

REDOX REACTION

Oxidation & Reduction

Let us do a comparative study of oxidation and reduction :

Oxidation

1. Addition of Oxygen e.g. $2\text{Mg} + \text{O}_2 \rightarrow 2\text{MgO}$
2. Removal of Hydrogen e.g. $\text{H}_2\text{S} + \text{Cl}_2 \rightarrow 2\text{HCl} + \text{S}$
3. Increase in positive charge e.g. $\text{Fe}^{2+} \rightarrow \text{Fe}^{3+} + \text{e}^-$
4. Increase in oxidation number
 (+2) (+4)
 e.g. $\text{SnCl}_2 \rightarrow \text{SnCl}_4$
5. Removal of electron e.g. $\text{Sn}^{2+} \rightarrow \text{Sn}^{4+} + 2\text{e}^-$

Reduction

1. Removal of Oxygen e.g. $\text{CuO} + \text{C} \rightarrow \text{Cu} + \text{CO}$
2. Addition of Hydrogen e.g. $\text{S} + \text{H}_2 \rightarrow \text{H}_2\text{S}$
3. Decrease in positive charge e.g. $\text{Fe}^{3+} + \text{e}^- \rightarrow \text{Fe}^{2+}$
4. Decrease in oxidation number
 (+7) (+2)
 e.g. $\text{MnO}_4^- \rightarrow \text{Mn}^{2+}$
5. Addition of electron e.g. $\text{Fe}^{3+} + \text{e}^- \rightarrow \text{Fe}^{2+}$

Oxidation Number

- It is an imaginary or apparent charge developed over atom of an element when it goes from its elemental free state to combined state in molecules.
- It is calculated on basis of an arbitrary set of rules.
- It is a relative charge in a particular bonded state.
- In order to keep track of electron-shifts in chemical reactions involving formation of compounds, a more practical method of using oxidation number has been developed.
- In this method, it is always assumed that there is a complete transfer of electron from a less electronegative atom to a more electronegative atom.

Rules governing oxidation number

The following rules are helpful in calculating oxidation number of the elements in their different compounds. it is to be remembered that the basis of these rule is the electronegativity of the element.

- **Fluorine atom :**
Fluorine is most electronegative atom (known). It always has oxidation number equal to -1 in all its compounds
- **Oxygen atom :**
In general and as well as in its oxides, oxygen atom has oxidation number equal to -2 .

In case of

- (i) peroxide (e.g. H_2O_2 , Na_2O_2) is -1 ,
- (ii) super oxide (e.g. KO_2) is $-1/2$
- (iii) ozonide (e.g. KO_3) is $-1/3$
- (iv) in OF_2 is $+2$ & in O_2F_2 is $+1$
- **Hydrogen atom :**
In general, H atom has oxidation number equal to $+1$. But in metallic hydrides (e.g. NaH , KH), it is -1 .
- **Halogen atom :**
In general, all halogen atoms (Cl, Br, I) have oxidation number equal to -1 .

Calculation of individual oxidation number

It is important to note that to calculate individual oxidation number of the element in its compound one should know the structure of the compound and use the following guidelines.

Formula :

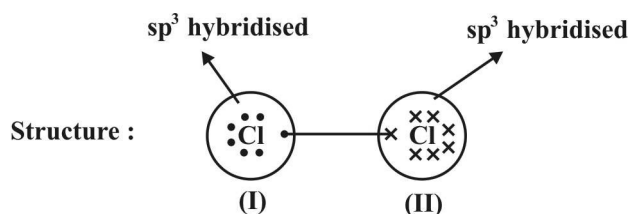
Oxidation Number = Number of electrons in the valence shell - Number of electrons taken up after bonding

Guidelines : It is based on electronegativity of elements.

- If there is a bond between similar type of atom and each atom has same type of hybridisation, then bonded pair electrons are equally shared by each element.

Example :

Calculate oxidation number of each Cl-atom in Cl_2 molecule



I : Number of electrons in the valence shell = 7
 Number of electrons taken up after bonding = 7.
 \therefore oxidation number = $7 - 7 = 0$.

II : similarly, oxidation number = $7 - 7 = 0$

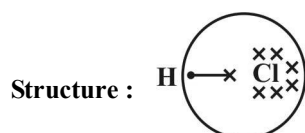
- If there is a bond between different type of atoms :

e.g. A – B (if B is more electronegative than A)

Then after bonding, bonded pair of electrons are counted with B-atom

Example :

Calculate oxidation number of each atom in HCl molecule



Note : Electron of H-atom is now counted with Cl-atom, because Cl-atom is more electronegative than H-atom

H : Number of electrons in the valence shell = 1
 Number of electrons taken up after bonding = 0
 Oxidation number of H = $1 - 0 = +1$

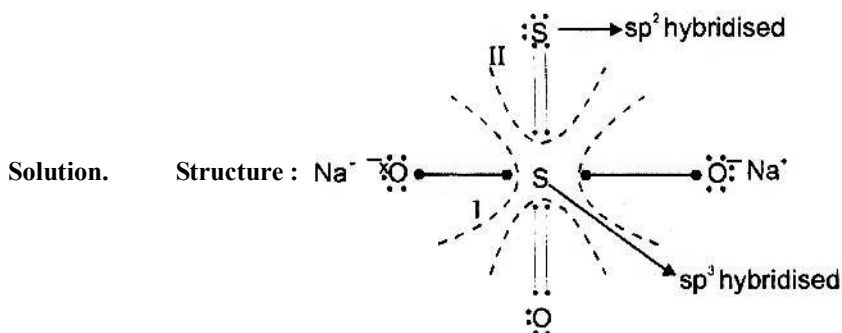
Cl : Number of electrons in the valence shell = 7
 Number of electrons taken up after bonding = 8
 Oxidation number of Cl = $7 - 8 = -1$

REDOX REACTION

Solved Examples

Example - 2

Calculate individual oxidation number of each S-atom in $\text{Na}_2\text{S}_2\text{O}_3$ (sodium thiosulphate) with the help of its structure.



