

level-01

$$(1) \quad M_1 V_1 + M_2 V_2 = M_f (V_1 + V_2)$$

$$25 \times 3 + 75 \times 4 = M_f (25 + 75)$$

$$M_f = 3.75 \text{ M}$$

$$(2) \quad 0.25 = \frac{W \times 100^4}{106 \times 250}$$

$$W = \frac{106}{16} = 6.625 \text{ gm}$$

$$(3) \quad N_1 V_1 = N_2 V_2$$

$$2 \times 1 = N_2 \times 6$$

$$N_2 = 0.33 \text{ N}$$

$$(4) \quad \chi_{\text{NaCl}} = \frac{\frac{5.85}{58.5}}{\frac{5.85}{58.5} + \frac{90}{18}} = \frac{0.1}{0.1 + 5} = 0.0196$$

~~$$(5) \quad M = \frac{0.006 \times 1000}{100} = 0.06 \text{ M}$$~~

~~$$(6) \quad M = \frac{98}{98 \times 2} = 0.05 \text{ M}$$~~

~~$$(7) \quad M = \frac{5 \times 1000}{40 \times 250} = 0.5$$~~

$$\begin{aligned} (6) \quad N &= M \times n_{\text{factor}} & n_{\text{factor}}(\text{H}_3\text{PO}_3) &= 2 \\ &= 0.3 \times 2 \\ &= 0.6 \text{ N} \end{aligned}$$

$$(7) \quad M_1 V_1 = M_2 V_2$$

$$M_1 \times 20 = 0.01 \times 19.85$$

$$M_1 = \frac{0.1985}{20} = 0.009925 \text{ M}$$

$$(8) \quad 4V_1 + 10V_2 = 6$$

$$V_1 + V_2 = 1$$

$$V_1 = \frac{2}{3} \text{ L}; \quad V_2 = \frac{1}{3} \text{ L}$$

$$V_1 = 0.67 \text{ L}; \quad V_2 = 0.33 \text{ L}$$

(11) ~~Molality is no. of mole of solute per 1000 gm of solvent.~~

$$(9) \quad M = \frac{22 \times 1.253 \times \frac{10}{1000}}{342 \times \frac{100}{1000}} = 0.805 \text{ M}$$

$$\begin{aligned} N &= 0.805 \times n_{\text{factor}} = 0.805 \times 6 \\ &= 4.830 \text{ N} \end{aligned}$$

$$m = \frac{22 \times 1000}{342 \times 78} = 0.825 \text{ m}$$

$$(10) \quad N_1 V_1 = N_2 V_2$$

$$0.4 \times 20 = N_2 \times 40$$

$$N_2 = 0.2$$

$$M_2 = 0.1 \text{ M}$$

$$(11) \quad c = \frac{6.02 \times 10^{20} \times 1000}{6.02 \times 10^{23} \times 100} = 0.01 M$$

$$(12) \quad \chi_{H_2O} = \frac{25/18}{25/18 + \frac{25}{46} + \frac{50}{60}} = 0.503$$

$$(13) \quad \frac{n_{\text{solute}}}{n_{\text{solute}} + n_{\text{solvent}}} = 0.2 \Rightarrow n_{\text{solute}} \times 0.8 = n_{\text{solvent}} \times 0.2$$
$$\frac{n_{\text{solute}}}{w_{\text{solvent}}} = \frac{1000}{78 \times 4}$$
$$m = 3.2 m$$

$$(14) \quad \frac{p^0 - p}{p^0} = \frac{w_B \times M_A}{M_B \times w_A}$$
$$0.00719 = \frac{71.5 \times 18}{M_B \times 1000}$$

$$M_B = 180$$

(15) Boiling point will be increased

$$(16) \quad p = p^0 N_1$$

(17) glucose

$$(18) \quad \chi_{\text{solute}} = \frac{50 - 45}{50} = 0.1$$

$$(19) \quad p_{\text{Total}} = p_{\text{Pentane}} + p_{\text{Hexane}} = \frac{1}{5} \times 440 + \frac{4}{5} \times 120 = 184$$

$$\chi_{\text{vapour (pentane)}} = \frac{\frac{1}{5} \times 440}{184} = 0.478$$

$$\begin{aligned} (20) \quad P_B &= P_B^\circ \times X_B \\ &= 75 \times \frac{1}{1.5} \\ &= 50 \end{aligned}$$

$$(21) \quad \frac{P^\circ - P}{23.8} = \frac{100 \times 18}{342 \times 1000}$$

$$P^\circ - P = 0.125 \text{ mmHg.}$$

(22) For BaCl_2 lowering of vapour pressure is highest because $i = 3$

$$(23) \quad P = \frac{P^\circ \times 9.9}{10.9} = 0.90 P^\circ$$

(24) same as ques. 19

(25) Benzene + methanol.

(26) $\text{CCl}_3 + \text{CHCl}_3$

(27) $\Delta S_{\text{mix}} > 0$ for ideal solution.

