

# Solutions

## Question1

A solution of two miscible liquids showing negative deviation from Raoult's law will have :

[27-Jan-2024 Shift 1]

Options:

- A.  
increased vapour pressure, increased boiling point
- B.  
increased vapour pressure, decreased boiling point
- C.  
decreased vapour pressure, decreased boiling point
- D.  
decreased vapour pressure, increased boiling point

**Answer: D**

**Solution:**

Solution with negative deviation has

$$P_T < P_A^0 X_A + P_B^0 X_B$$

$$P_A < P_A^0 X_A$$

$$P_B < P_B^0 X_B$$

If vapour pressure decreases so boiling point increases.

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## Question2

The quantity which changes with temperature is:

[27-Jan-2024 Shift 2]

Options:

- A.  
Molarity



B.

Mass percentage

C.

Molality

D.

Mole fraction

**Answer: A**

**Solution:**

$$1. \text{ Molarity} = \frac{\text{Moles of solute}}{\text{Volume of solution}}$$

Since volume depends on temperature, molarity will change upon change in temperature.

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### Question3

**A solution of  $\text{H}_2\text{SO}_4$  is 31.4%  $\text{H}_2\text{SO}_4$  by mass and has a density of 1.25g/mL. The molarity of the  $\text{H}_2\text{SO}_4$  solution is M (nearest integer) [Given molar mass of  $\text{H}_2\text{SO}_4 = 98\text{gmol}^{-1}$ ]**

**[29-Jan-2024 Shift 1]**

**Options:**

**Answer: 4**

**Solution:**

$$M = \frac{n_{\text{solute}}}{V} \times 1000$$

$$= \frac{\left(\frac{31.4}{98}\right)}{\left(\frac{100}{1.25}\right)} \times 1000$$

$$= 4.005 \approx 4$$

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### Question4

**The osmotic pressure of a dilute solution is  $7 \times 10^5 \text{Pa}$  at 273K. Osmotic pressure of the same solution at 283K is  $\times 10^4 \text{Nm}^{-2}$ .**

**[29-Jan-2024 Shift 1]**



**Answer: 72.56**

**Solution:**

$$\pi = CRT$$

$$\Rightarrow \frac{\pi_1}{\pi_2} = \frac{T_1}{T_2}$$

$$\Rightarrow \pi_2 = \frac{\pi_1 T_2}{T_1} = \frac{7 \times 10^5 \times 283}{273}$$

$$= 72.56 \times 10^4 \text{Nm}^{-2}$$

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## Question5

Molality of 0.8M $\text{H}_2\text{SO}_4$  solution (density 1.06gcm $^{-3}$ ) is \_\_\_\_  $\times 10^{-3}$ m.

[29-Jan-2024 Shift 2]

**Answer: 815**

**Solution:**

$$m = \frac{M \times 1000}{d_{\text{sol}} \times 1000 - M \times \text{Molar mass}_{\text{solute}}}$$

$$815 \times 10^{-3} \text{m}$$

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## Question6

What happens to freezing point of benzene when small quantity of naphthalene is added to benzene?

[30-Jan-2024 Shift 1]

**Options:**

A.

Increases

B.

Remains unchanged



C.

First decreases and then increases

D.

Decreases

**Answer: D**

**Solution:**

On addition of naphthalene to benzene there is depression in freezing point of benzene.

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## Question7

The mass of sodium acetate ( $\text{CH}_3\text{COONa}$ ) required to prepare 250mL of 0.35M aqueous solution is \_\_\_\_\_g. (Molar mass of  $\text{CH}_3\text{COONa}$  is  $82.02\text{gmol}^{-1}$ )

[30-Jan-2024 Shift 1]

**Answer: 7**

**Solution:**

$$\text{Moles} = \text{Molarity} \times \text{Volume in litres}$$

$$= 0.35 \times 0.25$$

$$\text{Mass} = \text{moles} \times \text{molar mass}$$

$$= 0.35 \times 0.25 \times 82.02 = 7.18\text{g}$$

Ans. 7

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## Question8

The solution from the following with highest depression in freezing point/lowest freezing point is

[30-Jan-2024 Shift 2]

**Options:**

A.

180g of acetic acid dissolved in water

B.

180g of acetic acid dissolved in benzene

C.

180g of benzoic acid dissolved in benzene

D.

180g of glucose dissolved in water

**Answer: A**

**Solution:**

$\Delta T_f$  is maximum when  $i \times m$  is maximum.

$$1) m_1 = \frac{180}{60} = 3, i = 1 + \alpha$$

Hence

$$\Delta T_f = (1 + \alpha) \cdot k_f = 3 \times 1.86 = 5.58^\circ\text{C} (\alpha < 1)$$

$$2) m_2 = \frac{180}{60} = 3, i = 0.5, \Delta T_f = \frac{3}{2} \times k_f' = 7.68^\circ\text{C}$$

$$3) m_3 = \frac{180}{122} = 1.48, i = 0.5, \Delta T_f = \frac{1.48}{2} \times k_f' = 3.8^\circ\text{C}$$

$$4) m_4 = \frac{180}{180} = 1, i = 1, \Delta T_f = 1 \cdot k_f' = 1.86^\circ\text{C}$$

As per NCERT,  $k_f'(\text{H}_2\text{O}) = 1.86 \text{ k} \cdot \text{kg mol}^{-1}$

$k_f'(\text{Benzene}) = 5.12 \text{ k} \cdot \text{kg mol}^{-1}$

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## Question9

**If a substance ' A ' dissolves in solution of a mixture of ' B ' and ' C ' with their respective number of moles as  $n_A$ ,  $n_B$  and  $n_C$ , mole fraction of C in the solution is:**

**[30-Jan-2024 Shift 2]**

**Options:**

A.

$$\frac{n_C}{n_A \times n_B \times n_C}$$

B.

$$\frac{n_C}{n_A + n_B + n_C}$$

C.



$$\frac{n_C}{n_A + n_B + n_C}$$

D.

$$\frac{n_B}{n_A + n_B}$$

**Answer: B**

**Solution:**

$$\text{Mole fraction of } C = \frac{n_C}{n_A + n_B + n_C}$$

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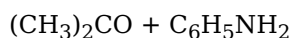
## Question 10

**Identify the mixture that shows positive deviations from Raoult's Law**

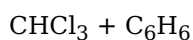
**[31-Jan-2024 Shift 1]**

**Options:**

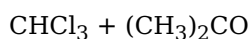
A.



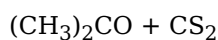
B.



C.



D.



**Answer: D**

**Solution:**

$(\text{CH}_3)_2\text{CO} + \text{CS}_2$  Exhibits positive deviations from Raoult's Law

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## Question 11

**We have three aqueous solutions of NaCl labelled as ' A ', ' B ' and ' C ' with concentration 0.1M, 0.01M & 0.001M, respectively. The value of**



van t' Haft factor (i) for these solutions will be in the order.

[1-Feb-2024 Shift 1]

Options:

A.

$$i_A < i_B < i_C$$

B.

$$i_A < i_C < i_B$$

C.

$$i_A = i_B = i_C$$

D.

$$i_A > i_B > i_C$$

Answer: A

Solution:

Salt	Values of i (for different conc. of a Salt)		
	0.1 M	0.01M	0.001M
NaCl	1.87	1.94	1.94

i approach 2 as the solution become very dilute.

## Question12

Mass of ethylene glycol (antifreeze) to be added to 18.6kg of water to protect the freezing point at  $-24^{\circ}\text{C}$  is \_\_\_ kg (Molar mass in  $\text{gmol}^{-1}$  for ethylene glycol 62,  $K_f$  of water =  $1.86\text{K kgmol}^{-1}$ )

[1-Feb-2024 Shift 2]

Answer: 15

Solution:

$$\Delta T_f = iK_f \times \text{molality}$$

$$24 = (1) \times 1.86 \times \frac{W}{62 \times 18.6}$$

$$W = 14880 \text{ gm}$$

$$= 14.880 \text{ kg}$$



## Question13

In the depression of freezing point experiment

- A. Vapour pressure of the solution is less than that of pure solvent
- B. Vapour pressure of the solution is more than that of pure solvent
- C. Only solute molecules solidify at the freezing point
- D. Only solvent molecules solidify at the freezing point

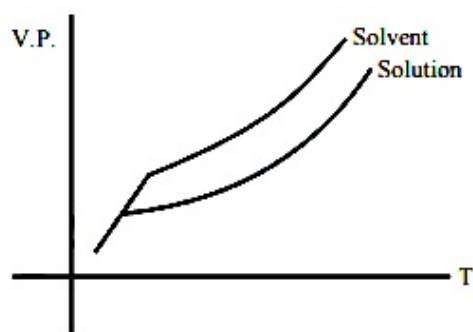
[24-Jan-2023 Shift 1]

Options:

- A. A and D only
- B. B and C only
- C. A and C only
- D. A only

Answer: A

Solution:



Vapour pressure (V.P.) of solvent is greater than vapour pressure (V.P.) of solution.  
Only solvent freezes.

## Question14

The Total pressure observed by mixing two liquid A and B is 350 mm Hg when their mole fractions are 0.7 and 0.3 respectively.

The Total pressure becomes 410 mm Hg if the mole fractions are changed to 0.2 and 0.8 respectively for A and B. The vapour pressure of pure A is \_\_\_ mm Hg. (Nearest integer)

Consider the liquids and solutions behave ideally.

[24-Jan-2023 Shift 2]

Answer: 314





## Solution:

Let V.P. of pure A be  $P_A^0$

Let V.P. of pure B be  $P_B^0$

When  $X_A = 0.7$  &  $X_B = 0.3$

$$P_s = 350$$

$$\Rightarrow P_A^0 \times 0.7 + P_B^0 \times 0.3 = 350 \dots (i)$$

When  $X_A = 0.2$  &  $X_B = 0.8$

$$P_s = 410$$

$$\Rightarrow P_A^0 \times 0.2 + P_B^0 \times 0.8 = 410 \dots (ii)$$

Solving (i) and (ii)

$$P_A^0 = 314 \text{ mm Hg}$$

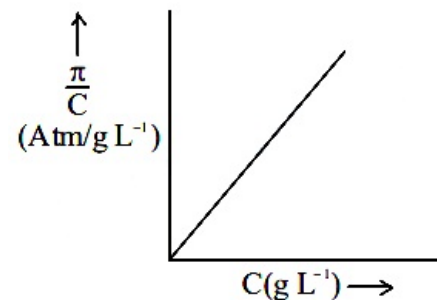
$$P_B^0 = 434 \text{ mm Hg}$$

$$= (314)$$

## Question 15

The osmotic pressure of solutions of PVC in cyclohexanone at 300K are plotted on the graph.

The molar mass of PVC is \_\_\_\_\_  $\text{g mol}^{-1}$   
(Nearest integer)



(Given :  $R = 0.083 \text{ L atm K}^{-1} \text{ mol}^{-1}$ )

[25-Jan-2023 Shift 1]

**Answer: 41500**

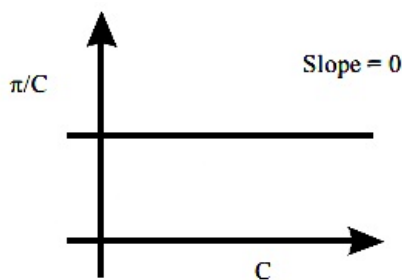
## Solution:

$$\pi = M' RT = \left( \frac{W/M}{V} \right) RT$$

$$\Rightarrow \pi = \left( \frac{W}{V} \right) \left( \frac{1}{M} \right) RT = C \left( \frac{RT}{M} \right)$$

$$\Rightarrow \frac{\pi}{C} = \frac{RT}{M} \neq f(c)$$

If we assume graph between  $\frac{\pi}{C}$  and C



Assuming  $\pi$  vs  $C$  graph

$$\text{Slope} = \frac{RT}{M} = \frac{0.083 \times 300}{M} = 6 \times 10^{-4}$$

$$\therefore M = \frac{0.083 \times 300}{6 \times 10^{-4}} = \frac{830 \times 300}{6}$$

$$= 41,500 \text{ gm / mole}$$

## Question16

The number of pairs of the solution having the same value of the osmotic pressure from the following is \_\_\_\_\_.

(Assume 100% ionization)

- A. 0.500M  $C_2H_5OH(aq)$  and 0.25M  $KBr(aq)$
- B. 0.100M  $K_4[Fe(CN)_6](aq)$  and 0.100M  $FeSO_4(NH_4)_2SO_4(aq)$
- C. 0.05M  $K_4[Fe(CN)_6](aq)$  and 0.25M  $NaCl(aq)$
- D. 0.15M  $NaCl(aq)$  and 0.1M  $BaCl_2(aq)$
- E. 0.02M  $KCl \cdot MgCl_2 \cdot 6H_2O(aq)$  and 0.05M  $KCl(aq)$

[25-Jan-2023 Shift 2]

**Answer: 4**

**Solution:**

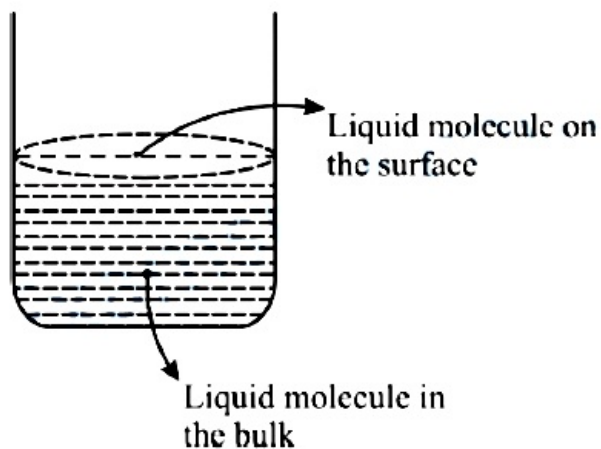
$$\Pi = iCRT$$

$$\Pi \propto iC$$

A, B, D and E have same value of osmotic pressure.

## Question17

Based on the given figure, the number of correct statement / s is/are \_\_\_\_\_



- A. Surface tension is the outcome of equal attractive and repulsion forces acting on the liquid molecule in bulk.**
- B. Surface tension is due to uneven forces acting on the molecules present on the surface.**
- C. The molecule in the bulk can never come to the liquid surface.**
- D. The molecules on the surface are responsible for vapour pressure if the system is a closed system.**
- [25-Jan-2023 Shift 2]

**Answer: 2**

**Solution:**

**Solution:**

B and D options are correct.

## Question 18

Solid Lead nitrate is dissolved in 1 litre of water. The solution was found to boil at  $100.15^{\circ}\text{C}$ . When 0.2 mol of NaCl is added to the resulting solution, it was observed that the solution froze at  $-0.8^{\circ}\text{C}$ . The solubility product of  $\text{PbCl}_2$  formed is \_\_\_\_\_  $\times 10^{-6}$  at 298K. (Nearest integer)

Given :  $K_b = 0.5\text{K kg mol}^{-1}$  and  $K_f = 1.8\text{ kg mol}^{-1}$ .

Assume molality to be equal to molarity in all cases.

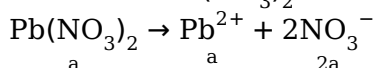
[29-Jan-2023 Shift 1]

**Answer: 13**

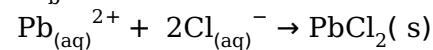
**Solution:**



Let a mole  $\text{Pb}(\text{NO}_3)_2$  be added



$$\Delta T_b = 0.15 = 0.5[3a] \Rightarrow a = 0.1$$



$$t = 0 \quad 0.1 \quad 0.2$$

$$t = \infty \quad (0.1 - x) \quad (0.2 - 2x)$$

In final solution

$$\Delta T_f = 0.8 = 1.8 \left[ \frac{0.3 - 3x + 0.2 + 0.2}{1} \right]$$

$$\Rightarrow x = \frac{2.3}{27}$$

$$\Rightarrow K_{sp} = \left( 0.1 - \frac{2.3}{27} \right) \left( 0.2 - \frac{4.6}{27} \right)^2 = 13 \times 10^{-6}$$

## Question 19

Match List I with List II.

List-I	List-II
A. van't Hoff factor, $i$	I. Cryoscopic constant
B. $k_f$	II. Isotonic solutions
C. Solutions with same osmotic pressure	III. $\frac{\text{Normal molar mass}}{\text{Abnormal molar mass}}$
D. Azeotropes	IV. Solutions with same composition of vapour above it

Choose the correct answer from the options given below :  
[29-Jan-2023 Shift 2]

Options:

A. A-III, B-I, C-II, D-IV

B. A-III, B-II, C-I, D-IV

C. A-III, B-I, C-IV, D-II

D. A-I, B-III, C-II, D-IV

Answer: A

Solution:

(A) van't Hoff factor,  $i$

$$i = \frac{\text{Normal molar mass}}{\text{Abnormal molar mass}}$$

(B)  $k_f$  = Cryoscopic constant

(C) Solutions with same osmotic pressure are known as isotonic solutions.

(D) Solutions with same composition of vapour over them are called Azeotrope.

## Question 20

## Match List-I and List-II.

List-I	List-II
A. Osmosis	I. Solvent molecules pass through semi permeable membrane towards solvent side.
B. Reverse osmosis	II. Movement of charged colloidal particles under the influence of applied electric potential towards oppositely charged electrodes.
C. Electro osmosis	III. Solvent molecules pass through semi permeable membrane towards solution side.
D. Electrophoresis	IV. Dispersion medium moves in an electric field.

**Choose the correct answer from the options given below:  
[29-Jan-2023 Shift 2]**

**Options:**

A. A-I, B-III, C-IV, D-II

B. A-III, B-I, C-IV, D-II

C. A-III, B-I, C-II, D-IV

D. A-I, B-III, C-II, D-IV

**Answer: B**

**Solution:**

A. Osmosis III

B. Reverse osmosis I

C. Electro osmosis IV

D. Electrophoresis II

## Question21

**A solution containing 2g of a non-volatile solute in 20g of water boils at 373.52K. The molecular mass of the solute is \_\_\_\_\_  $\text{gmol}^{-1}$ . (Nearest integer)**

**Given, water boils at 373K,  $K_b$  for water =  $0.52\text{K kg mol}^{-1}$**

**[30-Jan-2023 Shift 1]**

**Answer: 100**

**Solution:**

$$\Delta T_b = 373.52 - 373$$

$$= 0.52$$

$$\Delta T_b = K_b \cdot m$$

$$0.52 = 0.52 \times \frac{2}{\text{Molar Mass}} \times \frac{1}{20 \times 10^{-3}}$$

$$\text{Molar Mass} = 100 \text{g/mol}$$

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## Question22

A 300 mL bottle of soft drink has 0.2M CO<sub>2</sub> dissolved in it. Assuming CO<sub>2</sub> behaves as an ideal gas, the volume of the dissolved CO<sub>2</sub> at STP is \_\_\_\_\_ mL. (Nearest integer)

Given: At STP, molar volume of an ideal gas is 22.7 L mol<sup>-1</sup>

[30-Jan-2023 Shift 1]

**Answer: 1362**

**Solution:**

$$\text{Mole of CO}_2 = 0.2 \text{M} \times (300 \times 10^{-3}) \text{L}$$

$$= 0.06 \text{ Mole}$$

$$\text{Volume of 0.06 mole CO}_2 \text{ at S.T.P}$$

$$= 0.06 \times 22.7$$

$$= 1.362 \text{L}$$

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## Question23

Lead storage battery contains 38% by weight solution of H<sub>2</sub>SO<sub>4</sub>. The van't Hoff factor is 2.67 at this concentration. The temperature in Kelvin at which the solution in the battery will freeze is \_\_\_\_\_ (Nearest integer).