Note : I (central S-atom) is sp^3 hybridised (25% s-character) and II (terminal S-atom) is sp^2 hybridised (33% s-character). Therefore, terminal sulphur atom is more electronegative than central sulphur atom. Now, the shared pair of electrons are counted with terminal S-atom.

\therefore I, S-atom : Number of electrons in the valence shell = 6

Number of electrons left after bonding = 0

Oxidation number of central S-atom = $6 - 0 = +6$

II, S-atom : Number of electrons in the valence shell = 6

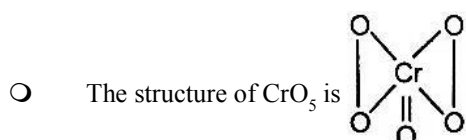
Number of electrons left after bonding = 8

Oxidation number of terminal S-atom = $6 - 8 = -2$

Now, you can also calculate Average Oxidation number of S = $\frac{6 + (-2)}{2} = +2$ (as we have calculated before)

Miscellaneous Examples :

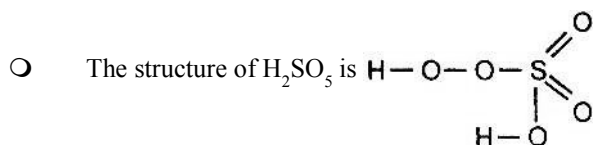
In order to determine the exact or individual oxidation number we need to take help from the structures of the molecules. Some special cases are discussed as follows :



From the structure, it is evident that in CrO_5 there are two peroxide linkages and one double bond. The contribution of each peroxide linkage is -2 . Let the oxidation number of Cr is x .

$\therefore x + (-2) \cdot 2 + (-2) = 0$ or $x = 6$

\therefore Oxidation number of Cr = $+6$ **Ans.**



From the structure, it is evident that in H_2SO_5 , there is one peroxide linkage, two sulphur-oxygen double bonds and one OH group. Let the oxidation number of S = x .

$$\therefore (+1) + (-2) + x + (-2) 2 + (-2) + 1 = 0$$

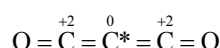
$$\text{or } x + 2 - 8 \quad \text{or } x - 6 = 0 \quad \text{or } x = 6$$

\therefore Oxidation number of S in H_2SO_5 is + 6 Ans.

Paradox of fractional oxidation number

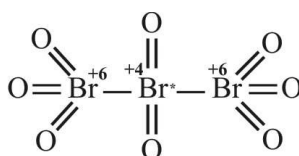
Fractional oxidation number is the average of oxidation state of all atoms of element under examination and the structural parameters reveal that the atoms of element for whom fractional oxidation state is realised are actually present in different oxidation states. Structure of the species C_3O_2 , Br_3O_8 and $\text{S}_4\text{O}_6^{2-}$ reveal the following bonding situations :

- The element marked with asterisk (*) in each species is exhibiting different oxidation number from rest of the atoms of the same element in each of the species. This reveals that in C_3O_2 , two carbon atoms are present in +2 oxidation state each whereas the third one is present in zero oxidation state and the average is +4/3. However, the realistic picture is +2 for two terminal carbons and zero for the middle carbon.



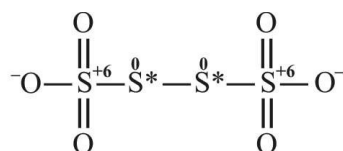
Structure of C_3O_2 (Carbon suboxide)

- Likewise in Br_3O_8 , each of the two terminal bromine atoms are present in +6 oxidation state and the middle bromine* is present in +4 oxidation state. Once again the average, that is different from reality, is + 16/3.



Structure of Br_3O_8 (Tribromooxide)

- In the same fashion, in the species $\text{S}_4\text{O}_6^{2-}$, average oxidation number of S is + 2.5, whereas the reality being + 5, 0*, 0* and +5 oxidation number respectively for respective sulphur atoms.



Structure of $\text{S}_4\text{O}_6^{2-}$, (tetrathionate ion)

In general, the conclusion is that the idea of fractional oxidation state should be taken with care and the reality is revealed by the structures only.

Oxidising and reducing agent

- **Oxidising agent or Oxidant :**

Oxidising agents are those compounds which can oxidise others and reduce itself during the chemical reaction. Those reagents in which for an element, oxidation number decreases or which undergoes gain of electrons in a redox reaction are termed as oxidants.

e.g. KMnO_4 , $\text{K}_2\text{Cr}_2\text{O}_7$, HNO_3 , conc. H_2SO_4 etc are powerful oxidising agents.

REDOX REACTION

○ Reducing agent or Reductant :

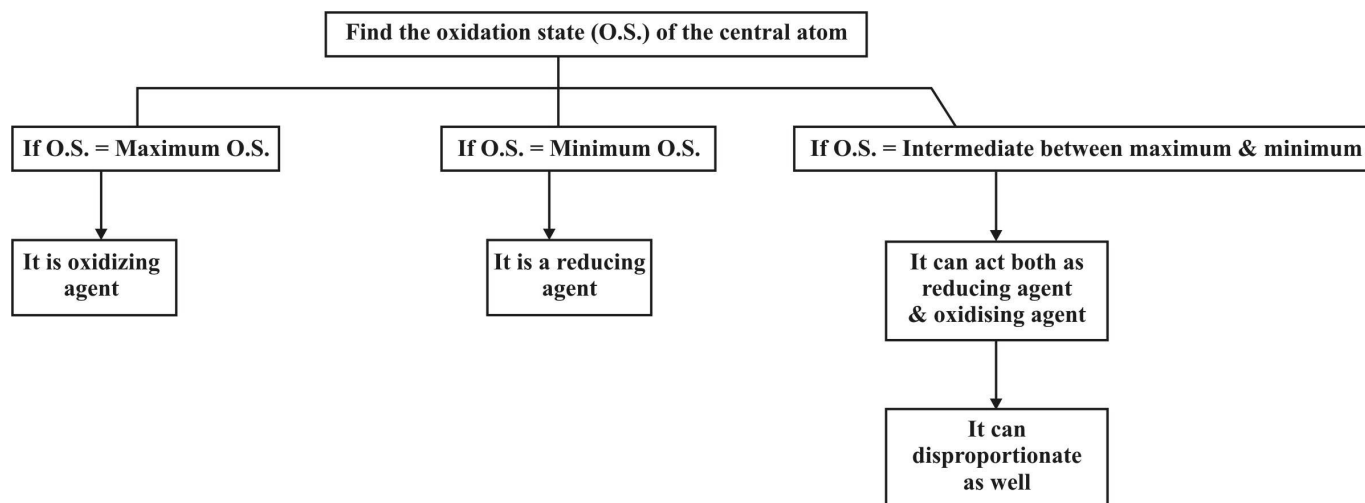
Reducing agents are those compounds which can reduce other and oxidise itself during the chemical reaction. Those reagents in which for an element, oxidation number increases or which undergoes loss of electrons in a redox reaction are termed as reductants.

