# Analytical Chemistry

Mosul University / College of Science New & Renewable Energy Dep

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1<sup>st</sup> Class



# **General Introduction**

# **Define Chemistry:**

Chemistry is the science which deals with materials (elements or compounds) and their reactions.

- Classified the Chemistry:
- **1.** Analytical chemistry: is the science which deal with the analysis of substances qualitatively and quantitatively.
- 2. *Biochemistry:* interest with studying the bio compounds which exist inside the living organisms.
- 3. Organic chemistry: studying of carbon compounds and their reactions.
- 4. Inorganic chemistry: studying of all elements in periodic table except carbon.
- **5.** *Physical chemistry:* deals with physical properties substances during reaction such as: heat, solubility, and conductivity...etc.
- 6. *Industrial chemistry:* this branch deals with manufacturing of materials and their use in our life.

# **Analytical Chemistry**

#### **Define Analytical Chemistry**

Is a branch of chemistry science which defined as the means that are det ected elements and materials and the method of separated and know the co mpounds of those substances in a mixture of them in addition to determined of those components quantitatively.

#### **Classification of Analytical Chemistry**

The following scheme for the classification of analytical chemistry.





#### **Classification of Analytical Chemistry**





- Analytical chemistry deals with separating, identifyi ng, and quantifying the relative amounts of the com ponents of an analyte.
- *Analyte* : the thing to analyzed; the component(s) of a sample that are to be determine





Figure 1-1 The relationship between analytical chemistry, other branches of chemistry, and the other sciences. The central location of analytical chemistry in the diagram signifies its importance and the breadth of its interactions with many other disciplines.



# Steps of Analysis:

1. The aim of analysis (determination, identification, separation)

- 2. Select the analysis method (which depends on: Accuracy in analysis, time, amount of sample to be determined.
- 3. Prepare the sample.
- 4. Using separation technique if it's necessary.

5. Analysis.

6. Results and discussion.









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## **Solutions**

A homogeneous mixture of two or more of substances. That is mean: overlapping molecules or ions of solute between molecules or ions of solvent, the product called *Solution*.

#### *Solution = Solute + Solvent*

There are different types of solution:



## A. According to the size of atoms or molecules:

- **1.** *Real Solution*: The solution which it's molecules pass through the filter paper fully and easily.
- 2. Suspension solution: In this solution some of the molecules don't pass through the filter paper fully and easily.
- **3.** *Colloidal solution*: In this solution the molecules don't pass through the filter paper as well as don't precipitate if let the solution to stable.

# **B.** According to the concentration of solute:

- Unsaturated Solution (Normal solution):

The solution which can added more amount of solute in it at room temperature.

- *Saturated Solution:* The solution which contain the solute in dynamic equilibrium state with solvent at room temp.
- *Super Saturated Solution:* The solution that contains more amount of solute than those found in saturated solution at high temperature when return to room temperature some of the solute will be precipitate.

#### **Concentrated Solution:**

That is mean the solution contains amount of solute.

#### **Diluted Solution:**

It is opposite the concentrated solution that is mean the solution contains less amount of solute.





## C. According to the Nature of Solvent:

Liquid Solution , Organic Solution

How can change concentrated solution to the diluted one :





## **Standard Solutions:**

Is a reagent of exactly known concentration which come from dissolve exactly weight of primary standard material in known volume of solvent.

There are two types of standard materials:





#### I. Primary Standard Material:

Is a highly purified compound that serves as reference material for a titrimetric method of analysis and which can directly prepared primary standard solution.

Such as: Na<sub>2</sub>CO<sub>3</sub>, KCl, NaCl, Na<sub>2</sub>SO<sub>4</sub>, MgSO<sub>4</sub>.7H<sub>2</sub>0...ect.

#### **II. Secondary Standard Material:**

Is a compound whose purity has been established by chemical analysis that serves as the reference material for a titrimetric method analysis. And can't prepared primary standard directly from these compound unless standardization it .

