - a) 3 b) 5 c) 1 d) 7

#### ANSWERS

1. = b, 2 = c, 3 = a, 4 = b

## Passage 4.

## Read the passage given below and answer the following questions:

The transition elements have incompletely filled d subshells in their ground state or in any of their oxidation states. The transition elements occupy position between s & p bblock in groups 3 - 12 of the periodic table. Starting from fourth period transition elements consists of four series: Sc to Zn , Y to Cd , La Hf to Hg and Ac Rf to Cn. In general electronic configuration of outer orbitalsis (n- $10d^{1-10}ns^{1-2}$ . The electronic configuration of Zn , Cd . Hg are represented by general formula (n-1) $d^{10}ns^2$ . All the transition elements have typical metallic properties liketensile strength , ductility , malleability. Except mercury which is a liquid at room temperature all other elements have typical metallic structure. Transition metals and their compounds exhibit catalytic property ,& paramagnetic behaviour , alloy formation

- 1. Which of the following characteristics of transition metals is associated catalytic activity
  - a) High enthalpies of atomisation b) Varuable oxidation state

c)Paramagnetic behaviour d) colour of hydrated ion

2. Transition elements forms alloys because

- a) Same atomic number b) same electronic configuration c) nearly same atomic size d) same oxidation state
- 3. Transition elements forms coloured compounds because
  - a) nearly same atomic size b) d-d transition

c) high charge on transition metal ion d) presence of paired electrons

- 4. Which of the following characteristics of transition metals is associated Complex formation
  - a) nearly same atomic size b) d-d transition c) high charge on transition metal ion d) presence of paired electrons

Answers 1-b, 2-c, 3-b, 4-c

#### Passage 5.

#### Read the passage given below and answer the following questions:

The stable halides of the 3d series of transition metals. The highest oxidation numbers are achieved in TiX4 (tetrahalides), VF5 and CrF6 . The +7 state for Mn is not represented in simple halides but MnO3F is known, and beyond Mn no metal has a trihalide except FeX3 and CoF3 . The ability of fluorine to stabilise the highest oxidation state is due to either higher lattice energy as in the case of CoF3 , or higher bond enthalpy terms for the higher covalent compounds, e.g., VF5 and CrF6. Although V+5 is represented only by VF5, the other halides, however, undergo hydrolysis to give oxohalides, VOX3 . Another feature of fluorides is their instability in the low oxidation states e.g., VX2 (X = Cl, Br or I) 4.3.6 Trends in the M3+/M2+ Standard Electrode Potentials 4.3.7 Trends in Stability of Higher Oxidation States + 6 CrF6 + 5 VF5 CrF5 + 4 TiX4 VXI 4 CrX4 MnF4 + 3 TiX3 VX3 CrX3 MnF3 FeXI 3 CoF3 + 2 TiX2 III VX2 CrX2 MnX2 FeX2 CoX2 NiX2 CuX2 II ZnX2 + 1 CuXIII ever, many copper (I) compounds are unstable in aqueous solution and undergo disproportionation. 2Cu+ Æ Cu2+ + Cu The stability of Cu2+ (aq) rather than Cu+ (aq) is due to the much more negative DhydH o of Cu2+ (aq) than Cu+ , which more than compensates for the second ionisation enthalpy of Cu.

- 1. Why is the highest oxidation state of a metal exhibited in its oxide or fluoride only?
  - a) Fluorine and oxygen large in size
  - b) Small size and high electronegativity of fluorine and oxygen
  - c) large size and low electronegativity of fluorine and oxygen
  - d) larger size and high electropositivity of fluorine and oxygen
- 2. Which is a stronger reducing agent Cr2+ or Fe2+ and why ?
  - a) Fe2+ is stronger reducing agent than Cr2+ Reason: d 4  $\rightarrow$  d 3 occurs in case of Cr2+ to Cr3+ But d 6  $\rightarrow$  d 5 occurs in case of Fe2+ to Fe3+
  - b) Cr2+ is stronger reducing agent than Fe2+ Reason: d 4  $\rightarrow$  d 3 occurs in case of Cr2+ to Cr3+ But d 6  $\rightarrow$  d 5 occurs in case of Fe2+ to Fe3+
  - c) Cr2+ is equal reducing agent like Fe2+ Reason: d 4  $\rightarrow$  d 3 occurs in case of Cr2+ to Cr3+ But d 6  $\rightarrow$  d 5 occurs in case of Fe2+ to Fe3+
  - d) None of the above
- 3. Explain why Cu+ ion is not stable in aqueous solutions?
  - a) Cu+ in aqueous solution undergoes disproportionation
  - b) Cu2+ in aqueous solution undergoes disproportionation
  - c) Cu in aqueous solution undergoes disproportionation
  - d) None of above

- 4. The lowest oxide of transition metal is basic where as the highest is acidic.because
  - a) The lower oxides are ionic while higher oxides are covalent.
  - b) The lower oxides are covalentwhile higher oxides are ionic
  - c) The lower oxides are ionic & higher oxides are ionic
  - d) The lower oxides are covalent & higher oxides are covalent ANSWER: 1 B, 2 B, 3 A, 4 B

## <u>CHAPTER 5</u> COORDINATION COMPOUNDS MCQS

- 1. Trunbull's blue is
  - (a) Ferricyanide
  - (b) Ferrous ferricyanide
  - (c) Ferrous cyanide
  - (d)  $Fe_3[Fe(CN)_6]_4$

2. The complex ions  $[Co(NH_3)_5(NO_2)]^{2+}$  and  $[Co(NH_3)_5(ONO)]^{2+}$  are called

- (a) Ionization isomers
- (b) Linkage isomers
- (c) Co-ordination isomers
- (d) Geometrical isomers

#### 3. Which of the following has square planar structure?

- (a)  $[NiCl_4]^{2-}$
- (b) [Ni(CO)<sub>4</sub>]
- (c) [Ni(CN)4]<sup>2-</sup>
- (d) None of these
- 4. Which of the following has magnesium?
- (a) Chlorophll
- (b) Haemocyanin
- (c) Carbonic anhydrate
- (d) Vitamin B<sub>12</sub>

5. Mohr's salt is (a)  $Fe_2(SO_4) \ 3 \ (NH_4)_2SO_4 \ 6H_2O$ (b)  $FeSO_4 \ (NH_4)_2 \ SO_4 \ 6H_2O$ (c)  $MgSO_4 \ 7H_2O$ (d)  $FeSO_4 \ 7H_2O$ 

6. Which of the following shall form an octahedral complex?

- (a)  $d^4$  (low spin)
- (b)  $d^8$  (high spin)
- (c)  $d^6$  (low spin)
- (d) All of these
- 7. EDTA is used for the estimation of
- (a)  $Na^+$  and  $K^+$  ions
- (b)  $Cl^{-}$  and  $Br^{-}$  ions
- (c)  $Cu^{2+}$  and  $Cs^+$  ions
- (d)  $Ca^{2+}$  and  $Mg^{2+}$  ions

8. The solution of the complex [Cu(NH<sub>3</sub>)<sub>4</sub>] SO<sub>4</sub> in water will

- (a) give the tests of  $Cu^{2+}$  ion
- (b) give the tests of  $NH_3$
- (c) give the tests of  $SO_4^{2-}$  ions
- (d) not give the tests of any of the above

9. The sum of coordination number of oxidation number of the metal M in the complex  $\left[M(en)_2\,C_2O_4\right]\,Cl$  are

- (a) 7
- (b) 8
- (c) 9
- (d) 6

#### 10. Which of the following will not give test for $Cl^-$ with AgNO<sub>3</sub>(aq) at 25°C?

- (a) COCl<sub>3</sub>.5NH<sub>3</sub>
- (b) COCl<sub>3</sub>.6NH<sub>3</sub>
- (c) COCl<sub>3.</sub>3NH<sub>3</sub>
- (d) COCl<sub>3</sub>.4NH<sub>3</sub>
- 11. Which of these statements about  $[Co(CN)_6]^{3-}$  is true?
- (a) It has 4 unpaired electron, high spin
- (b) No unpaired electron, high spin
- (c) No unpaired electron, low spin
- (d) 4 unpaired electron, low spin

12. The correct order of the stoichiometries of AgCl formed when AgNO<sub>3</sub> in excess is treated with complexes: CoCl<sub>3</sub>.6NH<sub>3</sub>, CoCl<sub>3</sub>.5NH<sub>3</sub>, CoCl<sub>3</sub>.4NH<sub>3</sub> respectively is

- (a) 3AgCl, lAgCl, 2AgCl
- (b) 3AgCl, 2AgCl, 1AgCl
- (c) 2AgCl, 3AgCl, 2AgCl
- (d) lAgCl, 3AgCl, 2AgCl
- 13. Correct increasing order of wavelength of absorption in visible region for complex of  $Co^{3+}$  is
- (a)  $[Co(H_2O)_6]^{3+}$ ,  $[Co(en)_3]^{3+}$ ,  $[Co(NH_3)_6]^{3+}$
- (b)  $[Co(H_2O)_6]^{3+}$ ,  $[Co(NH_3)_6]^{3+}$ ,  $[Co(en)_3]^{3+}$
- (c)  $[Co(NH_3)_6]^{3+}$ ,  $[Co(en)_3]^{3+}$ ,  $[Co(H_2O)_6]^{3+}$
- (d)  $[Co(en)_3]^{3+}$ ,  $[Co(NH_6)_6]^{3+}$ ,  $[Co(H_2O)_6]^{3+}$
- 14. Pick out the correct statement with respect to  $[Mn(CN)_6]^{3-1}$
- (a) It is sp<sup>2</sup>d<sup>2</sup> hybridised, tetrahedral
- (b) It is d<sup>2</sup>sp3 hybridised, octahedral
- (c) It is dsp<sup>2</sup> hybridised, square planar,
- (d) It is  $sp^3d^2$  hybridised octahedral

15. Facial and meridional isomerism will be shown by (a)  $[Co(NH_3)_3Cl_3]$ (b)  $[Co(NH_3)_4Cl_2]$  Cl (c)  $[Co(en)_3] Cl_3$ (d)  $[Co(NH_3)_5Cl] Cl_2$ 16. Which one has highest molar conductivity? (a)  $[Pt(NH_3)_2 Cl_2]$ (b)  $[CO(NH_3)_4 Cl_2] Cl$ (c)  $K_4[Fe(CN)_6]$ (d)  $[Cr(H_2O)_6] Cl_3$ 17. Which one will show optical isomerism? (a)  $[Co(NH_3)_3Cl_3]$ (b) cis- $[Co(en)_2 Cl_2] Cl$ (c) trans- $[Co(en)_2 Cl_2] Cl$ (d)  $[Co(NH_3)_4 Cl_2] Cl_3$ 18. The pair having the same magnetic moment is (At No. Cr = 24, Mn = 25, Fe = 26, Co = 27) (a)  $[Cr(H_2O)_6]^{2+}$  and  $[CoCl_4]^{2-}$ (b)  $[Cr(H_2O)_6]^{2+}$  and  $[Fe(H_2O)_6]^{2+}$ (c)  $[Mn(H_2O)_6]^{2+}$  and  $[Cr(H_2O)_6]^{2+}$ (d)  $[CoCl_4]^{2-}$  and  $[Fe(H_2O)_6]^{2+}$ 19. On treatment of 100 mL of 0.1 M CoCl<sub>3.6</sub>H<sub>2</sub>O with excess of AgNO<sub>3</sub>  $1.2 \times 10^2$  ions are precipitated. The complex is (a)  $[Co(H_2O)_4Cl_2]Cl_2H_2O$ (b)  $[Co(H_2O)_3Cl_3]_3H_2O$ (c)  $[Co(H_2O)_6] Cl_3$ (d) [Co(H<sub>2</sub>O)<sub>5</sub> Cl] Cl<sub>2</sub>.H<sub>2</sub>O 20. Among the ligands  $NH_3$  en, CN- and CO, the correct order of field strength is (a)  $NH_3 < en < CN^- < CO$ (b)  $CN^{-} < NH_{3} < CO < en$ (c) en < CN $^{-}$ < NH $_{3}$  < CO (d)  $CO < NH_3 < en < CN^-$ 21. Which cf the following complexes formed by  $Cu^{2+}$  ions is most stable? (a)  $\operatorname{Cu}^{2+} + 4\operatorname{NH}_3 \rightleftharpoons [\operatorname{Cu}(\operatorname{NH}_3)_4]^{2+}$ , logK = 11.6(b)  $\operatorname{Cu}^{2+} + 4\operatorname{CN}^{-} \rightleftharpoons [\operatorname{Cu}(\operatorname{CN})_4]^{2-}$ , logK = 27.3(c)  $Cu^{2+} + 2en [Cu(en),]^{2+}$ , logK = 15.4(d)  $Cu^{2+} + 4H_2O \rightleftharpoons [Cu(H_2O)_4]^{2+}$ ,  $\log K = 8.9$ 

22. The stabilisation of coordination compounds due to chelation is called the chelate effect. Which of the following is the most stable complex species?

(a)  $[Fe(CO)_5]$ 

(b)  $[Fe(CN)_{61}^{3-}]$ 

(c)  $[Fe(C_2O_4)_3]^{3-}$ (d)  $[Fe(H_2O)_6]^{3+}$ 

23. Indicate the complex ion which shows geometrical isomerism. (a)  $[Cr(H_2O)_4Cl_2]^+$ 

(b)  $[Pt(NH_3)_3 Cl]$ 

(c)  $[Co(NH_3)_6]^{3+}$ 

(d)  $[Co(CN)_5(NC)]^3$ -

24. In which one of the following species does the transition metal ion have  $d^3$  electronic configuration?  $\begin{array}{l} [A] \ [Cr(NH_3)_6]^{3+} \\ [B] \ [Co(H_2O)_6]^{2+} \end{array}$ 

 $[C] [CoF_6]^{3-}$ 

 $[D] [Fe(CN)_6]^{3-}$ 

25. (Crystal Field Theory) When the valence d orbitals of the central metal ion are split in energy in an octahedral ligand field, which orbitals are raised least in energy?

[A] dxy and  $dx^2-y^2$ 

[B] dxy, dxz and dyz

[C] dxz and dyz

[D] None of the above

26. The number of ions formed, when bis (ethane-1,2-diamine)copper (II) sulpahte is dissolve in water will be:

[A] 1

[B] 2

[C] 3

[D] None of the above

27. Among the following ions which one has the highest paramagnetism

 $[A] [Cr(H_2O)_6]^{3+}$  $[B] [Fe(H_2O)_6]^{2+}$  $[C] [Cu(H_2O)_6]^{2+}$  $[D] [Zn(H_2O)_6]^{2+}$ 

28. A complex of certain metal has the magnetic moment of 4.91 BM whereas another complex of the same metal with same oxidation state has zero magnetic moment. The metal ion could be [A] Co<sup>2+</sup>

[B] Mn<sup>2+</sup>

[C] Fe<sup>2+</sup>

[D] Fe<sup>3+</sup>

- 29. The CFSE for octahedral  $[CoCl_6]^{4-}$  is 18,000 cm<sup>-1</sup>. The CFSE for tetrahedral  $[CoCl_4]^{2-}$  will be (a) 18,000 cm<sup>-1</sup>.
- (b) 16,000 cm<sup>-1</sup>
- (c)  $8,000 \text{ cm}^{-1}$
- (d) 20,000 cm<sup>-1</sup>

30. Due to the presence of ambidentate ligands coordination compounds show isomerism. Palladium complexes of the type  $[Pd(C_6H_5)_2(SCN)_2]$  and  $[Pd(C_6H_5)_2(NCS)_2]$  are

- (a) linkage isomers
- (b) coordination isomers
- (c) ionisation isomers
- (d) geometrical isomers

31. The compounds [Co(SO<sub>4</sub>)(NH<sub>3</sub>)<sub>5</sub>]Br and [Co(SO<sub>4</sub>)(NH<sub>3</sub>)<sub>5</sub>]Cl represent

- (a) linkage isomerism
- (b) ionisation isomerism
- (c) coordination isomerism
- (d) no isomerism

32. A chelating agent has two or more than two donor atoms to bind to a single metal ion. Which of the following is not a chelating agent?