e.g. KI, $\text{Na}_2\text{S}_2\text{O}_3$ etc are the powerful reducing agents.

Note : There are some compounds also which can work both as oxidising agent and reducing agent

e.g. H_2O_2 , NO_2^-

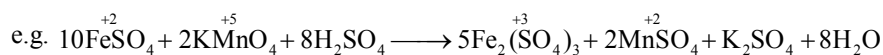
HOW TO IDENTIFY WHETHER A PARTICULAR SUBSTANCE IS AN OXIDISING OR A REDUCING AGENT



Redox reaction

A reaction in which oxidation and reduction simultaneously take place is called a redox reaction

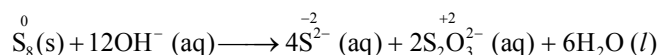
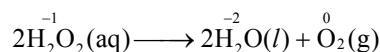
In all redox reactions, the total increase in oxidation number must be equal to the total decrease in oxidation number.



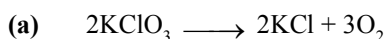
Disproportionation Reaction :

A redox reaction in which same element present in a particular compound in a definite oxidation state is oxidized as well as reduced simultaneously is a disproportionation reaction.

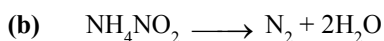
Disproportionation reactions are a special type of redox reactions. One of the reactants in a disproportionation reaction always contains **an element that can exist in at least three oxidation states**. The element in the form of reacting substance is in the intermediate oxidation state and both higher and lower oxidation states of that element are formed in the reaction. For example :



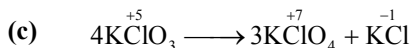
Consider the following reactions :



KClO_3 plays a role of oxidant and reductant both. Here, Cl present in KClO_3 is reduced and O present in KClO_3 is oxidized. Since same element is not oxidized and reduced, so it is not a disproportionation reaction, although it looks like one.



Nitrogen in this compound has -3 and +3 oxidation number, which is not a definite value. So it is not a disproportionation reaction. It is an example of comproportionation reaction, which is a class of redox reaction in which an element from two different oxidation state gets converted into a single oxidation state.

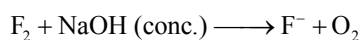
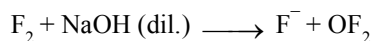


It is a case of disproportionation reaction and Cl atom is disproportionating.

List of some important disproportionation reactions

- $\text{H}_2\text{O}_2 \longrightarrow \text{H}_2\text{O} + \text{O}_2$
- $\text{X}_2 + \text{OH}^- (\text{dil.}) \longrightarrow \text{X}^- + \text{XO}^- \text{ (X = Cl, Br, I)}$
- $\text{X}_2 + \text{OH}^- (\text{conc.}) \longrightarrow \text{X}^- + \text{XO}_3^-$

F₂ does not undergo disproportionation as it is the most electronegative element.



- $(\text{CN})_2 + \text{OH}^- \longrightarrow \text{CN}^- + \text{OCN}^-$
 - $\text{P}_4 + \text{OH}^- \longrightarrow \text{PH}_3 + \text{H}_2\text{PO}_2^-$
 - $\text{S}_8 + \text{OH}^- \longrightarrow \text{S}^{2-} + \text{S}_2\text{O}_3^{2-}$
 - $\text{MnO}_4^{2-} \longrightarrow \text{MnO}_4^- + \text{MnO}_2$
 - $\text{NH}_2\text{OH} \longrightarrow \text{N}_2\text{O} + \text{NH}_3$
 $\text{NH}_2\text{OH} \longrightarrow \text{N}_2 + \text{NH}_3$
 - Oxyacids of Phosphorus (+1, +3 oxidation number)
 $\text{H}_3\text{PO}_2 \longrightarrow \text{PH}_3 + \text{H}_3\text{PO}_3$
 $\text{H}_3\text{PO}_3 \longrightarrow \text{PH}_3 + \text{H}_3\text{PO}_4$
 - Oxyacids of Chlorine (Halogens) (+1, +3, +5 Oxidation number)
 $\text{ClO}^- \longrightarrow \text{Cl}^- + \text{ClO}_2^-$
 $\text{ClO}_2^- \longrightarrow \text{Cl}^- + \text{ClO}_3^-$
 $\text{ClO}_3^- \longrightarrow \text{NO} + \text{HNO}_3$
 - $\text{HNO}_2 \longrightarrow \text{NO} + \text{HNO}_3$
- Reverse of disproportionation is called Comproportionation. In some of the disproportionation reactions, by changing the medium (from acidic to basic or reverse), the reaction goes in backward direction and can be taken as an example of Comproportionation reaction.
- $$\text{I}^- + \text{IO}_3^- + \text{H}^+ \longrightarrow \text{I}_2 + \text{H}_2\text{O}$$

REDOX REACTION

Balancing of redox reactions

All balanced equations must satisfy two criteria.