(28) We cannot obtain HCl or H_2O in their pure states.

(29) Minimum boiling azeotropic solution shows positive deviation from Raoult's law.

(30) Azeotropic mixture are constant temperature boiling mixture.

(31) Cane sugar Urea
 $C_1 = C_2$

$$\frac{2}{342} = \frac{6}{60 \times 1}$$

$$2 = 34.2$$

(32) $\pi = CRT$

$$C = \frac{\pi}{RT} = \frac{0.0821}{0.0821 \times 300} = 0.33 \times 10^{-2}$$

(33) $\pi = CRT$

$$= \frac{5}{342} \times 0.0821 \times 423 \times 10$$

$$= 5.078 \text{ atm}$$

(34) $P \propto \frac{1}{\text{Molar mass}}$

$$P_2 > P_1 > P_3$$

(35) Osmosis is the movement of solvent molecules from lower to higher concentration.

(36) same as (35).

(37) $C_1 = C_2$

$$\frac{5}{342} = \frac{1}{x}$$

$$x = 68.4$$

(38) Same as ques 32

(39) Blood is isotonic with 0.16M NaCl

(40) copper ferrocyanide

$$(41) \quad \pi V = \frac{m}{M} R T$$

$$6 \times 10^{-4} \times 1 = \frac{4}{M} \times 0.082 \times 300$$

$$M = 1.6 \times 10^5$$

(42) Boiling point will be increased.

$$(43) \quad \Delta T_b = K_b \times m$$

$$0.52 = \frac{0.52 \times 6 \times 1000}{M \times 100}$$

$$M = 60$$

$$(44) \quad \Delta T_b = K_b \times m$$

$$= \frac{0.513 \times 0.1 \times 1000}{200}$$

$$\Delta T_b = 0.2565$$

$$T_1 - T_0 = 0.2565$$

$$T_1 = 100.2565^\circ$$

$$(45) \quad \Delta T_b = i K_b m$$

$$= \frac{3 \times 0.52 \times 13.44}{134.4 \times 1}$$

$$= 1.56 \times 0.1$$

$$= 0.156 \sim 0.16$$

$$(1) \Delta T_b = K_b m$$

$$1 = \frac{2.53 \times 10 \times 1000}{M \times 100}$$

$$M = 253$$

$$(2) \Delta T_b = K_b m$$

$$\frac{0.25}{\Delta T_b} = \frac{1 \times 180}{60 \times 3} = 1$$

$$\Delta T_b = 0.25$$

$$T_1 = 100 - 0.25^\circ\text{C}$$

$$(3) \Delta T_b = i K_b m$$

$$= 1 \times 1.86 \times 0.05$$

$$= \frac{1.86 \times 5}{10000}$$

$$= 0.093$$

$$T_1 = -0.093^\circ\text{C}$$

$$(4) \frac{\Delta T_b}{0.186} = \frac{0.521}{1.86}$$

$$\Delta T_b = 0.0521$$

(5) on adding non volatile solute freezing point decreases.

(6) same as ques 1.

(7) $\Delta T_f \propto i K_f m$

for urea. $i=1$

(8) same as ques. 7.

(9)
$$\Delta T_f = \frac{1.86 \times 17 \times 1000}{46 \times 1000}$$

$\Delta T_f = 0.69^\circ\text{C}$

$T_i = -0.69^\circ\text{C}$

(10) KCl solution > phenol solution > glucose solution.

(11) for 0.1 M $\text{Al}_2(\text{SO}_4)_3$ $i=5$

(12) same as ques. 11

(13) $\Delta T_f \propto i$

	Urea	common salt (NaCl)	Na_2SO_4
i	1	2	3
	1:2:3		

(14) same as ques. 8

(15) same as ques. 8.

(16) same as ques. 8

(17) same as ques. 8

(18) same as ques. 8

(19) same as ques. 8

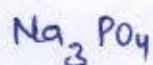
(20) same as ques. 19

(21) same as ques. 19

(22) for 0.1M $K_4[Fe(CN)_6]$

$$i = 5$$

(23) for sodium phosphate

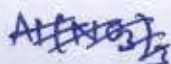


$$i = 4$$

(24) for $K_4[Fe(CN)_6]$; for $Al_2(SO_4)_3$

$$i = 5$$

$$i = 5$$



$$i = 4$$

(25) $Ca(NO_3)_2$

$$i = 3$$

(26) for urea (non electrolyte)

$$i = 1$$

(27) $\Delta T_f = i K_f m$

$$\alpha = \frac{i-1}{n-1}$$

$$0.2 = \frac{i-1}{1}$$

$$i = 1.2$$

$$\Delta T_f = 1.2 \times 1.86 \times 0.2$$

$$= 0.24 \times 1.86$$

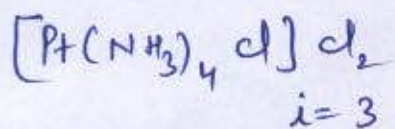
$$= 0.45$$

$$T_1 = -0.45^\circ C$$

$$(28) \Delta T_f = i K_f m$$

$$0.0054 = i \times 1.8 \times 0.001$$

$$i = 3$$



(29) Same as ques 28.

$$(30) i_{KNO_3} > i_{CH_3COOH}$$

$P_1 > P_2$

$$P_1 > P_2$$

(31) Same as ques. 20

$$(32) \alpha = \frac{i-1}{x+y-1}$$

$$(33) Y_A = \frac{P_A^0 X_A}{P_{Total}}$$

graph of Y_A Vs $\frac{1}{P_{Total}}$ is linear.

