# DIRECTORATE OF EDUCATION <br> Govt. of NCT, Delhi 

## SUPPORT MATERIAL 2023-2024

## Class: XI

## CHEMISTRY

Under the Guidance of

## Shri Ashok Kumar

Secretary (Education)
Shri Himanshu Gupta
Director (Education)

Dr. Rita Sharma
Addl. DE (School \& Exam.)

## Coordinators

| Mr. Sanjay Subhas Kumar | Mrs. Ritu Singhal | Mr. Raj Kumar | Mr. Krishan Kumar |
| :---: | :---: | :---: | :---: |
| DDE (Exam) | OSD (Exam) | OSD (Exam) | OSD (Exam) |

Production Team<br>Anil Kumar Sharma

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अशोक कुमार,भा.प्र.से सचिव ( शिक्षा)
ASHOK KUMAR, IAS
Secretary (Education)


## Message

राष्ट्रीय राजधानी क्षेत्र, दिल्ली सरकार
पुराना सचिवालय, दिल्ली-110054
दूरभाष : 23890187 टेलीफैक्स: 23890119
Government of National Capital Territory of Delhi
Old Secretariat, Delhl-110054
Phone : 23890187, Telefax : 23890119
e-mail : secyedu@nic.In
D.O. NO: :DE.5/228) (Exam/Message/SM

Dated : $24 \cdot 11,2023 \quad / 2018 / 1095$


#### Abstract

"Children are like wet cement, whatever falls on them makes an impression." Haim Ginott


Embracing the essence of this quote, the Directorate of Education, GNCT of Delhi is unwavering in its commitment to its core mission of delivering high-quality education to all its students. With this objective in mind, DoE annually develops support materials meticulously tailored to suit the learning needs of students from classes IX to XII.

Every year, our expert faculty members shoulder the responsibility of consistently reviewing and updating the Support Material to synchronize it with the latest changes introduced by CBSE. This continuous effort is aimed at empowering students with innovative approaches and techniques, fostering their problem-solving skills and critical thinking abilities. I am confident that this year will be no exception, and the Support Material will greatly contribute to our students' academic success.

The support material is the result of unwavering dedication of our team of subject experts. The Support Material has been specially curated for our students, with the belief that its thoughtful and intelligent utilization will undoubtedly elevate the standards of learning and will continue to empower our students to excel in their examinations.

I wish to congratulate the entire team for their invaluable contribution in creating a highly beneficial and practical Support Material for our students.

I extend my best wishes to all our students for a promising and bright future.

(Ashok Kumar)



सत्यमेव जयते

Directorate of Education
Govt. of NCT of Delhi

## Room No. 12, Civil Lines

Near Vidhan Sabha,
Delhi-110054
Ph.: 011-23890172
E-mail: diredu@nic.in

## MESSAGE

It brings me immense pleasure to present the support material for students of classes IX to XII, meticulously crafted by our dedicated subject experts. Directorate of Education is committed to empower educators and students alike by providing these resources free of cost for students of all government and government aided schools of Delhi.

The support material is an appreciable effort to align the content with the latest CBSE patterns. It has been carefully designed as a resource to facilitate the understanding, acquisition and practice of essential skills and competencies outlined in the curriculum.

The core of this support material lies in providing a framework for adopting an analysis-based approach to learning and problem-solving. It aims to prompt educators to reflect on their teaching methodologies and create an interactive pathway between the child and the text.

In the profound words of Dr A.P.J. Abdul Kalam, "Educationists should build the capacities of the spirit of inquiry, creativity, entrepreneurial and moral leadership among students and become their role model."

The journey of education is ongoing; it's the process, not just the outcome, which shapes us. This support material endeavours to be that catalyst of change for eachstudent of Directorate of Education.

Let us embark on this transformative journey together, ensuring that every student feels equipped not only with the knowledge but also, with the skills and mindset to thrive in the 21 st century.

I wish you all the best for all your future endeavours.

Dr. RITA SHARMA
Additional Director of Education (School/Exam)


Govt. of NCT of Delhi Directorate of Education Old Secretariat, Delhi-110054 Ph.: 23890185
D.O. No.DE:5/22.8.|Exam|Merage/sM|

Dated: .... 24.111 .2023

## MESSAGE

The persistent efforts of the Directorate in making the course material more accessible and student-friendly are evident in the conscientious preparation of the Support Material. Our team consistently adapts to the evolving educational landscape, ensuring that the Support Material for the various subjects of classes 9 to 12 align with the latest CBSE guidelines and syllabi prescribed for the annual examinations.

The Support Material encapsulates crucial subject-specific points and facts, tailored to suit the students, all presented in a lucid language. It is our firm belief that these resources will significantly augment the academic prowess of our students, empowering them to excel in their upcoming examinations.
I extend my heartfelt congratulations to the diligent officials and teachers whose dedication and expertise have played a pivotal role in crafting this invaluable content/resource.
I convey my best wishes to all our students for a future brimming with success. Remember, every page you read is a step towards an enlightened tomorrow.

(Dr Rita Sharma)

# DIRECTORATE OF EDUCATION <br> Govt. of NCT, Delhi 

## SUPPORT MATERIAL <br> 2023-2024

## CHEMISTRY

## Class: XI

## NOT FOR SALE

## PUBLISHED BY : DELHI BUREAU OF TEXTBOOKS

# भारत का संविधान 

## भाग 4क

## नागरिकों के मूल कर्तव्य

## अनुच्छेद 51 क

मूल कर्तव्य-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;
(ख) स्वतंत्रता के लिए हमारे राष्ट्रीय आंदोलन को प्रेरित करने वाले उच्च आदर्शों को हृदय में संजोए रखे और उनका पालन करें;
(ग) भारत की संप्रभुता, एकता और अखंडता की रक्षा करे और उसे अक्षुणण बनाए रखे;
(घ) देश की रक्षा करे और आह्वान किए जाने पर राष्ट्र की सेवा करे;
(ङ) 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;
(छ) 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;
(ज) 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 सकें; और
(ट) 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

It shall be the duty of every citizen of India -
(a) to abide by the Constitution and respect its ideals and institutions, the National Flag and the National Anthem;
(b) to cherish and follow the noble ideals which inspired our national struggle for freedom;
(c) to uphold and protect the sovereignty, unity and integrity of India;
(d) to defend the country and render national service when called upon to do so;
(e) 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;
(f) to value and preserve the rich heritage of our composite culture;
(g) to protect and improve the natural environment including forests, lakes, rivers, wildlife and to have compassion for living creatures;
(h) to develop the scientific temper, humanism and the spirit of inquiry and reform;
(i) to safeguard public property and to abjure violence;
(j) 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;
*(k) 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.

[^0]


## Support Material Preparation Team Class XI - Chemistry

| S.No. | Subject Expert | Designation | School |
| :---: | :--- | :--- | :--- |
| 1. | Ashok Kumar | Vice Principal | SBS AFPS SOSE <br> Jharoda Kalan |
| 2. | Sandeep | PGT (Chem.) | GBSSS Alipur |
| 3. | Amit Kumar | PGT (Chem.) | S.V. Sec. 8, Rohini |
| 4. | Manjeet Singh Hooda | PGT (Chem.) | SBV Keshav Puram |
| 5. | Mohit Garg | PGT (Chem.) | RPVV, Shalimar Bagh <br> BT-Block, New Delhi |

## CHEMISTRY (CODE NO. 043)

## Rationale

Higher Secondary is the most crucial stage of school education because at this juncture specialized discipline based, content -oriented courses are introduced. Students reach this stage after 10 years of general education and opt for Chemistry with a purpose of pursuing their career in basic sciences or professional courses like medicine, engineering, technology and study courses in applied areas of science and technology at tertiary level. Therefore, there is a need to provide learners with sufficient conceptual background of Chemistry, which will make them competent to meet the challenges of academic and professional courses after the senior secondary stage.

The new and updated curriculum is based on disciplinary approach with rigour and depth taking care that the syllabus is not heavy and at the same time it is comparable to the international level. The knowledge related to the subject of Chemistry has undergone tremendous changes during the past one decade. Many new areas like synthetic materials, bio -molecules, natural resources, industrial chemistry are coming in a big way and deserve to be an integral part of chemistry syllabus at senior secondary stage. At international level, new formulations and nomenclature of elements and compounds, symbols and units of physical quantities floated by scientific bodies like IUPAC and CGPM are of immense importance and need to be incorporated in the updated syllabus. The revised syllabus takes care of all these aspects. Greater emphasis has been laid on use of new nomenclature, symbols and formulations, teaching of fundamental concepts, application of concepts in chemistry to industry/ technology, logical sequencing of units, removal of obsolete content and repetition, etc.

## Objectives

The curriculum of Chemistry at Senior Secondary Stage aims to:

- promote understanding of basic facts and concepts in chemistry while retaining the excitement of chemistry.
- make students capable of studying chemistry in academic and professional courses (such as medicine, engineering, technology) at tertiary level.
- expose the students to various emerging new areas of chemistry and apprise them with their relevance in future studies and their application in various spheres of chemical sciences and technology.
- equip students to face various challenges related to health, nutrition, environment, population, weather, industries and agriculture.
- develop problem solving skills in students.
- expose the students to different processes used in industries and their technological applications.
- apprise students with interface of chemistry with other disciplines of science such as physics, biology, geology, engineering etc.
- acquaint students with different aspects of chemistry used in daily life.
- develop an interest in students to study chemistry as a discipline.
- integrate life skills and values in the context of chemistry.


# Course Structure <br> <br> Class : XI (Theory) (2023-24) <br> <br> Class : XI (Theory) (2023-24) <br> Chemistry <br> Total period (Theory $160+$ Practical 60) 

Time : 3 Hours
Unit No. Title

| Unit I Some Basic Concepts of Chemistry | 12 | 07 |
| :--- | :--- | :--- | :--- |


| Unit II | Structure of Atom | 14 | 09 |
| :--- | :--- | :--- | :--- |

$\begin{array}{llll}\text { Unit III Classification of Elements and } & 08 & 06\end{array}$ Periodicity in Properties

| Unit IV Chemical Bonding and Molecular | 14 | 07 |
| :--- | :--- | :--- | :--- | Structure

$\begin{array}{llll}\text { Unit } V & \text { Chemical Thermodynamics } & 16 & 09\end{array}$
$\begin{array}{llll}\text { Unit VI Equilibrium } & 14 & 07\end{array}$
Unit VII Redox Reactions 06
Unit VIII Organic Chemistry: Some Basic 14
Principles and Techniques
$\begin{array}{lll}\text { Unit IX Hydrocarbons } & 12 & 10\end{array}$
Total $110 \quad 70$

Unit I : Some Basic Concepts of Chemistry
12 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.

Discovery of Electron, Proton and Neutron, Atomic Number, Isotopes and Isobar. Thomson's model and its limitations. Rutherford's model and its limitations, 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 $\mathrm{s}, \mathrm{p}$ 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

08 Periods
Significance of classification, Brief History of the Development of periodic table, 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 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 $\Delta \mathrm{U}$ and $\Delta \mathrm{H}$, 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).

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

06 Periods
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.

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.

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 | $\mathbf{3 0}$ |

## PRACTICAL SYLLABUS

Total Periods 60
Micro-chemical methods are available for several of the 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
5. Determination of melting point of an organic compound.
6. Determination of boiling point of an organic compound
7. Crystallization of impure sample of any one of the following: Alum, Copper Sulphate, Benzoic Acid.

## C. Experiments based on $\mathbf{p H}$

(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.

## One of the following experiments:

(a) Study the shift in equilibrium between ferric ions and thiocyanate ions by increasing/decreasing the concentration of either of the ions.
(b) Study the shift in equilibrium between $\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$ and chloride ions by changing the concentration of either of the ions.

## E. Quantitative Estimation

(i) Using a mechanical balance/electronic balance
(ii) Preparation of standard solution of Oxalic acid.
(iii) Determination of strength of a given solution of Sodium hydroxide by titrating it against standard solution of Oxalic acid.
(iv) Preparation of standard solution of Sodium carbonate.
(v) Determination of strength of a given solution of hydrochloric acid by titrating it against standard Sodium Carbonate solution.

## F. Qualitative Analysis

(a) Determination of one anion and one cation in a given salt

Cations- $\mathrm{Pb}^{2+}, \mathrm{Cu}^{2+}, \mathrm{As}^{3+}, \mathrm{Al}^{3+}, \mathrm{Fe}^{3+}, \mathrm{Mn}^{2+}, \mathrm{Ni}^{2+}, \mathrm{Zn}^{2+}, \mathrm{Co}^{2+}, \mathrm{Ca}^{2+}, \mathrm{Sr}^{2+}$, $\mathrm{Ba}^{2+}, \mathrm{Mg}^{2+}, \mathrm{NH}^{4+}$
Anions $-\mathrm{CO}_{3}{ }^{2-}, \mathrm{S}^{2-}, \mathrm{NO}^{2-}, \mathrm{SO}_{3}{ }^{2-}, \mathrm{SO}_{4}{ }^{2-}, \mathrm{NO}_{3}^{-}, \mathrm{Cl}^{-}, \mathrm{Br}^{-}, \mathrm{I}^{-}, \mathrm{PO}_{4}{ }^{3-}$, $\mathrm{CH}_{3} \mathrm{COO}^{-}$
(Note: Insoluble salts excluded)
(b) Detection of -Nitrogen, Sulphur, Chlorine in organic compounds.
(c) Projects

Scientific investigations involving laboratory testing and collecting information from other sources.

A few suggested Projects

- Checking the bacterial contamination in drinking water by testing sulphide ion
- Study of the methods of purification of water
- Testing the hardness, presence of Iron, Fluoride, Chloride, etc., depending upon the regional variation in drinking water and study of causes of presence of these ions above permissible limit (if any).
- Investigation of the foaming capacity of different washing soaps and the effect of addition of Sodium carbonate on it
- Study the acidity of different samples of tea leaves.
- Determination of the rate of evaporation of different liquids Study the effect of acids and bases on the tensile strength of fibers.
- Study of acidity of fruit and vegetable juices.

Note: Any other investigatory project, which involves about 10 periods of work, can be chosen with the approval of the teacher.

## Practical Examination for Visually Impaired Students Class XI

Note: Same Evaluation scheme and general guidelines for visually impaired students as given for Class XII may be followed.

## A. List of apparatus for identification for assessment in practicals (All experiments)

Beaker, tripod stand, wire gauze, glass rod, funnel, filter paper, Bunsen burner, test tube, test tube stand, dropper, test tube holder, ignition tube, china dish, tongs, standard flask, pipette, burette, conical flask, clamp stand, dropper, wash bottle

- Odour detection in qualitative analysis
- Procedure/Setup of the apparatus


## B. List of Experiments

(a) Characterization and Purification of Chemical Substances

1. Crystallization of an impure sample of any one of the following: copper sulphate, benzoic acid
(b) Experiments based on $\mathbf{p H}$
2. Determination of pH of some solutions obtained from fruit juices, solutions of known and varied concentrations of acids, bases and salts using pH paper
3. Comparing the pH of solutions of strong and weak acids of same concentration.

## C. Chemical Equilibrium

1. Study the shift in equilibrium between ferric ions and thiocyanate ions by increasing/decreasing the concentration of either ion.
2. Study the shift in equilibrium between $\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$ and chloride ions by changing the concentration of either of the ions.

## D. Quantitative estimation

1. Preparation of standard solution of oxalic acid.
2. Determination of molarity of a given solution of sodium hydroxide by titrating it against standard solution of oxalic acid.

## E. Qualitative Analysis

1. Determination of one anion and one cation in a given salt
2. Cations $-\mathrm{NH}_{4}^{+}$ Anions - $\left(\mathrm{CO}_{3}\right)^{2-}, \mathrm{S}^{2-},\left(\mathrm{SO}_{3}\right)^{2-}, \mathrm{Cl}^{-}, \mathrm{CH}_{3} \mathrm{COO}^{-}$
(Note: insoluble salts excluded)
3. Detection of Nitrogen in the given organic compound.
4. Detection of Halogen in the given organic compound.

Note: The above practicals may be carried out in an experiential manner rather than recording observations.

## Prescribed Books:

1. Chemistry Part - I, Class-XI, Published by NCERT.
2. Chemistry Part - II, Class-XI, Published by NCERT,

## Chemistry - XI <br> Index

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## Some Basic Concepts of Chemistry

## FAST TRACK: QUICK REVISION

- Matter : Anything that has mass and occupies space.
- Precision : It 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 : $1 \mathrm{~L}=1 \mathrm{dm}^{3}=10^{3} \mathrm{~cm}^{3}=10^{-3} \mathrm{~m}^{3}$
- Temperature : $\mathrm{K}={ }^{\circ} \mathrm{C}+273.15 ; \frac{{ }^{\circ} \mathrm{F}-32}{9}=\frac{{ }^{\circ} \mathrm{C}}{5}$
- Standard Temperature Pressure (STP) : $0^{\circ} \mathrm{C}$ (273.15 K) temperature and 1 atm pressure.
- Normal Temperature Pressure (NTP) : $20^{\circ} \mathrm{C}$ (293.15 K) temperature and 1 atm pressure.
- Standard Ambient Temperature Pressure (SATP) : $25^{\circ} \mathrm{C}$ (298.15 K) temperature and 1 atm pressure
- Scientific Notation : Expressing a number in the form $\mathrm{N} \times 10^{n}$, 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 of their volume provided all gases are under the same temperature and pressure.

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- 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).
$1 u=1.66056 \times 10^{-24} \mathrm{~g}$
- Molecular mass : It is algebraic the sum of the atomic mass of the elements present in the molecule.

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

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

Avogadro number $\left(\mathrm{N}_{\mathrm{A}}\right)=6.022 \times 10^{23} \mathrm{~mol}^{-1}$

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

For example : Molar mass of $\mathrm{CH}_{4}=(1 \times 12)+(4 \times 1)=16 \mathrm{~g} \mathrm{~mol}^{-1}$

- 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 $=22.4 \mathrm{~L}$ volume at STP of gaseous substance
e.g., 1 mole of $\mathrm{CH}_{4}=16 \mathrm{~g}$ of $\mathrm{CH}_{4}=6.022 \times 10^{23}$ molecules of $\mathrm{CH}_{4}=22.4 \mathrm{~L}$ at STP
$n=\frac{w g}{\mathrm{M}_{m}}=\frac{\mathrm{VL}(\text { at STP) }}{22.4 \mathrm{~L}}=\frac{x \text { particles }}{\mathrm{N}_{\mathrm{A}}}=\frac{\mathrm{MV}}{1000}$
- Molar Volume $\left(\mathbf{V}_{m}\right)$ : It is volume occupied by one mole of gas at STP.

Molar volume of a gas $=22.4 \mathrm{~L}$ at $\mathrm{STP}(273 \mathrm{~K}, 1 \mathrm{~atm})$ or 22.7 L at STP $(273$ K, 1 bar)

Calculating Molar Volume: $\mathrm{PV}=n \mathrm{RT}$

$$
\therefore \mathrm{V}=\frac{n \mathrm{RT}}{\mathrm{P}}=\frac{1 \mathrm{~mol} \times 0.082 \mathrm{~L} \mathrm{~atm} \mathrm{~K}^{-1} \mathrm{~mol}^{-1} \times 273 \mathrm{~K}}{1 \mathrm{~atm}}=22.4 \mathrm{~L}
$$

Or

$$
\mathrm{V}=\frac{n \mathrm{RT}}{\mathrm{P}}=\frac{1 \mathrm{~mol} \times 0.083 \mathrm{~L} \mathrm{bar} \mathrm{~K}^{-1} \mathrm{~mol}^{-1} \times 273 \mathrm{~K}}{1 \mathrm{bar}}=22.7 \mathrm{~L}
$$

- 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., $\mathrm{C}_{6} \mathrm{H}_{6}$ is the molecular formula of benzene.
- Relationship between empirical and molecular formulae :

Molecular formula $=n \times$ Empirical formula
Where;

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

- Information Conveyed by a chemical equation :

| $\mathbf{N}_{\mathbf{2}}(\boldsymbol{g})$ | $+\mathbf{3} \mathbf{H}_{\mathbf{2}}(\boldsymbol{g})$ | $\rightarrow \mathbf{2 \mathbf { N H } _ { \mathbf { 3 } } ( \boldsymbol { g } )}$ |  |
| :--- | :--- | :--- | :--- |
| (i) 1 molecule of $\mathrm{N}_{2}+3$ molecules of $\mathrm{H}_{2}$ | $\rightarrow$ | 2 molecules of $\mathrm{NH}_{3}$ |  |
| (ii) 1 mole of $\mathrm{N}_{2}+3$ mole of $\mathrm{H}_{2}$ | $\rightarrow$ | 2 mole of $\mathrm{NH}_{3}$ |  |
| (iii) $1 \times 28 \mathrm{~g}$ of $\mathrm{N}_{2}$ | $+3 \times 2 \mathrm{~g}$ of $\mathrm{H}_{2}$ | $\rightarrow$ | $2 \times 17 \mathrm{~g}$ of $\mathrm{NH}_{3}$ |
| (iv) $1 \times 22.4 \mathrm{~L}$ of $\mathrm{N}_{2}$ | $+3 \times 22.4 \mathrm{~L}$ of $\mathrm{H}_{2}$ | $\rightarrow$ | $2 \times 22.4 \mathrm{~L}$ of $\mathrm{NH}_{3}$ |
|  | 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.

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

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

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

- Molarity (M) : It is number of moles of solute dissolved per litre ( $\mathrm{dm}^{3}$ ) of the solution.

Molarity $=\frac{\text { No. of moles of solute }}{\text { Volume of solution in } L}$
Molarity equation: $\quad \mathrm{M}_{1} \mathrm{~V}_{1}=\mathrm{M}_{2} \mathrm{~V}_{2}$ (Before dilution) (After Dilution)

Molarity of a solution decreases on increasing temperature.
Molarity of pure water is $55.56 \mathrm{~mol} \mathrm{~L}^{-1}$

- Molality (m)—It is number of moles of solute dissolved per $1000 \mathrm{~g}(1 \mathrm{~kg})$ of solvent.

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

- Mole Fraction $(\boldsymbol{x})$ is the ratio of number of moles of one component to the total number of moles (solute and solvents) present in the solution.

$$
x_{1}=\frac{n_{1}}{n_{1}+n_{2}} \text { and } x_{2}=\frac{n_{2}}{n_{1}+n_{2}}
$$

The sum of all the mole fractions in a solution is equal to one. i.e., $x_{1}+x_{2}=1$

| MIND MAP |
| :---: |
| SOME BASIC OF CONCEPTS OF CHEMISTRY |



## CASE BASE : QUESTIONS

## 1. Read the passage given below and answer the following questions:

The ideas underlying our modern understanding of thermodynamics and kinetic theory were developed during the nineteenth century. Central to these developments was the discovery that matter reacting chemically does not do so simply between equal masses of the samples involved. We now call the study of this phenomenon 'stoichiometry', defined as: 'the relationship between the amounts of substance that react together, and the products that are formed'.

Another development during the nineteenth century that was central to our modern understanding of the chemical nature of matter was the observation by Avogadro that 'equal volumes of ideal or perfect gases, at the same temperature and pressure, contain the same number of particles, or molecules'. This is now known as Avogadro's law. It provides the motivation to formulate expressions for the quantity of a sample that reacts with another sample. The most notable example of such a formulation is the gram-molecule, which has been used to refer to both a unit and a quantity.
(Reference: Milton Martin J. T. 2011A new definition for the mole based on the Avogadro constant: a journey from physics to chemistry Phil. Trans. R. Soc. A.3693993-4003)

The following questions are multiple choice questions. Choose the most appropriate answer:
I. The concept of stoichiometry mentioned in the study is based on the
a. formation of chemical bonds.
b. amount of reactant and product involved in a chemical reaction.
c. idea of temperature and pressure required for the reaction to occur.
d. oxidation states of reactant and product involved.
II. How much gram-molecules of $\mathrm{H}_{2} \mathrm{O}$ are produced on combustion of 32 g of methane in excess oxygen?
a. $\quad 72$
b. 4
c. 2
d. 36
III. When an antacid tablet is used, $\mathrm{Ca}(\mathrm{OH})_{2}$ reacts with HCl in the stomach to form inert $\mathrm{CaCl}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$. If the molar mass of $\mathrm{Ca}(\mathrm{OH})_{2}$ is $75 \mathrm{~g} / \mathrm{mol}$, how many moles of HCl are required to fully react with 150 g of $\mathrm{Ca}(\mathrm{OH})_{2}$ ?
a. 4
b. 1
c. 8
d. 2
IV. What must be held constant when applying Avogadro's law?
a. pressure and temperature
b. volume and temperature
c. moles and temperature
d. pressure and volume

ANS:- I-B, II-B, III-A, IV-A

## 2. Read the passage given below and answer the following questions:

The goal of this study was to examine the means used by textbook authors to introduce, define, and explain the mole concept in high school and introductory college chemistry textbooks. The analysis was framed by four questions:

1. How is the mole defined?
2. What concepts about the atom are introduced prior to the mole?
3. Is Avogadro's constant presented as an experimentally determined value?
4. What is the context for introducing the mole?

Twenty nine high school and introductory college level chemistry texts were examined. After independent reading of appropriate sections of each text, discussion of differences, second or third readings of texts, and subsequent discussions, both authors reach $100 \%$ agreement concerning the results. Major conclusions were

1. Two ways of defining the mole dominate the texts. One way defines the mole as Avogadro's number ( $6.02 \times 10^{23}$ ) particles; the other method defines the mole in terms of carbon 12.
2. All texts that present a definition in terms of C-12 introduce and define concepts about the atom prior to introducing the mole.
3. Most texts at all levels point out that the value $6.02 \times 10^{23}$ is an experimentally determined quantity.
4. Nearly all texts discuss the mole in relation to die problem of finding a way to count particles that are too small to be directly weighed. Most texts also use a familiar counting unit, such as the dozen, to introduce the mole by analogy.
(Reference: John R. Staver, Andrew T. Lumpe, A content analysis of the presentation of the mole concept in chemistry textbooks, Journal of Research in Science Teaching).

In these questions (Q. No. (i) to (iv), a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.
a) Assertion and reason both are correct statements and reason is correct explanation for assertion.
b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
c) Assertion is correct statement but reason is wrong statement.
d) Assertion is wrong statement but reason is correct statement
i. ASSERTION: Equal moles of different substances contain same number of constituent particles.

REASON: Equal weights of different substances contain the same number of constituent particles.
ii. ASSERTION: Both 106 g of sodium carbonate and 12 g of carbon have same number of carbon atoms.

REASON: Both contain 1 g -atom of carbon which contains $6.02 \times 10^{23}$ carbon atoms
iii. ASSERTION: Both $32 \mathrm{~g} \mathrm{SO}_{2}$ and $8 \mathrm{~g} \mathrm{CH}_{4}$ have same number of molecules. REASON: Equal moles of substances have equal number of molecules.
iv. ASSERTION: Average atomic mass of an element depends mainly on the heavier isotope.

REASON: The average atomic mass of an element is the sum of the masses of its isotopes, each multiplied by its natural abundance

ANS:- I-C, II-A, III-A, IV-D

## MULTIPLE CHOICE QUESTIONS (MCQ)

1. Which of the following is dependent of temperature?
(a) Molarity
(b) Molality
(c) Mole fration
(d) Mass percentage
2. 4 g of NaOH dissolved in 100 ml solution. Molarity of the solution is
(a) 1 M
(b) 10 M
(c) 0.1 M
(d) 4 M
3. Which has the maximum number of molecules among the following ?
(a) $44 \mathrm{~g}^{\text {of } \mathrm{CO}_{2}}$
(b) $44 \mathrm{~g}^{\text {of }} \mathrm{O}_{2}$
(c) 8 g of $\mathrm{H}_{2}$
(d) 64 g of $\mathrm{SO}_{2}$
4. $\quad 10 \mathrm{~mol}$ of Zn react with 10 mol of HCl . Calculate the number of moles of $\mathrm{H}_{2}$ produced.
(a) 5 mol
(b) 10 mol
(c) 20 mol
(d) 2.5 mol
5. The number of oxygen atoms in 4.4 g of $\mathrm{CO}_{2}$ is approximately
(a) $1.2 \times 10^{23}$
(b) $6 \times 10^{22}$
(c) $6 \times 10^{23}$
(d) $12 \times 10^{23}$
6. The molarity of a solution obtained by mixing 750 mL of 0.5 M HCl with 250 ml of 2 M HCl will be
(a) 0.975 M
(b) 0.875 M
(c) $\quad 1.00 \mathrm{M}$
(d) 1.175 M
7. Number 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. $6.02 \times 10^{20}$ molecules of urea are present in 100 mL of its solution. The concentration of the solution is
(a) $\quad 0.02 \mathrm{M}$
(b) 0.01 M
(c) $\quad 0.001 \mathrm{M}$
(d) 0.1 M
9. A gaseous hydrocarbons gives upon combustion, 0.72 g of water and 3.08 g of $\mathrm{CO}_{2}$. The empirical formula of the hydrocarbon is :
(a) $\mathrm{C}_{6} \mathrm{H}_{5}$
(b) $\mathrm{C}_{7} \mathrm{H}_{8}$
(c) $\mathrm{C}_{2} \mathrm{H}_{4}$
(d) $\mathrm{C}_{3} \mathrm{H}_{4}$
10. The density of solution prepared by dissolving 120 g of urea (Mol. mass $=60 \mathrm{u}$ ) in 1000 g of water is $1.15 \mathrm{~g} / \mathrm{mL}$. The molarity of the solution is
(a) $\quad 0.50 \mathrm{M}$
(b) 1.78 M
(c) $\quad 1.02 \mathrm{M}$
(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. $\quad 17 \mathrm{~g}$ of $\mathrm{NH}_{3}$ gas will occupy a volume of $\qquad$ $\mathrm{cm}^{3}$ at NTP.
2. The number of Li atoms in $\qquad$ g. is $6.022 \times 10^{24}$ atoms.
3. $(1 / 12)$ th of the mass of carbon atom is $\qquad$
4. Number of atoms of oxygen in 24 g of $\mathrm{O}_{3}$ is $\qquad$
5. The number of moles of barium carbonate which contains 1.5 moles of oxygen atoms is $\qquad$
6. A mixture having 2 g of $\mathrm{H}_{2}$ and 32 g of oxygen occupies a volume of
$\qquad$ at NTP.
7. If the phosphate of a metal has the formula $\mathrm{MPO}_{4}$ the formula of the metallic sulphate is $\qquad$
8. At NTP, the mass of 1 litre of gas is 3 g . Molecular mass of the gas is
$\qquad$
9. The percentage mass of magnesium in chlorophyll is $2.68 \%$ The number of magnesium atoms in 2 g of chlorophyll is $\qquad$
10. The mass of one molecule of carbon dioxide is $\qquad$
11 Percentage of nitrogen in urea is $\qquad$
11. Number of carbon atoms present in 18 g of glucose $\left(\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}\right)$
12. 0.5 mole of triatomic gas contains $\qquad$ atoms.
13. A binary compound contains $50 \% \mathrm{~A}$ (at. mass $=16$ ) and $50 \% \mathrm{~B}$ (at. mass 32). The empirical formula of the compound is $\qquad$ .
14. The number of hydrogen atoms in 60 u of ethane is $\qquad$
Ans: 1. 22400
15. 70 g
16. 1 u
17. $9.033 \times 10^{23}$
18. 0.5
19. 44.8 litre
20. $\mathrm{M}_{2}\left(\mathrm{SO}_{4}\right)_{3}$
$8 \quad 67.2$
21. $1.34 \times 10^{21}$
22. $7.3 \times 10^{-23}$
23. 46.67
24. $3.61 \times 10^{23}$
25. $9.033 \times 10^{23}$
26. $\mathrm{A}_{2} \mathrm{~B}$
27. $7.226 \times 10^{24}$

## 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 $\mathrm{C}_{12} \mathrm{H}_{22} \mathrm{O}_{11}$ 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 $=(\text { Molecular formula })_{n}$.
7. Gram-atomic mass of an element may be defnined 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)

## MATCH THE COLUMNS

1. 

Column X Column Y
a. $\quad 8 \mathrm{~g} \mathrm{CH}_{4}$
b. $\quad 1.7 \mathrm{~g} \mathrm{NH}_{3}$
c. $\mathrm{CH}_{3} \mathrm{OH}$
d. $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$

## Column Z

p. Emp. formulation $\rightarrow \mathrm{CH}_{2} \mathrm{O}$
q. $50 \%$ oxygen
r. $1.806 \times 10^{23}$ atoms of hydrogen
s. $25 \%$ hydrogen
2.

## Column X

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

Column X
a. 40 g of He
b. 35 g of Li
c. 40 u of He
d. 16 g of $\mathrm{O}_{2}$

## Column Y

i. $\quad 3.011 \times 10^{23}$ atoms
ii. $\quad 10$ atoms
iii. $\quad 6.022 \times 10^{24}$ atoms
iv. $\quad 3.011 \times 10^{24}$ atoms

## Column Z

p. 0.5 moles
q. $1.67 \times 10^{-23}$
r. $\quad 10$ moles
s. 5 moles
4.

## Column X

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

## Column Y

i. For very dilute solution
ii. No units
iii. $\mathrm{Mol} \mathrm{L}^{-1}$
iv. independent of temperature
3.

Ans: 1. a.(ii). (s), b.(i). (r), c.(iv). (q), d.(iii). (p)
2. a.(iii), b.(iv), c.(ii), d.(i)
3. a.(iii). (r), b.(iv). (s), c.(ii). (q), d.(i). (p)
4. a.(iv), b.(iii), c.(ii), d.(i)

## ASSERTION AND REASON TYPE QUESTIONS

## Directions for Q. No.1-10

A Both Assertion \& Reason are true and the reason is the correct explanation of the assertion.
B Both Assertion \& Reason are true but the reason is not the correct explanation of the assertion.
C Assertion is true statement but Reason is false.
D Assertion is false but Reason is true.

1. 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 $\mathrm{H}_{2}$ in 0.224 L of hydrogen is 0.01 mole.

Reason : 22.4 L of $\mathrm{H}_{2}$ at STP contain $6.023 \times 10^{23}$ moles.
4. Assertion : Atomic mass of Na is 23.

Reason : An atom of sodium is 23 times heavier than $1 / 12$ th mass of $\mathrm{C}-12$ isotope.
5. Assertion : Number of atoms of He in 60 u of He is 15.

Reason : Atomic weight of He is 4 u .
6. Assertion : In a gaseous reaction, the ratio of volumes of reactants and products is in agreement with their molar ratio.
Reason : Volume of gas is inversely proportional to its moles at particular temperature and pressure.
7. Assertion : The Empirical mass of ethane is half of its molecular mass.

Reason : The empirical formula represents the simplest whole number ratio of various atoms present in a compound.
8. Assertion : Significant figures for 0.200 is 3 whereas for 200 is 1

Reason : Zero at the end or right of a number are significant provided they are not on the right side of the decimal point.
9. Assertion : One molar aqueous solution has always higher concentration than One molal aqueous solution
Reason : The molality of solution depends upon the density of solution whereas molarity does not.
10. Assertion : In a combustion reaction in the air, oxygen is the limiting reagent
Reason : Limiting reagent is the reactant in a chemical reaction that limits the amount of product that can be formed.

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

## 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 .
4. What is the volume of 17 g of $\mathrm{NH}_{3}$ gas at STP?
[Ans. : 223.4 L]
5. How many molecules of $\mathrm{SO}_{2}$ are present in 11.2 L at STP?
[Ans. : $3.011 \times 10^{23}$ ]
6. Which has more number of atoms ? 1.0 g Na or 1.0 g Mg
[Ans. : 1.0 g Na ]
7. How many oxygen atoms are present in 16 g of ozone $\left(\mathrm{O}_{3}\right)$ ?
[Ans. : $2.007 \times 10^{23}$ ]
8. Calculate the number of molecules present in 22.0 g of $\mathrm{CO}_{2}$.
[Ans. : $3.011 \times 10^{23}$ ]
9. A substance has molecular formula $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$. What is its empirical formula.
10. Empirical formula of a compound X (Molar mass $=78 \mathrm{~mol}^{-1}$ ) is CH . Write its molecular formula.

## 1-MARK QUESTIONS

1. Classify following as pure substances and mixtures : air, glucose, gold, sodium and milk.
2. Which measurement is more precise 4.0 g or 4.00 g ?
[Ans. 4.00 g ]
3. How many significant figures are there in (i) 3.070 and (ii) 0.0025 ?
[Ans. (i) 4 (ii) 2]
4. Express the following in the scientific notation : (i) 0.0048 (ii) 234,000
5. 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]
6. At STP, what will be the volume of $6.022 \times 10^{23}$ molecules of $\mathrm{H}_{2}$ ?
[Ans. 22.4L]
7. 1 L of a gas at STP weighs 1.97 g . What is molecular mass ?
[Ans. $44.128 \mathrm{~g} \mathrm{~mol}^{-1}$ ]
8. Write the relationship between empirical formula and molecular formula.
9. Which is more informative? Empirical formula or Molecular formula.
10. How are $0.5 \mathrm{~mol} \mathrm{Na}_{2} \mathrm{CO}_{3}$ and $0.5 \mathrm{M} \mathrm{Na}_{2} \mathrm{CO}_{3}$ different from each other ?
11. Why molality is preferred over molarity of a solution?
12. Define molarity of a solution.
13. What is the effect of temperature on molarity of solution?
14. 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 $37^{\circ} \mathrm{C}$. Calculate its value in ${ }^{\circ} \mathrm{F}$.
3. Convert 5 L into $\mathrm{m}^{3}$.
4. What does the following prefixes stand for :
(a) pico
(b) nano
(c) micro
(d) deci
5. How many significant figures are present in the following :
(i) 4.00005
(ii) 0.004
6. Convert ' 450 pm ' into SI unit and write the answer in scientific notation upto 2 significant figures.

$$
\text { [Ans. } 4.5 \times 10^{-10} \mathrm{~m} \text { ] }
$$

7. Hydrogen peroxide and water contain $5.93 \%$ and $11.2 \%$ of hydrogen respectively. Show that the data illustrate law of multiple proportions.
8. The density ( in $g \mathrm{~mL}^{-1}$ ) of a 3.60 M sulphuric acid solution that is $29 \%$ $\mathrm{H}_{2} \mathrm{SO}_{4}\left(\right.$ Molar mass $\left.=98 \mathrm{~g} \mathrm{~mol}^{-1}\right)$ by mass will be $\qquad$
[Ans. $1.21 \mathrm{~g} / \mathrm{mL}$ ]
9 The cost of table salt ( NaCl ) is Rs. 10 per Kg . calculate its cost per mole. ( Molar mass of NaCl is $58.5 \mathrm{gmol}^{-1}$ )
[Ans. 0.58 Rs ]
10 Calculate the mole fraction of the solute in a 1.00 molal aqueous solution.
[Ans. 0.0177]
11 Dissolving 120 g of urea ( Molar mass of urea $=60 \mathrm{~g} \mathrm{~mol}^{-1}$ ) in 1000 of water gave a solution of density $1.15 \mathrm{~g} / \mathrm{mL}$. Calculate the molarity of the solution.
[Ans. 2.05 M]
12 Calculate the percentage of N in urea. (Molar mass of urea $=60 \mathrm{~g}$ $\mathrm{mol}^{-1}$ )
[Ans. 46.66]
1325 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 ]

14 How many atoms and molecules are present in 124 gm of phosphorus $\left(\mathrm{P}_{4}\right)$
[Ans. Atoms $=4 \mathrm{~N}_{\mathrm{A}}$ \& Molecules $=\mathrm{N}_{\mathrm{A}}$ ]
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 : $2 \mathrm{~N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{~N}_{2} \mathrm{O}(\mathrm{g})$
Which law is being obeyed in this experiment? Write the statement of the law.

