

CHEMISTRY



SOLUTIONS

Introduction:

In this chapter, we will discuss about liquid solutions and their formation. This will be followed by studying the properties of solutions, like vapour pressure and colligative properties. We will begin with types of solutions and expressions for concentration of solutions in different units.

Thereafter, we will state and explain Henry's law and Raoult's law, distinguish between ideal and non-ideal solution and deviation of real solutions from Raoult's law. We will also discuss abnormal colligative properties along with association and dissociation of solute.

Types of Solutions

All the three states of matter (solid, liquid and gas) may behave either as solvent or solute. When a solution is composed of only two chemical substances, it is termed as binary solution. Depending upon the state of solute or solvent, binary solutions can be classified as

Type of Solutions	Solute	Solvent	Common Examples
<i>Gaseous Solutions</i>	Gas	Gas	Mixture of oxygen and nitrogen gases
	Liquid	Gas	Chloroform mixed with nitrogen gas
	Solid	Gas	Camphor in nitrogen gas
<i>Liquid Solutions</i>	Gas	Liquid	Oxygen dissolved in water
	Liquid	Liquid	Ethanol dissolved in water
	Solid	Liquid	Glucose dissolved in water
<i>Solid Solutions</i>	Gas	Solid	Solution of hydrogen in palladium
	Liquid	Solid	Amalgam of mercury with sodium
	Solid	Solid	Copper dissolved in gold

Some Important Definitions

- **Mixture** - When two or more chemically non-reacting substances are mixed, they form mixture.
- **Heterogeneous Mixture** - It consists of distinct phases, and the observed properties are just the sum of the properties of individual phases.

- **Homogeneous Mixture** - It consists of a single phase which has properties that may differ from one of the individual components.
- **Solution** - The homogeneous mixture of two or more components such that at least one component is a liquid is called solution.
- **Solvent** - It is the constituent of solution which has same physical state as that of solution and generally present in greater amount than all the other components.
- **Solute** - The component of a solution other than solvent is called solute, may or may not have same physical state as that of solution. Generally it is in smaller amount.

Example - In a sugar syrup (liquid solution) containing 60% sugar (solid) and 40% water (liquid), water is termed as solvent, due to same physical state as that of solution.

Expressing the Strength of Solution

For a given solution the amount of solute dissolved per unit volume of solution is called concentration of solute. Strength of solution is the amount of solute in grams dissolved in one litre of solution. It is generally expressed in g/litre.

Other methods of expressing the strength of solution are:

1. Mass percentage –

$$\text{Mass \% of solute} = \frac{\text{Mass of solute}}{\text{Total mass of solution}} \times 100$$

$$\text{Mass \% of solvent} = \frac{\text{Mass of solvent}}{\text{Total mass of solution}} \times 100$$

2. Volume percentage –

$$\text{Volume \% of solute} = \frac{\text{Volume of solute}}{\text{Total volume of solution}} \times 100$$

$$\text{Volume \% of solvent} = \frac{\text{Volume of solvent}}{\text{Total volume of solution}} \times 100$$

3. Molality (m) - It is no. of moles of solute dissolved in 1 kg of the solvent.

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

4. **Molarity (M)** - It is no. of moles of solute dissolved in 1 litre of solution.

$$M = \frac{(\text{Number of moles of solute})}{(\text{Mass of solution \{in litre\}})}$$

5. **Normality (N)** - It is no. of gram-equivalents of solute dissolved in 1 litre of solution

$$N = \frac{(\text{Number of gram equivalents of solute})}{(\text{Volume of solution \{in litre\}})}$$

6. **Formality** - Ionic solutes do not exist in the form of molecules. Their molecular mass is expressed as Gram-formula mass. Molarity for ionic compounds is actually called as **formality**.

7. **Mole fraction –**

$$\text{Mole fraction of solute} = \frac{(\text{Number of moles of solute})}{(\text{Total moles of solution})}$$

$$\text{Mole fraction of solvent} = \frac{(\text{Number of moles of solvent})}{(\text{Total moles of solution})}$$

For a binary solution,

$$\text{mole fraction of solute} + \text{mole fraction of solvent} = 1.$$

8. **Parts per million (ppm) –**

It is defined in two ways

$$\text{ppm} = \text{mass fraction} \times 10^6$$

$$\text{ppm} = \text{mole fraction} \times 10^6$$

Solubility

Solubility of a substance is its maximum amount that can be dissolved in a specified amount of solvent at a specified temperature. It depends upon the nature of solute and solvent as well as temperature and pressure. Let us consider the effect of these factors in solution of a **solid** or a **gas** in a **liquid**.

1. Solubility of Solid in Liquid

A solute dissolves in a solvent if the intermolecular interactions are similar in them, i.e., like dissolves like. Polar solute dissolves in polar solvent and non-polar solute in non-polar solvent. For e.g., sodium chloride and sugar dissolves readily in water and naphthalene and anthracene dissolves readily in benzene.

Solute + Solvent $\rightarrow$ Solution

- i. **Dissolution:** When a solid solute is added to the solvent, some solute dissolves and its concentration increases in solution. This process is called **dissolution**.
- ii. **Crystallization:** Some solute particles collide with solute particles in solution and get separated out. This process is called **crystallization**.
- iii. **Saturated solution:** Such a solution in which no more solute can be dissolved at the same temperature and pressure is called a **saturated solution**.
- iv. **Unsaturated solution:** An unsaturated solution is one in which more solute can be dissolved at the same temperature.
- v. **Effect of temperature:** In general, if in a nearly saturated solution, the dissolution process is endothermic, the solubility should increase with rise in temperature, if it is exothermic, the solubility should decrease with rise in temperature.
- vi. **Effect of pressure:** Solids and liquids are highly incompressible, so pressure does not have any significant effect on solubility of solids and liquids.
- vii. **Supersaturated solution:** When more solute can be dissolved at higher temperature in a saturated solution, then the solution becomes supersaturated.

