

## Some Basic Concepts of Chemistry.

Molecular Weight =

$$\text{NaCl} - 23 + 35.5 = 58.5$$

Atomicity  $\text{H}_2\text{SO}_4 - 7$

Mole Concept -

$$1 \text{ mole} - N_A - 6.023 \times 10^{23}$$

GAM, GMM:

$$\begin{aligned} \text{GAM of Oxygen} &= \text{mass of 1g atom of Oxygen} \\ &= \text{mass of 1 mole atom of Oxygen} \\ &= \left( \frac{16}{N_A} \text{ g} \times N_A \right) = 16 \text{ g} \end{aligned}$$

$$\text{GMM of } \text{H}_2\text{SO}_4 = \frac{98}{N_A} \times N_A = 98 \text{ g}$$

Formula:

$$\text{Mole} = \frac{\text{weight}}{\text{Mw}}$$

$$\checkmark \text{ Mole of molecules} = \frac{\text{weight}}{\text{GMMW}}$$

$$\text{Mole atoms} = \text{atomicity} \times \text{mole molecules}$$

... .. given Volume ( ... )

$$\text{Mole of gas} = \frac{\text{given Volume (STP)}}{22.4}$$

$$\text{Mole Particles} = \frac{\text{No. of Particles}}{N_A}$$

Cal. the no. of molecules and atoms present in 2g of nitrogen? [N=14]

$$\text{No. of moles} = \frac{2}{28} = \frac{1}{14}$$

$$\text{No. of molecules} = \frac{1}{14} \times N_A$$

$$\text{No. of atoms} = 2 \times \frac{N_A}{14} = \frac{N_A}{7} \text{ atoms}$$

Vapour density is Unitless

$$\text{Relative density} = \frac{\text{density of gas A}}{\text{density of gas B.}}$$

$$\text{Specific gravity} = \frac{\text{Density of Solution}}{\text{Density of water.}}$$

$(N_2)_x \Rightarrow$  V.P To  
?

$$\begin{aligned} \text{Mol. wt} &= 2 \times \text{V.P} \\ &= 2 \times 70 = 140 \end{aligned}$$

$$140 = x \times 28$$
$$\boxed{x = 5}$$

$$140 = x \dots$$

$$x = 5$$

$$\% \text{ mass} = \frac{\text{No. of atoms} \times \text{at. mass}}{\text{MM}} \times 100$$

$$n = \frac{\text{M.F}}{\text{Em.F}}$$

Eg:  $\text{MM} = 92$

		At. weight	$\frac{\%}{\text{At. we}}$	Simplest ratio
Carbon	52.2%	12	$\frac{52.2}{12} = 4.4$	$\frac{4.4}{2.2} = 2$
Hydrogen	13%	1	$\frac{13}{1} = 13$	$\frac{13}{2.2} = 5.9$
Oxygen	34.8%	16	$\frac{34.8}{16} = 2.2$	$\frac{2.2}{2.2} = 1$

$\text{C}_2\text{H}_6\text{O} \Rightarrow$  Empirical formula.

$$\text{Em. formula mass} = 12 \times 2 + 1 \times 6 + 16 \times 1 = \underline{46}$$

$$\frac{92}{46} = 2$$

$$\text{M.F} = 2(\text{em.F}) = 2 \times \text{C}_2\text{H}_6\text{O} = \text{C}_4\text{H}_{12}\text{O}_2$$

For Complete Combustion of 3g ethane. find required moles of  $\text{O}_2$  and formed moles of  $\text{CO}_2$

Volume of  $\text{O}_2$  &  $\text{CO}_2$  at STP.