16 Give one example each of a molecule in which empirical formula and molecular formula is
(i) Same (ii) Different.

17 Calculate the number of moles in the following masses :
(i) 7.85 g of Fe ;
(ii) 7.9 mg of Ca

18 Calculate the percent of carbon, hydrogen and oxygen in ethanol ( $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ )
[Ans. 52.14\%, 13.13\%, 34.73\%]
19 How much copper can be obtained from 100 g of $\mathrm{CuSO}_{4}$ ? [Ans. 39.8g]
20 Calculate the amount of water (g) produced by the combustion of 16 g of methane.
[Ans. 36g]
21 How many moles of methane are required to produce $22 \mathrm{~g} \mathrm{CO}_{2}(\mathrm{~g})$ after combustion?
[Ans. 0.5 mol$]$
22 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\%]
23 Calculate molarity of water if its density is $1.00 \mathrm{~g} \mathrm{~mL}^{-1}$.
[Ans. 55.56 M ]
24 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 ]
25 The density of 3 M solution of NaCl is $1.25 \mathrm{~g} \mathrm{~mL}^{-1}$. Calculate molality of the solution.
[Ans. 2.8m]
$26 \mathrm{NH}_{3}$ gas can be prepared by Haber's process as, $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow$ $2 \mathrm{NH}_{3}(\mathrm{~g})$. At a particular moment concentration of all the species is 2 moles; calculate the concentration of $\mathrm{N}_{2}$ and $\mathrm{H}_{2}$ taken initially.
[Ans. 3 mole, 5 moles]

## 3-MARKS QUESTIONS

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

|  | \% Natural Abudance | Molar mass |
| :--- | :--- | :--- |
| ${ }^{24} \mathrm{Mg}$ | 80 | 24 |
| ${ }^{25} \mathrm{Mg}$ | 10 | 25 |
| ${ }^{26} \mathrm{Mg}$ | 10 | 26 |

2. The following data are obtained when dinitrogen and dioxygen react together to form different compounds :
(i)
(ii) (iii) (iv)
$\begin{array}{lllll}\text { Mass of dinitrogen } & 14 & 14 & 28 & 28\end{array}$
$\begin{array}{lllll}\text { Mass of dioxygen } & 16 & 32 & 32 & 80\end{array}$
Which law of chemical combination is obeyed by the above experimental data? Give its statement.
3. Calculate :
(i) Mass in gram of $5.8 \mathrm{~mol} \mathrm{~N}_{2} \mathrm{O}$
(ii) Number of moles in 8.0 g of $\mathrm{O}_{2}$
(iii) Molar mass if 11.2 L at STP weigh 8.5 g .
[Ans. (i) 255.2 g (ii) 0.25 mol (iii) $17 \mathrm{~g} \mathrm{~mol}^{-1}$ ]
4. In three moles of ethane $\left(\mathrm{C}_{2} \mathrm{H}_{6}\right)$, 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) $1.81 \times 10^{24}$ ]
5. 16 g of an ideal gas $\mathrm{SO}_{\mathrm{x}}$ occupies 5.6 L at STP. What is its molecular mass ? What is the value of X ?
[Ans. 64u, $\mathrm{x}=2$ ]
6. Calculate the number of moles :
(i) 5.0 L of $0.75 \mathrm{M} \mathrm{Na}_{2} \mathrm{CO}_{3}$
(ii) 7.85 g of Fe
(iii) 34.2 g of sucrose $\left(\mathrm{C}_{12} \mathrm{H}_{22} \mathrm{O}_{11}\right)$
[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) 52 u of He (iii) 52 g of He .
[Ans. (i) $3.13 \times 1025$ (ii) 13 (iii) $7.83 \times 1024]$
8. Vitamin C is essential for the prevention of scurvy. Combustion of 0.2000 g of vitamin C gives 0.2998 g of $\mathrm{CO}_{2}$ and 0.819 g of $\mathrm{H}_{2} \mathrm{O}$. What is the empirical formula of vitamin C ?
[Ans. $\mathrm{C}_{3} \mathrm{H}_{4} \mathrm{O}_{3}$ ]
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. $\mathrm{CH}_{2} \mathrm{C} 1, \mathrm{C}_{2} \mathrm{H}_{4} \mathrm{Cl}_{2}$ ]
10. A compound made up of two elements A and B has $\mathrm{A}=70 \%, \mathrm{~B}=30 \%$. 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) $\mathrm{A}_{2} \mathrm{~B}_{3}$ ]
11. The reaction $2 \mathrm{C}+\mathrm{O}_{2} \longrightarrow 2 \mathrm{CO}$ is carried out by taking 24.0 g of carbon and 96.0 g of $\mathrm{O}_{2}$. 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) $\mathrm{O}_{2}$, (ii) 64 g , (iii) 72]
12. A 10 g sample of a mixture of calcium chloride and sodium chloride is treated with $\mathrm{Na}_{2} \mathrm{CO}_{3}$ to precipitate calcium as calcium carbonate. This $\mathrm{CaCO}_{3}$ 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 $\mathrm{H}_{2}$ react with 29.0 g of $\mathrm{O}_{2}$ yield $\mathrm{H}_{2} \mathrm{O}$.
(i) Which is the limiting reagent.
(ii) Calculate the maximum amount of $\mathrm{H}_{2} \mathrm{O}$ that can be formed
(iii) Calculate the amount of reactant left unreacted
[Ans. $\mathrm{H}_{2}, 26.8 \mathrm{~g} \mathrm{H}_{2} \mathrm{O} \& 5.2 \mathrm{~g} \mathrm{O}_{2}$ ]

14 Zinc and hydrochloric acid react according to the reaction:
$\mathrm{Zn}(\mathrm{s})+2 \mathrm{HCl}(\mathrm{aq}) \longrightarrow \mathrm{ZnCl}_{2}(\mathrm{aq})+\mathrm{H}_{2}(\mathrm{~g})$
If 0.30 mol Zn are added to hydrochloric acid containing 0.52 mol of HCl , How many moles of $\mathrm{H}_{2}$ are produced ?
[ HCl is limiting reagent; $\mathrm{H}_{2}$ formed $\left.=0.36 \mathrm{~mol}\right]$
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 $\mathrm{Pb}=207 \mathrm{U}$ ]
[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 $\mathrm{O}_{2} \& 400 \mathrm{~mL}$ of $\mathrm{CO}_{2}$ ]

## 5-MARKS QUESTIONS

1 (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 $=10^{-18} \mathrm{~g}$ ]
[Ans. $5.02 \times 10^{4}$ ]
(ii) Which one of the following will have largest number of atoms?
(a) $\lg \mathrm{Au}(\mathrm{s})$
(b) 1 g Na (s)
(c) $1 \mathrm{gLi}(\mathrm{s})$
(d) 1 g of $\mathrm{Cl}_{2}(\mathrm{~g})$
[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) $26 \mathrm{~g} \mathrm{~mol}^{-1}$, (iii) $\mathrm{C}_{2} \mathrm{H}_{2}$ ]
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 $1.02 \mathrm{~g} \mathrm{~cm}^{-3}$ ).
[Ans.. 1.52 m ]
4. (i) Define : (a) Mole fraction (b) Mass percentage.
(ii) If the density of methanol is $0.793 \mathrm{~kg} \mathrm{~L}^{-1}$, 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 $\mathrm{C}_{\mathrm{x}} \mathrm{H}_{\mathrm{y}} \mathrm{O}_{\mathrm{z}}$, 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 $\mathrm{C}_{\mathrm{x}} \mathrm{H}_{\mathrm{y}}$. Find the empirical formula of the compound.

$$
\left[\text { Ans. } \mathrm{C}_{2} \mathrm{H}_{4} \mathrm{O}_{3}\right]
$$

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
$\mathrm{Mg}=20.0 \%, \mathrm{~S}=26.6 \%, \mathrm{O}=53.33 \%$
Calculate the molecular formula of the anhydrous salt and the crystalline salt. Molecular weight of the anhydrous salt is 120 .
[Ans. $\mathrm{MgSO}_{4} \cdot 7 \mathrm{H}_{2} \mathrm{O}$ ]
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 $27^{\circ} \mathrm{C}$, the weight of cylinder is reduced to 23.2 kg . Find the volume of n-butane in cubic meters used up at $27^{\circ} \mathrm{C}$ and 1 atm (Molecular weight of n-butane $=58$ ).
[Ans. $2.463 \mathrm{~m}^{3}$ ]
$42.5 \mathrm{~g}^{2}$ of $\mathrm{CaCO}_{3}$ was placed in 50 ml of a solution of HCl .1 .05 g of $\mathrm{CaCO}_{3}$ 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-I

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. If 30 mL of $\mathrm{H}_{2}$ and 20 mL of $\mathrm{O}_{2}$ react to form water, what is left at the end of the reaction?
(a) 10 mL of $\mathrm{H}_{2}$
(b) 5 mL of $\mathrm{H}_{2}$
(c) 10 mL of $\mathrm{O}_{2}$
(d) 5 mL of $\mathrm{O}_{2}$
2. 7.5 grams of a gas occupy 5.6 litres of volume at STP the gas is
(a) NO
(b) $\mathrm{N}_{2} \mathrm{O}$
(c) CO
(d) $\mathrm{CO}_{2}$
3. Write the relationship between empirical formula and molecular formula.
4. Why molarity is preferred over molarity in expressing the concentration of solution?
5. Which has more number of atoms? 1.0 g Na or 1.0 g Mg ?
6. How many atoms and molecules are present in 124 g of phosphorus $\left(\mathrm{P}_{4}\right)$ ? (2)
7. (a) How many significant figures are present in 0.0102 .
(b) Write the number in scientific notation 1013.6.
8. A sample of drinking water was found to be severely contaminated with chloroform $\mathrm{CHCl}_{3}$. 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.
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 formula?
10. (a) Explain the following terms:
(i) Gay Lussac's law
(ii) Limiting reagent
(b) 3.0 g of $\mathrm{H}_{2}$ react with 30.0 g of $\mathrm{O}_{2}$ yield $\mathrm{H}_{2} \mathrm{O}$.
(i) Which is the limiting reagent?
(ii) Calculate the maximum amount of $\mathrm{H}_{2} \mathrm{O}$ that can be formed.
(iii) Calculate the amount of reactant left unreacted.

## UNIT TEST-II

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. One mole of oxygen gas at STP is equal to
(a) $6.022 \times 10^{23}$ molecules of oxygen
(b) $\quad 6.022 \times 10^{23}$ atoms of oxygen
(c) 16 g of oxygen
(d) 32 g of oxygen
2. 1 g of $\mathrm{M}_{2} \mathrm{CO}_{3}$ on treatment with excess HCl produces 0.01186 moles of $\mathrm{CO}_{2}$. The molar mass of $\mathrm{M}_{2} \mathrm{CO}_{3}$ in $\mathrm{g} \mathrm{mol}^{-1}$ is ?
(1)
(a) 1186
(b) 84.3
(c) 118.6
(d) 11.86
3. How many atoms are present in 16 g of ozone?

In following questions a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.
(a) Assertion and Reason are true and Reason is the correct explanation of Assertion
(b) Assertion and Reason are true but Reason is not the correct explanation of Assertion
(c) Assertion is true but Reason is false
(d) Both Assertion and Reason are false
4. Assertion : The empirical mass of ethene is half of its molecular mass. (1)

Reason : The empirical formula represents the simplest whole number ratio of various atoms present in a compound.
5. Assertion : Combustion of 16 g of methane gives 18 g water.

Reason: In the combustion of methane, water is one of the products.
6. If 2 litres of $\mathrm{N}_{2}$ is mixed with 2 litres of $\mathrm{H}_{2}$ at a constant temperature and pressure, then what will be the volume of $\mathrm{NH}_{3}$ formed?
7. Calculate the percentage of Copper in a sample of $\mathrm{CuCl}_{2}$
(Atomic mass of $\mathrm{Cu}=63.5 \mathrm{u}, \mathrm{Cl}=35.5 \mathrm{u}$ )
8. In an experiment, when HCl was reacted with $\mathrm{CaCO}_{3}$ at $\mathrm{STP}, 48 \mathrm{Cm}^{3}$ of $\mathrm{CO}_{2}$ was formed. Calculate the number of mole of $\mathrm{CO}_{2}$ and number of molecules.
9. In the reaction $2 \mathrm{~A}+4 \mathrm{~B} \rightarrow 3 \mathrm{C}+4 \mathrm{D}$, when 5 moles of A react with 6 moles of B, then
(i) Which is the limiting reagent
(ii) Calculate the amount of C formed
(iii) Calculate the amount of excess reagent left after reaction
10. (a) How many grams atoms are there in 8.0 g of S ?
(b) The molarity of solution of $\mathrm{H}_{2} \mathrm{SO}_{4}$ is 1.35 M . Calculate its molality. (The density of solution is $1.02 \mathrm{~g} \mathrm{~cm}^{-3}$ )

## Chapter - 2

## Structure of Atom

## FAST TRACK : QUICK REVISION

- Information about fundamental particles of atom

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

- 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 ' $\lambda$ '.
(b) Frequency: Number of waves passing through a given point in one second.


Denoted by v. $\quad\left[\begin{array}{l}v=\frac{1}{t} \Rightarrow \sec ^{-1} \text { or } \mathrm{Hz} \\ t=\text { Time period }\end{array}\right]$
(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 $\bar{v}$

$$
\bar{v}=\frac{1}{\lambda}=\mathrm{cm}^{-1}\left(\text { or m }^{-1}\right)
$$

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

$$
\begin{aligned}
\text { velocity of light } c & =\frac{\text { Distance }}{\text { Time }}=\lambda \times \frac{1}{t}=v \lambda \\
\therefore \quad v & =\frac{c}{\lambda}
\end{aligned}
$$

- Energywise order for EM radiation.
cosmic $<\gamma$ rays $<$ X rays $<\mathrm{UV}<$ VIBGYOR $<\mathrm{IR}<$ Microwaves $<$ Radiowaves
$\xrightarrow[\lambda \text { (Increases) } \quad v \text { (Decreases) } \quad \text { Energy (Decreases) }]{ }$
- 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 $\alpha v$

$$
\begin{array}{lr}
\mathrm{E}=h v & (h=\text { Planck's constant }) \\
\mathrm{E}=n h v & \left(h=6.626 \times 10^{-34} \mathrm{~J} \mathrm{sec}\right) \\
\text { If ' } n \text { ' photons are emitted } \mathrm{E}=n h v &
\end{array}
$$

- Photo electric effect : The phenomenon of ejection of electrons from a metal surface when a light of suitable frequency falls on metal surface.
$h \mathrm{v}-h \mathrm{v}_{0}=\frac{1}{2} \mathrm{mv}^{2}$
$h \nu \Rightarrow$ Energy of incident light on metal surface.
$h v_{0} \Rightarrow$ Work function of metal.
$\frac{1}{2} \mathrm{mv}^{2}=$ Kinetic energy by which $e^{-}$is emitted from metal surface.
- de Broglie equation : All material particles in motion also exhibit wave like properties.

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

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

For macroscopic particles mass is large, $\lambda$ 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.

$$
\begin{aligned}
& \Delta x \times m \Delta v \geq \frac{h}{4 \pi} \\
& \Delta x=\text { uncertainty in position } \\
& \Delta v=\text { uncertainty in velocity }
\end{aligned}
$$

## Bohr's theory for $\mathbf{H}\left[\mathbf{H}\right.$ like one $\mathrm{e}^{-}$systems $\mathrm{He}^{+} ; \mathrm{Li}^{\mathbf{2 +}}$ ]

(1) $e^{-}$revolving round the nucleus in circular path [stationary state; SHELL] with a definite angular momentum $\frac{n h}{2 \pi}\left[\right.$ Here $n=$ no. of shell of $\left.e^{-}\right]$and with definite energy
$\mathrm{E}_{\mathrm{n}}=\left[\frac{-2 \pi^{2} m e^{4} z^{2}}{n^{2} h^{2}}\right] \Rightarrow-2.18 \times 10^{-18} \frac{\mathrm{Z}^{2}}{n^{2}} \mathrm{~J} /$ 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 $\Delta \mathrm{E}=\mathrm{E}_{f}-\mathrm{E}_{i}$.

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 ' $\lambda$ ' of light radiations do not merge with each other like in VIBGYOR (Continous Spectra).

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

$$
\begin{aligned}
& \frac{1}{\lambda}=1,09,678\left[\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right] \mathrm{Z}^{2} \mathrm{R}=\text { Rydberg constant }=109678 \mathrm{~cm}^{-1} \\
& n_{i}=1, n_{f}=2,3,4, \ldots . \quad[\text { Lyman series] (UV) } \\
& n_{i}=2, n_{f}=3,4,5, \ldots . \quad[\text { Balmer series] (VIBGYOR) } \\
& n_{i}=3, n_{f}=4,5,6 \quad \text { [Paschen series] IR. } \\
& n_{i}=4, n_{f}=5,6,7 \quad[\text { Bracket series }] \text { IR. } \\
& n_{i}=5, n_{f}=6,7,8 \quad \text { [Pfund series] IR. }
\end{aligned}
$$

Quantum numbers : The numbers which completely define the state of $e^{-}$in an atom.
(1) Principal Quantum No. : It describes the distance of $e^{-}$from nucleus ' $\boldsymbol{n}$ ' i.e., defines the shell no. It is denoted by ' $n$ '.

$$
\begin{aligned}
n= & 1,2,3,4,5, \ldots . . \\
& \quad \text { K, L, M, N, O } \ldots . .
\end{aligned}
$$

(2) Azimuthal ( $l$ ) 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 $n-1$.

| $n$ | $l(0$ to $n-l)$ |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| 1 | 0 | $l=0$ | $' s '$ | subshell |
| 2 | 0,1 | $l=1$ | $' p \prime$ | subshell of |
| 3 | $0,1,2$ | $l=2$ | $' d '$ | subshell of |
| 4 | $0,1,2,3$ | $l=3$ | $' f '$ | subshell of |

All subshells are wave functions for locating $e^{-}$.
In the same shell energy increase $s<p<d<f$.
(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. $m_{s}=-l \ldots \ldots . .0 . \ldots \ldots .+l=(2 l+1)$.
(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 $e^{-}$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) $1 s<2 s<2 p<3 s<3 p<4 s<3 d<4 p<5 s<4 d<5 p<6 s<4 f<5 d<6 p$ $<7 s<5 f<6 d<7 p$
(B) Half filled and completely filled subshells have more stability than incompletely filled subshells.
$\mathrm{Cr}=[\mathrm{Ar}] 4 s^{1} 3 d^{5}$
$\mathrm{Cu}=[\mathrm{Ar}] 4 s^{1} 3 d^{10}$
(C) As the shell no. inc. size of subshell increases e.g., size of $(2 s>1 s) ;(3 p$ $>2 p) ;(4 d>3 d)$
(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 $=\mathrm{n}-1$.
(E) $\psi(\mathbf{p s i})$

A wave function for locating an electron

## $\psi^{2}(p s i$ square $)$

The square of wave function where the probability of finding the $e^{-}$is maximum.
[Each value of $\psi^{2}$ is a region and defines one orbital]
(F) Orbit
(1) A definite distance from the nucleus for finding the $e^{-}$[ $e^{-}$as a particle].
(1) It has definite size and $e^{-}$in this orbit has definite energy.

## Orbital

(1) A probability region for locating the $e^{-}$around the nucleus. It is a wave function [ $e^{-}$as a wave]
(2) It does not define definite size. But only a boundary region diagram of a wave for locating the $e^{-}$.


## CASE BASED QUESTIONS

## 1. Read the passage given below and answer the following questions:

The capacities of shells with a given principal quantum number are fixed by (1) the rules governing the permitted values of the quantum numbers and (2) the Pauli Exclusion Principle. The permitted values of the quantum numbers are :

| Principal quantum number | $n$ | 1 to $\infty$ |
| :--- | :--- | :--- |
| Azimuthal quantum number | $l$ | 0 to $n-1$ ( $n$ values $)$ |
| Magnetic quantum number | $m_{l}$ | $-l$ to $+l,(2 l+1$ values $)$ |
| Spin quantum number | $m_{s}$ | $-1 / 2$ or $+1 / 2(2$ values $)$ |

The Pauli Exclusion Principle states that no two electrons in the same atom may have the same values of all four quantum numbers. It follows that, for a given value of $n$, there are $2 n^{2}$ different sets of values for the quantum numbers, because $l$ may have the values $\mathrm{O}, 1, \ldots, n-1$, and for each value of $l$ there are $2 l+1$ values of $m_{l}$ and for each set of values of $l$ and $m$ there are just two choices for $\mathrm{m}_{s}$.
(Reference : Thomas H. Hazlehurst, J. Chem. Educ. 1941, 18, 12, 580 Publication Date: December 1, 1941, Journal of American Chemical Society).

The following questions are multiple choice questions. Choose the most appropriate answer:
(i) Based on the concept of quantum numbers mentioned in the study, which of the following value sets of quantum numbers are not possible?
(a) $\quad \mathrm{n}=0, \quad \mathrm{l}=0, \quad \mathrm{~m}_{1}=0, \quad \mathrm{~m}_{\mathrm{s}}=-1 / 2$
(b) $\quad \mathrm{n}=5, \quad \mathrm{l}=3, \quad \mathrm{~m}_{1}=-4, \quad \mathrm{~m}_{\mathrm{s}}=+1 / 2$
(c) $\quad \mathrm{n}=3, \quad \mathrm{l}=1, \quad \mathrm{~m}_{1}=-1, \quad \mathrm{~m}_{\mathrm{s}}=-1 / 2$
(d) $\quad \mathrm{n}=6, \quad \mathrm{l}=1, \quad \mathrm{~m}_{1}=0, \quad \mathrm{~m}_{\mathrm{s}}=+1 / 2$
(ii) What will be the maximum possible number of electrons having $m_{s}=-1 / 2$ for $\mathrm{n}=5$ ?
(a) 50
(b) 25
(c) 32
(d) 72
(iii) Which of the following quantum numbers can distinguish between two electrons present in the same orbital?
(a) Azimuthal quantum number
(b) Principal quantum number
(c) Magnetic quantum number
(d) Spin quantum number
(iv) Maximum number of electrons having $\mathrm{n}=3$ and $1=1$ is-
(a) 14
(b) 6
(c) 10
(d) 2

ANS: I-B, II-B, III-D, IV-B

## 2. Read the passage given below and answer the following questions:

A hydrogen atom consists of an electron orbiting its nucleus. The electromagnetic force between the electron and the nuclear proton leads to a set of quantum states for the electron, each with its own energy. These states were visualized by the Bohr model of the hydrogen atom as being distinct orbits around the nucleus. Each energy state, or orbit, is designated by an integer, $n$ as shown in the figure. The Bohr model was later replaced by quantum mechanics in which the electron occupies an atomic orbital rather than an orbit, but the allowed energy levels of the hydrogen atom remained the same as in the earlier theory.


Spectral emission occurs when an electron transitions, or jumps, from a higher energy state to a lower energy state. To distinguish the two states, the lower energy state is commonly designated as $n^{\prime}$, and the higher energy state is designated as $n$. The energy of an emitted photon corresponds to the energy difference between the two states. Because the energy of each state is fixed, the energy difference between them is fixed, and the transition will always produce a photon with the same energy.
(Reference : Andrew, A. V. (2006). "2. Schrödinger equation". Atomic spectroscopy Introduction of theory to Hyperfine Structure p. 274 ISBN 978-0-387-255736. https://en.wikipedia.org/wiki/Hydrogen_spectral_series).
In these questions (Q. No. (i) to (iv), a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices :
(a) Assertion and reason both are correct statements and reason is correct explanation for assertion.
(b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
(c) Assertion is correct statement but reason is wrong statement.
(d) Assertion is wrong statement but reason is correct statement
i. ASSERTION: The energy states of H -atom are independent of azimuthal quantum number.
REASON: H-atom does not have any inter-electronic repulsion as it has only $1 \mathrm{e}^{-}$.
ii. ASSERTION: A spectral line will be seen for a $2 p_{x}-2 p_{y}$ transition.

REASON: Energy is released when electron drops to lower stationary state.
iii. ASSERTION: For Balmer series of hydrogen spectrum, the value $n_{1}=2$ and $\mathrm{n}_{2}=3,4,5 \ldots$.
REASON: The value of $\mathrm{n}_{2}$ for a line in Balmer series of hydrogen spectrum having the highest wavelength is 6 .
iv. ASSERTION: Electromagnetic radiations of fixed wavelengths are absorbed by the H -atom.
REASON: Radiations corresponding to the energy difference between the two stationary states are absorbed.

ANS:- I-A, II-D, III-C, IV-A

## MULTIPLE CHOICE QUESTIONS (MCQ)

1. Packet of energy is called
(a) Electron
(b) Photon
(c) Position
(d) Proton
2. Orbital which is not possible
(a) $2 p$
(b) 3 d
(c) 3 s
(d) 3 f
3. the magnetic quantum number of an atom 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 atom is related to the
(a) size of the orbital
(b) spin angular momentum
(c) orbital angular momentum
(d) orientation of the orbital in space
5. The designation of an orbital with in $=4$ and $1=3$
(a) 4 s
(b) 4 p
(c) 4 d
(d) 4 f
6. What transition in the hydrogen spectrum would have the same wavelength as the Balmer transition $\mathrm{n}=4$ to $\mathrm{n}=2$ in the $\mathrm{He}^{+}$spectrum?
(a) $\mathrm{n}=4$ to $\mathrm{n}=1$
(b) $\mathrm{n}=3$ to $\mathrm{n}=2$
(c) $\mathrm{n}=3$ to $\mathrm{n}=1$
(d) $\mathrm{n}=2$ to $\mathrm{n}=1$
7. The wave number of first line of Balmer series of hydrogen in $15200 \mathrm{~cm}-1$. The wave number of the first Balmer line of $\mathrm{Li}^{2+}$ ion is
(a) $15200 \mathrm{~cm}^{-1}$
(b) $60800 \mathrm{~cm}^{-1}$
(c) $76000 \mathrm{~cm}^{-1}$
(d) $136,800 \mathrm{~cm}^{-1}$
8. An electron is moving in Bohr's orbit. Its de Broglie wavelength is $\lambda$. What is the circumference of the forth orbit?
(a) $2 / \lambda$
(b) $2 \lambda$
(c) $3 \lambda$
(d) $3 / \lambda$
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) 3 s and 3 p orbitals all have the same energy.
(d) $3 \mathrm{~s}, 3 \mathrm{p}$ and 3 d 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 $\mathrm{Fe}^{2+}(\mathrm{At}$. no. of $\mathrm{Fe}=26)$ 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 $\qquad$ of radiation.
2. The angular momentum of the electron in the 4th energy shell in the hydrogen atom is $\qquad$ .
3. Lines of Balmer series appear in $\qquad$ region.
4. The maximum number of electrons in $\mathrm{Fe}^{3+}$ (At. No. 26) is $\qquad$ .
5. $\mathrm{Li}^{2+}$ and $\mathrm{He}^{+}$ions have spectrum similar to $\qquad$ atom.
6. Bohr's atomic theory is not able to explain the atomic spectra of atoms containing $\qquad$ electron.
7. An electron in the first shell will have $\qquad$ stability and $\qquad$ 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 $\qquad$
9. According to $\qquad$ no two electrons in an atom will have all the four quantum numbers $\qquad$
10. When there are two electrons in the same orbital they have $\qquad$ spins.
11. The s-subhells have $\qquad$ shape and the p-subshells have $\qquad$
12. The maximum number of electrons on a subshell is equal to $\qquad$ where $1=$ $\qquad$
Ans. 1. Planck's theory
13. $\frac{2 h}{\pi}$
14. Visible
15. 23
16. H -atom
17. more than 1
18. Larger, lower
19. orbital
20. Pauli exclusion principle; similar
21. 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. $\mathrm{Fe}^{2+}$ ion has more number of unpaired electrons than $\mathrm{Fe}^{3+}$.
5. The outer electronic configuration of chromium atom is $3 \mathrm{~d}^{4} 4 \mathrm{~s}^{2}$.
6. The designation of an orbital $n=4$ and $l=0$ is $4 s$.
7. All photons of light have same energy.
8. $\mathrm{Fe}^{3+}$ has $3 \mathrm{~d}^{5}$ 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 $\mathrm{He}^{+}$ion has radius as the first orbit in hydrogen atom.
12. Heisenberg principle is applicable to microscopic particles.
13. 3 s orbital has 2 radial nodes.
Ans. 1. (T)
14. (T)
15. (T)
16. (F)
17. (F)
18. (T)
19. (F)
20. $(\mathrm{T}) \quad$ 9. $(\mathrm{F}) \quad$ 10. $(\mathrm{T}) \quad$ 11. $(\mathrm{T}) \quad$ 12. $(\mathrm{T}), \quad 13 .(\mathrm{T})$

## MATCH THE COLUMNS

1. Match the following

## List-I

a. Lyman series
b. Balmer series
c. Paschen series
d. Brackett series

## List-II

p. Visible region
q. Infrared region
r. Absorption spectrum
s. Ultraviolet region
2. Match the following

## List-I

a. Principal quantum number
b. Azimuthal quantum number
c. Magnetic quantum number
d. Spin quantum number

## List-II

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

## List-III

i. $-l$ to $+l$
ii. 0 to $\infty$
iii. $\pm 1 / 2$
iv. 0 to $(\mathrm{n}-1)$
3. Match the following

## List-I

a. 2 s
b. $2 \mathrm{p}_{\mathrm{x}}$
c. $3 \mathrm{~d}_{\mathrm{xy}}$
d. $3 \mathrm{~d}_{\mathrm{z}^{2}}$

## List-II

p. DOUGHNUT shape i. along z -axis
q. Spherical
r. Dumb bell
s. Double dumb bell

## List-III

ii. In between $\mathrm{x} \& \mathrm{y}$-axis
iii. non-directional
iv. along x -axis
4. Match the following

## List-I

a. 2 s
b. $\psi^{2}$
c. Heisenberg's uncertainty
d. $3 \mathrm{~d}_{\mathrm{yz}}$

## List-II

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

Ans.: 1. a. (s), b. (p), c. (q), d. (q)
2. a. (q). (ii), b. (s). (iv), c. (r). (i), d. (p). (iii)
3.
a. (q). (iii), b. (r). (iv),
c. (s). (ii), d. (p).(i)
4. a. (q), b. (r), c. (s), d. (p)

## ASSERTION AND REASON TYPE QUESTIONS

## Directions: (Questions 1 to 10)

A. Both Assertion \& Reason are true and the reason is the correct explanation of the assertion.
B. Both Assertion \& Reason are true but the reason is not the correct explanation of the assertion.
C. Assertion is true statement but Reason is false.
D. Assertion is false but Reason is true.

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 $3 \mathrm{~d}_{\mathrm{xy}}$.

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.
5. Assertion: Black body is an ideal body that emits and absorbs radiations of all frequencies.

Reason: The frequency of radiation emitted by a body goes from a lower frequency to higher frequency with an increase in temperature.
6. Assertion: 2 p orbitals do not have any radial nodes.

Reason: The number of radial nodes in p -orbitals is given by ( $\mathrm{n}-2$ ) where n is the principal quantum number.
7. Assertion: The opposite lobes of a p-orbital have opposite sign whereas opposite lobes of d-orbital have the same sign.

Reason: The opposite lobes of a p-orbital have opposite charge whereas the opposite lobes of d-orbital have the same charge.
8. Assertion: Electronic configurations of $\mathrm{Cr}^{3+}$ (containing 21 electrons) is same as that of $\mathrm{Sc}(\mathrm{Z}=21)$ i.e., isoelectronic species have the same electronic configuration.

Reason: Orbitals of atoms are filled in order of increasing energy following aufbau principle.
9. Assertion: Hydrogen has one electron in its orbit but it produces several spectral lines.

Reason: There are many excited energy levels available.
10. Assertion: The free gaseous Cr atom has six unpaired electrons.

Reason: Half-filled d-orbitals have greater stability.
$\begin{array}{lllllllll}\text { Ans. 1.A } & \text { 2. } & \text { 3.A } & \text { 4.A } & \text { 5.B } & \text { 6.D } & \text { 7. } \mathrm{C} & \text { 8.D } & \text { 9.A }\end{array}$ 10.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?
8. Which of the following orbitals are possible ?
$1 \mathrm{p}, 2 \mathrm{~s}, 2 \mathrm{p}$ and 3 f
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. What is uncertain in uncertainty principle ?
9. Can a moving cricket ball have a wave character ? Justify your answer.
10. Heisenberg uncertainty principle has no significance in our everyday life. Explain.
11. Why uncertainty in position is more when uncertainty in velocity is less for an electron?
12. What are the four quantum numbers of 19th electron of copper?
$($ Given : Atomic number of copper $=29)$
13. How many electrons will be present in the sub-shells having ms, value of $-1 / 2$ for $n=4$ ?
14. Write the electronic configuration of $\mathrm{Ni}^{3+}$. (At. No . of $\mathrm{Ni}=28$ )
15. How many radial and angular nodes are present in 2 p orbital.
16. Mention the physical significance of $\Psi$ and $\Psi^{2}$.
[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 $\mathrm{He}^{+}$spectrum?
[Ans. $n_{1}=1$ and $n_{2}=2$ ]
Q. 5. What transition of $\mathrm{Li}^{2+}$ spectrum will have the same wavelength as that of the second line of Balmer series in $\mathrm{He}^{+}$spectrum ?
[Ans. $n_{2}=6$ to $\left.n_{1}=3\right]$
Q. 6. Calculate the energy required for the process

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

The ionization energy for the H atom in the ground state is $2.18 \times$ $10^{-18} \mathrm{~J} \mathrm{atom}^{-1}$ ]
[Ans. $8.72 \times 10^{-18} \mathrm{~J}$ ]
Q. 7. Calculate the wave number for the longest wavelength transition in the Balmer series of atomic hydrogen. [Ans. $1.523 \times 10^{6} \mathrm{~m}^{-1}$ ]
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 $\mathrm{He}^{+}$. What is the radius of this orbit? [Ans. - $54.38 \mathrm{eV}, 0.2645 \AA$ ]
Q. 10. What is the distance of separation between 3 rd and 4 th 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 $6000 \AA$.
Q. 13. Which one has a higher energy, a photon of violet light with wavelength $4000 \AA$ or a proton of red light with wavelength $7000 \AA$ ?

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

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 $10^{15} \mathrm{~Hz}$ ?
Q. 17. Find the energy of electron in 4th shell of $\mathrm{Li}^{2+}$ ion.
Q. 18. What is the wave number of an electron with shortest wavelength radiation in Lyman spectrum of $\mathrm{He}^{+}$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 $3.6 \AA$.
[Ans. $6.135 \times 10^{-29} \mathrm{~kg}$ ]
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 $=10^{-15} \mathrm{~m}$ )
[Hint: Taking $10^{-15} \mathrm{~m}$ as $\Delta x$, the $\Delta v$ 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 $2 s$ and $2 p_{x}$ orbitals and 1 s and $2 s$ orbitals?
Q. 28. Draw shape of $d_{x^{2}-y^{2}}$ 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) $2 p$
(b) $4 f$
Q. 32. Write down the electronic configuration of $\mathrm{Fe}^{3+}$ and $\mathrm{Ni}^{2+}$. How many unpaired electrons are present? (Given Atomic number, $\mathrm{Fe}=26$, $\mathrm{Ni}=28$ ).
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.
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 2 p subshell of nitrogen remain unpaired.
(b) Cr has configuration $3 d^{5} 4 s^{1}$ and not $3 d^{4} 4 s^{2}$.
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 $\mathrm{N}^{3-}$ ion.
Q. 37. Which one is having higher energy?
(a) Last electron of $\mathrm{Cl}^{-}$or last electron of $\mathrm{O}^{2-}$.
(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 $-2.18 \times 10^{-18} \mathrm{~J}$ atom $^{-1}$. What is the energy associated with the fourth orbit?
(ii) Calculate the radius of Bohr's third orbit for hydrogen atom.

$$
\left[\text { Ans. }-1.36 \times 10^{-19} \mathrm{~J} \text { atom }^{-1} \cdot 4.761 \mathrm{~nm}\right]
$$

Q. 2. A bulb emits light of wave length $4500 \AA$. 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 $1.05 \times 10^{5} \mathrm{~J} \mathrm{~mol}^{-1}$ 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. $a=3.2255 \times 10^{19} \mathrm{~J}, b=616 \mathrm{~nm}$ ]
Q. 4. Compare the frequency of light radiations emitted when electron falls from 5th shell to the 2nd shell in $\mathrm{Li}^{2+}$ ion and electron falls from 4th shell to the 1st shell in $\mathrm{He}^{+}$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 $\mathrm{CH}_{4}$ and $\mathrm{O}_{2}$ 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 $\left.=1.6 \times 10^{-19} \mathrm{~J}\right]$
[Ans. $3.87 \times 10^{-11} \mathrm{~m}$ ]
Q. 8. (i) Discuss the similarities and differences between a 1 s and 2 s orbital.
(ii) Draw the shape of $\mathrm{d}_{\mathrm{z}^{2}}$.
Q. 9. Calculate the wavelength of a tennis ball of mass 60 gm moving with a velocity of 10 m per second. $\quad\left(h=6.626 \times 10^{-34} \mathrm{~kg} \mathrm{~m}^{2} \mathrm{~s}^{-1}\right)$
[Ans. $10^{-3}$ metre]
Q. 10. Calculate the wavelength of 1000 kg rocket moving with a velocity of $3000 \mathrm{~km} / \mathrm{hr} . \quad\left(h=6.626 \times 10^{-34} \mathrm{~kg} \mathrm{~m}^{2} \mathrm{~s}^{-1}\right)$
[Ans. $7.9512 \times 10^{-40} \mathrm{~m}$ ]
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 $1 \AA$.
[Ans. $3.5 \times 10^{-24} \mathrm{~m} \mathrm{~s}^{1}$ ]
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 $\lambda_{A}=5 \times 10^{-10} \mathrm{~m}$ and $\lambda_{B}=$ $10 \times 10^{10} \mathrm{~m}$. 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 $\mathrm{He}^{+}$has angular momentum of $5 h / 2 \pi$. Find its energy and wavelength associated with it. Find the kinetic energy of this electron.
Q. 16 (i) An atomic orbital has $n=2$. What are the possible values of $l$ and $m_{l}$ ?
(ii) List the quantum numbers ( $m_{1}$ 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) $\mathrm{H}^{-}$
(b) $\mathrm{Na}^{+}$(c) $\mathrm{O}^{2-}$
(d) $\mathrm{F}^{-}$
(ii) What are the atomic numbers of elements whose outermost electrons are represented by (a) $3 s^{1}$ (b) $2 p^{3}$ and (c) $3 p^{5}$ ?
(iii) Which atoms are indicated by the following configurations?
(a) $[\mathrm{He}] 2 s^{1}$
(b) $[\mathrm{Ne}] 3 s^{2} 3 p^{3}$
(c) $[\mathrm{Ar}] 4 s^{2} 3 d^{1}$.
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 $6800 \AA$. Calculate threshold frequency $\left(v_{0}\right)$ and work function $\left(\mathrm{W}_{\mathrm{o}}\right)$ of the metal.
[Ans. $v_{o}=4.41 \times 10^{14} \mathrm{~s}^{-1} \mathrm{~W}_{\mathrm{o}}=2.92 \times 10^{-19} \mathrm{~J}$ ]
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 $\mathrm{He}^{+}$and $\mathrm{Li}^{2+}$ ions ?
[Ans. $\mathrm{E}_{1}$ of $\mathrm{He}^{+}=-54.4 \mathrm{eV}, \mathrm{E}_{1}$ of $\mathrm{Li}^{2+}=-122.4 \mathrm{eV}$ ]
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 $\left(\lambda_{0}\right)$ and threshold frequency $\left(v_{0}\right)$ of this light radiation. If $\mathrm{C} s$ 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 $2.0 \times 10^{3} \mathrm{~m} \mathrm{~s}^{-1}$. Find the wavelength of the particle constituting the beam

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

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 $5.5 \times 10^{-20} \mathrm{~ms}^{-1}$.
[Ans. $9.55 \times 10^{10} \mathrm{~m}$ ]
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) $\mathrm{Fe}^{3+}$ (ii) $\mathrm{Cu}^{+}$
[Given Atomic number of Fe and Cu are 26 \& 29]
Q. 8.(a) Draw the shapes of the following orbitals.
(i) $3 d_{x y}$ (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) $n=1, l=0$
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:

$$
(2 l+1) \text { or }(4 l+1) \text { or } n^{2}
$$

Q. 10.(a) A neutral atom has $2 \mathrm{~K}, 8 \mathrm{~L}$ 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 :
$\mathrm{Al}^{+}, \mathrm{Cr}^{2+}, \mathrm{Co}^{3+}$ and $\mathrm{Mn}^{2+}$
(Given Atomic number : $\mathrm{Al}=13, \mathrm{Cr}=24, \mathrm{Co}=27 \& \mathrm{Mn}=25$ )
(c) One electron is present in 4 f 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 $14^{\text {th }}$ electron in an atom.
(b) Increasing order of filling electron in $4 f, 5 p$ and $6 d$ 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-I

Time allowed : 1 Hour

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
(a) 4 s
(b) 4 p
(c) 4 d
(d) 4 f
2. Maximum number of unpaired electrons in chromium is (Given: Atomic number of $\mathrm{Cr}=24$ )
(a) 4
(b) 5
(c) 6
(d) 7
3. Which series of lines of the hydrogen spectrum lie in the visible region'?
4. Why de-Broglie's wavelength is not significant for macroscopic objects.(1)
5. Which of the following is not possible ?
(a) 2 p
(b) 3 d
(c) 3 f
(d) 4 p
6. Write two difference between orbit and orbital.
7. Calculate the wave number for the longest wavelength transition in the paschen series of atomic hydrogen.
8. (a) How many orbitals are associated with $n=4$ ?
(b) How many electrons will be present in the sub-shells having ms value of $-1 / 2$ for $n=3$ ?
(c) Draw the shape of $\mathrm{d}_{\mathrm{z}}^{2}$.
9. Calculate the uncertainty in the position of a dust particle with mass equal to 1 mg if the uncertainty in its velocity is $5.5 \times 10^{-20} \mathrm{~ms}^{-1}$.
10. (i) The energy associated with the first orbit in the hydrogen atom is $-2.18 \times 10^{-18} \mathrm{~J}^{2}$ atom $^{-1}$. What is the energy associated with the fifth orbit?
(ii) Calculate the radius of Bohr's fifth orbit for hydrogen atom.
(iii) Calculate the radial and angular nodes in 2 p orbital.
(iv) Define the black body and black body radiations.

## UNIT TEST-II

Time allowed : 1 Hour

General instructions :
(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. The de-Broglie wavelength associated with a ball of mass 1 kg having kinetic energy 0.5 j is
(a) $6.626 \times 10^{-34} \mathrm{~m}$
(b) $13.20 \times 10^{-34} \mathrm{~m}$
(c) $10.38 \times 10^{-21} \mathrm{~m}$
(d) $6.626 \times 10^{-34} \AA$
2. The radius of which of the following orbit is same as that of first orbit of hydrogen atom?
(a) $\mathrm{He}^{+}(\mathrm{n}=2)$
(b) $\mathrm{Li}^{2+}(\mathrm{n}=2)$
(c) $\mathrm{Li}^{2+}(\mathrm{n}=3)$
(d) $\mathrm{Be}^{3+}(\mathrm{n}=2)$
3. Which series of hydrogen spectrum lies in the UV region'?