1. Atom balance (mass balance) :

There should be the same number of atoms of each kind on reactant and product side.

2. Charge balance :

The sum of actual charges on both sides of the equation must be equal.

There are two methods for balancing the redox equations :

1. Oxidation – number change method
2. Ion electron method or half cell method

- Since First method is not very much fruitful for the balancing of redox reactions, students are advised to use second method (Ion electron method) to balance the redox reactions

Ion electron method :

By this method redox equations are balanced in two different medium.

- (a) Acidic medium (b) Basic medium

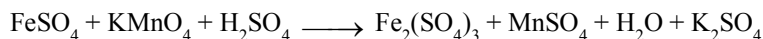
○ Balancing in acidic medium

Students are advised to follow the following steps to balance the redox reactions by ion electron method in acidic medium

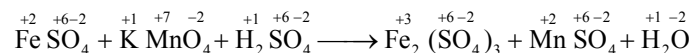
Solved Examples

Example - 3 :

Balance the following redox reaction :

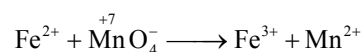


Sol. **Step-I** Assign the oxidation number to each element present in the reaction



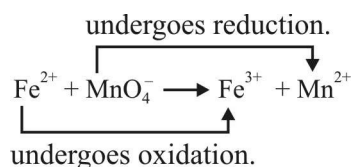
Step II :

Now convert the reaction in Ionic form by eliminating the elements or species, which are not undergoing either oxidation or reduction.



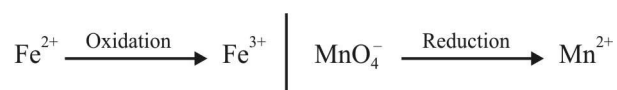
Step III :

Now identify the oxidation/reduction occurring in the reaction



Step IV :

Split the Ionic reaction in two half, one for oxidation and other for reduction.



Step V :

Balance the atom other than oxygen and hydrogen atom in both half reactions



Fe & Mn atoms are balanced on both side.

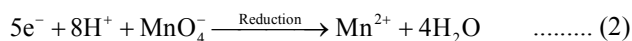
Step VI :

Now balance O & H atom by H₂O & H⁺ respectively by the following way : For one excess oxygen atom, add one H₂O on the other side and two H⁺ on the same side.



Step VII :

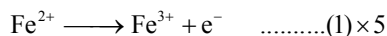
Equation (i) & (ii) are balanced atomwise. Now balance both equations chargewise. To balance the charge, add electrons to the electrically positive side.



Step VIII :

The number of electrons gained and lost in each half-reaction are equalised by multiplying both the half reactions with a suitable factor and finally the half reactions are added to give the overall balanced reaction.

Here, we multiply equation (1) by 5 and (2) by 1 and add them :

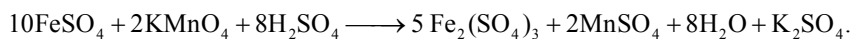
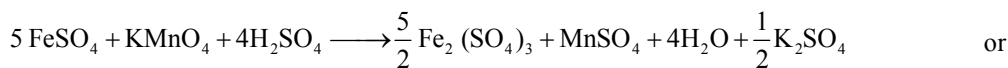


(Here, at his stage, you will get balanced redox reaction in Ionic form)

Step IX :

Now convert the ionic reaction into molecular form by adding the elements or species, which are removed in step (2).

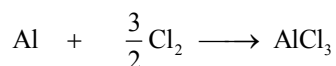
Now, by some manipulation, you will get :



○ **Balancing in basic medium :**

In this case, except step VI, all the steps are same. We can understand it by the following example :

$$\therefore \text{Equivalent weight of Zn} = \frac{65.5}{2} = 32.75\text{g}$$



$$27\text{ g} \quad \frac{3}{2} \times 71\text{ g}$$

\therefore 111.5 g chlorine reacts with 27 g of Al.

$$\therefore 35.5\text{ chlorine reacts with } \frac{27 \times 35.5}{111.5} = 9.0\text{ g of Al}$$

$$\therefore \text{Equivalent weight of aluminium} = \frac{27}{3} = 9.0$$

As we can see from the above examples that equivalent weight is the ratio of atomic weight and a factor (say n-factor or valency factor) which is in above three cases is their respective valencies.

Equivalent weight (E) :

$$\text{In general, Eq. wt. (E)} = \frac{\text{Atomic weight or molecular weight}}{\text{valency factor (v.f)}} = \frac{\text{Mol. wt.}}{\text{n - factor}} = \frac{M}{x}$$

$$\text{Number of Equivalents} = \frac{\text{mass of species}}{\text{eq. wt. of that species}}$$

For a solution, Number of equivalents = $N_1 V_1$, where N is the normality and V is the volume in litres

- Equivalent mass is a pure number which, when expressed in gram, is called gram equivalent mass.
- The equivalent mass of substance may have different values under different conditions.
- There is no hard and fast rule that equivalent weight will be always less than the molecular mass.

Valency factor calculation :

○ For Elements :

Valency factor = valency of the element.

○ For Acids :

Valency factor = number of replaceable H^+ ions per acid molecule

Solved Examples

Example - 5 : NaOH, KOH

Sol. v.f. \rightarrow 1 1

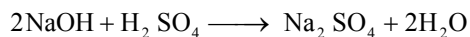
Eq. wt. $\frac{M}{1}$ $\frac{M}{1}$

- Bases may be defined as the substances in which OH group is/are directly attached with group I elements (Li, Na, K, Rb, Cs), group II elements (Be, Mg, Ca, Ba) or group III elements (Al, Ga, In, Tl), transition metals, non-metallic cations like PH_4^+ , NH_4^+ etc.
- **Acid - base reaction :**
In case of acid base reaction, the valence factor is the actual number of H^+ or OH^- replaced in the reaction. The acid or base may contain more number of replaceable H^+ or OH^- than actually replaced in reaction.
- **v. f.** for base is the number of H^+ ion from the acid replaced by each molecule of the base

REDOX REACTION

Solved Examples

Example - 6 :



Base Acid

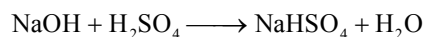
Sol. Valency factor of base = 1

Here, two molecule of NaOH replaced 2H^+ ion from the H_2SO_4 . Therefore, each molecule of NaOH replaced only one H^+ ion of acid, so v.f. = 1.