Mass-Volume



$$\text{no. of moles} = \frac{3}{30 \text{ (M.w of ethane)}} = \frac{1}{10} = 0.1$$

$$\text{Required mole of } O_2 = 0.35 \text{ moles}$$

$$\begin{aligned} \text{Volume of } O_2 &= 0.35 \times 22.4 \\ &= 7.84 \text{ L} \end{aligned}$$

$$\begin{array}{r} * \quad 2 \quad 7 \\ 0.1 \quad \times \\ \hline 0.1 \times 7 \\ \hline 0.7 \end{array}$$

$$\text{Produced moles of } CO_2 = 0.2 \text{ moles}$$

$$\begin{aligned} \text{Volume of } CO_2 &= 0.2 \times 22.4 \\ &= 4.48 \text{ L} \end{aligned}$$

$$\begin{array}{r} 2 \quad 4 \\ 0.1 \quad \times \\ \hline 0.1 \times 4 \\ \hline 0.4 \end{array}$$

L.R  $\Rightarrow$  which will get Consumed first

Type-I Mass-Mass

How much iron can be ther. obtained in the reduction of 1 kg of  $Fe_2O_3$ ? M.w = 160 g/mol.



$$n = \frac{1000}{160} = \frac{100}{16} \text{ mol.}$$

$$\text{no. of Fe} = 2 \times \frac{100}{16} = \frac{100}{8} \text{ mol.} = 12.5 \text{ mol}$$

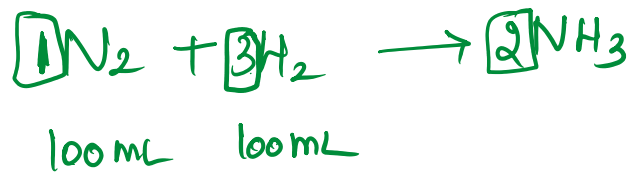
$$\text{Weight} = \text{no. of moles} \times \text{M.w}$$

$$= 12.5 \times 56 = 700 \text{ g} = 0.7 \text{ kg}$$

$$\begin{aligned} \text{weight} &= \text{no. of moles} \times \text{m.w.} \\ &= 12.5 \times 56 = 700 \text{ g} = 0.7 \text{ kg} \end{aligned}$$

Type-II Mass - Vol.

III  
100 mL N<sub>2</sub> 100 mL of H<sub>2</sub> mixed together to form NH<sub>3</sub>. Volume of NH<sub>3</sub> Produced.



$$\frac{100}{1} = 100 \quad \frac{100}{3} = \underline{\underline{33.3}} \quad \xrightarrow{\text{L.R.}}$$

$\frac{3}{100} \quad 2$

$$\begin{aligned} \text{For 100 mL of H}_2, \text{ Producing NH}_3 &= \frac{100 \times 2}{3} \\ &= \underline{\underline{66.6 \text{ mL}}} \end{aligned}$$

$$\text{Equivalent weight} = \frac{\text{At. weight}}{\text{Valency factor}}$$

Conc. Terms:

$$\text{Molarity (M)} = \frac{\text{No. of moles of Solute}}{\text{Volume of Solution (L)}}$$

$$\text{Molality (m)} = \frac{\text{No. of moles of Solute}}{\text{weight of Solvent (kg)}}$$

$$\text{Normality (N)} = \frac{\text{Mass of Solute (g)}}{\text{Equivalent mass} \times \text{Vol. of Solution (L)}}$$

$$\text{Mole fraction } X_A = \frac{\text{moles of Solute (A)}}{\text{moles of Solute (A) + moles of Solvent (B)}}$$

$$X_B = \frac{\text{moles of Solvent (B)}}{\text{moles of Solute (A) + moles of Solvent (B)}}$$

$$\boxed{X_A + X_B = 1}$$

$6.02 \times 10^{23}$  molecules of urea are dissolved in 1L of water. Cal. molarity (M) of soln if 40% of urea undergo dimerisation.



- a) 0.6 M    b) 0.7 M    c) 0.8 M    d) 0.9 M.

$$\text{Initial Molecules} = 6.02 \times 10^{23}$$

40% of Urea dimer.

$$\begin{aligned} \text{dimer molecule} &= 0.4 \times 6.02 \times 10^{23} / 2 \\ \text{remaining} &= 0.6 \times 6.02 \times 10^{23} \end{aligned}$$

Total molecule after dimerisation

$$= 0.6 \text{ NA} + 0.2 \text{ NA} = 0.8 \text{ NA}$$

$$\text{Molarity} = \frac{0.8 \text{ moles}}{1 \text{ (L)}} = 0.8 \text{ molarity}$$

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0.8M