



Liquid Solution PYQS-2022

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1. The depression in freezing point observed for a formic acid solution of concentration 0.5 mL L^{-1} is 0.0405°C . Density of formic acid is 1.05 g mL^{-1} . The Van't Hoff factor of the formic acid solution is nearly : (Given for water $k_f = 1.86 \text{ K kg mol}^{-1}$)

(A) 0.8

(B) 1.1

(C) 1.9

(D) 2.4

$$\Delta T_f = i \times K_b \times m$$

$$\begin{aligned} \text{mass of formic acid solution} &= V \times d \\ &= 0.5 \times 1.05 \\ &= 0.525 \text{ g in } 1 \text{ L} \end{aligned}$$

$$\text{molality} = \frac{0.525}{46}$$

$$\Delta T_f = 0.0405 = \frac{i \times 1.86 \times 0.525}{46}$$

$$i = \frac{0.0405 \times 460}{0.525 \times 1.86} = \frac{405 \times 460}{525 \times 186}$$

$$i = \underline{\underline{1.90}}$$

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2. Two solutions A and B are prepared by dissolving 1 g of non-volatile solutes X and Y, respectively in 1 kg of water. The ratio of depression in freezing points for A and B is found to be 1:4. The ratio of molar masses of X and Y is:

(A) 1:4

✓ (B) 1:0.25

(C) 1:0.20

(D) 1:5

$$\Delta T_{fA} = i \times K_{fA} \times m_A$$

$$\Delta T_{fB} = i \times K_{fA} \times m_B$$

$$\frac{1}{4} = \frac{m_A}{m_B} = \frac{\cancel{w_A}}{m_A} \times \frac{m_B}{\cancel{w_B}}$$

↳ molality

$$\frac{1}{4} = \frac{m_B}{m_A} \rightarrow \text{molar mass.}$$

$$\frac{1}{4} = \frac{Y}{X}$$

$$\frac{X}{Y} = \frac{4}{1}$$

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3. Boiling point of a 2% aqueous solution of a nonvolatile solute A is equal to the boiling point of 8% aqueous solution of a non-volatile solute B. The relation between molecular weights of A and B is.

(A) $M_A = 4M_B$

☒ (B) $M_B = 4M_A$

(C) $M_A = 8M_B$

(D) $M_B = 8M_A$

$$\begin{aligned} \cancel{\Delta T_A} &= \cancel{i} \times \cancel{K_b} \times \frac{2}{M_A \times 98} \times 1000 \\ \cancel{\Delta T_B} &= \cancel{i} \times \cancel{K_b} \times \frac{8}{M_B \times 92} \times 1000 \\ \frac{4}{M_B \times 92} \times 1000 &= \frac{2}{M_A \times 98} \times 1000 \\ \frac{4}{23} &= \frac{M_A \times 98}{M_B \times 92} \\ M_B \times 23 &= M_A \times 98 \\ B &= \underline{\underline{4M_A}} \text{ (nearest)} \end{aligned}$$

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4. Solute A associates in water. When 0.7 g of solute A is dissolved in 42.0 g of water, it depresses the freezing point by 0.2°C . The percentage association of solute A in water, is [Given : Molar mass of A = 93 g mol^{-1} . Molal depression constant of water is $1.86 \text{ K kg mol}^{-1}$]
- (A) 50% (B) 60% (C) 70% ☒ (D) 80%

$$\Delta T_f = i \times K_f \times m$$

$$0.2 = i \times 1.86 \times \frac{0.7}{42 \times 93} \times 1000$$

$$i = \frac{42 \times 93 \times 0.2}{0.7 \times 1.86 \times 1000}$$

$$= \frac{84 \times 93}{7 \times 186} \times 0.1$$

$$i = \underline{\underline{0.6}}$$



$$1 - \alpha + \alpha/2 = 0.6$$

$$1 - \alpha/2 = 0.6$$

$$\alpha = \underline{\underline{0.80}}$$

$$\alpha = \underline{\underline{80\%}}$$

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5. When a certain amount of solid A is dissolved in 100 g of water at 25°C to make a dilute solution, the vapour pressure of the solution is reduced to one-half of that of pure water. The vapour pressure of pure water is 23.76 mmHg. The number of moles of solute A added is .
(Nearest Integer) Assume moles of A to be less than moles of B