2. Solubility of Gas in Liquid

All gases are soluble in water as well as in other liquids to a greater or lesser extent. The solubility of a gas in liquid depends upon the following factors Nature of the gas, Nature of solvent, Temperature and Pressure.

Generally, the gases which can be easily liquified are more soluble in common solvents. For e.g., CO₂ is more soluble than hydrogen or oxygen in water. The gases which are capable of

forming ions in aqueous solutions are much more soluble in water than other solvents. For e.g., HCl and NH₃ are highly soluble in water but not in organic solvents (like benzene) in which they do not ionize.

- i. **Effect of temperature:** Solubility of most of the gases in liquids decreases with rise in temperature. In dissolution of a gas in liquid, heat is evolved and thus this is an exothermic process. The dissolution process involves dynamic equilibrium and thus follows **Le Chatelier's** principle. As dissolution is exothermic the solubility of gas should decrease with rise in temperature.
- ii. **Effect of pressure: Henry's law:** At constant temperature, the solubility of a gas in a liquid is directly proportional to the pressure of the gas.

$$p = K_H \times x,$$

K_H = Henry's law constant.

Applications of Henry's law

1. In manufacture of soft drinks and soda water, CO₂ is passed at high pressure to increase its solubility.
2. To minimise the painful effects accompanying the decompression of deep sea divers. O₂ diluted with less soluble. He gas is used as breathing gas.
3. At high altitudes, the partial pressure of O₂ is less than that at the ground level. This leads to low concentrations of O₂ in the blood of climbers which causes 'anoxia'.

Vapour Pressure of Solution

It is the pressure exerted by vapour on the surface layer of liquid at equilibrium between vapour and liquid.

Factors affecting Vapour Pressure

- i. **Nature of liquid** - Liquid with higher intermolecular attraction forces form less amount of vapour and hence lower vapour pressure and vice-versa.
- ii. **Temperature** - Vapour pressure increases with temperature of liquid. This is because, as temperature increases, kinetic energy of the molecules increases, hence, more molecules leave the surface of the liquid and come into vapour phase.

Raoult's Law

According to Raoult's law, for a solution of volatile liquids, the relative lowering of vapour pressure of solution is directly proportional to its mole fraction of dissolved solvent in solute.

$$\frac{P^\circ - P}{P^\circ} = X_A$$

Ideal Solution

The solution which obeys **Raoult's law** at all compositions of solute and solvent and at all temperature is called an **ideal solution**. Ex- Benzene and Toluene, n-hexane and n-heptane.

Characteristics of an ideal Solution

1. Raoult's law is obeyed by it.
2. $\Delta H_{\text{mixing}} = 0$ i.e., no heat should be absorbed or evolved during mixing.
3. $\Delta V_{\text{mixing}} = 0$, i.e., no change in volume (expansion or contraction) on mixing.

Non-ideal Solution

Those solutions which deviate from ideal behaviour are called **non-ideal solutions** or real solutions. Acetone and CS_2 , Acetone and $\text{C}_2\text{H}_5\text{OH}$

Characteristics of a non-ideal Solution

1. Raoult's law is not obeyed by it.
2. $\Delta H_{\text{mixing}} \neq 0$ i.e., solution may absorb or release heat.
3. $\Delta V_{\text{mixing}} \neq 0$ i.e., solution may expand or contract on mixing of solute and solvent.

Azeotropic Mixture

At the constant boiling temperature, liquid mixture vapourises without change in composition and the condensate contains same composition, i.e., mixture distills like a pure liquid, which has same composition. At this point, solution or mixture is called an **azeotropic mixture**.

Colligative Properties

The properties of dilute solution which depends only on number of particles of solute (molecules or ions) present in the solution and not on their nature, are called colligative properties. The important **colligative properties** are;

1. Relative lowering of vapour pressure
2. Elevation of boiling point
3. Depression in freezing point
4. Osmotic pressure

i. Relative Lowering of Vapour Pressure

When a non-volatile solute is added to a solvent, its vapour pressure gets **lowered**. If this pressure is divided by pressure of pure solvent, this is called **relative lowering of vapour pressure**.

According to **Raoult's law**,

$$\frac{P^\circ - P}{P^\circ} = X_A$$

where, P° = V.P. of pure solvent

$P^\circ - P$ = Lowering in vapour pressure

$$\frac{P^\circ - P}{P^\circ} = \frac{n_A}{n_A + n_B} \quad \left[X_A = \frac{n_A}{n_A + n_B} \right]$$

For dilute solution $n_A + n_B \approx n_B$

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

$$\frac{P^\circ - P}{P^\circ} = \frac{W_A}{M_A} \times \frac{M_B}{W_B}$$

where, W_A = weight of solute

W_B = weight of solvent

M_A = molecular weight of solute

M_B = molecular weight of solvent

ii. Relative Elevation of boiling point

A liquid boils at the temperature at which its vapour pressure is equal to the atmospheric pressure. The boiling point of a solution of non-volatile solute is always higher than that of the boiling point of pure solvent in which the solution is prepared. Similar to lowering of vapour pressure, the elevation of boiling point also depends on the number of solute particles rather than their nature.

Let T° be the boiling point of pure solvent and T be the boiling point of solution. The increase in boiling point $\Delta T_b = T - T^\circ$ is known as **elevation in boiling point**.

For dilute solutions, the ΔT_b is directly proportional to the molal concentration of the solute in a solution. Thus

$$\Delta T_b \propto m$$

$$\Delta T_b = K_b m$$

K_b is molal **elevation constant** (Ebullioscopic constant). The unit of K_b is $K \text{ kg mol}^{-1}$.