○ v. f. for acid is the number of OH^- replaced from the base by each molecule of acid

Solved Examples

Example - 7 :



Base Acid

Sol. Valency factor of acid = 1

Here, one of molecule of H_2SO_4 replaced one OH^- from NaOH. Therefore, valency factor for H_2SO_4 is one

$$\therefore \text{Eq. wt. of } \text{H}_2\text{SO}_4 = \frac{\text{Mol. wt}}{1}$$

○ Salts :

(a) In non-reacting condition

○ Valency factor = Total number of positive charge or negative charge present in the compound.

Solved Examples

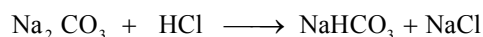
Example - 8 :

		Na_2CO_3 ,	$\text{Fe}_2(\text{SO}_4)_3$	$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$
Sol.	v.f.	2	$2 \times 3 = 6$	2
	Eq. wt.	$\frac{M}{2}$	$\frac{M}{6}$	$\frac{M}{2}$

(b) In reacting condition

Solved Examples

Example - 9 :



Base Acid

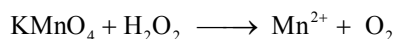
Sol. It is an acid base reaction, therefore valency factor for Na_2CO_3 is one while in non-reacting condition, it will be two.

(c) Equivalent weight of oxidising / reducing agents in a redox reaction

In case of redox change, v.f. = Total change in oxidation number per molecule.

Solved Examples

Example - 10 :



Sol. Mn in KMnO_4 is going from +7 to +2, so change in oxidation number per molecule of KMnO_4 is 5. So the valency factor of KMnO_4 is 5 and equivalent weight is $\frac{M}{5}$.

Normality :

Normality of a solution is defined as the number of equivalents of solute present in one litre (1000 mL) solution.

Let V mL of a solution is prepared by dissolving W g of solute of equivalent weight E in water.

○ Number of equivalents of solute = $\frac{W}{E}$

V mL of solution contain $\frac{W}{E}$ equivalents of solute

∴ 1000 mL solution will contain $\frac{W \times 1000}{E \times V}$ equivalents of solute.

○ **Normality (N)** = $\frac{W \times 1000}{E \times V}$

○ **Normality (N) = Molarity × Valency factor**

$N \times V$ (in mL) = $M \times V$ (in mL) × n

or

○ milliequivalents = millimoles × n

Solved Examples

Example 11 :

Calculate the normality of a solution containing 15.8 g of KMnO_4 in 50 mL acidic solution.

Sol : **Normality (N)** = $\frac{W \times 1000}{E \times V}$

Here $W = 15.8$ g, $V = 50$ mL $E = \frac{\text{molar mass of } \text{KMnO}_4}{\text{Valency factor}} = 158/5 = 31.6$

So, normality = 10 N

Example 12 :

Calculate the normality of a solution containing 50 mL of 5 M solution of $\text{K}_2\text{Cr}_2\text{O}_7$ in acidic medium.

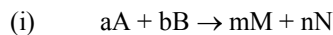
Sol : Normality (N) = Molarity × valency factor = $5 \times 6 = 30$ N

REDOX REACTION

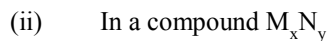
Law of Equivalence

The law states that one equivalent of an element combine with one equivalent of the other. In a chemical reaction, equivalents and milli equivalents of reactants react in equal amount to give same number of equivalents or milli equivalents of products separately.

Accordingly



$$\text{meq of A} = \text{meq of B} = \text{meq of M} = \text{m.eq. of N}$$

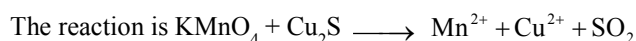


$$\text{meq of } M_xN_y = \text{meq of M} = \text{meq of N}$$

Solved Examples

Example 13 :

Find the number of moles of KMnO_4 needed to oxidise one mole Cu_2S in acidic medium.



Sol. From law of equivalence,

$$\text{equivalents of } \text{Cu}_2\text{S} = \text{equivalents of } \text{KMnO}_4$$

$$\text{moles of } \text{Cu}_2\text{S} \times \text{v.f.} = \text{moles of } \text{KMnO}_4 \times \text{v.f.}$$

$$1 \times 8 = \text{moles of } \text{KMnO}_4 \times 5 \quad \Rightarrow \text{moles of } \text{KMnO}_4 = 8/5$$

$$(\therefore \text{v.f. of } \text{Cu}_2\text{S} = 2(2-1) + 1(4 - (-2)) = 8 \text{ and v.f. of } \text{KMnO}_4 = 1(7-2) = 5)$$

Example 14 :

The number of moles of oxalate ions oxidized by one mole of MnO_4^- ion in acidic medium are :

(a) $\frac{5}{2}$

(b) $\frac{2}{5}$

(c) $\frac{3}{5}$

(d) $\frac{5}{3}$

Sol. Equivalents of $\text{C}_2\text{O}_4^{2-} = \text{equivalents of } \text{MnO}_4^-$

$$x (\text{mole}) \times 2 = 1 \times 5$$

$$(\therefore \text{v.f. of } \text{C}_2\text{O}_4^{2-} = 2(4-3) = 2 \text{ and v.f. of } \text{MnO}_4^- = 1(7-2) = 5.$$

$$x = \frac{5}{2} \text{ mole of } \text{C}_2\text{O}_4^{2-} \text{ ions.}$$

Solved Examples

Example : 15

How many millilitres of 0.02 M KMnO_4 solution would be required to exactly titrate 25 mL of 0.2 M $\text{Fe}(\text{NO}_3)_2$ solution in acidic medium ?

