DIRECTORATE OF EDUCATION Govt. of NCT, Delhi

SUPPORT MATERIAL (2020-2021)

Class : XI

CHEMISTRY

Under the Guidance of

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Published at Delhi Bureau of Text Books , 25/2 Institutional Area, Pankha Road, New Delhi-110058 by **Prabhjot Singh,** Secretary, Delhi Bureau of Text Books and Printed by Arihant Offset, New Delhi-110043

MANISHA SAXENA



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DO No. DE. 5/228/Exam/Message/S.M./2018 Dated

MESSAGE

The importance of adequate practice during examinations can never be overemphasized. I am happy that support material for classes IX to XII has been developed by the Examination Branch of Directorate of Education. This material is the result of immense hard work, co-ordination and cooperation of teachers and group leaders of various schools. The purpose of the support material is to impart ample practice to the students for preparation of examinations. It will enable the students to think analytically & rationally and test their own capabilities and level of preparation.

The material is based on latest syllabus prepared by the NCERT and adopted by the CBSE for the academic session 2020-21 and covers different levels of difficulty. I expect that Heads of Schools and Teachers will enable and motivate students to utilize this material during zero periods, extra classes and regular classes best to their advantage.

I would like to compliment the team of Examination Branch for their diligent efforts of which made it possible to accomplish this work in time. I also take this opportunity to convey my best wishes to all the students for success in their endeavours.

24.6 2020

(MANISHA SAXENA)

BINAY BHUSHAN, IAS



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D.O. No.

Date :

Dear Students,

Directorate of Education is committed to providing qualitative and best education to all its students. The Directorate is continuously engaged in the endeavor to make available the best study material for uplifting the standard of its students and schools.

Every year, the expert faculty of Directorate reviews and updates Support Material. The expert faculty of different subjects incorporates the changes in the material as per the latest amendments made by CBSE to make its students familiar with new approaches and methods so that students do well in the examination.

The book in your hand is the outcome of continuous and consistent efforts of senior teachers of the Directorate. They have prepared and developed this material especially for you. A huge amount of money and time has been spent on it in order to make you updated for annual examination.

Last, but not the least, this is the perfect time for you to build the foundation of your future. I have full faith in you and the capabilities of your teachers. Please make the fullest and best use of this Support Material.

Y BHUSHAN DIRECTOR (EDUCATION)

Dr. (Mrs.) Saroj Bala Sain Addl. Director of Education (School / Exam / EVGB//EB/ VOC.)



Govt. of NCT of Delhi Directorate of Education Old Secretaria, Delhi-110054 Tel.: 23890023, 23890093 D.O. No.<u>PA/AdU-06(s</u>uh)/86 Date : <u>03-10-2019</u>

I am very much pleased to forward the Support Material for classes IX to XII. Every year, the Support Material of most of the subjects is updated/revised as per the most recent changes made by CBSE. The team of subject experts, officers of Exam Branch, members of Core Academic Unit and teachers from various schools of Directorate has made it possible to make available unsurpassed material to students.

Consistence use of Support Material by the students and teachers will make the year long journey seamless and enjoyable. The main purpose to provide the Support Material for the students of government schools of Directorate is not only to help them to avoid purchasing of expensive material available in the market but also to keep them updated and well prepared for exam. The Support Material has always been a ready to use material, which is matchless and most appropriate.

I would like to congratulate all the Team Members for their tircless, unremitting and valuable contributions and wish all the best to teachers and students.

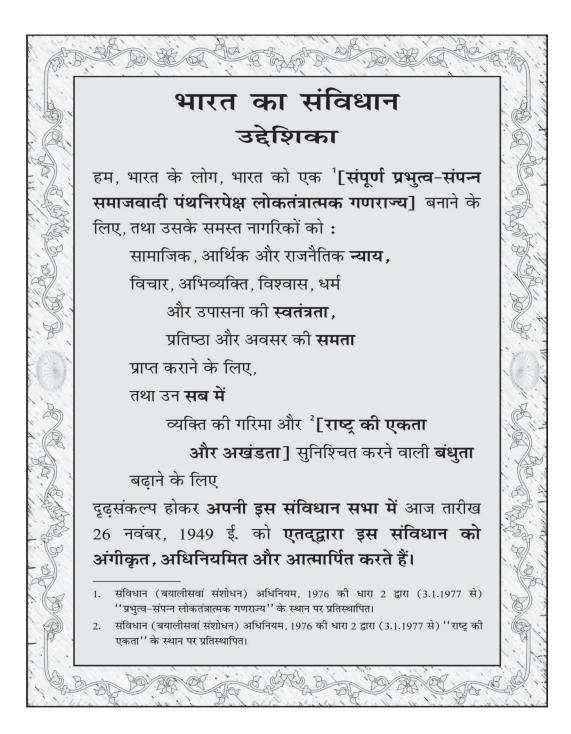
(Dr. Saroj Bala Sain) Addl.DE (School/Exam)

भारत का संविधान _{भाग 4क}				
	नागरिकों के मूल कर्तव्य			
अनु	च्छेद ५१क			
मूल व	फर्तव्य - भारत के प्रत्येक नागरिक का यह कर्तव्य होगा कि वह -			
(क)	संविधान का पालन करे और उसके आदर्शों, संस्थाओं, राष्ट्रध्वज और राष्ट्रगान का आदर करे;			
(폡)	स्वतंत्रता के लिए हमारे राष्ट्रीय आंदोलन को प्रेरित करने वाले उच्च आदर्शों को हृदय में संजोए रखे और उनका पालन करे;			
(ग)	भारत की संप्रभुता, एकता और अखंडता की रक्षा करे और उसे अक्षुण्ण बनाए रखे;			
(घ)	देश की रक्षा करे और आह्वान किए जाने पर राष्ट्र की सेवा करे;			
(퍟)	भारत के सभी लोगों में समरसता और समान भ्रातृत्व की भावना का निर्माण करे जो धर्म, भाषा और प्रदेश या वर्ग पर आधारित सभी भेदभावों से परे हो, ऐसी प्रथाओं का त्याग करे जो महिलाओं के सम्मान के विरुद्ध हों;			
(च)	हमारी सामासिक संस्कृति की गौरवशाली परंपरा का महत्त्व समझे और उसका परिरक्षण करे;			
(평)	प्राकृतिक पर्यावरण की, जिसके अंतर्गत वन, झील, नदी और वन्य जीव हैं, रक्षा करे और उसका संवर्धन करे तथा प्राणिमात्र के प्रति दयाभाव रखे;			
(ज)	वैज्ञानिक दृष्टिकोण, मानववाद और ज्ञानार्जन तथा सुधार की भावना का विकास करे;			
(झ)	सार्वजनिक संपत्ति को सुरक्षित रखे और हिंसा से दूर रहे;			
(ञ)	व्यक्तिगत और सामूहिक गतिविधियों के सभी क्षेत्रों में उत्कर्ष की ओर बढ़ने का सतत् प्रयास करे, जिससे राष्ट्र निरंतर बढ़ते हुए प्रयत्न और उपलब्धि की नई ऊँचाइयों को छू सके; और			
(5)	यदि माता-पिता या संरक्षक है, छह वर्ष से चौदह वर्ष तक की आयु वाले अपने, यथास्थिति, बालक या प्रतिपाल्य को शिक्षा के अवसर प्रदान करे।			

CONSTITUTION OF INDIA Part IV A (Article 51 A) Fundamental Duties

Fundamental Duties: It shall be the duty of every citizen of India —

- 1. to abide by the Constitution and respect its ideals and institutions, the National Flag and the National Anthem;
- 2. to cherish and follow the noble ideals which inspired our national struggle for freedom;
- 3. to uphold and protect the sovereignty, unity and integrity of India;
- 4. to defend the country and render national service when called upon to do so;
- 5. to promote harmony and the spirit of common brotherhood amongst all the people of India transcending religious, linguistic and regional or sectional diversities; to renounce practices derogatory to the dignity of women;
- 6. to value and preserve the rich heritage of our composite culture;
- 7. to protect and improve the natural environment including forests, lakes, rivers and wild life, and to have compassion for living creatures.
- 8. to develop the scientific temper, humanism and the spirit of inquiry and reform;
- 9. to safeguard public property and to adjure violence;
- 10. to strive towards excellence in all spheres of individual and collective activity so that the nation constantly rises to higher levels of endeavour and achievement.
- 11. who is a parent or guardian to provide opportunities for education to his child or, as the case may be, ward between the age of six and fourteen years.



THE CONSTITUTION OF INDIA PREAMBLE WE, THE PEOPLE OF INDIA, having solemnly resolved to constitute India into a ¹[SOVEREIGN SOCIALIST SECULAR **DEMOCRATIC REPUBLIC**] and to secure to all its citizens : JUSTICE, social, economic and political; LIBERTY of thought, expression, belief, faith and worship; **EQUALITY** of status and of opportunity; and to promote among them all **FRATERNITY** assuring the dignity of the individual and the ²[unity and integrity of the Nation]; IN OUR CONSTITUENT ASSEMBLY this twenty-sixth day of November, 1949 do HEREBY ADOPT, ENACT AND GIVE TO **OURSELVES THIS CONSTITUTION.** Subs. by the Constitution (Forty-second Amendment) Act, 1976, Sec.2, for "Sovereign Democratic Republic" (w.e.f. 3.1.1977) Subs. by the Constitution (Forty-second Amendment) Act, 1976, Sec.2, for "Unity of the Nation" (w.e.f. 3.1.1977)

DIRECTORATE OF EDUCATION Govt. of NCT, Delhi

SUPPORT MATERIAL (2020-2021)

CHEMISTRY Class : XI

NOT FOR SALE

PUBLISHED BY : DELHI BUREAU OF TEXTBOOKS

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Course Structure Class : XI (Theory) (2020-21) Chemistry

Time : 3 Hours]

Total period (Theory 160 + Practical 60)

Total Marks : 70

Time . 5 H	lours	IOIdI Mid	IKS.10
Unit No.	Title	No. of Periods	Marks
Unit I	Some Basic Concepts of Chemistry	12	11
Unit II	Structure of Atom	14	
Unit III	Classification of Elements and	08	04
	Periodicity in Properties		
Unit IV	Chemical Bonding and Molecular	14	21
	Structure		
Unit V	States of Matter: Gases, Liquids and solid	ls 12	
Unit VI	Chemical Thermodynamics	16	
Unit VII	Equilibrium	14	
Unit VIII	Redox Reactions	06	16
Unit IX	Hydrogen	08	
Unit X	s -Block Elements	10	
Unit XI	p -Block Elements	14	
Unit XII	Organic Chemistry: Some Basic	14	18
	Principles and Techniques		
Unit XIII	Hydrocarbons	12	
Unit XIV	Environmental Chemistry	06	
	Total	160	70



Unit I : Some Basic Concepts of Chemistry

8 Periods

10 Periods

General Introduction: Importance and scope of chemistry.

Nature of matter, laws of chemical combination, Dalton's atomic theory: concept of elements, atoms and molecules.

Atomic and molecular masses, mole concept and molar mass, percentage composition, empirical and molecular formula, chemical reactions, stoichiometry and calculations based on stoichiometry.

Unit II : Structure of Atom

Bohr's model and its limitations, concept of shells and subshells, dual nature of matter and light, de Broglie's relationship, Heisenberg uncertainty principle, concept of orbitals, quantum numbers, shapes of s, p and d orbitals, rules for filling electrons in orbitals - Aufbau principle, Pauli's exclusion principle and Hund's rule, electronic configuration of atoms, stability of half-filled and completely filled orbitals.

Unit III : Classification of Elements and Periodicity in Properties

06 Periods

Modern periodic law and the present form of periodic table, periodic trends in properties of elements -atomic radii, ionic radii, inert gas radii, Ionization enthalpy, electron gain enthalpy, electronegativity, valency. Nomenclature of elements with atomic number greater than 100

Unit IV : Chemical Bonding and Molecular Structure 14 Periods

Valence electrons, ionic bond, covalent bond, bond parameters, Lewis structure, polar character of covalent bond, covalent character of ionic bond, valence bond theory, resonance, geometry of covalent molecules, VSEPR theory, concept of hybridization, involving s, p and d orbitals and shapes of some simple molecules, molecular orbital theory of homonuclear diatomic molecules(qualitative idea only), hydrogen bond.

Unit V : States of Matter: Gases, Liquids and Solids 18 Period

Three states of matter, intermolecular interactions, types of bonding, melting and boiling points, role of gas laws in elucidating the concept of the molecule, Boyle's law, Charles law, Gay Lussac's law, Avogadro's law, ideal behaviour, empirical derivation of gas equation,



Avogadro's number, ideal gas equation. Deviation from ideal behaviour, liquefaction of gases, critical temperature, kinetic energy and molecular speeds (elementary idea)

Liquid State: vapour pressure, viscosity and surface tension (qualitative idea only, no mathematical derivations)

Solid state: Classification of solids based on different binding forces: molecular, ionic, covalent and metallic solids, amorphous and crystalline solids (elementary idea). Unit cell in two dimensional and three dimensional lattices, calculation of density of unit cell, packing in solids, packing efficiency, voids, number of atoms per unit cell in a cubic unit cell, point defects, electrical and magnetic properties.

Unit VI : Chemical Thermodynamics

16 Periods

Concepts of System and types of systems, surroundings, work, heat, energy, extensive and intensive properties, state functions. First law of thermodynamics -internal energy and enthalpy, heat capacity and specific heat, measurement of ΔU and ΔH , Hess's law of constant heat summation, enthalpy of bond dissociation, combustion, formation, atomization, sublimation, phase transition, ionization, solution and dilution. Second law of Thermodynamics (brief introduction). Introduction of entropy as a state function, Gibb's energy change for spontaneous and non- spontaneous processes, criteria for equilibrium. Third law of thermodynamics (brief introduction).

Unit VII : Equilibrium

Equilibrium in physical and chemical processes, dynamic nature of equilibrium, law of mass action, equilibrium constant, factors affecting equilibrium- Le Chatelier's principle, ionic equilibrium- ionization of acids and bases, strong and weak electrolytes, degree of ionization, ionization of poly basic acids, acid strength, concept of pH, Henderson Equation, hydrolysis of salts (elementary idea), buffer solution, solubility product, common ion effect (with illustrative examples).

Unit VIII: Redox Reactions

Concept of oxidation and reduction, redox reactions, oxidation number, balancing redox reactions, in terms of loss and gain of electrons and change in oxidation number, applications of redox reactions.



06 Periods

14 Periods

Unit IX: Hydrogen

08 Periods

Position of hydrogen in periodic table, occurrence, isotopes, preparation, properties and uses of hydrogen, hydrides-ionic covalent and interstitial; physical and chemical properties of water, heavy water, hydrogen peroxide -preparation, reactions and structure and use; hydrogen as a fuel.

Unit X : s-Block Elements (Alkali and Alkaline Earth Metals) 10 Periods

Group 1 and Group 2 Elements General introduction, electronic configuration, occurrence, anomalous properties of the first element of each group, diagonal relationship, trends in the variation of properties (such as ionization enthalpy, atomic and ionic radii), trends in chemical reactivity with oxygen, water, hydrogen and halogens, uses. Preparation and Properties of Some Important Compounds: Sodium Carbonate, Sodium Chloride, Sodium Hydroxide and Sodium Hydrogencarbonate, Biological importance of Sodium and Potassium. Calcium Oxide and Calcium Carbonate and their industrial uses, biological importance of Magnesium and Calcium

Unit XI : *p*-Block Elements

18 Periods

General Introduction to *p*-Block Elements :

Group 13 Elements : General introduction, electronic configuration, occurrence, variation of properties, oxidation states, trends in chemical reactivity, anomalous properties of first element of the group, Boron - physical and chemical properties, some important compounds, Borax, Boric acid, Boron Hydrides, Aluminium: Reactions with acids and alkalies, uses.

Group 14 Elements : General introduction, electronic configuration, occurrence, variation of properties, oxidation states, trends in chemical reactivity, anomalous behaviour of first elements. Carbon-catenation, allotropic forms, physical and chemical properties; uses of some important compounds: oxides. Important compounds of Silicon and a few uses: Silicon Tetrachloride, Silicones, Silicates and Zeolites, their uses.

Group 15 Elements : General introduction, electronic configuration, occurrence, oxidation states, trends in physical and chemical properties; Nitrogen preparation properties and uses; compounds of Nitrogen, preparation and properties of Ammonia and Nitric Acid, Oxides of



Nitrogen(Structure only); Phosphorus - allotropic forms, compounds of Phosphorus: Preparation and Properties of Phosphine, Halides and Oxoacids (elementary idea only).

Unit XII : Organic Chemistry -Some Basic Principles and Technique 14 Periods

General introduction, methods of purification, qualitative and quantitative analysis, classification and IUPAC nomenclature of organic compounds. Electronic displacements in a covalent bond: inductive effect, electromeric effect, resonance and hyper conjugation. Homolytic and heterolytic fission of a covalent bond: free radicals, carbocations, carbanions, electrophiles and nucleophiles, types of organic reactions.

Unit XIII: Hydrocarbons

12 Periods

Classification of Hydrocarbons Aliphatic Hydrocarbons:

Alkanes - Nomenclature, isomerism, conformation (ethane only), physical properties, chemical reactions including free radical mechanism of halogenation, combustion and pyrolysis.

Alkenes - Nomenclature, structure of double bond (ethene), geometrical isomerism, physical properties, methods of preparation, chemical reactions: addition of hydrogen, halogen, water, hydrogen halides (Markownikov's addition and peroxide effect), ozonolysis, oxidation, mechanism of electrophilic addition.

Alkynes - Nomenclature, structure of triple bond (ethyne), physical properties, methods of preparation, chemical reactions: acidic character of alkynes, addition reaction of - hydrogen, halogens, hydrogen halides and water.

Aromatic Hydrocarbons: Introduction, IUPAC nomenclature, benzene: resonance, aromaticity, chemical properties: mechanism of electrophilic substitution. Nitration, sulphonation, halogenation, Friedel Craft's alkylation and acylation, directive influence of functional group in monosubstituted benzene. Carcinogenicity and toxicity.

Unit XIV : Environmental Chemistry

06 Periods

Environmental pollution - air, water and soil pollution, chemical reactions in atmosphere, smog, major atmospheric pollutants, acid rain, ozone and its reactions, effects of depletion of ozone layer, greenhouse effect and global warming- pollution due to industrial wastes, green chemistry as an alternative tool for reducing pollution, strategies for control of environmental pollution.



PRACTICALS

Evaluation Scheme for Examination	Marks
Volumetric Analysis	08
Salt Analysis	08
Content Based Experiment	06
Project Work	04
Class record and viva	04
Total	30

PRACTICAL SYLLABUS

Total Periods 60

Micro-chemical methods are available for several of teh practical experiments. Wherever possible such techniques should be used :

A. Basic Laboratory Techniques

- 1. Cutting glass tube and glass rod
- 2. Bending a glass tube
- 3. Drawing out a glass jet
- 4. Boring a cork

B. Characterization and Purification of Chemical Substances

- 1. Determination of melting point of an organic compound.
- 2. Determination of boiling point of an organic compound
- 3. Crystallization of impure sample of any one of the following: Alum, Copper Sulphate, Benzoic Acid.

C. Experiments based on pH

- (a) Any one of the following experiments :
 - Determination of pH of some solutions obtained from fruit juices, solution of known and varied concentrations of acids, bases and salts using pH paper or universal indicator.
 - Comparing the pH of solutions of strong and weak acids of same concentration.
 - Study the pH change in the titration of a strong base using universal indicator.

(b) Study the pH change by common-ion in case of weak acid and weak bases.

D. Chemical Equilibrium.



CHEMISTRY (Code No. 043) Question Paper Design

Class-XI (2020-21)

Time : 3 Hours

Max. Marks: 70

S. No.	Typology of Questions	Very short Answer (VSA) (1 marks)	Short Answer-I (SA–I) (2 marks)	Short Answer-II (SA-II) (3 marks)	Long Answer (LA) (5 marks)	Total Marks	% Weightage
1.	Remembering : Exhibit memory of previously learned material by recalling facts, terms, basic concepts and answers.	2	1	1	-	7	10%
2.	Understanding : Demonstrate understanding of facts and ideas by organizing, comparing, translating, inter- preting, giving descriptions and stating main ideas.	6	2	2	1	21	30%
3.	Applying : Solve problems to new situ- ations by applying acquired knowledge, facts, techniques and rules in a different way.	6	2	2	1	21	30%
4.	Analyzing : Examine and break information into parts by identifying motives or causes. Make inferences and find evidence to support gener- alizations.	6	1	2	-	14	20%
5.	Evaluating : Present and defend opinions by making judgments about information, validity of ideas or quality of work based on a set of criteria.	1	2	2	-	11	16%
6.	Creating : Compile information together in a different way by combing elements in a new pattern or proposing alternative solutions.	-	1	-	1	7	10%
	TOTAL	20×1=20	7×2=14	7×3=21	3×5=15	70(37)	100%



Type of Ques.	Mark per Ques.	Total No. of Ques.	Total Marks
VSA / Objective	1	20	20
SA-I	2	7	14
LA-I	3	7	21
LA-II	5	3	15
Total		37	70

Question Wise Break Up

1. No chapter was weightage. Care to be taken to cover all the chapters.

2. Suitable internal variations may be made for generating various templates keeping the overall weightage to different form of questions and typology of questions same.

Choice(s):

There will be no overall choice in the question paper.

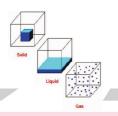
However, 33% internal choices will be given in all the sections.



Chemistry - XI Index

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Some Basic Concepts of Chemistry

FAST TRACK : QUICK REVISION

• Matter : Anything that has mass and occupies space.

Chapter - 1

- **Precision :** If refers to the closeness of various measurements for the same quantity.
- Accuracy : It refers to the agreement of a particular value to the true value of the result.
- Mass and weight : Mass of a substance is the amount of matter present in body, while weight is the force exerted by gravity on an object. The mass of a substance is constant whereas its weight may vary from one place to another due to change in gravity.
- Volume : $1 L = 1 dm^3 = 10^3 cm^3 = 10^{-3} m^3$

• Temperature :
$$K = {}^{\circ}C + 273.15; \frac{{}^{\circ}F - 32}{9} = \frac{{}^{\circ}C}{5}$$

- **Standard Temperature Pressure (STP) :** 0°C (273.15 K) temperature and 1 atm pressure.
- Normal Temperature Pressure (NTP) : 20°C (293.15 K) temperature and 1 atm pressure.
- Standard Ambient Temperature Pressure (SATP) : 25°C (298.15 K) temperature and 1 atm pressure
- Scientific Notation : Expressing a number in the form N × 10ⁿ, and N can vary between 1 to 9.99.
- **Significant figures :** These are meaningful digits which are known with certainty.
- Laws of Chemical Combination :
 - Law of Conservation of Mass (Antonie Lavoisier) : Mass can neither be created nor be destroyed.
 - Law of Definite Proportions (Joseph Proust) : A given compound always contains the same elements in the same proportion by mass.

- Law of Multiple Proportions (John Dalton) : When two elements combine to form two or more compounds, then the different masses of one element, which combine with a fixed mass of the other, bear a simple ratio to one another.
- Gay Lussac's Law : When gases combine or are produced in a chemical reaction, they do so in a simple ratio provided all gases are in the same temperature and pressure.

e.g.,
$$2H_2(g) + O_2(g) \rightarrow 2H_2O(g)$$

2 Vol 1 Vol 2 Vol
(at same T, P)

• Atomic Mass : It is defined as the average relative mass of an atom of an element as compared to the mass of an atom of carbon – 12 taken as 12.

Atomic mass is represented by 'u' (unified mass).

 $1u = 1.66056 \times 10^{-24} \,\mathrm{g}$

• **Molecular mass :** It is algebraic the sum of the atomic mass of the elements present in the molecule.

For example : Molecular mass of $CH_4 = (1 \times 12) + (4 \times 1) = 16 \text{ u}$

• Avogadro Number : It is the amount of atoms or molecules present in one mole of a substance.

Avogadro number (N_A) = $6.022 \times 10^{23} \text{ mol}^{-1}$

• **Molar Mass :** The mass of one mole of a substance in grams is called its molar mass.

For example : Molar mass of $CH_4 = (1 \times 12) + (4 \times 1) = 16g \text{ mol}^{-1}$

• Mole (*n*) : It is amount of a substance that contains as many particles or entities as the number of atoms in exactly 12 grams of pure C-12.

1 mole of a substance = Molar mass of substance = Avogadro's Number of chemical units = 22.4L volume at STP of gaseous substance

e.g., 1 mole of $CH_4 = 16g$ of $CH_4 = 6.022 \times 10^{23}$ molecules of $CH_4 = 22.4L$ at STP

$$n = \frac{wg}{M_m} = \frac{VL \text{ (at STP)}}{22.4L} = \frac{x \text{ particles}}{N_A} = \frac{MV}{1000}$$

• Molar Volume (V_m) : It is volume occupied by one mole of gas at STP. Molar volume of a gas = 22.4L at STP (273 K, 1atm) or 22.7L at STP (273 K, 1 bar)



Chemistry Class XI



Calculating Molar Volume: PV = nRT

:. V =
$$\frac{nRT}{P} = \frac{1 \text{mol x } 0.082 \text{L atm } \text{K}^{-1} \text{mol}^{-1} \text{ x } 273 \text{K}}{1 \text{ atm}} = 22.4 \text{L}$$

0r

$$\mathbf{V} = \frac{nRT}{P} = \frac{1 \text{ mol} \times 0.083 \text{ L} \text{ bar } \text{K}^{-1} \text{ mol}^{-1} \times 273 \text{ K}}{1 \text{ bar}} = 22.7 \text{ L}$$

• Percentage Composition : Mass % of the element

$$= \frac{\text{Mass of element in a molecule of the compound \times 100}}{\text{Molecular mass of the compound}}$$

- **Empirical Formula :** It represents the simplest whole number ratio of various atoms present in a compound. For *e.g.*, CH is the empirical formula of benzene.
- **Molecular Formula** : It shows the exact number of different of atoms present in a molecule of a compound. For *e.g.*, C₆H₆ is the molecular formula of benzene.
- **Relationship between empirical and molecular formulae :** Molecular formula = *n* × Empirical formula

Where; $n = \frac{\text{Molar mass}}{\text{Empirical formula mass}}$

• Information Conveyed by a chemical equation :

	N ₂ (g)	+	$3H_2(g)$	\rightarrow	2NH ₃ (g)
(i)	1 molecule of N_2	+	3 molecules of H_2	\rightarrow	2 molecules of NH_3
(ii)	1 mole of N ₂	+	3 mole of H_2	\rightarrow	2 mole of NH ₃
(iii)	$1 \times 28 \text{g of N}_2$	+	3×2 g of H ₂	\rightarrow	2×17 g of NH ₃
(iv)	1×22.4 L of N ₂	+	$3 \times 22.4 L$ of H_2	\rightarrow	2×22.4 L of NH ₃
	at STP		at STP		at STP

- Limiting Reagent : It is the reactant which gets consumed first or limits the amount of product formed.
- Mass Percent : It is the mass of the solute in grams per 100 grams of the solution.

Some Basic Concepts of Chemistry

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Mass percent= $\frac{\text{Mass of solute in } g \times 100}{\text{Mass of solution in } g}$

• **Parts per million (ppm) :** It is part of solute per million part of solution by mass.

 $ppm = \frac{Parts of solute (by mass) \times 10^{6}}{Parts of solution (by mass)}$

• Molarity (M) : It is number of moles of solute dissolved per litre (dm³) of the solution.

Molarity = $\frac{\text{No. of moles of solute}}{\text{Volume of solution in L}}$

Molarity equation : $M_1V_1 = M_2V_2$ (Before dilution) (After Dilution)

Molarity of a solution decreases on increasing temperature.

Molarity of pure water is $55.56 \text{ mol } \text{L}^{-1}$

• Molality (*m*)—It is number of moles of solute dissolved per 1000g (1kg) of solvent.

 $Molality = \frac{No. of moles of solute}{Mass of solvent in kg}$

Molality is independent of temperature.

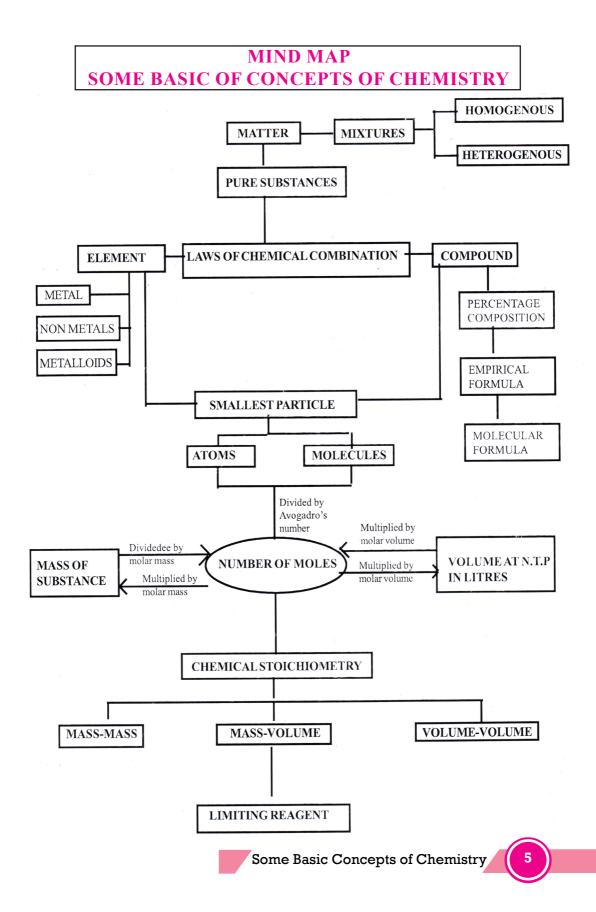
• Mole Fraction(x) is the ratio of number of moles of one component to the total number of moles (solute and solvents) present in the solution.

$$x_1 = \frac{n_1}{n_1 + n_2}$$
 and $x_2 = \frac{n_2}{n_1 + n_2}$

The sum of all the mole fractions in a solution is equal to one. *i.e.*, $x_1 + x_2 = 1$







		MULTIPLE CHOICE Q	UEST	FIONS (MCQ)
1.	Whic	h of the following is dependen	t of te	emperature ?
	(a)	Molarity	(b)	Molality
	(c)	Mole fration	(d)	Mass percentage
2.	4 g o	f NaOH dissolved in 100 ml so	lutior	n. Molarity of the solution is
	(a)	1 M	(b)	10 M
	(c)	0.1 M	(d)	4 M
3.	Whic	h has the maximum number of	mole	ecules among the following?
	(a)	44g of CO ₂	(b)	44g of O ₂
	(c)	8g of H ₂	(d)	64g of SO ₂
4.		ol of Zn react with 10 mol of H produced.	HC1. (Calculate the number of moles
	(a)	5 mol	(b)	10 mol
	(c)	20 mol	(d)	2.5 mol
5.	The r	number of oxygen atoms in 4.4	g of C	CO_2 is approximately
	(a)	1.2×10^{23}	(b)	6×10^{22}
	(c)	6×10^{23}	(d)	12×10^{23}
6.		nolarity of a solution obtained b nl of 2 M HCl will be	oy miz	xing 750 mL of 0.5 M HCl with
	(a)	0.975 M	(b)	0.875 M
	(c)	1.00 M	(d)	1.175 M
7.	Num	ber of atoms of He in 100 u of	He (/	Atomic mass of He is 4 u)
	(a)	25	(b)	50
	(c)	100	(d)	400
8.		$\times 10^{20}$ molecules of urea are pr entration of the solution is	resent	in 100 mL of its solution. The
	(a)	0.02 M	(b)	0.01 M
	(c)	0.001 M	(d)	0.1 M

Chemistry Class XI

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- 9. A gaseous hydrocarbons gives upon combustion, 0.72 g of water and 3.08 g of CO₂. The empirical formula of the hydrocarbon is :
 - (a) C_6H_5 (b) C_7H_8
 - (c) C_2H_4 (d) C_3H_4
- 10. The density of solution prepared by dissolving 120 g of urea (Mol. mass = 60 u) in 1000 g of water is 1.15 g/mL. The molarity of the solution is
 - (a) 0.50 M (b) 1.78 M
 - (c) 1.02 M (d) 2.05 M

Ans: 1. (a), 2. (a), 3. (b), 4. (a), 5. (a), 6. (b), 7. (a), 8. (b), 9. (b), 10. (d)

FILL IN THE BLANKS

1. 17 g of NH₃ gas will occupy a volume of _____ cm³ at NTP.

- 2. The number of Li atoms in ______ g. is 6.022×10^{24} atoms.
- 3. (1/12)th of the mass of carbon atom is _____

4. Number of atoms of oxygen in 24 g of O₃ is _____

- 5. The number of moles of barium carbonate which contains 1.5 moles of oxygen atoms is _____
- 6. A mixture having 2 g of H_2 and 32 g of oxygen occupies a volume of at NTP.
- 7. If the phosphate of a metal has the formula MPO₄ the formula of the metallic sulphate is _____
- 8. At NTP, the mass of 1 litre of gas is 3 g. Molecular mass of the gas is
- 9. The percentage mass of magnesium in chlorophyll is 2.68% The number of magnesium atoms in 2 g of chlorophyll is _____
- 10. The mass of one molecule of carbon dioxide is
- 11 Percentage of nitrogen in urea is
- 12 Number of carbon atoms present in 18 g of glucose ($C_6H_{12}O_6$)

Some Basic Concepts of Chemistry

13.	0.5 mole of triatomic gas contains					atoms.	
14.	A binary compound contains 50% A (at. mass = 16) and 50% B (at. mass 32). The empirical formula of the compound is						
15.	The number of hydrogen atoms in 60 u of ethane is						
Ans:	1. 22400 2. 70 g			70 g	3.	1 u	
	4.	9.033×10^{23}	5.	0.5	6.	44.8 litre	
	7.	$M_2(SO_4)_3$	8	67.2	9.	1.34×10^{21}	
	10.	7.3×10 ⁻²³	11.	46.67	12.	3.61×10 ²³	
	13.	9.033×10^{23}	14.	A ₂ B	15.	7.226×10^{24}	

TRUE AND FALSE TYPE QUESTIONS

Write true or false for the following statements

- 1. Equal volumes of different gases under similar conditions of temperature and pressure contain equal number of molecules.
- 2. 1 mole of $C_{12}H_{22}O_{11}$ contain 22 hydrogen atoms.
- 3. Nitrogen forms five oxides. It proves the law of multiple proportions.
- 4. The atomicity of phosphorus is four.
- 5 Molarity change with change in temp.
- 6. Empirical formula = $(Molecular formula)_n$.
- 7. Gram-atomic mass of an element may be defined as the mass of Avogadro's number of atoms.
- 8. Gay-Lussac's law of chemical combination is valid for all substances.
- 9. Avogadro's number varies with temperature and pressure.
- 10. 18 g of water vapour and 18 g of ice will contain the same number of molecules.

Ans:	1. (T)	2. (F)	3. (T)	4. (T)	5. (T)
	6. (F)	7. (T)	8. (F)	9. (F)	10. (T)



Chemistry Class XI



MATCH THE COLUMNS

1.

Column X

- 8 g CH_{4} a. 1.7 g NH₃ b.
- HCHO c.
- $C_{6}H_{12}O_{6}$ d.

2.

Column X

- Molarity a. Molality b.
- mole fraction c.
- d. ppm
- 3.

Column X

a.	40 g of He	i.	6.022×10^{23} atoms
b.	35 g of Li	ii.	10 atoms
c.	40 u of He	iii.	6.022×10^{24} atoms
d.	16 g of O ₂	iv.	3.011×10^{24} atoms

4.

Column X

- Petrol a. Graphite b.
- Sucrose c.
- d. Milk

1. a.(ii), b.(i), c.(iv), d.(iii) Ans: 3. a.(iii), b.(iv), c.(ii), d.(ii)

- 2. a.(iii), b.(iv), c.(ii), d.(i)
- 4. a.(iv), b.(iii), c.(ii), d.(i)

Some Basic Concepts of Chemistry

Column Y

- i. 0.1 mol
- ii. 0.5 mol
- iii. 40% carbon
- iv. Vapour density = 15

Column Y

- For very dilute solution i.
- ii. No units
- iii. Mol L^{-1}
- iv. independent of temperature

Column Y

- **Column Y**
- Heterogenous mixture i.
- Compound ii.
- iii. Element
- Hamogeneous mixture iv.
- 9

ASSERTION AND REASON TYPE QUESTIONS

Directions for Q. No.1-5

- A If both Assertion & Reason are true and the reason is the correct explanation of the assertion.
- B If both Assertion & Reason are true but the reason is not the correct explanation of the assertion.
- C If Assertion is true statement but Reason is false.
- D If both Assertion and Reason are false statements.
- Assertion : A solution of table salt in a glass of water is homogeneous
 Reason : A solution having same composition throughout is heterogeneous
- 2. Assertion : The molecular weight of oxygen is 32 amu.
 - Reason : The atomic weight of oxygen is 16 amu
- 3. Assertion : No of moles of H_2 in 0.224 L of hydrogen is 0.01 mole. Reason : 22.4 L of H_2 at STP contain 6.023 × 10²³ moles.
- 4. Assertion : Atomic mass of Na is 23.
- Reason : An atom of sodium is 23 times heavier than 1/12th mass of C-12 isotope.
- 5. Assertion : Number of atoms of He in 60 u of He is 15.
 - Reason : Atomic weight of He is 4 u.

Ans: 1.C 2.A 3.C 4.A 5.A

ONE WORD ANSWER TYPE QUESTIONS

- 1. What is the SI unit of density?
- 2. What is the SI unit of molarity?
- 3. Calculate the number of atoms in 32 u of He. [Ans. : 8]
- 4. What is the volume of 17 g of NH_3 gas at STP? [Ans. : 223.4 L]
- 5. How many molecules of SO_2 are present in 11.2 L at STP?

[Ans. : 3.011×10^{23}]

6. Which has more number of atoms ? 1.0 g Na or 1.0 g Mg

[Ans. : 1.0 g Na]



Chemistry Class XI



7. How any oxygen atoms are present in 16 g of ozone (O_3) ?

[Ans. : 2.007×10^{23}]

- 8. Calculate the number of molecules present in 22.0 g of CO_2 . [Ans. : 3.011×10^{23}]
- 9. A substance has molecular formula $C_6H_{12}O_6$. What is its empirical formula.
- 10. Empirical formula of a compound X (Molar mass = 78 mol^{-1}) is CH. Write its molecular formula.

1-MARK QUESTIONS

- 1. Name two chemical compounds used in treatment of cancer.
- 2. What is AZT ? Mention its use in medical science.
- 3. Classify following as pure substances and mixtures : air, glucose, gold, sodium and milk.
- 4. Which measurement is more precise 4.0g or 4.00g? [Ans. 4.00 g]
- 5. How many significant figures are there in (i) 3.070 and (ii) 0.0025 ?

[**Ans.** (i) 4 (ii) 2]

- 6. Express the following in the scientific notation : (i) 0.0048 (ii) 234,000
- 7. If ten volumes of dihydrogen gas react with five volumes of dioxygen gas, how much volume of water vapour would be produced ? [Ans. 10 volumes]
- 8. Define unified mass (u).
- 9. Define molar volume of a gas.
- 10. At STP, what will be the volume of 6.022×10^{23} molecules of H₂?

[Ans. 22.4L]

11. 1L of a gas at STP weighs 1.97g. What is molecular mass?

 $[Ans. 44.128 \text{ g mol}^{-1}]$

- 12. Write the relationship between empirical formula and molecular formula.
- 13. Which is more informative ? Empirical formula or Molecular formula.

Some Basic Concepts of Chemistry

- 14. How are 0.5 mol Na₂CO₃ and 0.5 M Na₂CO₃ different from each other ?
- 15. Why molality is preferred over molarity of a solution ?
- 16. Define molarity of a solution.
- 17. What is the effect of temperature on molarity of solution ?
- 18. What is limiting reactant in a reaction ?

2-MARKS QUESTIONS

- 1. Classify following substances as element, compounds and mixtures : water, tea, silver, steel, carbon dioxide and platinum.
- 2. The body temperature of a normal healthy person is 37°C. Calculate its value in°F.
- 3. At what temperature will both the Celsius and Fahrenheit scales read the same value?
- 4. Convert 5L into m^3 .
- 5. What does the following prefixes stand for :(a) pico (b) nano (c) micro (d) deci
- 6. How many significant figures are present in the following :
 - (i) 4.00005
 - (ii) 0.004
- 7. Convert '450 pm' into SI unit and write the answer in scientific notation upto 2 significant figures.

[Ans. 4.5×10^{-10} m]

- 8. Hydrogen peroxide and water contain 5.93% and 11.2 % of hydrogen respectively. Show that the data illustrate law of multiple proportions.
- 9. The density (in g mL⁻¹) of a 3.60 M sulphuric acid solution that is 29% H_2SO_4 (Molar mass = 98 g mol⁻¹) by mass will be

[**Ans.** 1.21 g/mL]

- 10 The cost of table salt (NaCl) is Rs. 10 per Kg. calculate its cost per mole.
 (Molar mass of NaCl is 58.5 gmol⁻¹) [Ans. 0.58 Rs]
- 11 Calculate the mole fraction of the solute in a 1.00 molal aqueous solution. [Ans. 0.0177]



Chemistry Class XI



- Dissolving 120 g of urea (Molar mass of urea = 60 g mol⁻¹) in 1000 of water gave a solution of density 1.15 g/mL. Calculate the molarity of the solution. [Ans. 2.05 M]
- 13 Calculate the percentage of N in urea. (Molar mass of urea = 60 g mol⁻¹)[Ans. 46.66]
- 14 25 ml of 3.0 M HCl are mixed with 75 mL 0f 4.0 M HCl. If the volumes are additive, the molarity of the final mixture will be. [Ans. 3.75 M]
- 15 How many atoms and molecules are present in 124 gm of phosphorus (P4) [Ans. Atoms = $4 N_A \& Molecules = N_A$]
- 45.4 L of dinitrogen reacted with 22.7 L of dioxygen and 45.4 L of nitrous oxide was formed.

The reaction is given below : $2N_2(g) + O_2(g) \longrightarrow 2N_2O(g)$

Which law is being obeyed in this experiment? Write the statement of the law.

17 Give one example each of a molecule in which empirical formula and molecular formula is

(i) Same (ii) Different.

- 18 Calculate the number of moles in the following masses :
 - (i) 7.85g of Fe;
 - (ii) 7.9mg of Ca
- 19 Calculate the percent of carbon, hydrogen and oxygen in ethanol (C_2H_5OH) [Ans. 52.14%, 13.13%, 34.73%]
- 20 How much copper can be obtained from 100 g of CuSO₄? [Ans. 39.8g]
- 21 Calculate the amount of water (g) produced by the combustion of 16 g of methane. [Ans. 36g]
- 22 How many moles of methane are required to produce 22 g CO₂ (g) after combustion? [Ans. 0.5 mol]
- A solution is prepared by adding 2 g of a substance A to 18 g of water.Calculate the mass per cent of the solute. [Ans. 10%]
- 24 Calculate molarity of water if its density is 1.00 g mL⁻¹.

[Ans. 55.56 M]

Some Basic Concepts of Chemistry

- 25 Calculate the molarity of NaOH in the solution prepared by dissolving its 4 g in enough water to form 250 mL of the solution. [Ans. 0.4 M]
- 26 The density of 3 M solution of NaCl is 1.25 g mL⁻¹. Calculate molality of the solution. [Ans. 2.8m]
- 27 NH₃ gas can be prepared by Haber's process as, $N_2(g) + 3H_2(g) \rightarrow 2NH_3(g)$. At a particular moment concentration of all the species is 2 moles; calculate the concentration of N₂ and H₂ taken initially.

[Ans. 3 mole, 5 moles]

3-MARKS QUESTIONS

1. Calculate the average atomic mass of Mg using the followinf data:

	% Natural Abudance	Molar mass		
24^{Mg}	80	24		
25^{Mg}	10	25		
26^{Mg}	10	26		

2. The following data are obtained when dinitrogen and dioxygen react together to form different compounds :

	(i)	(ii)	(iii)	(iv)
Mass of dinitrogen	14	14	28	28
Mass of dioxygen	16	32	32	80

Which law of chemical combination is obeyed by the above experimental data ? Give its statement.

- 3. Calculate :
 - (i) Mass in gram of 5.8 mol N_2O
 - (ii) Number of moles in 8.0 g of O_2
 - (iii) Molar mass if 11.2 L at STP weigh 8.5 g.

[Ans. (i) 255.2 g (ii) 0.25 mol (iii) 17 g mol⁻¹]

- 4. In three moles of ethane (C_2H_6) , calculate the following :
 - (i) Number of moles of carbon atom,
 - (ii) Number of moles of hydrogen atoms,
 - (iii) Number of molecules of ethane.

[Ans. (i) 6 moles, (ii) 18 moles, (iii) 1.81×10²⁴]



Chemistry Class XI



- 5. 16 g of an ideal gas SOx occupies 5.6 L at STP. What is its molecular mass ? What is the value of X ? [Ans. 64u, x = 2]
- 6. Calculate the number of moles :
 - (i) 5.0 L of 0.75 M Na₂CO₃
 (ii) 7.85 g of Fe
 - (iii) 34.2 g of sucrose $(C_{12}H_{22}O_{11})$

[Ans. (i) 3.75, (ii) 0.14, (iii) 0.1]

7. Calculate the number of atoms in each of the following :

(i) 52 moles of Ar. (ii) 52u of He (iii) 52g of He.

[**Ans.** (i) 3.13 × 1025 (ii) 13 (iii) 7.83 × 1024]

- 8. Vitamin C is essential for the prevention of scurvy. Combustion of 0.2000g of vitamin C gives 0.2998g of CO_2 and 0.819g of H_2O . What is the empirical formula of vitamin C? [Ans. $C_3H_4O_3$]
- 9. A compound contains 4.07% hydrogen, 24.27% carbon and 71.65% chlorine. Its molar mass is 98.96 g. What are its empirical and molecular formulas? [Ans. CH₂C1, C₂H₄Cl₂]
- 10. A compound made up of two elements A and B has A = 70%, B = 30%. Their relative number of moles in the compound is 1.25 and 1.88, calculate :
 - (i) Atomic masses of the elements A and B
 - (ii) Molecular formula of the compound , if its molecular mass is found to be 160.
 [Ans. (i) 56 and 16, (ii) A₂B₃]
- 11. The reaction $2C + O_2 \longrightarrow 2CO$ is carried out by taking 24.0 g of carbon and 96.0 g of O_2 . Find out.
 - (i) Which reactant is left in excess ?
 - (ii) How much of it is left?
 - (iii) How many grams of the other reactant should be taken so that nothing is left at the end of the reaction ? [Ans. (i) O₂, (ii) 64 g, (iii) 72]
- 12. A 10 g sample of a mixture of calcium chloride and sodium chloride is treated with Na_2CO_3 to precipitate calcium as calcium carbonate. This $CaCO_3$ is heated to convert all the calcium to CaO and the final mass of CaO is 1.62 g. Calculate % by mass of NaCl in original solution.

[**Ans.** 67.9%]



- 13. 3.0 g of H_2 react with 29.0 g of O_2 yield H_2O .
 - (i) Which is the limiting reagent.
 - (ii) Calculate the maximum amount of H₂O that can be formed
 - (iii) Calculate the amount of reactant left unreacted

$$[\mathbf{Ans.}~\mathbf{H}_2$$
 , 26.8g $\mathbf{H}_2\mathbf{O}$ & 5.2 g $\mathbf{O}_2]$

14 Zinc and hydrochloric acid react according to the reaction:

 $Zn(s) + 2HCl(aq) \longrightarrow ZnCl_2(aq) + H_2(g)$

If 0.30 mol Zn are added to hydrochloric acid containing 0.52 mol of HCl, How many moles of H_2 are produced ?

[HCl is limiting reagent; H_2 formed = 0.36 mol]

- How many moles of Lead (II) chloride will be formed from a reaction between 6.5 g of PbO and 3.2 g of HCl? [Atomic mass of Pb = 207 U][Ans. 0.029 mole]
- 16 What volume of oxygen at N.T.P is needed to cause the complete combustion of 200 ml of acetylene ?Also calculate the volume of carbon dioxide formed. [Ans. 500 mL of $O_2 & 400 \text{ mL of } CO_2$]

5-MARKS QUESTIONS

- (i) A black dot used as a full stop at the end of a sentence has a mass of about one attogram. Assuming that the dot is made up of carbon, calculate the approximate number of carbon atoms present in the dot.
 [Hint : 1 attogram = 10⁻¹⁸g]
 - (ii) Which one of the following will have largest number of atoms ?
 - (a) 1g Au (s) (b) 1g Na (s) (c) 1g Li (s) (d) 1g of Cl₂(g) [**Ans.** (i) 39.81 g (ii) 1 g of Li]
- 2. (i) What is the difference between empirical formula and molecular formula ?
 - (ii) A welding fuel gas contains carbon and hydrogen only. Burning a small sample of it in oxygen gas 3.38 g carbon dioxide, 0.690 g of water and no other products. A volume of 10.0 L (measured at STP) of this welding gas is found to weigh 11.6 g. Calcuate
 - (i) Empirical formula, (ii) molar mass of the gas, and (iii) Molecular formula.
 [Ans. (i) CH, (ii) 26 g mol⁻¹, (iii) C₂H₂]





- 3. (i) What is the difference between Molarity and Molality.
 - (ii) The Molarity of a solution of sulphuric acid is 1.35 M. Calculate its molality. (The density of acid solution is 1.02 g cm⁻³).

[Ans.. 1.52 m]

- 4. (i) Define : (a) Mole fraction (b) Mass percentage.
 - (ii) If the density of methanol is 0.793 kg L⁻¹, what is its volume needed for making 2.5 L of its 0.25 M solution ? [Ans. 0.0025 L]

HOTS QUESTIONS

1 In a compound $C_x H_y O_z$, the mass % of C and H is 6:1 and the amount of oxygen present is equal to the half of the oxygen required to react completely $C_x H_y$. Find the empirical formula of the compound.

 $[Ans. C_2H_4O_3]$

2 A crystalline salt when heated becomes anhydrous and loses 51.2 % of its weight. The anhydrous salt on analysis gave the following percentage composition

Mg = 20.0%, S = 26.6%, O = 53.33%

Calculate the molecular formula of the anhydrous salt and the crystalline salt. Molecular weight of the anhydrous salt is 120.

[Ans. MgSO₄.7H₂O]

- 3 An LPG cylinder weighs 14.8 Kg when empty. When full, it weighs 29.0 kg and shows a pressure of 2.5 atm. In the course of use at 27°C, the weight of cylinder is reduced to 23.2 Kg. Find the volume of n-butane in cubic meters used up at 27°C and 1 atm (Molecular weight of n-butane = 58).
 [Ans. 2.463 m³]
- 4 2.5 g of CaCO_3 was placed in 50 ml of a solution of HCl.1.05 g of CaCO₃ was left after the reaction. Calculate:
 - (a) the weight of HCl per litre
 - (b) the Molarity of HCl [Ans. (a) 21.17 g, (b) 0.58 M]

UNIT TEST

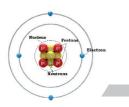
Maximum Marks : 20

Time allowed : 1 hour

General instructions : (i) All questions are compulsory. (ii) Maximum marks carried by each question are indicated against it. 1. If 30 mL of H_2 and 20 mL of O_2 react to form water, what is left at (1)the end of the reaction? (a) 10 mL of H_{2} (b) 5 mL of H_2 (d) $5 \text{ mL of } O_2$ (c) $10 \text{ mL of } O_2$ 7.5 grams of a gas occupy 5.6 litres of volume at STP the gas is 2. (1)(b) N₂O (c) CO (a) NO (d) CO_{2} What is AZT? Write its use. 3. (1)Why molarity is preferred over molarity in expressing the concen-4. (1)tration of solution? Which has more number of atoms ? 1.0 g Na or 1.0g Mg? 5. (1)How many atoms and molecules are present in 124 g of phosphorus (P_4)? (2) 6. (a) Write the name of two life saving drugs. 7. (2)(b) Define accuracy and precision. A sample of drinking water was found to be severely contaminated with 8. chloroform CHCl₂. The level of contamination was 15 ppm (by mass). (a) Express this in percent by mass. (b) Determine the molarity of chloroform in the water sample. (3)A compound contains 4.07% hydrogen, 24.27% carbon and 71.65% 9. chlorine. Its molar mass is 98.96 g. What are its empirical and molecular formula? (3)10. (a) Explain the following terms: (i) Gay Lussac's law (ii) Limiting reagent (b) 3.0 g of H_2 react with 30.0 g of O_2 yield H₂O. (i) Which is the limiting reagent? (ii) Calculate the maximum amount of H₂O that can be formed. (iii) Calculate the amount of reactant left unreacted. (5) ****







Chapter - 2

Structure of Atom

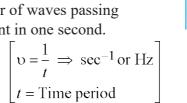
FAST TRACK : QUICK REVISION

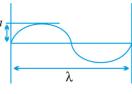
• Information about fundamental particles of atom

Name of Constant	UNIT	Electron	Proton	Neutron
Mass	amu	0.000546	1.00728	1.008665
	kg	9.109×10^{-31}	1.673×10^{-27}	1.675×10^{-27}
Charge	Coloumbs	-1.602×10^{-19}	$+ 1.602 \times 10^{-19}$	Zero
	esu	-4.8×10^{-10}	$+ 4.8 \times 10^{-10}$	Zero
	Relative Charge	- 1	+ 1	Zero

- **Electromagnetic radiations :** Energy emitted from any source (in forms of waves) in which electric and magnetic fields oscillated perpendicular to each other and travelling with a velocity of light is known as EM radiation.
- Characteristics of waves :
 - (a) Wavelength : The distance of one crest and one trough in a wave. Denoted by ' λ '.
 - (b) Frequency : Number of waves passing through a given point in one second.

Denoted by v.





- (c) Amplitude : The height of crest or depth of a trough denoted by 'a'.
- (d) Wave Number : Number of waves per unit length denoted by $\overline{\upsilon}$

$$\overline{\upsilon} = \frac{1}{\lambda} = \mathrm{cm}^{-1} \; (\mathrm{or} \; \mathrm{m}^{-1})$$

(e) Velocity : Linear distance travelled by a wave in one second.

velocity of light
$$c = \frac{\text{Distance}}{\text{Time}} = \lambda \times \frac{1}{t} = \upsilon \lambda$$

 $\therefore \quad \upsilon = \frac{c}{\lambda}$

• Energywise order for EM radiation.

 $cosmic < \gamma \, rays < X \, rays < UV < VIBGYOR < IR < Microwaves < Radiowaves$

λ (Increases) υ (Decreases) Energy (Decreases)

- **Photon :** A packet or particle of light energy is knows as **Photon.**
- **Planck's quantum theory :** The energy emitted or absorbed by a source is discontinuous in form of small packet of energy, called **quantum**. Quantum of light is called **photon**.

E
$$\alpha v$$

E = hv (h = Planck's constant)
E = nhv (h = 6.626×10⁻³⁴ J sec)
If ' n ' photons are emitted E = nhv

• **Photo electric effect :** The phenomenon of ejection of electrons from a metal surface when a light of suitable frequency falls on metal surface.

$$h\upsilon - h\upsilon_0 = \frac{1}{2} \mathrm{mv}^2$$

 $h\upsilon \Rightarrow$ Energy of incident light on metal surface.

 $hv_0 \Rightarrow$ Work function of metal.

 $\frac{1}{2}$ mv² = Kinetic energy by which e^{-} is emitted from metal surface.

• **de Broglie equation :** All material particles in motion also exhibit wave like properties.

$$\lambda = \frac{h}{mv} = \frac{h}{p}$$

For microscopic particles mass is very less therefore Wavelength of wave associated with it can be detected.



For macroscopic particles mass is large, λ of wave associated with it can not be detected. Hence dominant wave character.

Hence microscopic bodies have dual nature, where as macroscopic bodies have particle nature.

Heisenberg's Uncertainty Principle

It is impossible to determine the exact position and velocity of a moving subatomic particle simultaneously with accuracy.

$$\Delta x \times m \Delta v \geq \frac{h}{4\pi}$$

 Δx = uncertainty in position

 $\Delta v =$ uncertainty in velocity

Bohr's theory for H [H like one e⁻ systems He⁺; Li²⁺]

(1) e^- revolving round the nucleus in circular path [stationary state; SHELL] with a definite angular momentum $\frac{nh}{2\pi}$ [Here $n = \text{no. of shell of } e^-$] and with definite energy

$$\mathbf{E}_{\mathbf{n}} = \left[\frac{-2\pi^2 m e^4 z^2}{n^2 h^2}\right] \Longrightarrow -2.18 \times 10^{-18} \frac{\mathbf{Z}^2}{n^2} \,\mathrm{J/Atom}.$$

(2) As *n* increases, Energy of e^- becomes less – ve [Due to less force of Proton attraction]

As *n* decreases, Energy of e^- becomes More – ve [Due to more force of attraction by protons]

- (3) In infinity shell e^- has zero force of attraction therefore zero energy.
- (4) Electron energy only changes by definite values $\Delta E = E_f E_{i^*}$.

Hydrogen spectrum : When e^- in hydrogen atom is provided energy it gets excited to higher shell from ground state, it comes back to ground state by emitting energy in definite values.

"Quanta": The emission of light energy is known as emission spectra. It corresponds to each atom depending upon which energy shell e^- is excited.



It is **discontinuous** spectra as ' λ ' of light radiations do not merge with each other like in VIBGYOR (Continous Spectra).

When e^{-} falls from any excited state to

$\frac{1}{\lambda} = 1,09,678 \left[\frac{1}{n_f^2} - \frac{1}{n_i^2} \right]$	$\int Z^2 R = Rydberg constant = 109678 cm^{-1}$
$n_i = 1, n_f = 2, 3, 4, \dots$	[Lyman series] (UV)
$n_i = 2, n_f = 3, 4, 5, \dots$	[Balmer series] (VIBGYOR)
$n_i = 3, n_f = 4, 5, 6$	[Paschen series] IR.
$n_i = 4, n_f = 5, 6, 7$	[Bracket series] IR.
$n_i = 5, n_f = 6, 7, 8$	[Pfund series] IR.

Quantum numbers : The noumbers which completely define the state of e^{-} in an atom.

- (1) Principal Quantum No. : It describes the distance of *e*⁻ from nucleus '*n*' *i.e.*, defines the **shell** no. It is denoted by '*n*'.
 - п = 1, 2, 3, 4, 5, K, L, M, N, O

(2) Azimuthal (1) Quantum No. : It defines the path of e^- decided by angular momentum of e^- . Each angular momentum value corresponds to one subshell. The no. of subshells in a shell is 0 to n - 1.

п	<i>l</i> (0 to <i>n</i> - <i>l</i>)			
1	0	l = 0	<i>'s</i> '	subshell 🔵
2	0, 1	l = 1	<i>`p</i> '	subshell 🔶
3	0, 1, 2	l = 2	ʻd'	subshell 🔶
4	0, 1, 2, 3	l = 3	f'	subshell 🤸

All subshells are wave functions for locating e^{-} . In the same shell energy increase s .



- (3) Magnetic Quantum No. : It gives the no. of magnetic orientations an e^- can have in a subshell. That is number of orbitals in a sub-shell. $m_s = -l.....0.....+ l = (2l + 1).$
- (4) Spin Quantum No. : An e^- is continuously spinning on its own axis.

The value of $s = \frac{1}{2}$ or $-\frac{1}{2}$

An orbital can have maximum two e^- one with clockwise and other with anticlockwise spin.

Aufbau principle

- (a) Electrons are filled in increasing order of energy of sub-shell.
- (b) As 'n + l'value increases energy of e^{-} increases in that sub-shell.
- (c) For two sub-shells with same n + l value, as n value increases energy of e^{-1} increases.

Pauli's principle

No two electrons can have same set of four quantum numbers in an atom.

Hund's rule of maximum multiplicity

The pairing of e^- in degenerate orbitals (different orbitals with same energy) will get paired only once they have been singly occupied with same spin.

