

## Henry law problems

- ①  $H_2S$  a toxic gas with rotten egg like smell is used for the qualitative analysis, if the solubility of  $H_2S$  in water, at STP is  $0.195 \text{ mole} \cdot \text{kg}^{-1}$  the Henry law constant is - - -

Given  $n_{H_2S} = 0.195 \text{ moles}$

$n_{H_2O} = 55.55 \text{ moles}$

$$\chi_{H_2S} = \frac{n_{H_2S}}{n_{H_2S} + n_{H_2O}} = \frac{0.195}{0.195 + 55.55} = \frac{0.195}{55.695} = 0.0035$$

at STP pressure in bar =  $0.987 \text{ bar}$

Henry Law

$$P = K_H \chi$$

$$K_H = \frac{P}{\chi} = \frac{0.987}{0.0035} = \underline{\underline{285 \text{ bar}}}$$

- Q9) air contains  $O_2$  and  $N_2$  in the ratio  $0.2 : 0.8$ . if Henry law constant for  $O_2$  &  $N_2$  are  $3.3 \times 10^7 \text{ torr}$  and  $6.6 \times 10^7 \text{ torr}$  respectively, then the ratio of mole fractions of  $O_2$  &  $N_2$  dissolved in water at 1 bar - pressure is

Given  $\frac{P_{O_2}}{P_{N_2}} = \frac{0.2}{0.8} = \frac{1}{4}$

$$\frac{K_{H_{O_2}}}{K_{H_{N_2}}} = \frac{3.3 \times 10^7}{6.6 \times 10^7} = \frac{1}{2}$$

$$\therefore \chi = \frac{P_{O_2}}{P_{N_2}} \cdot \frac{K_{H_{N_2}}}{K_{H_{O_2}}}$$

$$\frac{\chi_{O_2}}{\chi_{N_2}} = \frac{P_{O_2} / K_{H_{O_2}}}{P_{N_2} / K_{H_{N_2}}} = \frac{\frac{1}{3.3 \times 10^7}}{\frac{1}{6.6 \times 10^7}} = \underline{\underline{2:1}}$$

3) The quantity of  $\text{CO}_2$  in 500ml of soda water when packed under 2.5 atm  $\text{CO}_2$  pressure at 298 K is — gm ( $K_H = 1.67 \times 10^8 \text{ Pa}$  at 298 K)

$$P_{\text{CO}_2} = 2.5 \text{ atm} = 2.5 \times 10^5 \text{ Pascal} \quad (1 \text{ atm} = 10^5 \text{ Pascal})$$

Henry law

$$P_{\text{CO}_2} = K_H \cdot X_{\text{CO}_2}$$

$$X_{\text{CO}_2} = \frac{P_{\text{CO}_2}}{K_H} = \frac{2.5 \times 10^5}{1.67 \times 10^8} = 1.497 \times 10^{-3}$$

moles of water :-

$$\text{water volume} = 500 \text{ ml}$$

$$\text{water mass} = 500 \times 1 = 500 \text{ g} \quad \text{density}$$

$$\text{water moles (n)} = \frac{500}{18} = \underline{\underline{27.78}}$$

$$X_{\text{CO}_2} = \frac{n_{\text{CO}_2}}{n_{\text{CO}_2} + n_{\text{water}}} = \frac{n_{\text{CO}_2}}{n_{\text{water}}} \quad (\text{moles of } \text{CO}_2 \text{ negligible})$$

$$n_{\text{CO}_2} = X_{\text{CO}_2} \times n_{\text{water}} = 1.497 \times 10^{-3} \times 27.78 = 0.042 \text{ moles}$$

∴ 1 mole has 44g of  $\text{CO}_2$

∴ 0.042 mole has — g of  $\text{CO}_2$

$$0.042 \times 44 = \underline{\underline{1.86 \text{ g}}}$$

Henry law constant for the solubility of methane in benzene at 298K is  $4.27 \times 10^5$  mmHg then the solubility of methane in benzene at 298K under 760 mmHg in mol/kg is

$$K_H = 4.27 \times 10^5 \text{ mmHg}$$

$$P = 760 \text{ mmHg}$$

$$P = K_H \times X$$

$$X = \frac{P}{K_H} = \frac{760}{4.27 \times 10^5} = 1.78 \times 10^{-5} = \underline{\underline{1.78 \times 10^{-3}}}$$