In following questions a statement of Assertion followed by a statement of Reason is given. Choose the correct answer out of the following choices.
(a) Assertion and Reason are true and Reason is the correct explanation of Assertion.
(b) Assertion and Reason are true but Reason is not the correct explanation of Assertion.
(c) Assertion is true but Reason is false.
(d) Both Assertion and Reason are false.
4. Assertion : It is impossible to determine the exact position and exact momentum of an electron simultaneously.

Reason : The path of an electron in an atom is clearly defined.
5. Assertion : All isotopes of a given element show the same type of chemical behaviour.

Reason: The chemical properties of an atom are controlled by the number of electrons in the atom.
6. Calculate the number of angular nodes and radial nodes in $3 p$ orbital. (2)
7. Calculate the mass of photon with wavelength $3.6 \mathrm{~A}^{\circ}$.
8. What transition in the hydrogen spectrum would have the same wavelength as the Balmer transition $\mathrm{n}=4$ to $\mathrm{n}=2$ of $\mathrm{He}^{+}$spectrum?
9. (a) The energy associated with Bohr's first orbit is $-2.18 \times 10^{-18} \mathrm{~J}^{2}$ atom $^{-1}$. What is the energy associated with fifth orbit?
(b) The work function for Caesium atom is 1.9 eV . Calculate the threshold wavelength.

$$
\left[\text { Given : lev }=1.6 \times 10^{-19} \mathrm{~J}\right]
$$

(c) How many sub-shells are associated with $\mathrm{n}=4$ ?
10. (i) How many electrons will present in sub-shell having spin quantum number value of $-1 / 2$ for $n=4$ ?
(ii) Which of the following transition will have minimum wavelength and why?

$$
\begin{equation*}
\mathrm{n}_{4} \rightarrow \mathrm{n}_{1}, \quad \mathrm{n}_{4} \rightarrow \mathrm{n}_{2}, \mathrm{n}_{2} \rightarrow \mathrm{n}_{1} \tag{5}
\end{equation*}
$$

(iii) Give the number of radial nodes for 3 s and 2 p orbitals.

# 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- 2 are called Alkaline earth metals. Group-15 are called Pnicogens Group-16 are called Chalcogens Group-17 are called Halogens Group-18 are called Noble gases
3. $1^{\text {st }}$ period -2 elements
$4^{\text {th }}$ and $5^{\text {th }}$ period -18 elements
$2^{\text {nd }}$ and $3^{\text {rd }}$ period -8 elements
$6^{\text {th }}$ period -32 elements
$7^{\text {th }}$ period - Incomplete ( 32 elements)
4. Groups

1 and 2 - ' $s$ ' block elements last electron entered in ' $s$ ' subshell $\left[s^{1}, s^{2}\right]$ 3 to 12 - ' $d$ ' block elements last electrons entered in ' $d$ ' subshell [ $d^{1}$ to $d^{10}$ ]. 13 to 18 - ' $p$ ' block elements last electrons enter in ' $p$ ' subshell $\left[p\right.$ ' to $\left.p^{6}\right]$. 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 [ $4 f$ lanthanoids, $5 f$ actinoids]
6. General outer electronic configuration
' $\boldsymbol{s}$ ' block : $n s^{1}, n s^{2}$ [Group 1 to 2]
' $p$ ' block: $n s^{1} n p^{1}$ to $n s^{2} n p^{6}$ Group 13 to 18
' $d$ ' block : $n s^{0-2}(n-1) d^{1 \text { to }}{ }^{10}$ Group 3 to 12
' $f$ 'block : $(n-2) f^{1 \text { to }}{ }^{14}(n-1) d^{0,1} n s^{2}$
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) Cation radius < Atomic radius - due to more no. of protons than number of electron coloumbic force increases, size decreases.

$$
\left[\mathrm{Mg}^{2+}<\mathrm{Mg}^{+}<\mathrm{Mg}\right]
$$

(B) Anion radius > Atomic radius - Due to more number of electron than number of protons

$$
\left[\mathrm{N}^{3-}>\mathrm{O}^{2-}>\mathrm{F}^{-}\right]
$$

Electron-Electron repulsion increase, coloumbic force of attraction decreases.
(C) For Isoelectronic species - More is the charge of cation lesser the size. More is the charge of anion, more is the size.
(D) Order of size $-\mathrm{O}^{2-}>\mathrm{F}^{-}>\mathrm{Na}>\mathrm{Na}^{+}>\mathrm{Mg}^{2+}$

## 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.

$$
\begin{aligned}
\mathrm{M}(\mathrm{~g})+\text { Energy } & \longrightarrow \mathrm{M}^{+}+e^{-} \\
\mathrm{IE}_{3}>\mathrm{IE}_{2} & >\mathrm{IE}_{1}
\end{aligned}
$$

(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.
I ionisation enthalpy $\mathrm{Li}<\mathrm{B}<\mathrm{Be}<\mathrm{C}<\mathrm{O}<\mathrm{N}<\mathrm{F}<\mathrm{Ar}$
II ionisation enthalpy $\mathrm{Be}<\mathrm{C}<\mathrm{B}<\mathrm{N}<\mathrm{F}<\mathrm{O}<\mathrm{Ne}$

## (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.

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 $\mathbf{s}>\mathbf{p}>\mathbf{d}>\mathbf{f}$ $\mathrm{d} \& \mathrm{f}$ 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.

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

## 11. Electron gain enthalpy:

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

$$
\mathrm{E}(\mathrm{~g})+e^{-} \longrightarrow \mathrm{E}^{-}(\mathrm{g})
$$

- 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 $\mathrm{F}>\mathrm{O}>\mathrm{N}$.

## Maximum electronegative Assign to $\mathbf{F}$.

* 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: $\mathbf{Z n}, \mathbf{C d}, \mathbf{H g}$
* 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

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

$$
\mathrm{Na}_{3}\left[\mathrm{Al}(\mathrm{OH})_{6}\right] \text { exists but } \mathrm{Na}\left[\mathrm{~B}(\mathrm{OH})_{4}\right] \text { not exists. }
$$

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.

| Elements with number of $\mathrm{e}^{-}$ | in valance shell |
| :--- | :--- |
| (a) $1,2,3$ | metals |
| (b) 4 | metalloids |
| (c) $5,6,7$ | non-metals |
| (d) 8 | noble gas |



| MIND MAP |
| :---: |
| CLASSIFICATION OF ELEMENTS AND PERIODICITY IN PROPERTIES |



## CASE BASED STUDY QUESTIONS

## 1. Read the passage given below and answer the following questions:

A period is a horizontal row in the periodic table. Although groups generally have more significant periodic trends, there are regions where horizontal trends are more significant than vertical group trends, such as the f-block, where the lanthanides and actinides form two substantial horizontal series of elements.

Elements in the same period show trends in atomic radius, ionization energy, electron affinity, and electronegativity. Moving left to right across a period, atomic radius usually decreases. This occurs because each successive element has an added proton and electron, which causes the electron to be drawn closer to the nucleus. This decrease in atomic radius also causes the ionization energy to increase when moving from left to right across a period. The more tightly bound an element is, the more energy is required to remove an electron. Electronegativity increases in the same manner as ionization energy because of the pull exerted on the electrons by the nucleus. Electron affinity also shows a slight trend across a period. Metals (left side of a period) generally have a lower electron affinity than non-metals (right side of a period), with the exception of the noble gases.
(Reference: https://en.wikipedia.org/wiki/Periodic_table)
The following questions are multiple choice questions. Choose the most appropriate answer:
(i) The atomic radii of Elements Z and X are compared. Element Z is having larger radius than Element X . (Both the elements does not have noble gas configuration and exist in same period in the periodic table)

Based on this you can say that the:
A) Element Z is located to the left side of Element X in the periodic table
B) Element Z is located to the right side of Element X in the periodic table
C) Element Z and X are probably in the same group
D) None of the above
(ii) In which of the following atoms is the 3 s orbital closest to the nucleus?
A) Br
B) Cl
C) I
D) Same distance in all of these atoms
(iii) $\qquad$ have the lowest first ionization energies of the groups listed.
A) Alkali metals
B) Transition metals
C) Halogens
D) Noble gases
(iv) The correct order of electronegativity is
A) $\mathrm{Cl}>\mathrm{F}>\mathrm{O}>\mathrm{Br}$
B) $\mathrm{F}>\mathrm{O}>\mathrm{Cl}>\mathrm{Br}$
C) $\mathrm{F}>\mathrm{Cl}>\mathrm{Br}>\mathrm{O}$
D) $\mathrm{O}>\mathrm{F}>\mathrm{Cl}>\mathrm{Br}$

ANS:- I-A, II-B, III-C, IV-B

## 2. Read the passage given below and answer the following questions:

As the number of protons increase within a period (or row) of the periodic table, the first ionization energies of the transition-metal elements are relatively steady, while that for the main-group elements increases. The effective nuclear charge mirrors and may explain the periodic trends in the first ionization energies of the transition-metal and main-group elements. The differing periodic trends in the effective nuclear charge are due to a greater increase in shielding in the transitionmetal elements than in the main-group elements. The difference in shielding is due to the entry of electrons into an inner-shell orbital for the transition-metal elements, while electrons enter an outer-shell orbital for the main-group elements.
(Reference: Paul S. Matsumoto J. Chem. Educ. 2005, 82, 11, 1660 Publication Date: November 1, 2005, Journal of the American Chemical Society)
Q.1. Why the first Ionisation energy of $B e$ is greater than that of $B$ ?
Q.2. Why the ionisation of s-electron require more energy than iorisation of p-electron of the same shell.
Q.3. The first iorisation enthalpy of aluminium is lower than that of magnesium. Why?
Q.4. Why the first iorisation energies of transition metal elements are relatively steady?

## 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 $\mathrm{N}^{3-}, \mathrm{O}^{2-}, \mathrm{Na}^{+}$and $\mathrm{F}^{-}$is
(a) $\mathrm{Na}^{+}<\mathrm{F}^{-}<\mathrm{O}^{2-}>\mathrm{N}^{3-}$
(b) $\mathrm{F}^{-}<\mathrm{O}^{2-}<\mathrm{N}^{3}>\mathrm{Na}^{+}$
(c) $\mathrm{O}^{2-}<\mathrm{N}^{3-}<\mathrm{F}^{-}>\mathrm{Na}^{+}$
(d) $\mathrm{N}^{3-}<\mathrm{Na}^{+}<\mathrm{F}^{-}>\mathrm{O}^{2-}$
4. The basic strength of the oxides follows the order
(a) $\mathrm{Al}_{2} \mathrm{O}_{3}>\mathrm{MgO}>\mathrm{Na}_{2} \mathrm{O}$
(b) $\mathrm{Al}_{2} \mathrm{O}_{3}<\mathrm{MgO}<\mathrm{Na}_{2} \mathrm{O}$
(c) $\mathrm{Na}_{2} \mathrm{O}_{3}<\mathrm{MgO}>\mathrm{Al}_{2} \mathrm{O}_{3}$
(d) $\mathrm{Al}_{2} \mathrm{O}_{3}>\mathrm{MgO}>\mathrm{Na}_{2} \mathrm{O}$
5. The correct order of the size of C, N, P, S follows the order
(a) $\mathrm{N}<$ 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) $\mathrm{Na}_{2} \mathrm{O}$
(b) $\mathrm{Al}_{2} \mathrm{O}_{3}$
(c) $\mathrm{P}_{2} \mathrm{O}_{5}$
(d) $\mathrm{SO}_{3}$
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 electronegativity of the following elements increase in the order
(a) $\mathrm{C}, \mathrm{N}, \mathrm{Si}, \mathrm{P}$
(b) N, Si, C, P
(c) $\mathrm{Si}, \mathrm{P}, \mathrm{C}, \mathrm{N}$
(d) $\mathrm{P}, \mathrm{Si}, \mathrm{N}, \mathrm{C}$
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 $\qquad$ .
2. In the periodic table, horizontal rows are known as $\qquad$ .
3. Elements of s-blocks and p-blocks are collectively called $\qquad$ .
4. Most electropositive elements belong to $\qquad$ group.
5. Most electronegative elements belong to $\qquad$ group.
6. The elements above atomic number 92 are called $\qquad$ .
7. The inner-transition elements belong to $\qquad$ block of the periodic table and are shown separately at the $\qquad$ of the periodic table.
8. An element having electronic configuration [Ar] $3 \mathrm{~d}^{5}, 4 \mathrm{~s}^{2}$ belongs to
$\qquad$ block.
9. $\mathrm{Ca}^{2+}$ has smaller ionic radius than $\mathrm{K}^{+}$ion because it has $\qquad$ .
10. The maximum electronegativity is shown by $\qquad$ .

11 The maximum ionisation enthalpy is shown by $\qquad$ .

12 The cation is $\qquad$ and the anion is $\qquad$ than the parent atom.

Ans: 1. Lithium
7. $\mathrm{F}-$, bottom
2. periods
3. normal elements or
8. $\mathrm{s}-$
9. more protons representative elements
4. $1^{\text {st }}$
10. F-
5. $17^{\text {th }}$
11. H
6. transuranic elements
12. smaller, bigger

## 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.
$5 \quad \mathrm{Mg}^{2+}$ ion has smaller size than Mg .
5. Electronegativity of F is larger than that of Cl but electron gain enthalpy of Cl is larger than of F .
6. The decreasing order of electronegativity of $\mathrm{F}, \mathrm{O}$ and N is $\mathrm{F}>\mathrm{O}>\mathrm{N}$.
7. Group-18 contain maximum gaseous elements.
8. $\mathrm{Al}_{2} \mathrm{O}_{3}$ is an amphoteric oxide.
9. Helium has the highest ionisation enthalpy.

| Ans: | 1. (T) | 2. $(\mathrm{T})$ | 3. (T) | 4. $(\mathrm{F})$ | 5. (T) |
| :--- | :--- | :--- | :--- | :--- | :--- |
|  | 6. (T) | 7. $(\mathrm{T})$ | 8. (T) | 9. $(\mathrm{T})$ | 10. $(\mathrm{T})$ |
|  |  |  | MATCH THE COLUMNS |  |  |

1. 

## Column A

a. Lightest element
b. Lightest metal
c. Heaviest metal
d. Most reactive metal

## Column B

i. Caesium
ii. Osmium
iii. Lithium
iv. Hydrogen
p. $\mathrm{Is}^{1}$

## Column C

q. $[\mathrm{He}] 2 \mathrm{~s}^{1}$
r. $\quad[\mathrm{Xe}] 6 \mathrm{~s}^{1}$
s. d-block element

## Column A

a. Fluorine
b. Helium
c. Chlorine
d. Caesium

## Column B

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

Column C
p. $\quad[\mathrm{Xe}] 6 \mathrm{~s}^{1}$
q. $\quad[\mathrm{He}] 2 \mathrm{~s}^{2} 2 \mathrm{p}^{5}$
r. $\quad \mathrm{Is}^{2}$
s. $[\mathrm{Ne}] 3 \mathrm{~s}^{2} 3 \mathrm{p}^{5}$
3.

## Column A

a. $\quad \mathrm{Na}_{2} \mathrm{O}$
b. $\mathrm{Cl}_{2} \mathrm{O}_{7}$
c. $\mathrm{Al}_{2} \mathrm{O}_{3}$
d. CO
4.

## Column A

a. s \& p-block
b. d-block
c. f-block
d. group-1 and group-2

## Column B

i. Amphoteric oxide
ii. Acidic oxide
iii. Neutral oxide
iv. Basic oxide
Ans: 1.
a. (iv). (p), b.
. (iii). (q), c. (ii). (s), d. (i). (r)
2. a. (iii). (q), b. (iv). (r), c. (i). (s), d. (ii). (p)
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.1-10

A Both Assertion \& Reason are true and the reason is the correct explanation of the assertion.
B Both Assertion \& Reason are true but the reason is not the correct explanation of the assertion.
C Assertion is true statement but Reason is false.
D Assertion is false but Reason is true.

1. Assertion : Ionic radius of $\mathrm{Na}^{+}$is smaller than Na

Reason : Effective nuclear charge of $\mathrm{Na}^{+}$is higher than Na
2. Assertion : First ionisation enthalpy of N is higher than O .

Reason : Extra stability of fully filled up 2 p 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.
5. Assertion: Noble gases have positive electron gain enthalpy.

Reason: Noble gases have stable closed shell electronic configuration.
6. Assertion: F is more electronegative than Cl .

Reason: F has more electron affinity than Cl .
7. Assertion: The ionic size of $\mathrm{O}^{2-}$ is bigger than that of $\mathrm{F}^{-}$ion.

Reason: $\mathrm{O}^{2-}$ and $\mathrm{F}^{-}$are isoelectronic ions.
8. Assertion: The ionic radii follows the order: $\mathrm{I}^{-}<\mathrm{I}<\mathrm{I}^{+}$.

Reason: Smaller the value of z/e, larger the size of the species.
9. Assertion: The first ionisation enthalpy of aluminium is lower than that of magnesium.
Reason: Ionic radius of aluminium is smaller than that of magnesium.
10. Assertion: First ionisation energy for nitrogen is higher than that of oxygen. Reason: Across a period effective nuclear charge decreases.

Ans: | $1 . \mathrm{A}$ | 2. A | 3. B | 4. A | 5.A | 6.C | 7.B | 8.D | 9.B |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | 10.C

## 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 $\mathrm{Na}^{+}$?
[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 $\left[\mathrm{AlCl}_{4}\right]^{-}$?
[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

1. 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 $\mathrm{ns}^{2}, \mathrm{np}^{5}$ configuration is non-metal or metal?
5. Define van der Waal's radius.
6. Write the outer shell configuration of atomic number 31. [Ans. $4 \mathrm{~s}^{2}, \mathrm{p}^{1}$ ]
7. Find the group number and period number of element having atomic number 52.
[Ans. Period $=5$ th, Group $=16$ th $]$
8. Arrange $\mathrm{O}^{2-}, \mathrm{O}^{-1}, \mathrm{O}$ in decreasing radius (size). [Ans. $\mathrm{O}^{2-}>\mathrm{O}^{-1}>\mathrm{O}$ ]
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 4 f series elements.
[Ans. $6 \mathrm{~s}^{2}, 5 \mathrm{~d}^{0-1}, 4 \mathrm{f}^{1}$ to 14 ]
12. Write two isoelectronic species with Br (35).
[Ans. $\mathrm{Kr}^{+}, \mathrm{Se}^{-1}$ ]
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 $\mathrm{F}>\mathrm{Cl}>\mathrm{Br}>\mathrm{I}$, 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 $\mathrm{O}^{-}$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?
7. Name a halogen, a metal and a group 13 element which are liquid at $30^{\circ} \mathrm{C}$.
[Ans. $\mathrm{Br}, \mathrm{Hg}, \mathrm{Ga}$ ]
8. The reducing power of elements increases down the group but reverse is true for oxidising power along a period. Why ?
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, $\mathrm{F}, \mathrm{Br}, \mathrm{Cl}$
11. Oxygen is more non-metallic than nitrogen but less than fluorine why?
12. $\mathrm{LiCl}, \mathrm{LiBr}, \mathrm{LiI}$ are covalent as well as ionic why?
13. $\mathrm{PbCl}_{2}$ is more stable than $\mathrm{PbCl}_{4}$. Why ?
[Ans. Inert pair effect]
14. [Magnesium and Lithium both form nitrides why?
15. Which has least I.E. $\left[3 p^{3}, 3 p^{6}, 2 p^{3}, 2 p^{6}\right]$ ?
16. (a) I.E. of sulphur is lower than chlorine.
(b) Arrange the following in decreasing order of their electro-negativity: $\mathrm{F}, \mathrm{O}, \mathrm{N}, \mathrm{Cl}, \mathrm{C}, \mathrm{H}$.
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 $\mathrm{Mg}, \mathrm{Na}, \mathrm{Al}, \mathrm{Si}$.
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 $4 s^{2} 4 p^{3}$. 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 $\mathrm{LiF}, \mathrm{NaF}, \mathrm{KF}, \mathrm{RbF}, \mathrm{CsF}$ are ionic ? But LiF is less ionic than CsF .
7. (a) Why Ca has larger atomic radius than Al ?
(b) Why $2 s^{2}$ electron is difficult to remove than $2 p$ electron?
8. (a) Why the compounds of group 17 with group 13 elements are more ionic and stable than with (group 1) elements? (b) $\mathrm{Na}_{2} \mathrm{O}$ is more ionic than $\mathrm{Li}_{2} \mathrm{O}$. why?
9. Explain the following data : Ionisation energy $\mathrm{Cl}<\mathrm{H}<\mathrm{O}<\mathrm{N}<\mathrm{F}$.
10. $\mathrm{IE}_{2}$ of $3^{\text {rd }}$ period elements is as follows. Why ?
$\mathrm{Mg}<\mathrm{Si}<\mathrm{Al}<\mathrm{P}<\mathrm{S}<\mathrm{Cl}<\mathrm{Ar}<\mathrm{Na}$.
11. Account fot the following:
(a) Halogens have very high negative electron gain enthalpy
(b) The electron gain enthalpy of $\mathrm{Cl}(\mathrm{Z}=17)$ 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 $\mathrm{Li}<\mathrm{B}<\mathrm{Be}<\mathrm{C}<\mathrm{O}<\mathrm{N}<\mathrm{F}<\mathrm{Ne}$. Explain why?
(i) Be has higher $\left(\Delta_{\mathrm{i}} \mathrm{H}\right)_{1}$ than B
(ii) O has lower $\left(\Delta_{\mathrm{i}} \mathrm{H}\right)_{1}$ 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) $\mathrm{F}^{-}$
(ii) Ar
(iii) $\mathrm{Ca}^{2+}$
(iv) $\mathrm{Rb}^{+}$
16. (a) Show by a chemical reaction with water that $\mathrm{Na}_{2} \mathrm{O}$ is a basic oxide and $\mathrm{Cl}_{2} \mathrm{O}_{7}$ is an acidic oxide.
(b) Name a species that will be isoelectronic with each of the following atoms or ions, (i) $\mathrm{F}^{-} \quad$ (ii) $\mathrm{Ca}^{2+}$
17. The first ionisation enthalpy values (in $\mathrm{kJmol}^{-1}$ ) of group- 13 elements are:

| B | Al | Ga | In | Tl |
| :---: | :---: | :---: | :---: | :---: |
| 801 | 577 | 579 | 558 | 589 |

How would you explain this deviation from the general trend?
18. The first $\left(\mathrm{IE}_{1}\right)$ and the second $\left(\mathrm{IE}_{2}\right)$ ionisation enthalpies $\left(\mathrm{kJ} \mathrm{mol}^{-1}\right)$ of three elements are given below:

|  | I | II | III |
| :---: | :---: | :---: | :---: |
| $\mathrm{IE}_{1}$ | 403 | 549 | 1142 |
| $\mathrm{IE}_{2}$ | 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) $\mathrm{B}, \mathrm{Al}, \mathrm{Ga}$ (decreasing order of atomic radii).
(ii) $\mathrm{C}, \mathrm{S}, \mathrm{N}\left(\right.$ decreasing order of $\left.(\Delta \mathrm{Heg})_{1}\right)$
(iii) Al forms amphoteric oxide. Why?
(iv) $\mathrm{Mg}^{2+}$ ion is smaller than $\mathrm{O}^{2-}$ ion although both have the some electronic configuration.
2. Element

I
II
III
(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-I

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. Which of the following show the weakest shielding effect?
(a) s
(b) p
(c) d
(d) f
2. Which has highest electronegativity?
(a) Cl
(b) O
(c) N
(d) S
3. Which pair of elements has similar properties?

13, 31, 11, 21
4. Write general outer electronic configuration of 4 f series elements.
5. Write the IUPAC symbol for the element having atomic number 120.
6. (a) Explain why cation are smaller and anions larger in radii than their parent atoms?
(b) Explain why ianisation enthalpy of nitrogen is more than that of oxygen.
7. The first ionisation enthalpy values (in $\mathrm{kJ} \mathrm{mol}^{-1}$ ) of group-13
elements are :

| B | 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 $\mathrm{Na}_{2} \mathrm{O}$ is a basic oxide and $\mathrm{Cl}_{2} \mathrm{O}_{7}$ is an acidic oxide.
(b) Name a species that will be isoelectronic with each of the following atoms or ions. (i) $\mathrm{F}^{-}$(ii) $\mathrm{Ca}^{2+}$
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?
(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 $\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right)^{2+}$ same?
(d) Why are there fourteen elements in the Lanthanide series?

## UNIT TEST-II

Time allowed : 1 Hour
General instructions :
(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. In the $\mathrm{P}^{3-}, \mathrm{S}^{2-}$ and $\mathrm{Cl}^{-}$ions, the increasing order of size is
(a) $\mathrm{Cl}^{-}, \mathrm{S}^{2-}, \mathrm{P}^{3-}$
(b) $\mathrm{P}^{3-}, \mathrm{S}^{2-}, \mathrm{Cl}^{-}$
(c) $\mathrm{S}^{2-}, \mathrm{Cl}^{-}, \mathrm{P}^{3-}$
(d) $\mathrm{S}^{2-}, \mathrm{P}^{3-}, \mathrm{Cl}^{-}$
2. The element with positive electron gain enthalpy is
(a) hydrogen
(b) sodium
(c) oxygen
(d) neon
3. Write the IUPAC name and symbol for the element with atomic number 118.

In following questions a statement of question followed by a statement of reason is given. Choose the correct answer out of the following choices :
(a) Both Assertion and Reason are true and Reason is the correct explanation of Assertion.
(b) Both Assertion and Reason are true but Reason is not the correct explanation of Assertion.
(c) Assertion is true but Reason is false.
(d) Both Assertion and Reason are false.
4. Assertion : Electron gain enthalpy becomes less negative as we go down a group.
Reason : Size of the atom increases on going down the group and the added electron would be farther from the nucleus.
5. Assertion : Boron has a smaller first ionisation enthalpy than Beryllium.(1) Reason: The penetration of 2 s electron to the nucleus is more than 2 p electron hence $2 p$ electron is more shielded by the inner core than 2 s electron.
6. Out of O and S , which has higher negative electron gain enthalpy and why?
7. Assign the position of elements having outer electronic configuration :
(i) $\mathrm{ns}^{2} \mathrm{np}^{4}$ for $\mathrm{n}=3$
(ii) $(\mathrm{n}-1) \mathrm{d}^{2} \mathrm{~ns}^{2}$ for $\mathrm{n}=4$
8. Consider the element $\mathrm{N}, \mathrm{P}, \mathrm{O}$ and S and arrange them is order of :
(i) increasing 1 st I.E.
(ii) increasing negative electron gain enthalpy
(iii) increasing non-metallic character
9. The first $\left(\mathrm{IE}_{1}\right)$ and second $\left(\mathrm{IE}_{2}\right)$ ionisation enthalpies $\left(\mathrm{kJmol}^{-1}\right)$ of three elements I, II and III are given below :

| Element | $\mathrm{IE}_{1}$ | $\mathrm{IE}_{2}$ |
| :--- | :--- | :--- |
| I | 403 | 2640 |
| II | 549 | 1060 |
| III | 1142 | 2080 |

Identify the element which is likely to be
(i) non-metal
(ii) an alkali metal
(ii) an alkaline earth metal
10. (a) Lithium shows diagonal relationship with which element and why?
(b) Among the elements of second period Li to Ne , pick out element:
(i) with the highest 1 st I.E.
(ii) with the highest electronegativity
(iii) with largest atomic radius
(iv) most reactive non-metal

## Chemical Bonding and Molecular Structure

## FAST TRACK : QUICK REVISION

- 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. $\mathrm{BF}_{3}, \mathrm{NO}_{2}, \mathrm{PCl}_{5}, \mathrm{SF}_{6}$ 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) - $1 / 2$ (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., $\mathrm{H}-\mathrm{F}<\mathrm{H}-\mathrm{Cl}<\mathrm{H}-\mathrm{Br}<\mathrm{H}-\mathrm{I}$
(iii) Greater the s character shorter the bond length.
e.g., $\mathrm{C}_{\mathrm{sp}}{ }^{3}-\mathrm{H}>\mathrm{C}_{\mathrm{sp}}{ }^{2}-\mathrm{H}>\mathrm{C}_{\mathrm{sp}}-\mathrm{H}>$
(iv) Bond length decreases with increase in bond order.
e.g., $\mathrm{C}-\mathrm{C}>\mathrm{C}=\mathrm{C}>\mathrm{C} \equiv \mathrm{C}$

## - 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., $\mathrm{NH}_{3}>\mathrm{PH}_{3}$
(iii) Greater the number of lone pair around central atom smaller the bond angle. e.g., $\mathrm{CH}_{4}>\mathrm{NH}_{3}>\mathrm{H}_{2} \mathrm{O}$

- 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 $\alpha$ Bond order $\alpha 1 /(B o n d ~ l e n g t h)$
- Resonance:
(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 $A-B$ :
$=16\left(\mathrm{X}_{\mathrm{A}}-\mathrm{X}_{\mathrm{B}}\right)+3.5\left(\mathrm{X}_{\mathrm{A}}-\mathrm{X}_{\mathrm{B}}\right)^{2}$,
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 $\alpha$ Charge density [(Charge )/Radius)].
(iv) Polarisability of anion $\alpha$ size of anion.
- Dipole moment:
(i) Dipole moment $(\mu)=$ charge $(Q) \times$ distance of separation (d)
(ii) Unit: Debye (D), 1D $=3.33564 \times 10^{-30} \mathrm{Cm}$
(iii) Being vector quantity, dipole moment of polyatomic molecule is taken as the resultant of all the bond moments.
(iv) If $\mu=0$, 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. $\mathrm{F}, \mathrm{O}$ 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 ( $\sigma$ ) and pi ( $\pi$ ) 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 $=1 \sigma$ bond

Double covalent bond $=1 \sigma$ bond $+1 \pi$ bond
Triple covalent bond $=1 \sigma$ bond $+2 \pi$ 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., $\quad \mathrm{p}-\mathrm{lp}>\mathrm{lp}-\mathrm{bp}>\mathrm{bp}-\mathrm{bp}$


## - 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.

$$
\text { i.e., } \quad \begin{aligned}
& \sigma 1 \mathrm{~s}, \sigma^{*} 1 \mathrm{~s}, \sigma 2 \mathrm{~s}, \sigma^{*} 2 \mathrm{~s}, \pi 2 p_{\mathrm{x}}=\pi 2 \mathrm{p}_{\mathrm{y}}, \sigma 2 \mathrm{p}_{\mathrm{z}}, \pi^{*} 2 \mathrm{p}_{\mathrm{x}}=\pi^{*} 2 \mathrm{p}_{\mathrm{y}} \\
& \sigma^{*} 2 \mathrm{p}_{\mathrm{z}} \text { (upto } 14 \text { electrons) } \\
& \sigma 1 \mathrm{~s}, \sigma^{*} 1 \mathrm{~s}, \sigma 2 \mathrm{~s}, \sigma^{*} 2 \mathrm{~s}, \sigma 2 \mathrm{p}_{\mathrm{z}}, \pi 2 \mathrm{p}_{\mathrm{x}}=\pi 2 \mathrm{p}_{\mathrm{y}}, \pi^{*} 2 \mathrm{p}_{\mathrm{x}}=\pi^{*} 2 \mathrm{p}_{\mathrm{y}} \\
& \sigma^{*} 2 \mathrm{p}_{\mathrm{z}} \text { (For more than } 14 \text { electrons) }
\end{aligned}
$$

(iii) Bond order $=1 / 2\left(\mathrm{~N}_{\mathrm{b}}-\mathrm{N}_{\mathrm{a}}\right)$
$\mathrm{N}_{\mathrm{a}}=$ No of electrons in anti-bonding molecular orbitals
$\mathrm{N}_{\mathrm{b}}=$ No of electrons in bonding molecular orbitals

| Total electron pairs | Bond <br> pairs | Lone pairs | Type of hybridization | Geometry due to repulsion | Bond angle | Example |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2 | 2 | 0 | sp | Linear | $180^{\circ}$ | $\mathrm{BeCl}_{2}$ |
| 3 | 3 | 0 | $\mathrm{sp}^{2}$ | Non-polar Planar | $120^{\circ}$ | $\mathrm{BF}_{3}$ |
| 3 | 2 | 1 | $\mathrm{sp}^{2}$ | Angular | <1200 | $\mathrm{SO}_{2}$ |
| 4 | 4 | 0 | $\mathrm{sp}^{3}$ or dsp ${ }^{2}$ | Tetrahedral | 109028' | $\mathrm{CH}_{4}$ |
| 4 | 3 | 1 | $\mathrm{sp}^{3}$ or $\mathrm{dsp}^{2}$ | Pyramidal | <109028' | $\mathrm{NH}_{3}$ |
| 4 | 2 | 2 | $\mathrm{sp}^{3}$ or $\mathrm{sp}^{2}$ | Bent | $<109{ }^{\circ} 8^{\prime}$ | $\mathrm{H}_{2} \mathrm{O}$ |
| 5 | 5 | 0 | $\mathrm{sp}^{3} \mathrm{~d}$ | Trigonal bipyramidal | $120^{\circ}$ \& $90^{\circ}$ | $\mathrm{PCl}_{5}$ |
| 5 | 4 | 1 | $s p^{3} d$ | See Saw | $<120{ }^{\circ}$ \& $<90^{\circ}$ | $\mathrm{SF}_{4}$ |
| 5 | 3 | 2 | $s p^{3} d$ | Bent T-shaped | $<90^{\circ}$ | $\mathrm{ClF}_{3}$ |
| 5 | 2 | 3 | $\mathrm{sp}^{3} \mathrm{~d}$ | Linear | $180^{\circ}$ | $\mathrm{I}_{3}{ }^{-}$ |
| 6 | 6 | 0 | $\mathrm{sp}^{3} \mathrm{~d}^{2}$ | Octahedral | $90^{\circ}$ | $\mathrm{SF}_{6}$ |
| 6 | 5 | 1 | $\mathrm{sp}^{3} \mathrm{~d}^{2}$ | Square pyramidal | $<90^{\circ}$ | $\mathrm{BrF}_{5}$ |
| 6 | 4 | 2 | $\mathrm{sp}^{3} \mathrm{~d}^{2}$ | Square planar | $90^{\circ}$ | $\mathrm{XeF}_{4}$ |
| 7 | 7 | 0 | $\mathrm{sp}^{3} \mathrm{~d}^{3}$ | Pentagonal bipyramidal | $90^{\circ}$ \& $72^{\circ}$ | $\mathrm{IF}_{7}$ |
| 7 | 6 | 1 | $\mathrm{sp}^{3} \mathrm{~d}^{3}$ | Pentagonal pyramidal | $<90^{\circ}$ \& $<72{ }^{\circ}$ |  |
| 7 | 5 | 2 | $\mathrm{sp}^{3} \mathrm{~d}^{3}$ | Pentagonal planar | $72^{\circ}$ | $\mathrm{XeF}_{5}{ }^{-}$ |



## CASE BASED STUDY-QUESTION

## PASSAGE - 1

Pauling introduced the concept of hybridisation. According to him the atomic orbitals combine to forms new set of equivalent orbitals known as hybrid orbitals. Unlike pure orbitals, the hybrid orbitals are used in bond formation.The phenomenon is known as hybridisation which can be defined as the process of intermixing of the orbitals of slightly different energies so as to redistribute their energies, resulting in the formation of new set of orbitals of equivalent energies and shape.

## Source: NCERT

The following questions are multiple choice questions. Choose the most appropriate answer:
I. In $\mathrm{SF}_{6}$ molecules , which sets of d-orbitals is involved ?
(A) $\mathrm{dx}^{2}-\mathrm{y}^{2}, \mathrm{dz}^{2}$
(B) $\mathrm{dz}^{2}, \mathrm{dxy}$
(C) dxy, dyz
(D) $\mathrm{dx}^{2}-y^{2}, d x y$
II. In $\mathrm{IF}_{7}$ molecule, which orbitals are involved
(A) $d x^{2}-y^{2}, d z^{2}, d x y$
(B) dxy, dyz, dzx
(C) $\mathrm{dx}^{2}-\mathrm{y}^{2}, \mathrm{dxy}, \mathrm{dxz}$
(D) $\mathrm{dz}^{2}$, dyz, dzx
III. In $\mathrm{PCl}_{5}$ molecule, d-orbitals involved is
(A) dxy
(B) dyz
(C) $d x^{2}-y^{2}$
(D) $\mathrm{dz}^{2}$
IV. Which of the following orbitals cannot undergoes hybridisation amongst themselves
(A) $3 \mathrm{~d}, 4 \mathrm{~s}$
(B) $2 \mathrm{~s}, 2 \mathrm{p}$
(C) $4 \mathrm{~s}, 4 \mathrm{~d}$
(D) $3 \mathrm{~s}, 3 \mathrm{p}, 4 \mathrm{~s}$

ANS. : I-A, II-A, III-D, IV-D

## PASSAGE -2

It can be said that covalent compound has partial ionic character due to the electronegativity difference of the two elements in a covalent bond .Dipole moment parameter is used to find the $\%$ of ionic character in the covalent compound. We can also said that ionic compounds do have some covalent character which can be explained qualitatively on the basis of the Fajan's Rule. According to Fajan rule a cation which has smaller size, high ionic charge must have large polarising power and the anion which has large size and high ionic charge must have large polarisibility. Both these factors help to develop covalent character in the molecule.

## In these questions (Q. No V-VIII, a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.

(a) Assertion and reason both are correct statements and reason is correct explanation for assertion.
(b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
(c) Assertion is correct statement but reason is wrong statement.
(d) Assertion is wrong statement but reason is correct statement
V. ASSERTION: $\mathrm{CO}_{2}$ molecule has zero dipole moment despite polar bond. REASON: $\mathrm{CO}_{2}$ molecule is linear in nature.
VI. ASSERTION: The dipole moment of Dichlorobenzene molecule can be zero.

REASON:The Dichlorobenzene must be 1, 4 Dichlorobenzene.
VII. ASSERTION: LiF has less covalent character than LiI

REASON: Halogen molecule are short of one electron to complete the octet.
VIII. ASSERTION: LiCl is more covalent than KCl

REASON: The polarising power of K is more than Li
ANS: V-A, VI-A, VII-B, VIII-C

## MULTIPLE CHOICE QUESTIONS (MCQ)

1. Which of the following molecules has both covalent and ionic bond
(a) $\mathrm{CH}_{3} \mathrm{Cl}$
(b) $\mathrm{NH}_{4} \mathrm{Cl}$
(c) HCl
(d) $\mathrm{BeCl}_{2}$
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 following molecules has maximum bond angle
(a) $\mathrm{NH}_{3}$
(b) $\mathrm{CH}_{4}$
(c) $\mathrm{H}_{2} \mathrm{O}$
(d) $\mathrm{CO}_{2}$
4. Identify correct statement regarding $\mathrm{NH}_{3}$ and $\mathrm{BF}_{3}$
(a) Both are Lewis acid
(b) Both are iso structural
(c) Both are Lewis base
(d) Have different values of dipole moment
5. Identify the molecule having sideways overlapping of atomic orbitals
(a) $\mathrm{CH}_{4}$
(b) $\mathrm{CO}_{2}$
(c) $\mathrm{NH}_{3}$
(d) $\mathrm{H}_{2} \mathrm{O}$
6. Which of the following chemical species is most stable?
(a) $\mathrm{O}_{2}$
(b) $\mathrm{O}_{2}^{+}$
(c) $\mathrm{O}_{2}^{-}$
(d) $\mathrm{O}_{2}{ }^{2-}$
7. Which of the following d orbitals involved in $\mathrm{sp}^{3} \mathrm{~d}$ hybridization?
(a) $d_{x y}$
(b) $d_{x z}$
(c) $d_{x^{2}-y^{2}}$
(d) $\mathrm{d}_{\mathrm{z}}{ }^{2}$
8. Which of the following molecule has net dipole moment?
(a) $\mathrm{CO}_{2}$
(b) $\mathrm{H}_{2} \mathrm{O}$
(c) $\mathrm{BF}_{3}$
(d) $\mathrm{CH}_{4}$
9. Which of the following compound has highest covalent character
(a) LiCl
(b) LiBr
(c) LiF
(d) LiI
10. The shape of $\mathrm{XeF}_{4}$ molecule according to VSEPR theory is
(a) Square planar
(b) Square pyramid
(c) Tetrahedral
(d) Pyramidal

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

## 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 ions $\qquad$ ion 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 $\mathrm{PCl}_{5}$ molecule the two equivalent axial $\mathrm{P}-\mathrm{Cl}$ bonds are $\qquad$ .than three equivalent equatorial $\mathrm{P}-\mathrm{Cl}$ bonds.
(vi) The state of hybridization of sulphur in $\mathrm{SF}_{6}$ is
(vii) The maximum number of Hydrogen bonds formed by a single $\mathrm{H}_{2} \mathrm{O}$ molecule is $\qquad$
(viii) A triple covalent bond consists of......sigma and.......pi bonds.
(ix) ...........bond is directional in nature.
(x) Atomic orbitals are $\qquad$ .centric and molecular orbitals are centric.