IMPORTANT POINTS

The filling of e^- in subshells follows this order. (As per Aufbau principle)

- (A) 1s < 2s < 2p < 3s < 3p < 4s < 3d < 4p < 5s < 4d < 5p < 6s < 4f < 5d < 6p < 7s < 5f < 6d < 7p
- (B) Half filled and completely filled subshells have more **stability** than incompletely filled subshells.

$$Cr = [Ar] 4s^1 3d^5$$

 $Cu = [Ar] 4s^1 3d^{10}$

(C) As the shell no. inc. size of subshell increases *e.g.*, size of (2s > 1s); (3p > 2p); (4d > 3d)



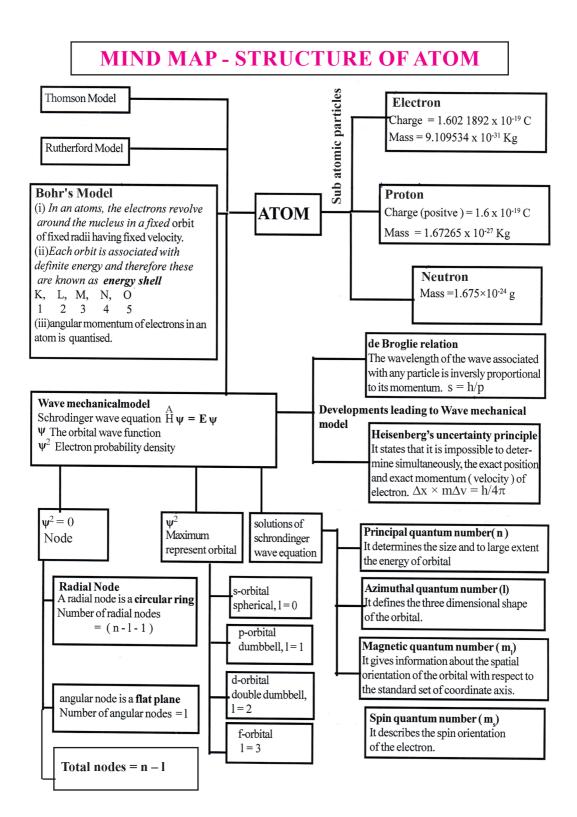
(D) The region in an orbital where probability of finding the e^- is zero is known as **Nodal plane** (or Node).

The no. of [radial nodes] = n - l - 1 and Angular Nodes = l, Total nodes = n-1.

(E)	ψ (psi)	ψ ² (psi square)
	A wave function for locating an electron	The square of wave function where the probability of finding the e^- is maximum. [Each value of ψ^2 is a region and defines one orbital]
(F)	Orbit	Orbital
	(1) A definite distance from the nucleus for finding the $e^{-}[e^{-}$ as a particle].	 (1) A probability region for locating the e⁻ around the nucleus. It is a wave function [e⁻ as a wave]
	 (1) It has definite size and e⁻ in this orbit has definite energy. 	 (2) It does not define definite size. But only a boundary region diagram of a wave for locating the <i>e</i>⁻.









MULTIPLE CHOICE QUESTIONS (MCQ)

				(-/
1.	Pack	ket of energy is called		
	(a)	Electron	(b)	Photon
	(c)	Position	(d)	Proton
2.	Orbi	ital which is not possible		
	(a)	2p	(b)	3d
	(c)	3s	(d)	3f
3.	the r	magnetic quantum number of a	an ato	m is related to the
	(a)	size of the orbital	(b)	spin angular momentum
	(c)	orbital angular momentum	(d)	orientation of the orbital in space
4.	The	principal quantum number of	an ato	om is related to the
	(a)	size of the orbital	(b)	spin angular momentum
	(c)	orbital angular momentum	(d)	orientation of the orbital in Spence
5.	The	designation of an orbital with	in =	4 and $1 = 3$
	(a)	4s	(b)	4p
	(c)	4d	(d)	4f
6.		at transition in the hydrogen space R at transition $n = 4$ to n		n would have the same wavelength n the He ⁺ spectrum?
	(a)	n = 4 to $n = 1$	(b)	n = 3 to $n = 2$
	(c)	n = 3 to $n = 1$	(d)	n = 2 to $n = 1$
7.		wave number of first line of B wave number of the first Baln		series of hydrogen in 15200 cm-1. le of Li^{2+} ion is
	(a)	15200 cm ⁻¹	(b)	60800 cm ⁻¹
	(c)	76000 cm ⁻¹	(d)	136,800 cm ⁻¹
8.	An e	electron is moving in Bohr's or		s de Broglie wavelength is λ . What

is the circumference of the forth orbit?

(a) $2/\lambda$ (b) 2λ

(c) 3λ (d) $3/\lambda$

Chemistry Class XI

- 9. Which of the following statements in relation to the hydrogen atom is correct?
 - (a) 3s-orbital is lower in energy than 3p-orbital
 - (b) 3p-orbital is lower in energy than 3-d-orbital
 - (c) 3s and 3p orbitals all have the same energy.
 - (d) 3s, 3p and 3d orbitals all have the same energy.
- 10. For principle quantum number, n = 4, the total number of orbitals having 1 = 3 is
 - (a) 3 (b) 7
 - (c) 5 (d) 9

11. The number of d-electrons retained in Fe^{2+} (At. no. of Fe = 26) ion is

- (a) 3 (b) 4
- (c) 5 (d) 6

12. Pauli exclusion principle helps to calculate the maximum number of electrons that can be accommodated in any

- (a) orbital (b) subsell
- (c) shell (d) All of these

Ans. 1. (b), 2. (d), 3. (d), 4. (a), 5. (d), 6. (d), 7. (d), 8. (c), 9. (d),

10. (b), 11. (d), 12. (a)

FILL IN THE BLANK

- 1. Bohr's theory is based on ______ of radiation.
- 2. The angular momentum of the electron in the 4th energy shell in the hydrogen atom is _____.
- 3. Lines of Balmer series appear in _____ region.
- 4. The maximum number of electrons in Fe^{3+} (At. No. 26) is _____.
- 5. Li^{2+} and He^+ ions have spectrum similar to _____ atom.
- 6. Bohr's atomic theory is not able to explain the atomic spectra of atoms containing ______ electron.
- 7. An electron in the first shell will have ______ stability and ______ energy than an electron in the third shell.

- 8. The space or three-dimensional region round the nucleus where there is maximum probability of finding an electron of specific energy is called an_____
- 9. According to _____ no two electrons in an atom will have all the four quantum numbers _____
- 10. When there are two electrons in the same orbital they have _____ spins.
- 11. The s-subhells have _____ shape and the p-subshells have _____
- 12. The maximum number of electrons on a subshell is equal to _____ where $l = ____$

Ans. 1. Planck's theory 2. $\frac{2h}{\pi}$

- 3. Visible 4. 23
- 5. H-atom 6. more than 1
- 7. Larger, lower 8. orbital
- 9. Pauli exclusion principle; similar
- 10. Opposite
- 11 Spherical, dumb bell shape.
- 12. 21 + 1; azimuthal quantum numbers

TRUE AND FALSE TYPE QUESTIONS

Write true or false for the following statements

- 1. Bohr's theory cannot explain the spectra of multi-electron atoms.
- 2. Bohr's theory based on the Planck's quantum theory.
- 3. Size of orbital is determined by principal quantum number.
- 4. Fe^{2+} ion has more number of unpaired electrons than Fe^{3+} .
- 5. The outer electronic configuration of chromium atom is $3d^44s^2$.
- 6. The designation of an orbital n=4 and l=0 is 4s.
- 7. All photons of light have same energy.
- 8. Fe^{3+} has $3d^5$ configuration.





- 9. The number of subshells is always equal to the order of the orbit.
- 10. Two electrons in the same orbital has antiparallel spin.
- 11. The second orbit in He^+ ion has radius as the first orbit in hydrogen atom.
- 12. Heisenberg principle is applicable to microscopic particles.
- 13. 3s orbital has 2 radial nodes.

Ans. 1. (T)	2. (T)	3. (T)	4. (F)	5. (F)	6. (T)	7. (F)
8. (T)	9. (F)	10. (T)	11. (T)	12. (T),	13. (T)	

MATCH THE COLUMNS

1. Match the following

List-I

- a. Lyman series
- b. Balmer series
- c. Paschen series
- d. Brackett series
- 2. Match the following

List-I

- a. Principal quantum number
- b. Azimuthal quantum number
- c. Magnetic quantum number
- d. Spin quantum number
- 3. Match the following

List-I

- a. 2s
- b. 2p_x
- c. 3d_{xy}
- d. 3d_z²

List-II

- p. Visible region
- q. Infrared region
- r. Absorption spectrum
- s. Ultraviolet region

List-II

- p. Spin of electrons
- q. Size of orbital
- r. Orientation of the orbital
- s. Shape of the orbital

List-II

- p. Dough not shape
- q. Spherical
- r. Dumb bell
- s. Double dumb bell

Structure of Atom

4. Match the following

List-I

- a. 2s
- b. ψ^2
- c. Heisenberg's uncertainty
- d. 3d_{yz}

List-II

- p. Two nodal planes
- q. One radial node
- r. Electron probability density principle
- s. Microscopic particles

ASSERTION AND REASON TYPE QUESTIONS

Directions: (Questions 1 to 4)

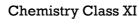
- A. If both Assertion & Reason are true and the reason is the correct explanation of the assertion.
- B. If both Assertion & Reason are true but the reason is not the correct explanation of the assertion.
- C. If Assertion is true statement but Reason is false.
- D. If both Assertion and Reason are false statements.
- 1. Assertion : Number of orbitals in 3rd shell is 9. Reason : Number of orbitals for a particular value of $n = n^2$.
- 2. Assertion : Two nodal planes are present in $3d_{xy}$. Reason : Number of nodal planes = 1
- 3. **Assertion :** The energy of an electron is largely determined by its principal quantum number.

Reason : The principal quantum number is a measure of the most probable distance of finding the electrons around the nucleus.

4. **Assertion :** An orbital cannot have more than two electrons, moreover, if an orbital has two electrons they must have opposite spins.

Reason : No two electrons in an atom can have same set of all the four quantum numbers.

Ans. 1. A 2. A 3. A 4. A





ONE WORD ANSWER TYPE QUESTIONS

- 1. Write the name of the theory which explain the wave nature of light.
- 2 Write the name of the theory which explain the Black body radiations and photo electric effect
- 3 If the length of the crest of a wave is 4 pm. Write the wavelength of this wave. [Ans.8 pm]
- 4. A radiation emitted from a hot iron is photon or quantum ?
- 5. Out of the d orbitals which does not have four lobes ?
- 6. What is the lowest value of n that allows g orbitals to exist ?
- 7. Which quantum number is not obtained from solution of Schrödinger wave equation ?
- Which of the following orbitals are possible ?
 1p, 2s, 2p and 3f
- 9. Write the name of non-directional subshell.
- 10. Write the name of quantum number which determines the orientation of orbitals ?
- 11. Write the name of quantum number which determines the shape of orbitals.
- 12. How many orbitals are present in 'g' subshell ?

1-MARK QUESTIONS

- 1 Write the relation between frequency and wave number.
- 2 Cs shows maximum photoelectric effect, why?
- 3 Distinguish between a photon and a quantum.
- 4 The line spectrum of an element is known as fingerprints of its atom. Comment.
- 5 What is the value of the Bohr's radius for the third orbit of hydrogen atom?
- 6 What type of metals are used in photoelectric cell ? Give one example. [Ans. With large size, less work function.]
- 7 Which series of lines of the hydrogen spectrum lie in the visible region'?
- 8 Mention the physical significance of ψ and ψ^2 .
- 9 Why did Heisenberg's uncertainty principle replace the concept of definite orbit by the concept of probability?





- 10. What is uncertain in uncertainty principle ?
- 11. Can a moving cricket ball have a wave character ? Justify your answer.
- 12. Heisenberg uncertainty principle has no significance in our everyday life. Explain.
- 13. Write the Schrodinger wave equation.
- 14. Why uncertainty in position is more when uncertainty in velocity is less for an electron ?
- 15. What are the four quantum numbers of 19th electron of copper ?(Given : Atomic number of copper = 29)
- 16. How many electrons will be present in the sub-shells having ms, value of -1/2 for n = 4 ?
- 17. Write the electronic configuration of Ni^{3+} . (At. No. of Ni = 28)
- 18. How many radial and angular nodes are present in 2p orbital.

[**Ans.** Radial nodes = 0, Angular nodes = 1]

2-MARKS QUESTIONS

- **Q. 1.** Define black body and black body radiations.
- **Q. 2.** Give the essential postulates of Bohr's model of an atom. How did it explain?
 - (i) the stability of the atom ?
 - (ii) origin of the spectral lines in H-atom ?
- **Q.3.** What is quantisation ? How quantisation of energy was introduced in Bohr's model ?
- **Q. 4.** What transition in the hydrogen spectrum would have the same wavelength as the Balmer transition n = 4 to n = 2 of He⁺spectrum?

[Ans.
$$n_1 = 1$$
 and $n_2 = 2$]

Q. 5. What transition of Li²⁺ spectrum will have the same wavelength as that of the second line of Balmer series in He⁺spectrum ?

[**Ans.** $n_2 = 6$ to $n_1 = 3$]

Q. 6. Calculate the energy required for the process

 $\operatorname{He}^+(g) \longrightarrow \operatorname{He}^{2+}(g) + e^-$

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The ionization energy for the H atom in the ground state is $2.18 \times 10^{-18} \text{ J atom}^{-1}$] [Ans. $8.72 \times 10^{-18} \text{ J}$]

- **Q. 7.** Calculate the wave number for the longest wavelength transition in the Balmer series of atomic hydrogen. [Ans. $1.523 \times 10^6 \text{ m}^{-1}$]
- Q. 8. To which orbit the electron in H atom will jump on absorbing 12.1 eV energy ? [Ans. 3rd orbit]
- Q. 9. Calculate the energy associated with the first orbit of He⁺. What is the radius of this orbit? [Ans. 54.38 eV, 0.2645 Å]
- Q. 10. What is the distance of separation between 3rd and 4th orbit of H-atom? [Ans. 3.703 Å]
- **Q. 11.** The energy of electron in the first Bohr's orbit is -13.6 eV. Calculate the energy of electron in the first excited state. [Ans. -3.4 eV]
- **Q. 12.** Calculate the number of protons emitted in 10 hours by a 60 W sodium lamp emitting radiations of wavelength 6000 Å.
- **Q. 13.** Which one has a higher energy, a photon of violet light with wavelength 4000 Å or a proton of red light with wavelength 7000 Å ?

[Given. $h = 6.62 \times 10^{-34} J sec.$]

Q. 14. A 100 watt bulb emits monochromatic light of wavelength 400 nm. Calculate the number of protons emitted per second by the bulb.

[Ans. $2.012 \times 10^{20} \, s^{-1}$]

- **Q. 15.** What are the maximum number of emission lines when the excited electron of a H atom in n = 4 drops to the ground state ? [Ans. 6]
- **Q. 16.** Which has more energy, light radiation of wavelength 400 pm or light radiation of frequency 10¹⁵ Hz ?
- **Q. 17.** Find the energy of electron in 4th shell of Li^{2+} ion.
- **Q. 18.** What is the wave number of an electron with shortest wavelength radiation in Lyman spectrum of He⁺ ion?
- **Q. 19.** Write short note on :
 - (a) Continuous and discontinuous spectrum.
 - (b) Absorbtion and emission spectrum.
- **Q. 20.** Calculate the mass of the photon with wavelength of 3.6 Å.

[Ans. 6.135×10^{-29} kg]

Structure of Atom



- **Q. 21.** Calculate the mass of the photon with wavelength of 5 pm.
- **Q. 22.** On the basis of uncertainty principle show that an electron cannot exist with in atomic nucleus. (*Given : Nuclear radius* = 10^{-15} m) [*Hint : Taking* 10^{-15} m as Δx , the Δv comes much higher than the velocity of light and hence is not possible]
- **Q. 23.** Explain why the uncertainty principle is significant only from the motion of subatomic particles and is negligible for macroscopic particles?
- Q. 24. List two differences between orbit and orbital.
- **Q. 25.** Show that the circumference of the Bohr orbit for the hydrogen atom is an integral multiple of the de Broglie wavelength associated with the electron revolving around the orbit
- **Q. 26.** Comment on "Bohr's model is against the Heisenberg uncertainty principle".
- **Q. 27.** What are the similarities and difference in 2s and $2p_x$ orbitals and 1s and 2s orbitals ?
- **Q. 28.** Draw shape of $d_{x^2-v^2}$ orbital.
- **Q. 29.** On the basis of Pauli's exclusion principle show that the maximum number of electrons in the M -shell (n = 3) of any individual atom is 18.
- **Q. 30.** Designate each subshell with n = 4.
- **Q. 31.** List the possible values for all the quantum numbers for the following subshell.

(a) 2p (b) 4f

- **Q. 32.** Write down the electronic configuration of Fe^{3+} and Ni^{2+} . How many unpaired electrons are present? (Given Atomic number, Fe = 26, Ni = 28).
- **Q. 33.** Out of principal, angular, magnetic and spin quantum number, which quantum number determines the ?
 - (a) Shape of the orbital
 - (b) Number of orbitals in an orbit
 - (c) Size of the orbital
 - (d) Spin orientation of the electron.

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- **Q. 34.** What is the Hund's rule of maximum multiplicity ? Explain with suitable example.
- Q. 35. Explain why :
 - (a) The three electrons present in 2p subshell of nitrogen remain unpaired.
 - (b) Cr has configuration $3d^5 4s^1$ and not $3d^4 4s^2$.
- **Q. 36.** (a) What is difference between '*l*' and 'L'?
 - (b) Nitrogen has 7 proton, 7 electron and 7 neutrons. Calculate the number of electron, protons and neutrons in N^{3-} ion.
- Q. 37. Which one is having higher energy?
 - (a) Last electron of Cl^- or last electron of O^{2-} .
 - (b) n = 4, l = 3 or n = 5, l = 2.

3-MARKS QUESTIONS

- **Q. 1.**(i) The energy associated with the first orbit in the hydrogen atom is $-2.18 \times 10^{-18} \text{ J}$ atom⁻¹. What is the energy associated with the fourth orbit ?
 - (ii) Calculate the radius of Bohr's third orbit for hydrogen atom.

[Ans.– $1.36 \times 10^{-19} \text{ J atom}^{-1}.4.761 \text{ nm}$]

- **Q. 2.** A bulb emits light of wave length 4500Å. The bulb is rated as 150 watt and 8% of the energy is emitted as light. How many photons are emitted by the bulb per second ? $[Ans. n = 27.2 \times 10^{18}]$
- **Q. 3.** When light with a wavelength of 400 nm falls on the surface of sodium, electrons with a kinetic energy of 1.05×10^5 J mol⁻¹ are emitted.
 - (a) What is the minimum energy needed to remove an electron from sodium ?
 - (b) What is the maximum wavelength of light that will cause a photoelectron to be emitted ?

[Ans. $a = 3.2255 \times 10^{19}$ J, b = 616 nm]

Q. 4. Compare the frequency of light radiations emitted when electron falls from 5th shell to the 2nd shell in Li^{2+} ion and electron falls from 4th shell to the 1st shell in He⁺ ion.





- Q. 5. Calculate the number of waves made by Bohr electron in one complete revolution in its third orbit. [Ans. 3]
- **Q. 6.** What should be the ratio of velocities of CH_4 and O_2 molecules so that they are associated with de Broglie waves of equal wavelength? [Ans. 2]
- **Q. 7.** Calculate the wavelength of an electron that has been accelerated in a particle accelerator through a potential difference of 1 kv.

[Given $1eV = 1.6 \times 10^{-19} J$] [Ans. $3.87 \times 10^{-7} m$]

- Q. 8. (i) Discuss the similarities and differences between a 1s and 2s orbital.
 (ii) Draw the shape of d₂.
- **Q. 9.** Calculate the wavelength of a tennis ball of mass 60 gm moving with a velocity of 10 m per second. $(h = 6.626 \times 10^{-34} \text{ kg m}^2 \text{ s}^{-1})$

Q. 10. Calculate the wavelength of 1000 kg rocket moving with a velocity of 3000 km/hr. $(h = 6.626 \times 10^{-34} \text{ kg m}^2 \text{ s}^{-1})$

```
[Ans. 7.9512 \times 10^{-40} m]
```

Q. 11. Calculate the uncertain it in the velocity of a cricket ball of mass 150 g, if uncertainity in its position is of the order of 1 Å.

[Ans. $3.5 \times 10^{-24} \text{ m s}^1$]

- **Q. 12.** (a) What is de-Broglie wavelength for an electron moving with velocity of light?
 - (b) What is the angular momentum of electron in 5th shell?
- **Q. 13.** Two particles A and B have wavelength $\lambda_A = 5 \times 10^{-10}$ m and $\lambda_B = 10 \times 10^{10}$ m. Find their frequency, wave number and energies. Which has more penetrating power and why ?
- **Q. 14.** (a) Which has max. uncertainty regarding position and why? Electron, proton and neutron.
 - (b) Find the number of waves associated with a light radiation of time period 5 ns.
- **Q. 15.** If an electron in He⁺ has angular momentum of $5h/2\pi$. Find its energy and wavelength associated with it. Find the kinetic energy of this electron.





[[]**Ans.** 10^{-3} metre]

- **Q. 16**.(i) An atomic orbital has n = 2. What are the possible values of *l* and m_l ?
 - (ii) List the quantum numbers $(m_1 \text{ and } l)$ of electrons for 3*d* orbital.
 - (iii) Which of the following orbitals are possible ?

2*d*, 1*s*, 2*p* and 3*f*.

- **Q. 17.** (a) Write the maximum number of electron in a subshell with l = 3 and n = 4.
 - (b) Write the maximum number of electron that can be associated with the following set of quantum numbers ?

n = 3, l = 1 and $m_l = -1$

- (c) Write the maximum number of electron that can be accomodated in an atom in which the highest principal quantum number value is 4.
- Q. 18. (i) Write the electronic configurations of the following ions :

(a) $H^{-}(b)Na^{+}(c)O^{2-}(d) F^{-}$

- (ii) What are the atomic numbers of elements whose outermost electrons are represented by (a) $3s^{1}(b) 2p^{3}$ and (c) $3p^{5}$?
- (iii) Which atoms are indicated by the following configurations ?

(a) [He] $2s^1$ (b) [Ne] $3s^2 3p^3$ (c) [Ar] $4s^2 3d^1$.

Q. 19. Calculate:

- (a) Total number of spherical nodes in 3*p* orbital.
- (b) Total number of nodal planes in 3*p* orbital.
- (c) Nodal planes in 3*d* orbital.

5-MARKS QUESTIONS

- **Q. 1.**(a) Define Photoelectric effect ? Mention its one practical application in daily life.
 - (b) Electrons are emitted with zero velocity from a metal surface when it is exposed to radiation of wavelength 6800 Å. Calculate threshold frequency (v_0) and work function (W_0) of the metal.

[Ans. $v_o = 4.41 \times 10^{14} \text{ s}^{-1} \text{ W}_o = 2.92 \times 10^{-19} \text{ J}$]

- **Q. 2.**(a) The electronic energy in Bohr's orbit is negative .How will you account for it?
 - (b) The ionisation energy of hydrogen atom is 13.6 eV. What will be the energy of the first orbit of He^+ and Li^{2+} ions ?

[Ans. E_1 of $He^+ = -54.4 \text{ eV}$, E_1 of $Li^{2+} = -122.4 \text{ eV}$]

Structure of Atom



Q. 3.(a) Define the following terms :

- (i) Threshold frequency (ii) Work function.
- (b) The work function for Cs atom is 1 .9 eV. Find threshold wavelength (λ_0) and threshold frequency (ν_0) of this light radiation. If Cs metal is irradiated with a radiation of wavelength 500 nm find kinetic energy and velocity of emitted electron.
- **Q. 4.**(a) State de Broglie equation. Write its significance.
 - (b) A beam of helium atoms moves with a velocity of 2.0×10^3 m s⁻¹. Find the wavelength of the particle constituting the beam

 $(h = 6.626 \times 10^{-34} \text{ J s})$ [Ans. 49.9 pm]

- **Q. 5.**(a) State Heisenbergs uncertainty principle. Give its mathematical expression. Also give its significance.
 - (b) Calculate the uncertainity in the position of a dust particle with mass equal to 1 mg if the uncertainity in its velocity is $5.5 \times 10^{-20} \text{ms}^{-1}$.

[Ans. 9.55×10^{10} m]

- **Q. 6.**(a) Cricket ball, a tennis ball and a proton which has more uncertainity in velocity and which follows Heisenberg uncertainity principle maximum.
 - (b) What is the similarity in de-Broglie and Heisenberg principle? Which is different from Bohr theory for structure of atom?
 - (c) Why energy in a given subshell is negative?
- **Q. 7.**(a) Write short notes on:
 - (i) Aufbau principle (ii) Pauli's principle (iii) Hund's rule.
 - (b) Write the electronic configuration of the following ions :
 - (i) Fe³⁺(ii) Cu⁺ [Given Atomic number of Fe and Cu are 26 & 29]
- **Q. 8.**(a) Draw the shapes of the following orbitals.

(i) $3d_{xy}$ (ii) d_{z^2}

- (b) What is the total number of orbitals associated with the principal quantum number n = 3 ?
- (c) Using *s*, *p*, *d*, *f* notations, describe the orbital with the following quantum numbers:-

(a)
$$n = 3$$
, $l = 0$, (b) $n = 4$, $l = 2$, (c) $n = 5$, $l = 3$, (d) $n = 1$, $l = 0$

Chemistry Class XI



- **Q.9.** Explain the following :
 - (i) Energy of electron is not decided by : n, l, m and s.
 - (ii) Maximum number of electron with -1/2 spin for n = 3 is 6,9,12 or none.
 - (iii) Maximum number of electron can be present for n + l = 4.
 - (iv) 3*f* subshell is not possible.
 - (v) Maximum number of electrons in a subshell is : (2l+1) or (4l+1) or n^2
- **Q. 10.**(a) A neutral atom has 2K, 8L and 15 M electrons. Find the total numbers of electrons in s, p, d and f subshell.
 - (b) How many unpaired electrons are present in the following ions : Al^+ , Cr^{2+} , Co^{3+} and Mn^{2+}

(Given Atomic number : Al=13, Cr = 24, Co = 27 & Mn = 25)

- (c) One electron is present in 4f subshell. What is the sum of $n + l + m_1 + m_s$ values assuming 'f' subshell follows 3 to + 3 order of filling electron.
- **Q. 11.** Answer the following :
 - (a) n + l value for 14th electron in an atom.
 - (b) Increasing order of filling electron in 4f, 5p and 6d subshells.
 - (c) '*m*' and '*l*' value for last electron of Mg atom.

(Given atomic number of Mg is 12)

(d) Subshell in which last electron is present in Ga.

(Given Atomic number of Ga is 31)

(e) Sum of spin of all the electron in element having atomic number 14.





UNIT TEST

Time allowed : 1 hour

Maximum Marks : 20

General instructions : (i) All questions are compulsory. (ii) Maximum marks carried by each question are indicated against it. 1. Designation for an orbital with n = 4 and l = 3 is (1)(b) 4p (c) 4d (d) 4f (a) 4s 2. Maximum number of unpaired electrons in chromium is (1)(Given: Atomic number of Cr = 24) (a) 4 (b) 5(c) 6 (d) 7 3. Which series of lines of the hydrogen spectrum lie in the visible region'? (1)4. Write the Schrodinger wave equation. (1)5. Which of the following is not possible? (a) 2p (b) 3d (c) 3f (d) 4p (1)Write four difference between orbit and orbital. 6. (2)Calculate the wave number for the longest wavelength transition in 7. the paschen series of atomic hydrogen. (2)8. (a) How many orbitals are associated with n = 4? (3) How many electrons will be present in the sub-shells having (b) ms value of -1/2 for n = 3? Draw the shape of d_{z}^{2} . (c) Calculate the uncertainty in the position of a dust particle with mass 9. equal to 1 mg if the uncertainty in its velocity is 5.5×10^{-20} ms⁻¹. (3) The energy associated with the first orbit in the hydrogen atom is 10. (i) -2.18×10^{-18} J atom⁻¹. What is the energy associated with the fifth orbit? Calculate the radius of Bohr's fifth orbit for hydrogen atom. (ii) (iii) Calculate the radial and angular nodes in 2p orbital. (iv) Define the black body and black body radiations. (5)*****

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Classification of Elements _{Chapter -} 3 and Periodicity in Properties

FAST TRACK : QUICK REVISION

• The first systematic classification of elements was provided by Russian chemist D.I. Mendeleev.

1. Mendeleev's periodic law

"The physical and chemical properties of elements are periodic functions of their atomic weight."

2. It was modified to Modern Periodic law :

"The physical and chemical properties of elements are periodic functions of their atomic numbers."

It is the long form of periodic table :

7 Horizontal rows are called Periods and 18 Vertical columns are called Group

Group-1 are called **Alkali metals** Group-15 are called **Pnicogens** Group-17 are called **Halogens**

Group-16 are called Chalcogens

Group-2 are called Alkaline earth metals.

Group-18 are called Noble gases

- **3.** 1^{st} period 2 elements 2^{nd} and 3^{rd} period 8 elements 4^{th} and 5^{th} period 18 elements 6^{th} period 32 elements 7^{th} period Incomplete (32 elements)
- 4. Groups

1 and 2 – 's' block elements last electron entered in 's' subshell $[s^1, s^2]$ 3 to 12 – 'd' block elements last electrons entered in 'd' subshell $[d^1 \text{ to } d^{10}]$. 13 to 18 – 'p' block elements last electrons enter in 'p' subshell $[p^1 \text{ to } p^6]$. Two *f*-block series lanthanoids and actinoids are placed in the bottom of periodic table.

- 5. (A) In 's' and 'p' block elements the electrons enters in outer most shell. In 'd' block elements the electron enters in the penultimate shell (n-1). 'f' block elements last electron enter the antepenultimate shell (n-2).
 - (B) 'f' block elements are placed in between 'd' block elements.'f' block elements in 2 rows [4f lanthanoids, 5f actinoids]

6. General outer electronic configuration

's' block : ns^1 , ns^2 [Group 1 to 2] **'p' block :** ns^1np^1 to $ns^2 np^6$ Group 13 to 18 **'d' block :** $ns^{0-2} (n-1) d^{1 \text{ to } 10}$ Group 3 to 12 **'f' block :** $(n-2)f^{1 \text{ to } 14} (n-1)d^{0, 1} ns^2$

7. General periodic trends in properties of elements

• ATOMIC RADIUS

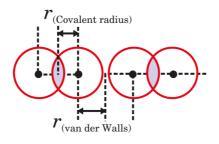
- (A) Left to right decreases due to effect of successive increasing nuclear change without addition of a new shell.
- (B) From top to bottom atomic radius increases due to successive addition of shell.
- (C) Noble gases have large radius than **group 17** due to complete filling of electron in outer shell electron-electron repulsion mildy increases.

• COVALENT RADIUS

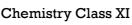
It is half of the distance between the centre of nuclei of two adjacent similar atoms which are bonded to each other by single covalent bond.

• van der Waal's Radius

van der Waal's radius is defined as one-half the distance between the centres of nuclei of two nearest like atoms belonging to two adjacent molecules of the element in the solid state.









• METALLIC RADIUS

Half of the distance between the centres of the nuclei of two adjacent atoms in the metallic crystal. A comparison of the three atomic radii show that van der Waal's radius is maximum while the covalent radius has the least value.

van der Waal's radius > Metallic radius > Covalent radius

• IONIC RADIUS

(A) <u>Cation radius < Atomic radius</u> – due to more no. of protons than number of electron coloumbic force increases, size decreases.

 $[Mg^{2+} < Mg^{+} < Mg]$

(B) <u>Anion radius > Atomic radius</u> – Due to more number of electron than number of protons

 $[N^{3-} > O^{2-} > F^{-}]$

Electron-Electron repulsion increase, coloumbic force of attraction decreases.

- (C) <u>For Isoelectronic species</u> More is the charge of cation lesser the size. More is the charge of anion, more is the size.
- (D) <u>Order of size</u> $O^{2-} > F^- > Na > Na^+ > Mg^{2+}$

8. (A) Ionisation enthalpy :

The minimum amount of energy which is required to remove the most loosely bound electron from an isolated atom in the gaseous state is called Ionisation enthalpy.

 $\begin{array}{rrrr} \mathrm{M}(\mathrm{g}) &+ & \mathrm{Energy} & \longrightarrow \mathrm{M}^+ &+ & e^- \\ & & \mathrm{IE}_3 &> & \mathrm{IE}_2 &> & \mathrm{IE}_1 \end{array}$

(B) Variation of I.E along a period:

Ionisation enthalpy increase along the period because atomic radii decrease and nuclear charge increase along the period.

 $\begin{array}{ll} I & \text{ionisation enthalpy} \\ II & \text{ionisation enthalpy} \end{array} \\ \begin{array}{ll} Li < B < Be < C < O < N < F < Ar \\ Be < C < B < N < F < O < Ne \end{array}$

(C) Variation down the group:

Ionisation enthalpy decrease down the group because atomic radius increase down the group.

Metallic behaviour : Decrease from left to right due to increase in ionisation enthalpy.

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Classification Of Elements And Periodicity In Properties

Non metallic behaviour : Increase from left to right due to more number of electron in outershell and added electron goes towards nucleus.

9. Screening effect or shielding effect:-

It is the decrease in the force of attraction between nucleus and outermost electron due to presence of inner shell electrons. As a result, the outer most electrons does not feel full charge of the nucleus. The actual charge felt by an electron is called effective Nuclear charge.

Shielding effect is in the following order s > p > d > f

d & f subshell show weak sheilding effect because their orbital size are large and are more diffused.

10. Isoelectronic species:

Ions of different elements which have the same number of electrons but different no. of protons are called isoelectronic ions.

	Na^+	Mg^{2+}	Al^{3+}	N ³⁻	O ^{2–}	F^-
No. of Protons	11	12	13	7	8	9
No. of electrons	10	10	10	10	10	10
Ionic Radii Al ³	$^{+} < Mg^{2+} <$	Na ⁺ <	F^{-} <	O ²⁻ <	N ³⁻	

11. Electron gain enthalpy:

The enthalpy change when an extra electron is added to neutral gaseous atom to form anion.

 $E(g) + e^- \longrightarrow E^-(g)$

- **Trends : From left to right** Increase due to decrease in size, more attraction of added electron by nucleus.
- From top to bottom—Decreases as the added electron is away from nucleus due to increase in size.
- *Cl* has more negative electron gain enthalpy than fluorine Due to small size of fluorine extra added electron has more inter electronic repulsion than chlorine which has large size.
- Similarly Phosphorus and Sulphur have negative electron gain enthalpy than nitrogen and oxygen respectively.
- Maximum electron gain enthalpy Chlorine (in periodic table)





Electron gain enthalpy –

Halogen > Oxygen > Nitrogen > Metal of group 1 and 13 and non metal of group 14 > metal of group 2.

• 2nd electron gain enthalpy is always positive.

12. Electro negativity:

The tendency of an atom to attract the shared pair of electron towards itself in a bonded state.

- Fluorine is the most electronegative element in the periodic table.
- Cesium is the least electronegative element in the periodic table.
- Electro-negativity decreases down the group and increases along the period

Difference between electron gain enthalpy and Electronegativity.

Electron gain enthalpy is the energy, but electronegativity is not the energy, it is only the tendency of an atom in a molecule to attract the shared pair of electrons. Three highest electronegative atoms F > O > N.

Maximum electronegative Assign to F.

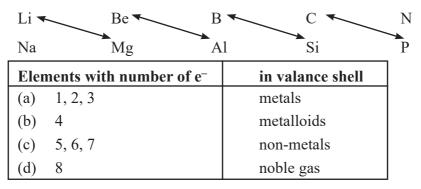
- * Lightest element : Hydrogen
- * Lightest metal : Lithium
- * Heaviest metal (highest density) : Osmium
- * Most reactive metal : Caesium
- * Most reactive nonmetal : Fluorine
- * Most malleable metal : Gold
- * Electrically best conductor : Silver
- * Metals which are relatively volatile : Zn, Cd, Hg
- * Strongest reducing agent in aqueous solution : Lithium
- * Strongest oxidising agent : Fluorine
- * The element of lowest ionisation energy : Caesium
- * The element of highest ionisation energy : Helium
- * The most electronegative element : Fluorine
- * The element of highest electron gain enthalpy : Chlorine
- * The group containing most electropositive metals : Group 1
- * The group containing most electronegative metals : Halogens Group 17
- * The group containing maximum number of gaseous elements : Group 18

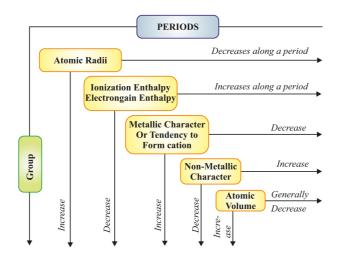
Classification Of Elements And Periodicity In Properties

13. Second period element—Show different behaviour that I group element— Due to (a) small size (b) High electron negativity (C) High polarising power (d) absence of 'd' orbital.

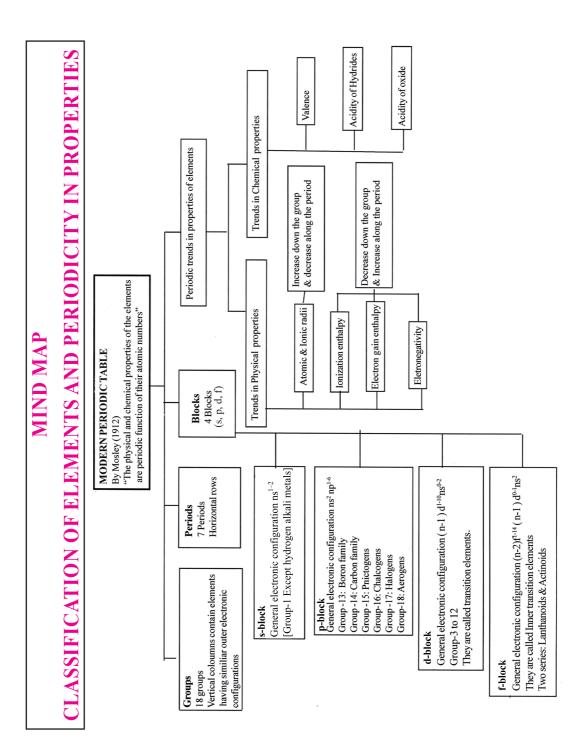
 $Na_{3}[Al(OH)_{6}]$ exists but $Na[B(OH)_{4}]$ not exists.

14. The similarities in properties of first member of a group to second member of just next higher group due to comparable atomic radius, nearly same polarising power of ions is known as **diagonal relationship**.









Classification Of Elements And Periodicity In Properties

MULTIPLE CHOICE QUESTIONS (MCQ)

- 1. According to modern periodic law, the physical and chemical properties of elements are the periodic functions of their ?
 - (a) Density (b) Atomic Number
 - (c) Mass Number (d) Atomic Mass
- 2. Highest electropositive element in the periodic table is
 - (a) Cs
 (b) Rb
 (c) K
 (d) Na
- 3. The correct order of ionic radii of the species N^{3-} , O^{2-} , Na^+ and F^- is
 - (a) $Na^+ < F^- < O^{2-} > N^{3-}$ (b) $F^- < O^{2-} < N^3 > Na^+$
 - (c) $O^{2-} < N^{3-} < F^- > Na^+$ (d) $N^{3-} < Na^+ < F^- > O^{2-}$
- 4. The basic strength of the oxides follows the order
 - (a) $Al_2O_3 > MgO > Na_2O$ (b) $Al_2O_3 < MgO < Na_2O$ (c) $Na_2O_3 < MgO > Al_3O_2$ (d) $Al_2O_3 > MgO > Na_2O$
- 5. The correct order of the size of C, N, P, S follows the order
 - (a) N < C < P < S(b) C < N < S < P(c) C < N < P < S(d) N < C < S < P

6. Which of the following oxide is most acidic?

- (a) Na_2O (b) Al_3O_2
- (c) P_2O_5 (d) SO_3
- 7. Downward in a group, electropositive character of elements
 - (a) increases (b) decreases
 - (c) remains same (d) none of these
- 8. Element which has more negative electron gain enthalpy is
 - (a) F (b) O
 - (c) Cl (d) S
- 9. The electronegivity of the following elements increase in the order
 - (a) C, N, Si, P (b) N, Si, C, P
 - (c) Si, P, C, N (d) P, Si, N, C

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- 10. The ionisation enthalpy of nitrogen is more than that of oxygen molecules because of
 - (a) greater attraction of electrons by the nucleus
 - (b) extra stability of the half filled p-orbitals
 - (c) smaller size of nitrogen
 - (d) more penetrating effect

Ans: 1. (b), 2. (a), 3. (a), 4. (b), 5. (d), 6. (d), 7. (a), 8. (c), 9. (c), 10. (d)

FILL IN THE BLANKS

1. Lightest metal in s-block elements is In the periodic table, horizontal rows are known as 2. 3. Elements of s-blocks and p-blocks are collectively called . Most electropositive elements belong to group. 4. 5. Most electronegative elements belong to group. The elements above atomic number 92 are called . 6. The inner-transition elements belong to _____ block of the periodic 7. table and are shown separately at the _____ of the periodic table. An element having electronic configuration [Ar] 3d⁵, 4s² belongs to 8. block. 9. Ca^{2+} has smaller ionic radius than K^+ ion because it has . 10. The maximum electronegativity is shown by . 11 The maximum ionisation enthalpy is shown by . The cation is _____ and the anion is _____ than the parent atom. 12 1. Lithium 7. F–, bottom Ans: 8. 2. periods s – 3. normal elements or 9. more protons representative elements 4. 1st 10. F-5. 17th 11. H 12. smaller, bigger 6. transuranic elements

Classification Of Elements And Periodicity In Properties

TRUE AND FALSE TYPE QUESTIONS

Write true or false for the following statements

- 1. First ionisation enthalpy of Be is higher than B.
- 2. Every period of the periodic table (except first period) starts with a member of alkali metal.
- 3. The energy liberated during the removal of one electron from an atom is called its ionisation potential.
- 4. Flourine has more negative electron gain enthalpy than chlorine.
- 5 Mg^{2+} ion has smaller size than Mg.
- 6. Electronegativity of F is larger than that of Cl but electron gain enthalpy of Cl is larger than of F.
- 7. The decreasing order of electronegativity of F, O and N is F > O > N.
- 8. Group-18 contain maximum gaseous elements.
- 9. Al_2O_3 is an amphoteric oxide.
- 10. Helium has the highest ionisation enthalpy.

Ans:	1. (T)	2. (T)	3. (T)	4. (F)	5. (T)
	6. (T)	7. (T)	8. (T)	9. (T)	10. (T)

MATCH THE COLUMNS

1	
T	٠

	Column A	(Column B
a.	Highest element	i.	Cesium
b.	Highest metal	ii.	Osmium
c.	Heaviest metal	iii.	Lithium
d.	Most reactive metal	iv.	Hydrogen

2.

Column A

- a. Fluorine
- b. Helium
- c. Chlorine
- d. Cesium

Column B

- i. High negative electron gain enthalpy
- ii. Most electropositive element
- iii. Most electronegative element
- iv. Highest ionisation enthalpy



	Column A	(Column B
a.	Na ₂ O	i.	Amphoteric oxide
b.	Cl ₂ O ₇	ii.	Acidic oxide
c.	Al ₂ O ₃	iii.	Neutral oxide
d.	СО	iv.	Basix oxide
	Column A	(Column B
a.	s & p-block	i.	Inner transition elements
b.	d-block	ii.	s-block elements
c.	f-block	 111.	Transition elements
d.	group-1 and group-2	iv.	Representative elements
An	s: 1. a.(iv), b.(iii), c.(ii), d	.(i)	
	3. a.(iv), b.(ii), c.(i), d.((iii)	4. a.(iv), b.(iii), c.(i), d.(ii)

ASSERTION AND REASON TYPE QUESTIONS

Directions for Q. No.1-5

- A If both Assertion & Reason are true and the reason is the correct explanation of the assertion.
- B If both Assertion & Reason are true but the reason is not the correct explanation of the assertion.
- C If Assertion is true statement but Reason is false.

D If both Assertion and Reason are false statements.

- 1. Assertion : Ionic radius of Na⁺ is smaller than Na
 - Reason : Effective nuclear charge of Na^+ is higher than Na
- 2. Assertion : First ionisation enthalpy of N is higher than O.
 - Reason : Extra stability of fully filled up 2p subshell of N atom
- 3. Assertion : Electron gain enthalpy of Cl is more negative than F atom.Reason : F is more electronegative than Cl atom.
- 4. Assertion : First ionisation enthalpy of Galium is higher than aluminium.Reason : Weak sheliding effect of 3d subshell is Galium.

Ans: 1. A 2. A 3. B 4. A

Classification Of Elements And Periodicity In Properties

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4.

ONE WORD ANSWER TYPE QUESTIONS

- 1. Metals are placed on which side of modern periodic table?
- 2. Which block of modern periodic table represent inner transition elements?
- 3. Name a halogen which has more negative electron gain enthalpy value?
- 4. Which element is iso-electronic with Na⁺? [Ans. Ne] [Given a atomic number of Sodium (Na) : 11]
- 5. An element is placed in 5th period and 3rd group what is its atomic number? [Ans. 39]
- 6. What is covalency of Al in $[AlCl_4]$? [Ans. 4]
- 7. Write the IUPAC Symbol for the element having atomic number 120. [Ans. Ubn]
- 8. Write the name of the group containing maximum number of gaseous elements.
- 9. Write the name of the subshell which show weakest sheilding effect.
- 10. Write the name of most electropositive element in the periodic table.
- 11. In what period and group will an element with Z = 118 will be present.

1-MARK QUESTIONS

- Which pair of elements has similar properties?
 13, 31, 11 & 21
- 2. Name the element which exhibit diagonal relationship with Be.

[**Ans.** 13, 31]

- 3. Which group elements are known as halogens?
- 4. The element with ns^2 , np^5 configuration is non-metal or metal?
- 5. Define van der Waal's radius.
- 6. Write the outer shell configuration of atomic number 31. [Ans. $4s^2$, p^1]
- 7. Find the group number and period number of element having atomic number 52.[Ans. Period = 5th, Group = 16th]
- 8. Arrange O^{2-} , O^{-1} , O in decreasing radius (size). [Ans. $O^{2-} > O^{-1} > O$]





- 9. Why noble gas have bigger size than halogens?
- 10. Why first electron gain enthalpy of sulphur is more negative then oxygen?
- 11. Write general outer electronic configuration of 4f series elements.

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[Ans. 6s<sup>2</sup>, 5d<sup>0–1</sup>, 4f<sup>1</sup> to 14]
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- 12. Write two isoelectronic species with Br (35). [Ans. Kr^+ , Se^{-1}]
- 13. Show that 4th period can have maximum 18 elements in it.
- 14. Second I.E. is always more than first I.E., why?
- 15. Electronegativity of F > Cl > Br > I, why?
- 16. Arrange F and Cl in terms of increasing chemical reactivity?
- 17. Second I.E. of Na is more than second IE of Mg. Why?
- 18. I.E. for cation is more than neutral atom. Why?
- 19. Define diagonal relationship with the help of an example.
- 20. Out of O^- and O, which has more negative electron gain enthalpy?
- 21. Mention any two anomalous properties of second period elements.

2-MARKS QUESTIONS

- 1. Cations are smaller than their parent atom whereas anions are larger in size than their parent atom. Explain.
- 2. Ionisation energy of nitrogen is more than 'O' and 'C' both, why ?
- **3.** First ionisation energy of boron is less than Be but size of Be is less than Boron. Why ?
- 4. Electron gain enthalpy of Mg is positive. Explain.
- **5.** Define co-valency.
- **6.** The reactivity of halogens decrease down the group but of alkali metals increases down the group. Why?
- Name a halogen, a metal and a group13 element which are liquid at 30°C. [Ans. Br, Hg, Ga]
- **8.** The reducing power of elements increases down the group but reverse is true for oxidising power along a period. Why ?

Classification Of Elements And Periodicity In Properties

- 9. What is the formula of binary compound formed between :
 - (a) 1st element of I group and iodine ?
 - (b) 2nd element of II group and 1st element of 17th group ?
- **10.** Arrange in the following in increasing order of property indicated:
 - (a) Size I, F, Cl, Br
 - (b) Oxidising power I, F, Br, Cl
- 11. Oxygen is more non-metallic than nitrogen but less than fluorine why?
- 12. LiCl, LiBr, LiI are covalent as well as ionic why?
- **13.** PbCl₂ is more stable than PbCl₄. Why ? [Ans. Inert pair effect]
- 14. [Magnesium and Lithium both form nitrides why?
- **15.** Which has least I.E. $[3p^3, 3p^6, 2p^3, 2p^6]$?
- **16.** (a) I.E. of sulphur is lower than chlorine.
 - (b) Arrange the following in decreasing order of their electro-negativity: F, O, N, Cl, C, H.
- 17. Element 'A' in group 17 (2nd period)'B' in group 16 (2nd period)

'C' in group 15 (2nd period)

Arrange 'A', 'B' and 'C' in their decreasing order of electro-negativity and ionisation enthalpy.

18. Element 'A' 13 group forms ionic compounds. Write the :

- (a) Formula of its oxide.
- (b) Arrange the following in their decreasing electro-positive character Mg, Na, Al, Si.
- 19. Write the atomic number of element place diagonally to :
 - (a) Group 14, period 4 (b) Group 2, period 5
 - (c) Group 17, period 4
- **20.** An element has outer shell electronic configuration $4s^2 4p^3$. Find :-
 - (a) The atomic number of element place next below it.
 - (b) Atomic number of next noble gas.





3-MARKS QUESTIONS

- 1. What is metallic radius, Covalent radius, van der waal's radius. Give one example for each.
- **2.** Oxygen has first electron gain enthalpy exothermic while second endothermic still a large number of ionic oxides are formed. Why ?
- **3.** In some properties Boron shows different properties with respect to rest of the membering the group. Justify.
- 4. Out of group 17, 18 and I, predict:-
 - (a) Which has most negative first electron gain enthalpy ?
 - (b) Which shows most metallic behaviour ?
 - (c) Which has highly positive electron gain enthalpy?
- 5. What are (a) representative elements, (b) Transition elements, (c) Lanthanoid and actinoids. Give their positions in modern periodic table.
- 6. Why LiF, NaF, KF, RbF, CsF are ionic ? But LiF is less ionic than CsF.
- 7. (a) Why Ca has larger atomic radius than Al?
 - (b) Why $2s^2$ electron is difficult to remove than 2p electron ?
- 8. (a) Why the compounds of group 17 with group 13 elements are more ionic and stable than with (group 1) elements? (b) Na₂O is more ionic than Li₂O. why?
- Explain the following data : Ionisation energy Cl < H < O < N < F.
- 10. IE_2 of 3^{rd} period elements is as follows. Why ? Mg < Si < Al < P < S < Cl < Ar < Na.
- **11.** Account fot the following:
 - (a) Halogens have very high negative electron gain enthalpy
 - (b) The electron gain enthalpy of Cl (Z = 17) is more negative than that of Fluorine (Z = 9).
 - (c) Ionisation enthaply of Nitrogen (Z = 7) is more than oxygen (Z = 8).
- **12.** What are the d- block elements? Write any four properties of d block elements and give their general outer electronic configuration.

- **13.** Explain the following:
 - (a) Modern Periodic law
 - (b) Electro-negativity
 - (c) Shielding effect
- **14.** Among the second period elements the actual ionisation enthalpies are in the order Li < B < Be < C < O < N < F < Ne. Explain why?
 - (i) Be has higher $(\Delta_i H)_1$ than B
 - (ii) O has lower $(\Delta_i H)_1$ than N and F?
- **15.** What do you understand by the isoelectronic species ? Name a species that will be isoelectronic with each of the following atoms or ions.
 - (i) F^- (ii) Ar (iii) Ca^{2+} (iv) Rb^+
- 16. (a) Show by a chemical reaction with water that Na_2O is a basic oxide and Cl_2O_7 is an acidic oxide.
 - (b) Name a species that will be isoelectronic with each of the following atoms or ions, (i) F⁻ (ii) Ca²⁺

17. The first ionisation enthalpy values (in $kJmol^{-1}$) of group-13 elements are:

В	Al	Ga	In	Tl
801	577	579	558	589

How would you explain this deviation from the general trend ?

18. The first (IE_1) and the second (IE_2) ionisation enthalpies (kJ mol⁻¹) of three elements are given below:

	Ι	II	III
IE ₁	403	549	1142
IE ₂	2640	1060	2080

Identify the element which is likely to be:-

- (a) a non metal
- (b) an alkali metal
- (c) an alkaline earth metal





5-MARKS QUESTIONS

- 1. (A) Which of the following have same chemical properties :
 - (a) Atomic number 17, 53
 - (b) Atomic number 8, 52
 - (c) Both
 - (d) None
 - (B) Answer the following :
 - (i) B, Al, Ga (decreasing order of atomic radii).
 - (ii) C, S, N (decreasing order of $(\Delta Heg)_1$)
 - (iii) Al forms amphoteric oxide. Why?
 - (iv) Si is a semiconductor while 'C' is a non-metal, why?

2.	Element	$\Delta_i \mathbf{H}^{\Theta}{}_1$	$\Delta_i \mathbf{H}^{\Theta}{}_{2}$	$\Delta eg H_{1}^{0}$
	Ι	1681	3374	- 328
	II	1008	1846	- 295
	III	2372	5251	+ 48

- (a) The most reactive non-metal.
- (b) The least reactive non-metal.
- (c) The least reactive element. Give reasons also.

[Ans. (a) 1 (b) II (c) III]

UNIT TEST

Maximum Marks : 20

Time allowed : 1 hour

General instructions :

(i) All questions are compulsory.

(ii) Maximum marks carried by each question are indicated against it.

1.	Which of the following show the weakest sheilding effect ?				ffect ?	(1)	
	(a) s	(b) p	(c)	d	(d)	f	
2.	Which has high	nest electronegativ	ity ?				(1)
	(a) Cl	(b) O	(c)	Ν	(d)	S	
3	Write the name	of the group cont	ainin	a maximu	m m111	mber of	(1)

- 3. Write the name of the group containing maximum number of (1) gaseous elements.
- 4. Write general outer electronic configuration of 4f series elements. (1)
- 5. Write the IUPAC symbol for the element having atomic number 120. (1)
- 6. (a) Explain why cation are smaller and anions larger in radii than (2) their parent atoms?
 - (b) Define accuracy & precision.
- 7. The first ionisation enthalpy values (in kJ mol⁻¹) of group-13 (2) elements are :

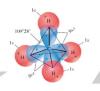
В	Al	Ga	In	Tl
801	577	579	558	589

How would you explain this deviation from the general trend?

- 8. (a) Show by a chemical reaction with water than Na_2O is a basic (3) oxide and Cl_2O_7 is an acidic oxide.
 - (b) Name a species that will be isoelectronic with each of the following atoms or ions. (i) F^- (ii) Ca^{2+}
- 9. Explain the following :
 - (a) Shielding effect
 - (b) Diagonal relationship
 - (c) Anomalous behavior of second period elements.
- 10. (a) Alkali metals do not form dis-positive ions. Why?
- (5)
- (b) Why is the IUPAC name and symbol of the element having atomic number 117.
- (c) Are the oxidation state and covalency of Al in $[Al(H_2O)_6]^{2+}$ same?
- (d) Why are there fourteen elements in the Lanthanide series?







Chemical Bonding and Molecular Structure

FAST TRACK : QUICK REVISION

Chapter - 4

- **Kossel-Lewis Concept:** Atoms take part in chemical combination to complete octet in their valence shell. This is known as octet rule.
- Limitation of Octet Rule: The octet rule, though useful but have some exceptions e.g. BF₃, NO₂, PCl₅, SF₆ etc.
- Lewis Symbol or Electron Dot Structure: Representing valence electrons by dots placed around the letter symbol of the element.

Types of Chemical Bonds:

- (i) Covalent Bond:
 - (a) Formed by sharing of electrons.
 - (b) It may be polar and nonpolar.
 - (c) It is directional in nature.
- (ii) Ionic Bond:
 - (a) Formed by transfer of electrons.
 - (b) Formation of ionic bond is favored by high lattice enthalpy, Low ionization enthalpy of metal atom and more negative electron gain enthalpy of nonmetal atom.
 - (c) It is non directional in nature.
 - Formal Charge (F.C.):
 - (i) It is charge appeared on individual atom in covalent molecule.
 - (ii) F.C. = (Total No. of valence electrons in free atom) (Total No. of unshared electrons) ½ (Total No. of shared electrons)
 Greater the F.C on atoms lesser the stability of that Lewis structure.
 - Lattice Enthalpy: Energy released when one mole of a crystalline solid is formed constituent gaseous ions.

Bond length:

- (i) It is equilibrium distance between the nuclei of two bonded atoms in a molecule.
- (ii) Greater the size of bonded atoms shorter the bond length. e.g., H-F < H-Cl < H-Br < H-I
- (iii) Greater the s character shorter the bond length. e.g., $C_{sp^3}-H > C_{sp^2}-H > C_{sp}-H >$
- (iv) Bond length decreases with increase in bond order. e.g., $C-C > C = C > C \equiv C$

Bond angle:

- (i) It is angle between the orbitals containing bonding electron pairs around central atom in a molecule or complex ion.
- (ii) Greater the electronegativity of central atom larger the bond angle e.g., $NH_3 > PH_3$
- (iii) Greater the number of lone pair around central atom smaller the bond angle. e.g., $CH_4 > NH_3 > H_2O$

Bond Enthalpy:

- (i) It is defined as amount of energy required to break one mole of bonds of a particular type between two atoms in gaseous state.
- (ii) For diatomic molecules, Bond enthalpy = Bond dissociation enthalpy
- (iii) For polyatomic molecules, Bond enthalpy = Average of all possible bond dissociation enthalpies.
- (iv) Bond enthalpy α Bond order α 1/(Bond length)

• Resonance:

60

- (i) According to the concept of resonance, whenever a single Lewis structure cannot describe a molecule accurately, a number of structures with similar energy, position of nuclei, bonding and non-bonding pairs of electrons are taken as canonical structures of the resonance hybrid which describes the molecule accurately.
- (ii) Resonance averages the bond characteristics as a whole.

• Partial ionic character of covalent bond A–B:

 $= 16(X_{A} - X_{B}) + 3.5(X_{A} - X_{B})^{2},$

where X_A and X_B are electro-negativities of A & B.



• Partial covalent character in ionic bond (Fajan's rule):

- (i) Fajan's rule is used to predict partial covalent character in ionic bond.
- (ii) Gretaer the polarizing power of cation and polarisability of anion greater the covalent character in ionic bond.
- (iii) Polarising power of cation α Charge density [(Charge)/Radius)].
- (iv) Polarisability of anion α size of anion.

• Dipole moment:

- (i) Dipole moment (μ) = charge (Q) × distance of separation (d)
- (ii) Unit: Debye (D), $1D = 3.33564 \times 10^{-30}$ Cm
- (iii) Being vector quantity, dipole moment of polyatomic molecule is taken as the resultant of all the bond moments.
- (iv) If μ = 0, molecule is non polar or symmetric.
- (v) If $\mu \neq 0$, molecule is polar or asymmetric.

Hydrogen bond:

- (i) It is dipole-dipole interaction between molecules in which 'H' atom is inserted between two highly electronegative elements i. e. F, O or N only.
- (ii) Hydrogen bond may be intra-molecular (when present within single molecule) and intermolecular (when present b/w two same or different molecules).
- (iii) Hydrogen bonds are stronger intermolecular forces than van der Waal forces.

• Sigma (σ) and pi (π) bonds:

- (i) Sigma bond is formed by axial overlapping and pi bond is formed by sideways overlapping of atomic orbitals.
- (ii) Sigma bond is stronger than pi bond due to greater extent of overlapping.
- (iii) Single covalent bond = 1 σ bond

Double covalent bond = 1 σ bond + 1 π bond

Triple covalent bond = 1 σ bond + 2 π bond

• VSEPR Theory: (VSEPR = Valence Shell Electron Pair Repulsion): The shape of a molecule depends upon the number of valence shell electron pairs (lp and bp) around the central atom and magnitude of repulsive forces between them

i.e., lp-lp > lp-bp > bp-bp

Hybridisation:

- (i) It is the phenomena of mixing of atomic orbitals of nearly same energy to form the new orbitals of equal energy and identical shape.
- (ii) The new orbitals are called hybrid orbitals and determine the shape of molecules.