Substituting the value of molality in above equation, we get

$$\Delta T_b = \frac{K_b \times 1000 \times w_2}{M_2 \times w_1}$$

$$M_2 = \frac{K_b \times 1000 \times w_2}{\Delta T_b \times w_1}$$

Where, w_1 = mass of solvent, w_2 = mass of solute and M_2 = molar mass of solute

iii. Depression in freezing point

Freezing point is the temperature at which vapour pressure of liquid phase becomes same as that of solid phase. The decrease in freezing point of a solvent on the addition of a non-volatile solute is known as **depression in freezing point**.

Let T° be the freezing point of pure solvent and T be the freezing point of solution. The decrease in freezing point $\Delta T_f = T^\circ - T$ is known as depression in freezing point.

For dilute solutions, the ΔT_f is directly proportional to the molal concentration of the solute in a solution. Thus

$$\Delta T_f \propto m$$

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

Here K_f is molal depression constant or cryoscopic constant

Substituting the value of molality in above equation, we get

$$\Delta T_f = \frac{K \times w_2 \times 1000}{M_2 \times w_1}$$

$$M_2 = \frac{K \times w_2 \times 1000}{\Delta T_f \times w_1}$$

Where, w_1 = mass of solvent, w_2 = mass of solute and M_2 = molar mass of solute

iv. Osmotic pressure

Osmosis is the spontaneous flow of the solvent molecules from a less concentrated solution (dilute) to a more concentrated solution through a **semi-permeable membrane**. The driving force of osmosis is called **osmotic pressure**. Osmotic pressure may be defined as "the minimum excess pressure that has to be applied on the solution to prevent the osmosis".

Osmotic pressure of a solution $\propto$ molar concentration of solute in that solution

$$\pi \propto c$$

$$\pi = cRT$$

where, R = Gas constant = $0.0821 \text{ lit atm K}^{-1} \text{ mole}^{-1}$

T = Temperature

c = Molar concentration

$$\pi = \frac{n}{V} RT \dots [c = \frac{n}{V}]$$

$$\pi = \frac{W_B}{M_B} \frac{RT}{V}$$

W_B = wt. of solute

M_B = Molar mass of solute

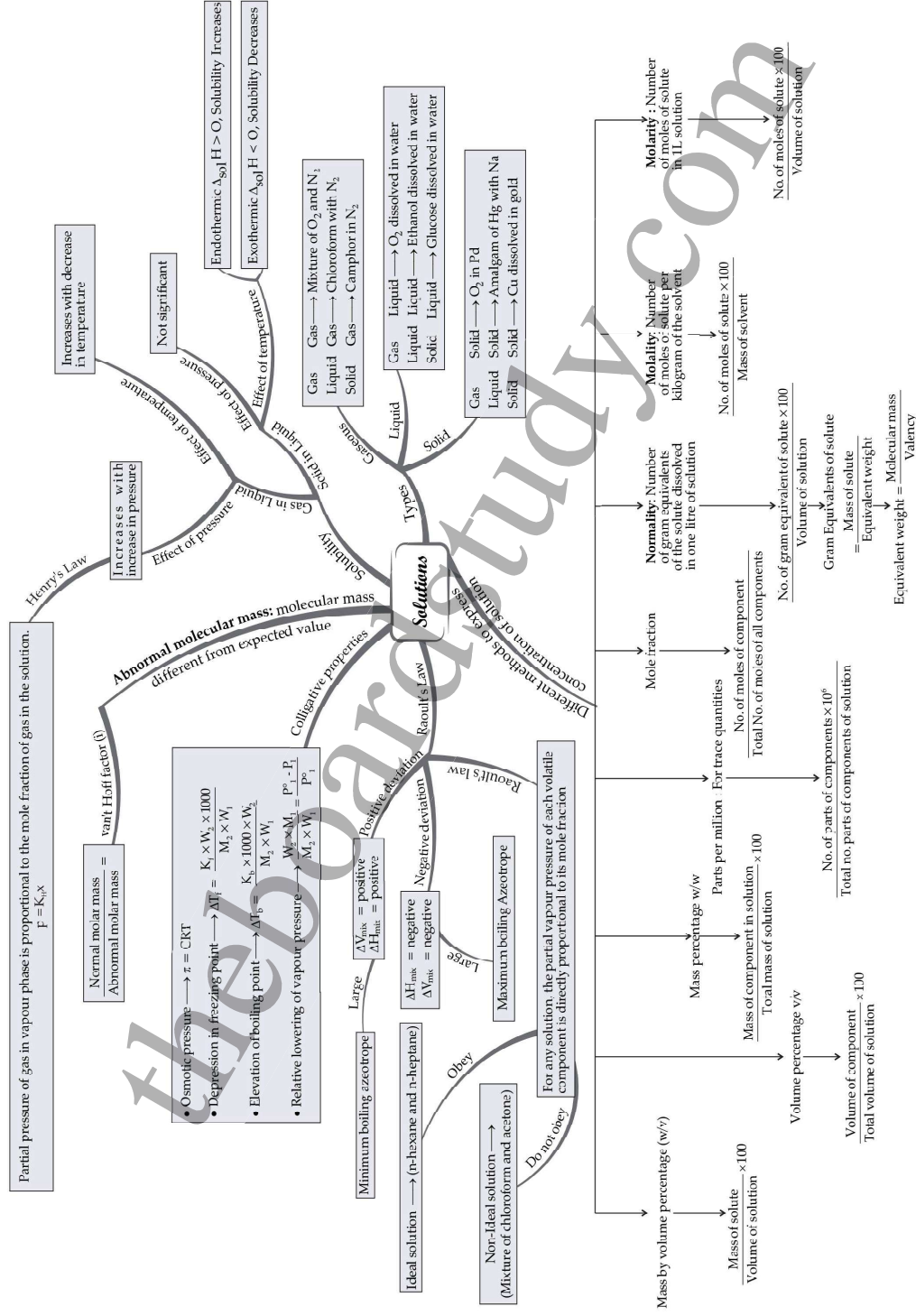
van't Hoff Factor

To calculate the extent of association or dissociation, van't Hoff in 1886 introduced a factor 'i' called van't Hoff factor. **van't Hoff factor 'i'** is defined as ratio of the experimental value of colligative property to the calculated value of colligative property.

i. e., $i = \frac{\text{Experiment colligative properties}}{\text{Calculated colligative properties}}$

MIND MAP : LEARNING MADE SIMPLE

CHAPTER - 2



Important Questions

Multiple Choice questions-

Question 1. Which of the following units is useful in relating concentration of solution with its vapour pressure?