Given K<sub>f</sub> = 1.8 K kg mol<sup>-1</sup>

[30-Jan-2023 Shift 2]

**Answer: 243**

**Solution:**

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

$$\Rightarrow \Delta T_f = 2.67 \times 1.8 \times \frac{38}{98} \times \frac{1000}{62}$$

$$\Rightarrow \Delta T_f = 30.05$$

$$\therefore \text{F.P.} = 243 \text{K}$$

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## Question24

The total pressure of a mixture of non-reacting gases X (0.6g) and Y (0.45g) in a vessel is 740 mm of Hg. The partial pressure of the gas X is mm of Hg. (Nearest Integer)

(Given : molar mass X = 20 and Y = 45gmol<sup>-1</sup> )

[31-Jan-2023 Shift 1]

**Answer: 555**

**Solution:**

$$P_X = \chi_X P_T$$
$$= \frac{\frac{0.6}{20}}{\frac{0.6}{20} + \frac{0.45}{45}} \times 740$$
$$P_X = 555 \text{ mm Hg}$$

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## Question25

At 27°C, a solution containing 2.5g of solute in 250.0 mL of solution exerts an osmotic pressure of 400 Pa. The molar mass of the solute is \_\_\_\_\_ g mol<sup>-1</sup> (Nearest integer)

(Given : R = 0.083L bar K<sup>-1</sup> mol<sup>-1</sup> )

[31-Jan-2023 Shift 1]

**Answer: 62250**

**Solution:**

$$\Pi = CRT$$
$$\frac{400 \text{ Pa}}{10^5} = \frac{\frac{2.5 \text{ g}}{M_0}}{250 / 1000 \text{ L}} \times 0.083 \frac{\text{L} \cdot \text{bar}}{\text{K} \cdot \text{mol}} \times 300 \text{ K}$$
$$M_0 = 62250$$



## Question26

Evaluate the following statements for their correctness.

(A) The elevation in boiling point temperature of water will be same for 0.1M NaCl and 0.1M urea.

(B) Azeotropic mixtures boil without change in their composition

(C) Osmosis always takes place from hypertonic to hypotonic solution

(D) The density of 32%  $\text{H}_2\text{SO}_4$  solution having molarity 4.09M is approximately  $1.26\text{gmL}^{-1}$

(E) A negatively charged sol is obtained when KI solution is added to silver nitrate solution.

Choose the correct answer from the options given below :

[31-Jan-2023 Shift 2]

Options:

A. B, D, and E only

B. A, B, and D only

C. A and C only

D. B and D only

Answer: D

Solution:

Solution:

(A)  $\Delta T_b \propto i \times c$

(B) Azeotropic mixtures have same composition in both liquid and vapour phase.

(C) Osmosis always takes place from hypotonic to hypertonic solution.

(D)  $M = \frac{30 \times 10 \times 1.26}{98} \approx 4.09\text{M}$

(E) When KI solutions is added to  $\text{AgNO}_3$  solution, positively charged solution results due to adsorption of  $\text{Ag}^{\text{ions}}$  ion dispersion medium

$\text{AgI} / \text{Ag}^+$

Positively charged

## Question27

20% of acetic acid is dissociated when its 5g is added to 500 mL of water. The depression in freezing point of such water is \_\_\_\_\_  $\times 10^{-3}$  °C. Atomic mass of C, H and O are 12,1 and 16 a.m.u. respectively.

[Given : Molal depression constant and density of respectively.

[1-Feb-2023 Shift 2]

Answer: 372



### Solution:

$$i = 1 + (n - 1)\alpha$$

$$(i = 1 + 0.2(2 - 1) = 1.2$$

$$\Delta T_f = iK_f m$$

$$\Delta T_f = 1.2 \times 1.86 \times \frac{5 \times 1000}{60 \times 500}$$

$$\Delta t_f = 3.72$$

$$\Delta T_f = 372 \times 10^{-2}$$

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## Question28

Mass of Urea ( $\text{NH}_2\text{CONH}_2$ ) required to be dissolved in 1000g of water to reduce the vapour pressure of water by 25% is.....g. (Nearest integer)

Given: Molar mass of N, C, O and H are 14, 12, 16 and 1 mol<sup>-1</sup> respectively.

[6-Apr-2023 shift 1]

Answer: 1111

### Solution:

$$\frac{P^0 - P_s}{P_s} = \frac{n_{\text{solute}}}{n_{\text{solvent}}} = \frac{\frac{x}{60}}{\frac{1000}{18}} = \frac{P^0 - 0.75P^0}{0.75P^0}$$

$$\Rightarrow x = \frac{10000}{9} = 1111 \text{ gm}$$

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## Question29

Consider the following pairs of solution which will be isotonic at the same temperature. The number of pairs of solutions is/are.....

- A. 1M aq. NaCl and 2M aq. Urea
- B. 1M aq.  $\text{CaCl}_2$  and 1.5M aq. KCl
- C. 1.5M aq.  $\text{AlCl}_3$  and 2M aq.  $\text{Na}_2\text{SO}_4$
- D. 2.5M aq. KCl and 1M aq.  $\text{Al}_2(\text{SO}_4)_3$

[6-Apr-2023 shift 2]

Answer: 4

### Solution:

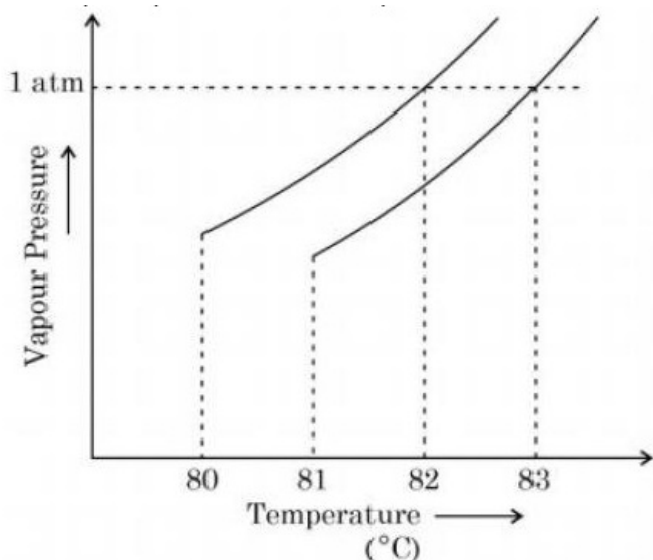
$$\pi = icRT$$

A, B, C and D are isotonic pairs.



## Question30

The vapour pressure vs. temperature curve for a solution solvent system is shown below:



The boiling point of the solvent is \_\_\_\_\_ °C  
[8-Apr-2023 shift 1]

Answer: 82

Solution:

Boiling point of solvent is 82°C

Boiling point of solvent is 83°C

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## Question31

If the boiling points of two solvents X and Y (having same molecular weights) are in the ratio 2 : 1 and their enthalpy of vaporizations are in the ratio 1 : 2, then the boiling point elevation constant of X is m times the boiling point elevation constant of Y. The value of m is \_\_\_\_\_ (nearest integer)

[8-Apr-2023 shift 2]

Options:

Answer: 8

Solution:

$$K_b = \frac{RT_b^2 m}{1000 \Delta H_{\text{vap}}}$$

$$\frac{(K_b)_x}{(K_b)_y} = \frac{(T_b^2 M)_x}{(T_b^2 M)_y} \times \frac{(\Delta H)_y}{(\Delta H)_x} = \left(\frac{2}{1}\right)^2 \times \left(\frac{2}{1}\right) = \frac{8}{1}$$

## Question32

If the degree of dissociation of aqueous solution of weak monobasic acid is determined to be 0.3 , then the observed freezing point will be \_\_\_\_\_ % higher than the expected/theoretical freezing point. (Nearest integer)  
[10-Apr-2023 shift 1]

**Answer: 30**

**Solution:**

For mono basic acid  $\rightarrow n = 2$

$$i = 1 + (n - 1)\alpha = 1 + (2 - 1)0.3$$

$$i = 1.3$$

$$\% \text{ increase} = \frac{(\Delta T_f)_{\text{obs}} - (\Delta T_f)_{\text{cal}}}{(\Delta T_f)_{\text{cal}}} \times 100$$

$$= \frac{K_f \times i \times m - K_f \times m}{K_f \times m} \times 100$$

$$= \frac{i - 1}{1} \times 100 = 30\%$$

## Question33

An aqueous solution of volume  $300\text{cm}^3$  contains 0.63g of protein. The osmotic pressure of the solution at 300 K is 1.29 mbar. The molar mass of the protein is \_\_\_\_\_  $\text{gmol}^{-1}$  Given :  $R = 0.083\text{L bar K}^{-1}\text{mol}^{-1}$   
[10-Apr-2023 shift 2]

**Answer: 40535**

**Solution:**

$$\because \pi = CRT$$

$$\pi = \frac{n}{V}RT$$

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

$$M = \frac{\omega RT}{\pi \times V}$$

$$M = \frac{0.63 \times 0.083 \times 300}{1.29 \times 10^{-3} \times 300 \times 10^{-3}}$$

$$M = 40535 \text{ gm / molL}$$

---

## Question34

**0.004 MK<sub>2</sub>SO<sub>4</sub> solution is isotonic with 0.01M glucose solution.**

**Percentage dissociation of K<sub>2</sub>SO<sub>4</sub> is (Nearest integer)**

**[11-Apr-2023 shift 1]**

**Answer: 75**

**Solution:**

For isotonic solution

$$(ic)_{\text{glucose}} = (ic)_{\text{K}_2\text{SO}_4}$$

$$0.01 = i(0.004)$$

$$i = \frac{0.01}{0.004} = \frac{10}{4} = \frac{5}{2}$$

$$1 + (n - 1)\alpha = \frac{5}{2}$$

$$1 + (3 - 1)\alpha = \frac{5}{2} \quad (\because n = 3 \text{ for } \text{K}_2\text{SO}_4)$$

$$2\alpha = \frac{3}{2}$$

$$\alpha = \frac{3}{4} \rightarrow 75\%$$

---

## Question35

**What weight of glucose must be dissolved in 100g of water to lower the vapour pressure by 0.20 mmHg ?**

**(Assume dilute solution is being formed)**

**Given : Vapour pressure of pure water is 54.2 mmHg at room temperature. Molar mass of glucose is 180gmol<sup>-1</sup>**

**[11-Apr-2023 shift 2]**

**Options:**

A. 2.59g

B. 3.59g

C. 3.69g

D. 4.69g

**Answer: C**

**Solution:**

$$\frac{P^0 - P_s}{P^0} = \frac{n}{N} \text{ (for dilute solution)}$$

$$\frac{0.2}{54.2} = \frac{n \times 18}{100}$$

$$n = \frac{100}{271 \times 18}$$

$$w = \frac{100 \times 180}{271 \times 18}; w = 3.69\text{g}$$

## Question36

**80 mole percent of  $\text{MgCl}_2$  is dissociated in aqueous solution. The vapour pressure of 1.0 molal aqueous solution of  $\text{MgCl}_2$  at  $38^\circ\text{C}$  is \_\_\_\_\_ mmHg. (Nearest integer)**

**Given : Vapour pressure of water at  $38^\circ\text{C}$  is 50 mm Hg.**

**[12-Apr-2023 shift 1]**

**Answer: 48**

**Solution:**



$$1 - \alpha \quad \alpha \quad 2\alpha$$

$$i = 1 + 2\alpha (\alpha = 0.8)$$

$$i = 2.6$$

$$\frac{\Delta p}{p^0} = \frac{i \times n_2}{n_1}$$

$$\Delta p = 2.34$$

$$p_s = 47.66$$

$$p_s \cong 48$$

## Question37

**Solution of 12g of non-electrolyte (A) prepared by dissolving it in 1000 mL of water exerts the same osmotic pressure as that of 0.05M glucose solution at the same temperature. The empirical formula of A is  $\text{CH}_2\text{O}$ . The molecular mass of A is \_\_\_\_\_ g. (Nearest integer)**

[13-Apr-2023 shift 1]

Answer: 240

Solution:

$$\Pi_A = \Pi_{\text{glucose}}$$
$$C_A RT = CRT$$

$$\frac{12 / M_A}{1} = 0.05$$

$$M_A (\text{ Molar mass of A}) = \frac{12}{0.05} = \frac{1200}{5} = 240 \text{ gm}$$

---

### Question38

Sea water contains 29.25%NaCl and 19%MgCl<sub>2</sub> by weight of solution.

The normal boiling point of the sea water is \_\_\_\_\_ °C (Nearest integer)

Assume 100% ionization for both NaCl and MgCl<sub>2</sub>

Given : K<sub>b</sub>(H<sub>2</sub>O) = 0.52K kg mol<sup>-1</sup>

Molar mass of NaCl and MgCl<sub>2</sub> is 58.5 and 95gmol<sup>-1</sup> respectively.

[13-Apr-2023 shift 2]

Answer: 116

Solution:

$$\text{Amount of solvent} = 100 - (29.25 + 19) = 51.75\text{g}$$

$$\Delta T_b = \left[ \frac{2 \times 29.25 \times 1000}{58.5 \times 51.75} + \frac{3 \times 19 \times 1000}{95 \times 51.75} \right] \times 0.52$$

$$\Delta T_b = 16.075$$

$$\Delta T_b = (T_b)_{\text{solution}} - (T_b)_{\text{solvent}}$$

$$(t)_{\text{solution}} = 100 + 16.07$$

$$= 116.07^\circ\text{C}$$

---

### Question39

The vapour pressure of 30%(w / v) aqueous solution of glucose is \_\_\_\_\_ mm Hg at 25°C.

[Given : The density of 30%(w / v), aqueous solutions of glucose is 1.2gcm<sup>-3</sup> and vapour pressure of pure water is 24 mm Hg.]



**(Molar mass of glucose is  $180\text{gmol}^{-1}$  )**  
**[15-Apr-2023 shift 1]**

**Answer: 23**

**Solution:**

$$\frac{24 - P_s}{P_s} = \frac{m \times 18}{1000}$$

$$\text{wt of solute} = 30 \text{ gm}$$

$$\text{Volume of solution} = 100 \text{ mL}$$

$$\text{wt. of solution} = 1.2 \times 100 = 120 \text{ gm}$$

$$\text{wt. of solvent} = 120 - 30 = 90 \text{ gm}$$

$$m \frac{30 \times 1000}{180 \times 90} = 185$$

$$\frac{24 - P_s}{P_s} = \frac{1.85 \times 18}{1000}$$

$$24 - P_s = 0.0333P_s$$

$$P_s(1.033) = 24$$

$$P_s = 23.22$$

---

## Question40

**The osmotic pressure of blood is 7.47 bar at 300K. To inject glucose to a patient intravenously, it has to be isotonic with blood. The concentration of glucose solution in  $\text{gL}^{-1}$  is**

**(Molar mass of glucose =  $180\text{gmol}^{-1}$ ,  $R = 0.083\text{Lbar}^{-1}\text{mol}^{-1}$  )**

**\_\_\_\_\_ (Nearest integer)**

**[24-Jun-2022-Shift-1]**

**Answer: 54**

**Solution:**

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

$$(\pi = CRT)$$

(Where C represents the concentration of glucose solution and  $\pi$  represents osmotic pressure)

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

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

$$= 54 \text{ gm / L}$$



## Question41

A company dissolves 'x' amount of  $\text{CO}_2$  at 298K in 1 litre of water to prepare soda water.  $X = \underline{\hspace{2cm}} \times 10^{-3}\text{g}$ . (nearest integer)

(Given : partial pressure of  $\text{CO}_2$  at 298K = 0.835 bar.

Henry's law constant for  $\text{CO}_2$  at 298K = 1.67 kbar.

Atomic mass of H, C and O is 1,12 , and 6 $\text{gmol}^{-1}$ , respectively)

[24-Jun-2022-Shift-2]

**Answer: 1221**

**Solution:**

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

$$P_g = (K_H)X_g$$

where  $X_g$  is mole fraction of gas in solution

$$0.835 = 1.67 \times 10^3 (X_{\text{CO}_2})$$

$$X_{\text{CO}_2} = 5 \times 10^{-4}$$

$$\text{Mass of CO}_2 \text{ in 1L water} = 1221 \times 10^{-3}\text{g}$$

---

## Question42

1L aqueous solution of  $\text{H}_2\text{SO}_4$  contains 0.02m mol  $\text{H}_2\text{SO}_4$ . 50% of this solution is diluted with deionized water to give 1L solution (A). In solution (A), 0.01m mol of  $\text{H}_2\text{SO}_4$  are added. Total m mols of  $\text{H}_2\text{SO}_4$  in the final solution is  $\underline{\hspace{2cm}} \times 10^3$  m mols

[25-Jun-2022-Shift-1]

**Answer: 0**

**Solution:**

$$n_{\text{H}_2\text{SO}_4} \text{ in Sol}^{\text{A}} = 50\% \text{ of original solution} = 0.01\text{m mol}$$

$$n_{\text{H}_2\text{SO}_4} \text{ in Final solution} = 0.01 + 0.01$$

$$= 0.02\text{m mol}$$

$$= 0.00002 \times 10^3\text{m mol}$$

The answer 0





## Question43

Solute A associates in water. When 0.7g of solute A is dissolved in 42.0g 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 = 93gmol<sup>-1</sup>. Molal depression constant of water is 1.86K kg mol<sup>-1</sup>.]

[25-Jun-2022-Shift-2]

Options:

- A. 50%
- B. 60%
- C. 70%
- D. 80%

Answer: D

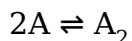
Solution:

$$\Delta T = i k_f \times m$$

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

$$i = \frac{0.2 \times 93 \times 6}{1.86 \times 100}$$

$$i = 0.60$$



$$1 - \alpha \quad \frac{\alpha}{2}$$

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

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

$$1 - \frac{\alpha}{2} = 0.60$$

$$1 - 0.60 = \frac{\alpha}{2}$$

$$\alpha = 0.80$$

## Question44

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 1.80K kg mol<sup>-1</sup> and molar mass of KCl is 74.6gmol<sup>-1</sup>)

[26-Jun-2022-Shift-1]



**Answer: 98**

**Solution:**

$$\Delta T_f = iK_b m$$

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

$$= 1.98$$

$$\alpha = \frac{i-1}{n-1} = \frac{0.98}{1} = 0.98 = 98\%$$

---

## Question45

The osmotic pressure exerted by a solution prepared by dissolving 2.0g of protein of molar mass  $60 \text{ kg mol}^{-1}$  in  $200 \text{ mL}^m$  of water at  $27^\circ\text{C}$  is \_\_\_\_\_ Pa. [integer value] (use  $R = 0.083 \text{ L}_{\text{bar}} \text{ mol}^{-1} \text{ K}^{-1}$ )

[26-Jun-2022-Shift-2]

**Answer: 415**

**Solution:**

$$\Pi = iCRT$$

$$= \frac{1 \times 2}{60000 \times 0.2} \times 0.083 \times 300$$

$$= 0.00415 \text{ bar } (\because 1 \text{ bar} = 10^5 \text{ Pa})$$

$$\text{So, } 0.00415 \times 10^5 \text{ Pa} = 415 \text{ Pa}$$

---

## Question46

Given below are two statements : one is labelled as Assertion (A) and the other is labelled as Reason (R).

**Assertion (A) :** At  $10^\circ\text{C}$ , the density of a 5M solution of KCl [atomic masses of K& Cl are 39&35.5 $\text{gmol}^{-1}$  respectively], is ' x '  $\text{gml}^{-1}$ . The solution is cooled to  $-21^\circ\text{C}$ . The molality of the solution will remain unchanged.

