MOLE CONCEPT

CONTENTS

Particulars			Page No.
Theory			01 – 32
Exercise - 1			33 – 42
Part - I	:	Subjective Questions	
Part - II	:	Objective Questions	
Part - III	:	Match the Columns	
Exercise - 2			43 – 49
Part - I	:	Objective Questions	
Part - II	:	Numerical type questions	
Part - III	:	One or More Than One Options Correct Ty	ре
Part - IV	:	Comprehensions	
Exercise - 3			50 – 53
Part - I	:	JEE(ADVANCED) Problems (Previous Yea	rs)
Part - II	:	JEE(MAIN) / AIEEE Problems (Previous Ye	ars)
Answer Key			54 – 56
Reliable Ranker I	Problem	s (RRP)	57 – 70
PART- 1	:	Paper JEE (main) pattern	
Part - 2	:	Paper JEE (advanced) pattern	
Part - 3	:	OLYMPIAD (previous Years)	
Part - 4	:	Additional Problems	
RRP Answer Key			71 – 72
RRP Solutions			73 – 94

JEE (Advanced) Syllabus

Concept of atoms and molecules ; Delton's atomic theory ; Mole concept ; Chemical formulae ; Balanced chemical equations ; Calculations (based on mole concept) involving common oxidation reduction, neutralisation, and displacement reaction ; Concentration in terms of mole fraction, molarity, molality and normality.

JEE (MAIN) Syllabus

Mole concept : Matter and its nature, Dalton's atomic theory ; Concept of atom, molecule, element and compound ; Physical quantities and their measurements in chemistry, precision and accuracy, significant figures, S.I. units, dimensional analysis ; Laws of chemcial combination ; Atomic and moleclar masses, mole concept, molar mass, percentage composition, empirical and molecular formulae ; Chemical equations



Basic Definitions :

Relative atomic mass :

One of the most important concept come out from Dalton's atomic theory was that of relative atomic mass or relative atomic weight. This is done by expressing mass of one atom with respect to a fixed standard. Dalton used hydrogen as the standard (H = 1). Later on oxygen (O = 16) replaced hydrogen as the reference. Therefore relative atomic mass is given as

On hydrogen scale : Relative atomic mass (R.A.M) = $\frac{\text{Mass of one atom of an element}}{\text{mass of one hydrogen atom}}$

On oxygen scale : Relative atomic mass (R.A.M) = $\frac{\text{Mass of one atom of an element}}{\frac{1}{16} \times \text{mass of one oxygen atom}}$

• The present standard unit which was adopted internationally in 1961, is based on the mass of one carbon-12 atom.

Relative atomic mass (R.A.M) = $\frac{\text{Mass of one atom of an element}}{\frac{1}{12} \times \text{mass of one } C - 12 \text{ atom}}$

Atomic mass unit (or amu) :

The atomic mass unit (amu) is equal to $\left(\frac{1}{12}\right)^{\text{th}}$ mass of one atom of carbon-12 isotope.

- $\therefore \quad 1 \text{ amu} = \frac{1}{12} \times \text{mass of one C-12 atom}$ $\simeq \text{mass of one nucleon in C-12 atom.}$ $= 1.66 \times 10^{-24} \text{ g or } 1.66 \times 10^{-27} \text{ kg}$
- O one amu is also called one Dalton (Da).

O Today, amu has been replaced by 'u' which is known as unified mass

Atomic & molecular mass :

÷

Atomic mass is the mass of 1 atom of a substance, it is expressed in amu.

• Atomic mass = R.A.M × 1 amu

Molecular mass is the mass of 1 atom of a substance, it is expressed in amu.

O Molecular mass = Relative molecular mass × 1 amu

Note: Relative atomic mass is nothing but the number of nucleons present in the atom.

Ex. Find the relative atomic mass of 'O' atom and its atomic mass.

- Sol. The number of nucleons present in 'O' atom is 16.
 - relative atomic mass of 'O' atom = 16.

Atomic mass = R.A.M × 1 amu = 16 × 1 amu = 16 amu

Mole : The Mass / Number Relationship

Mole is a chemical counting SI unit and defined as follows :

A mole is the amount of a substance that contains as many entities (atoms, molecules or other particles) as there are atoms in exactly 0.012 kg (or 12 g) of the carbon-12 isotope.

From mass spectrometer we found that there are 6.023×10^{23} atoms present in 12 g of C-12 isotope.

The number of entities in 1 mol is so important that it is given a separate name and symbol known as Avogadro constant denoted by N_{a} .

i.e. on the whole we can say that 1 mole is the collection of 6.02×10^{23} entities. Here entities may represent atoms, ions, molecules or even pens, chair, paper etc also include in this but as this number (N_A) is very large therefore it is used only for very small things.

HOW BIG IS A MOLE ?

Amount of water in Age of earth (seconds) world's oceans (litres) Population of earth Avogadro's number 602,200,000,000,000,000,000 Distance from earth

to sun (centimeters)

O Note: In modern practice gram-atom and gram-molecule are termed as mole.

Gram Atomic Mass :

The atomic mass of an element expressed in gram is called gram atomic mass of the element.

It is also defined as mass of 6.02×10^{23} atoms.

or

or

It is also defined as the mass of one mole atoms.

For example for oxygen atom :

Atomic mass of 'O' atom = mass of one 'O' atom = 16 amu gram atomic mass = mass of 6.02×10^{23} 'O' atoms = 16 amu × 6.02×10^{23} = 16 × 1.66×10^{-24} g × 6.02×10^{23} = 16 g (:: 1.66 × 10^{-24} × 6.02×10^{23} ~ 1)

Solved Examples -

Ex. How many atoms of oxygen are their in 16 g oxygen.

Sol. Let x atoms of oxygen are present

So,
$$x = \frac{16}{32} \times 2N_A$$

 $\Rightarrow x = N_A$

Gram molecular mass :

The molecular mass of a substance expressed in gram is called the gram-molecular mass of the substance.

It is also defined as mass of 6.02 × 10²³ molecules

or

or

It is also defined as the mass of one mole molecules.

For example for O_2 molecule :

Molecular mass of O_2 'r	nolecule = mass of one O_2 molecule
_	= 2 × mass of one 'O' atom
	= 2 × 16 amu
	= 32 amu
gram molecular mass	= mass of 6.02×10^{23} 'O ₂ ' molecules = 32 amu × 6.02×10^{23}
	$= 32 \times 1.66 \times 10^{-24} \text{ g} \times 6.02 \times 10^{23} = 32 \text{ g}$

Solved Examples -

The molecular mass of H₂SO₄ is 98 amu. Calculate the number of moles of each element in 294 g of H₂SO₄. Ex. Sol.

Gram molecular mass of $H_2SO_4 = 98 g$

moles of $H_2SO_4 = \frac{294}{98} = 3$ moles					
H ₂ SO ₄	н	S	0		
One molecule	2 atom	one atom	4 atom		
1 × N _A	$2 \times N_A$ atoms	1 × N _A atoms	$4 \times N_A$ atoms		
∴ one mole	2 mole	onemole	4 mole		
∴ 3 mole	6 mole	3 mole	12 mole		

Gay-Lussac's Law of Combining Volume :

According to him elements combine in a simple ratio of atoms, gases combine in a simple ratio of their volumes provided all measurements should be done at the same temperature and pressure

$H_2(g)$	+	$Cl_2(g)$	\longrightarrow 2HCl
1 vol		1 vol	2 vol

Avogadro's hypothesis :

Equal volumes of all gases have equal number of molecules (not atoms) at the same temperature and pressure condition.

S.T.P. (Standard Temperature and Pressure)

At S.T.P. condition : temperature = 0°C or 273 K

pressure = 1 bar

and volume of one mole of gas at STP is found to be experimentally equal to 22.7 litres which is known as molar volume.

Note : Measuring the volume is equivalent to counting the number of molecules of the gas.

-Solved Examples -

Ex. Calculate the volume in litres of 20 g hydrogen gas at STP.

No. of moles of hydrogen gas = $\frac{Mass}{Molecular mass} = \frac{20 \text{ gm}}{2 \text{ gm}} = 10 \text{ mol}$ Sol.

volume of hydrogen gas at STP = 10 x 22.7 lt. = 227 lt.

Y-map : Interconversion of mole - volume, mass and number of particles :



Percentage Composition :

Here we are going to find out the percentage of each element in the compound by knowing the molecular formula of compound.

We know that according to law of definite proportions any sample of a pure compound always possess constant ratio with their combining elements.

Solved Examples

Ex. Every molecule of ammonia always has formula NH, irrespective of method of preparation or sources. i.e. 1 mole of ammonia always contains 1 mol of N and 3 mole of H. In other words 17 g of NH, always contains 14 g of N and 3 g of H. Now find out % of each element in the compound.

Sol.	Mass % of N in NH ₃ = $\frac{\text{Mass of N in 1 mol NH}_3}{\text{Mass of 1 mol of NH}_3} \times 100 = \frac{14 \text{ gm}}{17} \times 100 = 82.35 \%$
	Mass % of H in NH ₃ = $\frac{\text{Mass of H is 1 mol NH}_3}{\text{Mass of 1 mole of NH}_3} \times 100 = \frac{3}{17} \times 100 = 17.65 \%$

DENSITY :

- It is of two types.
- I. Absolute density
- II. Relative density

For liquids and solids :

Absolute density = volume

density of the substance

Relative density or specific gravity = $\frac{1}{\text{density of water at 4°C (1 gm ml^{-1})}}$

For gases :

PM mass Absolute density =

= RT volume

where P is pressure of gas, M = mol. wt. of gas, R is the gas constant, T is absolute temperature. Vapour Density :

Vapour density is defined as the density of the gas with respect to hydrogen gas at the same temperature and pressure.

Vapour density =
$$\frac{d_{gas}}{d_{H_2}} = \frac{PM_{gas}}{PM_{H_2/RT}}$$

V.D. = $\frac{M_{gas}}{M_{H_2}} = \frac{M_{gas}}{2} \implies \boxed{M_{gas} = 2 \times V.D.}$

molecular formula mass

where $n = \overline{empirical formula mass}$

Ex. A gaseous mixture of H_2 and NH_3 gas contains 68 mass % of NH_3 . The vapour density of the mixture is

Sol. No. of moles of NH₃ in 100g mixture =
$$\frac{68}{17} = 4$$

No. of moles of H₂ in 100g mixture =
$$\frac{32}{2} = 16$$

$$M_{average} = \frac{Total mass}{Total moles} = \frac{100}{4+16} = 5$$

V.d =
$$\frac{5}{2}$$
 = 2.5

Empirical and molecular formula :

We have just seen that knowing the molecular formula of the compound we can calculate percentage composition of the elements. Conversely if we know the percentage composition of the elements initially, we can calculate the relative number of atoms of each element in the molecules of the compound. This gives us the empirical formula of the compound. Further if the molecular mass is known then the molecular formula can easily be determined.

The empirical formula of a compound is a chemical formula showing the relative number of atoms in the simplest ratio. An empirical formula represents the simplest whole number ratio of various atoms present in a compound.

The molecular formula gives the actual number of atoms of each element in a molecule. The molecular formula shows the exact number of different types of atoms present in a molecule of a compound. The molecular formula is an integral multiple of the empirical formula.

i.e. molecular formula = empirical formula × n

Solved Examples –

Ex. Acetylene and benzene both have the empirical formula CH. The molecular masses of acetylene and benzene are 26 and 78 respectively. Deduce their molecular formulae.

Sol. : Empirical Formula is CH

Step-1 The empirical formula of the compound is CH

 $\therefore \qquad \text{Empirical formula mass} = (1 \times 12) + 1 = 13.$ Molecular mass = 26

Step-2 To calculate the value of 'n'

$$n = \frac{Molecular mass}{Empirical formula mass} = \frac{26}{13} = 2$$

Step-3 To calculate the molecular formula of the compound.

```
Molecular formula = n \times (Empirical formula of the compound)
```

$$2 \times CH = C_2 H_2$$

Thus the molecular formula is $C_2 H_2$ Similarly for benzene

To calculate the value of 'n'

 $n = \frac{Molecular mass}{Empirical formula mass} = \frac{78}{13} = 6$

thus the molecular formula is $6 \times CH = C_6 H_6$

Sol.

An organic substance containing carbon, hydrogen and oxygen gave the following percentage composition. Ex. C = 40.684%; H = 5.085% and O = 54.228%

The molecular weight of the compound is 118 g. Calculate the molecular formula of the compound. Step-1

To calculate the empirical formula of the compound.

Element	Symbol	Percentage of element	At. mass of element	Relative no. of atoms = Percentage At. mass	Simplest atomic ratio	Simplest whole no. atomic ratio
Carbon	С	40.687	12	$\frac{40.687}{12} = 3.390$	3.390 3.389 =1	2
Hydrogen	н	5.085	1	$\frac{5.085}{1} = 5.085$	$\frac{5.085}{3.389}$ =1.5	3
Oxygen	ο	54.228	16	$\frac{54.228}{16} = 3.389$	3.389 3.389 =1	2

Empirical Formula is C₂H₃O₂

Step-2 To calculate the empirical formula mass.

The empirical formula of the compound is $C_{a}H_{a}O_{a}$.

Empirical formula mass =
$$(2 \times 12) + (3 \times 1) + (2 \times 16) = 59$$
.

Step-3 To calculate the value of 'n'

 $n = \frac{\text{Molecular mass}}{\text{Empirical formula mass}} = \frac{118}{59} = 2$ Molecular mass

Step-4 To calculate the molecular formula of the salt. = n × (Empirical formula) = 2 × $C_2 H_3 O_2 = C_4 H_6 O_4$ Molecular formula Thus the molecular formula is $C_A H_{e} O_A$

Chemical Reaction :

It is the process in which two or more than two substances interact with each other where old bonds are broken and new bonds are formed.

Chemical Equation :

All chemical reaction are represented by chemical equations by using chemical formula of reactants and products. Qualitatively a chemical equation simply describes what the reactants and products are. However, a balanced chemical equation gives us a lot of guantitative information. Mainly the molar ratio in which reactants combine and the molar ratio in which products are formed.

Attributes of a balanced chemical equation:

- (a) It contains an equal number of atoms of each element on both sides of equation.(POAC)
- (b) It should follow law of charge conservation on either side.
- (c) Physical states of all the reagents should be included in brackets.
- (d) All reagents should be written in their standard molecular forms (not as atoms)
- (e) The coefficients give the relative molar ratios of each reagent.

Write a balance chemical equation for following reaction : Ex.

When potassium chlorate (KCIO₃) is heated it gives potassium chloride (KCI) and oxygen (O₂).

 KCIO_3 (s) $\xrightarrow{\Delta}$ KCI (s) + O_2 (g) (unbalanced chemical equation) Sol. 2KCIO₂ (s) $\xrightarrow{\Lambda}$ 2 KCI (s) + 3 O₂ (g) (balanced chemical equation) Remember a balanced chemical equation is one which contains an equal number of atoms of each element on both sides of equation.

Interpretation of balanced chemical equations :

Once we get a balanced chemical equation then we can interpret a chemical equation by following ways

- Mass mass analysis •
- Mass volume analysis •
- Mole - mole analysis
- Vol Vol analysis (separately discussed as eudiometry or gas analysis) • Now you can understand the above analysis by following example
- Mass-mass analysis :

Consider the reaction

 $2\text{KCIO}_3 \longrightarrow 2\text{KCI} + 3\text{O}_2$ According to stoichiometry of the reaction mass-mass ratio: 2×122.5 : 2×74.5 : 3×32

 $\frac{\text{Mass of KCIO}_3}{\text{Mass of KCI}} = \frac{2 \times 122.5}{2 \times 74.5}$ or $\frac{\text{Mass of KCIO}_3}{\text{Mass of O}_2} = \frac{2 \times 122.5}{3 \times 32}$

Solved Examples -

- 367.5 gram KCIO, (M = 122.5) when heated. How many gram KCI and oxygen is produced. Ex.
- Sol. Balance chemical equation for heating of KCIO₃ is

 $2\text{KCIO}_3 \longrightarrow 2\text{KCI}$ + 30, 2×122.5 g : 2×74.5 g : 3×32 g mass-mass ratio : $\frac{\text{mass of KCIO}_3}{\text{mass of KCI}} = \frac{2 \times 122.5}{2 \times 74.5} \Rightarrow \frac{367.5}{W} = \frac{122.5}{74.5}$ $W = 3 \times 74.5 = 223.5 g$ $\frac{\text{Mass of KCIO}_3}{\text{Mass of O}_2} = \frac{2 \times 122.5}{3 \times 32} \Rightarrow \frac{367.5}{W} = \frac{2 \times 122.5}{3 \times 32}$ W = 144 a

Mass - volume analysis :

Now again consider decomposition of KCIO₃ $2\text{KCIO}_2 \longrightarrow 2\text{KCI} +$ 30₂ mass volume ratio : 2 × 122.5 g : 2 × 74.5 g : 3 × 22.7 lt. at STP we can use two relation for volume of oxygen $\frac{\text{Mass of KCIO}_3}{\text{volume of O}_2 \text{ at STP}} = \frac{2 \times 122.5}{3 \times 22.7 \text{ lt}}$...(i) $\frac{\text{Mass of KCI}}{\text{volume of O}_2 \text{ at STP}} = \frac{2 \times 74.5}{3 \times 22.7 \text{ lt}}$ and ...(ii)

 367.5 g KCIO_3 (M = 122.5) when heated, how many litre of oxygen gas is produced at STP. Ex.

Sol.

 $2\text{KCIO}_3 \longrightarrow 2\text{KCI} + 3\text{O}_2$ $\frac{\text{massof KCIO}_3}{\text{volume of O}_2 \text{ at STP}} = \frac{2 \times 122.5}{3 \times 22.7 \text{ lt}} \implies \frac{367.5}{V} = \frac{2 \times 122.5}{3 \times 22.7 \text{ lt}}$ $V = 3 \times 3 \times 11.35$ V = 102.15 lt \Rightarrow

Mole-mole analysis :

This analysis is very much important for quantitative analysis point of view. Students are advised to clearly understand this analysis.

Now consider again the decomposition of KCIO₂.

 $2\text{KCIO}_3 \longrightarrow 2\text{KCI} + 3\text{O}_2$

In very first step of mole-mole analysis you should read the balanced chemical equation like

2 moles KCIO₃ on decomposition gives you 2 moles KCI and 3 moles O₂ and from the stoichiometry of reaction we can write

$$\frac{\text{Moles of KCIO}_3}{2} = \frac{\text{Moles of KCI}}{2} = \frac{\text{Moles of O}_2}{3}$$

Now for any general balance chemical equation like

$$aA+bB \longrightarrow cC+dD$$

you can write.

Moles of A reacted moles of B reacted moles of C produced moles of D produced b d а С

Note : In fact mass-mass and mass-vol analysis are also interpreted in terms of mole-mole analysis you can use following chart also.



LIMITING REAGENT (L.R.):

- The reactant which is completely consumed when a reaction goes to completion is called Limiting (i) Reactant or Limiting reagent.
- The reactant whose Stoichiometric amount is least, is limiting reactant. (ii)

Given moles of reactant

Where ; Stoichiometric amount = Stoichiometric coefficient of reactant in balance Reaction

(iii) For calculation of moles of product, LR should be used. When amounts of two or more than two reactants are given :

Initial reacting mixture	aA n _a mol	+	bB n _B mol	\rightarrow	сC	+	dD
Stoichiometric amount	$\frac{n_A}{a}$		n _Β b				
If $\frac{n_A}{a} < \frac{n_B}{b} \Rightarrow A \text{ is limiting}$	reagent.						
If $\frac{n_A}{n_B} = \frac{a}{b}$ then reaction occurs to completion & no reactant is left at the end.							
If $\frac{n_A}{N_B} \rightarrow R$ is limiting	roagont						

 $\frac{1}{a} > \frac{1}{b} \Rightarrow B$ is limiting reagent.

28 gm Lithium is mixed with 48 gm O_2 to reacts according to the following reaction. Ex. $4Li + O_2 \longrightarrow 2Li_2O$ The mass of Li₂O formed is

2Li₂O

 $O_2 \longrightarrow$

4Li+

Sol. moles taken molestaken

48 28 7 32 =4 = 1.5 $\frac{\text{molestaken}}{\text{stoich.coeff.}} \qquad \frac{4}{4} = 1 \qquad \frac{1.5}{1} = 1.5$ Moles of Li_2O formed = $\frac{2}{4} \times 4 = 2$; Mass of Li_2O formed = $2 \times 30 = 60$ gm

- Ex. Calculate the mass of sucrose $C_{12}H_{22}O_{11}$ (s) produced by mixing 78 g of C(s), 11 g of $H_2(g)$ & 67.2 litre of O_2 (g) at 0°C and 1 atm according to given reaction of O₂ (g) (unbalanced)?
- **Sol.** 12C(s) + 11 H₂(g) + $\frac{11}{2}O_2 \rightarrow C_{12}H_{22}O_{11}(s)$ Moles taken $\frac{78}{12}$ $\frac{11}{2}$ $\frac{67.2}{22.4}$ = 6.5 = 5.5 = 3 $\frac{\text{molestaken}}{\text{stoich.coeff.}} = \frac{\frac{6.5}{12}}{0.54 = 0.5} = \frac{\frac{5.5}{11}}{0.545} = \frac{3}{5.5}$: Moles of $C_{12}H_{22}O_{11}$ formed = $\frac{5.5}{11} = 0.5$

Mass of sucrose obtained $= 0.5 \times 342 = 171$ grams.

PROBLEMS BASED ON MIXTURE :

The composition of any mixture may be determined by reacting the mixture with some substance, by which either one or more component of mixture may react.

1.5 gm mixture of SiO₂ and Fe₂O₃ on very strong heating leave a residue weighting 1.46 gm. The Ex. reaction responsible for loss of weight is

 $Fe_2O_3(s) \rightarrow Fe_3O_4(s) + O_2(g)$ What is the percentage by mass of Fe_2O_3 in original sample.

Sol.
$$3Fe_2O_3(s) \rightarrow 2Fe_3O_4 + \frac{1}{2}O_2$$

 $3 \times 160 \qquad \frac{1}{2} \times 32$
 $= 480 \text{ gm} \rightarrow = 16 \text{ gm}$
 $\log s \text{ of } 16 \text{ gm} \rightarrow 480 \text{ gm} \text{ Fe}_2O_3$
 $\log s \text{ of } 0.04 \text{ gm} \rightarrow 0.04 \times \frac{480}{16} = 1.2 \text{ gm} \text{ Fe}_2O_3$
 $\% \text{ by mass} = \frac{1.2}{1.5} \times 100 = 80\%$

In general, when a reaction is carried out in the laboratory we do not obtain actually the theoretical amount of the product. The amount of the product that is actually obtained is called the actual yield. Knowing the actual yield and theoretical yield the percentage yield can be calculate as :

% yield = $\frac{\text{Actual yield}}{\text{Theoretical yield}} \times 100$

The percentage yield of any product is always equal to the percentage extent of that reaction.

Ex. Aluminium reacts with sulphur to form aluminium sulphide. If 5.4 gm of Aluminium reacts with 12.8gm sulphure gives 12gm of aluminium sulphides, then the percent yield of the reaction is-

Sol 2AI + 3S
$$\longrightarrow$$
 Al₂S₃
Mole taken $\frac{5.4}{27}$ gm $\frac{12.8}{32}$ gm
= 0.2 = 0.4
 $\frac{\text{moles taken}}{\text{stoich.coeff.}} \frac{0.2}{2} \qquad \frac{0.4}{3}$
= 0.1 = 0.133
(L.R.)
moles of Al₂S₃ formed = $\frac{1}{2} \times 0.2 = 0.1$
mass = 0.1 × 150 = 15 gm
% yield = $\frac{\text{actual yield}}{\text{theoritical yield}} \times 100 = \frac{12}{15} \times 100 = 80$ %

PERCENTAGE PURITY:

The percentage of a specified compound or element in an impure sample may be given as

% purity = $\frac{\text{Actual mass of compound}}{\text{Total mass of sample}} \times 100$

If impurity is unknown, it is always considered as inert (unreactive) material.

Ex. A chalk sample exactly requires 17.52 gram HCl for complete reaction with all $CaCO_3$ present in it. If the chalk sample is 72% pure, the mass of sample taken is

Sol. $CaCO_3 + 2HCI \rightarrow CaCl_2 + H_2O + CO_2$

Moles of HCI =
$$\frac{17.52}{36.5}$$

Moles of CaCO₃ =
$$\frac{1}{2} \times \frac{17.52}{36.5}$$

Weight of CaCO₃ required = $\frac{1}{2} \times \frac{17.52}{36.5} \times 100$

Mass of sample taken :

 $= \frac{1}{2} \times \frac{17.52}{36.5} \times \frac{100 \times 100}{72} = 33.33 \text{ gm}$

PROBLEMS RELATED WITH SEQUENTIAL REACTION :

When one of products formed in previous reaction is consumed in the next one.

- $4NH_3(g) + 5O_2(g) \rightarrow 4NO(g) + 6H_2O(g)$ Ex. $2NO(g) + O_2(g) \longrightarrow 2NO_2(g)$ $3NO_2(g) + H_2O(\ell) \rightarrow 2HNO_3(aq.) + NO(g)$
- How many grams H₂SO₄ can be obtained from 1320 gm PbS as per reaction sequence? Ex. $\begin{array}{rcl} 2PbS &+ & O_2 & \longrightarrow & 2PbO + 2SO_2 \\ 3SO_2 &+ & 2HNO_3 + 2H_2O & \longrightarrow & 3H_2SO_4 + 2NO \\ [At. mass : Pb = 208, S = 32] \end{array}$
- **Sol.** Moles of PbS = $\frac{1320}{240}$ = 5.5 mol Moles of $SO_2 = 5.5 \text{ mol} = \text{moles of } H_2SO_4$ Mass of $H_2SO_4 = 5.5 \times 98 = 539$ gm [When amount of only one reactant is given generally other is assumed in excess.]

PROBLEM RELATED WITH PARALLEL REACTION :

When same two reactants form two or more products by independent reactions.

- $\begin{array}{ccc} \mathsf{KCIO}_3 & \longrightarrow & \mathsf{KCI} + \mathsf{O}_2 \\ \mathsf{KCIO}_3 & \longrightarrow & \mathsf{KCIO} + \mathsf{KCIO}_4 \end{array}$ Ex.
- Carbon reacts with oxygen forming carbon monoxide and/or carbon dioxide depending an availablity Ex. of oxygen. Find moles of each product obtained when 160 gm oxygen reacts with (a) 12 g carbon (b) 120 g carbon (c) 72 g carbon.

[initially use a reaction using lesser amount of oxygen]

Sol. (a)

 $C + \frac{1}{2}O_2 \longrightarrow CO$ t = 0 1mol 5mol **t =∞** 0 5-0.5 = 1mol (LR) 4.5mol Since CO & O₂ are left CO₂ is formed. $CO + \frac{1}{2}O_2 \longrightarrow CO_2$ 1mol 4.5mol t = 0 0 $\mathbf{t} = \infty \quad \mathbf{0}$ 4 mol 1 mol At end 1 mole CO₂ & no CO present $C + \frac{1}{2}O_2 \longrightarrow CO$ (b) t = 010mol 5mol 0 t =∞ 0 0 10mol At end only 10 mol CO present.

 $C + \frac{1}{2}O_2 \longrightarrow CO$ (C) t = 06mol 5mol 0 2mol t =∞ [LR] $CO + \frac{1}{2}O_2 \longrightarrow CO_2$

$$t = 0 \quad 6 \text{mol} \quad 2 \text{mol} \quad 0$$

$$t = \infty \quad 2 \text{mol} \quad 0 \text{ [LR]} \quad 4 \text{mol}$$

At end [2mol CO + 4 mol CO₂] left.

0

6mol

- Ex. 25.4 gm of iodine and 14.2 gm of chlorine are made to react completely to yield mixture of ICI and ICI₃. Ratio of moles of ICI & ICI₃ formed is (Atomic mass : I = 127, CI = 35.5)
- Sol.

 $\begin{array}{cccc} I_2 & + & CI_2 \longrightarrow & ICI + & ICI_3 \\ 0.1 \text{mol} & 0.2 \text{ mol} & x \text{ mol} & y \text{ mol} \\ \Rightarrow & 0.2 = x + y & \Rightarrow & x = 0.1 \\ 0.4 = x + 3y & \Rightarrow & y = 0.1 \\ \text{Ans. 1 : 1} \end{array}$

PRINCIPLE OF ATOM CONSERVATION (POAC)

POAC is nothing but the conservation of atoms of reactants and products involved in a chemical reaction. And if atoms are conserved, moles of atoms shall also be conserved. The principle is fruitful for the students when they don't get the idea of balanced chemical equation in the problem using POAC we do not need to balance a reaction and we can even add two or more reactions. This principle can be understood by the following example.

Consider the decomposition of $KCIO_3(s) \rightarrow KCI(s) + O_2(g)$ (unbalanced chemical reaction)

Apply the principle of atom conservation (POAC) for K atoms. or moles of K atoms in KCIO₃ = moles of K atoms in KCI Now, since 1 molecule of KCIO₃ contains 1 atom of K Thus, moles of K atoms in KCIO₃ = 1 × moles of KCIO₃ and moles of K atoms in KCI = 1 × moles of KCI

 \therefore moles of KCIO₃ = moles of KCI or

 $\frac{\text{wt.ofKCIO}_3 \text{ ing}}{\text{mol.wt.ofKCIO}_3} = \frac{\text{wt.ofKCI ing}}{\text{mol.wt.ofKCI}}$

The above equation gives the mass-mass relationship between KCIO₃ and KCI which is important in stoichiometric calculations. Again, applying the principle of atom conservation for O atoms, moles of O in KCIO₃ = 3 × moles of KCIO₃

moles of O in $O_2 = 3 \times \text{moles of P}$ moles of O in $O_2 = 2 \times \text{moles of } O_2$

 \therefore 3 × moles of KClO₃ = 2 × moles of O₂

or
$$3 \times \frac{\text{wt. of KCIO}_3}{\text{mol. wt. of KCIO}_3} = 2 \times \frac{\text{vol. of O}_2 \text{ at 1 atm and 0°C}}{\text{Molar vol. (22.4 lt)}}$$

- The above equations thus gives the mass-volume relationship of reactants and products.
- Ex. Calcium phosphide $Ca_{3}P_{2}$ formed by reacting magnesium with excess calcium orthophosphate $Ca_{3}(PO_{4})_{2}$, was hydrolysed by excess water. The evolved phosphine PH_{3} was burnt in air to yield phosphrous pentoxide ($P_{2}O_{5}$). How many gram of magnesium metaphosphate would be obtain if 192 gram Mg were used (Atomic weight of Mg = 24, P = 31)

$$Ca_{3}(PO_{4})_{2} + Mg \longrightarrow Ca_{3}P_{2} + MgO$$

$$Ca_{3}P_{2} + H_{2}O \longrightarrow Ca(OH)_{2} + PH_{3}$$

$$PH_{3} + O_{2} \longrightarrow P_{2}O_{5} + H_{2}O$$

$$MgO + P_{2}O_{5} \longrightarrow Mg(PO_{3})_{2}$$

magnesium metaphosphate.

Sol. POAC on Mg

$$1 \times n_{Mg} = 1 \times n_{Mg(PO_3)_2}$$
; $\frac{192}{24} = n_{Mg(PO_3)_2}$

$$W_{Mg(PO_3)_2} = \frac{192}{24} \times 182gm = 1456 \text{ gm}.$$

- Ex. 27.6 g K_2 CO₃ was treated by a series of reagents so as to convert all of its carbon to K_2 Zn₃[Fe(CN)₆]₂. Calculate the weight of the product. [mol. wt. of K_2 CO₃ = 138 and mol. wt. of K_2 Zn₃ [Fe(CN)₆]₂ = 698]
- Sol. Here we have no knowledge about series of chemical reactions but we know about initial reactant and final product, accordingly.

 $K_2CO_3 \xrightarrow{\text{Several}} K_2Zn_3[Fe(CN)_6]_2$ Since C atoms are conserved, applying POAC for C atoms, moles of C in K₂CO₃ = moles of C in K₂Zn₃[Fe(CN)₆]₂ 1 × moles of K₂CO₃ = 12 × moles of K₂Zn₃[Fe(CN)₆]₂ (∵ 1 mole of K₂CO₃ contains 1 moles of C) $\underbrace{\text{wt.of } K_2CO_3}_{\text{wt.of the product}} = 12 \times \text{wt.of the product}$

$$\overline{\text{mol. wt. of } K_2 CO_3} = 12 \times \frac{12 \times 1000}{\text{mol. wt. of product}}$$

wt. of
$$K_2 Zn_3 [Fe(CN)_6]_2 = \frac{27.6}{138} \times \frac{698}{12} = 11.6 g$$

Oxidation & Reduction

Let us do a comparative study of oxidation and reduction :

Oxidation	Reduction
1. Addition of Oxygen	1. Removal of Oxygen
e.g. $2Mg + O_2 \rightarrow 2MgO$	e.g. CuO + C \rightarrow Cu + CO
2. Removal of Hydrogen	2. Addition of Hydrogen
e.g. $H_2S + CI_2 \rightarrow 2HCI + S$	e.g. S + $H_2 \rightarrow H_2S$
3. Increase in positive charge e.g. $Fe^{2+} \rightarrow Fe^{3+} + e^{-}$	3. Decrease in positive charge e.g. Fe³+ + e⁻ → Fe²+
4. Increase in oxidation number	4. Decrease in oxidation number
(+2) (+4)	(+7) (+2)
e.g. $SnCl_2 \rightarrow SnCl_4$	e.g. $MnO_4^- \rightarrow Mn^{2+}$
5. Removal of electron	5. Addition of electron
e.g. $Sn^{2+} \rightarrow Sn^{4+} + 2e^{-}$	e.g. Fe³+ + e⁻ → Fe²+

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

In case of

• Oxygen atom :

In general and as well as in its oxides , oxygen atom has oxidation number equal to -2.

