



DPP SOLUTION

- **Subject – Physical Chemistry**
- **Chapter – Solutions**

DPP No.- 01



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Question



Homogeneous mixture of two or more than two components is called

- 1 Solute ✗
- 2 Solvent ✗
- 3 Both (A) & (B)
- ~~4~~ Solution

Sugar + water
↓
Homogeneous

Ans (D)

Which of the following is an example of gaseous solution?

- 1 Camphor in nitrogen gas ✓
solid *N₂ (g)*
- 2 Solution of hydrogen in palladium ✗
solute *solvent*
- 3 Chloroform mixed with nitrogen gas ✓
solute *solvent*
- ~~4 Both (A) & (C)~~

Ans (D)

In amalgam of mercury with sodium, solvent is

- ① Mercury
- ② Sodium
- ③ Amalgam
- ④ None of these

Na(Hg)
↓
Sodium alloy with Hg(l)
↓
solvent

↓
solute

Question



The unit of molality is

- ① mol L⁻¹
- ② mol kg⁻¹
- ③ mol⁻¹ L⁻¹
- ④ mol L

$$m = \frac{n_B}{w_A \text{ (in Kg)}} \rightarrow \frac{\text{mol}}{\text{Kg}}$$

Ans (B)

Question



The partial pressure of the gas in vapour phase is proportional to the mole fraction of the gas in the solution is given by

1 Raoult's law

2 Ostwald's law

3 Distribution law

~~4~~ Henry's law

$$P_A \propto x_A$$
$$P_A = \underline{K_H x_A}$$

Ans (4)

Question



Four gases like H_2 , He , CH_4 and CO_2 has Henry's constant values (K_H) are 69.16, 144.979, 0.413 and 1.67. The gas which is more soluble in liquid is

1 He

~~2 CH₄~~

3 H₂

4 CO₂

	H ₂	He	CH ₄	CO ₂
K _H	69.16	144.979	<u>0.413</u>	1.67

$K_H \propto \frac{1}{\text{Solubility}}$

Ans (2)

Question



The solubility of gas in a liquid increase with

- ① Increase of temperature
- ② Amount of liquid taken ✕
- ③ Decreases in temperature
- ④ Reduction of gas pressure

$T \uparrow$ solubility dec

$T \downarrow$ solubility inc

Question



→ solid or gas

Solubility of a substance is its maximum amount that can be dissolved in a specified amount of solvent. It depends upon

(i) Nature of solute ✓

(ii) Nature of solvent ✓

(iii) Temperature ✓

(iv) Pressure ✓

1 Only (i), (ii) and (iii) ✓

2 Only (i), (iii) and (iv)

3 Only (i) and (iv)

~~4~~ (i), (ii), (iii) and (iv)

Ans (4)

Question



During dissolution when solute is added to the solvent, some solute particles separate out from the solution as a result of crystallization. At the stage of equilibrium, the concentration of solute in the solution at given temperature and pressure.



2 Decreases

3 Remains constant

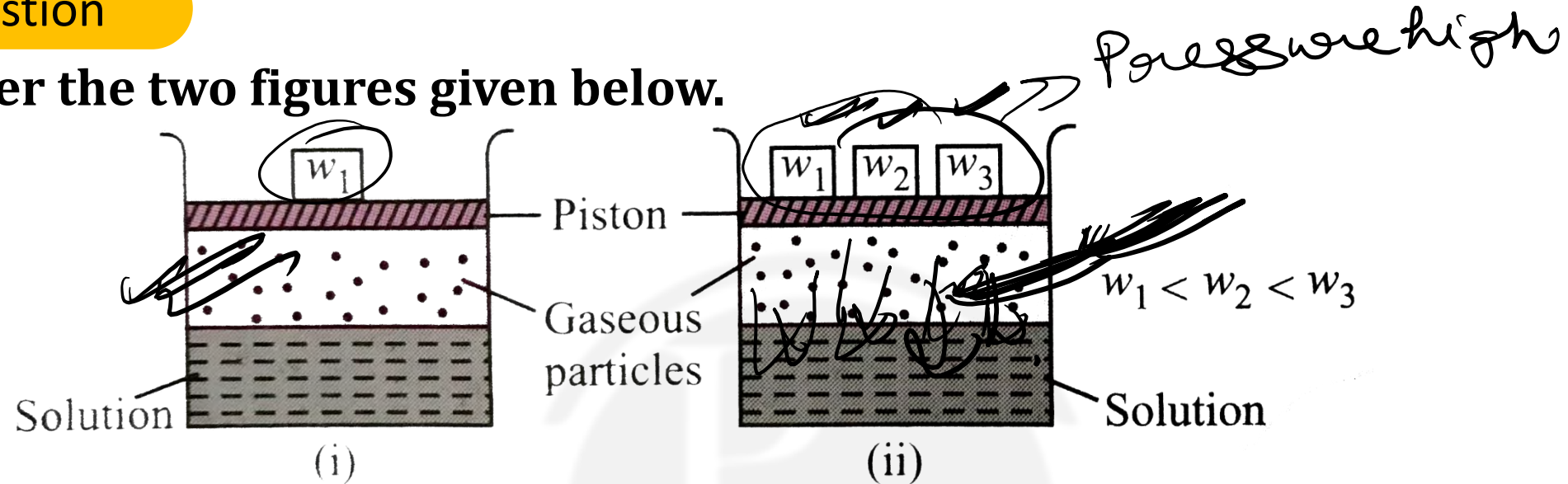
4 Keeps changing

$P, T, \text{Conc, density constt.}$

Question



Consider the two figures given below.



of the following statements regarding the experiment is true?