Ans. (i) Lattice enthalpy (ii) $\mathrm{Li}^{+}$(iii) more (iv) bond order (v) longer (vi) $\mathrm{sp}^{3} \mathrm{~d}^{2} \quad$ (vii) $4 \quad$ (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) $\mathrm{BeF}_{2}$ has more dipole moment than $\mathrm{BeCl}_{2}$.
(iii) In water two $\mathrm{O}-\mathrm{H}$ bond dissociation enthalpies are not identical.
(iv) 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) $\mathrm{BF}_{3}$ is stronger Lewis acid than $\mathrm{BCl}_{3}$.
(viii) Among alkali metal halides LiI has highest covalent character.
(ix) Resonating structures of a chemical species have no real existence.
(x) $\mathrm{XeF}_{2}$ and $\mathrm{ICl}_{2}^{-}$are iso structural.
Ans. (i) True
(ii) False
(iii) True
(iv) False
(v) True
(vi) False
(vii) False
(viii) True
(ix) True
(x) True

## MATCH THE COLUMNS

I. Match the species in Column I with the geometry/shape in Column II and Hybridisation in Column III
S.N. Column I Column II Column III

1. $\mathrm{BF}_{3}$
(a) Tetrahedral
(p) $\operatorname{sp}^{3} d^{2}$
2. $\mathrm{ClF}_{3}$
(b) Trigonal Planer
(q) $\mathrm{sp}^{3}$
$3 \quad \mathrm{NH}_{4}^{+}$
(c) Octahedral
(r) $\mathrm{Sp}^{3} \mathrm{~d}$
3. $\mathrm{SF}_{6}$
(d) Bent T-Shape
(s) $\mathrm{sp}^{2}$
II. Match the species in Column I with the hybridsation in Column II and geometry/shape in Column III
S.N. Column I Column II Column III
4. $\mathrm{BBr}_{3}$
(a) $\mathrm{sp}^{3}$
(p) Square Planer
5. $\mathrm{H}_{2} \mathrm{O}$
(b) $\mathrm{sp}^{3} \mathrm{~d}$
(q) Trigonal Planer
$3 \quad \mathrm{PCl}_{5}$
(c) $\mathrm{sp}^{2}$
(r) V-shape
6. $\mathrm{XeF}_{4}$
(d) $\mathrm{sp}^{3} \mathrm{~d}^{2}$
(s) Trigonal Bipyramidal

ANS : MATCH-I

1. b, s 2.d, r 3.a, q 4.c, p

MATCH-II

1. c, q 2. a, r 3.b, s 4. d, p

## 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.

1. Assertion (A): Among the two $\mathrm{O}-\mathrm{H}$ bonds in $\mathrm{H}_{2} \mathrm{O}$ molecule, the energy required to break the first $\mathrm{O}-\mathrm{H}$ bond and the other $\mathrm{O}-\mathrm{H}$ bond is the same. Reason (R): This is because the electronic environment around the oxygen is same after breakage of one $\mathrm{O}-\mathrm{H}$ bond.
2. Assertion (A): Though the central atom of both $\mathrm{NH}_{3}$ and $\mathrm{H}_{2} \mathrm{O}$ molecules are $\mathrm{sp}^{3}$ hybridised, yet $\mathrm{H}-\mathrm{N}-\mathrm{H}$ bond angle is greater than that of $\mathrm{H}-\mathrm{O}-\mathrm{H}$. Reason (R): This is because nitrogen atom has one lone pair and oxygen atom has two lone pairs.
3. Assertion (A): $\mathrm{SF}_{6}$ 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.
5. Assertion (A): Ionic compounds tend to be non-volatile.

Reason (R): Ionic compounds are solid.
6. ASSERTION : Bonding molecular orbital are more stable than Antibonding Molecular orbital

REASON : Electrons placed in Bonding molecular orbitals tend to hold the nuclei more together as compared to electrons placed in Antibonding molecular orbitals.
7. ASSERTION : Ortho-Nitrophenol has higher boiling point than ParaNitrophenol
REASON : Intramolecular Hydrogen bonding occur in Ortho-Nitrophenol as compare to intermolecular hydrogen bonding in Para-Nitro Phenol.
8. ASSERTION :When $\mathrm{p}_{\mathrm{x}}$ orbital combine with $\mathrm{p}_{\mathrm{y}}$ orbital than a sigma bond is produced.
REASON : Atomic orbitals of similar symmetry can results positive and negative overlap.
9. ASSERTION : $\mathrm{BF}_{3}$ molecule has zero dipole moment.

REASON : $\mathrm{BF}_{3}$ molecule shape is trigonal planer and symmetrical.
10. ASSERTION : O-O bond length in O3 molecule is identical.

REASON : Ozone molecule is angular in shape
Ans. 1. (iv) 2. (i) 3. (iv) 4. (iv) 5. (ii) 6.(i) 7. (iv) 8. (iv) 9. (i) 10. (iv)

## ONE WORD ANSWER TYPE QUESTIONS

1. Write the formal charge on central oxygen atom in $\mathrm{O}_{3}$ molecule?
2. Write the shape of $\mathrm{AB}_{2} \mathrm{E}_{3}$ 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 phenomenon used to describe a molecule whose single Lewis structure cannot describe it.
9. Name the geometry involved in $\mathrm{sp}^{3} \mathrm{~d}$ hybridization.
10. Name the molecular theory that can explain magnetic character of molecules.
Ans. 1. $+1, \quad$ 2. Linear, $\quad$ 3. Dipole moment, $\quad$ 4. Sigma bond, $\quad$ 5. $\mathrm{p}_{z}$
11. Antibonding molecular orbital, $\quad$ 7. Hydrogen bond, $\quad$ 8. Resonance,
12. Trigonal bipyramid, $\quad$ 10. Molecular orbital theory

## 1-MARK QUESTIONS

1. Why noble gases exist in mono atomic form?
2. Write the Lewis structure of $\mathrm{NO}_{2}{ }^{-}$.
3. Why $\mathrm{NH}_{3}$ and $\mathrm{BF}_{3}$ 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 $\mathrm{BeH}_{2}$ molecule has zero dipole moment although the $\mathrm{Be}-\mathrm{H}$ bonds are polar?
7. Which has highest bond angle? $\mathrm{NO}_{2}, \mathrm{NO}_{2}^{-}, \mathrm{NO}_{2}^{+}$
8. What is magnetic character of anion of $\mathrm{KO}_{2}$ ?
9. Why do atoms combine?
10. What is the significance of Lewis Symbols?
11. Why density of water is maximum at 277 K ?
12. Give structure of $\mathrm{BrF}_{5}$ according to VSEPR theory.
13. Why $\mathrm{NH}_{3}$ is liquid and $\mathrm{PH}_{3}$ is a gas?
14. Why $\mathrm{KHF}_{2}$ exist but $\mathrm{KHCl}_{2}$ and $\mathrm{KHBr}_{2}$ 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. $\mathrm{He}_{2}$ molecule does not exist. Give reason.
19. Why $\mathrm{PCl}_{5}$ dissociates to give $\mathrm{PCl}_{3}$ and $\mathrm{Cl}_{2}$ ?
20. Write the state of hybridization of O in $\mathrm{H}_{2} \mathrm{O}$.
21. Predict the shape of $\mathrm{ClF}_{3}$ according to VSEPR theory.
22. Why ice has less density than water?
23. Why the $\mathrm{H}-\mathrm{P}-\mathrm{H}$ bond angle in $\mathrm{PH}_{3}$ is less than $\mathrm{H}-\mathrm{N}-\mathrm{H}$ bond angle in $\mathrm{NH}_{3}$ ?
24. At room temperature $\mathrm{H}_{2} \mathrm{O}$ exist as liquid while $\mathrm{H}_{2} \mathrm{~S}$ exist as gas. Give reason.
25. $\mathrm{NH}_{3}$ has higher boiling point than $\mathrm{PH}_{3}$. Give reason.
26. Identify the chemical species having identical bond order: $\mathrm{O}_{2}{ }^{2+}, \mathrm{N}_{2}, \mathrm{O}_{2}$, $\mathrm{O}_{2}{ }^{2-}$.

## 2-MARKS QUESTIONS

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

$$
\begin{array}{ll}
\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightarrow \mathrm{H}(\mathrm{~g})+\mathrm{O}-\mathrm{H}(\mathrm{~g}) & \Delta_{\mathrm{a}} \mathrm{H}_{1}^{0}=493 \mathrm{~kJ} \mathrm{~mol}^{-1} \\
\mathrm{O}-\mathrm{H}(\mathrm{~g}) \rightarrow \mathrm{H}(\mathrm{~g})+\mathrm{O}(\mathrm{~g}) & \Delta_{\mathrm{a}} \mathrm{H}_{1}^{0}=424 \mathrm{~kJ} \mathrm{~mol}^{-1}
\end{array}
$$

What is the average bond enthalpy of $\mathrm{H}_{2} \mathrm{O}$ ?
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, $\mathrm{CO}_{2}$ or $\mathrm{N}_{2} \mathrm{O}$ ?
8. Discuss the partial ionic character of covalent bond by taking an example.
9. Draw the resonating structures of $\mathrm{O}_{3}$ 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 $\mathrm{F}_{2}$ is less than that of $\mathrm{Cl}_{2}$ ?
12. Define resonance. Draw resonating structures of $\mathrm{CO}_{2}$.
13. Assign reason for the following;
(i) $\mathrm{NH}_{3}$ is freely soluble in water while $\mathrm{PH}_{3}$ is not.
(ii) $\mathrm{B}_{2}$ is paramagnetic while $\mathrm{C}_{2}$ is not.
14. Out of $\mathrm{NH}_{3}$ and $\mathrm{NF}_{3}$ which is more polar. Explain with the help of dipole moment.
15. $\mathrm{N}_{2}$ is diamagnetic while $\mathrm{O}_{2}$ is paramagnetic. Explain on the basis of Molecular orbital theory.
16. $\mathrm{H}_{2}^{+}$and $\mathrm{H}_{2}^{-}$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 $(\mathrm{EN}=3.2)$ is more electronegative than Nitrogen ( $\mathrm{EN}=3.0$ ), yet chlorine does not form hydrogen bond while nitrogen does. Give reason. (Ans: larger atomic size of Cl ).
20. $\mathrm{ClF}_{3}$ is T shaped but $\mathrm{BF}_{3}$ is planar. Explain.
21. $\mathrm{N}\left(\mathrm{SiH}_{3}\right)_{3}$ and $\mathrm{N}\left(\mathrm{CH}_{3}\right)_{3}$ are not isostructural. Give reason.
22. Draw molecular orbital diagram for $\mathrm{N}_{2}{ }^{+}$molecule.

23 . HCl is a covalent compound but it ionises in the solution?
24. The molecule of $\mathrm{CO}_{2}$ is linear whereas that of $\mathrm{SnCl}_{2}$ is angular why?
25. Arrange the following in the order of property indicated for each set:
(i) $\mathrm{O}_{2}, \mathrm{O}_{2}^{+}, \mathrm{O}_{2}^{-}, \mathrm{O}_{2}^{2-}$ (increasing stability)
(ii) $\mathrm{LiCl}, \mathrm{NaCl}, \mathrm{KCl}, \mathrm{RbCl}$ (increasing covalent character)
(iii) $\mathrm{NO}_{2}, \mathrm{NO}_{2}^{+}, \mathrm{NO}_{2}^{-}$(decreasing bond angle)
(iv) $\mathrm{H}-\mathrm{F}, \mathrm{H}-\mathrm{Cl}, \mathrm{H}-\mathrm{Br}, \mathrm{H}-\mathrm{I}$ (increasing bond dissociation enthalpy)
26. Arrange the following in the order of property indicated for each set:
(i) $\mathrm{H}_{2} \mathrm{O}, \mathrm{NH}_{3}, \mathrm{H}_{2} \mathrm{~S}, \mathrm{HF}$ (increasing polar character)
(ii) $\mathrm{HF}, \mathrm{HCl}, \mathrm{HBr}, \mathrm{HI}$ ( decreasing dipole moment)
(iii) $\mathrm{NO}_{3}{ }^{-}, \mathrm{NO}_{2}^{-}, \mathrm{NO}$ (decreasing ' s ' character of hybridization)
(iv) $\mathrm{BeCl}_{2}, \mathrm{BCl}_{3}, \mathrm{CCl}_{4}, \mathrm{PCl}_{3}$ (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 BornHaber's Cycle.
Enthalpy of sublimation of $\mathrm{K}=89 \mathrm{~kJ} \mathrm{~mol}^{-1}$
Enthalpy of dissociation of $\mathrm{Cl}_{2}=244 \mathrm{~kJ} \mathrm{~mol}^{-1}$
Ionization enthalpy of potassium $=425 \mathrm{~kJ} \mathrm{~mol}^{-1}$
Electron gain enthalpy of chlorine $=-355 \mathrm{~kJ} \mathrm{~mol}^{-1}$
Enthalpy of formation of $\mathrm{KCl}=-438 \mathrm{~kJ}$ mol-1
3. What is meant by hybridization? Describe the shape of $\mathrm{sp}, \mathrm{sp}^{2}$ and $\mathrm{sp}^{3}$ hybridised orbitals.
4. Define bond order. Calculate the bond order in $\mathrm{N}_{2}$ and $\mathrm{O}_{2}$ molecules.
5. Give molecular orbital energy level diagram of $\mathrm{O}_{2}{ }^{2-}$. Write its electronic configuration, magnetic behaviour and bond order.
6. Which of the following in each pair has larger bond angle
(i) $\mathrm{CO}_{2}, \mathrm{BF}_{3}$
(ii) $\mathrm{H}_{2} \mathrm{O}, \mathrm{H}_{2} \mathrm{~S}$
(iii) $\mathrm{CH}_{4}, \mathrm{C}_{2} \mathrm{H}_{2}$
7. What is meant by resonance? Draw the resonating structures of carbonate ion and explain why all the $\mathrm{C}-\mathrm{O}$ bond lengths are identical in carbonate ion?
8. Compare relative stability of following species and predict their magnetic properties:
$\mathrm{O}_{2}, \mathrm{O}_{2}^{+}, \mathrm{O}_{2}^{-}$(superoxide), $\mathrm{O}_{2}{ }^{2-}$ (peroxide)
9. Draw the Lewis structure of the species as mentioned $\mathrm{BF}_{3}, \mathrm{SF}_{6}, \mathrm{NO}_{2}$ :
(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 $\mathrm{PCl}_{5}$ according to hybridization. Why all $\mathrm{P}-\mathrm{Cl}$ bonds lengths are not equivalent in $\mathrm{PCl}_{5}$ ?

## 5-MARKS QUESTIONS

1. Give reasons for the following:
(a) $\mathrm{NH}_{3}$ has higher boiling point than $\mathrm{PH}_{3}$.
(b) Ionic compounds do not conduct electricity in solid state.
(c) LiCl is more covalent than KCl .
(d) $\mathrm{NH}_{3}$ is more polar than $\mathrm{NF}_{3}$.
(e) $\mathrm{H}_{2} \mathrm{O}$ has bent structure.
2. (a) Define the term bond dissociation enthalpy. How is it related to bond order?
(b) Explain why $\mathrm{N}_{2}$ has greater bond dissociation enthalpy than $\mathrm{N}_{2}{ }^{+}$ while $\mathrm{O}_{2}$ has lesser bond dissociation enthalpy than $\mathrm{O}_{2}{ }^{+}$?
3. Draw the shape of following molecules according to VSEPR theory;
$\mathrm{XeO}_{3}, \mathrm{XeF}_{2}, \mathrm{XeOF}_{4}, \mathrm{SF}_{4}, \mathrm{XeF}_{4}$

## HOTS QUESTIONS

1. The bond angle of $\mathrm{H}_{2} \mathrm{O}$ is $104.5^{\circ}$ while that of $\mathrm{F}_{2} \mathrm{O}$ is $102^{\circ}$. Explain why? Solution: The bond pair of electrons are drawn more towards F in $\mathrm{F}_{2} \mathrm{O}$, whereas in $\mathrm{H}_{2} \mathrm{O}$ it is drawn towards O . So bp -bp repulsion in $\mathrm{H}_{2} \mathrm{O}$ is greater than that in $\mathrm{F}_{2} \mathrm{O}$.
2. Anhydrous $\mathrm{AlCl}_{3}$ is covalent. From the data given below, predict whether it would remain covalent or become ionic in aqueous solution.

$$
\begin{aligned}
& \Delta_{\mathrm{i}} \mathrm{H}\left(\mathrm{AlCl}_{3}\right)=5137 \mathrm{~kJ} \mathrm{~mol}^{-1}, \quad \Delta_{\mathrm{hyd}} \mathrm{H}\left(\mathrm{Al}^{3+}\right)=-4665 \mathrm{~kJ} \mathrm{~mol}^{-1} \\
& \Delta_{\mathrm{hyd}} \mathrm{H}\left(\mathrm{Cl}^{-}\right)=-381 \mathrm{~kJ} \mathrm{~mol}^{-1}
\end{aligned}
$$

Solution: Total energy released $=1 \Delta_{\text {hyd }} \mathrm{H}\left(\mathrm{Al}^{3+}\right)+3 \Delta_{\text {hyd }} \mathrm{H}\left(\mathrm{Cl}^{-}\right)$

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

Total energy required $=\Delta_{\mathrm{i}} \mathrm{H}\left(\mathrm{AlCl}_{3}\right)=5137 \mathrm{~kJ} \mathrm{~mol}^{-1}$
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: $\mu_{\text {cal }}=\mathrm{Q} \times \mathrm{r}=\left(1.6 \times 10^{-19} \mathrm{C}\right) \times\left(127 \times 10^{-12} \mathrm{~m}\right)=2.032 \times 10^{-29} \mathrm{C} \mathrm{m}$

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

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

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. Identify the molecule having sideways overlapping of atomic orbitals [1]
(a) $\mathrm{CH}_{4}$
(b) $\mathrm{CO}_{2}$
(c) $\mathrm{NH}_{3}$
(d) $\mathrm{H}_{2} \mathrm{O}$
2. The shape of $\mathrm{XeF}_{4}$ molecule according to VSEPR theory is
(a) Square planar
(b) Square pyramid
(c) Tetrahedral
(d) Pyramidal
3. Write the Lewis structure of $\mathrm{NO}_{2}^{-}$.
4. Which has highest bond angle? $\mathrm{NO}_{2}, \mathrm{NO}_{2}^{-}, \mathrm{NO}_{2}^{+}$[1]
5. Draw the resonating structures of $\mathrm{CO}_{2}$.
6. The enthalpy needed to break the two $\mathrm{O}-\mathrm{H}$ bonds in water are as follows:

$$
\begin{array}{ll}
\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) & \Delta_{\mathrm{a}} \mathrm{H}_{1}^{0}=493 \mathrm{~kJ} \mathrm{~mol}^{-1} \\
\mathrm{O}-\mathrm{H}(\mathrm{~g}) \longrightarrow \mathrm{O}-\mathrm{H}(\mathrm{~g}) & \Delta_{\mathrm{a}} \mathrm{H}_{2}^{0}=424 \mathrm{~kJ} \mathrm{~mol}^{-1}
\end{array}
$$

What is the average bond enthalpy of $\mathrm{H}_{2} \mathrm{O}$ ?

## 7. Out of $\mathrm{NH}_{3}$ and $\mathrm{NF}_{3}$ which is more polar. Explain with the help of dipole moment.

8. Compare relative stability of following species and predict their magnetic properties: $\mathrm{O}_{2}, \mathrm{O}_{2}^{+}, \mathrm{O}_{2}{ }^{-}$(superoxide), $\mathrm{O}_{2}{ }^{2-}$ (peroxide)
9. Explain the structure of $\mathrm{PCl}_{5}$ according to hybridization. Why all $\mathrm{P}-\mathrm{Cl}$ bonds lengths are not equivalent in $\mathrm{PCl}_{5}$ ?
10. (i) $\mathrm{N}_{2}$ is diamagnetic while $\mathrm{O}_{2}$ is paramagnetic. Explain on the basis of Molecular orbital theory.
(ii) Give reasons for the following:
(a) $\mathrm{NH}_{3}$ has higher boiling point than $\mathrm{PH}_{3}$.
(b) Ionic compounds do not conduct electricity in solid state.
(c) LiCl is more covalent than KCl .

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. Identify the molecule in which carbon has 'sp' hybridisation.
(a) $\mathrm{CO}_{2}$
(b) $\mathrm{CH}_{4}$
(c) $\mathrm{C}_{2} \mathrm{H}_{4}$
(d) $\mathrm{C}_{2} \mathrm{H}_{2}$
2. The shape of the molecule $\mathrm{SF}_{4}$ is
(a) Bent
(b) See-saw
(c) Tetrahedral
(d) Square Planer
3. Write the Lewis structure of $\mathrm{CO}_{3}{ }^{2-}$

In following questions a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices:
(a) Assertion and Reason both are correct statements and Reason is correct explanation for assertion.
(b) Assertion and Reason both are correct statement but Reason is not correct explanation for assertion.
(c) Assertion is correct statement but Reason is wrong statement.
(d) Assertion is wrong statement but Reason is correct statement.
4. Assertion : Pie $(\pi)$ bond are directional in nature.

Reason : Sigma bond are formed by axial approach of atomic orbitals.
5. Assertion : Boiling point of water is higher than $\mathrm{H}_{2} \mathrm{~S}$.

Reason : Hydrogen bonding is feasible in Water but in $\mathrm{H}_{2} \mathrm{~S}$ there is no Hydrogen bonding.
6. Why dipole moment of $\mathrm{BF}_{3}$ is zero but for $\mathrm{PCl}_{3}$ it is non zero?
7. Which one LiF or LiI is more ionic and why?
8. Explain with the help of labeled diagram the Valence Bond Theory for formation of $\mathrm{H}_{2}$ molecule.
9. Explain the Octet rule with relevant example. Write two limitation of Octet rule also.
10. Define Hybridisation. Write the salient features of hybridisation. Explain the hybridisation in $\mathrm{SF}_{6}$ molecule with relevant diagram.

Chapter - 5

## Chemical

## Thermodynamics

## FAST TRACK : QUICK REVISION

- System: Specific part of universe in which observations are made.
- Surroundings: Everything which surrounds the system.
- 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: $\Delta \mathrm{T}=0$
(ii) Adiabatic process: $\Delta \mathrm{q}=0$
(iii) Isobaric process: $\Delta \mathrm{P}=0$
(iv) Isochoric process: $\Delta \mathrm{V}=0$
(v) Cyclic process: $\Delta \mathrm{U}=0$
(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 $(\mathrm{P})$, volume $(\mathrm{V})$, temperature $(\mathrm{T})$, enthalpy $(\mathrm{H})$, free energy $(\mathrm{G})$, internal energy $(\mathrm{U})$, entropy $(\mathrm{S})$, 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: $\Delta \mathrm{U}=\mathrm{q}+\mathrm{w}$

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

$$
\begin{aligned}
& \omega_{\text {rev }} \\
&=-2.303 n R T \log V_{f} / V_{i} \\
& \text { or } \quad \omega_{\text {rev }}=-2.303 n R T \log P_{i} / P_{f}
\end{aligned}
$$

- Significance of $\Delta \mathbf{H} \& \Delta \mathbf{U}: \Delta H=q_{p}$ and $\Delta U=q_{v}$
- Relation between $\Delta \mathbf{H} \boldsymbol{\&} \Delta \mathbf{U}: \Delta \mathrm{H}=\Delta \mathrm{U}+\left(\mathrm{n}_{\mathrm{p}}-\mathrm{n}_{\mathrm{r}}\right) \mathrm{RT}$ for gaseous reaction
(i) $\Delta \mathrm{H}=\Delta \mathrm{U}$ if $\left(\mathrm{n}_{\mathrm{p}}-\mathrm{n}_{\mathrm{r}}\right)$ is zero; e.g. $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{HI}(\mathrm{g})$
(ii) $\Delta \mathrm{H}>\Delta \mathrm{U}$ if $\left(\mathrm{n}_{\mathrm{p}}-\mathrm{n}_{\mathrm{r}}\right)$ is positive; e.g. $\mathrm{PCl}_{5}(\mathrm{~g}) \longrightarrow \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})$
(iii) $\Delta \mathrm{H}<\Delta \mathrm{U}$ if $\left(\mathrm{n}_{\mathrm{p}}-\mathrm{n}_{\mathrm{r}}\right)$ is negative; e.g. $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$
- Heat Capacity (C): Amount of heat required to raise the temperature of a substance by $1^{\circ} \mathrm{C}$ or 1 K .

$$
\mathrm{q}=\mathrm{C} \Delta \mathrm{~T}
$$

- Specific Heat Capacity $\left(\mathbf{C}_{\mathbf{s}}\right)$ : Amount of heat required to raise the temperature of 1 g of a substance by $1^{\circ} \mathrm{C}$ or 1 K .

$$
\mathrm{q}=\mathrm{C}_{\mathrm{s}} \times \mathrm{m} \times \Delta \mathrm{T}
$$

- Molar Heat Capacity $\left(\mathbf{C}_{\mathrm{m}}\right)$ : Amount of heat required to raise the temperature of 1 mole of a substance by $1^{\circ} \mathrm{C}$ or 1 K .

$$
\mathrm{q}=\mathrm{C}_{\mathrm{m}} \times \mathrm{n} \times \Delta \mathrm{T}
$$

- 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 $\left(\Delta_{\mathrm{f}} \mathbf{H}^{\mathbf{}}\right)$ : 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).
- $\Delta_{\mathrm{f}} \mathrm{H}^{0}$ of an element in standard state is taken as zero.
- Compounds with -ve value of $\Delta_{\mathrm{f}} \mathrm{H}^{0}$ more stable than their constituents.
- $\Delta_{\mathrm{r}} \mathrm{H}^{0}=\sum \mathrm{a}_{\mathrm{i}} \Delta_{\mathrm{f}} \mathrm{H}^{0}$ (products) $-\sum \mathrm{b}_{\mathrm{i}} \Delta_{\mathrm{f}} \mathrm{H}^{0}$ (reactants); Where 'a' and ' b ' are coefficients of products and reactants in balanced equation.
- Standard Enthalpy of Combustion $\left(\Delta_{\mathbf{c}} \mathbf{H}^{\mathbf{0}}\right)$ : 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, $\quad \mathrm{Cl}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{Cl}(\mathrm{g}) ; \quad \Delta_{\mathrm{Cl}-\mathrm{Cl}} \mathrm{H}^{0}=242 \mathrm{~kJ} \mathrm{~mol}^{-1}$

- 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.
- $\quad \Delta_{\mathrm{r}} \mathrm{H}^{0}=\sum \Delta_{\text {bond }} \mathrm{H}^{0}$ (Reactants) $-\sum \Delta_{\text {bond }} \mathrm{H}^{0}$ (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.
- $\Delta \mathrm{S}_{\text {sys }}=\frac{\left(\mathrm{q}_{\text {rev }}\right)_{\text {sys }}}{\Delta \mathrm{T}}=\frac{\Delta \mathrm{H}_{\text {sys }}}{\Delta \mathrm{T}}$
- Unit of Entropy $=\mathrm{JK}^{-1} \mathrm{~mol}^{-1}$
- Second Law of Thermodynamics: For all the spontaneous processes totally entropy change must positive.

$$
\Delta \mathrm{S}_{\mathrm{total}}=\Delta \mathrm{S}_{\mathrm{sys}}+\Delta \mathrm{S}_{\mathrm{surr}}>0
$$

- Gibbs Helmholtz Equation for determination of Spontaneity:

$$
\Delta \mathrm{G}=\Delta \mathrm{H}-\mathrm{T} \Delta \mathrm{~S}
$$

(i) If $\Delta \mathrm{G}=-\mathrm{ve}$, the process is spontaneous
(ii) If $\Delta \mathrm{G}=+\mathrm{ve}$, the process is non-spontaneous
(iii) If $\Delta \mathrm{G}=0$, the process is in equilibrium

- Relation between Gibbs Energy Change and Equilibrium Constant:

$$
\Delta \mathrm{G}^{0}=-2.303 \mathrm{RT} \log \mathrm{~K}_{\mathrm{c}}
$$

- Third Law of Thermodynamics: The entropy of a perfectly crystalline solid at absolute zero $(0 \mathrm{~K})$ is taken to be zero.
MIND MAP : CHEMICAL THERMODYNAMICS



## CASE BASED STUDIES - QUESTIONS

## PASSAGE - $\mathbf{1}$

Heat of neutralization is defined as amount of heat released when one gram equivalent of a strong base reacts with one gram equivalent of strong acid. The heat of neutralization is come out to be as follows
$\mathrm{H}^{+}$(aq.) $+\mathrm{OH}^{-}$(aq.) $\rightarrow \mathrm{H}_{2} \mathrm{O}($ aq. $), \Delta \mathrm{H}(\mathrm{Neu})=$. ' X '
Since the basic reaction for neutralization reaction is same for all acids and bases, hence enthalpy of neutralization for strong acid and base is always same i.e., $\mathbf{- 5 7 . 1} \mathbf{k j}$. However, if any of the acid or base or both are weak than heat relased will be less. It is because some of heat released is utilized to ionize weak acid or base or both. In case of HF (weak acid), enthalpy of neutralization is more than that of strong acid and strong base due to high enthalpy of hydration of small size of ion.

## Answer the following question

1. Enthalpy of neutralization of HCN against NaOH is -51.4 kj . What is enthalpy of ionization of HCN ?
2. If heat of neutralization of $\mathrm{HA}, \mathrm{HB}, \mathrm{HC}, \mathrm{HD}$ acids follows the order $\mathrm{HA}>$ $\mathrm{HB}>\mathrm{HC}>\mathrm{HD}$ then arrange them in increasing order of acidic strength.
3. What will be heat change for complete neutralization of strong acid $\mathrm{R}(\mathrm{OH})_{2}$ by HCI.
4. What will heat released when equal volume of an aqueous solution of $0.5 \mathrm{M} \mathrm{HNO}_{3}$ is mixed with $0.3 \mathrm{M} \mathrm{NaOH}(\mathrm{aq})$

OR
What will be heat released when 100 ml of 0.2 M HCl is mixed 200 ml of 0.2 M KOH .

Ans: $1.5 .7 \mathrm{kj} \quad 2 . \mathrm{HA}>\mathrm{HB}>\mathrm{HC}>\mathrm{HD} \quad 3 .-114.2 \mathrm{kj}$ or 1.142 kj

## PASSAGE -2

According to first law of thermodynamics, the mathematical expression is as follows :

$$
\Delta \mathrm{U}=\mathrm{Q}+\mathrm{W}
$$

Where $\Delta U$ is the change in internal energy observed when a ' $Q$ ' amount of heat is involved and ' W ' amount of work is associated when the system moves from state ' $A$ ' to state ' $B$ '. The sign of ' $Q$ ' and ' $W$ ' is taken such that $\Delta U$ is measures accurately .

## Answer the following question:

1. What will be work done for isochoric process?
2. What is relationship between internal energy change and work done for an adiabatic process?
3. Define internal energy change?
4. A sample of gas present in a cylinder fitted with a frictionless pistion, expand against a constant pressure of 1 atm from volume of 5 L to 15 L . During the process it absorb 500J energy from surrounding. Calculate change in internal energy.

Or
60 Kj heat is supplied to a system at system at constant volume and temperature rises fro $\mathrm{pm} 20^{\circ} \mathrm{C}$ to $24^{\circ} \mathrm{C}$. Calculate the change in internal energy, work done and q.

ANS: 1. $02 . \Delta \mathrm{E}=\mathrm{w} 3$. Definition 4. -513 J or $\Delta \mathrm{E}=60 \mathrm{j}, \mathrm{w}=0, \mathrm{q}=60 \mathrm{~J}$

## 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 $25^{\circ} \mathrm{C}$ and 1.0 atm except:
(a) $\mathrm{N}_{2}(\mathrm{~g})$
(b) $\mathrm{Fe}(\mathrm{s})$
(c) Ne (g)
(d) $\mathrm{H}(\mathrm{g})$
3. For the following reaction at $25^{\circ} \mathrm{C}, \Delta \mathrm{H}^{\circ}=+115 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{\circ}=+125 \mathrm{~J} / \mathrm{K}$. Calculate $\Delta \mathrm{G}^{\circ}$ for the reaction at $25^{\circ} \mathrm{C}$ :

$$
\mathrm{SBr}_{4}(\mathrm{~g}) \longrightarrow \mathrm{S}(\mathrm{~g})+2 \mathrm{Br}_{2}(\mathrm{l})
$$

(a) +152 kJ
(b) -56.7 kJ
(c) +77.8 kJ .
(d) +37.1 kJ
4. Calculate $\Delta_{\mathrm{r}} \mathrm{H}^{0}$ for the following reaction at $25^{\circ} \mathrm{C}$ :

$$
\begin{array}{lllll} 
& \mathrm{Fe}_{3} \mathrm{O}_{4}(\mathrm{~s})+\mathrm{CO}(\mathrm{~g}) \longrightarrow & \longrightarrow & 3 \mathrm{FeO}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g}) \\
\Delta \mathrm{H}_{\mathrm{f}}^{\mathrm{o}}(\mathrm{~kJ} / \mathrm{mol}) & -1118 & -110.5 & -272 & -393.5
\end{array}
$$

(a) -263 kJ
(b) 54 kJ
(c) 19 kJ .
(d) -50 kJ
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 $\Delta \mathrm{H}^{0}$ for the following reaction at 298 K is -36.4 kJ .

$$
1 / 2 \mathrm{H}_{2}(\mathrm{~g})+1 / 2 \mathrm{Br}_{2}(\mathrm{l}) \longrightarrow \mathrm{HBr}(\mathrm{~g})
$$

Calculate $\Delta \mathrm{U}^{0}$ at 298 K . The universal gas constant, R , is $8.314 \mathrm{~J} / \mathrm{mol} \mathrm{K}$.
(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 $\Delta \mathrm{H}^{\circ}$ for the reaction be labeled $\Delta \mathrm{H}_{\mathrm{f}}^{0}$ ?
(a) $\mathrm{Al}(\mathrm{s})+3 / 2 \mathrm{H}_{2}(\mathrm{~g})+3 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{Al}(\mathrm{OH})_{3}(\mathrm{~s})$
(b) $\mathrm{PCl}_{3}(\mathrm{~g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{POCl}_{3}(\mathrm{~g})$
(c) $1 / 2 \mathrm{~N}_{2} \mathrm{O}(\mathrm{g})+1 / 4 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{NO}(\mathrm{g})$
(d) CaO (s) $+\mathrm{SO}_{2}$ (g) $\longrightarrow \mathrm{CaSO}_{3}$ (s)
8. Which statement is ture for reaction? $2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$
(a) $\Delta \mathrm{S}=+\mathrm{ve}$
(b) $\Delta \mathrm{H}>\mathrm{U}$
(c) $\Delta \mathrm{H}<\mathrm{U}$
(d) $\Delta \mathrm{H}=\mathrm{U}$
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) $\mathrm{q}+\mathrm{w}$
(d) None of these
12. According to second law of thermodynamics
(a) $\Delta \mathrm{S}_{\text {total }}=+\mathrm{ve}$
(b) $\Delta \mathrm{S}_{\text {total }}=-\mathrm{ve}$
(c) $\Delta \mathrm{S}_{\text {system }}=+\mathrm{ve}$
(d) $\Delta \mathrm{S}_{\text {system }}=-\mathrm{ve}$

Ans: 1.(d), 2.(d), 3.(c), 4.(c), 5.(b), 6.(d), 7.(a), 8. (c), 9.(a), 10.(c), 11.(c), 12.(a)

## FILL IN THE BLANKS

(i) $\qquad$ is a measure of the degree of randomness or disorder of a system.
(ii) A process which can take place either of its own or under some initiation is known as $\qquad$
(iii) For evaporation of water the sign of $\Delta \mathrm{H}$ is. $\qquad$ and sign of $\Delta \mathrm{S}$ is $\qquad$
(iv) The entropy of a perfectly crystalline solid is zero at $\qquad$
(v) The heat energy exchanged between the system and surroundings at constant temperature and pressure is known as. $\qquad$
$\qquad$ is the quantity of heat needed to raise the temperature of one mole of a substance by $1^{\circ} \mathrm{C}$
(vii) $\mathrm{C}_{\mathrm{p}}-\mathrm{C}_{\mathrm{v}}=$ $\qquad$
(viii) $\qquad$ $=\Delta \mathrm{H}-\mathrm{T} \Delta \mathrm{S}$.
(ix) According to .......... law of thermodynamics, $\Delta \mathrm{S}_{\text {total }}=+\mathrm{ve}$.
(x) If $\Delta \mathrm{H}=+\mathrm{ve}$ and $\Delta \mathrm{S}=+\mathrm{ve}$, the reaction is spontaneous at ......... temperature
Ans: (i) Entropy
(ii) spontaneous
(iii) $+\mathrm{ve},+\mathrm{ve}$
(iv) $-273^{\circ} \mathrm{C}$
(v) Enthalpy
(vi) molar heat capacity (vii) $R \quad$ (viii) $\Delta G$
(ix) second
(x) high

## TRUE AND FALSE TYPE QUESTIONS

## Write true or false for following statements:

(i) For every chemical reaction at equilibrium $\Delta \mathrm{G}^{0}$ 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 $\mathrm{q}+\mathrm{w}$ 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 $\Delta \mathrm{S}_{\text {system }}=+\mathrm{ve}$.
(vi) $\Delta \mathrm{H}$ is not a state function.
(vii) The energy of universe is conserved while its entropy is increasing.
(viii) Volume is extensive property while temperature is intensive property.
(ix) At $0^{\circ} \mathrm{C}$ 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

I. Match the sign of $\Delta \mathrm{H}$ in Column I with the sign of $\Delta \mathrm{S}$ Column II and sign of $\Delta \mathrm{G}$ in Column III for a Spontaneous \& Non spontaneous reaction.
S.N. Column I-[ $\Delta \mathrm{H}] \quad$ Column II-[ $\Delta \mathrm{S}] \quad$ Column III-[ $[\mathrm{G}]$

1. -Ve
(a) +Ve
(p) + Ve at low temperature
2. +Ve
(b) +Ve
(q) -Ve at High Temperature
$3+\mathrm{Ve}$
(c) +Ve
(r) -Ve at all temperature
3. +Ve
(d) -Ve
(s) +Ve at all Temperature
II. Match the Process in Column I with the parameters in Column II and parameters in Column III
S.N. Column I
4. Isochoric Process
5. Isothermal Process
6. Adiabatic Process
7. Isobaric Process

Column II
(a) Temperature constant
(b) Volume Constant
(c) Pressure Constant
(d) Heat Constant

Column III
(p) w is not zero
(q) T not constant
(r) $\Delta U=0$
(s) Work is zero

## ANS :

MATCH-I : 1.b, r 2. a, q 3. d, s 4. c, p
MATCH-II: 1.b, s 2.a, r 3.d, q 4. c, p

## 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.

1. Assertion (A): Enthalpy of graphite is lower than that of diamond.

Reason (R): Entropy of graphite is greater than that of diamond.
2. Assertion (A): Enthalpy of formation of $\mathrm{H}_{2} \mathrm{O}(1)$ is greater than that of $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$.
Reason (R): Enthalpy change is negative for condensation reaction, $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$ $\mathrm{H}_{2} \mathrm{O}(1)$
3. Assertion (A): $\Delta \mathrm{H}$ and $\Delta \mathrm{U}$ are same for the reaction $\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{NO}(\mathrm{g})$ Reason (R): All the reactants and products are gases.
4. Assertion (A): if both $\Delta \mathrm{H}^{0}$ and $\Delta \mathrm{S}^{0}$ are positive than the reaction will be spontaneous at high temperature
Reason (R): All processes with positive entropy change are spontaneous.
5. Assertion (A): Enthalpy of formation of HCl is equal to bond energy of HCl .
Reason (R): Enthalpy of formation and bond energy both involve the formation of one mole of HCl from the elements.
6. ASSERTION : The standard free energies changes for all the spontaneously occurring reaction are negative
REASON: The standard free energies of the elements in their standard states at 1 bar and 298 K are taken as zero
7. ASSERTION: Enthalpy and Entropy of any elementary substances in the standard states are taken as zero.

REASON: At absolute zero, particles of the perfectly crystalline substances become completely motionless.
8. ASSERTION : Enthalpy of Neutralization of $\mathrm{CH}_{3} \mathrm{COOH}$ by NaOH is less than that of HCl by NaOH

REASON : $\mathrm{CH}_{3} \mathrm{COOH}$ is a weak acid hence need energy to ionize completely.
9. ASSERTION : The heat absorbed during the isothermal expansion of an ideal gas against vacuum is zero
REASON: Internal energy of a gas depends upon Pressure of the gas.
10. ASSERTION : A reaction which is spontaneous and accompanied by decrease of randomness must be exothermic

REASON : All exothermic reactions are accompanied by decrease of randomness.

Ans: 1. (ii) 2. (i) 3. (ii) 4. (iii) 5. (i) 6. (ii) 7. (iv) 8. (i) 9. (iii) 10. (iii)

## 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 $\Delta \mathrm{G}^{0}$ for spontaneous reaction?
4. Write the relation between $\Delta \mathrm{H}$ and $\Delta \mathrm{U}$ for $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{HI}(\mathrm{g})$.
5. Write the SI unit of entropy.
6. Name the calorimeter used to measure $\Delta \mathrm{U}$.
7. What is the standard enthalpy of formation of graphite?
8. What is the sign of $\Delta \mathrm{H}$ for $\mathrm{H}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{H}(\mathrm{g})$ ?
9. If $\mathrm{K}_{\mathrm{c}}=1$, what will be the value of $\Delta \mathrm{G}$ ?
10. An exothermic reaction is spontaneous at all temperature. What is the sign of S ?
Ans: 1. Closed system
11. $\mathrm{W}=0$
12. $\Delta \mathrm{G}=-\mathrm{ve}$
13. $\Delta \mathrm{H}=\Delta \mathrm{U}$
14. $\mathrm{J} \mathrm{K}^{-1} \mathrm{~mol}^{-1}$
15. Bomb calorimeter
16. Zero
17. $\Delta H=+v e$
18. Zero
19. $\Delta \mathrm{S}=+\mathrm{ve}$

## 1-MARK QUESTIONS

1. 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, Gibb's 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 $\Delta \mathrm{H}$ is more significant than $\Delta \mathrm{U}$ ?
7. Write one example each of extensive and intensive properties.
8. Write a chemical equation in which $\Delta \mathrm{H}$ and $\Delta \mathrm{U}$ are equal.