- (a) mole fraction
- (b) parts per million
- (c) mass percentage
- (d) molality

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

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

Question 3. At equilibrium the rate of dissolution of a solid solute in a volatile liquid solvent is

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

Question 4. A beaker contains a solution of substance 'A'. Precipitation of substance 'A' takes place when small amount of 'A' is added to the solution. The solution is

- (a) saturated
- (b) supersaturated

(c) unsaturated

(d) concentrated

Question 5. Maximum amount of a solid solute that can be dissolved in a specified amount of a given liquid solvent does not depend upon .

(a) temperature

(b) nature of solute

(c) pressure

(d) nature of solvent

Question 6. Low concentration of oxygen in the blood and tissues of people living at high altitude is due to

(a) low temperature

(b) low atmospheric pressure

(c) high atmospheric pressure

(d) both low temperature and high atmospheric pressure

Question 7. Considering the formation, breaking and strength of hydrogen bond, predict which of the following mixtures will show a positive deviation from Raoult's law?

(a) Methanol and acetone.

(b) Chloroform and acetone.

(c) Nitric acid and water.

(d) Phenol and aniline.

Question 8. Colligative properties depend on

(a) the nature of the solute particles dissolved in solution.

(b) the number of solute particles in solution.

(c) the physical properties of the solute particles dissolved in solution.

(d) the nature of solvent particles.

Question 9. Which of the following aqueous solutions should have the highest boiling point?

(o) 1.0 M NaOH

(b) 1.0 M Na_2SO_4

(c) 1.0 M NH_4NO_3

(d) 1.0 M KNO_3

Question 10. The unit of ebullioscopic constant is

(a) K kg mol^{-1} or K (molality)^{-1}

(b) mol kg K^{-1} or $\text{K}^{-1} (\text{molality})$

(c) $\text{kg mol}^{-1} \text{K}^{-1}$ or $\text{K}^{-1} (\text{molality})^{-1}$

(d) K mol kg^{-1} or K (molality)

Question 11. In comparison to a 0.01 M solution of glucose, the depression in freezing point of a 0.01 M MgCl_2 solution is

(a) the same

(b) about twice

(c) about three times

(d) about six times

Question 12. An unripe mango placed in a concentrated salt solution to prepare pickle shrivels because

(a) it gains water due to osmosis.

(b) it loses water due to reverse osmosis.

(c) it gains water due to reverse osmosis.

(d) it loses water due to osmosis.

Question 13. At a given temperature, osmotic pressure of a concentrated solution of a substance

(a) is higher than that of a dilute solution.

(b) is lower than that of a dilute solution.

(c) is same as that of a dilute solution.

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

Question 14. Which of the following statements is false?

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

(b) The osmotic pressure of a solution is given by the equation $\pi = cRT$ (where c is the molarity of the solution).

(c) Decreasing order of osmotic pressure for 0.01 M aqueous solutions of barium chloride, potassium chloride, acetic acid and sucrose is $\text{BaCl}_2 > \text{KCl} > \text{CH}_3\text{COOH} > \text{sucrose}$.

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

Question 15. The values of Van't Hoff factors for KCl, NaCl and K_2SO_4 , respectively, are

(a) 2, 2 and 2

(b) 2, 2 and 3

(c) 1, 1 and 2

(d) 1, 1 and 1

Very Short Question:

Question 1. Define the term – solubility?

Question 2. What is the effect of pressure on solubility of a gas?

Question 3. State Henry's Law.

Question 4. State Raoult's Law.

Question 5. What are the factors on which vapour pressure depends?

Question 6. The vapour pressure of solvent gets lowered, when a non-volatile solute is added to it. Why?

Question 7. Name two ways by which vapour pressure of a liquid can be lowered.

Question 8. Define solution?

Question 9. Define the following terms:

(a) Molality

(b) Molarity

Question 10. How does change in temperature changes the molarity and molality values?

Short Questions:

Question 1 Find the molality and molarity of a 15% solution of H_2SO_4 when its density is

a) 1.10 g cm^{-3} & molar mass = 98 amu.

Question 2. Calculate the mole fraction of ethanol and water in a sample of rectified spirit which contains 46% ethanol by mass?

Question 3. Calculate the % composition in terms of mass of a solution obtained by mixing 300g of a 25% & 400 g of a 40% solution by mass?

Question 4. One litre of sea water weight 1030g and contains about $6 \times 10^{-3} \text{ g}$ of dissolved O_2 . Calculate the concentration of dissolved oxygen in ppm?

Question 5. The density of 85% phosphoric acid is 1.70 g cm^{-3} . What is the volume of a solution that contains 17g of phosphoric acid?

Question 6. Define the term azeotrope?

Question 7. Obtain a relationship between relative lowering of vapour pressure and mole fraction of solute?

Question 8. Draw the graphs of both deviations from ideal behaviours?

Question 9. A weak electrolyte AB is 5% dissociated in aqueous solution? What is the

freezing point of a 0.10 molar aqueous solution of AB? $K_f = 1.86 \text{ deg/molal}$?