- ☒ 1 The solubility of a gas in liquid in beaker (i) is greater than that in beaker (ii).
- ☒ 2 The solubility of gas in beaker (i) is less than that in beaker (ii)
- ☐ 3 The solubility of gas is equal in both beakers.
- ☐ 4 The solubility of gas remains unaffected by change in weights.

According to Henry's law the partial pressure of the gas in vapour phase (p) is proportional to the mole fraction of the gas (x) in the solution. For different gases the correct statements about Henry's constant is $P \propto x$

- ① ~~X~~ Higher the value of K_H at a given pressure, higher is the solubility of the gas.
- ② ~~X~~ Higher in the value of K_H at a given pressure, lower the solubility of the gas
- ③ ~~X~~ K_H is not a function of nature of gas
- ④ ~~X~~ K_H value for all gases is same at a given pressure.

Question



The value of Henry's law constant for some gases at 293 K is given below. Arrange the gases in the increasing order of their solubility.

✓ He : 144.97 kbar, H₂ : 69.16 kbar

✓ N₂ : 76.48 kbar, O₂ : 34.86 kbar

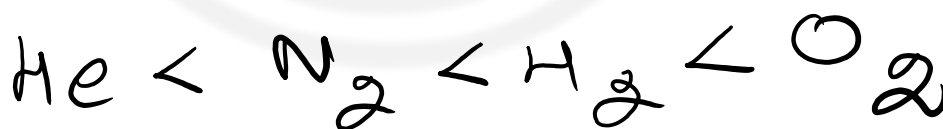
$$K_H \propto \frac{1}{\text{solubility}}$$

✓ 1 He < N₂ < H₂ < O₂

2 O₂ < H₂ < N₂ < He

3 H₂ < N₂ < O₂ < He

4 He < O₂ < N₂ < H₂



Ans (1)

Question

$$\%_B = \%_{H_2S}$$

$$M_A = 18g$$



H₂S is a toxic gas used in qualitative analysis. If solubility of H₂S in water at STP is 0.195 m, what is the value of K_H?

① 0.0263 bar

② 69.16 bar

③ 192 bar

④ 282 bar

$$m = 0.195m$$

$$P_{H_2S} = K_H \times \%_{H_2S}$$

$$1 = K_H \times \frac{3.51}{1003.51}$$

$$K_H = \frac{1003.51}{3.51} = 282 \text{ bar}$$

$$m = \frac{\%_B \times 1000}{\%_A \times M_A}$$

$$P = 1 \text{ bar}$$

$$0.195 = \frac{\%_B \times 1000}{(1 - \%_B) 18}$$

$$0.195 = \frac{1000 \%_B}{18 - 18 \%_B}$$

$$3.51 - 3.51 \%_B = 1000 \%_B$$

$$\%_B = \frac{3.51}{1003.51}$$

Ans (4)

Question



Henry's law constant for molality of methane in benzene at 298 K is 4.27×10^5 mm Hg. The mole fraction of methane in benzene at 298 K under 760 mm Hg is

~~1~~ 1.78×10^{-3} $K_H = 4.27 \times 10^5$ mm of Hg $P_{CH_4} = 760$ mm of Hg

2 17.43 $\%_{CH_4} = ?$

3 0.114 $P_{CH_4} = K_H \times \%_{CH_4}$

4 2.814 $760 = 4.27 \times 10^5 \times \%_{CH_4}$

$\frac{760}{4.27} \times 10^{-3} = \%_{CH_4}$

$\%_{CH_4} = 1.748 \times 10^{-3}$

Ans (1)

Question



When a gas is bubbled through water at 298 K, a very dilute solution of gas is obtained. Henry's law constant for the gas is 100 k bar. If gas exerts a pressure of 1 bar, the number of moles of gas dissolved in 1 litre of water is

