Chapter 10

Solutions

- 1. A binary liquid solution is prepared by mixing n-heptane and ethanol. Which one of the following statements is correct regarding the behaviour of the solution? [AIEEE-2009]
 - The solution is non-ideal, showing +ve deviation from Raoult's Law
 - (2) The solution is non-ideal, showing –ve deviation from Raoult's Law
 - (3) n-heptane shows +ve deviation while ethanol shows –ve deviation from Raoult's Law
 - (4) The solution formed is an ideal solution
- Two liquids X and Y from an ideal solution. At 300 K, vapour pressure of the solution containing 1 mol of X and 3 mol of Y is 550 mmHg. At the same temperature, if 1 mol of Y is further added to this solution, vapour pressure of the solution increases by 10 mmHg. Vapour pressure (in mmHg) of X and Y in their pure states will be, respectively

[AIEEE-2009]

- (1) 300 and 400
- (2) 400 and 600
- (3) 500 and 600
- (4) 200 and 300
- 3. If sodium sulphate is considered to be completely dissociated into cations and anions in aqueous solution, the change in freezing point of water (ΔT_f) , when 0.01 mol of sodium sulphate is dissolved in 1 kg of water, is $(K_f = 1.86 \text{ K kg mol}^{-1})$

[AIEEE-2010]

- (1) 0.0186 K
- (2) 0.0372 K
- (3) 0.0558 K
- (4) 0.0744 K
- 4. On mixing, heptane and octane from an ideal solution. At 373 K, the vapour pressures of the two liquid components (heptane and octane) are 105 kPa and 45 kPa respectively. Vapour pressure of the solution obtained by mixing 25.0 g of heptane and 35 g of octane will be (molar mass of heptane = 100 g mol⁻¹ and of octane = 114 g mol⁻¹)

[AIEEE-2010]

- (1) 144.5 kPa
- (2) 72.0 kPa
- (3) 36.1 kPa
- (4) 96.2 kPa

- 5. A 5% solution of cane sugar (molar mass 342 is isotonic with 1% of a solution of an unknown solute. The molar mass of unknown solute in g/mol is [AIEEE-2011]
 - (1) 34.2

- (2) 136.2
- (3) 171.2
- (4) 68.4
- 6. K_f for water is 1.86 K kg mol⁻¹. If your automobile radiator holds 1.0 kg of water, how many grams of ethylene glycol (C₂H₆O₂) must you add to get the freezing point of the solution lowered to 2.8°C?

[AIEEE-2012]

- (1) 93 g
- (2) 39 g
- (3) 27 g
- (4) 72 g
- 7. Consider separate solutions of 0.500 M $C_2H_5OH(aq)$, 0.100 M $Mg_3(PO_4)_2(aq)$, 0.250 M KBr(aq) and 0.125 M $Na_3PO_4(aq)$ at 25°C. Which statement is true about these solutions, assuming all salts to be strong electrolytes?

[JEE (Main)-2014]

- (1) They all have the same osmotic pressure.
- (2) 0.100 M ${\rm Mg_3(PO_4)_2(aq)}$ has the highest osmotic pressure.
- (3) 0.125 M Na₃PO₄(aq) has the highest osmotic pressure.
- (4) 0.500 M C₂H₅OH(aq) has the highest osmotic pressure.
- 8. The vapour pressure of acetone at 20°C is 185 torr. When 1.2 g of a non-volatile substance was dissolved in 100 g of acetone at 20°C, its vapour pressure was 183 torr. The molar mass (g mol⁻¹) of the substance is [JEE (Main)-2015]
 - (1) 32

(2) 64

(3) 128

- (4) 488
- 18 g glucose (C₆H₁₂O₆) is added to 178.2 g water. The vapor pressure of water (in torr) for this aqueous solution is [JEE (Main)-2016]
 - (1) 76.0
- (2) 752.4
- (3) 759.0
- (4) 7.6

- 10. The freezing point of benzene decreases by 0.45°C when 0.2 g of acetic acid is added to 20 g of benzene. If acetic acid associates to form a dimer in benzene, percentage association of acetic acid in benzene will be (K_f for benzene = 5.12 K kg mol^{-1}) [JEE (Main)-2017]
 - (1) 74.6%
- (2) 94.6%
- (3) 64.6%
- (4) 80.4%
- 11. Which one of the following statements regarding Henry's law is not correct? [JEE (Main)-2019]
 - (1) Different gases have different K_H (Henry's law constant) values at the same temperature
 - (2) The value of K_{\perp} increases with increase of temperature and K_H is function of the nature of the gas
 - (3) The partial pressure of the gas in vapour phase is proportional to the mole fraction of the gas in the solution
 - (4) Higher the value of K_H at a given pressure, higher is the solubility of the gas in the liquids.
- 12. Liquids A and B form an ideal solution in the entire composition range. At 350 K, the vapor pressures of pure A and pure B are 7 × 103 Pa and 12 × 103 Pa, respectively. The composition of the vapor in equilibrium with a solution containing 40 mole percent of A at this temperature is

[JEE (Main)-2019]

- (1) $x_A = 0.76$; $x_B = 0.24$ (2) $x_A = 0.37$; $x_B = 0.63$
- (3) $x_A = 0.28$; $x_B = 0.72$ (4) $x_A = 0.4$; $x_B = 0.6$
- 13. Elevation in the boiling point for 1 molal solution of glucose is 2 K. The depression in the freezing point for 2 molal solution of glucose in the same solvent is 2 K. The relation between K_b and K_f is

[JEE (Main)-2019]

- (1) $K_b = 0.5 K_f$ (2) $K_b = 2 K_f$
- (3) $K_b = 1.5 K_f$ (4) $K_b = K_f$
- 14. The freezing point of a diluted milk sample is found to be -0.2°C, while it should have been -0.5°C for pure milk. How much water has been added to pure milk to make the diluted sample?

[JEE (Main)-2019]

- (1) 3 cups of water and 2 cups of pure milk
- (2) 1 cup of water and 2 cups of pure milk
- (3) 2 cups of water to 3 cups of pure milk
- (4) 1 cup of water to 3 cups of pure milk

- 15. K₂Hgl₄ is 40% ionised in aqueous solution. The value of its van't Hoff factor (i) is [JEE (Main)-2019]
 - (1) 1.6

(2) 2.0

(3) 2.2

- (4) 1.8
- 16. Freezing point of a 4% agueous solution of X is equal to freezing point of 12% aqueous solution of Y. If molecular weight of X is A, then molecular weight of Y is [JEE (Main)-2019]
 - (1) 2A

(2) 3A

(3) A

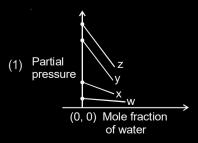
(4) 4A

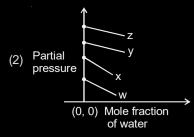
(Since density of solutions are not given therefore assuming molality to be equal to molarity and given % as % W/V)

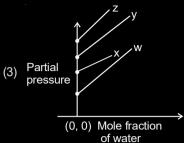
17. Molecules of benzoic acid (C_eH_eCOOH) dimerise in benzene. 'w' g of the acid dissolved in 30 g of benzene shows a depression in freezing point equal to 2 K. If the percentage association of the acid to form dimer in the solution is 80, then w is

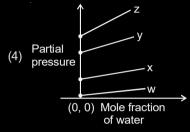
(Given that $K_f = 5 \text{ K kg mol}^{-1}$, Molar mass of benzoic acid = 122 g mol⁻¹) **[JEE (Main)-2019]**

- (1) 1.5 g
- (2) 2.4 g
- (3) 1.8 g
- (4) 1.0 g
- 18. The vapour pressures of pure liquids A and B are 400 and 600 mmHg, respectively at 298 K. On mixing the two liquids, the sum of their initial volumes is equal to the volume of the final mixture. The mole fraction of liquid B is 0.5 in the mixture. The vapour pressure of the final solution, the mole fractions of components A and B in vapour phase, respectively are [JEE (Main)-2019]
 - (1) 500 mmHg, 0.4, 0.6 (2) 500 mmHg, 0.5, 0.5
 - (3) 450 mmHg, 0.4, 0.6 (4) 450 mmHg, 0.5, 0.5
- 19. For the solution of the gases w, x, y and z in water at 298 K, the Henrys law constants (K_L) are 0.5, 2, 35 and 40 kbar, respectively. The correct plot for the given data is: [JEE (Main)-2019]









 Liquid 'M' and liquid 'N' form an ideal solution. The vapour pressures of pure liquids 'M' and 'N' are 450 and 700 mmHg, respectively, at the same temperature. Then correct statement is