9 Write the relationship between $\Delta \mathrm{H}$ and $\Delta \mathrm{U}$ for the reaction:

$$
\mathrm{C}(\mathrm{~s})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})
$$

10. Define standard enthalpy of formation.
11. Why is the standard enthalpy of formation of diamond not zero although it is an element?
12. The enthalpy of atomization of $\mathrm{CH}_{4}$ is $1665 \mathrm{~kJ} \mathrm{~mol}^{-1}$. What is the bond enthalpy of $\mathrm{C}-\mathrm{H}$ bond?
[Ans. 416.25 kJ ]
13. Identify the species for which $\Delta_{\mathrm{f}} \mathrm{H}^{\theta}=\mathrm{O}$, at $298 \mathrm{~K} ; \mathrm{O}_{3}(\mathrm{~g}), \mathrm{Br}_{2}(\mathrm{~g}), \mathrm{Cl}_{2}(\mathrm{~g})$ $\mathrm{CH}_{4}(\mathrm{~g})$.
14. For the reaction $2 \mathrm{Cl}(\mathrm{g}) \longrightarrow \mathrm{Cl}_{2}(\mathrm{~g})$; what are the sign of $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ ?
15. Out of Diamond and Graphite which has higher entropy?
16. Write an example of endothermic spontaneous reaction.
17. State second law of thermodynamics.
18. State third law of thermodynamics.
19. Which has more entropy? $1 \mathrm{~mol} \mathrm{H}_{2} \mathrm{O}(1)$ at $25^{\circ} \mathrm{C}$ or $1 \mathrm{~mol} \mathrm{H}_{2} \mathrm{O}(1)$ at $35^{\circ} \mathrm{C}$.
20. At what temperature the entropy of a perfectly crystalline solid is zero?
21. For a certain reaction $\Delta \mathrm{G}^{0}=0$, what is the value of $\mathrm{K}_{\mathrm{c}}$ ?
22. For a reaction both $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ 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 $\mathrm{q}+\mathrm{w}$ 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 $\Delta \mathrm{H}$ and $\Delta \mathrm{U}$.
5. Derive the relationship $C_{p}-C_{v}=R$.
6. State and explain Hess's Law of Constant Heat Summation with a suitable example.
7. Derive the relationship between $\Delta \mathrm{H}$ and $\Delta \mathrm{U}$.

Given, $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{NH}_{3}(\mathrm{~g}) ; \Delta_{\mathrm{r}} \mathrm{H}^{\theta}=-92.4 \mathrm{~kJ} \mathrm{~mol}^{-1}$;
What is the standard enthalpy of formation of $\mathrm{NH}_{3}$ gas?
[Ans. $-46.2 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
8. 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 $\mathrm{H}_{2}, \mathrm{Br}_{2}$ and HBr are $435 \mathrm{~kJ} \mathrm{~mol}^{-1}, 192 \mathrm{~kJ} \mathrm{~mol}^{-1}$ and $368 \mathrm{~kJ} \mathrm{~mol}^{-1}$ respectively.
[Ans. $-109 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
9. Is the bond dissociation enthalpy of all the four $\mathrm{C}-\mathrm{H}$ bonds in $\mathrm{CH}_{4}$ same? Give reason in support of your.
10. Define the term entropy. Write its unit. How does entropy of a system change on increasing temperature?
11. The equilibrium constant for a reaction is 10 . What will be the value of $\Delta \mathrm{G}^{\theta} ? \mathrm{R}=8.314 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}, \mathrm{~T}=300 \mathrm{~K}$. [Ans. $-5.744 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
12. The $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ for $2 \mathrm{Ag}_{2} \mathrm{O}(\mathrm{s}) \longrightarrow 4 \mathrm{Ag}(\mathrm{s})+\mathrm{O}_{2}(\mathrm{~g})$ are given +61.17 kJ $\mathrm{mol}^{-1}$ and $+132 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$ 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
2. Calculate the maximum work obtained when 0.75 mole of an ideal gas expands isothermally and reversibly at $27^{\circ} \mathrm{C}$ from a volume of 15 L to 25L.
[Ans. -955.7 J]
3. Calculate the number of kJ necessary to raise the temperature of 60 g of aluminium from 35 to $55^{\circ} \mathrm{C}$. Molar heat capacity of Al is $24 \mathrm{~J} \mathrm{~mol}^{-1} \mathrm{~K}^{-1}$.
[Ans. 1.067 kJ ]
4. The reaction of cyanamide, $\mathrm{NH}_{2} \mathrm{CN}$ (s), with Dioxygen was carried out in a bomb calorimeter, and $\Delta \mathrm{U}$ was found to be $-742.7 \mathrm{~kJ} \mathrm{~mol}^{-1}$ at 298 K . Calculate Enthalpy change for the reaction at 298K,

$$
\mathrm{NH}_{2} \mathrm{CN}(\mathrm{~s})+3 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{N}_{2}(\mathrm{~g})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})
$$

[Ans. $-741.5 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
5. The enthalpy of combustion of methane, graphite and dihydrogen at $298 \mathrm{~K}^{2}$ are $-890.3 \mathrm{~kJ} \mathrm{~mol}^{-1},-393.5 \mathrm{~kJ} \mathrm{~mol}^{-1}$ and $-285.8 \mathrm{~kJ} \mathrm{~mol}^{-1}$ respectively. Calculate enthalpy of formation of methane gas. [Ans. $-74.8 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
6. Explain the Born Haber Cycle to determine the lattice enthalpy of NaCl .
7. Enthalpies of formation of $\mathrm{CO}(\mathrm{g}), \mathrm{CO}_{2}(\mathrm{~g}), \mathrm{N}_{2} \mathrm{O}(\mathrm{g})$ and $\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})$ are -110 , $-393,81$ and 9.7 kJ mol-1 respectively. Find the value of $\Delta_{\mathrm{r}} \mathrm{H}$ for the reaction; $\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})+3 \mathrm{CO}(\mathrm{g}) \longrightarrow \mathrm{N}_{2} \mathrm{O}(\mathrm{g})+3 \mathrm{CO}_{2}(\mathrm{~g})$
[Ans. $-777.7 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
8. The combustion of 1 mol of benzene takes place at 298 K .After combustionCO $\mathrm{C}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ are formed and $3267 \mathrm{~kJ} \mathrm{~mol}^{-1}$ of heat is liberated. Calculate $\Delta_{\mathrm{f}} \mathrm{H}^{0}\left(\mathrm{C}_{6} \mathrm{H}_{6}\right)$.
Given: $\Delta_{\mathrm{f}} \mathrm{H}^{0}\left(\mathrm{H}_{2} \mathrm{O}\right)=-286 \mathrm{~kJ} \mathrm{~mol}^{-1}, \Delta_{\mathrm{f}} \mathrm{H}^{0}\left(\mathrm{CO}_{2}\right)=-393 \mathrm{~kJ} \mathrm{~mol}^{-1}$
[Ans. $48.51 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
9. Calculate the standard enthalpy of formation of $\mathrm{CH}_{3} \mathrm{OH}(1)$ from the following data:
$\mathrm{CH}_{3} \mathrm{OH}(\mathrm{l})+3 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) ; \Delta_{\mathrm{c}} \mathrm{H}^{\theta}=-726 \mathrm{~kJ} \mathrm{~mol}^{-1}$
$\mathrm{C}(\mathrm{g})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g}) ; \Delta_{\mathrm{f}} \mathrm{H}^{\theta}=-393 \mathrm{~kJ} \mathrm{~mol}^{-1}$
$\mathrm{H}_{2}(\mathrm{~g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) ; \Delta_{\mathrm{f}} \mathrm{H}^{\ominus}-286 \mathrm{~kJ} \mathrm{~mol}^{-1}$
[Ans. $-239 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
10. Give reasons:
(i) Evaporation of water is endothermic process but it is spontaneous.
(ii) A real crystal has more entropy than an ideal crystal.
(iii) Entropy of universe is increasing.
11. For the reaction at $298 \mathrm{~K}, 2 \mathrm{~A}+\mathrm{B} \longrightarrow \mathrm{C} ; \Delta \mathrm{H}=400 \mathrm{~kJ} \mathrm{~mol}^{-1}, \Delta \mathrm{~S}=0.2$ $\mathrm{kJ} \mathrm{K}^{-1} \mathrm{~mol}^{-1}$. At what temperature will the reaction become spontaneous considering $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ to be constant over the temperature range.
[Ans. T > 2000 K ]
12. Reaction $\mathrm{X}(\mathrm{s}) \rightarrow \mathrm{Y}(\mathrm{g}) \Delta \mathrm{H}=+\mathrm{ve}$ is spontaneous at temperature ' T '. Determine
(i) Sign of $\Delta \mathrm{S}$ for this reaction.
(ii) Sign of $\Delta \mathrm{G}$ for $\mathrm{Y} \longrightarrow \mathrm{X}$
(iii) Sign of $\Delta G$ 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) $\Delta \mathrm{P}=0$
(c) Water decomposes by absorbing 286.2 kJ of electrical energy per mole. When $\mathrm{H}_{2}$ and $\mathrm{O}_{2}$ combine to form one mole of $\mathrm{H}_{2} \mathrm{O}, 286.2 \mathrm{~kJ}$ of heat is produced. Which thermodynamic law is proved? Write its statement.
2. (a) Why work done is isothermal reversible process is always maximum?

$$
\left[\text { Hint: } \mathrm{q}_{\mathrm{v}}=\Delta \mathrm{U} \text { and } \mathrm{q}_{\mathrm{p}}=\Delta \mathrm{H}\right]
$$

(b) It has been found that 221.4 J is needed to heat 30 g of ethanol from $15^{\circ} \mathrm{C}$ to $18^{\circ} \mathrm{C}$. Calculate (a) specific heat capacity, and (b) molar heat capacity of ethanol.
[Ans. (a) $2.46 \mathrm{Jg}^{-1}{ }^{\circ} \mathrm{C}^{-1}$, (b) $113.2 \mathrm{~J} \mathrm{~mol}^{-1}{ }^{\circ} \mathrm{C}^{-1}$ ]
3. (a) Differentiate the terms Bond dissociation enthalpy \& Bond Enthalpy.
(b) Calculate enthalpy change for the process $\mathrm{CCl}_{4}(\mathrm{~g}) \longrightarrow \mathrm{C}(\mathrm{g})+4 \mathrm{Cl}(\mathrm{g})$ and calculate Bond enthalpy of $\mathrm{C}-\mathrm{Cl}$ bond in $\mathrm{CCl}_{4}$.
Given: $\Delta_{\text {vap }} \mathrm{H}^{\theta}=30.5 \mathrm{~kJ} \mathrm{~mol}^{-1} ; \Delta_{\mathrm{f}} \mathrm{H}^{\theta}\left(\mathrm{CCl}_{4}\right)=-135.5 \mathrm{~kJ} \mathrm{~mol}^{-1}$;
$\Delta_{\mathrm{a}} \mathrm{H}^{\theta}(\mathrm{C})=715 \mathrm{~kJ} \mathrm{~mol}^{-1}$ and $\Delta_{\mathrm{a}} \mathrm{H}^{\theta}\left(\mathrm{Cl}_{2}\right)=242 \mathrm{~kJ} \mathrm{~mol}^{-1}$
[Ans. $1304 \mathrm{~kJ} \mathrm{~mol}^{-1}, 326 \mathrm{~kJ} \mathrm{~mol}^{-1}$ ]
4. Predict the sign of $\Delta \mathrm{S}$ for the following changes:
(i) Freezing of water.
(ii) C (graphite) $\longrightarrow \mathrm{C}$ (diamond)
(iii) $\mathrm{H}_{2}(\mathrm{~g})$ at 298 k and $1 \mathrm{bar} \longrightarrow \mathrm{H}_{2}(\mathrm{~g})$ at 298 k and 10 bar
(iv) $\mathrm{H}_{2}$ (g) $+\mathrm{I}_{2}$ (g) $\longrightarrow 2 \mathrm{HI}$ (g)
(v) $2 \mathrm{NaHCO}_{3}$ (s) $\longrightarrow \mathrm{Na}_{2} \mathrm{CO}_{3}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$
5. (i) Define Gibbs Energy. Give its mathematical expression. What is Gibb's energy criteria of Spontaneity.
(ii) For the reaction:

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

Calculate $\Delta \mathrm{rG}^{\circ}$ for the reaction, and predict whether the reaction will occur spontaneously.
[Ans. $\Delta_{\mathrm{r}} \mathrm{G}^{\theta}=+0.16 \mathrm{~kJ}$, 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 $27^{\circ} \mathrm{C}$ from 10 L to 5 L . Calculate $\mathrm{q}, \mathrm{w}, \Delta \mathrm{U}$ and,$\Delta \mathrm{H}$.
Solution: $\mathrm{q}=-2.303 \mathrm{nRT} \log \mathrm{V}_{2} / \mathrm{V}_{1}=-2.303 \times 10 / 40 \mathrm{~mol} \times 2 \mathrm{Cal} \mathrm{K}^{-1}$ $\mathrm{mol}^{-1} \times 300 \mathrm{~K} \times \log 5 / 10=-103.635 \mathrm{Cal}$
For isothermal compression $\Delta \mathrm{U}=0$

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

Also when temperature is constant,

$$
\mathrm{PV}=\text { constant }, \Delta \mathrm{H}=\Delta \mathrm{U}+\Delta(\mathrm{PV})=0+0=0
$$

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: $\mathrm{w}_{(\text {exp. })}=-\left[2.303 \mathrm{nRT} \log \left(\mathrm{P}_{\mathrm{i}} / \mathrm{P}_{\mathrm{f}}\right)\right]$
$=-2.303 \times 1 \mathrm{~mol} \times 8.314 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1} \times 300 \mathrm{~K} \times \log 10 / 1$
$=5.74 \times 10^{3} \mathrm{~J}$
Now, $\mathrm{mgh}=5.74 \times 10^{3} \mathrm{~J}$ or $50 \mathrm{~kg} \times 9.81 \mathrm{~m} \mathrm{~s}^{-2} \times \mathrm{h}=5.744 \times 10^{3} \mathrm{~J}$
$\therefore \mathrm{h}=11.7 \mathrm{~m}$

## UNIT TEST-I

Time Allowed: $\mathbf{1} \mathbf{~ H r}$.
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 $2 \mathrm{Cl}(\mathrm{g}) \longrightarrow \mathrm{Cl}_{2}$ (g); what are the sign of $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ ? [1]
2. Write an example of endothermic spontaneous reaction.
3. ' $w$ ' 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?
4. 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?
5. State and explain Hess's Law of Constant Heat Summation with a suitable example.
6. Calculate the number of kJ necessary to raise the temperature of 60 g of aluminium from 35 to $55^{\circ} \mathrm{C}$. Molar heat capacity of Al is $24 \mathrm{~J} \mathrm{~mol}^{-1} \mathrm{~K}^{-1}$.
7. Calculate the standard enthalpy of formation of $\mathrm{CH}_{3} \mathrm{OH}$ (1) from the following data:
$\mathrm{CH}_{3} \mathrm{OH}(\mathrm{l})+3 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) ;$
$\Delta_{\mathrm{c}} \mathrm{H}^{\theta}=-726 \mathrm{~kJ} \mathrm{~mol}^{-1}$
$\mathrm{C}(\mathrm{g})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g}) ; \Delta_{\mathrm{f}} \mathrm{H}^{\theta}=-393 \mathrm{~kJ} \mathrm{~mol}^{-1}$
$\mathrm{H}_{2}(\mathrm{~g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) ; \Delta_{\mathrm{f}} \mathrm{H}^{\theta}=-286 \mathrm{~kJ} \mathrm{~mol}^{-1}$
8. (a) For oxidation of iron, $4 \mathrm{Fe}(\mathrm{s})+3 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{Fe}_{2} \mathrm{O}_{3}(\mathrm{~s})$ entropy change is $-549.4 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$ at 298 K . In spite of negative entropy change of this reaction, why is the reaction spontaneous? $\left(\Delta \mathrm{r} \mathrm{H}^{\circ}\right.$ for this reaction is $-1648 \mathrm{~kJ} \mathrm{~mol}^{-1}$ )
(b) For the reaction: $2 \mathrm{~A}(\mathrm{~g})+\mathrm{B}(\mathrm{g}) \longrightarrow 2 \mathrm{D}(\mathrm{g}), \Delta \mathrm{U}^{0}=-10.5 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{0}=-44.1 \mathrm{~J} \mathrm{~K}^{-1}$. Calculate $\Delta \mathrm{G}^{0}$ for the reaction, and predict whether the reaction will occur spontaneously.

## UNIT TEST-II

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. Among following the property which is intensive is
(a) Mass
(b) Volume
(c) Temperature
(d) Length
2. Heat of formation is zero for which substance.
(a) $\mathrm{CaCO}_{3}$
(b) HCl
(c) Carbon (Diamond)
(d) Carbon (Graphite)
3. What is an isochoric process and what is the work value associated with isochoric process?

In following questions a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.
(a) Assertion and Reason both are correct statements and reason is correct explanation for assertion.
(b) Assertion and Reason both are correct statements but reason is not correct explanation for assertion.
(c) Assertion is correct statement but reason is wrong statement.
(d) Assertion is wrong statement but reason is correct statement.
4. Name the law which is based on the statement "enthalpy is a state function."
5. Heat of combustion of ethane is $-1560 \mathrm{KJ} \mathrm{mol}^{-1}$ while that of acetylene is $-1296 \mathrm{KJ} \mathrm{mol}^{-1}$. Which is better fuel and why?
6. Derive the expression for work done in isothermal reversible expansion.
7. Define standard enthalpy of combustion and standard enthalpy of formation taking $\mathrm{C}_{2} \mathrm{H}_{6}$ (gas) molecule in both case.
8. State 'Hess law of Constant heat summation'. The molar heat of combustion of $\mathrm{C}_{2} \mathrm{H}_{2}(\mathrm{~g}), \mathrm{C}\left(\right.$ Graphite ) and $\mathrm{H}_{2}(\mathrm{~g})$ are $310.62 \mathrm{kcal}, 94.05 \mathrm{kcal}$ and 68.32 kcal respectively. Calculate the standard heat of formation of $\mathrm{C}_{2} \mathrm{H}_{2}(\mathrm{~g})$.
9. The reaction of Cyanamid, $\mathrm{NH}_{2} \mathrm{CN}(\mathrm{s})$, with dioxygen was carried out in a bomb calorimeter, and $\Delta \mathrm{U}$ was found to be $-742.7 \mathrm{kJmol}^{-1}$ at 298 K . Calculate enthalpy change for the reaction at 298 K .
$\mathrm{NH}_{2} \mathrm{CN}(\mathrm{s})+1.5 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{N}_{2}(\mathrm{~g})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
10. (i) State the First Law of Thermodynamics.
(ii) Define Entropy. What is the effect of temperature on Entropy?
(iii) Two moles of an ideal gas are held by a piston under 5 atm pressure at 300 K . The pressure is suddenly released to 0.4 atm and the gas is allowed to expand isothermally. Calculate $\mathrm{W}, \mathrm{q}, \Delta \mathrm{E}, \Delta \mathrm{H}$.


## Chapter - 6

## 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., $\Delta \mathrm{G}=0$.
- Equilibrium constant : For a general reaction :

$$
\begin{aligned}
& a \mathrm{~A}+b \mathrm{~B} c \mathrm{C}+d \mathrm{D} \\
& \mathrm{~K}_{c}=\frac{[\mathrm{C}]^{c}[\mathrm{D}]^{d}}{[\mathrm{~A}]^{a}[\mathrm{~B}]^{b}} \text { and } \mathrm{K}_{p}=\frac{\mathrm{P}_{\mathrm{C}}^{c} \times \mathrm{P}_{\mathrm{D}}^{d}}{\mathrm{P}_{\mathrm{A}}^{a} \times \mathrm{P}_{\mathrm{B}}^{b}}
\end{aligned}
$$

- Relationship between $K_{p}$ and $K_{c}$ :

$$
\begin{aligned}
\mathrm{K}_{p} & =\mathrm{K}_{c}(\mathrm{RT})^{\Delta n_{g}} \\
\Delta n_{g} & =n_{p}(g)-n_{r}(g)
\end{aligned}
$$

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

Chemical equation

$$
a \mathrm{~A}+b \mathrm{~B} \leftrightharpoons c \mathrm{C}+d \mathrm{D}
$$

$$
\mathrm{K}
$$

$$
c \mathrm{C}+d \mathrm{D} \leftrightharpoons a \mathrm{~A}+b \mathrm{~B}
$$

$$
K_{1}=\frac{1}{K}
$$

$$
n a \mathrm{~A}+n b \mathrm{~B} \leftrightharpoons n c \mathrm{C}+n d \mathrm{D}
$$

$$
\mathrm{K}_{2}=\mathrm{K}^{n}
$$

$\frac{1}{\mathrm{n}} \mathrm{aA}+\frac{1}{\mathrm{n}} \mathrm{bB} \rightleftharpoons \frac{1}{\mathrm{n}} \mathrm{cC}+\frac{1}{\mathrm{n}} \mathrm{dD}$

- Predicting the direction of reaction :

If $\mathrm{Q}_{c}=\mathrm{K}_{c} \Rightarrow$ The reaction is in a state of equilibrium. $\mathrm{Q}_{c}>\mathrm{K}_{c} \Rightarrow$ The reaction proceeds in reverse direction. $\mathrm{Q}_{c}<\mathrm{K}_{c} \Rightarrow$ The reaction proceeds in forward direction.

- Ostwald's dilution law : Degree of dissociation of weak electrolyte, $\alpha=\sqrt{\frac{\mathrm{K}}{\mathrm{C}}}$
- Ionic Product of water $\left(\mathrm{K}_{w}\right)=\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]\left[\mathrm{OH}^{-}\right]=10^{-14}$ at 298 K
- 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.
- Outcomes of Le-Chatelier's Principle

Change at equilibrium
Increase in temperature
Decrease in temperature
Increase in pressure
Decrease in pressure
Increase in Conc. of reactants Increase in Conc. of products

## Shift in equilibrium

Endothermic direction
Exothermic direction
Towards lesser gaseous moles
Towards greater gaseous moles
Forward direction
Reverse direction

- Conjugate Acid or Base : Acid-base pair which differ by $\mathbf{H}^{+}$ion.

Species $-\mathrm{H}^{+}=$Conjugate base
Species $+\mathrm{H}^{+}=$Conjugate acid

- $\mathbf{p H}$ of solution :
$\mathrm{pH}=-\log \left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$or $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}, \mathrm{pOH}=-\log \left[\mathrm{OH}^{-}\right]$
$\mathrm{pH}+\mathrm{pOH}=\mathrm{pK}_{w}=14$ at 298 K
- 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 $\mathrm{NH}_{4} \mathrm{OH}$ decreases in the presence of strong electrolyte $\mathrm{NH}_{4} \mathrm{Cl}$.
- 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., $\mathrm{CH}_{3} \mathrm{COONa}$ ) hydrolyse, $\mathrm{pH}>7$ (The anion acts as a base).

$$
\begin{array}{cc}
\mathrm{X}^{-}+\mathrm{H}_{2} \mathrm{O} \leftrightharpoons & \mathrm{HX} \\
& (\text { Weak acid }) \\
\mathrm{pH}=7+\frac{1}{2}\left(p \mathrm{~K}_{a}+\log \mathrm{C}\right) & \text { (Weak base) }
\end{array}
$$

3. Salt of strong acids and weak bases (e.g., $\left.\mathrm{NH}_{4} \mathrm{Cl}\right)$ hydrolyse, $\mathrm{pH}<7$. (The cation acts as an acid).

$$
\begin{aligned}
& \mathrm{M}^{+}+\mathrm{H}_{2} \mathrm{O} \leftrightharpoons \mathrm{MOH}+\mathrm{H}^{+} \\
& \mathrm{pH}=7-\frac{1}{2}\left(\mathrm{pK}_{b}+\operatorname{logC}\right)
\end{aligned}
$$

4. Salt of weak acids and weak base (e.g., $\mathrm{CH}_{3} \mathrm{COONH}_{4}$ ) 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 $\mathrm{K}_{a}$ and $\mathrm{K}_{b}$ for these ions.

$$
\begin{aligned}
& \mathrm{M}^{+}+\mathrm{X}^{-}+\mathrm{H}_{2} \mathrm{O} \leftrightharpoons \mathrm{MOH}+\mathrm{HX} \\
& \mathrm{pH}=7+\frac{1}{2}\left(\mathrm{pK}_{a}-\mathrm{pK}_{b}\right)
\end{aligned}
$$

- 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., $\mathrm{NH}_{4} \mathrm{OH}+\mathrm{NH}_{4} \mathrm{Cl}$
- Acidic buffer : Solution of weak acid and its salt with strong base, For e.g., $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{CH}_{3} \mathrm{COONa}$.
- Henderson Hasselbalch Equation for the pH of Buffer solution-

$$
\begin{array}{ll}
\mathrm{pH}=\mathrm{pK}_{\mathrm{a}}+\log \frac{[\text { Salt }]}{[\text { Acid }]} & \text { (for acidic buffer) } \\
\mathrm{pOH}=\mathrm{pK}_{\mathrm{a}}+\log \frac{[\text { Salt }]}{[\text { Base }]} & \text { (for basic buffer) }
\end{array}
$$

- Solubility Product $\left(\mathbf{K}_{s p}\right)$ : The equilibrium constant that represent the equilibrium between undissolved salt (solute) and its ions in a saturated solution is called solubility product constant $\left(\mathrm{K}_{s p}\right)$.

For $\mathrm{A}_{x} \mathrm{~B}_{y} \stackrel{a q}{\rightleftharpoons} x \mathrm{~A}^{y+}+y \mathrm{~B}^{x-}$

$$
\mathrm{K}_{s p}=\left[\mathrm{A}^{y+}\right]^{x}\left[\mathrm{~B}^{x-}\right]^{y}=(x s)^{x}(y s)^{y}=x^{x} \cdot y^{x} \cdot s^{(x+y)}
$$

where $s=$ Molar solubility
If ionic product $<\mathrm{K}_{s p}$; salt remain dissolve.
If ionic product $>\mathrm{K}_{s p}$; salt will be precipitated.

- Relationship between solubility $(\mathbf{s})$ and solubility product $\left(K_{s p}\right)$.

$$
\mathrm{K}_{s p}=x^{x} \cdot y^{y} \cdot \mathrm{~s}^{x+y}
$$

For binary salts (e.g., $\mathrm{AgCl}, \mathrm{AgBr}, \mathrm{AgI})$

$$
\mathrm{K}_{s p}=\mathrm{s}^{2}
$$

For Ternary salts (e.g., $\mathrm{PbI}_{2}$ )
$\mathrm{K}_{s p}=4 \mathrm{~s}^{3}$



## CASE BASED STUDY QUESTIONS

## PASSAGE - 1

According to Arrehenius theory, acids are substances that dissociates in water to give hydrogen ions and bases are substances that produce hydroxyl ions.

Bronsted-Lowry gave a more general definition of acids and bases. According to Bronsted-Lowry theory , acid is a substance that is capable of donating a hydrogen ion and bases are substance which are capable of accepting of a hydrogen ion.

According to Lewis Acids and Bases , an acid is a species which accepts electron pair and base which donates an electron pair

## Answer the following questions

1. Which of the following acts as both Bronsted acid as well as Bronsted base
(A) $\mathrm{HCO}_{3}$
(B) $\mathrm{O}_{2}$
(C) $\mathrm{HS}^{-}$
(D) $\mathrm{HPO}_{3}{ }^{2-}$
2. Arrange the following in order of increasing basic strength
$\mathrm{OH}^{-} ; \mathrm{CH}_{3} \mathrm{COO}^{-}, \mathrm{Cl}^{-}, \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{O}^{-}$
3. Define lewis acid, Give example.
4. All Arrhenius acid are Bronsted acid but Arrhenius bases are not Bronsted base. Explain.

OR
Classify the following as lewis acid and lewis base and show hot they acts as such
(A) $\mathrm{AlCl}_{3}$
(B) $\mathrm{H}_{2} \mathrm{O}$

ANS. 1. $\mathrm{HCO}_{3}^{-}, \mathrm{HS}_{2} \quad$ 2. $\mathrm{Cl}^{-}<\mathrm{CH}_{3} \mathrm{COO}^{-}<\mathrm{OH}^{-}<\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{O}^{-}$3. Definition
4. all Arrhenius acid gives $\mathrm{H}^{+}$in aqueous solution while Bronsted acids also gives $\mathrm{H}^{+}$and hence all Arrhenius acids are Bronsted acids but Arrhenius base given $\mathrm{OH}^{-}$in aqueous solution while Bronsted bases accept $\mathrm{H}^{+}$.
$\mathrm{AlCl}_{3}$ : lewis acid as it accept electron pair
$\mathrm{H}_{2} \mathrm{O}$ : Lewis base as it can donate lone pair of electron present on oxygen atom.

PASSAGE - 2
Common Ion effect is the phenomenon in which weak acid or weak base dissociation is suppressed due to the presence of the common ion provided by the strong electrolyte. For example the dissociation of $\mathrm{CH}_{3} \mathrm{COOH}$ is suppressed by the addition of the $\mathrm{CH}_{3} \mathrm{COONa}$. Similarly the dissociation of $\mathrm{NH}_{4} \mathrm{OH}$ is suppressed due to presence of $\mathrm{NH}_{4} \mathrm{Cl}$.This occurs due to the Le-Chatlier's principle .

Due to common ion effect there is decrease in solubility of an ionic precipitate by addition to the solution of a soluble compound with an ion common with then precipitate. In water treatment process highly soluble sodium carbonate salt is added to precipitate out sparingly soluble salt calcium carbonate. Salting out process used in manufacturing soap is also benefit from the common ion effect. Common ion effect also help in maintaining pH of a buffer solution.

## Answer the following questions:

1. What is common ion effect?
2. Why salt is added to soap during soap formation.
3. Name the principle common ion is based on.
4. Both cation of $2^{\text {th }}$ and $4^{\text {th }}$ group are precipitated as sulphides, but $4^{\text {th }}$ group radicals do no get precipitated when $\mathrm{H}_{2} \mathrm{~S}$ is passed through solution for $2^{\text {nd }}$ group why.

## Or

$3^{\text {rd }}$ group radicals are precipitated as hydroxides by addition of $\mathrm{NH}_{4} \mathrm{OH}$ in presence of $\mathrm{NH}_{4} \mathrm{Cl}$, what is the role of $\mathrm{NH}_{4} \mathrm{Cl}$.

## MULTIPLE CHOICE QUESTION (MCQ)

1. For the hypothetical reactions, the equilibrium constant $(\mathrm{k})$ values are given
$\mathrm{A} \rightleftharpoons \mathrm{B}: \mathrm{k}_{1}=2$
$\mathrm{B} \rightleftharpoons \mathrm{C}: \mathrm{K}_{2}=4$
$\mathrm{C} \rightleftharpoons \mathrm{D}: \mathrm{K}_{3}=8$
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
$\mathrm{SO}_{2}(\mathrm{~g})+\frac{1}{2} \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{SO}_{3}(\mathrm{~g})$ is $5 \times 10^{-2} \mathrm{~atm}^{-1 / 2}$
The equilibrium constant for the reaction
$2 \mathrm{SO}_{3}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})$ would be
(a) 100 atm
(b) $25 \times 10^{-4} \mathrm{~atm}$
(c) 400 atm
(d) $125 \times 19^{-6} \mathrm{~atm}^{-3 / 2}$
3. $\mathrm{A}(\mathrm{g})+3 \mathrm{~B}(\mathrm{~g}) \rightleftharpoons 4 \mathrm{C}(\mathrm{g})$ 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
$4 \mathrm{C}(\mathrm{g}) \rightleftharpoons \mathrm{A}(\mathrm{g})+3 \mathrm{~B}(\mathrm{~g})$
(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) $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HCl}(\mathrm{g})$
(b) $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})$
(c) $\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})$
(d) $\mathrm{H}_{2}(\mathrm{l})+\mathrm{CO}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2} \mathrm{CO}_{3}(\mathrm{l})$
5. For the reaction $\mathrm{CO}(\mathrm{g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{COCl}_{2}(\mathrm{~g})$, the value of $\mathrm{K}_{\mathrm{c}} / \mathrm{K}_{\mathrm{p}}$ is equal to
(a) RT
(b) $(\mathrm{RT})^{2}$
(c) $1 / \mathrm{RT}$
(d) 1.0
6. At $90^{\circ} \mathrm{C}$ pure water has $\mathrm{K}_{\mathrm{w}}=10^{-12}$. The solution with pH value 6.5 is
(a) Acidic
(b) Basic
(c) Amphoteric
(d) Data insufficient
7. 40 ml of $0.1 \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}$ is mixed with 20 mL of 0.1 M HCl . What is the pH of the mixture $?\left(\mathrm{pK}_{\mathrm{b}}\right.$ of ammonia solution $\left.=4.74\right)$
(a) 4.74
(b) 2.26
(c) 9.26
(d) 5
8. Identify Bronsted Lowry Acids in the reaction

$$
\underset{(\mathrm{X})}{\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}}+\underset{(\mathrm{Y})}{\mathrm{HCO}_{3}^{-}} \rightleftharpoons \underset{(\mathrm{P})}{\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{5}(\mathrm{OH})\right]^{2+}}+\underset{(\mathrm{Q})}{\mathrm{H}_{2} \mathrm{CO}_{3}}
$$

(a) $\mathrm{X}, \mathrm{Y}$
(b) Y, P
(c) P, Q
(d) $\mathrm{X}, \mathrm{Q}$
9. The $\mathrm{pK}_{\mathrm{a}}$ of weak acid HA is 4.80 and $\mathrm{pK}_{\mathrm{b}}$ 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 $\mathrm{KAl}\left(\mathrm{SO}_{4}\right)_{2}$, then $\mathrm{K}_{\text {sp }}$ is equal to
(a) $\mathrm{p}^{3}$
(b) $4 p^{4}$
(c) $\mathrm{p}^{4}$
(d) $4 \mathrm{p}^{3}$

## 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 $2 \mathrm{~A}(\mathrm{~g}) \rightleftharpoons \mathrm{B}(\mathrm{g}) \mathrm{Q}_{\mathrm{c}}>\mathrm{K}$ if ' A ' is added maintaining $\mathrm{Q}_{\mathrm{c}}>\mathrm{K}$, the reaction will move in backward direction.
3. For the reaction at equilibrium
$\mathrm{CaCO}_{3} \rightleftharpoons \mathrm{CaO}(\mathrm{s})+\mathrm{CO}_{2}(\mathrm{~g})$
What $\mathrm{CaO}(\mathrm{s})$ is removed reaction moves in forward direction.
4. For a reaction $\mathrm{aA}+\mathrm{bB} \rightleftharpoons \mathrm{cC}+\mathrm{dD}$ at equilibrium $\Delta \mathrm{G}^{0}=0$ always.
5. For a reaction at equilibrium $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HCl}(\mathrm{g})$ $\mathrm{K}=4$, the value of $\frac{\mathrm{K}_{\mathrm{b}}[\mathrm{HCl}]^{2}}{\mathrm{~K}_{\mathrm{f}}\left[\mathrm{H}_{2}\right]\left[\mathrm{Cl}_{2}\right]}$ is 1.
6. For the electrolyte $\mathrm{A}_{2} \mathrm{~B}$ if $\mathrm{K}_{\text {sp }}$ is solubility product then its solubling ' S ' M is $\left[\mathrm{K}_{\mathrm{sp}}\right]^{1 / 3} \div 4$.
7. $\mathrm{HCO}_{3}^{-}$is conjugate base of $\mathrm{H}_{2} \mathrm{CO}_{3}$.
8. $\mathrm{H}_{2} \mathrm{O}$ 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 $\mathrm{AB}_{2}(\mathrm{~s})$ solution if Ionic product (I.P) $>\mathrm{K}_{\text {sp }}$, 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. $\mathrm{K}_{\mathrm{p}} \& \mathrm{~K}_{\mathrm{c}}$ are ....... for reaction at equilibrium of type $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{Br}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HBr}(\mathrm{g})$.
3. If $\mathrm{c}_{\mathrm{c}}$ for reaction $\mathrm{CH}_{3} \mathrm{COOH}(\mathrm{l})+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH} \rightleftharpoons \mathrm{CH}_{3} \mathrm{COOC}_{2} \mathrm{H}_{5}(\mathrm{l})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ is 4. Then $Q_{c}$ and $K_{c}$ are $\qquad$
4. If $\mathrm{A}+\mathrm{B}-70 \mathrm{~J} / \mathrm{mol} \rightleftharpoons \mathrm{D}$, reaction temperature is increased then reaction moves in $\qquad$ direction.
5. Presence of catalyst will ...... the equilibrium constant.
6. The conjugate acid of $\mathrm{H}_{2} \mathrm{O}$ is $\qquad$
7. On dilution, the degree of dissociation of acetic acid will $\qquad$
8. The presence of $\mathrm{NH}_{4} \mathrm{Cl}$ in $\mathrm{NH}_{4} \mathrm{OH}$ solution will $\qquad$ the degree of dissociation of $\mathrm{NH}_{4} \mathrm{OH}$.
9. If Ionic product (IP) $<\mathrm{K}_{\mathrm{sp}}$ for a salt solution of AB , then addition of AB further $\qquad$ lead to precipitation initially.
10. $K_{p}$ is always equal to $K_{c}$ if $\Delta n_{g}$ is $\qquad$
Ans. 1. rate of backward reaction, 2. equal, 3. equal, 4. backward direction forward, 5. not change, $6 . \mathrm{H}_{3} \mathrm{O}^{+}$, 7. increase, 8. decrease, 9. will not, 10. zero.