exact

$$\frac{P_A^\circ - P_A}{P_A^\circ} = \frac{n_B}{n_A}$$

$$\frac{23.76 - \frac{23.76}{2}}{23.76} = \frac{n_B \times 18}{100}$$

$$n_B = \frac{100}{18} = \underline{\underline{5.55}}$$

$$\frac{P_A^\circ - P_A}{P_A^\circ} = \frac{n_B}{n_A} \quad (\text{not exact})$$

$$\frac{23.76 - \frac{23.76}{2}}{23.76} = \frac{n_B \times 18}{100}$$

$$\frac{23.76}{2 \times 23.76} = \frac{n_B}{5.55}$$

$$n_B = \frac{5.55}{2} = 2.78$$

$$= \underline{\underline{3}}$$

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6. 150 g of acetic acid was contaminated with 10.2 g ascorbic acid ($C_6H_8O_6$) to lower down its freezing point by $(x \times 10^{-1})^\circ C$. The value of x is (Nearest integer) [Given $K_f = 3.9 K kg mol^{-1}$; Molar mass of ascorbic acid = $176 g mol^{-1}$]

15

$$\Delta T_f = i \times K_f \times m$$

$$= 1 \times 3.9 \times \frac{10.2}{176 \times 150} \times 1000$$

$$= 1.50 = \underline{\underline{15 \times 10^{-1}}}$$

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7. A gaseous mixture of two substances A and B, under a total pressure of 0.8 atm is in equilibrium with an ideal liquid solution. The mole fraction of substance A is 0.5 in the vapour phase and 0.2 in the liquid phase. The vapour pressure of pure liquid A is atm. (Nearest integer)

5

$$P_T = 0.8 \text{ atm}$$

$$y_A = 0.5$$

$$y_B = 0.5$$

$$x_A = 0.2$$

$$x_B = 0.8$$

$$y_A = \frac{x_A P_A^\circ}{P_T}$$

$$0.5 = \frac{0.2 \times P_A^\circ}{0.8}$$

$$P_A^\circ = \frac{0.5 \times 0.8}{0.2} = \underline{\underline{2 \text{ atm}}}$$

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8. If O_2 gas is bubbled through water at 303 K, the number of millimoles of O_2 gas that dissolve in 1 litre of water is . (Nearest Integer) (Given : Henry's Law constant for O_2 at 303 K is 46.82 k bar and partial pressure of $O_2 = 0.920$ bar) (Assume solubility of O_2 in water is too small, nearly negligible)

$$p_A = K_H \times X_A$$

$$0.920 = 46.82 \times \frac{n_{O_2}}{n_{H_2O}}$$

$$.920 = 46.82 \times \frac{n_{O_2}}{\frac{1000}{18}}$$

$$\begin{aligned} n_{O_2} &= \frac{.920 \times 1000}{46.82 \times 18} \\ &= 1.09 \times 10^{-3} \text{ moles} \\ &= \underline{\underline{1 \text{ millimole}}} \end{aligned}$$

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9. 'x' g of molecular oxygen (O_2) is mixed with 200 g of neon (Ne). The total pressure of the non-reactive mixture of O_2 and Ne in the cylinder is 25 bar. The partial pressure of Ne is 20 bar at the same temperature and volume. The value of 'x' is [Given: Molar mass of $O_2 = 32 \text{ g mol}^{-1}$. Molar mass of Ne = 20 g mol^{-1}]

80

$$P_{Ne} = 20 \text{ bar}$$

$$P_T = 25 \text{ bar}$$

$$P_{O_2} = 25 - 20 = 5 \text{ bar}$$

$$P_{O_2} = \chi_{O_2} \times P_T$$

$$5 = \left(\frac{\frac{x}{32}}{\frac{x}{32} + \frac{200}{20}} \right) \times 25$$

$$1 = \frac{\frac{x}{32}}{\frac{x}{32} + \frac{200}{20}} \times 5$$

$$x + 320 = 5x$$

$$x = \underline{\underline{80 \text{ g}}}$$

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10. 1.80 g of solute A was dissolved in 62.5 cm³ of ethanol and freezing point of the solution was found to be 155.1 K. The molar mass of solute A is g mol^{-1} .

[Given: Freezing point of ethanol is 156.0 K. Density of ethanol is 0.80 g cm^{-3} .