Molecular Orbital Theory (MOT):

- (i) The overlap of atomic orbitals of same symmetry to form bonding and antibonding molecular orbitals by addition and substraction of their wave functions is known as MO theory.
- (ii) The electrons are filled in molecular orbitals in order of their increasing energy.

i.e.,

 $\sigma 1s, \sigma^* 1s, \sigma 2s, \sigma^* 2s, \pi 2p_x = \pi 2p_y, \sigma 2p_z, \pi^* 2p_x = \pi^* 2p_y, \sigma^* 2p_z$ (upto 14 electrons)

σ1s, σ*1s, σ2s, σ*2s, $σ2p_z$, $π2p_x = π2p_y$, $π*2p_x = π*2p_y$, $σ*2p_z$ (For more than 14 electrons)

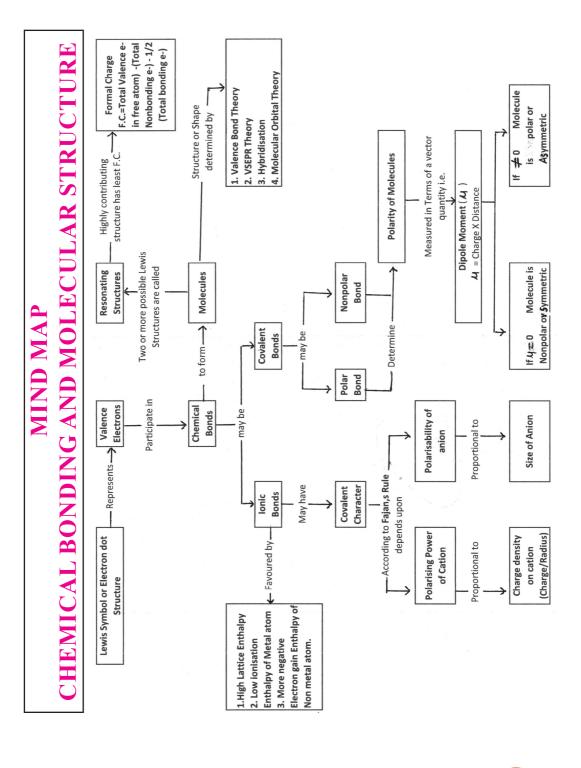
(iii) Bond order = $1/2 (N_b - N_a)$

 $N_a = No$ of electrons in anti-bonding molecular orbitals

 $N_{b} = No of electrons in bonding molecular orbitals$

Total	Bond	Lone	Type of	Geometry due to	Bond angle	Example
electron	pairs	pairs	hybridization	repulsion		
pairs						
2	2	0	sp	Linear	1800	BeCl ₂
3	3	0	sp ²	Non-polar Planar	1200	BF ₃
3	2	1	sp ²	Angular	<1200	SO ₂
4	4	0	sp ³ or dsp ²	Tetrahedral	109º28'	CH ₄
4	3	1	sp ³ or dsp ²	Pyramidal	<109º28'	NH ₃
4	2	2	sp ³ or sp ²	Bent	<109º28'	H ₂ O
5	5	0	sp ³ d	Trigonal bipyramidal	1200 & 900	PCI ₅
5	4	1	sp ³ d	See Saw	<120 º & <90º	SF ₄
5	3	2	sp ³ d	Bent T-shaped	<900	CIF ₃
5	2	3	sp ³ d	Linear	1800	I3 ⁻
6	6	0	sp ³ d ²	Octahedral	900	SF ₆
6	5	1	sp ³ d ²	Square pyramidal	<900	BrF ₅
6	4	2	sp ³ d ²	Square planar	900	XeF ₄
7	7	0	sp ³ d ³	Pentagonal bipyramidal	900 & 720	IF ₇
7	6	1	sp ³ d ³	Pentagonal pyramidal	<90º & <72º	
7	5	2	sp ³ d ³	Pentagonal planar	720	XeF5





MULTIPLE CHOICE QUESTIONS (MCQ)

1.	Which of the foll	lowing molecules ha	s both covale	nt and ionic bond		
	(a) CH ₃ Cl	(b) NH ₄ Cl	(c) HCl	(d) BeCl ₂		
2.	What is the maximum number of water molecules that can attach with one water molecule through intermolecular hydrogen bonds?					
	(a) 2	(b) 3	(c) 4	(d) 1		
3.	Which of the foll	lowing molecules ha	s maximum b	oond angle		
	(a) NH ₃	(b) CH ₄	(c) H ₂ O	(d) CO ₂		
4.	Identify correct s	tatement regarding N	NH_3 and BF_3			
	(a) Both are Lev	vis acid				
	(b) Both are iso	structural				
	(c) Both are Lew	vis base				
	(d) Have differen	nt values of dipole m	noment			
5.	Identify the mole	cule having sideway	s overlapping	g of atomic orbitals		
	(a) CH ₄	(b) CO ₂	(c) NH ₃	(d) H ₂ O		
6.	Which of the fol	lowing chemical spe	ecies is most s	stable?		
	(a) O ₂	(b) O ₂ ⁺	(c) O_2^{-}	(d) O_2^{2-}		
7.	Which of the foll	lowing d orbitals inv	olved in Sp ³ d	l hybridization?		
	(a) d _{xy}	(b) d _{xz}	(c) $d_{x^2-y^2}$	(d) d_{z^2}		
8.	Which of the foll	lowing molecule has	net dipole m	oment?		
	(a) CO ₂	(b) H ₂ O	(c) BF ₃	(d) CH ₄		
9.	Which of the foll	lowing compound ha	as highest cov	alent character		
	(a) LiCl	(b) LiBr	(c) LiF	(d) LiI		
10.	The shape of Xel	F ₄ molecule accordir	ng to VSEPR	theory is		
	(a) Square plana	r	(b) Square	pyramid		
	(c) Tetrahedral		(d) Pyrami	dal		

Ans. 1.(b) 2.(c) 3.(c) 4.(d) 5.(b) 6.(b) 7.(d) 8.(b) 9.(d) 10.(a)

Chemistry Class XI



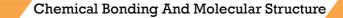
FILL IN THE BLANKS

- (i) The energy required to completely separate one mole of solid ionic compound into gaseous constituent ions is called.....
- (ii) Among alkali metal ionsion has highest polarizing power.
- (iii) According to molecular orbital theory molecules are said to be stable if the number of electrons in bonding molecular orbitals is the number of electrons in antibonding molecular orbitals.
- (iv) Isoelectronic molecules and ions have identical.....
- (v) In PCl_5 molecule the two equivalent axial P Cl bonds are.....than three equivalent equatorial P Cl bonds.
- (vi) The state of hybridization of sulphur in SF_6 is.....
- (vii) The strongest intermolecular hydrogen bonding in water is present at...°C.
- (viii) A triple covalent bond consists of.....sigma and.....pi bonds.
- (ix)bond is directional in nature.
- (x) Atomic orbitals are.....centric and molecular orbitals are.....
- Ans. (i) Lattice enthalpy (ii) Li⁺ (iii) more (iv) bond order (v) longer (vi) sp³d² (vii) 4 (viii) 1, 2 (ix) covalent (x) mono, poly

TRUE AND FALSE TYPE QUESTIONS

Write true or false for following statements:

- (i) Energy of resonance hybrid is less as compared to the contributing canonical structures.
- (ii) BeF_2 has more dipole moment than $BeCl_2$.
- (iii) In water two O–H bond dissociation enthalpies are not identical.
- (ix) Only the half filled orbitals of nearly same energy can participate in hybridization.
- (v) No bond is purely ionic or purely covalent.
- (vi) Chemical species having identical bond order have same bond dissociation enthalpies.



(vii) BF₃ is stronger Lewis acid than BCl₃.

(viii) Among alkali metal halides LiI has highest covalent character.

(ix) Resonating structures of a chemical species have no real existence.

- (x) XeF_2 and ICl_2^- are iso structural.
- Ans.(i) True(ii) False(iii) False(iv) False(v) True(vi) False(vii) False(viii) True(ix) True(x) True

MATCH THE COLUMNS

1. Match the species in Column I with the geometry/shape in Column II.

Column I	Column II
(i) PCl ₅	(a) Tetrahedral
(ii) ClF ₃	(b) Pyramidal
(iii) NH ₄ ⁺	(c) Bent T shape
(iv) H ₃ O ⁺	(d) Trigonal bipyramid

2. Match the species in Column I with the type of hybrid orbitals in Column II.

Column I	Column II		
(i) BF ₃	(a) sp^3d		
(ii) H ₂ O	(b) sp^2		
(iii) PCl ₅	(c) sp^3d^2		
(iv) SF ₆	(d) sp^3		

Ans. 1. (i) \rightarrow (d), (ii) \rightarrow (c), (iii) \rightarrow (a), (iv) \rightarrow (b). 2. (i) \rightarrow (b), (ii) \rightarrow (d), (iii) \rightarrow (a), (iv) \rightarrow (c).

ASSERTION AND REASON TYPE QUESTIONS

In the following questions a statement of assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given below for each question:

- (i) A and R both are correct, and R is correct explanation of A.
- (ii) A and R both are correct, but R is not the correct explanation of A.
- (iii) A is true but R is false.
- (iv) A and R both are false.



- Assertion (A): Among the two O-H bonds in H₂O molecule, the energy required to break the first O-H bond and the other O-H bond is the same. Reason (R): This is because the electronic environment around the oxygen is the same even after breakage of one O-H bond.
- Assertion (A): Though the central atom of both NH₃ and H₂O molecules are sp³ hybridised, yet H–N–H bond angle is greater than that of H–O–H. Reason (R): This is because nitrogen atom has one lone pair and oxygen atom has two lone pairs.
- Assertion (A): SF₆ molecule is unstable. Reason (R): A stable molecule must have 8 electrons around the central atom. i.e. octet rule should be satisfied.
- 4. Assertion (A): Pi bond is never formed alone. It is formed along with a sigma bond

Reason (R): Pi bond is formed by sideway overlap of p- orbitals only.

- Assertion (A): Ionic compounds tend to be non-volatile. Reason (R): Ionic compounds are solid.
- **Ans.** 1. (iv) 2. (i) 3. (iv) 4. (iii) 5. (ii)

ONE WORD ANSWER TYPE QUESTIONS

- 1. Write the formal charge on central oxygen atom in O_3 molecule?
- 2. Write the shape of AB_2E_3 type molecule.
- 3. Name the property used to measure the degree of polarity.
- 4. Name the covalent bond formed by axial overlapping of atomic orbitals.
- 5. Out of p_x , p_y , p_z orbitals which p orbital takes part in sp hybridization?
- 6. Name the molecular orbital having energy greater than that of combining atomic orbitals.
- 7. Name the intermolecular forces responsible for liquid state of water.
- 8. Name the phenomena used to describe a molecule whose single Lewis structure cannot describe it.
- 9. Name the geometry involved in sp^3d hybridization.
- 10. Name the molecular theory that can explain magnetic character of molecules.
- Ans. 1. +1, 2. Linear, 3. Dipole moment, 4. Sigma bond, 5. p_z
 6. Antibonding molecular orbital, 7. Hydrogen bond, 8. Resonance,
 9. Trigonal bipyramid, 10. Molecular orbital theory



Chemical Bonding And Molecular Structure

1-MARK QUESTIONS

- 1. Why noble gases exist in mono atomic form?
- 2. Write the Lewis structure of NO_2^{-} .
- 3. Why NH₃ and BF₃ have different shapes?
- 4. How many sigma and pi bonds are present in HCN molecule?
- 5. Why sigma bond is stronger than pi bond?
- 6. Explain why BeH₂ molecule has zero dipole moment although the Be-H bonds are polar?
- 7. Which has highest bond angle? NO_2 , NO_2^- , NO_2^+
- 8. What is magnetic character of anion of KO_2 ?
- 9. Why do atoms combine?
- 10. What is the significance of Lewis Symbols?
- 11. Why density of water is maximum at 277K?
- 12. Give structure of BrF_5 according to VSEPR theory.
- 13. Why NH_3 is liquid and PH_3 is a gas?
- 14. Why KHF₂ exist but KHCl₂ and KHBr₂ does not?

[Ans. HF...HF hydrogen bonding].

- 15. Boiling point of p-nitrophenol is more than O-nitrophenol why?
- 16. How paramagnetic character of a compound is related to the no. of unpaired electrons?
- 17. Define the term bond length.
- 18. He_2 molecule does not exist. Give reason.
- 19. Why PCl₅ dissociates to give PCl₃ and Cl₂?
- 20. Write the state of hybridization of O in H_2O .
- 21. Predict the shape of ClF_3 according to VSEPR theory.
- 22. Why ice has less density than water?
- 23. Why the H-P-H bond angle in PH₃ is less than H-N-H bond angle in NH₃?
- 24. At room temperature H_2O exist as liquid while H_2S exist as gas. Give reason.
- 25. NH_3 has higher boiling point than PH_3 . Give reason.
- 26. Identify the chemical species having identical bond order: $O_2^{2^+}$, N_2 , $O_2^{-2^-}$.



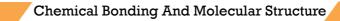


2-MARKS QUESTIONS

- 1. What is an Octet rule? What are its limitations?
- 2. The enthalpy needed to break the two O–H bonds in water are as follows:

$$\begin{split} H_2O(g) &\to H(g) + O - H(g) & \Delta_a H_1^{0} = 502 \text{ kJ mol}^{-1} \\ O - H(g) &\to H(g) + O(g) & \Delta_a H_1^{0} = 502 \text{ kJ mol}^{-1} \\ \end{split}$$
What is the average bond enthalpy of H_2O ?

- 3. Write two points of difference between sigma and pi bond.
- 4. Define Hydrogen bond. Is it weaker or stronger than van der Waal forces?
- 5. Define dipole moment. Give its significance.
- 6. Give applications of dipole moment.
- 7. Which is more polar and why, CO_2 or N_2O ?
- 8. Discuss the partial ionic character of covalent bond by taking an example.
- 9. Draw the resonating structures of O₃ and calculate formal charges on each O atom.
- 10. O-Nitrophenol is steam volatile while p-Nitrophenol is not. Give reason.
- 11. Define bond enthalpy. Why the bond enthalpy of F_2 is less than that of Cl_2 ?
- 12. Define resonance. Draw resonating structures of CO_2 .
- 13. Assign reason for the following;
 - (i) NH_3 is freely soluble in water while PH₃ is not.
 - (ii) B_2 is paramagnetic while C_2 is not.
- 14. Out of NH₃ and NF₃ which is more polar. Explain with the help of dipole moment.
- 15. N_2 is diamagnetic while O_2 is paramagnetic. Explain on the basis of Molecular orbital theory.
- 16. H_2^+ and H_2^- have same bond order. Which is more stable?
- 17. Differentiate between bonding and anti bonding molecular orbitals.
- 18. Discuss the conditions for the combination of atomic orbitals to form molecular orbitals.



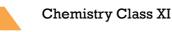
- 19. Although Chlorine (EN = 3.2) is more electronegative than Nitrogen (EN = 3.0), yet chlorine does not form hydrogen bond while nitrogen does. Give reason. (Ans: larger atomic size of Cl).
- 20. ClF_3 is T shaped but BF₃ is planar. Explain.
- 21. N $(SiH_3)_3$ and N(CH₃)₃ are not isostructural. Give reason.
- 22. Draw molecular orbital diagram for N_2^+ molecule.
- 23. HCl is a covalent compound but it ionises in the solution?
- 24. The molecule of CO_2 is linear whereas that of $SnCl_2$ is angular why?
- 25. Arrange the following in the order of property indicated for each set:
 - (i) $O_2, O_2^{+}, O_2^{-}, O_2^{2-}$ (increasing stability)
 - (ii) LiCl, NaCl, KCl, RbCl (increasing covalent character)
 - (iii) NO₂, NO₂⁺, NO₂⁻ (decreasing bond angle)
 - (iv) H-F, H-Cl, H-Br, H-I (increasing bond dissociation enthalpy)
- 26. Arrange the following in the order of property indicated for each set:
 - (i) H₂O, NH₃, H₂S, HF (increasing polar character)
 - (ii) HF, HCl, HBr, HI (decreasing dipole moment)
 - (iii) NO_3^{-} , NO_2^{-} , NO (decreasing 's' character of hybridization)
 - (iv) BeCl₂, BCl₃, CCl₄, PCl₃ (increasing bond angle)

3-MARKS QUESTIONS

- 1. How is ionic bond formed? On what factors it depends?
- 2. Calculate the lattice enthalpy of KCl from the following data by Born-Haber's Cycle.

Enthalpy of sublimation of K = 89 kJ mol⁻¹ Enthalpy of dissociation of Cl2 = 244 kJ mol⁻¹ Ionization enthalpy of potassium = 425 kJ mol⁻¹ Electron gain enthalpy of chlorine = -355 kJ mol⁻¹ Enthalpy of formation of KCl = -438 kJ mol⁻¹

3. What is meant by hybridization? Describe the shape of sp, sp² and sp³ hybridised orbitals.





- 4. Define bond order. Calculate the bond order in N_2 and O_2 molecules.
- 5. Give molecular orbital energy level diagram of CO. Write its electronic configuration,magnetic behaviour and bond order.
- 6. Which of the following in each pair has larger bond angle
 (i) CO₂, BF₃
 (ii) H₂O, H₂S
 (iii) CH₄, C₂H₂
- 7. What is meant by resonance? Draw the resonating structures of carbonate ion and explain why all the C–O bond lengths are identical in carbonate ion?
- 8. Compare relative stability of following species and predict their magnetic properties:

 O_2, O_2^+, O_2^- (superoxide), O_2^{2-} (peroxide)

- 9. Draw the Lewis structure of the species as mentioned below:
 - (i) In which the central atom has incomplete octet.
 - (ii) In which the central atom has an expanded octet,
 - (iii) An odd electron molecule is formed.
- 10. Explain the structure of PCl_5 according to hybridization. Why all P-Cl bonds lengths are not equivalent in PCl_5 ?

5-MARKS QUESTIONS

- 1. Give reasons for the following:
 - (a) NH_3 has higher boiling point than PH_3 .
 - (b) Ionic compounds do not conduct electricity in solid state.
 - (c) LiCl is more covalent than KCl.
 - (d) NH_3 is more polar than NF_3 .
 - (e) H_2O has bent structure.
- 2. (a) Define the term bond dissociation enthalpy. How is it related to bond order?
 - (b) Explain why N_2 has greater bond dissociation enthalpy than N_2^+ while O_2 has lesser bond dissociation enthalpy than O_2^+ ?
- Draw the shape of following molecules according to VSEPR theory; XeO₃, XeF₂, XeOF₄, SF₄, XeF₄

HOTS QUESTIONS

- 1. The bond angle of H_2O is 104.5° while that of F_2O is 102°. **Solution:** The bond pair of electrons are drawn more towards F in F_2O , whereas in H_2O it is drawn towards O. So bp-bp repulsion in H_2O is greater than that in F_2O .
- 2. Anhydrous $AlCl_3$ is covalent. From the data given below, predict whether it would remain covalent or become ionic in aqueous solution. $\Delta_i H (AlCl_3) = 5137 \text{ kJ mol}^{-1}, \quad \Delta_{hvd} H(Al^{3+}) = -4665 \text{ kJ mol}^{-1},$

 $\Delta_{\rm hyd} \, {\rm H}({\rm C1}^{-}) = -381 \, {\rm kJ} \, {\rm mol}^{-1}.$

Solution: Total energy released = $1\Delta_{hvd} H(Al^{3+}) + 3\Delta_{hvd} H(Cl^{-})$

$$= [(-4665) + (3 \times -381)] \text{ kJ mol}^{-1} = -5808 \text{ kJ mol}^{-1}$$

Total energy required = $\Delta_i H(AlCl_3) = 5137 \text{ kJ mol}^{-1}$

Since energy released is greater than the energy required, the compound will ionize in aqueous solution.

3. The dipole moment of HCl is 1.03 D, and the bond length is 127 pm. Calculate the percent ionic character of HCl molecule.

Solution: $\mu_{cal} = Q \times r = (1.6 \times 10^{-19} C) \times (127 \times 10^{-12} m) = 2.032 \times 10^{-29} C m$

$$= (2.032 \times 10^{-29} \text{C m}) \times \frac{1\text{D}}{3.336 \times 10^{-30} \text{Cm}} = 6.09 \text{ D}$$

% ionic character = $\frac{\mu_{obs.}}{\mu_{cal}} \times 100 = \frac{1.03\text{D}}{6.09\text{D}} \times 100 = 16.9\%$





UNIT TEST

Time Allowed: 1 hr

Maximum Marks : 20

General Instructions:

(i) All questions are compulsory.

1.	Identify the molecule (a) CH_4 (b)	e having sideway o) CO ₂	ys overlapping of (c) NH ₃	atomic orbitals (d) H ₂ O	[1]
2.	The shape of XeF ₄ n (a) Square planar (c) Tetrahedral	nolecule accordin	ng to VSEPR theo (b) Square pyras (d) Pyramidal		[1]
3.	Write the Lewis struc	ture of NO_2^{-} .			[1]
4.	Which has highest b	ond angle? NO ₂ ,	NO ₂ ⁻ , NO ₂ ⁺		[1]
5.	Draw the resonating s	structures of CO ₂			[1]
6.	$O-H(g) \longrightarrow$	H(g) + O - H(g) $H(g) + O(g)$	$\Delta_{a}H_{1}^{0} = 50$ $\Delta_{a}H_{2}^{0} = 42$		
	What is the average b	ond enthalpy of H	4 ₂ Ο?		[2]
7.	Out of NH ₃ and NF ₃ dipole moment.	which is more p	olar. Explain wit	h the help of	[2]
8.	Compare relative sta magnetic properties:	•	• • •		[3]
9.	Explain the structure P-Cl bonds lengths	5		on. Why all	[3]
10.	(i) N ₂ is diamagneti basis of Molecul	<u> </u>		ain on the	[2]
	Ð	her boiling point ounds do not con	duct electricity in	n solid state.	[3]



States of Matter : Gases, Liquids and Solids

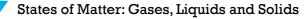
FAST TRACK : QUICK REVISION

- Magnitude of Intermolecular Forces: Hydrogen bonds > van der Waal forces (dipole-dipole > dipole-induced dipole > Dispersion forces)
- Gas Laws:
 - (i) **Boyle's Law:** $V \alpha 1/P$ or PV = constantor $P_1V_1 = P_2V_2$ (at constant n, T)
 - (ii) **Charles' Law:** V α T or $\frac{V_1}{T_1} = \frac{V_2}{T_2}$ (at constant n, P)
 - (iii) Avogadro's Law: V α n (at constant T, P)
 - (iv) Gay Lussac's Law: $P \alpha T$ (at constant n, V)

• Combined Gas Law:
$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$$

- Ideal Gas Equation: PV = nRT
- Values of gas constant R:
 - (i) $0.0821 \text{ L} \text{ atm } \text{K}^{-1} \text{ mol}^{-1}$
 - (ii) 0.083 L bar K⁻¹ mol⁻¹
 - (iii) 8.314 J K⁻¹ mol⁻¹
 - (iv) 1.99 Cal K⁻¹ mol⁻¹
- **Density of gas (d)**: d = PM/RT (M = molar mass of gas)
- Absolute zero or lowest possible temperature: -273°C or zero Kelvin, because at this temperature volume of gas becomes zero.

- Dalton's Law of partial pressure: P_{total} = p₁ + p₂ + p₃ +.... (at constant T&V) for non reacting gases.
- Ideal Gas: A gas which obeys ideal gas equation at all temperature and pressure.
- **Boyle's temperature**: Temperature at which a real gas behaves like an ideal gas over an appreciable range of pressure.
- **Compressibility factor (Z):** Z = PV/nRT
 - (i) For ideal gas Z = 1
 - (ii) For non ideal gas $Z \neq 1$
 - (a) **Positive deviation** (Z > 1): shows dominance of repulsive forces and hence less compressibility e.g. H₂, He etc.
 - (b) Negative deviation (Z < 1): shows dominance of attractive forces and hence more compressibility e.g. CH_4 , CO_2 etc.
- **Conditions under which real gases deviates from ideal behavior**: Low T and high P.
- **Cause of deviation from ideal behavior**: At low T and high P gas molecules are close enough and hence volume occupied by the gas molecules and attractive forces between them cannot be negligible.
- Conditions under which real gases behaves ideally: High T and low P.
- van der Waals' gas equation: $\left(P + \frac{an^2}{v^2}\right)(v nb) = nRT$
- van der Waals' constant 'a':
 - (i) It represents magnitude of attractive forces between gas molecules.
 - (ii) Ease of liquefaction of gas αT_c (αT_c = critical temperature) and Ease of liquefaction of gas αa
 - (iii) Unit of 'a' is L^2 atm mol⁻²
- Van der Waals' constant 'b':
 - (i) It represents co-volume or excluded volume i.e. effective volume of gas molecules.
 - (ii) It is four times of actual volume of gas
 - (iii) Unit of 'b' is $L^2 \mod^{-1}$

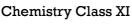


- **Critical temperature**: It is the temperature above which a gas can not be liquefied however large the pressure may be.
- Total Kinetic Energy of Gas = $\frac{3}{2}$ nRT
- Average Kinetic Energy of Gas = $\frac{3}{2}$ RT mol⁻¹ or $\frac{3}{2}$ KT molecule⁻¹ (k = Boltzmann constant = R/N_A)

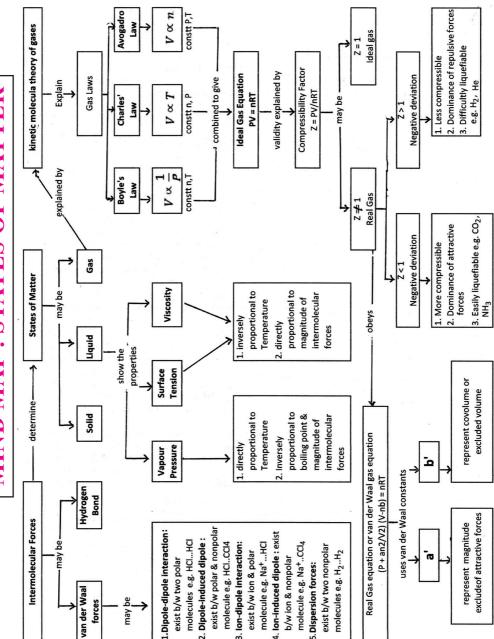
• Different types of Molecular speeds:

- (i) Most probable speed: Ump= $\sqrt{(2RT/M)}$
- (ii) Average speed: Uav = $\sqrt{(8RT/\pi M)}$
- (iii) Root mean square speed: Urms = $\sqrt{(3RT/M)}$ or $\sqrt{(3PV/M)}$
- (i) Vapour pressure α T
 - (ii) Vapour pressure $\alpha = \frac{1}{\text{Magnitude of intermolecular forces}}$
 - (iii) Vapour pressure $\alpha \frac{1}{\text{Boiling point}}$
- **Surface tension**: It is tangential force acting along the surface of a liquid perpendicularly on one centimeter length of it.
- **Viscosity**: It is internal resistance to the flow possessed by a liquid.
- Increase in temperature decreases surface tension and viscosity of liquid.









MIND MAP : STATES OF MATTER



MULTIPLE CHOICE QUESTIONS (MCQ)

1.	Which of the following property droplets:	explains the sph	erical shape of rain
	(a) Viscosity	(b) Critical phe	nomena
	(c) Surface tension	(d) Pressure	
2.	A gas would be most likely to obey	the ideal gas law	at
	(a) Low T and high P	(b) High T and	high P
	(c) Low T and low P	(d) High T and	low P
3.	If P is the pressure and d is the dense	ity of gas, then P	and d are related as:
	(a) $P \alpha 1/d$ (b) $P \alpha d$	(c) $P \alpha d^2$	(d) P $\alpha 1/d^2$
4.	A gas can be liquefied		
	(a) above its critical temperature	(b) at its critica	l temperature
	(c) below its critical temperature	(d) at any temp	erature
5.	Which of the following gas is experimental wall's constant 'a'	cted to have high	nest value of Van der
	(a) NH_3 (b) H_2	(c) N ₂	(d) He
6.	The compressibility factor (Z) for an	2	(.)
-	(a) 1.5 (b) 1.0	(c) 2.0	(d) zero
7.	Two separate bulbs contain ideal ga		
	that of B.The molecular mass of A is		-
	the same temperature, the ratio of pr		
0	(a) 2 (b) $1/2$	(c) 4	(d) $1/4$
8.	At which temperature the volume of $(a) 0.8C$ (b) 272 V	•	
0	(a) 0 °C (b) 273 K		(d) 273°C
9.	Dominance of strong attractive force (a) Depends on Z and indicates that	•	lecules of the gas:
	(a) Depends on Z and indicates that (b) Depends on Z and indicates that		
	(b) Depends on Z and indicates that(c) Depends on Z and indicates that		
	(d) Is independent of Z	$L \geq 1$	
10.		a of liquid inor	and an increasing
10.	Which of the following propertie temperature:	s of fiquid file	eases on mereasing
	(a) Vapour pressure	(b) Viscosity	
	(c) Surface tension	(d) Boiling Poi	nt
Ans.	1.(c), 2.(d), 3.(b), 4.(c), 5.(a),	6.(b), 7.(c), 8.	(c), 9.(c), 10.(a)

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FILL IN THE BLANKS

- 1. Pressure vs volume graph at constant temperature is known as.....
- 2. Surface tension of a liquidwith increase in magnitude of intermolecular forces.
- 3. is the temperature at which a real gas behave like an ideal gas over appreciable range of pressure.
- 4. Z > 1 indicates that the gas is compressible than expected from ideal gas behavior.
- 5. The average kinetic energy of gas molecules is directly proportional to the.....
- 6. Internal resistance in flow of liquids is called.....
- 7. is the temperature above which a gas cannot be liquefied however large the pressure may be.
- 8. Poise (P) is the unit of
- 9. The vapour pressure of any liquid is proportional to the magnitude of the intermolecular forces and is proportional to the temperature employed.
- 10 Van der Waal constant..... represent co-volume and represent magnitude of attractive forces.
- Ans. 1. Isotherm 2. increases 3. Boyle's temperature 4. less
 - 5. Kelvin temperature 6. viscosity 7. critical temperature
 - 8. viscosity 9. inversely, directly 10. 'b', 'a'

TRUE AND FALSE TYPE QUESTIONS

Write true or false for following statements:

- 1. At a given temperature and pressure the density of N_2 is more than that of O_2 .
- 2. Liquid at higher altitudes boil at a lower temperature.
- 3. Gases having Z < 1 cannot be liquefied easily.
- 4. According to the kinetic molecular theory, the collision between gas molecules is perfectly elastic.
- 5. Real gases deviate from ideal behavior at low temperature and high pressure.
- 6. A gas can be liquified above its critical temperature by applying high pressure.



States of Matter: Gases, Liquids and Solids

- 7. Mosquito cannot walk on kerosene oil because its surface tension is less than that of water.
- 8. No gas is ideal gas, all gases are real gases.
- 9. Surface tension increases on increasing temperature.
- 10. 0°C is known as absolute zero temperature.

Ans. 1. False	2. True	3. False	4. True	5. True
6. False	7. True	8. True	9. False	10. True

MATCH THE COLUMNS

1.

2.

Column-I

- (i) Boyle's Law
- (ii) Charle's Law
- (iii) Dalton's Law
- (iv) Avogadro Law
 - Column-I
- (i) Critical Temperature (a) Boiling Point
- (ii) Vapour Pressure
- (iii) Viscosity
- (iv) Surface Tension

- Column-II (a) V α n at constant T & P
- (b) $P_{total} = p_1 + p_2 + p_3 + \dots$ at constant T&V
- (c) V α T at constant n & p
- (d) $p \alpha 1/V$ at constant n & P
 - Column-II
- (b) Spherical shape of water droplet
- (c) Liquefaction of gases
- (d) Flow of Liquids

Ans. 1. (i) \rightarrow (d), (ii) \rightarrow (c), (iii) \rightarrow (b), (iv) \rightarrow (a).

2. (i) \rightarrow (c), (ii) \rightarrow (a), (iii) \rightarrow (d), (iv) \rightarrow (b).

ASSERTION AND REASON TYPE QUESTIONS

In the following questions a statement of assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given below for each question.

- A and R both are correct, and R is correct explanation of A. (i)
- (ii) A and R both are correct, but R is not the correct explanation of A.
- (iii) A is true but R is false.
- (iv) A is false but R is true.
- Assertion (A): Gases do not liquefy above their critical temperature even 1. on applying high pressure.

Reason (R): Above critical temperature, the molecular speed is high and intermolecular attractions can not hold the molecules together because the escape because of high speed.





2. Assertion (A): At constant temperature, pV vs V plot for real gases is not a straight line.

Reason (R): At high pressure all gases have Z > 1 but at intermediate pressure most gases have Z < 1.

3. Assertion (A): At zero degree Kelvin, the volume occupied by a gas is negligible.

Reason (R): All molecular motion ceases at 0 K.

- 4. Assertion (A): CO_2 has stronger intermolecular forces than CH_4 . Reason (R): Critical temperature of CO_2 is more.
- Assertion (A): Lower the critical temperature of the gas, more easily can it 5. be liquefied.

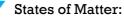
Reason (R): Critical temperature is the temperature above which a gas can not be liquefied depending upon the pressure.

Ans. 1. (i) 2. (ii) 3. (iii) 4. (i) 5. (iv)

ONE WORD ANSWER TYPE QUESTIONS

- 1. Write SI unit of pressure.
- 2. Write the value of lowest possible temperature.
- 3. Write the value of compressibility factor Z for ideal gas.
- 4. Write the unit of van der Waal constant which represent the magnitude of attractive forces between gas molecules.
- 5. Name the gas law which relates volume and pressure of gas at constant temperature.
- 6. Name the property responsible for spherical shape of water droplets.
- 7. Name the property which opposes the flow of liquids.
- 8. Two liquids A and B have vapour pressures 400 mm Hg and 450 mm Hg respectively at a given temperature. Which liquid has higher boiling point?
- Critical temperature of N₂ and O₂ are 126 K and 154.3 K respectively. 9. Which gas has greater magnitude of attractive forces?
- Mention the volume occupied by one mole of an ideal gas at STP. 10

3. Z = 1 4. L^2 atm mol⁻² 5. Boyle's law 2. −273°C Ans. 1. Pascal 6. Surface tension 7. Viscosity 8. Liquid A 9. O₂ 10. 22.7 L



States of Matter: Gases, Liquids and Solids

1-MARK QUESTIONS

- 1. Define Dalton's law of partial pressure of gases.
- 2. State Boyle's law.
- 3. Write van der Waal equation for n mol of gas.
- 4. Write the conditions in terms of temperature and pressure under which gases deviate from ideal behavior.
- 5. Write the relation between pressure and density of gas.
- 6. Write relation between average kinetic energy and temperature of a gas.
- 7. Define the term absolute zero.
- 8. In terms of Charles' law explain why -273°C is known as lowest temperature?
- 9. Write SI unit for quantity $(PV^2 T^2)/n^2$.
- 10. Define the term Critical temperature.
- 11. Define Boyle's temperature.
- 12. Define surface tension.
- 13. What is the value of normal boiling point and standard boiling point of water?
- 14 At a particular temperature vapour pressure of ethanol is more than that of water. Give reason.
- 15 Why vegetables are cooked with difficulty at a hill station?

2-MARKS QUESTIONS

- 1. Name the intermolecular forces present in: (i) H_2O (ii) HCl
- Critical temperature for carbon dioxide and methane are 31.1°C and -81.9°C respectively. Which of these has stronger intermolecular forces and why?
- 3. Explain the significance of van der Waal parameters.
- 4. A gas occupies 300 ml at 27°C and 730 mm pressure what would be its volume at STP. [Ans. 262.2 L]
- Calculate the temperature at which 28g N₂ occupies a volume of 10 litre at 2.46 atm. [Ans. 299.6 K]
- 6. Compressibility factor, Z of a gas is given as Z = PV/nRT
 - (i) What is the value of Z for an ideal gas?
 - (ii) For real gas, what is the value of Z above Boyle's temperature?

[**Ans.** (i) Z = 1 (ii) Z > 1]





- What will be the minimum pressure required to compress 500 dm³ of air at 1 bar to 200 dm³ at 30°C. [Ans. 2.5 bar]
- 8. Calculate the volume occupied by 8.8 g of CO_2 at 31.1 °C and 1 bar pressure. R = 0.083 L bar K⁻¹ mol⁻¹. [Ans. 5.05 L]
- 9. Calculate the temperature of 4 mol of a gas occupying in 5 dm³ at 3.32 bar. R = 0.083 bar dm³ K⁻¹ mol⁻¹. [Ans. 50K]
- 10 The pressure of the atmosphere is 2×10^{-6} mm at about 100 mile from the earth and temperature is -180° C. How many moles are there in 1 mL gas at this attitude? [Ans. 3.45×10^{-13} mol]
- 11. Calculate average kinetic energy of CO_2 molecules at 27°C.

[Ans. 3741.3 J mol⁻¹]

12. Calculate root mean square speed of methane molecules at 27°C.

[Ans. $6.84 \times 10^4 \text{ cm s}^{-1}$]

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- 13. Name two phenomena that can be explained on the basis of surface tension.
- 14. The van der Waal constants of two gases are as follows:

Gas	a (atm L mol ⁻¹)	b (L mol⁻¹)
А	1.39	0.0391
В	3.59	0.0427

Which of them is more easily liquefiable and which has greater molecular size?

- 15. Critical temperatures of NH_3 and SO_2 are 405.0 and 430.3 K respectively:
 - (i) Which one is easily liquefiable?
 - (ii) Which has higher value of van der Waal constant 'a'?
- 16. Arrange the following in the order of property indicated for each set:
 - (i) H₂O, NH₃, HCl, H₂ (increasing magnitude of intermolecular forces)
 - (ii) O₂, H₂, CO₂, SO₂ (ease of liquefaction)
 - (iii) O₂, He, CO₂, NH₃ (decreasing critical temperature)
 - (iv) O₂, He, CO₂, CH₄ (increasing value of van der Waal constant 'a')
- 17. Arrange Water, ethanol, ether and glycerine in the order of property given below:
 - (i) increasing order of vapour pressure
 - (ii) increasing order of boiling point
 - (iii) decreasing order of surface tension
 - (iv) increasing order of viscosity



States of Matter: Gases, Liquids and Solids

3-MARKS QUESTIONS

1. Explain the terms:

(i) Viscosity (ii) Vapour pressure (iii) Boiling point temperature

- 2. Calculate the total pressure in a mixture of 8 g of dioxygen and 4 g of dihydrogen confined in a vessel of 1 dm³ at 27°C. [Ans. 56.025 bar]
- 3. What will be the pressure exerted by a mixture of 3.2 g of methane and 4.4 g of carbon dioxide contained in a 9 dm³ flask at 27°C . [Ans. 0.82 atm]
- 4. Pressure of one gram of an ideal gas A at 27° C is found to be 2 bar. When 2 g of another ideal gas B is introduced in the same flask at same temperature, the pressure becomes 3 bar. Find the relationship between their molar masses. [Ans. $M_B = 4M_A$]
- 5. A 20g chunk of dry ice is placed in an empty 0.75 litre wire bottle tightly closed what would be the final pressure in the bottle after all CO_2 has been evaporated and temperature reaches to 25°C?

[Ans. Pressure inside the bottle = P + atm pressure = 14.828 + 1 = 15.828 atm]

- A gas at a pressure of 5 atm is heated from 0°C to 546°C and is simultaneously compressed to one third of its original volume. Find the final pressure of the gas. [Ans. 45 atm]
- 7. Calculate the compressibility factor for CO₂, if one mole of it occupies 0.4 litre at 300K and 40 atm. Comment on the result.

[Ans. 0.65, since Z < 1, CO₂ is more compressible than ideal gas]

8. Find the pressure of 4 g of O_2 and 2 g of H_2 confined in a bulb of 1 L at 0°C. [Ans. 25.215 atm]

5-MARKS QUESTIONS

- 1. Mention the intermolecular forces present between:
 - (i) H_2O and C_2H_5OH
 - (ii) Cl_2 and CCl_4
 - (iii) He and He atoms
 - (iv) Na^+ ion and H_2O
 - (v) HBr and HBr





- 2. (i) For Dalton's law of partial pressure derive the expression $P_{gas} = X_{gas} \cdot P_{total}$
 - (ii) A 2L flask contains 1.6 g of methane and 0.5 g of hydrogen at 27 °C. Calculate the partial pressure of each gas the mixture and calculate the total pressure.

[Ans.
$$P_{CH_4} = 1.23$$
 atm, $P_{H_2} = 3.079$ atm, $P_{total} = 4.31$ atm]

Using van der Waal's equation calculate the constant 'a' when 2 mole of a gas confined in a 4 L flask exerts a pressure of 11.0 atm at a temperature of 300K. The value of 'b' is 0.05 L mol⁻¹. [Ans. 6.49 atm L² mol⁻²]

HOTS QUESTIONS

1. A mixture of CO and CO_2 is found to have density of 1.50 g L⁻¹ at 20 °C and 740 mm pressure. Calculate the composition of mixture.

Solution:

Let the mol % of CO in mixture = x

- $\therefore \text{ mol } \% \text{ of } \text{CO}_2 = (100 \text{x})$ $\therefore \text{ Average molecular mass} = \frac{\left[(\text{x} \times 28) + (100 \text{x}) \times 24\right]}{100}$ $\therefore \frac{\left[(\text{x} \times 28) + (100 \text{x}) \times 24\right]}{100} = \frac{\text{dRT}}{\text{P}} \qquad \left(\text{Because, } \text{M} = \frac{\text{dRT}}{\text{P}}\right)$ $= \frac{(\text{x} \times 28) + (100 \text{x}) \times 24}{100} = 1.50 \text{gL}^{-1} \times 0.0821 \text{ L atm } \text{K}^{-1} \text{mol}^{-1} \times \frac{293 \text{K}}{100}$ $\therefore \text{ mol } \% \text{ of } \text{CO} = 43.38 \text{ and } \text{mol } \% \text{ of } \text{CO}_2 = (100 \text{x})$ = 100 43.38 = 56.62
- 2. A cylindrical balloon of 21 cm diameter is to be filled up with H_2 at NTP from a cylinder containing the gas at 20 atm at 27°C. The cylinder can hold 2.82 litre of water at NTP. Calculate the number of balloons that can be filled up.

Solution:

Volume of 1 balloon which has to be filled = $4/3 \pi (21/2)^3$

$$= 4851 \text{ mL} = 4.851 \text{ litre}$$

Let n balloons be filled, then volume of H_2 occupied by balloons = $4.851 \times n$ Also, cylinder will not be empty and it will occupy volume of $H_2 = 2.82$ litre.

- \therefore Total volume occupied by H₂ at NTP = 4.851 × n + 2.82 litre
- ∴ At STP

$$P_2 = 1 \text{ atm Available H}_2$$

$$V_1 = 4.851 \times n + 2.82$$

$$P_2 = 20 \text{ atm}$$

$$T_1 = 273 \text{ K}$$

$$T_2 = 300 \text{ K}$$

$$P_1 V_1 / T_2 = P_2 V_2 / T_2$$

$$V_2 = 2.82 \text{ litre}$$
or
$$1 \times (4.85 \text{ ln} + 2.82 / 273) = 20 \times 2.82 / 300$$
∴ n = 10

3. Calculate the temperature at which CO_2 has the same rms speed to that of O_2 at STP.

Solution:

rrms of O₂ = $\sqrt{3}$ RT / M at STP, urms of O₂ = $\sqrt{3}$ R × 273 / 32

For CO₂ urms CO₂ = $\sqrt{3}$ RT / 44

Given both are same; $3R \times 273 / 32 = 2RT / 44$

∴ T =375.38 K = 102.38°C

4. 50 litre of dry N_2 is passed through 36g of H_2O at 27°C. After passage of gas, there is a loss of 1.20g in water. Calculate vapour pressure of water.

Solution

The water vapours occupies the volume of N₂ gas i.e. 50 litre

- :. For H₂O vapour V = 50 litre, w = 1.20g, T = 300K, m = 18 g mol⁻¹ PV = w / m RT or P × 50 = 1.2 / 18 × 0.0821 × 300
- \therefore P = 0.03284 atm = 24.95 mm





UNIT TEST

Maximum Marks : 20

Time Allowed: 1 hr

General Instructions:

(i) All questions are compulsory.

(ii) Maximum marks carried by each question are indicated against it.

1.	If P is the pressure and d is the density of gas, then P and d are related as			l as:
		$P \alpha 1/d$	(b) $P \alpha d$	[1]
	(c) I	$P \alpha d^2$	(d) $P \alpha 1/d^2$	
2.	At w	At which temperature the volume of a gas is expected to be z		
	(a) 0	°C	(b) 273 K	
	(c) –	273 °C	(d) 273°C	
3.	Defi	ne Boyle's temperature.		[1]
4.	Write van der Waal equation for n mol of gas.		[1]	
5.	Why vegetables are cooked with difficulty at a hill station?		[1]	
6.	A gas occupies 300 ml at 27°C and 730 mm pressure what would be its volume at STP.			[2]
7.	Compressibility factor, Z of a gas is given as $Z = PV/nRT$			[2]
	(i)	What is the value of Z for an ide	-	
	(ii)	For real gas, what is the value of	•	
8.	Pressure of one gram of an ideal gas A at 27°C is found to be 2 bar. [3] When 2 g of another ideal gas B is introduced in the same flask at same temperature, the pressure becomes 3 bar. Find the relationship between their molar masses.			[3]
9.		ulate the total pressure in a mixtur	e .e	[3]
	4 g of dihydrogen confined in a vessel of 1 dm^3 at 27°C.			
	$\mathbf{R} = 0$	$0.083 \text{ bar } \text{dm}^3 \text{ K}^{-1} \text{ mol}^{-1}$		
10.	(a)	Critical temperatures of NH ₃ and respectively:	1 SO ₂ are 405.0 and 430.3 K	[2]
		(i) Which one is easily liquefia	ble?	
		(ii) Which has higher value of v	van der Waal constant 'a'?	
	(b)	Explain the terms:		[3]
	(i) Viscosity (ii) Vapour pressure (iii) Boiling point temperature			e



States of Matter: Gases, Liquids and Solids

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Chemical Thermodynamics

FAST TRACK : QUICK REVISION

- System: Specific part of universe in which observations are made.
- Surroundings: Everything which surrounds the system.

Chapter - 6

• Types of the System:

Open System: Exchange both matter and energy with the surroundings. For example – Reactant in an open test tube.

Closed System: Exchange energy but not matter with the surroundings. For example – Reactants in a closed vessel.

Isolated System: Neither exchange energy nor matter with the surroundings. For example – Reactants in a thermos flask.

Thermodynamic Processes:

- (i) Isothermal Process: $\Delta T = 0$
- (ii) Adiabatic process: $\Delta q = 0$
- (iii) Isobaric process: $\Delta P = 0$
- (iv) Isochoric process: $\Delta V = 0$
- (v) Cyclic process: $\Delta U = 0$
- (vi) Reversible process: Process which proceeds infinitely slowly by a series of equilibrium steps.
- (vii) Irreversible process: Process which proceeds rapidly and the system does not have chance to achieve equilibrium.
- Extensive Properties: Properties which depend upon the quantity or size of matter present in the system. For example mass, volume, internal energy, enthalpy, heat capacity, work etc.

- Intensive Properties: Properties which do not depend upon the quantity or size of matter present in the system. For example temperature, density, pressure, surface tension, viscosity, refractive index, boiling point, melting point etc.
- **State Functions**: The variables of functions whose value depend only on the state of a system or they are path independent.

For example – pressure (P), volume (V), temperature (T), enthalpy (H), free energy (G), internal energy (U), entropy (S), amount (n) etc.

- **Internal Energy**: It is the sum of all kind of energies possessed by the system.
- First Law of Thermodynamics: "The energy of an isolated system is constant."

Mathematical Form: $\Delta U = q + w$

- Sign Conventions for Heat (q) and Work (w):
 - (i) W = +ve, if work is done on system
 - (ii) W = -ve, if work is done by system
 - (iii) q = +ve, if heat is absorbed by the system
 - (iv) q = -ve, if heat is evolved by the system
- Work of Expansion/ compression: $\omega = -P_{ext} (V_f V_j)$
- Work done in Isothermal Reversible Expansion of an Ideal Gas:

 $\omega_{rev} = -2.303 \text{ nRT} \log V_f / V_i$

or $\omega_{rev} = -2.303 \text{ nRT} \log P_i / P_f$

- Significance of $\Delta H \& \Delta U$: $\Delta H = q_p$ and $\Delta U = q_v$
- **Relation between** $\Delta H \& \Delta U$: $\Delta H = \Delta U + (n_p n_r)RT$ for gaseous reaction
 - (i) $\Delta H = \Delta U$ if $(n_n n_r)$ is zero; e.g. $H_2(g) + I_2(g) \longrightarrow 2$ HI(g)
 - (ii) $\Delta H \ge \Delta U$ if $(n_p n_r)$ is positive; e.g. $PCl_5(g) \longrightarrow PCl_3(g) + Cl_2(g)$
 - (iii) $\Delta H \leq \Delta U$ if $(n_p n_r)$ is negative; *e.g.* $N_2(g) + 3H_2(g) \longrightarrow 2NH_3(g)$
- Heat Capacity (C): Amount of heat required to raise the temperature of a substance by 1°C or 1 K.

 $q = C \Delta T$

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• Specific Heat Capacity (C_s): Amount of heat required to raise the temperature of 1 g of a substance by 1°C or 1 K.

 $q = C_s \times m \times \Delta T$

• Molar Heat Capacity (C_m): Amount of heat required to raise the temperature of 1 mole of a substance by 1°C or 1 K.

$$q = C_m \times n \times \Delta T$$

- Standard State of a Substance: The standard state of a substance at a specified temperature is its pure form at 1 bar.
- Standard Enthalpy of Formation(Δ_fH⁰): Enthalpy change accompanying the formation of one mole of a substance from its constituent elements under standard condition of temperature (normally 298 K) and pressure (1bar).
 - $\Delta_f H^0$ of an element in standard state is taken as zero.
 - Compounds with –ve value of $\Delta_f H^0$ more stable than their constituents.
 - $\Delta_r H^0 = \sum a_i \Delta_f H^0$ (products) $-\sum b_i \Delta_f H^0$ (reactants); Where 'a' and 'b' are coefficients of products and reactants in balanced equation.
- Standard Enthalpy of Combustion(Δ_cH⁰): Enthalpy change accompanying the complete combustion of one mole of a substance under standard conditions (298 K, 1bar)
- Hess's Law of Constant Heat Summation: The total enthalpy change of a reaction remains same whether it takes place in one step or in several steps.
- **Bond Dissociation Enthalpy:** Enthalpy change when one mole of a gaseous covalent bond is broken to form products in gas phase.

For example, $\operatorname{Cl}_2(g) \longrightarrow 2\operatorname{Cl}(g); \Delta_{\operatorname{Cl-Cl}} H^0 = 242 \text{ kJ mol}^{-1}$

- For diatomic gaseous molecules; Bond enthalpy = Bond dissociation Enthalpy = Atomization Enthalpy
- For Polyatomic gaseous molecules; Bond Enthalpy = Average of the bond dissociation enthalpies of the bonds of the same type.
- $\Delta_{\rm r} {\rm H}^0 = \sum \Delta_{\rm bond} {\rm H}^0 ({\rm Reactants}) \sum \Delta_{\rm bond} {\rm H}^0 ({\rm Products})$
- **Spontaneous Reaction:** A reaction which can take place either an its own or under some initiation.





• Entropy(S): It is measure of degree of randomness or disorder of a system.

•
$$\Delta S_{sys} = \frac{(q_{rev})_{sys}}{\Delta T} = \frac{\Delta H_{sys}}{\Delta T}$$

- Unit of Entropy = JK^{-1} mol⁻¹
- Second Law of Thermodynamics: For all the spontaneous processes totally entropy change must positive.

$$\Delta S_{total} = \Delta S_{sys} + \Delta S_{surr} > 0$$

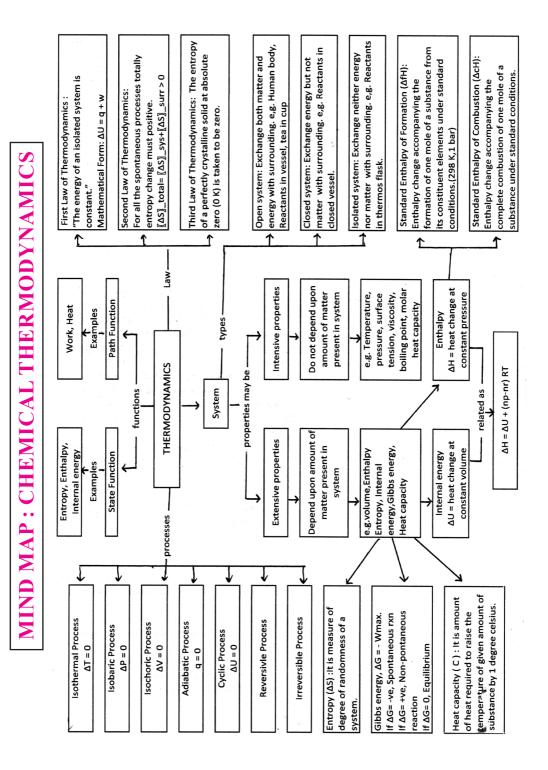
• Gibbs Helmholtz Equation for determination of Spontaneity:

$\Delta G = \Delta H - T \Delta S$

- (i) If $\Delta G = -ve$, the process is spontaneous
- (ii) If $\Delta G = +ve$, the process is non-spontaneous
- (iii) If $\Delta G = 0$, the process is in equilibrium
- Relation between Gibbs Energy Change and Equilibrium Constant: $\Delta G^{o} = -2.303 \text{ RT } \log K_{c}$
- Third Law of Thermodynamics: The entropy of a perfectly crystalline solid at absolute zero (0 K) is taken to be zero.



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MULTIPLE CHOICE QUESTIONS (MCQ)

- 1. Which one of the following thermodynamic quantities is not a state function?
 - (a) Gibbs free energy (b) Enthalpy
 - (c) Entropy (d) Work
- 2. All of the following have a standard heat of formation value of zero at 25°C and 1.0 atm except:
 - (a) $N_2(g)$ (b) Fe(s)
 - (c) Ne (g) (d) H(g)
- 3. For the following reaction at 25°C, $\Delta H^{\circ} = +115 \text{ kJ}$ and $\Delta S^{\circ} = +125 \text{ J/K}$. Calculate ΔG° for the reaction at 25°C :
 - $SBr_4(g) \longrightarrow S(g) + 2Br_2(l)$ (a) +152 kJ (b) -56.7 kJ

(c)
$$+77.8$$
 kJ. (d) $+37.1$ kJ

4. Calculate $\Delta_r H^0$ for the following reaction at 25°C :

	$Fe_3O_4(s)$	+ CO(g)	\longrightarrow	3FeO(s	s) + $CO_2(g)$
ΔH_{f}^{o} (kJ/mol)	-1118	-110.5	-2	72	-393.5
(a) -263 kJ			(b)	54 kJ	
(c) 19 kJ.			(d)	-50 kJ	ſ

- 5. A system suffers an increase in internal energy of 80 J and at the same time has 50 J of work done on it. What is the heat change of the system?
 - (a) +130 J (b) +30 J(c) -130 J (d) -30 J
- 6. The ΔH^0 for the following reaction at 298 K is -36.4 kJ.

 $1/2 \text{ H}_2(g) + 1/2 \text{ Br}_2(l) \longrightarrow \text{HBr}(g)$

Calculate ΔU^0 at 298 K. The universal gas constant, R, is 8.314 J/mol K.

- (a) -35.2 kJ (b) +35.2 kJ
- (c) -36.4 kJ (d) -37.6 kJ.
- 7. For which of the following reactions would the Ho for the reaction be labeled ΔH_f^{0} ?
 - (a) $Al(s) + 3/2 H_2(g) + 3/2 O_2(g) \longrightarrow Al(OH)_3(s)$
 - (b) $PCl_3(g) + 1/2 O_2(g) \longrightarrow POCl_3(g)$

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	(c) $1/2 N_2 O(g) + 1/4 O_2(g) \longrightarrow NO(g)$					
	(d) $\operatorname{CaO}(s) + \operatorname{SO}_2(g) \longrightarrow \operatorname{CaSO}_3(s)$					
8.	Which statement is ture for reaction? $2H_2(g) + O_2(g) \longrightarrow 2H_2O(g)$					
01	(a) $S = +ve$ (b) $H > U$					
	(c) $H < U$ (d) $H = U$					
9.	The heat of combustion of yellow phosphorous is –9.91 KJ and the red					
	phosphorous is -8.78 KJ. The heat of transition of yellow phosphorous to					
	red phosphorous is :					
	(a) -9.91 kJ (b) -8.78 kJ					
	(c) -9.34 kJ (d) -1.13 kJ					
10.	Entropy of universe is :					
	(a) Increasing (b) decreasing					
	(c) Constant (d) None of these					
11.	Which is state function?					
	(a) q (b) w					
	(c) $q + w(d)$ None of these					
12.	According to second law of thermodynamics					
	(a) $\Delta S_{total} = +ve$ (b) $\Delta S_{total} = -ve$					
	(c) $\Delta S_{system} = +ve$ (d) $\Delta S_{system} = -ve$					
Ans:	1.(d), 2.(d), 3.(c), 4.(c), 5.(b), 6.(d), 7.(a),					
	8. (a), 9.(d), 10.(c), 11.(d), 12.(a), 13.(c), 14.(a)					
	FILL IN THE BLANKS					
(i)						
(ii)	A process which can take place either of its own or under some initiation					
(;;;)	is known as					
(iii)	For evaporation of water the sign of ΔH is and sign of ΔS is					
(iv)	The entropy of a perfectly crystalline solid is zero at					
(v)	The heat energy exchanged between the system and surroundings at constant temperature and pressure is known as					
(vi)						
(1)	is the quantity of heat needed to raise the temperature of one mole of a substance by 10					
(vii)	i) $C_p - C_v = \dots$					
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(viii) $\ldots = \Delta H - T\Delta S$.

- (ix) According to law of thermodynamics, $\Delta S_{total} = +$ ve.
- (x) If $\Delta H = +ve$ and $\Delta S = +ve$, the reaction is spontaneous at temperature
- **Ans:** (i) Entropy (ii) spontaneous (iii) +ve, +ve (iv) -273°C
 - (v) Enthalpy (vi) molar heat capacity (vii) R (viii) ΔG
 - (ix) second (x) high

TRUE AND FALSE TYPE QUESTIONS

Write true or false for following statements:

- (i) For every chemical reaction at equilibrium ΔG^0 is zero.
- (ii) Entropy is not a state function because its value depends upon the condition of temperature and pressure.
- (iii) During isothermal expansion of an ideal gas, there is no change in internal energy.
- (iv) q and w are not state function but q+w is a state function.
- (v) The enthalpy of neutralization of a strong acid by a strong base is always constant.
- (v) For a spontaneous process $\Delta S_{system} = +ve$.
- (vii) The energy of universe is conserved while its entropy is increasing.
- (viii) Volume is extensive property while temperature is intensive property.
- (ix) At 0°C the entropy of a perfectly crystalline solid is zero.
- (x) Hess' law is a corollary of the first law of thermodynamics.
- Ans: (i) False(ii) False(iii) True(iv) True(v) True(vi) False(vii) True(viii) True(ix) False(x) True

MATCH THE COLUMNS

(a)

1. Column I

Column II

Pressure

- (i) State function
- (ii) Extensive property (b) Work
- (iii) Intensive property
- (iv) Path function
- (c) q + w(d) Heat capacity



Chemical Thermodynamics



Column I

Column II

Non-spontaneous at all temperatures

- $\Delta H = +ve; \Delta S = +ve$ Spontaneous at all temperatures (i) (a)
- (ii) $\Delta H = -ve; \Delta S = +ve$ (b)
- (iii) $\Delta H = +ve; \Delta S = -ve$
- (c) Non-spontaneous at high temperature (iv) $\Delta H = -ve; \Delta S = -ve$ Spontaneous at high temperature (d)
- Ans: 1. (i) \rightarrow (c), (ii) \rightarrow (d), (iii) \rightarrow (a), (iv) \rightarrow (b).