(34) same as ques. 28

$$(35) \quad Y_6 = \frac{408 \times 3}{408 \times 3 + 141 \times 7}$$

$$Y_6 = 0.554$$

$$Y_7 = 0.446$$

(36) same as 35

$$(37) \quad \Delta T_b = \text{£}Kb m$$

$$1.6 = \frac{\text{£} \times 2.34 \times 28 \times 76}{M \times 315}$$

$$(38) \quad i = \frac{5}{7} = 0.71$$

$$(39) \quad i = \frac{5.85}{3.2} = 1.82$$

$$d = \frac{i-1}{n-1} = \frac{1.82-1}{2-1} = 0.82$$

$$(40) \text{ weight (Benzene)} = v \times d = 5 \times 0.879 = 43.95 \text{ gm}$$

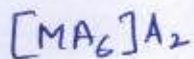
$$\Delta T_f = K_f \times m \Rightarrow 0.48 = \frac{5.12 \times 0.643 \times 1000}{M \times 43.95}$$

$$M = 156.05$$

$$(41) \Delta T_f = i K_f m$$

$$0.00558 = i \times 1.86 \times 0.001$$

$$i = 3$$



$$(42) \text{ for NaCl, } i = 2$$

$$\Delta T_f = i K_f m$$

$$\frac{\Delta T_f}{K_f} = 2 \times m$$

$$m = \frac{1}{1000} \sim 0.0585 \text{ gm NaCl}$$

(43) Non ideal solution with positive deviation -

(44) osmosis is the movement of solvent from low concentration to high concentration.

$$(45) P = (P_A^\circ - P_B^\circ) X_A + P_B^\circ$$

$$P_B^\circ = 140$$

Assertion & Reasoning type Questions

- (1) Molality is the no. of moles of solute dissolved in 1kg of solvent.
- (2) final Normality = $\frac{0.1 \times 100 + 0.2 \times 100}{200} = \frac{30}{200} = 0.15$
- (3) Both will form the vapours.
- (4) The composition of vapour phase is same as that of liquid phase.
- (5) Molecular mass of polymers cannot be calculated.
- (6) because water is polar and benzene is nonpolar.
- (7) Solutions having same osmotic pressure are called isotonic solutions.
- (8) Reverse osmosis is used in desalination of sea water.
- (9) Camphor is used as solvent in determination of molecular mass of naphthalene.
- (10) Both are colligative properties.
- (11) increase in surface area increases the rate of evaporation.
- (12) Due to strong intermolecular hydrogen bonding.
- (13) same as Ques. 10
- ~~(14) ~~Due to the addition of non-volatile solute vapour pressure~~~~
- (14) same as Ques. 3
- (15) Molar heat of vaporisation is the amount of heat required to vaporise one mole of liquid.
- (16) same as Ques. 14.
- (17) same as Ques. 6
- (18) use of pressure cooker reduces cooking time.
- (19) CCl_4 and H_2O are immiscible.
- (20) same as Ques. 7.
- (21) same as Ques. 13.

Previous year's Questions

Questions

$$(1) P_A = (P_A^{\circ} - P_B^{\circ}) X_A + P_B^{\circ}$$

$$(2) P = P_A^{\circ} X_A + P_B^{\circ} X_B$$

$$= \frac{200 \times 25.5}{119.5} + \frac{41.5 \times 40}{85}$$
$$= \frac{25.5}{119.5} + \frac{40}{85}$$

$$= 90.63 \text{ mmHg}$$

$$(3) \Delta T_b = i K_b m \quad \Delta T_b \propto i$$

NaCl will show higher elevation of b.p.

(4) Acetone + chloroform shows a negative deviation from Raoult's law.

$$(5) \frac{P^{\circ} - P}{P^{\circ}} = \frac{w}{40}$$

$$\frac{20}{100} = \frac{w}{40} + 1$$

or

$$\frac{5w}{40} = \frac{w}{40} + 1$$

$$\frac{4w}{40} = 1$$

$$w = 10 \text{ gm}$$

(6) Hypertonic

$$(7) \frac{P^{\circ} - P}{P^{\circ}} = \frac{w_B \times M_A}{M_B \times w_A}$$

$$\frac{P^{\circ} - P}{23.8} = \frac{100 \times 18}{342 \times 1000}$$

$$P^{\circ} - P = 0.125 \text{ mmHg}$$

$$(8) \begin{aligned} N &= M \times n \\ &= 3 \times 3 \\ &= 9 \end{aligned}$$

$$(9) \frac{\chi_{\text{Alcohol}}}{\chi_{\text{H}_2\text{O}}} = \frac{n_{\text{Alcohol}}}{n_{\text{H}_2\text{O}}} = \frac{138/46}{72/18} = \frac{3}{4}$$