- (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 (CI, Br, I) have oxidation number equal to -1. But if halogen atom is attached with a more electronegative atom than halogen atom, then it will show positive oxidation numbers.

e.g.
$$\mathsf{K} \overset{+5}{\mathsf{ClO}_3}$$
, $\overset{+5}{\mathsf{HIO}_3}$, $\overset{+7}{\mathsf{HCIO}_4}$, $\overset{+5}{\mathsf{KBrO}_3}$

• Metals :

(a) Alkali metal (Li , Na, K, Rb,) always have oxidation number +1

(b) Alkaline earth metal (Be, Mg, Ca......) always have oxidation number +2.

(c) Aluminium always has +3 oxidation number

Note : Metal may have negative or zero oxidation number

- Oxidation number of an element in free state or in allotropic forms is always zero
 - e.g. $\overset{0}{O_2}, \overset{0}{S_8}, \overset{0}{P_4}, \overset{0}{O_3}$
- Sum of the oxidation numbers of atoms of all elements in a molecule is zero.
- Sum of the oxidation numbers of atoms of all elements in an ion is equal to the charge on the ion .
- If the group number of an element in modern periodic table is n, then its oxidation number may vary from (n-10) to (n-18) (but it is mainly applicable for p-block elements).

e.g. N- atom belongs to $15^{\rm th}$ group in the periodic table, therefore as per rule, its oxidation number may vary from

 $-3 \text{ to } +5 \ (\ \overset{-3}{\mathsf{NH}}_3, \overset{+2}{\mathsf{NO}}, \overset{+3}{\mathsf{N}}_2\mathsf{O}_3, \overset{+4}{\mathsf{NO}}_2, \overset{+5}{\mathsf{N}}_2\mathsf{O}_5)$

• The maximum possible oxidation number of any element in a compound is never more than the number of electrons in valence shell.(but it is mainly applicable for p-block elements)

Calculation of average oxidation number :

Solved Examples -

Ex.	Calcu	Ilate oxidation number of underlined element :
		(a) Na ₂ S ₂ O ₃ (b) Na ₂ S ₄ O ₆
Sol.	(a)	Let oxidation number of S-atom is x. Now work accordingly with the rules given before .
		$(+1) \times 2 + (x) \times 2 + (-2) \times 3 = 0$
		x = + 2
	(b)	Let oxidation number of S-atom is x
		$\therefore \qquad (+1) \times 2 + (x) \times 4 + (-2) \times 6 = 0 \implies x = +2.5$
	0	It is important to note here that Na ₂ S ₂ O ₃ have two S-atoms and there are four S-atom in Na ₂ S ₄ O ₆
	Howe	ever none of the sulphur atoms in both the compounds have + 2 or + 2.5 oxidation number, it is the
	avera	ge of oxidation number, which reside on each sulphur atom. Therefore, we should work to calculate the
	individ	Jual oxidation number of each sulphur atom in these compounds.
		•

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. $KMnO_4$, $K_2Cr_2O_7$, HNO_3 , conc. H_2SO_4 etc are powerful oxidising agents.

• 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 , $Na_2S_2O_3$ 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.

e.g. $10 \stackrel{+2}{\text{Fe}} SO_4 + 2 \stackrel{+5}{\text{KMn}} O_4 + 8 H_2 SO_4 \longrightarrow 5 \stackrel{+3}{\text{Fe}}_2 (SO_4)_3 + 2 \stackrel{+2}{\text{Mn}} SO_4 + K_2 SO_4 + 8 H_2 O_4 +$

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 :

$$2H_{2}^{-1}O_{2}(aq) \longrightarrow 2H_{2}^{-2}O(\ell) + \overset{0}{O}_{2}(g)$$

$$\overset{0}{S_{8}}(s) + 12OH^{-}(aq) \longrightarrow 4\overset{-2}{S^{2-}}(aq) + 2\overset{+2}{S_{2}}\overset{-2}{O_{3}^{2-}}(aq) + 6H_{2}O(\ell)$$

$$\overset{0}{Cl_{2}}(g) + 2OH^{-}(aq) \longrightarrow C\overset{+1}{IO^{-}}(aq) + \overset{-1}{Cl^{-}}(aq) + H_{2}O(\ell)$$

Consider the following reactions :

 $2\text{KCIO}_3 \longrightarrow 2\text{KCI} + 3\text{O}_2$ (a)

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

(b) $NH_4NO_2 \longrightarrow N_2 + 2H_2O$ 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.

(c)
$$4KCIO_3 \longrightarrow 3KCIO_4 + KCI$$

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

List of some important disproportionation reactions

1.
$$H_2O_2 \longrightarrow H_2O + O_2$$

2.
$$X_2 + OH^-(dil.) \longrightarrow X^- + XO^-$$
 (X = Cl, Br, I)

 $X_2 + OH^{-}(conc.) \longrightarrow X^{-} + XO_2^{-}$ 3.

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

- 4. $(CN)_{2} + OH^{-} \longrightarrow CN^{-} + OCN^{-}$
- $P_{4} + OH^{-} \longrightarrow PH_{3} + H_{2}PO_{2}^{-}$ $S_{8} + OH^{-} \longrightarrow S^{2-} + S_{2}O_{3}^{2-}$ 5.
- 6.
- $\dot{MnO_4^{2-}} \longrightarrow MnO_4^{-} + \dot{MnO_2}$ 7.
- 8. Oxyacids of Phosphorus (+1, +3 oxidation number)

$$H_3PO_2 \longrightarrow PH_3 + H_3PO_3$$

$$H_{3}PO_{3} \longrightarrow PH_{3} + H_{3}PO_{3}$$

9. Oxyacids of Chlorine(Halogens)(+1, +3, +5 Oxidation number)

 $CIO^{-} \longrightarrow CI^{-} + CIO_{2}^{-}$

$$CIO_2^- \longrightarrow CI^- + CIO_3^-$$

$$CIO_{3}^{-} \longrightarrow CI^{-} + CIO_{4}^{-}$$

- 10. $HNO_2 \longrightarrow NO + HNO_3$
- 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.

$$I^{-} + IO_{3}^{-} + H^{+} \longrightarrow I_{2} + H_{2}O$$

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

lon 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 adviced to follow the following steps to balance the redox reactions by lon electron method in acidic medium

Ex. Balance the following redox reaction :

 $FeSO_4 + KMnO_4 + H_2SO_4 \longrightarrow Fe_2(SO_4)_3 + MnSO_4 + H_2O + K_2SO_4$

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

$$\overset{+2}{\mathsf{Fe}} \overset{+6-2}{\mathsf{SO}_4} + \overset{+1}{\mathsf{K}} \overset{+7-2}{\mathsf{MnO}_4} + \overset{+1}{\mathsf{H}_2} \overset{+6-2}{\mathsf{SO}_4} \longrightarrow \overset{+3}{\mathsf{Fe}_2} \overset{+6-2}{(\mathsf{SO}_4)_3} + \overset{+2}{\mathsf{Mn}} \overset{+6-2}{\mathsf{SO}_4} + \overset{+1}{\mathsf{H}_2} \overset{-2}{\mathsf{O}_4}$$

Step II :

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

$$Fe^{2+} + MnO_4^- \longrightarrow Fe^{3+} + Mn^{2+}$$

Step III :

. –

Now identify the oxidation / reduction occuring in the reaction

undergoes reduction.

$$Fe^{2+} + MnO_4^- \rightarrow Fe^{3+} + Mn^{2+}$$

undergoes oxidation.

Step IV : Spilt the lonic reaction in two half, one for oxidation and other for reduction.

$$Fe^{2+} \xrightarrow{\text{oxidation}} Fe^{3+} MnO_4^- \xrightarrow{\text{Reduction}} Mn^{2+}$$

Step V :

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

$$Fe^{2+} \longrightarrow Fe^{3+} MnO_4^{-} \longrightarrow Mn^{2+}$$

Fe & Mn atoms are balanced on both side.

Step VI :

Now balance O & H atom by $H_2O \& H^+$ respectively by the following way: For one excess oxygen atom, add one H_2O on the other side and two H^+ on the same side.

$Fe^{2+} \longrightarrow Fe^{3+}$	(no oxygen atom)	(i)
$8H^+ + MnO_4^- \longrightarrow$	$Mn^{2+} + 4H_{2}O$	(ii)

Step VII:

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

 $Fe^{2+} \xrightarrow{\text{oxidation}} Fe^{3+} + e^{-}$ (1)

 $5e^- + 8H^+ + MnO_4^- \xrightarrow{\text{Reduction}} Mn^{2+} + 4H_2O$ (2)

Step VIII:

Sol.

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 :

 $\underline{5e^- + 8H^+ + MnO_4^- \longrightarrow Mn^{2+} + 4H_2O} \quad(2) \times 1$

 $5Fe^{2+} + 8H^+ + MnO_4^- \longrightarrow 5Fe^{3+} + Mn^{2+} + 4H_2O$

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

Step IX :

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

Now, by some manipulation, you will get :

$$5 \operatorname{FeSO}_4 + \operatorname{KMnO}_4 + 4\operatorname{H}_2\operatorname{SO}_4 \longrightarrow \frac{5}{2} \operatorname{Fe}_2 (\operatorname{SO}_4)_3 + \operatorname{MnSO}_4 + 4\operatorname{H}_2\operatorname{O} + \frac{1}{2} \operatorname{K}_2\operatorname{SO}_4$$

or $10FeSO_4 + 2KMnO_4 + 8H_2SO_4 \longrightarrow 5Fe_2(SO_4)_3 + 2MnSO_4 + 8H_2O + K_2SO_4.$

Balancing in basic medium :

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

- **Ex.** Balance the following redox reaction in basic medium :
 - $CIO^- + CrO_2^- + OH^- \longrightarrow CI^- + CrO_4^{2-} + H_2O$ By using upto step V, we will get :

$$\overset{+1}{\mathsf{CI}} \overset{\mathsf{O}^{-}}{\underset{\mathsf{CI}}{\overset{\mathsf{H}^{-}}{\overset{\mathsf{Re} \ \mathsf{duction}}{\overset{\mathsf{CI}^{-}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}{\overset{\mathsf{O}^{-}}}}}}}}}}}}}}}}}}}}}}}}}}}}}$$

Now, students are advised to follow step VI to balance 'O' and 'H' atom.

$$2H^+ + CIO^- \longrightarrow CI^- + H_2O$$
 | $2H_2O + CrO_2^- \longrightarrow CrO_4^{2-} + 4H^+$

O Now, since we are balancing in basic medium, therefore add as many as OH⁻ on both side of equation as there are H⁺ ions in the equation.

 $2OH^{-} + 2H^{+} + CIO^{-} \longrightarrow CI^{-} + H_{2}O + 2OH^{-}$ Finally you will get $H_{2}O + CIO^{-} \longrightarrow CI^{-} + 2OH^{-}$ Now see equation (i) and (ii) in which O and H atoms are balanced by OH^{-} and H_{2}O
Now from step VIII $2e^{-} + H_{2}O + CIO^{-} \longrightarrow CI^{-} + 2OH^{-}$ $(i) \times 3$ $4OH^{-} + CrO_{2}^{-} \longrightarrow CrO_{4}^{2-} + 2H_{2}O \dots (i) \times 3$ $4OH^{-} + CrO_{2}^{-} \longrightarrow CrO_{4}^{2-} + 2H_{2}O + 3e^{-}$ $(ii) \times 2$ $Adding : 3CIO^{-} + 2CrO_{2}^{-} + 2OH^{-} \longrightarrow 3CI^{-} + 2CrO_{4}^{2-} + H_{2}O$

SOLUTIONS

A solution is a homogenous mixture of two or more pure substances whose composition may be altered within certain limits. Though the solution is homogenous in nature, yet it retains the properties of its constituents.

Generally solution is composed of two components, solute and solvent. Such type of solution is known as binary solutions.

Solvent is that component in solution whose physical state is the same as that of the resulting solution while other component is called as solute. If the physical state of both component is same, than the component in excess is known as solvent and other one is called as solute. Each component in a binary solution can be in any physical state such as liquid, solid and gaseous state.

CONCENTRATION TERMS :

The concentration of a solution is the amount of solute dissolved in a known amount of the solvent or solution. Solution can be described as dilute or concentrated solution as per their concentration. A dilute solution has a very small quantity of solute while concentrated solution has a large quantity of solute in solution. Various concentration terms are as follows.

Mass percentage :

It may be defined as the number of parts of mass of solute per hundred parts by mass of solution.

% by mass
$$\left(\frac{W}{W}\right)$$
 := $\frac{Wt. \text{ of solute}}{Wt. \text{ of solution}} \times 100$

[X % by mass means 100 gm solution contains X gm solute ; \therefore (100 – X) gm solvent] Mass-volume percentage (W/V %):

It may be defined as the mass of solute present in 100 cm³ of solution. For example, If 100 cm³ of solution contains 5 g of sodium hydroxide, than the mass-volume percentage will be 5% solution.

$$\% \left(\frac{w}{V}\right) = \frac{wt. \text{ of solute}}{\text{volume of solution}} \times 100 \text{ [for liq. solution]}$$

$$[X \% \left(\frac{w}{V}\right)]$$
 means 100 ml solution contains X gm solute]

Volume Percent :

It can be represented as % v/v or % volume and used to prepare such solutions in which both components are in liquids state. It is the number of parts of by volume of solute per hundred parts by volume of solution. Therefore,

$$\% \left(\frac{v}{V}\right) = \frac{volume \text{ of solute}}{volume \text{ of solution}} \times 100$$

Mole % = \frac{\text{Moles of solute}}{\text{Total moles}} \times 100

For gases % by volume is same as mole %

Mole Fraction (X) :

Mole fraction may be defined as the ratio of number of moles of one component to the total number of moles of all the components (solute and solvent) present in solution. It is denoted by letter X and the sum of all mole fractions in a solution always equals one.

Mole fraction (X) =
$$\frac{\text{Moles of solute}}{\text{Total moles}}$$

Mole fraction does not depend upon temperature and can be extended to solutions having more than two components.

Molarity (M) :

Molarity is most common unit for concentration of solution. It is defined as the number of moles of solute present in one litre or one dm³ of the solution or millimol of solute present in one mL of solution.

Molarity (M) = $\frac{\text{Mole of solute}}{\text{volume of solution in litre}}$

Molality (m) : The number of gram mole of the solute present in 1000 g of the solvent is known as molality of solution. It represented by letter 'm'.

Molality (m) = $\frac{\text{Moles of solute}}{\text{Mass of solvent (in kg)}}$

The unit of molality is mol/kg and it does not effect by temperature.

Parts per million (ppm): The very low concentration of solute in solution can be expressed in ppm. It is the numbers of parts by mass of solute per million parts by mass of the solution.

Parts per million (ppm) = $\frac{\text{Mass of solute}}{\text{Mass of solvent}} \times 10^6 \cong \frac{\text{Mass of solute}}{\text{Mass of solution}} \times 10^6$

Get yourselves very much confortable in their inter conversion. It is very handy.

Concentration Type	Mathematical Formula	Concept
Percentage by mass	$\% \left(\frac{w}{w}\right) = \frac{\text{Mass of solute} \times 100}{\text{Mass of solution}}$	Mass of solute present in 100 gm of solution.
Volume percentage	$\% \left(\frac{v}{v}\right) = \frac{\text{Volume of solute} \times 100}{\text{Volume of solution}}$	Volume of solute present in 100 cm ³ of solution.
Mass-volume percentage	$\% \left(\frac{w}{v}\right) = \frac{\text{Mass of solute} \times 100}{\text{Volume of solution}}$	Mass of solute present in 100 cm ³ of solution.
Parts per million	$ppm = \frac{Mass of solute \times 10^6}{Mass of solution}$	Parts by mass of solute per million parts by mass of the solution
Mole fraction	$X_A = \frac{\text{Mole of A}}{\text{Mole of A} + \text{Mole of B} + \text{Mole of C} + \dots}$	Ratio of number of
	$X_{B} = \frac{\text{Mole of B}}{\text{Mole of A} + \text{Mole of B} + \text{Mole of C} + \dots}$	moles of one component
		to the total number of moles.

Mole concept

	Molarity	$M = \frac{Mole \text{ of solute}}{Volume \text{ of solution}(in L)}$		L)	Moles of solute		
					in one litre of solution.		
	Molality	m= Molar n	Mass of solute hass of solute×N	\times 1000 fass of solvent(g)	Moles of solute in one		
					kg of solvent		
Ex.	Calculate the mole frac water ?(M (water) = 18	tions of the co ; M (glycero	omponents of the ol) = 92)	e solution composed	by 92 g glycerol and 90 g		
Sol.	Moles of water = 90 g / 1 Moles of glycerol = 92 g Total moles in solution = Mole fraction of water = 5	8 g = 5 mol wa / 92 g = 1 mol g 5 + 1 = 6 mol 5 mol / 6 mol =	iter glycerol 0.833				
	Mole fraction of glycerol	= 1 mol / 6 mol	= 0.167				
Ex. Sol.	What will be the Molarity of solution when water is added to 10 g CaCO_3 to make 100 mL of solution? Mol of $CaCO_3 = 10 / 100 = 0.1$ Molarity = Mole of solute / Volume of solution (L) = 0.10 mol / 0.10 L Therefore : Molarity of given solution = 1.0 M						
Ex. Sol.	Calculate the molality of a solution containing 20 g of sodium hydroxide (NaOH) in 250 g of water? Moles of sodium hydroxide = 20 / 40 = 0.5 mol NaOH 250 gm = 0.25 kg of water Hence molality of solution = Mole of solute / Mass of solvent (kg)= 0.5 mol / 0.25 kg or Molality(m) = 2.0 m						
Ex.	Calculate the grams of	copper sulph	ate (CuSO ₄) need	ed to prepare 250.0 m	L of 1.00 M CuSO ₄ ?		
Sol.	Moles of $CuSO_4 = M \times V$ Molar mass of copper sul	$r' = 1 \times \frac{250}{1000}$ lphate = 159.6	g/mol - Moles of CuSO	× Molar mass of conn	ar sulphata		
			$= 1 \times \frac{250}{1000} \times \frac{2}{1000} \times \frac{2}{100$	159.6 g/mol			
			= 39.9 gm of C	opper sulphate			
Ex.	How many grams of H ₂	SO₄ are prese	ent in 500 ml of 0.	2MH ₂ SO ₄ solution ?			
Sol.	$M = \frac{\text{moles}}{\text{vol.}} \Rightarrow \text{moles of}$	$fH_2SO_4 = M$	\times V = 0.2 \times $\frac{500}{1000}$	$\frac{1}{2}L = 0.1$			
	Mass of $H_2SO_4 = 0.1 \times 9$	9.8 g					
Ex.	Calculate the ppm of m	nercury in wat	er in given samp	le contain 30 mg of He	g in 500 ml of solution.		
Sol.	Parts per million = $\frac{Mass}{Ma}$	of solute $\times 1$ ass of solution	$\frac{0^6}{n}$				
	Mass of Hg = 30 mg						

Mass of water = $500/1 = 500g = 50 \times 10^4 \text{ mg}$ (density = mass / volume ; density of water 1 g / ml) $W = \frac{V}{d}$ Therefore, ppm of mercury = $\frac{30 \times 10^6}{50 \times 10^4}$ = 60 ppm of mercury **MIXING OF SOLUTIONS :** It is based on law of conservation of moles. M₁ M_2 Two solutions having same solute (i) = + V_1 ٧, Final molarity = $\frac{\text{Total moles}}{\text{Total volume}} = \frac{M_1V_1 + M_2V_2}{V_1 + V_2}$ NaCl NaCl $V_1 + V_2$ NaCl (ii) Dilution Effect : When a solution is diluted, the moles of solute do not change but molarity changes while on taking out a small volume of solution from a larger volume, the molarity of solution do not change but moles change proportionately. Final molarity = $\frac{M_1V_1}{V_1 + V_2}$ M_1 = V_1 H₂O NaCl n-fold or n-times dilution Final volume = $V_1 + V_2 = n(V_1)$ ⇒ 50 ml 0.2 M H_2SO_4 is mixed with 50 ml 0.3M H_2SO_4 . Find molarity of final solution. Ex. $M_{f} = \frac{\text{Total moles of } H_{2}\text{SO}_{4}}{\text{Total volume}} = \frac{50 \times 0.2 \times 10^{-3} + 50 \times 10^{-3} \times 0.3}{(50 + 50) \times 10^{-3}} = \boxed{0.25 \text{ M}}$ Sol. Find final molarity in each case : Ex. Sol. (i) 500 ml 0.1 M HCI + 500 ml 0.2M HCI $M_{f} = \frac{500 \times 0.1 + 500 \times 0.2}{500 + 500} = 0.15 \,\text{M}$ 50 ml 0.1M HCl + 150 ml 0.3MHCl + 300 ml H₂O (ii) $M_{f} = \frac{50 \times 0.1 + 150 \times 0.3}{50 + 150 + 300} = \frac{50}{500} = 0.1 \text{ M}$ (iii) 4.9g H₂SO₄ + 250 ml H₂O + 250 ml 0.1 M H₂SO₄ $M_{f} = \frac{\frac{4.9}{98} + \frac{250}{1000} \times 0.1}{\left(\frac{250 + 250}{1000}\right)} = \frac{50 + 25}{500} = \boxed{0.15 \text{ M}}$

Ex. How much water should be added to 2M HCI solution to form 1 litre of 0.5 M HCI ?

Sol. Let V be initial volume

Then mol of HCI = constant

 $2 \times V = 1 \times 0.5 \Longrightarrow V = 0.25 L$

Volume of water added = 1 - 0.25 = 0.75 L

- Ex. Find number of Na⁺ & PO₄⁻³ ions in 250 ml of 0.2M Na₃PO₄ solution.
- **Sol.** $Na_3PO_4 + aq. \longrightarrow 3Na^+(aq) + PO_4^{-3}(aq)$ [lonic compound when added to water ionize completely]. 50 millimoles (m.m.) 150 mm 50 mm No. of Na⁺ ions = 150 × 10⁻³ × N_A; No. of PO₄^{-3} ions = 50 × 10⁻³ × N_A
- Ex. 1.11g $CaCl_2$ is added to water forming 500 ml of solution. 20 ml of this solution is taken and diluted 10 folds. Find moles of C Γ ions in 2 ml of diluted solution.



- Ex. What volumes of 1M & 2M H_2SO_4 solution are required to produce 2L of 1.75M H_2SO_4 solution?
- **Sol.** Let XL be vol. of 1M solution.

 \therefore (2 – X)L is vol. of 2M solution.

Moles of $H_2SO_4 = 2 \times 1.75 = 1(X) + (2 - X)2$

$$3.5 = 4 - X$$
; X = 0.5 L

i.e. 0.5L of 1M & 1.5 L of 2M solution required.

Ex. 80g NaOH was added to 2L water. Find molality of solution if density of water = 1g/mL

- **Sol.** $m = \frac{\text{moles of NaOH}}{\text{mass of H}_2\text{O}} \times 1000 = \frac{80/40}{2 \times 1000} \times 1000 = \boxed{1 \text{ molal}}$
- Ex. A 100g NaOH solution has 20g NaOH. Find molality.

Sol.
$$m = \frac{20/40}{100-20} \times 1000 = \frac{500}{80} = \frac{6.25 \text{ mol/kg}}{6.25 \text{ mol/kg}}$$

Ex. Find molality of aqueous solution of CH_3COOH whose molarity is 2M and density d = 1.2 g/mL.

Hint :
$$\frac{1000 \times M}{1000 \times d - MMs}$$

where d = density in gL^{-1} , M = Molarity, m = molality, M_s = molar mass of solute.

Sol.
$$m = \frac{2}{1200 - 2 \times 60} \times 1000 = \boxed{1.85 m}$$

Ex. A solution is made by mixing 300 ml 1.5M $Al_2(SO_4)_3 + 300$ ml 2M $CaSO_4 + 400$ ml 3.5M $CaCl_2$ Find final molarity of (1) SO_4^{-2} , (2) Ca^{2+} , (3) $C\Gamma$. [Assume complete dissociation of these compounds].

Sol. (1)
$$[SO_4^{-2}]_f = \frac{\text{Total moles}}{\text{Total volume}} = \frac{300 \times 1.5 \times 10^{-3} \times 3 + 300 \times 2 \times 10^{-3}}{(300 + 300 + 400) \times 10^{-3}} = 1.95\text{ M}$$

(2)
$$[Ca^{+2}]_{f} = \frac{300 \times 2 + 400 \times 3.5}{1000} = 2M$$

(3) $[CI]_{f} = \frac{400 \times 3.5 \times 2}{1000} = 2.8M$

Ex. A solution has 80% $\frac{W}{W}$ NaOH with density 2gL⁻¹. Find (a) Molarity (b) Molality of solution.

Sol. Let V_{lit} be vol. of solution

Mass of solute =
$$(d \times V) \times \frac{\binom{\% \frac{W}{W}}{100}}{100} = 2 \times V \times \frac{80}{100} = 1.6V$$

(a) $M = \frac{1.6V/40}{V} = \boxed{0.04M}$ (b) $m = \frac{1.6V/40}{2V - 1.6V} \times 1000 = \boxed{100 \text{ mol kg}^{-1}}$

Ex. 4.450 g 100 per cent sulphuric acid was added to 82.20 g water and the density of the solution was found to be 1.029 g/cc at 25°C and 1 atm pressure. Calculate (a) the weight percent, (b) the mole fraction, (c) the mole percent, (d) the molality, (e) the molarity of sulphuric acid in the solution under these conditions.

Sol. Sulphuric acid = 4.450 g , Water = 82.20 g
$$\Rightarrow$$
 Wt. of solution = 86.65 g

 \therefore Density of solution = 1.029 g/cc.

(a) Weight percent =
$$\frac{\text{wt. of solute}}{\text{wt. of solution}} \times 100 = \frac{4.450}{86.65} \times 100 = 5.14$$

(b) Mole fraction :

Mole of solute =
$$\frac{\text{wt. of solute}}{\text{mol wt. of solute}} = \frac{4.45}{98} = 0.0454$$

$$\text{Mole of solvent} = \frac{82.20}{18} = 4.566$$

Total moles in solution = 0.0454 + 4.566 = 4.6114

Mole fraction of solute
$$=\frac{0.0454}{4.6114} = 0.0098$$

(c) Mole percent = $\frac{\text{moles of solute}}{\text{Total moles in solution}} \times 100$

= mole fraction of solute × 100 = 0.0098 × 100 = 0.98

(d) Molality=
$$\frac{\text{moles of solute}}{\text{mass of solvent (in gm)}} \times 1000$$

= $\frac{0.0454 \times 1000}{0.0454 \times 1000} = 0.552$

(e) Molarity $=\frac{\text{moles of solute}}{1000}$

litre of solution

Volume of solution = $\frac{Mass}{Density} = \frac{86.65}{1.029} ml$

$$=\frac{86.65}{1.029\times1000}$$
 litre

Molarity =
$$\frac{0.0454}{\frac{86.54}{1.029 \times 1000}} = \frac{0.0454 \times 1000 \times 1.029}{86.65} = 0.539$$

Ex. A solution of KCl has a density of 1.69 g mL^{-1} and is 67% by weight. Find the density of the solution if it is diluted so that the percentage by weight of KCl in the diluted solution is 30%.

Sol. Let the volume of the KCl solution be 100 mL,

100 g of solution contains = 67 g of KCl

169 g of solution =
$$\frac{67}{100} \times 169 = 113.23$$
 g

Lex x mL of H_2O be added.

New volume of solution = (100 + x) mL

New weight of solution = (169 + x) g

(Since x mL of $H_2O = x g$ of H_2O , $d_{H_2O} = 1$)

New percentage of the solution = 30%

% by weight =
$$\frac{\text{weight of solute} \times 100}{\text{weight of solution}}$$

$$30 = \frac{113.23}{(169 + x)} \times 100$$

x = 208.43 mL = 208.43 g

New density
$$= \frac{\text{New weight of solution}}{\text{New volume of solution}}$$
$$= \frac{(169 + x)}{(100 + x)}$$
$$\frac{(169 + 208.43)}{(100 + 208.43)} = \frac{377.43}{308.43}$$

∴ d=1.224

Quantitative Estimation of Elements

PERCENTAGE DETERMINATION OF ELEMENTS IN ORGANIC COMPOUNDS :

All these methods are applications of POAC

Do not remember the formulas, derive them using the concept, its easy. **Liebig's method :** (*for Carbon and hydrogen*)

(w)Organic Compound
$$\xrightarrow{\Delta}_{CuO}$$
 (w₁)CO₂ + H₂O(w₂)

% of C =
$$\frac{w_1}{44} \times \frac{12}{w} \times 100$$

% of H =
$$\frac{W_2}{18} \times \frac{2}{W} \times 100$$

where $w_1 = wt.$ of CO₂ produced, $w_2 = wt.$ of H₂O produced,

w = wt. of organic compound taken

Estimation of nitrogen : There are two methods for the estimation of nitrogen

(i) Dumas method

(ii) Kjedahl's method.

(i) Duma's method :

A known mass of compound is heated with copper oxide (CuO) in an atomsphere of CO_2 , which gives free nitrogen along with CO_2 and H_2O .

 $C_xH_vN_z + (2x + y/2) CuO \rightarrow xCO_2 + y/2 (H_2O) + z/2 (N_2) + (2x + y/2) Cu.$

The gaseous mixture is passed over a heated copper gauze which converts traces of nitrogen oxides formed to N_2 . The gaseous mixture is collected over an aqueous solution of KOH which absorbs CO_2 , and nitrogen is collected in the upper part of the graduated tube.

(w) Organic Compound $\xrightarrow{\Delta}$ N₂ \rightarrow (P, V, T given)

use PV = nRT to calculate moles of N_2 , n.

$$\therefore$$
 % of N = $\frac{n \times 28}{W} \times 100$

(ii) Kjeldahl's method :

A known mass of organic compound (0.5 gm) is mixed with K_2SO_4 (10 gm) and $CuSO_4$. (1.0 gm) or a drop of mercury (Hg) and conc. H_2SO_4 (25 ml), and heated in Kjeldahl's flask. $CuSO_4$ or Hg acts as a catalyst, while K_2SO_4 raises the boiling point of H_2SO_4 . The nitrogen in the organic compound is quantitatively converted to ammonium sulphate. The resulting mixture is then distilled with excess of NaOH solution and the NH₃ evolved is passed into a known but excess volume of standard HCI or H_2SO_4 . The acid left unused is estimated by titration with some standard alkali. The amount of acid used against NH₃ can thus be known and from this the percentage of nitrogen is calculated.

(a) C + H + S
$$\xrightarrow{\text{conc.}}_{\text{H}_2\text{SO}_4}$$
 CO₂ + H₂O + SO₂

(b) N
$$\xrightarrow{\text{conc.}}_{\text{H}_2\text{SO}_4}$$
 (NH₄)₂SO₄

(c)
$$(NH_4)_2SO_4 + 2NaOH \rightarrow Na_2SO_4 + 2NH_3 + 2H_2O_4$$

(d)
$$2NH_3 + H_2SO_4 \rightarrow (NH_4)_2SO_4$$

 $(w)O.C.+H_2SO_4 \rightarrow (NH_4)_2SO_4 \xrightarrow{NaOH} NH_3 + H_2SO_4 \rightarrow (molarity and volume (V litre) consumed given)$

$$\Rightarrow \qquad \% \text{ of } N = \frac{MV \times 2 \times 14}{W} \times 100$$

where M = molarity of H_2SO_4 . Some N containing compounds do not give the above set of reaction as in Kjeldahl's method.

Estimation of sulphur: A known mass of compound is heated with fuming HNO_3 or sodium peroxide (Na_2O_2) in the presence of $BaCl_2$ solution in Carius tube. Sulphur is oxidised to H_2SO_4 and precipitated as $BaSO_4$. It is filerted, dried and weighed.

$$(\mathsf{w}) \, \mathsf{O.C.} + \mathsf{HNO}_3 \mathop{\rightarrow} \mathsf{H}_2\mathsf{SO}_4 + \mathsf{BaCl}_2 \mathop{\rightarrow} (\mathsf{w}_1) \, \mathsf{BaSO}_4$$

$$\Rightarrow \qquad \% \text{ of } S = \frac{W_1}{233} \times \frac{1 \times 32}{W} \times 100\%$$

where $w_1 = wt$. of $BaSO_4$, w = wt. of organic compound

Estimation of phosphorous :

A known mass of compound is heated with fuming HNO₃ or sodium peroxide (Na_2O_2) in Carius tube which converts phosphorous to H_3PO_4 . Magnesia mixture ($MgCI_2 + NH_4CI$) is then added, which gives the precipitate of magnesium ammonium phosphate ($MgNH_4.PO_4$) which on heating gives magnesium pyrophosphate ($Mg_2P_2O_7$), which is weighed. O.C (w) + HNO₃ \rightarrow H₃PO₄ + [NH₃ + magnesia mix-

ture ammonium molybdate]

$$\rightarrow$$
 MgNH₄PO₄ $\xrightarrow{\Lambda}$ Mg₂P₂O₇(w₁)

% of P =
$$\frac{W_1}{222} \times \frac{2 \times 31}{W} \times 100$$

Estimation of halogens :

Carius method : A known mass of compound is heated with conc. HNO_3 in the presence of $AgNO_3$ contained in a hard glass tube known as carius tube in a furnce. C and H are oxidised to CO_2 and H_2O . The halogen forms the corresponding AgX. It is filtered, dried, and weighed.