Question 10. Henry's law constant for the molality of methane in benzene at 298 K is $4.27 \times 10^5 \text{ mmHg}$. Calculate the solubility of methane in benzene at 298 K under 760 mm Hg.

Long Questions:

Question 1.

The vapour pressure of CS_2 at 50°C is 854 mm Hg. A solution of 2.0g sulphur in 100g of CS_2 has a vapour pressure of 848.9 mm Hg. Calculate the formula of sulphur molecule.

Question 2. Calculate the mass percentage of benzene (C_6H_6) and carbon tetrachloride (CCl_4) if 22 g of benzene is dissolved in 122 g of carbon tetrachloride.

Question 3. Calculate the mole fraction of benzene in solution containing 30% by mass in carbon tetrachloride.

Question 4. Calculate the molarity of each of the following solutions: (a) 30 g of $\text{CO}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ in 4.3 L of solution (b) 30 mL of 0.5 M H_2SO_4 diluted to 500mL.

Question 5. Calculate (a) molality (b) molarity and (c) mole fraction of KI if the density of 20% (mass/mass) aqueous KI is 1.202 g mL^{-1} .

Assertion and Reason Questions:

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

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

Assertion: Camphor is used as a solvent in the determination of molecular masses of naphthalene, anthracene, etc.

Reason: Camphor has high molal elevation constant.

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

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

Assertion: Reverse osmosis is used in the desalination of sea water.

Reason: When pressure more than osmotic pressure is applied, pure water is squeezed out of the sea water through the membrane.

Case Study Questions:

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

The solubility of gases increases with increase of pressure. William Henry made a systematic investigation of the solubility of a gas in a liquid. According to Henry's law "the mass of a gas dissolved per unit volume of the solvent at constant temperature is directly proportional to the pressure of the gas in equilibrium with the solution". Dalton during the same period also concluded independently that the solubility of a gas in a liquid solution depends upon the partial pressure of the gas. If we use the mole fraction of gas in the solution as a measure of its solubility, then Henry's law can be modified as "the partial pressure of the gas in the vapour phase is directly proportional to the mole fraction of the gas in the solution"

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

- (i) Henry's law constant for the solubility of methane in benzene at 298K is $4.27 \times 10^5 \text{ mm Hg}$. The solubility of methane in benzene at 298K under 760mm Hg is:
 - a) 4.27×10^{-5}
 - b) 1.78×10^{-3}
 - c) 4.27×10^{-3}
 - d) 1.78×10^{-5}
- (ii) The partial pressure of ethane over a saturated solution containing $6.56 \times 10^{-2} \text{ g}$ of ethane is 1 bar. If the solution contains $5.00 \times 10^{-2} \text{ g}$ of ethane then what will be the partial pressure (in bar) of the gas?
 - a) 0.762
 - b) 1.312
 - c) 3.81
 - d) 5.0

(iii) K_H (K bar) values for $\text{Ar}_{(g)}$, $\text{CO}_{2(g)}$, $\text{HCHO}_{(g)}$ and $\text{CH}_{4(g)}$ are 40.39, 1.67, 1.83×10^{-5} and 0.413 respectively. Arrange these gases in the order of their increasing solubility. Arrange these gases in the order of their increasing solubility.

- a) $\text{HCHO} < \text{CH}_4 < \text{CO}_2 < \text{Ar}$
- b) $\text{HCHO} < \text{CO}_2 < \text{CH}_4 < \text{Ar}$
- c) $\text{Ar} < \text{CO}_2 < \text{CH}_4 < \text{HCHO}$
- d) $\text{Ar} < \text{CH}_4 < \text{CO}_2 < \text{HCHO}$

(iv) When a gas is bubbled through water at 298K, a very dilute solution of the gas is obtained. Henry's law constant for the gas at 298K is 150 kbar. If the gas exerts a partial pressure of 2 bar, the number of millimoles of the gas dissolved in 1L of water is:

- a) 0.55
- b) 0.87
- c) 0.37
- d) 0.66

(v) Which of the following statements is correct?

- a) K_H increases with increase of temperature.
- b) K_H decreases with increase of temperature.
- c) K_H remains constant with increase of temperature.
- d) K_H first increases then decreases, with increase of temperature.

2. Read the passage given below and answer the following questions

Few colligative properties are:

1. Relative lowering of vapour pressure: depends only on molar concentration of solute (mole fraction) and independent of its nature.
2. Depression in freezing point: it is proportional to the molal concentration of solution.
3. Elevation of boiling point: it is proportional to the molal concentration of solute.
4. Osmotic pressure: it is proportional to the molar concentration of solute

A solution of glucose is prepared with 0.052 g of glucose in 80.2 g of water. ($K_f = 1.86 \text{ K kg mol}^{-1}$ and $K_b = 5.2 \text{ K kg mol}^{-1}$)

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

(i) Molality of the given solution is.

- a) 0.0052m

- b) 0.0036m
- c) 0.0006m
- d) 1.29m

(ii) Boiling point for the solution will be.

- a) 373.05K
- b) 373.15K
- c) 373.02K
- d) 373.02K

(iii) The depression in freezing point of solution will be.

- a) 0.0187K
- b) 0.035K
- c) 0.082K
- d) 0.067K

(iv) Mole fraction of glucose in the given solution is.

- a) 6.28×10^{-5}
- b) 6.28×10^{-4}
- c) 0.00625
- d) 0.00028

(v) If same amount of sucrose ($C_{12}H_{22}O_{11}$) is taken instead of glucose, then.

- a) Elevation in boiling point will be higher.
- b) Depression in freezing point will be higher.
- c) Depression in freezing point will be lower.
- d) Both (a) and (b).