(10) 1 M NaCl because $i = 2$

$$(11) \begin{aligned} \Delta T_f &= i K_f m \\ &= 2 \times 1.86 \times 1 \\ &= 3.72^{\circ}\text{C} \\ T_f &= -3.72^{\circ}\text{C} \end{aligned}$$

$$(6) \quad m = \frac{450 \times 10^{-3} \times 1000}{180 \times 100} = 0.025 \text{m}$$

$$(7) \quad N_1 V_1 = N_2 V_2$$

$$0.4 \times 2.5 = N_2 \times 0.5$$

$\times 10^{-3}$

$$N_2 = 0.002 \text{N}$$

(8) When 0.1 N NaOH is used

$$N_1 V_1 = N_2 V_2$$

(Hcl) (NaOH)

$$0.2 \times V_1 = 50 \times 0.1$$

$$V_1 = 25 \text{cm}^3$$

When 0.5 N KOH is used

$$N_1 V_1 = N_2 V_2$$

(remaining Hcl) (KOH)

$$0.2 \times 25 = 0.5 \times V_2$$

$$V_2 = 10 \text{cm}^3$$

(9) $N = M \times \text{m factor}$

$$N = \frac{4 \times 1000}{40 \times 100} = 1 \text{N}$$

(10) $N = M \times \text{m factor}$

$$0.2 = M \times 2$$

$$M = 0.1 \text{M}$$

(11) $0.25 = \frac{W}{60 \times 2.5}$

$$W = 37.5 \text{gm}$$

$$(12) \quad x_{C_6H_6} = \frac{n_{C_6H_6}}{n_{C_6H_6} + n_{CCl_4}} = \frac{\frac{30}{78}}{\frac{30}{78} + \frac{70}{154}} = 0.459$$

$$(13) \quad M = \frac{5 \times 1000}{40 \times 250} = 0.5M$$

$$(14) \quad N = \frac{1.575 \times 1000}{63 \times 250} = 0.1N$$

$H_2C_2O_4 \cdot 2H_2O$ is an organic acid which contains 2 molecules of water of crystallisation.

$$(15) \quad M_1V_1 + M_2V_2 = M_3V_3$$

$$M_3 = \frac{0.5 \times 800 + 1 \times 200}{1000}$$

$$M_3 = 0.6M$$

$$(16) \quad W = \frac{E \times N \times V}{1000}$$

$$E = \frac{0.126 \times 1000}{0.1 \times 20} = 63$$

$$(17) \quad N = \frac{N_1V_1 + N_2V_2 + N_3V_3}{V_1 + V_2 + V_3} = \frac{60 \times \frac{N}{5} + 10 \times \frac{N}{2} + 30 \times \frac{N}{10}}{60 + 10 + 30} = 0.2N$$

$$(18) \text{ Total no. of molecules} = \frac{95}{100} \times \frac{6.023 \times 10^{23}}{58.5} \times 10 = 0.98 \times 10^{23} \sim 10^{23}$$

(19) Molality and mole fraction depends only upon weight but not on volume.

$$(20) \% \text{ weight of NaOH} = \frac{W_{\text{NaOH}}}{W_{\text{solution}}} \times 100$$
$$= \frac{50}{1050} \times 100 = 4.76\%$$

1.25 molar NaOH means
1.25 moles of NaOH in 1000 gm
of H_2O
 $W_{\text{NaOH}} = 50 \text{ gm}$
 $W_{\text{solution}} = 1000 + 50$
 $= 1050 \text{ gm}$

$$(21) M = \frac{25}{250} \times \frac{1000}{100} = 1 \text{ M}$$

$$m = \frac{25}{250} \times \frac{1000}{1.25 \times 100} = 0.8 \text{ m}$$

$$(22) m = \frac{180}{180 \times 1} = 1 \text{ molar}$$

$$(23) N_1 V_1 = N_2 V_2$$

$$0.1 \times 10 = N_2 \times 15$$

$$N_2 = \frac{1}{15} = 0.066 \text{ N}$$

(24) Total volume of 1m solution $>$ 1000 cc due to extra 1 mole H_2SO_4 .
1m is less concentrated than 1M.