 $(x_{M} = Mole fraction of 'M' in solution;$

 x_N = Mole fraction of 'N' in solution;

 y_M = Mole fraction of 'M' in vapour phase;

 y_N = Mole fraction of 'N' in vapour phase)

[JEE (Main)-2019]

$$(1) \quad \frac{x_M}{x_N} = \frac{y_M}{y_N}$$

$$(2) \quad \frac{x_M}{x_N} > \frac{y_M}{y_N}$$

$$(3) \quad \frac{x_M}{x_N} < \frac{y_M}{y_N}$$

(4)
$$(x_M - y_M) < (x_N - y_N)$$

- 21. The osmotic pressure of a dilute solution of an ionic compound XY in water is four times that of a solution of 0.01 M BaCl₂ in water. Assuming complete dissociation of the given ionic compounds in water, the concentration of XY (in mol L⁻¹) in solution is [JEE (Main)-2019]
 - (1) 16×10^{-4}
- $(2) 4 \times 10^{-4}$
- (3) 6×10^{-2}
- $(4) 4 \times 10^{-2}$

22. Molal depression constant for a solvent is 4.0 K kg mol⁻¹. The depression in the freezing point of the solvent for 0.03 mol kg⁻¹ solution of $\rm K_2SO_4$ is

(Assume complete dissociation of the electrolyte)

[JEE (Main)-2019]

- (1) 0.36 K
- (2) 0.18 K
- (3) 0.12 K
- (4) 0.24 K
- 23. At room temperature, a dilute solution of urea is prepared by dissolving 0.60 g of urea in 360 g of water. If the vapour pressure of pure water at this temperature is 35 mmHg, lowering of vapour pressure will be: (molar mass of urea = 60 g mol⁻¹)

[JEE (Main)-2019]

- (1) 0.031 mmHg
- (2) 0.017 mmHg
- (3) 0.028 mmHg
- (4) 0.027 mmHg
- 24. 1 g of a non-volatile non-electrolyte solute is dissolved in 100 g of two different solvents A and B whose ebullioscopic constants are in the ratio of 1 : 5. The ratio of the elevation in their boiling

points,
$$\frac{\Delta T_{b}\left(A\right)}{\Delta T_{b}\left(B\right)}$$
, is : [JEE (Main)-2019]

- (1) 1:5
- (2) 10:1
- (3) 5:1
- (4) 1:0.2
- 25. A solution is prepared by dissolving 0.6 g of urea (molar mass = 60 g mol⁻¹) and 1.8 g of glucose (molar mass = 180 g mol⁻¹) in 100 mL of water at 27°C. The osmotic pressure of the solution is

 $(R = 0.08206 \text{ L atm } \text{K}^{-1} \text{ mol}^{-1})$ [JEE (Main)-2019]

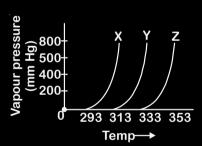
- (1) 1.64 atm
- (2) 2.46 atm
- (3) 8.2 atm
- (4) 4.92 atm
- 26. A solution containing 62 g ethylene glycol in 250 g water is cooled to -10°C. If K_f for water is 1.86 K kg mol⁻¹, the amount of water (in g) separated as ice is [JEE (Main)-2019]
 - (1) 64

(2) 32

(3) 16

- (4) 48
- 27. At 35°C, the vapour pressure of CS₂ is 512 mm Hg and that of acetone is 344 mm Hg. A solution of CS₂ in acetone has a total vapour pressure of 600 mm Hg. The false statement amongst the following is [JEE (Main)-2020]

- (1) Raoult's law is not obeyed by this system
- (2) A mixture of 100 mL CS₂ and 100 mL acetone has a volume < 200 mL
- (3) Heat must be absorbed in order to produce the solution at 35°C
- (4) CS₂ and acetone are less attracted to each other than to themselves
- 28. Two open beakers one containing a solvent and the other containing a mixture of that solvent with a non volatile solute are together sealed in a container. Over time [JEE (Main)-2020]
 - (1) The volume of the solution and the solvent does not change
 - (2) The volume of the solution increases and the volume of the solvent decreases
 - (3) The volume of the solution does not change and the volume of the solvent decreases
 - (4) The volume of the solution decreases and the volume of the solvent increases.
- 29. A graph of vapour pressure and temperature for three different liquids X, Y, and Z is shown below



The following inferences are made

- (A) X has higher intermolecular interactions compared to Y.
- (B) X has lower intermolecular interactions compared to Y
- (C) Z has lower intermolecular interactions compared to Y.

The correct inferences is/are [JEE (Main)-2020]

- (1) (B)
- (2) (C)
- (3) (A) and (C)
- (4) (A)

- 30. An open beaker of water in equilibrium with water vapour is in a sealed container. When a few grams of glucose are added to the beaker of water, the rate at which water molecules [JEE (Main)-2020]
 - (1) Leaves the solution increases
 - (2) Leaves the vapour increases
 - (3) Leaves the vapour decreases
 - (4) Leaves the solution decreases
- 31. The size of a raw mango shrinks to a much smaller size when kept in a concentrated salt solution. Which one of the following processes can explain this?

 [JEE (Main)-2020]
 - (1) Osmosis
 - (2) Reverse osmosis
 - (3) Diffusion
 - (4) Dialysis
- 32. Henry's constant (in kbar) for four gases α , β , γ and δ in water at 298 K is given below

(density of water = 10^3 kg m⁻³ at 298 K)

This table implies that [JEE (Main)-2020]

- (1) The pressure of a 55.5 molal solution of γ is 1 har
- (2) Solubility of γ at 308 K is lower than at 298 K
- (3) α has the highest solubility in water at a given pressure
- (4) The pressure of a 55.5 molal solution of δ is 250 bar
- 33. A set of solutions is prepared using 180 g of water as a solvent and 10 g of different non-volatile solutes A, B and C. The relative lowering of vapour pressure in the presence of these solutes are in the order [Given, molar mass of A = 100 g mol⁻¹; B = 200 g mol⁻¹; C = 10,000 g mol⁻¹]

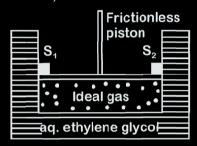
[JEE (Main)-2020]

- (1) A > C > B
- (2) C > B > A
- (3) A > B > C
- (4) B > C > A

34.	How much amount of NaCl should be added to 60 g of water (ρ = 1.00 g/mL) to decrease th freezing point of water to -0.2°C?			
	(The freezing point depression constant for water = 2 K kg mol ⁻¹) [JEE (Main)-2020]			
35.	A cylinder containing an ideal gas (0.1 mol of			

1.0 dm 3) is in thermal equilibrium with a large volume of 0.5 molal aqueous solution of ethylene glycol at its freezing point. If the stoppers \mathbf{S}_1 and \mathbf{S}_2 (as shown in the figure) are suddenly withdrawn, the volume of the gas in litres after equilibrium is achieved will be ______.

(Given, K_f (water) = 2.0 K kg mol⁻¹, R = 0.08 dm³ atm K^{-1} mol⁻¹)



[JEE (Main)-2020]

- 36. If 250 cm³ of an aqueous solution containing 0.73 g of a protein A is isotonic with one litre of another aqueous solution containing 1.65 g of a protein B, at 298 K, the ratio of the molecular masses of A and B is _____ × 10⁻² (to the nearest integer). [JEE (Main)-2020]
- 37. At 300 K, the vapour pressure of a solution containing 1 mole of n-hexane and 3 moles of n-heptane is 550 mm of Hg. At the same temperature, if one more mole of n-heptane is added to this solution, the vapour pressure of the solution increases by 10 mm of Hg. What is the vapour pressure in mm Hg of n-heptane in its pure state _____? [JEE (Main)-2020]
- 38. The osmotic pressure of a solution of NaCl is 0.10 atm and that of a glucose solution is 0.20 atm. The osmotic pressure of a solution formed by mixing 1 L of the sodium chloride solution with 2 L of the glucose solution is $x \times 10^{-3}$ atm. x is _____. (nearest integer)

[JEE (Main)-2020]

39. The elevation of boiling point of 0.10 m aqueous CrCl₃.xNH₃ solution is two times that of 0.05 m aqueous CaCl₂ solution. The value of x is _____.

[Assume 100% ionisation of the complex and CaCl₂, coordination number of Cr as 6, and that all NH₃ molecules are present inside the coordination sphere]

[JEE (Main)-2020]

40. A soft drink was bottled with a partial pressure of CO₂ of 3 bar over the liquid at room temperature. The partial pressure of CO₂ over the solution approaches a value of 30 bar when 44 g of CO₂ is dissolved in 1 kg of water at room temperature. The approximate pH of the soft drink is _____ × 10⁻¹.