- (a) thiosulphato
- (b) oxalato
- (c) glycinato
- (d) ethane-1, 2-diamine

33. The solution of the complex [Cu(NH<sub>3</sub>)<sub>4</sub>] Cl in water will

- (a) give the tests of  $Cu^{2+}$  ion
- (b) give the tests of NH<sub>3</sub>
- (c) give white ppt with AgNO<sub>3</sub>
- (d) not give the tests of any of the above

34. IUPAC name of [Pt(NH<sub>3</sub>)<sub>3</sub> Br (NO<sub>2</sub>) Cl] Cl is

(a)triamminechlorodibromidoplatinum (IV) chloride

(b)triamminechloridobromidonitrochloride- platinum (IV) chloride

(c)triamminebromidochloridonitroplatinum (IV) chloride

(d)triamminenitrochlorobromoplatinum (IV) chloride

35. Primary and secondary valency of Pt in [Pt(en)<sub>2</sub>Cl<sub>2</sub>] are

- (a) 4, 4
- (b) 4, 6
- (c) 6, 4
- (d) 2, 6

36. The name of complex  $[Fe(CN)_6]^{3-}$  is

- (a) Tricyanido ferrate (III) ion
- (b) Hexacyanido ferrate (III) ion

(c) Hexacyanido iron (III)

(d) Hexacyanido ferrate (II) ion

### Answer key

(1) b	(2) b	(3) c	(4) a	(5) b	(6) b
(7) d	(8) c	(9) c	(10) c	(11) c	(12) b
(13) d	(14) b	(15) a	(16) c	(17) b	(18) b
(19) d	(20) a	(21) b	(22) c	(23) a	(24) a
(25) b	(26) b	(27) b	(28) c	(29) c	(30) a
(31) b	(32) a	(33) c	(34) c	(35) d	(36) b

### Assertion Reason type questions

Choose the correct option:

(a) Assertion and reason both are correct and the reason is the correct explanation of assertion.

(b) Both assertion and reason are true but the reason is not the correct explanation of assertion.

(c) Assertion is correct but Reason is incorrect

(d) Assertion is a wrong statement but the reason is a correct statement

(e) Both Assertion and Reason are incorrect

1. Assertion: Toxic metal ions are removed by the chelating ligands. **Reason**: Chelate complexes tend to be more stable.

2. Assertion: Linkage isomerism arises in coordination compounds containing ambidentate ligand. **Reason:** Ambidentate ligand has two different donor atoms.

3. Assertion: Complexes of MX<sub>6</sub>, MX<sub>5</sub>L type (X and L are unidentate) do not show geometrical isomerism.

Reason: Geometrical isomerism is not shown by complexes of coordination number 6.

4. Assertion:  $[Fe(CN)_6]^{3-}$  ion shows magnetic moment corresponding to two unpaired electrons. **Reason:** Because it has  $d^2sp^3$  type hybridization.

5. Assertion:  $[Ni(CN)_4]^2$  is square planar and diamagnetic

**Reason**: It has no unpaired electrons due to the presence of a strong ligand.

6. Assertion: Oxalate ion is a bidentate ligand. Reason: Oxalate ion has two donor atoms

7. Assertion:  $[Ni(CO)_4]$  has square planar geometry while  $[Ni(CN)_4]^{4-}$  has tetrahedral geometry **Reason:** The geometry of any complex depends upon the nature of the ligands attached.

8. Assertion:  $[FeF_6]^{3-}$  is a low spin complex. **Reason**: Low spin complexes have a lesser number of unpaired electrons.

9. Assertion:  $F^-$  ion is a weak ligand and forms an outer orbital complex. **Reason**:  $F^-$  ion cannot force the electrons of  $dz^2$  and  $dx^2 - y^2$  orbitals of the inner shell to occupy  $d_{xy}$ ,  $d_{yz}$ , and  $d_{zx}$  orbitals of the same shell.

10. **Assertion:** Potassium ferrocyanide and potassium ferricyanide both are diamagnetic. **Reason**: Both have unpaired electrons.

11. Assertion: The  $[Ni(en)_3]Cl_2$  (en = ethylene diamine) has lower stability than  $[Ni(NH_3)_6]Cl_2$ **Reason**: In  $[Ni(en)_3]Cl_2$  the geometry of Ni is trigonal bipyramidal.

12. Assertion: The ligands nitro and nitrito are called ambidentate ligands. **Reason:** These ligands give linkage isomers.

13. **Assertion**: Geometrical isomerism is also called cis-trans isomerism. Reason : Tetrahedral complexes show geometrical isomerism.

14. Assertion:  $[Fe(CN)_6]^{3-}$  is weakly paramagnetic while  $[Fe(CN)_6]^{4-}$  is diamagnetic. **Reason**:  $[Fe(CN)_6]^{3-}$  has +3 oxidation state while  $[Fe(CN)_6]^{4-}$  has +2 oxidation state.

15. Assertion: Potassium ferrocyanide is diamagnetic whereas potassium ferricyanide is paramagnetic. **Reason**: Crystal field splitting in ferrocyanide ion is greater than that of ferricyanide ion.

16. Assertion:  $NF_3$  is a weaker ligand than  $N(CH_3)_3$ . **Reason**:  $NF_3$  ionizes to give  $F^-$  ions in aqueous solution.

17. Assertion:  $[Fe(CN)_6]^{3-}$  is weakly paramagnetic while  $Fe(CN)_6]^{4-}$  is diamagnetic.

**Reason**:  $[Fe(CN) ^{3-}$  has +3 oxidation state while  $[Fe(CN) ]^{4-}$  has +2 oxidation state.

18. Assertion:  $[Ti(H_2O)_6]^{3+}$  is coloured while  $[Sc(H_2O)_6]^{3+}$  is colourless. **Reason**: d-d transition is not possible in  $[Sc(H_2O)_6]^{3+}$ .

### Answer key

(1) a	(2) a	(3) c	(4) e	(5) a	(6) a
(7) d	(8) d	(9) a	(10) e	(11) e	(12) a
(13) c	(14) b	(15) c	(16) c	(17) b	(18) a

# <u>CHAPTER 6</u> HALOALKANES AND HALOARENES

- 1. Which of the following is the right name for the compound H<sub>3</sub>C-CHCl<sub>2</sub>?
- a) 1,2-Dichloroethane
- b) Ethylene dichloride
- c) Ethylidene chloride
- d) Vic-dichloride

2. Which of the following substances has the highest melting point?

- a) Chloromethane
- b) Tetrachloromethane
- c) Trichloromethane
- d) Dichloromethane

3. Which sequence should isomeric dichlorobenzenes be boiled in?

- a) para>ortho>meta
- b) meta>ortho>para
- c) ortho>meta>para
- d) para>meta>ortho

4. Which of the following undergoes nucleophilic substitution exclusively by SN 1 mechanism?

- (a) Benzyl chloride
- (b) Ethyl chloride
- (c) Chlorobenzene
- (d) Isopropyl chloride
- 5. The increasing order of nucleophilicity would be
- $\begin{array}{ll} (a) \ Cl^- < Br^- < l^- \\ (b) \ l^- < Cl^- < Br^- \\ (c) \ Br^- < Cl^- < F^- \\ (d) \ l^- < Br^- < Cl^- \end{array}$

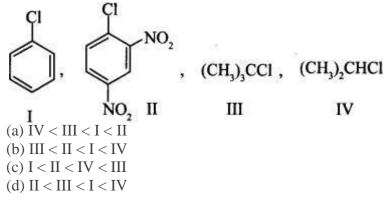
6.  

$$CH_3 - CH - CH = CH_2 + HBr \longrightarrow A$$
; 'A' is  
 $CH_3$   
 $Br$   
 $(a) CH_3 - C - CH_2 - CH_3$   
 $CH_3$   
 $CH_3$   
 $(b) CH_3 - CH - CH - CH_3$   
 $CH_3$   
 $(c) CH_3 - CH - CH_2 - CH_2Br$   
 $CH_4$   
 $(d) BrCH_2 - CH - CH_2 - CH_3$   
 $(d) BrCH_2 - CH - CH_2 - CH_3$   
 $(d) BrCH_2 - CH - CH_2 - CH_3$   
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 $(d) CH_3 - CH_$ 

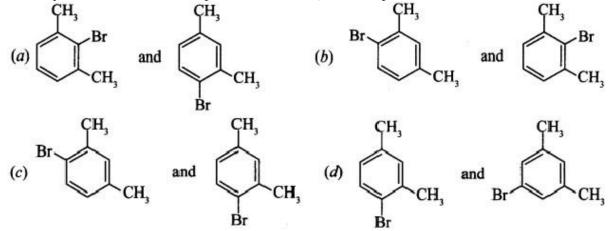
7. Which of the following is most reactive towards SN1 reaction?

- (a)  $C_6H_5C(CH_3)C_6H_5Br$
- (b) C<sub>6</sub>H<sub>5</sub>CH<sub>2</sub>Br
- (c) C<sub>6</sub>H<sub>5</sub>CH(C<sub>6</sub>H<sub>5</sub>)Br
- (d) C<sub>6</sub>H<sub>5</sub>CH(CH<sub>3</sub>)Br

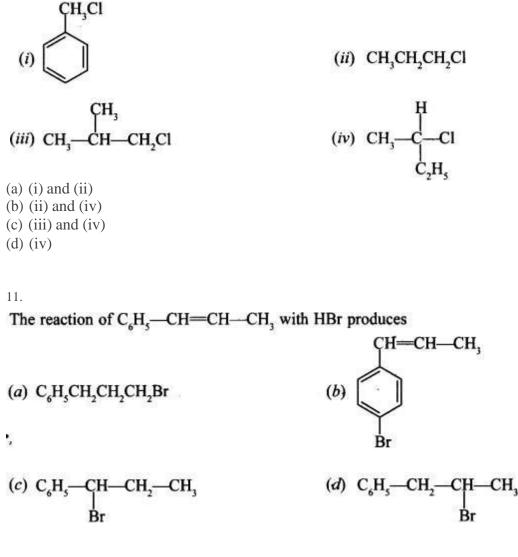
8. The correct order of increasing the reactivity of C—X bond towards nucleophile in following compounds



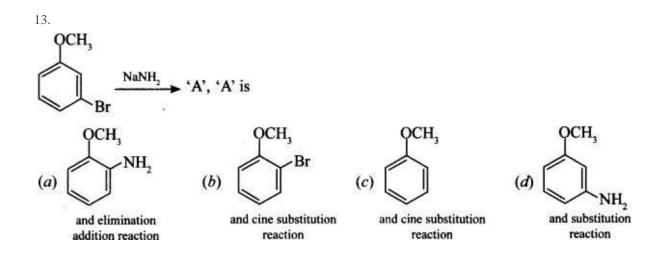
9. m-Xylene reacts with  $Br_2$  in presence of FeBr<sub>3</sub>, what are products formed







- 12.  $CH_3CH_2CH_2Br + NaCN \rightarrow CH_3CH_2CH_2CN + NaBr$ , will be fastest in
- (a) ethanol
- (b) methanol
- (c) N, N dimethyl formamide
- (d) Water



14. A dihalogen derivative 'X' of a hydrocarbon with three carbon atoms react with ale. KOH and produces hydrocarbon which forms red ppt. with ammonical  $Cu_2Cl_2$ . 'X' gives an aldehyde on reaction with aq. KOH. The compound 'X' is

- (a) 1, 3-Dichloropropane
- (b) 1, 2-Dichloropropane
- (c) 2, 2-Dichloropropane
- (d) 1, 1-Dichloropropane

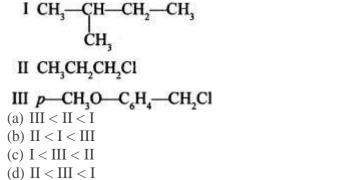
15. The synthesis of alkyl fluoride is best accomplished by

- (a) Finkelstein reaction
- (b) Swartz reaction
- (c) Free radical fluorination
- (d) Sandmeyers reaction

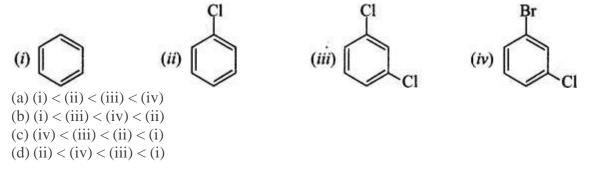
16. How many chiral compounds are possible on monochlorination of 2-methyl butane?

- (a) 2
- (b) 4
- (c) 6
- (d) 8

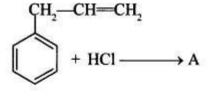
17. The increasing order of reactivity towards SN1 mechanism is

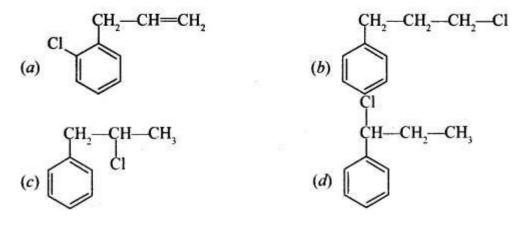


18. Arrange the following compounds in the increasing order of their densities.



19. What is 'A' in the following reaction?



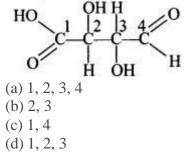


- 20. Which of the following alkyl halides will undergo SN1 reaction most readily?
- (a) (CH<sub>3</sub>)<sub>3</sub> C—F
- (b) (CH<sub>3</sub>)<sub>3</sub> C—Cl

(c) (CH<sub>3</sub>)<sub>3</sub> C—Br

(d) (CH<sub>3</sub>)<sub>3</sub> C—I

21. Which of the carbon atoms present in the molecule given below are asymmetric?



22. Which of the following compounds will give racemic mixture on nucleophilic substitution by OHion?

(a) 
$$CH_3$$
— $CH$ — $Br$  (b)  $CH_3$ — $C$ — $CH_3$  (c)  $CH_3$ — $CH$ — $CH_2Br$   
 $C_2H_5$   $C_2H_5$   $C_2H_5$ 

23. Which of the following compounds will be least reactive towards nucleophilic substitution by OH-ion?

1-Bromoethane, 1-Bromopropane, 1-Bromobutane, Bromobenzene

- (a) Bromobenzene
- (b) 1-Bromoethane
- (c) 1-Bromopropane
- (d) 1-Bromobutane
- 24. SN1 reaction of alkyl halides lead to
- (a) Retention of configuration
- (b) Racemisation
- (c) Inversion of configuration
- (d) None of these
- 25. p-dichlorobenzene has higher melting point than its o- and m- isomers because
- (a) p-dichlorobenzene is more polar than o- and m- isomer.
- (b) p-isomer has a symmetrical crystalline structure.
- (c) boiling point of p-isomer is more than o- and m-isomer.
- (d) All of these are correct reasons.
- 26. Chloropicrin is formed by the reaction of
- (a) steam on carbon tetrachloride.
- (b) nitric acid on chlorobenzene.