Sol. Method -1 : Mole concept method

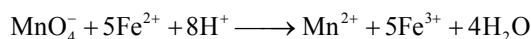
Starting with 25 mL of 0.2 M Fe^{2+} , we can write :

$$\text{Millimoles of } \text{Fe}^{2+} = 25 \times 0.2 \quad \dots\dots (1)$$

and in volume V (in milliliters) of the KMnO_4 ,

$$\text{Millimoles of } \text{MnO}_4^- = V(0.02) \quad \dots\dots (2)$$

The balanced reaction is :



This requires that at the equivalent point,

$$\frac{\text{m.moles of MnO}_4^-}{1} = \frac{\text{m.moles of Fe}^{2+}}{5}$$

$$\therefore \frac{V(0.02)}{1} = \frac{(25)(0.2)}{5} \text{ (from (1) \& (2))}$$

$$\therefore V = 50 \text{ mL.}$$

Method -2 : Equivalent Method :

At the equivalence point,

$$\text{milliequivalents of MnO}_4^- = \text{milliequivalents of Fe}^{2+}$$

$$M_1 \times v f_1 \times V_1 = M_2 \times v f_2 \times V_2$$

$$0.02 \times 5 \times V_1 = 0.2 \times 1 \times 25 \quad \because \text{MnO}_4^- \rightarrow \text{Mn}^{2+}; v.f. = 5, \text{ Fe}^{2+} \rightarrow \text{Fe}^{3+}; v.f. = 1)$$

$$\therefore V_1 = 50 \text{ mL.}$$

Titration

Titration is procedure for determining the concentration of a solution by allowing a carefully measured volume to react with a standard solution of another substance, whose concentration is known.

Standard solution - It is a solution whose concentration is known and is taken in burette. It is also called **Titrant**.

There are two type of **titrants** :

- **Primary titrants/standard** - These reagents can be accurately weighed and their solutions are not to be standardised before use.

Ex : Oxalic acid, $\text{K}_2\text{Cr}_2\text{O}_7$, AgNO_3 , CuSO_4 , ferrous ammonium sulphate, hypo etc.

- **Secondary titrants/standard** : These reagents cannot be accurately weighed and their solutions are to be standardised before use.

Ex : NaOH , KOH , HCl , H_2SO_4 , I_2 , KMnO_4 , etc.

Titrate : Solution consisting of substance to be estimated, generally taken in a beaker.

Equivalence point : It is the point when number of equivalents of titrant added becomes equal to number of equivalents of titrate.

At equivalence point :

$$n_1 V_1 M_1 = n_2 V_2 M_2$$

Indicator : An auxiliary substance added for physical detection of completion of titration at equivalence point. It generally show colour change on completion of titration.

Types of Titrations :

- Acid-base titrations (to be studied in Ionic equilibrium)

REDOX REACTION

○ Redox Titrations

Some Common Redox Titrations

Table of Redox Titrations : (Excluding Iodometric/Iodimetric titrations)

Estimation of	By titrating with	Reactions	Relation *between OA and RA
1. Fe ²⁺	MnO ₄ ⁻	$\text{Fe}^{2+} \rightarrow \text{Fe}^{3+} + \text{e}^{-}$ $\text{MnO}_4^{-} + 8\text{H}^{+} + 5\text{e}^{-} \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O}$	$5\text{Fe}^{2+} \equiv \text{MnO}_4^{-}$ $\text{Eq. wt. of Fe}^{2+} = \text{M}/1$
2. Fe ²⁺	Cr ₂ O ₇ ²⁻	$\text{Fe}^{2+} \rightarrow \text{Fe}^{3+} + \text{e}^{-}$ $\text{Cr}_2\text{O}_7^{2-} + 14\text{H}^{+} + 6\text{e}^{-} \rightarrow 2\text{Cr}^{3+} + 7\text{H}_2\text{O}$	$6\text{Fe}^{2+} \equiv \text{Cr}_2\text{O}_7^{2-}$ $\text{Eq. wt. of Cr}_2\text{O}_7^{2-} = \text{M}/6$
3. C ₂ O ₄ ²⁻	MnO ₄ ⁻	$\text{C}_2\text{O}_4^{2-} \rightarrow 2\text{CO}_2 + 2\text{e}^{-}$ $\text{MnO}_4^{-} + 8\text{H}^{+} + 5\text{e}^{-} \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O}$	$5\text{C}_2\text{O}_4^{2-} \equiv 2\text{MnO}_4^{-}$ $\text{Eq. wt. of C}_2\text{O}_4^{2-} = \text{M}/2$
4. H ₂ O ₂	MnO ₄ ⁻	$\text{H}_2\text{O}_2 \rightarrow 2\text{H}^{+} + \text{O}_2 + 2\text{e}^{-}$ $\text{MnO}_4^{-} + 8\text{H}^{+} + 5\text{e}^{-} \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O}$	$5\text{H}_2\text{O}_2 \equiv 2\text{MnO}_4^{-}$ $\text{Eq. wt. of H}_2\text{O}_2 = \text{M}/2$
5. As ₂ O ₃	MnO ₄ ⁻	$\text{As}_2\text{O}_3 + 5\text{H}_2\text{O} \rightarrow 2\text{AsO}_4^{3-} + 10\text{H}^{+} + 4\text{e}^{-}$ $\text{MnO}_4^{-} + 8\text{H}^{+} + 5\text{e}^{-} \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O}$	$\text{Eq. wt. of As}_2\text{O}_3 = \text{M}/4$
6. AsO ₃ ³⁻	BrO ₃ ⁻	$\text{AsO}_3^{3-} + \text{H}_2\text{O} \rightarrow \text{AsO}_4^{3-} + 2\text{H}^{+} + 2\text{e}^{-}$ $\text{BrO}_3^{-} + 6\text{H}^{+} + 6\text{e}^{-} \rightarrow \text{Br}^{-} + 3\text{H}_2\text{O}$	$\text{Eq. wt. of AsO}_3^{3-} = \text{M}/2$ $\text{Eq. wt. of BrO}_3^{-} = \text{M}/6$