$O_2$  is bubbled through water at 293K, assuming that  $O_2$  exerts a partial pressure of 0.98 bar the solubility of  $O_2$  in  $g \cdot L^{-1}$  is — ( $K_H = 34 \text{ Kbar}$ )

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

$$K_H = 34 \text{ Kbar} = 34 \times 10^3 \text{ bar}$$

$$P_{O_2} = K_H \times X_{O_2}$$

$$X_{O_2} = \frac{P_{O_2}}{K_H} = \frac{0.98}{34 \times 10^3} = 0.028 \times 10^{-3}$$

$$X_{O_2} = \frac{n_{O_2}}{n_{O_2} + n_{H_2O}} = \frac{n_{O_2}}{n_{H_2O}} = \frac{n_{O_2}}{55.55}$$

↓  
negligible

$$n_{O_2} = X_{O_2} \times 55.55$$

$$n_{O_2} = 0.028 \times 10^{-3} \times 55.55 = 1.56 \times 10^{-3} \text{ moles}$$

1 moles has ~~1.56 x 10~~ 32g  $O_2$

$1.56 \times 10^{-3}$  mole has — g  $O_2$

$$1.56 \times 10^{-3} \times 32$$

$$= \underline{\underline{0.05 \text{ g/L}}}$$

## Henry Law problems

① If  $N_2$  gas is bubbled through water at 293 K, how many moles of  $N_2$  gas would dissolve in 1L of water. Assume that  $N_2$  exerts a partial pressure of 0.987 bar. ( $K_H = 76.48 \text{ kbar}$ )  
( $7.16 \times 10^{-4} \text{ mol}$ )

② Henry's Law constant for oxygen dissolved in water is  $4.34 \times 10^4 \text{ atm}$  at  $25^\circ\text{C}$ . If the partial pressure of oxygen in air is 0.4 atm. Calculate moles of  $O_2$ ? and concentration (in moles per Lit) of the dissolved  $O_2$  in water.  
( $5.11 \times 10^{-4} \text{ mol}$ ,  $5.11 \times 10^{-4} \text{ mol}$ )

③ Henry's Law constant for the solubility of  $N_2$  gas in water at 298 K is  $1.0 \times 10^5 \text{ atm}$ . The mole fraction of  $N_2$  in air is 0.6. The number of moles of  $N_2$  from air dissolved in 10 moles of water at 298 K and 5 atm pressure is  
( $3 \times 10^{-4}$ )

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for a solution of acetone in chloroform,  
Henry's law constant is 150 torr at a temp  
of 300K. <sup>So</sup> Calculate the pressure of  
acetone when the mole fraction is 0.12  
(18 torr)

## RLVP problems

- ① At  $20^{\circ}\text{C}$ , the vapour pressure of diethyl ether is  $442\text{ mmHg}$ . When  $6.4\text{ g}$  of a non-volatile solute is dissolved in  $50\text{ g}$  of ether, the vapour pressure falls to  $410\text{ mmHg}$ . The molecular weight of the solute is — (molar mass of diethyl ether is  $= 74.8/\text{mol}$ )  
(130.8)
- ②  $139.18\text{ g}$  of glucose is added to  $178.2\text{ g}$  of water. The vapour pressure of water for this aqueous solution at  $10^{\circ}\text{C}$  is — (700 Torr)
- ③ The vapour pressure of acetone at  $20^{\circ}$  is  $185\text{ torr}$ . When  $1.2\text{ g}$  of a non-volatile substance was dissolved in  $100\text{ g}$  of acetone at  $20^{\circ}\text{C}$ , its vapour pressure was  $183\text{ Torr}$ . The molar mass of the substance is — (58 g/mol)  
(64)
- ④ The vapour pressure of pure water at  $25^{\circ}\text{C}$  is  $30\text{ mm}$ . The vapour pressure of  $10\%$  (w/w) glucose solution at  $25^{\circ}\text{C}$  is — (29.69 mmHg)

5) The weight of urea to be dissolved in 100g of water to decrease the vapour pressure of water by 5% is (16.66g)

6) 18g of Glucose is dissolved in 90g of water. The Relative lowering of vapour pressure of solution is — (1/51)