O.C. + HNO₃ + AgNO₃ \rightarrow AgX If X is Cl then colour = white If X is Br then colour = dull yellow If X is I then colour = bright yellow Flourine can't be estimated by this

% of X = $\frac{\text{Atomic mass of X}}{\text{Molecular mass of AgX}} \times \frac{\text{Wt. of AgX}}{\text{Wt. of Organic halide}} \times 100$

Ex. A sample of 0.5 gm of an organic compound was treated according to Kjeldahl's method. The ammonia evolved was absorbed by 2.45 gm of H₂SO₄. The residual acid required solution containing 0.6 gm. NaOH for neutralisation. Find the percentage composition of nitrogen in the compound ?

m mol of H_2SO_4 used to react with NaOH = $\frac{0.6}{40}$ = 15mmol.

Remaining mmol of $H_2SO_4 = \frac{2.45}{98} \times 10^3 - 15 = 10$ mmol of NH₃ used = 10 × 2 = 20

% N in sample = $\frac{20 \times 10^{-3} \times 14}{0.5} \times 100 = 56\%$

Ex. Calculate the molar mass of a compound in the Dumas method at 100°C for which volume of experimental container was 452 ml and the pressure was 745.1 torr. The difference in mass between the empty container and the final measurement was 1.129 gm.

Sol.
$$n = \frac{PV}{RT} = \frac{745.1}{760} \times \frac{452 \times 10}{0.0821 \times 373} = 0.01448 \text{ mol}$$

molar mass (M) = $\frac{1.129}{0.01448}$ = 78.0 gm/mol .

MISCELLANEOUS SOLVED PROBLEMS (MSPS)

- Find the relative atomic mass, atomic mass of the following elements.
 (i) Na (ii) F (iii) H (iv) Ca (v) Ag
- **Sol.** (i) 23, 23 amu (ii) 19, 19 amu (iii) 1, 1.008 amu, (iv) 40, 40 amu, (v) 108, 108 amu.
- 2. A sample of (C_2H_6) ethane has the same mass as 10^7 molecules of methane. How many C_2H_6 molecules does the sample contain ?

Sol. Moles of $CH_4 = \frac{10^7}{N_A}$

Mass of
$$CH_4 = \frac{10^7}{N_A} \times 16 = \text{mass of } C_2H_6$$

So Moles of $C_2H_6 = \frac{10^7 \times 16}{N_A \times 30}$
So No. of molecules of $C_2H_6 = \frac{10^7 \times 16}{N_A \times 30} \times N_A = 5.34 \times 10^6$

3. From 160 g of SO₂ (g) sample, 1.2046×10^{24} molecules of SO₂ are removed then find out the volume of left over SO₂ (g) at STP.

Sol. Given moles =
$$\frac{160}{64}$$
 = 2.5.

Removed moles = $\frac{1.2046 \times 10^{24}}{6.023 \times 10^{23}}$ = 2. so left moles = 0.5. volume left at STP = 0.5 × 22.7 = 11.35 lit.

- 4. 14 g of Nitrogen gas and 22 g of CO₂ gas are mixed together. Find the volume of gaseous mixture at STP.
- **Sol.** Moles of $N_2 = \frac{14}{28} = 0.5$.

moles of $CO_2 = \frac{22}{44} = 0.5$. so total moles = 0.5 + 0.5 = 1. so vol. at STP = 1 × 22.7 = 22.7 lit.

- 5. Show that in the reaction $N_2(g) + 3H_2(g) \rightarrow 2NH_3(g)$, mass is conserved.
- $\begin{array}{lll} \mbox{Sol.} & \mbox{N}_2 \ (g) + 3 \mbox{H}_2 (g) \rightarrow \ 2 \mbox{NH}_3 \ (g) \\ \mbox{moles before reaction} & 1 & 3 & 0 \\ \mbox{moles after reaction} & 0 & 0 & 2 \\ \mbox{Mass before reaction} = \mbox{mass of 1 mole N}_2 (g) + \mbox{mass of 3 mole H}_2 (g) \\ & = \ 14 \ x \ 2 + 3 \ x \ 2 = \ 34 \ g \\ \mbox{mass after reaction} = \ \mbox{mass of 2 mole NH}_3 \\ & = \ 2 \ x \ 17 = \ 34 \ g. \end{array}$
- 6. When x gram of a certain metal burnt in 1.5 g oxygen to give 3.0 g of its oxide. 1.20 g of the same metal heated in a steam gave 2.40 g of its oxide. shows the these result illustrate the law of constant or definite proportion
- **Sol.** Wt. of metal = 3.0 1.5 = 1.5 g so wt. of metal : wt of oxygen = 1.5 : 1.5 = 1 : 1similarly in second case, wt. of oxygen = 2.4 - 1.2 = 1.2 g so wt. of metal : wt of oxygen = 1.2 : 1.2 = 1 : 1so these results illustrate the law of constant proportion.
- 7. Find out % of O & H in H₂O compound.

Sol. % of O =
$$\frac{16}{18} \times 100 = 88.89\%$$
 and % of H = $\frac{2}{18} \times 100 = 11.11\%$

8. Acetylene & butene have empirical formula CH & CH₂ respectively. The molecular mass of acetylene and butene are 26 & 56 respectively deduce their molecular formula.

Ans.
$$C_2H_2 \& C_4H_8$$

Sol. $n = \frac{Molecular mass}{\Gamma = 1}$

Sol. $n = \overline{Empirical formula mass}$

For Acetylene : $n = \frac{26}{13} = 2$ \therefore Molecular formula = $C_2 H_2$ For Butene : $n = \frac{56}{14} = 4$ \therefore Molecular formula = $C_4 H_8$.

9. An oxide of nitrogen gave the following percentage composition : N = 25.94 and O = 74.06Calculate the empirical formula of the compound.

Ans.
$$N_2O_5$$

Sol

Element	% / Atomic mass	Simple ratio	Simple intiger rat
Ν	$\frac{25.94}{14} = 1.85$	1	2
0	$\frac{74.06}{16} = 4.63$	2.5	5

So empirical formula is N_2O_5 .

10. Find the density of CO₂(g) with respect to N₂O(g).
Sol. R.D. =
$$\frac{M.wt.of CO_2}{M.wt.of N_2O} = \frac{44}{44} = 1$$
.
11. Find the vapour density of N₂O₅
Sol. V.D. = $\frac{Mol.wt.of N_2O_5}{2} = 54$.
12. Write a balance chemical equation for following reaction :
When ammonia (NH₃) decompose into nitrogen (N₂) gas & hydrogen (H₂) gas.
Sol. NH₃ → $\frac{1}{2}$ N₂ + $\frac{3}{2}$ H₂ or 2NH₃ → N₂ + 3H₂.
13. When 170 g NH₃ (M =17) decomposes how many grams of N₂ & H₂ is produced.
Sol. NH₃ → $\frac{1}{2}$ N₂ + $\frac{3}{2}$ H₂

$$\frac{molesof NH_3}{1} = \frac{molesof N_2}{1/2} = \frac{molesof H_2}{3/2}$$
.
So moles of N₂ = $\frac{1}{2} \times \frac{170}{17} = 5$. So wt. of N₂ = 5 × 28 = 140 g.
Similarly moles of H₂ = $\frac{3}{2} \times \frac{170}{17} = 15$.
So wt. of H₂ = 15 × 2 = 30 g.
14. 340 g NH₃ (M =17) when decompose how many litres of nitrogen gas is produced at STP.
Sol. NH₃ → $\frac{1}{2}$ N₂ + $\frac{3}{2}$ H₂
moles of N₃ = $\frac{340}{17} = 20$.
So moles of N₂ = $\frac{1}{2} \times 20 = 10$.
∴ vol. of N₂ at STP = 10 × 22.7 = 227 lit.
15. 4 mole of MgCO₃ is reacted with 6 moles of HCI solution. Find the volume of CO₂ gas proc

- **15.** 4 mole of $MgCO_3$ is reacted with 6 moles of HCI solution. Find the volume of CO_2 gas produced at STP, the reaction is
- $\label{eq:MgCO_3} \begin{array}{l} \mathsf{H} \mathsf{gCO_3} + \mathsf{2HCI} \to \mathsf{MgCI_2} + \mathsf{CO_2} + \mathsf{H_2O}. \\ \mbox{Sol.} & \mbox{Here HCI is limiting reagent. So moles of CO_2 formed = 3.} \\ & \mbox{So vol. at STP = 3 \times 22.7 = 68.1 lit.} \end{array}$
- 16. 117 g NaCl is dissolved in 500 ml aqueous solution. Find the molarity of the solution.

Sol. Molarity =
$$\frac{117/58.5}{500/1000}$$
 = **4M.**

- **17.** 0.32 mole of LiAlH₄ in ether solution was placed in a flask and 74 g (1 moles) of t-butyl alcohol was added. The product is LiAlHC₁₂H₂₇O₃. Find the weight of the product if lithium atoms are conserved. [Li = 7, Al = 27, H = 1, C = 12, O = 16]
- Sol. Applying POAC on Li $1 \times \text{moles of LiAlH}_4 = 1 \times \text{moles of LiAlH } C_{12}H_{27}O_3$ $254 \times 0.32 = 1 \times \text{wt. of LiAlH } C_{12}H_{27}O_3$. wt. of LiAlH $C_{12}H_{27}O_3 = 81.28 \text{ g}$.

18.	Balance the following equations :					
	(a) $H_2O_2 + MnO_4^- \longrightarrow Mn^{+2} + O_2$ (acidic medium)					
	(b) $Zn + HNO_3$ (dil) $\longrightarrow Zn(NO_3)_2 + H_2O + NH_4NO_3$					
	(c) $CrI_3 + KOH + CI_2 \longrightarrow K_2CrO_4 + KIO_4 + KCI + H_2O.$					
	(d) $P_2H_4 \longrightarrow PH_3 + P_4$ (e) $Ca_2(PO_4)_2 + SiO_2 + C \longrightarrow CaSiO_2 + P_4 + CO$					
Ans.	(a) $6H^{+} + 5H_{2}O_{2} + 2MnO_{4}^{-} \longrightarrow 2Mn^{+2} + 5O_{2}^{+} + 8H_{2}O_{4}$					
	(b) $4Zn + 10HNO_3$ (dil) $\xrightarrow{4} 4Zn(NO_3)_2 + 3H_2O + NH_4NO_3$					
	(c) $2CrI_3 + 64KOH + 27CI_2 \longrightarrow 2K_2CrO_4 + 6KIO_4 + 54KCI + 32H_2O.$					
	(d) $6P_2H_4 \longrightarrow 8PH_3 + P_4$					
	(e) $2Ca_{_3}(PO_{_4})_2 + 6SiO_2 + 10C \longrightarrow 6CaSiO_3 + P_4 + 10CO$					
19.	Calculate the resultant molarity of following:					
	(a) 200 ml 1M HCl + 300 ml water (b) 1500 ml 1M HCl + 18.25 g HCl					
_	(c) 200 ml 1M HCl + 100 ml $0.5 \text{ M H}_2\text{SO}_4$ (d) 200 ml 1M HCl + 100 ml 0.5 M HCl					
Ans.	(a) 0.4 M (b) 1.33 M (c) 1 M (d) 0.83 M.					
Sol	(a) Final molarity = $\frac{200 \times 1 + 0}{200} = 0.4 \text{ M}$					
0011	(a) Final molancy = $\frac{1}{200 + 300} = 0.4$ W.					
	1500×1 18.25×1000					
	(b) Final molarity = $\frac{36.5}{36.5}$ = 1.33 M					
	1500					
	(c) Final molarity of H ⁺ = $\frac{200 \times 1 + 100 \times 0.5 \times 2}{1 \times 100 \times 0.5 \times 2} = 1 \text{ M}$					
	200 + 100					
	(d) Final molarity $= \frac{200 \times 1 + 100 \times 0.5}{200 \times 1 + 100 \times 0.5} = 0.83 \text{ M}$					
	200 + 100 200 + 100					
20.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the solution					
	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the	ne solution.				
	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the second	ne solution.				
Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$.	ne solution.				
Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$.	ne solution.				
Sol. 21.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution.	ne solution. ubstance in				
Sol. 21. Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$.	ne solution. ubstance in				
Sol. 21. Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$.	ne solution. ubstance in				
Sol. 21. Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. so % (w/w) = $\frac{0.25}{6.50} \times 100 = 3.8\%$.	ne solution. ubstance in				
Sol. 21. Sol. 22.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. so % (w/w) = $\frac{0.25}{6.50} \times 100 = 3.8\%$. 518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the	ne solution. ubstance in ne solution.				
Sol. 21. Sol. 22.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. so % (w/w) = $\frac{0.25}{6.50} \times 100 = 3.8\%$. 518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the	ne solution. ubstance in ne solution.				
Sol. 21. Sol. 22. Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. so % (w/w) = $\frac{0.25}{6.50} \times 100 = 3.8\%$. 518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$.	ne solution. ubstance in ne solution.				
Sol. 21. Sol. 22. Sol. 23.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. so % (w/w) = $\frac{0.25}{6.50} \times 100 = 3.8\%$. 518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of the wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s	ne solution. ubstance in ne solution. ubstance in				
Sol. 21. Sol. 22. Sol. 23.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. so % (w/w) = $\frac{0.25}{6.50} \times 100 = 3.8\%$. 518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of th wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution.	ne solution. ubstance in ne solution. ubstance in				
Sol. 21. Sol. 22. Sol. 23. Sol.	518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of t wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$. $so \% (w/w) = \frac{0.25}{6.50} \times 100 = 3.8\%$. 518 g of an aqueous solution contains 18 g of glucose (mol.wt. = 180). What is the molality of th wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solvent = $518 - 18 = 500$ g. \Rightarrow so molarity = $\frac{18/180}{500/1000} = 0.2$. 0.25 of a substance is dissolved in 6.25 g of a solvent. Calculate the percentage amount of the s the solution. wt. of solution = $0.25 + 6.25 = 6.50$.	ne solution. ubstance in ne solution. ubstance in				

Exercise-1

> Marked Questions may have for Revision Questions.

PART - I : SUBJECTIVE QUESTIONS

MOLE-I : Law of Chemical Combination

Section (A) : Molar volume of ideal gases at STP, Average molar mass



- **A-1.** Calculate volume of H_2 gas kept at STP if it contains as many H atoms as in 98 gm H_3PO_4 . [Atomic mass of P = 31]
- **A-2.** 40 mg of gaseous substance (X₂) occupies 4.8 mL of volume at 1 atm and 27°C. Atomic mass of element X is : (R : 0.08 atm L/mole-K)
- A-3. A gaseous mixture of H_2 and N_2O gas contains 66 mass % of N_2O . What is the average molecular mass of mixture :

Section (B) : Empirical Formula, % Composition of a given compound by mass, % By mole, Minimum molecular mass determination.

Commit to memory :

The molecular formula is an integral multiple of the empirical formula.

- **B-1.** Haemoglobin contains 0.25% iron by mass. The molecular mass of of Haemoglobin is 89600 then the number of iron atoms per molecule of Haemoglobin (Atomic mass of Fe = 56) -
- **B-2.** A compound has 52.17 % carbon, 13.05 % hydrogen and 34.78 % oxygen. If molar mass of compound is 46, find number of H-atoms per molecule of the compound.
- **B-3.** The empirical formula of a compounds is CH_2O . 0.25 mole of this compound contains 1 gm hydrogen. The molecular formula of compound is -
- **B-4.** In a gaseous mixture 2mol of CO₂, 1 mol of H₂ and 2 mol of He are present than determine mole percentage of CO₂.

1

MOLE-II : Basic Stoichiometry

Section (C) : Stoichiometry, Equation based calculations (Elementary level single equation or 2)

Commit to memory :			
Now for any general balance chemic	cal equation like		
a A + b B -	\longrightarrow c C + d D		
you can write.			
Moles of A reacted	moles of B reacted	moles of C produced	moles of D produced
a	b	C C	=d

C-1. Calculate the residue obtained on strongly heating $2.76 \text{ g Ag}_2\text{CO}_3$.

$$Ag_2CO_{3(s)} \xrightarrow{\Delta} 2Ag_{(s)} + CO_{2(g)} + \frac{1}{2}O_{2(g)}$$

C-2. Calculate the weight of iron which will be converted into its oxide by the action of 18g of steam.

Unbalanced reaction : Fe + $H_2O \longrightarrow Fe_3O_4 + H_2$.

- **C-3.** How many gm of HCl is needed for complete reaction with 43.5 gm MnO_2 ? (Mn = 55) HCl + MnO₂ \rightarrow MnCl₂ + H₂O + Cl₂ (unbalance)
- **C-4.** Nitric acid is manufactured by the Ostwald process, in which nitrogen dioxide reacts with water. $3NO_2(g) + H_2O(I) \rightarrow 2 HNO_3(aq) + NO(g)$ How many grams of nitrogen dioxide are required in this reaction to produce 25.2 gm HNO₃?
- **C-5.** What total volume, in litre at 627°C and 0.821 atm, could be formed by the decomposition of 16 gm of NH_4NO_3 ? Reaction : $2 NH_4NO_3 \rightarrow 2N_2 + O_2 + 4H_2O_{(q)}$.
- **C-6.** Calculate mass of phosphoric acid (H_3PO_4) required to obtain 53.4g pyrophosphoric acid. $2H_3PO_4 \rightarrow H_4P_2O_7 + H_2O$

Section (D) : Limiting reagent, % Excess, % Yield / Efficiency

- **D-1.** 50 g of $CaCO_3$ is allowed to react with 73.5 g of H_3PO_4 . Calculate : (i) Amount of $Ca_3(PO_4)_2$ formed (in moles) (ii) Amount of unreacted reagent (in moles)
- **D-2.** Carbon reacts with chlorine to form CCl_4 . 36 gm of carbon was mixed with 142 g of Cl_2 . Calculate mass of CCl_4 produced and the remaining mass of reactant.
- **D-3.** Potassium superoxide, KO₂, is used in rebreathing gas masks to generate oxygen : $KO_2(s) + H_2O(I) \rightarrow KOH(s) + O_2(g)$ (unbalanced) If a reaction vessel contains 0.158 mol KO₂ and 0.10 mol H₂O, how many moles of O₂ can be produced ?
- **D-4.** A chemist wants to prepare diborane by the reaction $6 \text{ LiH} + 8BF_3 \longrightarrow 6Li BF_4 + B_2H_6$ If he starts with 2.0 moles each of LiH & BF₃. How many moles of B₂H₆ can be prepared.

Section (E) : Principle of atom conservation (POAC), Reactions in sequence & parallel, Mixture analysis, % Purity

- E-1. What amount of CaO will be produced by 1 g of Calcium ?
- **E-2.** 6 gm nitrogen on successive reaction with different compounds gets finally converted into 30 gm $[Cr(NH_3)_xBr_2]$ Value of x is [Atomic mass of Cr = 52, Br = 80]
- **E-3.** All carbon atom present in 762 gm $KH_3(C_2O_4)_2$. 2H₂O is converted into CO_2 . How many gm of CO_2 were obtained.
- **E-4.** KCIO₃ decomposes by two parallel reaction

(i) $2\text{KCIO}_3 \xrightarrow{\Delta} 2\text{KCI} + 3\text{O}_2$ (ii) $4\text{KCIO}_3 \xrightarrow{\Delta} 3\text{KCIO}_4 + \text{KCI}$ If 3 moles of O_2 and 1 mol of KCIO_4 is produced along with other products then determine initial moles of KCIO_3 .

- **E-5.** When 4 gm of a mixture of NaHCO₃ and NaCl is heated, 0.66 gm CO_2 gas is evolved. Determine the percentage composition (by mass) of the original mixture.
- **E-6.** When 80 gm CH_4 is burnt completely, CO and CO_2 gases are formed in 1 : 4 mole ratio. What is the mass of O_2 gas used in combustion.

MOLE-III : Oxidation Reduction & Balancing Redox Equations

Section (F) : Basics of oxidation number

- F-1. Calculate the oxidation number of underlined elements in the following compounds :
 - (a) $K[Co(C_2O_4)_2(NH_3)_2]$ (d) $Na_2[Fe(CN)_5(NO^+)]$ (g) $[Fe(NO^+) (H_2O)_5]SO_4$

(b) $K_4 \underline{P}_2 O_7$	(c) <u>Cr</u> O ₂ Cl ₂
(e) $\underline{Mn_3O_4}$	(f) Ca(<u>C</u> ĪO ₂) ₂
(h) ZnO_{2}^{2}	(i) <u>Fe_{0.93}</u> O

Section (G) : Balancing redox reactions

- G-1. Write balanced net ionic equations for the following reactions in acidic solution :
 - (a) $S_4O_6^{2-} + AI \longrightarrow H_2S + AI^{3+}$ (b) $Fe^{2+} + Cr_2O_7^{2-} \rightarrow Fe^{3+} + Cr^{3+}$ (c) $IO_4^- + I^- \rightarrow I_2$ (d) $MnO_4^{2-} \rightarrow MnO_4^- + MnO_2$ (e) $HNO_2 \longrightarrow NO_3^- + NO$
- G-2. Write balanced net ionic equations for the following reactions in basic solution :
 - (a) $\text{ClO}^- + \text{CrO}_2^- \rightarrow \text{Cl}^- + \text{CrO}_4^{2-}$
 - (b) $Zn + NO_3^- \rightarrow NH_3 + ZnO_2^{2-}$
 - (c) $S \rightarrow S^{2-} + S_2 O_3^{2-}$
 - (d) $Mn(OH)_2 + MnO_4^- \longrightarrow MnO_2$
 - (e) $MnO_4^- + SO_3^{2-} \rightarrow MnO_4^{2-} + SO_4^{2-}$
MOLE-IV : Concentration Measurement

Section (H) : Units of concentration measurement, Interconversion of concentration units Commit to memory :

Molarity of solution =
$$\frac{\text{number of moles of solute}}{\text{volume of solution in litre}}$$

molality = $\frac{\text{number of moles of solute}}{\text{mass of solvent in gram}} \times 1000$
Let number of moles of solute in solution = n
Number of moles of solvent in solution = N
 \therefore Mole fraction of solute $(x_1) = \frac{n}{n+N}$
 \therefore Mole fraction of solvent $(x_2) = \frac{N}{n+N}$
% w/w = $\frac{\text{mass of solute in gm}}{\text{mass of solute in gm}} \times 100$
% w/v = $\frac{\text{mass of solute in gm}}{\text{volume of solution in ml}} \times 100$
% v/v = $\frac{\text{volume of solute in ml}}{\text{volume of solution in ml}} \times 100$
ppm_A = $\frac{\text{mass of A}}{\text{Total mass}} \times 10^6$ = mass fraction $\times 10^6$

- H-1. Find the mass of KOH needed to prepare 100 ml 1 M KOH solution. [At. mass K = 39]
- **H-2.** Calculate the molality of KCI solution prepared by dissolving 7.45 g of KCI to make 500 mL of the solution. $(d_{sol} = 1.2 \text{ g mL}^{-1})$
- H-3.(i) If you are given a 2M NaOH solution having density 1 g/mL, then find the molality of solution.
 - (ii) Find the molarity of 5m (molal) NaOH solution having density 1.5 g/ml.
 - (iii) Find the mole fraction of solute in problem (i)
 - (iv) Find the mole fraction of solute in problem (ii)
 - (v) Find the % (w/w) of NaOH in solution in problem (i)
 - (vi) Find the % (w/w) of NaOH in solution in problem (ii)
 - (vii) Find the % (w/v) of NaOH in solution in problem (ii)

(viii) A 300 g, 30% (w/w) NaOH solution is mixed with 500 g, 40% (w/w) NaOH solution. Find the mass percentage (w/w) of final solution.

- (ix) What is % (w/v) NaOH in problem (viii) if density of final solution is 2 g/ml?
- (x) What is the molality of final solution obtained in problem (viii)?
- H-4. Calculate molarity of all the ions produces by following aq. solution of electrolytes. (Assuming 100% dissociation)(i) 0.2M NaCl solution

(ii) $1.2M \text{Al}_2(\text{SO}_4)_3$ solution (iii) $1.2M \text{H}_2\text{SO}_4$ solution (iv) $2.1M \text{MgCl}_2$ solution

Section (I) : Dilution & Mixing of two liquids

- **I-1.** Find the Cl⁻ concentration in solution which is obtained by mixing one mole each of $BaCl_2$, NaCl and HCl in 500 ml water.
- I-2. What maximum volume of 3 M solution of KOH can be prepared from 1 L each of 1 M KOH and 6 M KOH solutions by using water ?
- I-3. Find the resultant molarity obtained by mixing the following.
 - (i) 2 litre, 0.5M HCl + 3 litre, 0.2 M HCl.
 - (ii) 500 ml, 1M NaCl + 200 ml, 2M NaCl
 - (iii) 100 ml, 0.1 M NaOH + 400 ml, 0.2 M NaOH
 - (iv) 2 litre, 0.2M HCl + 500 ml , 0.4 M HCl.

Section (J) : Percentage Determination of Elements in Organic Compounds

- **J-1.** In carius method of estimation of halogen, 0.15g of an organic compound gave 0.12g of AgBr. What is the percentage of bromine in the compound.
- **J-2.** 0.124 gm of an organic compound containing phosphorus gave 0.222 gm of $Mg_2P_2O_7$ by the usual analysis. Calculate the percentage of phosphorous in the compound.(Mg = 24, P = 31)
- **J-3.** 0.28 g of a nitrogenous compound was subjected to Kjeldahl's process to produce 0.17g of NH₃. The percentage of nitrogen in the organic compound is :
- J-4. In Dumas' method for estimation of nitrogen, 0.3g of an organic compound give 50 mL of nitrogen collected at 300K temperature and 700 mm pressure. What will be the percentage composition of nitrogen in the compound.
- **J-5.** In sulphur estimation, 0.157 g of an organic compound gave 0.4813 g of barium sulphate. What is the percentage of bromine in the compound.

PART - II : OBJECTIVE QUESTIONS

Single Choice Objective, straight concept/formula oriented

MOLE-I : Law of Chemical Combination

Section (A) : Molar volume of ideal gases at STP, Average molar mass

A-1.	Under the same conditions, two gases hav	e the same number of molecules. They must
	(A) be noble gases	(B) have equal volumes
	(C) have a volume of 22.4 dm ³ each	(D) have an equal number of atoms

A-2. 80gm of SO, gas occupies 14 litre at 2atm & 273K. The value of x is :

(B) x = 2

(A) x = 3

(C) x = 4

(D) none

Section (B) : Empirical Formula, % Composition of a given compound by mass, % By mole, Minimum molecular mass determination.

B-1. The empirical formula of a compound of molecular mass 120 is CH_2O . The molecular formula of the compound is :

(A) $C_2 H_4 O_2$ (B) $C_4 H_8 O_4$ (C) $C_3 H_6 O_3$ (D) all of these

B-2. Cortisone is a molecular substance containing 21 atoms of carbon per molecule. The mass percentage of carbon in cortisone is 69.98%. Its molar mass is :
 (A) 176.5 (B) 252.2 (C) 287.6 (D) 360.1

B-3.	-3. A compound contains 69.5% oxygen and 30.5% nitrogen and its molecular weight is 92. The formula compound is :-					
	(A) N_2O	(B) NO ₂	(C) N ₂ O ₄	(D) N ₂ O ₅		
B-4.	A compound of X and Y h	has equal mass of them. If	their atomic weights are	30 and 20 respectively. The		
	(A) X_2Y_2	(B) X_3Y_3	(C) X ₂ Y ₃	(D) X ₃ Y ₂		
		MOLE-II : Bas	ic Stoichiomet	ry		
Sect or 2)	ion (C) : Stoichiomet	ry, Equation based	calculations (Eleme	ntary level single equation		
C-1.ඍ	For the reaction 2P + C (A) 8 mol of R	$Q \rightarrow R$, 8 mol of P and ex (B) 5 mol of R	cess of Q will produce : (C) 4 mol of R	(D) 13 mol of R		
C-2.	If 1.5 moles of oxygen ((A) 27 g	combine with Al to form A (B) 40.5 g	I_2O_3 , the weight of Al user (C) 54g	d in the reaction is : (D) 81 g		
C-3.	How many liters of CO_2	$_{2}$ at STP will be formed wh	nen 0.01 mol of H_2SO_4 rea	acts with excess of Na_2CO_3 .		
	$Na_2CO_3 + H_2SO_4$	Na ₂ SO ₄ + CO ₂ + H ₂ O (B) 2.24 L	(C) 0.227 L	(D) 1.12 L		
C-4. ଅ	How many moles of po	tassium chlorate need to	be heated to produce 11.	2 litre oxygen at 0°C and 1 atm.		
	(A) $\frac{1}{2}$ mol	(B) $\frac{1}{3}$ mol	(C) $\frac{1}{4}$ mol	(D) $\frac{2}{3}$ mol		
C-5.	How many litres of oxyge (A) 11.2 L	en at 1atm & 273K will be (B) 22.4 L	e required to burn compl (C) 5.6 L	etely 2.2 g of propane (C ₃ H ₈) (D) 44.8 L		
C-6.	Volume of CO_2 obtained a	at STP by the complete d	ecomposition of 9.85 g B	aCO ₃ is		
	(A) 2.24 lit	(B) 1.135 lit	(C) 0.135 lit	(D) 2.27 lit		
Sect	ion (D) : Limiting rea	agent, % Excess, %	Yield / Efficiency			
D-1.	How many mole of Zn(I (A) 2 mole	FeS ₂) can be made from 2 (B) 3 mole	2 mole zinc, 3 mole iron a (C) 4 mole	and 5 mole sulphur. (D) 5 mole		
D-2. ব	 Equal weight of 'X' (At. wt. = 36) and 'Y' (At. wt. = 24) are reacted to form the compound X₂Y₃. Then : (A) X is the limiting reagent (B) Y is the limiting reagent (C) No reactant is left over and mass of X₂Y₃ formed is double the mass of 'X' taken (D) none of these 					
D-3.	0.5 mole of H_2SO_4 is mi (A) 0.2	xed with 0.2 mole of Ca (0 (B) 0.5	DH) ₂ . The maximum num (C) 0.4	ber of moles of $CaSO_4$ formed is (D) 1.5		
D-4.	If 1/2 moles of oxygen	combine with aluminium	to form Al ₂ O ₃ then weigl	nt of Aluminium metal used in the		
	(A) 27 g	(B) 18 g	(C) 54 g	(D) 40.5 g		
D-5.	The mass of P_4O_{10} pro-	duced if 440 gm of P_4S_3 is	s mixed with 384 gm of O	₂ is		
	$P_4S_3 + O_2 \longrightarrow P_4O_{10}$	+ SO ₂				
	(A) 568 gm	(B) 426 gm	(C) 284 gm	(D) 396 gm		

D-6. The mass of Mg_3N_2 produced if 48 gm of Mg metal is reacted with 34 gm NH_3 gas is

 $Mg + NH_3 \longrightarrow Mg_3N_2 + H_2$

(A) $\frac{200}{3}$ gm (B) $\frac{100}{3}$ gm (C) $\frac{400}{3}$ gm (D) $\frac{150}{3}$ gm

Section (E) : Principle of atom conservation (POAC), Reactions in sequence & parallel, Mixture analysis, % Purity

- E-1. 25.4 g of iodine and 14.2g of chlorine are made to react completely to yield a mixture of ICl and ICl_a. Calculate the number of moles of ICI and ICI, formed. (A) 0.1 mole, 0.1 mole (B) 0.1 mole, 0.2 mole (C) 0.5 mole, 0.5 mole (D) 0.2 mole, 0.2 mole
- E-2. What weight of CaCO₂ must be decomposed to produce the sufficient quantity of carbon dioxide to convert 21.2 kg of Na₂CO₃ completely in to NaHCO₃. [Atomic mass Na = 23, Ca = 40]

 $CaCO_3 \longrightarrow CaO + CO_2$ $Na_{2}CO_{3} + CO_{2} + H_{2}O \xrightarrow{2} 2NaHCO_{3}$ Kg (B) 20 Kg (C) 120 Kg (A) 100 Kg (D) 30 Kg

- E-3.2. NX is produced by the following step of reactions
 - $M + X_2 \longrightarrow M X_2$ $3MX_{a} + X_{a} \longrightarrow M_{a}X_{a}$

$$M_3 X_8 + N_2 CO_3 \longrightarrow NX + CO_2 + M_3O_4$$

How much M (metal) is consumed to produce 206 g of NX. (Take at wt of M = 56, N=23, X = 80)

11

(A) 42 g (B) 56 g (C)
$$\frac{14}{3}$$
 g (D) $\frac{7}{4}$ g

E-4.2. The following process has been used to obtain iodine from oil-field brines in California. $Nal + AgNO_3 \longrightarrow Agl + NaNO_3$ $2Agl + Fe \longrightarrow Fel_{2} + 2Ag$;

 $2\text{Fel}_2 + 3\text{Cl}_2 \longrightarrow 2\text{FeCl}_3 + 2\text{I}_2$ How many grams of AgNO, are required in the first step for every 254 kg I, produced in the third step. (A) 340 kg (B) 85 kg (C) 68 kg (D) 380 kg

- Mixture of MgCO₃ & NaHCO₃ on strong heating gives CO₂ & H₂O in 3 : 1 mole ratio. The weight % of NaHCO₃ E-5. present in the mixture is: (A) 30% (B) 80% (C) 40% (D) 50%
- E-6. An impure sample of CaCO₃ contains 38% of Ca. The percentage of impurity present in the sample is :