Such as: *NaOH*, *H*<sub>2</sub>*SO*<sub>4</sub>, *KOH*, *HCl*, *KMnO*<sub>4</sub>...ect..



Important requirements for a primary standard material are the following:

- 1. High purity establish methods for confirming purity should be available.
- 2. Atmospheric stability.
- 3. Absence of hydrate water so that the confirming of the solid does not change with variations in humidity.

4. Modest cost.

- 5. Reasonable solubility in the titration medium.
- 6. Reasonably large molar mass so that the relative error associated with weighing the standard is minimized.





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# **Aqueous Solution and Chemical Equilibria**

*Electrolysis:* are solutes which are ionized in solution to produce an electrically conducting medium. There are two types of electrolytes.

Strong electrolytes: the compounds which are completely ionized or dissociate in solution.  $HCl \rightarrow H^+ + Cl^-$ 

Weak electrolytes: the compounds which are partially dissociate in solution.

 $CH_3COOH = CH_3COO^- + H^+$ 

**Conjugate Acids and Bases** 

Acid<sub>1</sub>  $\longrightarrow$  Base<sub>1</sub> + H<sup>+</sup> (Conjugate base of the parent acid)

 $Base_2 + H^+ \longrightarrow Acid_2 \qquad (Conjugate acid of the parent base)$ 

The result in an acid /base or neutralization reaction :

#### $Acid_1 + Base_2 = Base_1 + Acid_2$

A conjugate acid: is formed when a base accepts aproton. A conjugate base: is formed when an acid losses a proton. e.g. acetate ion is conjugate base of acetic acid. **Neutralization Reaction:** 

 $HCl + NaOH \longrightarrow NaCl + H_2O$   $HCl \longrightarrow H^+ + Cl^ NaOH \longrightarrow Na^+ + OH^ NH_3 + H_2O \longrightarrow NH_4^+ + OH^ Conjugated of NH_3 Conjugated base of H_2O$ 

 $NH_3$ ,  $NH_4^+$  are conjugate pair.



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# **Chemical Equilibrium**

Many reaction used in analytical Chemistry never result in complete conversion of reactants to products.

Instead, they proceed to a state of chemical equilibrium that describe the concentrations of reactants existing among reactants product is constant. Equilibrium constant expressions are algebraic equation that describe the concentration relationships existing among reactants and products at equilibrium.



# **Equilibrium-** Constant Expressions:

A generalized equation for a chemical equilibrium is:

# $aA + bB \longrightarrow cC + dD$

where the capital letters represent the formulas of participating chemical species and the lower case are the small whole numbers required to balance the equation.

a,b,c,d = mole of A,B,C,D.

The equilibrium –constant expression of the above reaction is :

$$\boldsymbol{K}_{eq.} = \frac{[C]^{c}[D]^{d}}{[A]^{a}[B]^{b}}$$



 $S_2O8 + 3I^- \implies 2SO_4^{-2} + I_3^-$ 

#### H.W

*Ex.2.* :

Calculate the concentration of each of A,B of equilibrium state of 0.1 M AB solution (AB: weak electrolyte).  $K_{eq} = 3 \times 10^{-6}$ .

Х

 $AB \longrightarrow A^+ + B^-$ 0.1-X Х  $Keq. = \frac{[A^+][B^-]}{[AB]}$  $3 \times 10^{-6} = \frac{X^2}{0.1 - X}$ Neglect  $\sqrt{X^2} = \sqrt{3 \times 10^{-7}}$  $X = 5.5 \times 10^{-4}$  M

So  $[A^+] = [B^-]$ at equilibrium



## Solubility Product Constant

*Solubility:* is the amount of solute can be dissolved in a volume of solvent.

When one substance (solute) dissolves in any solvent, it is said to be *soluble*. When one substance does not dissolve in another it is said to be *insoluble*. Solubility depends on type of solvent, temperature and acidity (pH) of media.

The solubility can be determined by solubility product constant  $(K_{SP})$ .