(25) Molarity (volume is temperature dependent)

$$(26) \quad \begin{array}{l} N_1 V_1 = N_2 V_2 \\ \text{Acid} \quad \quad \quad \text{Base} \end{array}$$

$$\frac{6.3 \times 100}{63 \times 250} \times 10 = 0.1 \times V_2$$

$$V_2 = 40 \text{ ml.}$$

$$(27) \quad N_1 V_1 = N_2 V_2$$

$$0.5 \times 100 = 0.1 \times V_2$$

$$V_2 = 500 \text{ cm}^3$$

$$V_{\text{H}_2\text{O}} = 500 - 100 = 400 \text{ cm}^3$$

$$(28) \quad N = \frac{98 \times 1.8 \times 10}{49} = 36 \text{ N}$$

$$N_1 V_1 = N_2 V_2$$

$$36 \times V_1 = 0.2 \times 1000$$

$$V_1 = 5.55 \text{ ml}$$

$$(29) \quad 10\% \text{ strength} = \frac{10 \text{ mole in } 100 \text{ cc.}}{180}$$

$$1 \text{ mole of glucose will be present in } = \frac{0.1 \times 180}{10} = 1.8 \text{ L}$$

$$(30) \quad N = M \times \text{m factor}$$

$$= 3 \times 3$$

$$= 9 \text{ N}$$

$$(31) N_1 V_1 + N_2 V_2 = N V$$

$$10V + 4(1-V) = 7 \times 1$$

$$V = 0.5L$$

$$\text{Volume of } 10N \text{ HCl} = 0.5L$$

$$\text{" " } 4N \text{ " " } = 1 - 0.5 = 0.5L$$

$$(32) X_{H_2O} = \frac{18/18}{\frac{18}{18} + \frac{414}{46}} = 0.1$$

$$(33) \text{ volume strength} = \frac{22400 \times 25.5}{68}$$
$$= 8400 \text{ ml}$$
$$= 8.4 \%$$

$$(34) N_1 V_1 = N_2 V_2$$
$$2 \times V = 1 \times 20$$
$$V = 10 \text{ ml}$$

$$(35) \text{ ppm} = \frac{25 \times 10^{-3} \times 10^6}{5000}$$
$$= 5$$

$$(36) 0.03 = \frac{W}{60}$$
$$W = 1.8 \text{ gm}$$

$$(37) \quad N \times V = \frac{w}{\epsilon}$$

$$0.1 \times V = \frac{1}{5}$$

$$V = \frac{1}{5} \times 1000 = 200 \text{ cm}^3$$

$$(38) \quad 0.1 = \frac{w \times 1000}{40 \times 250}$$

$$w = 1 \text{ gm}$$

$$(39) \quad M_1 V_1 = M_2 V_2$$

$$12 \times V_1 = 18 \times 240$$

$$V_1 = 360 \text{ ml}$$

$$V_1 = 0.36 \text{ L}$$

Topic - 2

Raoult's law (Ideal and Non ideal solution)

$$(1) P = P_A + P_B$$

$$P = P_A^\circ \chi_A + P_B^\circ \chi_B$$

$$P = P_A^\circ \chi_A + P_B^\circ - P_B^\circ \chi_A$$

$$P = P_B^\circ + \chi_A (P_A^\circ - P_B^\circ)$$

(2) Solution which shows negative deviation from Raoult's law are called maximum boiling azeotropes.

(3) When concentration of salt solution increases, vapour pressure decreases and Boiling point increases.

(4) With the addition of H_2O , concentration decreases thus vapour pressure increases.

(5) Only nonideal solution forms azeotropic mixtures - ($CCl_4 + CHCl_3$)

$$(6) \frac{\text{Loss in weight of solvent}}{\text{gain in weight of } CaCl_2 \text{ tube}} = \frac{P^\circ - P}{P^\circ}$$

(7) $He + O_2$ mixture is used by deep sea divers because its very low solubility in blood.

$$(8) \frac{P - P_s}{P} = \frac{n}{n + N}$$

$$(9) \quad \frac{P^{\circ} - P_s}{P^{\circ}} = X_{\text{solute}}$$

$$\frac{0.8 - 0.6}{0.8} = X_{\text{solute}}$$

$$X_{\text{solute}} = \frac{0.2}{0.8} = 0.25$$

(10) $\text{CH}_3\text{CH}_2\text{OH} + \text{CH}_3\text{COCH}_3$ is nonideal & shows positive deviation.

$$(11) \quad P = P_A^{\circ} X_A + P_B^{\circ} X_B$$

$$84 = 0.8 \times 70 + 0.2 \times P_B^{\circ}$$

$$P_B^{\circ} = 140 \text{ mm}$$

$$(12) \quad 500 = \frac{1}{3} P_A^{\circ} + \frac{2}{3} P_B^{\circ}$$

$$P_A^{\circ} + 2P_B^{\circ} = 1500$$

$$500 + 25 = \frac{1}{4} P_A^{\circ} + \frac{3}{4} P_B^{\circ}$$

$$P_A^{\circ} + 3P_B^{\circ} = 2100$$

$$P_A^{\circ} = 300 \text{ mmHg}$$

$$P_B^{\circ} = 600 \text{ mmHg}$$

$$(12) \quad C = K_H \times P$$

$$= 1.4 \times 10^{-3} \times 0.5 = 7 \times 10^{-4}$$

$$\text{in } 100 \text{ ml} = 7 \times 10^{-5}$$

$$\begin{aligned} \text{weight of } \text{O}_2 \text{ in } 100 \text{ ml} &= 7 \times 10^{-5} \times 32 \\ &= 2.24 \times 10^{-3} \text{ gm} \end{aligned}$$