**Reason (R):** The molality of a solution does not change with temperature as mass remains unaffected with temperature.

In the light of the above statements, choose the correct answer from the options given below :

[27-Jun-2022-Shift-1]

Options:



- A. Both (A) and (R) are true and (R) is the correct explanation of (A).
- B. Both (A) and (R) are true but (R) is not the correct explanation of (A).
- C. (A) is true but (R) is false.
- D. (A) is false but (R) is true.

**Answer: A**

**Solution:**

Molality and Mass are temperature Independent so on changing temp., molality and mass remain unchanged.

### Question47

**2g of a non-volatile non-electrolyte solute is dissolved in 200g 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)**

**[27-Jun-2022-Shift-1]**

**Answer: 8**

**Solution:**

$$\Delta T_b = k_b m$$

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

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

$$\therefore y = 8$$

### Question48

**A solution containing  $2.5 \times 10^{-3}$  kg of a solute dissolved in  $75 \times 10^{-3}$  kg of water boils at 373.535K. The molar mass of the solute is \_\_\_\_g mol<sup>-1</sup>. [nearest integer] (Given :  $K_b(\text{H}_2\text{O}) = 0.52\text{K kg mol}^{-1}$  and boiling point of water = 373.15K )**

**[27-Jun-2022-Shift-2]**

**Answer: 45**

**Solution:**

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

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

$$\Delta T_b = 373.535 - 373.15 \\ = 0.385 \text{ K}$$

$$K_b(\text{H}_2\text{O}) = 0.52 \text{ K kg mol}^{-1}$$

$$\Delta T_b = \frac{K_b \times 10^3 \times W_{\text{solute}}}{M_{\text{solute}} \times W_{\text{solvent}}}$$

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

$$= 45.02$$

$$\approx 45$$

## Question 49

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 \_\_\_

[28-Jun-2022-Shift-1]

**Answer: 14**

**Solution:**

$$\frac{y_B}{1 - y_B} = \frac{P_B^0}{P_A^0} \left[ \frac{X_B}{1 - X_B} \right]$$

$$\Rightarrow \frac{y_B}{1 - y_B} = \frac{100}{50} \left[ \frac{0.7}{0.3} \right] = \frac{14}{3}$$

$$\Rightarrow y_B = \frac{14}{17}$$

## Question 50

1.2 mL of acetic acid is dissolved in water to make 2.0L 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 gmL<sup>-1</sup>, Molar mass of acetic acid is 60 gmol<sup>-1</sup>, K<sub>f</sub>(H<sub>2</sub>O) = 1.85 K kg mol<sup>-1</sup> ]



[29-Jun-2022-Shift-1]

Answer: 5

Solution:

$$M = d \times V = 1.02 \times 1.2 = 1.224 \text{ gm}$$

Moles of acetic acid = 0.0204 moles in 2L

So molality = 0.0102 mol / kg

$$\text{Now } \Delta T_f = i \times K_f \times M$$

$i = 1 + \alpha$  for acetic acid

$$0.0198 = (1 + \alpha) \times 1.85 \times 0.0102$$

$$\alpha = 0.04928$$

$$\cong 5\%$$

---

## Question51

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

[29-Jun-2022-Shift-2]

Answer: 3

Solution:

$$\Delta T_b = iK_b m$$

$$\Delta T_f = iK_f m$$

$$\frac{4}{4} = \frac{K_b 1.5}{K_f 4.5}$$

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

---

## Question52

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

[25-Jul-2022-Shift-1]

**Options:**

- A. 0.8
- B. 1.1
- C. 1.9
- D. 2.4

**Answer: C****Solution:****Solution:**

$$\Delta T_f \text{ of formic acid} = 0.0405^\circ\text{C}$$

$$\text{Concentration} = 0.5 \text{ mL / L}$$

$$\text{and density} = 1.05 \text{ g / mL}$$

$$\therefore \text{Mass of formic acid in solution} = 1.05 \times 0.5 \text{ g}$$

$$= 0.525 \text{ g}$$

$\therefore$  According to Van't Hoff equation,

$$\Delta T_f = i k_f \cdot m$$

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

(Assuming mass of 1L water = kg )

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

---

## Question53

**Two solutions A and B are prepared by dissolving 1g 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 [25-Jul-2022-Shift-2]**

**Options:**

- A. 1 : 4
- B. 1 : 0.25
- C. 1 : 0.20
- D. 1 : 5

**Answer: B****Solution:****Solution:**

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

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



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

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

$$M_A : M_B = 4 : 1$$

## Question54

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

[26-Jul-2022-Shift-2]

**Answer: 1**

**Solution:**

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

Elevation in boiling point is given by

$$\Delta T_b = K_b m$$

$$3 = K_b \times 1$$

Molality of (A) in the same solvent = 2

Depression in freezing point is given by

$$\Delta T_f = K_f m$$

$$6 = K_f \times 2 \dots (2)$$

Dividing (1) by (2)

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

$$\therefore X = 1$$

## Question55

Boiling point of a 2% aqueous solution of a non-volatile 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

[27-Jul-2022-Shift-1]

**Options:**

A.  $M_A = 4M_B$

B.  $M_B = 4M_A$

C.  $M_A = 8M_B$



$$D. M_B = 8M_A$$

**Answer: B**

**Solution:**

For A : 100 gm solution  $\rightarrow$  2 gm solute A

$$\therefore \text{Molality} = \frac{2 / M_A}{0.098}$$

For B : 100 gm solution  $\rightarrow$  8 gm solute B

$$\therefore \text{Molality} = \frac{8 / M_B}{0.092}$$

$$\therefore (\Delta T_B)_A = (\Delta T_B)_B$$

$\therefore$  Molality of A = Molality of B

$$\therefore \frac{2}{0.098M_A} = \frac{8}{0.092M_B}$$

$$\frac{2}{98} \times \frac{92}{8} = \frac{M_A}{M_B}$$

$$\frac{1}{4.261} = \frac{M_A}{M_B}$$

$$\therefore M_B = 4.261 \times M_A$$

---

## Question56

**When a certain amount of solid A is dissolved in 100g 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)**

**[27-Jul-2022-Shift-2]**

**Answer: 5.55**

**Solution:**

$\therefore$  Dilute solution given:

$$\frac{P^0 - P_s}{P^0} \sim \frac{n \text{ solute}}{n \text{ solvent}}$$

$$\frac{P^0 - P^0 / 2}{P^0} = \frac{n \text{ solute}}{n \text{ solvent}}$$

$$n \text{ Solute} \sim \frac{n \text{ solvent}}{2} = \frac{100}{18 \times 2} = 2.78 \text{ mol}$$

More accurate approach:

$$\frac{P^0 - P_s}{P_s} = \frac{n \text{ Solute}}{n \text{ solvent}}$$

$$\frac{P^0 - P^0 / 2}{P^0 / 2} = \frac{n \text{ solute}}{n \text{ solvent}}$$

$$n \text{ Solute} = n \text{ solvent} = \frac{100}{18} = 5.55 \text{ mol}$$



## Question57

150g of acetic acid was contaminated with 10.2g 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  $K_f = 3.9K kg mol^{-1}$ ; molar mass of ascorbic acid . =  $176gmol^{-1}$  ]

[28-Jul-2022-Shift-1]

Answer: 15

Solution:

M.wt. of Acetic acid = 60g

M.wt. of Ascorbic acid = 176g

$$\Delta T_f = K_f m$$

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

$$\Delta T_f = 1.506$$

$$= 15.06 \times 10^{-1}$$

$$= 15$$

---

## Question58

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)

[28-Jul-2022-Shift-2]

Answer: 2

Solution:

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

We know that  $P_A = Y_A \times P_T$

$$P_A = 0.5 \times 0.8 = 0.4$$

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

## Question59

If  $O_2$  gas is bubbled through water at 303K, 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 303K is 46.82k bar and partial pressure of  $O_2 = 0.920$  bar)

(Assume solubility of  $O_2$  in water is too small, nearly negligible)

[29-Jul-2022-Shift-1]

**Answer: 1**

**Solution:**

$$P = K_H \times X$$

$$0.920 \text{ bar} = 46.82 \times 10^3 \text{ bar} \times \frac{\text{mol of } O_2}{\text{mol of } O_2 + O}$$

$$0.920 = 46.82 \times 10^3 \times \frac{\text{mol of } O_2}{1000 / 18}$$

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

$$P = \frac{0.920}{46.82 \times 18} = n_{O_2}$$

$$\Rightarrow 1.09 \times 10^{-3} n_{O_2}$$

$$\Rightarrow m \text{ mol of } O_2 = 1$$

---

## Question60

'x' g of molecular oxygen ( $O_2$ ) is mixed with 200g of neon (Ne). The total pressure of the nonreactive 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. \_\_\_\_\_.

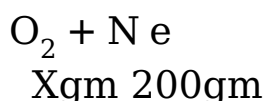
[Given: Molar mass of  $O_2 = 32 \text{ g mol}^{-1}$ .

Molar mass of Ne =  $20 \text{ g mol}^{-1}$  ]

[29-Jul-2022-Shift-2]

**Answer: 80**

**Solution:**



$$P_{\text{total}} = 25 \text{ bar} ; P_{\text{Ne}} = 20$$

$$P_{\text{O}_2} + P_{\text{Ne}} = 25$$

$$P_{\text{O}_2} = 25 - 20 = 5 \text{ bar}$$

$$5 = \frac{\frac{x}{32}}{\frac{x}{32} + \frac{200}{20}} \times 25$$

$$\frac{1}{5} = \frac{\frac{x}{32}}{\frac{x}{32} + 10}$$

$$\frac{1}{5} = \frac{x \times 32}{32(x + 320)}$$

$$5x = x + 320$$

$$4x = 320$$

$$x = \frac{320}{4} = 80 \text{ gm}$$

---

## Question61

1.80g of solute A was dissolved in  $62.5\text{cm}^3$  of ethanol and freezing point of the solution was found to be 155.1K. The molar mass of solute A is  $\text{g mol}^{-1}$  \_\_\_\_\_.

[Given : Freezing point of ethanol is 156.0K.

Density of ethanol is  $0.80\text{gcm}^{-3}$ .

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

[29-Jul-2022-Shift-2]

**Answer: 80**

**Solution:**

**Solution:**

$$\text{Mass of solvent} = d \times v = 0.8 \times 62.5 = 50 \text{ gram}$$

$$\Delta T_f = k_f \times m$$

$$0.9 = 2 \left[ \frac{1.8 \times 1000}{M_{\text{Solute}} \times 50} \right]$$

$$M_{\text{Solute}} = \left( \frac{2 \times 1.8 \times 1000}{0.9 \times 50} \right) = 80$$

---

## Question62

$\text{C}_6\text{H}_6$  freezes at  $5.5^\circ\text{C}$ . The temperature at which a solution 10g of  $\text{C}_4\text{H}_{10}$  in 200g of  $\text{C}_6\text{H}_6$  freeze is .....  $^\circ\text{C}$ . (The molal freezing point depression constant of  $\text{C}_6\text{H}_6$  is  $5.12^\circ\text{C} / \text{m}$ .)

[24 Feb 2021 Shift 2]

**Answer: 1**

**Solution:**

Pure solvent :  $C_6H_6(l)$

Given,  $T_f^\circ = 5.5^\circ C$

$K_f = 5.12^\circ C / m \Rightarrow m = 200g$

$m_{solute} = 10g$

Molar mass of solute  $C_4H_{10} = 12 \times 4 + 10 = 58$

Solute ( $C_4H_{10}$ ) is non-dissociative;

$\therefore i = 1$

$\therefore \Delta T_f = iK_f m$

$$\Rightarrow (T_f^\circ - T_f^1) = 1 \times 5.12 \times \frac{(10 / 58)}{(200 / 1000)}$$

$$5.5 - T_f^1 = 5.12 \times 5 \times 10$$

58

or  $T_f^1 \approx 1^\circ C$

## Question63

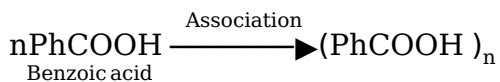
When 12.2g of benzoic acid is dissolved in 100g of water, the freezing point of solution was found to be  $-0.93^\circ C$  ( $K_f(H_2O) = 1.86K kg mol^{-1}$ ).

The number (n) of benzoic acid molecules associated (assuming 100% association) is .....

[26 Feb 2021 Shift 2]

**Answer: 2**

**Solution:**



Assuming 100% association ( $\alpha = 1$ ),

$$\Rightarrow i = 1 - \alpha \left( 1 - \frac{1}{n} \right) = \frac{1}{n} [\because \alpha = 1]$$

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

$$0 - (0.93) = 1.86 \times \frac{w_B \times 1000}{w_A \times M_B} \times \frac{1}{n}$$

[ $\because w_B = \text{mass of PhCOOH} = 12.2g$

$w_A = \text{mass of } H_2O = 100g$

$M_B = \text{molar mass of PhCOOH}$  ]

$= 122g mol^{-1}$

$$= 186 \times \frac{12.2 \times 1000}{100 \times 122} \times \frac{1}{n}$$

$$\Rightarrow n = \frac{1.86 \times 12.2 \times 1000}{0.93 \times 100 \times 122} = 2$$

$\therefore$  Number of benzoic acid molecules associated,  $n = 2$

## Question64

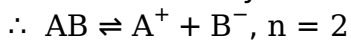
If a compound AB dissociates to the extent of 75% in an aqueous solution, the molality of the solution which shows a 2.5K rise in the boiling point of the solution is ..... molal. (Rounded off to the nearest integer) [ $K_b = 0.52\text{K kgmol}^{-1}$ ]

[25 Feb 2021 Shift 2]

**Answer: 3**

**Solution:**

As AB is a binary electrolyte,



$$i = 1 + \alpha(n - 1) = 1 + \frac{75}{100}(2 - 1) = 1.75$$

$$\text{Given, } \Delta T_b = 2.5\text{K}$$

$$K_b = 0.52\text{K kgmol}^{-1}$$

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

$$m = \frac{\Delta T_b}{K_b \times i} = \frac{2.5}{0.52 \times 1.75}$$

$$\therefore = 2.74 \sim 3\text{mol / kg}$$

---

## Question65

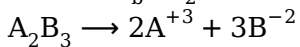
1 molal aqueous solution of an electrolyte  $A_2B_3$  is 60% ionised. The boiling point of the solution at 1atm is ..... K (Rounded off to the nearest integer). [Given,  $K_b$  for ( $H_2O$ ) =  $0.52\text{K kgmol}^{-1}$  ]

[25 Feb 2021 Shift 1]

**Answer: 375**

**Solution:**

$$\text{Given, } K_b(H_2O) = 0.52\text{K kgmol}^{-1}$$



No. of ions,  $n = 5$ , concentration,  $m = 1$  molal (Given)

$$\alpha = \frac{60}{100} = 0.6$$

$$\Delta T_b = i \cdot K_f \cdot m$$

$$= [(1 + (n - 1)\alpha) \times K_f \times m]$$

$$\Delta T_b = [(1 + (5 - 1)0.6) \times 0.52 \times 1]$$



$$\begin{aligned}
 &= (1 + 2.4) \times 0.52 \\
 \Delta T_b &= 1.768 \\
 T_b &= 1.768 + 373.15 \\
 &= 374.91 = 375\text{K}
 \end{aligned}$$


---

## Question66

When 9.45g of  $\text{ClCH}_2\text{COOH}$  is added to 500 mL of water, its freezing point drops by  $0.5^\circ\text{C}$ . The dissociation constant of  $\text{ClCH}_2\text{COOH}$  is  $x \times 10^{-3}$ . The value of  $x$  is \_\_\_\_\_ (Rounded off to the nearest integer)

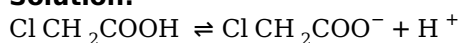
$$[K_{f(\text{H}_2\text{O})} = 1.86\text{K kgmol}^{-1}]$$

[24 Feb 2021 Shift 1]

**Answer: 35**

**Solution:**

**Solution:**



$$\text{Total dissociated} = 1 + \alpha$$

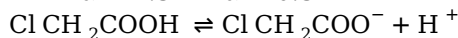
$$\Rightarrow i = 1 + \alpha$$

$$\Delta T_f = ik_1m$$

$$\Rightarrow 0.5 = (1 + \alpha)(1.86) \times \frac{9.45}{94.5} \times \frac{1000}{500}$$

$$\Rightarrow 0.5 = (1 + \alpha)(1.86)(0.2)$$

$$\Rightarrow 1 + \alpha = 1.34 \Rightarrow \alpha = 0.34$$



$$\text{Att} = 0 \quad \alpha \quad 0 \quad 0$$

$$\text{Attimet} \quad C - C\alpha \quad C\alpha \quad C\alpha$$

$$K_a = \frac{(C\alpha)^2}{C - C\alpha} = \frac{C\alpha^2}{1 - \alpha}$$

$$\Rightarrow K_a = \frac{0.2 \times (0.34)^2}{1 - 0.34} = 0.035$$

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


---

## Question67

A solute A dimerises in water. The boiling point of a 2 molar solution of A is  $100.52^\circ\text{C}$ .

The percentage association of A is (Round off to the nearest integer)

[Use :  $K_b$  for water =  $0.52\text{K kgmol}^{-1}$ , boiling point of water =  $100^\circ\text{C}$  ]

[18 Mar 2021 Shift 2]



**Answer: 100**

**Solution:**

Given :  $T_b = 100.52^\circ\text{C}$

$T_b^\circ = 100^\circ\text{C}$

$\Rightarrow K_b = 0.52\text{K kgmol}^{-1}$

Elevation in boiling point,

$$\Delta T_b = T_b - T_b^\circ$$

$$\Delta T_b = 100.52 - 100$$

$$\Delta T_b = 0.52^\circ\text{C}$$

For dimerisation (van't Hoff factor)  $i = \left(1 - \frac{\alpha}{2}\right)$

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

$$0.52 = \left(1 - \frac{\alpha}{2}\right) \times 0.52 \times 2$$

$$\alpha = 1$$

So, percentage association = 100%

---

## Question68

**2 molal solution of a weak acid HA has a freezing point of  $3.885^\circ\text{C}$ . The degree of dissociation of this acid is .....  $\times 10^{-3}$ . (Round off to the nearest integer).**

**[Given : Molal depression constant of water =  $1.85\text{K kgmol}^{-1}$ , freezing point of pure water =  $0^\circ\text{C}$  ]**

**[18 Mar 2021 Shift 1]**

**Answer: 50**

**Solution:**

Given

Depression in freezing point,

$$K_f = 1.85\text{K kgmol mol}^{-1}$$

$$T_f^\circ = 0^\circ\text{C}$$

$$T_f = 3.885^\circ\text{C}$$

$$\Delta T_f = T_f - T_f^\circ$$

$$= 3.885 - 0 = 3.885^\circ\text{C}$$

$$\Delta T_f = 3.885^\circ\text{C}$$

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

$$3.885 = i \times 1.85 \times 2$$

$$\text{but, } i = (1 + \alpha)$$

$$3.885 = (1 + \alpha) \times 1.85 \times 2$$

$$\Rightarrow \alpha = \frac{0.185}{3.7} \Rightarrow \alpha = 0.05$$

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

## Question69

A 1 molal  $K_4Fe(CN)_6$  solution has a degree of dissociation of 0.4. Its boiling point is equal to that of another solution which contains 18.1 weight per cent of a non-electrolytic solute A. The molar mass of A is ..... u. (Round off to the nearest integer).

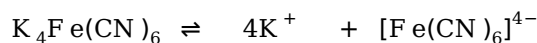
[Density of water =  $1.0\text{gcm}^{-3}$  ]

[17 Mar 2021 Shift 2]

**Answer: 85**

**Solution:**

**Solution:**



Initial conc.	1m	0	0
Final conc.	$(1 - 0.4)m$ = 0.6m	$4 \times 0.4$ = 1.6m	0.4m

Effective molality =  $0.6 + 1.6 + 0.4 = 2.6m$

As elevation in boiling point is a colligative property which depends on the amount of solute. So, to have same boiling point, the molality of two solutions should be same.