(c) chlorine on picric acid.

- (d) nitric acid on chloroform.
- 27. Fitting reaction can be used to prepare
- (a) Toluene
- (b) Acetophenone
- (c) Diphenyl
- (d) Chlorobenzene

28. Identify the end product (C) in the following sequence:

$C_2H_5OH \xrightarrow{SOCl_2}{Pyridine} A$	KCN (alc.)
	$B \xrightarrow{2H_2 O/H^+} C$
(a) $C_2H_5CH_2NH_2$	(b) $C_2H_5CONH_2$
(c) $C_2H_5COOH$	(d) $C_2H_5NH_2$ + HCOOH

29.

 $\begin{array}{c} CH_{3}CH_{2}CH_{2}Cl \xrightarrow{alc. KOH} B \xrightarrow{HBr} \\ C \xrightarrow{Na/ether} D \\ In the above reaction, the product D is \\ (a) Propane \end{array}$ 

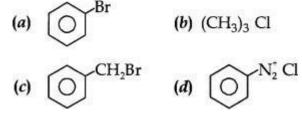
- (b) 2, 3-Dimethylbutane
- (c) Hexane
- (d) Allyl bromide

30. Identify X and Y in the following sequence  $C_2H_5 \text{ Br} \xrightarrow{X} \text{Product} \xrightarrow{Y} C_3H_7\text{NH}_2$ (a) X = KCN, Y = LiAlH4 (b) X = KCN, Y = H3O+ (c) X = CH3Cl, Y = AlCl3 HCl (d) X = CH3NH2, Y = HNO2

31. In the following sequence of reactions:

C<sub>2</sub>H<sub>5</sub>Br <u>AgCN</u> X <u>Reduction</u> Y; Y is (a) n-propylamine (b) isopropylamine (c) ethylamine (d) ethylmethylamine 32.

 $X \xrightarrow{\text{AgNO}_3} \text{Yellow or While ppt}$ Which of the following cannot be X?



33. Identify Z in the series  $CH_2 = CH_2 \xrightarrow{HBr} X \xrightarrow{aq. KOH} Y$ 

$$\xrightarrow{Na_2CO_3} Z$$

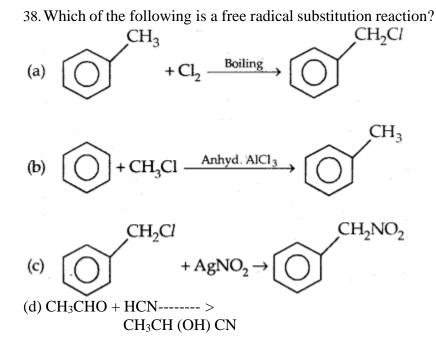
- (a)  $C_2H_5I^{--}$
- (b)  $C_2H_5OH$
- (c) CHI<sub>3</sub>
- (d) CH<sub>3</sub>CHO
- 34. The most reactive nucleophile among the following is
- (a) CH<sub>3</sub>O<sup>-</sup>
- (b)  $C_6H_5O^-$
- (c)  $(CH_3)_2 CHO^-$
- (d)  $(CH_3)_3 CO^-$
- 35. S<sub>N</sub>2 reaction of alkyl halides leads to
- (a) retention of configuration
- (b) racemisation
- (c) inversion of configuration
- (d) none of these.

36. Nucleophilicity order is correctly represented by

- (a)  $CH_3^- < \overline{N}H_2 < H\overline{O} < F^-$
- (b)  $CH_3^- \simeq N \overline{H}_2 > OH^- \simeq \overline{F}$
- (c)  $CH_3^- > \overline{N}H_2 > H\overline{O} > F^-$
- (d)  $NH_2^- > \overline{F} > HO^- > CH_3^-$

37. Which of the following are arranged in the decreasing order of dipole moment?

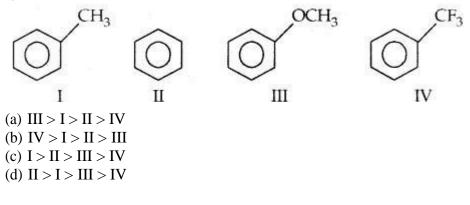
- (a)  $CH_3Cl$ ,  $CH_3Br$ ,  $CH_2F$
- (b) CH<sub>3</sub>Cl, CH<sub>3</sub>F, CH<sub>3</sub>Br
- (c) CH<sub>3</sub>Br, CH<sub>3</sub>Cl, CH<sub>3</sub>F
- (d)  $CH_3Br$ ,  $CH_3F$ ,  $CH_3Cl$



39. The reactivity order of halides for dehydrohalogenation is
(a) RF > RCl > RBr > RI
(b) RI > RBr > RCl > RF
(c) RI > RCl > RBr > RF
(d) RF > RI > RBr > RCl

40. The addition of HBr is easiest with
(a) CH<sub>2</sub> = CHCl
(b) ClCH = CHCl
(c) CH<sub>3</sub>-CH = CH<sub>2</sub>
(d) (CH<sub>3</sub>)<sub>2</sub>C = CH<sub>2</sub>

41. Among the following compounds, the decreasing order of reactivity towards electrophilic substitution is



42. Unpleasant smelling carbylamines are formed by heating alkali and chloroform with

(a) Any aliphatic amine

(b) Any aromatic amine

(c) Any amine

(d) Any primary amine

43. Which among MeX, R-CH<sub>2</sub>X, R<sub>2</sub>CHX, R<sub>3</sub>CX is most reactive towards  $S_N2$  reaction

(a) MeX

(b) RCH<sub>2</sub>X

(c) R<sub>3</sub>CHX

(d) R<sub>3</sub>CX

44. Number of stereoisomers of the compound 2-chloro-4- methylhex-2-ene is /are

(a) 1

(b) 2

- (c) 4
- (d) 16

45. In chlorination of benzene, the reactive species is

(a) Cl<sup>+</sup>

(b)  $Cl^{-}$ 

(c) Cl<sub>2</sub>

(d)  $Cl^{-2}$ 

46. An organic compound, on treatment with Br<sub>2</sub> in CCl<sub>4</sub> gives bromo derivative of an alkene. The compound will be
(a) CH<sub>3</sub>-CH = CH<sub>2</sub>
(b) CH<sub>3</sub>-CH = CH-CH<sub>3</sub>

(c)  $HC \equiv CH$ 

(d)  $H_2C = CH_2$ 

47. If one H is replaced by a Cl atom in C<sub>4</sub>H<sub>9</sub>Cl, the total number of structural isomers will be

(a) 4

(b) 5

(c) 7

(d) 10

48. Anti-Markovnikov addition of HBr is not observed in

(a) Propene

(b) 1-Butene

(c) But-2-ene

(d) Pent-2-ene

49. Among the following, the molecule with the highest dipole moment is

(a) CH<sub>3</sub>Cl

(b) CH<sub>2</sub>Cl<sub>2</sub>

(c) CHCl<sub>3</sub>

(d) CCl<sub>4</sub>

# 50. The number of isomers for the compound with molecular formula $C_2BrClFI$ is

(a) 3

(b) 4

(c) 5

(d) 6

# **ASSERTION REASON TYPE**

**Directions**: These questions consist of two statements, each printed as Assertion and Reason. While answering these questions, you are required to choose any one of the following four responses. (a) If both Assertion and Reason are correct and the Reason is a correct explanation of the Assertion.

(b) If both Assertion and Reason are correct but Reason is not a correct explanation of the Assertion.

- (c) If the Assertion is correct but Reason is incorrect.
- (d) If both the Assertion and Reason are incorrect.
- 51. Assertion: SN2 reaction of an optically active aryl halide with an aqueous solution of KOH always gives an alcohol with opposite sign of rotation.
   Beagen: SN2 reactions always present with inversion of configuration.

**Reason**: SN2 reactions always proceed with inversion of configuration.

- 52. **Assertion**: Alkylbenzene is not prepared by Friedel-Crafts alkylation of benzene. **Reason**: Alkyl halides are less reactive than acyl halides.
- 53. Assertion: Exposure of ultraviolet rays to human causes the skin cancer, disorder and disrupt the immune system.

Reason: Carbon tetrachloride is released into air, it rises to the atmosphere and depletes the ozone layer.

- 54. **Assertion**: CHCl<sub>3</sub> is stored in dark bottles. **Reason**: CHCl<sub>3</sub> is oxidised in dark.
- 55. Assertion: CCl4 is not a fire extinguisher. Reason: CCl4 is insoluble in water.
- 56. Assertion: Treatment of chloroethane with a saturated solution of AgCN gives ethyl isocyanide as a major product.

**Reason:** Cyanide ion (CN<sup>-</sup>) is an ambident nucleophile.

- 57. Assertion: KCN reacts with methyl chloride to give methyl isocyanide. **Reason:** CN<sup>-</sup> is an ambident nucleophile.
- 58. Assertion: tert-Butyl bromide undergoes Wurtz reaction to give 2, 2, 3,3-tetramethylbutane. Reason: In Wurtz reaction, alkyl halides react with sodium in dry ether to give hydrocarbon containing double the number of carbon atoms present in the halide.
- 59. Assertion: Presence of a nitro group at ortho or para position increases the reactivity of haloarenes towards nucleophilic substitution.
  - **Reason:** Nitro group, being an electron withdrawing group decreases the electron density over the benzene ring.
- 60. Assertion: Aryl iodides can be prepared by reaction of arenes with iodine in the presence of an oxidising agent

Reason: Oxidising agent oxidises L2 into HI.

#### Answer Key:

1. C	11. C	21. B	31. D	41. A	51. D
2. B	12. C	22. A	32. A	42. D	52. C
3. A	13. D	23. A	33. C	43.A	53. B
4. A	14. D	24. B	34. A	44. C	54. C
5. A	15. B	25. B	35. B	45. A	55. D
6. A	16. A	26. D	36. C	46. C	56. B
7. A	17. B	27. C	37. B	47. D	57. D
8. C	18. A	28. C	38. A	48. C	58. A
9. C	19. C	29. B	39. B	49. A	59. A
10. D	20. D	30. A	40. D	50. D	60. C

# **CHAPTER -7**

# ALCOHOLS, ETHERS AND PHENOLS

**Propene**,  $CH_3 - CH = CH_2$  can be converted to 1-propanol by oxidation. Which set of 1. reagents among the following is ideal to effect the conversion (a) Alkaline  $KMnO_4$  $B_2H_6$  and alkaline  $H_2O_2$ (b) (c)  $O_3/Z_n$  dust (d)  $OsO_4 / CH_4, Cl_2$  $A \xrightarrow{K_2C_{r_2}O_7} B \xrightarrow{CH_3M_{gI}} CH_3 \xrightarrow{CH_3} CH_3$ dil.  $H_2SO_4$   $H_2O$   $H_2O$   $H_3$  The reactant A is 2. (a) CH<sub>3</sub>CHOHCH<sub>3</sub> (b)  $CH_3COCH_3$ (c)  $C_2H_5OH$ (d) CH<sub>3</sub>COOH 3. Ortho-nitrophenol is steam volatile whereas para-nitrophenol is not. This is due to (a) Intramolecular hydrogen bonding present in ortho-nitrophenol (b) Intermolecular hydrogen bonding (c) Intramolecular hydrogen bonding present in para-nitrophenol (d) None of these 4. The strongest acid among the following aromatic compounds is (a) ortho-nitrophenol (b) para-chlorophenol (c) para-nitrophenol (d) meta-nitrophenol The alcohol that produces turbidity immediately with 5.  $ZnCl_2$  + conc. *HCl* at room temperature (a) 1-hydroxybutane 2-hydroxybutane (b) (c) 2-hydroxy-2-methylpropane (d) 1-hydroxy-2methylpropane The compound which gives the most stable carbonium ion on dehydration is 6. (a)  $CH_3 - CH - CH_2OH$  $CH_3$  $CH_3$ (b)  $CH_3 - C - OH$  $CH_3$ (c)  $CH_3 - CH_2 - CH_2 - CH_2OH$ 

(d)  $CH_3 - CH - CH_2 - CH_3$ 

7.	Phenol is treated with bromine water and sh during the process is	aken well.	The white precipitate formed
	(a) <i>m</i> -bromophenol	(b)	2, 4-dibromophenol
	(c) 2, 4, 6-tribromophenol	(d)	A mixture of o- and p-
	bromophenols		
8.	In the sequence of the following reactions		
	$CH_{3}CH_{2}OH \xrightarrow{[O]} X \xrightarrow{[O]} CH_{3}COOH$ ChromicAcid ChromicAcid		
	X is		
	(a) $CH_3COCH_3$	(b)	CH <sub>3</sub> CHO
	(c) $CH_3OCH_3$	(d)	CH <sub>3</sub> CH <sub>2</sub> COOH
9.	At low temperature phenol reacts with $Br_2$ in	CS <sub>2</sub> to form	n
	(a) <i>m</i> -bromophenol	(b)	o-and p-bromophenol
	(c) <i>p</i> -bromophenol	(d)	2,4,6-tribromophenol
10.	What amount of bromine will be required	to conver	t 2 g of phenol into 2, 4, 6-
	tribromophenol		
	(a) 4.00	(b)	6.00
	(c) 10.22	(d)	20.44
11.	Phenol reacts with CO <sub>2</sub> in presence of aqueo	us alkali a	and forms a product which on
	hydrolysis gives		
	(a) Salicylaldehyde	(b)	Salicylic acid
	(c) Benzaldehyde	(d)	Benzoic acid
12.	In presence of NaOH, phenol react with CHCl	<sub>3</sub> to form	o-hydroxy benzaldehyde. This
	reaction is called		
	(a) Riemer-Tiemann's reaction	(b)	Sandmeyer's reaction
	(c) Hoffmann's degradation reaction	(d)	Gattermann's aldehyde
synt	hesis		
13.	Methyl-terbutyl ether on heating with HI of o	ne molar c	oncentration gives
	(a) $CH_3I + (CH_3)_3COH$	(b)	$CH_3OH + (CH_3)_3CI$
	(c) $CH_3I + (CH_3)_3Cl$	(d)	None of the above
14.	Amongst the following, HBr reacts fastest with	h	
	(a) Propane-1-ol	(b)	Propane-2-ol
	(c)2-methyl propane-1-ol	(d)	2-methyl propane-2-ol
15.	The best reagent to convert pent-3-en-2-ol into	. ,	
	(a) Acidic permanganate	(b)	Acidic dichromate
	(c) Chromic anhydride in glacial acetic acid	(d)	Pyridinium chloro-chromate

**Read the assertion and reason carefully to mark the correct option out of the options given below :** 

- (a) If both assertion and reason are true and the reason is the correct explanation of the assertion.
- (b) If both assertion and reason are true but reason is not the correct explanation of the assertion.
- (c) If assertion is true but reason is false.
- (d) If assertion is false but reason is true.