Answers key

MCQ answers:

1. Answer: (a) mole fraction
2. Answer: (d) Powdered sugar in hot water.
3. Answer: (c) equal to the rate of crystallisation
4. Answer: (b) supersaturated

5. Answer: (c) pressure
6. Answer: (b) low atmospheric pressure
7. Answer: (a) Methanol and acetone.
8. Answer: (b) the number of solute particles in solution.
9. Answer: (b) 1.0 M Na_2SO_4
10. Answer: (a) K kg mol^{-1} or K (molality)^{-1}
11. Answer: (c) about three times
12. Answer: (d) it loses water due to osmosis.
13. Answer: (a) is higher than that of a dilute solution.
14. Answer: (a) Two different solutions of sucrose of same molality prepared in different solvents will have the same depression in freezing point.
15. Answer: (b) 2, 2 and 3

Very Short Answers:

1. The maximum amount of a substance that can be dissolved in a specified amount of solvent is called its solubility.
2. The solubility of a gas increases with increases of pressure.
3. Answer: Henry's Law states that at a constant temperature the solubility of a gas in a liquid is directly proportional to the pressure of the gas.
4. Answer: Raoult's Law states that for a solution of volatile liquids, the partial vapour pressure of each component in the solution is directly proportional to its mole fraction.
5. Answer: The factors on which vapour pressure depends are –
1) Temperature of the liquid. 2) Nature of the liquid.
6. Answer: When a non-volatile solute is added to a solvent, the surface area for escape of solvent molecules decreases and vapour pressure gets lowered.
7. Answer: The two ways by which vapour pressure can be lowered are –
1) By decreasing the temperature.

- 2) By adding a non- volatile solute.
8. Solutions are homogeneous mixtures of two or more than two components..
9. (a) Molality is defined as the number of moles of the solute per kilogram of solvent.

$$\text{Molality (m)} = \frac{\text{Moles of solutes}}{\text{Mass of solvent in Kg}}$$

(b) Molarity (M) = Number of moles of solute dissolved in one litre of solution.

$$\text{Molarity (M)} = \frac{\text{No. of Moles of solute}}{\text{Volume of solution in litre}}$$

10. As the temperature increases, volume increases and molarity decreases whereas molality does not change with any change in temperature.

Short Answers:

1. Answer

Volume = mass/density

$$= 100 \text{ g} / 1.10 \text{ g/cm}^3 = 90.9 \text{ cm}^3$$

$$\text{Molarity} = \frac{\text{no. of moles of H}_2\text{SO}_4}{\text{volume of solution}} \times 1000$$

$$= \frac{(15/98)}{90.9} \times 1000 = 1.68 \text{ m}$$

$$\text{Molality} = \frac{\text{no. of moles of solute}}{\text{mass of solvent in kg}}$$

$$\frac{(15/98)}{85 \text{ g}} \times 1000$$

$$= 1.8 \text{ M.}$$

2. Answer:

Mass of ethanol = 46g

Mass of water = 100 – 46 = 54g

$$X_A = \frac{X_A}{X_A + X_B}$$

Mole fraction of ethanol,

$$\frac{\frac{46}{46}}{\frac{46}{46} + \frac{54}{18}} = \frac{1}{1+3} = \frac{1}{4} = 0.25.$$

Mole fraction of water = $1 - 0.25 = 0.75$

Mole fraction of water = $1 - 0.25 = 0.75$

3. Answer:

mass of solute in 400g of 40%

$$\frac{40}{100} \times 400 = 160\text{g}$$

Total mass of solute = $160 + 75 = 235\text{g}$

Total mass of solution = $400 + 300 = 700\text{g}$

$$\text{Mass\% of solute} = \frac{\text{mass of solute}}{\text{Total mass of solution}} \times 100$$

$$= \frac{235}{700} \times 100 = 33.57\%$$

Mass % of solvent = $100 - 33.57 = 66.43\%$

4. Answer:

mass of $\text{O}_2 = 6 \times 10^{-3} \text{g}$

$$\text{ppm of } \text{O}_2 \text{ in } 1030 \text{ g sea water} = \frac{\text{mass of } \text{O}_2}{\text{mass of sea water}} \times 10^6$$

$$= \frac{6 \times 10^{-3}}{1030} \times 10^6 = 5.8 \text{ ppm.}$$

5. Answer :

Ans. 85g phosphoric acid is present in 100g of solution.

17g of phosphoric acid is present in

$$\frac{100}{85} \times 17 = 20\text{g of solution}$$

$$\begin{aligned}\text{Volume of 17g of 85\% acid} &= \frac{\text{mass}}{\text{density}} \\ &= \frac{20\text{g}}{1.70\text{g/cm}^3} = 11.8\text{ cm}^3\end{aligned}$$

6. Answer

A solution at certain concentration when continues to boil at constant temperature without change in its composition in solution & in vapour phase is called an azeotrope.