2. (i)
$$\rightarrow$$
 (d), (ii) \rightarrow (a), (iii) \rightarrow (b), (iv) \rightarrow (c).

ASSERTION AND REASON TYPE QUESTIONS

In the following questions a statement of assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given below for each question.

- (i) A and R both are correct, and R is correct explanation of A.
- A and R both are correct, but R is not the correct explanation of A. (ii)
- (iii) A is true but R is false.
- (iv) A and R both are false.
 - 1. Assertion (A): Enthalpy of graphite is lower than that of diamond. Reason (R): Entropy of graphite is greater than that of diamond.
 - Assertion (A): Enthalpy of formation of $H_2O(l)$ is greater than that of 2. $H_2O(g)$.

Reason (R): Enthalpy change is negative for condensation reaction, $H_2O(g)$ $H_2O(1)$

- Assertion (A): ΔH and ΔU are same for the reaction $N_2(g) + O_2(g) \longrightarrow 2NO(g)$ 3. Reason (R): All the reactants and products are gases.
- Assertion (A): if both ΔH^0 and ΔS^0 are positive than the reaction will be 4. spontaneous at high temperature Reason (R): All processes with positive entropy change are spontaneous.
- Assertion (A): Enthalpy of formation of HCl is equal to bond energy of 5. HC1.

Reason (R): Enthalpy of formation and bond energy both involve the formation of one mole of HCl from the elements.

Ans: 1. (ii) 2. (i) 3. (ii) 4. (iii) 5. (i)





ONE WORD ANSWER TYPE QUESTIONS

- 1. 'w' amount of work is done by the system and 'q' amount of heat is supplied to the system. What type of system would it be?
- 2. What is the work done in free expansion of an ideal gas?
- 3. What is the sign of ΔG^{Θ} for spontaneous reaction?
- 4. Write the relation between ΔH and ΔU for $H_2(g) + I_2(g) \rightarrow 2HI(g)$.
- 5. Write the SI unit of entropy.
- 6. Name the calorimeter used to measure ΔU .
- 7. What is the standard enthalpy of formation of graphite?
- 8. What is the sign of ΔH for $H_2(g) \longrightarrow 2H(g)$?
- 9. If $K_c = 1$, what will be the value of ΔG ?
- 10. An exothermic reaction is spontaneous at all temperature. What is the sign of S?
- Ans: 1. Closed system2. W = 03. $\Delta G = -ve$ 4. $\Delta H = \Delta U$ 5. J K⁻¹ mol⁻¹6. Bomb calorimeter7. Zero8. $\Delta H = -ve$ 9. Zero10. $\Delta S = +ve$

1-MARK QUESTIONS

- Day temp Name the thermodynamic system to which following belong:
 (i) Human body (ii) Milk in Thermos flask (iii) Tea in steel kettle
- 2. Identify State functions out of the following: Enthalpy, Entropy, Heat, Temperature, Work, Free energy
- 3. Give two examples of state functions.
- 4. Write the mathematical statement of first law of thermodynamics.
- 5. Predict the internal energy change for an isolated system? [Ans. Zero]
- 6. Why ΔH is more significant than ΔU ?
- 7. Write one example each of extensive and intensive properties.
- 8. Write a chemical equation in which ΔH and ΔU are equal.
- 9 Write the relationship between ΔH and ΔU for the reaction:

$$C(s)+O_2(g) \longrightarrow CO_2(g)$$

10. Define standard enthalpy of formation.



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- 11. Why is the standard enthalpy of formation of diamond not zero although it is an element?
- 12. The enthalpy of atomization of CH_4 is 1665 kJ mol⁻¹. What is the bond enthalpy of C – H bond? [Ans. 416.25 kJ]
- 13. Identify the species for which $\Delta_f H^{\theta} = O_3$, at 298 K; Br₂, Cl₂, CH₄. [Hint: Cl₂ (Br₂ is liquid at 298K)]
- 14. For the reaction $2Cl(g) \longrightarrow Cl_2(g)$; what are the sign of ΔH and ΔS ?
- 15. For an isolated system $\Delta U= 0$, what will be ΔS ?
- 16. Why entropy of steam is more than that of water at its boiling point?
- 17. Out of Diamond and Graphite which has higher entropy?
- 18. Write an example of endothermic spontaneous reaction.
- 19. State second law of thermodynamics.
- 20. State third law of thermodynamics.
- 21. Which has more entropy? 1 mol H_2 O(1) at 25°C or 1 mol H_2 O(1) at 35°C.
- 22. At what temperature the entropy of a perfectly crystalline solid is zero?
- 23. For a certain reaction $\Delta G^{\theta} = 0$, what is the value of K_c?
- 24. How can a non spontaneous reaction be made spontaneous?
- 25. For a reaction both ΔH and ΔS are negative. Under what conditions does the reaction occur.

2-MARKS QUESTIONS

1. In a process 701 J of heat is absorbed by a system and 394 J work is done by the system. What is the change in internal energy for the process?

[Ans. 307 J]

- 2. Neither q nor w is state functions but q + w is a state function. Explain.
- 3. Classify the following as extensive or intensive properties :-Heat capacity, Density, Temperature, Molar heat capacity.
- 4. Derive the relationship between ΔH and ΔU .
- 5. Derive the relationship $C_p C_v = R$.
- 6. A 1.25g sample of octane (C_8H_{18}) is burnt in excess of oxygen in a bomb calorimeter. The temperature of the calorimeter rises from 294.05 to





300.78K.If heat capacity of the calorimeter is 8.93 kJ K⁻¹. Find the heat transferred to calorimeter. [Ans. 0.075 kJ]

- 7. Show that for an ideal gas, the molar heat capacity under constant volume conditions is equal to 3/2 R.
- 8. Expansion of a gas in vacuum is called free expansion. Calculate the work done and change in internal energy when 1 mol of an ideal gas expands isothermally from I L to 5 L into vacuum.
- 9. State and explain Hess's Law of Constant Heat Summation with a suitable example.
- 10. Derive the relationship between ΔH and ΔU . Given, N₂(g) + 3H₂(g) \longrightarrow 2NH₃(g); $\Delta_r H^{\theta} = -92.4 \text{ kJ mol}^{-1}$; What is the standard enthalpy of formation of NH₃ gas? [Ans. -46.2 kJ mol⁻¹]
- 11. Calculate the enthalpy change for the reaction:

 $\mathrm{H}_{2}(\mathrm{g}) + \mathrm{Br}_{2}(\mathrm{g}) \longrightarrow 2 \ \mathrm{HBr}(\mathrm{g}).$

Given the bond enthalpies H_2 , Br_2 and HBr are 435 kJ mol⁻¹, 192 kJ mol⁻¹ and 368 kJ mol⁻¹ respectively. [Ans. -109 kJ mol⁻¹]

- Is the bond dissociation enthalpy of all the four C H bonds in CH₄ same? Give reason in support of your.
- 13. Define the term entropy. Write its unit. How does entropy of a system change on increasing temperature?
- 14. Dissolution of ammonium chloride in water is endothermic but still it dissolves in water readily. Why?
- 15. Calculate the entropy change in the surroundings when 1.00 mol of $H_2O(l)$ is formed under standard conditions; $\Delta_f H^{\theta} = -286 \text{ kJ mol}^{-1}$.

- 16. The enthalpy of vaporization of a liquid is 30 kJ mol⁻¹ and entropy of vaporization is 75 J K⁻¹ mol⁻¹. Calculate the boiling point of liquid at 1 atm. [Ans. 400 K]
- 17. The equilibrium constant for a reaction is 10. What will be the value of ΔG^{θ} ? R = 8.314J K⁻¹ mol⁻¹, T = 300K. [Ans. -5.527 kJ mol⁻¹]
- 18. Derive the relationship, $\Delta G = -T\Delta S_{total}$ for a system.

[[]**Ans.** 959.7 J K⁻¹ mol⁻¹]

19. The ΔH and ΔS for $2Ag_2 O(s) \longrightarrow 4Ag(s) + O_2(g)$ are given + 61.17kJ mol⁻¹ and +132 J K⁻¹mol⁻¹ respectively. Above what temperature will the reaction be spontaneous? [Ans. >463.4 K]

3-MARKS QUESTIONS

- 1. Differentiate between the following (with examples)
 - (i) Open and Closed System.
 - (ii) Adiabatic and Isothermal process.
 - (iii) State function and path function
- Calculate the maximum work obtained when 0.75 mole of an ideal gas expands isothermally and reversibly at 27°C from a volume of 15L to 25L. [Ans. -955.7 J]
- Calculate the number of kJ necessary to raise the temperature of 60 g of aluminium from 35 to 55°C. Molar heat capacity of Al is 24 J mol⁻¹ K⁻¹.
 [Ans. 1.067 kJ]
- 4. The reaction of cyanamide, NH_2CN (s), with Dioxygen was carried out in a bomb calorimeter, and ΔU was found to be -742.7 kJ mol⁻¹ at 298K. Calculate Enthalpy change for the reaction at 298K,

- The enthalpy of combustion of methane, graphite and dihydrogen at 298 K are -890.3 kJ mol⁻¹, -393.5 kJ mol⁻¹ and -285.8 kJ mol⁻¹ respectively. Calculate enthalpy of formation of methane gas. [Ans. -74.8 kJ mol⁻¹]
- 6. Explain the Born Haber Cycle to determine the lattice enthalpy of NaCl.
- 7. Enthalpies of formation of CO(g), CO₂(g), N₂O(g) and N₂O₄ (g) are -110, -393, 81 and 9.7 kJ mol-1 respectively. Find the value of $\Delta_r H$ for the reaction; N₂O₄(g) + 3 CO (g) \longrightarrow N₂O (g) + 3CO₂ (g)

 $[Ans. -777.7 \text{ kJ mol}^{-1}]$

8. The combustion of 1 mol of benzene takes place at 298K .After combustionCO₂ and H₂O are formed and 3267 kJ mol⁻¹ of heat is liberated. Calculate $\Delta_{\rm f}$ H⁰ (C₆H₆).

Given: $\Delta_{f} H^{0}(CO_{2}) = -286 \text{ kJ mol}^{-1}, \Delta_{f} H^{0}(H_{2}O) = -393 \text{ kJ mol}^{-1}$ [Ans. 48.51 kJ mol}^{-1}]





9. Calculate the standard enthalpy of formation of CH₂OH(1) from the following data:

- For oxidation of iron, 4 Fe(s) + 3 $O_2(g) \rightarrow 2$ Fe₂ $O_3(s)$ entropy change is 10. -549.4 J K⁻¹ mol⁻¹ at 298 K. In spite of negative entropy change of this reaction, why is the reaction spontaneous? ($\Delta_r H^{\theta}$ for this reaction is -1648 $[Ans. \Delta S_{total} = +4980.6 \text{ J } \text{K}^{-1} \text{ mol}^{-1}]$ $kJ mol^{-1}$)
- 11. Give reasons:
 - Evaporation of water is and endothermic process but it is spontaneous. (i)
 - (ii) A real crystal has more entropy than an ideal crystal.
 - (iii) Entropy of universe is increasing.
- For the reaction at 298 K, $2A + B \longrightarrow C$; $\Delta H = 400 \text{ kJ mol}^{-1}$, $\Delta S = 0.2$ 12. kJ K⁻¹ mol⁻¹. At what temperature will the reaction become spontaneous considering ΔH and ΔS to be constant over the temperature range.

[Ans. T > 2000 K]

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- Reaction $X \rightarrow Y$; $\Delta H = +ve$ is spontaneous at temperature 'T'. Determine 13.
 - Sign of ΔS for this reaction. (i)
 - (ii) Sign of ΔG for $Y \longrightarrow X$
 - (iii) Sign of ΔG at a temperature < T

5-MARKS QUESTIONS

- 1. (a) What is reversible process in Thermodynamics?
 - (b) Name the thermodynamic processes for which : (i) q = 0 (ii) $\Delta U = 0$ (iii) $\Delta V = 0$ (iv) $\Delta P = 0$
 - (c) Water decomposes by absorbing 286.2 kJ of electrical energy per mole. When H₂ and O₂ combine to form one mole of H₂O, 286.2 kJ of heat is produced. Which thermodynamic law is proved? Write its statement.
- (a) Although heat is a path function but heat absorbed by the system under 2. certain specific conditions is independent of path. What are those conditions? Explain. [Hint: $q_v = \Delta U$ and $q_n = \Delta H$]

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- (b) It has been found that 221.4 J is needed to heat 30g of ethanol from 15°C to 18°C. Calculate (a) specific heat capacity, and (b) molar heat capacity of ethanol. [Ans. (a) $2.46 \text{ Jg}^{-1} \text{ °C}^{-1}$, (b) $113.2 \text{ Jmol}^{-1} \text{ °C}^{-1}$]
- 3. (a) Differentiate the terms Bond dissociation enthalpy & Bond Enthalpy.
 - (b) Calculate enthalpy change for the process $CCl_4(g) \longrightarrow C(g) + 4Cl(g)$ and calculate Bond enthalpy of C–Cl bond in CCl_4 . Given: $\Delta_{vap} H^{\theta} = 30.5 \text{ kJ mol}^{-1}$; $\Delta_f H^{\theta}(CCl_4) = -135.5 \text{ kJ mol}^{-1}$; $\Delta_a H^{\theta}(C) = 715 \text{ kJ mol}^{-1}$ and $\Delta_a H^{\theta}(Cl_2) = 242 \text{ kJ mol}^{-1}$ [**Ans.** 1304 kJ mol}^{-1}, 326 kJ mol}^{-1}]
- 4. Predict the sign of ΔS for the following changes:
 - (i) Freezing of water.
 - (ii) $C(graphite) \longrightarrow C(diamond)$
 - (iii) $H_2(g)$ at 298 k and 1 bar $\longrightarrow H_2(g)$ at 298 k and 10 bar
 - (iv) $H_2(g) + I_2(g) \longrightarrow 2HI(g)$
 - (v) $2NaHCO_3(s) \longrightarrow Na_2CO_3(s) + CO_2(g) + H_2O(g)$
- 5. (i) Define Gibbs Energy. Give its mathematical expression. What is Gibb's energy criteria of Spontaneity.
 - (ii) For the reaction:

 $2A(g) + B(g) \rightarrow 2D(g), \Delta U^{\theta} = -10.5 \text{ kJ and } \Delta S^{\theta} = -44.1 \text{ J K}^{-1}.$

Calculate ΔrG° for the reaction, and predict whether the reaction will occur spontaneously.

[Ans. $\Delta_r G^{\theta} = +0.16 \text{ kJ}$, Non-spontaneous]

HOTS QUESTIONS

1. Does entropy increase or decrease when egg is boiled?

Ans.:On boiling egg, entropy decreases as due to denaturation, the helical structure of protein become more complicated and random coiled structure.

2. 10 g of argon is compressed isothermally and reversibly at a temperature of 27°C from 10 L to 5 l. Calculate q, w, ΔU and ΔH .

Solution: q = -2.303 nRT log V₂ / V₁ = -2.303 × 10/40 mol × 2 Cal K⁻¹ mol⁻¹ × 300 K × log 5/10 = -103.635 Cal





For isothermal compression $\Delta U = 0$

$$W = \Delta U - q = 0 - (-103.635) = +103.635$$
 Cal

Also when temperature is constant,

 $PV = constant, \Delta H = \Delta U + \Delta (PV) = 0 + 0 = 0$

3. 1 mole of an ideal gas expand isothermally and reversibly from a pressure of 10 atm to 1 atm at 300 K. Calculate the height to which an object of 50 kg can be lifted by this expansion.

Solution: $w_{(exp.)} = -2.303 \text{ nRT}$ log $P_i / P_f = -2.303 \times 1 \text{ mol} \times 8.314 \text{ J K}^{-1} \text{mol}^{-1} \times 300 \text{ K} \times \log 10/1$ $= 5.74 \times 10^3 \text{ J}$

Now, mgh = 5.74×10^3 J or $50 \text{ kg} \times 9.81$ m s⁻² × h = 5.744×10^3 J \therefore h = 11.7 m



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UNIT TEST

Time Allowed: 1 hr

Maximum Marks : 20

General Instructions:

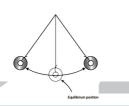
- (i) All questions are compulsory.
- (ii) Maximum marks carried by each question are indicated against it.
 - 1. For the reaction $2Cl(g) \longrightarrow Cl_2(g)$; what are the sign of ΔH and ΔS ? [1]
 - 2. Write an example of endothermic spontaneous reaction. [1]
 - 3. 'w' amount of work is done by the system and 'q' amount of heat [1] is supplied to the system. What type of system would it be?
 - 4. In a process 701 J of heat is absorbed by a system and 394 J work [2] is done by the system. What is the change in internal energy for the process?
 - 5. State and explain Hess's Law of Constant Heat Summation with [2] a suitable example.
 - 6. Calculate the number of kJ necessary to raise the temperature of [3]
 60 g of aluminium from 35 to 55°C. Molar heat capacity of Al is 24 J mol⁻¹ K⁻¹.
 - 7. Calculate the standard enthalpy of formation of CH3OH (l) from the following data:

$$\begin{split} & \operatorname{CH}_{3}\operatorname{OH}(l) + 3/2 \operatorname{O}_{2}(g) \longrightarrow \operatorname{CO}_{2}(g) + 2\operatorname{H}_{2}\operatorname{O}(l); \\ & \Delta_{c} \operatorname{H}^{\theta} = -726 \text{ kJ mol}^{-1} \\ & \operatorname{C}(g) + \operatorname{O}_{2}(g) \longrightarrow \operatorname{CO}_{2}(g); \Delta_{f} \operatorname{H}^{\theta} = -393 \text{ kJ mol}^{-1} \\ & \operatorname{H}_{2}(g) + \frac{1}{2} \operatorname{O}_{2}(g) \longrightarrow \operatorname{H}_{2}\operatorname{O}(l); \Delta_{f} \operatorname{H}^{\theta} = -286 \text{ kJ mol}^{-1} \end{split}$$

- 8. (a) For oxidation of iron, 4 Fe(s) + 3 O₂(g) → 2 Fe₂O₃(s) entropy change is -549.4 J K⁻¹ mol⁻¹ at 298 K. In spite of negative entropy change of this reaction, why is the reaction spontaneous? (Δr H° for this reaction is -1648 kJ mol⁻¹) [2]
 - (b) For the reaction: 2A (g) + B (g) \longrightarrow 2D (g), $\Delta U^{\circ} = -10.5$ kJ and $\Delta S^{\circ} = -44.1$ J K⁻¹. Calculate ΔrG° for the reaction, and predict whether the reaction will occur spontaneously. [3]







Equilibrium

FAST TRACK : QUICK REVISION

- Equilibrium : It is a state in a process when two opposing processes (forward and reverse) occur simultaneously at the same rate. The free energy change at equilibrium state is zero *i.e.*, $\Delta G = 0$.
- Equilibrium constant : For a general reaction :

Chapter - 7

$$aA + bB \iff cC + dD$$

 $K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b} \text{ and } K_p = \frac{P_C^c \times P_D^d}{P_A^a \times P_B^b}$

Relationship between K_p and K_c:

$$\begin{split} \mathbf{K}_p &= \mathbf{K}_c \, (\mathrm{RT})^{\Delta n_g} \\ \Delta n_g &= n_p(g) - n_r(g) \end{split}$$

• Magnitude of equilibrium constant depends upon the way in which a reaction is written :

Chemical equation	Equilibrium constant
$aA + bB \iff cC + dD$	K
$cC + dD \implies aA + bB$	$K_1 = \frac{1}{K}$
$naA + nbB \implies ncC + ndD$	$K_2 = K^n$
$\frac{1}{n}aA + \frac{1}{n}bB \xrightarrow{\longrightarrow} \frac{1}{n}cC + \frac{1}{n}dD$	$K_3 = K^{1/n}$

• Predicting the direction of reaction :

- If $Q_c = K_c \Rightarrow$ The reaction is in a state of equilibrium.
 - $Q_c > K_c \Rightarrow$ The reaction proceeds in reverse direction.

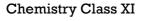
 $Q_c < K_c \Rightarrow$ The reaction proceeds in forward direction.

- Ostwald's dilution law : Degree of dissociation of weak electrolyte, $\alpha = \sqrt{\frac{K}{C}}$
- Ionic Product of water $(K_{\mu}) = [H_3O^+] [OH^-] = 10^{-14}$ at 298K
- Le-Chatelier's Principle : When a system of equilibrium is subjected to a change in temperature, pressure or concentration, the equilibrium shifts itself in such a way so as to undo or nullify the effect of change.

Change at equilibriumShift in equilibriumIncrease in temperatureEndothermic directionDecrease in temperatureExothermic directionIncrease in pressureTowards lesser gaseous molesDecrease in pressureTowards greater gaseous molesIncrease in Conc. of reactantsForward directionIncrease in Conc. of productsReverse direction

• Outcomes of Le-Chatelier's Principle

- Conjugate Acid or Base : Acid-base pair which differ by H⁺ ion.
 Species H⁺ = Conjugate base
 Species + H⁺ = Conjugate acid
- pH of solution : pH = -log [H₃O⁺] or [H⁺] = 10^{-pH}, pOH = -log [OH⁻] pH + pOH = pK_w = 14 at 298K
- **Common ion effect** : The depression of ionisation of weak electrolyte by the presence of common ion from a strong electrolyte is called common ion effect. For example degree of dissociation of NH_4OH decreases in the presence of strong electrolyte NH_4Cl .





- **Hydrolysis of salts and pH of their solutions :** Hydrolysis of salt is defined as the reaction of cation or anion with water as a result of which the pH of water changes.
 - 1. Salts of strong and strong bases (*e.g.*, NaCl) do not hydrolyse. The solution pH will be 7.
 - 2. Salts of weak acids and strong bases (*e.g.*, CH_3COONa) hydrolyse, pH > 7 (The anion acts as a base).

$$X^{-} + H_{2}O \iff HX + OH^{-}$$
(Weak acid) (Weak base)

$$pH = 7 - \frac{1}{2} (pK_{a} + \log C)$$

3. Salt of strong acids and weak bases (*e.g.*, NH_4Cl) hydrolyse, pH < 7. (The cation acts as an acid).

$$M^{+} + H_{2}O \implies MOH + H^{+}$$
$$pH = 7 - \frac{1}{2} (pK_{b} + logC)$$

4. Salt of weak acids and weak base $(e.g., CH_3COONH_4)$ hydrolyse. The cation acts as an acid and anion as a base but whether the solution is acidic or basic depends upon the relative values of K_a and K_b for these **ions**.

$$M^{+} + X^{-} + H_{2}O \implies MOH + HX$$
$$pH = 7 + \frac{1}{2} (pK_{a} - pK_{b})$$

- **Buffer solutions :** The solutions, which resist the change in pH on dilution or addition of small amounts of acid or base, are called buffer solutions.
- **Basic buffer :** Solution of weak base and its salt with strong acid, For *e.g.*, NH₄OH + NH₄Cl
- Acidic buffer : Solution of weak acid and its salt with strong base, For *e.g.*, CH₃COOH + CH₃COONa.
- Henderson Hasselbalch Equation for the pH of Buffer solution—

$$pH = pK_{a} + \log \frac{[Salt]}{[Acid]}$$

$$pOH = pK_{a} + \log \frac{[Salt]}{[Base]}$$
(for acidic buffer)
(for basic buffer)

Equilibrium



• Solubility Product (K_{sp}) : The equilibrium constant that represent the equilibrium between undissolved salt (solute) and its ions in a saturated solution is called solubility product constant (K_{sp}) .

For
$$A_x B_y \xrightarrow{aq} x A^{y+} + y B^{x-}$$

 $K_{sp} = [A^{y+}]^x [B^{x-}]^y = (xs)^x (ys)^y = x^x y^x s^{(x+y)}$

where s = Molar solubility

If ionic product $< K_{sp}$; salt remain dissolve.

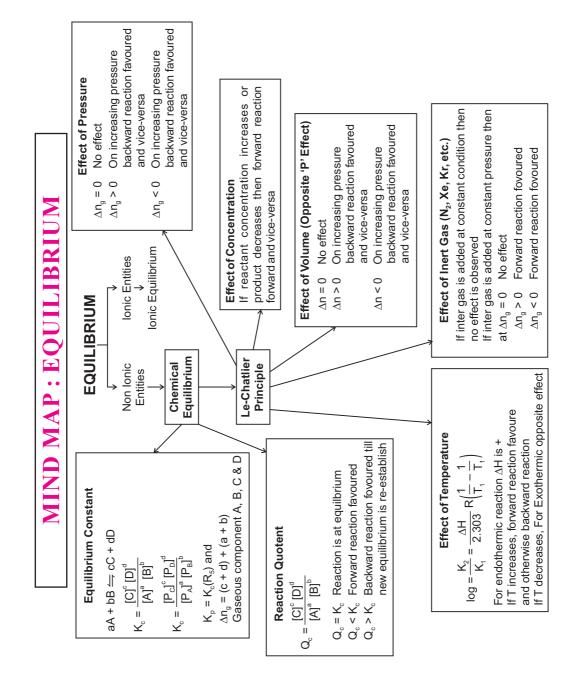
If ionic product $> K_{sp}$; salt will be precipitated.

Relationship between solubility (s) and solubility product (K_{sp}).

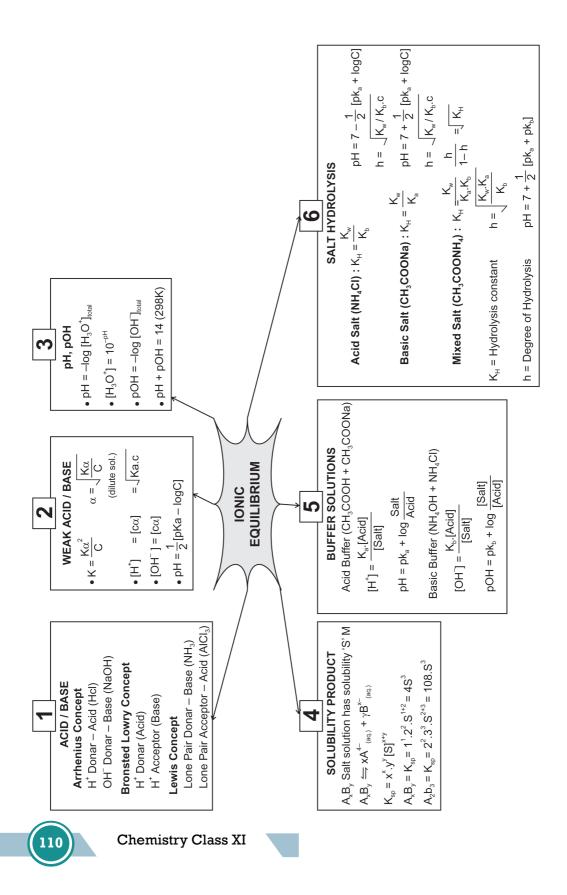
 $K_{sp} = x^x . y^y . s^{x+y}$ For binary salts (e.g., AgCl, AgBr, AgI) $K_{sp} = s^2$ For Ternary salts (e.g., PbI2) $K_{sp} = 4s^3$











MULTIPLE CHOICE QUESTION (MCQ)

1. For the hypothetical reactions, the equilibrium constant (k) values are given

$$A \rightleftharpoons B : k_1 = 2$$
$$B \rightleftharpoons C : K_2 = 4$$
$$C \rightleftharpoons D : K_3 = 8$$

The equilibrium constant (K) for the reaction $A \rightleftharpoons D$ is

(a) 48 (b) 24 (c) 12 (d) 64

2. The equilibrium constant for the reaction

$$SO_2(g) + \frac{1}{2} O_2(g) \rightleftharpoons SO_3(g) \text{ is } 5 \times 10^{-2} \text{ atm}^{-1/2}$$

The equilibrium constant for the reaction

$$2SO_3(g) \rightleftharpoons 2SO_2(g) + O_2(g)$$
 would be
(a) 100 atm (b) 25×10^{-4} atm (c) 400 atm (d) 125×19^{-6} atm^{-3/2}

3. A(g)+3B(g) ⇒ 4C(g) initial concentration of A is equal to that of B. The equilibrium concentrations of A and C are equal. What is the equilibrium constant for

$$4C(g) \rightleftharpoons A(g) + 3B(g)$$

(a) 4 (b) 1/8 (c) B (d) 16

- 4. The equilibrium reaction that is not affected by volume change at constant temperature is
 - (a) $H_2(g) + Cl_2(g) \rightleftharpoons 2HCl(g)$ (b) $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$

(c)
$$PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$$
 (d) $H_2(l) + CO_2(g) \rightleftharpoons H_2CO_3(l)$

- 5. For the reaction CO(g) + Cl₂ (g) ⇒ COCl₂(g), the value of K_c/K_p is equal to
 (a) RT (b) RT (c) 1/RT (d) 1.0
- 6. At 90°C pure water has $K_w = 10^{-12}$. The solution with pH value 6.5 is (a) Acidic (b) Basic (c) Amphoteric (d) Data insufficient



Equilibrium

7. 40 ml of 0.1 M NH_4OH is mixed with 20 mL of 0.1 M HCl. What is the pH of the mixture? (pK_b of ammonia solution = 4.74)

(a) 4.74 (b) 2.26 (c) 9.26 (d) 5

8. Identify Bronsted Lowry Acids in the reaction $\begin{bmatrix} A1(H \ O) \ 1^{3-} + HCO \longrightarrow \begin{bmatrix} A1(H \ O) \ (OH) \ 1^{2+} + HCO \end{bmatrix}$

$$\begin{array}{ccc} [\operatorname{AI}(\operatorname{H}_{2}\operatorname{OI}_{6}]^{\circ} &+\operatorname{HCO}_{3} \rightleftharpoons [\operatorname{AI}(\operatorname{H}_{2}\operatorname{O})_{5}(\operatorname{OH})]^{\circ} &+\operatorname{H}_{2}\operatorname{CO}_{3} \\ (X) & (Y) & (P) & (Q) \end{array}$$

$$(a) X, Y & (b) Y, P & (c) P, Q & (d) X, Q \end{array}$$

- 9. The pK_a of weak acid HA is 4.80 and pK_b of a weak base BOH is 4.78. The pH of an aqueous solutions of corresponding salt BA will be
 - (a) 7.01 (b) 4.79 (c) 9.22 (d) 10.0
- 10 If 'P' M is the solubility of $KAl(SO_4)_2$, then K_{sp} is equal to

(a)
$$p^3$$
 (b) $4p^4$ (c) p^4 (d) $4p^2$

TRUE AND FALSE TYPE QUESTIONS

- 1. Equilibrium state can be achieved if a reversible reaction is carried out in closed or open container.
- 2. For a reaction $2A(g) \rightleftharpoons B(g) Q_c > K$ if 'A' is added maintaining $Q_c > K$, the reaction will move in backward direction.
- 3. For the reaction at equilibrium

 $CaCO_3 \rightleftharpoons CaO(s) + CO_2(g)$

What CaO(s) is removed reaction moves in forward direction.

- 4. For a reaction $aA + bB \rightleftharpoons cC + dD$ at eqilibrium $G^{\theta} = 0$ always.
- 5. For a reaction at equilibrium $H_2(g) + Cl_2(g) \rightleftharpoons 2HCl(g)$ K = 4, the value of $\frac{K_b[HCl]^2}{K_c[H_2][Cl_2]}$ is 1.
- 6. For the electrolyte A_2B if K_{sp} is solubility product then its solubling 'S' M is $[K_{sp}]^{\frac{1}{3}} \div 4$.
- 7. HCO_3^{-} is conjugate base of $H_2CO_3^{-}$.
- 8. H_2O can act as acid as well as base.
- 9. The pH of buffer solution remain same when any amount of dilution is done.



10. For a salt AB₂(s) solution if Ionic product (I.P) > K_{sp}, then precipitation will take place.

Ans. 1. False	2. True	3. False	4. False	5. True
6. True	7. True	8. True	9. False	10. True

FILL IN THE BLANKS

- 1. At equilibrium rate of forward reaction is always equal to
- 2. $K_p \& K_c$ are for reaction at equilibrium of type $H_2(g) + Br_2(g) \rightleftharpoons 2HBr(g)$.
- 3. If K_c for reaction $CH_3COOH(l) + C_2H_5OH \rightleftharpoons CH_3COOC_2H_5(l) + H_2O(l)$ is 4. Then Q_c and K_c are
- 4. If A+B −70J/mol ⇒ D, reaction temperature is increased then reaction moves in direction.
- 5. Presence of catalyst will the equilibrium constant.
- 6. The conjugate acid of H₂O is
- 7. On dilution, the degree of dissociation of acetic acid will
- 8. The presence of NH₄Cl in NH₄OH solution will the degree of dissociation of NH₄OH.
- 9. If Ionic product (IP) < K_{sp} for a salt solution of AB, then addition of AB further lead to precipitation initially.
- 10. K_p is always equal to K_c if Δn_g is
- Ans. 1. rate of backward reaction, 2. equal, 3. equal, 4. forward,
 5. not change, 6. H₃O⁺, 7. increase, 8. decrease, 9. will not, 10. zero

MATCH THE COLUMNS

1.

Column-I

- A) $Na(g) + 3H_2(g) \rightleftharpoons 2NH_3(g); \Delta H = -ve$
- B) $2N_2(g) + 2O_2(g) \rightleftharpoons 4 \operatorname{NO}(g) : \Delta H = +ve$
- C) $X(g) \rightleftharpoons 2Y(g) \Delta H = +ve$
- D) $PC1_5(g) \rightleftharpoons PC1_3 + Cl_2; \Delta H = +ve$

Ans. (A) Q (B) P, R (C) P, S (D) P, S

Column-II

- P) K increases with increase in temp
- Q) K decreases with increase in temperature
- R) Pressure has no effect
- S) Product moles increases due to addition of inert gast at constant pressure.

Equilibrium



2.

Coloumn-I

- A) Salt of weak acid and weak base
- B) Salt of weak Acid and strong Base
- C) Salt of strong acid and strong base
- D) Salt of strong acid and weak base

 $\textbf{Ans.} (A) \ Q \quad (B) \ P \quad (C) \ S \quad (D) \ R \\$

Column-II

- P) $pH = \frac{1}{2} (pK_w + pK_a + \log_c)$
- Q) $pH = \frac{1}{2} (pK_w + pK_a pk_b)$
- R) $pH = \frac{1}{2} (pK_w pK_b \log_c)$
- S) $pH = \frac{1}{2} (pK_w)$

ASSERTION - REASON TYPE QUESTION

Each question contains statement-1 (assertion) and Statement-2 (Reason) Examine the statements carefully and mark the correct answer according to the instruction given below :

- A. If both the statements are true and statement -2 is the correct explanation of statement-I
- B. If both the statements are true but statement-2 is not the correct explanation of statement-I
- C. If statement-I is true and statement-2 is false
- D. If statement-I is false and statement-2 is true.
- Statement-1 : The endothermic reactions are favoured at lower temperature and the exothermic reactions are favoured at higher temperature. Statement-2 : when a system in equilibrium is disturbed by changing the temperature, it will tend to adjust itself so as to overcome the effect of change.
- 2. Statement-1 : The melting point of ice decreases with increase of pressure Statement-2 : Ice contracts on melting.
- Statement -1 : The gas phase reaction PCl₃(g) + Cl₂(g) ⇒ PCl₅(g) shifts to the right on increasing pressure.
 Statement-2 : When pressure increase, equilibrium shifts towards more number of moles.
- 4. Statement-1 : The physical equilibrium is not static but dynamic in nature. Statement-2: The physical equilibrium is a state in which two opposing process are proceeding at the same rate.
- 5. Statement-1 : The catalyst does not after the equilibrium constant. Statement-2 : Because for the catalysed reaction and uncatalysed reaction ΔH remains same and equilibrium constant depends on ΔH .





- 6. Statement-1 : If water is heated to 59°C, the pH will increase. Statement-2 : K_w increases with increase in temperature.
- Statement-1: Addition of HCl(aq.) to CH₃COOH (aq.) decrease the ionisation of CH₃COOH (aq.).
 Statement-2 : Due to common ion effect H⁺, ionisation of CH₃COOH decreases.
- Statement-1: Sparingly soluble salts AB and XY₂ with the same solubility product, will have different solubility.
 Statement 2: Solubility of sparingly soluble salts depends upon solubility product.
- Statement-1 : The ionisation constants of weak diprotic acid are in the order of Ka₁ > Ka₂. Statement-2 : Removal of H⁺ from anion is difficult as compared to neutral atom.
- 10. Statement-1 : In a titration of weak acid with strong base, the pH at the half equivalence point is pK_a.
 Statement-2 : At half equivalence point it will form acidic buffer at its

Statement-2 : At half equivalence point, it will form acidic buffer at its maximum capacity where [Acid] = [Salt].

Ans.: 1. D, 2. A, 3. C, 4. A, 5. A, 6. D, 7. A, 8. B, 9. A, 10. A

ONE WORD ANSWER TYPE QUESTIONS

1. What is sum of pH + pOH at 25°C?

[**Ans.** 14]

2. Write the Henderson Hasselbalch equation for acidic buffer

Ans. pH = pka + $10g \frac{[SALT]}{[ACID]}$

- 3. How is degree of dissociation related with concentration terms and Ka, for weak electrolyte. Ans. $\alpha = \sqrt{Ka/c}$
- 4. How NH_3 is defined as Lewis base?

[Ans. It contain Lone paid of electrons]

- 5. How are K_p and K_c related? [Ans. $K_p = K_c (RT)^{\Delta n}$]
- 6. How does K affected for endothermic reaction if temperature is increased? [Ans. K get decreased]
- 7. What is the effect of catalyst on K? [Ans. K remains unaffected]
- 8. How is pH scale affected by increasing temperature?

[Ans. pH scale gets contracted]

- 9. What is the conjugate base of HCO_3^{-2} ? [Ans. CO_3^{2-}]
- 10. What is the nature of CH_3COOH in conc. HCl solution?

[Ans. Bronsted Base]

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Equilibrium

1-MARK QUESTIONS

- 1. Define physical equilibrium. Give an example also.
- 2. Fizz is observed when soda water bottle is opened. Why ?
- **3.** Justify the statement : 'Both physical and chemical equilibrium are dynamic in nature'
- 4. State Henry's law.
- **5.** In a reversible reaction, the two substances are in equilibrium. If the concentration of each one is reduced to half, then what is the effect on the equilibrium constant ?
- **6.** K_1 and K_2 are equilibrium constant for reactions (1) and (2)

(i)
$$N_2(g) + O_2(g) \rightleftharpoons 2 NO(g)$$

(ii) NO(g)
$$\rightleftharpoons 1/2$$
 N₂(g) + 1/2 O₂(g)

Calculate the relation between K_1 and K_2 .

- 7. Write the equilibrium constant expression for the following reaction : $3 \operatorname{Fe}(s) + 4 \operatorname{H}_2O(g) \rightleftharpoons \operatorname{Fe}_3O_4(s) + 4 \operatorname{H}_2(g)$
- 8. Classify the equilibrium as homogeneous or heterogeneous :

$$CH_{3}COOC_{2}H_{5}(aq.) + H_{2}O(1) \rightleftharpoons CH_{3}COOH(aq.) + C_{2}H_{5}OH(aq.)$$

$$(P_{NH})$$

9.
$$K_p = \frac{(\Gamma_{NH_3})}{(P_{H_2})^{3/2} (P_{N_2})^{1/2}}$$

Write the balanced chemical equation corresponding to the above expression.

- 10. Give the direction in which the reaction would proceed if $Q_c > K_c$.
- **11.** $Hb(s) + O_2(g) \rightleftharpoons HbO_2(s)$

Predict the direction in which equilibrium gets shifted if partial pressure of $O_2(g)$ is lowered.

12. Discuss the position of equilibrium if the following reaction is carried out in the presence of catalyst.

$$2SO_2(g) + O_2(g) \rightleftharpoons 2SO_3(g)$$

13. Which of the following are Lewis acids ?
 H₂O, BF₃, H⁺, NH₄⁺





- 14. Write the conjugate acids for the following Bronsted bases. C_6H_5OH, H_2O
- Write the conjugate bases for the following Bronsted acids.
 H₂O, CH₃COOH.
- **16.** Which of the following are Lewis acids ?

(a)
$$H_2O$$
 (b) $AlCl_3$ (c) NH_4^+

- 17. Define Ostwald's dilution law.
- **18.** SO_3^{2-} is Bronsted base or acid and why ?
- **19.** Why pH of our blood remains almost constant at 7.4 though we quite often eat spicy food ?
- 20. pH of black coffee is 5.0 at 25°C. Is black coffee acidic or basic ?[Ans. Acidic]
- **21.** What will be the value of $(pK_a + pK_b)$ at 25°C.
- 22. What will be the pH of 1 M KNO₃ solutions at 25°C?
- **23.** $CaCl_2(s) + H_2O(l) \rightleftharpoons CaCl_2(aq.) + Heat$

Discuss the solubility of CaCl₂ if temperature is increased.

- **24.** Why does the solubility of CO_2 decrease with rise in temperature ?
- **25.** The solubility of $A_2 X_3$ is y mol dm⁻³. Calculate its solubility product.
- **26.** Write the K_{sp} expression for Al (OH)₃.
- 27. What is the condition for precipitation of a salt?
- **28.** Pridict whether the solution is acidic, basic or natural when NH_4NO_3 undergo hydrolysis.
- **29.** Explain why pure NaCl precipitates out when HCl gas is passed through the solution of NaCl ?
- **30.** Give the Henderson's Hasselbalch equation for an acidic buffer solution.
- **31.** On which of the factors the equilibrium depend : Temperature, nature of reactant and product, initial concentration and pressure of the reactants.
- **32.** Arrange the following in increasing acidic strength HCl, HBr, HF, HI [Ans, HF < HCl < HBr < H]

[Ans. HF < HCl < HBr < HI]





33. Arrange the following in increasing Lewis base strange NH₃, BiH₃, PH₃, AsH₃, SbH₃

[Ans. $BiH_3 < SbH_3 < ASH_3 < PH_3 < NH_3$]

34. Arrange following in increasing pH value $0.1M \text{ CH}_3\text{COOH}, 0.1 \text{ M NaCl}, 0.1\text{MHCl}, 0.1\text{MNaOH}, 0.1\text{MNH}_4\text{OH}$ [Ans. $0.1\text{MHCl} < 0.1\text{M CH}_3\text{COOH} < 0.1\text{M NaCl} < 0.1\text{ NH}_4\text{OH} < 0.1\text{M NaOH}$]

35. Arrange following in increasing order of degree of hydrolysis.
0.1M NH₄OH, 0.01 M NH₄OH, 10⁻⁵ M NH₄OH, 10⁻³ M NH4OH, 10⁻⁶ M NH₄OH

[**Ans.** 0.1M NH₄OH < 10⁻² M NH₄OH, 10⁻³M NH₄OH < 10⁻⁵MNH₄OH < 10⁻⁶ M NH₄OH]

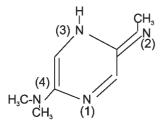
36. Arrange following in increasing order of acidic strengh

 $\label{eq:coop} \begin{array}{l} \mbox{CH}_3\mbox{COOH},\mbox{CH}_3\mbox{COOH},\mbox{CH}_2\mbox{COOH},\mbox{CH}_2\mbox{COOH}\\ \mbox{[Ans. CH}_3\mbox{COOH} < \mbox{C}_6\mbox{H}_5\mbox{COOH} < \mbox{HCOOH} < \mbox{CH}_2\mbox{FCOOH}\\ \end{array}$

37. Arrange following in increasing order of basic strength in gas phase NH₃, (CH₃)₂NH, (CH₃)₃N, CH₃NH₂

 $[\mathbf{Ans.} \ \mathrm{NH}_3 < \mathrm{NH}_3 \mathrm{NH}_2 < (\mathrm{CH}_3)_2 \mathrm{NH} < (\mathrm{CH3})_3 \mathrm{N}]$

38. Arrange the following pkb in increasing order



[**Ans.**pK₂ < pK₁ < pK₄ < pK₃]

39. Arrange the basic strength of following F⁻, Br⁻, Cl⁻, I⁻

[Ans. $I^- < Br^- < Cl^- < F^-$]

40. Arrange the following in increasing base strength CH₃⁻, NH₂⁻, OH⁻, F⁻

[Ans.
$$F^- < OH^- < NH_2 < CH_3$$
]



2-MARKS QUESTIONS

- The standard Gibbs energy change at 300 k for the reaction 2A ⇒ B + C is 2494. 2 J. At a given temperature, and time. the composition of the reaction mixture is [A] = ½, [B] = 2, [C] = ½. The reaction proceed in the (R = 8.314J/K/mo1, = 2.718) [Ans. Reverse direction]
- 2. The equilibrium constant for

 $N_2(g) + O_2(g) \rightleftharpoons 2NO(g)$ is K, then calculate equilibrium constant for $\frac{1}{2}N_2(g) + \frac{1}{2}O_2(g) \rightleftharpoons NO(g)$. [Ans. \sqrt{K}]

3. For the reversible reaction $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ at 500°C, the value of Kp is 1.44×10^{-5} atn⁻². Find the K_c value.

[Ans. $1.44 \times 10^{-5} / (0.082 \times 773)^{-2}$]

- 4. The equilibrium constant at 298 K for the reaction A + B ⇒ C + D is 100. If the initial concentration of all the four species were 1M each, then equilibrium concentration of D will be [Ans. 1.818]
- 5. For the reaction

 $NH_4COO NH_2(g) \rightleftharpoons 2NH_3(g) + CO_2(g)$

If equilibrium pressure is 3 atm. Find the value of Kp [Ans. 4]

6. A buffer solution with pH 9 is to be prepared by mixing NH_4Cl that should be added to one litre of 1.0m NH_4OH kb 1.8×10^{-5}

[Ans. $NH_4Cl = 1.8 M$]

- 7. Calculate the solubility of silver chloride in water at room temperature if the K_{sp} of AgCl is 1.6×10^{-10} . [Ans. 1.26×10^{-5} M]
- 8. Calculate the molar solubility of Ni(OH)₂ in 0.10m NaOH. The ionic product of Ni(OH)₂ is 2.0×10^{-15} . [Ans. 2.0×10^{-13} M]
- 9. Calculate the pH of 10⁻⁸ M HCl solution. [Ans. 6.96]
- 10. How many grams of NaOH must be dissolved in IL of the solution to give it a pH value of L²? [Ans. 0.4g]





3-MARKS QUESTIONS

- 1. The equilibrium constant for the reaction $H_2(g) + Br_2(g) \rightleftharpoons 2HBr(g)$ at 1024 K is 1.6×10^5 . Find the equilibrium pressure of all gases if 10.0 bar of HBr is introduced into a sealed container at 1024K. [Ans. 10 bar]
- 2. For the reaction $2BrCl(g) \rightleftharpoons Br_2(g) + Cl_2(g) K_c$ is 32 at 500 K. If initially pure BrCl is present at a concentration of 3.30×10^{-3} M, what is its molar concentration in the mixture at equilibrium? [Ans. 3.0×10^{-4} M]
- 3. What is the equilibrium constant K_p and K_c for the reaction $PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$ if the pressure is 1.0 atm in 8.0L container at equilibrium. [Ans. $K_c = 0.04 K_p = 1.77$]
- 4. The K_p for the reaction, $N_2O_4(g) \rightleftharpoons NO_2(g)$ is 640 mm at 775 K. Calculate the percentage dissociation of N_2O_4 at equilibrium pressure of 160mm. At what pressure the dissociation will be 50%.

[**Ans.** 70.7%, P = 480 mm]

- 5. Show that degree of dissociation (α) for the dissociation of PCl₅ into PCl₃ and Cl₂ at pressure P is given by $\alpha = \left[\frac{Kp}{P+kp}\right]^{1/2}$
- 6. How much of $0.3M \text{ NH}_4\text{OH}$ should be mixed with 30 mL of 0.2m solution of NH_4Cl to give a buffer solution of pH 10. pk_b for NH_4OH is 4.75.

[**Ans.** V = 112.5 mL]

- 7. Predict whether a precipitate will be formed or not on mixing 20 mL of 0.001 N NaCl solution with 80 mL of 0.01 M AgNO₃ solution. K_{sp} for AgCl is 1.5×10^{-10} . [Ans. Yes, ppt will formed.]
- 8. The values of Ksp of two sparingly soluble salts $Ni(OH)_2$ and AgCN are 2.0×10^{-15} and 6.0×10^{-17} respectively. Which salt is more soluble. Explain

[Ans. $S_{Ni(OH)_2} = 5.8 \times 10^{-5}M : S_{(Ag CN)} = 7.8 \times 10^{-9}M . Ni(OH)_2$ is more soluble]

9. The ionization constant of propanoic acid is 1.32×10^{-15} . Calculate the degree of ionization if its solution is 0.05 M. What will be its degree of ionization if the solution is 0.01 M in HCl solution.

[Ans. 1.62×10^{-2} , 1.32×10^{-3}]

10. Calculate the pH of a solution obtained by mixing 50ml of 0.2M HCl with
49.9 mL of 0.2m NaOH solution.[Ans. 3.699]





HOTS QUESTIONS

1. The molar solubility of $Cd(OH)_2$ is 1.84×10^{-5} M. Calculate the expected solubility of $Cd(OH)_2$ in a buffer solution of pH = 12.

Ans.
$$Cd(OH)_2 \rightleftharpoons Cd_{(aq.)}^{2+} + 2OH_{(aq.)}^{-}$$

 $S = 10^{-2}$
 $2.49 \times 10^{-14} = S(10^{-2})^2 \qquad \therefore S = 2.49 \times 10^{-10M}$

2. An aqueous solution contains an unknown concentration of Ba^{2+} . When 50 ml of a 1M solution of Na_2SO_4 is added. $BaSO_4$ just begins to precipitate. The final volume is 500ml. The solubility product of $BaSO_4$ is 1×10^{-10} . Find the original concentration.

Ans.
$$K_{sp} = \left[Ba^{2+} \right] \left[SO_4^{2-} \right] = \left[Ba^{2+} \right] \left[\frac{50 \times 1}{500} \right] = 10^{-9} \times 500$$

 $Ba^{2+} = 10^{-9} M$
 $10^{-9} \times 500 = 450 \times M$ $\therefore M = 1.11 \times 10^{-9} M$

3. An aqueous solution contains 0.10 M H_2S and 0.20 M HCl. If the equilibrium constants for the formation of HS⁻ from H_2S is 1.0×10^{-7} and that of S²⁻ from 4S⁻ ions is 1.2×10^{-13} , then find the concentration of S⁻² ions in aqueous solution.

Ans.
$$H_2S(aq.) \rightleftharpoons 2H^+ + S^{2-}$$

 $(0.1 - x) \qquad (2x + 0.29) \qquad x$
 $K_a = K_{a_1} \times K_{a_2} = 1.2 \times 10^{-20}$
 $1.2 \times 10^{-20} = \frac{(0.2)^2 [S^{2-}]}{0.1}, [S^{2-}] = 3 \times 10^{-20}$

4. How many litres of water must be added to 1 litre of an aqueous solution of HCl with a pH of 1 to create an aqueous solution with pH of 2?

Ans. $0.1 \times 1 = (1 + v) (0.01) \implies v = 9L$

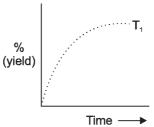
5. A certain buffer solution contains equal concentration of X⁻ and HX. The $K_{\rm b}$ for X⁻ is 10⁻¹⁰. Find the pH of the buffer .

Ans.
$$k_a \cdot k_b = 10^{-14}$$
 $\therefore k_a = \frac{10^{-14}}{10^{-10}} = 10^{-4}$
 $pH = pka + \log \frac{[x^-]}{[Hx]}$
 $\therefore pH = 4 + \log \frac{1}{1} = 4$ $\therefore pH = 4$

Equilibrium

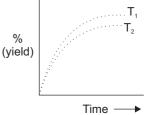


6. The % yield of Ammonia as a function of time in the reaction $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g), \Delta H < O$ at (P, T) is given below:



If this reaction is conducted at $T_2 > T_1$, then plot the % yield of NH_3 as a function of time on same graph

Ans. Initially on increasing temperature the rate of reaction increases, however since the reaction is exothermic therefore % yield of NH_3 get decreased overall after a certain interval of time.



7. Consider the reaction $NH_4COONH_2(s) \rightleftharpoons 2NH_3(g) + CO_2(g)$ at a certain temperature, the equilibrium pressure of the system is 0.318 atm. Find K_p of the decomposition of ammonium carbonate.

Ans.
$$P_{total} = 3P$$
 : $P = 0.318/3 = 0.106$
 $Kp = 4P^3 = 4.76 \times 10^{-3}$

8. The equilibrium constant for the reaction $CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g)$ is 5. How many moles of CO_2 must be added to 1 litre container already containing 3 moles each of CO and H_2O to make 2M equilibrium concentration of CO?

Ans. $CO + H_2O \rightleftharpoons CO_2 + H_2$ $t = 0 \quad 3 \quad 2 \quad x \quad 0$ At equilibrium $2 \quad 2 \quad x+1 \quad 1$ $\therefore S = \frac{x+1}{4} \Longrightarrow x = 19$

9. At constant temperature, the equilibrium constant K_p

$$N_2O_4 \rightleftharpoons 2NO_2$$
 is given by
 $k_p = \frac{4x^2P}{1-x}$ where, P = Pressure and X = Extent of reaction



How does the value of K_p change on following changes

- (a) 'P' increases
- (b) X changes
- (c) 'P' decreases

Ans. K_p is equilibrium constant which does not change on changing the P, x.K_p depends on temperature.

10. When two reactants A and B are mixed to give product 'c' and 'p' the reaction quotient 'Q' at the initial stages of the reaction will be?

Ans. In the beginning of the reaction Q = 0. As the reaction proceeds in the forward direction Q starts increasing.

At chemical equilibrium Q = K





UNIT TEST

Time Allowed: 1 hr

General Instructions:

(i) All questions are compulsory.

(ii) Maximum marks carried by each question are indicated against it.

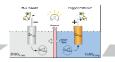
1.	What is the pH 10^{-3} M HCl solution ?					
	(a) 1	(b) 11	(c) 3	(d) 14		
2.	Which one ca	n act as Arrh	enius Acid ?		[1]	
	(a) NH ₃	(b) H ₂ O	(c) HCl	(d) C_6H_5OH		
3.	Write the conjugate base of $CH_3COOH + H_2$.					
4.	Write the relation between K_p and K_c .					
5.	What is the nature of following reaction Exothermic or endothermic $A + B - 70J \longrightarrow C$					

- The pKa of CH₃COOH and pKb of NH₄OH are 4.76 and 4.75 respectively. Calculate the pH of CH₃COONH₄. [2]
- 7. What is a buffer solution. Calculate the pH of the solution obtained by adding 4mol of CH_3COOH with 3 mol of NaOH in 1 litre container. [2] pKa, $CH_3COOH = 4.74 \log 2 = 0.3010 \log 3 = 0.4771$
- 8. Calculate the molar solubility of Ni(OH)₂ in 0.1M KOH solution. The K_{sp} for Ni(OH)₂ is 2.0×10^{-15} . [3]
- 9. K_p = 0.04 atm at 899 K for the equilibrium shown below. What is the equilibrium concentration of 2H₆ when it is placed in a flask at 4.0 atm pressure and allowed to come to equilibrium. [3]
 C₂H₆(g) ⇒ C₂H₄(g) + H₂(g)
- 10. Ionization constant of Benzoic acid is 6.46×10^{-5} and K_{sp} for silver benzoate is 2.5×10^{-13} . How many times is silver benzoate more soluble in buffer of pH 3.19 compared to its solubility in pure water.

$$[\mathrm{H}^+] = 6.46 \times 10^{-4}$$
 [5]









Redox Reactions

FAST TRACK : QUICK REVISION

Oxidation and Reduction :

Oxidation

- 1. Addition of oxygen
- 2. Removal an Hydrogen
- **3.** Addition of an electronegative element.
- 4. Removal of an electropositive element
- 5. Loss of electron(s)
- 6. Increase in oxidation number.

Reduction

- 1. Removal of oxygen
- 2. Addition of Hydrogen
- **3.** Removal of an electronegative element.
- 4. Addition of an electropositive element.
- 5. Gain of electron(s)
- 6. Decrease in oxidation number.
- **Reducing Agent :** Donor of electrons.
- Oxidising Agent : Acceptor of electrons.
- **Redox Reaction :** Reactions in which oxidation and reduction takes place simultaneously.
- **Oxidation Number :** It is charge that an atom appears to have in a given species when the bonding electron are counted towards more electronegative atom.
- Calculation of Oxidation Number :
 - (a) Oxidation number of all the elements in their elemental form (in standard state) is taken as zero. Oxidation number of element in a molecule Cl₂, F₂, O₂, P₄, O₃, Fe, H₂, N₂, C (graphite) is zero.
 - (b) Common Oxidation number of elements of first group is +1. Common Oxidation number of elements of second group + 2.
 - (c) For ions composed of only one atom, the oxidation number is equal to the charge on the ion.

- (d) The oxidation number of oxygen in most compounds is -2. While in peroxides (*e.g.*, H₂O₂, Na₂O₂), each oxygen atom is assigned an oxidation number of -1, in super oxides (*e.g.*, KO₂, RbO₂) each oxygen atom is assigned an oxidation number of $-(\frac{1}{2})$.
- (e) In oxygen difluoride (OF_2) and dioxygen difluoride (O_2F_2) , the oxygen is assigned an oxidation number of +2 and +1, respectively.
- (f) The oxidation number of hydrogen is + 1 but in metal hydride its oxidation no. is 1.
- (g) In all its compounds, fluorine has an oxidation number of -1.
- (h) The algebraic sum of the oxidation number of all the atoms in a compound must be zero.

(i) In polyatomic ion, the algebraic sum of all the oxidation numbers of atoms of the ion must equal the charge on the ion.

• Types of Redox Reactions:

(i) Combination Reaction : 0 0 +2 -3 $3 \text{ Mg}(s) + N_2(g) \xrightarrow{>} \text{ Mg}_3 N_2(s)$ (ii) Decomposition Reaction : +1 +5 -2 +1 -1 0 $2\text{KClO}_3(s) \xrightarrow{>} 2 \text{ KCl}(s) + 3O_2(g)$ (iii) Metal Displacement : +2 + 6 -2 0 +2 + 6 -2 0 $\text{CuSO}_4(aq) + \text{Zn}(s) \xrightarrow{>} \text{ZnSO}_4(aq) + \text{Cu}(s)$ (iv) Non-metal displacement : 0 +1 -2 +2 -2 +1 0 $\text{Ca}(s) + 2 \text{ H}_2O(1) \xrightarrow{>} \text{Ca}(\text{OH})_2 + \text{H}_2(g)$

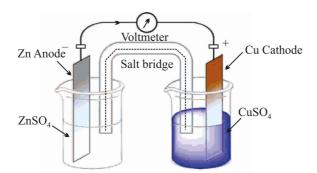
(v) Disproportionation reactions : It is a reaction in which same element is reduced and oxidized simultaneously.

0 -1 + 1C1₂ (g) + 2 OH⁻ (aq) \rightarrow Cl⁻ (aq) + ClO⁻ (aq) + H₂O (1)

- Stock Notation : Representing oxidation number of metal in Roman numerals within parenthesis after the symbol or name of metal in the molecular formula or name of a compound. For *e.g.*, Stock Notation of Ferric oxide is Fe₂(III)O₃ or Iron (III) oxide.
- Fractional Oxidation Number : When two or more atoms of an element are present in different oxidation states, then calculated oxidation number may comes out as fractional due to average of all the different oxidation states.

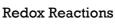
In reality no element can have a fractional oxidation state.

- Balancing of Redox Reactions :
 - (A) Oxidation number method
 - (B) Half reaction method
- Electrode Potential (E) : Potential difference between electrode and electrolytic solution due to charge separation.
- Standard Electrode Potential (E^θ): Electrode Potential measured at 298
 K and 1M concentration of metal ions (or 1 bar pressure of gas).
- Electrochemical Cell : A device in which chemical energy of a spontaneous redox reaction is converted into electrical energy.