**16. Assertion** : Phenol is more reactive than benzene towards electrophilic substitution reaction

Reason : In the case of phenol, the intermediate carbocation is more resonance stabilized

- **17. Assertion** : Lucas reagent is a mixture of anhydrous  $Z_nCl_2$  and concentrate HCl**Reason** : Primary alcohol produce ppt. with Lucas reagents
- **18.Assertion** : Phenol undergo Kolbe reaction, ethanol does not. **Reason**: Phenoxide ion is more basic than ethoxide ion
- **19.Assertion** : Alcohol and phenol can be distinguished by sodium hydroxide. **Reason** : Phenol is acidic while alcohol is neutral.
- **20.** Assertion : The major products formed by heating  $C_6H_5CH_2OCH_3$  with *HI* are  $C_6H_5CH, I$  and  $CH_3OH$ .

**Reason** :Benzyl cation is more stable than methyl cation.

- **21. Assertion** : Acid catalysed dehydration of *t*-butanol is slower than *n*-butanol. **Reason** : Dehydration involves formation of the protonated alcohol, *ROH*<sup>+</sup><sub>2</sub>.
- 22. Assertion : 4-nitrophenol is more acidic than 2, 4, 6-trinitrophenol.Reason : Phenol is a weaker acid than carbonic acid.
- **23. Assertion** : Phenol reacts with acyl halides in presence of pyridine to form phenyl acetate. **Reason** : Benzoylation of phenol is carried out in the presence of *NH*<sub>4</sub>*OH* .

**24. Assertion** : Both symmetrical and unsymmetrical ethers can be prepared by Williamson's synthesis.

**Reason** : Williamson's synthesis is an example of nucleophilic substitution reaction.

**25.** Assertion : *t*-Butyl methyl ether is not prepared by the reaction of *t*-butyl bromide with sodium methoxide.

**Reason** : Sodium methoxide is a strong nucleophile.

#### Read the passage given below and answer the following questions:

An organic compound (A) having molecular formula C<sub>6</sub>H<sub>6</sub>O gives a characteristic colour with

aqueous FeCl<sub>3</sub> solution. (A) on treatment with  $CO_2$  and NaOH at 400 K under pressure gives (B), which on acidification gives a compound (C). The compound (C) reacts with acetyl chloride to give (D) which is a popular pain killer.

# The following questions are multiple choice questions. Choose the most appropriate answer:

### (26) Compound (A) is

(a) 2-hexanol	(b) dimethyl ether	r (c) phenol	(d) 2-me	ethyl pentanol.		
(27) Number	of carbon atoms in	n compound (D) is				
(a) 7	(b) 6	(c) 8	(d) 9			
<ul> <li>(28) The conversion of compound (A) to (C) is known as</li> <li>(a) Reimer- Tiemann reaction (b) Kolbe's reaction (c) Schimdt reaction (d) Swarts reaction</li> </ul>						
<ul> <li>(29) Compound (A) on heating with compound (C) in presence of POCl<sub>3</sub> gives a compound</li> <li>(D) which is used</li> </ul>						

(a) in perfumery as a flavouring	(b) as an	(c) as an	(d) as an intestinal
agent	antipyretic	analgesic	antiseptic.

# Read the passage given below and answer the following questions:

Both alcohols and phenols are acidic in nature, but phenols are more acidic than alcohols. Acidic strength of alcohols mainly depends upon the inductive effect. Acidic strength of phenols depends upon a combination of both inductive effect and resonance effects of the substituent and its position on the benzene ring. Electron withdrawing groups increases the acidic strength of phenols whereas electron donating groups decreases the acidic strength of phenols. Phenol is a weaker acid than carboxylic acid.

# The following questions are multiple choice questions. Choose the most appropriate answer:

### (30) Phenols are highly acidic as compare to alcohols due to

(a) the higher molecular mass of phenols (b) the stronger hydrogen bonds in phenols

(c) alkoxide ion is a strong conjugate base (d)phenoxide ion is resonance stabilised.

# (31) The correct decreasing order of $pK_a$ value is

# 32) The compound that does not liberate CO<sub>2</sub>, on treatment with aqueous sodium bicarbonate solution is

(a) benzoic acid (b) benzenesulphonic acid (c) salicylic acid <sup>(d)</sup> carbolic acid.

# (33) Most acidic amongst the following is

(a) o-nitro phenol (b) p-nitro phenol (c) m-nitro phenol (d) phenol

# Read the passage given below and answer the following questions:

Although chlorobenzene is inert to nucleophilic substitution, however it gives quantitative yield of phenol when heated with aq. NaOH at high temperature and under high pressure. As far as electrophilic substitution in phenol is concerned the - OH group is an activating group, hence, its presence enhances the electrophilic substitution at o- and p-positions.

The following questions are multiple choice questions. Choose the most appropriate answer:

(34)	Conversion	of chlorobenzene	into pheno	ol involves
	Conversion	or emoropenzene	meo prient	

(a) modified $S_N 1$ mechanism	(b) modified $S_N 2$ mechanism
(c) both (a) and (b)	(d) elimination-addition mechanism.

# (35) Phenol undergoes electrophilic substitution more readily than benzene because

(a) the intermediate carbo cation is a resonance hybrid of more resonating structures than that from benzene

(b) the intermediate is more stable as it has positive charge on oxygen, which can be better accommodated than on carbon

(c) in one of the canonical structures, every atom (except hydrogen) has complete octet (d) the -OH group is o, p-directing which like all other o, p-directing group, is activating

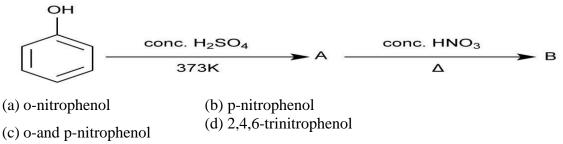
# (36) Phenol on treatment with excess of conc. HNO<sub>3</sub> gives

(a) o-nitrophenol (b) p-nitrophenol

(c) o-and p-nitrophenol

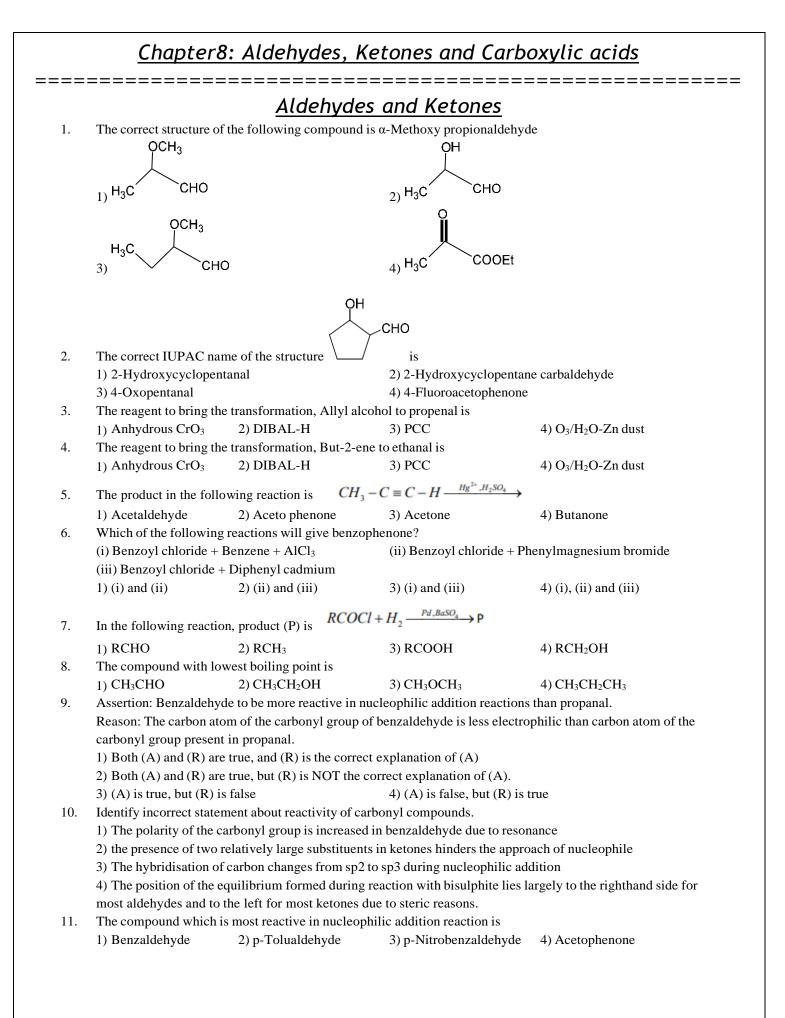
(d) 2,4,6-trinitrophenol

# (37) The major product (B) of the following reaction is



# **ANSWERS**:

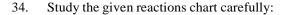
(1) b	(2) a	(3) a	(4) c	(5) c	(6) b
(7) c	(8) b	(9) b	(10) c	(11) b	(12) a
(13) b	(14) d	(15) d	(16) a	(17) c	(18) c
(19) a	(20) a	(21) d	(22) d	(23) c	(24) b
(25) b	(26) c	(27) d	(28) b	(29) d	(30) d
(31) a	(32) d	(33) b	(34) d	(35) a	(36) d
(37) d					

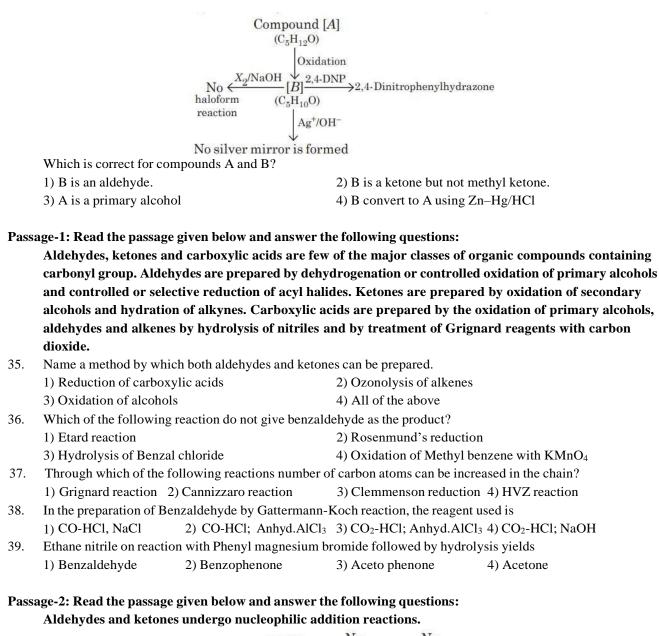


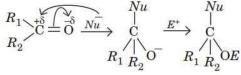
12.	Arrange the following compounds in i	ncreasing order of their reacti	vity in nucleophilic addition reactions.
	Ethanal, Propanal, Propanone, Butano	_	
	1) Butanone < Propanole < Propanal		< Butanone < Ethanal > Propanal
	3) Propanal < Ethanal < Propanone < 2	Butanone 4) Ethanal < P	ropanal < Propanone < Butanone
	[Hint: Electron deficiency at C=O car	bon (electrophilicity) and Ste	ric factors]
13.	Which among the following is most re	active to give nucleophilic ad	dition?
	1) FCH <sub>2</sub> CHO 2) ClCH <sub>2</sub> CH	IO 3) BrCH <sub>2</sub> CHC	4) ICH <sub>2</sub> CHO
	[Hint: Electron deficiency at C=O carb	oon (electrophilicity) and Elec	tronegativity of halogen]
14.	Formaldehyde reacts with methyl mag	nesium bromide followed by	hydrolysis to form
	1) Methanol2) Ethanol	3) Propanol	4) Butanol
15.	Predict the product of the following re	action:	$H_3 \xrightarrow{(i) \mathrm{NH}_2 - NH_2; (ii) KOH/Glycol, \Delta} ?$
	1) CH <sub>3</sub> CH <sub>2</sub> CH <sub>3</sub> 2) CH <sub>3</sub> CHO	HCH <sub>3</sub> HCH <sub>3</sub> $CH_3 - CO - C$	$\Pi_3 \longrightarrow \ell$
	$3) CH_3 CH_2 CHO \qquad 4) CH_3 CON$	HCH <sub>3</sub> [Hint: Wolf-Kishner re	duction]
16.	The reaction of carbonyl compounds w	vith ammonia derivatives give	es N-substituted derivatives. Find the
	incorrect match of product.		
	1) Ammonia-Imine		nine-Hydrazone
	3) 2,4-DNP-2,4-DN phenyl hydrazone	e 4) Semi carbaz	zide-semi carbazone
17.	Which of the following compounds wa	-	with alkaline KMnO <sub>4</sub> solution?
	1) Butan-1-ol2) Butan-2-o	, ·	4) But-2-ene
18.	Assertion (A): Oxidation of ketones is	-	
	Reason (R): C–C bond of ketones is st	•	•
	1) Both (A) and (R) are true, and (R) i	•	
	2) Both (A) and (R) are true, but (R) is	=	
10	3) (A) is true, but (R) is false		but (R) is true
19.	. In Clemmensen reduction, carbonyl		
	<ol> <li>zinc amalgam + HCl</li> <li>Phenyl hydrazine, glycol</li> </ol>	2) sodium ama 4) Dil. NaOH	iigaiii + HCi
20.	Hydrocarbons are formed when aldehy	·	vith Hydrazine KOH and Glycol The
20.	reaction is called	, act and notonos are reacted ,	
	1) Cannizzaro reaction 2) Clemmen	sen reduction 3) Rosenmund	reduction 4) Wolf-Kishner reduction
21.	Which of the following reagents are no	ot correctly matched with the	reaction?
	1) CH <sub>3</sub> CH=CHCHO $\rightarrow$ CH <sub>3</sub> CH=CHCO	OOH: Ammonical AgNO <sub>3</sub>	
	2) CH <sub>3</sub> CH=CHCHO→ CH <sub>3</sub> CH=CHCH	I <sub>2</sub> OH: H <sub>2</sub> /Pt	
	3) R—COOH $\rightarrow$ R—CH <sub>2</sub> OH: NaBH <sub>4</sub>		
	4) $CH_3CH_2COCl \rightarrow CH_3CH_2CHO: H_2,$		
22.	Assertion (A): Aromatic aldehydes an		
	Reason (R): Aromatic aldehydes and t		
	1) Both (A) and (R) are true, and (R) i	•	
	2) Both (A) and (R) are true, but (R) is 2) (A) is $(R) = \frac{1}{2} 1$	=	
22	3) (A) is true, but (R) is false		but (R) is true
23.	Which of the following will not give a 1) Phenyl acetaldehyde 2) 2-Methyl		(d) 1 Dhanylpropona
24.	The reagent used to distinguish acetal		de 4) 1-Phenylpropanone
24.	1) Sodium hydrogen sulphite	2) Phenyl hydr	azine
	3) Fehling's solution	3) Tollen's rea	
25.	Assertion (A): Aldehydes and ketones		-
	Reason (R): Both aldehydes and keton		
	1) Both (A) and (R) are true, and (R) i		A)
	2) Both (A) and (R) are true, but (R) is	=	
		1	