(First dissociation constant of $H_2CO_3 = 4.0 \times 10^{-7}$; log 2 = 0.3; density of the soft drink = 1 g mL⁻¹) [JEE (Main)-2020]

- 41. 10.30 mg of O₂ is dissolved into a liter of sea water of density 1.03 g/mL. The concentration of O₂ in ppm is ______ . [JEE (Main)-2020]
- 42. When 9.45 g of $CICH_2COOH$ is added to 500 mL of water, its freezing point drops by 0.5°C. The dissociation constant of $CICH_2COOH$ is x × 10^{-3} . The value of x is _____. (Rounded off to the nearest integer)

$$[K_{f(H_2O)} = 1.86 \text{ K kg mol}^{-1}]$$
 [JEE (Main)-2021]

43. C₆H₆ freezes at 5.5°C. The temperature at which a solution of 10 g of C₄H₁₀ in 200 g of C₆H₆ freeze is ____ °C. (The molal freezing point depression constant of C₆H₆ is 5.12°C/m.)

[JEE (Main)-2021]

44. 1 molal aqueous solution of an electrolyte A₂B₃ is 60% ionised. The boiling point of the solution at 1 atm is _____ K. (Rounded-off to the nearest integer)

[Given K_b for $(H_2O) = 0.52 \text{ K kg mol}^{-1}$]

[JEE (Main)-2021]

45. If a compound AB dissociates to the extent of 75% in an aqueous solution, the molality of the solution which shows a 2.5 K rise in the boiling point of the solution is ____ molal. (Rounded-off to the nearest integer)

 $[K_b = 0.52 \text{ K kg mol}^{-1}]$ [JEE (Main)-2021]

[JEE (Main)-2021]

47.	of water, the freezing point of solution was found to be -0.93° C ($K_{\rm f}$ ($H_{\rm p}$ O) = 1.86 K kg mol ⁻¹). The	53.	point of 3.885° C. The degree of dissociation of this acid is × 10^{-3} .
	number (n) of benzoic acid molecules associated		(Round off to the Nearest Integer).
	(assuming 100% association) is [JEE (Main)-2021]		[Given : Molal depression constant of water = 1.85 K kg mol ⁻¹
48.	${\rm AB_2}$ is 10% dissociated in water to ${\rm A^{2+}}$ and ${\rm B^-}.$		Freezing point of pure water = 0° C]
	The boiling point of a 10.0 molal aqueous solution		[JEE (Main)-2021]
	of AB ₂ is°C. (Round off to the Nearest Integer)	54.	A solute A dimerizes in water. The boiling point of a 2 molal solution of A is 100.52°C. The percentage
	[Given : Molal elevation constant of water $K_b = 0.5$ K kg mol ⁻¹ boiling point of pure water = 100° C]		association of A is (Round off to the Nearest Integer).
	[JEE (Main)-2021]		[Use : K_b for water = 0.52 K kg mol ⁻¹
49.	At 363 K, the vapour pressure of A is 21 kPa and		Boiling point of water = 100°C]
	that of B is 18 kPa. One mole of A and 2 moles		[JEE (Main)-2021]
	of B are mixed. Assuming that this solution is ideal, the vapour pressure of the mixture is kPa.	55.	At 20°C, the vapour pressure of benzene is 70 torr and that of methyl benzene is 20 torr. The mole fraction of benzene in the vapor phase at 20°C
	(Round off to the Nearest Integer]		above an equimolar mixture of benzene and methyl benzene is \times 10 ⁻² . [JEE (Main)-2021]
	[JEE (Main)-2021]		(Nearest integer)
50.	The mole fraction of a solute in a 100 molal aqueous solution is $___$ × 10^{-2} .	56.	The vapour pressures of A and B at 25°C are 90
	(Round off to the Nearest Integer).		mm Hg and 15 mm Hg respectively. If A and B are mixed such that the mole fraction of A in the
	[Given : Atomic masses : H : 1.0 u, O : 16.0 u]		mixture is 0.6, then the mole fraction of B in the
E 1	[JEE (Main)-2021]		vapour phase is $x \times 10^{-1}$. The value of x is (Nearest integer) [JEE (Main)-2021]
51.	The oxygen dissolved in water exerts a partial pressure of 20 kPa in the vapour above water. The molar solubility of oxygen in water is	57.	Which one of the following 0.06 M aqueous solutions has lowest freezing point?
	10 ⁻⁵ mol dm ⁻³ .		[JEE (Main)-2021]
	(Round off to the Nearest Integer).		(1) KI (2) $Al_2(SO_4)_3$
	[Given : Henry's law constant = $K_H = 8.0 \times 10^4 \text{ kPa}$		(3) $C_6H_{12}O_6$ (4) K_2SO_4
	for O ₂ .	58.	2 0
	Density of water with dissolved oxygen = 1.0 kg dm ⁻³]		drink manufacturing process at 298 K. If CO ₂ exerts a partial pressure of 0.835 bar then x m mol of CO ₂ would dissolve in 0.9 L of water. The value
	[JEE (Main)-2021]		of x is (Nearest integer)
52.	A 1 molal K ₄ Fe(CN) ₆ solution has a degree of dissociation of 0.4. Its boiling point is equal to that		(Henry's law constant for CO_2 at 298 K is 1.67 × 10 3 bar) [JEE (Main)-2021]
	of another solution which contains 18.1 weight percent of a non electrolytic solute A. The molar mass of A is u. (Round off to the Nearest Integer).	59.	When 3.00 g of a substance 'X' is dissolved in 100 g of CCl ₄ , it raises the boiling point by 0.60 K. The molar mass of the substance 'X' is g mol ⁻¹ . (Nearest integer)
	[Density of water = 1.0 g cm ⁻³]		[Given K _b for CCl ₄ is 5.0 K kg mol ⁻¹]
	[JEE (Main)-2021]		[.JFF (Main)-2021]

monomer in acetone. The boiling point of the solution in acetone increases by 0.17° C. The increase in boiling point of solution in benzene in $^{\circ}$ C is $x \times 10^{-2}$. The value of x is (Nearest integer)		water at 300 K exerted an osmotic pressure of 2.42 \times 10 ⁻³ bar. The molar mass of the biopolymer is \times 10 ⁴ g mol ⁻¹ . (Round off to the Nearest Integer)	30.	
[Atomic mass : C = 12.0, H = 1.0, O = 16.0]		[Use : R = 0.083 L bar mol ⁻¹ K ⁻¹]		
[JEE (Main)-2021]		[JEE (Main)-2021]	31.	
The osmotic pressure of blood is 7.47 bar at 300 K.	68.	In a solvent 50% of an acid HA dimerizes and the rest dissociates. The van't Hoff factor of the acid is \times 10 ⁻² . (Round off to the Nearest Integer).		
To inject glucose to a patient intravenously, it has				
to be isotonic with blood. The concentration of	[JEE (Main)-2021]			
glucose solution in gL^{-1} is (Molar mass of		Of the following four aqueous solutions, total		
glucose = 180 g mol ⁻¹		number of those solutions whose freezing point is lower than that of 0.10 M C_2H_5OH is		
$R = 0.083 L bar K^{-1} mol^{-1})$ (Nearest integer)		(Integer answer) [JEE (Main)-2021] (i) 0.10 M Ba ₃ (PO ₄) ₂ (ii) 0.10 M Na ₂ SO ₄		
[JEE (Main)-2022]		(iii) 0.10 M KCI (iv) 0.10 M Li ₃ PO ₄		
A company dissolves 'x' amount of ${\rm CO_2}$ at 298 K in	69.	83 g of ethylene glycol dissolved in 625 g of water. The freezing point of the solution is K. (Nearest integer) [JEE (Main)-2021]	63.	
1 litre of water to prepare soda water. X =				
× 10 ⁻³ g. (nearest integer)		[Use : Molal freezing point depression constant		
(Given: partial pressure of CO ₂ at 298 K = 0.835 bar.		of water = 1.86 K kg mol ⁻¹ ,		
Henry's law constant for CO ₂ at 298 K = 1.67 kbar.		Freezing point of water = 273 K,		
Atomic mass of H, C and O is 1, 12, and 6 g mol^{-1} ,		Atomic masses : C : 12.0 u, O : 16.0 u, H : 1.0 u]		
respectively) [JEE (Main)-2022]	70.	1 kg of 0.75 molal aqueous solution of sucrose can be cooled up to -4°C before freezing. The amount of ice (in g) that will be separated out is		
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		(Nearest integer) [JEE (Main)-2021]		
of solute A in water is:		[Given : K _f (H ₂ O) = 1.86 K kg mol ⁻¹]		
[Given : Molar mass of A = 93 g mol ⁻¹ . Molal		40 g of glucose (Molar mass = 180) is mixed with 200 mL of water. The freezing point of solution is	35.	
depression constant of water is 1.86 K kg mol ⁻¹ .]		K. (Nearest integer) [JEE (Main)-2021]		
[JEE (Main)-2022]		[Given : $K_f = 1.86 \text{ K kg mol}^{-1}$; Density of water = 1.00 g cm ⁻³ ; Freezing point of water = 273.15 K]		
(1) 50% (2) 60%			36.	
(3) 70% (4) 80%		solutions will exhibit the largest freezing point depression? [JEE (Main)-2021]		
A 0.5 percent solution of potassium chloride was	71.	(1) Glycine (2) KHSO ₄		
found to freeze at -0.24°C. The percentage dissociation of potassium chloride is		(3) Hydrazine (4) Glucose		
(Nearest integer)		1.22 g of an organic acid is separately dissolved in	37.	
(Molal depression constant for water is 1.80 K kg		100 g of benzene ($K_b = 2.6 \text{ K kg mol}^{-1}$) and 100		
mol ⁻¹ and molar mass of KCl is 74.6 g mol ⁻¹)		g of acetone ($K_b = 1.7 \text{ K kg mol}^{-1}$). The acid is known to dimerize in benzene but remain as a		
[JEE (Main)-2022]		- Known to dimenze in benzene but femalif as a		