- ① 0.555
- ② 55.55×10^{-5}
- ③ 55.55×10^{-3}
- ④ 5.55×10^{-5}

$$K_H = 100 \text{ Kbar} \\ = 100000 \text{ bar}$$

$$P_{\text{gas}} = 1 \text{ bar}$$

$$n_{\text{gas}} = ?$$

1 Litre of water \rightarrow solvent

$$1000 \text{ ml of } H_2O = V$$

$$1 \text{ g/ml of } H_2O = d$$

$$W_A = 1000 \times 1 = 1000 \text{ g}$$

$$n_A = \frac{1000}{18} = 55.55$$

Ans (2)

$$P_{\text{gas}} = K_H(\text{gas}) \%_{\text{gas}}$$

$$1 = \frac{10^5 n_{\text{gas}}}{n_{\text{gas}} + 55.55}$$

$$n_{\text{gas}} + 55.55 = 10^5 n_{\text{gas}}$$

$$55.55 = 10^5 n_{\text{gas}} - n_{\text{gas}} \approx 10^5 n_{\text{gas}}$$

$$n_{\text{gas}} = \frac{55.55}{10^5} = 55.55 \times 10^{-5}$$

Question



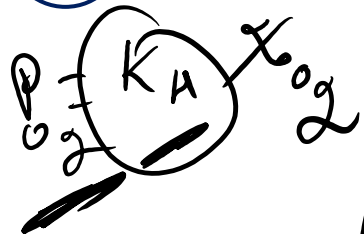
How much oxygen is dissolved in 100 mL water at 298 K if partial pressure of oxygen is 0.5 atm and $K_H = 1.4 \times 10^{-3} \text{ mol/L/atm}$?

1 22.4mg

2 22.4 g

3 2.24 g

4 2.24mg



$$w_{O_2} = ?$$

$$V_{\text{of } H_2O} = 100 \text{ ml}$$

$$T = 298 \text{ K}$$


$$P_{O_2} = 0.5 \text{ atm}$$

$$K_H = 1.4 \times 10^{-3} \text{ mol L}^{-1} \text{ atm}^{-1}$$

$$S_{O_2} = K_H P_{O_2}$$

$$K = \frac{S_{O_2}}{P_{O_2}} \Rightarrow \frac{\text{mol}}{\text{L atm}}$$

Ans (4)


$$S_{O_2} = K_H P_{O_2}$$
$$= 1.4 \times 10^{-3} \times 0.5 = 0.7 \times 10^{-3} = 7 \times 10^{-4} \text{ mol/L}$$

$$\begin{array}{l} 1000 \text{ ml has moles of } O_2 = 7 \times 10^{-4} \text{ mol} \\ 100 \text{ ml} \end{array} \quad \underbrace{\hspace{1cm}} = \frac{7 \times 10^{-4} \times 100}{1000} = 7 \times 10^{-5} \text{ mol}$$

$$M_{O_2} = 32 \text{ g}$$

$$\begin{aligned} \text{Mass of } O_2 &= 7 \times 10^{-5} \times 32 = 224 \times 10^{-5} \text{ g} \\ &= 2.24 \times 10^{-3} \text{ g} \\ &= 2.24 \text{ mg} \end{aligned}$$

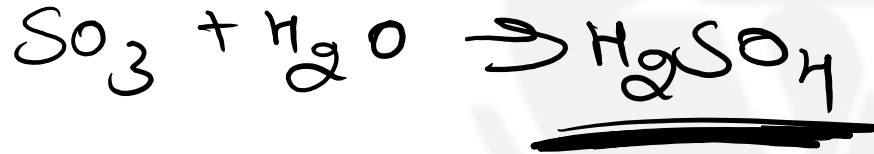
Henry's law is not applicable for aqueous solution of

1 O_2

2 N_2

~~3 SO_3~~

4 He



Question



On increasing temperature, the solubility of NaNO_3 in water



endothermic

$T \uparrow$ solubility \uparrow

☒ 1 Increases

☐ 2 Decreases

☐ 3 Remains unaffected

☐ 4 Cannot be determined

Ans (1)

Question



O₂ is bubbled through water at 293 K. Assume that O₂ exerts a partial pressure of 0.98 bar, find the solubility of O₂ in g L⁻¹. The value of Henry's Law constant K_H for O₂ is 34.84 k bar.

~~1~~ 0.05

$$P_{O_2} = 0.98 \text{ bar}$$

$$K_H = 34.84 \text{ k bar}$$

$$= 34840 \text{ bar}$$

2 0.08

3 0.07

4 0.01

$$P_{O_2} = K_H \cdot x_{O_2}$$

$$0.98 = \frac{34840 \cdot n_{O_2}}{n_{O_2} + n_{H_2O}}$$

$$\text{volume water}^{(A)} = 1 \text{ L} = 1000 \text{ ml}$$

$$W_A = 1000 \text{ g} \quad M_A = 18 \text{ g}$$

$$n_{H_2O} = \frac{W_A}{M_A} = \frac{1000}{18} = 55.55$$

$$0.98 = \frac{34840 \cdot n_{O_2}}{n_{O_2} + 55.55}$$

$$0.98 n_{O_2} + 54.439 = 34840 n_{O_2}$$

Ans (1)

$$\underline{34840\text{g}}_{\text{O}_2} - \underline{0.98\text{g}}_{\text{O}_2} = 54.43\text{g}$$

$$34839\text{g}_{\text{O}_2} = 54.43\text{g}$$

$$n_{\text{O}_2} = \frac{54.43\text{g}}{34839} = 0.0015\text{ mol}$$

$$\begin{aligned} n_{\text{O}_2} &= 0.0015 \times 32\text{ g} \\ &= 0.048\text{ g} \end{aligned}$$



Thank

You...