Molality of non-electrolyte solution = molality of  $K_4[Fe(CN)_6] = 2.6m$

Now, 18.1 weight per cent solution means 18.1g solute is present in 100g solution and hence,  $(100 - 18.1 = 81.9)$ g water.

$$\text{Molality} = \frac{(\text{Mass of solute} / \text{Molar mass of solute})}{\text{Mass of solvent (g)}} \times 1000$$

$$\text{Now, } 2.6 = \frac{18.1/M}{81.9/1000}$$

where, M is the molar mass of non-electrolyte solute

Molar mass of solute, M = 85

---

## Question70

The exact volumes of 1M NaOH solution required to neutralise 50mL of 1M  $H_3PO_3$  solution and 100mL of 2M  $H_3PO_2$  solution, respectively, are [16 Mar 2021 Shift 2]

**Options:**

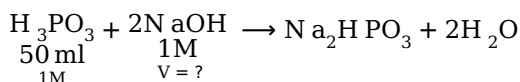
- A. 100mL and 100mL
- B. 100mL and 50mL
- C. 100mL and 200mL
- D. 50mL and 50mL

**Answer: C**

**Solution:**







Millimoles of  $\text{H}_3\text{PO}_3 = M \times V = 1 \times 50 = 50$

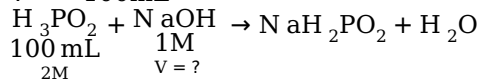
For 1 millimole of  $\text{H}_3\text{PO}_3$ , we require 2 millimoles of  $\text{NaOH}$ .

For 50 millimole of  $\text{H}_3\text{PO}_3$ , we require  $(2 \times 50) = 100$  millimoles of  $\text{NaOH}$ .

Millimoles of  $\text{NaOH} = M \times V = 100$

$$1 \times V = 100$$

$$V = 100\text{mL}$$



Millimoles of  $\text{H}_3\text{PO}_2 = M \times V = 2 \times 100 = 200$

For 1 millimole of  $\text{H}_3\text{PO}_2$ , we require 1 millimoles of  $\text{NaOH}$ .

For 200 millimole of  $\text{H}_3\text{PO}_2$ , we require 200 millimoles of  $\text{NaOH}$ .

So, volume of  $\text{NaOH} = 200\text{mL}$

## Question71

A 6.50 molal solution of  $\text{KOH}$  (aq) has a density of  $1.89\text{gcm}^{-3}$ . The molarity of the solution is .....  $\text{mol dm}^{-3}$  (Round off to the nearest integer).

[Atomic masses: K : 39.0u, O : 16.0u, H : 1.0u ]

[16 Mar 2021 Shift 1]

**Answer: 9**

**Solution:**

**Solution:**

$$\text{Molality (m)} = \frac{M \times 1000}{\{(1000d) - M \times M_{\text{solute}}\}}$$

where, M = molarity

d = density of solution

$M_{\text{solute}}$  = molar mass of solute

Putting the values,

$$6.50 = \frac{M \times 1000}{(1000 \times 1.89) - (M \times 56)}$$

$$\text{On solving, } M = \frac{12285}{1364}$$

$$M = 9 \text{ molar}$$

## Question72

At  $20^\circ\text{C}$ , the vapour pressure of benzene is 70 torr and that of methyl benzene is 20 torr. The mole fraction of benzene in the vapour phase at  $20^\circ\text{C}$  above an equimolar mixture of benzene and methyl benzene is

\_\_\_\_\_  $\times 10^{-2}$ . (Nearest integer)

[20 Jul 2021 Shift 1]

**Answer: 78**

**Solution:**

$$P_B^\circ = 40$$

$$P_T^\circ = 20$$

$$K_B = 0.5 = K_T$$

$$\begin{aligned} \text{Now } y_B &= \frac{K_B P_B^\circ}{K_B P_B^\circ + K_T P_T^\circ} \\ &= \frac{70 \times 0.5}{70 \times 0.5 + 20 \times 0.5} \end{aligned}$$

---

## Question73

1.46g of a biopolymer dissolved in a 100mL water at 300K exerted an osmotic pressure of  $2.42 \times 10^{-3}$  bar

The molar mass of the biopolymer is \_\_\_\_\_  $\times 10^4 \text{ gmol}^{-1}$ . (Round off to the Nearest Integer)

[Use :  $R = 0.083 \text{ L bar mol}^{-1} \text{ K}^{-1}$ ]

[27 Jul 2021 Shift 1]

**Answer: 15**

**Solution:**

**Solution:**

$$\pi = CRT$$

$\pi$  = osmotic pressure

C = molarity

T = Temperature of solution

let the molar mass be M gm / mol

$$2.42 \times 10^{-3} \text{ bar} =$$

$$\left( \frac{1.46 \text{ g}}{M \text{ gm / mol}} \right) \times \left( \frac{0.083 \text{ l - bar}}{\text{mol - K}} \right) \times (300 \text{ K})$$

$$\Rightarrow M = 15.02 \times 10^4 \text{ g / mol}$$

---

## Question74

When 3.00g of a substance 'X' is dissolved in 100g of  $\text{CCl}_4$ , it raises the boiling point by 0.60K. The molar mass of the substance 'X' is \_\_\_\_\_  $\text{g mol}^{-1}$ . (Nearest integer).

[ Given  $K_b$  for  $\text{CCl}_4$  is  $5.0 \text{ K kg mol}^{-1}$  ]

[25 Jul 2021 Shift 2]

**Answer: 250**

**Solution:**

$$\Delta T_b = K_b \times \text{molality}$$

$$0.60 = 5 \times \left( \frac{3 / M}{100 / 100} \right)$$

$$M = 250$$

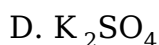
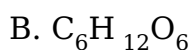
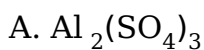
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## Question75

**Which one of the following 0.06M aqueous solutions has lowest freezing point?**

**[22 Jul 2021 Shift 2]**

**Options:**



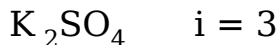
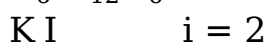
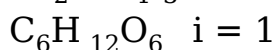
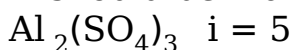
**Answer: A**

**Solution:**

$$T_f - T_f' = iK_f m$$

For minimum  $T_f'$

'i' should be maximum.



## Question76

**The water having more dissolved  $\text{O}_2$  is :**

**[22 Jul 2021 Shift 2]**

**Options:**

A. boiling water

B. water at  $80^\circ\text{C}$

C. polluted water

D. water at 4°C

**Answer: D**

**Solution:**

**Solution:**

On heating concentration of O<sub>2</sub> in water decreases.

So boiling water and water at 80°C having less O<sub>2</sub> concentration. Polluted water also having less O<sub>2</sub> concentration. So water at 4°C having maximum O<sub>2</sub> concentration.

---

## Question77

The density of NaOH solution is 1.2gcm<sup>-3</sup>. The molality of this solution is \_\_\_\_\_ m.

(Round off to the Nearest Integer)

[Use : Atomic masses : Na : 23.0u O : 16.0u H : 1.0u

Density of H<sub>2</sub>O : 1.0gcm<sup>-3</sup> ]

[27 Jul 2021 Shift 1]

**Answer: 5**

**Solution:**

Consider 1l solution

mass of solution = (1.2 × 1000)g

= 1200gm

Neglecting volume of NaOH

Mass of water = 1000gm

⇒ Mass of NaOH = (1200 – 1000)gm

= 200gm

⇒ Moles of NaOH =  $\frac{200\text{g}}{50\text{g/mol}} = 5\text{mol}$

⇒ molality =  $\frac{5\text{mol}}{1\text{kg}} = 5\text{m}$

---

## Question78

CO<sub>2</sub> gas is bubbled through water during a soft drink manufacturing process at 298K. If CO<sub>2</sub> exerts a partial pressure of 0.835 bar then xm mol of CO<sub>2</sub> would dissolve in 0.9L of water. The value of x is \_\_\_\_\_.

(Nearest integer)

(Henry's law constant for CO<sub>2</sub> at 298K is 1.67 × 10<sup>3</sup> bar )



## [25 Jul 2021 Shift 1]

**Answer: 25**

**Solution:**

From Henry's law

$$P_{\text{gas}} = K_H \cdot X_{\text{gas}}$$

$$0.835 = 1.67 \times 10^3 \times \frac{n(\text{CO}_2)}{\frac{0.9 \times 1000}{18}}$$

$$n(\text{CO}_2) = 0.025$$

$$\text{Millimoles of CO}_2 = 0.025 \times 1000 = 25$$

---

## Question79

The vapour pressures of A and B at 25°C are 90mmHg and 15mmHg respectively. If A and B are mixed such that the mole fraction of A in the mixture is 0.6, then the mole fraction of B in the vapour phase is  $x \times 10^{-1}$ . The value of x is \_\_\_\_\_. (Nearest integer)

[20 Jul 2021 Shift 2]

**Answer: 1**

**Solution:**

Given  $P_A^\circ = 90\text{mmHg}$

, at 25°C

$P_B^\circ = 15\text{mmHg}$

$$\text{and } \left. \begin{array}{l} X_A = 0.6 \\ X_B = 0.4 \end{array} \right\} P_T = X_A P_A^\circ + X_B P_B^\circ$$

$$= (0.6 \times 90) + (0.4 \times 15)$$

$$= 54 + 6 = 60\text{mm}$$

Now mol fraction of B in the vapour phase

$$\text{i.e. } Y_B = \frac{P_B}{P_T} = \frac{X_B P_B^\circ}{60} = 0.1 = 1 \times 10^{-1}$$

therefore:  $x = 1$

---

## Question80

40g of glucose (Molar mass = 180) is mixed with 200 mL of water. The freezing point of solution is ..... K. (Nearest integer)



**[Given,  $K_f = 1.86 \text{ K kg mol}^{-1}$ , density of water =  $1.00 \text{ g cm}^{-3}$ , freezing point of water =  $273.15 \text{ K}$ ]  
[27 Aug 2021 Shift 2]**

**Answer: 271**

**Solution:**

40 g of glucose mixed with 200 mL of water  
180 g of glucose = 1 moles of glucose  
40 g of glucose mol = 0.22 mol  
1 mL of water = 1g of water [ $d = 1 \text{ g / cm}^3$ ,  $1 \text{ mL} = 1 \text{ cm}^3$ ]  
200 mL of water = 200g of water

$$\Delta T_f = K_f m$$

where,  $\Delta T_f$  = depression in freezing point,

$K_f$  = molal elevation constant =  $1.86 \text{ K kg mol}^{-1}$  and

$m$  = molality of solution

$$\Delta T_f = \frac{1.86 \times 0.22}{200} \times 1000, (\because 1000 \text{ g} = 1 \text{ kg})$$

$$\Delta T_f = 2 \text{ K}$$

$$\therefore (T_f - T'_f) = 2 \left[ \begin{array}{l} T_f = \text{freezing point of water}(273 \text{ K}) \\ T'_f = \text{freezing point of solution} \end{array} \right]$$

$$273 \text{ K} - T'_f = 2 \text{ K}$$

$$T'_f = (273 - 2) \text{ K} = 271 \text{ K}$$

## Question 81

**1 kg of 0.75 molar aqueous solution of sucrose can be cooled up to  $-4^\circ \text{C}$  before freezing. The amount of ice (in g) that will be separated out is .....** (Nearest integer)

**[Given,  $K_f(\text{H}_2\text{O}) = 1.86 \text{ K kg mol}^{-1}$ ]**

**[27 Aug 2021 Shift 1]**

**Answer: 518**

**Solution:**

Let mass of water (initially present) =  $x \text{ g}$

Mass of sucrose =  $(1000 - x) \text{ g}$

$$\text{Moles of sucrose} = \frac{1000 - x}{342}$$

$$\text{Molality} = \frac{\text{moles of sucrose}}{\text{mass of water (initially)}}$$

$$0.75 = \frac{\left(\frac{1000-x}{342}\right)}{\left(\frac{x}{1000}\right)}$$

$$\frac{x}{1000} = \frac{1000-x}{342 \times 0.75}$$

$$256.5x = 10^6 - 1000x$$

$$\Rightarrow x = 795.86 \text{ g}$$

$$\text{Moles of sucrose} = 0.5969$$

$$\text{New mass of H}_2\text{O} = a \text{ kg}$$

$$\text{Depression in freezing point } \Delta T_f = K_f \times m$$

$$4 = \frac{0.5969}{a} \times 1.86$$

$$\Rightarrow a = 0.2775 \text{ kg}$$

$$\text{Ice separated} = 795.86 - 277.5$$

$$= 518.3 \text{ g} \approx 518 \text{ g.}$$

## Question82

If the degree of dissociation of aqueous solution of weak monobasic acid is determined to be 0.3, then the observed freezing point will be \_\_\_\_\_ % higher than the expected/theoretical freezing point. (Nearest integer) [27 Aug 2021 Shift 1]

**Answer: 30**

**Solution:**

For mono basic acid  $\rightarrow n = 2$

$$i = 1 + (n-1)\alpha = 1 + (2-1)0.3$$

$$i = 1.3$$

$$\% \text{ increase} = \frac{(\Delta T_f)_{\text{obs}} - (\Delta T_f)_{\text{cal}}}{(\Delta T_f)_{\text{cal}}} \times 100$$

$$= \frac{K_f \times i \times m - K_f \times m}{K_f \times m} \times 100$$

$$= \frac{i-1}{1} \times 100 = 30\%$$

## Question83

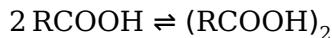
1.22g of an organic acid is separately dissolved in 100g of benzene ( $K_b = 2.6 \text{ K kg mol}^{-1}$ ) and 100g of acetone ( $K_b = 1.7 \text{ K kg mol}^{-1}$ ). The acid is known to dimerise in benzene but remain as a monomer in acetone. The boiling point of the solution in acetone increases by  $0.17^\circ\text{C}$ . The increase in boiling point of solution in benzene in  $^\circ\text{C}$  is  $x \times 10^{-2}$ . The value of  $x$  is..... (Nearest integer)

**[Atomic mass: C = 12.0, H = 1.0, O = 16.0]  
[31 Aug 2021 Shift 2]**

**Answer: 13**

**Solution:**

With benzene as solvent, RCOOH dimerises.



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

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

where,  $\Delta T_b$  = boiling point elevation,

$K_b$  = ebulliscopic constant

$m$  = molality

$i$  = van't Hoff factor

$$\Delta T_b = \frac{1}{2} \times 2.6 \times \frac{1.22 / M_w}{100 / 1000} \dots(i)$$

With acetone as solvent, no dimerisation.

$$\therefore i = 1$$

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

$$0.17 = 1 \times 1.7 \times \frac{1.22 / M_w}{100 / 1000} \dots(ii)$$

Eq. (i) divide by Eq. (ii),

$$\frac{\Delta T_b}{0.17} = \frac{\frac{1}{2} \times 2.6 \times \frac{1.22 / M_w}{100 / 1000}}{1 \times 1.7 \times \frac{1.22 / M_w}{100 / 1000}}$$

$$\Rightarrow \Delta T_b = \frac{0.26}{2} = 13 \times 10^{-2}$$

$$x = 13$$

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## Question84

**Which one of the following 0.10M aqueous solutions will exhibit the largest freezing point depression?**

**[31 Aug 2021 Shift 1]**

**Options:**

- A. Hydrazine
- B. Glucose
- C. Glycine
- D.  $\text{KHSO}_4$

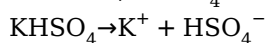
**Answer: D**

**Solution:**





The van't Hoff factor (i) is highest for  $\text{KHSO}_4$  as it dissociates into its ions. van't Hoff factor is the ratio of total number of ions produced after complete dissociation to before dissociation. Hydrazine, glucose and glycine will not dissociate to produce ions. Whereas,  $\text{KHSO}_4$  dissociates into ions.



Hence, i will be highest for  $\text{KHSO}_4$ .

$\therefore$  Colligative property i.e. freezing point depression will be highest for  $\text{KHSO}_4$  as  $\Delta T_f \propto i$  [ $\because$  m is constant i.e. 0.1]

Hence, correct option is (d).

## Question 85

In a solvent 50% of an acid HA dimerizes and the rest dissociates. The van't Hoff factor of the acid is \_\_\_\_\_  $\times 10^{-2}$ .

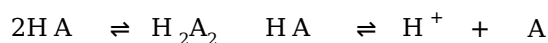
(Round off to the nearest integer)

[27 Jul 2021 Shift 2]

**Answer: 125**

**Solution:**

**Solution:**



Initial moles	$a \times \frac{50}{100}$	0	$a \times \frac{50}{100}$	0	0
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Final moles	0	0.25a	0	0.5a	0.5a
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$$\text{Now, } i = \frac{\text{final moles}}{\text{initial moles}} = \frac{0.25a + 0.5a + 0.5a}{0.5a + 0.5a}$$

$$= 1.25 = 125 \times 10^{-2}$$

## Question 86

83g of ethylene glycol dissolved in 625g of water. The freezing point of the solution is .....K. (Nearest integer)

[Use, molal freezing point depression constant of water

=  $1.86 \text{K kg mol}^{-1}$ , Freezing point of water = 273K and Atomic masses :

C = 12.0u, O = 16.0u, H = 1.0u ]

[26 Aug 2021 Shift 2]

**Answer: 269**

**Solution:**

$$\Delta T_f = iK_f \times m$$

For ethylene glycol,  $i = 1$

$$\text{Molality (m)} = \frac{\text{Number of moles of solute}}{\text{Weight of solvent(kg)}}$$

$$\Delta T_f = 1 \times 1.86 \times \frac{83 \times 1000}{62 \times 625} \approx 3.98$$

$$\begin{aligned} \text{Freezing point of solution} &= \text{Freezing point} - \Delta T_f \text{ of water} \\ &= 273 - 3.98 \approx 269. \end{aligned}$$

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## Question87

**Of the following four aqueous solutions, total number of those solutions whose freezing point is lower than that of 0.10M**

**$C_2H_5OH$  is ..... . (Integer answer)**

**(i) 0.10M  $Ba_3(PO_4)_2$**

**(ii) 0.10M  $Na_2SO_4$**

**(iii) 0.10M  $KCl$**

**(iv) 0.10M  $Li_3PO_4$**

**[26 Aug 2021 Shift 1]**

**Answer: 4**

**Solution:**

**Solution:**

As 0.10M  $C_2H_5OH$  is non-dissociative in nature and all salts

[0.10M  $Ba_3(PO_4)_2$ , 0.10M  $Na_2SO_4$ ,

0.10M  $KCl$ , 0.10M  $Li_3PO_4$ ] are electrolyte and dissociative in nature.

$$\Delta T_f = iK_f m$$

Where  $i$  is van't Hoff factor.

$i > 1$  for all other electrolytes where dissociation takes place.

So, for all these electrolytes effective molarity is more than 0.10. So, all the electrolytes given will have low freezing point.