(A) 5%	(B) 95%	(C) 10%	(D) 2.5%

		MOLE-III : Balancing	Oxidation Redu Redox Equation	ction & 1s			
Section (F) : Basics of oxidation number							
F-1.	The oxidation nu (A) + 1	umber of Oxygen in Na ₂ C (B) + 2	0₂ is : (C) − 2	(D) – 1			
F-2.	The oxidation nu (A) + 3	umber of Phosphorus in N (B) + 2	/lg ₂ P ₂ O ₇ is : (C) + 5	(D) – 3			

F-3.১	The oxidation states of Sulphur in the anions SO ₃ (A) $S_2O_6^{2-} < S_2O_4^{2-} < SO_3^{2-}$				$_{3}^{2^{-}}$, S ₂ O ₄ ²⁻ and S ₂ O ₆ ²⁻ follow the order : (B) S ₂ O ₄ ²⁻ < SO ₃ ²⁻ < S ₂ O ₆ ²⁻						
	(C) $SO_3^{2-} < S_2O_4^{2-} < S_2O_6^{2-}$				(D) S ₂ C	$D_4^2 < S_2^0$	₆ ²⁻ < SO ₃	2- 3			
F-4.	Match L below t List-I (a) (b) (c) (d) Code :	List-I (Co he lists : NaN ₃ N ₂ H ₂ NO N ₂ O ₅	impound	ls) with L	.ist-II (Ox	List-II (1) (2) (3) (4)	+5 +2 −1/3 −1	Nitrogen) and sel	ect ans	wer using the codes given
	(A) (C)	(a) 3 3	(b) 4 4	(c) 2 1	(d) 1 2		(B) (D)	(a) 4 4	(b) 3 3	(c) 2 1	(d) 1 2
F-5.	1 mole of N_2H_4 loses ten moles of electrons to f appears in the new compound, what is the oxid oxidation state of hydrogen). (A) - 1 (B) - 3			ons to for ne oxidat	m a nev ion state (C) + 3	w compo e of nitro	und Y. As gen in Y	ssuming ? (There (D) + 5	g that all the nitrogen e is no change in the		
F-6.	The average oxidation state of Fe in Fe ₃ (A) $- 8/3$ (B) $8/3$			Fe in Fe ₃ 0	0 ₄ is :	(C) 2			(D) 3		
Section	on (G)	: Balar	ncing r	edox r	eactio	ns					
G-1.	In the re (A) x =	eaction 3, y = 2	xHI + yF	INO ₃ — (B) x=	→ NC 2, y = 3) + I ₂ + ⊦	l₂O, upo (C) x =	on balan = 6, y = 2	cing with	whole r (D) x =	number coefficients : = 6, y = 1
G-2.১	For the the corr (A) 2, 5	redox re ect who , 16	eaction le numbe	MnO₄ [−] + er stoich (B) 16,	$C_2O_4^{2-}$ iometric 5, 2	+ H⁺ — coefficie	\rightarrow Mr nts of M (C) 5, \hat{c}	n²+ + CC InO₄⁻, C₂ 16, 2	$D_{2} + H_{2}O_{4}$ O_{4}^{2-} and	, H⁺ are r (D) 2,	espectively: 16, 5
G-3.	For the coefficient (A) x =	redox re ents: 1, y = 5	eaction of	хР₄ + уН (В) х	INO ₃ — = 2, y =	$\rightarrow H_{3}P$ 10	$O_4 + NC$ (C) x =	D ₂ + H ₂ O, = 1, y = 2	upon ba	alancing (D) x =	y with whole number = 1, y = 15
G-4.	In the reaction $X^- + XO_3^- + H^+ \longrightarrow X_2 + H_2C$ (A) 1 : 5 (B) 5 : 1		+ H ₂ O, t	the molar ratio in which λ (C) 2 : 3		which X	⁻ and X (D) 3 :	O₃⁻ react is : 2			
G-5.	CN⁻ is o	oxidised	by NO ₃ -	in prese	ence of a	cid :					
		a CN⁻ -	+ b NO ₃ -	+ c H⁺ -	→ (a	1 + b) NC) + a CC	$D_2 + \frac{c}{2}H_2$	2 ⁰		
	What a (A) 3, 7	re the w 7, 7	hole nun	nber valı (B) 3, 1	ues of a, 10, 7	b, c in th	at order (C) 3, ²	·: 10, 10		(D) 3,	7, 10
			MOL	.E-IV	: Co	ncen	tratic	on Me	easur	eme	nt
Sectio	on (H) :	Units	of con	centra	tion m		ment,	Interco	onvers	ion of	concentration units
H-1.	500 mL (A) 0.1	. of a glu M	cose sol	(B) 1.0	ntains 6. M	02×10^{2}	- moleci (C) 0.2	uies. The M	econcen	(D) 2.0	of the solution is

H-2. Equal moles of H_2O and NaCl are present in a solution. Hence, molality of NaCl solution is : (A) 0.55 (B) 55.5 (C) 1.00 (D) 0.18

H-3.æ	What is the molarity of l	H_2SO_4 solution that has a $S = 32$	density of 1.84 g/cc and	contains 98% by mass of H_2SO_4 ?
	(A) 4.18 M	(B) 8.14 M	(C) 18.4 M	(D) 18 M
H-4.	1000 g aqueous solution (A) 10 ppm	on of Ca(NO ₃) ₂ contains (B) 100 ppm	10 g of calcium nitrate. (C) 1000 ppm	Concentration of the solution is : (D) 10,000 ppm
H-5.	171 g of cane sugar (C	12H22O11) is dissolved in	1 litre of water. The mol	arity of the solution is :
	(A) 2.0 M	(B) 1.0 M	(C) 0.5 M	(D) 0.25 M
H-6.	Molality of 20% (w/w) a	aq.glucose solution is :		
	(A) $\frac{25}{18}$ m	(B) $\frac{10}{9}$ m	(C) $\frac{25}{9}$ m	(D) $\frac{5}{18}$ m
H-7.	Molarity of liquid HCl, i (A) 36.5 M	f density is 1.17 g/cc. : (B) 18.25 M	(C) 32.05 M	(D) 42.10 M
Sect	ion (I) : Dilution & M	ixing of two liquids		
I -1 .	If 500 ml of 1 M solution will be :	of glucose is mixed with	500 ml of 1 M solution of	f glucose final molarity of solution
	(A) 1 M	(B) 0.5 M	(C) 2 M	(D) 1.5 M
-2 .	The volume of water 0.2 M HCl to obtain 0.2	that must be added t 5 M solution of HCl is :	o a mixture of 250 ml	of 0.6 M HCl and 750 ml of
	(A) 750 mi	(B) 100 mi	(C) 200 mℓ	(D) 300 mℓ
I -3.	What volume of a 0.8 M (A) 100 mL	l solution contains 100 m (B) 125 mL	illi moles of the solute? (C) 500 mL	(D) 62.5 mL
-4. &	2M of 100 ml Na_2SO_4 is	s mixed with 3M of 100 m	INaCI solution and 1M o	f 200 ml CaCl ₂ solution. Then the
	ratio of the concentratio (A) 1/2	n of cation and anion. (B) 2	(C) 1.5	(D) 1
I -5 .	How much volume of 3 (A) 300 ml	.0 M H ₂ SO ₄ is required f (B) 320 ml	or the preparation of 1.0 (C) 333.3 ml	litre of 1.0 M solution? (D) 350.0 ml
-6.	Equal weight of NaCl an will be –	d KCl are dissolved separ	ately in equal volumes of	solutions. Molarity of the solutions
	(A) Equal (C) Greater for KCl		(B) Greater for NaCl (D) Uncomparable.	
I -7.	Equal volumes of 10% ((A) basic	(w/v) of HCl is mixed with (B) neutral	10% (w/v) NaOH solutior (C) acidic	n. The resultant solution be. (D) can't be predicted.
Sect	ion (J) : Percentage	Determination of E	lements in Organic	Compounds
J-1.	In kieldahl's method nit	togen present in estimate	ed as :	
	(A) N ₂	(B) NH ₃	(C) NO ₂	(D) None of these
J-2.	An organic compound ha	as 6.2 % of phosphorus.	On sequence of reaction	, the phosphorous present in the

J-2. An organic compound has 6.2 % of phosphorus. On sequence of reaction, the phosphorous present in the 10gm of organic compound is converted to Mg₂P₂O₇. Find the weight of Mg₂P₂O₇ formed.
 (A) 2.22 gm
 (B) 10.0 gm
 (C) 4.44 gm
 (D) 1.11 gm

1.

PART - III : MATCH THE COLUMN

	Column - I		Column - II
(A)	A gaseous organic compound containing C = 52.17%, H = 13.04% & O = 34.78% (by weight) having molar mass 46 g/mol.	(p)	One mole of compound contains 4N _A atoms of Hydrogen.
(B)	0.3 g of an organic compound containing C, H and O on combustion yields 0.44 g of CO_2 and 0.18 g of H ₂ O, with two O atoms per molecule.	(q)	The empirical formula of the compound is same as its molecule formula.
(C)	A hydrocarbon containing C = 42.857% and H = 57.143% (by mole) containing 3C atoms per molecule.	(r)	Combustion products of one mole of compound contains larger number of moles of CO_2 than that of H_2O .
(D)	A hydrocarbon containing 10.5 g carbon per gram of hydrogen having vapour density 46.	(s)	CO ₂ gas produced by the combustion of 0.25 mole of compound occupies a volume of 11.2 L at 0°C and 1 atm.

2. 🙇 Column I

- (A) $Zn(s) + 2HCI(aq) \rightarrow ZnCI_2(s) + H_2(g)$ above reaction is carried out by taking 2 moles each of Zn and HCI
- (B) AgNO₃(aq) + HCl(aq) \rightarrow AgCl(s) + HNO₃(g) above reaction is carried out by taking 170 g AgNO₃ and 18.25 g HCl (Ag = 108)
- (C) $CaCO_3(s) \rightarrow CaO(s) + CO_2(g)$ 100 g CaCO₃ is decomposed
- (D) $2\text{KCIO}_3(s) \rightarrow 2\text{KCI}(s) + 3\text{O}_2(g)$ 2/3 moles of KCIO_3 decomposed

3. Column-l

- (A) 120 g CH₃COOH in 1 L solution ($d_{sol} = 1.2$ g/mL)
- (B) 120 g glucose dissolved in 1 L solution $(d_{sol} = 1.2 \text{ g/mL})$
- (C) $X_{NH_2CONH_2} = 1/31$ (aqueous solution)
- (D) 19.6% (w/v) H_2SO_4 solution \rightarrow (d_{solution} = 1.2 g/mL)

Column II

- (p) 50% of excess reagent left
- (q) 22.7 L of gas at STP is liberated
- (r) 1 moles of solid (product) obtained.
- (s) HCl is the limiting reagent

Column-II

- (p) M = 2
- (q) 10% w/w solution
- (r) 12% w/v solution
- (s) m = 1.85
- (t) m = 0.617



12.	$xNO_{3}^{-} + yl^{-} + zH^{+} \rightarrow 2NO_{3}^{-}$ (A) 2, 6, 8	O + 3I ₂ + 4H ₂ O x , y, z re (B) 1, 6 , 4	spectively in the above e (C) 0, 6 , 8	equation are : (D) 2 , 3 , 4	
13.১	A solution of glucose red (m) at 10°C. When you v (A) mole fraction (x) and (C) mole fraction ($x/2$) at	eived from some researc will calculate its molality a molality (m) nd molality (m/2)	ch laboratory has been marked mole fraction x and molality and mole fraction in your laboratory at 24° C you will find (B) mole fraction (2x) and molality (2m) (D) mole fraction (x) and (m ± dm) molality		
14.	36.5 % HCl has density (A) 15.7, 15.7	equal to 1.20 g mL⁻¹. The (B) 12, 12	molarity (M) and molality (C) 15.7, 12	y (m), respectively, are (D) 12, 15.7	
15.	An aqueous solution of (A) 1.79	ethanol has density 1.028 (B) 2.143	5 g/mL and it is 2M. Wha (C) 1.951	t is the molality of this solution ? (D) None of these.	
16.১	Mole fraction of ethyl alc alcohol by weight is : (A) 54%	cohol in aqueous ethyl alc (B) 25%	ohol (C ₂ H ₅ OH) solution is (C) 75%	0.25. Hence percentage of ethyl (D) 46%	
17.	Calculate the mass perc in a 100 ml sulphuric acid ml. (molecular weight of (A) 80.8%	ent (w/w) of sulphuric acid d solution containing 80 m f $H_2SO_4 = 98$). Take react (B) 84%	d in a solution prepared by hass percent (w/w) of H_2SG tion $SO_3 + H_2O \rightarrow H_2SO$ (C) 41.65%	dissolving 4 g of sulphur trioxide D_4 and having a density of 1.96 g/ (D) None of these	
18.	Molarity and Molality of a of solution. (A) 1 g/cc	a solute (M. wt = 50) in aq (B) 0.95 g/cc	ueous solution is 9 and 18 (C) 1.05 g/cc	respectively. What is the density (D) 0.662 g/cc	
19.2	What volume (in ml) of 0	$0.2 \mathrm{M}\mathrm{H_2SO_4}$ solution shoul	d be mixed with the 40 ml	of 0.1 M NaOH solution such that	
	the resulting solution ha	s the concentration of H_2	SO_4 as $\frac{6}{55}$ M.		
	(A) 70	(B) 45	(C) 30	(D) 58	

PART - II : SUBJECTIVE QUESTIONS

- 1. How many gram ions of SO_4^{-2} are present in 1.25 mole of K_2SO_4 . $AI_2(SO_4)_3$. $24H_2O$:
- 2. A certain organic substance used as a solvent in many reactions contains carbon, hydrogen, oxygen and sulphur. Weight % of hydrogen in the compound is 7.7. The weight ratio C : O : S = 3 : 2 : 4. What is the least possible molar mass (in g) of the compound ?
- 3. Calculate maximum mass of CaCl₂ produced when 2.4×10^{24} atoms of calcium is taken with 96 litre of Cl₂ gas at 380 mm pressure and at 27°C. [R : 0.08 atm L/mole-K & N_A = 6 × 10²³]
- **4.** A 2 g sample containing Na_2CO_3 and $NaHCO_3$ losses 0.248 g when heated to $300^\circ C$, the temperature at which $NaHCO_3$ decomposes to Na_2CO_3 , CO_2 and H_2O . What is the percentage of Na_2CO_3 in the given mixture?
- **5.** A sample of chalk contains clay as impurity. The clay impurity loses 11% of its weight as moisture on prolong heating. 5 gram sample of chalk on heating shows a loss in weight (due to evolution of CO_2 and water) by 1.1 g. Calculate % of chalk (CaCO₃) in the sample. [Hint : Chalk (CaCO₃) release CO_2 on heating]
- 6. A fluorine disposal plant was constructed to carryout the reactions :

 $F_2 + 2NaOH \longrightarrow \frac{1}{2}O_2 + 2NaF + H_2O$

 $2NaF + CaO + H_2O \longrightarrow CaF_2 + 2NaOH$

As the plant operated, excess lime was added to bring about complete precipitation of the fluoride as CaF_2 . Over a period of operation, 1900 kg of fluorine was fed into a plant and 10,000 kg of lime was required. What was the percentage utilisation of lime ? [Lime : CaO]

7. $Cl_2 + KOH \xrightarrow{60\%} KCl + KClO + H_2O$

 $\mathsf{KCIO} \xrightarrow{50\%} \mathsf{KCI} + \mathsf{KCIO}_3$

 $\text{KCIO}_3 \xrightarrow{80\%} \text{KCIO}_4 + \text{KCI}$

113.5 L Cl₂ gas at STP is passed in 10 L KOH solution, containing 1 mole of potassium hydroxide per liter. Calculate the total moles of KCl produced, rounding it off to nearest whole number. (Yield of chemical reactions are written above the arrow (\rightarrow) of respective reaction)

8. If 240 g of carbon is taken in a container to convert it completely to CO_2 but in industry it has been found that 280 g of CO was also formed along with CO_2 . Find the mole percentage yield of CO_2 . The reactions occurring

are $C + O_2 \longrightarrow CO_2$; $C + \frac{1}{2} O_2 \longrightarrow CO$

- 9. Among the following compounds given below, what is the sum of the oxidation states of all underlined elements? \underline{CO}_2 , $K_2\underline{Mn}O_4$
- **10.** Find the sum of average oxidation number of S in H_2SO_5 (peroxy monosulphuric acid) and $Na_2S_2O_3$ (sodium thiosulphate).
- **11.** The reaction $Cl_2(g) + S_2O_3^{2-} \longrightarrow SO_4^{2-} + Cl^-$ is to be carried out in basic medium. Starting with 1.5 mole of $Cl_2, 0.1$ mole $S_2O_3^{2-}$ and 3 mole of OH^- . How many moles of OH^- will be left in solution after the reaction is complete. Assume no other reaction occurs.
- 12. What volume of water should be added to 50 ml of HNO₃ having density 1.5 g ml⁻¹ and 63.0% by weight to have one molar solution.
- **13.** What is the quantity of water (in g) that should be added to 16 g. methanol to make the mole fraction of methanol as 0.25 :
- **14.** What volume (in mL) of 90% alcohol by weight (d = 0.8 g mL^{-1}) must be used to prepare 80 mL of 10% alcohol by weight (d = 0.9 g mL^{-1})?
- **15.** A solution containing 0.1 mol of a metal chloride MCl_x requires 500 ml of 0.8 M AgNO₃ solution for complete reaction $MCl_x + xAgNO_3 \rightarrow xAgCl + M(NO_3)_x$. Then the value of x is :
- **16.** 60 ml of a "**x**" % w/w alcohol (d = 0.6 g/cm³) must be used to prepare 200 cm³ of 12% w/w alcohol by weight (d = 0.90 g/cm³). Calculate the value of "**x**"?
- **17.** During estimation of nitrogen present in an organic compound by Kjeldahl's method, the ammonia evolved from 0.5g of the compound in Kjeldahl's estimation of nitrogen, neutralized 10 mL of $1M H_2SO_4$. What is the percentage of nitrogen in the compound :
- **18.** In a gravimetric determination of P, an aqueous solution of dihydrogen phosphate ion $H_2PQ_4^-$ is treated with a mixture of ammonium and magnesium ions to precipitate magnesium ammonium phosphate, $Mg(NH_4)PO_4.6H_2O$. This is heated and decomposed to magnesium pyrophosphate, $Mg_2P_2O_7$, which is weighed. A solution of $H_2PO_4^-$ yielded 1.054 g of $Mg_2P_2O_7$. What weight of NaH_2PO_4 was present originally?

PART - III : ONE OR MORE THAN ONE OPTIONS CORRECT TYPE

- 1. Which is/are correct statements about 1.7 g of NH_3 :
 - (A) It contain 0.3 mol H atom
 - (C) Mass % of hydrogen is 17.65%
- (B) it contain 2.408×10^{23} atoms (D) It contains 0.3 mol N-atom

- **2.** Select the correct statement(s) for $(NH_4)_3PO_4$.
 - (A) Ratio of number of oxygen atoms to number of hydrogen atoms is 1:3
 - (B) Ratio of number of cations to number of anions is 3 : 1
 - (C) Ratio of number of gm-atoms of nitrogen to gm-atoms of oxygen is 3 : 2
 - (D) Total number of atoms in one mole of $(NH_4)_3PO_4$ is 20.
- **3.** If 27 g of Carbon is mixed with 88 g of Oxygen and is allowed to burn to produce CO_2 , then :
 - (A) Oxygen is the limiting reagent.
 - (B) Volume of CO_2 gas produced at STP is 51.075 L.
 - (C) C and O combine in mass ratio 3:8.
 - (D) Volume of unreacted O_2 at STP is 11.35 L.
- 4. The density of air is 0.001293 g/cm³ at STP. Identify which of the following statement is correct (A) Vapour density is 14.67
 - (B) Molecular weight is 29.35
 - (C) Vapour density is 0.001293 g/cm³
 - (D) Vapour density and molecular weight cannot be determined.
- 5. 12 g of Mg was burnt in a closed vessel containing 32 g oxygen. Which of the following is /are correct.(A) 2 gm of Mg will be left unburnt.
 - (B) 0.75 gm-molecule of O₂ will be left unreacted.
 - (C) 20 gm of MgO will be formed.
 - (D) The mixture at the end will weight 44 g.
- 6. (i) $K_4 Fe(CN)_6 + 3H_2SO_4 \longrightarrow 2K_2SO_4 + FeSO_4 + 6HCN$ (ii) $6HCN + 12H_2O \longrightarrow 6HCOOH + 6NH_3$

(iii) (a) $6NH_3 + 3H_2SO_4 \longrightarrow 3(NH_4)_2SO_4$ (b) $6HCOOH \xrightarrow{H_2SO_4} 6CO + 6H_2O$ Above steps of reactions occur in a container starting with one mole of $K_4[Fe(CN)_6]$, 5 mole of H_2SO_4 and enough water. Find out the limiting reagent in step (i) and calculate maximum moles of CO gas and $(NH_4)_2$ SO_4 that can be produced.

(A) $LR = H_2SO_4$

```
(C) 6 moles of CO, 2 moles of (NH_4)_2SO_4
```

(B) LR = $K_4 Fe(CN)_6$, (D) 5 moles of CO, 2.5 moles of $(NH_4)_2 SO_4$

- **7.** A sample of a mixture of $CaCl_2$ and NaCl weighing 4.44 g was treated to precipitate all the Ca as $CaCO_3$,
which was then heated and quantitatively converted to 1.12g of CaO. (At . wt. Ca = 40, Na = 23, Cl = 35.5)
(A) Mixture contains 50% NaCl
(C) Mass of $CaCl_2$ is 2.22 g(B) Mixture contains 60% $CaCl_2$
(D) Mass of $CaCl_2$ 1.11 g
- 8. For the following reaction : $Na_2CO_3 + 2HCI \longrightarrow 2NaCI + CO_2 + H_2O$ 106.0 g of Na_2CO_3 reacts with 109.5 g of HCI. Which of the following is/are correct. (A) The HCI is in excess. (B) 117.0 g of NaCI is formed. (C) The volume of CO_2 produced at 0°C and 1 atm is 22.4 L (D) None of these
- 9. (a) $A + B \rightarrow A_3B_2$ (unbalanced)
 $A_3B_2 + C \rightarrow A_3B_2C_2$ (unbalanced)
Above two reactions are carried out by taking 3 moles each of A and B and one mole of C. Then which option
is/are correct ?(A) 1 mole of $A_3B_2C_2$ is formed
(C) 1/2 mole of $A_3B_2C_2$ is formed(B) 1/2 mole of $A_3B_2C_2$ is formed
(D) 1/2 mole of A_3B_2 is left finally

10. 'A' reacts by following two parallel reactions to give B & C If half of 'A' goes into reaction I and other half goes to reaction-II. Then, select the correct statement(s) $A + N \xrightarrow{I} B + L$ $A + N \xrightarrow{II} \frac{1}{2}B + \frac{1}{2}(C) + L$ (A) B will be always greater than C (B) If 2 mole of C are formed then total 2 mole of B are also formed (C) If 2 mole of C are formed then total 4 mole of B are also formed (D) If 2 mole of C are formed then total 6 mole of B are also formed 11. Which of the following are examples of disproportionation reaction : (B) $\text{KCIO}_3 \longrightarrow \text{KCI} + \text{O}_2$ (D) $\text{CI}_2 + \text{OH}^- \longrightarrow \text{CIO}^- + \text{CI}^- + \text{H}_2\text{O}$ (A) HgO \longrightarrow Hg + O₂ (C) $KClO_3 \longrightarrow KClO_4 + KCl$ 12.2 Consider the redox reaction $2S_2O_3^{2-} + I_2 \longrightarrow S_4O_6^{2-} + 2I^-$: (B) $S_2O_3^{2-}$ gets oxidised to $S_4O_6^{2-}$ (A) $S_2O_3^{2-}$ gets reduced to $S_4O_6^{2-}$ (D) I, gets oxidised to I-(C) I₂ gets reduced to I⁻ 13. Which of the following statements is/are correct? In the reaction $xCu_3P + yCr_2O_7^{2-} + zH^+ \longrightarrow Cu^{2+} + H_3PO_4 + Cr^{3+}$ (A) Cu in Cu₃P is oxidised to Cu²⁺ whereas P in Cu₃P is also oxidised to PO₄³⁻ (B) Cu in Cu₃P is oxidised to Cu²⁺ whereas P in Cu₃P is reduced to H₃PO₄ (C) In the conversion of Cu₃P to Cu²⁺ and H₃PO₄, 11 electrons are involved (D) The value of x is 6. 14. Solutions containing 23 g HCOOH is/are : (A) 46 g of 70% $\left(\frac{w}{v}\right)$ HCOOH (d_{solution} = 1.40 g/mL) (B) 50 g of 10 M HCOOH ($d_{solution} = 1 \text{ g/mL}$) (C) 50 g of 25% $\left(\frac{w}{w}\right)$ HCOOH (D) 46 g of 5 M HCOOH ($d_{solution} = 1 \text{ g/mL}$) 15. Which of the following solutions contains same molar concentration ? (B) 33.0 g (NH_{a})₂ SO₄ in 200 mL solution (A) 166 g. KI/L solution (D) 27.0 mg Al³⁺ per mL solution (C) 25.0 g CuSO₄.5H₂O in 100mL solution 16. If 100 ml of 1M H_2SO_4 solution is mixed with 100 ml of 9.8%(w/w) H_2SO_4 solution (d = 1 g/ml) then : (B) volume of solution become 200 ml (A) concentration of solution remains same (C) mass of H_2SO_4 in the solution is 98 g (D) mass of H_2SO_4 in the solution is 19.6 g 17.2 Equal volume of 0.1M NaCl and 0.1M FeCl, are mixed with no change in volume due to mixing. Which of the following will be true for the final solution. (No precipitation occurs). Assume complete dissociation of salts and neglect any hydrolysis.

(A) $[Na^+] = 0.05 \text{ M}$ (B) $[Fe^{2+}] = 0.05 \text{ M}$ (C) $[Cl^-] = 0.3 \text{ M}$ (D) $[Cl^-] = 0.15 \text{ M}$

PART - IV : COMPREHENSION

Read the following comprehension carefully and answer the questions.

Comprehension #1

A chemist decided to determine the molecular formula of an unknown compound. He collects following informations :

(I) Compounds contains 2 : 1 'H' to 'O' atoms(number of atoms).

(II) Compounds has 40% C by mass

(III) Approximate molecular mass of the compound is 178 g

(IV) Compound contains C, H and O only.

1.	What is the % by	What is the % by mass of oxygen in the compound							
	(A) 53.33%	(B) 88.88%	(C) 33.33%	(D) None of these					
2.	What is the empir (A) CH ₂ O	ical formula of the compo (B) CH ₂ O	und (C) C.H.O	(D) CH ₂ O ₂					
3.	Which of the follow	wing could be molecular fo	ormula of compound	(-)3 - 2					
-									

(A) $C_6 H_6 O_6$ (B) $C_6 H_{12} O_6$ (C) $C_6 H_{14} O_{12}$ (D) $C_6 H_{14} O_6$

Comprehension # 2

According to the Avogadro's law, equal number of moles of gases occupy the same volume at identical condition of temperature and pressure. Even if we have a mixture of non-reacting gases then Avogadro's law is still obeyed by assuming mixture as a new gas.

Now let us assume air to consist of 80% by volume of Nitrogen (N_2) and 20% by volume of oxygen (O_2). If air is taken at STP then its 1 mol would occupy 22.7 L. 1 mol of air would contain 0.8 mol of N_2 and 0.2 mol of

 $O^{}_{_2}$ hence the mole fractions of $N^{}_{_2}$ and $O^{}_{_2}$ are given by $\,X^{}_{N^{}_2}=0.8$, $\,X^{}_{O^{}_2}=0.2$

- 4.Volume occupied by air at STP containing exactly 11.2 g of Nitrogen :
(A) 22.4 L(B) 8.96 L(C) 11.35 L(D) 2.24 L
- 5. If air is treated as a solution of O_2 and N_2 then % W/W of oxygen is :

(A) 10	(B) $\frac{200}{9}$	(C) 700	(D) <u>350</u>
9		9	9
Density of air at \$ (A) 1 g/L	STP is : (B) 1.26 g/L	(C) 2.52 g/L	(D) can't be determined

Comprehension #3

6.

The concentrations of solutions can be expressed in number of ways; viz : mass fraction of solute (or mass percent), Molar concentration (Molarity) and Molal concentration (molality). These terms are known as concentration terms and also they are related with each other i.e. knowing one concentration term for the solution, we can find other concentration terms also. The definition of different concentration terms are given below :

Molarity : It is number of moles of solute present in one litre of the solution. Molality : It is the number of moles of solute present in one kg of the solvent

molesof solute

Mole Fraction = moles of solute + moles of solvent

If molality of the solution is given as 'a' then mole fraction of the solute can be calculated by

Mole Fraction =
$$\frac{a}{a + \frac{1000}{M_{solvent}}}$$
; = $\frac{a \times M_{solvent}}{(a \times M_{solvent} + 1000)}$

where a = molality and $M_{solvent}$ = Molar mass of solvent

We can change : Mole fraction \leftrightarrow Molality \leftrightarrow Molarity

7.	60 g of	60 g of solution containing 40% by mass of NaCl are mixed with 100 g of a solution containing 15% by mass							
	NaCl. Determine the mass percent of sodium chloride in the final solution.								
	(A) 24.	4%	(B) 78%	6	(C) 48.8%	(D) 19	.68%		
8.	Whati	s the molality of	the above	e solutio	n.				
	(A) 4.4	m	(B) 5.5	m	(C) 24.4 m	(D) no	ne		
9.	What i	s the molarity o	f solution	if density	y of solution is 1.6 g/ml				
	(A) 5.8	5 M	(B) 6.6	7 M	(C) 2.59 M	(D) no	ne		
Compr	rehensi	on # 4							
	Colum	in-l		Colum	n-ll	Colur	nn-III		
	(A)	2 M - aqueous NaOH solution	5 1	(i)	2 mole solute/litre solution	(I)	6 % (w/v) solution		
	(density = 1.25 gm/ml)								
	(B)	1.5 m - aqueo NaOH solution	us 1 6 am/ml)	(ii)	1.5 mole solute/litre solution	(II)	8 % (w/v) solution		
	(C) 0.5 M aqueous		s gm/m)	(iii)	0.5 mole solute/litre	(111)	9 % (w/v) solution		
	()	Glucose solut (density = 1.0	ion 9 gm/ml)	()	solution				
	(D)	1.5 M aqueou Urea solution (density = 1.1	s 5 gm/ml)	(iv)	1.5 mole solute/kg solvent	(IV)	9 gm solute per 100 gm solvent		
10.	Which (A) A –	of the following i – II	is correct (B) B –	match?	, (C) C – iii – IV	(D) D -	– iv – iii		
11.	Which of the following is correct i (Δ) $\Delta = i = 1$ (B) B = i		match?	(C) C – iii – I	ם (ח)	- ii - 1			
	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~		(2)2						
12.	Which	of the following	is correct	match?					
	(A) A - ii - III $(B) B -$		ii — III	(C) C – ii – III	(D) D ·	— ii — III			

Exercise-3

PART - I : JEE (ADVANCED) / IIT-JEE PROBLEMS (PREVIOUS YEARS)

* Marked Questions may have more than one correct option.

1. Reaction of Br_2 with Na_2CO_3 in aqueous solution gives sodium bromide and sodium bromate with evolution of CO_2 gas. The number of sodium bromide molecules involved in the balanced chemical equation is

[JEE 2011, 4/180]

- Dissolving 120 g of urea (mol. wt. 60) in 1000 g of water gave a solution of density 1.15 g/mL. The molarity of the solution is : [JEE 2011, 3/160]
 (A) 1.78 M
 (B) 2.00 M
 (C) 2.05 M
 (D) 2.22 M
- **3.** 29.2% (w/w) HCl stock solution has a density of 1.25 g mL⁻¹. The molecular weight of HCl is 36.5 g mol⁻¹. The volume (mL) of stock solution required to prepare a 200 mL solution of 0.4 M HCl is :

[JEE 2012, 4/136]

4.*For the reaction : $I^- + CIO_3^- + H_2SO_4 \longrightarrow CI^- + HSO_4^- + I_2$
The correct statement(s) in the balanced equation is/are :[JEE(Advanced) 2014, 3/120]
(A) Stoichiometric coefficient of HSO_4^- is 6.
(B) lodide is oxidized.
(C) Sulphur is reduced.(D) H_2O is one of the products.