In saturated solution of AxBy salt.

$$A_x B_y = x A^+ + y B^-$$

 $\mathbf{K} = \frac{[A^+]^x [B^-]^y}{[A_x B_y]}$ 

at equilibrium state

:: Conc. of solid compound is constant

 $\therefore \mathbf{K} = [\mathbf{A}^+]^{\mathbf{x}} \ [\mathbf{B}^-]^{\mathbf{y}}$ 

For  $AgCl_{(s)} \longrightarrow Ag^+ + Cl^ \therefore Ksp = [Ag^+] [Cl^-]$ For  $CaF_2 \longrightarrow Ca^{+2} + 2F^ \therefore Ksp = [Ca^{+2}] [F^-]^2$ 

#### *Ex.1*

Calculate the amount of solubility in mole/liter of AgCl?  $K_{sp} = 1.8 \times 10^{-10}$ . *H.W* 

#### *Ex.2*

The amount of solubility of  $BaSO_4$  is  $1.14 \times 10^{-5}$ M.Calculate the solubility product constant?

 $BaSO_4 \implies Ba^{+2} + SO_4^{-2}$ Ksp =  $[Ba^{+2}] [SO_4^{-2}]$   $[Ba^{+2}] = 1.14 \times 10^{-5} \text{ M} = [SO_4^{-2}]$ Ksp =  $[1.14 \times 10^{-5}] [1.14 \times 10^{-5}]$ Ksp =  $1.30 \times 10^{-10} \text{ M}$ 



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# **Dissociation Constant**

# Applying the Ion- Product for Water

Aqueous solutions contain small concentrations of hydronium and hydroxide ion as a consequence of the dissociation.

# $2H_2O \longrightarrow H_3O^+ + OH^-$ or $H_2O \longrightarrow H^+ + OH^-$



$$K = \frac{[H_3 O^+] [OH^-]}{[H_2 O]^2}$$

 $K[H_2O]^2 = Kw = [H_3O^+][OH^-]$ At: 25C°  $K_w = 1 \times 10^{-14}$ 

At pure  $H_2O$   $[H^+] = [OH^-] = 10^{-7}$ 

 $pK_w = pH + pOH = 14$ 

 $-\log[H^+] = -\log[OH^-] = 7$ pH = pOH = 7



# **Example:**

Calculate the hydrogen or hydronium ion concentration in solution contain  $[OH^-]$  (a). 0.01mole/L, (b)  $2 \times 10^{-9}$  mole/L, and describe the solution?

 $K_w = [H^+] [OH^-] = 1 \times 10^{-14}$ 

(a). 
$$[H^+] = \frac{Kw}{[OH^-]} \rightarrow [H^+] = \frac{1 \times 10^{-14}}{0.01}$$
  
 $[H^+] = 1 \times 10^{-12} \text{ M} , \text{ pH} = -\log [H^+] \rightarrow \text{pH} = -\log 1 \times 10^{-12} , \text{ pH} = 12$ 

. The solution is Alkaline  $[OH^-] > [H^+]$ 

(b). 
$$[H^+] = \frac{1 \times 10^{-14}}{2 \times 10^{-9}}$$
  
 $[H^+] = 5 \times 10^{-6} \text{ M}$ ,  $pH = -\log [H^+] \rightarrow pH = -\log 5 \times 10^{-6}$ ,  $pH = 5.3$ 

• The solution is Acidic  $[H^+] > [OH^-]$ 

Dissociation of weak acid and weak base:

### $CH_3COOH \longrightarrow CH_3COO^{-} + H^+$

 $pK_w = pH + pOH$  pH = pKw - pOH  $pH = \frac{1}{2} pKa - \frac{1}{2} \log C$   $pOH = \frac{1}{2} pKb - \frac{1}{2} \log C$   $pH = pKw - \frac{1}{2} pKb + \frac{1}{2} \log C$  $pOH = pKw - \frac{1}{2} pKa + \frac{1}{2} \log C$ 