	3) (A) is true, but (R) is false	4) (A) is false, but	t (R) i	s true	
26.	Compounds A and C in the following reaction are				
	$CH_{3}CHO \xrightarrow{(i)CH_{3}MgBr,(ii)H_{2}O} (A) \xrightarrow{H_{2}SO_{4},\Lambda} (B) -$	Hydroboration,oxidaton	(C)		
	1)Identical 2) Position isomers			4) Optical isomers	
	[Hint: A and C are secondary and primary alcohols	·		/ I	
27.	What are the correct steps to convert acetaldehyde to				
	1) CH <sub>3</sub> MgBr, H <sub>2</sub> O, Oxidation	2) Oxidation, Ca(C	OH)2,	Heat	
	3) Reduction, KCN, Hydrolysis	4) Oxidation, $C_2H$			
28.	Alkene (X) ( $C_5H_{10}$ ) on ozonolysis gives a mixture of	two compounds (	Y) and	l (Z). Compound (Y) gives positiv	ve .
	Fehling's test and iodoform test. Compound (Z) doe	s not give Fehling's	s test	but give iodoform test. Compound	ls
	(X), (Y) and (Z) are				
	1) X=C <sub>6</sub> H <sub>5</sub> COCH <sub>3</sub> Y=CH <sub>3</sub> CHO Z=CH <sub>3</sub> COCH <sub>3</sub>	2) X= CH <sub>3</sub> -CH=C	CH(CH	(3) <sub>2</sub> Y=CH <sub>3</sub> CHO Z=CH <sub>3</sub> COCH <sub>3</sub>	
	3) X=CH <sub>3</sub> CH <sub>2</sub> CH=CH <sub>2</sub> Y=CH <sub>3</sub> CH <sub>2</sub> CHO Z=HCHO	4) X=CH <sub>3</sub> CH=CH	I-CH <sub>3</sub>	Y=CH <sub>3</sub> CHO Z=CH <sub>3</sub> CHO	
29.	A compound (X) with a molecular formula $C_5H_{10}O_5$	gives a positive 2,4	-DNP	test but a negative Tollen's test.	
	On oxidation it gives a carboxylic acid (Y) with a mo	olecular formula C <sub>3</sub>	$_{3}H_{6}O_{2}$	Potassium salt of (Y) undergoes	
	Kolbe's reaction and gives a hydrocarbon (Z). (X),	Y) and (Z) respect	ively	are	
	1) pentan-3-one, propanoic acid, butane	2) pentanal, pentan	noic a	cid, octane	
	3) 2-methylbutanone, butanoic acid, hexane	• •		one, propanoic acid, hexane	
30.	An organic compound (X) with molecular formula	• •			
	undergoes Cannizzaro reaction and on vigorous oxid	ation it gives 1,4-be	enzen	edicarboxylic acid. Compound (X	)
	is				
	1) benzaldehyde 2) o-methylbenzaldehyde		•	•	
31.	Compound (X) with molecular formula $C_3H_8O$ is tre		-	-	t
	(Y) with molecular formula $C_3H_6O_{-}(Y)$ does not for				
	AgNO <sub>3</sub> . (Y) when treated with an aqueous solution $f(T)$	of NH <sub>2</sub> CONHNH <sub>2</sub>	$_2$ . HCl	and sodium acetate to give a	
	product (Z). The structure of (Z) is		CON	T	
	1) $CH_3CH_2CH=NNHCONH_2$ 3) $(CH_3)_2C=NCONHNH_2$	2) (CH <sub>3</sub> ) <sub>2</sub> C=NNH 4) CH <sub>3</sub> CH <sub>2</sub> CH=N			
32.	Aldehydes and ketones are isomers as they have sat				
52.	these functional groups can be distinguished by var	-		• •	
	two isomers P and Q which undergo ozonolysis to g				
	$P \xrightarrow{(i) O_3/CH_2CI_2} R+HCHO Q \xrightarrow{(i) O_3/CH_2O} R+IO R+IO R+IO R+IO R+IO R+IO R+IO R+IO$	<u>, , , , , , , , , , , , , , , , , , , </u>	ICHO	)	
	- ,	-			
	Which of the given options cannot be correct for P, o	-			
	I. If P is 4-vinyl toluene then R gives Cannizzaro rea			eaction.	
	II. If Q is 4-vinyl toluene then S gives haloform reac				
	III. If Q is 2-phenylpropene then S gives haloform re				
	IV. If P is 2-phenylpropene then R gives both Cannix 1) I and II only 2) I and III only		react		
33.	1) I and II only2) I and III onlyMatch the column I with column II and mark the app	3) II and III only		4) II and IV only	
55. Г	11				
H			G	COLUMN-II Kolbe's reaction	
Ľ	(A) $RCOCH_3 \xrightarrow{Zn-Hg;HCl} RCH_2CH_3$		(i)	Kolbe s reaction	
	(B) $2C_6H_5CHO \xrightarrow{NaOH} C_6H_5CH_2OH + C_6H_5CH_5CH_2OH + C_6H_5CH_5CH_5CH_5OH + C_6H_5CH_5CH_5OH + C_6H_5CH_5CH_5OH + C_6H_5CH_5OH + C_6H_5CH_5OH + C_6H_5CH_5OH + C_6H_5CH_5OH + C_6H_5OH $	6H5COONa	(ii)	Clemmensen	
L			(115	reduction	
(	$(C)  C_6H_6 + CH_3COCl \xrightarrow{Anhyd \ AlCl_3} C_6H_5C$	OCH <sub>3</sub>	(iii)	Friedel-Craft's	
			0.0	reaction	
(	(D) $C_6H_5OH + CO_2 + NaOH \longrightarrow HOC_6H$	I <sub>4</sub> COONa	(iv)	Cannizzaro reaction	
_	-				

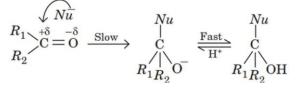
 $1) (A) \rightarrow (ii), (B) \rightarrow (iv), (C) \rightarrow (iii), (D) \rightarrow (i)$  $2) (A) \rightarrow (i), (B) \rightarrow (iii), (C) \rightarrow (ii), (D) \rightarrow (iv)$  $3) (A) \rightarrow (iii), (B) \rightarrow (ii), (C) \rightarrow (i), (D) \rightarrow (iv)$  $4) (A) \rightarrow (iv), (B) \rightarrow (i), (C) \rightarrow (ii), (D) \rightarrow (iii)$ 







Carbonyl carbon is electron deficient hence acts as an electrophile. Nucleophile attacks on the electrophilic carbon atom of the carbonyl group from a direction perpendicular to the plane of the molecule.



	formed as intermediate. This intermediate captures a proton from the reaction medium to give the neutral product. Aldehydes are generally more reactive than ketones in nucleophilic addition reactions.
40	Aldehydes other than formaldehyde react with Grignard's reagent to give addition products which on hydrolysi
40.	
	give
41.	1) tertiary alcohols       2) secondary alcohols       3) primary alcohols       4) Alkanes         Which of the following compounds will up dense Compigance resetion?
41.	Which of the following compounds will undergo Cannizzaro reaction?1) CH <sub>3</sub> CHO2) CH <sub>3</sub> COCH <sub>3</sub> 3) C <sub>6</sub> H <sub>5</sub> CHO4) C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> CHO
40	1) $CH_3CHO$ 2) $CH_3COCH_3$ 3) $C_6H_5CHO$ 4) $C_6H_5CH_2CHO$ Propanal on treatment with dilute sodium hydroxide gives
42.	1) CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CHO 2) CH <sub>3</sub> CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CHO
43.	3) CH <sub>3</sub> CH <sub>2</sub> CH(OH)CH(CH <sub>3</sub> ) CHO 4) CH <sub>3</sub> CH <sub>2</sub> COOH Various products formed on oxidation of 2,5-dimethylhexan-3-one are
45.	
	[Hint: When 2,5–dimethylhexan–3–one undergoes oxidation according to Popoff's rule, the products obtained
4.4	are 2-methylpropanoic acid, acetic acid and $CO_2$ .]
44.	Assertion: Benzaldehyde is more reactive than ethanal towards nucleophilic attack.
	Reason: The overall effect of $-I$ and $+R$ effect of phenyl group decreases the electron density on the carbon store of $C=0$ group in hangeldahuda
	atom of C=O group in benzaldehyde. 1) Both (A) and (B) are true, and (B) is the correct explanation of (A)
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A). 2) (A) is the set of
<b>D</b>	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true
Pass	age-3: Read the following passage and answer the questions given below:
	Test for Aldehydes and Ketones:
	(i) Both give iodoform test when one a-hydrogen is present.
	(ii) Fehling's test: Aliphatic aldehydes reduce the Fehling's solution to red cuprous oxide.
	$R - CHO + 2CuO + 5OH^{-} \rightarrow RCOOH + Cu_2O \downarrow + 3H_2O$ (Red Ppt.)
	Aromatic aldehydes do not respond to this test.
	(iii)Ketones are not oxidised by Tollen's reagent. Aldehydes form silver mirror with ammonical silver
	nitrate (Tollen's reagent) solution.
	$R - CHO + 2[Ag(NH_3)_2]^+ \xrightarrow{OH^-} RCOOH + 2Ag_{(Silver mirror)} \downarrow + H_2O + 4NH_3$
45.	Which of the following statements is incorrect?
	1) FeCl <sub>3</sub> is used in the detection of phenols.
	<ul><li>2) Fehling solution is used in the detection of glucose.</li></ul>
	3) Tollens' reagent is used in the detection of unsaturation.
	4) NaHSO <sub>3</sub> is used in the detection of carbonyl compounds.
46.	Which of the following compounds will give a coloured crystalline compound with 2,4-DNP?
40.	1) $CH_3COCl$ 2) $CH_3COOC_2H_5$ 3) $CH_3COCH_3$ 4) $CH_3CONH_2$
47.	How will you distinguish between aldehydes and ketones?
<b>ч</b> /.	1) Fehling's test       2) Lucas test       3) Iodoform test       4) Hinsberg reagent
48.	To differentiate between pentan-2-one and pentan-3-one a test is carried out. Which of the following is the
40.	correct answer?
	1) Pentan-2-one will give silver mirror test       2) Pentan-2-one will give iodoform test.         2) Pentan-2-one will give iodoform test.       4) Pentan-2-one will give Fahling's test
40	3) Pentan-3-one will give iodoform test 4) Pentan-3-one will give Fehling's test
49.	Assertion: Methyl Ketones can be converted into acids by haloform reaction.
	Reason: Addition of Grignard reagents to HCHO followed by hydrolysis gives Primary Alcohols.
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
	-
	<ul> <li>2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).</li> <li>3) (A) is true, but (R) is false</li> </ul>

4) (A) is false, but (R) is true

#### **SUBJECTIVE QUESTIONS:**

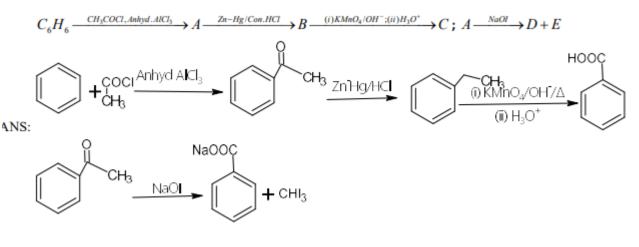
50. In industry aldehydes are being prepared by controlled oxidation of primary alcohol using acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> or aqueous or alkaline KMnO<sub>4</sub> as oxidant. Mohan suggested the owner of factory to use Collin's reagent instead of acidic potassium dichromate. The yield of factory increased sharply. Now answer the following questions:

(i) What is Collin's reagent?

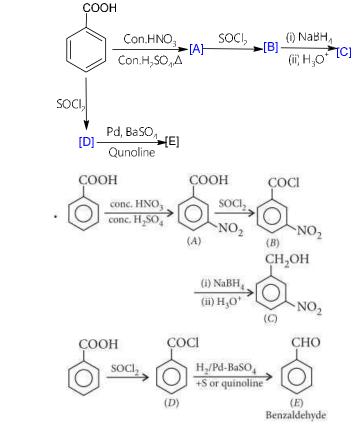
#### (ii) What are the advantages of using Collin's reagent over conventional oxidising agent?

ANS: (i) Collin's reagent is a mixture of pyridine (C<sub>5</sub>H<sub>5</sub>N) and CrO<sub>3</sub> in dichloromethane (CH<sub>2</sub>Cl<sub>2</sub>).
(ii) Collin's reagent is a mild oxidant. It oxidises 1°-alcohols to aldehydes and 2°-alcohols are oxidised to ketones. In case of using acidic K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> as oxidant, the aldehydes and ketones formed by the oxidation of alcohols undergo oxidation to give carboxylic acids

51. Write the structures of A, B, C, D and E in the following reactions:



52. Identify A to E in the following reactions:



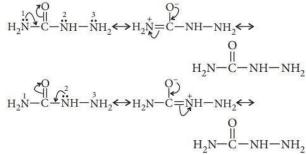
ANS:

53. Give a plausible explanation for each one of the following:

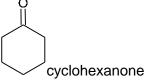
(i) There are two –NH<sub>2</sub> groups in semicarbazide. However, only one such group is involved in the formation of semicarbazones.

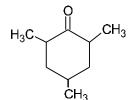
#### (ii) Cyclohexanone forms cyanohydrin in good yield but 2,4,6-trimethylcyclo-hexanone does not.

ANS: (i) Semicarbazide has the following resonance structures arising due to the electron withdrawing nature of the O atom. Lone pairs of N-1 and N-2 are involved in conjugation with C O group while that of N-3 is not involved in resonance thus, it is involved in the formation of semicarbazone.



(ii) Formation of cyanohydrin involves the nucleophilic attack of cyanide ions ( $CN^{-}$ ) at the carbonyl carbon. In cyclohexanone, reaction proceeds but in 2,4,6-trimethylcyclohexanone, the methyl groups cause steric hindrance and yields are poor.





2,4,6-trimethylcyclohexan-1-one

- 54. An organic compound with molecular formula C<sub>9</sub>H<sub>10</sub>O forms 2,4-DNP derivative, reduces Tollens' reagent and undergoes Cannizzaro reaction. On vigorous oxidation it gives 1,2-benzene-dicarboxylic acid. Identify the compound.
- ANS: The compound forms 2,4-DNP derivative. It shows that it is a carbonyl compound. Further it reduces Tollens' reagent which shows that it contains aldehydic group. It undergoes Cannizzaro reaction indicating that aldehyde group is without any a-hydrogen. On vigorous oxidation, it gives 1,2-benzenedicarboxylic acid which shows that there are two carbon residues on benzene ring. Since the molecular formula is  $C_9H_{10}O$ , it fits into the structure, 2-ethylbenzaldehyde.