5. A compound H_2X with molar weight of 80 g is dissolved in a solvent having density of 0.4 g ml⁻¹. Assuming no change in volume upon dissolution, the **molality** of a 3.2 molar solution is

[JEE(Advanced) 2014, 3/120]

6. The mole fraction of a solute in a solution is 0.1. At 298 K, molarity of this solution is the same as its molality. Density of this solution at 298 K is 2.0 g cm⁻³. The ratio of the molecular weights of the solute

and solvent,
$$\left(\frac{MW_{solute}}{MW_{solvent}}\right)$$
, is

7. The order of the oxidation state of the phosphorus atom in H_3PO_2 , H_3PO_4 , H_3PO_3 and $H_4P_2O_6$ is [JEE(Advanced) 2017]

(A) $H_3PO_4 > H_4P_2O_6 > H_3PO_3 > H_3PO_2$ (B) $H_3PO_3 > H_3PO_2 > H_3PO_4 > H_4P_2O_6$ (C) $H_3PO_2 > H_3PO_3 > H_4P_2O_6 > H_3PO_4$ (D) $H_3PO_4 > H_3PO_2 > H_3PO_3 > H_4P_2O_6$

8. The mole fraction of urea in an aqueous urea solution containing 900 g of water is 0.05. If the density of the solution is 1.2 g cm⁻³, the molarity of urea solution is _____ (Given data : Molar masses of urea and water are 60 g mol⁻¹ and 18 g mol⁻¹, respectively)

[JEE(Advanced) 2019]

[JEE(Advanced) 2016]

PART - II : JEE (MAIN) / AIEEE PROBLEMS (PREVIOUS YEARS)

1.	A 5.2 molal aqueous alcohol in the solution	solution of methyl alc n?	ohol, CH_3OH , is supplied. WI	hat is the mole fraction of methyl [AIEEE-2011, 3/120]
	(1) 0.100	(2) 0.190	(3) 0.086	(4) 0.050
2.	The molality of a ure is :	a solution in which 0.0	100 g of urea, $[(NH_2)_2CO]$ is a	dded to 0.3000 dm ³ of water at STP [Re. Paper AIEEE-2011, 3/120]
	(1) 5.55 × 10 ^{-₄}	(2) 33.3 m	(3) 3.33 × 10⁻² m	(4) 0.555 m

Mole <u>concept</u>

3.	The density of a solution 1.15 g/mL. The molarity (1) 0.50 M	on prepared by dissolving of this solution is : (2) 1.78 M	g 120 g of urea (mol. ma (3) 1.02 M	ass = 60 u) in 1000 g of water is [AIEEE-2012, 4/120] (4) 2.05 M			
4.	The ratio of number of oxygen atoms (O) in 16.0 g ozone (O ₃), 28.0 g carbon monoxide (CO) and 16.0 g oxygen (O ₂) is :-						
	(Atomic mass : $C = 12$,	O = 16 and Avogadro's	constant $N_A = 6.0 \times 10^2$	[AIEEE 2012 (Online)]			
	(1) 3 : 1 : 1	(2) 1 : 1 : 2	(3) 3 : 1 : 2	(4) 1 : 1 : 1			
5.	The molarity of a solution	n obtained by mixing 750) mL of 0.5(M) HCl with 2	50 mL of 2(M)HCI will be :			
	(1) 0.875 M	(2) 1.00 M	(3) 1.75 M	(4) 0.975 M			
6.	Consider the following re	eaction :					
	xMnO ₄ ⁻ + yC ₂ O ₄ ²⁻ + zH ⁺	\rightarrow xMn ²⁺ + 2yCO ₂ + $\frac{z}{2}$ H	٥				
	The values of x, y and z (1) 5, 2 and 16	in the reaction are, respectively 2, 5 and 8	ectively : (3) 2, 5 and 16	[JEE(Main)-2013, 4/120] (4) 5, 2 and 8			
7.	10 mL of 2(M) NaOH solu	ution is added to 200 mL of	0.5 (M) of NaOH solution	n. What is the final concentration			
	? (1) 0.57 M	(2) 5.7 M	(3) 11.4 M	[JEE(Main-online)-2013] (4) 1.14 M			
8.	The density of 3M solut	ion of sodium chloride is	s 1.252 g mL ^{-1} . The mola	ality of the solution will be (molar			
	mass, NaCl = 58.5 g m (1) 2.18 m	ol ⁻¹) (2) 3.00 m	(3) 2.60 m	[JEE(Main-online)-2013] (4) 2.79 m			
9.	In which of the following (a) $H_2O_2 + 2H^+ + 2e^{-1}$ (b) $H_2O_2 - 2e^{-1}$ (c) $H_2O_2 + 2e^{-1}$ (d) $H_2O_2 + 2OH^ 2e^{-1}$	$\begin{array}{l} \text{reactions H}_2O_2 \text{ acts as a} \\ e^- \longrightarrow 2H_2O \\ \Rightarrow O_2 + 2H^+ \\ \Rightarrow 2OH^- \\ 2e^- \longrightarrow O_2 + 2H_2O \end{array}$	reducing agent?	[JEE(Main)-2014, 4/120]			
	(1) (a), (b)	(2) (c), (d)	(3) (a), (c)	(4) (b), (d)			
10.	The amount of $BaSO_4$ for will be : (Ba = 137, Cl = 35.5, Section 2014)	rmed upon mixing 100 mL =32. H = I and O = 16)	of 20.8% BaCl ₂ solution v	vith 50 mL of 9.8% H ₂ SO ₄ solution [JEE(Main-online)-2014]			
	(1) 33.2 g	(2) 11.65 g	(3) 23.3 g	(4) 30.6 g			
11.	For the estimation of nitro	ogen, 1.4 g of an organic c	ompound was digested by	Kjeldahl method and the evolved			
	ammonia was absorbed	l in 60 mL of $\displaystyle rac{\mathrm{M}}{\mathrm{10}}$ sulphuri	c acid. The unreacted ac	id required 20 mL of $\frac{M}{10}$ sodium			
	hydroxide for complete	neutralizaton. The perce	ntage of nitrogen in the o	compound is : [JEE(Main-online)-2014]			
	(1) 3%	(2) 5%	(3) 6%	(4) 10%			
12.	The molecular formula (Mol. wt. 206). What wo gram resin?	of a commercial resin us uld be the maximum upta	sed for exchanging ions i ake of Ca ²⁺ ions by the re	n water softening is C ₈ H ₇ SO ₃ Na sin when expressed in mole per [JEE(Main)-2015, 4/120]			
	(1) $\frac{1}{103}$	(2) <u>1</u> 206	(3) $\frac{2}{309}$	(4) $\frac{1}{412}$			
13.	A sample of a hydrate of The dried sample weig Cl = 35.5 amu)	barium chloride weighing hed 52 g. The formula	61 g was heated until all t of the hydrated salt is :	he water of hydration is removed. (Atomic mass , Ba = 137 amu, [JEE(Main)-2015 (online)]			
	(1) BaCl ₂ .H ₂ O	(2) BaCl ₂ .4H ₂ O	(3) BaCl ₂ .3H ₂ O	(4) $BaCl_2.2H_2O$			

14.	At 300 K and 1 atm, 15 mL of a gaseous hydrocarbon requires 375 mL air containing 20% O ₂ by volume for complete combustion. After combustion the gases occupy 330 mL. Assuming that the water formed is in liquid form and the volumes were measured at the same temperature and pressure, the formula of the hydrocarbon is :- [JEE(Main)-2016]						
	(1) C ₄ H	H ₁₀	(2) C ₃ H ₆	(3) C ₃ H ₈	(4) C ₄ H ₈		
15.	The no	on-metal that doe	es not exhibit positive oxi	dation state is :	[JEE(Main)-2016 (online)]		
	(1) Fluo	orine	(2) Oxygen	(3) Chlorine	(4) lodine		
16.	5 L of a temper	n alkane requires rature and pressu	25 L of oxygen for its con ure, the alkane is :	nplete combustion. If all v	olumes are measured at constant [JEE(Main)-2016 (online)]		
	(1) But	ane	(2) Isobutane	(3) Ethane	(4) Propane		
17.	The m	ost abundant ele	ments by mass in the bo	ody of a healthy human a	adult are :		
	Oxyge would	n (61.4%) ; Carbo gain if all ¹ H ator	n (22.9%), Hydrogen (10. ms are replaced by ² H a	0%) ; and Nitrogen (2.6%) toms is:	. The weight which a 75 kg person [JEE(Main)-2017]		
	(1)	15 kg	(2) 37.5 kg	(3) 7.5 kg	(4) 10 kg		
18.	1 gram of M ₂ C	of a carbonate (N O ₃ in g mol ⁻¹ is	I_2CO_3) on treatment with C_3 :-	excess HCl produces 0.01	186 mole of CO ₂ . the molar mass [JEE(Main)-2017]		
	(1)	1186	(2) 84.3	(3) 118.6	(4) 11.86		
19.	Excess of NaOH (aq) was added to 100 mL of $FeCl_3(aq)$ resulting into 2.14 g of $Fe(OH)_3$. The molarit $FeCl_3(aq)$ is : [JEE(Main)-2017 (online)]						
	(Given molar mass of Fe = 56 gmol ^{-1} and molar mass of Cl = 35.5 g mol ^{-1})						
	(1) 1.8	Μ	(2) 0.2 M	(3) 0.6 M	(4) 0.3 M		
20.	The pa	ir of compounds	having metals in their h	ighest oxidation state is :			
	(1) Mn	O_2 and CrO_2Cl_2		(2) $[\text{FeCl}_4]^-$ and Co_2O_3	[JEE(Main)-2017 (online)]		
	(3)[Fe	(CN) ₆] ⁻³ and [Cu([CN) ₄] ²⁻	(4) $[NiCl_4]^{2-}$ and $[CoCl_4]$	2-		
21.	The rat compo comple	tio of mass percer und $(C_XH_YO_Z)$ co ately to CO ₂ and	nt of C and H of an organic ontains half as much oxy H_2O . The empirical form	c compound $(C_xH_YO_z)$ is (gen as required to burn c ula of compound C_xH_YO	6: 1. If one molecule of the above one molecule of compound C_XH_Y is : [JEE(Main)-2018]		
	(1) C ₂ H	I ₄ O	(2) C ₃ H ₄ O ₂	(3) C ₂ H ₄ O ₃	(4) C ₃ H ₆ O ₃		
22.	An unk atom c	nown chlorohydro nly; chlorine atoi	ocarbon has 3.55% of chlo ms present in 1 g of chlo	orine. If each molecule of rohydrocarbon are :	the hydrocarbon has one chlorine		
	(Atomi	c wt. of Cl = 35.5	5 u; Avogadro constant =	= 6.023 × 10 ²³ mol−1)	[JEE(Main)-2018 (online)]		
	(1) 6.0	23 × 10 ²¹	(2) 6.023 × 10 ²³	(3) 6.023 × 10 ²⁰	(4) 6.023 × 10 ⁹		
23.	For the	following reaction $\Omega(s) + 163\Omega(s)$	on, the mass of water protection $\rightarrow 114CO$ (d) + 110 H (oduced from 445 g of C_{53}	₇ H ₁₁₀ O ₆ is : [IFE(Main)-2019 (Ian)]		
	(1) 495	i g	(2) 490 g	(3) 890 g	(4) 445 g		
24.	The an	nount of sugar (C	$C_{12}H_{22}O_{11}$) required to pr	epare 2 L of its 0.1 M aq	ueous solution is :		
	(1) 68.4	4 g	(2) 17.1 g	(3) 34.2 g	(4)136.8 g [JEE-Main(Jan.)-2019]		

Mole concept

25.	A solution of sodium s solution in mol kg ⁻¹ i	ater. The molality of Na+ ions in that [JEE-Main(Jan.)-2019]								
	(1) 16	(2) 8	(3) 4	(4) 12						
26.	For a reaction,	[JEE(Main)-2019 (-April)]								
	$N_2(g) + 3H_2(g) \rightarrow 2N$	lH ₃ (g) ;								
	identify dihydrogen (H_2) as a limiting reagent in the following reaction mixtures.									
	(1) 14g of N ₂ + 4g of	H ₂	(2) 28g of N ₂ + 6g of	(2) 28g of N_2 + 6g of H_2						
	(3) 56g of N ₂ + 10g o	fH ₂	(4) 35g of N ₂ + 8g of	(4) 35g of N_2 + 8g of H_2						
27.	5 moles of AB ₂ weigh and molar mass of B	× 10 ⁻³ kg. The molar mass of A(M _A) [JEE(Main)-2019 (-April)]								
	(1) M _A = 50 × 10 ^{−3} a	nd $M_B = 25 \times 10^{-3}$	(2) M _A = 25 × 10 ^{−3} a	and $M_B = 50 \times 10^{-3}$						
	(3) $M_A = 5 \times 10^{-3}$ an	d M _B = 10 × 10 ⁻³	(4) $M_A = 10 \times 10^{-3} a$	and $M_B = 5 \times 10^{-3}$						
28.	What would be the molality of 20% (mass/mass) aqueous solution of KI?									
	(molar mass of KI =	= 166 g mol ^{_1})		[JEE-Main(April)-2019]						
	(1) 1.08	(2) 1.48	(3) 1.51	(4) 1.35						
29.	Oxidation number of	potassium in K ₂ O, K ₂ O ₂	and KO_2 , respectively,	is :						
				[JEE-Main(Jan)-2020]						
	(1) +1, +4 and +2	(2) +1, +2 and +4	(3)+1, +1 and +1	(4) +2 , +1 and $+\frac{1}{2}$						

30. Ferrous sulphate heptahydrate is used to fortify foods with iron. The amount (in grams) of the salt required to achieve 10 ppm of iron in 100 kg of wheat is _____. [JEE-Main(Jan)-2020]
 Atomic weight : Fe = 55.85 ; S = 32.0 ;
 O = 16.00

31. NaClO₃ is used, even in spacecrafts, to produce O₂. The daily consumption of pure O₂ by a person is 492L at 1 atm, 300K. How much amount of NaClO₃, in grams, is required to produce O₂ for the daily consumption of a person at 1 atm, 300 K ? [JEE-Main(Jan)-2020] NaClO₃(s) + Fe(s) \rightarrow O₂(g) + NaCl(s) + FeO(s) R = 0.082 L atm mol⁻¹ K⁻¹

-	ANSWER KEY											
	EXERCISE # 1											
	PART - I											
A-1.	(34.05 L	_)	A-2.	(100)	A-3.	5.40	B-1.	(4)	B-2.	(6)	B-3.	(C ₂ H ₄ O ₂)
B-4.	% CO ₂	$=\frac{2}{2+1+}$	<u>-</u> 2 × 10	0 = 40%.	C-1.	2.16 g	C-2.	42 g	C-3.	73 gm	C-4.	27.6 gm
C-5. D-2.	63 L 154 gm	, 24 gm	C-6. D-3.	58.8 gm 0.1185)	D-1. D-4.	(i) 1/6 n 0.25 ma	nole (ii) : ble	5/12 mol E-1.	le m = 1.4	łg	
E-2.	4		E-3.	$W_{CO_2} =$	= 528 grr	n E-5.	63%, 3 [°]	7%	E-6.	304 gm		
F-1.	(a) +3 (f) +3		(b) +5 (g) +1	-	(c) +6 (h) +2		(d) +2 (i) 200/9	93 = 2.15	(e) +8/3 5	}		
F-2.	(a)	(+7) KMnO∠	4 + KCI	+ H ₂ SO	$_{4} \longrightarrow$	(+2) MnSO ₄	+ K ₂ SC	$D_4 + H_2O$	+ $CI_{2}^{(0)}$.			
		(+7) KMnO4	ı (oxidaı	nt)	\longrightarrow	(+2) MnSO ₄	(reduct	ion half).				
	$ \overset{(-1)}{K} \overset{(-1)}{Cl} \text{ (reductant)} \longrightarrow \overset{(0)}{Cl_2} \text{ (oxidation half).} $											
	(b)	$FeCl_2 + H_2O_2 + HCl \longrightarrow FeCl_3 + H_2O $ (oxidation half)										
		(+2) FeCl ₂ (reductar	nt)→	(+3) • FeCl ₃	(oxidatio	on half).					
		$H_2^{(-1)}$ (c	oxidant)	\longrightarrow F	l ₂ O ²⁻ (rec	duction h	alf).					
	(c)	⁽⁰⁾ Cu + ⊦	(+5) 1NO ₃ (d	il) —→	²⁺ Cu (NC	$(D_3)_2 + H_2$	0 + NO					
		⁽⁰⁾ Cu (re	ductant)	$\longrightarrow C^2$	2+ Cu (NO ₃) ₂	oxidati	on half).					
		HNO_3	(oxidant)	NO (red	uction ha	alf).					
	(d)	Na ₂ HA	3 sO ₃ +	KBrO ₃ +	HCI —	\rightarrow Na	CI + KB	r + H ₃ A	sO ₄			
		Na ₂ HA	3 .sO ₃ (rea	ductant)	\longrightarrow F	H_3^{+5} AsO $_4$	(oxidatio	on half).				
		KBrO_3	(oxidar	ıt) →	_1 KBr.							
	(e)	$\stackrel{0}{\mathrm{I}_{2}}$ + N	$a_{2}^{+2}S_{2}O_{3}$	\longrightarrow N	+2.5 Na ₂ S ₄ O ₆	₀ + NaI						
		$\stackrel{0}{\mathrm{I}_2}$ (oxi	dant) —	\longrightarrow NaI	(reducti	on half).						
		$Na_{2}^{+2}S_{2}^{+2}C$	D ₃ (redu	ctant) —	$\rightarrow Na_2$	^{+2.5} S ₄ O ₆ (c	oxidation	half).				

G-1.	(a) S ₄ ($O_6^{2-} + 6A$	AI + 20 F	$d^+ \longrightarrow d^+ \to d^+$	$4H_2S +$	$6AI^{3+} + 6$	6H₂O						
	(D) OF	e +C	$_{2}O_{7} + 1^{4}$	$4\Pi \rightarrow 0$	ле + 2	2Cr + /	п ₂ 0						
	(c) IO	$_{4} + /1 +$	$-8H^{\circ} \rightarrow$	$4I_2 + 4F_2$	I ₂ 0								
	(d) 3N (e) 3H	$\ln O_4^{2-} + c$ $NO_2 - c$	$4H^+ \rightarrow 2$ $\rightarrow NO_3$	2MnO ₄ ⁻ + ₃ ⁻ + 2NO	$MnO_2 - H_2O$	+2H ₂ O + H⁺							
G-2.	(a) 3C	$210^{-} + 20$	$CrO_{2}^{-} + 2$	$2OH^{-} \rightarrow$	$3Cl^{-} + 2$	$2 CrO_4^{2-} +$	H ₂ O						
	(b) 4Z	$2n + NO_3$		$\rightarrow \mathrm{NH}_{2}$	$_3 + 4$ ZnO	$D_2^{2-} + 2H_2$	$_{2}O$						
	(c) 4S (d) 3M	+6OH ⁻ In(OH) -	$\rightarrow 2S^{2-}$ + 2MnO	$+S_2O_3^{2-}$	$+3H_2O$	+ 2H O -	+ 20H ⁻						
	(e) 2N	$4 n \Omega^{-} + S$	$SO^{2-} + 2$	₄ ′ ∩H⁻ →′	$2MnO^{2-}$	$\frac{1}{2} + SO^{2-}$	- H O						
H-1. H-3.	(i) 2.17 (i) 2.17 (vii) 25	7 m , 5%,	(ii) 6.2 (viii) 3	H-2. 25 M, 6.25%,	0.168 (iii) 0. (ix) 72	9 m 0376, 2.5%,	(iv) 0.0 (x) 14.)826, 2 m.	(v) 8%		(vi) 16	67%.	
H-4.	(i)	Na(0.2	$CI \rightarrow 2$	Na ⁺ + 0.2	CI⁻ 0.2								
	(ii)	Al ₂ ((SO ₄) .2	→ 2A 2.	1 ³⁺ + 4	3SO ₄ ⁻² 3.6							
	(iii)	H ₂ S	$O_4 \rightarrow 2$	2H⁺ 2.4	+ SC 1.) ₄ ⁻² 2							
	(iv)	Mg(2.	$CI_2 \rightarrow 1$	Mg ²⁺ 2.1	+ 20	CI⁻ ⊧.2							
I-1. I-3.	8 M (i)	0.32 N	1	I-2.	2.33	L							
	(ii)	9/7 M											
	(iii)	, 0.18 N	1										
J-1.	(iv) 34	0.24 N J-2.	1 50	J-3.	50	J-4.	17.46	J-5.	42.1				
						PA	RT - II						
A-1.	(B)	A-2.	(B)	B-1.	(B)	B-2.	(D)	B-3.	(C)	B-4.	(C)	C-1.	(C)
C-2.	(C)	C-3.	(C)	C-4.	(B)	C-5.	0	C-6.	(B)	D-1.	(A)	D-2.	(C)
D-3. F-4	(Α) (Δ)	D-4. E-5	(B) (D)	D-5. E-6	(B) (A)	D-6. F-1	(A) (D)	E-1. F-2	(A) (C)	E-2. F-3	(B) (B)	E-3. F-4	(A) (A)
F-5.	(C)	F-6.	(B)	G-1.	(C)	G-2.	(A)	G-3.	(C)	G-4.	(B)	G-5.	(D)
H-1.	(C)	H-2.	(B)	H-3.	(C)	H-4.	(D)	H-5.	(C)	H-6.	(A)	H-7.	(C)
I -1.	(A)	I -2 .	(C)	I -3.	(B)	I -4.	(D)	I -5 .	(C)	-6.	(B)	I -7.	(C)
J-1.	(B)	J-2.	(A)										

PART - III

- (A q,s); (B p, s); (C p, q, r); (D q, r) (A p,q,r,s; (B p,s; (C q,r); (D q) 1.
- 2.
- (A)-(P,Q,R,S) ; (B)-(Q,R,T) ; (C)-(Q,S) ; (D)-(P) 3.

	EXERCISE # 2												
	 PART - I												
1. 8. 15.	(C) (B) (B)	2. 9. 16.	(A) (B) (D)	3. 10. 17.	(C) (D) (A)	4. 11. 18.	(A) (A) (B)	5. 12. 19.	(A) (A) (A)	6. 13.	(C) (A)	7. 14.	(A) (D)
						PA	RT - II						
1. 7. 12	5 4 700 ml	2. 8. 13	78 50 27	3. 9. 14	(222 gr 10 10	n)	4. 10. 15	66.4 % 8 4	%. 5. 11. 16	33.33 2 60	8%	6. 17	28 56
18.	01.14	10.	21	14.	10		10.	-	10.	00			50
	PART - III												
1. 7. 13.	(ABC) (AC (ACD)	2. 8. 14.	(AB) (ABC) (AB)	3. 9. 15.	(BCD) (BD) (ACD)	4. 10. 16.	(AB) (AD) (ABD)	5. 11. 17.	(BCD) (CD) (ABD)		6. 12.	(BC) (BC)	
						PA	RT - IV						
1. 8.	(A) (B)	2. 9.	(B) (B)	3. 10.	(B) (A)	4. 11.	(C) (B)	5. 12.	(B) (D)	6.	(B)	7.	(A)
					E	XER	CISE #	± 3					
						PA	RT - I						
1.	5	2.	(C)	3.	8 mL.	4.	(ABD)	5.	8	6.	(9)	7.	(A)
8.	2.98						от и						
						PA	R I - II						
1. 8	(3) (D)	2. 9	(1) (4)	3. 10	(4) (B)	4. 11	(4) (D)	5. 12	(1) (4)	6. 13	(3) (4)	7. 14	(A) (3)
15.	(1)	э. 16.	(4)	17.	(3)	18.	(2)	19.	(2)	20.	(3)	21.	(3)
22.	(3)	23.	(1)	24.	(1)	25.	(3)	26.	(3)	27.	(3)	28.	(3)
29.	(3)	30.	4.95 to	4.97	31.	2120	to 2140						

Reliable Ranker Problems

This Section is not meant for classroom discussion. It is being given to promote self-study and self testing amongst the Resonance students.

Self Assessment Test

PART- 1 : PAPER JEE (MAIN) PATTERN

		SECTION-I : (Max	ximum Marks : 80)					
•	This section contains TWENTY questions.							
•	Each question has FOUR options (A), (B), (C) and (D). ONLY ONE of these four options is correct.							
•	For each question, dar	ken the bubble correspo	onding to the correct opt	tion in the ORS.				
•	For each question, ma	rks will be awarded in <u>c</u>	one of the following cate	gories :				
	Full Marks : +4If only Zero Marks : 0 If pope	the bubble corresponding of the bubbles is dark	ng to the correct option i	is darkened.				
	Negative Marks : -1 Ir	all other cases	eneu.					
1.	112.0 mL of NO ₂ at 0°C and 1 atm was liquefied, the density of the liquid being 1.15 g mL ⁻¹ . Calculate the volume and the number of molecules in the liquid NO ₂ . (A) 0.10 mL and 3.01×10^{22} (B) 0.20 mL and 3.01×10^{21} (C) 0.20 mL and 6.02×10^{23} (D) 0.40 mL and 6.02×10^{21}							
2.	Common salt obtained f of NaCl present in 10.0	rom sea - water contains g of the common salt is :	96% NaCl by mass. The a (At. wt. Na = 23)	pproximate number of molecules				
	(A) 10 ²¹	(B) 10 ²²	(C) 10 ²³	(D) 10 ²⁴				
3.	X and Y are two elements which form $X_2 Y_3$ and $X_3 Y_4$. If 0.20 mol of $X_2 Y_3$ weighs 32.0 g and 0.4 mol of $X_3 Y_4$							
	weighs 92.8 g, the atomic weights of X and Y are respectively (A) 16.0 and 56.0 (B) 8.0 and 28.0 (C) 56.0 and 16.0 (D) 28.0 and 8.0							
4.	Consider the following s 1. If all the reactants are 2. 2 moles of $H_2(g)$ and 3. equal wt. of carbon are The above statements 1	tatements : a not taken in their stoichi 3 moles of $O_2(g)$ produce nd oxygen are taken to pr I, 2, 3 respectively are (T	ometric ratio, then at leas 2 moles of water. roduce CO_2 then O_2 is lim = True, F = False)	t one reactant will be left behind. iting reagent.				
	(A) T T T	(B) F T F	(C) F F F	(D) T F T				
5.	Which of the following equations is a balanced one : (A) $5BiO_3^- + 22H^+ + Mn^{2+} \longrightarrow 5Bi^{3+} + 7H_2O + MnO_4^-$ (B) $5BiO_3^- + 14H^+ + 2Mn^{2+} \longrightarrow 5Bi^{3+} + 7H_2O + 2MnO_4^-$ (C) $2BiO_3^- + 4H^+ + Mn^{2+} \longrightarrow 2Bi^{3+} + 2H_2O + MnO_4^-$							
	(D) $6BIO_3^- + 12H^+ + 3IM$	$0^{2*} \longrightarrow 6Bl^{3*} + 6H_2O$	$+ 3MnO_4^-$					
6.	During the disproportion alkaline medium is :	ation of lodine to iodide a	nd iodate ions, the ratio o	f iodate and iodide ions formed in				
	(A) 1 : 5	(B) 5 : 1	(C) 3 : 1	(D) 1 : 3				
7.	How much NaNO $_3$ mus per mL?	t be weighed out to mak	e 50 ml of an aqueous s	olution containing 70 mg of Na⁺				
	(A) 12.394 g	(B) 1.29 g	(C) 10.934 g	(D) 12.934 g				

8.	The strength of 10^{-2} M Na ₂ CO ₃ solution in terms of molality will be (density of solution = 1.10 g mL ⁻¹). (Molecular weight of Na ₂ CO ₃ = 106 g mol ⁻¹) (A) 9.00 x 10^{-3} (B) 1.5 x 10^{-2} (C) 5.1 x 10^{-3} (D) 11.2 x 10^{-3}					
•				(D) 11.2 × 10		
9.	(A) 273 K	(B) 298 K	(C) 277 K	(D) None		
10.	What is the molarity of H (A) 4.18 M	H_2SO_4 solution that has a (B) 8.14 M	density 1.84 g/cc at 35ºC (C) 18.4 M	and contains 98% by weight- (D) 18 M		
11.	5.85 g of NaCl is dissolv (A) 6.02 × 10 ¹⁹	red in 1 L of pure water. T (B) 1.2 × 10 ²²	he number of ions in 1 m (C) 1.2×10^{20}	L of this solution is (D) 6.02 × 10 ²⁰		
12.	A compound is compose	ed of 74% C, 8.7% H and	17.3% N by mass. If the n	nolecular mass of the compound		
	IS 162, what is its molec (A) C_5H_7N	$(B) C_{10}H_{16}N_2$	(C) $C_8 H_{14} N_3$	(D) $C_{10}H_{14}N_2$		
13.	Calculate the volume of (A) 1891.67 L O_2	O_2 needed for combustic (B) 3733.33 L O_2 .	on of 1 kg of carbon at ST (C) 933.33 L O ₂ .	$P. C + O_2 \xrightarrow{\Delta} CO_2.$ (D) 4666.67 L O ₂ .		
14.	Li metal is one of the few reaction is :	v substances that reacts c	lirectly with molecular nitr	ogen. The balanced equation for		
	How many grame of the	$6\text{Li}(s) + N_2(g) -$	$\longrightarrow 2Li_3N(s)$	of lithium motal and 8.4 g of		
	molecular nitrogen ?	product, infinant fininde, c	ande prepared nom 5.5g	y of human metal and 0.4 g of		
	(A) 21.00 g of Li ₃ N.	(B) 2.91 g of Li ₃ N.	(C) 5.83 g of Li ₃ N.	(D) 10.50 g of Li ₃ N.		
15.	Potassium super oxide, $0.15 \text{ mol KO}_2 \text{ and } 0.10 \text{ m}$ $2\text{KO}_2 + 2\text{H}_2\text{O} \longrightarrow 2\text{K}$ (A) H_2O limiting reagent	KO_2 , is used in rebreathin nol H_2O , what is the limitin $COH + H_2O_2 + O_2$ c, 0.05 mol of O_2 .	ng gas masks to generate g reactant ? How many m (B) KO ₂ limiting reagent,	O_2 . If a reaction vessel contains oles of oxygen can be produced? 0.05 mol of O_2 .		
	(C) H_2O limiting reagent	t, 0.075 mol of O ₂ .	(D) KO_2 limiting reagent	:, 0.075 mol of O ₂ .		
16.	A 1 g sample of $KCIO_3$ v equation. (i) $2KCIO_3 \longrightarrow 2KCI$ and the remaining und (ii) $4KCIO_3 \longrightarrow 3KCI$ If the amount of O_2 evolv	vas heated under such co + $3O_2$ derwent change according IO_4 + KCI ved was 146.8 mL at 0°C	onditions that a part of it d g to the equation and 1 atm, calculate the p	ecomposed according to the bercentage by weight of $KCIO_4$ in		
	the residue.	(B) 49.8 %	(C) 62 5 %	(D) 87 1 %		
17.	A piece of aluminium we mass). After the metal is H ₂ SO, in the resulting so	eighing 2.7 g is heated wire carefully dissolved the solution.	th 75.0 ml of H_2SO_4 (sp. golution is diluted to 400m	gr. 1.2 containing 25% H_2SO_4 by I. What is the molarity of the free		
	(Ă) 1.056 M	(B) 0.560 M	(C) 0.312 M	(D) 0.198 M		
18.	100 ml of 0.15 M solution weight.	n of $Al_2(SO_4)_3$, the density	of the solution is 1.5 g/ml	. Report the no. of AI^{3+} ions in this		
	(A) 1.8 × 10 ²⁵ ions	(B) 6 × 10 ²² ions	(C) 1.8 × 10 ²³ ions	(D) 1.8 × 10 ²² ions		
19.	5 g sample of $CuSO_4$. 5 precipitate (BaSO ₄) obtathe sample.	H_2O was dissolved in wate ained was washed and dr	er. BaCl ₂ solution was mix ied, it weighed 4.66 g. W	ted in excess to this solution. The hat is the % of SO_4^{2-} by weight in		
	(A) /6.8%	(B) 38.4%	(C) 51%	(D) 19.2%		
20.	At room temperature, th of ethanol contains the s (A) 418.95 ml.	e density of water is 1.0 g same number of molecule (B) 736.33 ml.	/mI and the density of eth es as are present in 175 n (C) 566.82 mI.	anol is 0.789 g/ml. What volume nl of water ? (D) 911.84 ml.		

	SECTION-II : (Maximum Marks: 20)
~	This section contains FIVE questions.
~	The answer to each question is a NUMERICAL VALUE.
~	For each question, enter the correct numerical value (If the numerical value has more than two decimal places, truncate/round-off the value to TWO decimal places; e.g. 6.25, 7.00, -0.33,30, 30.27, -127.30, if answer is 11.36777 then both 11.36 and 11.37 will be correct) by darken the corresponding bubbles in the ORS.
	For Example : If answer is -77.25, 5.2 then fill the bubbles as follows.
~	Answer to each question will be evaluated according to the following marking scheme: <i>Full Marks</i> : +4 If ONLY the correct numerical value is entered as answer.
21.	The sodium salt of methyl orange has 7% sodium. What is the minimum molecular weight of the compound?
22.	A 10.0 g sample of a mixture of calcium chloride and sodium chloride is treated with Na_2CO_3 solution. This calcium carbonate is heated to convert all the calcium to calcium oxide and the final mass of calcium oxide is 1.62 g. The percentage by mass of calcium chloride in the original mixture is ?
23.	$2KI + I_2 + 22 HNO_3 \longrightarrow 2HIO_3 + 2KIO_3 + 22NO_2 + 10H_2O$ If 3 mole of KI & 2 moles I ₂ are reacted with excess of HNO ₃ . Volume (in lit.) of NO ₂ gas evolved at 0°C and 1 atm is
24.	Minimum amount (in gm) of Ag ₂ CO ₃ (s) required to produce sufficient oxygen for the complete combustion of C_2H_2 which produces 11.2 ltr of CO ₂ at 0°C and 1 atm after combustion is: [Ag = 108] Ag ₂ CO ₃ (s) \longrightarrow 2Ag (s) + CO ₂ (g) + 1/2O ₂ (g) $C_2H_2 + 5/2O_2 \longrightarrow 2CO_2 + H_2O$
25.	In the reaction $4A + 2B + 3C \longrightarrow A_4B_2C_3$ what will be the number of moles of product formed, starting from 2 moles of A, 1.2 moles of B & 1.44 moles of C?