72.	I he osmotic pressure exerted by a solution prepared	78.	Elevation in boiling point for 1.5 molal solution of
	by dissolving 2.0 g of protein of molar mass 60 kg		glucose in water is 4 K. The depression in freezing
	mol ⁻¹ in 200 mL of water at 27°C is Pa.		point for 4.5 molal solution of glucose in water is
	[Integer value]		4 K. The ratio of molal elevation constant to molal
	(use R = 0.083 L bar mol ⁻¹ K ⁻¹) [JEE (Main)-2022]		depression constant (K_b/K_f) is
73.	2 g of a non-volatile non-electrolyte solute is dissolved in 200 g of two different solvents A and B		[JEE (Main)-2022]
	whose ebullioscopic constants are in the ratio of	79.	The depression in freezing point observed for a
	1:8. The elevation in boiling points of A and B are in		formic acid solution of concentration 0.5 mL L ⁻¹ is
	the ratio $\frac{x}{y}$ (x:y). The value of y is (Nearest		0.0405°C. Density of formic acid is 1.05 g mL ⁻¹ . The Van't Hoff factor of the formic acid solution is
	Integer) [JEE (Main)-2022]		nearly (Given for water $K_r = 1.86 \text{ k kg mol}^{-1}$)
7/	A solution containing 2.5×10^{-3} kg of a solute		[JEE (Main)-2022]
74.	dissolved in 75×10^{-3} kg of water boils at		(1) 0.8 (2) 1.1
			(3) 1.9 (4) 2.4
	373.535 K. The molar mass of the solute is g mol ⁻¹ . [nearest integer] (Given : $K_b(H_2O) = 0.52$ K kg	80.	Two solutions A and B are prepared by dissolving
	mol ⁻¹ and boiling point of water = 373.15 K)		1 g of non-volatile solutes X and Y, respectively in
	[JEE (Main)-2022]		1 kg of water. The ratio of depression in freezing
75.	The vapour pressures of two volatile liquids A and B		points for A and B is found to be 1 : 4. The ratio of molar masses of X and Y is
	at 25°C are 50 Torr and 100 Torr, respectively. If the		[JEE (Main)-2022]
	liquid mixture, contains 0.3 mole fraction of A, then the mole fraction of liquid B in the vapour phase is		(1) 1:4 (2) 1:0.25
			(3) 1:0.20 (4) 1:5
	$\frac{x}{17}$. The value of x is [JEE (Main)-2022]	81.	The elevation in boiling point for 1 molal solution
76.	2.5 g of protein containing only glycine (C ₂ H ₅ NO ₂)		of non-volatile solute A is 3 K. The depression in
	is dissolved in water to make 500 mL of solution.		freezing point for 2 molal solution of A in the same
	The osmotic pressure of this solution at 300 K is		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]
	found to be 5.03×10^{-3} bar. The total number of		[JEE (Main)-2022]
	glycine units present in the protein is	82	Boiling point of a 2% aqueous solution of a non-
		02.	volatile solute A is equal to the boiling point of 8%
	(Given : R = 0.083 L bar K ⁻¹ mol ⁻¹)		aqueous solution of a non-volatile solute B. The
	[JEE (Main)-2022]		relation between molecular weights of A and B is
77.	1.2 mL of acetic acid is dissolved in water to make		[JEE (Main)-2022]
	2.0 L of solution. The depression in freezing point		(1) $M_A = 4M_B$ (2) $M_B = 4M_A$
	observed for this strength of acid is 0.0198°C. The		(3) $M_A = 8M_B$ (4) $M_B = 8M_A$
	percentage of dissociation of the acid is	83.	When a certain amount of solid A is dissolved in 100 g of water at 25°C to make a dilute solution,
	(Nearest integer)		the vapour pressure of the solution is reduced to
	[Given: Density of acetic acid is 1.02 g mL ⁻¹		one-half of that of pure water. The vapour pressure
	Molar mass of acetic acid is 60 g mol ⁻¹		of pure water is 23.76 mmHg. The number of
	$K_f(H_2O) = 1.85 \text{ K kg mol}^{-1}$		moles of solute A added is (Nearest
	[JEE (Main)-2022]		Integer) [JEE (Main)-2022]

84.	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 × 10 ⁻¹)°C. The value of x is (Nearest integer) (Given: $K_f = 3.9 \text{ K kg mol}^{-1}$; molar mass of ascorbic		(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 ${\rm O_2}$ in water is too small, nearly			
			negligible) [JEE (Main)-2022]			
	acid = 176 g mol ⁻¹) [JEE (Main)-2022]	87.	1.80 g of solute A was dissolved in 62.5 $\mbox{cm}^{\mbox{\tiny 3}}$ of			
85.	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		ethanol and freezing point of the solution was			
			found to be 155.1 K. The molar mass of solute A			
			is g mol ⁻¹ .			
	the liquid phase. The vapour pressure of pure liquid A is atm. (Nearest integer)		[Given : Freezing point of ethanol is 156.0 K.			
	[JEE (Main)-2022]		Density of ethanol is 0.80 g cm ⁻³ .			
86.	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)		Freezing point depression constant of ethanol is			
			2.00 K kg mol ⁻¹] [JEE (Main)-2022]			

Chapter 10

Solutions

1. Answer (1)

Ethanol has H-Bonding, n-heptane tries to break the H-bonds of ethanol, hence, V.P. increases. Such a solution shows positive deviation from Raoult's Law.

2. Answer (2)

Let V. P. of pure X = x

and V. P. of pure Y = y

Then,
$$\frac{1}{4}x + \frac{3}{4}y = 550$$
 ...(i)

and
$$\frac{1}{5}x + \frac{4}{5}y = 560$$
 ...(ii)

Solving (i) and (ii), we get

$$x = 400 \text{ mm}$$

and v = 600 mm

3. Answer (3)

$$\Delta T_f = i K_f m$$

i for Na₂SO₄ is 3(100% ionisation)

$$\Delta T_f = 3 \times 1.86 \times \frac{0.01}{1}$$

 $\Delta T_f = 0.0558 \text{ K}$

4. Answer (2)

$$P = X_A P_A^o + X_B P_B^o$$

$$n_{heptane} = \frac{25}{100} = 0.25$$

$$n_{\text{octane}} = \frac{35}{114} = 0.307$$

$$P = \frac{0.25}{0.25 + 0.307} \times 105 + \frac{0.307}{0.25 + 0.307} \times 45$$

$$=47.127 + 24.84 = 71.96$$

≈ 72 kPa

5. Answer (4)

For isotonic solution

$$\pi_1 = \pi_2$$

$$C_1 = C_2$$

$$\frac{W_1}{M_1V_1} = \frac{W_2}{M_2V_2}$$

$$\frac{5}{342 \times 100} = \frac{1}{M_2 \times 100}$$

$$M_2 = 68.4$$

- 6. Answer (1)
- 7. Answer (1)

$$\pi = iCRT$$

$$\pi_{C_2H_sOH} = 1 \times 0.500 \times R \times T = 0.5RT$$

$$\pi_{Mg_{0}(PO_{4})_{0}} = 5 \times 0.100 \times R \times T = 0.5 RT$$

$$\pi_{KBr} = 2 \times 0.250 \times R \times T = 0.5 RT$$

$$\pi_{\text{Na},\text{PO}_4} = 4 \times 0.125 \times \text{RT} = 0.5 \,\text{RT}$$