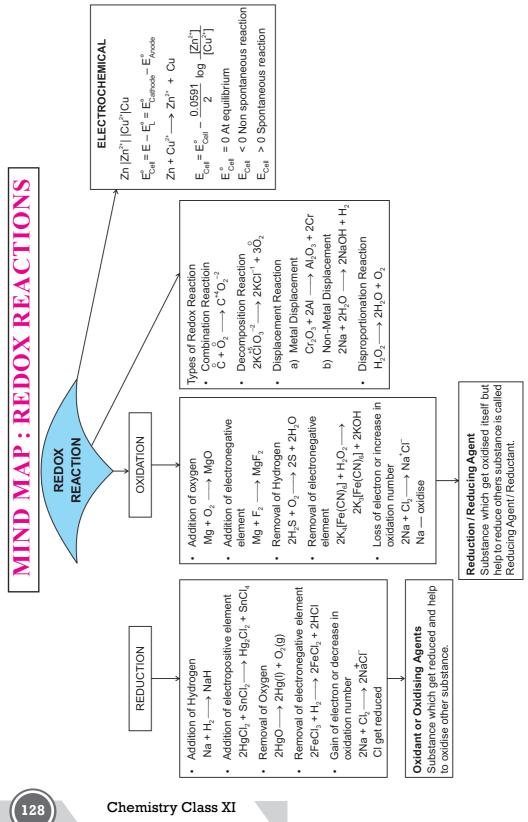


	Cell diagram: Zn Zn ²⁺ Cu ²⁺ Cu						
	LHS oxid	ation,	$Zn \rightarrow Zn^{2+} +$	2 <i>e</i> -			
	RHS redu	ction	$Cu^{2+} + 2e^- \rightarrow Cu$				
	Overall re	eaction $Zn(s) + Cu^{2+}$	$(aq) \rightarrow \operatorname{Zn}^{2+}(aq) +$	- Cu(s)			
•	Represent	tation of an Electroch	emical cell :				
	Flow of electrons						
	← Flow of current —						
		$\operatorname{Zn}(s) \operatorname{Zn}^{2+}$	$(aq) \parallel \mathrm{Cu}^{2+}(aq) \mid \mathrm{Cu}$	(s)			
		Left Electrode	Salt Bridge	Right Electrode			
	LOAN	Oxidation		Reduction			
		Anode		Cathode			
		Negative		Positive			

• Functions of Salt Bridge : (i) To complete inner circuit. (ii) To maintain electrical neutrality around electrodes.







MULTIPLE CHOICE QUESTIONS (MCQ)

1.	The oxidation state of Fe in Fe_3O_4 is					
	(a) +2	(b) + 3				
	(c) $\frac{8}{3}$	(d) $+2, +3$				
2.	The oxidation state of 'S' in KAl(S	$SO_4)_2$.12H ₂ O is				
	(a) –2	(b) – 1				
	(c) 2	(d) + 6				
3.	Oxidation state carbon in C_3O_2 is					
	(a) $\frac{4}{3}$	(b) 0				
	(c) 2	(d) 0, 2				
4.	The reaction $S_8 + 12OH^- \longrightarrow 4S_8$	$S^{2-} + 2S_2O_3^{2-} + 6H_2O$ is				
	(a) Combination reaction	(b) Decomposition reaction				
	(c) Non-metal displacement	(d) Disproportionation reaction				
5.	E^0 for H^+/H_2 is					
	(a) 0	(b) +1V				
	(c) -1.0V	(d) –2.0V				
6.	Which one act as strong oxidising	gagent				
	$K^+/K = -2.9$ 3V, $Ag^+/Ag = 0.80$, H	$Hg^{2+}/Hg = 0.79V$				
	(a) K ⁺	(b) K				
	(c) Hg^{2+}	(d) Ag^+				
7.	The coefficient of HCl in balance	reaction is				
	$Pb_{3}O_{4} + HCl \longrightarrow PbCl_{2} + Cl_{2} +$	H ₂ O				
	(a) 1	(b) 8				
	(c) 3	(d) 4				
8.	Sum of oxidation numbers of all I	Bromine atoms in Br ₃ O ₈ is				
	(a) 6	(b) 4				
	(c) 16	(d) 20				
9.	In the reaction $6ClO_2^- \longrightarrow 4ClO_2^- \longrightarrow 4ClO$	$D_3^- + 2Cl^-, Cl^-$ ion is				
	(a) Oxidised Reduced	(b) Reduced				
	(c) Odixised and	(d) Neither Oxidised nor reduced				

Redox Reactions

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10. 'I' can not act as reducing agent in following state

- (a) -1 (b) +1
- (c) +7 (d) +5

Ans: 1. (d) 2. (d) 3. (d) 4. (d) 5. (a) 6. (d) 7. (b) 8. (c) 9. (c) 10. (d)

FILL IN THE BLANKS

- (i) Oxidation is _____ of electrons.
- (ii) S.H.E. stands for _____.
- (iii) Oxidation state of Oxygen in O_2F_2 is _____.
- (iv) Disproportionation is a type of _____ reaction.
- (v) Oxidant is one which ______ electron..

(vi) $Cl_2 + 2OH^- \longrightarrow ClO^- + C1^-$ is a _____ type of reaction.

(vii) Oxidation state of F is always either _____ or _____.

- (viii) Oxidation state of Oxygen in O₃ is _____.
- (ix) Reducing agent are also called _____.
- (x) Hydrogen economy is use of Hydrogen as _____.
- Ans: (i) loss, (ii) standard hydrogen electrode, (iii) +1, (iv) redox, (v) gain, (vi) disproportionation, (vii) 0, -1, (viii) zero, (ix) reductant, (x) fuel

TRUE AND FALSE TYPE QUESTIONS

- (i) In Redox reaction first oxidation take place.
- (ii) Oxidising agents are also called reductant.
- (iii) Fluorine cannot have +1 oxidation state.
- (iv) O_2^+ has oxidation state of oxygen as $+\frac{1}{2}$.
- (v) If for the reaction $Ca^{2+} + 2e^- \longrightarrow Ca(s)$; $E^{\Theta} = -2.87$ Then for the reaction $2Ca^{2+} + 4e^- \longrightarrow 2Ca(s)$; $E^{\Theta} = 2(-2.87)V$
- (vi) Salt bridge is used for enhancing E° values of individual half reaction.
- (vii) Anode is -ve charged in Galvanic cell.
- (viii) KCl can be use in salt bridge.
- (ix) Current flows in galvanic cell from Anode to cathode.
- (x) MnO_4^{-} is colourless in basic medium.
- Ans: (i) False (ii) False (iii) True (iv) True (v) False (vi) False (vii) True (viii) True (ix) False (x) False 130 Chemistry Class XI

MATCH THE COLUMNS

1. Column-I Column-II (a) $CH_4 + 2O_2 \longrightarrow CO_2 + 2H_2O$ (p) Disproportionation (b) $Cr_2O_3 + 2A1 \longrightarrow Al_2O_3 + 2Cr$ (q) Non-metal displacement (c) Fe +2HCl \longrightarrow FeCl₂ + H₂ (r) Metal displacement (d) $P_4 + 3OH^- + 3H_2O \rightarrow PH_3 + 3H_2PO_2^-$ (s) Metal displacement **Ans.** (a) - (s) (b) - (r) (c) - (q) (d) - (p)2. Column-I Column-II (Oxidation sate of N) (a) NO (p) + 5(b) NO_2 (q) + 3(c) NO_2^{-} (r) + 4(d) NO_3^{-} (s) +2

Ans. (a) - (s) (b) - (r) (c) - (q) (d) - (p)

ASSERTION AND REASON TYPE QUESTIONS

Each question contain statement-1 (Assertion) and statement-2 (Reason)

Examine the statements carefully and work the correct answer according to the instruction given below :

- (a) If both the statements are True and Statement-2 is the correct explanation of the statement-1
- (b) If both the statements are True and statement-2 is not the correct explanation of statement-1
- (c) If statement-1 is true and statement-2 is False.
- (d) If statement-1 is false and statement-2 is True.
- 1. Statement 1 : In HF, the oxidation state of 'F' is -1
 - Statement 2 : 'F' being most electronegative, will have -1 oxidation in its compound.
- 2. Statement 1 : Oxygen has zero oxidation state in O_2 .
 - Statement 2 : Element in their elemental form have zero oxidation state.
- 3. Statement 1 : Oxidation state of Oxygen in H_2O_2 is -1.
 - Statement 2 : H_2O_2 has peroxide linkage.

Redox Reactions



For the reaction $Zn + Cu^{2+} \longrightarrow Zn^{2+} + Cu$; E_{cell} is +ve. 4. Statement - 1 : For standard Hydrogen electrode $E^{\circ} = 0$ Statement - 2 : $2H_2O_2 \longrightarrow 2H_2O + O_2$ is Decomposition reaction 5. Statement - 1 : (Redox). Oxygen has -2 oxidation state in H₂O. Statement - 2 : $C + O_2 \longrightarrow CO_2$; carbon get oxidised. 6. Statement - 1 : Statement - 2 : Gain of Hydrogen is reduction. $CaCO_3 \longrightarrow CaO + CO_2$ is not redox reaction. 7. Statement - 1 : C, Ca, O do not change their oxidation number in the Statement - 2 : reaction. 8. Statement - 1 : Oxidation also occurs when decrease in electron density is observed. Statement - 2 : Oxidation is gain of electro-positive element. $Cr_2O_7^{2-}$ is a self indicator. 9. Statement - 1 : MnO_4^{-} acts as a self indicator. Statement - 2 : Equivalence point comes first before end point. 10. Statement - 1 : Equivalence point cannot be obtained even by graphical Statement - 2 : method.

Ans: 1. (a) 2. (a) 3. (a) 4. (b) 5. (d) 6. (b) 7. (a) 8. (c) 9. (d) 10. (d)

ONE WORD ANSWER TYPE QUESTIONS

- 1. What is the oxidation number of S in S_8 .
- 2. What is the oxidation state of Oxygen in H_2O_2 .
- 3. Name the substance used in salt-bridge.
- 4. Name an indicator which can act as self-indicator.
- 5. When a substance gains electron, it is called :
- 6. Name the ion which is used for balancing the hydrogen atom in acidic medium.
- 7. In the reaction $3Mg + N_2 \longrightarrow Mg_3N_2$, Nitrogen is oxidised or reduced.

Ans: 1. zero2. 13. $NH_4Cl \text{ or } KCl$ 4. $KMnO_4$

5. Reduction $6. H^+$ 7. Reduced



- 1. Define oxidation and reduction according to electronic concept.
- 2. Define oxidation and reduction according to oxidation number.
- **3.** A freshly cut apple is almost white but it turns reddish brown after sometime. Give reason.
- 4. Define oxidation number.
- 5. Write oxidation number of Mn in $KMnO_4$.
- 6. Write oxidation number of Cr in $Cr_2O_7^{2-}$.
- 7. Write Stock notation of MnO_2 and $AuCl_3$.
- 8. Define redox reaction with example.
- 9. Define disproportionation reaction. Give one example.
- **10.** Define the term redox titration.
- 11. Name the indicator used in redox titration involving $K_2Cr_2O_7$ as an oxidizing agent.
- 12. At what concentration of Cu²⁺ (aq.) will electrode potential become equal to its standard electrode potential ? [Ans. 1 M]
- 13. The standard reduction potentials of three metals cations X, Y and Z are + 0.52, 3.03 and 1.18 V respectively. Arrange X, Y and Z in order of increasing reducing power. [Ans. X < Z < Y]
- **14.** An electrochemical cell consists of two electrodes *i.e.*, Anode and Cathode. What is the direction of flow of electrons in this cell ?
- 15. Why anode is negatively charged in an electrochemical cell?
- **16.** Out of Zn and Cu vessel one will be more suitable to store 1 M HCl?

[Ans. Cu]

Given
$$E_{Zn^{2+}/Zn}^{\theta} = -0.76 \text{ V}, \ E_{Cu^{2+}/Cu}^{\theta} = +0.34 \text{ V}.$$

15. Is it safe to stir 1 M AgNO₃ solution with copper spoon ?

Given
$$E_{Ag^+/Ag}^{\theta} = + 0.80 \text{ V}, \quad E_{Cu^{2+}/Cu}^{\theta} = + 0.34 \text{ V}.$$
 [Ans. No]

1. Identify oxidant and reductant in the reaction :

 $I_2 (aq) + 2S_2O_3^{2-} (aq) \longrightarrow 2 I^- (aq) + S_4O_6^{2-} (aq).$

- 2. Calculate oxidation number of Fe in Fe_3O_4 and write a suitable justification of your answer.
- 3. Oxidation-reduction reactions are complementary. Explain.
- 4. Write formula for the following compounds :
 - (i) Mercury (II) chloride
 - (ii) Nickel (II) sulphate
 - (iii) Iron (III) sulphate
 - (iv) Chromium (III) oxide
- 5. Justify that the reaction : $H_2O(s) + F_2 \longrightarrow HF + HOF$ is a redox reaction.
- **6.** A decomposition reaction may or may not be a redox reaction. Write two decomposition reactions in support of the statement.
- Split the reaction 2 K (s) + C1₂ (g) → 2 KC1 (s) into oxidation and reduction half reactions.
- 8. Calculate the oxidation number of underlined elements in following compounds:

(i) CaO_2 (ii) $H_2S_2O_7$ (iii) K_2MnO_4 (iv) KI_3

- 9. Write the functions of salt bridge in an electrochemical cell.
- **10.** Define the term redox couple. Write the practical application of redox couple.
- The standard reduction potentials of two metals A and B are 0.76 V and + 0.34 V respectively. An electrochemical cell is formed using electrodes of these metals.
 - (i) Identify the cathode and anode.
 - (ii) Write the direction of flow of electron.



- 1. Calculate oxidation number of :
 - (i) Cr in $Cr_2O_4^{2-}$
 - (ii) $O in KO_2$
 - (iii) Na in Na₂O₂.
- **2.** Account for the following :
 - (i) HNO_3 acts as oxidizing agent while HNO_2 can act both as reducing and oxidizing agent.
 - (ii) AgF_2 is unstable compound and act as a strong oxidizing agent.
 - (iii) Ozone acts as an oxidising agent.
- Permanent ion (MnO₄⁻) reacts with sulfur dioxide gas in acidic medium to produce Mn²⁺ ion and hydrogen sulphate ion. Write ionic equation and balance by ion electron method.
- 4. Balance the following equation by oxidation number method : $P_4(s) + OH^-(aq) \longrightarrow PH_3 + H_2PO_2^-(aq)$ [*Basic Medium*]
- 5. Balance the following equation by ion electron method :

$$C1_2O_7(g) + H_2O_7(1) \longrightarrow ClO_2^-(aq) + O_2(g)$$
 [Basic medium]

6. Depict the galvanic cell in which the reaction

 $Zn(s) + 2Ag^{+}(aq) \longrightarrow Zn^{2+}(aq) + 2Ag(s)$ takes place. Further show :

- (i) Which electrode is negatively charged ?
- (ii) The carriers of the current in the cell
- (iii) Individual reaction at each electrode.
- 7. Explain with suitable reasons :
 - (i) Reaction $FeSO_4(aq) + Cu(s) \longrightarrow CuSO_4(aq) + Fe$ does not occur.
 - (ii) Zinc can displace copper from aqueous CuSO₄ solution but Ag cannot.
 - (iii) Solution of AgNO₃ turns blue when copper rod is immersed in it.



- 1. (i) MnO_4^{2-} undergoes disproportionation reaction in acidic medium but MnO_4^{-} does not. Give reason.
 - (ii) Give one example each of the following redox reactions:
 - (a) Combination reaction
 - (b) Decomposition reaction
 - (c) Metal displacement reaction
- 2. Consider the cell reaction of an electrochemical cell : Ni(s) + 2 Ag⁺(aq) \rightarrow Ni²⁺ (aq) + 2 Ag (s) and answer the following questions :
 - (i) Write anode and cathode half reactions.
 - (ii) Mention the direction of flow of electrons.
 - (iii) How is the electrical neutrality maintained in the solutions of the two half cells ?
 - (iv) Write the formula for calculating standard emf of this cell.
 - (v) How does the emf change when the concentration of silver ions is decreased ?
- 3. Justify the reason that following reactions are redox reactions.
 - (a) CuO (s) + $H_2(g) \longrightarrow Cu(s) + H_2O(g)$
 - (b) $\operatorname{Fe}_2O_3 + 3\operatorname{CO}(g) \longrightarrow 2 \operatorname{Fe}(g) + 3\operatorname{CO}_2(g)$
 - (c) $\operatorname{NH}_3(g) \operatorname{5O}_2(g) \longrightarrow \operatorname{4NO}(g) + \operatorname{5H}_2\operatorname{O}(g)$
 - (d) $BCl_3(g) + 3 LiAlH_4 \longrightarrow B_2H_6 + LiCl + AlCl_3$
 - (e) $2K + F_2 \longrightarrow 2KF$
- [Hints:- CuO is oxidizing agent, H_2 is acting as reducing agent because Cu (II) is changing to Cu (0) by gain of $e^- H_2$ is getting oxidised to H_2O (g), its oxidations sate is changing from 0 to +1, by loss of electrons.
 - (ii) It is redox reaction: Fe_2O_3 is getting reduced to fe. CO is getting oxidised to CO_2 .]



- **4.** Using standard electrode : Predict if the reaction between as the following is feasible.
 - (i) Fe^{3+} (aq) and I^{-} (aq)
 - (ii) Ag^+ and Cu
 - (iii) Fe^{3+} and $Br^{-}(aq)$
 - (iv) Ag and Fe^{3+} (aq)
 - (iv) Br_2 (aq) and Fe^{2+} (aq)
- Hint: $-E_{I_2/\Gamma}^{\theta} = 0.541 \text{ V}, E_{Cu^{2+}/Cu}^{\theta} = 0.34 \text{ V}, E_{Br_2/Br^-}^{\theta} = 1.09 \text{ V}, E_{Ag^+/Ag}^{\theta} = 0.80 \text{ V}, E_{Fe^{3+}/Fe^{2+}}^{\theta} = 0.77 \text{ V}.$
- **5.** Draw the diagram for the galvanic cell which would have overall chemical reaction as

$$Zn+2Ag^{+} \longrightarrow Zn^{2+}+2Ag.$$

Answer the following :

- (i) Write the reactions occurring at each electrode.
- (ii) In which directions do the electrons flow in the external circuit?
- (iii)Name the salt to be taken in salt bridge.
- (iv)Label the anode and cathode.
- (v) How does the EMF change when the concentration of solvers ions is decreased?

HOTS QUESTIONS

1. 6×10^{-3} mole K₂Cr₂O₇ reacts completely with 9×10^{-3} mole Xⁿ⁺ to give XO₃⁻ and Cr³⁺. Find the value of X.

Ans.
$$K_2 Cr_2 O_7 + X^{n+} \longrightarrow X^{+5} O_3^{-} + Cr^{3+}$$

 $6 \times 10^{-3} \times 6 = (5-n) \times 9 \times 10^{-3} \longrightarrow n = 1$

2. For the redox reaction

 $K_2Cr_2O_7 + XH_2SO_4 + YSO_2 \longrightarrow K_2SO_4 + Cr_2(SO_4)_3 + ZH_2O$ What is the sum of x + y + z

Ans.
$$K_2Cr_2O_7 + H_2SO_4 + 3SO_2 \longrightarrow K_2SO_4 + Cr_2(SO_4)_3 + H_2O$$

 $\therefore x = 1 \quad y = 3 \quad z = 1 \quad \therefore x + y + z = 5$

Redox Reactions



3. An aqueous solution containing 1M each of Au⁺³, Cu⁺², Ag⁺, Li⁺ is being electrolysed using inert electrodes the value of standard potentials are $E^{\theta}_{Ag^+/Ag} = 0.80 \text{ V}, \quad E^{\theta}_{Cu^2+Cu} = 0.34 \text{ V}, \quad E^{\theta}_{Au^{3+}/Au} = 1.50 \text{ V}, \quad E^{\theta}_{Li^+/Li} = -3.03 \text{ V}$ With increasing voltage, find the sequence of deposition of metals on the cathode.

Ans. Only Au^{3+} , Ag^+ and Cu^{2+} will deposit at cathode.

 Li^+ will not deposit at cathode be cause SRP of water is -0.8274V So after Cu^{2+} ; H₂ will evolve at cathode.

4.
$$E^{\theta}$$
 for $Cl_2(g) + 2l^- \longrightarrow 2Cl^-(aq.)$ is 1.36 V, then calculate.
 E^{θ} for $4Cl^-(aq.) \longrightarrow 2Cl_2(g) + 4e^-$
Ans. $E^{\theta}_{Cl^-/Cl_2} = -1.36 E^{\theta}$ is independent of amount of substance

5. Why salt bridge is made up of saturated solution of KNO₃ in agar–agar.
 Ans. Velocities of both K⁺ and NO₃⁻ are nearly the same.





UNIT TEST

Maximum Marks : 20

Time Allowed: 1 hr

General Instructions:

(i) All questions are compulsory.

1.	Identify the oxidised and Reduced species in the following reaction [1] $H_2S + Cl_2 \longrightarrow 2HCl + S$				
	2	2	(c) Both H_2 , Cl_2	(d) None of the	ese
2.	What is the (a) +1	e oxidation sta (b) +3	te of Br in BrO ₃ ⁻ ? (c) +4	(d) +5	[1]
3.	•	• •	ion in Redox Reaction $T^- \longrightarrow PH_3 + 3H_2PC$		[1]
4.	What is a 1	edox couple?	Give one example.		
5.	•		on given below : \longrightarrow Cu(s) + H ₂ O(g))	[1]
6.	Assign oxi (a) NaH ₂ <u>P</u>		to the underlined element $I_4 P_2 O_7$ (c) $K_2 Mn O_7$		[2]
7.	-		olysis in following cas CuCl ₂ with platinum		[2]
8.	Answer fo (i) Wh (ii) Wh	llowing : ich electrode i at are carrier c	$(s) + 2Ag^+ (aq.) \longrightarrow$ s negatively charged for current in the cell ? n at each electrode.		s) [3]
9.	$Hg^{2+}/Hg =$ (i) Wh	$0.79V \text{ Mg}^{2+}/$ ich one is stro	$K = -2.93V, Ag^{+}/Ag =$ Mg = -2.37V, Cr ³⁺ /Cr ng reducing agent ? ng oxidising agent ?		[3]
	(iii) Wh	ich redox coup	ole is a stronger reduct	ing agent than H ⁺ /H	I ₂ ?
10.			-electron or oxidation $\longrightarrow PH_3(g) + H_2Pe_{*****}$		[5] edium]
			Red	ox Reactions	139

Chapter - 9

Hydrogen

FAST TRACK : QUICK REVISION

• Hydrogen is the first element in the periodic table and also the lightest element known. Electronic configuration of Hydrogen is 1s¹.

• Isotopes of hydrogen :

- (i) Protium $\binom{1}{1}$ H)
- (ii) Deuterium $\binom{2}{1}$ H or $\binom{2}{1}$ D)
- (iii) Tritium $\binom{3}{1}$ H or $\binom{3}{1}$ T)
- Preparation of Dihydrogen :
 - (i) Laboratory preparation : $Zn + 2H^+ \rightarrow Zn^{2+} + H_2$.
 - (ii) Commercial preparation : By electrolysis of acidified water.
 - (iii) High purity dihydrogen is obtained by electrolysing warm aqueous barium hydroxide.
- Properties :
 - * Reaction with halogen: $H_2 + X_2 \longrightarrow 2HX [X = F, Cl, Br, I]$

* Reaction with oxygen: $H_2(g) + O_2(g) \xrightarrow{\Delta} 2H_2O(l);$ $\Delta H^{\circ} = -285.9 \text{ kJ mol}^{-1}$

* Reaction with nitrogen: $3H_2(g) + N_2(g) \xrightarrow{\Delta} 2NH_3(g);$ $\Delta H^{\sigma} = -92 \text{ kJ mol}^{-1}$

* Reaction with alkali metals: $H_2(g) + 2M(g) \xrightarrow{\Delta} 2MH(s)$

It is relatively inert at room temperature due to the high H-H bond enthalpy.

• Uses of Dihydrogen :

(i) For synthesis of Ammonia (NH₃)

- (ii) For production of Methanol (CH₃OH)
- (iii) In oxyhydrogen torches
- (iv) In a fuel cell.

• Hydrides

- (i) **Ionic or salt like or saline hydrides** are formed with most of the *s*-block elements. Significant covalent character is found in LiH, BeH₂ and MgH₂.
- (ii) **Covalent or Molecular hydrides** are formed with most of the *p*-block elements. There are further classified as :
- (a) Electron deficient hydrides are formed by group 13 elements *e.g.*, B_2H_6 . They acts as Lewis acid.
- (b) Electron Precise hydrides are formed by group 14 elements *e.g.*, CH_4 .
- (c) **Electron rich hydrides** have lone pair of electrons on central atoms of the molecules. Elements of group 15-17 form these types of hydrides.

 $\rm NH_3, \rm HF$ has high m.p./b.p. due to presence of intermolecular hydrogen bonding.

(iii) Metallic or Non-stoichiometric or Interstitial hydrides are formed by d and f-block elements. For example La H_{2.87} or NiH_{0.6-0.7}.

• **Water :** (H₂O)

Hard water : Hard water contains calcium and magnesium salts in the form of hydrogencarbonate, chloride and sulphate. Hard water does not give lathers with soap.

Soft water : Water free from soluble salts of calcium and magnesium is soft water.

Types of Hardness :

Temporary hardness is due to presence of calcium or magnesium hydrogen carbonate in water. Temporary hardness can be removed by :

(i) Boiling

(ii) Clark's Method

Permanent hardness :

Such hardness is due to presence of calcium or magnesium chlorides and sulphates.



Hydrogen



Permanent hardness can be removed by :

- (i) Treatment with washing soda
- (ii) Calgon's method

(iii) Ion exchange method.

Demineralised or Deionised water : Water free from all soluble mineral salts is known as **demineralised water.**

• Hydrogen Peroxide (H₂O₂)

Preperation :

(i) By electrolytic oxidation of acidified sulphate solutions at high current density.

(ii) 2-Ethylanthraquinol $\underbrace{O_2 \text{ (air)}}_{H_2/Pd}$ H₂O₂ + (Oxidised product)

Physical Properties

- (i) Miscible with water in all proportions.
- (ii) A 30% of H_2O_2 solution is marked as '100 volume' hydrogen peroxide.

• Chemical Properties :

- (i) It acts as an oxidising as well as reducing agent.
- (ii) Oxidising action in acidic medium :

 $2\mathrm{Fe}^{2+}(\mathrm{aq}) + 2\mathrm{H}^{+}(\mathrm{aq}) + \mathrm{H}_{2}\mathrm{O}_{2}(\mathrm{aq}) \longrightarrow 2\mathrm{Fe}^{3+}(\mathrm{aq}) + 2\mathrm{H}_{2}\mathrm{O}(l)$

(iii) Reducing action in acidic medium :

 $2MnO_4^- + 6H^+ + 5H_2O_2 \longrightarrow 2Mn^{2+} + 8H_2O + SO_2$

- Storage of H₂O₂:
 - (i) Stored in wax-linked glass or plastic vessels in dark. Urea can be added as a stabiliser.
 - (ii) It is kept away from dust because dust can induce explosive decomposition of the compound.
- Uses of H_2O_2 :
 - (i) As an antiseptic it is sold in the market name **perhydrol**.
 - (ii) In synthesis of hydroquinone.
 - (iii) As a bleaching agent.





 Auto-protolysis of water: Water accepts a proton from other water molecule to from H₃O⁺ and OH⁻ this porous is called auto – protolysis of water

$$H_2O(l) + H_2O(l) \Longrightarrow H_3O^+(aq) + OH^-(aq)$$

Its significance is that water can act as acid as well as base i.e. it is amphoteric in nature.

- 2. Hydrogen economy: It is transportation and storage of energy in the form of liquid or gaseous hydrogen. Advantage of hydrogen economy is that energy is transmitted in the form of dihydrogen and not as electric power
- **3.** Hydrogenation: It is a process of converting polyunsaturated oils in edible fats.

Vegetable oil + H₂ $\frac{\text{Ni}}{473\text{K}}$ Vanaspati ghee (fat).

4. Syngas: It is a mixture of CO and H₂ in 1:1 ratio and also known as water gas or syntnesis gas.

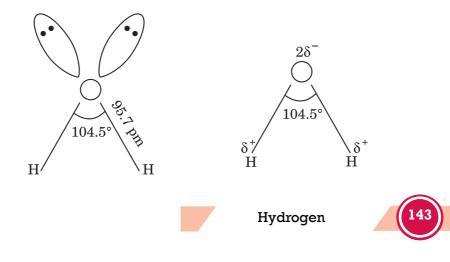
$$C(s) + H_2O(g) \xrightarrow{1270 \text{ K}} CO_2 + H_2(g)$$

Carbon Carbon monoxide

5. Water gas shift reaction.

$$CO + H_2O \xrightarrow{673 \text{ K}} CO_2 + H_2$$

- 6. Fuel-cell:- Fuel cell is a cell in which chemical energy of fuel is converted into electrical energy.
- 7. Structure of water: It is bent molecule in gas phase with HOH bond angle 104.5° and O–H bond length of 95.7 pm as shown if figure

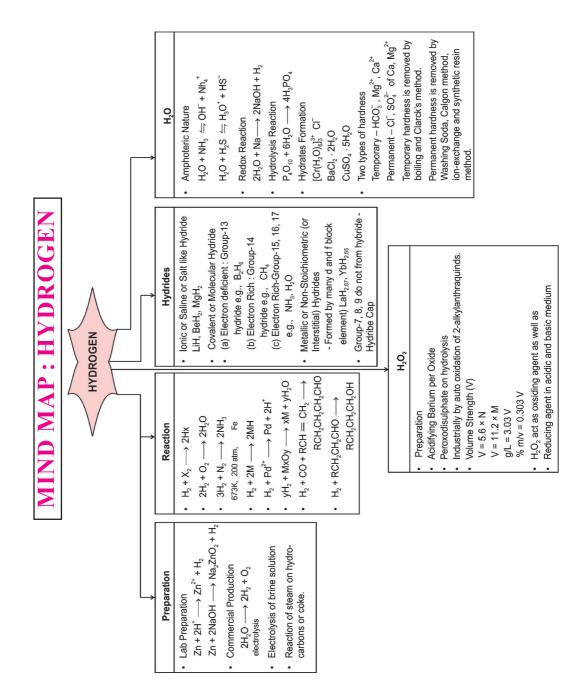


- 8. Calgon:- It is sodium polymetaphosphate (NaPO₃)_n it is used to remove. Permanent hardness of water.
- **9. De-ionized water:** Pure di-mineralised (ionized water) free from all soluble mineral matter is obtained by passing water successively through a cation exchanger (in the H⁺ form) and an anion exchanger for removal by cation and anions

 $2RH + M^{2+} \longrightarrow MR_{2} + 2H^{+}$ (R is the resin anion M^{2+} is cation) $RNH_{2} + H_{2}O \longrightarrow RNH_{3}^{+} OH^{-}$ $RNH_{3}^{+} + OH^{-} + X^{-} \longrightarrow RNH_{3}^{+} . X^{-} + OH^{-}$ $H^{+} + OH^{-} \longrightarrow H_{2}O$











MULTIPLE CHOICE QUESTIONS (MCQ)

1.	What is the strength of 20 volume solution of H_2O_2 ?						
	(a) 5g/L	(b) 10g/L	(c) 30g/L	(d) 60g/L			
2.	The number of H_2O molecules which are involve in Hydrozen bonding in $CuSO_4$ ·5 H_2O molecule itself is						
	(a) H ₃ PO	(b) H_3PO_3	(c) H_3PO_4	(d) PH ₃			
3.	P_4O_{10} on h	ydrolysis produc	es				
	(a) H ₃ PO	(b) H_3PO_3	(c) H_3PO_4	(d) PH ₃			
4.	D ₂ O has hi	igher value of foll	owing physical pa	rameters than H ₂ O, excep	t		
	(a) Molec	cular Mass	(b) Melting I	Point			
	(c) Density	у	(d) Dielectric	e Constant			
5.	Which one	e is ionic hydride	in nature?				
	(a) CrH	(b) NH ₃	(c) H ₂ O	(d) NaH			
6.	Syn-gas is	a mixture of					
	(a) CO+N	N_2 (b) O_3	(c) $\mathrm{CO} + \mathrm{H}_2$	(d) $CO + H_2CO_3$			
7.	The electro	onic conguration	of 'D' (Isotope of I	Hydrogen)			
	(a) $1s^2$	(b) $1s^22s^1$	(c) $1s^1$	(d) $1s^22s^22p^1$			
8.	Which gro	oup forms hydride	?				
	(a) 6	(b) 7	(c) 8	(d) 9			
9.	When Mn are involve	•	${}_{2}O_{2}$ in basic medi	um then following species	S		
	(a) MnO ₂	(b) O ₂	(c) OH ⁻	(d) Mn^{2+}			
10	. Select the	incorrect stateme	nt for H_2O_2 structu	ire			
	(a) It is not	on planar					
	(b) O-O b	oond length is mo	re in gaseous state	than in solid phase			
	(c) Both O	H bond are in dif	ferent plane				
	(d) O-O-	H bond angle in g	gas phase is more	than in solid phase			
ns:	1. (d) 2. (d	3.(c) 4.(d)	5. (d) 6. (c) 7. ((c) 8. (a) 9. (d) 10. (d)			

Ans: 1. (d) 2. (d) 3. (c) 4. (d) 5. (d) 6. (c) 7. (c) 8. (a) 9. (d) 10. (d)





TRUE AND FALSE TYPE QUESTIONSE

- **1.** H_2O_2 decomposes slowly on exposure to light.
- **2.** H_2O_2 on reaction with Pbs convert Pbs into Pb.
- **3.** Chemically calgon is sodium hexametaphosphate $Na_6P_6O_{18}$.
- **4.** NH_3 is electron rich hydride.
- 5. Phosphorus form PH₅.
- 6. H_2 gas cannot reduce Pb²⁺ ion.
- 7. Hydroformylation of olefins yields aldehydes which futher undergoes reduction to give alcohols.
- 8. Hydrogenation of vegetable oils using nickel as catalyst gives edible fats.
- **9.** Ice has cage like structure with air spaces.
- **10.** Soft water gives lather with Soap.

Ans: 1. True	2. False	3. True	4. True	5. False
6. False	7. True	8. True	9. True	10. True

FILL IN THE BLANKS

1. Cation exchange resin contain large organic molecule with _____ group.

2. At atmospheric pressure ice crystallises in for.

- **3.** Due to high ______ of H₂O, H₂O has a very strong hydrating tendency.
- 4. Water is present in $[Cr(H_2O)_6]^{3+}.3Cl^-$ in the form of _____.
- The H-H bond dissociation enthalpy of H₂ is ______, is the highest 5. for a single bond between two atoms of any elements.

6. In the Clark's method for softening compound is used.

- 7. During photosynthesis H₂O is _____.
- 8. BeH₂ and MgH₂ are ionic and _____ in nature.
- 9. When NaH is electrolysed, then ______ is released at anode.

10. Hydrated sodium aluminium silicate is _____.

Ans:	1. SO ₃ H	2. Hexag
	5 10 5 00 1 ¹	11 6 9 (11)

9. H₂

gonal 3. Dielectric 4. Coordinated water 5. 435.88 kj mol⁻¹ 6. Ca(HCO₃)₂ 7. Oxidised 8. Polymeric

Hydrogen

10. Zeolite/Permutit



MATCH THE COLUMNS

1.	Column -I				Column-II		
	А.	Boiling			p.	CaC	CO ₃
	В.	Clark's M	Method		q.	Mg	(OH) ₂
	C.	Washing	g soda		r.	NaA	AISiO ₄
	D.	Ion-excl	nange meth	od	s.	NaC	21
2.	Column -I				Column-II		
	A.	$H_2O + N$	H ₃ ≕OH	$+ \mathrm{NH_4}^+$		p.	Hydroformylation
	В.	$2H_2O +$	$2Na \longrightarrow 2$	2NaOH + H	[₂	q.	Acid base reaction
	C.	$P_4O_{10} +$	6Н ₂ О —	→ 4H ₃ PO ₄		r.	Redox Reaction
	D.	$2H_{2} + C$	O + RCH =	= CH ₂		s.	Hydrolysis reaction
			→ RCH ₂ CH	I ₂ CH ₂ O ₄			
An	s. 1.	$A \rightarrow q$,	$B \rightarrow p$,	$C \rightarrow s$,	D	ightarrow	r
	2.	$A \rightarrow q$,	$B \rightarrow r$,	$C \rightarrow s$,	D	\rightarrow]	0

ONE WORD ANSWER TYPE QUESTIONS

- 1. Name the gas release when zinc reacts with NaOH.
- 2. When brine solution is electrolysed then nature of solution will be?
- **3.** What happens when Al_4C_3 reacts with D_2O ?
- **4.** In which medium H_2O_2 act as reducing agent?
- 5. What is the chemical name of calgon's?
- 6. What happens when LiH reacts with Al_2C_{16} ?
- 7. What happens when warm aqueous Barium hydroxide solution is electrolysed?
- 8. What type of particles are emitted by Tritium?
- 9. What is the name for the following chemical reaction?

 $CO(g) + H_2O(g) \longrightarrow CO_2(g) + H_2(g)$





- **10.** What is the term used to refer "Transportation and storage of energy in the form of liquid or gaseous dihydrogen"
- Ans: 1. H₂ 2. Basic 3. $Al_4C_3 + 12D_2O \longrightarrow 3CD_4 + 4Al(OD)_3$ 4. Acidic, Basic 5. Sodium hexameter phosphate (Na₆P₆O₁₈)
 - 6. $8LiH + Al_2Cl_6 \longrightarrow 2LiAlH_4 + 6LiCl$
 - 7. H_2 gas is produced 8. β^- (beta negative)
 - 9. Water gas shift reaction 10. Hydrogen economy

ASSERTION AND REASON TYPE QUESTIONS

Each question contain statement-1 (Assertion) and statement-2 (Reason). Examine the statements carefully and mark the correct answer according to the instruction given below :

- (A) If both the statements are True and statement-2 is the correct explanation of statement-1
- (B) If both the statements are True and statement-2 is not the correct explanation of statement-1
- (C) If statement-1 is true and statement-2 is false
- (D) If statement-1 is false and statement-2 is true
- 1. Statement-1 : HF form extensive hydrogen bonding.
- Statement-2 : F has highest tendency to form hydrogen bonding.
- 2. Statement-1 : $MgCl_2$ solution produces lather with soap.
- Statement-2 : Hard water does not produce lather with soap.
- Statement -1 : Density of ice is less than water.
 Statement-2 : Ice has open cage structure.
- 4. Statement-1 : Water can act as acid as well as base.
 Statement-2 : Water can accept as well as donate H⁺ ion.
- 5. Statement-1 : $KMnO_4$ act as self indicator.
- Statement-2 : $KMnO_4$ only act as reducing agent.
- 6. Statement-1 : Washing soda (Na_2CO_3) is use to remove temporary hardness.

Statement-2 : Clark's method is used to remove temporary hardness.





7.	Statement-1	:	In cation exchange process, H^+ exchanges for Na ⁺ , Ca ²⁺ , Mg^{2+} .
	Statement-2	:	In anion exchang process OH^- exchanges for anion like Cl^- , HCO_3^- , SO_4^{-2-} .
8.	Statement-1	:	When Na reacts with H_2O , H_2 gas is release.

- Statement-2 : P_4O_{10} on hydrolysis produce H_3PO_3 .
- 9. Statement-1 : CH₄ is a covalent hydrides.
 Statement-2 : CH₄ is dectron precise type hydrides.
- Statement-1 : H₂ gas is use in metallurigical process.
 Statement-2 : H₂ gas is use as fuel.

Ans: 1. (a) 2. (d) 3. (a) 4. (a) 5. (c) 6. (d) 7. (b) 8. (c) 9. (a) 10. (b)

1-MARK QUESTIONS

- 1. Name the isotope of hydrogen which is radioactive in nature. [Ans. Tritium]
- 2. H⁺ ions does not exist freely and is always associated with other atoms or molecule. Explain.
- **3.** Give the composition of water gas. [Ans. CO, H₂]
- 4. Name the compound whose electrolysis in aqueous state, give high purity (99.95%) dihydrogen. [Ans. aq Ba(OH)₂ solution]
- 5. Give the main purpose of water gas shift reaction.
- 6. Write the chemical reaction occuring during coal gasification.
- 7. Name the element used in fuel cell for generating electricity. $[Ans. H_2]$
- **8.** Give an example of electron deficient covalent hydride. $[Ans. B_2H_6]$
- 9. Name the hydrides which have high potential for hydrogen storage. [Ans. Metallic hydrides]
- 10. Name the groups in *d*-block elements which do not form metallic hydrides.

[**Ans.** 7, 8, 9]

- **11.** H_2 is relatively inert at room temperature. Explain.
- 12. Complete the reaction : $C(s) + H_2O(g) \xrightarrow{1270 \text{ K}} (A) \underbrace{(g)}_{(g)} + (B) \underbrace{(g)}_{(g)} . \quad [Ans. CO, H_2]$





- 13. Name the phenomenon as a reason of which water has unusual boiling point.[Ans. Extensive hydrogen bonding]
- **14.** Draw structure of water.
- **15.** At atmospheric pressure ices crystallised in the form but at very low temperature it condenses to form. [**Ans.** Hexagonal, cubic]
- 16. Mention the temperature at which density of ice is maximum.[Ans. 4°C]
- **17.** Density of ice is than density of liquid water. [Ans. Less]
- **18.** Complete the reaction :

 $2H_2O(l) + 2Na(s) \longrightarrow$

- 19. How many hydrogen-bonded water molecules (s) are associted in CuSO₄.5H₂O. [Ans. One]
- 20. Name the compound used in Clark's method to remove temporary hardness of water. [Ans. Lime]
- **21.** Write the chemical formula of "Calgon". [Ans. $Na_4P_6O_{18}$]
- 22. A 30% solution of H₂O₂ is marketed as volume.[Ans. 100 volume]
- **23.** Draw gas phase structure of H_2O_2 .
- 24. Name the organic compound whose auto-oxidation is used to produce H₂O₂ commercially or industrially. [Ans. 2-Ethylanthraquinol]
- 25. How is heavy water obtained from ordinary water?

2-MARKS QUESTIONS

1. Complete the following reactions :

(i) $CO(g) + H_2(g) \xrightarrow{\Delta} Catalyst$ (ii) $Zn(s) + NaOH(aq) \xrightarrow{\Delta}$

- 2. Among NH₃, H₂O and HF which would you except to have highest magnitude of hydrogen bonding and why ?
- **3.** How do you except the metallic hydrides to be useful for hydrogen storage? Explain.
- **4.** How can the production of dihydrogen obtained from "Coal gasification" be increased ?

Hydrogen



- 5. Write the name of isotopes of hydrogen. What is the mas ratio of these isotopes ?
- **6.** Complete the reactions :

(i)
$$\operatorname{CO}(g) + 2\operatorname{H}_2(g) \xrightarrow[\text{Cobalt}]{\text{Catalyst}}$$

(ii) $\operatorname{CH}_4(g) + \operatorname{H}_2\operatorname{O}(g) \xrightarrow[\text{Ni}]{270K}$

7. Comment on the reactions of dihydrogen with :

(i) Chlorine, (ii) Sodium.

- **8.** Arrange the following :
 - (i) LiH, NaH, CsH (In increasing order of ionic character)

(ii) H—H, D—D, F—F (In decreasing order of bond dissociation enthalpy)

- 9. List two uses of dihydrogen.
- **10.** Complete the reactions :

(i) $H_2 + CO + RCH = CH_2 \longrightarrow$

(ii) $H_2 + RCH_2CH_2CHO \longrightarrow$

- 11. Give two reactions to show amphoteric nature of water.
- **12.** Complete the reactions :
 - (i) $2F_2(g) + 2H_2O(l) \longrightarrow$
 - (ii) $6CO_2(g) + 12H_2O(l) \longrightarrow$
- **13.** What is the difference between the term hydrolysis and hydration.
- **14.** What do you understand by term 'autoprotolysis' of water ? What is its significance ?
- **15.** What causes the temporary and permanent harness of water ?
- **16.** Is demineralised or distill water useful for drinking purposes ? If not, how can it be made useful ?
- **17.** Explain the terms :
 - (i) Hydrogen economy.
 - (ii) Fuel cell.
- **18.** Write chemical reactions to justify that hydrogen peroxide can function as an oxidising as well as reducing agent.





- **19.** Compare the structure of H_2O and H_2O_2 .
- **20.** How does H_2O_2 behaves as a bleaching agent ?
- **21.** H_2O_2 acts as an oxidizing as well as reducing agent. Why?

1. Complete the chemical reactions :

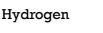
(i) $8LiH + Al_2Cl_6 \longrightarrow$

(ii) $2\text{LiH} + \text{B}_2\text{H}_6 \longrightarrow$

- What do you understand by : (i) Electron deficient, (ii) Electron precise, (iii) Electron rich compounds of hydrogen ? Provide justifications with suitable examples.
- **3.** What do you understand by the term "non-stoichiometric hydrides"? Do you expect this type of the hydrides to be formed by alkali metals. Explain and Justify your answer.
- **4.** Arrange the following :

(i) CaH₂, BeH₂, TiH₂ (in order of increasing electrical conductance)
(ii) NaH, MgH₂, H₂O (in order of increasing bond dissociation enthalpy)
(iii) Li, F, H (in order of increasing ionisation enthalpy)

- 5. What do you understand by the terms :
 - (i) Syn gas (ii) Water gas shift reaction (iii) Producer gas.
- 6. Would gas except the hydrides of N, O and F to have lower boiling point than the hydrides of their subsequent group members ? Give reasons.
- 7. Can phosphorous with outer electronic configuration $3s^23p^3$ form PH₅? Explain.
- 8. Why and how the hydrogen is regarded as a fuel of future ? Explain.
- **9.** Write the reactions when dihydrogen reacts with (i) O₂ (ii) N₂ (iii) Cl₂ under specific conditions.
- **10.** Name the hydrides :
 - (i) Which is non stoichiometric in nature ?
 - (ii) Which are stoichiometric compounds?
 - (iii) Which has electron rich type hydrides ?





- **11.** Complete the reactions :
 - (i) $CaO(s) + H_2O(g) \longrightarrow$ (ii) $AlCl_3(g) + H_2O(l) \longrightarrow$ (iii) $Ca_3N_2(s) + H_2O(l) \longrightarrow$
- **12.** Discuss the principle and method of softening of hard water by synthetic exchange of resin method.
- 13. What is meant by 'demineralised' water and how can it be obtained ?
- 14. What properties of water make it useful as a solvent ? What types of compound can it (i) dissolved (ii) hydrolyse ?
- **15.** Calculate the strength of 10 volume solution of H_2O_2 .
- **16.** Complete the reactions :
 - (i) $2Fe^{2+}(aq) + 2H^{+}(aq) + H_2O_2(aq) \longrightarrow$ (ii) $HOCl + H_2O_2 \longrightarrow$ (iii) $Mn^{2+} + H_2O_2 \longrightarrow$
- **17.** Give three uses of H_2O_2 .
- **18.** Complete the reactions :

(i)
$$CaC_2 + 2D_2O \longrightarrow$$

(ii)
$$SO_3 + D_2O \longrightarrow$$

(iii)
$$Al_4C_3 + 12D_2O \longrightarrow$$

- **19.** Give the limitations of using H_2 as a fuel.
- **20.** H_2O_2 is stored in a wax lined glass or plastic vessels. Explain an equation showing decomposition of H_2O_2 on exposure to light.

- 1. Answer the following :
 - (a) Name the most abudant form of hydrogen isotope. $[Ans. 1^{1}H]$
 - (b) Name the particles emitted by tritium. [Ans. β^{-}]
 - (c) Mixture of CO and H_2 is used for preparation [Ans. Methanol]
 - (d) Name the catalyst used in Haber's Process for manufacture of $NH_3(g)$.

[Ans. Fe]

[Ans. NH₃, H₂O]



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(e) Name two electron rich hydrides.



- **2.** Answer the following :
 - (a) During Clark's method. Name the compound in which Mg is precipitated out. [Ans. Magnesium Hydroxide]
 - (b) Give the formula of Zeolite used in ion exchange method to remove permanent hardness of water. [Ans. NaAlSiO₄]
 - (c) Complete the reaction :

 $BaO_2.8H_2O(s) + H_2SO_4(aq) \rightarrow$

- (d) H_2O_2 is miscible with water. Assign reason.
- (e) Name the compound when can be used as a hair beach, mild antiseptic in the form of perhydrol. [Ans. H₂O₂]
- **3.** (a) Complete the following chemical equations

(b)
$$_$$
 + water \longrightarrow CaCO₃ + NH₃ (Ammonia)

- (c) _____ + Hydrogen peroxide $\xrightarrow{H^+}$ CrO₅+_____
- (d) $Na_2O + H_2O \longrightarrow$
- (e) $D_2O + Na_3As \longrightarrow$
- 4. Describe the usefulness of water in biosphere and biological systems.

HOTS QUESTIONS

- 1. Calculate the hardness of water sample which contains 0.001 mole of MgSO₄ dissolved per litre of water.
- Ans. 1 mole MgSO₄ = 1 mole CaCO₃ 10^{-3} Mole MgSO₄ = 10^{-3} mol CaCO₃ $\therefore 0.120$ g MgSO₄ = 0.1g CaCO₃ in 1000 mL \therefore Hardness = =100ppm
- 2. 2g of Al is treated separately with excess dilute H_2SO_4 and excess NaOH. The ratio of volumes of Hydrogen evolved under similar condition is $\frac{x}{y}$. Find $\frac{x}{y}$
- Ans. $2A1 + 3H_2SO_4 \longrightarrow Al_2(SO_4)_3 + 3H_2$ $2A1 + 2NaOH + 2H_2) 2NaAlO_2 + 3H_2$ \therefore Ratio is 1 : 1



- 3. What mass of CaO will be required to remove the hardness of 1000 litres of water containing 1.62g of Ca(HCO₃)₂ per litre?
- Ans. $Ca(HCO_3)_2 + CaO \longrightarrow 2CaCO_3 + H_2O + CO_2$ 162g 56g $\therefore 5.6 \times 10^2$ g be cause solution has 1620 g Ca(HCO₃)₂ What is the volume of O₂ liberated at N.T.P. by complete decomposition 4. of 100mL of 2m solution of H₂O₂? $= 11.2 \times M = 22.4$ Ans. Volume strength = 0.1L i.e. $22.4 \times 0.1 = 2.24L$ O₂ released
- Mention an example in which H₂O acts as reducing agent. 5.

Ans. $2F_2 + 2H_2O \longrightarrow O_2 + 4HF$

100mL





UNIT TEST

Tim	e Allowed: 1 hr Maximum Mark	ks : 20					
(i)	<i>eral Instructions:</i> All questions are compulsory. Maximum marks carried by each question are indicated against it.						
1.	Hydrogen has maximum oxidation state in [1]						
	(a) NaH (b) MgH_2 (c) H_2O (d) C & H						
2.	Which one does not cause hardness of water?						
	(a) $MgCl_2$ (b) $CaCl_2$ (c) $MgSO_4$ (d) $AlCl_3$						
3.	Give one reaction for preparation of hydrogen gas in laboratory.	[1]					
4.	What causes the hardness of water?	[1]					
5.	Draw the structure of H_2O_2 .	[1]					
6.	Complete the reaction with balancing	[2]					
	(a) $\operatorname{Fe}^{2+}(\operatorname{aq.}) + \operatorname{H}^{+}(\operatorname{aq.}) + \operatorname{H}_{2}\operatorname{O}_{2}(\operatorname{aq.}) \longrightarrow$						
	(b) $HOCl + H_2O_2 \longrightarrow$						
7.	What is Hydrogen Economy. What are its advantage?	[2]					
8.	Explain the following	[3]					
	(i) Atomic hydrogen or oxy-hydrogen torch function for cutting and welding purposes. Why?						
	(ii) CaH_2 , BeH_2 and TiH_2 arrange in order of increasing electrical conductance and give reason.						
	(iii) Water shows amphoteric behaviour, support by giving appropriate example.						
9.	What are different types of hydrides? Give example.	[3]					
10.	Discuss the principle and method of softening of hard water by [5] synthetic ion-exchange resins.						



Hydrogen





Chapter - **10**

s-Block Elements

FAST TRACK : QUICK REVISION

- s-block elements consists of group-I (Alkali metals) and group-2 (Alkaline earth metals).
- Group 1st elements Li, Na, K, Rb, Cs, Fr.
- Group 2nd elements Be, Mg, Ca, Sr, Ba, Ra.
- Atomic radius : Atomic radius of alkali metals are greater than alkaline earth metals.
- Hydration enthalpy : Decreases with increases in ionic sizes.
- **Ionic mobility :** Smaller the size of ion, more highly it is hydrated and hence lower is its ionic mobility.

 $Li^+ < Na^+ < K^+ < Rb^+ < Cs^+$

- **Ionisation enthalpies :** 1st I.E. of group 1st is smaller than group 2nd elements but 2nd I.E. of group 2nd is smaller than group 1st elements.
- Flame colouration : Due to low I.E., *s*-block elements and their salts imparts characteristics colour of oxidising flame (except Be and Mg). Be and Mg do not show flame colouration because they have small size and very high ionisation enthalpy.
- **Reducing character :** Due to large negative electrode potentials alkali metals are stronger reducing agent than alkaline earth metal.
- Reactivity towards air :

 $4\text{Li} + \text{O}_2 \longrightarrow 2\text{Li}_2\text{O} \text{ (Lithium oxide)}$ $2\text{Na} + \text{O}_2 \longrightarrow \text{Na}_2\text{O}_2 \text{ (Sodium peroxide)}$ $M + \text{O}_2 \longrightarrow \text{MO}_2 \text{ (M = K, Rb, Cs metal superoxide)}$

Alkaline earth metals being smaller in size do not from superoxides.

• Reactivity towards H₂O :

 $2M + 2H_2O \longrightarrow 2MOH + H_2$ (Alkali metal) $M + 2H_2O \longrightarrow M(OH)_2 + H_2$

(Alkaline earth metals)

• Reactivity towards hydrogen :

$$\begin{split} & 2M + H_2 \longrightarrow 2MH \qquad (M = \text{Li, Na, K, Rb, Cs}) \\ & M + H_2 \longrightarrow MH_2 \qquad (M = \text{Mg, Ca, Sr, Ba}) \\ & 2\text{BeCl}_2 + \text{LiAlH}_4 \longrightarrow 2\text{BeH}_2 + \text{LiCl} + \text{AlCl}_3. \end{split}$$

• Reactivity towards halogens :

 $2M + X_2 \longrightarrow 2MX (M = Li, Na, K, Rb, Cs)$ $M + X_2 \longrightarrow MX_2 (M = Mg, Ca, Sr, Ba)$ $BeO + C + Cl_2 \xleftarrow{600-800 \text{ K}} BeCl_2 + CO$

- Solution in liquid ammonia : The fresh solution of alkali metals and alkaline earth metals (except Be and Mg) is deep blue, paramagnetic and highly reducing due to presence of ammoniated electrons.
- Solubility of alkaline earth metal carbonate in water : Li₂CO₃ < Na₂CO₃ < K₂CO₃ < RbCO₃ < Cs₂CO₃
- Solubility of alkaline earth metal carbonates in water. BaCO₃ < SrCO₃ < CaCO₃ < MgCO₃ < BeCO₃
- Solubility of alkaline earth metal sulphates in water : BaSO₄ < SrSO₄ < CaSO₄ < MgSO₄ < BeSO₄
- Thermal stability of alkali metal carbonates : Li₂CO₃ < Na₂CO₃ < K₂CO₃ < Rb₂CO₃ < Cs₂CO₃
- Thermal stability of alkaline earth metal carbonates : BeCO₃ < MgCO₃ < CaCO₃ < SrCO₃ < BaCO₃
- Anamolous behaviour of Li and Be : It is due to very small size, high I.E. and high polarising power (*i.e.*, charge/radius)
- Diagonal relationship (similarities) between Li and Mg : (i) Both Li and Mg are hard.



(ii) Both react with N₂ to form nitrides.

 $6Li + N_2 \longrightarrow 2Li_3N$ $3Mg + N_2 \longrightarrow Mg_3N_2$

(iii) Decomposition of carbonates :

 $\begin{array}{c} \mathrm{Li_2CO_3} \longrightarrow \mathrm{Li_2O} + \mathrm{CO_2} \\ \mathrm{MgCO_3} \overset{\Delta}{\longrightarrow} \mathrm{MgO} + \mathrm{CO_2} \end{array}$

- (iv) Both LiCl and $MgCl_2$ are deliquescent. They form hydrates salts LiCl.2H₂O and MgCl₂.6H₂O.
- (v) **Decomposition of nitrates :**

 $4\text{LiNO}_{3} \xrightarrow{\Delta} 2\text{Li}_{2}\text{O} + 4\text{NO}_{2} + \text{O}_{2}$ $2\text{Mg(NO}_{3})_{2} \xrightarrow{\Delta} 2\text{MgO} + 4\text{NO}_{2} + \text{O}_{2}$

- Diagonal relationship (similarities) between Be and Al :
 - (i) Both are passive to acids due to formation of oxide layer.
 - (ii) Hydroxides of both dissolve in alkali to form $[Be(OH)_4]^{2-}$ and $[Al(OH)_4]^{-}$.
 - (iii) Chloride of both has bridged structure.
 - (iv) Both have tendency to form complexes of BeF_4^{2-} , AlF_6^{3-} .
- Manufacturing of washing soda (Na₂CO₃.10H₂O) :

Solvay process :

$$\begin{split} \mathrm{NH}_3(g) + \mathrm{CO}_2(g) + \mathrm{H}_2\mathrm{O}(l) &\longrightarrow \mathrm{NH}_4\mathrm{HCO}_3(\mathrm{aq}) \\ \mathrm{NH}_4\mathrm{HCO}_3(\mathrm{aq}) + \mathrm{NaCl}(\mathrm{aq}) &\longrightarrow \mathrm{NaHCO}_3(\mathrm{s}) + \mathrm{NH}_4\mathrm{Cl}(\mathrm{aq}) \\ \mathrm{2NaHCO}_3 & \stackrel{\Delta}{\longrightarrow} \mathrm{Na}_2\mathrm{CO}_3 + \mathrm{H}_2\mathrm{O}(l) + \mathrm{CO}_2(g) \\ \mathrm{2NH}_4\mathrm{Cl}(\mathrm{aq}) + \mathrm{Ca}(\mathrm{OH})_2 &\to \mathrm{CaCl}_2(\mathrm{s}) + \mathrm{2H}_2\mathrm{O}(l) + \mathrm{2NH}_3(g) \end{split}$$

• Manufacturing of caustic soda (NaOH) : Castner-Kellner cell.