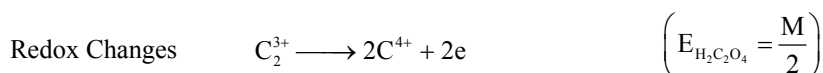
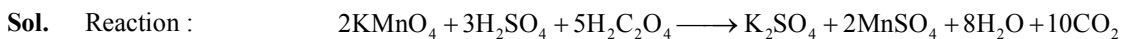
Permanganate Titrations :

- KMnO₄ is generally used as oxidising agent in acidic medium, generally provided by dilute H₂SO₄.
- KMnO₄ works as self indicator persistent pink color is indication of end point.
- Mainly used for estimation of Fe²⁺, oxalic acid, oxalates, H₂O₂ etc.

Solved Examples

Example 16 :

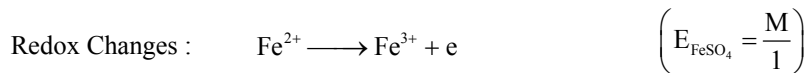
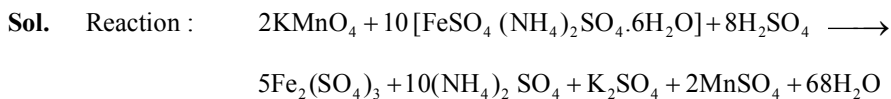
Write the balanced reaction of titration of KMnO₄ Vs oxalic acid in presence of H₂SO₄.



Indicator : KMnO₄ acts as self indicator.

Example : 17

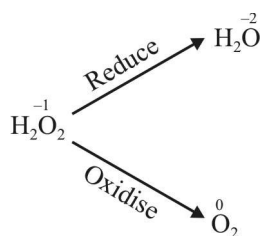
Write the balanced reaction of titration of KMnO_4 Vs ferrous ammonium sulphate in presence of H_2SO_4 .



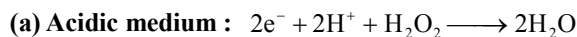
Indicator : KMnO_4 acts as self indicator.

Hydrogen peroxide (H_2O_2)

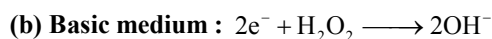
H_2O_2 can behave both like oxidising and reducing agent in both the mediums (acidic and basic).



○ **Oxidising agent : ($\text{H}_2\text{O}_2 \rightarrow \text{H}_2\text{O}$)**



v.f = 2

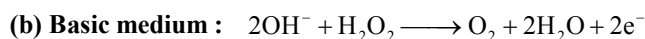


v.f = 2

○ **Reducing agent : ($\text{H}_2\text{O}_2 \rightarrow \text{O}_2$)**



v.f = 2



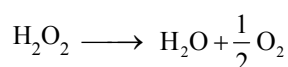
v.f = 2

Note : Valency factor of H_2O_2 is always equal to 2.

Volume strength of H_2O_2 : Strength of H_2O_2 is represented as 10V, 20V, 30V etc.

20V H_2O_2 means **one litre** of this sample of H_2O_2 on decomposition gives **20L of O_2** gas of **STP**.

Decomposition of H_2O_2 is given as :



REDOX REACTION

$$1 \text{ mole} \quad \frac{1}{2} \times 22.4 \text{ L O}_2 \text{ at STP}$$

$$= 34\text{g} \quad = 11.2 \text{ L O}_2 \text{ at STP}$$

To obtain 11.2 litre O₂ at STP, at least 34 g H₂O₂ must be decomposed.

$$\text{For 20 L O}_2, \text{ we should decompose atleast } \frac{34}{11.2} \times 20 \text{ g H}_2\text{O}_2$$

$$\therefore \quad 1\text{L solution of H}_2\text{O}_2 \text{ contains } \frac{34}{11.2} \times 20 \text{ g H}_2\text{O}_2$$

$$\therefore \quad 1\text{L solution of H}_2\text{O}_2 \text{ contains } \frac{34}{11.2} \times \frac{20}{17} \text{ equivalents of H}_2\text{O}_2 \quad (E_{\text{H}_2\text{O}_2} = \frac{M}{2} = \frac{34}{2} = 17)$$

$$\text{Normality of H}_2\text{O}_2 = \frac{34}{11.2} \times \frac{20}{17} = \frac{20}{5.6}$$

$$\textcircled{\circ} \quad \text{Normality of H}_2\text{O}_2 \text{ (N)} = \frac{\text{Volume strength of H}_2\text{O}_2}{5.6}$$

$$\therefore \quad M_{\text{H}_2\text{O}_2} = \frac{N_{\text{H}_2\text{O}_2}}{\text{v.f.}} = \frac{N_{\text{H}_2\text{O}_2}}{2}$$

$$\textcircled{\circ} \quad \text{Molarity of H}_2\text{O}_2 \text{ (M)} = \frac{\text{Volume strength of H}_2\text{O}_2}{11.2}$$

Strength (in g/L) : Denoted by S

$$\text{Strength} = \text{Molarity} \times \text{Mol. wt} = \text{Molarity} \times 34$$

$$\text{Strength} = \text{Normality} \times \text{Eq. weight} = \text{Normality} \times 17$$

Solved Examples

Example 18 :

20 mL of H₂O₂ after acidification with dilute H₂SO₄ required 30 mL of $\frac{N}{12}$ KMnO₄ for complete oxidation. Find the strength of H₂O₂ solution. [Molar mass of H₂O₂ = 34]

Sol. meq. of KMnO₄ = meq. of H₂O₂

$$30 \times \frac{1}{12} = 20 \times N'$$

$$N' = \frac{30}{12 \times 20} = \frac{1}{8} N$$

$$\therefore \quad \text{strength} = N' \times \text{equivalent mass} = \frac{1}{8} \times 17 = 2.12 \text{ g/L.}$$

Hardness of water (Hard water does not give lather with soap)

Temporary hardness - due to bicarbonates of Ca & Mg

Permanent hardness - due to chlorides & sulphates of Ca & Mg. There are some method by which we can soften the water sample.