8. Answer (2)

Vapour pressure of pure acetone $P_A^\circ = 185$ torr Vapour pressure of solution, $P_S = 183$ torr Molar mass of solvent, $M_A = 58$ g/mole

as we know
$$\frac{P_A^{\circ} - P_S}{P_S} = \frac{n_B}{n_A}$$

$$\Rightarrow \frac{185 - 183}{183} = \frac{W_B}{M_B} \times \frac{M_A}{W_A}$$

$$\Rightarrow \frac{2}{183} = \frac{1.2}{M_B} \times \frac{58}{100}$$

$$\Rightarrow M_B = \frac{1.2}{2} \times \frac{58}{100} \times 183$$

=63.68 g/mole

$$\frac{p^{o}-p_{s}}{p_{s}} = \frac{n_{solute}}{n_{solvent}} = \frac{\frac{18}{180}}{\frac{178.2}{18}} = \frac{18}{17.82}$$

At normal boiling point of water V.P. = p^{o} = 760 torr

$$\therefore \frac{760 - p_s}{p_s} = \frac{18}{1782}$$

or,
$$1800 p_s = 760 \times 1782$$

$$p_s = 752.4 \text{ torr}$$

10. Answer (2)

$$0.45 = i(5.12) \frac{0.2/60}{20} \times 1000$$

$$\Rightarrow$$
 $i = 1 - \frac{\alpha}{2}$

$$\Rightarrow$$
 0.527 = $1 - \frac{\alpha}{2}$

$$\Rightarrow \frac{\alpha}{2} = 0.473$$

$$\Rightarrow \alpha = 0.946$$

∴ % association = 94.6%

11. Answer (4)

Solubility decreases with the increase in value of K_{H} .

12. Answer (3)

$$P_{\Lambda}^{\circ} = 7 \times 10^{3}$$

$$P_{\rm p}^{\circ} = 12 \times 10^3$$

$$X_{A} = 0.4$$

$$X_{D} = 0.6$$

$$\therefore$$
 P = $(7 \times 0.4 + 12 \times 0.6) \times 10^3 = 10^4$

$$P_A = 2.8 \times 10^3, P_B = 7.2 \times 10^3$$

$$Y_A = 0.28, Y_B = 0.72$$

13. Answer (2)

$$\Delta T_b = k_b m \implies k_b(1) = 2 \implies k_b = 2 \text{ km}^{-1}$$

$$\Delta T_f = k_f m \implies k_f(2) = 2 \implies k_f = 1 \text{ km}^{-1}$$

$$k_f = 0.5 \text{ k}_b \implies k_b = 2 \text{ k}_f$$

14. Answer (1)

Freezing point of diluted milk = -0.2°C

$$\Delta T_f' = 0.2$$
°C

Freezing point of pure milk = -0.5°C

$$\Delta T_f = 0.5^{\circ}C$$

$$\frac{\Delta T_f}{\Delta T_f'} = \frac{k_f}{k_f} \frac{m}{m'}$$

$$\Rightarrow \frac{0.5}{0.2} = \frac{w_1'}{w_1}$$

$$\Rightarrow \frac{W_1'}{W_1} = \frac{5}{2}$$

2 cups of pure milk mixed with 3 cups of water overall 5 cups of diluted milk.

15. Answer (4)

$$K_2HgI_4 \Longrightarrow 2K^+ + [HgI_4]^{2-}$$

$$n = 3$$

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

$$0.4 = \frac{i-1}{3-1}$$

$$i = 1.8$$

16. Answer (2)

$$\frac{4}{M_x} = \frac{12}{M_y}$$

$$\Rightarrow$$
 M_v = 3M_x

$$M_v = 3A$$

(Since density of solutions are not given therefore assuming molality to be equal to molarity and given % as % W/V)

17. Answer (2)

$$2C_6H_5COOH \longrightarrow (C_6H_5COOH)_2$$

$$1 \qquad 0$$

t = 0

 $1-2\alpha$ t

Moles at equilibrium = $1 - 2\alpha + \alpha = 1 - \alpha$

$$2\alpha = 0.8$$
, $\alpha = 0.4$

Moles at equilibrium = 0.6

i = 0.6

$$\Delta T_f = ik_f m \implies 2 = 0.6 \times 5 \times \left(\frac{w}{122}\right) \times 1000$$

 $w = 2.4 \text{ g}$

18. Answer (1)

$$P = x_B p_B^\circ + x_A p_A^\circ$$

= 0.5 × 600 + 0.5 × 400 = 300 + 200 = 500
$$p_B = y_B P_{Total}$$

$$y_B = \frac{p_B}{P_{Total}} = \frac{300}{500} = \frac{3}{5} = 0.6$$

$$y_A = \frac{p_A}{P_{Tatal}} = \frac{200}{500} = \frac{2}{5} = 0.4$$

19. Answer (1)

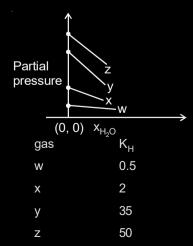
According to Henry's law,

$$P = K_H \cdot X_{gas}$$

$$\therefore$$
 $x_{gas} = 1 - x_{H_2O}$

$$P = K_H - K_H \cdot X_{H_2O}$$

$$y = C + mx$$



20. Answer (2)

$$P_M^o = 450 \text{ mmHg}, P_N^o = 700 \text{ mmHg}$$

$$P_M = P_M^o X_M = Y_M P_T$$

$$\Rightarrow P_M^o = \frac{Y_M}{X_M}(P_T)$$

Similarly,
$$P_N^o = \frac{Y_N}{X_N}(P_T)$$

Given,
$$P_M^o < P_N^o$$

$$\Rightarrow \frac{Y_M}{X_M} < \frac{Y_N}{X_N}$$

$$\Rightarrow \frac{Y_M}{Y_N} < \frac{X_M}{X_N}$$

21. Answer (3)

$$\pi_{XY} = 4\pi_{BaCl_a}$$

$$\therefore$$
 2[XY] = 4 × (0.01) × 3

[XY] =
$$0.06 = 6 \times 10^{-2} \frac{\text{mol}}{\text{I}}$$

22. Answer (1)

$$K_2SO_4 \longrightarrow 2K^+ + SO_4^{2-}$$

i (van't Hoff Factor) = 3

$$\triangle T_f = iK_f m$$

$$= 3 \times 4 \times 0.03$$

$$= 0.36 \text{ K}$$

23. Answer (2)

Relative lowering of VP is given by

$$\frac{P_B^{\circ} - P_B}{P_B^{\circ}} = X_A = \frac{n_A}{n_A + n_B} \simeq \frac{\textbf{n}_A}{\textbf{n}_B}$$

$$\frac{P_{B}^{\circ} - P_{B}}{35} = \frac{0.6 \times 18}{60 \times 360} = \frac{1}{2000}$$

On solving,
$$\Delta P_B = P_B^{\circ} - P_B = 0.017$$

24. Answer (1)

$$\Delta T_b = k_b \times m$$

$$\frac{\left(k_{b}\right)_{A}}{\left(k_{b}\right)_{B}}=\frac{1}{5}$$

$$\therefore \frac{(\Delta T_{b})_{A}}{(\Delta T_{b})_{B}} = \frac{(k_{b})_{A}}{(k_{b})_{B}} = \frac{1}{5}$$

25. Answer (4)

Osmotic pressure (π) = CRT

Solute: urea and glucose

$$\pi = (C_1 + C_2) RT$$

$$= \left(\frac{0.6}{60 \times 0.1} + \frac{1.8}{180 \times 0.1}\right) \times 0.0821 \times 300$$

$$= 0.2 \times 0.0821 \times 300$$

$$= 4.926 \text{ atm}$$

26. Answer (1)

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

$$\Rightarrow 10 = 1.86 \times \frac{62 \times 1000}{62 \times W_A}$$

$$\Rightarrow W_A = \frac{1.86 \times 1000}{10}$$
$$= 186 g$$

Given amount of H₂O is 250 g

∴ The amount of water separated as ice
 = 250 – 186 = 64 q

27. Answer (2)

Since the vapour pressure of the solution is greater than individual vapour pressure of both pure components, the solution shows a positive deviation from Raoult's law.

28. Answer (2)

There will be lowering in vapour pressure for solution containing non-volatile solute. So, there will be transfer of solvent molecules from pure solvent to solution and hence, volume of beaker containing solvent (pure) will decrease and volume of beaker containing solution will increase.

29. Answer (1)

Vapour pressure of a liquid at a given temperature is inversely proportional to intermolecular force of attraction. At the same temperature, vapour pressure of X is higher than that of Y.