Cathode : Na⁺ + $e^- \xrightarrow{Hg}$ Na-Hg Anode : Cl⁻ $\longrightarrow \frac{1}{2}$ Cl₂ + e^- 2Na-Hg + 2H₂O \longrightarrow 2NaOH + 2Hg + H₂ Plaster of paris : (CaSO₄.¹/₂H₂O) 2(CaSO₄.2H₂O) $\xrightarrow{\Delta}$ 2(CaSO₄).H₂O + 3H₂O Gypsum Chemistry Class XI

MULTIPLE CHOICE QUESTIONS (MCQ)

- 1. The alkali metals are low melting. Which of the following alkali metal is expected to melt if the room temperature rises to 30°C?
 - (a) Na (b) K
 - (c) Rb (d) Cs
- 2. Alkali metals react with water vigorously to form hydroxides and dihydrogen. Which of the following alkali metals reacts with water least vigorously?
 - (a) Li (b) Na
 - (c) K (d) Cs
- 3. The reducing power of a metal depends on various factors. Suggest the factor which makes Li, the strongest reducing agent in aqueous solution.
 - (a) Sublimation enthalpy (b) Ionisation enthalpy
 - (c) Hydration enthalpy (d) Electron-gain enthalpy
- 4. Metal carbonates decompose on heating to give metal oxide and carbon dioxide. Which of the metal carbonates is most stable thermally?
 - (a) MgCO₃ (b) CaCO₃
 - (c) SrCO₃ (d) BaCO₃
- 5. Which of the carbonates given below is unstable in air and is kept in CO₂ atmosphere to avoid decomposition.
 - (a) BeCO₃ (b) MgCO₃
 - (c) $CaCO_3$ (d) $BaCO_3$
- 6. Metals form basic hydroxides. Which of the following metal hydroxide is the least basic?
 - (a) $Mg(OH)_2$ (b) $Ca(OH)_2$
 - (c) $Sr(OH)_2$ (d) $Ba(OH)_2$
- 7. Some of the Group 2 metal halides are cavalent and soluble in organic solvents. Among the following metal halides, the one which is soluble in ethanol is
 - (a) BeCl₂ (b) MgCl₂
 - (c) CaCl₂ (d) SrCl₂



s-Block Elements



- 8. The order of decreasing ionisation enthalpy in alkali metals is
 - (a) Na > Li > K > Rb (b) Rb > Na > K > Li
 - (c) Li > Na > K > Rb (d) K > Li > Na > Rb
- 9. The solubility of metal halides depends on their nature, lattice enthalpy and hydration enthalpy of the individual ions. Amongst fluorides of alkali metals, the lowest solubility of LiF in water is due to
 - (a) Ionic nature of lithium fluoride
 - (b) High Lattice enthalpy
 - (c) High hydration enthalpy of lithium atom
 - (d) Low ionisation enthalpy of lithium atom
- 10. Amphoteric hydroxides react with both alkalies and acids. Which of the following Group 2 metal hydroxides is soluble in sodium hydroxide?
 - (a) $\operatorname{Be(OH)}_2$ (b) $\operatorname{Mg(OH)}_2$
 - (c) $Ba(OH)_2$ (d) $Ca(OH)_2$

ASSERTION-REASON TYPE QUESTIONS

The question given below contains statement -1 (Assertion) and Statement-2 (Reason) Each question has four choice (a), (b), (c) and (d) out of which only one is correct. Choice the correct option as under.

- (a) Statement-1 is true, statement-2 is true.Statement-2 is a correct explanation for statement-1.
- (b) Statement-1 is true, statement-2 is true ;Statement-2 is not a correct explanation of statement-1.
- (c) Statement -1 is true, statement-2 is false.
- (d) Statement-1 is false, statement-2 is true.
- 1. Statement-1: Sodium metal is softer than potassium metal.

Statement-2: Metallic bonding in Potassium is weaker than in Sodium.

2. Statement-1 : Be(OH)₂ is soluble in HCl and NaOH.

Statement-2 : $Be(OH)_2$ is amphoteric in nature.

3. Statement-1 : Metallic character of alkali metals increases on going down a group from top to bottom.





Statement-2 : Ionisation enthalpy of alkali metals increases on going down from top to bottom.
4. Statement-1 : Super oxides of alkali metals are diamagnetic.

Statement-2 : Super oxides contain the ion O_2 -which has one unpaired electron.

5. Statement-1 : Alkali metals do not impart colour to the flame.

Statment-2: Their ionization enthalpies are very low.

- 6. Statement-1 : Sodium cannot be obtained by chemical reduction of its ore.
 - Statment-2: Sodium is one of the strongest reducing agent.
- 7. Statement-1 : Beryllium hydroxide becomes soluble in excess alkali forming beryllate ion $[Be(OH)_4]^{2-}$.

Statment-2: Beryllium ion has greater tendency to form complexes.

FILL IN THE BLANKS

- 1. In the synthesis of sodium carbonate, the recovery of ammonia is done by treating NH_4Cl with $Ca(OH)_2$. The by product obtained in this process is
 - (a) NaCl (b) NaOH
 - (c) CaCl₂ (d) NaHCO₃
- 2. When sodium is dissolved in liquid ammonia, a solution of deep blue colour is obtained. The colour of the solution is due to.....
 - (a) Sodium ion (b) Ammoniated electron
 - (c) Sodium amide (d) Ammoniated sodium ion
- 3. By adding gypsum to cement.....
 - (a) Setting time of cement becomes less.
 - (b) Setting time of cement increases
 - (c) Colour of cement becomes light
 - (d) Shining surface is obtain.
- 4. A substance which gives crimson red flame and breaks on heating to give oxygen and a brown gas is
 - (a) Magnesium nitrate (b)
 - (b) Calcium nitrate
 - (c) Barium nitrate
- (d) Strontium nitrate



s-Block Elements



5.	The hydration energy of Mg^{2+} is greater than that of						
	(a) Al^{3+}				(d) Mg^{3+}		
6.	The active c	onstituent of bl	eaching po	wer is			
	(a) Ca(OCl))2	(b)	Ca(OCl)	Cl		
	(c) Ca(ClO	$_{2})_{2}$	(d)	Ca(ClO ₂))C1		
7.	2	${\rm CO}_2$ and incre	-		marines because it		
	(c) Produce	s ozone	(d)	None of	the above		
8.	hydrozide ar		which become	nes inert o le M is	te MSO ₄ , water insoluble on heating. The hydroxide		
9.		ic effect is max	Č,) 51		
9.	(a) Cs	(b) K		(d)	Ti		
10							
10.					or the following statement. eption of are ionic		
	(a) Barium	halide	(b)	Strontiun	n halide		
	(c) Berylliu	m halide	(d)	Calcium	halide		
11.	Flame test is	not given by					
	(a) Be	(b) Sr	(c) K	(d) Ca			
		MATC	H THE CO	OLUMNS	5		
In th	In the following questions more than one ention of column L and II may be						

In the following questions more than one option of column I and II may be correlated.

- **Column-I** 1. (i)
- **Column-II**

- (ii) Na
- (a) Insoluble sulphate Strongest monoacidic base (b)
- (iii) Ca

Li

- Most negative E° value among alkali metals. (c)
- (iv) Ba
- (d) Insoluble oxalate
- 6s² outer electronic configuration (e)





	Column-I		Column-II
(i)	CaCO ₃	(a)	Dentistry, ornamental work
(ii)	Ca(OH) ₂	(b)	Manufacture of sodium carbonate from caustic soda
(iii)	CaO	(c)	Manufacture of high quality paper
(iv)	CaSO ₄	(d)	Used in white washing.
	Column-I		Column-II
(i)	Column-I Cs	(a)	Column-II Apple green
(i) (ii)	-	(a) (b)	
. /	Cs		Apple green
(ii)	Cs Na	(b)	Apple green Violet

- (v) Sr (e) Crimson red
- (vi) Ba (f) Blue

Column-II

- (a) NaOH (a) Photo electric cells
 - (b) Na_2CO_3

Column-I

Liquid Na

- (b) Coolant in nuclear reactors
- (c) SO_2 absorber
- (d) Caesium (d) Detergent

1-MARK QUESTIONS

- **1.** What is the oxidation state of K in KO_2 ?
- 2. Why are group I element called alkali metals?
- 3. Potassium carbonate cannot be prepared by solvay process. Why ?
- 4. LiCl is soluble in organic solvent. Why?
- 5. Why are group I elements called alkali metals?
- 6. Alkali metals are strong reducing agents. Why?
- 7. Why do alkali metals give characteristics flame colouration ?
- 8. Arrange the following in order of increasing covalent character : MCl, MBr, MF, MI (where M = Alkali metal) [Ans. MF < MCl < MBr < MI]
- 9. Alkali metals can not be obtained by chemical reduction method. Explain.
- **10.** Why is sodium metal kept under kerosene oil ?





4.

(c)

3.

2.

- **11.** Why Be and Mg do not give characteristics colour to the flame ?
- **12.** Arrange the alkaline earth metal carbonate in the decreasing order of thermal stability.
- **13.** Why do alkaline earth metals not form any superoxide ?
- 14. Why gypsum is added to cement?
- **15.** How plaster of paris is obtained from gypsum ?
- **16.** BeO is insoluble in water but BeSO₄ is soluble in water ? Why ?
- 17. Why second I.E. of group II elements is less than group I elements?
- **18.** What is quick lime ? How is it prepared ?
- **19.** Why does Be show similarities with Al?
- **20.** Name the alkaline earth metal hydroxide which is amphoteric.

2-MARKS QUESTIONS

- 1. Why are alkali metals soft and have low melting points ?
- 2. Write any four similarities between Li and Mg.
- **3.** Why are potassium and caesium rather than Lithium used in photoelectric cells ?
- **4.** Why is Li₂CO₃ decomposed at a lower temperature whereas Na₂CO₃ at higher temperature ?
- 5. Among the alkali metals which has :
 - (i) Highest melting point.
 - (ii) Most electropositive character
 - (iii) Lowest size of ion.
 - (v) Strongest reducing character. [Ans. (i) Li (ii) Cs (iii) Li (iv) Li]
- **6.** Why does the solubility of alkali earth metal carbonates and sulphates decrease down the group ?
- 7. Draw the structure of BeCl₂ in (i) Vapour phase (ii) Solid state.
- 8. When CO₂ gas is passed in lime water it turns milky but in case of excess CO₂ milkiness disappears. Support the statement by giving suitable reaction equations.
- 9. (i) E^{θ} for M^{2+} (aq) + 2 $e^{-} \longrightarrow M(s)$ (where M = Ca, Sr, Ba) is nearly constant.





(ii) What is dead burnt plaster ? How is it obtained from gypsum?

10. Write two important uses of (i) Limestone (ii) Quick lime.

3-MARKS QUESTIONS

- **1.** Assign reason for the following :
 - (i) Compounds of lithium are generally covalent.
 - (ii) Alkali metals are strong reducing agent.
 - (iii) LiCl is more covalent than NaCl.
- 2. Discuss the various reactions that occur in Solvay process.
- **3.** Explain why ?
 - (i) Lithium salts are commonly hydrated.
 - (ii) Sodium peroxide is widely used as oxidising agent.
 - (iii) Sodium wire is used to remove moisture from benzene but can't be used for drying alcohol.
- **4.** Sodium hydroxide is generally prepared by electrolysis of brine solution in the Castner-Kellner cell :
 - (i) Write the reactions that occur in the cell.
 - (ii) Write any two uses of NaOH.
- 5. Explain with suitable reasons :
 - (a) A solution of Na_2CO_3 is alkaline.
 - (b) Alkali metals are prepared by electrolysis of their fused chlorides.
 - (c) Sodium is found to be more useful than potassium ?
- 6. Arrange the following in order of property mentioned against each :
 - (i) BaCl₂, MgCl₂, BeCl₂, CaCl₂ (Increasing ionic character)
 - (ii) Mg(OH)₂, Sr(OH)₂, Ba(OH)₂, Ca(OH)₂ (Increasing solubility in water)
 - (iii) BeO, MgO, BaO, CaO (Increasing basic strength)
- 7. What happens when :
 - (i) Mg is burnt in air.
 - (ii) Quick lime is heated with silica.
 - (iii) Chlorine is heated with slaked lime.
- 8. Write the raw material required for the manufacture of portland cement ? Why gypsum is added into it ?
- 9. (i) Why alkaline earth metals cannot be obtained by reduction of their oxides?



s-Block Elements



- (ii) Why the elements of group 2 are known as alkaline earth metals?
- 10. (i) Alkaline earth metals forms ionic salt having bivalent cations. Explain. Why?
 - (ii) A piece of magnesium ribbon continues to burn in SO₂. Why ?

5-MARKS QUESTIONS

- 1. Explain the following observation :
 - (a) LiI is more soluble than KI in ethanol.
 - (b) Sodium reacts with water less vigorously than potassium.
 - (c) LiF is insoluble in water.
 - (d) The mobilities of the alkali metal ions in aqueous solution are $Li^+ < Na^+ < K^+ < Rb^+ < Cs^+$.
 - (e) Lithium is the only alkali metal to form a nitride directly.
- 2. Complete the following reaction equations :
 - (i) $BeCl_2 + LiAlH_4 \longrightarrow$
 - (ii) $CaO + SiO_2 \longrightarrow$
 - (iii) $Ca(OH)_2 + Cl_2 \longrightarrow$
 - (iv) CaO + $P_4O_{10} \longrightarrow$
 - (v) $Ca(OH)_2 + CO_2 \longrightarrow$
- **3.** Compare the solubility and thermal stability of the following :

Compounds of the alkali metals with those of alkaline earth metals (a) nitrates (b) carbonates (c) sulphates.

- **4.** Explain the significance of Sodium (Na), Potassium (K), Magnesium (Mg) and Calcium(Ca) in biological fluids.
- **5.** Explain the significance of Sodium Potassium, Magnesium and Calcium biological fluids.
- **6.** (i) A solutions of Na_2CO_3 is alkaline why?
 - (ii) BeO insoluble but BeSO₄ in soluble in water. Why?
 - (iii) Lithium salts are commonly hydrated and those of other alkali metal ions are usually anhydrous give reasons.
 - (iv) What is the importance of cement?
 - (v) What happen when quick lime is heated with silica?





UNIT TEST

e Allowed: 1 hr Maximum M	Iarks : 20
All questions are compulsory.	
Photoelectric effect is maximum in	[1]
(a) Cs (b) K (c) Na (d) Li	
Flame test is not shown by	[1]
(a) Be (b) Sr (c) K (d) Ca	
Oxidation state of K in KO ₂ is	[1]
LiCl is soluble in organic solvent. Why?	[1]
Why is sodium kept under kerosene?	[1]
Write only four similarities in properties of Li and Mg.	[2]
Write two important uses of CaO and CaCO ₃ .	[2]
Discuss the various reactions that occur in solvay process.	[3]
What happens when :	[3]
(a) Mg is burnt in air.	
(b) Quick lime is heated with silica.	
(c) Chlorine is heated with slaked lime.	
Complete the following chemical reactions :	[5]
(a) $CaO + SiO_2 \longrightarrow$	
(b) $Ca(OH)_2 + CO_2 \longrightarrow$	
(d) $3Mg + N_2 \xrightarrow{\Delta}$	
(e) $NH_4HCO_3 + NaCl \longrightarrow$	
	(a) Cs (b) K (c) Na (d) Li Flame test is not shown by (a) Be (b) Sr (c) K (d) Ca Oxidation state of K in KO ₂ is LiCl is soluble in organic solvent. Why? Why is sodium kept under kerosene? Write only four similarities in properties of Li and Mg. Write two important uses of CaO and CaCO ₃ . Discuss the various reactions that occur in solvay process. What happens when : (a) Mg is burnt in air. (b) Quick lime is heated with silica. (c) Chlorine is heated with silica. (c) Chlorine is heated with slaked lime. Complete the following chemical reactions : (a) CaO + SiO ₂ \longrightarrow (b) Ca(OH) ₂ + CO ₂ \longrightarrow (c) 2Mg(NO ₃) ₂ \longrightarrow (d) $3Mg + N_2 \xrightarrow{\Delta}$







p-Block Elements

FAST TRACK : QUICK REVISION

General outer Electronic configuration : ns^2np^{1-6} . Inert Pair Effect:

- Reluctance of *ns*² electrons of valence shell to participate in bond formation is termed as inert pair effect.
- It arises due to poor or insufficient shielding of *ns*² electrons by intervening d- or f-electrons & hence increases down the group.

Causes of Anomalous Behaviour of First Element in groups of p-Block:

- (i) Very small size
- (ii) Unavailability of vacant d-orbital
- (iii) Tendency to form $p_{\pi} p_{\pi}$ multiple bonds.

Group No-13 Elements: (B, Al, Ga, In, Tl, Nh)

- General Electronic Configuration: ns² np¹
- Atomic radius: B < Ga < Al < In <TI

 $r_{Ga}^{} < r_{Al}^{}$ due to ineffective shielding of valence electrons by intervening 3d-electrons in Ga.

- **Ionization Enthalpies:** B > Tl > Ga > Al > In
- **Electronegativity:** B > Tl > In > Ga > Al
- Oxidation States: B (+3), Al (+3), Ga (+3, +1), In (+3, +1), Tl (+1, +3)

Tl (+1) is more stable than Tl (+3) due to inert pair effect.

• Nature of Compounds: Compounds of group 13 elements are electron deficient i.e. Lewis Acid and hence used as industrial catalyst *e.g.* BF₃, AlCl₃.

•	Oxides:	B_2O_3	Al_2O_3, Ga_2O_3	ln_2O_3	Tl ₂ O
		Acidic	Amphoteric	Basic	Strongly Basic

- Halides: MX₃ type, Electron deficient (Lewis acid), AICI₃ exist as dimer
- **Borax:** $Na_2B_4O_7.10H_2O$. On heating it form transparent glassy bead consisting of $NaBO_2 + B_2O_3$.
- **Boric acid:** H₃BO₃, It acts as a Lewis acid by accepting electron pair from OH⁻ ions of water.
- **Diborane:** B₂H₆, Colourless & toxic gas, acts as Lewis acid due to having electron deficient 3c-2e⁻ bonds. Obtained by treating BF₃ with LiAIH₄ or NaH, Also obtained by treating NaBH₄ with l₂.
- **Borazine:** $B_3N_3H_6$, It is isostructural with benzene and hence known as inorganic benzene. Prepared by heating B_2H_6 with NH₃

Group -14 Elements: (C, Si, Ge, Sn, Pb,Fl)

- General Electronic Configuration: ns² np²
- Atomic radius: C < Si < Ge < Sn < Pb
- **lonisation Enthalpy:** LiH_r : C > Si > Ge > Sn < Pb
- Oxidation States: C (+4), Si (+4), Ge (+4, +2), Sn (+4, +2), Pb (+4, +2) Pb (+2) is more stable than Pb (+4) due to inert pair effect.
- Oxides: Form di oxides (MO₂) & mono oxides (MO).
 PbO₂ is powerfull oxidizing agent because Pb stabilizes in +2 oxidation state due to inert pair effect. CO₂ is gas while SiO₂ is network solid because C has ability to form p_π p_π multiple bonds.
- **Halides:** Form tetra halides (MX_4) & dihalides (MX_2) .

Tetra halides are more covalent due to greater polarizing power of cation.

 CCI_4 is not hydrolysed with water as C has no vacant d-orbital to accept electron pair from water.

- Catenation: $C >> Si > Ge \approx Sn >> Pb$
- Allotrops of carbon: Diamond (sp³), Graphite (sp²), Fullerenes (sp²)
- Silicones: Silicones are synthetic organosilicon compounds containing R₂SiO repeating units. Silicones are water repellent, heat resistant, chemically inert, electrical insulators, resistant to oxidation.
- Silicates: Silicates are compounds in which anions are derived from Si-o-si- tetrahedral units.
- Zeolites: Zeolites are 3D silicates in which some of the Si atoms are replaced by Al³⁺ ions and negative charge is balanced by cations such as Na⁺, K⁺, Ca²⁺ etc.
- ZSM-5 is used in petrochemical industries to convert methanol into petrol.





MULTIPLE CHOICE QUESTIONS (MCQ)

- 1. The element which exists in liquid state for a wide range of temperature and can be used for measuring high temperature is
 - (i) B (ii) Al
 - (iii) Ca (iv) Ga
- 2. Which of the following is Lewis acid?
 - (i) AlCl₃ (ii) MgCl₂
 - (iii) CaCl₂ (iv) BaCl₂
- 3. The geometry of a complex species can be understood from the knowledge of type of hybridisation of orbitals of central atom. The hybridisation of orbitals of central atom in $[Be(OH_4]^-$ and the geometry of the complex are respectively
 - (i) sp^3 , tetrahedral (ii) sp^3 , square planar
 - (iii) sp^3d^2 , octahedral (iv) dsp^2 , square planar
- 4. Which of the following oxides is acidic in nature?
 - (i) B_2O_3 (ii) Al_2O_3
 - (iii) Ga_2O_3 (iv) In_2O_3
- 5. The exhibition of highest co-ordination number depends on the avilability of vacant orbitals in the central atom. Which of the following elements is not likely to act as central atom in $MF_6^{-3-?}$
 - (i) B (ii) Al
 - (iii) Ga (iv) In
- 6. Boric acid is an acid because its molecule
 - (i) Contains replaceable H^+ ion
 - (ii) Gives up a proton
 - (iii) Accepts OH^- from water releasing proton
 - (iv) Combines with proton from water molecule
- 7. Catenation i.e., linking of similar atoms depends on size and electronic configuration of atoms. The tendency of catenation in Group 14 elements follows the order:
 - (i) C > Si > Ge > Sn (ii) C >> Si > Ge = Sn
 - (iii) Si > C > Sn > Ge (iv) Ge > Sn > Si > c





- 8. Silicon has a strong tendency to form polymers like silicones. The chain length of silicone plymer can be controlled by adding
 - (i) MeSiCl₃ (ii) Me₂SiCl₂
 - (iii) Me₃SiCl (iv) Me₄Si
- 9. Ionisation enthalpy $(\Delta_1 H_1 \text{ kJ mol}^{-1})$ for the elements of Group 13 follows the order
 - $(i) \qquad B > Al > Ga > In > T1 \qquad (ii) \qquad B < Al < Ga < In < T1$
 - (iii) B < Al > Ga < T1 (iv) B > Al < Ga > In < T1

10. In the structure of diborane

- (i) All hydrogen atoms lie in one plane and boron atoms lie in a plane perpendicular to this plane.
- (ii) 2 boron atoms and 4 terminal hydrogen atoms lie in the same plane and 2 bridging hydrogen atoms lie in the perpendicular plane.
- (iii) 4 bridging hydrogen atoms and boron atoms lie in one plane and two terminal hydrogen atoms lie in a plane perpendicular to this plane.
- (iv) All the atoms are in the same plane.
- 11. A compound X, of boron reacts with NH_3 on heating to give another compound Y which is called inorganic benzene. The compound X can be prepared by treating BF_3 with Lithium aluminum hydride. The compounds X and Y are represented by the formulas.

(i)	$\mathrm{B_2H_6}, \mathrm{B_3N_3H_6}$	(ii)	$B_2O_3, B_3N_3H_6$
(iii)	BF ₃ , B ₃ N ₃ H ₆	(iv)	$B_{3}N_{3}H_{6}, B_{2}H_{6}$

12. Quartz is extensively used as a piezoelectric material, it contains

- (i) Pb (ii) Si
- (iii) Ti (iv) Sn
- 13. The most commonly used reducing agents is
 - (i) AlCl₃ (ii) PbCl₂
 - (iii) SnCl₄ (iv) SnCl₂
- 14. Dry ice is
 - (i) Solid NH_3 (ii) Solid SO_2
 - (iii) Solid CO_2 (iv) Solid N_2



p-Block Elements

- 15. Cement, the important building material is a mixture of oxides of several elements. Besides calcium, iron and sulphur, oxides of elements of which of the group(s) are present in the mixture?
 - (i) Group 2 (ii) Groups 2, 13 and 14
 - (iii) Groups 2 and 13 (iv) Groups 2 and 14

MATCH THE COLUMNS

In the following questions more than one correlation is possible between options of Column I and Column II. Make as many correlation as you can

1.	Column I			Column II			
	(i)	BF ₄	(a)	Oxidation state of central atom is +4			
	(ii)	AlCl ₃	(b)	Strong oxidishing agent			
	(iii)	SnO	(c)	Lewis acid			
	(iv)	PbO ₂	(d)	Can be further oxidised			
			(e)	Tetrahedral shape			
2.	(Column I		Column II			
	(i)	Diborane	(a)	Used as a flux for soldering metals			
	(ii)	Galluim	(b)	Crystalline form of silica			
	(iii)	Borax	(c)	Banana bonds			
	(iv)	Aluminosilicate	(d)	Low melting, high boiling, useful for measuring high temperature			
	(v)	Quartz	(e)	Used as catalyst in petrochemical industries			
3.	(Column I		Column II			
	(i)	Boron in $[B(OH)_4]^-$		(a) Sp^2			
	(ii)	Aluminium in [Al(H	₂ O) ₆	$]^{3+}$ (b) Sp^{3}			
	(iii)	Boron in B_2H_6		(c) Sp^3d^2			
	(iv)	Carbon in Buckmins	terfu	llerene			
	(v)	Silicon in SiO ₄ ^{4–}					
	(vi)	Germanium in [GeC	1 ₆] ^{2–}				





ASSERTION-REASON TYPE QUESTIONS

In the following questions a statement of Assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given below each question

1.	Assertion (A)	:	If aluminium atoms replace a few silicon atoms in
			three dimensional network of silicon dioxide, the
			overall structure acquires a negative charge

Reason (R) : Aluminium is trivalent while silicon is tetravalent.

- (i) Both A and R are correct and R is the correct explanation of A.
- (ii) Both A and R are correct but R is not the correct explanation of A.
- (iii) Both A and R are not correct
- (iv) A is not correct but R is correct.
- 2. Assertion (A) : Silicon is water repelling in nature.

Reason (R) : Silicon is organosilicon polymers, which have (-R_2SiO-) as repeating unit

- (i) A and R both are correct and R is the correct explanation of A.
- (ii) Both A and R are correct but R is not the correct explanation of A
- (iii) A and R both are not true.
- (iv) A is not true but R is true.

1-MARK QUESTIONS

- 1. Mention two important ores of boron.
- 2. Name the elements of group 13 which forms only covalent compounds.
- 3. Why the atomic radius of gallium is less than that of Al ?
- 4. Why does Boron form electron deficient compounds ?
- **5.** Boron does not exist as B^{3+} ion. Why ?
- 6. Why the trihalide of group 13 elements fume in moist air ?
- 7. Aluminum form $[AIF_6]^{3-}$ but boron does not form $[BF_6]^{3-}$.
- 8. Why boric acid is a monobasic acid ?
- 9. White fumes appear around the bottle of anhydrous AlCl₃. Give reason.
- **10.** AlCl₃ exist as dimer while BCl₃ exist as monomer, why ?



p-Block Elements



- **11.** Mention the type of hybridization of Boron in B_2H_6 .
- **12.** Write the formula of inorganic benzene.
- 13. Why aluminum utensils should not be kept in water overnight.

[Ans. sp^3]

- **14.** Explain what happens when boric acid is heated.
- 15. BCl₃ exists but BH₃ does not. Explain.
- **16.** Why SnCl₄ is more covalent than SnCl₂?
- **17.** Why PbCl₄ is good oxidising agent ?
- **18.** What are germanes and plumbanes ?
- **19.** Give one example of zeolite.
- **20.** Mention the type of hybridization of carbon in diamond and graphite.
- **21.** Why CCl_4 is insoluble in water but $SiCl_4$ is soluble in water? Explain.
- 22. Give two uses of silicones.
- 23. Why graphite is used as lubricant?
- **24.** Lead (Pb) do not form PbI₄. Why ?
- **25.** CO_2 is gas while SiO₂ is solid at room temperature. Explain why ?
- **26.** Explain why silicon shows a higher covalency than carbon?
- 27. Out of carbon and silicon which can form multiple bonds and why ?
- **28.** Write the formula of dry ice.
- **29.** Mention the basic building unit of all silicates.
- **30.** Graphite is a good conductor of electricity, but diamond is not. Why ?

2-MARKS QUESTIONS

- 1. Draw the structure of diborane.
- 2. What happens when :
 - (a) Borax is heated strongly.
 - (b) Boric acid is added to water.
- **3.** Write balanced chemical equations for :
 - (a) $BF_3 + LiH \longrightarrow$

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(b) $B_2H_6 + NH_3 \longrightarrow$



- 4. Write chemical reactions to justify amphoteric nature of Al.
- 5. Suggest reason why the B-F bond length in BF_3 and BF_4^- differ.
- **6.** Give reason for the following :
 - (i) BF_3 act as weak Lewis acid.
 - (ii) Boron cannot show covalency more than four.
- 7. How can you explain higher stability of BCl₃ as compared to TlCl₃?
- 8. Give reason for the following :
 - (i) Aluminium alloys are used to make aircraft body.
 - (ii) Aluminium wire is used to make transmission cables.
- 9. Describe the shapes of BF_3 and BH_4^- . Assign the hybridization of boron in these species.
- **10.** Explain the chemistry of borax bead test.
- **11.** $[SiF_6]^{2-}$ is known whereas $[SiCl_6]^{2-}$ not. Give reason.
- **12.** Hydrolysis of $SiCl_4$ take place but of CCl_4 does not. Why ?
- **13.** Account for the following :
 - (a) CO_2 is gas while SiO₂ is solid at room temperature.
 - (b) Solid CO₂ is known as dry ice.
- Elemental silicon does not form graphite like structure as carbon does. Give reason.
- 15. Suggest a reason as to why CO is poisonous?
- **16.** How is excessive content of CO₂ responsible for global warming ?
- 17. What is allotropy ? Name two elements which exhibit allotropy.
- **18.** Write equations for the production of water gas and producer gas from coke.
- **19.** Define zeolite. Name the zeolite which converts alcohols directly into gasoline.
- 20. Arrange the hybrides of group 14 elements in increasing order of :
 - (a) Thermal stability
 - (b) Reducing power.





3-MARKS QUESTIONS

1. Give reasons of the following :

(i) In diborane, two B - H - B bonds are different from common covalent bonds.

- (ii) Aluminium metal shows amphoteric behaviour.
- (iii) Quartz is used to develop extremely accurate clocks.
- 2. A certain salt X gives the following results :
 - (i) Its aqueous solution is alkaline to litmus.
 - (ii) It swells up to a glassy material Y on strong heating.
 - (iii) When conc. H_2SO_4 is added to a hot solution of X, white crystal of an acid Z separates out. Write equations for all the above reactions and identify X, Y and Z.
- **3.** Write balanced chemical equation for :
 - (i) $B_2H_6 + H_2O \longrightarrow$
 - (ii) Al + NaOH \longrightarrow
 - (iii) NaOH + $B_2H_6 \longrightarrow$
- **4.** List two important properties in which boron differs from the rest of the members of group. Mention the main reasons for the difference.
- 5. What are electron deficient compounds? Are BCl₃ and SiCl₄ electron deficient species ? Explain.
- 6. Select the member(s) of group 14 that :
 - (i) Forms the most acidic dioxide.
 - (ii) Is commonly found in + 2 oxidation state.
 - (iii) Used as semiconductor.
- 7. What are allotropes ? Sketch the structure of two allotropes of carbon namely diamond and graphite.
- 8. Give suitable reasons for the following :
 - (a) CO_2 turns lime water milky, but if passed for a long time, the solution become transparent again.
 - (b) Graphite is a good conductor of electricity but diamond is insulator.
 - (c) Lead (IV) chloride is highly unstable towards heat.





- 9. (i) Write the resonance structure of CO_3^{2-} and HCO_3^{-} .
 - (ii) Write the name of thermodynamically most stable form of carbon.
- **10.** (i) Explain why is there a phenomenal decreases in ionisation enthalpy from carbon to silicon ?
 - (ii) Write an industrial application of silicones.

5-MARKS QUESTIONS

- When metal X is treated with NaOH, a white precipitate 'A' is obtained, which is soluble in excess of NaOH to give soluble complex (B). Compound 'A' is soluble in dilute HCl to form compound 'C'. The compound 'A' when heated strongly gives 'D', which is used to extract metal. Identify X, A, B, C and D. Write suitable equations to support their identities.
- 2. (i) If B-Cl bond has dipole moment explain why BCl₃ molecules has zero dipole moment.
 - (ii) A mixture of dil. NaOH and aluminium pieces is used to open drain. Give reason.
 - (iii) Aluminium wire is used to make transmission cables. Why ?
- **3.** (i) Identify the compounds X and Y in the following reactions :
 - (a) $Na_2B_4O_7 + 2HCl + 5H_2O \rightarrow 2NaCl + X$
 - (b) $X \xrightarrow{370 \text{ K}} \text{HBO}_2 \xrightarrow{> 370 \text{ K}} Y.$
 - (ii) Write the name of group 13 element which is used to measure high temperature.
 - (iii) Why in case of thallium + 1 oxidation state is more stable than + 3?
- **4.** Compare the general trend in the following properties of the elements of group 13 and 14 :
 - (a) Atomic size, (b) Ionisation enthalpy, (c) Metallic character,
 - (d) Oxidation states, (e) Nature of halides.





- 5. Name the following :
 - (a) The crystalline form of silica used in modern radio and T.V. broadcasting and mobile radio communication.
 - (b) The oxides of carbon which form a complex with haemoglobin 300 times more faster than oxygen.
 - (c) The allotrope of carbon which has $\Delta_f H^{\theta} = 0$.
 - (d) A type of polymer is semiorganic in nature.
 - (e) Two man made silicates.
- 6. Explain the formation of (i) water gas (ii) producer gas. Give their uses. What happens when CO₂ is passed through limewater?
 - (i) for short duration
 - (ii) for long duration.



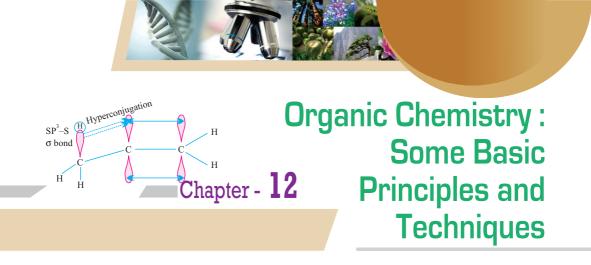


UNIT TEST

Gen (i)	ne Allowed: 1 hrMaximum Markeral Instructions:All questions are compulsory.Maximum marks carried by each question are indicated against it.	ks : 20					
1.	Which of the following is Lewis acid?	[1]					
	(a) $AlCl_3$ (b) $MgCl_2$ (c) $CaCl_2$ (d) $BaCl_2$						
2.	Dry ice is	[1]					
	(a) Solid CO_2 (b) Solid SO_2 (c) Solid N_2 (d) Solid NH_3						
3.	Chemical formula of diborane is	[1]					
4.	Write an example of shape relative catalyst.	[1]					
5.	Allotrope of carbon with sp ³ hybridisation state is?	[1]					
6.	Complete the following chemical equations :	[2]					
	(a) $B_2H_6 + NH_3 \longrightarrow$ (b) $BF_3 + LiH \longrightarrow$						
7.	Write the chemical reactions involved in borax bead test.	[2]					
8.	What are allotrope? Sketch the structure of two allotrope of carbon namely diamond and graphite.						
9.	Compare the properties of the elements of group-13 and 14	[3]					
	(a) Atomic size						
	(b) Ionisation enthalpy						
	(c) Oxidation state						
10.	(a) Explain the formation of (i) water gas (ii) producer gas						
	(b) Identify the compound X and Y in the following reactions						
	(i) $Na_2B_4O_7 + 2HCl + 5H_2O \longrightarrow 2NaCl + X$						
	(ii) $X \xrightarrow{370 \text{ K}} \text{HBO}_2 \xrightarrow{>370 \text{ K}} Y$						
	(c) Write two important applications of silicons.						

p-Block Elements





FAST TRACK : QUICK REVISION

ORGANIC CHEMISTRY

It deals with the study of hydrocarbons (compounds of carbon and hydrogen elements) and their derivatives.

Some organic compounds may also contain nitrogen, oxygen, sulphur, phosphorus, halogens, etc.

Berzelius, proposed that a 'vital force' was responsible for the formation of organic compounds.

This was rejected by F. Wohler who synthesised first organic compound urea from an inorganic compound.

$$\begin{array}{ccc} \mathrm{NH}_4\mathrm{CNO} & \stackrel{\Delta}{\longrightarrow} & \mathrm{NH}_2\mathrm{CONH}_2\\ \mathrm{Ammonium\ cyanate} & & \mathrm{Urea} \end{array}$$

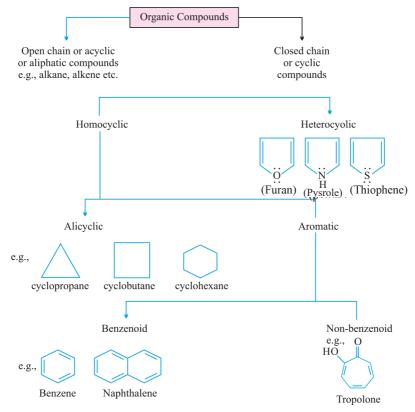
- Acetic acid was synthesised by Kolbe and methane by Berthelot.
- Types of hybridisation of C-atom :

Hybridisation	Structure	Bond angle	Examples
sp ³	Tetrahedral	109°28'	Ethane, Methane
sp ²	Trigonal	120°	Ethene, Propene
sp	Linear	180°	Ethyne, Propyne

- Reasons for existence of large number of organic compounds:
- **Catenation :** The property of atoms of an element to link with one another forming chains of identical atoms is called *catenation*. Carbon exhibits catenation to the maximum extent.
- **Isomerism :** It is the property by virtue of which two or more compounds have the same molecular formula but different physical or chemical properties.

• Formation of multiple bonds : Because of its small size carbon atom is capable of forming multiple bonds with other atoms and this gives a variety of compounds.





• CLASSIFICATION OF CARBON ATOMS

On the basis of number of C attached

- (i) **Primary carbon atom :** when carbon atom is attached with one other carbon atom only, it is called **primary or 1°** carbon atom.
- (ii) Secondary carbon atom : When carbon atom is attached with two other carbon atoms, it is called **secondary or 2°** carbon atom.
- (iii) **Tertiary carbon atom :** When carbon atom is attached with three other carbon atoms, it is called **tertiary or 3°** carbon atom.
- (iv) **Quaternary carbon atom :** When carbon atom is attached with four other carbon atoms, it is called **quarterly or 4°** carbon atom.
- Functional Group : The atom *e.g.*, -Cl, -Br, etc., or group of atoms *e.g.*, -COOH, -CHO, which is responsible for the chemical properties of the molecule, is called functional group.

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Organic Chemistry: Some Basic Principles and Techniques

• Homologous Series : The series in which the molecular formula of adjacent members differ by a – CH₂ unit, is called homologous series and the individual members are called homologous, *e.g.*, The homologous series of alkene group is

$$\begin{array}{c} C_{2}H_{4} \\ C_{3}H_{6} \\ C_{4}H_{8} \\ C_{5}H_{10} \end{array} \right\} difference of -CH_{2} unit or 14 unit mass$$

The general characteristics of this series are :

1. All the homologues contain same functional group. That's why their chemical properties are almost similar.

Series	General Formula
Alkanes	C _n H _{2n+2}
Alkenes	C _n H _{2n}
Alkynes	C _n H _{2n-2}
Alcohol and ether	$C_nH_{2n+2}O$
Aldehyde and ketone	C _n H _{2n} O
Acid and ester	$C_nH_{2n}O_2$

2. All the members of a series have same general formula, *e.g.*,

- 3. All the members can be prepared by almost similar methods.
- 4. With increase in the molecular weight of a series, the physical properties vary gradually.

• Representation of Organic Compounds :

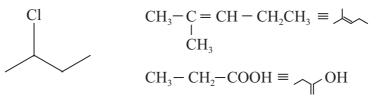
Organic compounds can be represented by the following ways:

(i) **Complete Structural Formula :** All the bonds present between any two atoms are shown clearly. *e.g.*,





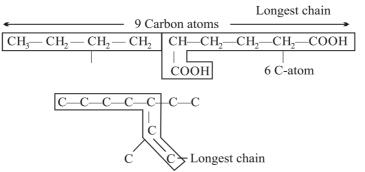
- (ii) Condensed Formula : All the bonds are not shown clearly. *e.g.*, CH₃CHCH₂CH₃
 Cl
 CH₃CH(CI)CH₂CH₃
- (iii) **Bond Line Formula :** Every fold and free terminal represents a carbon and lines represent the bond. *e.g.*,



• **IUPAC Nomenclature of Organic Compounds :** Following rules are used to write the IUPAC name of an organic compound.

Rule 1. : Longest chain rule : The chain containing the principal functional group, secondary functional group and multiple bonds as many as possible is the longest possible chain.

In the absence of functional group, secondary group and multiple bonds, the chain containing the maximum number of C-atoms will be the longest possible chain *e.g.*,



Word Root for Carbon Chain

Chain length	Word root	Chain length	Word root
C ₁	Meth-	C ₇	Hept
C ₂	Eth-	C ₈	Oct
C ₃	Prop-	C ₉	Non
C ₄	But-	C ₁₀	Dec
C ₅	Pent-	C ₁₁	Undec
C ₆	Hex-	C ₁₂	Dodec

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Rule 2 : Lowest number rule : Numbering is done in such a way so that

- (i) branching if present gets the lowest number.
- (ii) the sum of numbers of side chain is lowest.
- (iii) principal functional group gets the lowest number.

Select the principal functional group from the preference series :

Functional group other than the principal functional group are called substituents.

Rule 3: Naming the prefix and suffixes : Prefix represents the substituent and suffix is used for principal functional group.

Primary suffix are **ene**, **ane** or **yne** used for double, single and triple bonds respectively.

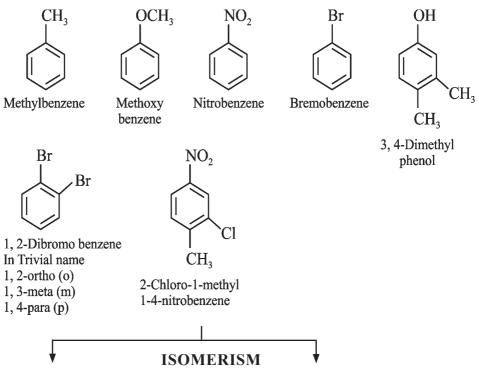
No.	Class	Formula	Prefix	Suffix
1.	Acid halides		halocarbonyl	—oyl halide —carbonyl halide
2.	Alcohols	—OH	hydroxy	—ol
3.	Aldehydes	—CHO	formyl	—al
				—carbaldehyde
4.	Ketones	ightarrow C = O	oxo (keto)	—one
5.	Amides	-CONH ₂	carbamoyl	—amide
6.	Amine	-NH ₂	amino	—amine
7.	Carboxylic acid	—СООН	carboxy	—carboxylic acid
8.	Ester	—COOR	alkoxy carbonyl	—alkyl alkan oate
9.	Nitriles	—CN	cyano	—nitrile
10.	Sulphonic acid	—SO ₂ —OH	sulpho	—sulphonic acid

Secondary suffixes are tabulated below :





• Nomenclature of substituted benzene compounds :



Structural isomerism

Same molecular formula but different structures

Types

1. Chain Isomerism

- *e.g.*, Pentane and 2-Methylbutane
- 2. Position Isomerism *e.g.*, But-1-ene
 - But-2-ene

3. Functional Isomerism *e.g.*, Propanal & Propanone

Ethanol & Methoxymethane

4. **Metamerism** *e.g.*, Pentan-2-one and Pentan-3-one

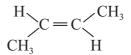
Stereo isomerism

Same molecular and structural formula but different configuration

e.g., Geometrical isomerism (cis-trans isomerism)

$$\underset{H}{\overset{H_3C}{\longrightarrow}} C = C \underset{H}{\overset{CH_3}{\swarrow}}$$

Cis.But-2-ene



Trans.But-2-ene

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• Fission of a Covalent Bond :

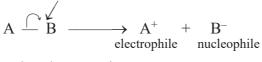
(i) **Homolytic Fission :** In this one of the electrons of the shared pair in a covalent bond goes with each of the bonded atoms. The neutral chemical species thus formed, are called free radicals. Generally, homolytic fission takes place in non-polar covalent molecules in the presence of sunlight or high temperature.

$$A \xrightarrow{\frown} B \xrightarrow{\text{Sunlight}} \underbrace{A^* + B^*}_{\text{free radicals}}$$
$$Cl_2 \xrightarrow{\text{Sunlight}} 2Cl^*$$

Free radicals are highly reactive, neutral and electron deficient species.

(ii) Heterolytic Fission : The covalent bond breaks in such a fashion that the shared pair of electrons goes with one of the fragments.

more electronegative



less electronegative

$$A \longrightarrow A^- + B^+$$

nucleophile electrophile

Heterolytic fission generally takes place in polar covalent molecules but in non-polar molecules, it takes place in the presence of catalyst like AiCl₃ (anhy.), FeCl₃ (anhy.) etc.

• Attacking Reagents :

These are of two types

(i) Electrophiles or Electrophilic Reagents

These are electron deficient species, i.e., behave as Lewis acids.

e.g., Cl^+ , NO_2 , CH_3CO^+ etc.

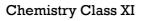
BF₃, ZnCl₂ (anhydrous), FeCl₃ (anhydrous), AlCl₃ (anhydrous)

(ii) Nucleophiles or Nucleophilic Reagents

These are negatively charged or neutral molecules with unshared electron pair.

e.g., $O\overline{H}$, CN^{-} , $R^{\circ\circ}NH_2$, $\overset{\circ\circ}NH_3$

- Reaction Intermediates :
 - (i) Free radicals : These are the product of homolysis and contain an odd electron. These are highly reactive planar species with sp² hybridisation.

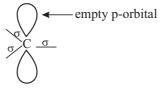




Their order of stability is

$$\begin{aligned} (C_{6}H_{5})_{3}\mathring{C} &> (C_{6}H_{5})_{2}\,\mathring{C}H > C_{6}H_{5}\mathring{C}H_{2} \\ &> CH_{2} = CH - \mathring{C}H_{2} > 3^{\circ} > 2^{\circ} > 1^{\circ} > CH_{2} = \mathring{C}H \end{aligned}$$

(ii) Carbocations : These are the product of heterolysis and contain a carbon bearing positive charge. These are electron deficient species. These are also polar chemical species i.e., sp² hybridised with an empty p-orbital.



Stability order of carbocation is

$$(C_6H_5)_3\overset{+}{C} > (C_6H_5)_2\overset{+}{C}H > C_6H_5\overset{+}{C}H_2$$

> $CH_2 = CH - \overset{+}{C}H_2 > 3^\circ > 2^\circ > 1^\circ > CH_2 = \overset{+}{C}H$

(iii) **Carbanions :** These are the product of heterolysis and contain a carbon bearing negative charge and 8 electrons in its valence shell.

These have pyramidal shape with sp³ hybridised carbon (having one lone pair) order of stability of carbanions is

$$(C_6H_5)_3\overline{C} > (C_6H_5)_2\overline{C}H > C_6H_5\overline{C}H_2$$

> $\overline{C}H_3 > 1^\circ > 2^\circ > 3^\circ$ carbanions

Electron Displacement in Covalent Bond

1. Inductive Effect : If shared pair is more shifted towards more electronegative atom, the less electronegative atom acquires slight positive charge and more electronegative atom acquires partial negative charge,

$$e.g., \quad \overset{+\delta}{\operatorname{CH}}_{3} \longrightarrow \overset{-\delta}{\operatorname{Cl}}$$

Permanent effect and propagates through carbon chain.

Atoms or groups having greater electron affinity than hydrogen are said to have electron attracting or negative inductive effect (-l) while that having, smaller electron affinity than hydrogen are said to have electron releasing or positive inductive effect (+l).

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e.g.,
$$\overset{+\delta\delta}{CH_3} \rightarrow \overset{+\delta}{CH_2} \rightarrow \overset{-\delta}{Cl}$$

 $\overset{+\delta\delta\delta}{CH_3} \rightarrow \overset{+\delta\delta}{CH_2} \rightarrow \overset{-\delta}{CH_2} \rightarrow \overset{-\delta}{Cl}$
 $\overset{1^{\circ} alkyl halide}$

Cl has -l effect and alkyl group has +I effect.

Order of groups producing –I effect is

$$\begin{split} R_{3}N > &- NO_{2} > - CN > - SO_{3}H > - CHO > - CO > - COOH > - F \\ &> - Cl > - Br > - I > - OH > - OR > - NH_{2} > - C_{6}H_{5} > - H \end{split}$$

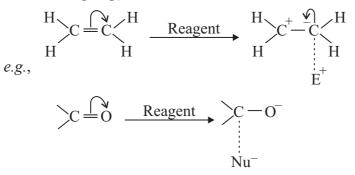
Order of groups producing +1 effect is

3° alkyl group > 2° alkyl group > 1° alkyl group > – $CH_3 > -H$

• Applications of Inductive Effect

- (i) Presence of groups showing +I effect increases the stability of carbocation while presence of groups showing –I effect decreases their stability.
- (ii) Strength of acid increases with the attachment of group showing –I effect and decreases with the attachment of group showing +I effect.
- (iii) Presence of +I showing groups increases the basic strength of amines.
- 2. Electromeric Effect : Defined as the polarity produced in a multiple bonded compound as a reagent approaches it. In the presence of attacking reagent, the two π electrons are completely transferred to any of the one atom. This effect is temporary.

It may be of +E type (when displacement of electron pair is away from the atom of group) or of -E type (when displacement is towards the atom or group).



3. Hyper-conjugation : It involves delocalisation of σ electron of a C – H bond of an alkyl group attached directly to an atom of unsaturated system or to an atom with an unshared p-orbital.

e.g.,
$$\stackrel{\mathrm{H}}{|}_{\mathrm{CH}_2} \xrightarrow{\mathrm{CH}_2} \stackrel{\mathrm{H}^+}{\longleftrightarrow}_{\mathrm{CH}_2} \operatorname{CH} - \bar{\mathrm{CH}}_2$$

This effect is also called no bond resonance or Baker Nathan effect.





Applications of Hyper-conjugation

Stability of alkenes : More the number of α -hydrogen atoms, more stable is the alkene.

$$H_{3}\overset{\alpha}{C} - \underbrace{C}_{H_{2}} \overset{\alpha}{C} - \underbrace{C}_{CH_{3}} \overset{\alpha}{C} + \underbrace{C}_{CH_{3}}$$

Stability of Carbocation : Greater the number of alkyl groups attached to positively charged carbon atom, the greater is the stability.

e.g.,
$$(CH_3)_2^+CH > CH_3 - CH_3^+CH > CH_3^+$$

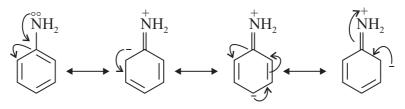
4. **Resonance Effect :** When the properties of a molecule cannot be shown by a single structure and two or more structures are required to show all the properties of that molecule, then the structures are called resonating structures or canonical forms and the molecule is referred as resonance hybrid. This phenomenon is called resonance.

Conditions for resonance

- (i) The arrangement of atoms must be identical in all the formula.
- (ii) The energy content of all the canonical forms must be nearly same.
- (iii) Each canonical of π electrons. This effect may be of +R type or -R type.

Positive Resonance Effect (+R)

Electron donating groups with respect to conjugated system show +R effect. Central atom of functional groups should be more electronegative than the surrounding atoms or groups to show +R effect. *e.g.*, halogens, –OH, –OR, –NH₂, NHCOR, etc.

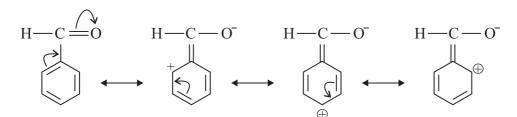


Negative Resonance Effect (-R)

Electron withdrawing groups with respect to conjugate system show –R effect. Central atom of functional groups should be less electronegative than surrounding atoms or groups to show –R effect. *e.g.*, halogens, –COOH, –COOR, CHO, –CN, –NO₂, etc.

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Organic Chemistry: Some Basic Principles and Techniques



Methods of Purification of Organic Compounds •

Method	Principle	Applications
Crystallization	Different solubilities of a given organic compound and its impurities in the same solvent.	 Crystallization of sugar (containing an impurity of common salt) is achieved by shaking the impure solid with hot ethanol at 348K (sugar dissolves whereas common salt remains insoluble).
Sublimation	Some solid substances change from solid to vapour state without passing through liquid state. Sublimable compounds get separated from non-sublimable impurities.	 Iodine from sodium chloride (as iodine sublimes readily leaving behind sodium chloride). Camphor, naphtalene, anthracence, benzoic Acid, etc. are purified.
Distillation	 It is used to separate Volatile liquids from non-volatile impurities. Liquids having sufficient difference in their boiling points. 	 Hexane (b.p. 342K) and toluene (b.p. 384K) Chloroform (b.p. 334K) and aniline (b.p. 457K)
– Fractional Distillation	If the difference in boiling points of two liquids is not much, this method is used.	• Crude oil in petroleum industry is separated into various useful fractions such as gasoline, kerosene oil, diesel oil, lubricating oil, etc.
– Steam Distillation	This method is used to separate substances which are (i) steam volatile, (ii) immiscible with water, (iii) posses a vapour pressure of 10-15 mm Hg and (iv) contain non- volatile impurities.	 Aniline is separated from aniline water mixture. Essential oils, turpentine oil, o-nitrophenol, bromobenzene nitrobenenze, etc. can be purified.
Differential Extraction	By shaking an aqueous solution of an organic compound with an organic solvent in which the organic compound is more soluble than in water. The organic solvent and the aqueous solution should be immiscible with each other so that they can form two distinct layers which can be separated by using separating funnel.	• Benzoic acid can be extracted from its water solution using benzene.



Chromatography	Differential movement of individual components of a mixture through a stationary phase under the influence of a mobile phase.	•	Widely used for separation purification, identification and characterization of the components of a mixture, whether coloured or colourless.
– Adsorption Chromatography	Differential adsorption of the various components of a mixture on a suitable adsorbent such as silica get or alumina.		
– Column Chromatography	The mixture is passed through adsorbent packed in glass tube.	•	Mixture of naphthalene and benzophenone.
– Thin Layer Chromatography	The mixture is passed over adsorbent on a thin glass plate.	•	Amino acids can be detected by spraying the plate with ninhydrin solution.
– Partition Chromatography	Differential partitioning of components of a mixture between stationary and mobile phases.		
– Paper Chromatography	A special quality paper known as chromatography paper is used. It contains water trapped in it, which acts as the stationary phase.	•	For separation of sugars and amino acids.

Types of Chromatography	Mobile / Stationary Phase
Column Chromatography	Liquid / Solid
Thin Layer Chromatography	Liquid / Solid
High Performance Liquid Chromatography (HPLC)	Liquid / Solid
Gas Liquid Chromatography (GLC)	Gas / Solid
Partition or Paper Chromatography	Liquid / Solid

Element	Detection	Confirmatory Test	Reactions
Carbon	$2CuO + C \xrightarrow{\Delta} 2Cu + CO_2$	CO ₂ gas turns lime water milky.	$\begin{array}{c} \text{CO}_2 + \text{Ca(OH)}_2 \longrightarrow \text{CaCO}_3 \downarrow + \text{H}_2\text{O} \\ \text{Lime water} \text{Milkiness} \end{array}$
Hydrogen	$CuO + 2H \xrightarrow{\Lambda} Cu + H_2O$	Water droplets appear on the cooler part of the ignition tube and also turns anhydrous CuSO ₄ blue.	$\begin{array}{c} \mathrm{CuSO}_4 + 5\mathrm{H}_2\mathrm{O} \longrightarrow \mathrm{CuSO}_4.5\mathrm{H}_2\mathrm{O} \\ \mathrm{White} & \mathrm{Blue} \end{array}$
Nitrogen	Lassaigne's extract (L.E.) Na + C + N $\xrightarrow{\Delta}$ NaCN (L.E.)	L.E. + FeSO ₄ + NaOH, boil and cool + FeCl ₃ + conc. HCl. Gives blue or green colour.	$\begin{array}{l} \operatorname{FeSO}_4 + 2\operatorname{NaOH} \longrightarrow \operatorname{Fe}(\operatorname{OH})_2 + \operatorname{Na}_2\operatorname{SO}_4 \\ \operatorname{Fe}(\operatorname{OH})_2 + 6\operatorname{NaCN} \longrightarrow \\ \operatorname{Na}_4[\operatorname{Fe}(\operatorname{CN})_6] + 2\operatorname{NaOH} \\ \operatorname{3Na}_4[\operatorname{Fe}(\operatorname{CN})_6] + 4\operatorname{FeCl}_3 \longrightarrow \\ \operatorname{Fe}_4[\operatorname{Fe}(\operatorname{CN})_6]_3 + 12\operatorname{NaCl} \\ \operatorname{Prussian blue} \end{array}$



Sulphur	$2Na + S \xrightarrow{\Delta} Na_2S$ (L.E.)	 L.E. + sodium nitroprusside A deep violet colour. L.E. + CH₃COOH + (CH₃COO)₂Pb Gives a black ppt. 	$\begin{array}{c} Na_{2}S + Na_{2}[Fe(CN)_{5}NO] \longrightarrow \\ Sodium nitroprusside \\ Na_{4}[Fe(CN)_{5}NOS] \\ Deep violet \\ Na_{2}S + (CH_{3}COO)_{2}Pb \xrightarrow{CH_{3}COOH} \\ Pbs \downarrow + 2CH_{3}COONa \\ Black ppt. \end{array}$
Halogens	$Na + X \xrightarrow{\Delta} NaX$ (L.E.)	 L.E. + HNO₃ + AgNO₃ - White ppt. soluble in aq. NH₃ (or NH₄OH) confirms Cl. - Yellow ppt. partially soluble in aq. NH₃ (or NH₄OH) confirms Br. - Yellow ppt. insoluble in aq. NH₃ (or NH₄OH) confirms I. 	$\begin{array}{c} NaX + AgNO_{3} \xrightarrow{HNO_{3}} AgX \downarrow \\ ppt. \\ AgCl + 2NH_{3(aq.)} \longrightarrow [Ag(NH_{3})_{2}]Cl \\ White \\ ppt. \\ \end{array}$
Nitrogen and sulphur together	$\begin{array}{c} Na + C + N + S \xrightarrow{\Delta} \\ NaSCN \\ Sodium thiocyanate \\ (L.E.) \end{array}$	As in test for nitrogen; instead of green or blue colour, blood red colouration confirms presence of N and S both.	$\begin{array}{c} \text{NaSCN} + \text{FeCl}_3 \longrightarrow \\ & [\text{Fe}(\text{SCN})\text{Cl}_2 + \text{NaCl} \\ & \text{Blood red colour} \end{array}$
Phospho- rus	$P \xrightarrow{Na_2O_2, \text{ boil}} Na_3PO_4$	Solution is boiled with nitric acid and then treated with ammonium molybdate $(N H_4)_2 M \circ O_4$. Formation of yellow ppt. indicates presence of phosphate (hence, phosphorus) in organic compound.	$\begin{split} Na_{3}PO_{4}+3HNO \longrightarrow H_{3}PO_{4}+3NaNo_{3} \\ H_{3}PO_{4}+12(NH_{4})_{2}MoO_{4}+21HNO_{3} \\ \longrightarrow (NH_{4})_{3}PO_{4}.12MoO_{3} \\ Ammonium phosphomolybalate \\ (yellow ppt.) \\ +21NH_{4}NO_{3}+12H_{2}O \end{split}$

• **Quantitative analysis of organic compounds :** The percentage composition of elements presence an organic compound is determined by the methods based on the following principles :





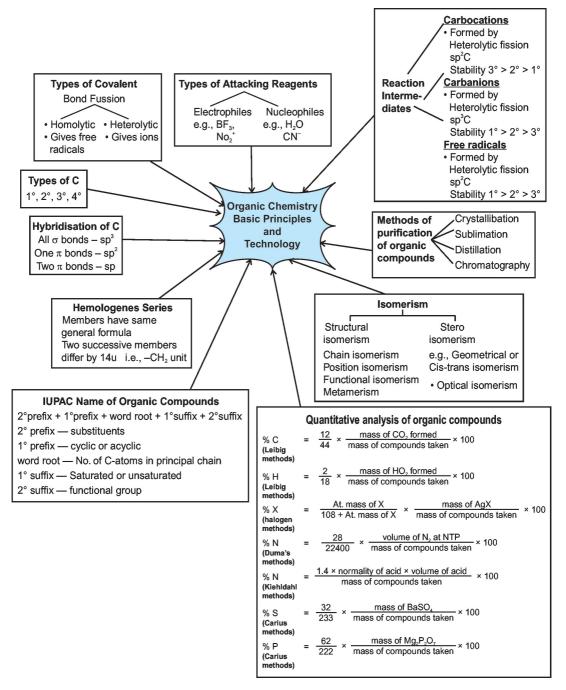
Elements	ents Method		
Carbon and Hydrogen	Liebig's Combustion method : A known mass of an organic compound is burnt in the presence of excess of O ₂ and CuO. $C_xH_v + (x + \frac{y}{4})O_2 \xrightarrow{\Lambda} xCO_2 + \frac{y}{2}H_2O$		
	$CO_{2} \text{ evolved is absorbed by conc. solution of KOH or ascarite (NaOH + CaO).H_{2}O produced is absorbed by anhydrous CaCl2 or Mg(ClO4)2.Increase in masses of these absorbing compounds gives the masses of CO2 and H2O produced.% of C = \frac{12}{44} \times \frac{\text{mass of CO}_{2} \text{ formed}}{\text{mass of compound taken}} 100;$		
	% of H = $\frac{2}{18} \times \frac{\text{mass of H}_2\text{O formed}}{\text{mass of compound taken}} 100$		
Halogens	Carius method :Halogen in organic compound is precipitated as silver halide by boilingwith conc. NHO3 and then adding AgNO3. $X \frac{HNO3, \Delta}{AgNO3} AgX \downarrow$ % of $X = \frac{At. mass of X}{108 + At. mass of X} \times \frac{mass of AgX formed}{mass of compound taken} \times 100$		
Nitrogen	Dumas method : Nitrogen containing organic compound is heated with CuO in an atmosphere of CO ₂ . $C_xH_yN_z + (2x + \frac{y}{2})CO_2 \longrightarrow xCO_2 + \frac{y}{2}H_2O + N_2 + (2x + \frac{y}{2})Cu$ N_2 evolved gets collected over conc. KOH solution which absorbs all other gases. % of N = $\frac{28}{22400} \times \frac{Vol. of N_2 at STP}{mass of compound taken} \times 100$		
	Kjeldahl's method :Organic compound + $H_2SO_4 (conc.) \longrightarrow (NH_4)_2SO_4 \xrightarrow{2NaOH} Na_2SO_4 + 2NH_3 + 2H_2O + 2NH_3 + H_2SO_4 \longrightarrow (Na_4)_2SO_4$ % of N = $\frac{1.4 \times molarity of acid \times vol. of acid used \times basicity of acid}{mass of compound taken}$		

Sulphur	Carius method :Sulphur in organic compound is converted into H_2SO_4 by boiling with Na_2O_2 or conc. HNO3 and is precipitated as $BaSO_4$ by adding excess of $BaCl_2$ solution in water. $S \frac{(i) HNO_3, \Delta}{(ii) BaCl_2} \rightarrow BaSO_4 \downarrow$ white ppt.% of $S = \frac{32}{233} \times \frac{mass of BaSO_4 formed}{mass of compound taken} \times 100$	
Phosphorus	Ignition method : $P \xrightarrow{HNO_{3}}_{heat} H_{3}PO_{4}$ $H_{3}PO_{4} + Mg^{2+} + NH_{4}CL \xrightarrow{\Delta} MgNH_{4}PO_{4} + HCl$ $Magnesium ammonium phosphate (white ppt.)$ $2MgNH_{4}PO_{4} \xrightarrow{\Delta} Mg_{2}P_{2}O_{7} + 2NH_{3} + H_{2}O$ $Magnesium pyrophosphate$ $\% \text{ of } P = \frac{62}{222} \times \frac{mass \text{ of } Mg P_{2}O_{7} \text{ formed}}{mass \text{ of compound taken}} \times 100$	





MIND MAP : ORGANIC CHEMISTRY



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MULTIPLE CHOICE QUESTIONS (MCQ)

1. Homolytic fission of C-C bond in ethane gives an intermediate in which carbon is: a. sp³ hybridised b. sp² hybridised c. sp-hybridised d. sp³d- hybridized 2. The kind of delocalization involving sigma bond in conjugation with pi electrons is called: a. Inductive effect b. Hyperconjugation effect c. Electromeric effect d. Mesomeric effect 3. Which organic species has only one type of hybridized carbon? b. $CH_3 - CH - CH_3$ a. $CH_2 = C = CH_2$ d. $CH_2 = CH - CH_2$ c. $CH_2 - C = CH$ 4. Which of the following can act as an electrophile? a. CNb. OH⁻ c. H₂O d. BF, 5. Which of the following is correct about the species: $(CH_3)_3 - C^+$ b. Its C+ is sp² hybridised a. It is planar d. All of these c. A nucleophile can attack on its C^+ 6. Which of the following has all the effects namely Inductive, Mesomeric and Hyperconjugative ? b. $CH_3CH = CH_2$ a. CH₃Cl d. $CH_2 = CH - CH = CH_2$ c. CH₃CH=CHCOCH₂Cl 7. The most stable free radical among the following is: a. C₆H₅CH₂CH₂ b. C₆H₅CHCH₃ c. CH₃CH₂ d. CH₃CHCH₃ 8. Isomers of a compound must have : a. Same physical properties b. Same chemical properties d. Same molecular weight c. Same structural properties 9. The type of isomerism not found in alkenes is : a. Chain isomerism b. Geometrical isomerism c. Metamerism d.Position isomerism





10. Which of the following species have six π conjugated electrons?

a. (b.
$$CH_2 = CH - CH = CH - CH_2^-$$
 c. (N) d. All of these

11. The correct decreasing order of priority for the functional groups of organic compounds in the IUPAC system of nomenclature is:

a. -COOH, -SO₃H, -CONH₂, -CHO b. -SO₃H, -COOH, -CONH₂, -CHO c. -CHO, -COOH, -SO₂H, -CONH₂ d. -CONH₂, -CHO, -SO₂H, -COOH

- 12. The IUPAC name of CH_3 -CH=CH-C=CH is: a. pent-3-en-1-yne b. pent-3-en-4-yne c. pent-2-en-4-yne d. pent-2-en-3-yne
- 13. Different structures generated due to rotation about C-C axis of an organic molecule are example of:
 - b. Conformational isomerism a. Geometrical isomerism
 - c. Optical isomerism d. Structural isomerism
- 14. Which of the following process is not used for the preparation of solid impurities?
 - a. Distillation b. Sublimation c. Crystallisation d. Vapouristaion
- 15. Quantitative measurement of nitrogen in an organic compound is done by the method:
 - a. Berthelot method b. Lassaigne method d. Kjehldahl method
 - c. Carius method

ANSWERS

1. a 2. b 3. d 4. d 5. d 6. c 7. b 8. d 9. c 10. d 14. d 15. a 11. a 12. a 13. b

FILL IN THE BLANKS

- The CH_4 molecule has _____ 1. structure.
- The property of carbon responsible for the large number of carbon compounds 2. is called .
- A triple bond between two carbon atoms is composed of one 3. and bonds.