$$(13) \quad \chi_B = \frac{1}{56.5} \Rightarrow \frac{\Delta P}{P} = \chi_B$$

$$\Delta P = P \chi_B$$

$$= \frac{760 \times 1}{56.5} = 13.45 \text{ torr}$$

(14) In equimolar mixture

$$\chi_A = 0.5, \chi_B = 0.5$$

$$P = 0.5 \times 150 + 0.5 \times 150$$

$$P = 125$$

$$\gamma_B = \frac{\chi_B P_B^0}{P} = \frac{0.5 \times 100}{125} = 0.4$$

(15) Chloroform-acetone solution shows negative deviation from Raoult's law.

(16) Methanol is a volatile substance, so its aqueous solution in water has more vapour pressure than methanol.

$$(17) \quad \frac{P^0 - P}{P^0} = \chi_B = 0.2$$

$$(18) \quad 184 = 200 \times \frac{3}{3+2} + P_B^0 \times \frac{2}{3+2}$$

$$P_B^0 = 160 \text{ torr.}$$

(19) Chloroform + Acetone ~~is~~ shows negative deviation from Raoult's law. $\Delta V_{\text{mix}} = -ve$.

$$(20) \quad \chi_A = \frac{P_A^0 \chi_A}{P_A^0 \chi_A + P_B^0 \chi_B}$$
$$= \frac{1 \times 1}{1 \times 1 + 2 \times 2} = \frac{1}{5} = 0.2$$

(21) An ideal solution have zero volume change and zero heat of mixing.

Topic-3

Colligative properties, Abnormal Molecular mass and Distribution law

$$(1) \quad \Delta T_f = i K_f m$$

$$\Delta T_f = \frac{i \times K_f \times w_B \times 1000}{M_B \times w_A}$$

$$3.82 = \frac{i \times 1.86 \times 5 \times 1000}{142 \times 45}$$

$$i = 2.63$$

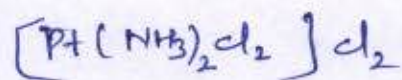
(2) Dissociation, $i > 1$

Association, $i < 1$

$$(3) \quad \Delta T_f = i K_f m$$

$$0.0054 = i \times 1.86 \times 0.001$$

$$i = 3$$



$$(4) \quad \pi_1 = \pi_2$$

$$\frac{w_1}{M_1} = \frac{w_2}{M_2}$$

$$\frac{3}{M_1} = \frac{0.05 \times 180}{180}$$

$$M_1 = 60$$

$$n = \frac{60}{30} = 2 \quad (\text{CH}_2\text{O})_2 = \text{C}_2\text{H}_4\text{O}_2$$

(5) Two sucrose solution of same molality will have different depression of freezing point.

$$(6) \quad \Delta T_f = \frac{1000 \times K_f \times w_2}{M_2 \times w_1} = \frac{1000 \times 1.86 \times 68.5}{342 \times 1000} = 0.372^\circ\text{C}$$

$$\Delta T_f = T^\circ - T_1 \\ = 0 - 0.372 = -0.372^\circ\text{C}$$

(7) Concentration of BaCl_2 solution is lower, thus solvent moves from BaCl_2 to NaCl solution.

(8) $i < \frac{1}{\text{Concentration}}$

$$(9) \quad \Delta T_f = \frac{5.12 \times 1000 \times 0.223}{36 \times 4.4} = 1.91^\circ\text{C}$$

(Theoretical)

$$\text{(Actual)} \Delta T_f = 5.3 - 4.47 \\ = 0.83^\circ\text{C}$$

$$i = \frac{0.83}{1.91} = 0.43$$

$$i < 1$$

Association will take place.

(10) Sucrose \rightarrow no ion

$\text{NaCl} \rightarrow \text{Na}^+ + \text{Cl}^-$

$\text{CaCl}_2 \rightarrow \text{Ca}^{2+} + 2\text{Cl}^-$

glucose \rightarrow no ion.