7. Answer:

According to Raoult's Law –

$$P_1 = x_1 p_1^\circ$$

$$\Delta P = P_1^\circ - P = P_1^\circ - P_1^\circ X_1$$

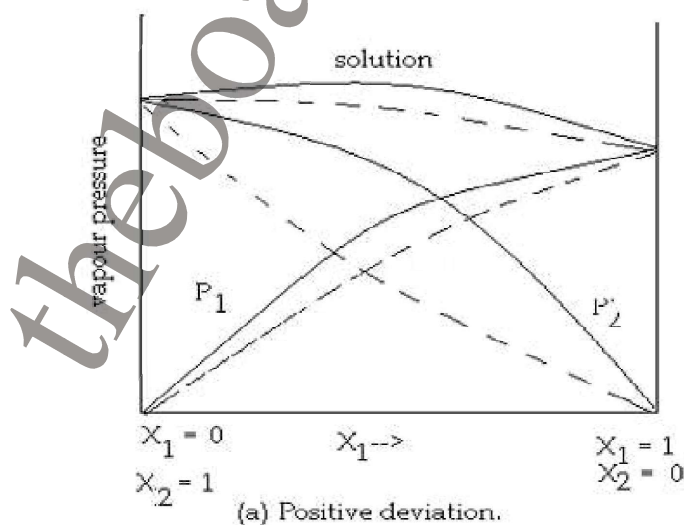
$$= \Delta P_1 = P_1^\circ (1 - X_1)$$

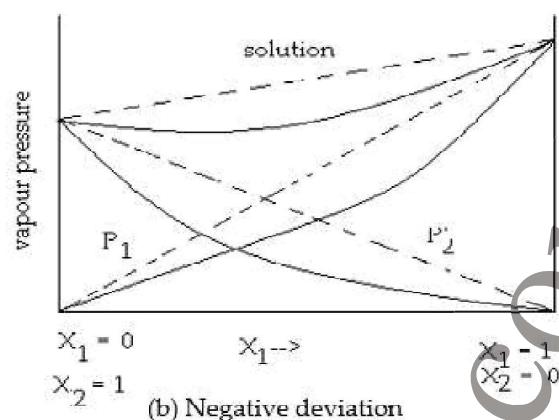
$$\Delta P_1 = P_1^\circ X_2 \quad (X_1 + X_2 = 1)$$

$$\frac{\Delta P_1}{P_1^\circ} = X_2$$

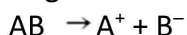
$$= \frac{P_1^\circ - P_1}{P_1^\circ} = X_2$$

Relative lowering of vapour pressure.

8. Answer:

**9. Answer :**

Degree of dissociation α of AB = $\frac{.5}{100} = 0.05$



M 0 0

No. of moles dissolved

No. of moles after dissociations

$m(1-\alpha)$ $m\alpha$ $m\alpha$

$0.1(1-0.05)$ 0.1×0.05 0.1×0.05

Total moles = $0.1(1-0.05) + (0.1 \times 0.05) + (0.1 \times 0.05)$

= $0.095 + 0.005 + 0.005 = 0.105m$

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

$$= 1.86 \times 0.105$$

$$= 0.1953 \text{ deg.}$$

$$T_f = 0^\circ\text{C} - 0.1953 = -0.1953^\circ\text{C}$$

10. Answer :

$$p = 760 \text{ mm Hg}$$

$$k_H = 4.27 \times 10^5 \text{ mm Hg}$$

According to Henry's law,

$$p = k_H x$$

$$x = \frac{p}{k_H}$$

$$= \frac{760 \text{ mm Hg}}{4.27 \times 10^5 \text{ mm Hg}}$$

$$= 177.99 \times 10^{-5}$$

$$= 178 \times 10^{-5} \text{ (approximately)}$$

Hence, the mole fraction of methane in benzene is 178×10^{-5} .

Long Answers:

1. Answer:

$$P_A^\circ = 854 \text{ mm } P_A = 848.9 \text{ mm}, W_B = 2.0 \text{ g},$$

$$W_A = 100 \text{ g } M_B = ?$$

$$M_A = 12 + 2(32) = 76 \text{ g/mol (CS}_2\text{)}$$

$$\frac{P_A^\circ - P_A}{P_A^\circ} = X_B = \frac{W_B}{M_B} \times \frac{W_A}{M_A}$$

$$M_B = W_B \times \frac{W_A}{M_A} \times \frac{P_A^\circ}{P_A^\circ - P_A}$$

$$= 2 \times \frac{100}{76} \times \frac{854}{854 - 848.9}$$

$$= 254.5 \text{ g/mol.}$$

Let the formula = S_x

$$X \times 32 = 254.5 \text{ g/mol}$$

$$X = \frac{254.5}{32}$$

$$= 7.95$$

$$= 8.$$

$$= \text{Formula} = S_8$$

2. Answer:

$$\text{Mass percentage of } C_6H_6 = \frac{\text{Mass of } C_6H_6}{\text{Total mass of the solution}} \times 100\%$$

$$= \frac{\text{Mass of } C_6H_6}{\text{Mass of } C_6H_6 + \text{Mass of } CCl_4} \times 100\%$$

$$= \frac{22}{22 + 122} \times 100\%$$

$$= 15.28\%$$

$$\text{Mass percentage of } \text{CCl}_4 = \frac{\text{Mass of } \text{CCl}_4}{\text{Total mass of the solution}} \times 100\%$$

$$= \frac{\text{Mass of } \text{CCl}_4}{\text{Mass of } \text{C}_6\text{H}_6 + \text{Mass of } \text{CCl}_4} \\ = \frac{122}{22 + 122} \times 100\%$$

$$= 84.72\%$$

Alternatively,

$$\text{Mass percentage of } \text{CCl}_4 = (100 - 15.28) \%$$

$$= 84.72\%$$

3. Answer:

Let the total mass of the solution be 100 g and the mass of benzene be 30g.

$$\therefore \text{Mass of carbon tetrachloride} = (100 - 30) \text{ g} \\ = 70 \text{ g}$$

$$\text{Molar mass of benzene (C}_6\text{H}_6) = (6 \times 12 + 6 \times 1) \text{ g mol}^{-1} \\ = 78 \text{ g mol}^{-1}$$

$$\therefore \text{Number of moles of } \text{C}_6\text{H}_6 = \frac{30}{78} \text{ mol}$$

$$= 0.3846 \text{ mol}$$

$$\text{Molar mass of carbon tetrachloride (CCl}_4) = 1 \times 12 + 4 \times 35.5 \\ = 154 \text{ g mol}^{-1}$$

$$\therefore \text{Number of moles of } \text{CCl}_4 = \frac{70}{154} \text{ mol}$$

$$= 0.4545 \text{ mol}$$

Thus, the mole fraction of C_6H_6 is given as:

$$\frac{\text{Number of moles of } C_6H_6}{\text{Number of moles of } C_6H_6 + \text{Number of moles of } CCl_4}$$

$$= \frac{0.3846}{0.3846 + 0.4545}$$

$$= 0.458$$

4. Answer:

Molarity is given by:

$$\text{Molarity} = \frac{\text{Moles of solute}}{\text{Volume solution in litre}}$$