Therefore (X) has lower intermolecular interactions compared to Y. Statement (B) is correct.

30. Answer (4)

The rate at which water molecules leaves the solution decreases.

31. Answer (1)

Osmosis can explain the given process. There are many phenomena which we observed in nature or at home. Raw mango shrivel when pickled in brine.

The solvent molecules will flow through the membrane from pure solvent to the solution. This process of flow of the solvent is called osmosis.

32. Answer (2)

With temperature, the value of $K_{\rm H}$ (Henry's constant) increases and solubility of gas in liquid decreases

Ideally Henry's law is applicable for dilute solutions.

:. 55.5 molal solution of δ at 250 bar will not follow Henry's law.

33. Answer (3)

Relative lowering in vapour pressure (RLVP)

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

n → moles of solute

 $N \rightarrow moles of solvent$

$$n_A = \frac{10}{100}, n_B = \frac{10}{200}, n_C = \frac{10}{10000}$$

From the above relation

RLVP (A) > RLVP (B) > RLVP (C)

A > B > C

34. Answer (1.76)

$$\Delta T_f = 0.2^{\circ}C$$

$$\therefore \Delta T_f = ik_f \cdot m$$

i = 2 for NaCl

$$0.2 = 2 \times 2 \times \frac{W_{NaCl} \times 1000}{58.5 \times 600}$$

$$\Rightarrow W_{NaCl} = \frac{58.5 \times 600 \times 0.2}{4 \times 1000}$$
= 1.755
= 1.76

35. Answer (2.18)

For aqueous solution

$$\Delta T_f = K_f \cdot m = 2 \times 0.5$$

∴ Final volume of ideal gas =
$$\frac{nRT}{P}$$

$$= \frac{0.1 \times 0.08 \times 272}{1}$$

$$\approx 2.18 L$$

36. Answer (177)

Isotonic solutions have the same osmotic pressure.

$$\pi_A = C_A RT; C_A = \frac{0.73 \times 1000}{M_A \times 250}$$

$$\pi_{B} = C_{B}RT; C_{B} = \frac{1.65}{M_{P}}$$

$$\pi_A = \pi_B \Rightarrow C_A = C_B$$

$$\therefore \quad \frac{0.73 \times 1000}{M_A \times 250} = \frac{1.65}{M_B}$$

$$\frac{M_A}{M_B} = 1.77 = 177 \times 10^{-2}$$

37. Answer (600)

If X_1 and P_1° are the mole fraction and vapour pressure of n-hexane in solution and X_2 and P_2° are the mole fraction and vapour pressure of n-heptane in solution then

$$550 = X_1 P_1^{\circ} + X_2 P_2^{\circ}$$

$$= \frac{P_1^{\circ}}{4} + \frac{3P_2^{\circ}}{4} \Rightarrow P_1^{\circ} + 3P_2^{\circ} = 2200 \qquad ...(1)$$

On addition of 1 more mole of n-heptane

$$560 = X'_{1}P_{1}^{\circ} + X'_{2}P_{2}^{\circ}$$

$$= \frac{P_{1}^{\circ}}{5} + \frac{4P_{2}^{\circ}}{5} \implies P_{1}^{\circ} + 4P_{2}^{\circ} = 2800 \qquad ...(2)$$

From (1) and (2), $P_2^{\circ} = 600 \text{ mm Hg}$

38. Answer (167.00)

For NaCl:
$$\pi_1 = iC_1RT \Rightarrow C_1 = \frac{0.10}{2 RT}$$

For Glucose:
$$\pi_2 = C_2RT \Rightarrow C_2 = \frac{0.20}{RT}$$

When 1 L of NaCl solution and 2 L glucose solution are mixed.

$$C_1' = \frac{0.10}{6 \text{ RT}} \text{ and } C_2' = \frac{0.20 \times 2}{3 \text{ RT}} = \frac{0.40}{3 \text{ RT}}$$

$$\therefore \quad \pi_{Total} = iC_1'RT + C_2'RT = \frac{0.10}{3} + \frac{0.40}{3} = \frac{0.50}{3}$$

$$\pi_{\text{Total}} \simeq 167 \times 10^{-3} \text{ atm}$$

39. Answer (5.00)

Molality of CaCl₂ solution = 0.05 m

$$\Delta T_{b} = i K_{b} m = 3 \times K_{b} \times 0.05 = 0.15 K_{b}$$

Molality of CrCl₃.xNH₃ = 0.10 m

$$\Delta T_{b}' = i K_{b} \times 0.10; \Delta T_{b}' = 2\Delta T_{b}$$

$$iK_{h} \times 0.10 = 2 \times 0.15 K_{h} \Rightarrow i = 3$$

Since, co-ordination number of Cr is 6.

- \therefore The complex is $[Cr(NH_3)_5Cl]Cl_2$
- ∴ x = 5
- 40. Answer (37)

At 30 bar pressure mass of CO_2 in 1 kg water = 44 gm At 3 bar pressure mass of CO_2 in 1 kg water = 4.4 gm

Moles of CO_2 in 1 kg water = 0.1

$$H_2CO_3 \rightleftharpoons H^+ + HCO_3^-$$

0.1(1- α) 0.1 α 0.1 α

$$Ka_1 = \frac{[H^+][HCO_3^-]}{[H_2CO_3]}$$

$$4 \times 10^{-7} = \frac{0.1\alpha^2}{1-\alpha} \approx 0.1\alpha^2$$

$$\alpha = 2 \times 10^{-3}$$

[H⁺] = 0.1
$$\alpha$$
 = 2 × 10⁻⁴; pH = 3.7
= 37 × 10⁻¹

41. Answer (10)

ppm of O₂ =
$$\frac{\text{wt. of O}_2}{\text{wt. of H}_2\text{O}} \times 10^6$$

= $\frac{10.3 \text{ mg}}{1.03 \times 10^6 \text{ mg}} \times 10^6$
= 10 ppm

42. Answer (35)

Moles of CICH₂COOH =
$$\frac{9.45}{94.5}$$
 = 0.1 moles

$$C = \frac{0.1}{0.5} = 0.2 M$$

Assuming molarity = molality

$$\Delta T_f = ik_f \cdot m = (1 + \alpha)1.86 \times 0.2$$

$$\Rightarrow$$
 $(1+\alpha) = \frac{0.5}{0.2 \times 1.86} = 1.344 \approx 1.34$

$$\Rightarrow \alpha = 0.34$$

$$K_a$$
 of (CICH₂COOH) = $\frac{C\alpha \cdot C\alpha}{C(1-\alpha)}$

$$=\frac{C\alpha^2}{1-\alpha}=\frac{0.2\times(0.34)^2}{1-0.34}$$

$$= 35 \times 10^{-3}$$

43. Answer (1)

$$\Delta T_f = ik_f m$$

$$i = 1 \text{ for } C_4 H_{10}; \ T_f^o = 5.5^{\circ} C$$

$$m = \frac{10 \times 1000}{58 \times 200} = \frac{50}{58}$$

$$K_f = 5.12^{\circ}C / m$$

$$\Delta T_f = 5.12 \times \frac{50}{58} = 4.41^{\circ}C$$

$$T_{\rm f}^{\rm o} - T_{\rm f} = 4.41$$

$$T_f = 5.50 - 4.41 = 1.09 \approx 1^{\circ}C$$

44. Answer (375)

$$A_2B_3 \longrightarrow 2A^{3+} + 3B^{2-}$$
 (Assuming A – cation,
1 – 0.6 2 × 0.6 3 × 0.6 B – anion)

$$\therefore$$
 van't Hoff factor = 1 - 0.6 + 1.2 + 1.8 = 3.4

$$\Delta T_b$$
 = ik_b · m
= 3.4 × 0.52 × 1
= 1.768
 T_b = 373 + 1.768
= 374.77
≈ 375

45. Answer (3)

AB
$$\underset{1-\alpha}{\longleftrightarrow} A^{n+} + B^{n-}$$

 $i = 1 + \alpha = 1 + 0.75 \ (\because \alpha = 0.75)$
 $= 1.75$
 $\Delta T_b = i K_b m$
 $m = \frac{2.5}{1.75 \times 0.52} = 2.75 \text{ mol / kg} \approx 3$

46. Answer (24)

$$n_{SO_2} = \frac{1 \times 0.224}{0.082 \times 298} \approx 0.0092 \approx 0.01 \,\text{moles}$$

$$n_{NaHSO_3} = 0.01$$

$$NaHSO_3 \rightarrow Na^+ + HSO_3^-$$

Ignoring the dissociation of HSO_3^- into H^+ and SO_3^{2-}

van't Hoff factor (i) = 2

$$\frac{P^{o}_{H_2O} - P_S}{P^{o}_{H_2O}} = \frac{in_{NaHSO_3}}{n_{H_2O} + in_{NaHSO_3}}$$

$$\left(\operatorname{asn}_{\operatorname{HSO}_{2}^{-}} << \operatorname{n}_{\operatorname{H}_{2}\operatorname{O}} \right)$$