80 Freezing point depression constant of ethanol is $2.00 \text{ K kg mol}^{-1}$]

$$\Delta T_f = i \times K_f \times m$$

$$0.9 = i \times 2 \times \frac{1.80 \times 1000}{m \times \underbrace{62.5 \times 0.8}_{\text{mass of solvent}}}$$

$$m = \frac{2 \times \overset{20}{\cancel{180}} \times 1000}{\cancel{9} \times 62.5 \times \cancel{8}} = \frac{\overset{5000}{\cancel{4000}}}{62.5 \times \cancel{8}} = \underline{\underline{80}}$$

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11. The osmotic pressure of blood is 7.47 bar at 300 K. To inject glucose to a patient intravenously, it has to be isotonic with blood. The concentration of glucose solution in g L^{-1} is 54 (Molar mass of glucose = 180 g mol^{-1} , $R = 0.083 \text{ L bar}^{-1} \text{ mol}^{-1}$) (Nearest integer)

$$\pi = i \times C \times R \times T$$

$$7.47 = 1 \times \frac{w}{180} \times 0.083 \times 300$$

$$w = \frac{7.47 \times 180}{0.083 \times 300} = \underline{\underline{54 \text{ g/L}}}$$

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12. A company dissolves 'X' amount of CO_2 at 298 K in 1 litre of water to prepare soda water $X = \underline{\hspace{1cm}} \times 10^{-3}$ g. (nearest integer) (Given: partial pressure of CO_2 at 298 K = 0.835 bar. Henry's law constant for CO_2 at 298 K = 1.67 k bar. Atomic mass of H, C and O is 1, 12 and 16 g mol^{-1} , respectively)

1223

$$p = K_H \times X$$

$$0.835 = 1.67 \times 10^3 \times \frac{n_{\text{CO}_2}}{n_{\text{CO}_2} + n_{\text{H}_2\text{O}}}$$

$$\cancel{0.835} = \cancel{1.67} \times 10^3 \times \frac{w}{44} \div \frac{w}{44} + 55.55$$

$$\frac{1}{2} = \frac{w \times 1000}{w + 44 \times 55.55}$$

$$2000w = w + 44 \times 55.55$$

$$1999w = 44 \times 55.55$$

$$w = \frac{2444.2}{1999} = 1.2227 \text{ g} \\ = 1222.7 \times 10^{-3} \text{ g} \\ = \underline{\underline{1223}}$$

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13. The elevation in boiling point for 1 molal solution of non-volatile solute A is 3 K. The depression in freezing point for 2 molal solution of A in the same solvent is 6 K. The ratio of K_b and K_f i.e., K_b/K_f is 1: X. The value of X is [nearest integer]

①

$$\Delta T_b = 3K = K_b \times 1$$

$$\Delta T_f = 6K = K_f \times 2$$

$$\frac{3}{6} = \frac{K_b}{K_f} \times \frac{1}{2}$$

$$K_f = K_b$$

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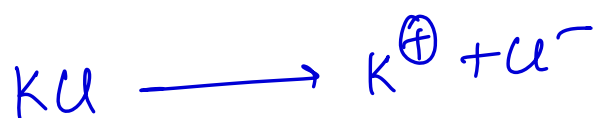
14. A 0.5 percent solution of potassium chloride was found to freeze at -0.24°C . The percentage dissociation of potassium chloride is (Nearest integer) (Molal depression constant for water is $1.80 \text{ K kg mol}^{-1}$ and molar mass of KCl is 74.6 g mol^{-1})

98%

$$\Delta T_f = i \times K_f \times m$$

$$0.24 = i \times 1.80 \times \frac{0.5}{74.6 \times 99.5} \times 1000$$

$$i = 1.98$$



$$\begin{array}{ccc} 1 & 0 & 0 \\ 1-\alpha & \alpha & \alpha \end{array}$$

$$1 + \alpha = 1.98$$

$$\alpha = 0.98$$

$$\alpha = \underline{\underline{98\%}}$$

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15. The osmotic pressure exerted by a solution prepared by dissolving 2.0 g of protein of molar mass 60 kg mol^{-1} in 200 mL of water at 27°C is Pa. [integer value]