## MATCH THE COLUMNS

I. Match the reaction in Column I with the parameters in Column II and unit (M=Molarity) of K eq in Column III
S.N. Column I Column II Column III

1. $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}), \Delta \mathrm{H}=$-ve $\quad(\mathrm{a}) \mathrm{T}$ increase then K increase $(\mathrm{p}) \mathrm{M}^{0}$
2. $2 \mathrm{~N}_{2}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 4 \mathrm{NO}(\mathrm{g}), \Delta \mathrm{H}=+\mathrm{ve}(\mathrm{b}) \mathrm{T}$ increase then K decrease $(\mathrm{q}) \mathrm{M}^{2-}$
3. $2 \mathrm{X}(\mathrm{g}) \rightleftharpoons \mathrm{Y}(\mathrm{g}), \Delta \mathrm{H}=+\mathrm{Ve}$
(c) P has not effect
(r) $\mathrm{M}^{-1}$
4. $\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}), \Delta \mathrm{H}=+\mathrm{ve}$
(d) Equilibrium move forward (s) M On addition of Xe Gas
II. Match the parameter in Column I with the pH expression in Column II and examples in Column III
S.N. Column I
5. Salt of weak acid and weak base
6. Salt of weak acid and strong base

3 Salt of strong acid and strong base
4. Salt of strong acid and weak base

Column II-pH
(a) $7+0.5(\mathrm{pka}+\log \mathrm{C})$
(b) 7+ $0.5(\mathrm{pka}-\mathrm{pkb})$
(q) NaCl
(c)7-0.5(pkb $+\operatorname{logC}$ )
(r) $\mathrm{CH}_{3} \mathrm{COONa}$
(d) 0.5 (pkw)

ANS. :
MATCH-I: 1. b, q 2. c, p 3. a, r 4. d, s
MATCH-II : 1. b, s 2. a, r 3. d, q 4. c, p

## 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.

1. 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.
3. Statement -1: The gas phase reaction $\mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{5}(\mathrm{~g})$ shifts to the right on increasing pressure.
Statement-2 : When pressure increase, equilibrium shifts towards more number of moles.
4. Statement-1: The chemical equilibrium is not static but dynamic in nature. Statement-2: The chemical equilibrium is a state in which two opposing process are proceeding at the same rate.
5. Statement-1 : The catalyst does not change the equilibrium constant. Statement-2 : Because for the catalysed reaction and uncatalysed reaction $\Delta \mathrm{H}$ remains same and equilibrium constant depends on $\Delta \mathrm{H}$.
6. Statement-1 : If water is heated to $59^{\circ} \mathrm{C}$, the pH will increase. Statement-2 : $\mathrm{K}_{\mathrm{w}}$ increases with increase in temperature.
7. Statement-1: Addition of HCl (aq.) to $\mathrm{CH}_{3} \mathrm{COOH}$ (aq.) decrease the ionisation of $\mathrm{CH}_{3} \mathrm{COOH}$ (aq.).
Statement-2 : Due to common ion effect $\mathrm{H}^{+}$, ionisation of $\mathrm{CH}_{3} \mathrm{COOH}$ decreases.
8. Statement-1: Sparingly soluble salts AB and $\mathrm{XY}_{2}$ with the same solubility product, will have different solubility.
Statement 2: Solubility of sparingly soluble salts depends upon solubility product.
9. Statement-1 : The ionisation constants of weak diprotic acid are in the order of $\mathrm{Ka}_{1}>\mathrm{Ka}_{2}$.
Statement-2 : Removal of $\mathrm{H}^{+}$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 $\mathrm{pK}_{\mathrm{a}}$.
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
11. What is sum of $\mathrm{pH}+\mathrm{pOH}$ at $25^{\circ} \mathrm{C}$ ?
[Ans. 14]
12. Write the Henderson Hasselbalch equation for acidic buffer Ans. $\mathrm{pH}=\mathrm{pka}+10 \mathrm{~g} \frac{[\mathrm{SALT}]}{[\text { ACID }]}$
13. How is degree of dissociation related with concentration terms and Ka , for weak electrolyte.

$$
\text { Ans. } \alpha=\sqrt{\mathrm{Ka} / \mathrm{c}}
$$

4. How $\mathrm{NH}_{3}$ is defined as Lewis base?
[Ans. It contain Lone paid of electrons]
5. How are $\mathrm{K}_{\mathrm{p}}$ and $\mathrm{K}_{\mathrm{c}}$ related? [Ans. $\mathrm{K}_{\mathrm{p}}=\mathrm{K}_{\mathrm{c}}(\mathrm{RT})^{\Delta \mathrm{n}}$ ]
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 $\mathrm{HCO}_{3}^{-}$?
[Ans. $\mathrm{CO}_{3}{ }^{2-}$ ]
10. What is the nature of $\mathrm{CH}_{3} \mathrm{COOH}$ in conc. HCl solution?
[Ans. Bronsted Base]

## 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 Law of Chemical equilibrium.
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. $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$ are equilibrium constant for reactions (1) and (2)
(i) $\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{g})$
(ii) $\mathrm{NO}(\mathrm{g}) \rightleftharpoons 1 / 2 \mathrm{~N}_{2}(\mathrm{~g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g})$

Calculate the relation between $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$.
7. Write the equilibrium constant expression for the following reaction :
$3 \mathrm{Fe}(\mathrm{s})+4 \mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \rightleftharpoons \mathrm{Fe}_{3} \mathrm{O}_{4}(\mathrm{~s})+4 \mathrm{H}_{2}(\mathrm{~g})$
8. Classify the equilibrium as homogeneous or heterogeneous :
$\mathrm{CH}_{3} \mathrm{COOC}_{2} \mathrm{H}_{5}($ aq. $)+\mathrm{H}_{2} \mathrm{O}(1) \rightleftharpoons \mathrm{CH}_{3} \mathrm{COOH}($ aq. $)+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ (aq.)
9. $\mathrm{K}_{p}=\frac{\left(\mathrm{P}_{\mathrm{NH}_{3}}\right)}{\left(\mathrm{P}_{\mathrm{H}_{2}}\right)^{3 / 2}\left(\mathrm{P}_{\mathrm{N}_{2}}\right)^{1 / 2}}$

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. $\mathrm{Hb}(\mathrm{s})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{HbO}_{2}(\mathrm{~s})$

Predict the direction in which equilibrium gets shifted if partial pressure of $\mathrm{O}_{2}(\mathrm{~g})$ is lowered.
12. Discuss the position of equilibrium if the following reaction is carried out in the presence of catalyst.
$2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g})$
13. Which of the following are Lewis acids?
$\mathrm{H}_{2} \mathrm{O}, \mathrm{BF}_{3}, \mathrm{H}^{+}, \mathrm{NH}_{4}^{+}$
14. Write the conjugate acids for the following Bronsted bases.
$\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{OH}, \mathrm{H}_{2} \mathrm{O}$
15. Write the conjugate bases for the following Bronsted acids. $\mathrm{H}_{2} \mathrm{O}, \mathrm{CH}_{3} \mathrm{COOH}$.
16. Which of the following are Lewis acids?
(a) $\mathrm{H}_{2} \mathrm{O}$
(b) $\mathrm{AlCl}_{3}$
(c) $\mathrm{NH}_{4}^{+}$
17. Define Ostwald's dilution law.
18. $\mathrm{SO}_{3}{ }^{2-}$ 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 $25^{\circ} \mathrm{C}$. Is black coffee acidic or basic ? [Ans. Acidic]
21. What will be the value of $\left(\mathrm{pK}_{\mathrm{a}}+\mathrm{pK}_{\mathrm{b}}\right)$ at $25^{\circ} \mathrm{C}$.
22. What will be the pH of $1{\mathrm{M} \mathrm{KNO}_{3} \text { solutions at } 25^{\circ} \mathrm{C} \text { ? }}^{\text {? }}$
23. $\mathrm{CaCl}_{2}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{O}(l) \rightleftharpoons \mathrm{CaCl}_{2}($ aq. $)+$ Heat

Discuss the solubility of $\mathrm{CaCl}_{2}$ if temperature is increased.
24. Why does the solubility of $\mathrm{CO}_{2}$ decrease with rise in temperature?
25. The solubility of $\mathrm{A}_{2} \mathrm{X}_{3}$ is $\mathrm{y} \mathrm{mol} \mathrm{dm}{ }^{-3}$. Calculate its solubility product.
26. Write the $\mathrm{K}_{\text {sp }}$ expression for $\mathrm{Al}(\mathrm{OH})_{3}$.
27. What is the condition for precipitation of a salt ?
28. Pridict whether the solution is acidic, basic or natural when $\mathrm{NH}_{4} \mathrm{NO}_{3}$ 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 $\mathrm{HCl}, \mathrm{HBr}, \mathrm{HF}, \mathrm{HI}$
[Ans. $\mathrm{HF}<\mathrm{HCl}<\mathrm{HBr}<\mathrm{HI}$ ]
33. Arrange the following in increasing Lewis base strength $\mathrm{NH}_{3}, \mathrm{BiH}_{3}, \mathrm{PH}_{3}, \mathrm{AsH}_{3}, \mathrm{SbH}_{3}$

$$
\text { [Ans. } \mathrm{BiH}_{3}<\mathrm{SbH}_{3}<\mathrm{ASH}_{3}<\mathrm{PH}_{3}<\mathrm{NH}_{3} \text { ] }
$$

34. Arrange following in increasing pH value $0.1 \mathrm{M} \mathrm{CH} 33 \mathrm{COOH}, 0.1 \mathrm{M} \mathrm{NaCl}, 0.1 \mathrm{MHCl}, 0.1 \mathrm{MNaOH}, 0.1 \mathrm{MNH}_{4} \mathrm{OH}$

$$
\text { [Ans. } 0.1 \mathrm{MHCl}<0.1 \mathrm{M} \mathrm{CH}_{3} \mathrm{COOH}<0.1 \mathrm{M} \mathrm{NaCl}<
$$

$$
\left.0.1 \mathrm{NH}_{4} \mathrm{OH}<0.1 \mathrm{M} \mathrm{NaOH}\right]
$$

35. Arrange following in increasing order of degree of hydrolysis.
$0.1 \mathrm{M} \mathrm{NH} 44,0 H, 0.01 \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}, 10^{-5} \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}, 10^{-3} \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}, 10^{-6}$ $\mathrm{M} \mathrm{NH}_{4} \mathrm{OH}$

$$
\begin{array}{r}
{[\text { Ans. } 0.1 \mathrm{M} \mathrm{NH}+4 \mathrm{OH}<} \\
10^{-2} \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}, 10^{-3} \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}< \\
\\
\left.10^{-5} \mathrm{MNH}_{4} \mathrm{OH}<10^{-6} \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}\right]
\end{array}
$$

36. Arrange following in increasing order of acidic strength $\mathrm{CH}_{3} \mathrm{COOH}, \mathrm{HCOOH}, \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{COOH}, \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOH}, \mathrm{CH}_{2} \mathrm{COOH}$
[Ans. $\mathrm{CH}_{3} \mathrm{COOH}<\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOH}<\mathrm{HCOOH}<\mathrm{CH}_{2} \mathrm{FCOOH}$ ]
37. Arrange the basic strength of following $\mathrm{F}^{-}, \mathrm{Br}^{-}, \mathrm{Cl}^{-}, \mathrm{I}^{-}$

$$
\left[\text { Ans. } \mathrm{I}^{-}<\mathrm{Br}^{-}<\mathrm{Cl}^{-}<\mathrm{F}^{-}\right]
$$

38. Arrange the following in increasing base strength $\mathrm{CH}_{3}{ }^{-}, \mathrm{NH}_{2}{ }^{-}, \mathrm{OH}^{-}, \mathrm{F}^{-}$
[Ans. $\mathrm{F}^{-}<\mathrm{OH}^{-}<\mathrm{NH}_{2}<\mathrm{CH}_{3}{ }^{-}$]

## 2-MARKS QUESTIONS

1. Calculate pH of 0.001 M CH 3 COOH having $3 \%$ dissociation.
[Ans. 45229]
2. The equilibrium constant for
$\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{g})$ is K , then calculate equilibrium constant for $1 / 2 \mathrm{~N}_{2}(\mathrm{~g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{NO}(\mathrm{g})$.
3. For the reversible reaction $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})$ at $500^{\circ} \mathrm{C}$, the value of Kp is $1.44 \times 10^{-5} \mathrm{atn}^{-2}$. Find the $\mathrm{K}_{\mathrm{c}}$ value.

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

4. The equilibrium constant at 298 K for the reaction $\mathrm{A}+\mathrm{B} \rightleftharpoons \mathrm{C}+\mathrm{D}$ is 100. If the initial concentration of all the four species were 1 M each, then equilibrium concentration of $D$ will be
[Ans. 1.818]
5. For the reaction
$\mathrm{NH}_{4} \mathrm{COO} \mathrm{NH}_{2}(\mathrm{~s}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})+\mathrm{CO}_{2}(\mathrm{~g})$
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 $\mathrm{NH}_{4} \mathrm{Cl}$ that should be added to one litre of $1.0 \mathrm{~m} \mathrm{NH}_{4} \mathrm{OH} \mathrm{kb} 1.8 \times 10^{-5}$
[Ans. $\mathrm{NH}_{4} \mathrm{Cl}=1.8 \mathrm{M}$ ]
7. Calculate the solubility of silver chloride in water at room temperature if the $\mathrm{K}_{\text {sp }}$ of AgCl is $1.6 \times 10^{-10}$.
[Ans. $1.26 \times 10^{-5} \mathrm{M}$ ]
8. Calculate the molar solubility of $\mathrm{Ni}(\mathrm{OH})_{2}$ in 0.10 m NaOH . The ionic product of $\mathrm{Ni}(\mathrm{OH})_{2}$ is $2.0 \times 10^{-15}$.
[Ans. $2.0 \times 10^{-13} \mathrm{M}$ ]
9. Calculate the pH of $10^{-8} \mathrm{M} \mathrm{HCl}$ 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 12 ?
[Ans. 0.4 g ]

## 3-MARKS QUESTIONS

1. The equilibrium constant for the reaction $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{Br}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HBr}(\mathrm{g})$ at 1024 K is $1.6 \times 10^{5}$. Find the equilibrium pressure of all gases if 10.0 bar of HBr is introduced into a sealed container at 1024 K . [Ans. 10 bar ]
2. For the reaction $2 \mathrm{BrCl}(\mathrm{g}) \rightleftharpoons \mathrm{Br}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \mathrm{K}_{\mathrm{c}}$ is 32 at 500 K . If initially pure BrCl is present at a concentration of $3.30 \times 10^{-3} \mathrm{M}$, what is its molar concentration in the mixture at equilibrium? [Ans. $3.0 \times 10^{-4} \mathrm{M}$ ]
3. When 0.02 M HCl solution is added to $0.01 \mathrm{M} \mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}$, will a precipitate of $\mathrm{PbCl}_{2}$ be formed or not. $\mathrm{K}_{\text {sp }}=1.6 \times 10^{-5}$
[Ans: No]
4. Find the value of Kc for the reaction

$$
2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{2}(\mathrm{~g}) \mathrm{K}_{\mathrm{p}}=3.4 \mathrm{bar}^{-1} \text { at } 1000^{\circ} \mathrm{C}
$$

[Ans. 359.24]
5. Ammonium hydrogen sulphide dissociate according to equation

$$
\mathrm{NH}_{4} \mathrm{HS}(\mathrm{~s}) \rightleftharpoons \mathrm{NH}_{3}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{~S}(\mathrm{~g})
$$

If the observed pressure at equilibrium is 1.12 atm at 380 K . What is Kp for the reaction
[Ans. 0.3136]
6. How much of $0.3 \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}$ should be mixed with 30 mL of 0.2 m solution of $\mathrm{NH}_{4} \mathrm{Cl}$ to give a buffer solution of $\mathrm{pH} 10 . \mathrm{pk}_{\mathrm{b}}$ for $\mathrm{NH}_{4} \mathrm{OH}$ is 4.75 .
[Ans. $\mathrm{V}=112.5 \mathrm{~mL}$ ]
7. Predict whether a precipitate will be formed or not on mixing 20 mL of 0.001 N NaCl solution with 80 mL of $0.01 \mathrm{M} \mathrm{AgNO}_{3}$ solution. $\mathrm{K}_{\mathrm{sp}}$ for AgCl is $1.5 \times 10^{-10}$.
[Ans. Yes, ppt will formed.]
8. The values of Ksp of two sparingly soluble salts $\mathrm{Ni}(\mathrm{OH})_{2}$ and AgCN are $2.0 \times 10^{-15}$ and $6.0 \times 10^{-17}$ respectively. Which salt is more soluble. Explain

$$
\begin{array}{r}
{\left[\text { Ans. } \mathrm{S}_{\mathrm{Ni}(\mathrm{OH})_{2}}=7.93 \times 10^{-6} \mathrm{M}: \mathrm{S}_{(\mathrm{Ag} \mathrm{CN})}=7.8 \times 10^{-9} \mathrm{M} \cdot \mathrm{Ni}(\mathrm{OH})_{2}\right.} \\
\text { is more soluble] }
\end{array}
$$

9. The ionization constant of propanoic acid is $1.32 \times 10^{-5}$. 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. $1.62 \times 10^{-2}, 1.32 \times 10^{-3}$ ]
10. Calculate the pH of a solution obtained by mixing 50 ml of 0.2 M HCl with 49.9 mL of 0.2 m NaOH solution.
[Ans. 3.699]

## HOTS QUESTIONS

1. The molar solubility of $\mathrm{Cd}(\mathrm{OH})_{2}$ is $1.84 \times 10^{-5} \mathrm{M}$. Calculate the expected solubility of $\mathrm{Cd}(\mathrm{OH})_{2}$ in a buffer solution of $\mathrm{pH}=12$.

Ans. $\mathrm{Cd}(\mathrm{OH})_{2} \rightleftharpoons \mathrm{Cd}_{\text {(aq.) }}^{2+}+2 \mathrm{OH}_{\text {(aq.) }}^{-}$

$$
\mathrm{S} \quad 10^{-2}
$$

$2.49 \times 10^{-14}=\mathrm{S}\left(10^{-2}\right)^{2} \quad \therefore \mathrm{~S}=2.49 \times 10^{-10 \mathrm{M}}$
2. An aqueous solution contains an unknown concentration of $\mathrm{Ba}^{2+}$. When 50 ml of a 1 M solution of $\mathrm{Na}_{2} \mathrm{SO}_{4}$ is added. $\mathrm{BaSO}_{4}$ just begins to precipitate. The final volume is 500 ml . The solubility product of $\mathrm{BaSO}_{4}$ is $1 \times 10^{-10}$. Find the original concentration.

Ans. $\mathrm{K}_{\text {sp }}=\left[\mathrm{Ba}^{2+}\right]\left[\mathrm{SO}_{4}^{2-}\right]=\left[\mathrm{Ba}^{2+}\right]\left[\frac{50 \times 1}{500}\right]=10^{-9} \times 500$

$$
\begin{aligned}
& \mathrm{Ba}^{2+}=10^{-9} \mathrm{M} \\
& 10^{-9} \times 500=450 \times \mathrm{M} \quad \therefore \mathrm{M}=1.11 \times 10^{-9} \mathrm{M}
\end{aligned}
$$

3. An aqueous solution contains $0.10 \mathrm{M} \mathrm{H}_{2} \mathrm{~S}$ and 0.20 M HCl . If the equilibrium constants for the formation of $\mathrm{HS}^{-}$from $\mathrm{H}_{2} \mathrm{~S}$ is $1.0 \times 10^{-7}$ and that of $\mathrm{S}^{2-}$ from $4 \mathrm{~S}^{-}$ions is $1.2 \times 10^{-13}$, then find the concentration of $\mathrm{S}^{-2}$ ions in aqueous solution.

Ans. $\quad \mathrm{H}_{2} \mathrm{~S}($ aq. $) \rightleftharpoons 2 \mathrm{H}^{+} \quad+\mathrm{S}^{2-}$

$$
\begin{aligned}
&(0.1-\mathrm{x}) \\
& \mathrm{K}_{\mathrm{a}}= \mathrm{K}_{\mathrm{a}_{1}} \times \mathrm{K}_{\mathrm{a}_{2}}=1.2 \times 10^{-20} \\
& 1.2 \times 10^{-20}=\frac{(0.2)^{2}\left[\mathrm{~S}^{2-}\right]}{0.1},\left[\mathrm{~S}^{2-}\right]=3 \times 10^{-20}
\end{aligned}
$$

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. $\quad 0.1 \times 1=(1+\mathrm{v})(0.01) \Rightarrow \mathrm{v}=9 \mathrm{~L}$
5. A certain buffer solution contains equal concentration of $\mathrm{X}^{-}$and HX . The $\mathrm{K}_{\mathrm{b}}$ for $\mathrm{X}^{-}$is $10^{-10}$. Find the pH of the buffer .

Ans. $\mathrm{k}_{\mathrm{a}} \cdot \mathrm{k}_{\mathrm{b}}=10^{-14} \quad \therefore \mathrm{k}_{\mathrm{a}}=\frac{10^{-14}}{10^{-10}}=10^{-4}$

$$
\begin{aligned}
\mathrm{pH} & =\mathrm{pka}+\log \frac{\left[\mathrm{x}^{-}\right]}{[\mathrm{Hx}]} \\
\therefore \mathrm{pH} & =4+\log \frac{1}{1}=4 \quad \therefore \mathrm{pH}=4
\end{aligned}
$$

6. The $\%$ yield of Ammonia as a function of time in the reaction $\mathrm{N}_{2}(\mathrm{~g})+$ $3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}), \Delta \mathrm{H}<\mathrm{O}$ at $(\mathrm{P}, \mathrm{T})$ is given below:


If this reaction is conducted at $\mathrm{T}_{2}>\mathrm{T}_{1}$, then plot the $\%$ yield of $\mathrm{NH}_{3}$ 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 $\mathrm{NH}_{3}$ get decreased overall after a certain interval of time.

7. Consider the reaction $\mathrm{NH}_{4} \mathrm{COONH}_{2}(\mathrm{~s}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})+\mathrm{CO}_{2}(\mathrm{~g})$ at a certain temperature, the equilibrium pressure of the system is 0.318 atm . Find $\mathrm{K}_{\mathrm{p}}$ of the decomposition of ammonium carbonate.

Ans. $\quad P_{\text {total }}=3 P \quad \therefore \mathrm{P}=0.318 / 3=0.106$

$$
\mathrm{Kp}=4 \mathrm{P}^{3}=4.76 \times 10^{-3}
$$

8. The equilibrium constant for the reaction $\mathrm{CO}(\mathrm{g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \rightleftharpoons \mathrm{CO}_{2}(\mathrm{~g})+$ $\mathrm{H}_{2}(\mathrm{~g})$ is 5 . How many moles of $\mathrm{CO}_{2}$ must be added to 1 litre container already containing 3 moles each of CO and $\mathrm{H}_{2} \mathrm{O}$ to make 2 M equilibrium concentration of CO ?

Ans.

$$
\mathrm{CO}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{CO}_{2}+\mathrm{H}_{2}
$$

$$
\mathbf{t}=\mathbf{0} \quad 3 \quad 2 \quad \begin{array}{lllll} 
& 2 & x & 0
\end{array}
$$

$\begin{array}{lllll}\text { At equilibrium } & 2 & 2 & \mathrm{x}+1 & 1\end{array}$

$$
\therefore S=\frac{x+1}{4} \Rightarrow x=19
$$

9. At constant temperature, the equilibrium constant $\mathrm{K}_{\mathrm{p}}$ $\mathrm{N}_{2} \mathrm{O}_{4} \rightleftharpoons 2 \mathrm{NO}_{2}$ is given by $\mathrm{k}_{\mathrm{p}}=\frac{4 \mathrm{x}^{2} \mathrm{P}}{1-\mathrm{x}} \quad$ where, $\mathrm{P}=$ Pressure and $\mathrm{X}=$ 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 $\mathrm{Q}=0$. As the reaction proceeds in the forward direction $Q$ starts increasing.

At chemical equilibrium $\mathrm{Q}=\mathrm{K}$

## UNIT TEST-I

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. What is the $\mathrm{pH} 10^{-3} \mathrm{M} \mathrm{HCl}$ solution ?
(a) 1
(b) 11
(c) 3
(d) 14
2. Which one can act as Arrhenius Acid ?
(a) $\mathrm{NH}_{3}$
(b) $\mathrm{H}_{2} \mathrm{O}$
(c) HCl
(d) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{OH}$
3. Write the conjugate base of $\mathrm{CH}_{3} \mathrm{COOH}, \mathrm{H}_{2} \mathrm{O}$.
4. Write the relation between $\mathrm{K}_{\mathrm{p}}$ and $\mathrm{K}_{\mathrm{c}}$.
5. What is the nature of following reaction

Exothermic or endothermic
$\mathrm{A}+\mathrm{B}-70 \mathrm{~J} \longrightarrow \mathrm{C}$
6. The pka of $\mathrm{CH}_{3} \mathrm{COOH}$ and pkb of $\mathrm{NH}_{4} \mathrm{OH}$ are 4.76 and 4.75 respectively. Calculate the pH of $\mathrm{CH}_{3} \mathrm{COONH}_{4}$.
7. What is a buffer solution. Calculate the pH of the solution obtained by adding 4 mol of $\mathrm{CH}_{3} \mathrm{COOH}$ with 3 mol of NaOH in 1 litre container. [2] pka, $\mathrm{CH}_{3} \mathrm{COOH}=4.74 \log 2=0.3010 \log 3=0.4771$
8. Calculate the molar solubility of $\mathrm{Ni}(\mathrm{OH})_{2}$ in 0.1 M KOH solution. The $\mathrm{K}_{\text {sp }}$ for $\mathrm{Ni}(\mathrm{OH})_{2}$ is $2.0 \times 10^{-15}$.
9. $\mathrm{K}_{\mathrm{p}}=0.04 \mathrm{~atm}$ at 899 K for the equilibrium shown below. What is the equilibrium concentration of $\mathrm{H}_{2}$ when it is placed in a flask at 4.0 atm pressure and allowed to come to equilibrium.
$\mathrm{C}_{2} \mathrm{H}_{6}(\mathrm{~g}) \rightleftharpoons \mathrm{C}_{2} \mathrm{H}_{4}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g})$
10. The first ionization constant of $\mathrm{H}_{2} \mathrm{~S}$ is $9.1 \times 10^{-4}$. Calculate the concentration of $\mathrm{HS}^{-1}$ in its 0.1 M solution. How will this concentration be effected if the solution is 0.1 M HCl also? If the second dissociation constant of $\mathrm{H}_{2} \mathrm{~S}$ is $1.2 \times 10^{-12}$. Calculate the concentration of $\mathrm{S}^{2-}$ in both conditions. [5]

## UNIT TEST-II

Time Allowed: $\mathbf{1} \mathbf{H r}$.
Maximum Marks : 20
General Instructions:
(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. What is the $\mathrm{pH} 10^{-8} \mathrm{MHCl}$ solution ?
(a) $\mathrm{pH}>7$
(b) $\mathrm{pH}<7$
(c) $\mathrm{pH}=7$
(d) Cannot be defined
2. What is the conjugate acid for the $\mathrm{NH}_{3}$ ?
(a) $\mathrm{NH}_{2}^{-}$
(b) $\mathrm{NH}^{2-}$
(c) $\mathrm{N}^{3-}$
(d) $\mathrm{NH}_{4}^{+}$
3. Define Lewis acid and base with one example each.

In following questions a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.
(a) Assertion and Reason both are correct statement and reason is correct explanation for assertion.
(b) Assertion and Reason both are correct statements but reason is not correct explanation for assertion.
(c) Assertion is correct statement but reason is wrong statement.
(d) Assertion is wrong statement but reason is correct statement.
4. Assertion : HI is stronger acid than HCl

Reason : Size of I is more than Cl and hence H-I bond strength is less than HCl .
5. Assertion : $\mathrm{BF}_{3}$ is Lewis acid and $\mathrm{NH}_{3}$ is Lewis base.

Reason : $\mathrm{NH}_{3}$ is short of octet and $\mathrm{BF}_{3}$ molecule contain lone pair of electron.
6. Arrange the following in increasing acidic strength. Give reason also $\mathrm{CH}_{4}, \mathrm{NH}_{3}, \mathrm{H}_{2} \mathrm{O}, \mathrm{HF}$
7. $\mathrm{K}_{\mathrm{c}}$ for the reaction $\mathrm{SO}_{2}+0.5 \mathrm{O}_{2} \rightarrow \mathrm{SO}_{3}$ at $600^{\circ} \mathrm{C}$ is 61.7. Calculate $\mathrm{K}_{\mathrm{p}}$.
8. 25.4 ml of hydrogen and 20.4 ml of iodine when heated in a closed container, produced 30.8 mL of HI at equilibrium. Calculate the degree of dissociation of HI at same temperature.
9. Define common ion effect. The solubility of $\mathrm{CaF}_{2}$ in water at T K is $2 \times 10^{-4} \mathrm{moles} / \mathrm{L}$. Calculate (i) $\mathrm{K}_{\text {sp }}$, and (ii) Solubility in 0.01 M NaF solution.
10. (i) What is a buffer solution? What are its types?
(ii) Derive the Henderson-Hasselbalch equation for an Acidic buffer with the help of relevant example.
(ii) 8 g of NaOH was dissolved in one litre of a solution containing one mole of $\mathrm{CH}_{3} \mathrm{COOH}$ and one mole of $\mathrm{CH}_{3} \mathrm{COONa}$. Find the pH of the resulting solution. (The $\mathrm{pK}_{\mathrm{a}}$ of $\mathrm{CH}_{3} \mathrm{COOH}$ is 4.74).

## 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 : Reduce other substance and oxidise itself.
- Oxidising Agent : Oxidise other substance but reduce itself.
- 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 $\mathrm{Cl}_{2}, \mathrm{~F}_{2}, \mathrm{O}_{2}, \mathrm{P}_{4}, \mathrm{O}_{3}, \mathrm{Fe}, \mathrm{H}_{2}, \mathrm{~N}_{2}, \mathrm{C}$ (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., $\mathrm{H}_{2} \mathrm{O}_{2}, \mathrm{Na}_{2} \mathrm{O}_{2}$ ), each oxygen atom is assigned an oxidation number of -1 , in super oxides (e.g., $\mathrm{KO}_{2}, \mathrm{RbO}_{2}$ ) each oxygen atom is assigned an oxidation number of $-(1 / 2)$.
(e) In oxygen difluoride $\left(\mathrm{OF}_{2}\right)$ and dioxygen difluoride $\left(\mathrm{O}_{2} \mathrm{~F}_{2}\right)$, 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: $0 \quad 0 \quad+2-3$

$$
3 \mathrm{Mg}(\mathrm{~s})+\mathrm{N}_{2}(\mathrm{~g}) \xrightarrow{\Delta} \mathrm{Mg}_{3} \mathrm{~N}_{2}(\mathrm{~s})
$$

(ii) Decomposition Reaction : +1+5-2 $+\mathbf{1 - 1} 0$

$$
2 \mathrm{KClO}_{3}(\mathrm{~s}) \xrightarrow{\Delta} \quad 2 \mathrm{KCl}(\mathrm{~s})+3 \mathrm{O}_{2}(\mathrm{~g})
$$

(iii) Metal Displacement : +2+6-2 0 $\mathbf{0}+2+6-2 \quad 0$

$$
\mathrm{CuSO}_{4}(\mathrm{aq})+\mathrm{Zn}(\mathrm{~s}) \rightarrow \mathrm{ZnSO}_{4}(\mathrm{aq})+\mathrm{Cu}(\mathrm{~s})
$$

(iv) Non-metal displacement :0 +1-2 $\mathbf{0}+2-2+1 \quad 0$

$$
\mathrm{Ca}(\mathrm{~s})+2 \mathrm{H}_{2} \mathrm{O}(1) \rightarrow \mathrm{Ca}(\mathrm{OH})_{2}+\mathrm{H}_{2}(\mathrm{~g})
$$

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

$$
\begin{aligned}
& \mathbf{0} \\
& \mathrm{Cl}_{2}(\mathrm{~g})+2 \mathrm{OH}^{-}(\mathrm{aq}) \rightarrow \mathrm{Cl}^{-}(\mathrm{aq})+\mathrm{ClO}^{-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(1)
\end{aligned}
$$

- 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 $\mathrm{Fe}_{2}$ (III) $\mathrm{O}_{3}$ 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 ( $\mathbf{E}^{\boldsymbol{\theta}}$ ) : Electrode Potential measured at 298 K and 1 M 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: $\mathbf{Z n}\left|\mathbf{Z n}^{2+} \| \mathbf{C u}^{\mathbf{2 +}}\right| \mathbf{C u}$
LHS oxidation, $\mathrm{Zn} \rightarrow \mathrm{Zn}^{2+}+2 e^{-}$ RHS reduction $\mathrm{Cu}^{2+}+2 e^{-} \rightarrow \mathrm{Cu}$
Overall reaction $\mathrm{Zn}(\mathrm{s})+\mathrm{Cu}^{2+}(a q) \rightarrow \mathrm{Zn}^{2+}(a q)+\mathrm{Cu}(\mathrm{s})$

- Representation of an Electrochemical cell :
$\qquad$
$\longleftarrow-$ Flow of current

$$
\mathrm{Zn}(s)\left|\mathrm{Zn}^{2+}(a q) \| \mathrm{Cu}^{2+}(a q)\right| \mathrm{Cu}(s)
$$

| LOAN | Left Electrode | Salt Bridge | Right Electrode |
| :--- | :--- | :--- | :--- |
|  | Oxidation |  | Reduction |
|  | Anode |  | Cathode |
|  | Negative |  | Positive |

- Functions of Salt Bridge : (i) To complete inner circuit. (ii) To maintain electrical neutrality around electrodes.
MIND MAP : REDOX REACTIONS



## CASE BASED STUDY - QUESTIONS

1. Read the given passage and answer the questions.

Redox reactions are reactions in which oxidation and reduction takes place simultaneously. Oxidation number are assigned in accordance with the set of rules. Oxidation number and ion electron methods both are used in balancing ionic equations. Redox reactions are classified as combination, decomposition, displacement and disproportionation reactions. The concept of redox couple and electrode processes is basis of electrolysis and electrochemical cells.
(a) What are oxidation number of each individual Br in $\mathrm{Br}_{3} \mathrm{O}_{8}$

Ans. $+6,+4,+6$

(b) If electrolysis of $\mathrm{CuSO}_{4}$, solution is carried out using Cu electrodes, what will be reaction taking place at anode.

Ans. $\mathrm{Cu} \rightarrow \mathrm{Cu}^{2+}+2 \mathrm{e}^{-}$
(c) What is oxidation number of Cr in $\mathrm{CrO}_{5}$ ?

Ans. +6

$\because$ It has peroxide linkage.
(d) Give one example of disproportionation reaction.

Ans. $2 \mathrm{Cu}^{+} \rightarrow \mathrm{Cu}^{2+}+\mathrm{Cu}$
(e) $\mathrm{MnO}_{4}^{2-}+\mathrm{H}^{+} \rightarrow \mathrm{MnO}_{4}^{-}+\mathrm{MnO}_{2}+\mathrm{H}_{2} \mathrm{O}$ [Balance this reaction]

Ans. $\mathrm{MnO}_{4}^{2-} \rightarrow \mathrm{MnO}_{4}^{-}+\mathrm{e}^{-}$

$$
\begin{align*}
& 2 \mathrm{e}^{-}+4 \mathrm{H}^{+}+\mathrm{MnO}_{4}^{2-} \rightarrow \mathrm{MnO}_{2}+2 \mathrm{H}_{2} \mathrm{O}  \tag{ii}\\
& 3 \mathrm{MnO}_{4}^{2-}+4 \mathrm{H}^{+} \rightarrow 2 \mathrm{MnO}_{4}^{-}+\mathrm{MnO}_{2}+2 \mathrm{H}_{2} \mathrm{O}
\end{align*}
$$

## 2. Redox Reactions : Passage Based Question (Assertion and Reason)

Passage : Redox reactions are those reactions in which, there is a simultaneous oxidation and reduction taking place. There is an addition of oxygen and removal of hydrogen taking place in oxidation reactions. In Reduction, hydrogen gets added and oxygen gets removed. Redox reactions are also used to determine the strength of reductant/oxidant. In oxidation, there is a decrease in electron density while in reduction, there is an increase in electron density around the atom.
(Q1-Q4) There are assertion and reason which have been put forward. Read the given statement and choose correct alternative from the following :
(Note : A-Assertion \& R-Reason)
(a) Both A and R are true and R is the correct explanation of A .
(b) Both A and R are true but R is not the correct explanation of A .
(c) A is true but R is false.
(d) A is false but R is true.

Q1. A: Oxidation-Reduction (Redox) couple is the combination of oxidized and reduced form of a substance that is involved in Oxidation-Reduction half cell.
R: As in representation $\mathrm{E}^{\circ} \mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}$ and $\mathrm{E}^{\circ} \mathrm{Cu}^{2+} / \mathrm{Cu}^{+}$are two Redox couples.

Q2. A: In Formaldehyde (HCHO) oxidation no. of carbon is 0 .
R: Formaldehyde is a covalent compound.
Q3. A: Oxidation state of hydrogen is water is +1 and $\mathrm{CaH}_{2}$ is -1 .
R: $\mathrm{CaH}_{2}$ is metal hydride and for hydrogen, it assigned the oxidation state of -1 .

Q4. A: Redox reactions are also called neutralization reactions.
R: As the number of electron gained/lost in the reaction are balanced.

## Or

A: Substances which get reduced an act as reducing agent.
R: Oxidizing agent itself gets reduced.
Ans. : Q1. (c), Q2. (b), Q3. (a), Q4. (d) or (d)

## MULTIPLE CHOICE QUESTIONS (MCQ)

1. The oxidation state of Fe in $\mathrm{Fe}_{3} \mathrm{O}_{4}$ is
(a) +2
(b) +3
(c) $\frac{8}{3}$
(d) $+2,+3$
2. The oxidation state of ' S ' in $\mathrm{KAl}\left(\mathrm{SO}_{4}\right)_{2} \cdot 12 \mathrm{H}_{2} \mathrm{O}$ is
(a) -2
(b) -1
(c) 2
(d) +6
3. Oxidation state carbon in $\mathrm{C}_{3} \mathrm{O}_{2}$ is
(a) $\frac{4}{3}$
(b) 0
(c) 2
(d) 0,2
4. The reaction $\mathrm{S}_{8}+12 \mathrm{OH}^{-} \longrightarrow 4 \mathrm{~S}^{2-}+2 \mathrm{~S}_{2} \mathrm{O}_{3}{ }^{2-}+6 \mathrm{H}_{2} \mathrm{O}$ is
(a) Combination reaction
(b) Decomposition reaction
(c) Non-metal displacement
(d) Disproportionation reaction
5. $\mathrm{E}^{0}$ for $\mathrm{H}^{+} / \mathrm{H}_{2}$ is
(a) 0
(b) +1 V
(c) -1.0 V
(d) -2.0 V
6. Which one act as strong oxidising agent
$\mathrm{K}^{+} / \mathrm{K}=-2.93 \mathrm{~V}, \mathrm{Ag}^{+} / \mathrm{Ag}=0.80, \mathrm{Hg}^{2+} / \mathrm{Hg}=0.79 \mathrm{~V}$
(a) $\mathrm{K}^{+}$
(b) K
(c) $\mathrm{Hg}^{2+}$
(d) $\mathrm{Ag}^{+}$
7. The coefficient of HCl in balance reaction is
$\mathrm{Pb}_{3} \mathrm{O}_{4}+\mathrm{HCl} \longrightarrow \mathrm{PbCl}_{2}+\mathrm{Cl}_{2}+\mathrm{H}_{2} \mathrm{O}$
(a) 1
(b) 8
(c) 3
(d) 4
8. Sum of oxidation numbers of all Bromine atoms in $\mathrm{Br}_{3} \mathrm{O}_{8}$ is
(a) 6
(b) 4
(c) 16
(d) 20
9. In the reaction $6 \mathrm{ClO}_{2}^{-} \longrightarrow 4 \mathrm{ClO}_{3}^{-}+2 \mathrm{Cl}^{-}, \mathrm{Cl}^{-}$ion is
(a) Oxidised Reduced
(b) Reduced
(c) Odixised and
(d) Neither Oxidised nor reduced
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 $\qquad$ of electrons.
(ii) S.H.E. stands for $\qquad$ .
(iii) Oxidation state of Oxygen in $\mathrm{O}_{2} \mathrm{~F}_{2}$ is $\qquad$ .
(iv) Disproportionation is a type of $\qquad$ reaction.
(v) Oxidant is one which $\qquad$ electron.
(vi) $\mathrm{Cl}_{2}+2 \mathrm{OH}^{-} \longrightarrow \mathrm{ClO}^{-}+\mathrm{Cl}^{-}$is a $\qquad$ type of reaction.
(vii) Oxidation state of F is always either $\qquad$ or $\qquad$ .
(viii) Oxidation state of Oxygen in $\mathrm{O}_{3}$ is $\qquad$ .
(ix) Reducing agent are also called $\qquad$ .
(x) Hydrogen economy is use of Hydrogen as $\qquad$ .

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) $\mathrm{O}_{2}^{+}$has oxidation state of oxygen as $+1 / 2$.
(v) If for the reaction $\mathrm{Ca}^{2+}+2 \mathrm{e}^{-} \longrightarrow \mathrm{Ca}(\mathrm{s}) ; \mathrm{E}^{\Theta}=-2.87$

Then for the reaction $2 \mathrm{Ca}^{2+}+4 \mathrm{e}^{-} \longrightarrow 2 \mathrm{Ca}(\mathrm{s}) ; \mathrm{E}^{\Theta}=2(-2.87) \mathrm{V}$
(vi) Salt bridge is used for enhancing $\mathrm{E}^{\circ}$ 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) $\mathrm{MnO}_{4}^{-}$is colourless in basic medium.
$\begin{array}{rllll}\text { Ans: (i) False } & \text { (ii) False } & \text { (iii) True } & \text { (iv) True } & \text { (v) False } \\ \text { (vi) False } & \text { (vii) True } & \text { (viii) True } & \text { (ix) False } & \text { (x) False }\end{array}$

## MATCH THE COLUMNS

## Column-I

## Column-II

(a) $\mathrm{CH}_{4}+2 \mathrm{O}_{2} \longrightarrow \mathrm{CO}_{2}+2 \mathrm{H}_{2} \mathrm{O}$
(p) Disproportionation
(b) $\mathrm{Cr}_{2} \mathrm{O}_{3}+2 \mathrm{Al} \longrightarrow \mathrm{Al}_{2} \mathrm{O}_{3}+2 \mathrm{Cr}$
(q) Non-metal displacement
(c) $\mathrm{Fe}+2 \mathrm{HCl} \longrightarrow \mathrm{FeCl}_{2}+\mathrm{H}_{2}$
(r) Metal displacement
(d) $\mathrm{P}_{4}+3 \mathrm{OH}^{-}+3 \mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{PH}_{3}+3 \mathrm{H}_{2} \mathrm{PO}_{2}^{-}$
(s) Metal displacement
Ans. (a) - (s)
(b) $-(\mathrm{r})$
(c) $-(q)$
(d) $-(\mathrm{p})$

## Column-I

(a) NO
(p) +5
(b) $\mathrm{NO}_{2}$
(q) +3
(c) $\mathrm{NO}_{2}^{-}$
(r) +4
(d) $\mathrm{NO}_{3}^{-}$
(s) +2
Ans. (a) - (s)
(b) $-(\mathrm{r})$
(c) $-(q)$
(d) $-(\mathrm{p})$

## Column-I

(A) Increase in oxidation number
(B) Reducing agent
(C) $\mathrm{MnO}_{2}+4 \mathrm{HCl} \rightarrow \mathrm{MnCl}_{2}+\mathrm{Cl}_{2}+\mathrm{H}_{2} \mathrm{O}$
(D) Photosynthesis
Ans. (A) - (b)
(B) $-(\mathrm{a})$
(C) - (d)
(d) (D) - (c)

Column-I
(A) Decrease in oxidation number
(B) Oxidizing agent
(C) $2 \mathrm{Cu}^{+} \rightarrow \mathrm{Cu}^{2+}+\mathrm{Cu}$
(D) $\mathrm{Mn}_{3} \mathrm{O}_{4}$
(a) Disproportionation
(b) Fractional oxidation number
(c) Reduction
(d) Gain of electron

Column-II
(Oxidation sate of N )
Ans. (A) - (c)
(B) $-(\mathrm{d})$
(C) $-(a)$
5.

## Column-I

Column-II
(A) $\mathrm{H}_{2} \mathrm{O}_{2}$
(a) -1
(B) $\mathrm{MnSO}_{4}$
(b) +3
(C) $\mathrm{AlCl}_{3}$
(c) +5
(D) $\mathrm{P}_{2} \mathrm{O}_{5}$
(d) +6
Ans. (A) - (a)
(B) $-(\mathrm{d})$
(C) - (b)
(D) - (c)

## 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 ' $\mathrm{F}^{\prime}$ is -1

Statement - 2: 'F' being most electronegative, will have -1 oxidation in its compound.
2. Statement - 1: Oxygen has zero oxidation state in $\mathrm{O}_{2}$.

Statement - 2: Element in their elemental form have zero oxidation state.
3. Statement - 1: Oxidation state of Oxygen in $\mathrm{H}_{2} \mathrm{O}_{2}$ is -1 .