- (a) By boiling : $2\text{HCO}_3^- \longrightarrow \text{H}_2\text{O} + \text{CO}_2 + \text{CO}_3^{2-}$ or
 By Slaked lime : $\text{Ca}(\text{HCO}_3)_2 + \text{Ca}(\text{OH})_2 \longrightarrow \text{CaCO}_3 + 2\text{H}_2\text{O}$
 $\text{Ca}^{2+} + \text{CO}_3^{2-} \longrightarrow \text{CaCO}_3$
- (b) By Washing Soda : $\text{CaCl}_2 + \text{Na}_2\text{CO}_3 \longrightarrow \text{CaCO}_3 + 2\text{NaCl}$
- (c) By ion exchange resins : $\text{Na}_2\text{R} + \text{Ca}^{2+} \longrightarrow \text{CaR} + 2\text{Na}^+$
- (d) By adding chelating agents like $(\text{PO}_3^-)_3$ etc.

Parts Per Million (ppm)

When the solute is present in very less amount, then this concentration term is used. It is defined as the number of parts of the solute present in every 1 million parts of the solution. ppm can both be in terms of mass or in terms of moles. If nothing has been specified, we take ppm to be in terms of mass. Hence, a 100 ppm solution means that 100 g of solute is present in every 1000000g of solution.

$$\text{ppm}_A = \frac{\text{mass of A}}{\text{Total mass}} \times 10^6 = \text{mass fraction} \times 10^6$$

Measurement of Hardness :

Hardness is measured in terms of ppm (parts per million) of CaCO_3 or equivalent to it.

$$\text{Hardness in ppm} = \frac{\text{mass of CaCO}_3}{\text{Total mass of solution}} \times 10^6$$

Solved Examples

Example 19 :

0.00012% MgSO_4 and 0.000111% CaCl_2 is present in water. What is the measured hardness of water and millimoles of washing soda required to purify water 1000 L water ?

Sol. Basis of calculation = 100 g hard water

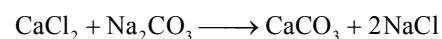
$$\text{MgSO}_4 = 0.00012\text{g} = \frac{0.00012}{120} \text{ mole}$$

$$\text{CaCl}_2 = 0.000111\text{g} = \frac{0.000111}{111} \text{ mole}$$

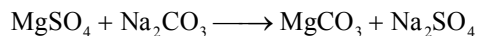
$$\therefore \text{equivalent moles of CaCO}_3 = \left(\frac{0.00012}{120} + \frac{0.000111}{111} \right) \text{ mole}$$

$$\therefore \text{mass of CaCO}_3 = \left(\frac{0.00012}{120} + \frac{0.000111}{111} \right) \times 100 = 2 \times 10^{-4} \text{ g}$$

$$\text{Hardness (in terms of ppm of CaCO}_3) = \frac{2 \times 10^{-4}}{100} \times 10^6 = 2 \text{ ppm}$$



REDOX REACTION



$$\therefore \text{ Required Na}_2\text{CO}_3 \text{ for 100g of water} = \left(\frac{0.00012}{120} + \frac{0.000111}{111} \right) \text{ mole}$$

$$= 2 \times 10^{-6} \text{ mole}$$

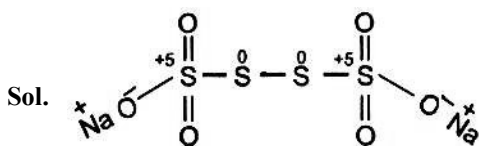
$$\therefore \text{ Required Na}_2\text{CO}_3 \text{ for 1000 litre water} = \frac{2 \times 10^{-6}}{100} \times 10^6 = \frac{2}{100} \text{ mole} \quad (\because d = 1 \text{g/mL})$$

$$= \frac{20}{1000} \text{ mole} = 20 \text{ m mole}$$

SOLVED EXAMPLES

Example – 1

Calculate individual oxidation number of each S-atom in $\text{Na}_2\text{S}_4\text{O}_6$ (sodium tetrathionate) with the help of its structure.



Example – 2

Find the average and individual oxidation number of Fe & Pb in Fe_3O_4 & Pb_3O_4 , which are mixed oxides.

Sol. (i) Fe_3O_4 is mixture of FeO & Fe_2O_3 in 1 : 1 ratio
so, individual oxidation number of Fe = +2 & +3

$$\text{ \& average oxidation number} = \frac{1(+2) + 2(+3)}{3} = 8/3$$

(ii) Pb_3O_4 is a mixture of PbO & PbO_2 in 2 : 1 molar ratio
so, individual oxidation number of Pb are +2 & +4

$$\text{ \& average oxidation number of Pb} = \frac{2(+2) + 1(+4)}{3} = 8/3$$

Example – 3

Balance the following equations :

- $\text{H}_2\text{O}_2 + \text{MnO}_4^- \longrightarrow \text{Mn}^{+2} + \text{O}_2$ (acidic medium)
- $\text{Zn} + \text{HNO}_3$ (dil) $\longrightarrow \text{Zn}(\text{NO}_3)_2 + \text{H}_2\text{O} + \text{NH}_4\text{NO}_3$
- $\text{CrI}_3 + \text{KOH} + \text{Cl}_2 \longrightarrow \text{K}_2\text{CrO}_4 + \text{KIO}_4 + \text{KCl} + \text{H}_2\text{O}$
- $\text{P}_2\text{H}_4 \longrightarrow \text{PH}_3 + \text{P}_4$
- $\text{Ca}_3(\text{PO}_4)_2 + \text{SiO}_2 + \text{C} \longrightarrow \text{CaSiO}_3 + \text{P}_4 + \text{CO}$