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## Question88

**The molarity of the solution prepared by dissolving 6.3g of oxalic acid ( $H_2C_2O_4 \cdot 2H_2O$ ) in 250 mL of water in  $mol L^{-1}$  is  $x \times 10^{-2}$ . The value of  $x$**

**is ..... (Nearest integer)**

**[Atomic mass H = 1.0, C = 12.0, O = 16.0]**

**[31 Aug 2021 Shift 1]**

**Answer: 20**

**Solution:**



$$\begin{aligned} \text{Molarity} &= \frac{\text{number of moles of solute}}{\text{volume of solution(in litre)}} \\ &= \frac{\text{weight of solute / molecular mass of solute}}{\text{volume of solution(in litre)}} \\ \text{Molecular mass of solute [H}_2\text{C}_2\text{O}_4\cdot 2\text{H}_2\text{O}] &= 126\text{g} \\ x \times 10^{-2} &= \frac{6.3\text{g} / 126\text{g}}{250\text{ mL}} \times 1000 \\ x \times 10^{-2} &= 0.2 \\ x \times 10^{-2} &= 20 \times 10^{-2} \\ \therefore x &= 20 \end{aligned}$$


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## Question89

**100 mL of Na<sub>3</sub>PO<sub>4</sub> solution contains 3.45g of sodium. The molarity of the solution is ..... × 10<sup>-2</sup> mol L<sup>-1</sup>. (Nearest integer)  
[ Atomic masses – Na = 23.0u, O = 16.0u,, P = 31.0u ]  
[26 Aug 2021 Shift 2]**

**Answer: 50**

**Solution:**

**Solution:**

Number of mole of Na<sup>+</sup> ions

$$= \frac{\text{mass of sodium}}{\text{molar mass of sodium}} = \frac{3.45}{23}$$

So, number of mole of Na<sub>3</sub>PO<sub>4</sub> =  $\frac{\text{moles of sodium}}{3}$

{as 1 molecule of Na<sub>3</sub>PO<sub>4</sub> contains 3 atoms of Na }

$$= \frac{3.45}{3 \times 23} = 0.05$$

$$\text{Molarity} = \frac{\text{Moles of Na}_3\text{PO}_4}{\text{Volume of solution(in L)}} = \frac{0.05}{100} \times 1000$$

∴ Answer is 50

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## Question90

**If 80g of copper sulphate CuSO<sub>4</sub>·5H<sub>2</sub>O. is dissolved in deionised water to make 5L of solution. The concentration of the copper sulphate solution is x × 10<sup>-3</sup> mol L<sup>-1</sup>. The value of x is ..... .  
[Atomic masses Cu = 63.54u, S = 32u, O = 16u, H = 1u ]  
[1 Sep 2021 Shift 2]**

**Answer: 64**

**Solution:**



Given, mass of  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O} = 80\text{g}$

The concentration of copper sulphate solution is  $x \times 10^{-3} \text{ mol / L}$ .

$$\text{Molarity} = \frac{\text{Number of moles of solute}}{\text{Volume of solution(L)}} \dots(i)$$

$$\begin{aligned} \text{Molar mass of } \text{CuSO}_4 \cdot 5\text{H}_2\text{O} &= 63.54 + 32 + 16 \times 4 \\ &= 5 \times 18 = 249.54 \text{ g/mol} \end{aligned}$$

$$\begin{aligned} \text{Number of moles of solute} &= \frac{\text{Weight of solute}}{\text{Molecular mass of solute}} \\ &= \frac{80\text{g}}{249.54\text{g / mol}} = 0.32 \text{ mol} \end{aligned}$$

Volume of solution = 5 L

From Eq. (i),

$$\text{Molarity} = \frac{0.3205}{5} = 64.11 \times 10^{-3} \text{ mol / L}$$

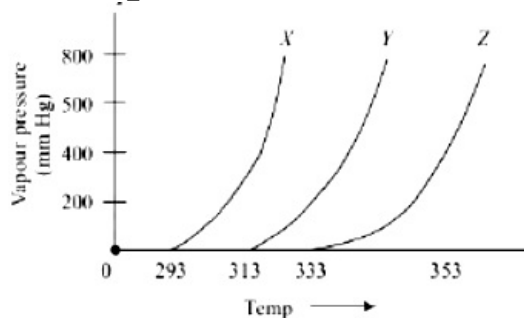
$$\therefore x = 64.11$$

or  $x \approx 64$

Hence, answer is 64.

## Question91

**The vapour pressure of solution will be less than the vapour pressure of pure solvent, so some vapour molecules will get condensed to maintain new equilibrium.**



**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 inference(s) is/are:**

**[Jan. 08, 2020(I)]**

**Options:**

A. (A) and (C)

B. (A)

C. (B)

D. (C)

**Answer: C**

**Solution:**

**Solution:**

At a particular temperature as intermolecular force of attraction increases vapour pressure decreases. Thus, intermolecular forces are inversely proportional to vapour pressure and directly proportional to temperature. Therefore X has lower intermolecular interactions compared to Y.

## Question92

At 35°C, the vapour pressure of CS<sub>2</sub> is 512mmHg and that of acetone is 344mmHg. A solution of CS<sub>2</sub> in acetone has a total vapour pressure of 600mmHg. The false statement amongst the following is:  
[Jan. 07,2020(I)]

Options:

- A. Raoult's law is not obeyed by this system
- B. a mixture of 100mLCS<sub>2</sub> and 100mL acetone has a volume <200mL
- C. CS<sub>2</sub> and acetone are less attracted to each other than to themselves
- D. heat must be absorbed in order to produce the solution at 35°C

Answer: B

Solution:

Solution:

Mixture of carbon disulphide and acetone will show positive deviation from Raoult's Law. The dipolar interaction between solute (CS<sub>2</sub>) solvent (acetone) molecules in solution are weaker. So the vapour pressure of solution will be greater than the individual vapour pressure of pure components.

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## Question93

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:  
[Jan. 07,2020 (II)]

Options:

- A. the volume of the solution increases and the volume of the solvent decreases
- B. the volume of the solution decreases and the volume of the solvent increases
- C. the volume of the solution and the solvent does not change
- D. the volume of the solution does not change and the volume of the solvent decreases

Answer: A

Solution:

Solution:

There will be lowering in vapour pressure in second beaker.



## Question94

How much amount of NaCl should be added to 600g of water ( $\rho = 1.00\text{g / mL}$ ) to decrease the freezing point of water to  $-0.2^\circ\text{C}$ ? \_\_\_\_\_ . (The freezing point depression constant for water =  $2\text{K kgmol}^{-1}$ )  
[NV, Jan. 09,2020 (I)]

Answer: 1.75

Solution:

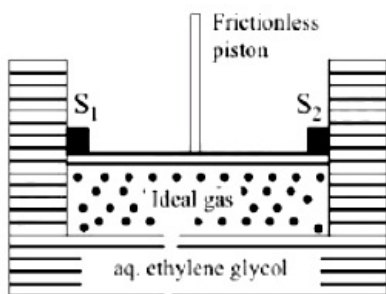
$$\Delta T_f = iK_f m$$

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

$$w = \frac{0.2 \times 58.5 \times 600}{1000 \times 4} = \frac{1.2 \times 58.5}{40} = 1.75\text{g}$$

## Question95

A cylinder containing an ideal gas ( $0.1\text{mol}$  of  $1.0\text{d m}^3$ ) is in thermal equilibrium with a large volume of  $0.5\text{mol}$  aq. ethylene glycol at its freezing point. If the stoppers  $S_1$  and  $S_2$  (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) =  $2.0\text{K kgmol}^{-1}$ ,  $R = 0.08\text{d m}^3\text{aamK}^{-1}\text{mol}^{-1}$ )

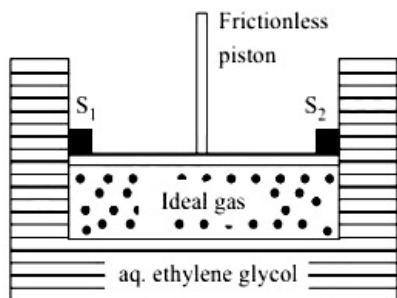


[NV, Jan.09,2020 (II)]

Answer: 2.18

Solution:





$$K_f = 2.0 \text{ K kg mol}^{-1}; m = 0.5$$

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

$$T_{\text{initial}} = 272 \text{ K}$$

$$n = 0.1 \text{ mol } V = 1 \text{ d m}^3$$

$$P_{\text{gs}} = \frac{nRT}{V} = \frac{0.1 \times 0.08 \times 272}{1} = 2.176 \text{ atm}$$

After releasing piston

$$P_1 V_1 = P_2 V_2$$

$$2.176 \times 1 = 1 \times V_2$$

$$V_2 = 2.18 \text{ d m}^3$$

## Question 96

A set of solutions is prepared using 180g of water as a solvent and 10g 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 gmol<sup>-1</sup>; B = 200 gmol<sup>-1</sup>; C = 10,000 gmol<sup>-1</sup>]

[Sep. 06, 2020(II)]

**Options:**

A. B > C > A

B. C > B > A

C. A > B > C

D. A > C > B

**Answer: C**

**Solution:**

**Solution:**

Relative lowering in vapour pressure (RLVP)

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

n → moles of solute

N → moles of solvent

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

$$\text{Moles of solvent (H}_2\text{O)} = \frac{180}{18} = 10 \text{ mol}$$

$$\text{RLV } P_A = \frac{0.1}{10.1} = \frac{1}{101}$$

$$\text{RLV } P_B = \frac{0.05}{10.05} = \frac{1}{201}$$

$$\text{RLV } P_C = \frac{10^{-3}}{10} = 10^{-4}$$

## Question97

At 300K, the vapour pressure of a solution containing 1 mole of n - hexane and 3 moles of n -heptane is 550mm 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 10mm of H g. What is the vapour pressure in mmH g of n -heptane in its pure state \_\_\_\_\_ ?  
[NV, Sep. 04, 2020(I)]

**Answer: 600**

**Solution:**

$$P_{\text{total}} = P_{\text{hexane}}^{\circ} \cdot X_{\text{hexane}} + P_{\text{heptane}}^{\circ} \cdot X_{\text{heptane}}$$

$$550 = [P_{\text{hexane}}^{\circ}] \times \frac{1}{4} + [P_{\text{heptane}}^{\circ}] \times \frac{3}{4}$$

$$\Rightarrow 2200 = P_{\text{hexane}}^{\circ} + 3P_{\text{heptane}}^{\circ} \dots\dots(i)$$

$$560 = [P_{\text{hexane}}^{\circ}] \times \frac{1}{5} + [P_{\text{heptane}}^{\circ}] \times \frac{4}{5}$$

$$\Rightarrow 2800 = P_{\text{hexane}}^{\circ} + 4P_{\text{heptane}}^{\circ} \dots\dots(ii)$$

From equation (i) and (ii),

$$P_{\text{heptane}}^{\circ} = 600\text{mm of H g}$$

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## Question98

Henry's constant (in k bar) for four gases  $\alpha$ ,  $\beta$ ,  $\gamma$  and  $\delta$  in water at 298K is given below:

	$\alpha$	$\beta$	$\gamma$	$\delta$
$K_H$	50	2	$2 \times 10^{-5}$	0.5

(density of water =  $10^3 \text{kgm}^{-3}$  at 298K )

This table implies that:

[Sep. 03, 2020(I)]

**Options:**

- A.  $\alpha$  has the highest solubility in water at a given pressure
- B. solubility of  $\gamma$  at 308K is lower than at 298K
- C. The pressure of a 55.5 molal solution of  $\gamma$  is 1 bar
- D. The pressure of 55.5 molal solution of  $\delta$  is 250 bar



**Answer: D**

**Solution:**

(a) From Henry's law  $p = K_H(x)$

Higher the value of  $K_H$  smaller will be the solubility of the gas, so  $\gamma$  is more soluble.

(b) Though solubility of gases will decrease with increase in temperature but this conclusion can not be drawn from the given table.

(c) For  $\gamma$

$$(p)_\gamma = (K_H)_\gamma \cdot (x)_\gamma$$

$$= 2 \times 10^{-5} \left[ \frac{55.5}{55.5 + \frac{1000}{18}} \right] = 10^{-5} \text{ k bar} = 10^{-2} \text{ bar}$$

(d) For  $\delta \Rightarrow$

$$(p)_\delta = (K_H)_\delta \cdot (x)_\delta$$

$$= 0.5 \left[ \frac{55.5}{55.5 + \frac{1000}{18}} \right]$$

$$= 0.5 \times 0.5 = 0.25 \text{ k bar} = 250 \text{ bar.}$$

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## Question99

**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: [Sep. 02,2020]**

**Options:**

- A. leaves the vapour increases
- B. leaves the solution increases
- C. leaves the solution decreases
- D. leaves the vapour decreases

**Answer: A**

**Solution:**

**Solution:**

The vapour pressure of solution will be less than the vapour pressure of pure solvent, so some vapour molecules will get condensed to maintain new equilibrium.

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## Question100

**The elevation of boiling point of 0.10m aqueous  $\text{CrCl}_3, x\text{N H}_3$  solution is two times that of 0.05m aqueous  $\text{CaCl}_2$  solution. The value of x is \_\_\_\_\_**

**[Assume 100% ionisation of the complex and  $\text{CaCl}_2$ , coordination**

number of Cr as 6, and that all  $\text{NH}_3$  molecules are present inside the coordination sphere]  
 [NV, Sep. 06, 2020 (I)]

**Answer: 5**

**Solution:**

Molality of  $\text{CaCl}_2$  solution = 0.05m

$$\Delta T_b = iK_b m = 3 \times K_b \times 0.05 = 0.15K_b$$

Molality of  $\text{CrCl}_3 \cdot x\text{NH}_3$  = 0.10m

$$\Delta T_b' = iK_b \times 0.10 \Rightarrow \Delta T_b' = 2 \Delta T_b$$

$$iK_b \times 0.10 = 2 \times 0.15K_b \Rightarrow i = 3$$

Since, co-ordination number of Cr is 6 .

$\therefore$  The complex is  $[\text{Cr}(\text{NH}_3)_5\text{Cl}]\text{Cl}_2$ .

$\therefore x = 5$ .

## Question101

The osmotic pressure of a solution of  $\text{NaCl}$  is 0.10atm and that of a glucose solution is 0.20atm. The osmotic pressure of a solution formed by mixing 1 L of the sodium chloride solution with 2L of the glucose solution is  $x \times 10^{-3}$ atm . x is \_\_\_\_\_. (nearest integer)  
 [NV, Sep.04, 2020 (II)]

**Answer: 167**

**Solution:**

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

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

When 1L of  $\text{NaCl}$  solution and 2L glucose solution are mixed.

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

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

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

## Question102

If  $250\text{cm}^3$  of an aqueous solution containing 0.73g of a protein A is



isotonic with one litre of another aqueous solution containing 1.65g of a protein B, at 298K, the ratio of the molecular masses of A and B is  $\underline{\hspace{2cm}} \times 10^{-2}$  (to the nearest integer).  
 [NV, Sep. 03, 2020 (II)]

**Answer: 177**

**Solution:**

$$\pi_A = iC_A RT, \pi_B = iC_B RT$$

For isotonic solution,

$$\pi_A = \pi_B \quad i_1 C_1 = i_2 C_2 \quad (\text{For protein } i = 1)$$

$$\Rightarrow C_1 = C_2$$

$$\Rightarrow \frac{0.73 \times 1000}{M_A \times 250} = \frac{1.65}{M_B \times 1}$$

$$\therefore \frac{M_A}{M_B} = \frac{0.73 \times 4}{1.65} = 1.77 = 177 \times 10^{-2}$$

## Question103

A solution of sodium sulfate contains 92g of  $\text{Na}^+$  ions per kilogram of water. The molality of  $\text{Na}^+$  ions in that solution in  $\text{mol kg}^{-1}$  is:  
 [Jan. 9, 2019(I)]

**Options:**

- A. 12
- B. 4
- C. 8
- D. 16

**Answer: B**

**Solution:**

$$\text{Number of moles in 92g of } \text{Na}^+ = \frac{92}{23} = 4\text{mol}$$

$$\text{Molality (m)} = \frac{\text{Number of moles}}{\text{Mass of solvent (in kg)}}$$

$$\therefore m = \frac{4}{1} = 4\text{mol kg}^{-1}$$

## Question104

Liquids A and B form an ideal solution in the entire composition range. At 350K, the vapor pressures of pure A and pure B are  $7 \times 10^3 \text{ Pa}$  and  $12 \times 10^3 \text{ Pa}$ , respectively. The composition of the vapour is in equilibrium with a solution containing 40 mole percent of A at this temperature is:  
[Jan. 10,2019(I)]

Options:

A.  $x_A = 0.37$ ;  $x_B = 0.63$

B.  $x_A = 0.28$ ;  $x_B = 0.72$

C.  $x_A = 0.4$ ;  $x_B = 0.6$

D.  $x_A = 0.76$ ;  $x_B = 0.24$

Answer: B

Solution:

$$P_A^\circ = 7 \times 10^3$$

$$P_B^\circ = 12 \times 10^3$$

$$x_A' = 0.4; x_B' = 1 - 0.4$$

$$x_B' = 0.6$$

$$P_{\text{total}} = P_A^\circ x_A' + P_B^\circ x_B'$$

$$= 7 \times 10^3 \times 0.4 + 12 \times 10^3 \times 0.6$$

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

$$\therefore x_A = \frac{P_A^\circ x_A'}{P_{\text{total}}} = \frac{7 \times 10^3 \times 0.4}{10^4}$$

$$\therefore x_A = 0.28, x_B = 1 - 0.28 = 0.72$$

## Question105

Which one of the following statements regarding Henry's law is not correct?

[Jan. 9,2019(I)]

Options:

A. Higher the value of  $K_H$  at a given pressure, higher is the solubility of the gas in liquids.

B. Different gases have different  $K_H$  (Henry's law constant) values at the same temperature.

C. The partial pressure of the gas in vapour phase is proportional to the mole fraction of the gas in the solution.

D. The value of  $K_H$  increases with increase of temperature and  $K_H$  is function of the nature of the gas.

Answer: A



## Solution:

### Solution:

The solubility of the gas in liquids decreases with the increase in value of  $K_H$  at a given pressure.

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## Question106

Freezing point of a 4% aqueous 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:

[Jan. 12,2019 (I)]

### Options:

A. 3A

B. 2A

C. A

D. 4A

**Answer: A**

### Solution:

$$(\Delta T_f)_x = (\Delta T_f)_y$$

$$K_f m_x = K_f m_y$$

$$m_x = m_y$$

$$\frac{\text{No. of moles of x}}{\text{wt. of solvent in kg}} = \frac{\text{No. of moles of y}}{\text{wt. of solvent in kg}}$$

Given, 4% aqueous solution of x = 4g of solute, x present in 100g of  $H_2O$

12% aqueous solution of y = 12g of solute, y present in 100g of  $H_2O$

$$\text{Now, } \frac{4g}{m_1} \times \frac{1000}{100} = \frac{12g}{m_2} \times \frac{1000}{100}$$

$$\frac{4g}{A} = \frac{12g}{m_2} \quad [m_1 = A]$$

$$m_2 = 3A$$

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## Question107

Molecules of benzoic acid ( $C_6H_5COOH$ ) dimerise in benzene. 'w' g of the acid dissolved in 30g of benzene shows a depression in freezing point equal to 2K . If the percentage association of the acid to form dimer in the solution is 80 , then w is :

(Given that  $K_f = 5K \text{ kgmol}^{-1}$ , Molar mass of benzoic acid =  $122\text{gmol}^{-1}$ )

[Jan. 12, 2019 (II)]

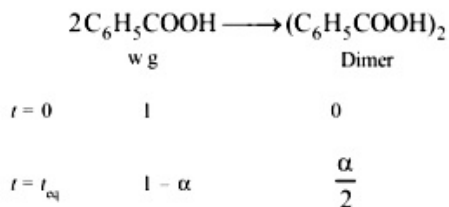
### Options:



- A. 2.4g
- B. 1.0g
- C. 1.5g
- D. 1.8g

**Answer: A**

**Solution:**



Moles at equilibrium =  $1 - \alpha + \frac{\alpha}{2} = 1 - \frac{\alpha}{2}$

$i = \frac{\text{Total moles at equilibrium}}{\text{Initial moles}} = \frac{1 - \frac{\alpha}{2}}{1}$

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

$\Delta T_f = iK_f m$

$2 = 0.6 \times 5 \times \left( \frac{w}{\frac{122}{30}} \right) \times 1000$

$w = 2.4\text{g}$

## Question 108

The freezing point of a diluted milk sample is found to be  $-0.2^\circ\text{C}$ , while it should have been  $-0.5^\circ\text{C}$  for pure milk. How much water has been added to pure milk to make the diluted sample?