Statement - 2: $\quad \mathrm{H}_{2} \mathrm{O}_{2}$ has peroxide linkage.
4. Statement - 1: For the reaction $\mathrm{Zn}+\mathrm{Cu}^{2+} \longrightarrow \mathrm{Zn}^{2+}+\mathrm{Cu}$; $\mathrm{E}_{\text {cell }}$ is +ve .

Statement - 2 : For standard Hydrogen electrode $\mathrm{E}^{\circ}=0$
5. Statement - 1: $2 \mathrm{H}_{2} \mathrm{O}_{2} \longrightarrow 2 \mathrm{H}_{2} \mathrm{O}+\mathrm{O}_{2}$ is Decomposition reaction (Redox).

Statement - 2: Oxygen has -2 oxidation state in $\mathrm{H}_{2} \mathrm{O}$.
6. Statement - 1: $\mathrm{C}+\mathrm{O}_{2} \longrightarrow \mathrm{CO}_{2}$; carbon get oxidised.

Statement - 2: Gain of Hydrogen is reduction.
7. Statement - 1: $\mathrm{CaCO}_{3} \longrightarrow \mathrm{CaO}+\mathrm{CO}_{2}$ is not redox reaction.

Statement - 2: $\mathrm{C}, \mathrm{Ca}, \mathrm{O}$ do not change their oxidation number in the reaction.
8. Statement - 1: Oxidation also occurs when decrease in electron density is observed.
Statement - 2: Oxidation is gain of electro-positive element.
9. Statement - 1: $\quad \mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}$ is a self indicator.

Statement - $2: \quad \mathrm{MnO}_{4}^{-}$acts as a self indicator.
10. Statement-1: Equivalence point comes first before end point.

Statement - 2 : Equivalence point cannot be obtained even by graphical 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 $\mathrm{S}_{8}$.
2. What is the oxidation state of Oxygen in $\mathrm{H}_{2} \mathrm{O}_{2}$.
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 $3 \mathrm{Mg}+\mathrm{N}_{2} \longrightarrow \mathrm{Mg}_{3} \mathrm{~N}_{2}$, Nitrogen is oxidised or reduced.
Ans: 1. zero
8. 1
9. $\mathrm{NH}_{4} \mathrm{Cl}$ or KCl
10. $\mathrm{KMnO}_{4}$
11. Reduction
12. $\mathrm{H}^{+} \quad$ 7. Reduced
1-MARK QUESTIONS
13. Define oxidation and reduction according to electronic concept.
14. Define oxidation and reduction according to oxidation number.
15. A freshly cut apple is almost white but it turns reddish brown after sometime. Give reason.
16. Define oxidation number.
17. Write oxidation number of Mn in $\mathrm{KMnO}_{4}$.
18. Write oxidation number of Cr in $\mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}$.
19. Write Stock notation of $\mathrm{MnO}_{2}$ and $\mathrm{AuCl}_{3}$.
20. Define redox reaction with example.
21. Define disproportionation reaction. Give one example.
22. Define the term redox titration.
23. Name the indicator used in redox titration involving $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$ as an oxidizing agent.
24. At what concentration of $\mathrm{Cu}^{2+}$ (aq.) will electrode potential become equal to its standard electrode potential?
[Ans. 1 M]
25. The standard reduction potentials of three metals cations $\mathrm{X}, \mathrm{Y}$ and Z are $+0.52,-3.03$ and -1.18 V respectively. Arrange $\mathrm{X}, \mathrm{Y}$ and Z in order of increasing reducing power.
[Ans. $\mathrm{X}<\mathrm{Z}<\mathrm{Y}$ ]
26. An electrochemical cell consists of two electrodes i.e., Anode and Cathode. What is the direction of flow of electrons in this cell?
27. Why anode is negatively charged in an electrochemical cell?
28. Out of Zn and Cu vessel one will be more suitable to store 1 M HCl ?
[Ans. Cu]
Given $\mathrm{E}_{\mathrm{Zn}^{2+} / \mathrm{Zn}}^{\theta}=-0.76 \mathrm{~V}, \mathrm{E}_{\mathrm{Cu}}{ }^{2-/} / \mathrm{Cu}=+0.34 \mathrm{~V}$.
29. Is it safe to stir $1 \mathrm{M} \mathrm{AgNO}_{3}$ solution with copper spoon?

Given $\mathrm{E}_{\mathrm{Ag}^{+} / \mathrm{Ag}}^{\theta}=+0.80 \mathrm{~V}, \mathrm{E}_{\mathrm{Cu}^{2-} / \mathrm{Cu}}^{\theta}=+0.34 \mathrm{~V}$.
[Ans. No]

## 2-MARKS QUESTIONS

1. Identify oxidant and reductant in the reaction :

$$
\mathrm{I}_{2}(\mathrm{aq})+2 \mathrm{~S}_{2} \mathrm{O}_{3}{ }^{2-}(\mathrm{aq}) \longrightarrow 2 \mathrm{I}^{-}(\mathrm{aq})+\mathrm{S}_{4} \mathrm{O}_{6}{ }^{2-}(\mathrm{aq}) .
$$

2. Calculate oxidation number of Fe in $\mathrm{Fe}_{3} \mathrm{O}_{4}$ 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 : $\mathrm{H}_{2} \mathrm{O}(\mathrm{s})+\mathrm{F}_{2} \longrightarrow \mathrm{HF}+\mathrm{HOF}$ 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.
7. Split the reaction $2 \mathrm{~K}(\mathrm{~s})+\mathrm{C1}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{KC1}(\mathrm{~s})$ into oxidation and reduction half reactions.
8. Calculate the oxidation number of underlined elements in following compounds:
(i) $\mathrm{CaO} \underline{O}_{2}$
(ii) $\mathrm{H}_{2} \underline{\mathrm{~S}}_{2} \mathrm{O}_{7}$
(iii) $\mathrm{K}_{2} \underline{\mathrm{MnO}}_{4}$ (iv) $\mathrm{KI}_{3}$
9. Write the functions of salt bridge in an electrochemical cell.
10. Define the term redox couple. Write the practical application of redox couple.
11. 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.

## 3-MARKS QUESTIONS

1. Calculate oxidation number of :
(i) Cr in $\mathrm{Cr}_{2} \mathrm{O}_{4}{ }^{2-}$
(ii) O in $\mathrm{KO}_{2}$
(iii) Na in $\mathrm{Na}_{2} \mathrm{O}_{2}$.
2. Account for the following :
(i) $\mathrm{HNO}_{3}$ acts as oxidizing agent while $\mathrm{HNO}_{2}$ can act both as reducing and oxidizing agent.
(ii) $\mathrm{AgF}_{2}$ is unstable compound and act as a strong oxidizing agent.
(iii) Ozone acts as an oxidising agent.
3. Permanent ion $\left(\mathrm{MnO}_{4}^{-}\right)$reacts with sulfur dioxide gas in acidic medium to produce $\mathrm{Mn}^{2+}$ ion and hydrogen sulphate ion. Write ionic equation and balance by ion electron method.
4. Balance the following equation by oxidation number method :

$$
\mathrm{P}_{4}(\mathrm{~s})+\mathrm{OH}^{-}(\mathrm{aq}) \longrightarrow \mathrm{PH}_{3}+\mathrm{H}_{2} \mathrm{PO}_{2}^{-}(\mathrm{aq})[\text { Basic Medium }]
$$

5. Balance the following equation by ion electron method :

$$
\mathrm{C1}_{2} \mathrm{O}_{7}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}_{2}(1) \longrightarrow \mathrm{ClO}_{2}^{-}(\mathrm{aq})+\mathrm{O}_{2}(\mathrm{~g})[\text { Basic medium }]
$$

6. Depict the galvanic cell in which the reaction
$\mathrm{Zn}(\mathrm{s})+2 \mathrm{Ag}^{+}(\mathrm{aq}) \longrightarrow \mathrm{Zn}^{2+}(\mathrm{aq})+2 \mathrm{Ag}(\mathrm{s})$ 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 $\mathrm{FeSO}_{4}(\mathrm{aq})+\mathrm{Cu}(\mathrm{s}) \longrightarrow \mathrm{CuSO}_{4}(\mathrm{aq})+\mathrm{Fe}$ does not occur.
(ii) Zinc can displace copper from aqueous $\mathrm{CuSO}_{4}$ solution but Ag cannot.
(iii) Solution of $\mathrm{AgNO}_{3}$ turns blue when copper rod is immersed in it.

## 5-MARKS QUESTIONS

1. (i) $\mathrm{MnO}_{4}{ }^{2-}$ undergoes disproportionation reaction in acidic medium but $\mathrm{MnO}_{4}^{-}$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 : $\mathrm{Ni}(\mathrm{s})+2 \mathrm{Ag}^{+}(\mathrm{aq}) \rightarrow$ $\mathrm{Ni}^{2+}(\mathrm{aq})+2 \mathrm{Ag}(\mathrm{s})$ 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) $\mathrm{CuO}(\mathrm{s})+\mathrm{H}_{2}(\mathrm{~g}) \longrightarrow \mathrm{Cu}(\mathrm{s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$
(b) $\mathrm{Fe}_{2} \mathrm{O}_{3}+3 \mathrm{CO}(\mathrm{g}) \longrightarrow 2 \mathrm{Fe}(\mathrm{g})+3 \mathrm{CO}_{2}(\mathrm{~g})$
(c) $\mathrm{NH}_{3}(\mathrm{~g}) 5 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 4 \mathrm{NO}(\mathrm{g})+5 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$
(d) $\mathrm{BCl}_{3}(\mathrm{~g})+3 \mathrm{LiAlH}_{4} \longrightarrow \mathrm{~B}_{2} \mathrm{H}_{6}+\mathrm{LiCl}+\mathrm{AlCl}_{3}$
(e) $2 \mathrm{~K}+\mathrm{F}_{2} \longrightarrow 2 \mathrm{KF}$
[Hints:- CuO is oxidizing agent, $\mathrm{H}_{2}$ is acting as reducing agent because Cu (II) is changing to $\mathrm{Cu}(0)$ by gain of $\mathrm{e}^{-} \mathrm{H}_{2}$ is getting oxidised to $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$, its oxidations sate is changing from 0 to +1 , by loss of electrons.
(ii) It is redox reaction: $\mathrm{Fe}_{2} \mathrm{O}_{3}$ is getting reduced to fe. CO is getting oxidised to $\mathrm{CO}_{2}$.]
4. Using standard electrode : Predict if the reaction between as the following is feasible.
(i) $\mathrm{Fe}^{3+}(\mathrm{aq})$ and $\mathrm{I}^{-}(\mathrm{aq})$
(ii) $\mathrm{Ag}^{+}$and Cu
(iii) $\mathrm{Fe}^{3+}$ and $\mathrm{Br}^{-}$(aq)
(iv) Ag and $\mathrm{Fe}^{3+}$ (aq)
(iv) $\mathrm{Br}_{2}$ (aq) and $\mathrm{Fe}^{2+}$ (aq)

Hint:- $\mathrm{E}_{\mathrm{I}_{2} / I^{-}}=0.541 \mathrm{~V}, \mathrm{E}_{\mathrm{Cu}^{2+} / \mathrm{Cu}},=0.34 \mathrm{~V}, \mathrm{E}_{\mathrm{Br}_{2} / \mathrm{Br}^{-}}=1.09 \mathrm{~V}, \mathrm{E}_{\mathrm{Ag}^{+} / \mathrm{Ag}}^{\theta}=0.80 \mathrm{~V}$, $\mathrm{E}_{\mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}}^{\theta}=0.77 \mathrm{~V}$.
5. Draw the diagram for the galvanic cell which would have overall chemical reaction as

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

## 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. $6 \times 10^{-3}$ mole $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$ reacts completely with $9 \times 10^{-3}$ mole $\mathrm{X}^{\mathrm{n}+}$ to give $\mathrm{XO}_{3}^{-}$and $\mathrm{Cr}^{3+}$. Find the value of X .
Ans. $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}+\mathrm{X}^{\mathrm{n}+} \longrightarrow \stackrel{+5}{\mathrm{X}} \mathrm{O}_{3}^{-}+\mathrm{Cr}^{3+}$

$$
6 \times 10^{-3} \times 6=(5-\mathrm{n}) \times 9 \times 10^{-3} \longrightarrow \mathrm{n}=1
$$

2. For the redox reaction

$$
\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}+\mathrm{X} \mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{Y} \mathrm{SO}_{2} \longrightarrow \mathrm{~K}_{2} \mathrm{SO}_{4}+\mathrm{Cr}_{2}\left(\mathrm{SO}_{4}\right)_{3}+\mathrm{ZH}_{2} \mathrm{O}
$$

What is the sum of $x+y+z$

$$
\begin{gathered}
\text { Ans. } \mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}+\mathrm{H}_{2} \mathrm{SO}_{4}+3 \mathrm{SO}_{2} \longrightarrow \mathrm{~K}_{2} \mathrm{SO}_{4}+\mathrm{Cr}_{2}\left(\mathrm{SO}_{4}\right)_{3}+\mathrm{H}_{2} \mathrm{O} \\
\therefore \mathrm{x}=1 \quad \mathrm{y}=3 \quad \mathrm{z}=1 \quad \therefore \mathrm{x}+\mathrm{y}+\mathrm{z}=5
\end{gathered}
$$

3. An aqueous solution containing 1 M each of $\mathrm{Au}^{+3}, \mathrm{Cu}^{+2}, \mathrm{Ag}^{+}, \mathrm{Li}^{+}$is being electrolysed using inert electrodes the value of standard potentials are

$$
\mathrm{E}_{\mathrm{Ag}^{+} / \mathrm{Ag}^{-}}^{\theta}=0.80 \mathrm{~V}, \quad \mathrm{E}_{\mathrm{Cu}^{2}+\mathrm{Cu}}^{\theta}=0.34 \mathrm{~V}, \quad \mathrm{E}_{\mathrm{Au}^{3+} / \mathrm{Au}}^{\theta}=1.50 \mathrm{~V}, \quad \mathrm{E}_{\mathrm{Li}^{+} / \mathrm{Li}}^{\theta}=-3.03 \mathrm{~V}
$$

With increasing voltage, find the sequence of deposition of metals on the cathode.

Ans. Only $\mathrm{Au}^{3+}, \mathrm{Ag}^{+}$and $\mathrm{Cu}^{2+}$ will deposit at cathode.
$\mathrm{Li}^{+}$will not deposit at cathode be cause SRP of water is -0.8274 V
So after $\mathrm{Cu}^{2+} ; \mathrm{H}_{2}$ will evolve at cathode.
4. $\mathrm{E}^{\theta}$ for $\mathrm{Cl}_{2}(\mathrm{~g})+21^{-} \longrightarrow 2 \mathrm{Cl}^{-}$(aq.) is 1.36 V , then calculate.
$\mathrm{E}^{\theta}$ for $4 \mathrm{C1}^{-}$(aq.) $\longrightarrow 2 \mathrm{Cl}_{2}(\mathrm{~g})+4 \mathrm{e}^{-}$
Ans. $\quad E^{\theta}{\mathrm{Cl} / \mathrm{Cl}_{2}}=-1.36 \mathrm{E}^{\theta}$ is independent of amount of substance
5. Why salt bridge is made up of saturated solution of $\mathrm{KNO}_{3}$ in agar-agar.

Ans. Velocities of both $\mathrm{K}^{+}$and $\mathrm{NO}_{3}{ }^{-}$are nearly the same.

## UNIT TEST-I

Time Allowed: $1 \mathbf{h r}$
Maximum Marks : 20
General Instructions:
(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. Identify the oxidised and Reduced species in the following reaction [1]

$$
\mathrm{H}_{2} \mathrm{~S}+\mathrm{Cl}_{2} \longrightarrow 2 \mathrm{HCl}+\mathrm{S}
$$

(a) $\mathrm{H}_{2} \mathrm{~S}$
(b) $\mathrm{Cl}_{2}$
(c) Both $\mathrm{H}_{2}, \mathrm{Cl}_{2}$
(d) None of these
2. What is the oxidation state of Br in $\mathrm{BrO}_{3}^{-}$?
[1]
(a) +1
(b) +3
(c) +4
(d) +5
3. Classify the type of reaction in Redox Reaction form :

$$
\begin{equation*}
3 \mathrm{H}_{2} \mathrm{O}+\mathrm{P}_{4}+3 \mathrm{OH}^{-} \longrightarrow \mathrm{PH}_{3}+3 \mathrm{H}_{2} \mathrm{PO}_{2}^{-} \tag{1}
\end{equation*}
$$

4. What is a redox couple? Give one example.
5. Identify oxidant in reaction given below :

$$
\begin{equation*}
\mathrm{CuO}(\mathrm{~s})+\mathrm{H}_{2}(\mathrm{~g}) \longrightarrow \mathrm{Cu}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \tag{1}
\end{equation*}
$$

6. Assign oxidation number to the underlined elements
(a) $\mathrm{NaH}_{2} \underline{\mathrm{PO}}_{4}$
(b) $\mathrm{H}_{4} \mathrm{P}_{2} \mathrm{O}_{7}$
(c) $\mathrm{K}_{2} \mathrm{MnO}_{4}$
(d) $\mathrm{H}_{2} \mathrm{~S}_{2} \mathrm{O}_{7}$
7. Predict product of electrolysis in following case

- An aqueous solution of $\mathrm{CuCl}_{2}$ with platinum electrodes.

8. Consider the reaction $\mathrm{Zn}(\mathrm{s})+2 \mathrm{Ag}^{+}$(aq.) $\longrightarrow \mathrm{Zn}^{2+}$ (aq.) $+2 \mathrm{Ag}(\mathrm{s})$

Answer following :
(i) Which electrode is negatively charged ?
(ii) What are carrier of current in the cell ?
(iii) Individual reaction at each electrode.
9. $\mathrm{E}^{\theta}$ values are given : $\mathrm{K}^{+} / \mathrm{K}=-2.93 \mathrm{~V}, \mathrm{Ag}^{+} / \mathrm{Ag}=0.80 \mathrm{~V}$
$\mathrm{Hg}^{2+} / \mathrm{Hg}=0.79 \mathrm{~V} \mathrm{Mg}^{2+} / \mathrm{Mg}=-2.37 \mathrm{~V}, \mathrm{Cr}^{3+} / \mathrm{Cr}=-0.74 \mathrm{~V}$
(i) Which one is strong reducing agent ?
(ii) Which one is strong oxidising agent?
(iii) Which redox couple is a stronger reducing agent than $\mathrm{H}^{+} / \mathrm{H}_{2}$ ?
10. Balance the reaction (ion-electron or oxidation number)
$\mathrm{P}_{4}(\mathrm{~s})+\mathrm{OH}^{-}(\mathrm{aq}.) \longrightarrow \mathrm{PH}_{3}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{PO}_{2}^{-}$(aq.) [Basic medium]

## UNIT TEST-II

Time Allowed: 1 Hr. (REDOX REACTIONS) Maximum Marks : 20
General Instructions:
(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. The average oxidation No. of Iodine is $\mathrm{I}_{3}{ }^{-}$ion is.
2. What is oxidation state of Cr in $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$ ?
3. Write the name of cell in which chemical energy is converted into Electrical energy.
4. Why is anode negatively charged in an electrochemical cell?
5. Identify the oxidised and Reduced species in the following reaction

$$
\begin{equation*}
\mathrm{H}_{2} \mathrm{~S}+\mathrm{Cl}_{2} \longrightarrow 2 \mathrm{HCl}+\mathrm{S} \tag{1}
\end{equation*}
$$

6. A decomposition reaction may or may not be a redox Reaction. Write two decomposition reactions in support of the statement.
7. Write the functions of salt bridge in a electrochemical cell.
8. Account for the following :
(i) $\quad \mathrm{HNO}_{3}$ acts as oxidizing agent while $\mathrm{HNO}_{2}$ can act both as Reducing and oxidizing agent.
(ii) $\mathrm{AgF}_{2}$ is unstable compound and act as a strong oxidizing agent.
(iii) Ozone acts as an oxidizing agent.
9. Explain with suitable reasons :
(i) Reaction $\mathrm{FeSO}_{4}(\mathrm{aq})+\mathrm{Cu}(\mathrm{s}) \rightarrow \mathrm{CuSO}_{4}(\mathrm{aq})+\mathrm{Fe}$ does not occur.
(ii) Zinc can displace copper from aqueous $\mathrm{CuSO}_{4}$ solution but Ag cannot.
(iii) Solution of $\mathrm{AgNO}_{3}$ turns blue when copper rod is immersed in it.
10. (i) Give one example each of the following redox reactions:
(a) Combination Reaction
(b) Decomposition Reaction
(c) Metal displacement Reaction
(ii) Remaining two are Assertion and Reason. Read the statement carefully and choose the correct alternative :
(a) Both Assertion and Reason are true and Reason is the correct explanation of Assertion.
(b) Both Assertion and Reason are true but Reason is not the correct explanation of Assertion.
(c) Assertion is true but Reason is false.
(d) Both Assertion and Reason are false.
(I) Assertion : Redox couple is the combination of oxidised and Reduced from of a substance involved in an oxidation or reduction half cell.

Reason: In the Representation $\mathrm{E}^{\theta} \mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}$ and $\mathrm{E}^{\theta} \mathrm{Cu}^{2+} / \mathrm{Cu}, \mathrm{Fe}^{3+} /$ $\mathrm{Fe}^{2+}$ and $\mathrm{Cu}^{2+} / \mathrm{Cu}$ are Redox couples.
(II) Assertion : Oxidation no. of C in HCHO is zero.

Reason : Formaldehyde is covalent compound.


## Organic Chemistry :

 Some Basic Principles and Techniques
## 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.


- Acetic acid was synthesised by Kolbe and methane by Berthelot.
- Types of hybridisation of C-atom :

| Hybridisation | Structure | Bond angle | Examples |
| :---: | :---: | :---: | :---: |
| $\mathrm{sp}^{3}$ | Tetrahedral | $109^{\circ} 28^{\prime}$ | Ethane, Methane |
| $\mathrm{sp}^{2}$ | Trigonal | $120^{\circ}$ | Ethene, Propene |
| sp | Linear | $180^{\circ}$ | 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 ORGANIC 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 $1^{\circ}$ carbon atom.
(ii) Secondary carbon atom : When carbon atom is attached with two other carbon atoms, it is called secondary or $2^{\circ}$ carbon atom.
(iii) Tertiary carbon atom : When carbon atom is attached with three other carbon atoms, it is called tertiary or $3^{\circ}$ carbon atom.
(iv) Quaternary carbon atom : When carbon atom is attached with four other carbon atoms, it is called quarterly or $4^{\circ}$ carbon atom.

- Functional Group : The atom e.g., $-\mathrm{Cl},-\mathrm{Br}$, etc., or group of atoms e.g., $-\mathrm{COOH},-\mathrm{CHO}$, which is responsible for the chemical properties of the molecule, is called functional group.
- Homologous Series : The series in which the molecular formula of adjacent members differ by a $-\mathrm{CH}_{2}$ unit, is called homologous series and the individual members are called homologous, e.g., The homologous series of alkene group is

$$
\left.\begin{array}{l}
\mathrm{C}_{2} \mathrm{H}_{4} \\
\mathrm{C}_{3} \mathrm{H}_{6} \\
\mathrm{C}_{4} \mathrm{H}_{8} \\
\mathrm{C}_{5} \mathrm{H}_{10}
\end{array}\right\} \text { difference of }-\mathrm{CH}_{2} \text { unit or } 14 \text { unit mass }
$$

The general characteristics of this series are :

1. All the homologues contain same functional group. That's why their chemical properties are almost similar.
2. All the members of a series have same general formula, e.g.,

| Series | General Formula |
| :--- | :--- |
| Alkanes | $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}+2}$ |
| Alkenes | $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}}$ |
| Alkynes | $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}-2}$ |
| Alcohol and ether | $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}+2} \mathrm{O}$ |
| Aldehyde and ketone | $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}} \mathrm{O}$ |
| Acid and ester | $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}} \mathrm{O}_{2}$ |

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., $\mathrm{CH}_{3} \mathrm{CHCH}_{2} \mathrm{CH}_{3}$
।
Cl $\mathrm{CH}_{3} \mathrm{CH}(\mathrm{CI}) \mathrm{CH}_{2} \mathrm{CH}_{3}$
(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 |
| :---: | :---: | :---: | :---: |
| $\mathrm{C}_{1}$ | Meth- | $\mathrm{C}_{7}$ | Hept |
| $\mathrm{C}_{2}$ | Eth- | $\mathrm{C}_{8}$ | Oct |
| $\mathrm{C}_{3}$ | Prop- | $\mathrm{C}_{9}$ | Non |
| $\mathrm{C}_{4}$ | But- | $\mathrm{C}_{10}$ | Dec |
| $\mathrm{C}_{5}$ | Pent- | $\mathrm{C}_{11}$ | Undec |
| $\mathrm{C}_{6}$ | Hex- | $\mathrm{C}_{12}$ | Dodec |

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 :
$-\mathrm{COOH}>-\mathrm{SO}_{3} \mathrm{H}>$ acid anhydride $-\mathrm{COOR}>-\mathrm{COX}>-\mathrm{CONH}_{2}$
$>-\mathrm{CN}>-\mathrm{NC}>-\mathrm{CHO}>\mathrm{C}=\mathrm{O}>-\mathrm{OH}>-\mathrm{SH}$
$>-\mathrm{NH}_{2}>=>\equiv>$ substituents
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.

Secondary suffixes are tabulated below :

| No. | Class | Formula | Prefix | Suffix |
| :--- | :--- | :--- | :--- | :--- |
| 1. | Acid halides | O <br> $\\|$ <br> $\mathrm{C}-\mathrm{X}$ | halocarbonyl | - oyl halide <br> -carbonyl halide |
| 2. | Alcohols | -OH | hydroxy | -ol |
| 3. | Aldehydes | -CHO | formyl | -al <br> - carbaldehyde |
| 4. | Ketones | $\rangle \mathrm{C}=\mathrm{O}$ | oxo (keto) | - one |
| 5. | Amides | $-\mathrm{CONH}_{2}$ | carbamoyl | - amide |
| 6. | Amine | $-\mathrm{NH}_{2}$ | amino | - amine |
| 7. | Carboxylic <br> acid | $-\mathrm{COOH}_{2}$ | carboxy | -carboxylic <br> acid |
| 8. | Ester | -COOR | alkoxy <br> carbonyl | -alkyl <br> alkanoate |
| 9. | Nitriles | $-\mathrm{CN}^{\text {conano }}$ | - nitrile |  |
| 10. | Sulphonic acid | $-\mathrm{SO}_{2}-\mathrm{OH}$ | sulpho | - sulphonic acid |

- Nomenclature of substituted benzene compounds :


Methylbenzene


Methoxy benzene


Nitrobenzene


Bremobenzene


3, 4-Dimethyl phenol

1, 2-Dibromo benzene In Trivial name
1, 2-ortho (0)
1, 3-meta (m)
1, 4-para (p)

2-Chloro-1-methyl 1-4-nitrobenzene


## 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)


Cis.But-2-ene


Trans.But-2-ene

## - 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.


$$
\mathrm{Cl}_{2} \xrightarrow{\text { Sunlight }} 2 \mathrm{Cl}^{*}
$$

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.
$\mathrm{A} \frown \mathrm{B} \xrightarrow[\text { electrophile }]{\text { more electronegative }} \mathrm{A}^{+}+\underset{\text { nucleophile }}{\mathrm{B}^{-}}$


Heterolytic fission generally takes place in polar covalent molecules but in non-polar molecules, it takes place in the presence of catalyst like $\mathrm{AiCl}_{3}$ (anhy.), $\mathrm{FeCl}_{3}$ (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., $\mathrm{Cl}^{+}, \mathrm{NO}_{2}, \mathrm{CH}_{3} \mathrm{CO}^{+}$etc.
$\mathrm{BF}_{3}, \mathrm{ZnCl}_{2}$ (anhydrous), $\mathrm{FeCl}_{3}$ (anhydrous), $\mathrm{AlCl}_{3}$ (anhydrous)
(ii) Nucleophiles or Nucleophilic Reagents

These are negatively charged or neutral molecules with unshared electron pair.
e.g., $\mathrm{O} \overline{\mathrm{H}}, \mathrm{CN}^{-}, \mathrm{R}^{\circ} \mathrm{NH}_{2}, \stackrel{\circ}{\mathrm{~N}} \mathrm{H}_{3}$

## - Reaction Intermediates :

(i) Free radicals : These are the product of homolysis and contain an odd electron. These are highly reactive planar species with $\mathrm{sp}^{2}$ hybridisation.

Their order of stability is

$$
\begin{aligned}
& \left(\mathrm{C}_{6} \mathrm{H}_{5}\right)_{3} \stackrel{\circ}{\mathrm{C}}>\left(\mathrm{C}_{6} \mathrm{H}_{5}\right)_{2} \stackrel{\circ}{\mathrm{C}} \mathrm{H}>\mathrm{C}_{6} \mathrm{H}_{5} \stackrel{\circ}{\mathrm{C}} \mathrm{H}_{2} \\
& \quad>\mathrm{CH}_{2}=\mathrm{CH}-\stackrel{\circ}{\mathrm{C}} \mathrm{H}_{2}>3^{\circ}>2^{\circ}>1^{\circ}>\mathrm{CH}_{2}=\stackrel{\circ}{\mathrm{C}} \mathrm{H}
\end{aligned}
$$

(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., $\mathrm{sp}^{2}$ hybridised with an empty p-orbital.


Stability order of carbocation is

$$
\begin{aligned}
& \left(\mathrm{C}_{6} \mathrm{H}_{5}\right)_{3} \stackrel{+}{\mathrm{C}}>\left(\mathrm{C}_{6} \mathrm{H}_{5}\right)_{2} \stackrel{+}{\mathrm{C}} \mathrm{H}>\mathrm{C}_{6} \mathrm{H}_{5} \stackrel{+}{\mathrm{C}} \mathrm{H}_{2} \\
& \quad>\mathrm{CH}_{2}=\mathrm{CH}-\stackrel{+}{\mathrm{C}} \mathrm{H}_{2}>3^{\circ}>2^{\circ}>1^{\circ}>\mathrm{CH}_{2}=\stackrel{+}{\mathrm{C}} \mathrm{H}
\end{aligned}
$$

(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 $\mathrm{sp}^{3}$ hybridised carbon (having one lone pair) order of stability of carbanions is

$$
\begin{aligned}
\left(\mathrm{C}_{6} \mathrm{H}_{5}\right)_{3} \overline{\mathrm{C}} & >\left(\mathrm{C}_{6} \mathrm{H}_{5}\right)_{2} \overline{\mathrm{C}} \mathrm{H}>\mathrm{C}_{6} \mathrm{H}_{5} \overline{\mathrm{C}} \mathrm{H}_{2} \\
>\overline{\mathrm{C}} \mathrm{H}_{3} & >1^{\circ}>2^{\circ}>3^{\circ} \text { carbanions }
\end{aligned}
$$

## - 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., ${\stackrel{+}{\mathrm{C}} \mathrm{H}_{3}} \rightarrow-\stackrel{-}{\mathrm{C}} \mathrm{l}$

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 $(+l)$.


Cl has -1 effect and alkyl group has +I effect.
Order of groups producing -I effect is

$$
\begin{aligned}
\mathrm{R}_{3} \mathrm{~N} & >-\mathrm{NO}_{2}>-\mathrm{CN}>-\mathrm{SO}_{3} \mathrm{H}>-\mathrm{CHO}>-\mathrm{CO}>-\mathrm{COOH}>-\mathrm{F} \\
& >-\mathrm{Cl}>-\mathrm{Br}>-\mathrm{I}>-\mathrm{OH}>-\mathrm{OR}>-\mathrm{NH}_{2}>-\mathrm{C}_{6} \mathrm{H}_{5}>-\mathrm{H}
\end{aligned}
$$

Order of groups producing +1 effect is
$3^{\circ}$ alkyl group $>2^{\circ}$ alkyl group $>1^{\circ}$ alkyl group $>-\mathrm{CH}_{3}>-\mathrm{H}$

## - 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 $\pi$ 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 $\sigma$ electron of a $\mathrm{C}-\mathrm{H}$ bond of an alkyl group attached directly to an atom of unsaturated system or to an atom with an unshared p-orbital.


This effect is also called no bond resonance or Baker Nathan effect.

## Applications of Hyper-conjugation

Stability of alkenes: More the number of $\alpha$-hydrogen atoms, more stable is the alkene.


Stability of Carbocation : Greater the number of alkyl groups attached to positively charged carbon atom, the greater is the stability.
e.g., $\left(\mathrm{CH}_{3}\right)_{2} \stackrel{+}{\mathrm{C}} \mathrm{H}>\mathrm{CH}_{3}-\stackrel{+}{\mathrm{C}} \mathrm{H}>\stackrel{+}{\mathrm{C}} \mathrm{H}_{3}$
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 $\pi$ 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, $-\mathrm{OH},-\mathrm{OR},-\mathrm{NH}_{2}, \mathrm{NHCOR}$, 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, $-\mathrm{COOH},-\mathrm{COOR}, \mathrm{CHO},-\mathrm{CN},-\mathrm{NO}_{2}$, etc.


## - 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 348 K (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). <br> - Camphor, naphtalene, anthracence, benzoic Acid, etc. are purified. |
| Distillation | It is used to separate <br> - Volatile liquids from non-volatile impurities. <br> - Liquids having sufficient difference in their boiling points. | - Hexane (b.p. 342 K ) and toluene (b.p. 384 K ) <br> - Chloroform (b.p. 334K) and aniline (b.p. 457 K ) |
| - 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. |
| $\begin{array}{\|l\|} \hline- \text { Steam } \\ \text { Distillation } \end{array}$ | This method is used to separate substances which are (i) steam volatile, (ii) immiscible with water, (iii) posses a vapour pressure of $10-15 \mathrm{~mm} \mathrm{Hg}$ and (iv) contain nonvolatile impurities. | - Aniline is separated from aniline water mixture. <br> - 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 <br> components of a mixture through a <br> stationary phase under the influence of a <br> mobile phase. | - Widely used for separation <br> purification, identification <br> and characterization of the <br> components of a mixture, <br> whether coloured or <br> colourless. |
| :--- | :--- | :--- |
| - Adsorption <br> Chromatography | Differential adsorption of the various <br> components of a mixture on a suitable <br> adsorbent such as silica get or alumina. |  |
| - Column <br> Chromatography | The mixture is passed through adsorbent <br> packed in glass tube. | - Mixture of naphthalene and <br> benzophenone. |
| - Thin Layer <br> Chromatography | The mixture is passed over adsorbent on <br> a thin glass plate. | Amino acids can be detected <br> by spraying the plate with <br> ninhydrin solution. |
| - Partition <br> Chromatography | Differential partitioning of components of <br> a mixture between stationary and mobile <br> phases. | A special quality paper known as <br> chromatography paper is used. It contains <br> water trapped in it, which acts as the <br> stationary phase. |
| - Paper <br> Chromatography |  |  |


| 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 | $2 \mathrm{CuO}+\mathrm{C} \xrightarrow{\Delta} 2 \mathrm{Cu}+\mathrm{CO}_{2}$ | $\mathrm{CO}_{2}$ gas turns lime water milky. | $\underset{\text { Lime water }}{\mathrm{CO}_{2}+\underset{\text { Milkiness }}{\mathrm{Ca}(\mathrm{OH})_{2}} \underset{\mathrm{CaCO}_{3} \downarrow}{ } \downarrow+\mathrm{H}_{2} \mathrm{O}}$ |
| Hydrogen | $\mathrm{CuO}+2 \mathrm{H} \xrightarrow{\Delta} \mathrm{Cu}+\mathrm{H}_{2} \mathrm{O}$ | Water droplets appear on the cooler part of the ignition tube and also turns anhydrous $\mathrm{CuSO}_{4}$ blue. | $\underset{\text { White }}{\mathrm{CuSO}_{4}}+5 \mathrm{H}_{2} \mathrm{O} \longrightarrow \underset{\text { Blue }}{\mathrm{CuSO}_{4} \cdot 5 \mathrm{H}_{2} \mathrm{O}}$ |
| Nitrogen | Lassaigne's extract (L.E.) <br> $\mathrm{Na}+\mathrm{C}+\mathrm{N} \xrightarrow{\Delta} \mathrm{NaCN}$ <br> (L.E.) | L.E. $+\mathrm{FeSO}_{4}+\mathrm{NaOH}$, boil and cool $+\mathrm{FeCl}_{3}+$ conc. HCl. Gives blue or green colour. | $\begin{gathered} \mathrm{FeSO}_{4}+2 \mathrm{NaOH} \longrightarrow \mathrm{Fe}(\mathrm{OH})_{2}+\mathrm{Na}_{2} \mathrm{SO}_{4} \\ \mathrm{Fe}(\mathrm{OH})_{2}+6 \mathrm{NaCN} \longrightarrow \\ \mathrm{Na}_{4}\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]+2 \mathrm{NaOH} \\ 3 \mathrm{Na}_{4}\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]+4 \mathrm{FeCl}_{3} \longrightarrow \\ \mathrm{Fe}_{4}\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]_{3}+12 \mathrm{NaCl} \\ \text { Prussian blue } \end{gathered}$ |


| Sulphur | $2 \mathrm{Na}+\mathrm{S} \xrightarrow{\Delta} \underset{\text { (L.E.) }}{\mathrm{Na}_{2} \mathrm{~S}}$ | $\begin{gathered} - \text { L.E. }+ \text { sodium } \\ \text { nitroprusside } \\ \text { A deep violet colour. } \\ - \text { L.E. }+\mathrm{CH}_{3} \mathrm{COOH}+ \\ \left(\mathrm{CH}_{3} \mathrm{COO}\right)_{2} \mathrm{~Pb} \\ \text { Gives a black ppt. } \end{gathered}$ | $\left\lvert\, \begin{gathered} \mathrm{Na}_{2} \mathrm{~S}+\underset{\text { Sodium nitroprusside }}{\mathrm{Na}_{2}\left[\mathrm{Fe}(\mathrm{CN})_{5} \mathrm{NO}\right] \longrightarrow} \\ \mathrm{Na}_{4}\left[\mathrm{Fe}(\mathrm{CN})_{5} \mathrm{NOS}\right] \\ \text { Deep violet } \\ \mathrm{Na}_{2} \mathrm{~S}+\underset{\left(\mathrm{CH}_{3} \mathrm{COO}\right)_{2} \mathrm{~Pb} \xrightarrow{\mathrm{CH}_{3} \mathrm{COOH}}}{\mathrm{Pbs} \downarrow+2 \mathrm{CH}_{3} \mathrm{COONa}} \\ \text { Black ppt. } \end{gathered}\right.$ |
| :---: | :---: | :---: | :---: |
| Halogens | $\begin{equation*} \mathrm{Na}+\mathrm{X} \xrightarrow{\Delta} \mathrm{NaX} \tag{L.E.} \end{equation*}$ | L.E. $+\mathrm{HNO}_{3}+\mathrm{AgNO}_{3}$ <br> - White ppt. soluble in aq. $\mathrm{NH}_{3}\left(\right.$ or $\left.\mathrm{NH}_{4} \mathrm{OH}\right)$ confirms Cl . <br> - Yellow ppt. partially soluble in aq. $\mathrm{NH}_{3}\left(\right.$ or $\left.\mathrm{NH}_{4} \mathrm{OH}\right)$ confirms Br. <br> - Yellow ppt. insoluble in aq. $\mathrm{NH}_{3}$ (or $\left.\mathrm{NH}_{4} \mathrm{OH}\right)$ confirms I. |  |
| Nitrogen <br> and <br> sulphur <br> together | $\mathrm{Na}+\mathrm{C}+\mathrm{N}+\mathrm{S} \xrightarrow{\Delta}$ <br> NaSCN <br> Sodium thiocyanate (L.E.) | As in test for nitrogen; instead of green or blue colour, blood red colouration confirms presence of N and S both. | $\begin{gathered} \mathrm{NaSCN}+\mathrm{FeCl}_{3} \longrightarrow \\ {\left[\mathrm{Fe}(\mathrm{SCN}) \mathrm{Cl}_{2}+\mathrm{NaCl}\right.} \\ \text { Blood red colour } \end{gathered}$ |
| Phosphorus | $\mathrm{P} \xrightarrow{\mathrm{Na}_{2} \mathrm{O}_{2}, \text { boil }} \mathrm{Na}_{3} \mathrm{PO}_{4}$ | Solution is boiled with nitric acid and then treated with ammonium molybdate $\left(\mathrm{NH}_{4}\right){ }_{2} \mathrm{MoO}_{4}$. Formation of yellow ppt. indicates presence of phosphate (hence, phosphorus) in organic compound. |  |

- 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 | Method |
| :---: | :---: |
| Carbon and Hydrogen | Liebig's Combustion method : <br> A known mass of an organic compound is burnt in the presence of excess of $\mathrm{O}_{2}$ and CuO . $\mathrm{C}_{\mathrm{x}} \mathrm{H}_{\mathrm{y}}+\left(\mathrm{x}+\frac{\mathrm{y}}{4}\right) \mathrm{O}_{2} \xrightarrow{\Delta} \mathrm{xCO}_{2}+\frac{\mathrm{y}}{2} \mathrm{H}_{2} \mathrm{O}$ <br> $\mathrm{CO}_{2}$ evolved is absorbed by conc. solution of KOH or ascarite $(\mathrm{NaOH}+$ $\mathrm{CaO}) \cdot \mathrm{H}_{2} \mathrm{O}$ produced is absorbed by anhydrous $\mathrm{CaCl}_{2}$ or $\mathrm{Mg}\left(\mathrm{ClO}_{4}\right)_{2}$. <br> Increase in masses of these absorbing compounds gives the masses of $\mathrm{CO}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ produced. $\begin{aligned} & \% \text { of } \mathrm{C}=\frac{12}{44} \times \frac{\text { mass of } \mathrm{CO}_{2} \text { formed }}{\text { mass of compound taken }} 100 \\ & \% \text { of } \mathrm{H}=\frac{2}{18} \times \frac{\text { mass of } \mathrm{H}_{2} \mathrm{O} \text { formed }}{\text { mass of compound taken }} 100 \end{aligned}$ |
| Halogens | Carius method : <br> Halogen in organic compound is precipitated as silver halide by boiling with conc. $\mathrm{NHO}_{3}$ and then adding $\mathrm{AgNO}_{3}$. $\left\{\begin{array}{l} \mathrm{X} \frac{\mathrm{HNO}_{3}, \Delta}{\mathrm{AgNO}_{3}} \mathrm{AgX} \downarrow \\ \% \text { of } \mathrm{X}=\frac{\text { At. mass of } \mathrm{X}}{108+\text { At. mass of } \mathrm{X}} \times \frac{\text { mass of } \mathrm{AgX} \text { formed }}{\text { mass of compound taken }} \times 100 \end{array}\right.$ |
| Nitrogen | Dumas method : <br> Nitrogen containing organic compound is heated with CuO in an atmosphere of $\mathrm{CO}_{2}$. $\mathrm{C}_{\mathrm{x}} \mathrm{H}_{\mathrm{y}} \mathrm{~N}_{\mathrm{z}}+\left(2 \mathrm{x}+\frac{\mathrm{y}}{2}\right) \mathrm{CO}_{2} \longrightarrow \mathrm{xCO}_{2}+\frac{\mathrm{y}}{2} \mathrm{H}_{2} \mathrm{O}+\mathrm{N}_{2}+\left(2 \mathrm{x}+\frac{\mathrm{y}}{2}\right) \mathrm{Cu}$ <br> $\mathrm{N}_{2}$ evolved gets collected over conc. KOH solution which absorbs all other gases. $\% \text { of } \mathrm{N}=\frac{28}{22400} \times \frac{\text { Vol. of } \mathrm{N}_{2} \text { at STP }}{\text { mass of compound taken }} \times 100$ <br> Kjeldahl's method : <br> Organic compound $+\mathrm{H}_{2} \mathrm{SO}_{4 \text { (conc.) }} \longrightarrow\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4} \xrightarrow{2 \mathrm{NaOH}} \mathrm{Na}_{2} \mathrm{SO}_{4}+$ $2 \mathrm{NH}_{3}+2 \mathrm{H}_{2} \mathrm{O}+2 \mathrm{NH}_{3}+\mathrm{H}_{2} \mathrm{SO}_{4} \longrightarrow\left(\mathrm{Na}_{4}\right)_{2} \mathrm{SO}_{4}$ $\% \text { of } \mathrm{N}=\frac{1.4 \times \text { molarity of acid } \times \text { vol. of acid used } \times \text { basicity of acid }}{\text { mass of compound taken }}$ |


| Sulphur | Carius method : <br> Sulphur in organic compound is converted into $\mathrm{H}_{2} \mathrm{SO}_{4}$ by boiling with $\mathrm{Na}_{2} \mathrm{O}_{2}$ or conc. $\mathrm{HNO}_{3}$ and is precipitated as $\mathrm{BaSO}_{4}$ by adding excess of $\mathrm{BaCl}_{2}$ solution in water. $\left\{\begin{array}{l} \mathrm{S} \xrightarrow[\text { (ii) } \mathrm{BaCl}_{2}]{\text { (i) } \mathrm{HNO}_{3}, \Delta} \xrightarrow{\mathrm{BaSO}_{4} \downarrow} \\ \text { white ppt. } \end{array}\right.$ |
| :---: | :---: |
| Phosphorus | Ignition method : $\left[\begin{array}{l} \mathrm{P} \xrightarrow[\text { heat }]{\mathrm{HNO}_{3}} \mathrm{H}_{3} \mathrm{PO}_{4} \\ \mathrm{H}_{3} \mathrm{PO}_{4}+\mathrm{Mg}^{2+}+\mathrm{NH}_{4} \mathrm{CL} \xrightarrow{\Delta} \xrightarrow[\begin{array}{c} \text { Magnesium ammonium } \\ \text { phosphate (white ppt.) } \end{array}]{\mathrm{MgNH}_{4} \mathrm{PO}_{4}+\mathrm{HCl}} \\ 2 \mathrm{MgNH}_{4} \mathrm{PO}_{4} \xrightarrow[\text { Magnesium pyrophosphate }]{\Delta} \mathrm{Mg}_{2} \mathrm{P}_{2} \mathrm{O}_{7}+2 \mathrm{NH}_{3}+\mathrm{H}_{2} \mathrm{O} \end{array}\right.$ |

## MIND MAP : ORGANIC CHEMISTRY



## CASE BASED STUDY - QUESTIONS

## PASSAGE -I

The resonance effect is defined as the polarity produced in the molecule by the interaction of two $\pi$ bonds as between a $\pi$ bond and lone pair of electrons present an on adjacent atom. The effect is transmitted through the chain. In positive resonance effect the transfer of electrons is away from on atom or subsitituent group attached to the conjugated system. The electron displacement makes certain positions in the molecule of high electron densities. In negative resonance effect the transfer of electrons is towards the atom or substituent group attached to the conjugated system.