7) at a certain Temp., the vapour pressure of water is 50mm. The relative lowering of vapour pressure of a solution containing 36g of Glucose in 900g. of water is (0.004)

8) 3g of urea is dissolved in 36g of boiling water. The lowering in vapour pressure is (19mm)

9) Calculate the vapour pressure of 10% (w/w) aqueous glucose solution at 20°C, if vapour pressure of pure water is 31.8 mm — (31.45mm)

10) Vapour pressure of solvent is 76 cmHg and vapour pressure of solution is 38 cmHg. find out  
a) Lowering in vapour pressure (38 cmHg)  
b) Relative lowering in vapour pressure. (0.5)

## Osmotic pressure problems

- 1) at 283K, the Osmotic pressure of 2% solution of "X" is  $7.87 \times 10^4 \text{ Nm}^{-2}$ . Calculate molar mass of X? (598)  
( $R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$ )
- 2) A 5% solution of cane sugar is isotonic with 0.5% of X. The molecular weight of substance "X" is — (34.2 g/mol)
- 3) at 300K, a one litre solution of Sucrose (342 g/mol) was prepared by dissolving 40g of sucrose. What is the approximate Osmotic pressure (in Kpa) of solution at the same temperature  
( $R = 8.314 \times 10^6 \text{ cm}^3 \text{ Pa K}^{-1} \text{ mol}^{-1}$ ) (292)
- 4) at 10°C, the Osmotic pressure of urea solution was found to be 500 mmHg. The solution is diluted "x" times and the temperature raised to 25°C when the Osmotic pressure was noticed to be 105.3 mmHg, then "x" is. (5)
- 5) The Osmotic pressure of 5% aqueous solution of sugar (342 g/mol) at 15°C is — (3.45 atm)



- (6) Osmotic pressure of a solution is  $0.0821 \text{ atm}$  at a temperature of  $300 \text{ K}$ . The concentration in mole/lit will be —  $(0.3 \times 10^2)$
- (7) The Osmotic pressure of blood at  $27^\circ \text{C}$  is  $8.21 \text{ atm}$ . The amount of Glucose (in gm) should be used per litre for an intravenous injection so that it is isotonic with blood is  $(60 \text{ gm})$
- (8) The Osmotic pressure of blood is  $7.40 \text{ atm}$  at  $27^\circ \text{C}$ . Number of mole of glucose to be used per liter for an intravenous injection that is to have the same Osmotic pressure as blood is  $(0.3)$
- (9) Two solution of Glucose have Osmotic pressures  $1.5$  and  $2.5 \text{ atm}$ . If of first is mixed with  $2 \text{ L}$  of second solution, the Osmotic pressure in  $\text{atm}$  of the solution will be —  $(2.16 \text{ atm})$
- (10) The Osmotic pressure of solution containing  $4.0 \text{ g}$  of solute (molar mass  $246$ ) per litre at  $27^\circ \text{C}$  is —  $(R = 0.082 \text{ Latm K}^{-1} \text{ mol}^{-1})$   $(0.4 \text{ atm})$

Solutions  
chapter

Van't Hoff's factor - problems

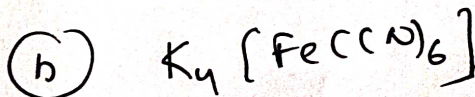
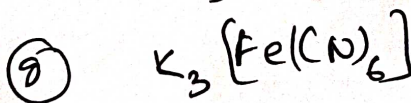
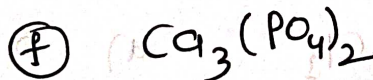
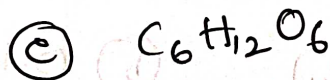
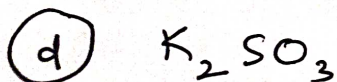
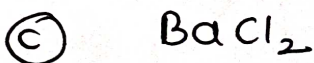
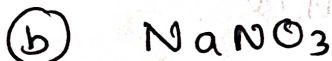
- ① If  $\text{BaCl}_2$  ionizes to an extent of 80% in aqueous solution, the value of Van't Hoff factor is — (2.6)
- ② a decimolar solution of  $\text{K}_4[\text{Fe}(\text{CN})_6]$  at 300K is 50% dissociated, then Osmotic pressure of the solution is (7.38 atm)
- ③ The experimental depression in freezing point of dilute solution is 0.025K. if the Van't Hoff factor is 2. the calculated depression in freezing point is — (0.0125K)
- ④ Which of the following aqueous solution has highest freezing point?
- a) 0.1M  $\text{Al}_2(\text{SO}_4)_3$       b) 0.1M  $\text{BaCl}_2$   
c) 0.1M  $\text{AlCl}_3$       d) 0.1M  $\text{NH}_4\text{Cl}$