# Example

Calculate the pH value at 25C° of (0.1M) solution (of weak base) tri methyl amine  $(CH_3)_3 N$ ? Kb =  $5.3 \times 10^{-5}$ .

```
pH = pKw - \frac{1}{2} pKb + \frac{1}{2} \log C

pH = 14 - \frac{1}{2}(-\log Kb) + \frac{1}{2} \log 0.1

pH = 14 - \frac{1}{2} (-\log 5.3 \times 10^{-5}) + \frac{1}{2} \log 0.1

pH = 14 - (2.14) + \frac{1}{2} (1)

pH = 11.36
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There are *Two* types of the methods for expressing the concentration:

First: Physical methods for expressing concentration:

I. Percentage %:

#### A. Weight percent : W/W %

The number of grams of solute in 100 gram of solution.

 $W/W \% = \frac{Weight (wt)solute (g)}{weight (wt) solute (g)} \times 100$ 

Weight solution would be more properly called mass percent and abbreviated m/m.

**B.** Volume percent : V/V %

The number of milliliters of solute in 100 milliliter of solution.

 $V/V \% = \frac{Volume \ solute \ (mL)}{Volume \ solution \ (mL)} \times 100$ 

*Volume of solution = V. of solute + V. of solvent* 

#### C. Weight – Volume percent : W/V%

The number of grams of solute in 100 mL of solution. It's used for two miscible and diluted liquids.

 $W/V \% = \frac{Weight \ solute \ (g)}{Volume \ solution \ (mL)} \times 100$ 

#### Example 1.:

How many grams of NaCl needed to prepare 500mL of normal saline solution in concentration W/V=70%?

#### *H.W*

#### Example 2.

Calculate the weight percent of solution prepared by dissolve 5g of  $AgNO_3$  in 100 mL of water? Note; water density =1.

 $W/W \% = \frac{Weight (wt)solute (g)}{weight (wt) solute (g)} \times 100$  $Mass = Density \times Volume$  $Mass = 1 \times 100 = 100$ 

Weight of solution = Wt.of Solute + Wt.of Solvent Weight of solution = 5 + 100 = 105

$$W/W \% = \frac{5}{105} \times 10$$
$$W/W \% = 4.76 \%$$

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#### Methods of Expressing Concentration:

Concentration: The amount of solute located in a certain solvent or solution.

Mole: Is Avokadro's number of particles. Particles: ions, electrons, atoms, molecules.

$$Mole = \frac{Weight(w)}{Molecular weight(M.wt.)} = \frac{W}{M.wt.}$$

Molecular weight = (M.wt): is a total of atomic weight in the compound.

$$\mathbf{M.wt.} = \frac{W}{Mole} \quad \text{g / mole}$$

#### **Example 1**

Calculate the M.wt. of benzoic acid  $C_7H_6O_2$ ? A.wt.: C=12, H= 1, O= 16

H.W





#### Example 2

How many grams of sodium ion Na<sup>+</sup> are contained in 20g. of sodium carbonate Na<sub>2</sub>CO<sub>3</sub> A.wt.: Na = 23, C=12, O = 16.

Mole of 
$$(Na_2CO_3) = \frac{wt}{M.wt} = \frac{20}{106} = 0.188$$
 Mole of  $Na_2CO_3$ 

$$Na_2CO_3 \rightarrow 2 Na^+ + CO_3^{-2}$$

Mole (Na<sup>+</sup>) = 2 × 0.188 = 0.376 mole Mole =  $\frac{wt}{M.wt}$  = 0.376 =  $\frac{wt}{23}$ Wt, = 8.679 g. *Or* 

Na	<u>Na<sub>2</sub>CO<sub>3</sub></u>	3
A.wt. $\times$ no. of mole	 M.wt	<u> </u>
$23 \times 2$	106	
g	20	