Organic Chemistry: Some Basic Principles and Techniques

- 4. An organic compound which decomposes below its boiling point can be purified by _____.
- 5. The central atom of $CH_2 = C = CH_2$ is ______ hybridized.
- 6. There is a difference of _____ mass units between two consecutive members of a homologous series.
- 7. Geomerical isomerism happens due to ______ around p bond.
- 8. Electrophiles are the species which attack the regions of ______ electron density.
- 9. Hyperconjugation effect is also known as _____ resonance.
- 10. In Duma's method, the nitrogen present in an organic compound is set free as _____.

ANSWERS

- 1. Tetrahedral 2. Catenation 3. σ , two π 4. Vacuum distillation 5. sp
- 6. 14 7. Restricted rotation 8. high 9. No-bond 10. Nitrogen

TRUE AND FALSE TYPE QUESTIONS

- 1. In homologous series all the members have the same physical properties.
- 2. IUPAC name of CH₃CN is Methanenitrile.
- 3. Cis-trans isomers have different dipole moments.
- 4. Ethanol and methoxymethane are position isomers.
- 5. A free radical is a species with an unpaired valence electron.
- 6. Acetylene is a linear molecule.
- 7. Resonance brings down the stability of molecule.
- 8. Inductive effect is observed in π bond in presence of attacking reagent.
- 9. The percentage of carbon and hydrogen are estimated simultaneously in an organic compound by Liebig method.
- 10. Chromatography is the method used to separate and purify compounds when present in small amounts.

ANSWERS

1. F 2. F 3. T 4. F 5. T 6. T 7. F 8. F 9. T 10. T

ASSERTION REASON TYPE QUESTIONS

The questions given below are Assertion (A) and Reason (R). Use the following key to select the correct answer.

(a) If both assertion and reason are correct and reason is correct explanation for _assertion.





- (b) If both assertion and reason are correct but reason is not correct explanation for assertion.
- (c) If reason is correct but assertion is incorrect.
- (d) If both assertion and reason are incorrect.
- 1. Assertion: But-1-ene and 2-Methylprop-1-ene are position isomers. Reason: Position isomers have same molecular formula but different arrangement of carbon atoms.
- 2. Assertion: All the carbon atoms of But-2-ene lie in one plane. Reason: All the carbon atoms in But-2-ene are sp2 hybridized.
- Assertion: Alkanes having more than three carbon atoms exhibit chain isomerism.
 Beason: All earbon atoms in alkanes are sp hybridised.

Reason: All carbon atoms in alkanes are sp-hybridised.

- 4. Assertion: In $CH_2 = C = CH_2$, all the carbon atoms are sp² hybridised. Reason: All the hydrogen atoms lie in one plane.
- Assertion: Butane and 2-Methylbutane are homologues. Reason: Butane is a straight chain alkane while 2-Methylbutane is branched chain alkane.
- Assertion: Tertiary carbocations are generally formed more easily than primary carbocations.
 Reason: Hyperconjugation as well as inductive effect due to additional alkyl groups stabilize tertiary carbocations.
- 7. Assertion: Alkyl carbanions like ammonia have pyramidal shape. Reason: The carbon atom carrying negative charge has an octet of electrons
- Assertion: Carbocations are planar in nature. Reason: Carbocations are sp² Hybridised.
- Assertion: IUPAC name of compound CH₃CH=CH-CHO is But-2-enal. Reason: Functional group gets preference over multiple in IUPAC name of a compound.
- Assertion: Compounds with difference in their boiling points by about 30°C can be separated by simple distillation.

Reason: All liquid mixture can be separated by distillation method.

ANSWERS

1. d 2. c 3. c 4. d 5. b 6. a 7. b 8. a 9. a 10. c

Organic Chemistry: Some Basic Principles and Techniques

MATCH THE COLUMNS

Match the statements (a,b,c,d) in column I with the statements (I,ii,iii,iv) in column II.

	• • • • • • • • • • • • • • • • • • • •	
1.	Column I	Column II
	a. Leibig method	i. N ₂
	b. Dumas method	ii. AgX
	c. Kjehldahl method	iii. CO ₂ and H ₂ O
	d. Carius method	iv. NH ₃
2.	Column I	Column II
	a. Nonbenzenoid aromatic compound	i. 50% s character
	b. Catenation	ii. Species containing single unpaired nonbonding electrons
	c. Free radical	iii. Chain-forming property of an element
	d. sp-hybridised carbon atom	iv. Tropolone
AN	SWERS	
1.	a. iiii b. iv c. i d. ii	
2.	a. iv b. iii c. ii d. i	

ONE WORD ANSWER TYPE QUESTIONS

- 1 Write the formula of next higher homologue of C_2H_5OH .
- 2. Mention the hybridisation of underlined carbon in $CH_3C \equiv N$.
- 3. What type of isomerism is shown by Pentane and 2-Methylbutane?
- 4. Nucleophiles are Lewis acids or Lewis bases?
- 5. What type of bond fission results in the formation of free radicals?
- 6. What is the number of electrons present in the outermost shell of carbon in the methyl radical?
- 7. What is the other name for no-bond resonance?
- 8. What is the name of the Prussian blue coloured compound formed in Lassaigne's test for nitrogen in an organic compound?
- 9. SO_3 is an electrophile or nucleophile in sulphonation reaction of benzene?
- 10. Name suitable technique of separation of the components from a mixture of calcium sulphate and camphor.





1-MARK QUESTIONS

- 1. Which unique property of carbon is responsible for the large number of carbon compounds?
- 2. How many σ and π bonds are there in propyne?
- 3. What is the hybridization of carbon in ethyne?
- 4. Which has the longest C-C bond length among ethane, ethene and ethyne.
- 5. How many secondary carbon atoms are present in 2-Methylpentane?
- 6. Draw structure of 3-Isopropyl-2-methylhexane.
- 7. Draw bond line structure of $CH_3(CH_2)_6CH = CH(CH_2)_2 COOH$
- 8. What are the bond angles in sp^3 , sp^2 and sp hybrid orbitals?.
- 9. Write the formulae of first four members of homologous series of alkyne family.
- 10. Write the correct of priority of the following functional groups: $-C \equiv N, \geq C = O, -OH, -COOH, -CONH_2$
- 11. Write IUPAC name of :
 - (i) $CH_3 CH_2 CN$
 - (ii) $CH_2 = CHCH_2OH$
 - (iii) CH₃CH₂CH(CH₃)-CO-CH₂CH₃

$$(iv) CH_3CH_2 - O - CH_2CH(CH_3)CH_3$$

(v) $Cl-CH_2-C\equiv CH$

- 12. What type of isomerism is exhibited by Propanal and Propanone?
- 13. What is the essential condition for a compound to exhibit geometrical isomerism?
- Classify the following into electrophiles and nucleophiles: H+, NH₃, AlCl₃, NO₂⁺, CN⁻, H₂O, ROH, RNH₂, Carbocation
- 15. What type of attacking reagents are produced by heterolytic cleavage of covalent bond?
- 16. Name each of the following species and indicate which member of each pair is more stable:

(i)
$$CH_3^+$$
, $CH_3CH_2^+$

(ii)
$$C_6H_5CHCH_3$$
, CH_3CHCH_3

(iii)
$$CH_2 = CH - CH_2$$
, $CH = CH - CH_3$

(iv)
$$CH_3 - \overline{CH}_2$$
, $CH_3 - \overline{CH} - CH_3$

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- 17. Identify electrophilic centre in CH₃CHO.
- 18. What is state oh hybridization of positively charged carbon atom in carbocation?
- 19. What is the effect of introducing an alkyl group on the stability of carbocation?
- 20. Out of Benzyl and ethyl carbocation which is more stable and why?
- 21. Arrange the following in increasing order of acidic strength: ClCH₂COOH, CH₃CH₂COOH, ClCH₂COOH
- 22. Name two solvents which are commonly used to dissolve organic solids.
- 23. Name the technique that can be used for purification of iodine that contains traces of NaCl.
- 24. A liquid (10 mL) has three components A, B, C. which technique is suitable to sparate A,B, C from such a small amount of mixture?
- 25. Name one commonly used adsorbent in column chromatography.
- 26. Under what condition do we use fractional distillation?
- 27. A liquid compound starts decomposing well before its boiling point under normal pressure. How will you purify it?
- 28. Which elements are normally not detected in an organic compound?
- 29. For which type of compounds Kjehldahl's method is not useful?
- 30. How do you precipitate sulphur in Carius method?
- 31. Which method is used to estimate carbon and hydrogen?
- 32. What do we notice in Lassaigne's test if the compound contains both nitrogen and sulphur?

2-MARKS QUESTION

- 1. How will you account for the presence of large number of organic compounds?
- 2. Draw the structural formulae of the following compounds:
 - (i) Ethoxypropane
 - (ii) But-1-en-3-yne
 - (iii) 3,4,4,-Trimethylhex-1-yne
 - (iv) sec-butyl alcohol
 - (v) But-2-enoic acid
- 3. How is alkyl group represented? Give the structure and the names of the alkyl groups which originate from (i) n-Butane (ii) isobutene





- 4. Give IUPAC name of the following compounds:
 - (i) $C_{6}H_{5}CH_{2}CH_{2}OH$ (ii) $(CH_{3})_{2}CH_{2}CH_{2}CHO$ (iii) $CH_{2}=CH-C\equiv N$ (iv) $CH_{3}-CH-C-CH-CH_{3}$ | || |Br O CH_{3} (v) $CH_{3}-CH-CH=C-CH_{2}-COOH$ | | |OH OH_{3}
- 5. What is functional isomerism? Give two examples.
- 6. Distinguish between position isomerism and functional isomerism.
- 7. What is metamerism? Give example.
- 8. How are free radicals formed?
- 9. What is the effect of introducing an alkyl group on the stability of a free radical?
- 10. Give two examples each of the groups exerting -I and + I effect when attached to a chain of carbon atoms.
- 11. A tertiary butyl carbocation is is more stable than isobutyl carbocation. Justify.
- 11. What do you understand by +R and -R effect?
- 12. Define hyperconjugation.
- 13. What is the difference between inductive and electromeric effect?
- 14. All electrophiles are Lewis acids while nucleophile are Lewis bases. Explain.
- 15. What is the purpose of filtration through hot water funnel?
- 16. What precautions are necessary while purifying an organic solid with the help of crystallization process?
- 17. Discuss the principle of steam distillation.
- 18. Discuss the role of fractionating column in fractional distillation.
- 19. How will you prepare Lassaigne's extract? Name the elements which can be detected from this extract?
- 20. Why do we boil Lassaigne's extract with conc. HNO_3 while detecting halogens in an organic compound?

Organic Chemistry: Some Basic Principles and Techniques

3-MARKS QUESTIONS

- 1. Discuss the orbital structure of ethene.
- 2. Define (i) Functional group (ii) Homologous series.
- 3. What do you understand by 1°, 2°, 3° and 4° carbon? Write one example having atleast one carbon of each type.
- 4. Why stability of carbocations follows the order: tertiary>secondary>primary?
- 5. What are the various conditions essential for resonance?
- 6. Write resonance structures of $CH_2 = CH CHO$. Indicate relative stability of the contributing structures.
- 7. Inductive effect is of permanent nature while electromeric effect is only temporary. Explain.
- 8. What is chromatography? Name different types of chromatographic processes.
- 9. You are given a mixture of methanol and acetone. Discuss the process which you will employ to separate them.
- 10. Explain the reason for the fusion of an organic compound with metallic sodium for testing nitrogen, sulphur and halogens.

5-MARK QUESTIONS

- 1. What are free radicals? Justify the stability of the aliphatic primary, secondary and tertiary free radicals.
- 2. What is a carbanion? How is it formed? Discuss relative stability of primary, secondary and tertiary carbanion.
- 3. Arrange the following in the order of property indicated against each set:
 - (i) $-C_6H_5$, $-NO_2$, -COOH, -I, -F, $-CH_3$, $-C_2H_5$ (In the increasing order of -I effect)
 - (ii) CH₃CH₂CH₂CH₂⁺, (CH₃)₃C⁺, CH₃CH₂CH₂CHCH₃ (In the order of increasing stability)
 - (iii) -Cl, -CONH₂, -CHO (In the increasing priority order if present in same molecule)





- 4. Draw the resonance strutuctures for the following compounds. Show the electron shift using curved arrow notation.
 - (i) $C_6H_5NO_2$
 - (ii) CH₃CH=CHCHO
 - (iii) C₆H₅OH
 - (iv) $C_6H_5CH_2^+$
 - (v) $CH_3CH = CHCH_2^+$
- 5. Suggest a method to separate the constituents from the following mixture:
 - (i) Mixture of two miscible liquids
 - (ii) A mixture of oil and water
 - (iii) A mixture of plant pigments
 - (iv) A mixture of solid benzoic acid and sodium chloride
 - (v) o-Nitrophenol and p-Nitrophenol present in the mixture.
- 6. 0.378g of an organic compound containing carbon and hydrogen was subjected to combustion by Leibig's method, the CO_2 and H_2O formed were passed through potash bulbs and anhydrous $CaCl_2$ tube. At the end of the experiment, the increase in the respective weights were 0.264g and 0.162g. Calculate the percentage of carbon and hydrogen.

(Ans: C = 19.05%, H = 4.76%)

UNIT TEST

Time Allowed: 1 hr Maximum Mar		s : 20	
(i)	neral Instructions: All questions are compulsory. Maximum marks carried by each question	on are indicated against it.	
1.	Write bond line formula for the followin HOCH ₂ CH ₂ CH ₂ CH(CH ₃)CH(CH ₃)COC		[1]
2.	Write IUPAC name of the following con $CH_3 - CH - C - CH - CH_3$ $NO_2 O CH_3$	npound:	[1]
3.	The central atom of compound $CH_2 = C$	=CH ₂ is hybridized.	[1]
4.	There is a difference of	mass units between two eries.	[1]
5	5		[1]
6.	 (i) What type of isomerism is exhibited compounds? CH₃CH₂CHO and CH₃-C-C 	by the following pair of	[1]
	0		

- (ii) Give one example each of nucleophile and electrophile.
- 7. (i) Arrange the following in increasing order of stability:

$$(CH_3)_3C^-$$
, $CH_3 - CH - CH_3$, $CH_3CH_2^-$

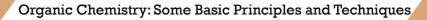
(ii) Differentiate between inductive and electromeric effect.

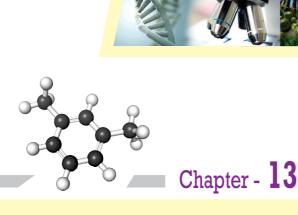




- 8. (i) When do we use hot water funnel for filtration?
 - (ii) How will you separate a mixture bof two organic compounds which have different solubilities in the same solvent?
 - (iii) An organic liquid decomposes below its boiling point. How will you purify it?
- 9. Draw the resonating structures of (a) Phenol (b) Benzaldehyde.
- 10. Arrange the following in the order of property indicated against each set:

- (iv) HCOOH, CH₃COOH, ClCH₂COOH (Increasing order of acidic strength)
- (v) O₂NCH₂CH₂O⁻, CH₃CH₂O⁻ (species having greater stability)





Hydrocarbons

FAST TRACK : QUICK REVISION

Hydrocarbons are the organic compounds containing carbon and hydrogen only. Depending upon the types of carbon-carbon bonds present, hydrocarbons can be classified into three categories- (i) Saturated (ii) Unsaturated (iii) Aromatic hydrocarbons.

Saturated hydrocarbons contain carbon-carbon muiltiple bonds–double bonds, triple bonds or both.

ALKANES : Saturated open chain hydrocarbons containing carbon-carbon single bonds. These are inert under normal conditions i.e. do not react with acids, bases and other reagents. Alkanes exhibit Chain isomerism, Position isomerism and conformational isomerism.

General methods of preparation of alkanes :

1. From Unsaturated hydrocarbons : By hydrogenation in the presence of platinium, palladium or nickel as catalyst.

General Chemical Equation : $R-CH=CH_{2}+H_{2} \xrightarrow{Ni} R - CH_{2} - CH_{3}$ [Where R is H or Alkyl group] *e.g.* $CH_{2}=CH_{2}+H_{2} \xrightarrow{Ni} CH_{3}-CH_{3}$ Ethene $CH_{3}-C \equiv CH+H_{2} \xrightarrow{Ni} CH_{3}-CH=CH_{2}+H_{2} \xrightarrow{Ni} CH_{3}-CH_{2}-CH_{3}$ Propyne

2. From alkyl halides : on reduction with Zinc and dilute hydrochloric acid

$$\begin{array}{ccc} CH_{3}-Cl+H_{2} & \xrightarrow{Zn, H^{+}} & CH_{4}+HCl \\ Chloromethane & & Methane \\ C_{2}H_{5}-Cl+H_{2} & \xrightarrow{Zn, H^{+}} & C_{2}H_{6}+HCl \\ Chloroethane & & Ethane \end{array}$$

 $\begin{array}{c} CH_{3}CH_{2}CH_{2}Cl+H_{2} & \xrightarrow{Zn, H^{+}} & CH_{3}CH_{2}CH_{3}+HCl \\ 1-Chloropropane & Propane \end{array}$

3. From alkyl halides by Wurtz reaction : Reaction of alkyl halide with sodium in dry ether, useful only for the preparation of symmetrical alkanes.

 $CH_{3}CH_{2}Br + 2Na + Br CH_{2}CH_{3} \xrightarrow{Dry Ether} CH_{3}CH_{2}CH_{2}CH_{3}$ Bromoethane Butane

 $CH_{3}CH_{2}CH_{2}Br+2Na+BrCH_{2}CH_{2}CH_{3} \xrightarrow{Dry \ Ether} CH_{3}CH_{2}CH_{2}CH_{2}CH_{2}CH_{2}CH_{2}CH_{2}CH_{3}$ 1-Bromopropane CH_{3}CH_{2}CH_{2}CH_{2}CH_{3}CH_{2}CH_{2}CH_{2}CH_{3}CH_{2}CH_{2}CH_{2}CH_{3}CH_{2}CH_{2}CH_{3}CH_{2}CH_{3}CH_{2}CH_{3}CH_{

 $\begin{array}{c} \text{CH}_{3}\text{CH}_{2}\text{ Br} + 2\text{Na} + \text{Br} \text{ CH}_{2}\text{CH}_{3} \xrightarrow{\text{Dry Ether}} \text{CH}_{3}\text{CH}_{2}\text{CH}_{2}\text{CH}_{2}\text{CH}_{3} \\ \text{1-Bromopropane} & \text{1-Bromoethane} & \text{n-Pentane (desired)} \end{array}$

4. From Carboxylic acids : By decarboxylation with soda lime

 $\begin{array}{c} CH_{3}COONa + NaOH \xrightarrow{\Delta} CH_{4} + Na_{2}CO_{3} \\ Sodium acetate & Sodium \\ Hydroxide \end{array}$

5. By Kolbe's electrolytic method : Electrolysis of an aqueous solution sodium or potassium salt of carboxylic acid. Alkane containing even number of carbon atoms is formed at anode.

$$2CH_3COO Na + 2H_2O \xrightarrow{\text{Electrolysis}} CH_3 - CH_3 + 2CO_2 + H_2 + 2NaOH$$

At Anode : (Oxidation)

$$\begin{array}{c} O \\ 2CH_{3} - \overset{I}{C} - O^{-} \xrightarrow{-2e^{-}} 2CH_{3} - \overset{O}{C} - O^{-} \longrightarrow 2CH_{3} + 2CO_{2} \\ 2CH_{3} \cdot \longrightarrow CH_{3} - CH_{3} \\ \text{At Cathode : (Reduction)} \\ 2H_{2}O \longrightarrow 2OH^{-} + 2H^{+} \\ 2H^{+} + 2e^{-} \longrightarrow H_{2} \end{array}$$

Physical Properties of alkanes :

- 1. Boiling point of alkanes decreases on branching due to decrease in surface ara of molecule with branching which decreases magnitude of van der Waal's forces of attraction.
- 2. Alkanes being non-polar in nature are soluble in non-polar solvents.

Hydrocarbons



Chemical properties of Alkanes :

Alkanes undergo substitution reactions.
 e.g., Halogenation, Nitration, Sulphonation.
 Halogenation : For example Chlorination of methane

 $\begin{array}{c} \mathrm{CH}_{4} + \mathrm{Cl}_{2} & \longrightarrow & \mathrm{CH}_{3}\mathrm{Cl} + \mathrm{HCl} \\ \mathrm{Chloromethane} \\ \mathrm{CH}_{3}\mathrm{Cl} + \mathrm{Cl}_{2} & \longrightarrow & \mathrm{CH}_{2}\mathrm{Cl}_{2} + \mathrm{HCl} \\ \mathrm{Dichloromethane} \\ \mathrm{CH}_{2}\mathrm{Cl}_{2} + \mathrm{Cl}_{2} & \longrightarrow & \mathrm{CHCl}_{3} + \mathrm{HCl} \\ \mathrm{Trichloromethane} \\ \mathrm{CHCl}_{3} + \mathrm{Cl}_{2} & \longrightarrow & \mathrm{CCl}_{4} + \mathrm{HCl} \\ \mathrm{Tetrachloromethane} \end{array}$

Rate of reaction of alkanes with halogens is $F_2 > Cl_2 > Br_2 > I_2$. Rate of replacement of hydrogen in alkanes is $3^\circ > 2^\circ > 1^\circ$. Fluorination is too violent to be controlled. Iodination is reversible and it is therefore carried out in the presence of oxidising agent like HNO₃.

Mechanism of halogenation : Free radical mechanism

 $Cl - Cl \xrightarrow{hv} Cl \cdot + \cdot Cl$ Initiation $Cl \cdot + CH_4 \longrightarrow H - Cl + \cdot CH_3$ Propagation $\cdot CH_3 + Cl - Cl \longrightarrow CH_3 Cl + \cdot Cl$ Propagation $Cl \cdot + \cdot CH_3 \longrightarrow CH_3 Cl$ Termination

Combustion : Complete combustion gives carbon dioxide and water.

 $CH_4 + 2O_2 \longrightarrow CO_2 + 2H_2O$

Isomerisation :

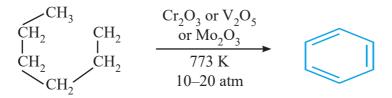
$$CH_{3} CH_{2}CH_{2}CH_{2}CH_{3} \xrightarrow{AlCl_{3} + HCl} CH_{3} CH_{3} CHCH_{3}$$

$$I = CH_{3}$$





Aromatisation :



(vi) Reaction with steam

 $CH_4 + H_2O \xrightarrow{Ni} CO + 3H_2$

Pyrolysis : Decomposition of higher alkanes to lower alkanes on heating.

$$C_{6}H_{14} \xrightarrow{773K} C_{6}H_{12} + H_{2}$$

 $C_{4}H_{8} + C_{2}H_{6}$
 $C_{3}H_{6} + C_{2}H_{4+}CH_{4}$

Conformations :

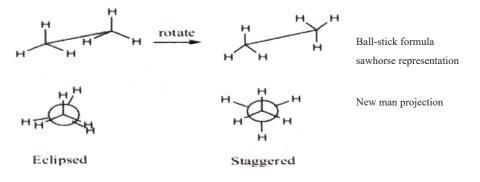
Different spatial arrangement of atoms arising due to rotation around C-C single bond.

Conformation of ethane, CH₃CH₃

Two conformational isomers or conformers.

Eclipsed form = all hydrogen atoms nearest to each other.

Staggered form = all hydrogen atoms are farthest apart.



Stability of eclipsed conformation is least while staggered conformation is most stable. The energy difference between two extreme forms is 12.5kJmol⁻¹. Due to this small energy difference the two forms are easily inter-convertible at ordinary temperature and cannot be separated and isolated.



Hydrocarbons



ALKENES

These are unsaturated non-cyclic hydrocarbons which have sp^2 -hybridisation with 120° bond angle.

Alkenes are also called olefins [oil.forming] which indicates their high reactive nature.

Alkenes have general formula $C_n H_{2n}$, where $n = 2, 3, 4, \dots$

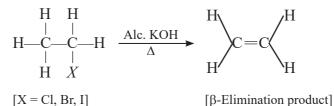
 C_2H_4 (ethene), C_3H_6 (propene), etc.

• Methods of Preparation of Alkenes

(i) From alkynes

$$R - C \equiv C - R' + H_2 \xrightarrow{Pd/C} \underset{H}{R - C} = C \xrightarrow{R}_{H}$$

(ii) From alkyl halide [Dehydrohalogenational]



(iii) From vicinal dihalides

$$\begin{array}{ccc} \mathrm{CH}_{2}\mathrm{Br}\mathrm{-\!\!-}\mathrm{CH}_{2}\mathrm{Br}+\mathrm{Zn} & \overset{\Delta}{\longrightarrow} & \mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{Zn}\mathrm{Br}_{2}\\ \mathrm{CH}_{3}\mathrm{-\!\!-}\mathrm{CH}\mathrm{-\!\!-}\mathrm{CH}_{2}+\mathrm{Zn} & \overset{\Delta}{\longrightarrow} & \mathrm{CH}_{3}\mathrm{-\!\!-}\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{Zn}\mathrm{Br}_{2}\\ \mathrm{Br} & \mathrm{Br} \end{array}$$

(iv) From alcohols by acidic dehydrogenation

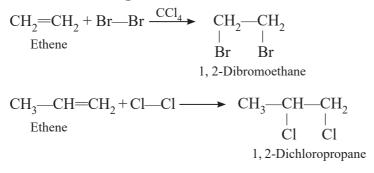
 $\begin{array}{ccccccc} H & H \\ H & -C & C \\ H & -C & -C \\ H & H \\ H & OH \\ Ethanol \end{array} \xrightarrow{\text{Conc. } H_2 \text{SO}_4} CH_2 = CH_2 + H_2 \\ CH_2 = CH_2 + H_2 \\ Ethene \\ Ethene \\ CH_2 = CH_2 + H_2 \\ Ethene \\ Ethene$





• Chemical Properties of Alkenes :

1. Addition of Halogens :



2. Addition of hydrogen halides HCl, HBr, Hl : Add up to alkenes to form alkyl halides as per their reactivity order in HI>HBr>HCl.

Addition reaction of HBr to unsymmetrical alkenes (Markownikov's rule) According to Markownikovs' rule, the negative part of the addendum (adding molecule) gets attached to that carbon atom which possesses lesser number of hydrogen atoms.

$$H \qquad Br \\ H - C - C = C - H + HBr \longrightarrow H_3C - C - CH_3 \\ H H H H H H H 2-Bromopropane$$

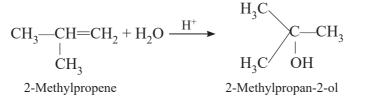
Anti Markownikov addition or peroxide effect or Kharasch effect in the presence of organic peroxide, addition of only HBr molecule on unsymmetrical alkene takes place contrary to the Markownikov's rule. Peroxide effect is not observed in case of HF, HCl and HI.

 $CH_{3} - CH = CH_{2} + HBr \xrightarrow{(C_{6}H_{5}CO)_{2}O_{2}} CH_{3}CH_{2}CH_{2}Br$ 1-bromopropane

3. Addition of sulphuric acid

 $\label{eq:ch3} \begin{array}{rcl} \mbox{CH}_3\mbox{--}\mbox{CH}_2\mbox{+}\mbox{HOSO}_2\mbox{OH} & \rightarrow &\mbox{CH}_3\mbox{CH}(\mbox{OSO}_3\mbox{H})\mbox{CH}_3 \\ \mbox{[cold and conc.]} & & \mbox{propyl hydrogen sulphate} \end{array}$

4. Addition of water



Hydrocarbons



5. Oxidation : Alkenes decolourise cold dilute aqueous solution potassium permanganate (Baeyer's reagent). It is used as a test for unsaturation.

$$CH_{2} = CH_{2} + H_{2}O + [O] \xrightarrow{Dil.KMnO_{4}} CH_{2} - CH_{2}$$
$$| \qquad | \qquad | \\OH OH$$
$$Ethane-1,2-diol or glycol$$

Acidic $KMnO_4$ or acidic $K_2Cr_2O_7$ oxidises alkenes to ketones and/or acids depending upon the nature of alkene and the experimental conditions.

 $(CH_{3})_{2}C=CH_{2} \xrightarrow{KMnO_{4}/H^{+}} (CH_{3})_{2}CO + CO_{2} + H_{2}O$ 2-Methyl propene Propane-2-one $CH_{3}-CH=CH-CH_{3} \xrightarrow{KMnO_{4}/H^{+}} 2CH_{3}COOH$ But-2-ene Ethanoic acid

6. Ozonolysis : Reaction of ozone with alkene to form ozonide which on subsequent reductive cleavage with zinc dust and water give carbonyl compounds (aldehydes & ketones).

$$H_{3}C$$

$$C=CH_{2}+O_{2} \xrightarrow{Zn/H_{2}O} H_{3}C$$

$$C=O+HCHO+H_{2}O_{2}$$

$$H_{3}C$$

$$Zn + H_2O_2 \longrightarrow ZnO + H_2O$$

7. Polymerization

$$n(CH_{2}=CH_{2}) \xrightarrow{\text{High temperature/pressure}}_{\text{catalyst}} - [CH_{2}-CH_{2}-]_{n}$$

$$polythene$$

$$n(CH_{3}-CH=CH_{2}) \xrightarrow{\text{High temperature/pressure}}_{\text{catalyst}} \left[-CH_{2}-CH_{2}- \right]_{n}$$





ALKYNES

These are unsaturated hydrocarbons with general formula C_nH_{2n-2} e.g., C_2H_2 (ethyne), C_3H_4 (propyne).

Alkynes also exhibit electrophilic addition reaction but less reactive than alkenes because the dissociation of π -electron cloud requires more energy.

H–C=C–H contins 3σ and 2π -bonds and bond length is 120 pm. In acetylene. H–C–C bond angle is 180°.

• Methods of Preparation of Alkynes

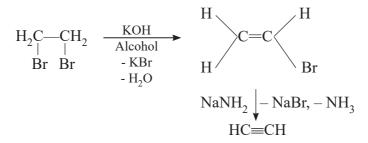
1. From calcium carbide

$$CaCO_{3} \longrightarrow CaO + CO_{2}$$

$$CaO + C \longrightarrow CaC_{2} + CO$$

$$CaC_{2} + 2H_{2}O \longrightarrow Ca(OH)_{2} + C_{2}H_{2}$$

2. From vicinal dihalides



3. From tetrahalides

 Br_2CH — $CHBr_2 + 2Zn \xrightarrow{CH_3OH} H$ — $C \equiv C$ — $H + 2ZnBr_2$

Physical Properties of Alkynes :

- 1. The first two members are gases next eight members $(C_5 C_{12})$ are liquids and higher members are solids.
- 2. They are all colourless and odourless with the exception of acetylene which has slightly garlic odour due to the presence of PH_3 and H_2S as impurities.
- 3. Alkynes are insoluble in water but soluble in organic solvents like ethers, carbon tetrachloride and benzene.
- 4. Melting point, boiling point and density increase with increase in molar mass.

Hydrocarbons

Chemical properties of Alkynes

Alkynes show electrophilic as well as nucleophilic addition reactions.

(i) Acidic character of alkyne

$$HC \equiv CH + Na \longrightarrow HC \equiv \overline{C}Na + \frac{1}{2}H_{2}$$

monosodium
acetylides
$$H - C \equiv CNa^{+} + Na \longrightarrow Na^{+}C \equiv \overline{C}Na^{+} + \frac{1}{2}H_{2}$$

disodium acetylide
$$CH_{3} - C \equiv C - H + NaNH_{2} \longrightarrow CH_{3} - C \equiv \overline{C}Na^{+} + NH_{3}$$

sodium propynide

These reactions are not shown by alkenes, alkanes and non-terminal alkynes, hence used for distinction between alkane, alkene and alkyne.

Acetylenic hydrogens are acidic in nature due to 50% s-character in sp-hybridised orbitals. Acidity of alkynes is lesser than water.

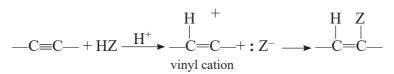
Acidic behaviour order

(i)
$$HC \equiv CH > CH_2 = CH_2 > CH_3 - CH_3$$

 sp sp^2 sp^3

(ii)
$$HC \equiv CH > CH_3 - C \equiv CH >> CH_3 - C \equiv C - CH_3$$

(ii) Electrophilic addition reactions



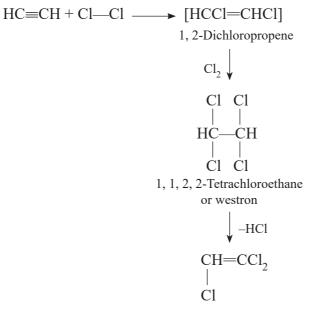
The addition product formed depends upon the stability of vinylic cation. Addition on unsymmetrical alkynes takes place according to Markovnikov's rule.

$$CH_{3} - C \equiv C - H + H_{2} \xrightarrow{Pt/Pd/Ni} [CH_{3} - CH = CH_{2}] \xrightarrow{H_{2}} CH_{3}CH_{2}CH_{3}$$
propene

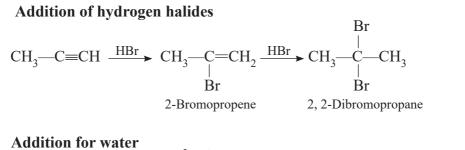


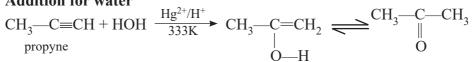


Addition of halogens

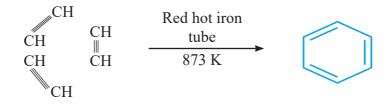


westrosol (1, 1, 2-Trichloroethene)





(iii) Cyclic polymerisation of ethyne





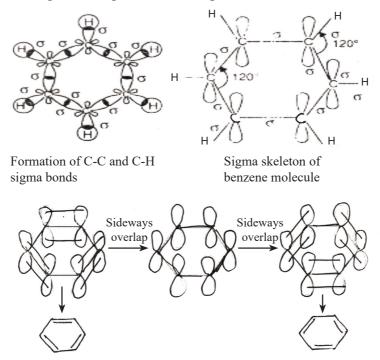
Hydrocarbons

AROMATIC HYDROCARBONS

These hydrocarbon are also known as arenes. The parent member of the family aromatic hydrocarbons is benzene.

Aromatic compounds containing benzene ring are known as benzenoids.

Structure of benzene : Hexagonal ring of carbon atoms with alternate single and double bonds. Each carbon atom is sp² hybridised. Planar ring, bond angle 120°. All C-C bond lengths are equal due to complete delocalisation of π electrons.



HUCKEL'S RULE

- Huckel's rule, (based on calculations) : a planar cyclic molecule with alternating double and single bonds has aromatic stability if it has $(4n + 2\pi)$ electrons (n is 0, 1, 2, 3, 4)
- For n = 1 : 4n+2 = 6; benzene is stable and electrons are delocalized.

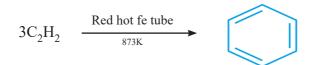
BenzeneThree double bonds;six π electrons



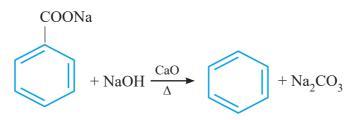


• METHODS OF PREPARATION

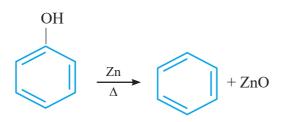
1. Cyclic polymerisation of ethyne



2. Decarboxylation of aromatic acids



3. Reduction of phenol



Physical Properties of Benzene :

- (i) Aromatic hydrocarbons are non-polar molecules and are usually colourless liquids or solids with a characteristic aroma.
- (ii) Aromatic hydrocarbons are immiscible with water but readily miscible with organic solvents.
- (iii) Aromatic compounds burn with sooty flame.

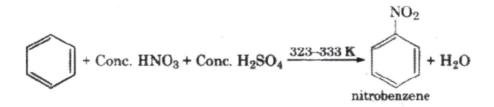
Chemical Reactions of Benzene :

- (i) Benzene gives electrophile substitution reactions.
- (ii) According to experimental evidences, electrophile substitution reaction involve following three steps :
 - Generation of electrophilie
 - Formation of carbocation intermediate.
 - Removal of proton from the carbocation intermediate.

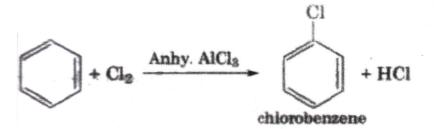


Hydrocarbons

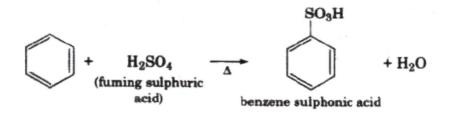




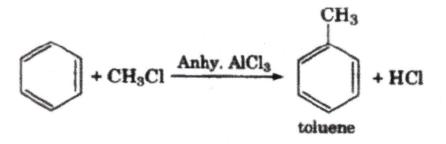
(ii) Halogenation



(iii) Sulphonation

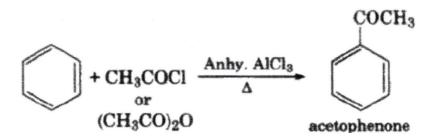


(iv) Friedel-Craft's alkylation reaction

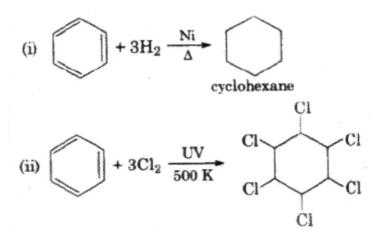








Benzene also undergoes addition reactions e.g.



benzene bexachloride or 666 (BHC or Gammexane or lindane)

• **COMBUSTION**

 $2C_6H_6 + 15O_2 \longrightarrow 12CO_2 + 6H_2O$

Directing influence of substituents in monosubstituted benzene

- (i) **Ortho and para directing groups :** Ring activating groups e.g., NH₂, CH₃, C₂H₅, OCH₃ etc. (+ R effect)
- (ii) Meta directing groups : Ring deactivating groups

e.g. - NO_2 , - CN, - CHO, - COOH, - SO_3H (- R effect).

MULTIPLE CHOICE QUESTIONS (MCQ)

1.	ro dipole moment?			
	(a.) cis-But-2-ene	(b) trans-But-2-ene		
	(c) But-1-ene	(d) 2-Methylprop-1-ene		
2.	Bond length of (I) ethane, (II) ethene, (III) Acetylene, (IV) Benzene follows the order:			
	(a) $I > II > III > IV$	(b) $I > II > IV > III$		
	(c) $I > IV > II > III$	(d) $III > IV > II > I$		
3. The methyl group in benzene ring is:				
	(a) Ortho directing	(b) Ortho and meta directing		
	(c) Para directing	(d) Ortho and para directing		
4. The dihedral angle in the staggered conformation of C_2H_6				
	(a) 120°	(b) 60°		
	(c) 0°	(d) 90°		
5.	On heating C_2H_2 in red hot tub	be copper tube, the compound formed is:		
	(a) Ethylene	(b) Benzene		
	(c) Ethane	(d) Methane		
6.	Action of acetylene on dilute H	I_2SO_4 gives:		
	(a) Acetic acid	(b) Acetone		
	(c) Acetaldehyde	(d) Ethyl alcohol		
7. Which of the following compounds will exhibit cis-trans (geomisomerism?				
	(a) Butanol	(b) 2-Butyne		
	(c) 2-Butenol	(d) 2-Butene		
8.	Hydrolysis of ozonide of But-1-ene gives			
	(a) Ethylene only			
	(b) Acetaldehyde and formaldehyde			
	(c) Propionaldehyde and formaldehyde			
	(d) Acetaldehyde only			
9.	Reaction of hydrogen bromide with propene in the absence of a peroxide is a/an			
	(a) free radical reaction	(b) nucleophilic substitution		
	(c) electrophilic addition	(d) nucleophilic substitution		
	()r	() ···································		





10.	Among the following compounds, the one which is most reactive towards electrophilic nitration is:		
	(a) Benzoic acid (b) Nitrobenzene		
	(c) Toluene (d) Benzene		
ANS	WERS: 1.b 2.c 3.d 4.b 5.b 6.c 7.d 8.c 9.c 10.c		
FILL IN THE BLANKS			
1.	Alkanes mainly undergo reaction.		
2.	Halogenation of alkanes does not occur in		
3.	The H – C – H bond angle in ethene is		
4.			
5.	Baeyer's reagent is used for testing		
6.	Benzene favours substitution reaction.		
7.	The Dipole moment of Benzene is		
8.	The nitro group in the benzene nucleus is directing. It the reactivity of the benzene ring.		
9.	Melting point and boiling point increase as the molar masses		
10. The reaction of solid calcium carbide with water produces, a flammable gas.			
ANS	WERS: 1. Substitution2. Dark3. 120°4. Markownikov's5. Unsaturation6. electrophilic7. Zero8. Meta, decreases9. Increase10. Ethyne		

TRUE AND FALSE TYPE QUESTIONS

- 1. Alkanes mainly undergo substitution reactions using the free-radical mechanism.
- 2. The decreasing order of boiling points among the isomeric pentanes is neo > iso > n.
- 3. The acidic character of three types of hydrocarbons follows the order alkanes > alkenes > alkynes.
- 4. The peroxide effect is observed only in addition of HBr, and not with HCl and HI.





- 5. Wurtz reaction is suitable for the preparation of both symmetrical and unsymmetrical alkanes.
- 6. For a compound to be aromatic it must have $(4n + 2)\pi$ electrons.
- 7. Benzene has planar structure.
- 8. The benzene molecule has two different carbon-carbon bond lengths, corresponding to alternate single and double bonds.
- 9. In Friedel-Crafts reaction, AlCl₃ is an electrophile.
- 10. An electron-donating substituent in benzene ring gives a meta product.

ANSWERS: 1. T 2. F 3. F 4. T 5. F 6. T 7. T 8. F 9. F 10. F

MATCH THE COLUMNS

Match the statements (a,b,c,d) in column I with the statements (i, ii, iii, iv) in column II.

1. Co	olumn I	Column II		
a.	C_2H_6	i. Ele	ctrophilic add	dition
b. C ₂ H ₄		ii. Electrophilic substitution		
c. C ₂ H ₂		iii. Free radical substitution		
d.	. C ₆ H ₆	iv. Fr	ee radical add	dition
2. Column I a. Alkanes b. Alkenes		Column II		
		i. Saturated nature		
		ii. Ozonolysis		
c.	Alkynes	iii. Geometrical isomeris		
d.	Arenes	iv. Aromatic character		cter
ANSWERS:	1. a. $\rightarrow iii$	$b. \rightarrow i$	$c. \rightarrow i$	$d. \rightarrow ii$
	2. a. $\rightarrow i$	b. → ii, iii	$c. \rightarrow ii$	$d. \rightarrow i, iv$

ASSERTION-REASON TYPE QUESTIONS

Type 1. The questions given below consist of Assertion(A) and Reason (R). Use the following key to select correct answer.

- (a) If both assertion and reason are correct and reason is correct explanation for assertion.
- (b) If both assertion and reason are correct but reason is not correct explanation for assertion.
- (c) If assertion is correct but a reason is incorrect.
- (d) If assertion and reason both are incorrect.

Chemistry Class XI



1. Assertion: The IUPAC name of $CH_3CH = CH - C \equiv CH$ is pent-3-en-1-yne and not pent-2-en-4-yne.

Reason: While deciding the locants of double and triple Bonds, lowest sum rule is always followed.

- Assertion: Tropylium cation is aromatic in character.
 Reason: The only property which decides the aromatic character is its planar nature.
- 3. Assertion: Friedel-craft reaction between benzene and acetic anhydride in the presence of anhydrous AlCl₃ yields acetophenone and not polysubstituted products.

Reason: Acetophenone formed poisons the catalyst preventing further reaction.

4. **Assertion:** But-1-ene on reacting with HBr in the presence of peroxide, products 1-bromobutane.

Reason: It involves the formation of a primary free radical.

5. Assertion: Alkanes with more than three carbon atoms exhibit chain isomerism.

Reason: Branching of the carbon atom chain is necessary for exhibiting chain isomerism.

- Assertion: Benzene reacts with chlorine in the form of light to form BHC.
 Reason: BHC is also called gammexane or 666.
- 7. Assertion: All the hydrogen atoms in $CH_2 = C = CH_2$ lie in one plane. Reason: All the carbon atoms in it are sp² hybridised.
- 8. **Assertion:** Propene reacts with HBr in the presence of benzoyl peroxide to yield 2-bromopropane.

Reason: In the presence of peroxide, the addition of HBr to propene follows ionic mechanism.

- Assertion: Benzene does not decolourise bromine water.
 Reason: Benzene is stabilised by resonance due to delocalisation of π electrons.
- 10. Assertion: Acidity of C-H bond decreases in the order: $HC \equiv CH > H_2C = CH_2 > H_3C - CH_3$

 $\label{eq:Reason: Greater the percentage s-character, more is the acidity of C-H bond.$

ANSWERS: 1.a 2.c 3.c 4.c 5.a 6.b 7.c 8.d 9.a 10.a





ONE WORD TYPE QUESTIONS

- 1. What is the state of hybridisation of Carbon atoms in alkanes?
- 2. What is self linking property of atoms called?
- 3. What is the general formula of alkenes?
- 4. What is the other name for Geometrical isomerism?
- 5. Which out of Ethane, Ethene and Ethyne has longest C-C bond?
- 6. What is the number of bonds in But-3-en-1-yne?
- 7. Name the product formed when Propyne is treated with aqueous H_2SO_4 in the presence of dil.HgSO₄.
- 8. What is C-C-C bond angle in benzene?
- 9. Name the product formed when Benzene reacts with CH_3Cl in the presence of anhydrous aluminium chloride.
- 10. -COOH is ortho, para directing or Meta directing group?

1-MARK QUESTIONS

- 1. Name the chain isomer of C_5H_{12} which has tertiary carbon atom.
- 2. Give the IUPAC name of the lowest molecular weight alkane that contains a quaternary carbon.
- 3. Write the reaction involved in Kolbe's electrolytic method to prepare ethane.
- 4. Define term decarboxylation.
- 5. Why dry ether and not water is used as a solvent in the preparation of alkane by Wurtz reaction?
- 6. Sodium salt of which carboxylic acid will be needed for the preparation of propane by decarboxylation method?
- 7. Complete the following reaction:

 $CH_3Cl + Na$ dry ether

- 8. Amongst the following which one has the maximum boiling point? n-Pentane, iso-pentane, neo-pentane.
- 9. Define the term conformation.
- 10. Write IUPAC name of $CH_3CH = CHCH_2CH = CCH_2CH = CH_2$

Chemistry Class XI



CH₂CH₂

- 11. Draw the cis and trans isomers of CHCl=CHCl.
- 12. What happens when 2-Bromobutane is being treated with alc. KOH?
- 13. Name the reagents used to carry out the following conversions: $CH_3-CH=CH_2 \rightarrow CH_3CH(OH)CH_2OH$
- 14. Complete the following reaction :

 $CH_2 - CH = CH_2 + HBr$ Organic peroxide

- An alkene A on ozonolysis gives a mixture of ethanol and pentan-3-one. Write IUPAC name of element.
- 16. When alkyne is treated with bromine water then what will be the colour of the product?
- 17. Why alkynes do not exhibit geometrical isomerism?
- 18. Complete the following reaction:
 - (i) $CH_3C \equiv CH \xrightarrow{H_2O, Hg^{2+}/H^+}$? (ii) $CaC_2 + 2H_2O \longrightarrow +$
- 19. How will you convert ethyne to benzene?
- 20. Write chemical equation for combustion reaction of Hexyne.
- 21. Write IUPAC name of $C_6H_5 CH_2 CH_2 CH = CH_2$.
- 22. Why is benzene extraordinarily stable although it contains three double bonds?
- 23. Write chemical reaction to exemplify Friedel-Crafts alkylation of benzene.
- 24. What is the nature of substitution in benzene?
- 25. Why $\sqrt{2}$ is not aromatic?
- 26. C-C bond length in benzene is intermediate between C-C and C=C. Why?
- 27. Starting from benzene, how would you synthesize m- Bromonitrobenzene.
- 28. Give one example each of o, p-directing group and m-directing group.
- 29. Complete the reaction:

 $C_6H_6 + CH_3COC1$ Anhy. AlCl₃

30. Write the electrophile which is involved in the nitration of benzene.





2-MARKS QUESTIONS

- 1. What effect does branching have on the boiling point of an alkane and why?
- 2. What is the difference between isomers and conformers?
- 3. Is it possible to isolate pure staggered ethane or pure eclipsed ethane at room temperature? Give reason.
- 4. Draw Newman projection formula for conformations of ethane.
- 5. How will you convert methyl bromide to ethane?
- 6. Wurtz reaction cannot be used for the preparation of unsymmetrical alkanes? Give reason.
- 7. How can ethene be prepared from (i) Ethanol (ii) Ethyl bromide?
- 8. Melting point of cis-But-2-ene is lower than that of trans-But-2-ene. Give reason.
- 9. Draw the structures of cis and trans Hex-2-ene.
- 10. Explain with the help of equation : Ozonolysis of propene.
- 11. Give a chemical test to distinguish between ethene and ethane.
- 12. What do you understand by peroxide effect (Kharasch effect)?
- 11. What factor determines the stability of alkene?
- 14. Arrange the following alkenes in decreasing order of stability : $CH_3-CH=CH-CH_3, CH_2=CH_2, CH_3-CH=CH_2$
- 15. Complete the reaction:

(i)
$$HBr \rightarrow ?$$

(ii) $HBr \rightarrow ?$
(iii) $HBr \rightarrow ?$
 $H_3COOH ? + ?$

16. An alkene on treatment with HBr in presence of peroxide can generate two types of free radicals $CH_3 - C \cdot - CH_2$ -Br and $CH_3 - C(CH_3) - CH_2 \cdot$

Predict the final product of the reaction and give reason. (Hint: Stability of free radicals)

17. What happens when But-2-ene reacts with acidified potassium permanganate solution?





- 18. You are provided with But-2-yne, how will you convert it into:
 - (i) cis-But-2-ene
 - (ii) trans-But-2-ene
- An alkene C4H8 reacts with HBr both in the presence and absence of peroxide 19. to give the same product. Identify the alkene.
- 20. Arrange ethane, ethene and ethyne in the order of increasing acidity.
- 21. Identify A and B in the following reaction:

A $\xrightarrow{\text{Na}}$ CH=CH $\xrightarrow{\text{Red hot iron tube}}$ B

Write the structures of the products A and B of the following reactions: 22.

(i)
$$HC \equiv CH \xrightarrow{Na} A \xrightarrow{CH_3Br} B$$

- $BrH_2C-CH_2Br \xrightarrow{Alc.KOH} A \xrightarrow{NaNH_2} B$ (ii)
- 23. Give a chemical test to distinguish between ethyne and ethene.
- Out of benzene and toluene ,which will undergo nitration easily and why? 24.
- 25. Why does presence of a nitro group make the benzene ring less reactive in comparison to the unsubstituted benzene ring? Explain.
- 26. What happens when Chlorine is passed through benzene in the presence of sunlight and absence of halogen carrier?

3-MARKS QUESTIONS

- Write the structures and name of products obtained in the reaction of sodium 1. with a mixture of 1-Iodo-2-methylpropane and 2-Iodopropane.
- 2. State Markownikov's rule. Using this rule, write the reaction of propene with (i) HCl & (ii) H_2O .
- Complete the following reactions: 3.

(i)
$$CH_3CH_2Br \xrightarrow{Alc.KOH}$$

- (ii) $CH_3CH = CH_2 + O_3 \xrightarrow{Zn/H_2O}$ (iii) $CH_2 = CH_2 + H_2O + [O] \xrightarrow{Dil.KMnO_4}$
- (i) Write the structure of 3, 4-Dimethylhept-3-ene. 4.
 - (ii) Name the compounds obtained by ozonolysis of 3-Methylpent-2-ene.



Hydrocarbons



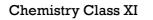
- 5. Complete the following reactions:
 - (i)
 - $CH \equiv CH \xrightarrow{\text{NaNH}_2, CH_3\text{Br}} ?$ $CH \equiv CH \xrightarrow{H_2\text{O}, \text{HgSO}_4/\text{H}_2\text{SO}_4} ?$ (ii)
 - (iii) $CH_3C \equiv CH + H_2 \xrightarrow{Pt} ? \xrightarrow{H_2} ?$
- Write the mechanism of nitration of benzene. 6.
- 7. Arrange in the order of decreasing relative reactivity with an electrophile and explain:

Toluene, p-Nitrotoluene, 1, 4-Dinitrobenzene

- What is meant by delocalization of π electrons? How does it affect stability 8. of benzene?
- What are the conditions for a compound/species to be aromatic according 9. to Huckel's rule?
- How will you convert benzene into 10.
 - Acetophenone (i)
 - (ii) m- Chloronitrobenzene?

5-MARKS QUESTIONS

- Define isomerism. Write all the structural isomers of $hexane(C_6H_{14})$ and 1. arrange them in increasing order of boiling points.
- 2. Write short note on (i) Wurtz reaction (ii) Kolbe's electrolysis (iii) Ozonolysis
- 3. Alkenes show geometrical isomerism while alkanes do not. Give a suitable explanation.
- 4. An alkene 'A' of molecular mass '28u' on treatment with bromine gives a product 'B'. The Compound 'B' on further dehalogenation with zinc gives back 'A'. Give the structures of 'A' and 'B' and also the sequence of reactions.
- 5. An organic compound 'A' with formula C_4H_9Br on treatment with KOH(alc.) gave two isomeric compounds 'B' and 'C' with formula C₄H₈. Ozonolysis of 'B' gave only one product CH3CHO while 'C' gave two different products. Identify A, B and C.
- How will you convert Ethyne into (i) 1, 1, 2, 2-Tetrachloroethane (ii) Ethene 6. (iii) Ethanal (iv) Benzene (v) Sodium ethynide
- Discuss the structure of benzene with an emphasis on resonance and orbital 7. pictures.



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UNIT TEST

Maximum Marks : 20

Time Allowed: 1 hr

General Instructions:

(i) All questions are compulsory.