(5) The freezing point of the solution containing 0.3 g of acetic acid (mass = 60 g/mol) in 30 g of benzene is lowered by  $0.42^\circ\text{C}$  then the Van't Hoff factor is ( $K_f = 5.12 \text{ K kg mol}^{-1}$ )

(6)  $\text{K}_2[\text{HgI}_4]$  is 40% ionised in aqueous solution. The value of its Van't Hoff factor is — (1.8).

(7) molal 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 of  $\text{K}_2\text{SO}_4$  is — (0.36 K).

(8) Find out  $i$  value for following



Find out moles & molecules

- ① 10g of  $\text{CaCO}_3$  = \_\_\_\_\_ moles = \_\_\_\_\_ molecules
- ② 0.4g of  $\text{NaOH}$  = \_\_\_\_\_ moles = \_\_\_\_\_ molecules
- ③ 1.1g of  $\text{CaCl}_2$  = \_\_\_\_\_ moles = \_\_\_\_\_ molecules
- ④  $49 \times 10^{-2}$  g of  $\text{H}_2\text{SO}_4$  = \_\_\_\_\_ moles = \_\_\_\_\_ molecules
- ⑤ 365g of  $\text{HCl}$  = \_\_\_\_\_ moles = \_\_\_\_\_ molecules
- ⑥ 5.85g of  $\text{NaCl}$  = \_\_\_\_\_ moles = \_\_\_\_\_ molecules

Find out mass & volume

- ⑦  $\frac{1}{4}$  moles of  $\text{CO}_2$  = \_\_\_\_\_ g ~~moles~~ = \_\_\_\_\_ lit
- ⑧ 2 moles of  $\text{CH}_4$  = \_\_\_\_\_ g = \_\_\_\_\_ lit
- ⑨ 10 moles of  $\text{CaCO}_3$  = \_\_\_\_\_ g = \_\_\_\_\_ lit
- ⑩ 0.6 moles of  $\text{KNO}_3$  = \_\_\_\_\_ g = \_\_\_\_\_ lit

Calculate molar masses & GEW

- ①  $H_2SO_4$
- ②  $HCl$
- ③  $HNO_3$
- ④  $H_2C_2O_4$
- ⑤  $H_2C_2O_4 \cdot 2H_2O$
- ⑥  $CH_3COOH$
- ⑦  $NaOH$
- ⑧  $KOH$
- ⑨  $Be(OH)_2$
- ⑩  $Mg(OH)_2$
- ⑪  $Ca(OH)_2$
- ⑫  $Al(OH)_3$
- ⑬  $Si(OH)_4$
- ⑭  $NaCl$
- ⑮  $KCl$
- ⑯  $MgCl_2$
- ⑰  ~~$MgCl_2$~~   
 $AlCl_3$
- ⑱  $Na_2CO_3$
- ⑲  $Mg(NO_3)_2$
- ⑳  $KNO_3$

WORK SHEET - I  
Solution

Topic: Solution  
( w/w %  
w/v %  
v/v % )

- 1 \* 20g of solute present in 180g of solvent  
find out its weight-weight percentage?
- 2 \* 150ml of Glucose present in 1250ml of  
solution find out v/v % ?
- 3 \* 10% (  $\frac{w}{v}$  ) of Sucrose means  
Solution.  
\_\_\_\_\_ g of Sucrose in \_\_\_\_\_ of water Solution.
- 4 \* 80g of NaOH present in 200g of solvent  
find out its w/w %.
- 5 \* 15% (  $\frac{w}{w}$  ) of NaCl present in 1000 (grams)  
of solution, what is the mass of NaCl \_\_\_\_\_
- 6 \* 75% (  $\frac{w}{v}$  ) of KOH solution has 25g of KOH.  
find out solution volume?