Lamya Sarsam Wt. = $\frac{20 \times 46}{106}$	= 8.679 g	



## Specific gravity (Sp.gr.):

#### It's the ratio between density of substance to the density of water

 $Sp.gr. = \frac{Density \ of \ Substance}{Density \ of \ Water}$ 

Density of Water = 1 g/L

$$d = \text{Sp.gr.} = \frac{1}{1}$$
$$= 1$$

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## II. Parts per thousand, Parts per million and Parts per billion:

 $Cppt = \frac{Wt.g}{V.mL} \times 10^3$ 

(which used specially in oceanography).

For very dilute solution part per million (ppm) is a convenient way to express concentration.

 $Cppm = \frac{Wt.g}{V.mL} \times 10^6$ 

For aqueous solution.

 $ppm = 1.0 mg/L \equiv 1.0 \mu g / mL$  $Cppb = \frac{Wt.g}{V.mL} \times 10^{9}$ 

#### **Example:**

Calculate the number of grams of KCl to prepare solution in volume 500mL, the concentration of KCl in the solution is 250 ppm?

H.W

#### **III.** P- Function:

Sometimes it's suitable to expressing concentration of ion by logarithm. The most well known p-function is pH which it is negative logarithm of [H<sup>+</sup>]. (H<sup>+</sup> concentration).

 $pH = -log [H^+]$  $pH = -\log[OH^{-}]$  $pH = -\log[X^{-}]$ Example: NaCl  $\rightarrow$  Na<sup>+</sup> + Cl<sup>-</sup> 0.1 M 0.1 M 0.1 M  $pNa^{+} = -\log [Na^{+}]$ Lamya Sarsanlog [0.1]

 $0.1 = 10^{-1}$ 

# Second: Chemical Methods for Expressing concentration

**I.** Molarity (M): Is the total number of moles of a solute in 1L of solution. Or the total number of millimoles in 1 mL.

 $M = \frac{no.moles (solute)}{V. Soln.L}$ 

No. = number.

Soln. = Solution.



 $M = \frac{W \times 1000}{M.Wt. \times VmL}$ 

This law used for *solid state* material

*mole/L, mmol/mL*  $\rightarrow$  *Molar* 

No. of moles =  $M \times V_{(L)}$ No. of millimoles =  $M \times V_{(mL)}$ 

 $M = \frac{Sp.gr. \times \% \times 10}{M.Wt.}$  Molarity of <u>*liquid state*</u> solution

Specific gravity :

 $Sp.gr. = \frac{Density \, of \, Substance}{density \, of \, Water}$ 

: Density of water (H<sub>2</sub>O)  $d(H_2O) = 1$ Sp.gr. = d

. .

### **Dilution Law**

Number of moles of concentration solution = number of moles of dilution solution.

Conc. Soln.		<u>dil. Soln,</u>
no. of moles	=	no. of moles
no. of millimoles	=	no. ofmillimoles

 $M_1 \cdot V_1 = M_2 \cdot V_2$ 

**Ex.1.** Prepare 0.1M of calcium carbonate  $CaCO_3$  in 2 litter?

A.wt. : Ca = 40, C= 12, O = 16.

 $M = \frac{W \times 1000}{M.Wt. \times VmL}$ 1L = 1000 mL

$$0.1 = \frac{W \times 1000}{100 \times 2000}$$

 $W = 20 g of CaCO_3 in 2 litter$ 

*Ex.2.* Calculate the volume of  $NH_3$  conc. which used to prepare 500 mL of 0.1 M  $NH_3$ , sp.gr. of concentrated solution = 0.9 and the percentage = 28%, M.wt. = 17.?

$$M = \frac{Sp.gr.\times\%\times10}{M.Wt.}$$

$$M = \frac{0.9.\times28\times10}{17}$$

$$M = 14.8 \text{ M of } \text{NH}_3 \text{ conc.}$$

$$M_1 V_1 = M_2 V_2$$

$$14.8 \times V_1 = 0.1 \times 500$$

$$V_1 = \frac{0.1.\times500}{14.8}$$