415

$$\pi = CRT$$

$$= \frac{2}{60 \times 10^3} \times \frac{1000}{200} \times 0.083 \times 300$$

$$= \frac{600}{60 \times 20} \times 0.083$$

$$\underline{1 \text{ bar} = 10^5 \text{ Pa}}$$

$$= 41.5 \times 10^{-4} \text{ bar}$$

$$= 41.5 \times 10^{-4} \times 10^5 \text{ Pa}$$

$$= \underline{\underline{415 \text{ Pa}}}$$

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16.
8

2 g of a non-volatile non-electrolyte solute is dissolved in 200 g of two different solvents A and B whose ebullioscopic constants are in the ratio of 1:8. The elevation in boiling points of A and B are in the ratio $\frac{x}{y}$ ($x:y$). The value of y is ___ (Nearest integer)

$$\Delta T_{bA} = 1 \times K_{bA} \times \frac{2}{200 \times m} \times 1000$$

$$\Delta T_{bB} = 1 \times K_{bB} \times \frac{2}{200 \times m} \times 1000$$

$$\frac{\Delta T_{bA}}{\Delta T_{bB}} = \frac{K_{bA}}{K_{bB}} = \frac{1}{8}$$

$$\frac{x}{y} = \frac{1}{8}$$

$$y = \underline{\underline{8}}$$

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17.

45

A solution containing 2.5×10^{-3} kg of a solute dissolved in 75×10^{-3} kg of water boils at 373.535 K. The molar mass of the solute is __ g mol^{-1} . [nearest integer] (Given: $K_b(\text{H}_2\text{O}) = 0.52 \text{ K kg mol}^{-1}$, boiling point of water = 373.15 K)

$$\Delta T_b = \frac{373.535 - 373.150}{0.385}$$

$$0.385 = \frac{2.5 \times 1000 \times 0.52}{75 \times M}$$

$$M = \frac{2.5 \times 0.52 \times 1000}{0.385 \times 75}$$

$$M = \frac{45.02}{1} \approx \underline{\underline{45}}$$

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18. The vapour pressures of two volatile liquids A and B at 25°C are 50 Torr and 100 Torr, respectively. If the liquid mixture contains 0.3 mole fraction of A, then the mole fraction of liquid B in the vapour phase is $\frac{x}{17}$. The value of x is

14

$$P_A^0 = 50 \text{ torr} \quad P_B^0 = 100 \text{ torr}$$

$$X_A = 0.3 \quad X_B = 0.7$$

$$Y_B = \frac{X_B P_B^0}{P_T}$$

$$= \frac{0.7 \times 100}{50 \times 0.3 + 0.7 \times 100} = \frac{70}{15 + 70} = \frac{14}{85}$$

$$Y_B = \frac{14}{17} \Rightarrow x = \underline{\underline{14}}$$

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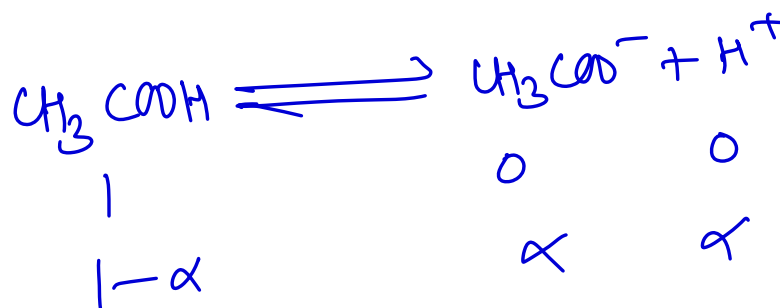
19. 1.2 mL of acetic acid is dissolved in water to make 2.0 L of solution. The depression in freezing point observed for this strength of acid is 0.0198°C . The percentage of dissociation of the acid is (Nearest integer) [Given : Density of acetic acid is 1.02 g mL^{-1} Molar mass of acetic acid is 60 g mol^{-1} $K_f(\text{H}_2\text{O}) = 1.85 \text{ K kg mol}^{-1}$]

5

$$0.0198 = i \times 1.85 \times \frac{1.2 \times 1.02}{60 \times 2}$$

$$i = \frac{0.0198 \times 600 \times 2}{1.85 \times 1.2 \times 1.02}$$

$$i = \frac{198 \times 600 \times 2}{185 \times 12 \times 102} = 1.05$$



$$1 + \alpha = 1.05$$

$$\alpha = .05$$

$$\alpha = \underline{\underline{5\%}}$$

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20. Elevation in boiling point for 1.5 molal solution of glucose in water is 4 K. The depression in freezing point for 4.5 molal solution of glucose in water is 4 K. The ratio of molal elevation constant to molal depression constant (K_b/K_f) is..

3

$$4K = K_b \times 1.5$$

$$4K = K_f \times 4.5$$

$$\frac{K_b}{K_f} \times \frac{1}{3} = 1$$

$$\frac{K_b}{K_f} = \underline{\underline{3}}$$