(ii) Maximum marks carried by each	question are indicated against it.
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1.	Amongst the following which one has the maximum boiling point	[1]
	and why? n-Pentane, iso-pentane, neo-pentane	

- 2. What is the number of σ and π bonds in But-3-en-1-yne? [1]
- Action of acetylene on dilute H₂SO₄/dil.HgSO₄ gives: [1]
 (a) Acetic acid (b) Acetone (c) Acetaldehyde (d) Ethyl alcohol
- 4. An alkene A on ozonolysis gives a mixture of ethanal and [1] pentan-3-one. Write IUPAC name of element.
- 5. The nitro group in the benzene nucleus is _____ directing. [1] It the reactivity of the benzene ring.
- 6. Arrange the following alkenes in decreasing order of stability and [1] give reason.

 $CH_3-CH=CH-CH_3$, $CH_2=CH_2$, $CH_3-CH=CH_2$

- 7. (i) Give a chemical test to distinguish between ethyne and ethene. [2]
 - (ii) Melting point of cis-But-2-ene is lower than that of trans-But-2-ene. Give reason.
- 8. Complete the following reactions: [3] (i) $CH_3CH_2Br \xrightarrow{Alc.KOH}$ (ii) $CH_3CH=CH_2+O_2 \xrightarrow{Zn/H_2O}$

(iii)
$$CH_2 = CH_2 + H_2O + [O]$$
 $\xrightarrow{Dil.KMnO_4}$

- 9. (i) What are the conditions for a compound/species to be aromatic [3] according to Huckel's rule?
 - (ii) How will you convert Benzene to acetophenone?
- 10. (i) An alkene 'A' of molecular mass '28u' on treatment with [5] bromine gives a product 'B'. The Compound 'B' on further dehalogenation with zinc gives back 'A'. Give the structures of 'A' and 'B' and also the sequence of reactions.
 - (ii) Why is benzene extraordinarily stable although it contains three double bonds?
 - (iii) How can we convert ethyne into benzene?

Hydrocarbons



FAST TRACK : QUICK REVISION

- Environmental pollution: Environmental pollution is the effect of undesirable changes in our surroundings that have harmful effects on plants, animals and human beings. A substance which causes pollution is known as pollutant. Pollutants can be degradable and non-degradable.
- Atmospheric pollution: Any undesirable changes in their atmosphere which adversely effect living beings is called air pollution. Air pollution is generally limited to troposphere and stratosphere.

Ozone is present in stratosphere and prevents UV radiations of sun from reaching the earth's surface.

Tropospheric pollution: It is due to gaseous and particulate pollutants.

1. Gaseous air pollutants:

Oxides of sulphur: Major sources of oxides of sulphur (mainly SO_2) are burning of fossil fuels containing sulphur. Sulphur dioxide is converted to sulphur trioxide in presence of particulat matter.

 $2SO_2 + O_2 \longrightarrow 2SO_3$

Sulphur dioxide is a corrosive gas which produces acid rain that causes damage and destruction of vegetation and degradation of soils, building materials and watercourses. SO_2 in ambient air is also associated with asthma and chronic bronchitis. It also causes irritation to eyes.

• **Oxides of nitrogen:** Major sources of nitrogen oxides are high temperature combustion processes, oxidation of nitrogen in the air and fuel respectively, denitrifying bacteria, etc.

 $N_{2} + O_{2} \longrightarrow 2NO$ $2NO + O_{2} \longrightarrow 2NO_{2}$ $NO + O_{3} \longrightarrow NO_{2} + O_{2}$

Finally, these gases are converted into nitric acid (HNO₃) which comes down to the surface of the earth in the form of acid rain.

 $-NO_2$ is a respiratory irritant,

- The oxides produce eye irritation, injury to liver and kidneys.
- **Hydrocarbons:** They are majorly produced naturally (e.g. marsh gas) as well as due to incomplete combustion of automobile fuel.

- Hydrocarbons are carcinogenic, these harm plants.

- **Oxides of carbon:** two major pollutants are oxides of carbon i.e., carbon monoxide and carbon dioxide.
- Carbon monoxide: Carbon monoxide (CO) is a toxic gas which is emitted into atmosphere by incomplete combustion of coal and firewood and by oxidation of hydrocarbons and other organic compounds.

CO may reduce the oxygen carrying capacity of the blood by combining with haemoglobin to produce carboxyhaemoglobin, This oxygen deficiency results in headache weak eyesight, choking and cardiovascular disorders.

- **Carbon dioxide:** CO₂ is released into atmosphere by respiration burning of fossil fuels, forest fire decomposition of limestone in cement industry, etc.
 - It is a greenhouse gas, the concentration of which is constantly raising.
 - In excess it causes headache and nausea.
- Greenhouse effect and global warming: The greenhouse effect is the process in which the emission of infrared radiation by the atmosphere warms the Earth surface.
 - Greenhouse gases include carbon dioxide, methane, ozone chlorofluorocarbons (CFCs) and water vapour.
 - Earth absorbs energy from sunlight entering the atmosphere and emit energy out to space in form of infrared rays. The outgoing radiation emitted by the surface is in the absorption range of many atmospheric gases, including carbon dioxide, methane, and water vapour. These radiations are thus locked in the earth's atmosphere. This results in the steady increase in the temperature of the earth resulting in global warming.
- Acid rain: Rainwater normally has a pH of 5.6 due to dissolution of CO₂ present in the atmosphere.

$$CO_2 + H_2O = H_2CO_3 = H^+ + HCO_3^-$$



Environmental Chemistry

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• When the pH Falls below 5.6, the rain water becomes acidic, It is caused due to presence of acidic gases into the atmosphere the common ones are sulphur dioxide and nitrogen oxides which are changed into sulphuric acid and nitric acid by combining with oxygen and water.

$$2SO_2 + O_2 + 2H_2O \longrightarrow 2H_2SO_4$$
$$4NO_2 + O_2 + 2H_2O \longrightarrow 4HNO_3$$

• Harmful effects of acid rain: It is causes extensive damage to buildings and statues made by marble, limestone due to the reaction,

 $CaCO_3 + H_2SO_4 \longrightarrow CaSO_4 + CO_2 + H_2O$

- It is toxic to vegetation and aquatic life.

— It corrodes water pipes resulting in the leaching of the heavy metals such as Fe, Pb, and Cu into the drinking water which have toxic effects.

2. **Particulate pollutants:** Particulate pollutants are small solid particles and liquid droplets suspended in air.

Smoke: it consists of solid or mixture of solid and liquid particles formed by combustion of organic matter. e.g., cigarette smoke, oil smoke, smoke from fossil fuel etc.

Dust: It consists of fine particles produced during crushing and grinding of solid materials. Common dust particulate emission include cement, fly ash, silica dust, from Industries, dust storm, ground limestone, etc.

Mist: These are formed by particles of spray liquids and condensation of sepals in air. For example, sulphuric acid, mist herbicide or insecticide that Miss their targets and travel through air from mist.

Fumes: These are produced by condensation of vapours. For example, metal fumes, metallurgical fumes and alkali fumes.

Smog: The word smog has its origin from smoke and fog. It is a major air pollutant.

Classical smog	Photochemical smog	
Also called as London smog.	Also called as Los Angeles smog.	
Form due to oxides of sulphur.	Formed due to oxides of Nitrogen.	
Contains primary pollutants.	Content secondary pollutants.	
Causes bronchitis and problem in lungs.	Causes irritation in Eyes.	
It is reducing in nature.	It is oxidising in nature.	





Formation of photochemical smog: It is formed through sequence of following reactions:

$$N_{2} + O_{2} \longrightarrow 2NO$$

(in gasoline)
$$2NO + O_{2} \longrightarrow 2NO_{2}$$
$$NO_{2} \longrightarrow NO + O$$
$$O + O_{2} \longrightarrow O_{3}$$
$$O_{3} + NO \longrightarrow NO_{2} + O_{2}$$

Ozone reacts with hydrocarbons to form Peroxyacetyl nitrate (PAN), formaldehyde, acrolein etc. Effects of photochemical smog:

Ozone and PAN are eye irritants. Photochemical smog also corrodes metals stones buildings materials rubber and painted surfaces.

Measures to control photochemical smog:

Catalytic converters are used in automobiles, which prevent the release of Nitrogen oxide and hydrocarbon to the atmosphere. Certain plants e.g. Pinus, Juniparus, Quercus, Pyrus and Vitis can metabolize nitrogen oxide and therefore their plantation could help in this matter.

► Stratospheric pollution: The upper stratosphere consists of considerable amount of ozone (O₃) which protects us from the harmful ultraviolet (UV) radiations coming from the sun. These radiations causes skin cancer in humans.

Depletion of Ozone layer : Nitric Oxide and chlorofluorocarbons are found to be most responsible for depletion of Ozone layer.

$$NO + O_{3} \longrightarrow NO_{2} + O_{2}$$

$$O_{2} \longrightarrow O + O$$

$$NO_{2} + O \longrightarrow NO + O_{2}$$

$$CF_{2}Cl_{2} \xrightarrow{hv} \cdot CF_{2}Cl + Cl^{*}$$
(free radical)
$$CFCl_{3} \xrightarrow{hv} CFCl_{2} + Cl^{*}$$

$$Cl^{*} + O_{3} \longrightarrow ClO^{*} + O_{2}$$

$$ClO^{*} + O \longrightarrow Cl + O_{2}$$

Environmental Chemistry

• Ozone depletion by oxides of Nitrogen:

Nitrous oxide is quite inert, in the stratosphere it is photochemically converted into more reactive nitric oxide.

$$NO + O_{3} \longrightarrow NO_{2} + O_{2}$$
$$O_{2} \xrightarrow{hv} O + O$$
$$NO_{2} + O \longrightarrow NO + O_{2}$$

Thus NO is regenerated in chain reaction.

• Effects of ozone depletion:

With the depletion of Ozone layer, more UV radiation filters into troposphere.

— UV radiations leads to ageing of skin, cataract, Sunburn, skin cancer, killing of many phyotplanktons, damage to fish productivity, etc.

Water pollution: Water pollution may be defined as any change in its physical, chemical, biological properties or contamination with foreign materials that can adversely affect human beings or reduce its utility for the intended use.

Pollutants	Major sources	
natural waste	Leaching of minerals slits from soil erosion falling of organic matter from bank,etc.	
organic Chemicals	Pesticides, surfactants, detergents, Industrial waste	
metals	Nuclear power plants, mining, metal plating industries.	
man-made wastes	Sewage domestic waste, soaps and detergents, waste from animal sheds and slaughter houses, run off from agricultural fields, industrial wastes, oil pollution.	

Major water pollutants and their sources:

• **Eutrophication:** The process in which nutrient enriched water bodies support a dense plant population, which kills animal life by depriving it of Oxygen and results in subsequent loss of biodiversity is known as eutrophication.





• **BOD:** The amount of oxygen required by bacteria to break down the organic matter present in a certain volume of a sample of water, is called Biochemical Oxygen Demand (BOD)

Element	Permissible limit	Effect of excess amount
Fluoride	1 ppm or 1mg dm ⁻³	Over 10 ppm causes harmful effect to bones and teeth.
Lead	50ppb	Excess amount can damage kidney, liver reproductive system, etc.
Nitrate	50ppm	Excess amount can cause blue baby syndrome.
Sulphate	500ppm	Excess can cause laxative effects.

International standards for drinking water:

► Soil pollution: Soil pollution is the addition of such chemical substances (in an indefinite proportion) which deteriorates the quality, texture and mineral content of the soil and disturbs the biological balance of the organisms in it and has little effect on the plant growth.

Some major soil pollutants and their sources:

Pollutants	Major sources	
Industrial wastes	Waste products from paper, sugar, chemical Industries dumped into the soil.	
Agricultural wastes	Chemical such as fertilizers pesticides extra used for killing insects fungi and weeds.	
Soil conditioners	Used to protect soil fertility but contains several toxic metals like Pb, As, Hg, Cd, etc	
Farm wastes	Wet slurry, faecal wastes are seeped into the soil.	
Radioactive pollutants	Dumping of nuclear waste into the soil.	

Remedial measures:

- Forestation should be done to check the spread of desert.
 - Use of chemical fertilizers should be minimised.
 - Recycling, digestion and incineration of urban waste and organic waste should be done.
 - The industrial effluents should not be allowed to discharge into fields.
 - The fertility of the soil can be improved by soil rotation and mixed farming.

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• Control of environmental pollution:

- Waste management: Environmental pollution can be controlled to a certain extent by managing the waste disposal in a proper way.
- **Recycling:** A large amount of disposed waste materials can be reduced by recycling the waste.

Thus, it reduces the landfill and converts waste into usable products.

Green chemistry: Green Chemistry may be defined as a strategy to design chemical process and products that reduces or eliminate the use and generation of hazardous substances.

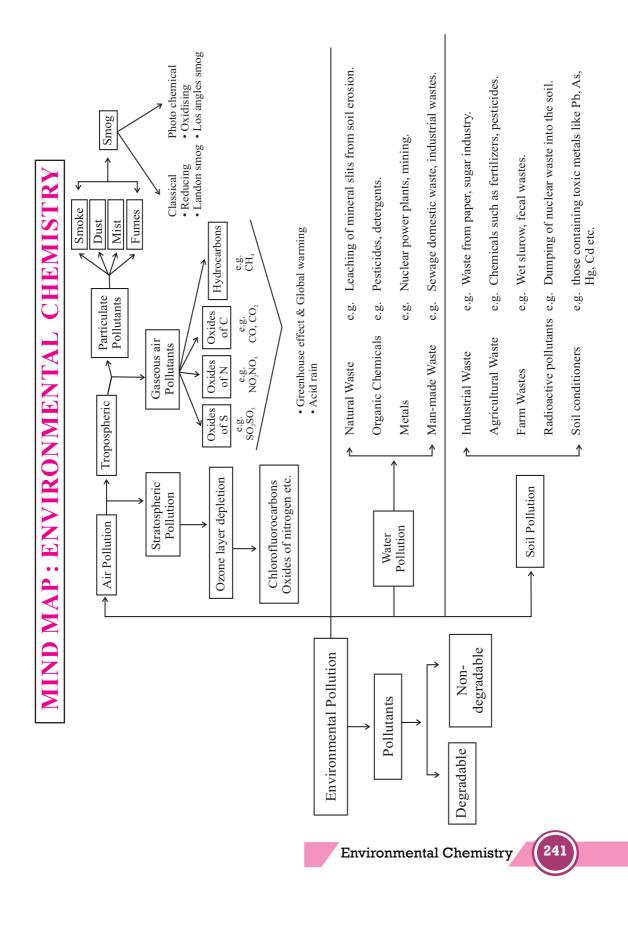
- It is an alternative tool for reducing pollution. Green Chemistry includes concepts such as waste minimization, solvent selection, atom utilisation, intensive processing and alternative synthetic routes from sustainable resources.
- Green chemistry in day-to-day life:
 - Dry cleaning of clothes: Tetra chloroethene $(Cl_2C = CCl_2)$ was earlier used as solvent for dry cleaning. The compound contaminates the ground water and is also a suspected carcinogen.

The process using this compound is now being replaced by a process, where liquefied carbon dioxide, with suitable present is used.

 $\begin{array}{ll} -- & \mbox{Bleaching of paper: Chlorine gas was used earlier for bleaching paper.} \\ & \mbox{These days, hydrogen peroxide (H_2O_2) with suitable catalyst, which} \\ & \mbox{promotes the bleaching action of hydrogen peroxide, is used.} \end{array}$







MULTIPLE CHOICE QUESTIONS (MCQ)

- 1. Which of the following is not the consequence of global warming?
 - (a) Increase in average temperature of Earth.
 - (b) Melting of Himalayan glaciers
 - (c) Rise in sea level
 - (d) Eutrophication

2. Which of the following statement is incorrect?

- (a) Oxidize of Nitrogen in atmosphere can cause depletion of Ozone layer
- (b) Ozone absorbs infrared rays
- (c) Depletion of O_2 is due to its chemical reaction with halo alkanes
- (d) None of these
- 3. Which of the following are the hazardous pollutants present in automobile exhaust gases?
 - (i) N_2 (ii) CO (iii) CH_4 (iv) Oxides of nitrogen
 - (a) ii and iii (b) i and ii
 - (c) ii and iv (d) i and iii

4. Which of the following gas causes green house effect to maximum extent?

- (a) CH₄ (b) Water vapour
- (c) N_2O (d) CO_2
- 5. The lowest region of atmosphere is?
 - (a) Stratosphere (b) Troposphere
 - (c) Mesosphere (d) Hydrosphere
- 6. Classical smog occurs in place of
 - (a) Excess of NO₂ (b) Warm dry climate
 - (c) Cool humid climate (d) All of these
- 7. When huge amount of sewage is dumped in a river, the BOD
 - (a) Will increase (b) Will remain unchanged
 - (c) Will in decrease (d) May increase or decrease
- 8. Which of the following practices will not come under Green chemistry?
 - (a) Use of CO_2 as solvent instead of Cl_2
 - (b) Use of H_2O_2 instead of Cl_2 for bleaching
 - (c) Synthesis of ethanal from ethane in one step
 - (d) Use of tetrachloroethene as a solvent for dry cleaning



- 9. Eutrophication causes reduction in
 - (a) Dissolved salts (b) Dissolved oxygen
 - (c) Nutrients (d) All of these

10. SO_2 and NO_2 cause pollution by increasing

- (a) Acidity (b) Alkalinity
- (c) Buffer action (d) Both (a) and (c)

ANSWERS: 1.d 2.b 3.c 4.d 5.b 6.c 7.a 8.d 9.d 10.a.

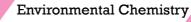
FILL IN THE BLANKS

- 1. Three substances normally considered as primary pollutant are _____, ____, and _____.
- 2. Sulphur dioxide is mainly produced by the burning of _____.
- 3. BOD stands for _____.
- 4. The single plant nutrient mainly responsible for eutrophication is _____.
- 5. The lowest layer of atmosphere is the _____.
- 6. A crucially important species formed by oxygen in stratosphere is _____.
- 7. _____ is a greenhouse gas.
- 8. A major class of organic halides that are thought to pose a threat to stratospheric ozone are _____.
- 9. The component that distinguishes classical smog from photochemical smog is _____.
- 10. Carbon dioxide traps heat in the atmosphere. This is called the _____.

Answers : 1. CO, NO_2 , SO_2 2. Coal 3. Biochemical oxygen demand 4. Phosphate 5. Troposphere 6. Ozone 7. Carbon dioxide 8. Chlorofluorocarbons 9. Sulphur dioxide 10. Greenhouse effect

TRUE AND FALSE TYPE QUESTIONS

- 1. The troposphere is the region above the stratosphere.
- 2. Dust, mist, fumes, smoke and smog are particulate pollutants.
- 3. Carboxyhaemoglobin is less stable than oxyhaemoglobin.
- 4. Ozone occurs naturally in the troposphere.



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- 5. Plants such as Pinus, Juniparus, Pyrus and Vitis metabolize nitrogen oxides.
- 6. Clean water should have BOD of 17 ppm or more.
- 7. Photochemical smog is a mixture of oxidising pollutants.
- 8. The exhaust from jet aeroplanes contains nitric oxide, which can destroy the ozone layer.
- 9. PAN is one of the constituents of photochemical smog.
- 10. DDT is non-biodegradable and persistent.

Answers: 1. F 2. T 3. F 4. F 5. T 6. F 7. T 8. T 9. T 10. T

MATCH THE COLUMNS

Match the column:

1.

Column I	Column II
a. CO ₂	i. a gas produced by the partial combustion of many fuels
b. CO	ii. a gas that occurs naturally in the atmosphere and which is needed for photosynthesis
c. O ₃	iii. a dangerous gaseous pollutant with corrosive properties formed by combustion of fossil fuels.
d. SO ₂	iv. an allotrope of oxygen found in the upper atmosphere

2. Match the pollutants (a, b, c, d) in column I with the effects (i, ii, iii, iv) in column II.

Column I		Column II	
a. Phosphate fertil	isers in water	i. BOD level of water increases	
b. Methane in air		ii. Acid rain	
c. PAN		iii. Global warming	
d. Nitrogen oxides	s in air	iv. Photochemical oxidant	
ANSWERS: 1. a. ii	b. i c. iv	d. iii	
2. a. i	b. iii c. iv	d. ii	





ASSERTION-REASON TYPE QUESTIONS

In the following questions a statement of Assertion (A) followed by reason(R) is given. Use the following key to select correct answer.

- (a) Both Assertion and Reason are correct and Reason is the correct explanation of Assertion.
- (b) Both Assertion and Reason are correct but Reason is not the correct explanation of Assertion.
- (c) Both Assertion and Reason are incorrect.
- (d) Assertion is not correct but Reason is correct.

1. **Assertion:** The pH of acid rain is less than 5.6.

Reason: Carbon dioxide present in the atmosphere dissolves in rain water and forms carbonic acid.

- Assertion: Photochemical smog is oxidizing in nature.
 Reason : Photochemical smog contains NO₂ and O₃, which are formed during the sequence of reactions.
- 3. **Assertion:** If BOD level of water in a reservoir is less than 5ppm, it is highly polluted.

Reason: High biological oxygen demand means low activity of bacteria in water.

- Assertion: Ozone is destroyed by solar radiation in upper atmosphere.
 Reason: Thinning of the ozone layer allows excessive UV radiations to reach the surface of earth.
- 5. Assertion: Carbondioxide is an of the important greenhouse gas.

Reason: It is largely produced by respiratory function of animals and plants.

ANSWERS : 1. b 2. a 3. c 4. d 5. b

ONE WORD TYPE QUESTIONS

- 1. Name the gas which reacts with haemoglobin in blood.
- 2. In which part of atmosphere is ozone layer present?
- 3. Besides CO_2 name one other green house gas.
- 4. Which acid is most abundant in acid rain?
- 5. Name the gas whose release was responsible for Bhopal gas tragedy.

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- 6. What is the nature of photochemical smog?
- 7. Write full form of BOD.
- 8. What is the lowest region of the atmosphere which extends upto a height of 10 km from sea level?
- 9. Name the gas which is produced by the incomplete combustion of gasoline.
- 10. What does CFC stand for?

1-MARK QUESTIONS

- 1. Name two gaseous pollutants.
- 2. What is the size range of particulates?
- 3. What are primary pollutants?
- 4. Name two greenhouse gases.
- 5. What is the composition of photochemical smog?
- 6. What is the composition of classical smog?
- 7. Which are the compounds responsible for ozone layerdepletion?
- 8. Name the pollutants which has affected Taj Mahal.
- 9. Name two strong acids present in acid rain.
- 10. What is PAN?
- 11. Name two sources of phosphate pollution.
- 12. When a huge amount of sewage is dumped into a river, what will be the effect on BOD?
- 13. How is NO formed in atmosphere?
- 14. In which season the depletion of ozone on Antarctica takes place and when is it replenished?
- 15. Define acid rain.

2-MARKS QUESTIONS

1. Explain giving reason: The presence of CO reduces the amount of haemoglobin available in the blood for carrying oxygen to body cells.





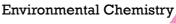
- 2. Oxygen plays a key role in the troposphere while ozone in stratosphere. Explain.
- 3. What is meant by Eutrophication?
- 4. Define BOD.
- 5. What is the importance of measuring BOD of a water body?
- 6. Why does water with excessive algae growth become polluted?
- 7. A person was using water supplied by municipality. Due to shortage of water, he started using underground water, he felt laxative effect. What could be the cause?
- 8. What do you understand by the term Green chemistry?
- 9. Give two examples each of biodegradable and non-biodegradable waste.
- 10. List few ways to control photochemical smog.

3-MARKS QUESTIONS

- 1. Give three harmful effects of oxides of sulphur.
- 2. What is Global warming? What is its cause?
- 3. What is photochemical smog and what are its harmful effects?
- 4. How is classical smog different from photochemical smog?
- 5. State briefly the reactions causing ozone layer depletion in the stratosphere.
- 6. How does rain water get contaminated with acidic impurities?
- 7. Discuss the harmful effects of acid rain.
- 8. Suggest any four methods for waste management.

5-MARKS QUESTIONS

- 1. What is Green house effect? How is it responsible for global warming?
- 2. How can you apply green chemistry in for the following?
 - (i) To control photochemical smog
 - (ii) To avoid use of halogenated solvents in dry cleaning
 - (iii) To reduce the use of detergents
 - (iv) To reduce the consumption of petrol and diesel.



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UNIT TEST

Time Allowed: 1 hr

Maximum Marks : 20

General Instructions:

(i) All questions are compulsory.

(ii) Maximum marks carried by each question are indicated against it.

1.	(a) S	lowest region of atmosphere is Stratosphere Mesosphere	(b) Troposphere(d) Hydrosphere	[1]
2.	(a) l	sical smog occurs in place of Excess of NO ₂ Cool humid climate	(b) Warm dry climate(d) All of these	[1]
3.	Nam	e two Greenhouse gases.		[1]
4.		ajor class of organic halides that a ratospheric ozone are	re thought to pose a threat	[1]
5.	· · · · · · · · · · · · · · · · · · ·			[1]
6.		ain giving reason: The presence of noglobin available in the blood fo		[2]
7.	How	v is classical smog different from j	photochemical smog?	[2]
8.		ne BOD. What is the importance body?	of measuring BOD of a	[3]
9.		v does rain water get contaminated e one harmful effect of acid rain.	l with acidic impurities?	[3]
10.	How	v can you apply green chemistry f	or the following?	[5]
	(i)	To control photochemical smog		
	(ii)	To avoid use of halogenated sol	vents in dry cleaning	
	(iii)	To reduce the use of detergents		
	(iv)	To reduce the consumption of p	etrol and diesel.	





SAMPLE PAPER - I SUB : CHEMISTRY CLASS - XI

Time Allowed: 1 hr

Maximum Marks: 70

General Instructions:

- All question are compulsory
- Q 1 to 20 are very short answer objective type and each carries 1 mark each.
- Q 21 to 27 are short answer question and carry 2 marks each.
- Q 28 to 34 are also short answer question and carry 3 marks each.
- Q 35 to 37 are long answer questions and carry 5 marks each.
- There is no overall choice. However internal choice have been provided in some questions.
- Use of log tables if necessary. Use of calculators is not allowed.
- 1. What are the oxygen moles in 0.5 mol of $CaCO_3$?
 - (a) 1 mol (b) 0.2 mol
 - (c) 1.5 mol (d) 3.0 mol
- 2. What is the unit of wave number (v) ?
- 3. The general configuration of 'f' block is
 - (a) $(n-1) f^{1-14} nd^{0-1} ns2$ (b) $(n-1)f^{0-1} nd^2 ns^2$
 - (c) (n-2) f^{1-14} (n-1) d^{0-1} ns² (d) (n-2) f^{1-14} (n-1) d^{0-2} ns⁰⁻¹
- 4. The shape of IBr_{2} is
 - (a) Tetrahedral (b) Planar
 - (c) Linear (d) V-shape
- 5. What is the pressure of the gas in tube



- 6. Arrange the following in order of decreasing boiling point : Be, Mg, Ca, Sr
- 7. $\Delta_f H^0$ for Graphite is _____.
- 8. Nature of NaH is _____.





Q.9 - Q.10 Assertion-Reason Type Questions

Each question contains statements-1 (Assertion) and Statement-2 (Reason) Examine the statements carefully and mark the correct answer according to the instruction given below:

- A. If both the statements are true and statement-2 is the correct explaination of statement-1.
- B. If both the statements are true but statement-2 is not the correct explanation of statement-1.
- C. If statement-1 is true and statement-2 is false.
- D. If statement-1 is false and statement-2 is true.
- 9. Statement-1 For reaction A + B ⇐ C, K = 4 on addition of catalyst K becomes more than 4.

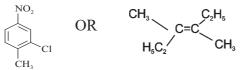
Statement-2 Catalyst only helps to attain the equilibrium faster from either end of reaction.

- Statement-1 Cl₂ + 2OH⁻ → ClO⁻ + Cl⁻ is a disproportion reaction.
 Statement-2 In disproportionation, the same element get oxidised as well as reduce.
- 11. Complete the reaction

$$H_{2}(g) + Pd^{2+} (aq.) \longrightarrow OR$$

$$CO(g) + H_{2}O(g) \xrightarrow{673K} Catalyst}$$

- 12. Why ammoniacal solution of alkali metal is blue in colour?
- 13. What is a producer gas?
- 14. Write the IUPAC name of following



- 15. What does B.O.D. stands for?
- 16. Calculate number of atoms in 52u of He.
- 17. Which series of lines of the hydrogen spectrum lie in the visible region?
- 18. Write the name of element with highest electron gain enthalpy.





- 19. Draw the shape of ClF_3 . OR, Draw the shape of SF_6 .
- 20. What is the unit of 'a' in van der Waal's equation?
- 21. (i) What are the number of waves made by a Bohr electron in an orbit of maximum magnetic quantum number 3?
 - (ii) If kinetic energy of a particle is doubled. What will happen to de Broglie wavelength as compared to previous de Broglie wavelength.
- 22. (i) Why $PbCl_2$ is more stable than $PbCl_4$?
 - (ii) Why Electron gain enthalpy of Mg is positive?

OR

- (i) Second I.E. (Ionisation Enthalpy) is always more than first Ionisation energy.
- (ii) Why first electron gain enthalpy of sulphur is more negative than oxygen.
- 23. Balance the following reaction by (ion-electron or oxidation number method)

 $Cl_2O_7(g) + H_2O_2(aq.) \longrightarrow ClO_2^{-}(aq.) + O_2(g) + H^+(Acidic medium)$

- 24. (i) What is the difference between hydrolysis and hydration?
 - (ii) Arrange the following in order of increasing electrical conductance CaH_2 , BeH_2 and TeH_2 .
- 25. A sample of 0.5g of an organic compound was treated according to Kjeldahl's method. The ammonia evolved was absorbed in 50 ml of 0.5 M H_2SO_4 . The residual acid requirede 60 mL of 0.5 solution of NaOH for neutralisation. Find the percentage composition of nitrogen in the compound.
- 26. (i) Out of benzene, m-dinitrobenzene and toluene, which will undergo nitration most easily and why?
 - (ii) What effect does branching of an alkane chain has on its boiling point?
- 27. (i) What is Eutrophication?
 - (ii) What is the action of F^- on enamel present on the surface of teeth? OR
 - (i) What is the upper limit concentration of lead in drinking water?
 - (ii) What is smog? Classify them as reducing smog or oxidising smog.





- 28. A crystalline salt on being rendered anhydrous loses 45.6% of its weight. The percentage composition of the anhydrous salt is A1 = 10.50%, K= 15.1%, S= 24.96%, O=49.92%
 Find the simplest formula of the anhydrous and crystalline salt. (Atomic Mass : K= 39, A1 = 27, S = 32, O = 16).
 - 29. Explain following with example :
 (i) Aufbau's Rule (ii) Hund's Rule (iii) Pauli's Exclusion Principle OR

An ion with mass number 81 contains 31.7% more neutrons as compared to protons. Assign the atomic symbol.

30. Explain the bonding in SF_6 using hybridisation concept and define what is hybridisation.

OR

On the basis of molecular orbital theory find the bond order, molecular orbital configuration and magnetic nature of O_2^{+} .

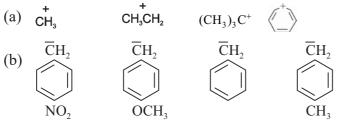
31. Derive van der Waal's equation :

$$\left(P + \frac{an^2}{v^2}\right)(v - nb) = nRT$$

32. Explain Born Haber cycle with by considering example of formation of MgCl₂ as given in the chemical reaction

 $Mg(s) + Cl_2(g) \longrightarrow MgCl_2(s)$

- 33. Give suitable reasons :
 - (i) A solution of Na₂CO₃ is alkaline why?
 - (ii) Beo insoluble but BeSO₄ soluble in water why?
 - (iii) Lithium salts are commonly hydrated as compare to other alkali metal ions. Why?
- 34. Arrange following in order of increasing stability



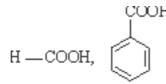




(c) $CH_3\dot{C}H_2$, $\dot{C}H_3$, $\dot{C}H_2$ -CH= CH_2 , $\dot{C}H$ = CH_2

OR

- (i) What is ambident nucleophile? Mention one example.
- (ii) Distinguish between homolytic and hetrolytic bond cleavage.
- (iii) Which one is stronger acid and why



- 35. (a) What is a buffer solution? Give example.
 - (b) What is common ion effect?
 - (c) Define Le-Chatlier principle and explain effect of following :
 - (i) Change of concentration (ii) Change of pressure

OR

- (a) Find out K_c for following reaction 2NOCl(g) \rightleftharpoons 2NO(g) + Cl₂(g); $K_p = 1.8 \times 10^{-4}$ at 500K
- (b) $K_p = 0.04$ atm at 899K. What is the equilibrium concentration of C_2H_6 where it is placed in a flask at 4.0 atm pressure and allow to come to equilibrium

 $C_2H_6 \rightleftharpoons C_2H_4(g) + H_2(g)$

- (c) What is the unit of K_p for the following chemical reaction? $2NH_3(g) \rightleftharpoons N_2(g) + 3H_2(g)$
- 36. (i) Show with the help of chemical reaction that Al shows amphoteric behaviour.
 - (ii) Draw the structure of (a) B_2H_6 (b) Boric acid.
 - (iii) Write the formula of Borax.

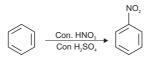
OR

(i) Explain Lewis acid strength $BF_3 < BCl_3 < BBr_3 < BI_3$

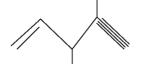




- (ii) What are silicones? Give reaction for formation of chain silicones.
- (iii) Why CO is poisionous in nature?
- 37. (i) Explain with the help of mechanism

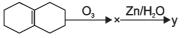


- (ii) Draw the Newman structure of (a) 2-Methyl butane(b) 1-Methyl prop-1-ene
- (iii) Calculate total number of σ and π bond(s) in



OR

- (i) Explain Kolbe's electrolysis with mechanism.
- (ii) State Huckel Rule's.Check whether is an aromatic or non aromatic, anti-aromatic.
- (iii) Write the product







SAMPLE PAPER - II SUB : CHEMISTRY CLASS - XI

Time Allowed: 1 hr

Maximum Marks: 70

General Instructions:

- All question are compulsory
- Q 1 to 20 are very short answer objective type and each carries 1 mark each.
- Q 21 to 27 are short answer question and carry 2 marks each.
- + Q 28 to 34 are also short answer question and carry 3 marks each.
- Q 35 to 37 are long answer questions and carry 5 marks each.
- There is no overall choice. However internal choice have been provided in some questions.
- Use of log tables if necessary. Use of calculators is not allowed.

SECTION-A

1.	The number	of nodal planes in p	x orbitals is	
	(a) 1	(b) 2	(c) 3	(d) 0
2.	Which of the	e following has smal	llest bond angle?	
	(a) H ₂ O	(b) H ₂ S	(c) NH ₃	(d) SO ₂
3.	(a) ΔG -ve (b) ΔG +ve (c) ΔG -ve	on to be spontaneous $\Delta H + ve, \Delta S + ve$ $\Delta H - ve, \Delta S + ve$ $\Delta H - ve, \Delta S - ve$ $\Delta H - ve, \Delta S - ve$ $\Delta H - ve, \Delta S + ve$	at all the temperatur	re:
4.	Which is is 1	most strongly hydrat	ted?	
	(a) Li ⁺	(b) Na ⁺	(c) K ⁺	(d) Rb^+
5.	Which of the	e following has large	est ionic radii?	
	(a) Na ⁺	(b) Mg^{2+}	(c) F ⁻	(d) O ^{2–}
6.	When carbo hybrid orbita		other atoms or gro	ups it uses
7.	Surface tens	ionv	with increase in temp	perature.
8.	The second	electron gain enthal	by is	





Directions for Question No. 9 and 10 : A statement of assertion (A) followed by a statement of reason (R) is given. Choose the correct option out of the choices given below for each question:

- (a) A and R both are correct and R is the correct explanation of A.
- (b) A and R both are correct but R is not the correct explanation of A.
- (c) A is true but R is false.
- (d) A and R both are false.
- 9. Assertion: The entropy of ice is less than that of water. Reason: Ice has a cage like structure.
- Assertion: London forces are much more stronger between Xenon atoms than between Helium atoms.

Reason: Xenon atom is bigger than Helium atom.

11. Write empirical formula of CH₃COOH and K₂CO₃

Or

Define mole fraction.

12. Mention the quantum number which determines the energy of electron in the H-atom.

Or

How many unpaired electrons are there in Ni^{2+} ion? (Given : Z = 28)

13. State the condition for the formation of precipitate.

Or

Write the conjugate acid and conjugate base of HSO₄⁻.

- 14. Using VSEPR theory draw the shape of XeF_4 molecule.
- 15. Write IUPAC name of $CH_2 = CH CH(OH)C \equiv CH$.
- 16. In a reaction between an oxidant and a reductant which will give up electrons and which will accept electrons?

Or

Calculate oxidation number of Cr in K₂Cr₂O₇ and Mn in KMnO₄.

17. Elements of which group form electron precise hydrides?

Or

Which part of periodic table is known as hydride gap?

18. State the reason of using certain alkali metals in photoelectric cells.

Or

Name those alkaline earth metals which do not impart colour to the flame.

- 19. Mention the compounds which are responsible for ozone layer depletion.
- 20. Cis But-2-ene has lower melting point that trans But-2-ene. Give reason.





SECTION: B

21. Balance the following redox reaction in acidic medium by ion electron method.

 $Zn(aq.) + NO_3^{-}(aq.) \longrightarrow Zn^{2+}(aq.) + N_2O(g) + H_2O$

- 22. (i) Mg²⁺ ion is smaller than O2-ion, although both are isoelectronic. Give reason.
 - (ii) Write IUPAC name and symbol for the element with atomic no. 120.
- 23. (i) Mention the number of radial nodes in 6s orbitals.
 - (ii) Write electronic configuration of Fe^{2+} ion. (Given, Z = 26)

Or

Calculate the wavelength of a ball of mass 0.1kg moving with a velocity of 10ms^{-1} . (Given, $h = 6.626 \times 10^{-34} \text{ Js}$)

24. Describe on one method to remove permanent hardness of water.

Or

Give one chemical reaction each to show that hydrogen peroxide can act as oxidising as well as reducing agent.

25. Write a short note on Greenhouse effect and global warming.

Or

State the point of differences between Classical and photochemical smog.

- 26. (i) Stability of carbocations follows the order $3^{\circ} > 2^{\circ} > {}^{\circ}1$. Explain this order of stability of carbocations.
 - (ii) In what manner is Electromeric effect different from Inductive effect?
- 27. (i) Mention the reason of not using Wurtz reaction for the preparation of unsymmetrical alkanes from alkyl halides.
 - (ii) How will you convert Benzene to p-Nitrobromobenzene?

SECTION-C

28. The density of 3M solution of NaCl is 1.25g/mL. Calculate the molality of the solution. (Given: Atomic masses: Na=23u, Cl=35.5u)

Or

Calculate the molarity of nitric acid (HNO_3) in a sample having a density 1.41g/mL and mass percent of nitric acid in it being 69%. (Atomic mass: N=14u, H=1u, O=16u)

Sample Paper



- 29. (i) The ball hit with a hockey by a player does not form a wave. State reason.
 - (ii) Write the possible values of 'm' for an electron with l=2.
 - (iii) Chromium has configuration $3d^54s^1$ and not $3d^44s^2$. Explain.
- 30. (i) Explain non linear shape H_2S and non-planar shape of PCl_3 using VSEPR theory.
 - (ii) Can we have a diatomic molecule with its ground state molecular orbitals full with electrons. Give reason for your answer.
- 31. Calculate enthalpy change for the reaction: $CH_4(g) + 2O_2(g) \longrightarrow CO_2(g) + 2H_2O(l)$ The enthalpy of formation of $CH_4(g)$, $CO_2(g)$ and $H_2O(l)$ are -74.8kJmol⁻¹, -393.5 kJmol⁻¹ and 285.8kJmol⁻¹ respectively.

Or

Calculate the bond enthalpy of Cl-Cl bond from the following data: $CH_4(g) + Cl_2(g) \rightarrow CH_3Cl (l) + HCl \quad \Delta H = -100.3 \text{ kJmol}^{-1}$ Given: bond enthalpies of C — H, C — Cl and H — Cl bonds are 413, 326 and 431 kJmol⁻¹ respectively.

- 32. A neon-dioxygen mixture contains 70.6 g dioxygen and 167.5 g neon. If pressure of the mixture of gases in the cylinder is 25 bar, what is the partial pressure of dioxygen and neon in the mixture? (Atomic mass: O = 16u, Ne = 20u)
- 33. (i) Compounds of beryllium are much more covalent than other group 2 elements. Give reason.
 - (ii) Why does the solubility of alkaline earth metal carbonates and sulphates in water decrease down the group?
 - (iii) When a metal of group 1 is dissolved in liquid ammonia a blue solution is obtained initially. How do you account for the blue colour of the solution?
- 34. (i) State the necessary compound to be aromatic according to Huckel's rule.
 - (iii) Explain why alkyl groups act as electron donors when attached to a π system.

Or

- (i) Draw the resonance structures of Phenol.
- (ii) Suggest a method used to purify the liquids which have high boiling points and decompose below their boiling points.





SECTION-D

- 35. (i) Boron trihalides(BX_3) act as Lewis acids. Why?
 - (ii) Conc. HNO₃ can be transported in aluminium containers. Give reason.
 - (iii) Pb(IV) is less stable tha Pb(II). Give reason.
 - (iv) Gallium has higher ionisation enthalpy than aluminium. Why?
 - (v) What do you understand by diagonal relationship?

Or

- (i) Why CCl_4 is resistant to hydrolysis but $SiCl_4$ is readily hydrolysed?
- (ii) Explain why there is a decrease in ionisation enthalpy from carbon to silicon?
- (iii) Boron does not form B^{3+} ion. Give reason.
- (iv) How can you explain greater stability of BCl₃ as compared to TlCl₃?
- (v) Define diagonal relationship between elements in modern period table.
- 36. (i) Explain common ion effect with example.
 - (ii) The concentration of hydrogen ion in a sample of soft drink is 4 x 10-3 M. Calculate its pH.
 - (iii) What is the effect of removal of CH_3OH on the equilibrium of the reaction, $2H_2(g) + CO(g) \rightleftharpoons CH_3OH(g)$.

Or

- (i) Calculate H_3O^+ ion concentration of a water sample having pH = 6.78.
- (ii) Define buffer solutions.
- (iii) State Lewis definition of acids and bases. Give one example of each.
- (i) An alkene A on ozonolysis gives a mixture of propanal and pentan-3one. Write the structural formula of A.
 - (ii) Complete the following reactions:
 - (a) $CH_3CH=CH_2 + HBr$ organic peroxide
 - (b) $CH_3CH_2Br + Na \longrightarrow$
 - (iii) Write a short note on Friedel Crafts alkylation.

Or

- (i) Why eclipsed form and staggered form of ethane cannot be isolated at room temperature?
- (ii) State Markovnikov's rule.
- (iii) Which out of Ethene or Ethyne is more acidic and why?
- (iv) What happens with 2-Bromobutane is being treated with KOH (alcoholic)?





SAMPLE PAPER - III SUB : CHEMISTRY CLASS - XI

Time Allowed: 1 hr

Maximum Marks : 70

General Instructions:

- All question are compulsory
- Q 1 to 20 are very short answer objective type and each carries 1 mark each.
- Q 21 to 27 are short answer question and carry 2 marks each.
- Q 28 to 34 are also short answer question and carry 3 marks each.
- Q 35 to 37 are long answer questions and carry 5 marks each.
- There is no overall choice. However internal choice have been provided in some questions.
- Use of log tables if necessary. Use of calculators is not allowed.
- 1. Which of the following d-orbital has double disc (body shooter shape) (a) d_{xy} (b) d_{z^2} (c) $d_{x^2-y^2}$ (d) d_{yz}
- 2. Among the following, the compound that contains, ionic, covalent and Coordinate linkage is

(a) NH_4Cl (b) NH_3 (c) NaCl (d) CO_2

- Which of the following is a state function
 (a) q (b) w (c) q + w (d) None of these
- 4. Which of the following alkali metal ion has highest ionic mobility (a) Li^+ (b) Na^+ (c) k^+ (d) Cs^+
- 5. The correct order of electron gain enthalpy with negative sign of F, Cl, Br and I having atomic number 9, 17, 35 and 53 respectively is
 (a) I > Br > Ci > F
 (b) F > Cl > Br > I
 (c) Cl > F > Br > I
 (d) Br > I > Cl > F
- 6. Arrange the following carbonations in order of their increasing stability $(CH_3)_3C^+$, $CH_3CH_2^+$, $(CH_3)_2CH^+$, CH_3^+
- 7. The IUPAC name and symbol of element with Z = 120 is _____.
- 8. Direction for Question No. 9 to 10 : A statement of assertion (A) followed by a statement of Reason (R) is given.
 - (i) A and R both are correct, and R is correct explanation of A.
 - (ii) A and R both are correct, but R is not correct explanation of A.
 - (iii) A is true but R is false.
 - (iv) A and R both are false.

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- 9. Assertion (A) : Enthalpy of graphite is lower than that of diamond. Reason (R) : Enthalpy of graphite is greater than that of diamond.
- 10. Assertion (A) : Lower the critical temperature of the gas; more easily can it be liquified.Reason (R) : Critical temperature is the temperature above which a gas cannot be liquified depending upon the pressure.
- 11. If ten volumes of dihydrogen gas react with five volumes of dioxygen gas, how much volume of water vapour would be produced?

OR

Calculate the number of atoms in 32.0 u of He.

12. Why are alkali metals used in photoelectric cells?

OR

Write electronic configuration of Cr^{3+} ion. (Atomic No. of Cr = 24)

- 13. Draw the shape of ClF₃ molecule according to VSEPR theory?
- 14. Write the conjugate acid and conjugate base of H_2O .

OR

Write the relation between solubility and solubility product of $Cr_2(SO_4)_3$.

15. Consider the given standard reduction potentials of following elements A, B, C & D and arrange them in order of their increasing reducing power.

A = -3.71V, B = -0.76V, C = +0.34V, D = +0.80VOR

- 16. Write the cause of temporary and permanent hardness of water.
- 17. Why do magnesium and beryllium not import colour to the flame in the flame test?
- 18. Write IUPAC name of $CH_3 CO CH_2 CHO$.
- 19. Out of Cis-But-2-ene and Trans-but-2-ene which has greater boiling point and why?
- 20. Define Biochemical oxygen demand.
- 21. (i) What is the lowest value of n that allows 'g' orbitals to exist?
 - (ii) Why 4s orbital is filled before 3d?

OR

Calculate wave number for the longest wavelength transition in the Balmer Series of hydrogen atom.



Sample Paper



- 22. Explain why?
 - (i) $\Delta_i H_1$ of 'N' is more than that of 'O'.
 - (ii) A cation is smaller than parent atom.
- 23. Balance the following redox reaction in acidic medium by ion electron method.

 $\operatorname{Cr}_2O_7^{2-}(\operatorname{aq.}) + \operatorname{SO}_2(g) \longrightarrow \operatorname{Cr}^{3+}(\operatorname{aq.}) + \operatorname{SO}_4^{2-}(\operatorname{aq.})$

24. Write chemical reactions to justify that H_2O_2 can act as an oxidising as well as reducing agent.

OR

- (i) What is water gast shift reaction?
- (ii) What do you mean by 20 vol. H_2O_2 ?
- 25. In sulphur estimate, 0.157 g of an organic empound gave 0.4813 g of barium sulphate. What is the percentage of sulphur in the compound?
- 26. Write chemical equations to covert:
 - (i) Ethyne to Ethanal
 - (ii) Benzene to m-Nitrotoluene
- 27. Define smog. How is classical smog different from photochemical smog?

OR

Explain the following terms:

- (a) Green house effect
- (b) Green chemistry
- 28. Calcium carbonate reacts with aqueous HCl according to the reaction: CaCO₃(s) + 2HCl(aq.) → CaCl₂(aq.) + CO₂(g) + H₂O(l) What mass of CaCO3 is required to react completely with 25 mL of 0.75 M HCl?
- 29. (i) State Hund's rule of maximum multiplicity of electrons.
 - (ii) The mass of an electron is 9.1×10^{-31} kg. If its K.E. 3.0×10^{-25} J, calculate its wavelength.
- 30. N_2 is diamagnetic while O_2 is paramagnetic. Explain on the basis of molecular orbital theory.

OR

Explain the structure of PCl_5 according to hybridization. Why all P — Cl bonds.





31. What will be the pressure exerted by a mixture of 3.2 g of methane and 4.4 g of carbon dioxide contained in 9 dm² flask at 27°C.

OR

Calculate the compressibility factor for CO_2 , if one mole of it occupies 0.4 litre at 300K and 40 atm. Comment on the result.

32. The combustion of 1 mol of benzene takes place at 298K. After combustion CO_2 and H_2O are formed and 3267 kJ mol⁻¹, $\Delta_f H^{\Theta}(H_2O) = -393$ kJ mol⁻¹.

OR

For the reaction : $2A(g) + B(g) \longrightarrow 2D(g)$, $\Delta U^{\Theta} = -10.5 \text{ kJ}$ and $\Delta S^{\circ} = -44.1 \text{ J K}^{-1}$. Calculate ΔG° for the reaction, and predict whether the reaction will occur spontaneously.

- 33. Complete the following reaction equations:
 - (i) $\text{Li}(\text{NO}_3)_2 \xrightarrow{\Delta}$
 - (ii) $\operatorname{BeCl}_2 + \operatorname{LiAlH}_4 \xrightarrow{\Delta}$
 - (iii) $Ca(OH)_2 + Cl_2 \xrightarrow{\Delta}$
- 34. (i) Out of $NO_2 CH_2 O$ and $CH_3 CH_2 O^-$ which is more stable and why?
 - (ii) Why is it necessary to prepare Lassaigne extract for detection of N, S and halogens?
 - (iii) Define the term hyperconjugation.
- 35. (i) State Le Chatelier's principle.
 - (ii) Calculate the pH of 10^{-8} M HCl.
 - (iii) A sample of HI (g) is placed in flask at a pressure of 0.2 atm. At equalibrium the partial pressure of HI (g) is 0.04 atm. What is the Kp for given equilibrium?

2HI (g) \rightleftharpoons H₂(g) + I₂(g)

OR

- (i) Define common ion effect.
- (ii) Write the relationship between K_p and K_c for the reaction:

 $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$

(iii) Equal volume of 0.002 M solutions of sodium iodate and cupric chlorate are mixed together. Will it lead to precipitation of copper iodate? (For cupric iodate $K_{sp} = 7.4 \times 10^{-8}$)





- 36. (i) Draw the structure of Diborane indicating electron precise and electron deficient bonds.
 - (ii) Give reason for the following :
 - (a) $PbCl_4$ is powerful oxidising agent.
 - (b) Fullerenes are the purest allotropes of carbon.
 - (c) N N bond dissociation enthalpy is less than P P bond dissociation enthalpy.

OR

- (i) What happens when (give chemical equation) :
 - (a) Borax is heated strongly.
 - (b) BF_3 is treated with LiH.
- (ii) Give reason for the following :
 - (a) $NaBiO_3$ is good oxidising agent.
 - (b) Graphite acts as a good lubricant.
 - (c) Boric and acid is a weak acid.
- 37. (i) Propanal and pentan-3-one are the ozonolysis product of an alkene. What is the structural formula and IUPAC name of alkene?
 - (ii) Give the main products of the reactions:

(a)
$$C_6H_6 + CH_3Cl$$
 Anhydrous AlCl₃

- (b) $CH_3Cl + Na$ Dry ether
- (c) $CH_3 CHCl CH_2 CH_3 + KOH(alc.) \rightarrow$

OR

- (i) Give chemical euqation for each of the following :
 - (a) Decarboxylation
 - (b) Friedel Craft acetylation
- (ii) Addition of HBr to propene yields 2-Bromopropane, while in the presence of benzoyl peroxide, the same reaction yields 1-Bromopropane. Explain and give mechanism.





SAMPLE PAPER - IV SUB : CHEMISTRY CLASS - XI

Time Allowed: 1 hr

Maximum Marks: 70

General Instructions:

- All question are compulsory
- Q 1 to 20 are very short answer objective type and each carries 1 mark each.
- Q 21 to 27 are short answer question and carry 2 marks each.
- Q 28 to 34 are also short answer question and carry 3 marks each.
- Q 35 to 37 are long answer questions and carry 5 marks each.
- There is no overall choice. However internal choice have been provided in some questions.
- Use of log tables if necessary. Use of calculators is not allowed.

SECTION-A

1. Number of matamers possible for molecular formula C_4H_{10} are _____.

OR

Functional isomer of CH₃CH₂CHO is _____.

- 2. Among the following, the least thermally stable is ? (a) K_2CO_3 (b) Li_2CO_3
 - (c) Na_2CO_3 (d) Rb_2CO_3
- 3. Van der waals constant is a maximum for
 - (a) He (b) Ne
 - (c) Ar (d) Kr
- 4. Orbital which is not possible
 - (a) 2p (b) 3d (c) 3s (d) 3f
- 5. Which of the following is paramagnetic
 - (a) CO (b) O_2^-
 - (c) N_2 (d) NO^+
- 6. Which of the following is not correct?
 - (a) ΔG is zero for a reversible reaction.
 - (b) ΔG is positive for a spontaneous reaction.
 - (c) ΔG is negative for a spontaneous reaction.
 - (d) ΔG is positive for a non-spontaneous reaction.



Sample Paper



Direction for Question No. 7 and 8 :

- (i) If both assertion & reason are true and the reason is the correct explanation of the assertion.
- (ii) If both assertion & reason are true but the reason is the correct explanation of the assertion.
- (iii) If assertion is true statement but reason is false.
- (iv) If both assertion and reason are false statements.
- 7. Assertion : Number of orbitals in 3rd shell is 9. Reason : Number of orbitals for a particular value of $n = n^2$.
- Assertion : Ionic radius of Na⁺ is smaller than Na.
 Reason : Effective nuclear charge of Na⁺ is higher than Na.
- 9. What is the IUPAC name of picric acid.
- 10. Why noble gases have positive electron gain enthalpy?
- 11. How many molecules of SO_2 are present in 11.2 L at STP?

OR

Calculate the number of atoms in 35 g of Li (Atomic mass of Li = 7 u).

12. For an isolated system, $\Delta U = 0$, what will be ΔS ? OR

Write the statement of third law of thermodynamics.

- 13. Calculate the number of sigma and pie bonds in C_2H_2 ?
- 14. Give the relation between K_a and K_b .

OR

Write the conjugate base of H_2CO_3 .

15. Write the oxidation number of Cr in $K_2Cr_2O_7$. OR

Write the stock notation MnO_2 .

16. Give an example of electron deficient covalent hydride.

OR

Name the isotope of hydrogen which is radioactive in nature.

- 17. Potassium carbonate cannot be prepared by solvay process. Why?
- 18. Define the Boyle temperature or Boyle point.
- 19. Out of staggered and eclipsed form which is more stable and why?
- 20. What is the meaning of the term eutrophication with regards to water pollution.

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SECTION-B

- 21. Explain why :
 - (a) The three electron present in 2p subshell of nitrogen remain upaired.
 - (b) Cr has configuration $3d^5 4s^1$ and not $3d^4 4s^2$.

OR

Calculate the radius of Bohr's fifth orbit for hydrogen atom.

- 22. (a) Show by a chemical reaction with water that Na_2O is a basic oxide and Cl_2O_7 is an acidic oxide.
 - (b) Name a species that will be isoelectronic with each of the following atoms or ions, (i) F^- (ii) Ca^{2+}
- 23. Balance the following redox reaction in acidic medium $MnO_4^- + H^+ + Fe^{2+} \longrightarrow Mn^{2+} + H_2O + Fe^{3+}.$
- 24. What do you understand by the terms :
 - (i) Syn gas
 - (ii) Hydrogen economy

OR

What causes the temporary and permanent harness of water?

- 25. 0.40 g of an organic compound gave 0.3 g of silver bromide of Carius method. Find the percentage of bromine in the compound. [Atomic mass of Ag & Br are 105 and 80 u]
- 26. Propanal and pentan-3-one ate the ozonolysis product of an alkene. What is the structural formula of alkene?

OR

Convert the following :

- (a) 1-Bromopropane to 2-Bromopropane
- (b) Ethanol to Glycol
- 27. Explain the following terms :
 - (a) Acid Rain (b) Global warming

OR

What are the reactions involved for ozone layer depletion in the stratosphere?

- (a) Green house effect (b) Green chemistry
- 28. In a compound $C_x H_y O_z$, the mass % of C and H is 6 : 1 and the amount of oxygen present is equal to the half of the oxygen required to react completely $C_x H_y$. Find the empirical formula of the compound.





- 29. When light with a wavelength of 400 nm falls on the surface of sodium, electrons with a kinetic energy of 1.05×10^5 J mol⁻¹ are emitted.
 - (a) What is the minimum energy needed to remove on electron from sodium?
 - (b) What is the maximum wavelength of light that will cause a photoelectrons to be emitted?
- 30. (a) Describe the hybridisation in case of C_2H_2 .
 - (b) Which out of NH_3 and NF_3 has higher dipole moment and why?

OR

- (a) Use molecular orbital theory to predict why the He₂ molecule does not exist?
- (b) Compare the stability of O_2 and O_2^+ on the basis of molecular theory.
- 31. Pressure of one gram of an ideal gas A at 27°C is found to be 2 bar. When 2g of another gas (ideal) B is introduced in the same flask at the same temperature the pressure become 3 bar. Find a relationship between their molecular masses.

OR

Explain the following terms :

- (a) Dalton's law of partial pressure
- (b) Surface tension
- (c) Coefficient of viscosity
- 32. The standard enthalpy of combustion of Surcose $(C_{12}H_{32}O_{11})$ at 298 K producing CO₂(g) and H₂O(*l*) is -5200.7 kJ mol⁻¹. If $\Delta_f H^\circ$ of CO₂(g) and H₂O(*l*) are -393.0 kJ mol⁻¹ and -286 kJ mol⁻¹.

OR

Explain the First law of thermodynamics and how can you prove the following relation :

 $\Delta H = \Delta U + P \Delta V$ from the first law of the thermodynamics ($\Delta U = q + w$)?

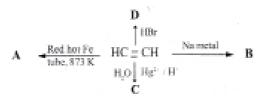
- 33. Assign reason for the following :
 - (a) Potassium carbonate cannot be prepared by solvay process.
 - (b) Be and Mg do not give characteristics colour to the flame.
 - (c) Alkali metals are strong reducing agent.
- 34. Explain the resonance effect and draw the resonance structures of $CH_2 = CH CHO$. Indicate the relative stability of the contributing structures.





SECTION-D

- 35. (a) Addition of HBr to propene yields-2-bromopropane, while in the presence of benzoylperoxide, the same reaction yields 1-bromopropane. Explain and give tis mechanism.
 - (b) Identify A, B, C & D



(c) Give a chemical test between ethen and ethane.

OR

- (a) Out of benzene and toluene, which will undergo nitration more easily and why?
- (b) Explain the following :
 - (i) Friedal Craft reaction
 - (ii) Wurtz reaction
- 36. (a) What is solubility product? How is it different from ionic product?
 - (b) Equal volume of 0.02 M solutions of sodium iodate and cupric chlorate are mixed together. Will it lead to precipitation of copper iodate? (For cupric iodate Ksp = 7.4×10^{-8}).

OR

- (a) Define Le-Chatelier principle.
- (b) Explain ionic product of water. What is the effect of temperature on ionic product of water?
- (c) Calculate the pH of 10^{-10} M NaOH solution.
- 37. (a) Draw the structure of diborane.
 - (b) Explain the following reactions :
 - (i) Silicon is heated with methyl chloride at high temperature in the presence of Cu.
 - (ii) Hydrate alumina is treated with aqueous NaOH solution.

OR

- (a) What are silicons. If the starting material for the manufacture of silicons is RSiCl₃, write the structure of the product formed.
- (b) Complete the following reactions :

$$B_2H_6 + NH_3 \longrightarrow$$
 (ii) $H_3BO_3 + H_2O \longrightarrow$

(c) Explain Inert pair effect.

(i)