Lowering in vapour presssure

$$= \frac{2 \times 0.01}{2 + 2 \times 0.01} \times 24$$

=
$$23.76 \times 10^{-2}$$
 mmHg $\approx 24 \times 10^{-2}$ mmHg

47. Answer (02.00)

$$\Delta T_f = iK_f m$$

$$0.93 = i \times 1.86 \times \frac{12.2 \times 1000}{122 \times 100}$$

i = 0.5

n(Benzoic acid) ---- (Benzoic acid)

 $i = \frac{\text{Total number of particles after association}}{\text{Number of particles before association}}$

$$0.5 = \frac{1}{n}$$

$$n = 2$$

48. Answer (106)

$$AB_2 \rightleftharpoons A^{2+} + 2B^{-}$$

$$1-\alpha$$
 α 2α

$$i = 1 + \alpha$$

$$i = 1.1$$

$$\Delta T_h = T_s - T^\circ = i \times K_h \times \text{molality}$$

$$T_s - 100 = 1.1 \times 0.5 \times 10$$

$$T_s = 105.5$$

49. Answer (19)

An ideal solution is prepared by mixing 1 mol of A and 2 moles of B

Using Raoult's law

$$P_s = \chi_A P_A^o + \chi_B P_B^o$$

$$=\frac{1}{3}\times 21+\frac{2}{3}\times 18$$

= 19 kPa

50. Answer (64)

Molality of an aqueous solution of a solute = 100 m

Number of moles of solvent = $\frac{1000}{18}$

Mole fraction of solute =
$$\frac{100}{100 + \frac{1000}{18}} = \frac{100 \times 18}{2800}$$

$$= 0.6428 = 64.28 \times 10^{-2}$$

⁻≃ 64

51. Answer (25)

$$P_{O_2}$$
 (over water) = 20 kPa

$$K_H$$
 for O_2 = 8.0 × 10⁴ kPa

If X_{O_2} is the mole fraction of O_2 in soution, then according to Henry's law

$$P_{O_2} = K_H(X_{O_2})$$

$$X_{O_2} = \frac{20}{8.0 \times 10^4} = 2.5 \times 10^{-4}$$

Mass of 1 kg of water containing $O_2 = 1 L$

∴ Molarity of O_2 in solution = 25 × 10⁻⁵ M

52. Answer (85)

i for
$$K_4[Fe(CN)_6] = 1 + (5 - 1) 0.4 = 2.6$$

$$\Delta T_{b_4} = 2.6 \times K_f \times 1$$

i for A = 1.

$$(\Delta T_b)_A = 1 \times \frac{18.1 \times 1000}{A \times 81.9} \times K_f$$

Equating these two

$$A = 85$$

53. Answer (50)

$$HA \rightleftharpoons H^+ + A^-$$

 $1-\alpha \qquad \alpha \qquad \alpha$

 $(\alpha - degree of dissociation)$

van't Hoff factor (i) = 1 + α

Assuming given freezing point is -3.885°C

$$\Delta T_f = ik_f \cdot m$$

$$\Rightarrow$$
 3.885 = (1 + α) × 1.85 × 2

$$\Rightarrow$$
 (1 + α) = 1.05

$$\alpha = 0.05$$

$$= 50 \times 10^{-3}$$

54. Answer (100)

$$\begin{array}{ccc}
2A & \rightarrow & A_2 \\
1-\alpha & & \frac{\alpha}{2}
\end{array}$$

$$i = 1 - \alpha + \frac{\alpha}{2}$$

Also

$$\Delta T_h = iK_h m$$

$$0.52 = i \times 0.52 \times 2$$

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

$$\frac{1}{2} = 1 - \alpha + \frac{\alpha}{2}$$

$$\Rightarrow 1-\frac{\alpha}{2}=\frac{1}{2}$$

% of association of A is 100%

55. Answer (78)

$$P_{Total} = X_{R}P_{R}^{\circ} + X_{MR}P_{MR}^{\circ}$$

 ${\rm X_B-mole}$ fraction of benzene in solution phase ${\rm X_{MB}-mole}$ fraction of methyl benzene in solution phase

$$Y_{R}P_{Total} = X_{R}P_{R}^{\circ}$$

Y_B - mole fraction of benzene in vapor phase

$$Y_{B} = \frac{0.5 \times 70}{0.5 \times 70 + 0.5 \times 20} = 0.7777$$
$$= 77.77 \times 10^{-2}$$
$$\approx 78 \times 10^{-2}$$

56. Answer (1)

$$x_A = 0.6$$
 $P_T = x_A P_A^o + x_B P_B^o$
 $= 0.6 \times 90 + 0.4 \times 15$
 $= 54 + 6 = 60$
 $x_A P_A^o = y_A P_T$
 $0.6 \times 90 = y_A (60)$
 $\Rightarrow y_A = 0.9$

$$y_B = 0.1 = 1 \times 10^{-1}$$

57. Answer (2)

 $\Delta T_F \propto i$ (for equimolar solutions)

Solute	İ
KI	2
$Al_2(SO_4)_3$	5
$C_6H_{12}O_6$	1
K₂SO₄	3

58. Answer (25)

According to Henry's law

$$P_{gas} = X_{gas} \times K_{H}$$

$$X_{gas} = \frac{0.835}{1.67 \times 10^3} = 5 \times 10^{-4}$$

$$X_{gas} = \frac{n_{gas}}{n_{gas} + n_{H_2O}} \approx \frac{n_{gas}}{n_{H_2O}}$$

$$n_{gas} = 5 \times 10^{-4} \times \frac{900}{18} = 0.025$$
 mole or 25 mmol

59. Answer (250)

$$\Delta T_b = K_b m$$

$$0.60 = \frac{5.0 \times 3.00 \times 1000}{M \times 100}$$

$$M = 250$$

Molecular weight of the substance = 250

60. Answer (15)

$$\pi = iCRT$$

$$2.42 \times 10^{-3} = \frac{1 \times 1.46 \times 1000 \times 0.083 \times 300}{M \times 100}$$

 $M = 150223 \text{ g mol}^{-1}$

 $M = 15.0223 \times 10^4 \text{ amol}^{-1}$

61. Answer (125)

 $i = \frac{total \, number \, of \, particle \, after \, dissociation \, / \, association}{total \, number \, of \, particle \, before \, dissociation \, / \, association}$

let 'a' is total moles of HA

$$HA \longrightarrow H^+ + A^-$$
0.5a
0.5a

$$HA \longrightarrow (HA)_2$$
 $0.5a \longrightarrow 0.5a$

$$i = \frac{\left(0.5 + 0.5 + \frac{0.5}{2}\right)a}{a} = 1.25 = 125 \times 10^{-2}$$

62. Answer (4)

i for C₂H₅OH is 1.

$$\Delta T_f = ik_f m$$

 $\Delta T_f \propto i$ (considering molarity = molality)

$$Ba_3(PO_4)_2$$

$$i = 5$$

$$i = 3$$

$$i = 2$$

63. Answer (269)

$$\Delta T_f = (T_0 - T_s) = i \times Molality \times k_f$$

$$273 - T_s = 1 \times \frac{83 \times 1000}{62 \times 625} \times 1.86$$

$$T_{s} = 269$$

64. Answer (518)

Molality of sucrose solution = 0.75 m

Mass of sucrose = $0.75 \times 342 \text{ g} = 256.5 \text{ g}$

Mass of solutions = 1256.5 g

Mass of sucrose in 1 kg solution

$$=\frac{256.5\times1000}{1256.5}=204.1\,\mathrm{g}$$

Mass of water in 1 kg solution = 1000 - 204.1

$$= 795.9 g$$

After colling the solution to -4°C

$$4 = \frac{1.86 \times 204.1 \times 1000}{342 \times w_{B}^{'}}; w_{B}^{'} = 277.5 g$$

($w_{B}^{'}$ is the mass of water left)

Mass of ice separated = 795.9 - 277.5

$$= 518.4 \simeq 518 \text{ g}$$

65. Answer (271)

Moles of glucose =
$$\frac{40}{180}$$

$$\Delta T_f = iK_f m \qquad (i = 1 \text{ for glucose})$$
$$= 1 \times 1.86 \times \frac{40 \times 1000}{180 \times 200}$$

∴ Freezing point =
$$T_f - \Delta T_f$$

= 273.15 - 2.06
= 271.09
 \simeq 271

66. Answer (2)

$$\Delta T_f \propto i \times m$$

greater the value of i, greater will be the ΔT_f value.