[Jan. 11, 2019(I)]

**Options:**

- A. 1 cup of water to 2 cups of pure milk
- B. 3 cups of water to 2 cups of pure milk
- C. 1 cup of water to 3 cups of pure milk
- D. 2 cups of water to 3 cups of pure milk

**Answer: B**

**Solution:**

Freezing point of diluted milk =  $-0.2^\circ\text{C}$

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

Freezing point of pure milk =  $-0.5^\circ\text{C}$



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

$$\frac{\Delta T_f}{\Delta T_f'} = \frac{K_f \times m}{K_f \times m'}; m = \frac{\text{mole of solute}}{\text{mass of solvent (kg)}}$$

Moles of solute are same in both samples.

$$\therefore \frac{0.5}{0.2} = \frac{W'}{W}$$

$$\frac{W'}{W} = \frac{5}{2}; W' = \frac{5}{2}W$$

2 cups of pure milk is mixed with 3 cups of water to make 5 cups of diluted milk.

---

## Question 109

**$\text{K}_2\text{HgI}_4$  is 40% ionised in aqueous solution. The value of its van't Hoff factor (i) is :**

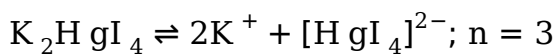
**[Jan. 11, 2019 (II)]**

**Options:**

- A. 1.6
- B. 1.8
- C. 2.0
- D. 2.2

**Answer: B**

**Solution:**



$$\therefore \alpha = \frac{i-1}{n-1}; 0.4 = \frac{i-1}{3-1}$$

$$i = 1.8$$

---

## Question 110

**Elevation in the boiling point for 1 molal solution of glucose is  $2K$ . The depression in the freezing point for 2 molal solution of glucose in the same solvent is  $2K$ . The relation between  $K_b$  and  $K_f$  is:**

**[Jan. 10, 2019 (II)]**

**Options:**

- A.  $K_b = 1.5K_f$
- B.  $K_b = K_f$
- C.  $K_b = 0.5K_f$
- D.  $K_b = 2K_f$



**Answer: D**

**Solution:**

According to the question we can write

$$\Delta T_b = K_b m \Rightarrow K_b(1) = 2 \Rightarrow K_b = 2 \text{K m}^{-1}$$

$$\Delta T_f = K_f m \Rightarrow K_f(2) = 2 \Rightarrow K_f = 1 \text{K m}^{-1}$$

$$\text{So, } \frac{K_b}{K_f} = \frac{2}{1}$$

$$\therefore K_b = 2K_f$$

---

## Question111

**A solution containing 62g ethylene glycol in 250g water is cooled to  $-10^\circ\text{C}$ . If  $K_f$  for water is  $1.86 \text{K kg mol}^{-1}$ , the amount of water (in g) separated as ice is:**

**[Jan. 9, 2019 (II)]**

**Options:**

A. 48

B. 32

C. 64

D. 16

**Answer: C**

**Solution:**

As we know,

$$\Delta T_f = K_f \cdot m$$

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

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

Total amount of water = 250g

$\therefore$  The amount of water separated as ice  
= 250 - 186 = 64g

---

## Question112

**The mole fraction of a solvent in aqueous solution of a solute is 0.8. The molality (in  $\text{mol kg}^{-1}$ ) of the aqueous solution is :**

**[April 12, 2019 (I)]**

**Options:**

A.  $13.88 \times 10^{-2}$





B.  $13.88 \times 10^{-1}$

C. 13.88

D.  $13.88 \times 10^{-3}$

**Answer: C**

**Solution:**

Let, 1 mole be present in the solution,

$$n_{\text{solute}} = 0.2\text{mol}$$

$$n_{\text{solvent}} = 0.8\text{mol}$$

$$= 0.8 \times 18\text{g molality}$$

$$= \frac{\text{Moles of solute}}{\text{Kilogram of solvent}}$$

$$\therefore m = \frac{0.2 \times 1000}{0.8 \times 18} = \frac{1000}{4 \times 18} \approx 13.88$$

---

## Question113

**What would be the molality of 20% (mass/mass) aqueous solution of KI ? (molar mass of KI =  $166\text{gmol}^{-1}$ )**

**[April9, 2019 (II)]**

**Options:**

A. 1.08

B. 1.35

C. 1.48

D. 1.51

**Answer: D**

**Solution:**

20% W/W KI solution (Given)

i.e. 100g solution contains 20gKI

$$\therefore \text{Mass of solvent} = 100 - 20 = 80\text{g}$$

$$\text{No. of moles of KI is } \frac{20}{166} = 0.121\text{mol}$$

$$\therefore \text{Molality} = \frac{0.121 \times 1000}{80} = 1.51\text{m}$$

---

## Question114

**1g of a non-volatile non-electrolyte solute is dissolved in 100g 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,  $\frac{\Delta T_b(A)}{\Delta T_b(B)}$ , is:

[April 10, 2019 (II)]

Options:

- A. 5 : 1
- B. 10 : 1
- C. 1 : 5
- D. 1 : 0.2

Answer: C

Solution:

Ebullioscopic constant (molal deviation const.) is given by,

$$k_b = \frac{\Delta T_b}{m} \Rightarrow \frac{(k_b)_A}{(k_b)_B} = \frac{1}{5}$$

$$\therefore \frac{(\Delta T_b)_A}{(\Delta T_b)_B} = \frac{(k_b)_A}{(k_b)_B} = \frac{1}{5}$$

---

## Question 115

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)

[April 9, 2019 (I)]

Options:

A.  $\frac{x_M}{x_N} = \frac{y_M}{y_N}$

B.  $(x_M - y_M) < (x_N - y_N)$

C.  $\frac{x_M}{x_N} < \frac{y_M}{y_N}$

D.  $\frac{x_M}{x_N} > \frac{y_M}{y_N}$

Answer: D

Solution:



$$P_M^0 = 450\text{mmHg}, P_N^0 = 700\text{mmHg}$$

$$P_M = P_M^0 x_M = y_M P_T$$

$$\Rightarrow P_M^0 = \frac{y_M}{x_M} (P_T) \text{ Similarly,}$$

$$P_N^0 = \frac{y_N}{x_N} (P_T)$$

$$\text{Given, } P_M^0 < P_N^0$$

$$\Rightarrow \frac{y_M}{x_M} < \frac{y_N}{x_N}$$

$$\Rightarrow \frac{y_M}{y_N} < \frac{x_M}{x_N}$$

## Question116

The vapour pressures of pure liquids A and B are 400 and 600mmHg, respectively at 298K. 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:

[April 8, 2019(I)]

Options:

A. 450mmHg, 0.4, 0.6

B. 500mmHg, 0.5, 0.5

C. 450mmHg, 0.5, 0.5

D. 500mmHg, 0.4, 0.6

Answer: D

Solution:

Solution:

$$P = x_B P_B^0 + x_A P_A^0$$

$$= 0.5 \times 600 + 0.5 \times 400 = 300 + 200 = 500$$

Using the relation  $p_i = y_i P_{\text{Total}}$ , we can calculate the mole fractions of the components in vapour phase.

$$P_B = y_B P_{\text{total}}$$

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

$$y_A = \frac{P_A}{P_{\text{Total}}} = \frac{200}{500} = \frac{2}{5} = 0.4$$

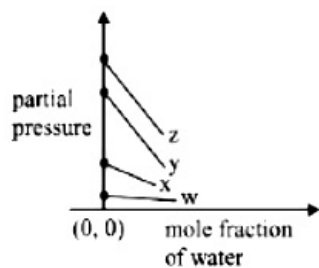
## Question117

For the solution of the gases w, x, y and z in water at 298K. the Henry's law constants ( $K_H$ ) are 0.5, 2, 35 and 40kbar. respectively. The correct plot for the given data is:

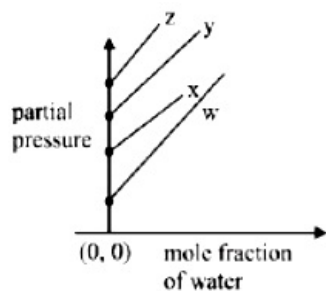
[April 8, 2019 (II)]

Options:

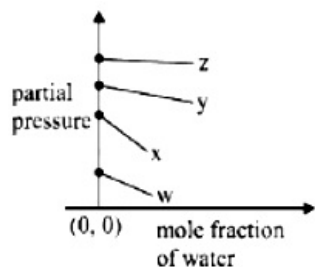
A.



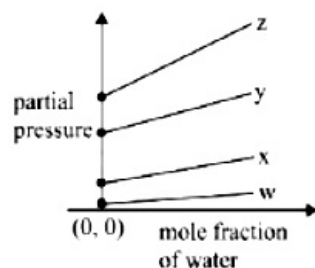
B.



C.



D.



Answer: A

Solution:

According to Henry's law

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

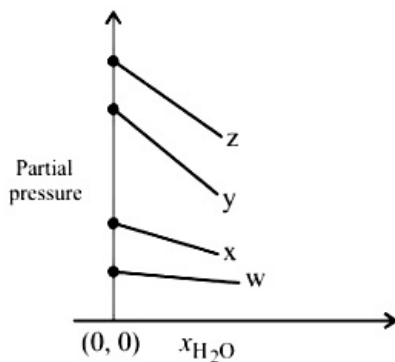
$$\because X_{\text{gas}} + X_{\text{H}_2\text{O}} = 1$$

$$\therefore X_{\text{gas}} = 1 - X_{\text{H}_2\text{O}}$$

$$\therefore P = K_H (1 - X_{\text{H}_2\text{O}})$$

$$\therefore P = K_H - K_H \cdot X_{\text{H}_2\text{O}}$$

$$y = c + mx; m = -K_H \Rightarrow \text{slope is negative.}$$



Gas	$K_H$
w	0.5
x	2
y	35
z	50

## Question 118

A solution is prepared by dissolving 0.6g of urea (molar mass =  $60 \text{ gmol}^{-1}$ ) and 1.8g of glucose (molar mass =  $180 \text{ gmol}^{-1}$ ) in 100mL, of water at  $27^\circ\text{C}$ . The osmotic pressure of the solution is:

( $R = 0.08206 \text{ LatmK}^{-1} \text{ mol}^{-1}$ )

[April 12, 2019 (II)]

Options:

- A. 8.2atm
- B. 2.46atm
- C. 4.92atm
- D. 1.64atm

**Answer: C**

**Solution:**

**Solution:**

Osmotic pressure ( $\pi$ ) = CRT

Since, there are two solutes i.e. urea and glucose.

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

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

$$= 0.2 \times 0.0821 \times 300 = 4.926 \text{ atm}$$

## Question 119



**At room temperature, a dilute solution of urea is prepared by dissolving 0.60g of urea in 360g of water. If the vapour pressure of pure water at this temperature is 35mmHg, lowering of vapour pressure will be : (molar mass of urea =  $60\text{g mol}^{-1}$ ) [April 10, 2019 (I)]**

**Options:**

- A. 0.027mmHg
- B. 0.028mmHg
- C. 0.017mmHg
- D. 0.031mmHg

**Answer: C**

**Solution:**

**Solution:**

Relative lowering of vapour pressure, is given by,

$$\frac{p^\circ - p}{p^\circ} = x_A = \frac{n_A}{n_A + n_B} = \frac{n_A}{n_B}$$

Given,  $p^\circ = 35\text{mmHg}$ ,  $n_{\text{urea}} = \frac{0.60}{60}$ ,  $n_{\text{water}} = \frac{360}{18}$

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

$$\Delta p = p^\circ - p = 0.017$$

---

## Question120

**The osmotic pressure of a dilute solution of an ionic compound XY in water is four times that of a solution of 0.01M  $\text{BaCl}_2$  in water. Assuming complete dissociation of the given ionic compounds in water, the concentration of XY (in  $\text{mol L}^{-1}$ ) in solution is: [April 9, 2019 (I)]**

**Options:**

- A.  $4 \times 10^{-2}$
- B.  $6 \times 10^{-2}$
- C.  $4 \times 10^{-4}$
- D.  $16 \times 10^{-4}$

**Answer: B**

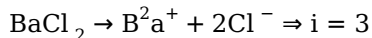
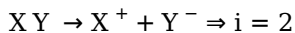
**Solution:**

**Solution:**

We know,  $\Pi = iCRT$ ;  $\Pi_{xy} = 4\Pi_{\text{BaCl}_2}$



Since both are ionic solute, i.e.,



Now,

$$2[XY] = 4 \times 3 \times [0.01]$$

$$[XY] = 0.06$$

$$= 6 \times 10^{-2} \text{ mol / L}$$

---

## Question 121

**Molar depression constant for a solvent is  $4.0 \text{ K kg mol}^{-1}$ . The depression in the freezing point of the solvent for  $0.03 \text{ mol kg}^{-1}$  solution  $K_2SO_4$  is:**

**(Assume complete dissociation of the electrolyte)**

**[April 9, 2019 (II)]**

**Options:**

A. 0.18K

B. 0.24K

C. 0.12K

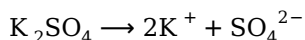
D. 0.36K

**Answer: D**

**Solution:**

**Solution:**

Dissociation of Potassium Sulphate ( $K_2SO_4$ ),



$i$  (Van't Hoff factor) = 3

We know that,  $\Delta T_f = iK_f m$

where,  $K_f$  is molal depression constant and  $m$  is molality.

$$= 3 \times 4 \times 0.03 = 0.36 \text{ K}$$

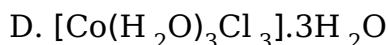
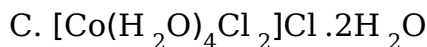
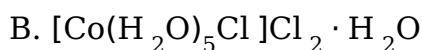
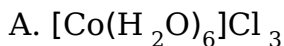
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## Question 122

**For 1 molal aqueous solution of the following compounds, which one will show the highest freezing point?**

**[2018]**

**Options:**



**Answer: D**

**Solution:**

**Solution:**

	Number of particles (i)
(a) $[Co(H_2O)_6]Cl_3$	4
(b) $[Co(H_2O)_5Cl]Cl_2 \cdot H_2O$	3
(c) $[Co(H_2O)_4Cl_2]Cl \cdot 2H_2O$	2
(d) $[Co(H_2O)_3Cl_3] \cdot 3H_2O$	1

z; where  $\Delta T_f = (T_f - T_f')$

Remember, greater the no. of particles, lower will be the freezing point. Compound (d) will have the highest freezing point due to least number of particles.

## Question123

**The mass of non-volatile, non-electrolyte solute (molar mass =  $50\text{g mol}^{-1}$ ) needed to be dissolved in 114g octane to reduce its vapour pressure to 75%, is [Online April 16,2018]**

**Options:**

- A. 37.5g
- B. 75g
- C. 150g
- D. 50g

**Answer: C**

**Solution:**

**Solution:**

Molar mass of octane = 114g / mol

From the lowering of vapour pressure we have,

$$\frac{\Delta P}{P} = \frac{\frac{W_2}{M_2}}{\frac{W_2}{M_2} + \frac{W_1}{M_1}}$$
$$\frac{75}{100} = \frac{\frac{W_2}{50\text{g/mol}}}{\frac{W_2}{50\text{g/mol}} + \frac{114\text{g}}{114\text{g/mol}}}$$
$$0.75 = \frac{\frac{W_2}{50}}{\frac{W_2}{50} + 1}$$





$$\frac{W_2}{50} + 1 = \frac{W_2}{50 \times 0.75}$$

$$W_2 = 150\text{g}$$

**Note :**  $W_2$  and  $M_2$  are mass and molar mass of solute whereas  $W_1$  and  $M_1$  are mass and molar mass of octane.

---

## Question124

Two 5 molal solutions are prepared by dissolving a non-electrolyte, non-volatile solute separately in the solvents X and Y. The molecular weights of the solvents are  $M_X$  and  $M_Y$ , respectively where  $M_X = \frac{3}{4}M_Y$ . The relative lowering of vapour pressure of the solution in X is "m" times that of the solution in Y. Given that the number of moles of solute is very small in comparison to that of solvent, the value of "m" is:  
[Online April 15, 2018 (II)]

Options:

A.  $\frac{3}{4}$

B.  $\frac{1}{2}$

C.  $\frac{1}{4}$

D.  $\frac{4}{3}$

**Answer: A**

**Solution:**

**Solution:**

The relationship between molar masses of the two solvents is

$$M_X = \frac{3}{4}M_Y \dots (i)$$

The relative lowering of vapour pressure of the two solutions is

$$\left(\frac{\Delta P}{P}\right)_X = m \left(\frac{\Delta P}{P}\right)_Y$$

But, the relative lowering of vapour pressure of solutions is directly proportional to the mole fraction of solute. Given 5 molal solution, means 5 moles of solute are dissolved in 1kg (or 1000g) of solvent.

$$\text{The number of moles of solvent} = \frac{1000\text{g}}{M}$$

$$\text{The mole fraction of solute} = \frac{5}{1000 / M}$$

$$= M \times \frac{5}{1000}$$

$$\text{Hence, } M_X \times \frac{5}{1000} = m \times M_Y \times \frac{5}{1000} \dots (ii)$$

Substitute equation (i) in equation (ii)

$$\frac{3}{4} \times M_Y \times \frac{5}{1000} = m \times M_Y \times \frac{5}{1000}$$

$$m = \frac{3}{4}$$

---

## Question125



**A solution is prepared by mixing 8.5g of  $\text{CH}_2\text{Cl}_2$  and 11.95 g of  $\text{CHCl}_3$ . If vapour pressure of  $\text{CH}_2\text{Cl}_2$  and  $\text{CHCl}_3$  at 298 K are 415 and 200mmHg respectively, the mole fraction of  $\text{CHCl}_3$  in vapour form is: (Molar mass of Cl =  $35.5\text{g mol}^{-1}$ ) [Online April 9, 2017]**

**Options:**

- A. 0.162
- B. 0.675
- C. 0.325
- D. 0.486

**Answer: C**

**Solution:**

**Solution:**

Molar mass of  $\text{CHCl}_3 = 119.5\text{g / mol}$

Molar mass of  $\text{CH}_2\text{Cl}_2 = 85\text{g / mol}$

Moles of  $\text{CHCl}_3 = \frac{11.95}{119.5} = 0.1\text{mol}$

Moles of  $\text{CH}_2\text{Cl}_2 = \frac{8.5}{85} = 0.1\text{mol}$

Mole fraction of  $\text{CHCl}_3 = \frac{0.1}{0.2} = 0.5\text{mol}$

Mole fraction of  $\text{CH}_2\text{Cl}_2 = \frac{0.1}{0.2} = 0.5\text{mol}$

(Given -

Vapour pressure of  $\text{CHCl}_3 = 200\text{mmHg} = 0.263\text{atm}$ .

Vapour pressure of  $\text{CH}_2\text{Cl}_2 = 415\text{mmHg} = 0.546\text{atm}$ .) (1atm = 760mmHg)

$\therefore P_{\text{total}} = \text{Mole fraction of } \text{CHCl}_3 \times$

(Vapour pressure of  $\text{CHCl}_3$ ) + Mole fraction of  $\text{CH}_2\text{Cl}_2$

$\times$  ( Vapour pressure of  $\text{CH}_2\text{Cl}_2$  )

$= 0.5 \times 0.263 + 0.5 \times 0.546 = 0.4045$

Mole fraction of  $\text{CHCl}_3$  in vapour form

$= \frac{\text{vapour pressure}}{P_{\text{total}}} = \frac{0.1315}{0.4045} = 0.325$ .