55. Identify A, B and C in the following sequence of reactions:  $CH_3CHO \xrightarrow{(i)C_2H_3MgCl,(ii)H_2O} [A] \xrightarrow{Con.H_2SO_4} [B] \xrightarrow{HBr+Peroxide} [C]$ 

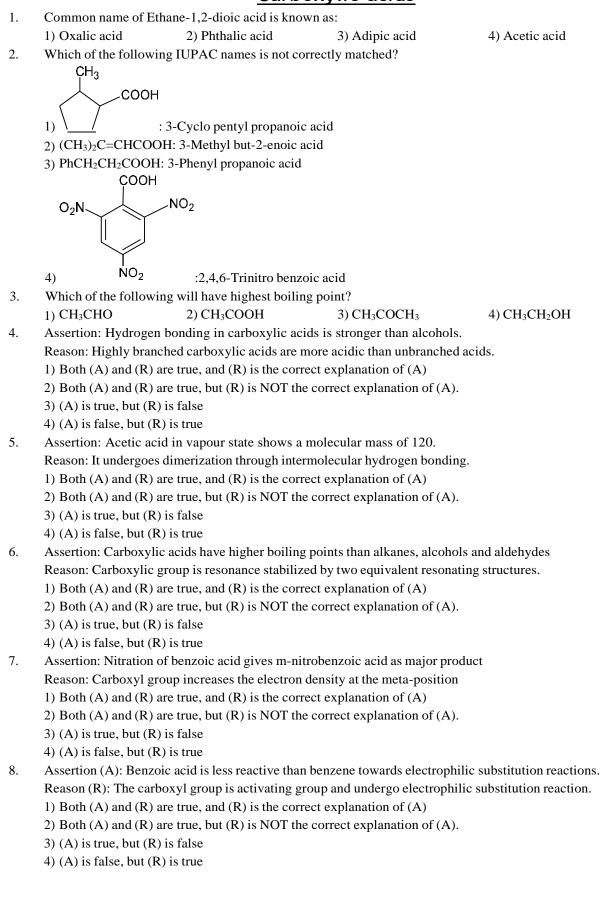
ANS:  $CH_3CHO \xrightarrow{C_2H_3MgCl} [CH_3CH(OMgCl)C_2H_5] \xrightarrow{H_2O} CH_3 - CH(OH) - C_2H_5 + Mg(OH)Cl$ 

$$CH_{3} - CH(OH) - C_{2}H_{5} \xrightarrow{Con.H_{2}SO_{4}} CH_{3} - CH = CH - CH_{3} \xrightarrow{HBr+Peroxide} CH_{3} - CH_{2} - CH(Br) - CH_{3} \xrightarrow{(Bur-2-ene)} CH_{3} - CH_{2} - CH(Br) - CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} - CH_{2} - CH(Br) - CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} - CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} - CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} - CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} - CH_{3} \xrightarrow{(Con.H_{2}SO_{4})} CH_{3} \xrightarrow{(Con.H_{2}SO_{$$

# 56. Explain: The reaction of carbonyl compound with pure HCN is very slow and becomes fast in presence of a base

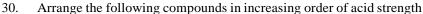
ANS: With pure HCN reaction occurs very slowly because it is a weak nucleophile. With base it produces CN<sup>-</sup> ion which is a strong nucleophile and readily adds to the carbonyl compound

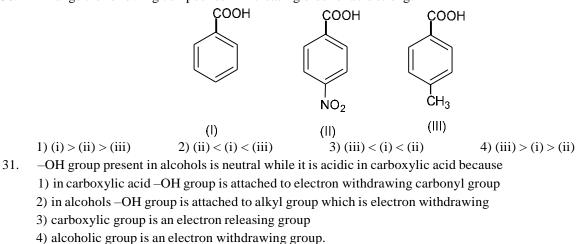
# Carboxylic acids



9.	Assertion (A): Aromatic carboxylic groups do not une	•	
	Reason (R): Carboxyl group is deactivating and the ca	atalyst aluminium chloride	gets bonded to the carboxyl
	group.		
	1) Both (A) and (R) are true, and (R) is the correct ex	planation of (A)	
	2) Both (A) and (R) are true, but (R) is NOT the corre	ect explanation of (A).	
	3) (A) is true, but (R) is false		
	4) (A) is false, but (R) is true		
10.	Carboxylic acids do not undergo Friedel Craft's react	ion because	
	1) —COOH group is meta directing	2) —COOH group is resor	nance stabilised
	3) carboxyl group is deactivating and gets bonded to 1	Friedel Craft's catalyst	
	4) all of above.		
11.	The carboxylic acid that does not undergo HVZ react	ion is:	
	1) CH <sub>3</sub> COOH 2) (CH <sub>3</sub> ) <sub>2</sub> CHCOOH	3) CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> COOH	I 4) (CH <sub>3</sub> ) <sub>3</sub> CCOOH
12.	Which of the following will undergo HVZ reaction?		
	1) Benzoic acid 2) Ethanoic acid	3) Methanoic acid	4) 2,2-Dimethylpropanoic acid
13.	Which of the following acids does not form anhydrid	e?	
		3) Propionic acid	4) n-butyric acid
14.	The reaction in which the aqueous solution of sodium	salt of carboxylic acids or	-
	•	2) Kolbe's electrolysis dec	
	•	4) Reduction of carboxylic	-
15.	Which of the following statements is correct regardin	•	
		2) It is a weaker acid than	
		4) Its calcium salt on heati	
16.	What happens when a carboxylic acid is treated with		-
	•••	2) Primary alcohol is form	
	•	4) Grignard reagent is form	
17.	The final product in the given reaction is: $CH_3COC$ 1) CH <sub>3</sub> CH <sub>2</sub> CN 2) CH <sub>3</sub> CH <sub>2</sub> CONH <sub>2</sub>	$\begin{array}{c} H_3 \longrightarrow [A] \\ \hline 3) \text{ CH}_3 \text{CONH}_2 \end{array}$	4) CH <sub>3</sub> COONH <sub>4</sub>
	$(A) + C_2 H_5 OH \rightarrow (B) + (C); (C) + HOH \xrightarrow{H^+}$		
	$(B) + Ca(OH)_2 \rightarrow Ca  salt + H_2O;  Ca  Salt - \frac{dyd}{dyd}$		
18.	$(B) + Ca(OII)_2 \rightarrow Ca sail + H_2O$ ; Ca sail — the above set of reactions are		(A), (B), (C) and (D) in
	(A) (B) (C)	(D)	
	$\frac{(1)}{1}  (CH_3CO)_2O  CH_3COOH  CH_3COOC_2H_5$		
	2 CH <sub>3</sub> COCl HCOOH CH <sub>3</sub> COOCH <sub>3</sub>	CH <sub>3</sub> OH	
	3 CH <sub>3</sub> COOH CH <sub>3</sub> OH CH <sub>3</sub> COOCH <sub>3</sub>	CH <sub>3</sub> OH	
	4 CH <sub>3</sub> NH <sub>2</sub> CH <sub>3</sub> COOH CH <sub>3</sub> COOCH <sub>3</sub>	CH <sub>3</sub> OH	
19.	Which of the following is the strongest acid?		
		3) Methyl alcohol	4) Water
20.	Assertion: m-Chlorobenzoic acid is a stronger acid th	-	
	Reason: In m-chlorobenzoic acid both – I-effect and -	R-effect of Cl operate but	in p-chlorobenzoic acid only
	+R-effect of Cl operates.		
	1) Both (A) and (R) are true, and (R) is the correct ex	•	
	2) Both (A) and (R) are true, but (R) is NOT the corre	-	
		4) (A) is false, but (R) is tr	ue
21.	Assertion: Formic acid is a stronger acid than benzoi		
	Reason: pKa of formic acid is lower than that of benz		
	1) Both (A) and (R) are true, and (R) is the correct ex		
	2) Both (A) and (R) are true, but (R) is NOT the corre	-	
	3) (A) is true, but (R) is false	4) (A) is false, but (R) is tr	ue

22.	2. Assertion: o-Substituted benzoic acids are generally stronger acids than benzoic acids.							
	Reason: Increased strength is due to ortho-effect by any group present at ortho postion.							
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).							
	3) (A) is true, but (R) is false							
	4) (A) is false, but (R) is true							
23.	Assertion (A): Carboxylic acids are more acidic than phenols.							
	Reason (R): -OH group in Phenols is ortho and para directing.							
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).							
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true							
_								
Pass	age-1: Read the passage given below and answer the following questions:							
	Carboxylic acids having an a-hydrogen atom when treated with chlorine or bromine in the presence of							
	small amount of red phosphorus gives a-halo carboxylic acids. The reaction is known as Hell-Volhard-							
	Zelinsky reaction. $RCH_2COOH + X_2 \xrightarrow{\text{Red}P} RCH(X)COOH_{(X=Cl,Br)}$							
	When sodium salt of carboxylic acid is heated with soda lime it loses carbon dioxide and gives							
	hydrocarbon with less number of C-atoms. $RCH_2COONa \xrightarrow{NaOH+CaO; \Lambda} RCH_3 + Na_2CO_3$							
24.	Assertion (A): (CH <sub>3</sub> ) <sub>3</sub> CCOOH does not give H.V.Z. reaction.Reason (R): (CH <sub>3</sub> ) <sub>3</sub> CCOOH does not have α-hydrogen atom	1						
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).							
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true							
25.	Assertion (A): H.V.Z. reaction involves the treatment of carboxylic acids having a-hydrogens with $Cl_2$ or $Br_2$ in							
	presence of small amount of red phosphorus.							
	Reason (R): Phosphorus reacts with halogens to form phosphorus trihalides.							
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).							
26	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true Assertion (A): $C_6H_5COCH_2COOH$ undergoes decarboxylation easily than $C_6H_5COCOOH$ .							
26.	Reason (R): $C_6H_5COCH_2COOH$ is a $\beta$ -keto acid.							
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).							
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true							
27.	Assertion (A): On heating 3-methylbutanoic acid with soda lime, isobutane is obtained.							
	Reason (R): Soda lime is a mixture of NaOH + CaO in the ratio $3:1$ .							
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).							
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true							
28.	The correct order of increasing acidic strength is							
	1) Phenol < Ethanol < Chloroacetic acid < Acetic acid							
	2) Ethanol < Phenol < Chloroacetic acid < Acetic acid							
	3) Ethanol < Phenol < Acetic acid < Chloroacetic acid							
	4) Chloroacetic acid < Acetic acid < Phenol < Ethanol							
29.	Which of the following is the correct order of relative strength of acids?							
	1) $CICH_2COOH > BrCH_2COOH > FCH_2COOH$ 2) $BrCH_2COOH > CICH_2COOH > FCH_2COOH$							
	3) $FCH_2COOH > ClCH_2COOH > BrCH_2COOH$ 4) $ClCH_2COOH > FCH_2COOH > BrCH_2COOH$							



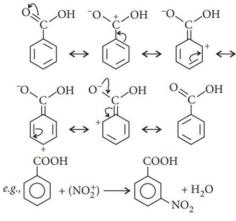


- 32. Which of the following orders is not correct for the decreasing order of acidic character?
  1) CH<sub>3</sub>CH<sub>2</sub>CH(Cl)COOH > CH<sub>3</sub>CH(Cl) CH<sub>2</sub>COOH > CH<sub>2</sub>(Cl)CH<sub>2</sub>CH<sub>2</sub>COOH > CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>COOH
  2) ICH<sub>2</sub>COOH > BrCH<sub>2</sub>COOH > ClCH<sub>2</sub>COOH > FCH<sub>2</sub>COOH
  - 3)  $CCl_3COOH > CHCl_2COOH > CH_2ClCOOH > CH_3COOH$
  - 4)  $HCOOH > CH_3COOH > C_2H_5COOH > (CH_3)_2CHCOOH$

#### **SUBJECTIVE OUESTIONS:**

#### 33. Give reasons:

- (i) Electrophilic substitution in benzoic acid takes place at meta-position.
- (ii) Carboxylic acids do not give the characteristic reactions of carbonyl group.
- (iii) pKa value of 4-nitrobenzoic acid is lower than that of benzoic acid.
- ANS: . (i) Electrophilic substitution in benzoic acid takes place at meta-position. Due to resonance in benzoic acid, there is high electron density at meta-position. Therefore, electrophilic substitution in benzoic acid takes place at meta-position.



(ii) The carbonyl group in —COOH is inert and does not show nucleophilic addition reaction like carbonyl compound. It is due to resonance stabilization of carboxylate ion :

$$\begin{array}{c} R-C=0 \longleftrightarrow R-C-0^{-1} \\ 1 \\ 0^{-1} \\ 0 \end{array}$$

(iii) Due to presence of strong electron withdrawing group (—NO<sub>2</sub>), 4-nitrobenzoic acid is more acidic than benzoic acid and therefore, pKa value is lower.

34. Which acid of each pair shown here would you expect to be stronger and why? (i) F—CH<sub>2</sub>—COOH or Cl—CH<sub>2</sub>—COOH (ii) Phenol or CH<sub>3</sub>COOH ANS: (i) F—CH<sub>2</sub>—COOH is more acidic than Cl—CH<sub>2</sub>—COOH because F is more electronegative or stronger -I group than Cl.

(ii) Phenol is less acidic than CH<sub>3</sub>COOH because phenoxide ion has non-equivalent resonating structures whereas Carboxylate ion has equivalent resonating structures which is more stable.

- 35. During practical exams, lab assistant provided two test tubes containing 5 mL benzoic acid and 5 mL acetaldehyde to every student. A student, Rahul found that test tubes given to him were unlabelled. He informed the teacher before performing any experiment with the given chemicals. How can the chemicals be distinguished for correct labelling?
- ANS: Chemicals can be distinguished by sodium bicarbonate test and iodoform test. Benzoic acid will give brisk effervescence due to evolution of carbon dioxide gas with sodium bicarbonate solution while acetaldehyde does not. Acetaldehyde will give yellow precipitate of iodoform with iodine and sodium hydroxide solution while benzoic acid does not

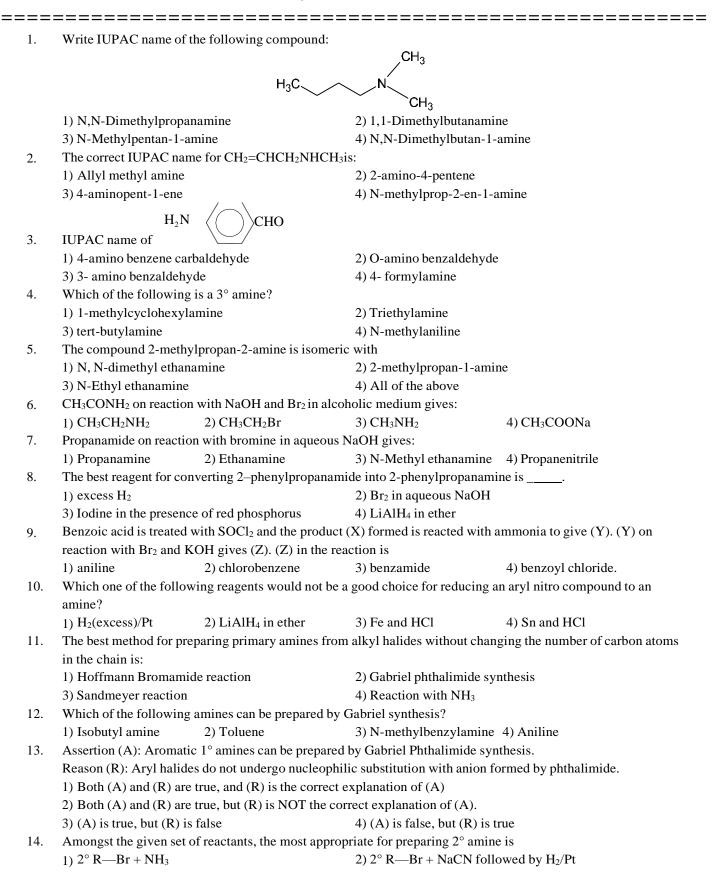
#### KEY-Aldehydes and Ketones

1	2	3	4	5	6	7	8	9	10
1	2	3	4	3	4	1	4	4	1
11	12	13	14	15	16	17	18	19	20
3	1	1	2	1	2	2	4	1	4
21	22	23	24	25	26	27	28	29	30
3	1	3	3	4	2	1	2	1	3
31	32	33	34	35	36	37	38	39	40
2	2	1	2	4	4	1	2	3	2
41	42	43	44	45	46	47	48	49	
3	2	1	4	3	3	1	2	2	