Solute	i
glycine	1
KHSO ₄	3
hydrazine	1
alucosa	1

67. Answer (13)

$$K_b$$
 (benzene) = 2.6 K kg mol⁻¹

$$K_b$$
 (acetone) = 1.7 K kg mol⁻¹

In acetone

$$0.17 = \frac{1 \times 1.7 \times 1.22 \times 1000}{M \times 100}$$

M = 122 g/mol

In benzene

$$\Delta T_{b} = \frac{1}{2} \times \frac{1.22}{122} \times 2.6 \times 10$$

$$= 0.13 \text{ °C}$$

$$= 13 \times 10^{-2} \text{ °C}$$

68. Answer (54)

$$7.47 = C \times 0.083 \times 300$$

$$(\pi = CRT)$$

(Where C represents the concentration of glucose solution and π represents osmotic pressure)

$$C = \frac{7.47}{0.083 \times 300}$$
(mol L⁻¹)

which in gm/L =
$$\frac{7.47}{0.083 \times 300} \times 180$$

= 54 gm/l

69. Answer (1221)

According to Henry's law, partial pressure of a gas is given by

$$P_g = (K_H) X_g$$

where X_{α} is mole fraction of gas in solution

$$0.835 = 1.67 \times 10^3 \left(X_{CO_2} \right)$$

$$X_{CO_0} = 5 \times 10^{-4}$$

Mass of CO_2 in 1 L water = 1221 × 10⁻³ g

70. Answer (4)

Since, $\Delta T_{\ell} = ik_{\ell}m$

$$m = \frac{0.7}{93} \times \frac{1000}{42}$$

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

$$i = 0.6$$

$$\alpha = \frac{i-1}{\frac{1}{2}-1} = \frac{0.6-1}{\frac{1}{2}-1} = 0.8$$

Hence, percentage association of solute A is 80%.

71. Answer (98)

$$\Delta T_f = iK_b m$$

$$i = \frac{0.24 \times 99.5 \times 74.6}{1.80 \times 0.5 \times 1000} = 1.98$$

$$\alpha = \frac{i-1}{n-1} = \frac{0.98}{1} = 0.98$$

72. Answer (415)

$$\pi$$
 = i CRT

$$(i = 1)$$

$$\pi = \frac{2 \times 1000}{60 \times 10^3 \times 200} \times .083 \times 300$$

$$\pi = .00415$$
 atm

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

73. Answer (8)

$$\Delta T_b = k_b m$$

$$\frac{(\Delta T_b)_A}{(\Delta T_b)_B} = \frac{(k_b)_A}{(k_b)_B}$$

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

74. Answer (45)

$$W_{\text{solute}} = 2.5 \times 10^{-3} \text{ kg}$$

$$W_{\text{solvent}} = 75 \times 10^{-3} \text{ kg}$$

$$\Delta T_{h} = 373.535 - 373.15$$

$$= 0.385 K$$

 $K_b(H_2O) = 0.52 \text{ K kg mol}^{-1}$

$$\Delta T_{b} = \frac{K_{b} \times 10^{3} \times W_{solute}}{M_{solute} \times W_{solvent}}$$

$$M_{\text{solute}} = \frac{0.52 \times 10^{3} \times 2.5 \times 10^{-3}}{75 \times 10^{-3} \times 0.385}$$
$$= 45.02$$
$$\approx 45$$

75. Answer (14)

$$P_{T} = P_{A}^{0}.x_{A} + P_{B}^{0}.x_{B}$$

$$= 50 \times 0.3 + 100 \times 0.7$$

= 85 mm Hg

$$y_B = \frac{70}{85} = \frac{x}{17}$$

$$\frac{X}{17} = 14$$

76. Answer (330)

Since,

$$\pi = icRT$$

$$5.03 \times 10^{-3} = \frac{2.5}{M} \times \frac{1000}{500} \times 0.083 \times 300$$

Molar mass of protein = 24751.5 g/mol

Number of glycine units in protein =
$$\frac{24751.5}{75}$$

$$= 330$$

$$\Delta T_{b} = i \times K_{b} \times m$$

Moles of solute (acetic acid) =
$$\frac{1.2 \times 1.02}{60}$$

As moles of solute are very less.

So, take molarity and molality same.

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

$$i = 1.05$$

$$\alpha = \frac{i-1}{n-1} = \frac{0.05}{1} = 0.05$$

78. Answer (3)

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

$$\Delta T_{\epsilon} = i \times K_{\epsilon} \times m$$

$$i = 1$$

$$4 = 1 \times K_{h} \times 1.5$$

$$4 = 1 \times K_{r} \times 4.5$$

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

79. Answer (3)

 ΔT_r of formic acid = 0.0405°C

Concentration = 0.5 mL/L

and density = 1.05 g/mL

∴ Mass of formic acid in solution = 1.05 × 0.5 g

$$= 0.525 q$$

.. According to Van't Hoff equation,

$$\Delta T_f = ik_f \cdot m$$

$$0.0405 = i \times 1.86 \times \frac{0.525}{46 \times 1}$$

(Assuming mass of 1 L water = kg)

$$i = \frac{0.0405 \times 46}{1.86 \times 0.525} = 1.89 \approx 1.9$$

80. Answer (2)

$$\Delta T_{f} = i k_{f} \times m$$

$$\frac{\Delta T_{f(A)}}{\Delta T_{f(B)}} = \frac{1}{4}$$

$$\frac{i \times K_f \times \frac{1}{M_A} \times 1}{i \times K_f \times \frac{1}{M_A} \times 1} = \frac{1}{4}$$

$$\frac{M_B}{M_A} = \frac{1}{4}$$

$$M_{A}: M_{B} = 4:1$$

81. Answer (1)

Molality of a solution of non volatile solute (A) = 1

Elevation in boiling point is given by

$$\Delta T_{b} = K_{b} m$$

$$3 = K_b \times 1$$
 ... (1)

Molality of (A) in the same solvent = 2

Depression in freezing point is given by

$$\Delta T_f = K_f m$$

$$6 = K_{f} \times 2$$
 ... (2)

Dividing (1) by (2)

$$\frac{K_b}{K_f} = \frac{1}{X} = \frac{1}{1}$$

82. Answer (2)

$$(\Delta T_b)_A = (\Delta T_b)_B$$

$$K_b \cdot M_A = K_b \cdot M_B$$

$$\Rightarrow$$
 M_A = M_B

$$\Rightarrow \ \frac{\frac{2}{M_A}}{100} \times 1000 = \frac{\frac{8}{M_B}}{100} \times 1000$$

$$\Rightarrow$$
 M_B = 4M_A

83. Answer (06.00)

$$\frac{P_o-P_s}{P_s}=\frac{n_A}{n_B}$$

$$1 = \frac{n_A}{n_B}$$

$$n_A = n_B$$

:. Moles of solute added considering it as a non- electrolyte

$$=\frac{100}{18} \approx 5.55$$

84. Answer (15)

M.wt. of Acetic acid = 60 g

M.wt. of Ascorbic acid = 176 g

$$\Delta T_f = K_f m$$

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

$$\Delta T_f = 1.506$$

= 15.06 × 10⁻¹
= 15

85. Answer (2)

Given that $X_A = 0.2$, $Y_A = 0.5$, $P_T = 0.8$ atm

We know that $P_A = Y_A \times P_T$ $P_A = 0.5 \times 0.8 = 0.4$

Now
$$P_A = X_A \times P_A^o \Rightarrow P_A^o = \frac{0.4}{0.2} = 2$$
 atm

86. Answer (1)

From Henry's law,

$$X(oxygen) = \frac{p(oxygen)}{K_H} = \frac{0.920}{46.82 \times 10^3} = 1.96 \times 10^{-5}$$

As 1 litre of water contains 55.5 mol of it, therefore, \rightarrow n represents moles of O_2 in solution.

$$X(oxygen) = \frac{n}{n + 55.5} \approx \frac{n}{55.5} (n \ll 55.5)$$

$$\frac{n}{55.5} = 1.96 \times 10^{-5}$$

 $n = 108.8 \times 10^{-5} = 1.08 \times 10^{-3}$ moles

m moles of oxygen = $1.08 \times 10^{-3} \times 10^{3} = 1$ m mole

87. Answer (80)

$$\Delta T_f = k_f m$$

$$0.9 = \frac{2 \times 1.8 \times 1000}{62.5 \times 0.8 \times M}$$

$$M = \frac{2 \times 1800}{62.5 \times 0.8 \times 0.9}$$

= 80 g/mol