## Question126

**The freezing point of benzene decreases by  $0.45^\circ\text{C}$  when 0.2g of acetic acid is added to 20g 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 =  $5.12\text{K kg mol}^{-1}$  )**

**[2017]**

**Options:**

- A. 64.6%

B. 80.4%

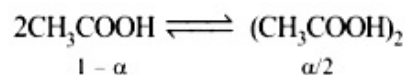
C. 74.6%

D. 94.6%

**Answer: D**

**Solution:**

In benzene,



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

Here  $\alpha$  is degree of association

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

$$0.45 = \left(1 - \frac{\alpha}{2}\right) (5.12) \frac{\left(\frac{0.2}{60}\right)}{\frac{20}{1000}}$$

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

$$\alpha = 0.945$$

$$\% \text{ degree of association} = 94.6\%$$

---

## Question127

5g of  $\text{Na}_2\text{SO}_4$  was dissolved in xg of  $\text{H}_2\text{O}$ . The change in freezing point was found to be  $3.82^\circ\text{C}$ . If  $\text{Na}_2\text{SO}_4$  is 81.5% ionised, the value of x

( $K_f$  for water =  $1.86^\circ\text{Ckgmol}^{-1}$ ) is approximately:

(molar mass of S =  $32\text{gmol}^{-1}$  and that of Na =  $23\text{gmol}^{-1}$ )

[Online April 8, 2017]

**Options:**

A. 15g

B. 25g

C. 45g

D. 65g

**Answer: C**

**Solution:**

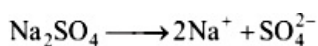
Molality (experimental)

$$= \frac{\Delta T_f}{K_f} = \frac{3.82}{1.86} = 2.054 \text{ mol} / 1000 \text{ g solvent}$$

$$\text{Molality (theoretical)} = \frac{\text{moles of solute}}{\text{wt. of solvent(g)}} \times 1000$$

$$= \frac{5 \text{ g} / 142 \text{ g/mol}}{x} \times 1000$$





Moles before dissociation	1	0	0
Moles after dissociation	$1 - \alpha$	$2\alpha$	$\alpha$

$$\text{Von't Hoff factor (i)} = \frac{\text{Moles after dissociation}}{\text{Moles before dissociation}}$$

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

$\text{Na}_2\text{SO}_4$  is ionised 81.5%

means  $\alpha = 0.815$

$$= \frac{(1 - 0.815) + 2 \times 0.815 + 0.815}{1}$$

$$= 2.63.$$

$$i = \frac{\text{Observed molality}}{\text{Calculated molality}}$$

$$2.63 = \frac{2.054}{\frac{0.0352}{x} \times 1000}$$

$$x = 45.07\text{g}$$

## Question128

The solubility of  $\text{N}_2$  in water at 300K and 500 torr partial pressure is  $0.01\text{gL}^{-1}$ . The solubility (in  $\text{gL}^{-1}$ ) at 750 torr partial pressure is: [Online April9, 2016]

Options:

- A. 0.0075
- B. 0.005
- C. 0.02
- D. 0.015

Answer: D

Solution:

Solution:

According to Henry's law

$$\frac{P_1}{P_2} = \frac{S_1}{S_2}$$

$$\frac{500}{750} = \frac{0.01}{S_2}$$

$$\therefore S_2 = \frac{750 \times 0.01}{500} = 0.015\text{g / L}$$

## Question129

An aqueous solution of a salt  $\text{M X}_2$  at certain temperature has a van't Hoff factor of 2 . The degree of dissociation for this solution of the salt is:



**[Online April 10,2016]**

**Options:**

- A. 0.50
- B. 0.33
- C. 0.67
- D. 0.80

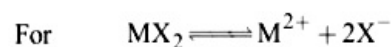
**Answer: A**

**Solution:**

**Solution:**

Van't Hoff factor (i)

$$= \frac{\text{No. of particles after association or dissociation}}{\text{No. of particles before association or dissociation}}$$



$t = 0$	1	0	0
at eq.	$1 - \alpha$	$\alpha$	$2\alpha$

$$\text{Total no. of particles} = 1 - \alpha + \alpha + 2\alpha = 1 + 2\alpha$$

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

$$\therefore \alpha = 0.50 \text{ or } 50\%$$

---

## Question130

**18 g glucose ( $\text{C}_6\text{H}_{12}\text{O}_6$ ) is added to 178.2g water. The vapour pressure of water (in torr) for this aqueous solution is: [2016, 2006]**

**Options:**

- A. 752.4
- B. 759.0
- C. 7.6
- D. 76.0

**Answer: A**

**Solution:**

**Solution:**

According to Raoult's Law

$$\frac{P^\circ - P_s}{P_s^\circ} = \frac{W_B \times M_A}{M_B \times W_A} \dots (i)$$

Here  $P^\circ$  = Vapour pressure of pure solvent,

$P_s$  = Vapour pressure of solution

$W_B$  = Mass of solute,  $W_A$  = Mass of solvent

$M_B$  = Molar mass of solute,  $M_A$  = Molar Mass of solvent



Vapour pressure of pure water at 100°C (by assumption = 760 torr )

By substituting values in equation (i) we get,

$$\frac{760 - P_s}{P_s} = \frac{18 \times 18}{180 \times 178.2} \dots (ii)$$

On solving (ii) we get

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

---

## Question 131

**A solution at 20°C is composed of 1.5mol of benzene and 3.5mol of toluene. If the vapour pressure of pure benzene and pure toluene at this temperature are 74.7 torr and 22.3 torr, respectively, then the total vapour pressure of the solution and the benzene mole fraction in equilibrium with it will be, respectively:**

**[Online April 10,2015]**

**Options:**

- A. 35.8 torr and 0.280
- B. 38.0 torr and 0.589
- C. 30.5 torr and 0.389
- D. 30.5 torr and 0.480

**Answer: B**

**Solution:**

**Solution:**

$$\text{Total V.P. of solution} = P_A^\circ X_A + P_B^\circ X_B$$

$$\text{Given, } P_A^\circ = 74.7 \text{ torr, } P_B^\circ = 22.3 \text{ torr}$$

$$n_{\text{benzene}} = 1.5 \text{ mol, } n_{\text{toluene}} = 3.5 \text{ mol}$$

$$n_{\text{solution}} = 1.5 + 3.5 = 5 \text{ mol}$$

$$x_A = \frac{n_{\text{benzene}}}{n_{\text{solution}}} = \frac{1.5}{5} = 0.3$$

$$x_B = \frac{n_{\text{toluene}}}{n_{\text{solution}}} = \frac{3.5}{5} = 0.7$$

$$\text{Total V.P. of solution} = (0.3 \times 74.7 + 0.7 \times 22.3) \text{ torr}$$

$$= (22.4 + 15.6) \text{ torr} = 38 \text{ torr}$$

$$\text{Mole fraction of benzene in vapour form} = \frac{22.4}{38} = 0.589$$

---

## Question 132

**The vapour pressure of acetone at 20°C is 185 torr. When 1.2g of a non-volatile substance was dissolved in 100g of acetone at 20°C, its vapour pressure was 183 torr. The molar mass ( $\text{gmol}^{-1}$ ) of the substance is :**

**[2015]**

**Options:**

- A. 128
- B. 488
- C. 32
- D. 64

**Answer: D**

**Solution:**

**Solution:**

Using relation,

$$\frac{P^\circ - P_s}{P^\circ} = \frac{w_2 M_1}{w_1 M_2}$$

where  $w_1, M_1$  = mass in g and mol. mass of solvent

$w_2, M_2$  = mass in g and mol. mass of solute

$P^\circ = 185$  torr,  $P_s = 183$  torr

$$\frac{185 - 183}{185} = \frac{1.2 \times 58}{100 \times M_2}$$

(Mol. mass of acetone = 58 )

$M_2 = 64.68 \approx 64$

$\therefore$  Molar mass of substance = 64

## Question 133

**Determination of the molar mass of acetic acid in benzene using freezing point depression is affected by:  
[Online April 11, 2015]**

**Options:**

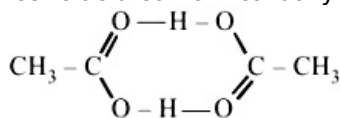
- A. partial ionization
- B. dissociation
- C. complex formation
- D. association

**Answer: D**

**Solution:**

**Solution:**

Acetic acid contain carboxylic group  $-\text{COOH}$  which can form H-bonding so acetic acid dimerises.



## Question 134

**Choose the correct statement with respect to the vapour pressure of a liquid among the following:**  
**[Online April 19, 2014]**

**Options:**

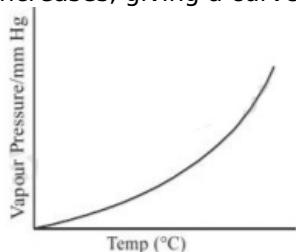
- A. Increases linearly with increasing temperature
- B. Increases non-linearly with increasing temperature
- C. Decreases linearly with increasing temperature
- D. Decreases non-linearly with increasing temperature

**Answer: B**

**Solution:**

**Solution:**

If we plot a graph between the vapour pressure and the temperature. We would get a curve that rises faster as T increases, giving a curved line.



---

## Question135

**For an ideal solution of two components A and B, which of the following is true?**

**[Online April 19,2014]**

**Options:**

- A.  $\Delta H_{\text{mixing}} < 0$  (zero)
- B.  $\Delta H_{\text{mixing}} > 0$  (zero)
- C. A – B interaction is stronger than A – A and B – B interactions
- D. A – A, B – B and A – B interactions are identical.

**Answer: D**

**Solution:**

**Solution:**

Solutions in which solute-solute and solvent-solvent interactions are almost similar to solute solvent interactions are known as ideal solution.

---





## Question 136

Consider separate solutions of  $0.500\text{M C}_2\text{H}_5\text{OH (aq)}$ ,  $0.100\text{M Mg}_3(\text{PO}_4)_2(\text{aq})$ ,  $0.250\text{M KBr (aq)}$  and  $0.125\text{M Na}_3\text{PO}_4(\text{aq})$  at  $25^\circ\text{C}$ . Which statement is true about these solutions, assuming all salts to be strong electrolytes?  
[2014]

Options:

- A. They all have the same osmotic pressure.
- B.  $0.100\text{M Mg}_3(\text{PO}_4)_2(\text{aq})$  has the highest osmotic pressure.
- C.  $0.125\text{M Na}_3\text{PO}_4(\text{aq})$  has the highest osmotic pressure.
- D.  $0.500\text{M C}_2\text{H}_5\text{OH (aq)}$  has the highest osmotic pressure.

Answer: A

Solution:

Solution:

$$\pi = iCRT \quad \pi_{\text{C}_2\text{H}_5\text{OH}} = 1 \times 0.500 \times R \times T = 0.5RT$$

$$\pi_{\text{Mg}_3(\text{PO}_4)_2} = 5 \times 0.100 \times R \times T = 0.5RT$$

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

$$\pi_{\text{Na}_3\text{PO}_4} = 4 \times 0.125 \times RT = 0.5RT$$

Since the osmotic pressure of all the given solutions is equal. Hence all are isotonic solutions.

## Question 137

The observed osmotic pressure for a  $0.10\text{M}$  solution of  $\text{Fe}(\text{NH}_4)_2(\text{SO}_4)_2$  at  $25^\circ\text{C}$  is  $10.8\text{atm}$ . The expected and experimental (observed) values of van't Hoff factor ( $i$ ) will be respectively:  
( $R = 0.082\text{LatmK}^{-1}\text{mol}^{-1}$ )  
[Online April 19, 2014]

Options:

- A. 5 and 4.42
- B. 4 and 4.00
- C. 5 and 3.42
- D. 3 and 5.42

Answer: A

Solution:



Given,  $\pi_{ob} = 10.8\text{atm}$

$$\pi_{nor} = CRT = 0.10 \times 0.0821 \times 298 = 2.446$$

Now experimental value of (i)

$$= \frac{\text{Observed osmotic pressure}}{\text{Normal osmotic pressure}} = \frac{10.8}{2.446} = 4.42$$

---

## Question138

**The molarity of a solution obtained by mixing 750mL of 0.5(M )H Cl with 250mL of 2(M )H Cl will be:  
[2013]**

**Options:**

- A. 0.875M
- B. 1.00M
- C. 1.75M
- D. 0.975M

**Answer: A**

**Solution:**

**Solution:**

From molarity equation :

$$M_1V_1 + M_2V_2 = M \times V$$

$$M = \frac{M_1V_1 + M_2V_2}{V} \text{ where } V = \text{total volume}$$

$$= \frac{750 \times 0.5 + 250 \times 2}{1000}$$

$$= 0.875\text{M}$$

---

## Question139

**Vapour pressure of pure benzene is 119 torr and that of toluene is 37.0 torr at the same temperature. Mole fraction of toluene in vapour phase which is in equilibrium with a solution of benzene and toluene having a mole fraction of toluene 0.50, will be :  
[Online April 23, 2013]**

**Options:**

- A. 0.137
- B. 0.237
- C. 0.435
- D. 0.205

**Answer: B**



## Solution:

### Solution:

$$P_A = P_A^\circ \times X_A = \text{total pressure} \times y_A$$

$$P_B = P_B^\circ \times X_B = \text{total pressure} \times y_B$$

where x and y represents mole fraction in liquid and vapour phase respectively.

$$\frac{P_B^\circ X_B}{P_A^\circ X_A} = \frac{y_B}{y_A}; \frac{P_B^\circ(1 - X_A)}{P_A^\circ X_A} = \frac{1 - y_A}{y_A}$$

$$\text{on putting values } \frac{119(1 - 0.50)}{37 \times 0.50} = \frac{1 - y_A}{y_A}$$

on solving  $y_A = 0.237$

---

## Question140

**12g of a nonvolatile solute dissolved in 108g of water produces the relative lowering of vapour pressure of 0.1. The molecular mass of the solute is :**

**[Online April9, 2013]**

**Options:**

A. 80

B. 60

C. 20

D. 40

**Answer: C**

### Solution:

#### Solution:

$$\frac{P^\circ - P_s}{P^\circ} = \frac{W_B \times M_A}{M_B \times W_A}$$

$$0.1 = \frac{12}{M_B} \times \frac{18}{108}$$

$$m_B = \frac{12 \times 18}{0.1 \times 108} = 20$$

---

## Question141

**A molecule M associates in a given solvent according to the equation  $M \rightleftharpoons (M)_n$ . For a certain concentration of M, the van't Hoff factor was found to be 0.9 and the fraction of associated molecules was 0.2. The value of n is:**

**[Online April 22, 2013]**

**Options:**



- A. 3
- B. 5
- C. 2
- D. 4

**Answer: C**

**Solution:**

**Solution:**

van't Hoff factor (i) and the degree of association are related as below:

$$i = 1 - \alpha \left( 1 - \frac{1}{n} \right)$$

$$0.9 = 1 - 0.2 \left( 1 - \frac{1}{n} \right)$$

On solving,

$$\left( 1 - \frac{1}{n} \right) = \frac{1}{2}$$

$$\frac{1}{n} = 1 - \frac{1}{2} = \frac{1}{2}$$

$$\therefore n = 2$$

## Question142

**How many grams of methyl alcohol should be added to 10 litre tank of water to prevent its freezing at 268K ?(  $K_f$  for water is  $1.86K \text{ kgmol}^{-1}$  )  
[Online April 25, 2013]**

**Options:**

- A. 880.07g
- B. 899.04g
- C. 886.02g
- D. 868.06g

**Answer: D**

**Solution:**

**Solution:**

$$\Delta T_f = K_f m$$

where m = molality

$$273 - 268 = 1.86 \times \frac{w}{M \times V}$$

$$5 = 1.86 \times \frac{w}{32 \times 10}$$

$$w = \frac{5 \times 32 \times 10}{1.86} = 860.2 \approx 868.06g$$



## Question143

The density of a solution prepared by dissolving 120g of urea (mol. mass = 60u ) in 1000g of water is 1.15g / mL. The molarity of this solution is :

[2012]

Options:

- A. 0.50M
- B. 1.78M
- C. 1.02M
- D. 2.05M

Answer: D

Solution:

$$\text{Molarity} = \frac{\text{Moles of solute}}{\text{Volume of solution (l)}}$$
$$\text{Mass of solution} = 1000 + 120 = 1120$$
$$d = \frac{M}{V}; V = \frac{M}{d} = \frac{1120}{1.15} \text{mL}$$
$$= \frac{120 \times 1.15}{60 \times 1120} \times 1000 = 2.05\text{M}$$

---

## Question144

$K_f$  for water is  $1.86\text{K kgmol}^{-1}$ . If your automobile radiator holds 1.0kg of water, how many grams of ethylene glycol ( $\text{C}_2\text{H}_6\text{O}_2$ ) must you add to get the freezing point of the solution lowered to  $-2.8^\circ\text{C}$ ?