$\% \text{CaCl}_2$ has least freezing point.

$$(11) \quad \Delta T_f = K_f \times m$$

$$0.01 = \frac{K_f \times 1.5 \times 1000}{60 \times 250}$$

$$K_f = 0.1$$

(12) isotonic solution have same osmotic pressure.

$$(13) \quad \Delta T_b = \frac{0.52 \times 0.6 \times 1000}{60 \times 100} = 0.052^\circ\text{C}$$

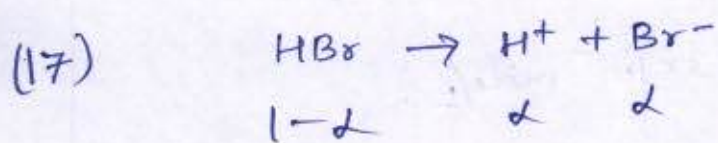
$$\Delta T_b = T_1 - T_0$$

$$0.052 = T_1 - 100 \Rightarrow T_1 = 100.052^\circ\text{C}$$

(14) egg will shrink because osmosis occurs. ~~shrink~~

$$(15) \quad \Delta T_b = \frac{1000 \times K_b \times 10}{100 \times 100} = K_b$$

(16) H_2SO_4 gives 3 ions & NaCl gives 2 ions. vapour pressure of solution of NaCl is higher.



$$\text{total ions} = 1-d + d + d = 1+d. \quad i = 1+d$$

$$= 1 + \frac{90}{100} = 1.9$$

$$\Delta T_f = i K_f m$$

$$= 1.9 \times 1.86 \times 8 \frac{1}{81}$$

$$= \frac{\quad}{100/1000}$$

$$= 3.534^\circ\text{C}$$

$$\Delta T_f = T_0 - T_1$$

$$3.534 = 0 - T_1 \Rightarrow T_1 = 3.534^\circ\text{C}$$

(18) freezing point of pure solvent decreases on the addition of solute.

$$\begin{aligned}(19) \quad \Delta T_b &= i K_b m \\ &= 2 \times 0.52 \times 1 \\ &= 1.04\end{aligned}$$

$$\Delta T_b = T_1 - T_0$$

$$1.04 = T_1 - 100$$

$$T_1 = 101.04^\circ\text{C}$$

(20) atm

(21) optical activity

$$(22) \quad \Delta T_f = i K_f m$$

$$0.00732 = i \times 1.86 \times 0.002$$

$$i = 2.$$

$$(23) \quad \pi = CRT$$

$$C = \frac{\pi}{RT} = \frac{0.0821}{0.0821 \times 300} = 0.33 \times 10^{-2} \text{ mole/L}$$

(24) no. of particles increases, boiling point increases & freezing point decreases
~~AB~~ $D < B > A < C$

(25) $\text{Ca}(\text{NO}_3)_2$ & Na_2SO_4 both will give equal ions.

(26)	KNO_3	2	$\Delta T_b = i K_b m$
	Na_3PO_4	4	
	BaCl_2	3	
	K_2SO_4	3	
		3	

$$(27) \quad \chi_{\text{glucose}} = \chi_{\text{urea}}$$

$$\frac{x \times 1000}{180 \times 100} = \frac{6 \times 1000}{60 \times 100}$$

$$x = 18 \text{ gm}$$

(28) CaCl_2 act as non volatile solute so results in depression in freezing point.

(29) When freezing starts, liquid solvent is in equilibrium with solid solvent.

$$(30) \quad \Delta T_f = i K_f m \\ = 1.2 \times 1.86 \times 0.2 = 0.45^\circ\text{C}$$

$$\Delta T_f = T_0 - T_1$$

$$0.45 = 0 - T_1$$

$$T_1 = -0.45^\circ\text{C}$$

$$(31) \quad i = \frac{40}{25}$$

$$1 + \alpha = \frac{40}{25}$$

$$\alpha = \frac{15}{25} = 0.6 \text{ or } 60\%$$

$$(32) \quad \Delta T_f = K_f \times m$$

$$1.01 = \frac{1.93 \times 20 \times 1000}{M \times 500}$$

$$M = 76.4$$

$$(33) \frac{\Delta T_f(\text{KCl})}{\Delta T_f(\text{BaCl}_2)} = \frac{2}{3}$$

$$\Delta T_f(\text{BaCl}_2) = \frac{3}{2} \times 2 = 3^\circ\text{C} \text{ v } -3^\circ\text{C}$$

(34) by this method lowering in vapour pressure, vapour pressure of solvent and relative lowering of vapour pressure all can be measured.

$$(35) \pi = \frac{wRT}{Mv}$$

$$T = \frac{16.4 \times 180 \times 100}{10 \times 1000 \times 0.082} = 360\text{K}$$

(36) $i > 1$ dissociation.

(37) A & B

A 0.1M glucose $\rightarrow 0.1 \times 1 = 0.1$

B 0.05M NaCl $\rightarrow 0.05 \times 2 = 0.1$

isotonic means same molar concentration.

$$(38) \frac{p^\circ - p}{p^\circ} = \frac{w \times M}{M \times w}$$

$$\frac{640 - 600}{640} = \frac{2.175 \times 78}{m \times 39.08}$$

$$m = 69.60$$



3 ions.

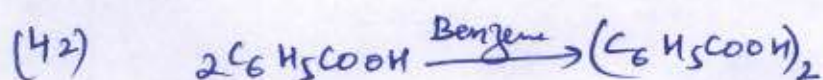
$$(40) \frac{P^{\circ} - P}{P^{\circ}} = i \chi_B$$

i is maximum for 0.1M BaCl_2

$$(41) \pi V = nRT$$

$$\pi V = \frac{W}{M} RT$$

$$M = \frac{4 \times 0.0821 \times 300}{4 \times 10^{-4}} = 2.4 \times 10^5$$



$$i = \frac{1}{2} = 0.5$$

$$(43) \pi V = nRT$$

$$\pi V = \frac{m_2}{M_2} RT$$

$$M_2 = \frac{m_2 RT}{\pi V}$$

(44) Freezing point will be highest due to lower concentration of NaCl.