PART - 2 : PAPER JEE (ADVANCED) PATTERN

SECTION-I : (Maximum Marks : 12)

- This section contains **FOUR** questions.
- Each question has **FOUR** options (A), (B), (C) and (D). **ONLY ONE** of these four options is correct.
- For each question, darken the bubble corresponding to the correct option in the ORS.
- For each question, marks will be awarded in <u>one of the following categories</u> :

Full Marks	+3	If only the bubble corresponding to the correct option is darkened.
Zero Marks	0	If none of the bubbles is darkened.
Negative Marks	—1	In all other cases

- 2.Equal weight of Zn metal and iodine are mixed together and the iodine is completely converted to ZnI_2 . What
fraction of weight of the original Zinc remains unreacted. (Atomic wt. Zn = 65)
(A) 0.500(B) 0.744(C) 0.488(D) 0.256
- One litre of a mixture of CO and CO₂ is passed through red hot charcoal in tube. The new volume becomes 1.4 litre. Find out % composition of mixture by volume. All measurements are made at same P and T. (A) CO₂ 40%, CO 60% (B) CO₂ 60%, CO 40% (C) CO₂ 25%, CO 75% (D) CO₂ 30%, CO 70%
- 4. Amongst the following, the pair having both the metals in their highest oxidation state is : (A) $[Fe(CN)_6]^3$ and $[Co(CN)_6]^3$ -(C) TiO_2 and MnO_2 (D) $[MnCl_4]^2$ - and $[NiF_6]^2$ -

SECTION-II: (Maximum Marks: 32)

This section contains **EIGHT** questions.

Each question has FOUR options for correct answer(s). ONE OR MORE THAN ONE of these four option(s) is (are) correct option(s).

For each question, choose the correct option(s) to answer the question.

Full Marks	:	+4	If only (all) the correct option(s) is (are) chosen.
Partial Marks	:	+3	If all the four options are correct but ONLY three options are chose
Partial Marks	:	+2	If three or more options are correct but ONLY two options are chos
	both	of which	n are correct options.
Partial Marks	:	+1	If two or more options are correct but ONLY one option is chosen
	and	t is a co	prrect option.
Zero Marks	:	0	If none of the options is chosen (i.e. the question is unanswered).
Negative Marks	s:	-1	In all other cases.
For Example :	If firs	st, third a	and fourth are the ONLY three correct options for a question with seco

option being an incorrect option; selecting only all the three correct options will result in +4 marks. Selecting only two of the three correct options (e.g. the first and fourth options), without selecting any incorrect option (second option in this case), will result in +2 marks. Selecting only one of the three correct options (either first or third or fourth option), without selecting any incorrect option (second option in this case), will result in +1 marks. Selecting any incorrect option(s) (second option in this case), with or without selection of any correct option(s) will result in -1 marks.

- 5. 1120 mL of ozonised oxygen at 0°C and 1 atm weigh 1.76 g. Calculate the volume of oxygen in the ozonised oxygen. (A) It contain 400 mL O_{2} (B) It contain 224 mL O₃ (C) It contain 400 mL O_3 (D) It contain 896 mLO₂
- 6. A 5L vessel contains 2.8 g of N₂. When heated to 1800 K, 30% molecules are dissociated into atoms. (A) Total no. of moles in the container will be 0.13
 - (B) Total no. of molecules in the container will be close to 0.421 x 10²³.
 - (C) Total no. of moles in the container will be 0.098.
 - (D) All of these are correct.
- 7. Equal masses of SO₂ and O₂ are placed in a flask at 0°C and 1 atm choose the correct statement. (A) The number of molecules of O₂ are more than SO₂
 - (B) Volume occupied at 0°C and 1 atm is more for O₂ than SO₂
 - (C) The ratio of number of atoms of SO_2 and O_2 is 3 : 4.
 - (D) Moles of SO₂ is greater than the moles of O₂.
- 8. For the reaction $2P + Q \rightarrow R$, 12 mol of P and 8 mol of Q are taken then (A) 3 mol of R is produced (B) 6 mol of R is produced
 - (C) 25% of Q is left behind (D) 25% of Q has reacted

9.
$$\begin{array}{l} C_{2}H_{5} \\ (CH-COOH)_{n} + AgNO_{3}(Excess) \longrightarrow Silver salt \longrightarrow Ag (metal) \\ C_{2}H_{5} \\ If 0.5 \text{ mole of silver salt is taken and weight of residue obtained is 216 g. (Ag = 108 g/n) \\ Then which the following is correct is$$

nol). Then which the following is correct :

(A) $n = 4$	(B) $n = 2$
(C) M.wt. of silver salt is 718 g/mol	(D) M.wt. of silver salt is 388 g/mol

10.为	21.2 g sample of impure Na_2CO_3 is dissolved and reacted with a solution of $CaCl_2$, the weight of precipitate of $CaCO_3$ is 10.0 g. Which of the following statements is/are correct? (A) The % purity of Na_2CO_3 is 50% (B) The percentage purity of Na_2CO_3 is 60% (C) The number of moles of $Na_2CO_3 = CaCO_3 = 0.1$ mol. (D) The number of moles of $NaCl$ formed is 0.1 mol.							
11.24	In the following reaction : $Cr(OH)_3 + OH^- + IO_3^-$ (A) IO_3^- is oxidising agent (C) 6e ⁻ are being taken per iodine atom	$T \rightarrow CrO_4^{2-} + H_2O + I^-$ (B) Cr(OH) ₃ is oxidise (D) None of these	d					
12.	Select dimensionless quantity(ies) : (A) vapour density (B) molality	(C) specific gravity	(D) mass fraction					
•	SECTION-III : (M This section contains SIX questions. The answer to each question is a NUMERIC For each question, enter the correct numeric second decimal place; e.g. 6.25, 7.00, -0.33 11.36 and 11.37 will be correct) by darken th For Example : If answer is -77.25, 5.2 the Answer to each question will be evaluated a <i>Full Marks</i> : +3 If ONLY the correct of <i>Zero Marks</i> : 0 In all other cases.	Maximum Marks: 18) AL VALUE. al value (in decimal nota 3,30, 30.27, -127.30, i ne corresponding bubble n fill the bubbles as follo ccording to the following numerical value is entere	ation, truncated/rounded-off to the f answer is 11.36777 then both as in the ORS. ows. g marking scheme: ed as answer.					
13.	Average atomic mass of magnesium is 24.31 a remaining 21 mole % of ²⁵ Mg and ²⁶ Mg. Calcula 0.1.	.m.u. This magnesium is c ate mole % of ²⁶ Mg. Repo	composed of 79 mole % of ²⁴ Mg and rt your answer after multiplying by					
14.	200 g impure CaCO $_{3}$ on heating gives 5.6 lt. CC sample.	P_2 gas at STP. Find the perc	centage of calcium in the lime stone					
15.	Molarity of H_2SO_4 is 18 M. Its density is 1.8 g/c answer x/500).	m³, hence molality is (If y	our answer is 'x' then, Report your					
16.	1 g of a mixture of equal number of moles of Li ₂ C of the reactions. $Li_2CO_3 + 2HCI \longrightarrow 2LiCI + H_2O + CC$ $M_2CO_3 + 2HCI \longrightarrow 2 MCI + H_2O + CC$ If the atomic mass of Li is 7, then find the Atom Report M – 16.	CO_3 and M_2CO_3 required 4 CO_2 CO_2 mic mass of M.	14.44 ml of 0.5 M HCl for completion					
17.æ	A 3 : 2 molar ratio mixture of FeO and Fe_2O_3 re and Fe_2O_3 . Find the mass (in g) of O_2 gas requi	act with oxygen to produc red per mole of the initial	e a 2 : 3 molar ratio mixture of FeO mixture.					
18.	92 g mixture of CaCO $_3$, and MgCO $_3$ heated structure carbonates it was found that the weight of reside the mixture.	ongly in an open vessel. A due left behind is 48 g. Fi	fter complete decomposition of the ind the mass of $MgCO_3$ in grams in					

	PAR	T-3:OLYMPI	IAD (PREVIOUS Y	EARS)	
1.	The volume of 16g of o $(A) 2.24 \text{ dm}^3$	xygen at S.T.P. is : (B) 11.2 dm ³	(C) 22.4 dm ³	(D) 8 dm ³	[NSEC-2000]
2.	When 25 g of Na $_2$ SO $_4$ is (A) 2.5 ppm	s dissolved in 10³ Kg o (B) 25 ppm	f solution, its concentration (C) 250 ppm	will be (D) 100 ppm	[NSEC-2000]
3.	Consider the following of Element A B	lata Ato	mic weight 12.01 35.5		[NSEC-2000]
	A and B combine to for then the weight of one r (A) 154.0 g	m new substance X. If nole of X is (B) 74.0 g	f 4 moles of B combines wit (C) 47.5 g	h 1 mole of A to g (D) 166.0 g	ive 1 mole of X, [NSEC-2000]
4.	Oxidation Number of M (A) –7	In in [MnO₄]⁻ is : (B) + 7	(C) + 2	(D) – 2	[NSEC-2001]
5.	The amount of salt requ (A) 0.05 mole	uired to prepare 10 dm (B) 0.02 mole	³ of decimolar solution is : (C) 0.01 mole	(D) 1.00 mole	[NSEC-2001]
6.	If 1 dm³ of a gas weight (A) 56 g	ts 2.5 g at STP, its grai (B) 11.2 g	m-molecular weight is : (C) 22.4 g	(D) 224 g	[NSEC-2001]
7.	How many moles of air the person is at 1.0 atm (A) 0.15 mol	are there in the lungs of pressure and has nor (B) 0.25 mol	of an average adult with a lu rmal body temperature at 3 (C) 1.15 mol	ng capacity of 3.8 7°C). (D) 2.25 mol.	L . (Assume that [NSEC-2002]
8.	The sterile saline soluti water and diluting to 10 (A) 0.00684 M	on used to rinse conta 0 mL. The molarity of (B) 0.09564 M	ct lenses can be made by d the solution will be of (C) 1.0684 M	issolving 400 mg (D) 0.0684 M	of NaCl in sterile [NSEC-2002]
9.	A molal solution contain (A) one litre of solution (C) one litre of the solve	ns one gram mole of s ent	olute in : (B) 1000 g of the solve (D) 22.4 litre of the sol	nt ution	[NSEC-2002]
10.	An average cup of coffe a cup ? (A) 8.33 × 10⁻³	ee contains about 125 (B) 6.44×10^{-4}	mg of caffeine, $C_8 H_{10} N_4 O_2$. (C) 6.234 × 10 ⁻²³	How many moles ((D) none of the	of caffeine are in [NSEC-2002] se
11.	Cystine has a sulphur c weight ? (A) 240	ontent of 26.7%. If its (B) 24	molecule contains two ator (C) 2400	ns of sulphur, wha (D) 120.	at is its molecular [NSEC-2002]
12.	Which of the following s (A) 0.46 g of C_2H_5OH in (C) 0.23 g of CH ₃ OH in	solutions are unimolar 10 mL of solution 100 mL of solution	solutions ? (B) 110.98 g of $CaCl_2$ i (D) 5.88 g of NaCl in 1	n 1000 mL of solu 000 mL of solutior	[NSEC-2002] tion n.
13.	1.00 g of a pure elemer (A) U	nt contains 4.39 × 10 ²¹ (B) Ce	atoms. The element is (C) Ba	(D) Au.	[NSEC-2003]
14.	The maximum amount	of $CH_{3}CI$ that can be p	prepared by reacting 20.0 g	of CH_4 with 10.0 g	of Cl ₂ is [NSEC-2003]
	(A) 30.0 g	(B) 7.1 g	(C) 63.1 g	(D) 31.6 g	
15.	Considering air as a 4: 5 m \times 5 m \times 5 m at STI (A) 160 g	I mixture of nitrogen a P is approximately (B) 160 kg	nd oxygen, the mass of air (C) 16 g	in a hall with dime (D) 1.60 kg.	ensions [NSEC-2003]

Mole concept

16.	The molar concentration (A) 1	n of pure water at 4ºC and (B) 5.5	1 atm pressure is (C) 18	(D) 55.5	[NSEC-2003]
17.	A mixture of aluminium a hydrogen gas (measure approximately	and zinc weighing 1.67 g v d at 273 K and 1 atm pres	was completely dissolved sure). The amount of alu	l in acid and evol minium in the ori	ved 1.69 L of ginal mixture is [NSEC-2004]
	(A) 1.8 g	(B) 2.0 g	(C) 1.2 g	(D) 2.2 g	
18.	The largest number of m (A) CO_2	nolecules is present in 1 g (B) H ₂ O	of (C) C_2H_5OH	(D) N ₂ O ₅ .	[NSEC-2004]
19.	20 g of solute X are disso of the solutes in these tw that of the solute Y is	lved in 50 g of water. 15 g vo solutions are the same	of solute Y are dissolved in the solute, the ratio of the solute of the	in 70 g of benzen molecular weigh	e. The molalities ts of solute X to [NSEC-2004]
	(A) 7:5	(B) 4:3	(C) 15:28	(D) 28:15	
20.	If 0.5 mol of $BaCl_2$ is mix formed is	ed with 0.2 mol of Na ₃ PO	₄ , the maximum number	of moles of $Ba_{_3}(F)$	$PO_4)_2$ that can be [NSEC-2004]
	(A) 0.1	(B) 0.2	(C) 0.5	(D) 0.7	
21.	The total number of elec (A) 4.8×10^{24}	etrons present in 8.0 g of r (B) 3.01 × 10 ²⁴	nethane is (C) 4.8 × 10 ²⁵	(D) 3.01 × 10 ²³ .	[NSEC-2004]
22.	The percentage abundat a.m.u) is	nces of ¹² C and ¹³ C are 98	.9 and 1.1 respectively. T	he average mass	s of carbon (in [NSEC-2005]
	(A) 12.111	(B) 12.981	(C) 12.011	(D) 12.891	
23.	1000 mL of a gas weigh (A) 22.4 g	s 1.5 g at NTP. Its gram n (B) 33.6 g	olecular weight is (C) 11.2 g	(D) 15 g.	[NSEC-2005]
24.	0.1 g of an element cont (A) Ga	ains 4.39 × 10²º atoms. T (B) Ce	he element is (C) Pb	(D) Ba.	[NSEC-2005]
25.	The percentages of C, H	and N in an organic comp	ound are 40%, 13.3% and	d 46.7%. The emp	pirical formula of
					[NSEC-2006]
	this compound is (A) CH_2N	(B) CH ₄ N	(C) CH ₅ N	(D) $C_3 H_9 N_3$.	[]
26.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25-25	(B) CH₄N otopes with masses 24 , 2 ge mass of the isotope mix	(C) CH_5N 25 and 26 with relative at ature would be (C) 24.50	(D) $C_3H_9N_3$.	, 15% and 5% [NSEC-2006]
26.	this compound is (A) CH_2N An element has three is respectively. The averag (A) 25.25 17.1 grams of eluminum	(B) CH ₄ N otopes with masses 24 , 2 ge mass of the isotope mix (B) 25.50	(C) CH ₅ N 25 and 26 with relative at cture would be (C) 24.50	(D) $C_3H_9N_3$. bundance of 80% (D) 24.25	, 15% and 5% [NSEC-2006]
26. 27.	this compound is (A) CH_2N An element has three is respectively. The averag (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 sulfate $AI_2(SO_4)_3$ is disso the ion the solution? (Negle (B) 5.00 x 10 ⁻² M	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 wed in enough water to p ect any hydrolysis) (C) 1.50 x 10^{-1} M	(D) C ₃ H ₉ N ₃ . bundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ I	5, 15% and 5% [NSEC-2006] solution .What is [NSEC-2007]
26. 27. 28.	this compound is (A) CH_2N An element has three is respectively. The averag (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M Chlorine can be prepare	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 sulfate Al ₂ (SO ₄) ₃ is disso te ion the solution? (Negle (B) 5.00 x 10^{-2} M d by reacting HCl with Mr	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 wed in enough water to p ect any hydrolysis) (C) 1.50 x 10^{-1} M inO ₂ . The reaction is repre	(D) $C_3H_9N_3$. bundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M sented by the eq	o, 15% and 5% [NSEC-2006] solution .What is [NSEC-2007] M uation:
26. 27. 28.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M Chlorine can be prepare $MnO_2(s) + 4HC$	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 a sulfate Al ₂ (SO ₄) ₃ is disso the ion the solution? (Negle (B) 5.00 x 10 ⁻² M d by reacting HCl with Mr $L_{(aq)} \longrightarrow Cl_2(g) + MnCl$	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (ved in enough water to p ect any hydrolysis) (C) 1.50 x 10^{-1} M 10_2 . The reaction is repre $_{2(aq)}$ + 2H ₂ O(I)	(D) $C_3H_9N_3$. bundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M esented by the eq	o, 15% and 5% [NSEC-2006] solution .What is [NSEC-2007] M uation:
26. 27. 28.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M Chlorine can be prepare MnO ₂ (s) + 4HC Assuming the reaction g needed to produce 2 50	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 sulfate Al ₂ (SO ₄) ₃ is disso the ion the solution? (Negle (B) 5.00 x 10 ⁻² M d by reacting HCl with Mr L _(aq) \longrightarrow Cl ₂ (g) + MnCl oes to completion, what r a of Cl	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (C) 24.50 (C) 1.50 x 10 ⁻¹ M (C) 1.50 x 10 ⁻¹ M (C) 1.50 x 10 ⁻¹ M (C) 1.50 x 10 ⁻¹ M	(D) $C_3H_9N_3$. bundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M esented by the eq	(NSEC-2006) [NSEC-2006] solution .What is [NSEC-2007] M uation: HCl by mass) is [NSEC-2007]
26. 27. 28.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M Chlorine can be prepare MnO ₂ (s) + 4HC Assuming the reaction g needed to produce 2.50 (A) 5.15 g	(B) CH ₄ N otopes with masses 24 , 2 ge mass of the isotope mix (B) 25.50 a sulfate Al ₂ (SO ₄) ₃ is disso a ion the solution? (Negle (B) 5.00 x 10 ⁻² M d by reacting HCl with Mr L _(aq) \longrightarrow Cl ₂ (g) + MnCl oes to completion, what r g of Cl ₂ (B) 14.3 g	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (Ved in enough water to p ect any hydrolysis) (C) 1.50 x 10^{-1} M $M_{2^{c}}$. The reaction is repre $M_{2(aq)} + 2H_2O(I)$ mass of concentrated HC (C) 19.4 g	(D) $C_3H_9N_3$. bundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ f sented by the eq I solution (36.0% (D) 26.4 g	, 15% and 5% [NSEC-2006] solution .What is [NSEC-2007] M uation: HCl by mass) is [NSEC-2007]
26. 27. 28. 29.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M Chlorine can be prepare MnO ₂ (s) + 4HC Assuming the reaction g needed to produce 2.50 (A) 5.15 g How many moles of Nat (A) 0.008	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 a sulfate Al ₂ (SO ₄) ₃ is disso the ion the solution? (Negle (B) 5.00 x 10 ⁻² M d by reacting HCl with Mr L _(aq) \longrightarrow Cl ₂ (g) + MnCl oes to completion, what r g of Cl ₂ (B) 14.3 g ions are there in 20mL of (B) 0.020	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (C) 24.50 (C) 1.50 x 10 ⁻¹ M M_{2} . The reaction is repre $2(aq) + 2H_2O(I)$ mass of concentrated HC (C) 19.4 g 5 0.40M solution of Na ₃ PC (C) 0.024	(D) $C_{3}H_{9}N_{3}$. pundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M esented by the eq I solution (36.0% (D) 26.4 g D_{4} ? (D) 0.008	(INSEC-2006) [NSEC-2006] solution .What is [NSEC-2007] wi uation: HCl by mass) is [NSEC-2007] [NSEC-2007]
26. 27. 28. 29.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10 ⁻² M Chlorine can be prepare MnO ₂ (s) + 4HC Assuming the reaction g needed to produce 2.50 (A) 5.15 g How many moles of Na ⁺ (A) 0.008 Oxalic acid ($H_2C_2O_4$) rea	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 a sulfate Al ₂ (SO ₄) ₃ is disso a ion the solution? (Negle (B) 5.00 x 10^{-2} M d by reacting HCl with Mr L _(aq) \longrightarrow Cl ₂ (g) + MnCl oes to completion, what r g of Cl ₂ (B) 14.3 g ions are there in 20mL of (B) 0.020 acts with permanganate io	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (Ved in enough water to p ect any hydrolysis) (C) 1.50 x 10^{-1} M 10_2 . The reaction is repre $2(aq) + 2H_2O(I)$ mass of concentrated HC (C) 19.4 g 10.40M solution of Na ₃ PC (C) 0.024 n according to the balance	(D) $C_{3}H_{9}N_{3}$. bundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M sented by the eq I solution (36.0% (D) 26.4 g D_{4} ? (D) 0.008 ced equation give	(
26. 27. 28. 29. 30.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10 ⁻² M Chlorine can be prepare MnO ₂ (s) + 4HC Assuming the reaction g needed to produce 2.50 (A) 5.15 g How many moles of Nat (A) 0.008 Oxalic acid (H ₂ C ₂ O ₄) reac 5H ₂ C ₂ O _{4(aq)} + 2N	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 a sulfate Al ₂ (SO ₄) ₃ is dissoned ion the solution? (Neglet (B) 5.00 x 10 ⁻² M d by reacting HCl with Mr L _(aq) \longrightarrow Cl ₂ (g) + MnCl oes to completion, what r g of Cl ₂ (B) 14.3 g ions are there in 20mL of (B) 0.020 acts with permanganate ion AnO_4 (aq) + $6H^+_{(aq)} \rightarrow 2M$	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (Ved in enough water to p ect any hydrolysis) (C) 1.50 x 10 ⁻¹ M $M_{2(aq)} + 2H_2O(I)$ mass of concentrated HC (C) 19.4 g 5 0.40M solution of Na ₃ PC (C) 0.024 n according to the balance n^{2+} (aq) + 10CO _{2(g)} + 8H ₂ C	(D) $C_{3}H_{9}N_{3}$. pundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M resented by the eq I solution (36.0% (D) 26.4 g D_{4} ? (D) 0.008 ced equation give $D_{(1)}$	(INSEC-2006) (INSEC-2006) solution .What is [INSEC-2007] M uation: HCl by mass) is [INSEC-2007] [INSEC-2007] en below :
26. 27. 28. 29. 30.	this compound is (A) CH_2N An element has three is respectively. The average (A) 25.25 17.1 grams of aluminum the molarity of the sulfat (A) 1.67 x 10^{-2} M Chlorine can be prepare MnO ₂ (s) + 4HC Assuming the reaction g needed to produce 2.50 (A) 5.15 g How many moles of Na ⁺ (A) 0.008 Oxalic acid (H ₂ C ₂ O ₄) rea $5H_2C_2O_{4(aq)} + 2N$ How many mL of 0.0154	(B) CH ₄ N otopes with masses 24, 2 ge mass of the isotope mix (B) 25.50 a sulfate Al ₂ (SO ₄) ₃ is disso e ion the solution? (Negle (B) 5.00 x 10 ⁻² M d by reacting HCl with Mr L _(aq) \longrightarrow Cl ₂ (g) + MnCl oes to completion, what r g of Cl ₂ (B) 14.3 g ions are there in 20mL of (B) 0.020 acts with permanganate ion MnO_4 (aq) + 6H ⁺ _(aq) \rightarrow 2Mi M KMnO ₄ solution are req	(C) CH ₅ N 25 and 26 with relative at ature would be (C) 24.50 (Ved in enough water to p ect any hydrolysis) (C) 1.50 x 10^{-1} M MO ₂ . The reaction is repre (2(aq) + 2H ₂ O(I) mass of concentrated HC (C) 19.4 g 5 0.40M solution of Na ₃ PO (C) 0.024 n according to the balance n ²⁺ (aq) + 10CO _{2(g)} + 8H ₂ O	(D) $C_3H_9N_3$. pundance of 80% (D) 24.25 repare 1.00 L of s (D) 2.50 x 10 ⁻¹ M resented by the eq I solution (36.0% (D) 26.4 g D_4^2 (D) 0.008 red equation give $D_{(0)}$ of 0.0208 M H ₂ C	$_{2}O_{4}$ solution ?

Mole concept

31.	What is the Na ⁺ ion cond mL of $0.30M Na_3PO_4$ so	centration in the solution f lution ?	ormed by mixing 20 mL c	of 0.10 M Na ₂ SO ₄	solution with 50 [NSEC-2008]
	(A) 0.15 M	(B) 0.24 M	(C) 0.48	(D) 0.70	
32.	A currency counting ma oxygen atoms in 24.8 g How many days would b (A) 9.33×10^{17}	chine counts 60 million n of Na ₂ S ₂ O ₃ .5H ₂ O (M.W. = be required to count these (B) 7.03 × 10^{10}	otes per day. A bank has = 248) = notes ? (C) 8.03 × 10 ¹⁵	an many notes a (D) 6.66 × 10 ⁻¹²	s number of [NSEC-2008]
33.	Which of the following e	quations represented an o	oxidation-reduction reaction	on?	[NSEC-2008]
	(A) $H_2SO_4 + 2NH_3$	\rightarrow (NH ₄) ₂ SO ₄			
	(B) $H_2SO_4 + Na_2CO_3$ —	\rightarrow Na ₂ SO ₄ + H ₂ O + CO	2		
	(C) $2K_2CrO_4 + H_2SO_4 -$	$\longrightarrow K_2 Cr_2 O_7 + K_2 SO_4 + H_2$	H ₂ O		
	(D) $2H_2SO_4 + Cu \longrightarrow$	$CuSO_4 + 2H_2O + SO_2$			
34.	Methyl-t-butyl ether, C_5 gas, O_2 are required to b	H_{12} O, is added to gasoline ourn 1.0 mol of this compo	to promote cleaner burni ound completely to form o	ng.How many me carbon dioxide ar	oles of oxygen nd water ?
	(A) 4.5 mol	(B) 6.0 mol	(C) 7.5 mol	(D) 8.0 mol	
35.	The hydrated salt Na_2SC	O ₄ .nH ₂ O loses all water of	f crystallization on heatin	g and is reduced	to 44.1% of its
	original weight. Therefor (A) 5	e, the value n is (B) 10	(C) 6	(D) 7	[NSEC-2008]
36.	The simplest formula of	a compound containing 50	0% of element 'A' (Atomic	weight = 10) and	50% of element
	'B' (Atomic weight = 20) (A) AB) is (B) A_B	(C) A B	(D) A B	[NSEC-2008]
37.	3.7 dm ³ of 1 M NaOH sc	(2) (2)	3 of 0.3 M NaOH solution	The molarity of	the resulting
•	solution is :		(O) 0 70 M	(D) 0 50 M	[NSEC-2009]
20		(B) 0.10 M	(C) 0.73 M	(D) 0.59 M	
38.	(A) 2.68×10^{18}	(B) 2.52×10^{21}	(C) 1.83×10^{24}	- (D) 2.4 × 10 ²¹	[NSEC-2009]
39.	The oxidation-reduction	reaction among the follow	wing is -		[NSEC-2009]
	$(A) H_2 SO_4 + 2NH_3 \longrightarrow$	• (NH ₄) ₂ SO ₄			
	(B) $H_2SO_4 + Na_2CO_3$ —	\rightarrow Na ₂ SO ₄ + H ₂ O + CO	2		
	$(C) 2K_2CrO_4 + H_2SO_4 - $	$\longrightarrow K_2 Cr_2 O_7 + K_2 SO_4 + H_2$	H ₂ O		
	(D) $2H_2SO_4 + Cu \longrightarrow$	$CuSO_4 + 2H_2O + SO_2$			
40.	Silver metal reacts with	nitric acid according to th	e equation		[NSEC-2009]
	3Ag (s) + 4HNC The volume of 1.15 M H (A) 4.74 mL	D_3 (aq) \longrightarrow 3AgNO ₃ (and NO	q) + NO (g) + 2H ₂ O (l) t with 0.784 g of silver is - (C) 8.43 mL	- (D) 25.3 mL	
41.	P_4 (s) + 3OH ⁻ (aq) + 3H ₂ In the above equation, the (A) P_4 and OH ⁻	$_{2}^{0}$ (I) \longrightarrow PH ₃ (g) + 3H he species getting oxidize (B) OH ⁻ and P ₄	$H_2PO_2^{-}$ (aq) ed and reduced respective (C) P_4 and H_2O	ely are : (D) P_4 and P_4	[NSEC-2009]
42.	A Compound Containing H = 8.1 %. One mole of molecular formula of the (A) $BeN_4H_{12}Cl_2$	ng beryllium has the fo the compound has mass compound is : (B) $BeN_2H_{10}CI$	Ilowing composition, Be of 148g and average ato (C) BeN ₄ H ₂ CI ₃	e = 6.1%, N = 3 mic mass of bery (D) Be ₂ N ₄ H ₁₀ Cl ₂	37.8% Cl=48%, /llium is 9. The [NSEC-2010]

Mole concept

43.	The molarity of 20% w/v (A) 2.32	v sulphuric acid of density (B) 2.02	y 1.14 g cm⁻³ is (C) 2.12	(D) 2.22	[NSEC-2010]
44.	An inorganic bromide im mL of 0.20 M AgNO $_3$ to α	purity in a sample is preci completely precipitate the	pitated as silver bromide. e impurity. The mass perc	2.00 g of the sam centage of the im	nple required 6.4 purity is INSEC-20101
	(A) 5.11	(B) 2.56	(C) 9.15	(D) 1.28	[
45.	The compound which c (A) HNO_2	an act as an oxidizing ag (B) HI	ent as well as reducing a (C) HCN	gent is (D) HCOOH	[NSEC-2010]
46.	Maximum number of mo mole of sodium phospha (A) 0.2	bles of barium phosphate ate is (B) 0 4	formed when 0.9 mole of	barium chloride i	is mixed with 0.4 [NSEC-2010]
47.	The largest number of m (A) 70g of Sulphur dioxid (C) 36g of Water	nolecules are present in de	(B) 64g of Nitrogen pent(D) 34g of Carbon dioxid	oxide le	[NSEC-2010]
48.	The number of water m	olecules present in 0.2	0 g sample of $CuSO_4$. 5	H ₂ O (Molar mas	ss = 249.7) is
	(A) 1.2 × 10 ²¹	(B) 2.14 × 10 ²¹	(C) 2.14 × 10 ²²	(D) 1.2 × 10 ²³	[NSEC-2011]
49.	The compound in which (A) KMnO ₄ (C) MnSO ₄	Mn has oxidation state c	of +3 is (B) K ₂ [Mn(CN) ₆] (D) CsMn(SO ₄) ₂ ,12H ₂ O		[NSEC-2011]
50.	Excess of silver nitrate i sample. 1.4 g of silver c Molar masses (g-mol ⁻¹) (A) 0.25 g	s added to a water sampl hloride is precipitated. Th : AgNO ₃ 169.91, AgCl (B) 0.35 g	e to determine the amou ne mass of chloride ion pr 143.25 (C) 0.50 g	nt of chloride ion resent in the sam (D) 0.75 g	present in the ple is : [NSEC-2011]
51.	The vapour density of ga is : (A) M	as A is four times that of B (B) 4M	. If the molecular mass of (C) M/4	^E B is M then mole (D) 2M	ecular mass of A [NSEC-2012]
52.	In the reaction, $2KCIO_3$ - N.T.P. will be : (A) 9.74 dm ³	→ 2KCl + 3O ₂ when 36.75 (B) 8.92 dm ³	5 g of KCIO ₃ is heated, the (C) 10.08 dm ³	e volume of oxyg (D) 22.4 dm ³	en evolved at [NSEC-2012]
53.	A 500 g toothpaste samp	le has 0.4 g fluoride conc	entration. The fluoride cor	ncentration in terr	ns of ppm will be
	: (A) 200	(B) 400	(C) 500	(D) 800	[NSEC-2012]
54.	In a nitration experimen	t, 10.0g of benzene gas a	nd 13.2 g of nitrobenzene	e. The percentag	e yield is :
	(A) 83.5%	(B) 62.7%	(C) 88.9%	(D) 26.7%	[NSEC-2012]
55.	In the compound Na_2S_2 (A) -2	D_{3} , the oxidation state of (B) +2	sulphur is (C) +4	(D) +6	[NSEC-2013]
56.	I. $5H_2O_2 + 2MnO_4^- + 6H^+$	$\rightarrow 2\mathrm{Mn}^{2+} + 5\mathrm{O}_2 + 8\mathrm{H}_2\mathrm{O}$			[NSEC-2014]
	II. $H_2O_2 + Ag_2O \rightarrow 2Ag$ The role of hydrogen per (A) oxidising in I and red (C) reducing in I as well	$+H_2O+O_2$ roxide in the above reaction lucing in II as in II	on is (B) reducing in I and oxi (D) oxidising in I as well	dising in II as in II	
57.	Aluminum carbide (AI_4C_3) to produce 11.2 L of met (A) 48) liberates methane on tre hane under STP conditio (B) 72	atment with water. The gra ons is : [Given : Al = 27] (C) 144	ams of aluminum (D) 24	carbide required [NSEC-2014]