$$V_1 = 3.33 \text{ mL dilute to } 500 \text{ mL by } \text{D.W to give } 0.1 \text{ M NH}_3$$



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**II.** Formality (F): It is number of formula weight solute in liter of solution.

$$F = \frac{W \times 1000}{F.Wt. \times VL} \qquad F = F.w/L, m.Fw./mL \rightarrow Formal$$

III. Normality (N): It is the number of equivalents of solute in liter of solution.

 $N = \frac{no.of \ equivalent}{Vol.of \ solution \ (L)}$ 

no.eq. =  $\frac{Wt.}{eq.Wt.}$  ....

The equivalent weight (eq.wt.) of a substance is not a constant quantity, but it's value depend upon the reaction, in which it is taken part.

Lamya Sarsam Since;  $Eq.wt = \frac{Mwt}{n}$ 

n : the reacting units



For solid state material.

$$N = eq/L, m.eq./mL \rightarrow Normal$$

$$N = \frac{Sp.gr.\times\%\times10}{eq.wt.}$$

For *liquid state* solution.





## To calculate the equivalent weight (eq.wt.):

$$eq.wt. = \frac{M.wt.}{n}$$

n =active unite.

- $n = H^+$  (acids).
- $n = OH^{-}$  (bases).
- $n = charge \times number of ions (salt).$
- n = no. of electrons lost or gained (oxidation –reduction).

# Calculate the equivalent weight:

A. Of Element

 $eq.wt. = \frac{A.wt.}{no.of \ oxidant}$ 

**Ex.** What is the eq.wt. of Mg? A.wt. = 24

Eq.wt. = 
$$\frac{A.wt.}{no.of \text{ oxidant}}$$
  
=  $\frac{24}{2}$   
= 12





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*Ex.1*. Calculate the equivalent weight of HCl? M.wt. = 36.5

$$HCl \rightarrow H^+ + Cl^-$$

eq.wt. of HCl =  $\frac{M.wt.}{no.of \text{ proton replacable of base}}$ eq.wt. of HCl =  $\frac{36.5}{1}$ = 36.5

*Ex.2.* Calculate the equivalent weight of  $H_2SO_4$ ? M.wt. = 98.

$$H_2SO_4 \rightarrow 2H^+ + SO_4^-$$

eq.wt. of  $H_2SO_4 = \frac{M.wt.}{no.of proton replacable of base}$ 

eq.wt. of 
$$H_2SO_4 = \frac{98}{2}$$
  
= 49

So,  $H_2SO_4$  has two reacting units of proton ; there are two equivalents of proton in each mole. While HCl has one reacting unit of proton, there is one equivalent of proton in each mole. So the normality of  $H_2SO_4$  solution is twice its molarity.

 $N_{of H2SO4} = M \times 2$ 

C. Of Base

# $eq.wt. = \frac{M.wt.}{no.of \ reactive hydroxil \ groups}$

*Ex.1*. Calculate the equivalent weight of NaOH? M.wt. = 40

eq.wt. of NaOH =  $\frac{M.wt.}{no.of reactive hydroxil groups}$ eq.wt. of NaOH =  $\frac{40}{1}$ = 40

*Ex.2.* Calculate the equivalent weight of  $Mg(OH)_2$ ? M.wt.58

eq.wt. of  $Mg(OH)_2 = \frac{M.wt.}{no.of reactive hydroxil groups}$ 

Lamya Sarsaeq.wt. of Mg(OH)<sub>2</sub> = 
$$\frac{58}{2}$$
 = 29



# D. Of Salt $eq.wt. = \frac{M.wt.}{number of metal atoms \times no.of charge or no.of oxidant}$

*Ex.:* Calculate the eq.wt. of  $Na_2CO_3$ ?M.wt. = 106

eq.wt. of Na<sub>2</sub>CO<sub>3</sub> = 
$$\frac{M.wt}{2\times(+1)}$$

$$= \frac{106}{2\times(+1)}$$
$$= 53$$




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### E. Of material that suffer oxidation reduction