#### **KEY-Carboxylic acids**

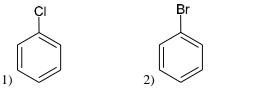
1	2	3	4	5	6	7	8	9	10
1	1	2	3	1	2	3	3	1	3
11	12	13	14	15	16	17	18	19	20
4	2	1	2	1	2	3	1	1	1
21	22	23	24	25	26	27	28	29	30
2	1	2	1	3	1	2	3	3	3
31	32								
1	2								

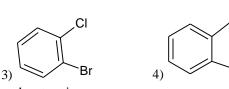
# Chapter 9: Amines



	3) 1° R—NH <sub>2</sub> + RCHO	) followed by H <sub>2</sub> /Pt						
		+ potassium phthalimide fo	llowed by H <sub>3</sub> O <sup>+</sup> /heat.					
15.			ases with increase in molar	mass.				
101	Reason (R): Intermolecular H bonds formed by the higher amines are weaker.							
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)							
		e true, but (R) is NOT the co	•					
	3) (A) is true, but (R) is $(R)$		4) (A) is false, but (R) is $(A = A + A)$					
16.				$H_2$ , $(C_2H_5)_2$ NH, $C_2H_5N(CH_3)_2$				
	1) $C_2H_5N(CH_3)_2 > (C_2H_3)_2$		2) $(C_2H_5)_2NH > C_2H_5N(C_2H_5)_2NH > C_2H_5N(C_5H_5)_2NH > C_2H_5N(C_5H_5)_2NH > C_2H_5N(C_5H_5)_2NH > C_$					
	3) $C_4H_9NH_2 > (C_2H_5)_2N_2$		4) $(C_2H_5)_2NH > C_4H_9NH$	$H_2 > C_2 H_5 N(CH_3)_2$				
17.		owest boiling points among						
	1) they have highest me	olecular mass	2) they do not form hydr	ogen bonds				
	3) they are more polar	in nature	4) they are most basic in	nature				
18.	Which of the following	should be most volatile?						
	(i) CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub>	(ii) (CH <sub>3</sub> ) <sub>3</sub> N	(iii) CH <sub>3</sub> CH <sub>2</sub> NHCH <sub>3</sub>	(iv) CH <sub>3</sub> CH <sub>2</sub> CH <sub>3</sub>				
	1) (ii)	2) (iv)	3) (i)	4) (iii)				
19.	The shape of (CH <sub>3</sub> ) <sub>3</sub> N i	s pyramidal because						
	1) nitrogen forms three	sp <sup>3</sup> hybridised sigma bond	s with carbon atoms of met	hyl groups and there is one non-				
	bonding electron pair	1 2 0						
		sp <sup>2</sup> hybridised sigma bond	s with carbon atoms of met	hyl groups and fourth orbital				
	forms pi bond	-F)8						
	-	encies which are arranged i	n pyramidal shape.					
	-	n present on nitrogen is delo						
20.	· •		he following compounds is					
		νΗ <sub>2</sub>	ŅН <sub>2</sub> ŅН <sub>2</sub>					
			$\Upsilon$ $\Upsilon$ $\Upsilon$ $\Theta_2$ $\Theta_2$					
		(1)	- 0113					
	1) 11 111 1		(II) (III)					
	1) II <iii< i<="" td=""><td>2) III <math>&lt;</math> I <math>&lt;</math> II</td><td>3) III &lt; II &lt; I</td><td>4) II <math>&lt;</math> I <math>&lt;</math> III A</td></iii<>	2) III $<$ I $<$ II	3) III < II < I	4) II $<$ I $<$ III A				
21.	Aniline is a resonance l	•						
	1) 3 resonating structur		2) 6 resonating structure					
	3) 2 resonating structur		4) 5 resonating structure	8				
22.	•	ent alkyl amines depends u						
	1) +I effect	2) steric effect	3) solvation effect	4) all of these				
23.	Methylamine reacts wi		_·					
	1) $CH_3 - O - N = O$	2) CH <sub>3</sub> — O—CH <sub>3</sub>	3) CH <sub>3</sub> OH	4) CH <sub>3</sub> CHO				
24.	•	methylamine reacts with ni						
	1) NH <sub>3</sub>	2) N <sub>2</sub>	3) H <sub>2</sub>	4) $C_2H_6$				
25.	Which of the following	species are involved in the	carbylamine test?					
	1) R—NC	2) COCl <sub>2</sub>	3) NaNO <sub>2</sub> + HCl	4) All of the above				
26.	Which of the following	amines will give carbylam	ine reaction?					
	1) $(C_2H_5)_3N$	2) $(C_2H_5)_2NH$	3) $C_2H_5NH_2$	$4) C_3 H_7 NHC_2 H_5$				
27.	Assertion (A): Acylatic	on of amines gives a monosi	ubstituted product whereas	alkylation of amines gives poly				
	substituted product.							
	Reason (R): Acyl group	p sterically hinders the appr	oach of further acyl groups					
	1) Both (A) and (R) are	e true, and (R) is the correct	explanation of (A)					
	2) Both (A) and (R) are	e true, but (R) is NOT the co	prrect explanation of (A).					
	3) (A) is true, but (R) is	s false	4) (A) is false, but (R) is	true				

- 28. Why is the activating effect of -NHCOCH<sub>3</sub> group in the above reaction less than the activating effect of amino group? 1) Due to mesomeric effect of benzene ring 2) Due to inductive effect of alkyl group 3) Due to resonance effect of acetanilide 4) Due to electromeric effect of alkyl group 29. Which of the following can exist as zwitter ion? 1) p-Aminoacetophenone 2) Sulphanilic acid 3) p-Nitroaminobenzene 4) p-Methoxyphenol 30. The reaction of benzenesulphonyl chloride with ethylamine yields 1) N-ethylbenzenesulphonamide, insoluble in alkali 2) N, N-diethylbenzenesulphonamide, soluble in alkali 3) N, N-diethylbenzenesulphonamide, insoluble in alkali 4) N-ethylbenzenesulphonamide, soluble in alkali. Hinsberg's reagent which is used to test amines is 31. 1) Benzene sulphonamide 2) Benzene diazonium chloride 3) Benzene sulphonyl chloride 4) Acetanilide 32. Assertion (A): N, N-Diethylbenzene sulphonamide is insoluble in alkali. Reason (R): Sulphonyl group attached to nitrogen atom is strong electron withdrawing group. 1) Both (A) and (R) are true, and (R) is the correct explanation of (A) 2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A). 3) (A) is true, but (R) is false 4) (A) is false, but (R) is true Amino group is o, p-directing for electrophilic substitution reaction. But, on nitration the major product is m-33. nitroaniline because 1) aniline gets protonated with strong acids to give anilinium ion which is m-directing 2) nitration requires nitric acid which oxidises -NH<sub>2</sub> to -NO<sub>2</sub> group 3) electrophile  $NO_2^+$  is a m-directing group 4) benzene ring exerts + I effect and deactivates the ring 34. Electrophilic substitution of aniline with bromine- water at room temperature gives 2) 3-bromoaniline 3) 2, 4, 6-tribromoaniline 4) 3, 5, 6-tribromoaniline 1) 2-bromoaniline
  - o-Chloroaniline is treated with a mixture of NaNO<sub>2</sub> and HCl and the product is reacted with cuprous bromide. 35. The final product in the reaction will be





4) Phenol

Br

Benzene diazonium chloride on reaction with H<sub>3</sub>PO<sub>2</sub> and water gives 36.

1) Chloro benzene 2) Benzene

3) Benzene phosphate 37. Which of the following is incorrectly matched with the product when benzene diazonium chloride is reacted with the reagent?

	Reagent	Product
1	CuBr, HBr	Bromo benzene
2	$NaBF_4, \Delta$	Nitro benzene
3	Cu/HCl	Chlorobenzene
4	KCN	Cyano benzene

Identify X, Y and Z in the given reaction:  $CH_2 = CH_2 \xrightarrow{Br_2/CCl_4} X \xrightarrow{NaCN} Y$ 38.

1) X=CH<sub>2</sub>Br–CH<sub>2</sub>Br; Y=CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CN; Z=CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>

2) X= CH<sub>2</sub>Br–CH<sub>2</sub>Br; Y=CH<sub>3</sub>CH<sub>2</sub>CN; Z=CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>

3) X=CH<sub>3</sub>CH<sub>2</sub>Br; Y=CH<sub>3</sub>CH<sub>2</sub>CN; Z=CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>

4) X=CH<sub>2</sub>Br–CH<sub>2</sub>Br; Y=NCCH<sub>2</sub>CH<sub>2</sub>CN; Z=H<sub>2</sub>NCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>

39.	$C_6H_5NO_2 \xrightarrow{S_{n+HCl}} A \xrightarrow{NaNO_2 + HCl;0^6C} B \xrightarrow{H_2O} C;$ Correct statement regarding A, B, C is 1)
	1)Compound A is aniline       2) Compound B is benzene diazonium chloride
	3) C is Phenol4) All of these
0.	$C_6H_5CONH_2 \xrightarrow{BP_2+KOH} A \xrightarrow{NaNO_2+HCl;0^{\circ}C} B \xrightarrow{KI} C;$ ; Compound C is the reaction is
	1) Iodo benzene2) Anilinium iodide3) Acetyl iodide4) Bromo benzene
ass	sage-1: Read the passage given below and answer the following questions:
	Benzene ring in aniline is highly activated. This is due to the sharing of lone pair of nitrogen with the ring which results in increase in the electron density on the ring and hence facilitates the electrophilic attack. The substitution mainly takes place at ortho and para positions because electron density is more at ortho and para positions. On reaction with aqueous bromine all the ortho and para positions get substituted resulting in the formation of 2,4,6-tribromoaniline. To get a mono bromo compound, the amino group is
	acetylated before bromination. After bromination, the bromo acetanilide is acid hydrolysed to give the
	desired halogenated amine.
1.	Assertion (A): Benzene ring of aniline is highly deactivated.
	Reason (R): In aniline, the sharing of lone pair of nitrogen with the ring increases the electron density on the
	ring.
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true
2.	Assertion (A): In aniline $-NH_2$ group facilitates the electrophilic attack.
	Reason (R): It is due to decrease in electron density on the ring.
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true
3.	Assertion (A): In aniline, the substitution mainly takes place at ortho and para positions.
	Reason (R): The electron density is more at ortho and para positions
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true
4.	Assertion (A): The amino group of aniline is acetylated before bromination.
	Reason (R): It is due to the strong deactivating effect of $-NH_2$ group.
	1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
	2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).
	3) (A) is true, but (R) is false 4) (A) is false, but (R) is true
ass	sage-2: Read the passage given below and answer the following questions:
	Coniine (X) is a poisonous chemical compound with molecular formula C <sub>8</sub> H <sub>17</sub> N. In 399 B.C Socrates was
	sentenced to death by drinking a coniine containing mixture of poison hemlock. It is the murder weapon
	in mystery novel "Five little Pigs" by Agatha Christie. This when treated with nitrous acid does not give
	Nitrogen. Also, another Organic compound 'Y" when treated with HNO <sub>2</sub> gives nitrogen. This compound
	Y when reacts with Hinsberg reagent forms a compound 'Z' which is soluble in NaOH. Such chemical
	reactions form the basis to differentiate organic compounds and to elucidate its structure.
5.	Which of the following statement is incorrect-
	1) Compound X and Y both are nitrogenous containing organic compounds
5.	2) Y is a Primary amine
	<ul><li>2) Y is a Primary amine</li><li>3) X might contain N in the ring</li></ul>
	<ul><li>3) X might contain N in the ring</li><li>4) X and Y both can give carbylamine reaction.</li></ul>
6.	3) X might contain N in the ring

47.	Hinsberg's reagent is	
	1) Phenylisocyanide	2) Benzenesulphonyl chloride
	3) p-toluenesulphonic acid	4) O-dichlorobenzene

48. Which of the following compounds will dissolve in an alkali solution after it has undergone reaction with Hinsberg reagent?
1) (C<sub>2</sub>H<sub>5</sub>)<sub>2</sub>NH
2) C<sub>6</sub>H<sub>5</sub>N(CH<sub>3</sub>)<sub>2</sub>
3) CH<sub>3</sub>NH<sub>2</sub>
4) C<sub>6</sub>H<sub>5</sub>NHC<sub>6</sub>H<sub>5</sub>

Passage-3: Read the passage given below and answer the following questions:

The amines are basic in nature due to the presence of a lone pair of electron on N-atom of the  $-NH_2$  group, which it can donate to electron deficient compounds. Aliphatic amines are stronger bases than NH<sub>3</sub> because of the +I effect of the alkyl groups. Greater the number of alkyl groups attached to N-atom, higher is the electron density on it and more will be the basicity. Thus, the order of basic nature of amines is expected to be  $3^{\circ}>2^{\circ}>1^{\circ}$ , however the observed order is  $2^{\circ}>1^{\circ}>3^{\circ}$ . This is explained on the basis of crowding on N-atom of the amine by alkyl groups which hinders the approach and bonding by a proton, consequently, the electron pair which is present on N is unavailable for donation and hence  $3^{\circ}$  amines are the weakest bases. Aromatic amines are weaker bases than ammonia and aliphatic amines. Electron-donating groups such as  $-CH_3$ ,  $-OCH_3$ , etc. increase the basicity while electron-withdrawing substitutes such as  $-NO_2$ , -CN, halogens, etc. decrease the basicity of amines. The effect of these substituents is more at p than at m-positions

3) ClCH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>

49. Which of the following has highest pKb value?1) FCH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>2) CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub>

50.Among the compounds:  $C_3H_7NH_2$ ,  $CH_3NH_2$ ,  $C_2H_5NH_2$  and  $C_6H_5NH_2$ . Which is the least basic compound?1)  $CH_3NH_2$ 2)  $C_2H_5NH_2$ 3)  $C_3H_7NH_2$ 4)  $C_6H_5NH_2$ 

- 51. Assertion (A): Acetanilide is less basic than aniline.
  Reason (R): Acetylation of aniline results in decrease of electron density on nitrogen.
  1) Both (A) and (R) are true, and (R) is the correct explanation of (A)
  2) Both (A) and (R) are true, but (R) is NOT the correct explanation of (A).
  3) (A) is true, but (R) is false
  4) (A) is false, but (R) is true
- 52.Arrange the following in increasing order of basic strength: Aniline, p-nitroaniline and p-toluidine<br/>1) Aniline < p-Nitroaniline < p-Toluidine<br/>3) p-Toluidine < p-Nitroaniline < Aniline</th>2) Aniline < p-Toluidine < p-Nitroaniline<br/>4) p-Nitroaniline < Aniline < p-Toluidine</th>
- 53. Which of the following statements is not correct?1) Methylamine is more basic than NH<sub>3</sub>
- 2) Primary amines form hydrogen bonds.