JEE	(Adv.)-Ch	emistry			Mole	concept			
58.	The specific	entration of HI	NO ₃ is : SEC-20141						
	(A) 15.8	(B) 31.6	(C) 11.1	(D) 14.2	[100				
		PART -	4 : ADDITIONAL	PROBLEMS					
		SL	JBJECTIVE QUE	STIONS					
1.	Carbon disu 5C How much (Iphide, CS_2 , can be r + $2SO_2 \longrightarrow CS_2 \cdot CS_2 \cdot CS_2$ can be produced	made from by-product SO + 4CO I from 440 kg of waste SO	$_{2}$. The overall reaction is $_{2}$ with 60 kg of coke if the	s $e SO_2$ convers	sion is 80%?			
2.2	The percent	t yield for the followin	ng reaction carried out in c	arbon tetrachloride (CC	I_4) solution is 8	30%			
	Br ₂ (a) What am (b) What am	$Br_2 + Cl_2 \longrightarrow 2BrCl$ (a) What amount of BrCl would be formed from the reaction of 0.025 mol Br_2 and 0.025 mol Cl_2 ? (b) What amount of Br_2 is left unchanged?							
3. 🗷	Write balan (a) $S_4O_6^{2-}$ (a (b) $S_2O_3^{2-}$ (ac (c) CIO_3^{-} (ac (d) IO_3^{-} (aq) (e) HSO_4^{-} (a (f) HNO_2 (aq	ced net ionic equation q) + AI(s) \longrightarrow H ₂ S q) + Cr ₂ O ₇ ²⁻ (aq) \longrightarrow q) + As ₂ S ₃ (s) \longrightarrow C + Re(s) \longrightarrow ReO aq) + As ₄ (s) + Pb ₃ O ₄ () \longrightarrow NO ₃ ⁻ + NO(ons for the following reacti $S(aq) + Al^{3+}(aq)$ → $S_4O_6^{2-}(aq) + Cr^{3+}(aq)$ $Cl^-(aq) + H_2AsO_4^-(aq) + H_4^-(aq) + I^-(aq)$ $(aq) + I^-(aq)$ $(aq) + I^-(aq)$ $(aq) + H_2AsO_4(a) + H_2A$ $(aq) + H_2AsO_4(a) + H_2A$	ons in acidic solution : HSO_4^- (aq) sO_4^- (aq)					
4.	Write balanced net ionic equations for the following reactions in basic solution : (a) $C_4H_4O_6^{2-}(aq) + CIO_3^{-}(aq) \longrightarrow CO_3^{2-}(aq) + CI^{-}(aq)$ (b) $AI(s) + BiONO_3(s) \longrightarrow Bi(s) + NH_3(aq) + [AI(OH)_4]^{-}(aq)$ (c) $H_2O_2(aq) + CI_2O_7(aq) \longrightarrow CIO_2^{-}(aq) + O_2(g)$ (d) $TI_2O_3(s) + NH_2OH(aq) \longrightarrow TIOH(s) + N_2(g)$ (e) $[Cu(NH_3)_4]^{2+}(aq) + S_2O_4^{2-}(aq) \longrightarrow SO_3^{2-}(aq) + Cu(s) + NH_3(aq)$ (f) $Mn(OH)_5(s) + MnO_5^{-}(aq) \longrightarrow MnO_5(s)$								
5.	Consider the XeF Determine t	e following reaction ir $F_6 + (-CH_2 - CH_2)_{h}$ - he moles of Xe F_6 req	hvolved in the preparation $\longrightarrow (-CF_2-CF_2)_n + HI$ juired for preparation of 10	of teflon polymer	² ₂−CF₂) _n .				
		ONLY O	NE OPTION CO	RRECT TYPE					
1.	In a certain obtained [A (A) 35 38 %	operation 358 g of Ti t. wt. Ti = 48, Mg = 2	iCl_4 is reacted with 96 g of 24] (B) 66.6 %	Mg. Calculate % yield c	of Ti if 32 g of ⊺ (D) 60 %	Fi is actually			

(B) 66.6 % (A) 35.38 % (C) 100 % Phosphoric acid $(H_{3}PO_{4})$ prepared in a two step process. 2. $\begin{array}{c} \mathsf{P}_4 + 5\mathsf{O}_2 \xrightarrow{\mathsf{G}} \mathsf{P}_4\mathsf{O}_{10} \\ \mathsf{P}_4\mathsf{O}_{10} + 6\mathsf{H}_2\mathsf{O} \xrightarrow{\mathsf{G}} \mathsf{4H}_3\mathsf{PO}_4. \end{array}$ (A) (B)

We allow 62 g of phosphorus to react with excess oxygen which form P_4O_{10} in 85% yield. In the step (B) reaction 90% yield of H_3PO_4 is obtained. Produced mass of H_3PO_4 is : (A) 37.48 g (B) 149.94 g (C) 125.47 g (D) 564.48 g

(D) 60 %

3.	For the the corr (A) (B) (C) (D)	redox reaction, rect coefficients of MnO ₄ ⁻ 2 16 5 2	$MnO_{4}^{-} + C_{2}O_{4}^{2-} + C_{2}O_{4}^{2-}$ of the reactions for $C_{2}O_{4}^{2-}$ 5 5 16 16	+ H ⁺ → N or the balance H ⁺ 16 2 2 5	$n^{2+} + CO_2 + H_2O$ d reaction are			
4.	A miner ammor (A) 5.8	ral water sample nia gas in one litro ×10 ⁻⁴ mol	was analysed an e water bottle is (B) 1 × 10 ⁻² mo	d found to con (d _{water} ≈ 1 g/m I (C)	tain 1 × 10⁻³ % am ll) 0.58 ×10⁻² mol	monia (w/v (D) san	w). The mole of dissolve ne as w/w	ю
	(A) 5.8	×10 ⁻⁴ mol	(B) 1 × 10 ⁻² mo	l (C)	0.58 ×10⁻² mol	(D) Hkj@	@Hkkj ds1 eku	
5.	(i) 2AI + (ii) AICI (iii) AI(C Above : g of Na amount each re (A) (B)	- 6HCl $_{3}$ + 3NaOH DH) ₃ + NaOH series of reaction OH is added for s t of NaAlO ₂ that of haction goes to 10 L.R. in step (I) Al Al		+ $3H_2$ $_3$ + $3NaCl$ $_2$ + $2H_2O$ starting with 1. Find out limiti in step (iii). (A step (II)	B g of Al and 109.5 ng reagent in each ssume reactions a L.R. in step (II Al(OH) ₃ Al(OH) ₂	g of HCI in a step and are taken ir I)	first step and further 10 calculate the maximum sequence and also tha Moles of NaAlO ₂ 0.66 0.5)0 า at
	(C)	Al	AICI	,	NaOH		0.5	
6.	(D) 12 g of (A) 12	alkaline earth m	etal gives 14.8 g (B) 20	of its nitride. (C)	Atomic weight of n	netal is - (D) 14.8	8	
7.2	When 1	100g of ethylene	polymerises enti	irely to polyeth	nene, the weight of	polyether	ne formed as per the	
	equatio (A) (n/2	on $n(C_2H_4) \rightarrow (-C_2)$:H ₂ –CH ₂ –) _n is : (B) 100g	(C)	(100/n)g	(D) 100	ng	
8.24	Calcula	ate the amount of Ni + 4CO sed in this proce (Ni = 59 u)	Ni needed in the Ni(CO)₄ ess is obtained th	e Mond's proce	ess given below ess, in which 6 g	of carbon	is mixed with 44 g CO	2
	(A) 14.6	675 g	(B) 29 g	(C)	58 g	(D) 28 g	g	
9.2	The ma (A) 49 g	ass of 70% H ₂ SO	₄ required for neu (B) 98 g	utralisation of (C)	l mol of NaOH. 70 g	(D) 34.3	3 g	
10.	0.05 m product H = 1, 0	ole of LiAIH ₄ in eft t LiAIHC ₁₂ H ₂₇ O ₃ v C = 12, O = 16).	ther solution was veighed 12.7 g. l	placed in a fl If Li atoms are	ask containing 74g conserved, the p	g (1 mole) o ercentage	of t-butyl alcohol. The yield is : (Li = 7, Al = 27	7,
	(A) 25%	0	(B) 75%	(C)	100%	(D) 15%	6	
11.	What w P, and	reights of P_4O_6 ar	P_4O_{10} will be p	roduced by th	e combustion of 31	$ g of P_4 in 3$	32g of oxygen leaving n	10
	(Å) 2.7	5 g, 219.5 g	(B) 27.5 g, 35.5	5 g (C)	55 g, 71 g	(D) 17.	5 g, 190.5 g	
12.	10 g of CO_3 is h in the o	a sample of a mi neated to convert riginal mixture is	xture of $CaCl_2$ and all the Ca to CaCl_2.	nd NaCl is trea D and the final	ated to precipitate mass of CaO is 1.6	all the cald 62 g . The p	cium as CaCO ₃ . This C bercent by mass of CaC	a ¦l₂
	(A) 32.1	1 %	181162%	((`)	Z1 X %	(1)) 11 (1 %	
4.0					21:0 /0 0 ·	(2) 11.		

14.2	If a piece of iron gains 10% of its weight due to partial rusting into Fe_2O_3 the percentage of total iron that has rusted is :								
	(A) 23	(B) 13	(C) 23.3	(D) 25.67					
15.๖	When x grams o Which of the foll	When x grams of carbon are heated with y grams of oxygen in a closed vessel, no solid residue is left behind. Which of the following statements is correct ?							
	(A) y/x must lie l	(A) y/x must lie between 1.33 and 2.67 (B) y/x must be greater than or equal 2.67.							
	(C) y/x must be	less than or equal 1.33	(D) y/x must	be greater than or equal 1.33.					
16.	On mixing 15.0 solution is found	ml of ethyl alcohol of density to have a density of 0.924 g	v 0.792 g ml ⁻¹ with ml ⁻¹ . The percent	15 ml of pure water at 4°C, the tage contraction in volume is :	eresulting				

- (A) 8 % (B) 2 %
 - MATCH THE COLUMN

(C) 3%

1. Column-I (A) Molarity

(B) Molality

(C) Mole fraction

(D) Mass %

Where M_A , M_B are molar masses, n_A , n_B are no of moles & X_A , X_B are mole fractions of solute and solvent respectively.

2.2 Column-I

(A) 100 ml of 0.2 M AICl₃ solution + 400 ml of 0.1 M HCl solution(p) Total concentration of cation(s) = 0.12 M (B) 50 ml of 0.4 M KCl + 50 ml H₂O (q) $[SO_4^{2-}] = 0.06 \text{ M}$ (C) 30 ml of 0.2 M K₂SO₄ + 70 ml H₂O (r) $[SO_{4}^{2-}] = 2.5 \text{ M}$ (D) 200 ml 24.5% (w/v) H₂SO₄ (s) $[CI^{-}] = 0.2 \text{ M}$

SINGLE AND DOUBLE VALUE INTEGER TYPE

- 1. The measured density at 0°C and 1 atm of He is 0.1784 g/L. What is the weight (in g) of one mole of He ?
- 2. The 'roasting' of 100.0 g of a copper ore yielded 71.8 g pure copper. If the ore is composed of Cu₂S and CuS with 4.5 % inert impurity, calculate the percent of Cu_2S in the ore. $Cu_2S + O_2 \longrightarrow 2Cu + SO_2^2$ and $CuS + O_2 \longrightarrow Cu + SO_2$ The reactions are :
- A piece of Al weighing 27 g is reacted with 200 ml of H₂SO₄ (specific gravity = 1.8 and 54.5 % by weight) After 3. the metal is completely dissolved 73 g HCl is added and solution is further diluted to 500 ml solution then find the concentration of H⁺ ion in mol/litre.
- 4. 1 g of dry green algae absrobs 4.7×10^{-3} mole of CO, per hour by photosynthesis. If the fixed carbon atoms were all stored after photosynthesis as starch (C₆H₁₀O₅)_n. Aproximately how long (in hour) would it take for the algae to double their own weight assuming phtosynthesis takes place at a constant rate?

Column-II

(s) $\frac{X_A}{X_B M_B} \times 1000$

Column-II

(D) 4 %

(p) Dependent on temperature

(q) $\frac{M_A \times n_A}{n_A M_A + n_B M_B} \quad x \ 100$

(r) Independent of temperature

5. CN^- ion is oxidised by a powerful oxidising agent to NO_3^- and CO_2 or CO_3^{2-} depending on the acidity of the reaction mixture.

 $CN^{-} \longrightarrow CO_2 + NO_3^{-} + H^+ + ne^-$

What is the number (n) of electrons per mole of $CN^{\scriptscriptstyle -}$ involved in the process ?

- 6. To 100 ml of 5 M NaOH solution (density 1.2 g/ml) were added 200 ml of another NaOH solution which has a density of 1.5 g/ml and contains 20 mass percent of NaOH. What will be the volume of the gas (at 0°C and 1 atm) in litres liberated when aluminium reacts with this (final) solution. The reaction is AI + NaOH + $H_2O \longrightarrow NaAIO_2 + H_2$
- 7. A drop (0.05 mL) of 12 M HCl is spread over a thin sheet of aluminium foild (thickness 1 mm and density of AI = 2.7 g/mL). Assuming whole of the HCl is used to dissolve. At what will be the maximum area of hole produced in foild (in cm²). [Report your answer after multiplying by 10].
- 8. When 1 mole of A reacts with $\frac{1}{2}$ mole of B₂ (A + $\frac{1}{2}$ B₂ \rightarrow AB), 100 Kcal heat is liberated and when 1 mole of

A reacted with 2 mole of $B_2 (A + 2B_2 \rightarrow AB_4)$, 200 Kcal heat is liberated. When 1 mole of A is completely reacted with excess, of B_2 to form AB as well as AB_4 , 140 Kcal heat is liberated calculate the mole of B_2 used. [Write your answer as number of mole of B_2 used × 10]

- 9. In the reaction : $2AI + Cr_2O_3 \longrightarrow Al_2O_3 + 2Cr$, 49.8 g of AI reacted with 200.0 g Cr_2O_3 . How much grams of reactant remains at the completion of the reaction ?
- 10. The following reaction

 $xZn + yHNO_3(dil) \longrightarrow aZn(NO_3)_2 + bH_2O + cNH_4NO_3$

What is the sum of the coefficients (a + b + c)?

- **11.** H_3PO_4 (98 g mol⁻¹) is 98% by mass of solution. If the density is 1.8 g/ml, calculate the molarity.
- 12. 3.0 litre of water are added to 2.0 litre of 5 M HCI. What is the molarity of HCI (in M) the resultant solution ?

ONE OR MORE THAN ONE OPTIONS CORRECT TYPE

- 1. In the reaction $I_2 + C_2H_5OH + OH^- \longrightarrow CHI_3 + HCOO^- + H_2O + I^-$ which of the following statements is/are correct ?
 - (A) The coefficients of OH^- and I^- in the given in balanced equation are, respectively, 6 and 5.
 - (B) The coefficients of OH⁻ and I⁻ in the given balanced equation are , respectively, 5 and 6.
 - (C) C_2H_2OH is oxidised to CHI₂ and HCOO⁻.
 - (D) The number of electrons in the conversion of C_2H_5OH to CHI_3 and $HCOO^-$ is 8.
- 2. One mole of a mixture of N_2 , NO_2 and N_2O_4 has a mean molar mass of 55.4. On heating to a temperature at which all the N_2O_4 may be presumed to have dissociated : $N_2O_4 \implies 2NO_2$, the mean molar mass tends to the lower value of 39.6. What is the mole ratio of N_2 : NO_2 : N_2O_4 in the original mixture ? (A) 0.5 : 0.1 : 0.4 (B) 0.6 : 0.1 : 0.3 (C) 0.5 : 0.2 : 0.3 (D) 0.6 : 0.2 : 0.2

3. Silver metal in ore is dissolved by potassium cyanide solution in the presence of air by the reaction

 $4 \text{ Ag} + 8 \text{ KCN} + \text{O}_2 + 2\text{H}_2\text{O} \longrightarrow 4 \text{ K}[\text{Ag} (\text{CN})_2] + 4 \text{ KOH}$

- (A) The amount of KCN required to dissolve 100 g of pure Ag is 120 g.
- (B) The amount of oxygen used in this process is 0.742 g (for 100 g pure Ag)
- (C) The amount of oxygen used in this process is 7.40 g (for 100 g pure Ag)
- (D) The volume of oxygen used at 0° C and 1 atm is 5.20 litres.
- 4. Crude calcium carbide, CaC_2 , is made in an electric furnace by the following reaction,

 $CaO + 3C \longrightarrow CaC_2 + CO$

The product contain 85% CaC_2 and 15% unreacted CaO.

- (A) 1051.47 kg of CaO is to be added to the furnace charge for each 1000 kg of CaC₂.
- (B) 893.8 kg of CaO is to be added to the furnace charge for each 1000 kg of crude product.
- (C) 708.2 kg of CaO is to be added to the furnace charge for each 1000 kg of CaC_2 .
- (D) 910.3 kg of CaO is to be added to the furnace charge for each 1000 kg of crude product.
- 5. Which of the following statement is/are correct ?

Excess of $H_2S(g)$ is bubbled into 1.0 L of 0.1 M CuCl₂ solution.

 $Cu^{2+} + H_2S(g) \longrightarrow CuS(s) + 2H^+$

(A) 9.55 g of CuS is produced.

- (B) The concentration of H⁺ ions is 0.2 M
- (C) The concentration of $\rm H^{+}$ ions is 0.1 M. (D) 95.5 g CuS is produced.
- 6. Which of the following statements is/are correct ? 1.0 g mixture of $CaCO_3(s)$ and glass beads liberate 0.22 g of CO_2 upon treatment with excess of HCI. Glass does not react with HCI.

 $CaCO_3 + 2HCI \longrightarrow CO_2 + H_2O + CaCl_2$

- [M.wt. of $CaCO_3 = 100$, M.wt. of $CO_2 = 44$, [Atomic weight of Ca = 40]
- (A) The weight of CaCO, in the original mixture is 0.5 g
- (B) The weight of calcium in the original mixture is 0.2 g
- (C) The weight percent of calcium in the original mixture is 40% Ca.
- (D) The weight percent of Ca in the original mixture is 20% Ca.
- **7.** 100 g sample of clay (containing 19% H₂O, 40% silica, and inert impurities as rest) is partially dried so as to contain 10% H₂O
 - Which of the following is/are correct statements (s)?
 - (A) The percentage of silica in paritially dried clay is 44.4%
 - (B) The mass of paritially dried clay is 90.0 g.
 - (C) The percentage of inert impurity in paritially dried clay is 45.6%
 - (D) The mass of water evaporated is 10.0 g

-				R	RP /	ANS	SWE	ER H	KEY				
	PART - 1												
1.	(2)	2.	(3)	3.	(3)	4.	(1)	5.	(2)	6.	(1)	7.	(4)
8.	(1)	9.	(3)	10.	(3)	11.	(3)	12.	(4)	13.	(1)	14.	(3)
15.	(1)	16.	(2)	17.	(4)	18.	(4)	19.	(2)	20.	(3)		
21.	328.57	22.	32.1	23.	739.2	24.	345	25.	0.48				
						ΡΑ	RT - 2						
1.	(B)	2.	(B)	3.	(A)	4.	(B)	5.	(BD)	6.	(AB)		
7.	(ABC)	8.	(BC)	9.	(AC)	10.	(AC)	11.	(ABC)	12.	(ACD)	
13.	1	14.	5	15.	1	16.	7	17.	2	18.	42		
						ΡΑ	RT - 3						
1.	(B)	2.	(B)	3.	(A)	4.	(B)	5.	(D)	6.	(A)	7.	(A)
8.	(D)	9.	(B)	10.	(B)	11.	(A)	12.	(B)	13.	(C)	14.	(B)
15.	(B)	16.	(D)	17.	(C)	18.	(B)	19.	(D)	20.	(A)	21.	(B)
22.	(C)	23.	(B)	24.	(D)	25.	(B)	26.	(D)	27.	(C)	28.	(B)
29.	(C)	30.	(A)	31.	(D)	32.	(C)	33.	(D)	34.	(C)	35.	(B)
36.	(B)	37.	(D)	38.	(D)	39.	(D)	40.	(C)	41.	(D)	42.	(A)
43.	(A)	44.	(A)	45.	(A)	46.	(A)	47.	(C)	48.	(B)	49.	(D)
50.	(B)	51.	(B)	52.	(C)	53.	(D)	54.	(A)	55.	(B)	56.	(C)
57.	(D)	58.	(A)										
						ΡΑ	RT - 4						
1.	76 kg c	of CS ₂											
2.	(a)				$Br_2 + C$	Cl ₂ —	$\rightarrow 2BrCl$						
					1 mol	1mol	2mol						
					0.025	0.025	0.050	mol					
	(b) The	oretica	al yield of		BrCl = 0.050 mol								
		actua	al yield is	$=\frac{0.05}{10}$	$\frac{\times 80}{0} = 0.0$)4 mole	9						
		Br ₂ le	ft unreac	ted = 0	.025 – 0.0	02 = 0.0	05 mol.						
3. (a) $S_4O_6^{2-}(aq) + 6 AI(s) + 20 H^+ \longrightarrow 4H_2S(aq) + 6AI^{3+}(aq) + 6H_2O$ (b) $6S_2O_3^{2-}(aq) + Cr_2O_7^{2-}(aq) + 14 H^+ \longrightarrow 3S_4O_6^{2-}(aq) + 2Cr^{3+}(aq) + 7H_2O$ (c) $14CIO_3^{-}(aq) + 3As_2S_3(s) + 18H_2O \longrightarrow 14CI^{-}(aq) + 6H_2AsO_4^{-}(aq) + 9HSO_4^{-}(aq) + 15H^+$ (d) $7IO_3^{-}(aq) + 6Re(s) + 3H_2O \longrightarrow 6ReO_4^{-}(aq) + 7I^{-}(aq) + 6H^+$ (e) $26H^+ + 30HSO_4^{-}(aq) + As_4(s) + 10 Pb_3O_4(s) \longrightarrow 30 PbSO_4(s) + 4H_2AsO_4^{-}(aq) + 24H_2O$ (f) $3HNO_2(aq) \longrightarrow NO_3^{-} + 2NO(g) + H_2O + H^+$

- 4. (a) $3C_4H_4O_6^{-2}(aq) + 5CIO_3^{-}(aq) + 18OH^{-} \longrightarrow 12CO_3^{-2}(aq) + 5CI^{-}(aq) + 15H_2O$ (b) $11AI(s) + 3BiONO_3(s) + 21H_2O + 11OH^{-} \longrightarrow 3Bi(s) + 3NH_3(aq) + 11AI(OH)_4^{-}(aq)$ (c) $4H_2O_2(aq) + CI_2O_7(aq) + 2OH^{-} \longrightarrow 2CIO_2^{-}(aq) + 4O_2(g) + 5H_2O$ (d) $TI_2O_3(s) + 4NH_2OH(aq) \longrightarrow 2TIOH(s) + 2N_2(g) + 5H_2O$ (e) $[Cu(NH_3)_4]^{2+}(aq) + S_2O_4^{-2-}(aq) + 4OH^{-} \longrightarrow 2SO_3^{-2-}(aq) + Cu(s) + 4NH_3(aq) + 2H_2O$ (f) $3Mn(OH)_2(s) + 2MnO_4^{-}(aq) \longrightarrow 5MnO_2(s) + 2H_2O + 2OH^{-}$
- **5.** 4

ONLY ONE OPTION CORRECT TYPE

1.	(A)	2.	(B)	3.	(A)	4.	(A)	5.	(C)	6.	(C)	7.	(B)
8.	(A)	9.	(C)	10.	(C)	11.	(B)	12.	(A)	13.	(A)	14.	(C)
15.	(D)	16.	(C)										

PART - III

1.	(A - p); (B - r,s); (C - r); (D - r,q)
----	--

2. (A - p,s); (B - s); (C - p,q); (D - r)

SINGLE AND DOUBLE VALUE INTEGER TYPE

1.	4	2.	62%	3.	6	4.	8	5.	10	6.	67	7.	2
8.	11	9.	60	10.	8	11.	18	12.	2				

ONE OR MORE THAN ONE OPTIONS CORRECT TYPE

1.	(ACD) 2.	(A) 3.	(ACD)	4.	(AB)	5.	(AB)	6.	(ABD)
7.	(ABCD)									

Reliable Ranker Problems

PART-1

1.	Mole of NO ₂ = $\frac{112}{22400}$ = 5 × 10 ⁻³
	Mass of NO ₂ = 5 × 10 ⁻³ × 46 = 0.23 gm
	Volume of NO ₂ = $\frac{\text{Mass}}{\text{Density}} = \frac{0.23}{1.15} = 0.2 \text{ ml}$
2.	Number of molecule = $5 \times 10^{-3} \times 6.023 \times 10^{23} = 3.1 \times 10^{21}$. Mass of NaCl = $10 \times 0.96 = 9.6$ g
	moles of NaCl = $\frac{9.6}{58.5}$
	no. of molecules = $\frac{9.6}{58.5} \times 6.023 \times 10^{23} \simeq 10^{23}$
3.	$\frac{32}{2x+3y} = 0.2$
	$\frac{92.8}{3x+4y} = 0.4$
	Hence $x = 56 \& y = 16$.
4.	(1) It is a fact. (2) $2H_2 + O_2 \longrightarrow 2H_2O$ Initial mole 2 3 0 final mole 0 $3-1=2$ 2 (3) $C + O_2 \longrightarrow CO_2$
	<u>w</u> <u>w</u>
	12 32
	Here C is limiting reagent.
	Reduction
5.	$BiO_{3}^{-} + Mn^{2+} \longrightarrow Bi^{3+} + MnO_{4}^{-}$
	(i) 2a + 6H + BiO = 2Bi3 + 2H O
	(i) $2e + 6n^{2} + 6iO_{3} \longrightarrow Bi^{0} + 3n_{2}O$ (ii) $4H_{2}O + Mn^{2+} \longrightarrow MnO_{4}^{-} + 8H^{+} + 5e$
	(i) × 5 + (ii) × 2, we get 14 H ⁺ + 5 BiO ₃ ⁻ + 5Mn ²⁺ \longrightarrow 5E

(i) x 5 + (ii) x 2, we get 14 H⁺ + 5 BiO₃⁻ + 5Mn²⁺ \longrightarrow 5Bi³⁺ + 2MnO₄⁻ + 7 H₂O Hence, (2) is the correct balanced reaction.

6. $3I_2 + OH^- \longrightarrow IO_3^- + 5I^-$ (balance reaction) So, ratio is 1 : 5.

7. Explanation : M. wt. of NaNO₃ = 85 70 mg of Na⁺ are present in 1 mL 50 ml of solution contains 50 x 70 = 3500 mg = 3.5 g Na⁺ ion 23 g of Na⁺ are present in 85 g of NaNO₃

3.5 g of Na⁺ are present in
$$\frac{85}{23} \times 3.5 = 12.934$$
 g of NaNO₃

8. Explanation :
$$m = \frac{M \times 1000}{(1000 \times d - M \times M.Wt.)}$$
 where 'm' is molality, M is molarity.

$$= \frac{10^{-2} \times 1000}{(1000 \times 1.1 - 10^{-2} \times 106)}$$
$$= \frac{10}{1100 - 1.6} = \frac{10}{1099.4} = 9.00 \times 10^{-3}$$
 [Take 1099.4 = 1100]

- 9. At 4°C i.e. 277 K density of water = 1 g/ml \therefore 1 kg water \Rightarrow 1000 ml water = 1 lit.
 - ... Molality & molarity remains same.

10. Molarity =
$$\frac{(\% w/w) \times density \times 10}{Molar mass of solute} = \frac{98 \times 1.84 \times 10}{98} = 18.4 \text{ M}$$

11. Mole of NaCl =
$$\frac{5.85}{58.5} = 0.1$$

Molarity =
$$\frac{0.1}{1}$$
 = 0.1 M

Moles in 1 ml of solution = MV = $0.1 \times 10^{-3} = 10^{-4}$ mole. Number of ions in 1 ml = $2 \times 10^{-4} \times 6.023 \times 10^{23} = 1.204 \times 10^{20}$.

12.	Element	Percent	r.a.m.	No. of atoms	atomic ratio
	С	74	12	74/12 = 6.16	6.16/1.23 = 5
	Н	8.7	1	8.7/1 = 8.7	8.7/1.123 = 7
	Ν	17.3	14	17.3/14 = 1.23	1.23/1.23 = 1

The ratio of atoms = C : H : N = 5 : 7 : 1 Empirical formula = C_5H_7N Empirical formula mass = 5 C + 7H + N = 5 x 12 + 7 x 1 + 14 = 81 Molecular mass = 162 (given)

No. of empirical units per molecule = $n = \frac{Molecularmass}{Empirical formula mass} = \frac{162}{81} = 2$ Molecular formula = (Empirical formula) × 2 = (C₅ H₇N) × 2 = C₁₀H₁₄N₂

13. C + O₂
$$\xrightarrow{\Delta}$$
 CO₂
12g C = 1 mol O₂ = 22.7 L O₂
∴ 1000 g C = $\frac{22.7}{12} \times 1000$ or 1891.67 L O₂

Mole concept

14.		6Li +	$N_2 \longrightarrow$		2Li₃N			
	Initial mole	$\frac{3.5}{7} = \frac{1}{2}$	$\frac{8.4}{28} = 0.3$		0			
	final mole	0	$0.3 - \frac{1}{12}$		$\frac{1}{2} \times \frac{1}{3}$	$=\frac{1}{6}$		
	mass of Li ₃ N	$=\frac{1}{6}\times 35=5.8$	3 g.					
15.	Initial mole final mole	$2KO_2 + 0.15$ (0.15 - 0.1) ∴ moles of $O_2 =$	$2H_{2}O \longrightarrow 0.1$ 0 $= 0.05$	2KOH 0 0.1	+	H ₂ O ₂ 0 0.05	+	O ₂ 0 0.05
16.	KCIO ₃ → KCI + Applying POAC moles of O in K 3 × mo 3 × $\frac{1}{mo}$	O_2 C for O atoms in the CIO ₃ = moles of les of KCIO ₃ = 2 <u>vt.of KCIO₃</u> = 2	the eqn.(i), O in O ₂ × moles of O ₂ 2 × $\frac{\text{volume at N}}{2240}$	<u>TP(mL)</u> 0				
	Wt . of	$\text{KCIO}_{3} = \frac{2 \times 146}{3 \times 10^{-3}}$	$\frac{3.8 \times 122.5}{22400} = 0.53$	58 g.				
	In the second reaction : The amount of KCIO ₃ left = $1 - 0.5358 = 0.4642$ g. We have, $KCIO_3 \rightarrow KCIO_4 + KCI$ 0.4642 g. Applying POAC for O atoms, moles of Q in KCIQ = moles of KCIQ							
	3 × mo	les of KCIO ₃ = 4 wt.of KCIO ₃ = 4	× moles of KCIO 4 ×	4				
	Wt. of $KCIO_4 =$ Wt. of residue	$\frac{3 \times 0.4642 \times 138.5}{122.5 \times 4}$ = 1 - wt. of Oxyg = 1 - $\frac{146.8}{24400}$ ×	⁵ mol. wt.of KCl ¹ ⁵ = 0.3937 g. gen 32 g = 0.7902 g.	04		(ii)		
	\therefore % of KCIO ₄ i	in the residue $=$	$\frac{0.3937}{0.7902} \times 100 = 4$	49.8 %.				
17.	Molarity of H_2S^{0}	$O_4 = \frac{\text{sp.gravity}}{\text{Molecut}}$	< % w/w×10 Ilar mass					
		$=\frac{1.2\times25\times10}{98}$	$\frac{10}{98} = \frac{12 \times 25}{98} = 3$.06 M				
	3H ₂ SO ₄ + 2AI -	$\longrightarrow Al_2(SO_4)_3 +$	3H ₂					
	$\frac{2.7}{27} = 0$	D.1						

Mole of H_2SO_4 used = $\frac{3}{2} \times 0.1 = 0.15$ Initial mole of $H_2SO_4 = 0.75 \times 3.06 = 0.2295$ Mole of H_2SO_4 remaining = 0.2295 - 0.15 Molarity of final $H_2SO_4 = \frac{0.0795}{0.4} = 0.198 \text{ M}.$ 18. Moles of $Al_2(SO_4)_3 = M \times V = 0.15 \times 0.1 = 0.015$ Mass of $Al_2(SO_4)_3 = Mole \times Molar mass = 0.015 \times 342 = 5.13 \text{ g}.$ Moles of $AI_{3^+} = 2 \times moles$ of $AI_2(SO_4)_3 = 2 \times 0.015 = 0.03$. No. of Al³⁺ ions = $0.03 \times 6.023 \times 10^{23} = 1.81 \times 10^{22}$ ions. $CuSO_{4}.5H_{2}O(aq) + BaCl_{2}(aq) \longrightarrow BaSO_{4}(s) + CaCl_{2}(aq)$ 19. 4.66 g Mass of $BaSO_4 = 4.66 g$ Mole of $BaSO_4 = \frac{4.66}{233} = \frac{2}{100}$:. Mole of $SO_4^{2-} = \frac{2}{100}$ Mass of $SO_4^{2-} = \frac{2}{100}$ (ionic mass of SO_4^{2-}) = 1.92 g % SO₄²⁻ = $\frac{1.92}{5} \times 100 = 38.4\%$.