 $eq.wt. of oxidation = \frac{M.wt.}{number of loss electrons}$ 

M.wt.  $eq.wt. of reduction = \frac{m.wt.}{number of gain electrons}$ 

**Ex.:** Calculate the eq.wt. of manganese  $Mn^{+2}$  and ferrous  $Fe^{+2}$  in the equation below?

 $MnO_{4}^{-} + Fe^{+2} + 8H^{+} + 5e^{-} \rightarrow Mn^{+2} + Fe^{+3} + 4H_{2}O$ 

eq.wt. of  $Mn^{+2} = \frac{M.wt.}{number of gain electrons}$ eq.wt. of  $Mn^{+2} = \frac{M.wt.}{5}$ eq.wt. of  $Fe^{+3} = \frac{M.wt.}{number of loss electrons}$ Eq. wt. of  $Fe^{+3} = \frac{M.wt.}{I}$ 

### F. Of Complex Formation Reaction

 $eq.wt. = \frac{A.wt.}{no.of \ oxidant \ of \ ion \ which \ contact \ with \ ligand}$ 

*Ex.*: Calculate the eq.wt. of Nickle  $Ni^{+2}$  in the equation below?

 $Ni^{+2} + 2DMG \rightarrow Ni (DMG)_2 \downarrow$ 

$$\begin{array}{c} CH_3 - CH - N - OH \\ DMG: \qquad | \\ CH_3 - CH - N - OH \end{array}$$

eq.wt. =  $\frac{\text{At.wt.}}{\text{no.of oxidant of ion which contact with ligand}}$ 

eq.wt. Ni = 
$$\frac{\text{At.wt.}}{2}$$



• What is the relationship between Molarity or Normality with part per million ppm?

$$C ppm = \frac{Wt.g}{V.mL} \times 10^{6}$$
$$M = \frac{W \times 1000}{M.Wt. \times VmL}$$

#### Multiple denominator and numerator by 10<sup>6</sup>

$$M = \frac{W \times 1000}{M.Wt.\times VmL} \times \frac{10^{6}}{10^{6}}$$
$$M = \frac{ppm \times 1000}{M.Wt.\times 10^{6}}$$
$$M = \frac{ppm}{M.Wt.\times 1000}$$
$$\therefore ppm = M \times M.wt. \times 1000$$
$$ppm_{Max} = N \times eq.wt. \times 1000$$





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**IV. Molality (m):** It is number of moles of solute in kilogram of solvent. Or no. of millimoles of solute in (g) of solvent.

 $m = \frac{W \times 1000}{M.wt. \times W \text{ solvent } kg}$ 

In dilute aqueous solution:

M = m

Weight of solution = Weight of solvent = Volume of solution.

Molar Fraction (X):

It's the ratio between numbers of solute moles to

number of moles of all solution contains.

Mole of Fraction of Solute(X) =  $\frac{\text{no.moles of(solute)}}{\text{no.moles of (solution)}}$ 

no. of moles of solution = no. of solute moles + no. of solvent moles

Mole of Fraction of  $solvent(X) = \frac{no.moles of (solvent)}{no.moles of (solution)}$ 

X solute + X solvent = 1

**Ex.:** Calculate the mole fraction of 5.8g acetone solution  $CH_3 - C - CH_3$  with 90g water? A.wt.: C = 12, O = 16, H = 1Mole of Fraction of Solute<sub>Acetone</sub>(X) =  $\frac{no.moles of(solute)}{no.moles of(solution)}$ Moles (solute) =  $\frac{Wt}{M_{.}wt}$ 5.8 *g*  $=\overline{(3\times12)+(1\times16)+(6\times1)}=58g/mol$  $Mole_{(solute)} = 0.1$  mole  $Mole_{(solvent)} = \frac{90}{18} = 5$  mole no.of solution moles = 5.0 + 0.1 = 5.1 mole

Mole Fraction of Solute =  $\frac{no.moles of(solute)}{no.moles of (solution)}$ 

X solute 
$$=\frac{0.1}{5.1}$$
  
X = 0.0196

Mole Fraction of Solvent = 
$$\frac{5.0}{5.1}$$
  
X = 0.9804

Total Mole Fraction = 0.9804 + 0.0196

= 1.0