4) All will have same value

3) Ethylamine has higher boiling point than propane 4) Dimethylamine is less basic than methyl amine

#### **SUBJECTIVE QUESTIONS:**

54. Account for the following-

# i) Nitrogen of Aniline act as deactivating group on reacting with RX in presence of Anhyd.AlCl<sub>3</sub>ii) Amines are less polar than alcohol.

- ANS: i) Aniline does not react with FC Reaction due to the catalyst Anhyd.AlCl<sub>3</sub> which is Lewis acid and Nitrogen acquires +ve charge and act as deactivating group
  - ii) Nitrogen is less electronegative than Oxygen
- 55. A colourless substance 'A' (C<sub>6</sub>H<sub>7</sub>N) is sparingly soluble in water and gives a water-soluble compound 'B' on treating with HCl. On reacting with CHCl3 and alcoholic potash 'A' produces an obnoxious smell due to the formation of compound 'C' How will you differentiate A with benzyl amine and N, N-dimethyl benzylamine

ANS: A=Aniline (C<sub>6</sub>H<sub>5</sub>NH<sub>2</sub>) B= Anilinium Chloride (C<sub>6</sub>H<sub>5</sub>NH<sub>3</sub>)<sup>+</sup> Cl<sup>-</sup>, C= Phenyl isocyanide (C<sub>6</sub>H<sub>5</sub>NC) Aniline gives red compound or dye with HONO (Cold) +  $\beta$ -Naphthol. Compound A (Aniline), Benzyl amine and N,N-dimethyl benzyl amine can be differentiated by Hinsberg test.

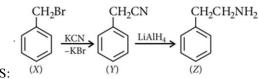
56. An organic solid compound (A) (C<sub>13</sub>H<sub>11</sub>ON) is insoluble in aqueous NaOH and aqueous HCl. However, its prolonged heating with aqueous NaOH gives a liquid (B) and a compound (C). Compound (B) can be

separated from (C) by steam distillation, Compound (B)on treatment with benzene diazoniumchloride gives a yellow dye (E). Acidification of C gives white precipitate of the compound (D)  $(C_7H_6O_2)$  which on distillation with soda lime gives benzene. Identify compound A, B, C, D and E.

ANS:  $A = C_6H_5CONHC_6H_5$   $B = C_6H_5NH_2$   $C = C_6H_5COONa$   $D = C_6H_5COOH$ 

E= p-amino azo benzene

57. A compound X (C<sub>7</sub>H<sub>7</sub>Br) reacts with KCN to give Y (C<sub>8</sub>H<sub>7</sub>N). Reduction of Y with LiAlH<sub>4</sub> yields Z (C<sub>8</sub>H<sub>11</sub>N). Z gives carbylamine reaction, reacts with Hinsberg's reagent in the presence of aq. KOH to give a clear solution. With NaNO<sub>2</sub> and HCl at 0°C (Z) gives a neutral compound which gives red colour with ammonium ceric nitrate. What are X, Y and Z?



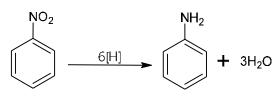
ANS:

58. Amit wants to manufacture aniline for the synthesis of dye stuff. For this he has selected tin and hydrochloric acid as reducing agent for the reduction of nitrobenzene. But his friend suggested to use iron scrap and hydrochloric acid as the reducing agent.

Now answer the following questions:

- (i) Write the chemical equation for the reduction of nitrobenzene to aniline.
- (ii) Why Amit's friends have suggested to use scrap iron and HCl in place of tin and HCl?

ANS: (i)  $[Fe+2HCl \rightarrow FeCl_2+2H]x3$ 

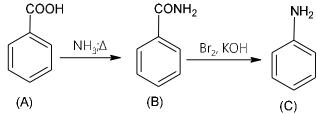


nitrobenzene

aniline

(ii) Amit's friend has suggested to use scrap iron and hydrochloric acid because in this reaction  $FeCl_2$  formed gets hydrolysed to release HCl during the reaction. Thus only small amount of HCl is required to initiate the reaction and scrap iron is also cheaper

- 59. An aromatic compound 'A' on treatment with aqueous ammonia and heating forms compound 'B' which on heating with Br<sub>2</sub> and KOH forms a compound 'C' of molecular formula C<sub>6</sub>H<sub>7</sub>N. Write the structures and IUPAC names of compounds A, B and C.
- ANS: Formula of the compound 'C' indicates it to be an amine. Since it is obtained by the reaction of Br<sub>2</sub> and KOH with the compound 'B' so compound 'B' can be an amide. It is also indicated because 'B' is obtained from compound 'A' by reaction with ammonia following by heating. So compound 'A' could be an aromatic acid. Formula of compound 'C' shows that it is aniline, then 'B' is benzamide and compound 'A' is benzoic acid. The sequence of reactions can be written as follows:



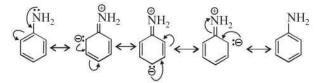
- 60. Give plausible explanation for each of the following:
  - (i) Why are amines less acidic than alcohols of comparable molecular masses?
  - (ii) Why do primary amines have higher boiling point than tertiary amines?
  - (iii) Why are aliphatic amines stronger bases than aromatic amines?

ANS: (i) In alcohols, the hydrogen atom is attached to more electronegative oxygen atom whereas nitrogen of amines is less electronegative. After the loss of H+ ion, the negative charge is more easily accommodated on oxygen than in case of nitrogen in amines. Hence, amines have lesser tendency to lose H+ ions, so they are less acidic than alcohols.

## $RNH_2 \rightarrow RNH^- + H^+$ ; $ROH \rightarrow RO^- + H^+$

(ii) Primary amines (R - NH2) have two hydrogen atoms on nitrogen which can undergo intermolecular hydrogen bonding whereas no such hydrogen bonding is present in tertiary amines (R3N). So primary amines boil at a higher temperature than tertiary amines.

(iii) In aromatic amines, the lone-pair of electrons on nitrogen atom is involved in resonance with the benzene ring as shown below for aniline



It shows that this pair of electrons is less available for protonation. In case of aliphatic amines electron releasing alkyl groups increase electron density on nitrogen atom. So, aliphatic amines are stronger bases than aromatic amines

1	2	3	4	5	6	7	8	9	10
4	4	1	2	4	3	2	4	1	2
11	12	13	14	15	16	17	18	19	20
2	1	4	3	3	3	2	2	1	4
21	22	23	24	25	26	27	28	29	30
4	4	3	2	1	3	3	3	2	4
31	32	33	34	35	36	37	38	39	40
3	1	1	3	1	2	2	4	4	1
41	42	43	44	45	46	47	<b>48</b>	49	50
4	3	1	3	4	2	2	3	1	4
51	52	53							
1	4	4							

#### **KEY-AMINES**

# 

	<u>Chapter 10 : BIOMOLECULES</u>							
1.	Glucose gives silver mirror with Tollen's reagen	t. It shov	vs the p	oresence of				
	(a) An acidic group	(b)	An al	coholic group				
	(c) A ketonic group	(d)	An al	dehydic group				
2.	On complete hydrolysis of starch, we finally get							
	(a) Glucose		(b)	Fructose				
	(c) Glucose and fructose		(d)	Sucrose				
3.	Which of the following statements is right							
	(a) Cellulose are linear polymers of $\beta$ – glucose mo	lecules w	vith $\beta$ –	1,4 – linkages				
	<ul> <li>(b) Starches are polymers of α – glucose molecules cross-linkages</li> </ul>	with β–	1,4 – lin	kages and some $\beta - 1, 6 -$				
	(c) Proteins are polyamides of $\beta$ – amino acids							
	(d) The structural information about their biosynthe called nucleic acids, e.g. RNA and DNA	esis is co	ntained	in a class of compounds				
4.	A sugar, that is not a disaccharide, among the fo	llowing i	is					
	(a)Lactose		(b)	Galactose				
	(c) Sucrose		(d)	Maltose				
5.	The helical structure of proteins is established b	y						
	(a) Peptide bonds		(b)	Dipeptide bond				
	(c) Hydrogen bond		(d)	Vander Waal's forces				
6.	Secondary structure of a protein refers to							
	(a) Mainly denatured proteins and structures of pro	-	-					
	(b) Three dimensional structure, specially the bon distant from each other in the polypeptide chain	l						
	(c) Linear sequence of amino acid residues in the p							
	(d) Regular folding patterns of continuous portions	of the po	lypepti	de chain				
7.	What is the monomer of polypeptide		(1-)	<u>C</u> 1				
	(a) Amino acid		(b)	Glucose				
0	(c) Nucleoside	tona Th	(d)	Nucleotide				
8.	Amino acids usually exist in the form of Zwitter		ns mea	iis that it consists of				
	(a) The basic group $-NH_2$ and the acidic group $-CO$							
	(b) The basic group $-NH_{3}^{+}$ and the acidic group $-CC$	-						
	(c) The basic group $-CO_{\frac{1}{2}}$ and the acidic group <i>NH</i>	+ 3						
	(d) No acidic or basic group							
9.	A nanopeptide containspeptide linkages							
	(a) 10		(b)	8				
	(c) 9		(d)	18				

# 10. Which of the following statements about the assembly of nucleotides in a molecule of deoxyribose nucleic acid (DNA) is correct

- (a)A pentose of one unit connects to a pentose of another
- (b) A pentose of one unit connects to the base of another
- (c) A phosphate of one unit connects to a pentose of another
- (d) A phosphate of one unit connects to the base of another

# 11. In DNA, the complementary bases are

- (a) Uracil and adenine; cytosine and guanine
- (b) Adenine and thymine; guanine and cytosine
- (c) Adenine and thymine; guanine and uracil
- (d) Adenine and guanine; thymine and cytosine

#### 12. The base present in DNA, but not in RNA is

(a)Guanine

(c) Uracil

(b) Adenine

(d) Thymine

#### 13. A nucleoside on hydrolysis gives

- (a) A heterocyclic base and orthophosphoric acid
- (b) An aldopentose, a heterocyclic base and orthophosphoric acid
- (c) An aldopentose and a heterocyclic base
- (d) An aldopentose and orthophosphoric acid

# **Read the assertion and reason carefully to mark the correct option out of the options given below :**

- (a) If both assertion and reason are true and the reason is the correct explanation of the assertion.
- (b) If both assertion and reason are true but reason is not the correct explanation of the assertion.
- (c) If assertion is true but reason is false.

#### (d) If assertion is false but reason is true.

- **14. Assertion**: Proteins on hydrolysis produce amino acids.**Reason**: Amino acids contain  $-NH_2$  and -COOH groups.
- **15. Assertion** : All Amino acids exist as Zwitter ions. **Reason** : Amino acids have both *–NH*<sup>2</sup> and *–COOH* group.
- 16. Assertion : Fructose reduces Fehling's solution and Tollen's reagent.Reason : Fructose does not contain any aldehyde group.

**17. Assertion** :Sequence of bases in DNA is TGAACCCTT and sequence of bases in *m*-RNA is CATTAAACC.

**Reason** : In DNA nitrogenous bases have hydrogen bonds.

**18. Assertion** : Sucrose is a non-reducing sugar.

## Reason : It has glycosidic linkage

# **19.** Assertion : $\beta$ -glycosidic linkage is present in maltose

**Reason** : Maltose is composed of two glucose units in which C-1 of one glucose unit is linked to C-4 of another glucose unit.

### 20. Assertion: Sucrose is called an invert sugar.

**Reason:** on hydrolysis, sucrose bring the change in the sign of rotation from dextro (+) to leavo (-)

## Read the passage given below and answer the following questions:

When a protein in its native form, is subjected to physical changes like change in temperature or chemical changes like change in pH, the hydrogen bonds are disturbed. Due to this, globules unfold and helix gets uncoiled and protein loses its biological activity. This is called denaturation of protein. The denaturation causes change in secondary and tertiary structures but primary structures remains intact.

Examples of denaturation of protein are coagulation of egg white on boiling, curding of milk, formation of cheese when an acid is added to milk.

# The following questions are multiple choice questions. Choose the most appropriate answer:

### (21) Mark the wrong statement about denaturation of proteins.

(a) The primary structure of the protein does not change.

(b) Globular proteins are converted into fibrous proteins.

- (c) Fibrous proteins are converted into globular proteins.
- (d) The biological activity of the protein is destroyed.

# (22) Which structure of protein remain(s) intact during denaturation process?

- (a) Both secondary and tertiary structures
- (b) primary structure only
- (c) secondary structure only
- (d) tertiary structure

# (23) $\alpha$ -helix and $\beta$ -pleated structures of proteins are classified as

- (a) primary structure
- (b) secondary structures
- (c) tertiary structure
- (d) quaternary structure

# (24) Cheese is a

- (a) globular protein
- (b) conjugated protein
- (c) denatured protein
- (d) derived protein

# (25) Secondary structure of protein refers to -

(a) mainly denatured of proteins and structures of prosthetic groups

(b) three-dimensional structure, especially the bond between amino acid residues that are distant from each other in the polypeptide chain

(c) linear sequence of amino acid residues in the polypeptide chain

(d) regular folding patterns of continuous portions of the polypeptide chain.

### Read the passage given below and answer the following questions:

Glucose ( $C_6H_{12}O_6$ ) contains six carbon atoms and an aldehyde group and is therefore referred to as an aldohexose. The glucose molecule can exist in an open-chain (acyclic) and ring (cyclic) form in equilibrium. The later being the result of an intramolecular reaction between the aldehyde C atom and the C-5 hydroxyl group to form an intramolecular hemiacetal. As the ring contain five carbon atoms and one oxygen atom. Which resembles the structure of pyran, the cyclic form of glucose is also referred to as glucopyranose. In this ring, each carbon is linked to a hydroxyl side group with the exception of the fifth atom, which links to a sixth carbon atom outside the ring, forming a CH<sub>2</sub>OH group.

#### 26. Glucose react with bromine water which product is formed

(a) saccharic acid	(b) gluconic acid	(c) hexanoic acid	(d) bromo
hexane			
<b>27. The</b> number of chir	al centers in the open chain s	tructure of glucose is	
(a) 3	(b) 4	(c) 5	(d) 6

(a) 3 (b) 4(c) 5

**28.**  $\alpha$ -D-Glucopyranose and  $\beta$ -D-Glucopyranose are

(a) Diastereomers (b)Meso compounds (c) Epimers (d) Anomers

- 29. Glycosidic linkage is
  - (a) an amide linkage (b) an ester linkage (c) an ether linkage (d) an amine linkage
- 30. Glucose and Fructose can be differentiated by
  - (a) Tollens reagent (b) Cold KMnO4 (c) bromine water (d) PCC

ANSWERS:- (1) d	(2) a	(3) b	(4) b	(5) c	(6) d
(7) a	(8) c	(9) b	(10) c	(11) b	(12) d
(13) c	(14) b	(15) a	(16) b	(17) d	(18) a
(19) d	(20) a	(21) c	(22) b	(23) b	(24) c
(25) d	(26) b	(27) b	(28) d	(29) c	(30) c