20. Let the volume of ethanol containing the same number of molecules as are present in 175 ml of H_2O be V ml. As given,

moles of C_2H_5OH in V ml = moles of H_2O in 175 ml

Now,
$$\frac{wt.of \ C_2H_5OH}{mol.wt.of \ C_2H_5OH} = \frac{wt.of \ H_2O}{mol.wt.of \ H_2O}$$

or,
$$\frac{0.789 \times V}{46} = \frac{1.0 \times 175}{18} \qquad \therefore \qquad V = 566.82 \text{ ml.}$$

21. % of Na =
$$\frac{mass \ of \ sodium}{molecular \ mass} \times 100 \implies 7 = \frac{23}{M} \times 100$$

M = $\frac{23 \times 100}{7}$ = 328.6

22.
$$CaCl_2 + Na_2CO_3 \longrightarrow CaCO_3 + 2 NaCl$$

 $CaCO_3 \xrightarrow{\Delta} CaO + CO_2$

Mole of CaCl₂ = mole of CaCO₃ = mole of CaO = $\left(\frac{1.62}{56}\right)$

Mass of CaCl₂ = $\left(\frac{1.62}{56}\right)$ Molar mass of CaCl₂ = $\left(\frac{1.62}{56}\right) \times 111$ g. % of CaCl₂ = $\frac{3.21}{10} \times 100 = 32.1$ %.

23.	KI is limiting reagent \therefore 3 mole of KI will give 33 mole of NO ₂ according to stoichiometry.
24.	$\begin{array}{l} Ag_{2}CO_{3}\left(s\right) \longrightarrow 2Ag\left(s\right) + CO_{2}\left(g\right) + 1/2O_{2}\left(g\right) \\ C_{2}H_{2} + 5/2O_{2} \longrightarrow 2CO_{2} + H_{2}O \\ By Stoichiometry of reaction \end{array}$
	Moles of CO_2 formed = $\frac{11.2}{22.4} = \frac{1}{2}$
	Moles of O ₂ required = $\frac{5}{4} \times \frac{1}{2} = \frac{5}{8}$
	$\therefore \text{ Moles of Ag}_2\text{CO}_3 \text{ required } = 2 \times \frac{5}{8} = \frac{5}{4} \qquad ; \qquad \text{Mass of Ag}_2\text{CO}_3 \text{ required } = \frac{5}{4} \times 276 = 345 \text{ g}$
25.	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$
	PART - 2
1.	mol. wt. $\operatorname{CaCl}_2 = 111 \text{ g}$ $\therefore 111 \text{ g CaCl}_2 \text{ has } N_A \text{ ions } \operatorname{Ca}^{+2} (N_A = \operatorname{Avogadro number})$ $\therefore 222 \text{ g CaCl}_2 \text{ has } N_A \text{ ions } \operatorname{Ca}^{+2} = \frac{N_A \times 222}{111} = 2 \text{ N}_A \text{ ions of } \operatorname{Ca}^{+2}$ $\therefore 111 \text{ g CaCl}_2 \text{ has } 2 \text{ N}_A \text{ ions of } \operatorname{Cl}^{-}$ $\therefore 222 \text{ g CaCl}_2 \text{ has } 2 \text{ N}_A \text{ ions of } \operatorname{Cl}^{-} = \frac{2 \times N_A \times 222}{111} \text{ ions of } \operatorname{Cl}^{-} = 4 \text{ N}_A \text{ ions of } \operatorname{Cl}^{-}$
2.	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$
	finally $\frac{x}{65} - \frac{x}{254} = 0$ $\frac{x}{254}$ Fraction of Zn unreacted $= \frac{\frac{x}{65} - \frac{x}{254}}{\frac{x}{65}} = 1 - \frac{65}{254} = 0.744$
3.	On passing through charcoal only CO_2 reduces to CO. $CO + C \longrightarrow No$ reaction Volume a
	$\begin{array}{ccc} & & CO_2 + C \longrightarrow 2 CO \\ \text{Volume before reaction} & & & 0 \\ \text{Volume after reaction} & & & 0 \\ \text{As given} & & a + b = 1 \\ & & \text{and} \\ \end{array} \begin{array}{c} & & a + 2b = 1.4 \end{array}$
	:. $b = 0.4$ litre :. % of $b = \frac{0.4}{1} \times 100 = 40$ %
	∴ $a = 0.6$ litre ∴ % of $a = \frac{0.6}{1} \times 100 = 60$ %

4. MnO₁⁻; x + 4(-2) = -1 or x = +7;
CrO₂Cl₂; x + 2(-2) + 2(-1) = 0 or x = +6.
5. Let the volume of oxygen in 1120 mL of ozonised oxygen be x mL at S.T.P.
∴ Volume of ozone = (1120 - x) mL at S.T.P.
We known that
vol. of mixture x its density = mass
= vol. of oxygen x its density + vol. of ozone x its density
Also, density =
$$\frac{mass}{volume}$$

∴ density of ozone = $\frac{48}{22400}$ g/mL (at S.T.P.)
and density of ozone = $\frac{48}{22400}$ g/mL (at S.T.P.)
Hence, x x $\frac{32}{22400}$ + (1120 - x) x $\frac{48}{22400}$ = 1.76
or, 2x + (1120 - x) x 3 = 1.76 × 1400
or, x = (3360 - 2464) mL = **996 mL O**₂.
6. (A) and (B) Explanation: 30% of molecule dissociated N₂ → 2N
Amount of N₂ left $= \frac{2.8}{2.8} \times \frac{70}{100} = 0.1 \times 0.7 = 0.07$
(in moles)
No. of moles of N atoms formed = 2x $\frac{30}{100} \times 0.1 = 0.06$
(A) Total no. of molecules = 0.07 × 0.023 × 10²³ = 4.2 × 10²² molecule = 0.421 × 10²³
∴ We have to calculate molecule of introgen not atoms.
7. Let W gas of SO₂ and O₂ are taken
molecules of O₂ = $\frac{WN_A}{32}$; molecules of SO₂ = $\frac{WN_A}{64}$
hence molecules of O₂ > molecules of SO₂
since moles of Q₂ = moles of SO₂, hence volume of O₂ at 0°C & 1 atm > volume of SO₂ at 0°C & 1 atm
8. $2P + Q \longrightarrow R$
initial mole 12 8 0
final mole 12 8 0
fi

n = 4M.wt = 58 + [165]n g/mol = 718 g/mol

Mw of $CaCO_3 = 100$, Mw of $Na_2CO_3 = 106$ 10. Mw of HNO₃ = 63 g mol⁻¹ $Na_2CO_3 + CaCO_3 \longrightarrow CaCO_3 + 2NaCl$ (a) moles of CaCO₃ = $\frac{10}{100}$ = 0.1 mol moles of $Na_2CO_3 = moles of CaCO_3 = 2 \times moles of NaCl$ Weight of $Na_{3}CO_{3} = 0.1 \times 106 = 10.6 \text{ g}$ % purity $Na_2CO_3 = \frac{10.6}{21.2} = 100 = 50\%$ (b) wrong (c) correct (d) moles of NaCl = 2 × 0.1 = 0.2 mol 11. Cr oxidises from +3 to +6 while I reduces from +5 to -1. One I atom gain 6 electron. 13. Let mole % of ²⁶Mg be x. $\frac{(21-x)25+x(26)+79(24)}{100}=24.31$ *.*... x = 10% Answer = 1 $CaCO_{3} \longrightarrow CaO + CO_{2}$ 14. $\frac{5.6}{22.4} = \frac{1}{4}$ mole mole of CaO = mole of Ca = $\frac{1}{4}$ mass of Ca = $\frac{1}{4} \times 40 = 10$ % of Ca in sample = $\frac{10}{200} \times 100 = 5\%$ 15. Let volume of solution is 1000 ml moles of $H_2SO_4 = 18$ mass of $H_2 SO_4 = 18 \times 98 = 1764 g$ mass of solution = $1000 \times 1.8 = 1800$ g mass of solvent = 1800 - 1764 = 36 g molality = $\frac{18}{\left(\frac{36}{1000}\right)}$ = 500 $\Rightarrow \frac{500}{500}$ = 1 Let each species be a moles, M be molecular mass of metal 16. $a \times [2 \times 7 + 12 + 48] + a [2 \times M + 12 + 48] = 1$ (1) and a moles of each carbonate reacts with 2a mole of HCI hence $4a = 44.44 \times 0.5 \times 10^{-3}$ or $a = 11.11 \times 0.5 \times 10^{-3}$(2) Thus M from solving the equation (1) and (2) is 23 g M = 23 gM - 16 = 7

18.

17. From one mole of initial mixture, some FeO must have reacted with oxygen and got converted into Fe_2O_3 .

 $4 \text{FeO} + \text{O}_2 \longrightarrow 2 \text{Fe}_2 \text{O}_3$ $\frac{3}{5}$ 2 5 Initial moles $\frac{2}{5} + \frac{x}{2}$ $\frac{3}{5} - x$ Final moles But, final moles ratio is 2 : 3. $\therefore \qquad \frac{\left(\frac{3}{5}-x\right)}{\left(\frac{2}{5}+\frac{x}{2}\right)} = \frac{2}{3}$ \therefore $x = \frac{1}{4}$ $\therefore \qquad \text{Moles of FeO reacted} = x = \frac{1}{4}$:. Moles of O_2 required = $\frac{1}{4}(x) = \frac{1}{16} = 0.0625$ *.*:. Mass of O_2 required = 0.0625 × 32 = 2 g Let x be the mass of $CaCO_3$ hence mass of MgCO₃ = 92 - x $CaCO_3$ + $MgCO_3$ $\frac{x}{100}$ $\frac{92-x}{84}$ $CaO + CO_2$ MgO + CO₂ x 100 <u>92 - x</u> 84 mass of residue = 48 g $\frac{x}{100} \times 56 + \frac{92 - x}{84} \times 40 = 48$ \Rightarrow $\frac{x}{100} + \frac{92 - x}{84} = \frac{6}{7}$ \Rightarrow x = 50 \Rightarrow mass of MgCO₃ = 92 - 50 = 42 g. *.*..

PART - 3

18. mole = $\frac{w}{m} = \frac{1}{m}$ for largest no. of molecule m should be lowest.

19. $X = \frac{20}{M_1 \times 50}$ $Y = \frac{15}{70 \times M_2}$

X = Y
$$\frac{20}{M_1 \times 50} = \frac{15}{70 \times M_2}$$
; $\frac{28}{15} = \frac{M_1}{M_2}$

20.
$$3 \operatorname{Bacl}_{2} + 2\operatorname{NO}_{3}\operatorname{PO}_{4} \longrightarrow \operatorname{Ba}_{4}(\operatorname{PO}_{4})_{2} + 6\operatorname{Nacl}$$

 $\frac{0.5}{3}$ $\frac{0.2}{2}$ 0.1
21. $\frac{1}{2} \times 6.023 \times 10^{23} = 3.0125 \times 10^{23}$
37. $\frac{3.7 \times 1 + 5 \times 0.3}{8.7} = \frac{3.7 + 1.5}{8.7} = \frac{5.2}{8.7} = \frac{52}{87}$
38. $\operatorname{H}_{2}C_{0}O_{2}\operatorname{H}_{2}O$
 $\frac{252 \times 10^{-3}}{126} \times 2 \times \operatorname{N}_{A} = 4 \times \operatorname{N}_{A} \times 10^{-3} = 4 \times 6.02 \times 10^{23} \times 10^{-3} = 24.092 \times 10^{23} = 2.4 \times 10^{21}.$
39. $\begin{array}{c} & & & \\ &$

48. Number of water molecules =
$$\frac{0.2}{249.7}$$
 N_A × 5 = 2.14 × 10²¹
49. Cs[Mn(SO₄)₂.12H₂O]
x - 4 = -1
x = + 3
50. AgNO₃ + Cl⁻ → AgCl ↓ + NO₃⁻
1.4 g
= $\frac{1.4}{143.5}$ mol
= $\frac{1.4}{143.25}$ × 35.5 gm Cl = 0.35 gm
51. VD_A = 4VD_B or $\frac{M_A}{2} = 4 \times \frac{M_B}{2}$, or M_A = 4M_B
52. 2KClO₃ → 2KCl + 3O₂
n_{KClO3} = $\frac{36.75}{122.5} = 0.3$
By mole-mole analysis $\frac{n_{KClO_3}}{2} = \frac{n_{O_2}}{3}$
or $n_{O_2} = \frac{3}{2} \times 0.3 = 0.45$
or volume of O₂ = 0.45 × 22.4
= 10.08 lit.
= 10.08 dm³
53. 500 gm toothpaste contains 0.4 g fluoride
∴ 10⁶ g toothpaste will contain $\frac{0.4}{500} \times 10^6 = 800$ g fluoride

 \therefore ppm of fluoride = 800

54.

78 gm benzene gives 123 gm nitrobenzene

$$\therefore \qquad 10 \text{ gm benzene} = \frac{123}{78} \times 10 = 15.77 \text{gm}(\text{should be})$$

% age yield = $\frac{1}{T}$

$$= \frac{\text{Actual yield}}{\text{Theoretical yield}} \times 100$$

$$=\frac{13.2}{15.77}\times 100 \simeq 83.5\%$$

56. I. 5H₂O₂ + 2MhO₁⁻ + 6H⁻ → 2Mh⁻² + 5O₂ + 8H₂O
II. H₂O₂ + Ag₂O → 2Ag + H₂O + O₂
57. Al₁C₄ + 12H₂O → 3CH₄ + 4Al(OH)₅

$$\frac{11.2}{22.4} = \frac{1}{2} mole$$
Mole of methane = $\frac{11.2}{22.4} = \frac{1}{2} mole$
Wt of Al₄C₅ = $\frac{1}{6} \times 144 = 24 g$
58. D = $\frac{M}{V} \Rightarrow 1.42 = \frac{100}{V}$
 $\Rightarrow V = \frac{100}{1.42}$
Molarity of HNO₂ = $\frac{70 \times 1.42 \times 1000}{63 \times 100} = 15.8$

PART - 4
1. SO₂ that converted = 440 × $\frac{80}{100}$ Kg = 352 kg
 $5C + 2SO2 \longrightarrow CS2 + 4CO$
Initial mole $5 \times 10^3 = \frac{352}{64} \times 10^3 = 0 = 0$
 $= 5.5 \times 10^5$
final mole $0 = 5.5 \times 10^3 = \frac{352}{5} \times 5 \times 10^3 = 1 \times 10^3 = 4 \times 10^3$
mole of CS₃ = 1000
mass = 1000 × 76 g = 76 Kg
2. (a) Br₂ + Cl₂ → 2BrCl
 $1 \mod 1 \mod 2 \mod 0$
(b) Theoretical yield of BrCl = 0.050 \mod 0

actual yield is = $\frac{0.05 \times 80}{100}$ = 0.04 mole

 Br_2 left unreacted = 0.025 - 0.02 = 0.005 mol.

5. Balanced chemical equation is

 $4nXeF_{6} + + CH_{2} - CH_{2} - CH_{2} - CF_{2} - CF_{2} - CF_{2} - H_{1} + 4nHF + 4nXeF_{4}$

$$n_{teflon} = \frac{100}{100 n} = \frac{1}{n}$$

 \therefore n_{XeF_6} required $=\frac{1}{n} \times 4n = 4$ moles

ONLY ONE OPTION CORRECT TYPE

1.		TiCl ₄ +	$2Mg \longrightarrow$	Ti	+	2MgCl ₂
	Initial mole	$\frac{358}{190} = 1.88$	$\frac{96}{24} = 4$			
	final mole	0	4 – 2 × 1.88	1.88		2 × 1.88
	wt of Ti obtaine	$ed = \frac{358}{190} \times 48$				
	% yield = $\frac{32}{358}$	$\frac{100}{\times 48} = 35.38 \%$	1			

2. Produced mass of
$$H_3PO_4 = \left(\frac{62}{4 \times 31}\right) \times 0.85 \times 0.9 \times 4 \times 98 = 149.94 \text{ g}$$

- 3. The balanced equation is $2MnO_4^- + 5C_2O_4^{2-} + 16H^+ \longrightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$
- 4. 10^{-3} g NH₃ in 100 g solution one litre water has mass = 1000×1 g As NH₃ is very less hence we can say 100 g water has 10^{-3} g NH₃

:. 1000 g water has =
$$\frac{10^{-3}}{100} \times 1000$$
 g = 10^{-2} g NH₃ = $\frac{10^{-2}}{17}$ mole NH₃ = 5.88×10^{-4} mole NH₃.

5.

6.

7.

8.

Mole of AI = $\frac{18}{27}$ = $\frac{2}{3}$ Mole of HCl = $\frac{109.5}{36.5}$ = 3 Moles of NaOH = $\frac{100}{40}$ = 2.5 $2 \text{ AI} + 6 \text{ HCI} \longrightarrow 2 \text{ AICI}_3 + 3 \text{H}_2$ 0 Initial mole 2/3 3 0 final mole 2/30 1 $AICI_3 + 3NaOH \longrightarrow AI(OH)_3 + 3NaCI$ 0 0 2/3 2.5 Initial mole final mole $0 \qquad 2.5 - 2/3 \times 3$ 2/3 = 0.5 $AI(OH)_3 + NaOH \longrightarrow NaAIO_2 + 2H_2O$ 2/3 0.5 Initial mole 0 0 0.5 final mole Ans. $NaAlO_2 = 0.5$ moles. $3M + N_2 \longrightarrow M_3N_2$ Let Atomic wt. of metal = a (3a + 28) g nitride contains metal = 3a gram so 14.8 g nitride contains metal = $\frac{3a}{3a+28} \times 14.8 = 12$ ÷ a = 40. so By applying POAC for C atoms moles of ethylene $\times 2 =$ mole of polythene $\times n \times 2$ $\frac{100g}{28} \times 2 = \frac{\text{wt. of polethene}}{28 \times n} \times n \times 2$ wt. of polyethene = 100 gC + $CO_2 \longrightarrow 2CO$ $\left(\frac{6}{12}\right) = 0.5 \qquad \left(\frac{44}{44}\right) = 1$ Given moles

So C is limiting reagent

 \therefore CO formed = 1 moles

Now moles of Ni need to react with 1 moles of CO are $\frac{1}{4} \times 58.7 = 14.675$ g.

9. $H_2SO_4 + 2NaOH \longrightarrow Na_2SO_4 + 2H_2O$ for 1 mole NaOH $\frac{1}{2}$ mole H₂SO₄ required 70 g H₂SO₄ in 100 g solution $\frac{98}{2}$ g H₂SO₄ in = $\frac{100}{70}$ × 49 = 70 g solution. Li AlH₄ + t-butyl alcohol $\xrightarrow{\text{Ether}}$ LiAlHC₁₂H₂₇O₃ (M.W. = 254) 10. 0.05 mole 12.7 gram $=\frac{12.7}{254}=0.05$ mole Li atom remain conserved so No. of mole of LiAlH₄ = No. of mole of LiAlHC₁₂H₂₇O₃ So No. of mole of $LiAIHC_{12}H_{27}O_3 = 0.05$ % yield = $\frac{0.05}{0.05} \times 100 = 100\%$ $\longrightarrow P_4O_6$ + 11. O₂ P_4O_{10} P₄ + 31 gram 32 gram According to question weight of P is conserved so Let Mole of $P_4O_6 = a$ Mole of $P_4O_{10} = b$ Initial weight of P = Final weight of P. $31 = [a \times 4] \times 31 + [b \times 4] \times 31$ 4 a + 4b = 1] $(1) \times 3$ Initial weight of oxygen = Final weight of oxygen $32 = [a \times 6] \times 16 + [a \times 10] \times 16$ 3a + 5b = 1] $(2) \times 4$ 12a + 20b = 412 a + 12 b = 3 So $b = \frac{1}{8}$ 8 b = 1 Similarly $a = \frac{1}{\alpha}$ $P_4O_6 = \frac{1}{8} \times 220 = 27.5$ $P_4O_{10} = \frac{284}{8} = 35.5.$ So weight of 12. $CaCl_{2} + NaCl = 10 g$ Let weight of $CaCl_2 = x g$ $CaCl \rightarrow CaCO_{3} \rightarrow CaO$ 1 mol 1 mol 1 mol

 $\frac{x}{111} \mod \frac{x}{111} = \frac{1.62}{56}$ x = 3.21 g% of $\operatorname{CaCl}_2 = \frac{3.21}{10} \times 100 = 32.1 \% = \frac{3.21}{10} \times 100 = 32.1 \%$

13. Mole fraction of A i.e.
$$X_A = \frac{n_A}{\text{Total moles}}$$

So
$$X_{H_2O} = \frac{n_{H_2O}}{\text{Total moles}}$$

Now
$$\frac{X_A}{X_{H_2O}} = \frac{n_A}{n_{H_2O}}$$

and molality =
$$\frac{n_A \times 1000}{n_{H_2O} \times 18} = \frac{X_A \times 1000}{X_{H_2O} \times 18} = \frac{0.2 \times 1000}{0.8 \times 18} = 13.9 \text{ Ans.}$$

14. Let wt. of Fe = 100 g so wt. of
$$O_2 = 10 \text{ g}$$

$$2\text{Fe} + \frac{3}{2}\text{O}_2 \longrightarrow \text{Fe}_2\text{O}_3$$

С

by the stoichiometry of the reaction $\frac{10}{32}$ mole of O₂ will combine with $\frac{10}{24}$ mole of Fe

wt. of Fe =
$$\frac{10}{24} \times 56 = 23.3$$
 g or 23.3%.

15.

+
$$\frac{1}{2}O_2 \longrightarrow CO \dots (1)$$

0

Initial mole

final mole 0
$$\frac{y}{32} - \left(\frac{x}{12}\right)\frac{1}{2}$$

 $\frac{x}{12}$ $\frac{y}{32}$

$$CO \qquad + \qquad \frac{1}{2}O_2 \qquad \longrightarrow \qquad CO_2 \qquad \dots (2)$$

For no solid residue C should be zero in eq. (1) For that $\frac{y}{32} - \frac{x}{12} \times \frac{1}{2} > 0$

$$\frac{y}{32} > \frac{x}{24}$$
$$\frac{y}{x} > \frac{32}{24}$$
$$\frac{y}{x} > 1.33$$

16. Mass of ethyl alcohol = 1.5×0.792 g Mass of water = 15×1 Total mass of solution = $15 + 0.792 \times 15 = 26.88$

Volume of solution = $\frac{\text{mass}}{\text{density}} = \frac{26.88}{0.924} = 29.09$

% decrease in volume =
$$\left(\frac{30-29.09}{30}\right) \times 100 \cong 3\%$$
.

PART - 3

2. (A) Molarity of cation =
$$\frac{M_1V_1 + M_2V_2}{V_1 + V_2} = \frac{0.2 \times 100 + 0.1 \times 400}{500} = \frac{0.6}{5} = 0.12$$

Molarity of Cl⁻ =
$$\frac{3(0.2)100 + 0.1 \times 400}{500} = \frac{0.6 + 0.4}{5} = 0.2$$

(B) Molarity of cation =
$$\frac{50 \times 0.4 + 0}{100} = 0.2$$

Molarity of Cl⁻ =
$$\frac{0.4 \times 50 + 0}{100} = 0.2$$

(C) Molarity of cation = $\frac{2(0.2)30 + 0}{100} = 0.12$

Molarity of
$$SO_4^{2-} = \frac{30 \times 0.2}{100} = 0.06$$

(D) 24.5 g H_2SO_4 in 100 ml solution

Molarity =
$$\frac{\frac{25.4}{98}}{0.1}$$
 = 2.5

$$\therefore \qquad \text{Concentration of cation} = 2 \times 2.5 \text{ M}$$

Concentration of SO,²⁻ = 2.5 M.

SINGLE AND DOUBLE VALUE INTEGER TYPE

- Density of He = 0.1784 g/lit.
 1 mole of He will occupy 22.4 lit. at 0°C and 1 atm
 ∴ Mass of 1 mole = V x d = 22.4 x 0.1784 = 3.99 = 4 g.
- 2. Mass of $Cu_2S \& CuS = 100 4.5 = 95.5 g$ Let mass of Cu_2S is x g. $Cu_2S + O_2 \longrightarrow 2Cu + SO_2$ $CuS + O_2 \longrightarrow Cu + SO_2$ Mass of Cu from $Cu_2S + Mass$ of Cu from CuS = 71.8

$$\frac{x}{159} \times 63.3 \times 2 + \frac{(95.5 - x)}{95.5} 63.5 = 71.8$$

$$x\left(\frac{127}{159} - \frac{63.5}{95.5}\right) = 8.3$$

$$x = \frac{8.3}{0.134} = 62.01 \text{ g}$$

$$\therefore$$
 % of Cu₂S is 62.

3. Molarity of
$$H_2SO_4 = \frac{1.8 \times 54.5 \times 10}{98} = 10$$

 $2AI + 3H_2SO_4 \longrightarrow Al_2 (SO_4)_3 + 3H_2$
1 moles 2 moles
(limiting)
Moles of H_2SO_4 left = 2 - 1.5 = 0.5 moles
moles of HCl added = 2 moles
final volume of the solution = 500 ml
moles of H⁺ ion = 3
concentration of H⁺ ion = 6 M

4. 1 × moles of
$$CO_2 = 6n \times moles$$
 of starch

$$= 6n \times \frac{1}{162n}$$

So moles of $CO_2 = \frac{6}{162}$

Now 4.7×10^{-3} moles of CO₂ are absorbed in 1 hr

So $\frac{6}{162}$ moles of CO₂ are absorbed in = $\frac{1}{4.7 \times 10^{-3}} \times \frac{6}{162}$ = 8 hrs.

- 5. Balanced the equation. $15H_2O + 3CN^- \longrightarrow 3CO_2 + 3NO_3^- + 30H^+ + 30e^-$
- 6. Mole of NaOH in 1^{st} solution = 0.5 moles

moles of NaOH addded = $\frac{200 \times 1.5 \times 0.2}{40}$ = 1.5

moles of NaOH in the final solution = 1.5 + 0.5 = 2 moles Al + NaOH + H₂O \longrightarrow NaAlO₂ + 3/2 H₂ moles of H₂ produced from 2 moles of NaOH = 3 moles volume of H₂ produced at 0°C and 1 atm = $3 \times 22.4 = 67.2$ litre **Ans.** 67

7. m moles of HCl =
$$12 \times 0.05 = 0.6$$

Now AI + 3HCI \rightarrow AICI₃ + $\frac{3}{2}$ H₂

so m moles of AI = $\frac{1}{3} \times 0.6$

or weight of AI =
$$\frac{1}{3} \times \frac{0.6 \times 27}{1000} = 0.0054$$
 gram

:. Volume of foil =
$$\frac{0.0054}{2.7}$$
 mL or cm³ = 0.002 cm³

Now, Area x thickness = Volume

Area = $\frac{0.002}{0.01}$ = 0.2 cm²

(thickness = 0.01 cm)

8. A +
$$\frac{1}{2}$$
 B₂ \longrightarrow AB, 100 Kcal
x x/2 x
A + 2B₂ \longrightarrow AB₄, 200 Kcal
(1-x) 2(1-x) (1-x)
100 x + 200 (1-x) = 140

200 - 100 x = 140 $x = \frac{60}{100} = 0.6$ $n_{B_2} \text{ used} = \frac{x}{2} + 2(1-x) = \frac{1}{2} \times 0.6 + 2(1-0.6) = 0.3 + 2 \times 0.4 = 1.1 \text{ mol}$ $Ans = 1.1 \times 10 = 11$

9. (Atomic weight of Al and Cr = 27 and 52, M.wt. of $Cr_2O_3 = 152$)

Moles of AI =
$$\frac{49.8 \text{ g}}{27 \text{ g AI}}$$
 = 18.4 mol

$$=\frac{18.4}{2}=9.2$$
 mol of Cr₂O₃

Moles of $Cr_2O_3 = \frac{200 \text{ g}}{152 \text{ g} Cr_2O_3} = 1.31 \text{ mol}$

Since 2 mol Al is required for 1 mol of Cr_2O_3 . So, Al is the limiting reagent and Cr_2O_3 is in excess. Moles of Cr_2O_3 is excess = (1.31 - .92) = 0.4 mol Weight of excess $Cr_2O_3 = 0.4 \times 152 = 60 \text{ g} Cr_2O_3$

10. Balance the equation by any method $4Zn + 10HNO_3 \longrightarrow 4Zn(NO_3)_2 + 3H_2O + NH_4NO_3$ $\therefore a + b + c = 4 + 3 + 1 = 8$

11. Molarity = $\frac{10 \times 1.8 \times 98}{98}$ = 18 M

12. Molarity of HCl =
$$\frac{\text{Total moles of HCl}}{\text{Total volume}} = \frac{5 \times 2}{2 + 3} = 2 \text{ M}$$

ONE OR MORE THAN ONE OPTIONS CORRECT TYPE

2.	Let	mol of $N_2 = x$,	mol of	$NO_2 = y,$	mol of $N_2O_4 = z$			
	therefore	$\frac{28x+46y+92z}{1}$	= 55.4	(1)				
	lf	$N_2O_4 \longrightarrow 2NO_4$	D ₂					
		$\frac{28x+(y+2z)46}{x+y+z+z}$	= 39.6					
	\Rightarrow	$\frac{28x+46y+92z}{1+z}$	= 39.6	(2)				
	By deviding equ	uation (1) by equa	tion (2)					
		$1 + z = \frac{55.4}{39.6} =$	1.4					
		z = 0.4 mol						
	Given	x + y + z = 1		(3)				
	Put the value of z in eq. (1)							
		28x + 46y + 92 ·	+ 0.4 = 55.4					
		28x + 46y = 18.0	6	(4)				
	By equation (3)	& (4)						
		y = 0.1						
		x = 0.5 ,	y = 0.1,	z = 0.4				
3.	(A) , (C) and (D)) Explanation :						
	4 Ag +	8 KCN + O ₂ + 2 F	$H_2O \longrightarrow 4 \text{ K}$	Ag(CN) ₂] + 4 KO	н			
	\Rightarrow 4 x 108 g of Ag reacts with 8 x 65 g of KCN							
	100 g c	of Ag reacts with						
		$\frac{8\times65}{4\times108}$	× 100 = 120					
	Hence ,, to diss	olve 100 g of Ag ,	the amount of	KCN required = 1	20 g			
	Hence, statem	ent (A) is correct.						
	\Rightarrow 4 × 108	3 g of Ag require 3	2 g of O_2					
	1 g of A	Ag require $\frac{32}{4 \times 108}$	= 0.0740 g					

 $\Rightarrow 100 \text{ g of Ag require} = 7.4 \text{ g}$ Hence, choice (C) is correct. Hence, volume of O_2 required = $\frac{7.4}{32} \times 22.4 = 5.20$ litre Hence, (A), (C), (D) are correct while (B) is incorrect.

$$CaO(s) + 3C(s) \longrightarrow CaC_2(s) + CO(g)$$

(A) Final product contain
$$85\%$$
 CaC₂ & 15% CaO
Let mass of product is 100 g

 $\therefore \qquad \text{Mass of } CaC_2 = 85 \text{ g}$ Mass of CaO = 15 g

Used mole of CaO = mole of CaC₂ produced = $\frac{85}{64}$

 \therefore mass of CaO for producing 85 g CaC₂ = $\frac{85}{64}$ × 56 = 74.375 g.

$$\therefore \qquad \text{Initial total mass of CaO} = 74.375 + 15 = 89.375$$
$$85 \text{ g CaC}_{2} \text{ obtained from} = 89.38 \text{ g CaO}$$

$$\therefore \qquad 1 \text{ g CaC}_2 \text{ obtained from} = \frac{89.38}{85} \text{ g CaO}$$

$$10^6 \text{ g CaC}_2 \text{ obtained from} = \frac{89.38}{85} \times 10^6 = 1051470 \text{ g}$$

For 1000 kg CaC_2 requires = 1051.47 kg CaO.

(B) 100 g product requires CaO = 89.38 g

1 g product requires =
$$\frac{89.38}{100}$$

$$10^6 \text{ g product requires} = \frac{89.38}{100} \times 10^6$$

For 1000 kg (crude) product = 893.8 kg CaO.

5. Mol of $Cu^{2+} = 1.0 L \times 0.1 M = 0.1 M Cu^{2+} = 0.1 \times 2 mol H^+$ (A) Weight of CuS = 0.1 × 95.5 = 9.55 g

(B) Concentration of H⁺ =
$$\frac{0.2 \text{ mol}}{1.0 \text{ L}} = 0.2 \text{ M}$$

(C) and (D) are wrong.

4.

6. (A) Weight of $CaCO_3 = (0.22 \text{ g } CO_2)$

$$\left(\frac{1 \operatorname{mol} \operatorname{CO}_2}{44 \operatorname{g} \operatorname{CO}_2}\right) \left(\frac{1 \operatorname{mol} \operatorname{CaCO}_3}{\operatorname{mol} \operatorname{CO}_2}\right) \left(\frac{100 \operatorname{g} \operatorname{CaCO}_3}{\operatorname{mol} \operatorname{CaCO}_3}\right) = \frac{0.22 \times 100}{44} = 0.5 \operatorname{g} \operatorname{CaCO}_3$$

(B) Moles of CaCO₃ = moles of Ca = $\left(\frac{0.22}{44}\right)$ = 0.005 mol

Weight of Ca = $0.005 \times 40 = 0.2$ g Ca

(D) % of Ca =
$$\frac{0.2}{1.0} \times 100 = 20\%$$
 Ca

Hence (C) is wrong.

7.

	Silica H	l ₂ O	Impurities
% in original clay \Rightarrow	40	19	100 - (40 + 19) = 41
% after partial drying \Rightarrow	а	10	100 – (a + 10) = 90 – a

On heating, only water evaporates from clay, whereas silica and impurities are left as it is. Therefore, % ratio of silica and impurities remains unchanged, i.e.

 $\frac{40}{a} = \frac{41}{90-a} \,, \ \therefore \ a = 44.4\%$

% of mipurities after partial drying = (90 - a) = (90 - 44.4) = 45.6%