(45) Camphor is used in molecular mass determination due to its volatile nature.

$$(46) \Delta \propto \frac{R_0 - P}{R_0}$$

$$\frac{30 - P}{30} = 0.9$$

$$P = 30 - 3 = 27$$

(47)

$$\frac{\Delta T_{f1}}{\Delta T_{f2}} = \frac{M_2}{M_1}$$

(Urea)
Molar concentration (M_1) = $\frac{10}{60} = \frac{1}{6}$

Non volatile solute

$$M_2 = \frac{50}{m}$$

$$\frac{1}{6} = \frac{50}{m}$$

$$m = 300 \text{ gm/mol}$$

(48)

$$\frac{\Delta T_{f1}}{\Delta T_{f2}} = \frac{M_2}{M_1}$$

$$\frac{2.15}{\Delta T_{f2}} = \frac{180}{342}$$

$$\Delta T_{f2} = 4.085\text{K or } 269.05\text{K}$$

(49) Osmotic pressure method is especially suitable for determination of molecular masses.

(50)

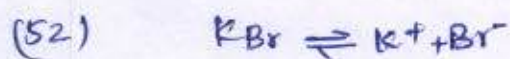
$$\frac{P - P_s}{P} = \frac{W_1 \times M_2}{M_1 \times W_2}$$

$$\frac{W_{\text{glucose}} \times 18}{50 \times 180} = \frac{W_{\text{urea}} \times 18}{50 \times 60}$$

$$W_{\text{glucose}} = 3$$

$$(51) \quad \frac{P - P_s}{P} \times \frac{1000}{M} = \text{molality (m)}$$

$$m = \frac{25 - 24.5}{25} \times \frac{1000}{18} = 1.11 \text{ m}$$



$$i = \frac{1 - 0.8 + 2 \times 0.8}{1} = 1.8$$

$$\Delta T_f = 1.8 \times 1.86 \times 0.5 = 1.674 \text{ K}$$

$$T_f = 271.326 \text{ K}$$

(53) Aqueous solution of any substance (non volatile) freezes below 0°C because vapour pressure of solution becomes lower than that of pure solvent.

$$(54) \quad \frac{\Delta T_f}{\Delta T_b} = \frac{K_f}{K_b}$$

$$\frac{\Delta T_f}{0.18} = \frac{1.86}{0.512}$$

$$\Delta T_f = 0.654$$

$$\Delta T_b - T_1 = 0.654$$

$$T_1 = -0.654^\circ\text{C}$$

$$(55) \quad \pi = CRT$$

$$C = \frac{7.8}{0.0821 \times 310} = 0.31 \text{ mole/l}$$

$$(56) \quad K_b = \frac{M_1 R T_0^2}{1000 \Delta_{\text{vap}} \times H}$$

$$= \frac{18 \times 0.002 \times (373.15)^2}{1000 \times 9.72} = 0.52 \text{ K/mole}$$

(57) Distribution law was given by Nernst.

$$(58) \quad \Delta T_f = \frac{1000 \times K_f \times W}{M \times W}$$

$$0.74 = \frac{1000 \times 1.86 \times 20}{M \times 500}$$

$$M = 100.$$

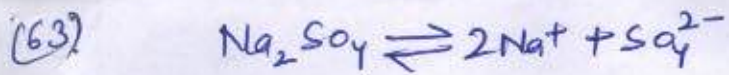
So, degree of ionisation of electrolyte 0%.

(59) The molal depression constant (K_f) for camphor is maximum. So, depression of freezing point will also be maximum.

$$(60) \quad \chi_{\text{solute}} = \frac{760 - 750}{760} = \frac{1}{76}$$

(61) Naphthalene is a nonvolatile solid, hence on addition in benzene decreases its vapour pressure.

$$(62) \quad \chi_A = \frac{0.8 - 0.2}{0.8} = \frac{0.6}{0.8} = \frac{3}{4} = 0.75$$



$$i = 1 - \alpha + 2\alpha + \alpha$$

$$i = 1 + 2\alpha$$

(64) At high altitude, the boiling point of water is lower because atmospheric pressure is low.

(65) In glucose $i=1$ while others are electrolyte $i > 1$

$$\Delta T_f = i K_f m$$

ΔT_f is lowest for glucose.

(66)
$$2.74 = \frac{0.1 - \alpha + \alpha + 2\alpha}{0.1}$$

$$\alpha = 0.087 \text{ or } 8.7\%$$

$$\text{degree of dissociation} = \frac{0.087}{0.1} \times 100 = 87\%$$