[2012]

Options:

- A. 72g
- B. 93g
- C. 39g
- D. 27g

Answer: B

Solution:

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

Given  $\Delta T_f = 2.8$ ,  $K_f = 1.86\text{K kgmol}^{-1}$ ,  $i = 1$   
(ethylene glycol is a non- electrolyte)



Wt. of solvent = 1kg  
Let of wt of solute = x  
Mol. wt of ethylene glycol = 62

$$2.8 = 1 \times 1.86 \times \frac{x}{62 \times 1}$$

$$\text{or } x = \frac{2.8 \times 62}{1.86} = 93\text{g}$$

---

## Question 145

A solution containing 0.85g of  $\text{ZnCl}_2$  in 125.0g of water freezes at  $-0.23^\circ\text{C}$ . The apparent degree of dissociation of the salt is ( $K_f$  for water =  $1.86\text{K kgmol}^{-1}$ , atomic mass:  $\text{Zn} = 65.3$  and  $\text{Cl} = 35.5$ )  
[Online May 12, 2012]

Options:

- A. 1.36%
- B. 73.5%
- C. 7.35%
- D. 2.47%

Answer: B

Solution:

$$\begin{aligned}\text{Mol. wt. (M}_B) &= \frac{K_f \times W_B \times 1000}{\Delta T_f \times W_A} \\ &= \frac{1.86 \times 0.85 \times 1000}{0.23 \times 125} \approx 55\text{g}\end{aligned}$$

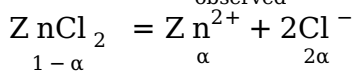
Where,

$$W_B = 0.85\text{g}$$

$$W_A = 125\text{g}$$

$$\Delta T_f = 0^\circ\text{C} - (-23^\circ\text{C}) = 23^\circ\text{C}$$

$$\text{Now, } i = \frac{M_{\text{normal}}}{M_{\text{observed}}} = \frac{136.3}{55} = 2.47$$



$$\text{Van't Hoff factor (i)} = \frac{1 - \alpha + \alpha + 2\alpha}{1} = 2.47$$

$$\therefore \alpha = 0.735 = 73.5\%$$

---

## Question 146

The freezing point of a 1.00m aqueous solution of HF is found to be  $-1.91^\circ\text{C}$ . The freezing point constant of water,  $K_f$  is  $1.86\text{K kgmo}^{-1}$ . The percentage dissociation of HF at this concentration is  
[Online May 26, 2012]



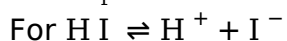
**Options:**

- A. 30%
- B. 10%
- C. 5.2%
- D. 2.7%

**Answer: D****Solution:**

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

$$i = \frac{\Delta T_f}{K_f \times m} = \frac{1.91}{1.86 \times 1} = 1.027$$



$$1 - \alpha + \alpha + \alpha = i = 1.027$$

$$1 + \alpha = 1.027$$

$$\alpha = 0.027 \text{ or } 2.7\%$$

**Question 147**

**Liquids A and B form an ideal solution. At 30°C, the total vapour pressure of a solution containing 1 mol of A and 2 mol of B is 250mmHg. The total vapour pressure becomes 300mmHg when 1 more mol of A is added to the first solution. The vapour pressures of pure A and B at the same temperature are [Online May 19, 2012]**

**Options:**

- A. 150, 450mmHg
- B. 125, 150mmHg
- C. 450, 150mmHg
- D. 250, 300mmHg

**Answer: C****Solution:****Solution:**

Let vapour pressure of A =  $P_A^\circ$

Vapour pressure of B =  $P_B^\circ$

In first solution,

$$\text{Mole fraction of A}(X_A) = \frac{1}{1+2} = \frac{1}{3}$$

$$\text{Mole fraction of B}(X_B) = \frac{2}{1+2} = \frac{2}{3}$$

According to Raoult's law,

$$\text{Total vapour pressure} = 250 = P_A^\circ X_A + P_B^\circ X_B$$



$$250 = \frac{1}{3}P_A^\circ + \frac{2}{3}P_B^\circ \dots(i)$$

In second solution

$$\text{Mole fraction of A}(X_A) = \frac{2}{2+2} = \frac{2}{4} = \frac{1}{2}$$

$$\text{Mole fraction of B}(X_B) = \frac{2}{4} = \frac{1}{2}$$

$$\therefore \text{Total vapour pressure} = 300 = P_A^\circ X_A + P_B^\circ X_B$$

$$300 = \frac{1}{2}P_A^\circ + \frac{1}{2}P_B^\circ \dots(ii)$$

Multiplying equation (i) by  $\frac{1}{2}$  and equation (ii) by  $\frac{1}{3}$

$$\frac{1}{6}P_A^\circ + \frac{2}{6}P_B^\circ = 125$$

$$\frac{1}{6}P_A^\circ + \frac{1}{6}P_B^\circ = 100$$

$$\frac{1}{6}P_B^\circ = 25$$

$P_B^\circ = 25 \times 6 = 150\text{mmHg}$  On substituting value of  $P_B^\circ$  in equation (ii) we get

$$300 = P_A^\circ \times \frac{1}{2} + 150 \times \frac{1}{2}$$

$$P_A^\circ = 450\text{mmHg}$$

## Question148

**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 :**

**[2011RS]**

**Options:**

A. 171.2

B. 68.4

C. 34.2

D. 136.2

**Answer: B**

**Solution:**

For isotonic solutions

$$\pi_1 = \pi_2$$

$$C_1 = C_2$$

$$\frac{5 / 342}{0.1} = \frac{1 / M}{0.1}$$

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

$$\Rightarrow M = \frac{342}{5} = 68.4\text{g / mol}$$

## Question149

**On mixing, heptane and octane form an ideal solution. At 373K, the**



vapour pressures of the two liquid components (heptane and octane) are 105kPa and 45kPa respectively. Vapour pressure of the solution obtained by mixing 25.0g of heptane and 35g of octane will be (molar mass of heptane = 100gmol<sup>-1</sup> and of octane = 114 gmol<sup>-1</sup>)  
[2010]

**Options:**

- A. 72.0kPa
- B. 36.1kPa
- C. 96.2kPa
- D. 144.5kPa

**Answer: A**

**Solution:**

$$\begin{aligned}P_{\text{Total}} &= P_A \cdot X_A + P_B \cdot X_B \\&= P_{\text{Heptane}} \cdot X_{\text{Heptane}} + P_{\text{Octane}} \cdot X_{\text{Octane}} \\&= 105 \times \frac{25 / 100}{\frac{25}{100} + \frac{35}{114}} + 45 \times \frac{35 / 114}{\frac{25}{100} + \frac{35}{114}} \\&= 105 \times \frac{0.25}{0.25 + 0.3} + 45 \times \frac{0.3}{0.25 + 0.3} \\&= \frac{105 \times 0.25}{0.55} + \frac{45 \times 0.3}{0.55} = \frac{26.25 + 13.5}{0.55} \\&= 72\text{kPa}\end{aligned}$$

---

## Question150

If sodium sulphate is considered to be completely dissociated into cations and anions in aqueous solution, the change in freezing point of water ( $\Delta T_j$ ), when 0.01 mol of sodium sulphate is dissolved in 1kg of water, is ( $K_j = 1.86\text{K kgmol}^{-1}$ )  
[2010]

**Options:**

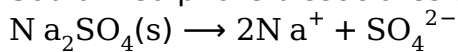
- A. 0.372K
- B. 0.0558K
- C. 0.0744K
- D. 0.0186K

**Answer: B**

**Solution:**



Sodium sulphate dissociates as



Hence van't Hoff factor,  $i = 3$

$$\text{Now } \Delta T_f = iK_f \cdot m$$

$$= 3 \times 1.86 \times 0.01 = 0.0558\text{K}$$

---

## Question151

Two liquids X and Y form an ideal solution. At 300K, vapour pressure of the solution containing 1mol of X and 3mol of Y is 550mmHg. At the same temperature, if 1 mol of Y is further added to this solution, vapour pressure of the solution increases by 10mmHg. Vapour pressure (in mmHg) of X and Y in their pure states will be, respectively:

[2009]

Options:

- A. 300 and 400
- B. 400 and 600
- C. 500 and 600
- D. 200 and 300

Answer: B

Solution:

$$P_{\text{total}} = P_x^\circ X_x + P_y^\circ X_y$$

$$550 = P_x^\circ \times \frac{1}{4} + P_y^\circ \times \frac{3}{4}$$

$$P_x^\circ + 3P_y^\circ = 550 \times 4 \dots (i)$$

In second case,

$$P_{\text{total}} = P_x^\circ \times \frac{1}{5} + P_y^\circ \times \frac{4}{5}$$

$$P_x^\circ + 4P_y^\circ = 560 \times 5 \dots (ii)$$

Subtract (i) from (ii)

$$\therefore P_y^\circ = 560 \times 5 - 550 \times 4 = 600$$

$$\therefore P_x^\circ = 400$$

---

## Question152

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?

[2009]

Options:

- A. The solution is non-ideal, showing -ve deviation from Raoult's Law.
- B. The solution is non-ideal, showing + ve deviation from Raoult's Law.



C. n-Heptane shows + ve deviation while ethanol shows - ve deviation from Raoult's Law.

D. The solution formed is an ideal solution.

**Answer: B**

**Solution:**

**Solution:**

For this solution intermolecular interactions between n -heptane and ethanol are weaker than n -heptane –n– heptane & ethanol-ethanol interactions, hence the solution of n -heptane and ethanol is non-ideal and shows positive deviation from Raoult'slaw.

---

## Question153

**At 80°C, the vapour pressure of pure liquid ' A ' is 520mm H g and that of pure liquid ' B ' is 1000mmH g. If a mixture solution of ' A ' and 'B' boils at 80°C and at 1 atm pressure, the amount of ' A ' in the mixture is (1atm = 760mmH g)**

**[2008]**

**Options:**

A. 52 mol percent

B. 34mol percent

C. 48 mol percent

D. 50mol percent

**Answer: D**

**Solution:**

**Solution:**

At 1 atmospheric pressure the boiling point of mixture is 80°C.

At boiling point the vapour pressure of mixture,  $P_T = 1 \text{ atmosphere} = 760\text{mmHg}$ .

Using the relation,

$P_T = P_A^\circ X_A + P_B^\circ X_B$ , we get

$P_B^\circ = 1000\text{mmHg}$ ,  $X_A + X_B = 1$  }

or  $760 = 520X_A + 1000 - 1000X_A$

or  $480X_A = 240$

or  $X_A = \frac{240}{480} = \frac{1}{2}$  or 50 mol. percent

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## Question154

**The vapour pressure of water at 20°C is 17.5mmHg. If 18 g of glucose ( $C_6H_{12}O_6$ ) is added to 178.2g of water at 20°C, the vapour pressure of the resulting solution will be**

**[2008]**



**Options:**

- A. 17.325mmHg
- B. 15.750mmHg
- C. 16.500mmHg
- D. 17.500mmHg

**Answer: A****Solution:****Solution:**

**Note:** On addition of glucose to water, vapour pressure of water will decrease. The vapour pressure of a solution of glucose in water can be calculated using the relation

$$\frac{P^\circ - P_s}{P^\circ} = \frac{\text{Moles of glucose in solution}}{\text{Moles of water in solution}}$$

$$\text{or } \frac{17.5 - P_s}{17.5} = \frac{18 / 180}{178.2 / 18} \quad [\because P^\circ = 17.5]$$

$$\text{or } 17.5 - P_s = \frac{0.1 \times 17.5}{9.9}$$

$$\text{or } P_s = 17.325\text{mmHg.}$$

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## Question155

**A mixture of ethyl alcohol and propyl alcohol has a vapour pressure of 290mm at 300K . The vapour pressure of propyl alcohol is 200mm. If the mole fraction of ethyl alcohol is 0.6, its vapour pressure (in mm ) at the same temperature will be [2007]**

**Options:**

- A. 360
- B. 350
- C. 300
- D. 700

**Answer: B****Solution:**

$$P_A^\circ = ?, \text{ Given } P_B^\circ = 200\text{mm}, X_A = 0.6,$$

$$X_B = 1 - 0.6 = 0.4, P = 290$$

$$P = P_A + P_B = P_A^\circ X_A + P_B^\circ X_B$$

$$290 = P_A^\circ \times 0.6 + 200 \times 0.4$$

$$\therefore P_A^\circ = 350\text{mm}$$



## Question156

Equal masses of methane and oxygen are mixed in an empty container at 25°C. The fraction of the total pressure exerted by oxygen is [2007]

Options:

- A. 1 / 2
- B. 2 / 3
- C.  $\frac{1}{3} \times \frac{273}{298}$
- D. 1 / 3

Answer: D

Solution:

Let the mass of methane and oxygen = mg.

Mole fraction of O<sub>2</sub>

$$= \frac{\text{Moles of O}_2}{\text{Moles of O}_2 + \text{Moles of CH}_4}$$

$$= \frac{m / 32}{m / 32 + m / 16} = \frac{m / 32}{3m / 32} = \frac{1}{3}$$

Partial pressure of O<sub>2</sub> = Total pressure × mole fraction of O<sub>2</sub>,

$$P_{\text{O}_2} = P \times \frac{1}{3} = \frac{1}{3}P$$

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## Question157

A 5.25% solution of a substance is isotonic with a 1.5% solution of urea (molar mass = 60gmol<sup>-1</sup>) in the same solvent. If the densities of both the solutions are assumed to be equal to 1.0gcm<sup>-3</sup>, molar mass of the substance will be [2007]

Options:

- A. 210.0gmol<sup>-1</sup>
- B. 90.0gmol<sup>-1</sup>
- C. 115.0gmol<sup>-1</sup>
- D. 105.0gmol<sup>-1</sup>.

Answer: A

Solution:



Osmotic pressure ( $\pi$ ) of isotonic solutions are equal. For solution of unknown substance

$$(\pi = CRT)$$

$$C_1 = \frac{5.25 / M}{V}$$

For solution of urea,

$$C_2(\text{ concentration } ) = \frac{1.5 / 60}{V}$$

$$\text{Given, } \pi_1 = \pi_2$$

$$\therefore \pi = CRT$$

$$\therefore C_1RT = C_2RT$$

$$\text{or } C_1 = C_2$$

$$\text{or } \frac{5.25 / M}{V} = \frac{1.5 / 60}{V}$$

$$\therefore M = 210 \text{ g / mol}$$

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## Question158

**Among the following mixtures, dipole-dipole as the major interaction, is present in [2006]**

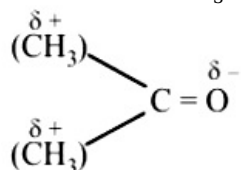
**Options:**

- A. KCl and water
- B. benzene and carbon tetrachloride
- C. benzene and ethanol
- D. acetonitrile and acetone

**Answer: D**

**Solution:**

Acetonitrile ( $\overset{\delta+}{\text{CH}_3} - \text{C} \equiv \overset{\delta-}{\text{N}}$ ) and acetone



both are polar molecules, hence dipole-dipole interaction exists between them. Between KCl and water, ion-dipole interaction is found and in benzene-ethanol and benzene- carbon tetrachloride dispersion force is present.

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## Question159

**Benzene and toluene form nearly ideal solution. At 20°C, the vapour pressure of benzene is 75 torr and that of toluene is 22 torr. The partial vapour pressure of benzene at 20°C for a solution containing 78g of benzene and 46g of toluene in torr is [2005]**

**Options:**



- A. 53.5
- B. 37.5
- C. 25
- D. 50

**Answer: D**

**Solution:**

Vapour pressure of benzene = 75 torr

Vapour pressure of toluene = 22 torr

mass of benzene in = 78g

Moles of benzene =  $\frac{78}{78} = 1\text{mol}$

∴ (Mol.wt of benzene = 78 )

Mass of toluene in solution = 46g

Hence moles of toluene =  $\frac{46}{92} = 0.5\text{mol}$

Now partial pressure of benzene

$$= P_b^0 \cdot X_b = 75 \times \frac{1}{1 + 0.5} = 75 \times \frac{1}{1.5}$$

$$= 75 \times \frac{2}{3} = 50$$

---

## Question160

**Equimolar solutions in the same solvent have [2005]**

**Options:**

- A. Different boiling and different freezing points
- B. Same boiling and same freezing points
- C. Same freezing point but different boiling points
- D. Same boiling point but different freezing points

**Answer: B**

**Solution:**

**Solution:**

Equimolar solutions of normal solutes in the same solvent will have the same b. pts and same f . pts.

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## Question161

**For which of the following parameters, the structural isomers  $\text{C}_2\text{H}_5\text{OH}$  and  $\text{CH}_3\text{OCH}_3$  would be expected to have the same values?(Assume ideal behaviour).**



[2004]

**Options:**

- A. Boiling points
- B. Vapour pressure at the same temperature
- C. Heat of vaporization
- D. Gaseous densities at the same temperature and pressure

**Answer: D**

**Solution:**

**Solution:**

Gaseous densities of ethanol and dimethyl ether would be same at same temperature and pressure. The heat of vaporisation, V.P. and b.pts will differ due to H bonding in ethanol.

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## Question162

**Which of the following liquid pairs shows a positive deviation from Raoult's law?**

[2004]

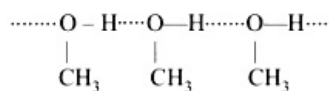
**Options:**

- A. Water - nitric acid
- B. Benzene-methanol
- C. Water - hydrochloric acid
- D. Acetone-chloroform

**Answer: B**

**Solution:**

**Note:** Positive deviations are shown by such solutions in which solvent-solvent and solute-solute interactions are stronger than the solvent-solute interactions. In such solutions, the interactions among molecules becomes weaker. Therefore, their escaping tendency increases which results in the increase in their partial vapour pressure. In a solution of methanol there exists intermolecular H-bonding.



In this solution benzene molecules come between methanol molecules which weaken intermolecular forces. This results in increase in vapour pressure.

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## Question163

**Which one of the following aqueous solutions will exhibit highest**





## boiling point? [2004]

### Options:

- A. 0.015M urea
- B. 0.01M  $KNO_3$
- C. 0.01M  $Na_2SO_4$
- D. 0.015M glucose

**Answer: C**

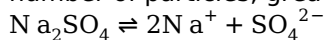
### Solution:

$$\because \Delta T_b = T_b - T^\circ$$

Where  $T_b =$  b. pt of solution

$$T_b^\circ = \text{b.pt of solvent or } T_b = T_b^\circ + \Delta T_b$$

**Note:** Elevation in boiling point is a colligative property, which depends upon the no. of particles. Thus greater the number of particles, greater is its elevation and hence greater will be its boiling point.



Since  $Na_2SO_4$  has maximum number of particles (3), hence has maximum boiling point.

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## Question 164

### Which one of the following statements is FALSE? [2004]

#### Options:

- A. The correct order of osmotic pressure for 0.01M aqueous solution of each compound is  $BaCl_2 > KCl > CH_3COOH > \text{sucrose}$
- B. The osmotic pressure ( $\pi$ ) of a solution is given by the equation  $\pi = MRT$ , where M is the molarity of the solution
- C. Raoult's law states that the vapour pressure of a component over a solution is proportional to its mole fraction
- D. Two sucrose solutions of same molality prepared in different solvents will have same freezing point depression

**Answer: D**

### Solution:

#### Solution:

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

Since  $K_f$  has different values for different solvents, hence even if m is same,  $\Delta T_f$  will be different.



## Question165

**If liquids A and B form an ideal solution [2003]**

**Options:**

- A. the entropy of mixing is zero
- B. the free energy of mixing is zero
- C. the free energy as well as the entropy of mixing is zero
- D. the enthalpy of mixing is zero

**Answer: D**

**Solution:**

**Solution:**

When A and B form an ideal solution,  $\Delta H_{\text{mix}} = 0$

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## Question166

**A pressure cooker reduces cooking time for food because [2003]**

**Options:**

- A. boiling point of water involved in cooking is increased
- B. the higher pressure inside the cooker crushes the food material
- C. cooking involves chemical changes helped by a rise in temperature
- D. heat is more evenly distributed in the cooking space

**Answer: A**

**Solution:**

**Solution:**

**Note:** On increasing pressure, the temperature is also increased. Thus in pressure cooker due to increase in pressure the b.p. of water increases.

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## Question167

**In a 0.2 molal aqueous solution of a weak acid HX, the degree of ionization is 0.3. Taking  $K_f$  for water as 1.85, the freezing point of the solution will be nearest to [2003]**



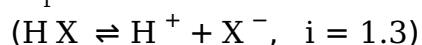
**Options:**

- A.  $-0.360^{\circ}\text{C}$
- B.  $-0.260^{\circ}\text{C}$
- C.  $+0.480^{\circ}\text{C}$
- D.  $-0.480^{\circ}\text{C}$

**Answer: D****Solution:**

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

$$\Delta T_f = 1.85 \times 0.2 \times 1.3 = 0.480^{\circ}\text{C}$$



$$1 - 0.3$$

$$\Delta T_f = T_f^{\circ} - T_f$$

$$\therefore T_f = 0 - 0.480^{\circ}\text{C} = -0.480^{\circ}\text{C}$$

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## Question168

**In mixture A and B components show -ve deviation as [2002]**

**Options:**

- A.  $\Delta V_{\text{mix}} > 0$  [2002]
- B.  $\Delta H_{\text{mix}} < 0$
- C. A – B interaction is weaker than A – A and B – B interaction
- D. A – B interaction is stronger than A – A and B – B interaction.

**Answer: D****Solution:****Solution:**

In solution containing A and B component showing negative deviation A-A and B-B interactions are weaker than that of A-B interactions. For such solutions,

$$\Delta H = -ve \text{ and } \Delta V = -ve$$

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## Question169

**Freezing point of an aqueous solution is  $(-0.186)^{\circ}\text{C}$ . Elevation of boiling point of the same solution is  $K_b = 0.512^{\circ}\text{C}$ ,  $K_f = 1.86^{\circ}\text{C}$ , find the increase in boiling point. [2002]**



**Options:**

- A. 0.186°C
- B. 0.0512°C
- C. 0.092°C
- D. 0.2372°C.

**Answer: B**

**Solution:**

$$\Delta T_b = K_b \frac{W_B}{M_B \times W_A} \times 1000$$

$$\Delta T_f = K_f \frac{W_B}{M_B \times W_A} \times 1000$$

$$\frac{\Delta T_b}{\Delta T_f} = \frac{K_b}{K_f}, \quad \frac{\Delta T_b}{0.186} = \frac{0.512}{1.86}$$
$$= 0.0512^\circ\text{C}.$$

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