(a) Molar mass of $CO(NO_3)_2 \cdot 6H_2O$ $59 + 2(14 + 3 \times 16) + 6 \times 18$
 $= 291 \text{ g mol}^{-1}$

Therefore, Moles of $CO(NO_3)_2 \cdot 6H_2O = \frac{30}{291} \text{ mol}$
 $= 0.103 \text{ mol}$

Therefore, $= \frac{0.103 \text{ mol}}{4.3 \text{ L}}$ molarity

$$= 0.023 \text{ M}$$

(b) Number of moles present in 1000 mL of 0.5 $H_2SO_4 = 0.5 \text{ mol M}$
 $\therefore$ Number of moles present in 30 mL of 0.5 M $H_2SO_4 = \frac{0.5 \times 30}{1000} \text{ mol}$
 $= 0.015 \text{ mol}$

Therefore, molarity $= \frac{0.015 \text{ mol}}{0.5 \text{ L}} = 0.03 \text{ M}$

5. Answer:

(a) Molar mass of $KI = 39 + 127 = 166 \text{ g mol}^{-1}$

20% (mass/mass) aqueous solution of KI means 20 g of KI is present in 100 g of solution.

That is,

20 g of KI is present in $(100 - 20) \text{ g of water} = 80 \text{ g of water}$

Therefore, molality of the solution = $\frac{\text{Moles of } KI}{\text{Mass of water in kg}}$

$$\frac{20}{\frac{166}{0.08} \text{ m}}$$

$$= 1.506 \text{ m}$$

$$= 1.51 \text{ m (approximately)}$$

(b) It is given that the density of the solution = 1.202 g ml^{-1}

Therefore, Volume of 100 g solution = $\frac{\text{Mass}}{\text{Density}}$

$$\frac{100 \text{ g}}{1.202 \text{ g ml}^{-1}}$$

$$= 83.19 \text{ mL}$$

$$= 83.19 \times 10^{-3} \text{ L}$$

Therefore, molarity of the solution = $\frac{\frac{20}{166} \text{ mol}}{83.19 \times 10^{-3} \text{ L}}$

(c) Moles of KI = $\frac{20}{166} = 0.12 \text{ mol}$

Moles of water = $\frac{80}{18} = 4.44 \text{ mol}$

Therefore, mole fraction of KI = $\frac{\text{Moles of KI}}{\text{Moles of KI} + \text{Moles of water}}$
 $= \frac{0.12}{0.12 + 4.44}$
 $= 0.0263$

Assertion and Reason Answers:

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

Explanation:

Camphor has high molal depression constant.

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

Explanation:

If a pressure larger than the osmotic pressure is applied to the solution side, the pure solvent flows out of the solution to the solvent through semi-permeable membrane and this phenomenon is called reverse osmosis.

Case Study Answers:

1. Answer :

i. (b) 1.78×10^{-3}

Explanation:

$$K_H = 4.17 \times 10^5 \text{ mm HG}$$

$$p = 760 \text{ mm Hg}$$

According to Henry's law, $P = K_H \times X_{\text{CH}_4}$

$$X_{\text{CH}_4} = \frac{P}{K_H} = \frac{760}{4.17 \times 10^5} = 1.78 \times 10^{-3}$$

ii. (a) 0.762

Explanation:

According to Henry's law, $m = K_H \times p$

$$6.56 \times 10^{-2} = K_H \times 1$$

For another case, $5 \times 10^{-2} = 6.56 \times 10^{-2} \times p$

$$p = \frac{5 \times 10^{-2}}{6.56 \times 10^{-2}} = 0.762 \text{ bar}$$

iii. (c) $\text{Ar} < \text{CO}_2 < \text{CH}_4 < \text{HCHO}$

Explanation:

Higher the value of K_H at a given pressure, the lower is the solubility of the gas.

iv. (c) 0.37

Explanation:

The mole fraction of the gas in solution,

$$x = \frac{p}{K_H} = \frac{1}{150 \times 10^3}$$

If n is the number of moles of gas in a solution of 1 L of water containing 55.5 mol then,

$$x = \frac{n}{n+55.5} \text{ or, } \frac{n}{55.5} = \frac{1}{150 \times 10^3}$$

$[n + 55.5 \approx 55.5, \text{ as } n \text{ is very small}]$

$$n = \frac{55.5}{150} \times 10^{-3} = 0.37 \text{ millimoles}$$

v. (a) K_H increases with increase of temperature.

2. Answer :

i. (b) 0.0036m

Explanation:

$$m = \frac{0.052}{180} \times \frac{1000}{80.2} = 0.0036$$

ii. (c) 373.02K

Explanation:

$$\Delta T_b = k_b \times m = 5.2 \times 0.0036 = 0.0187 \text{ K}$$

$$T_b = 373 + 0.0187 = 373.0187 \text{ K} \approx 373.02 \text{ K}$$

iii. (d) 0.067 K

Explanation:

$$\Delta T_f = k_f \times m = 1.86 \times 0.0036 = 0.067 \text{ K}$$

iv. (a) 6.28×10^{-5}

Explanation:

$$\text{Moles of water} = \frac{80.2}{18} = 4.455$$

$$\text{Mole fraction of glucose} = \frac{0.00028}{4.45 + 0.00028} = 6.28 \times 10^{-5}$$

v. (c) Depression in freezing point will be lower.

Explanation:

Depression in freezing point or elevation in boiling point is proportional to molarity, which is proportional to number of moles. For same amount, higher the molar mass of solute, lower will be number of moles. Hence, lower will be the colligative property.