1. Draw resonance structure of carboxylate ion
2. What do you understand by +R and -R effect.
3. Write resonance structures of $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CHO}$ indicate relative stability of the contributing structures.
4. Draw the resonating structures of (a) phenol (b) Benzoic acid
5. Out of the resonating structures of $\mathrm{CO}_{2}$ which structures is not correct for $\mathrm{CO}_{2}$ ?

$$
\begin{align*}
& : \ddot{\mathrm{O}}=\mathrm{C}=\ddot{\mathrm{O}}: \quad \stackrel{\ominus}{:} \stackrel{.}{\mathrm{O}}-\mathrm{C} \equiv \stackrel{+}{\mathrm{O}}: \\
& \text { (ii) }  \tag{iv}\\
& \text { (iii) } \tag{i}
\end{align*}
$$

## PASSAGE -II

The Lassaigne's extract is usually alkaline because excess of sodium reacts with water to form sodium hydroxide. If not it may be made alkaline by the addition of a few drops of a dilute solution of sodium hydroxide. To a part of the extract a small amount of a freshly prepared ferrous sulphate solution is added and the content are warmed. A few drops of ferric chloride solution are then added to the contents and the resulting solution is acidified with dilute hydrochloric acid the appearance of a bluish green colour due to the formation of ferric ferrocyanide confirm the presence of nitrogen in the organic compound.

1. In sodium fusion test of organic compound the nitrogen in an organic compound is converted into which compound.
2. In the Lassaigne's test for the detection of sulphure the purple colour is due to the for of which complex.
3. What is the name of the Prussian blue colour compound formed in Lassaigne's test for nitrogen in an organic compound.
4. Why do we boil Lassaigne's extract with conc. $\mathrm{HNO}_{3}$ while detecting halogens in an organic compound.
5. Explain the reason for the fusion of an organic compound with metallic sodium for testing nitrogen, sulphure and halogens.

## MULTIPLE CHOICE QUESTIONS (MCQ)

1. Homolytic fission of $\mathrm{C}-\mathrm{C}$ bond in ethane gives an intermediate in which carbon is:
a. $\mathrm{sp}^{3}$ hybridised
b. $\mathrm{sp}^{2}$ hybridised
c. sp-hybridised
d. $\mathrm{sp}^{3} \mathrm{~d}$ - 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?
a. $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{CH}_{2}$
b. $\mathrm{CH}_{3}-\stackrel{\oplus}{\mathrm{C}} \mathrm{H}-\mathrm{CH}_{3}$
c. $\mathrm{CH}_{3}-\mathrm{C}=\mathrm{CH}$
d. $\mathrm{CH}_{2}=\stackrel{\oplus}{\mathrm{C}} \mathrm{H}-\mathrm{CH}_{2}$
4. Which of the following can act as an electrophile?
a. $\mathrm{CN}^{-}$
b. $\mathrm{OH}^{-}$
c. $\mathrm{H}_{2} \mathrm{O}$
d. $\mathrm{BF}_{3}$
5. Which of the following is correct about the species: $\left(\mathrm{CH}_{3}\right)_{3}-\mathrm{C}^{+}$
a. It is planar
b. Its $\mathrm{C}+$ is $\mathrm{sp}^{2}$ hybridised
c. A nucleophile can attack on its $\mathrm{C}^{+}$
d. All of these
6. Which of the following has all the effects namely Inductive, Mesomeric and Hyperconjugative?
a. $\mathrm{CH}_{3} \mathrm{Cl}$
b. $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}$
c. $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CHCOCH}_{2} \mathrm{Cl}$
d. $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CH}=\mathrm{CH}_{2}$
7. The most stable free radical among the following is:
a. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2} \stackrel{\bullet}{\mathrm{C}} \mathrm{H}_{2}$
b. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CHCH}_{3}$
c. $\mathrm{CH}_{3} \mathrm{CH}_{2}$
d. $\mathrm{CH}_{3} \stackrel{\bullet}{\mathrm{CHCH}} 3$
8. Isomers of a compound must have :
a. Same physical properties
b. Same chemical properties
c. Same structural properties
d. Same molecular weight
9. Most stable carbanion among the following is
a.

b.

c.

d.

10. Which of the following species have six $\pi$ conjugated electrons?
a.

b. $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{2}{ }^{-}$
c.

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. $-\mathrm{COOH},-\mathrm{SO}_{3} \mathrm{H},-\mathrm{CONH}_{2},-\mathrm{CHO}$
b. $-\mathrm{SO}_{3} \mathrm{H},-\mathrm{COOH},-\mathrm{CONH}_{2},-\mathrm{CHO}$
c. $-\mathrm{CHO},-\mathrm{COOH},-\mathrm{SO}_{3} \mathrm{H},-\mathrm{CONH}_{2}$
d. $-\mathrm{CONH}_{2},-\mathrm{CHO},-\mathrm{SO}_{3} \mathrm{H},-\mathrm{COOH}$
12. The IUPAC name of $\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}-\mathrm{C} \equiv \mathrm{CH}$ 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. The IUPAC name of the following compound is

a. 4-Bromo-3- cyanophenol
b. 2-Bromo-5-hydroxy benzonitrile
c. 2-cyano-4-hydroxybromo benzene
d. 6-Bromo-3-hydroxy benzonitrile
14. The order of the stability of the following of carbocations is :

$$
\begin{array}{ll}
: \ddot{\mathrm{O}}=\mathrm{C}=\ddot{\mathrm{O}}: & \stackrel{\ominus}{:} \ddot{\mathrm{O}}-\mathrm{C} \equiv \stackrel{+}{\mathrm{O}}: \\
{ }^{\oplus} \ddot{\mathrm{O}}-\mathrm{C} \equiv \ddot{\mathrm{O}} & \ddot{\cdot}
\end{array} \quad \stackrel{+}{\mathrm{O}} \equiv \mathrm{C}-\ddot{\mathrm{O}^{\ominus}} \stackrel{\ominus}{:} .
$$

a. III $>$ I $>$ II
b. III $>$ II $>$ I
c. $\mathrm{II}>\mathrm{III}>\mathrm{I}$
d. I $>$ II $>$ III
15. Quantitative measurement of nitrogen in an organic compound is done by the method:
a. Berthelot method
b. Lassaigne method
c. Carius method
d. Kjehldahl method

## ANSWERS

| 1.a | 2.b | 3.d | 4.d | 5.d | 6.c | 7.b | 8.d | 9.d | $10 . \mathrm{d}$ |
| :--- | :---: | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 11.a | 12.a | 13.b | 14.a | 15.d |  |  |  |  |  |

## FILL IN THE BLANKS

1. A triple bond between two carbon atoms is composed of one $\qquad$ and
$\qquad$ bonds.
2. An organic compound which decomposes below its boiling point can be purified by $\qquad$ .
3. Electrophiles are the species which attack the regions of $\qquad$ electron density.
4. Hyperconjugation effect is also known as $\qquad$ resonance.
5. In Duma's method, the nitrogen present in an organic compound is set free as $\qquad$ .

## ANSWERS

1. $\sigma$, two $\pi \quad$ 2. Vacuum distillation 3. high 4. No-bond 5. Nitrogen

TRUE AND FALSE TYPE QUESTIONS

1. Ethanol and methoxymethane are position isomers.
2. A free radical is a species with an unpaired valence electron.
3. Inductive effect is observed in $\pi$ bond in presence of attacking reagent.
4. The percentage of carbon and hydrogen are estimated simultaneously in an organic compound by Liebig method.
5. Chromatography is the method used to separate and purify compounds when present in small amounts.

## ANSWERS

1. F 2. T 3.F $\quad$ 4. T $\quad$ 5. 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: Duma's method is more applicable to nitrogen containing organic compounds than the Kjeldahl's method.
Reason: Kjeldahl's method does not give satisfactory result in which nitrogen atom is directly attached to oxygen atom.
3. Assertion: Alkanes having more than three carbon atoms exhibit chain isomerism.
Reason: All carbon atoms in alkanes are sp-hybridised.
4. Assertion: In $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{CH}_{2}$, all the carbon atoms are $\mathrm{sp}^{2}$ hybridised.

Reason: All the hydrogen atoms lie in one plane.
5. Assertion: Butane and 2-Methylbutane are homologues.

Reason: Butane is a straight chain alkane while 2-Methylbutane is branched chain alkane.
6. 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
8. Assertion: Carbocations are planar in nature.

Reason: Carbocations are $\mathrm{sp}^{2}$ Hybridised.
9. Assertion: IUPAC name of compound $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}-\mathrm{CHO}$ is But-2-enal.

Reason: Functional group gets preference over multiple in IUPAC name of a compound.
10. Assertion: Glycerol is purified by distillation under reduce pressure. Reason: Organic compounds in liquid state are purified by distillation.
ANSWERS


## 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
a. Leibig method
b. Dumas method
c. Kjehldahl method
d. Carius method

Column II
i. $\mathrm{N}_{2}$
ii. $\operatorname{AgX}$
iii. $\mathrm{CO}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$
iv. $\mathrm{NH}_{3}$

## Column II

e) Aniline
f) Halogens
g) Schiff's Nitrometer
h) $\mathrm{CaCl}_{2}$ tube
2.

Column I
a. Nonbenzenoid aromatic compound
b. Catenation
c. Free radical
d. sp-hybridised carbon atom

## Column II

i. $50 \%$ s character
ii. Species containing single unpaired nonbonding electrons
iii. Chain-forming property of an element
iv. Tropolone

ANSWERS

1. a. iiii, h
b. i, g
c. iv, e
d. ii, f
2. 

a. iv
b. iii
c. ii
d. i

## ONE WORD ANSWER TYPE QUESTIONS

1 Name one common adsorbent in column chromatography.
2. Mention the hybridisation of underlined carbon in $\mathrm{CH}_{3} \underline{\mathrm{C}} \equiv \mathrm{N}$.
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. $\mathrm{SO}_{3}$ 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. Which has the longest $\mathrm{C}-\mathrm{C}$ bond length among ethane, ethene and ethyne.
3. How many secondary carbon atoms are present in 2-Methylpentane?
4. Draw structure of 3-Isopropyl-2-methylhexane.
5. Draw bond line structure of $\mathrm{CH}_{3}\left(\mathrm{CH}_{2}\right)_{6} \mathrm{CH}=\mathrm{CH}\left(\mathrm{CH}_{2}\right)_{2}-\mathrm{COOH}$
6. What are the bond angles in $\mathrm{sp}^{3}, \mathrm{sp}^{2}$ and sp hybrid orbitals?.
7. Write the correct of priority of the following functional groups:

$$
-\mathrm{C} \equiv \mathrm{~N},>\mathrm{C}=\mathrm{O},-\mathrm{OH},-\mathrm{COOH},-\mathrm{CONH}_{2}
$$

8. Write IUPAC name of :
(i) $\mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{CN}$
(ii) $\mathrm{CH}_{2}=\mathrm{CHCH}_{2} \mathrm{OH}$
(iii) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}\left(\mathrm{CH}_{3}\right)-\mathrm{CO}-\mathrm{CH}_{2} \mathrm{CH}_{3}$
(iv) $\mathrm{CH}_{3} \mathrm{CH}_{2}-\mathrm{O}-\mathrm{CH}_{2} \mathrm{CH}\left(\mathrm{CH}_{3}\right) \mathrm{CH}_{3}$
(v) $\mathrm{Cl}-\mathrm{CH}_{2}-\mathrm{C} \equiv \mathrm{CH}$
9. What type of isomerism is exhibited by Propanal and Propanone?
10. Classify the following into electrophiles and nucleophiles:
$\mathrm{H}+, \mathrm{NH}_{3}, \mathrm{AlCl}_{3}, \mathrm{NO}_{2}^{+}, \mathrm{CN}^{-}, \mathrm{H}_{2} \mathrm{O}, \mathrm{ROH}, \mathrm{RNH}_{2}$, Carbocation
11. What type of attacking reagents are produced by heterolytic cleavage of covalent bond?
12. Name each of the following species and indicate which member of each pair is more stable:
(i) $\mathrm{CH}_{3}{ }^{+}, \mathrm{CH}_{3} \mathrm{CH}_{2}{ }^{+}$
(ii) $\mathrm{C}_{6} \mathrm{H}_{5} \stackrel{+}{\mathrm{C}} \mathrm{HCH}_{3}, \mathrm{CH}_{3} \mathrm{CHCH}_{3}$
(iii) $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CH}_{2}, \mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{3}$
(iv) $\mathrm{CH}_{3}-\stackrel{-}{\mathrm{C}} \mathrm{H}_{2}, \mathrm{CH}_{3}-\overline{\mathrm{C}} \mathrm{H}-\mathrm{CH}_{3}$
13. Identify electrophilic centre in $\mathrm{CH}_{3} \mathrm{CHO}$.
14. What is state oh hybridization of positively charged carbon atom in carbocation?
15. What is the effect of introducing an alkyl group on the stability of carbocation?
16. Out of Benzyl and ethyl carbocation which is more stable and why?
17. Arrange the following in increasing order of acidic strength:
$\mathrm{ClCH}_{2} \mathrm{COOH}, \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{COOH}, \mathrm{ClCH}_{2} \mathrm{CH}_{2} \mathrm{COOH}$
18. Name two solvents which are commonly used to dissolve organic solids.
19. Name the technique that can be used for purification of iodine that contains traces of NaCl .
20. A liquid ( 10 mL ) has three components $\mathrm{A}, \mathrm{B}, \mathrm{C}$. which technique is suitable to sparate $\mathrm{A}, \mathrm{B}, \mathrm{C}$ from such a small amount of mixture?
21. Under what condition do we use fractional distillation?
22. A liquid compound starts decomposing well before its boiling point under normal pressure. How will you purify it?
23. For which type of compounds Kjehldahl's method is not useful?
24. How do you precipitate sulphur in Carius method?
25. Which method is used to estimate carbon and hydrogen?
26. 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. Give IUPAC name of the following compounds:
(i) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{OH}$
(ii) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CHO}$
(iii) $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{C} \equiv \mathrm{N}$
(iv) $\mathrm{CH}_{3}-\mathrm{CH}-\mathrm{C}-\mathrm{CH}-\mathrm{CH}_{3}$
(v) $\mathrm{CH}_{3}-\mathrm{CH}-\mathrm{CH}=\underset{\text { । }}{\mathrm{C}-\mathrm{CH}_{2}-\mathrm{COOH}}$
4. Indicate sigma ( $\sigma$ ) and pi $(\mathrm{p})$ bonds in the following molecules
(i) $\mathrm{CH}_{2} \mathrm{Cl}_{2}$
(ii) $\mathrm{HCONH} . \mathrm{CH}_{3}$
5. Write bond line formulas for: (i) isopropyl alcohol (ii) heptane-4-one
6. Which is suspected to more stable: $\mathrm{O}_{2} \mathrm{NCH}_{2} \mathrm{CH}_{2} \mathrm{O}^{-}$or $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{O}^{-}$and why?
7. What is the effect of introducing an alkyl group on the stability of a free radical?
8. Give two examples each of the groups exerting -I and +I effect when attached to a chain of carbon atoms.
9. A tertiary butyl carbocation is is more stable than isobutyl carbocation. Justify.
10. All electrophiles are Lewis acids while nucleophile are Lewis bases. Explain.
11. What is the purpose of filtration through hot water funnel?
12. What precautions are necessary while purifying an organic solid with the help of crystallization process?
13. Discuss the principle of steam distillation.
14. Discuss the role of fractionating column in fractional distillation.
15. How will you prepare Lassaigne's extract? Name the elements which can be detected from this extract?

## 3-MARKS QUESTIONS

1. Why stability of carbocations follows the order: tertiary>secondary>primary?
2. What are the various conditions essential for resonance?
3. Write resonance structures of $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CHO}$. Indicate relative stability of the contributing structures.
4. Inductive effect is of permanent nature while electromeric effect is only temporary. Explain.
5. What is chromatography? Name different types of chromatographic processes.
6. You are given a mixture of methanol and acetone. Discuss the process which you will employ to separate them.

## 5-MARK QUESTIONS

1. Classify the following reactions in one of the reaction type studies in this unit:
(i) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br} \longrightarrow \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{SH}+\mathrm{Br}^{-}$
(ii) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}=\mathrm{CH}_{2}+\mathrm{HCl} \longrightarrow\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}(\mathrm{Cl}) \mathrm{CH}_{3}$
(iii) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CCH}_{2} \mathrm{OH}+\mathrm{HBr} \longrightarrow\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CBrCH}_{2} \mathrm{CH}_{3}$
(iv) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br}+\mathrm{HO}^{-} \longrightarrow \mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{H}_{2} \mathrm{O}+\mathrm{Br}^{-}$
(d) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Cl}+\mathrm{KOH} \longrightarrow \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}+\mathrm{KCl}$
aq
2. (a) An organic compound contains $69 \%$ carbon and $4.8 \%$ hydrogen the remainder being oxygen. Calculate the masses of carbon dioxide and water produce when 0.20 g of this compound is subjected to complete combustion.
(b) 0.3780 of an organic compound gave 0.574 g of silver chloride in carious estimation. Calculate the percentage of chlorine in the compound.
3. Arrange the following in the order of property indicated against each set:
(i) $-\mathrm{C}_{6} \mathrm{H}_{5},-\mathrm{NO}_{2},-\mathrm{COOH},-\mathrm{I},-\mathrm{F},-\mathrm{CH}_{3},-\mathrm{C}_{2} \mathrm{H}_{5}$ ( In the increasing order of -I effect)
$+$
(ii) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CH}_{2}^{+},\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}^{+}, \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CHCH}_{3}$ (In the order of increasing stability)
(iii) $-\mathrm{Cl},-\mathrm{CONH}_{2},-\mathrm{CHO}$ (In the increasing priority order if present in same molecule)
4. Draw the resonance structures for the following compounds. Show the electron shift using curved arrow notation.
(i) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{NO}_{2}$
(ii) $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CHCHO}$
(iii) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{OH}$
(iv) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2}^{+}$
(v) $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CHCH}_{2}^{+}$
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. 0.378 g of an organic compound containing carbon and hydrogen was subjected to combustion by Leibig's method, the $\mathrm{CO}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ formed were passed through potash bulbs and anhydrous $\mathrm{CaCl}_{2}$ tube. At the end of the experiment, the increase in the respective weights were 0.264 g and 0.162 g . Calculate the percentage of carbon and hydrogen.
(Ans: $\mathrm{C}=19.05 \%, \mathrm{H}=4.76 \%$ )

## UNIT TEST-I

Time Allowed: $1 \mathbf{H r}$.
Maximum Marks : 20

## General Instructions:

(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. Write bond line formula for the following compound:
$\mathrm{HOCH}_{2} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CH}\left(\mathrm{CH}_{3}\right) \mathrm{CH}\left(\mathrm{CH}_{3}\right) \mathrm{COOH}$
2. Write IUPAC name of the following compound:

3. The central atom of compound $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{CH}_{2}$ is $\qquad$ hybridized. [1]

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 but 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.
4. Assertion: Carbocations are planar in nature.

Reason : Carbocations are $\mathrm{sp}^{2}$ Hybridized.
5. Assertion : All the carbon atoms of But-2-ene lie in are plane.

Reason : All the carbon atoms in But-2-ene are $\mathrm{sp}^{2}$ hybridized.
6. (i) What type of isomerism is exhibited by the following pair of compounds?
$\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CHO}$ and $\mathrm{CH}_{3}-\mathrm{C}-\mathrm{CH}_{3}$
$\|$
O
(ii) Give one example each of nucleophile and electrophile.
7. (i) Arrange the following in increasing order of stability:

(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:
(i) $-\mathrm{C}_{6} \mathrm{H}_{5},-\mathrm{NO}_{2},-\mathrm{COOH},-\mathrm{I},-\mathrm{F},-\mathrm{CH}_{3},-\mathrm{C}_{2} \mathrm{H}_{5}$
(In the increasing order of $-\mathbf{I}$ effect)
(ii) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CH}_{2}^{+},\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}^{+}, \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CHCH}_{3}$
(In the order of increasing stability)
(iii) $-\mathrm{COOH},-\mathrm{CONH}_{2},-\mathrm{CHO}$
(In the increasing priority order if present in same molecule)
(iv) $\mathrm{HCOOH}, \mathrm{CH}_{3} \mathrm{COOH}, \mathrm{ClCH}_{2} \mathrm{COOH}$
(Increasing order of acidic strength)
(v) $\mathrm{O}_{2} \mathrm{NCH}_{2} \mathrm{CH}_{2} \mathrm{O}^{-}, \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{O}^{-}$
(species having greater stability)

## UNIT TEST-II

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. Which of the following can act as an electropliile?
(a) $\mathrm{CN}^{-}$
(b) $\mathrm{OH}^{-}$
(c) $\mathrm{H}_{2} \mathrm{O}$
(d) $\mathrm{BF}_{3}$
2. The most stable free radical among the following is :
(a) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2} \mathrm{CH}_{2}$
(b) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CHCH}_{3}$
(c) $\mathrm{CH}_{3} \mathrm{CH}_{2}$
(d) $\mathrm{CH}_{3} \mathrm{CH} \mathrm{CH}_{3}$
3. What is the other name for no band resonance?

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 but 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.
4. Assertion : In $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{CH}_{2}$ all the carbon atoms are $\mathrm{sp}^{2}$ hybridized. [1] Reason: All the hydrogen atoms lie in one plane.
5. Assertion : Glycerol is purified by distillation under reduce pressure.

Reason : Organic compound in liquid state are purified by distillation.
6. Give IUPAC name of the following compounds :
(a) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{OH}$
(b) $\mathrm{CH}_{3}-\mathrm{CH}-\mathrm{C}-\mathrm{CH}-\mathrm{CH}_{3}$

7. Discuss the principle of steam distillation.
8. Explain the reason for the fusion of an organic compound with metallic sodium for testing nitrogen sulphur and halogens.
9. What are the various conditions essential for resonance?
10. Suggest a method to separate the constituents from the following mixture :
(a) Mixture of two miscible liquids.
(b) A mixture of oil and water.
(c) A mixture of plant pigments.


## Chapter-9

## 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 :
$\mathrm{R}-\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{H}_{2} \xrightarrow[\Delta]{\mathrm{Ni}} \mathrm{R}-\mathrm{CH}_{2}-\mathrm{CH}_{3}$
[Where R is H or Alkyl group]


2. From alkyl halides : on reduction with Zinc and dilute hydrochloric acid


Chloromethane Methane
$\mathrm{C}_{2} \mathrm{H}_{5}-\mathrm{Cl}+\mathrm{H}_{2} \xrightarrow{\mathrm{Zn}, \mathrm{H}^{+}} \mathrm{C}_{2} \mathrm{H}_{6}+\mathrm{HCl}$
Chloroethane
Ethane


3. From alkyl halides by Wurtz reaction : Reaction of alkyl halide with sodium in dry ether, useful only for the preparation of symmetrical alkanes.

 1-Bromopropane

4. From Carboxylic acids : By decarboxylation with soda lime

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.
$2 \mathrm{CH}_{3} \mathrm{COO} \mathrm{Na}+2 \mathrm{H}_{2} \mathrm{O} \xrightarrow{\text { Electrolysis }} \mathrm{CH}_{3}-\mathrm{CH}_{3}+2 \mathrm{CO}_{2}+\mathrm{H}_{2}+2 \mathrm{NaOH}$ At Anode: (Oxidation)

$2 \mathrm{CH}_{3}{ }^{-} \longrightarrow \mathrm{CH}_{3}-\mathrm{CH}_{3}$
At Cathode : (Reduction)
$2 \mathrm{H}_{2} \mathrm{O} \longrightarrow 2 \mathrm{OH}^{-}+2 \mathrm{H}^{+}$
$2 \mathrm{H}^{+}+2 \mathrm{e}^{-} \longrightarrow \mathrm{H}_{2}$

## 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.

## Chemical properties of Alkanes :

- Alkanes undergo substitution reactions.
e.g., Halogenation, Nitration, Sulphonation.

Halogenation : For example Chlorination of methane


Rate of reaction of alkanes with halogens is $\mathrm{F}_{2}>\mathrm{Cl}_{2}>\mathrm{Br}_{2}>\mathrm{I}_{2}$. 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 $\mathrm{HNO}_{3}$.

Mechanism of halogenation : Free radical mechanism

| $\mathrm{Cl}-\mathrm{Cl} \xrightarrow{h \mathrm{u}} \mathrm{Cl} \bullet+\cdot \cdot \mathrm{Cl}$ | Initiation |
| :--- | :--- |
| $\mathrm{Cl} \cdot+\mathrm{CH}_{4} \longrightarrow \mathrm{H}-\mathrm{Cl}+\cdot \mathrm{CH}_{3}$ | Propagation |
| $\cdot \mathrm{CH}_{3}+\mathrm{Cl}-\mathrm{Cl} \longrightarrow \mathrm{CH}_{3} \mathrm{Cl}+\cdot \mathrm{Cl}$ | Propagation |
| $\mathrm{Cl} \cdot+\cdot \mathrm{CH}_{3} \longrightarrow \mathrm{CH}_{3} \mathrm{Cl}$ | Termination |

Combustion : Complete combustion gives carbon dioxide and water.

$$
\mathrm{CH}_{4}+2 \mathrm{O}_{2} \longrightarrow \mathrm{CO}_{2}+2 \mathrm{H}_{2} \mathrm{O}
$$

## Isomerisation :



## Aromatisation :



$$
\mathrm{Cr}_{2} \mathrm{O}_{3} \text { or } \mathrm{V}_{2} \mathrm{O}_{5}
$$

 $10-20 \mathrm{~atm}$

(vi) Reaction with steam

$$
\mathrm{CH}_{4}+\mathrm{H}_{2} \mathrm{O} \xrightarrow[\Delta]{\mathrm{Ni}} \mathrm{CO}+3 \mathrm{H}_{2}
$$

Pyrolysis : Decomposition of higher alkanes to lower alkanes on heating.

$$
\mathrm{C}_{6} \mathrm{H}_{14} \xrightarrow{773 \mathrm{~K}} \xrightarrow{\longrightarrow} \mathrm{C}_{6} \mathrm{H}_{12}+\mathrm{H}_{2}
$$

## Conformations :

Different spatial arrangement of atoms arising due to rotation around $\mathrm{C}-\mathrm{C}$ single bond.

## Conformation of ethane, $\mathrm{CH}_{3} \mathrm{CH}_{3}$

Two conformational isomers or conformers.
Eclipsed form = all hydrogen atoms nearest to each other.
Staggered form $=$ all hydrogen atoms are farthest apart.


Eclipsed
Stability of eclipsed conformation is least while staggered conformation is most stable. The energy difference between two extreme forms is $12.5 \mathrm{kJmol}^{-1}$. Due to this small energy difference the two forms are easily inter-convertible at ordinary temperature and cannot be separated and isolated.

## ALKENES

These are unsaturated non-cyclic hydrocarbons which have.sp ${ }^{2}$-hybridisation with $120^{\circ}$ bond angle.
Alkenes are also called olefins [oil.forming] which indicates their high reactive nature.

Alkenes have general formula $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}}$, where $\mathrm{n}=2,3,4, \ldots \ldots$.
$\mathrm{C}_{2} \mathrm{H}_{4}$ (ethene), $\mathrm{C}_{3} \mathrm{H}_{6}$ (propene), etc.

## - Methods of Preparation of Alkenes

## (i) From alkynes


cis-alkene
(ii) From alkyl halide [Dehydrohalogenational]

(iii) From vicinal dihalides

$$
\begin{aligned}
& \mathrm{CH}_{2} \mathrm{Br}-\mathrm{CH}_{2} \mathrm{Br}+\mathrm{Zn} \xrightarrow{\Delta} \mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{ZnBr}_{2} \\
& \mathrm{CH}_{3}-\mathrm{CH}-\mathrm{CH}_{2}+\mathrm{Zn} \xrightarrow{\Delta} \mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{ZnBr}_{2}
\end{aligned}
$$

$\mathrm{Br} \quad \mathrm{Br}$
(iv) From alcohols by acidic dehydrogenation


## - Chemical Properties of Alkenes :

## 1. Addition of Halogens :



> 1, 2-Dibromoethane


1, 2-Dichloropropane
2. Addition of hydrogen halides $\mathbf{H C l}, \mathbf{H B r}, \mathbf{H I}:$ Add up to alkenes to form alkyl halides as per their reactivity order in $\mathrm{HI}>\mathrm{HBr}>\mathrm{HCl}$.
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.


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 $\mathrm{HF}, \mathrm{HCl}$ and HI .


## 3. Addition of sulphuric acid


4. Addition of water

5. Oxidation : Alkenes decolourise cold dilute aqueous solution potassium permanganate (Baeyer's reagent). It is used as a test for unsaturation.


Ethane-1,2-diol or glycol

Acidic $\mathrm{KMnO}_{4}$ or acidic $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$ oxidises alkenes to ketones and/or acids depending upon the nature of alkene and the experimental conditions.


2-Methyl propene
Propane-2-one


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).


## 7. Polymerization




## ALKYNES

These are unsaturated hydrocarbons with general formula $\mathrm{C}_{\mathrm{n}} \mathrm{H}_{2 \mathrm{n}-2}$ e.g., $\mathrm{C}_{2} \mathrm{H}_{2}$ (ethyne), $\mathrm{C}_{3} \mathrm{H}_{4}$ (propyne).

Alkynes also exhibit electrophilic addition reaction but less reactive than alkenes because the dissociation of $\pi$-electron cloud requires more energy.
$\mathrm{H}-\mathrm{C} \equiv \mathrm{C}-\mathrm{H}$ contins $3 \sigma$ and $2 \pi$-bonds and bond length is 120 pm . In acetylene. $\mathrm{H}-\mathrm{C}-\mathrm{C}$ bond angle is $180^{\circ}$.

## - Methods of Preparation of Alkynes

## 1. From calcium carbide

$$
\begin{aligned}
& \mathrm{CaCO}_{3} \longrightarrow \mathrm{CaO}+\mathrm{CO}_{2} \\
& \mathrm{CaO}+\mathrm{C} \longrightarrow \mathrm{CaC}_{2}+\mathrm{CO} \\
& \mathrm{CaC}_{2}+2 \mathrm{H}_{2} \mathrm{O} \longrightarrow \mathrm{Ca}(\mathrm{OH})_{2}+\mathrm{C}_{2} \mathrm{H}_{2}
\end{aligned}
$$

## 2. From vicinal dihalides



## 3. From tetrahalides

$$
\mathrm{Br}_{2} \mathrm{CH}-\mathrm{CHBr}_{2}+2 \mathrm{Zn} \xrightarrow[\Delta]{\mathrm{CH}_{3} \mathrm{OH}} \mathrm{H}-\mathrm{C} \equiv \mathrm{C}-\mathrm{H}+2 \mathrm{ZnBr}_{2}
$$

## Physical Properties of Alkynes:

1. The first two members are gases next eight members $\left(\mathrm{C}_{5}-\mathrm{C}_{12}\right)$ 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 $\mathrm{PH}_{3}$ and $\mathrm{H}_{2} \mathrm{~S}$ 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.

## - Chemical properties of Alkynes

Alkynes show electrophilic as well as nucleophilic addition reactions.

## (i) Acidic character of alkyne

$$
\begin{aligned}
& \mathrm{HC} \equiv \mathrm{CH}+\mathrm{Na} \longrightarrow \begin{array}{c}
\mathrm{HC} \equiv \overline{\mathrm{C}} \mathrm{C} \mathrm{Na}+\frac{1}{2} \mathrm{H}_{2} \\
\text { monotyodium } \\
\text { acetlides }
\end{array} \\
& \mathrm{H}-\mathrm{C} \equiv \mathrm{CNa}^{+}+\mathrm{Na} \longrightarrow \overline{\mathrm{Na}}{ }^{+} \mathrm{C} \equiv \overline{\mathrm{C}} \mathrm{Na}^{+}+\frac{1}{2} \mathrm{H}_{2} \\
& \mathrm{CH}_{3}-\mathrm{C} \equiv \mathrm{C}-\mathrm{H}+\mathrm{NaNH}_{2} \longrightarrow \longrightarrow \begin{array}{c}
\text { sodium acetylide propynide }
\end{array}
\end{aligned}
$$

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)

(ii)


## (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.


## Addition of halogens



Addition of hydrogen halides


## Addition for water



## (iii) Cyclic polymerisation of ethyne



## 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 $\mathrm{sp}^{2}$ hybridised. Planar ring, bond angle $120^{\circ}$. All C-C bond lengths are equal due to complete delocalisation of $\pi$ electrons.


Formation of C-C and C-H sigma bonds


Sigma skeleton of benzene molecule


## 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 $(4 n+2 \pi)$ electrons ( n is $0,1,2,3,4$ )
- For $\mathrm{n}=1: 4 \mathrm{n}+2=6$; benzene is stable and electrons are delocalized.


## Benzene



Three double bonds;
six $\pi$ 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.
(i) Nitration

(ii) Halogenation

(iii) Sulphonation

(iv) Friedel-Craft's alkylation reaction

(v) Friedel-Crafts acylation reaction


Benzene also undergoes addition reactions e.g.
(i)

(ii)


benzene bexachloride or 666
(BHC or Gammexane or lindane)

## - COMBUSTION

$2 \mathrm{C}_{6} \mathrm{H}_{6}+15 \mathrm{O}_{2} \longrightarrow 12 \mathrm{CO}_{2}+6 \mathrm{H}_{2} \mathrm{O}$

Directing influence of substituents in monosubstituted benzene
(i) Ortho and para directing groups : Ring activating groups e.g., $\mathrm{NH}_{2},-\mathrm{CH}_{3},-\mathrm{C}_{2} \mathrm{H}_{5},-\mathrm{OCH}_{3}$ etc. $(+\mathrm{R}$ effect $)$
(ii) Meta directing groups : Ring deactivating groups
e.g. $-\mathrm{NO}_{2},-\mathrm{CN},-\mathrm{CHO},-\mathrm{COOH},-\mathrm{SO}_{3} \mathrm{H}(-\mathrm{R}$ effect).

## MIND MAP : HYDROCARBON



## CASE BASED STUDY - QUESTIONS

## PASSAGE-I

Presence of double bond in an alkene is tested by reacting with either water or with dilute solution of Bayer's regent. The position of double bond is located with the help of ozonalysis. It is done by joining together the product of ozonalysis which are carbonyl compounds at their carbonyl carbon atoms by double bond. One mole of ozone is used in the ozonalysis reaction per mole of double bond in a particular alkene.

1. An alkene upon ozonalysis will give $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CHO}$ and $\mathrm{CH}_{3} \mathrm{COCH}_{3}$. Give IUPAC name and structure of the given alkene.
2. Complete the given chemical equation.

$$
\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{3}+\mathrm{O}_{3} \xrightarrow[\mathrm{H}_{2} \mathrm{O}]{\mathrm{Zn}} \mathrm{~A}
$$

3. On ozonolysis one molecule of a hydrocarbon produces two molecules of ethanal and one molecule of ethanedial. Give the structure on ozonolysis of hydrocarbon.
4. An alkene give 5-keto-2-methyl hexanal. Give IUPAC name of the given alkene.
5. (a) $\mathrm{A} \xrightarrow{\mathrm{O}_{3}} \mathrm{CH}_{3} \mathrm{CHO}$ ?
(b) $\mathrm{A} \xrightarrow[\mathrm{Zn} / \mathrm{H}_{2} \mathrm{O}]{\mathrm{O}_{3}} \mathrm{CH}_{3} \mathrm{CHO}+\mathrm{CH}_{3}-\stackrel{\text { - }}{\mathrm{C}}-\mathrm{CH}_{2}-\mathrm{CH}_{3}$

## PASSAGE-II

Benzene responds to electrophilic substitution reactions. However some reagents are needed in these reactions in order to generate the attaching electrohiles. The directive influence in disubstitution taking place in the ring depends upon the nature of the group already present. An activating group in creases the electron density at the ortho and para positions. The deactivating group decreases the same at these positions thereby healing the metal position comparatively a point of high electron density.

1. Complete the following chemical equation

2. Explain why $\left(-\mathrm{NO}_{2}\right)$ group is m -directing \& reactivating group.
3. Arrange the following compound in increase order of reactivity towards electrophilic substitution reaction




4. Write the product in the following reaction

5. Which of the following is most reactive to electrophilic attack.

(a)

(b)

(c)

(d)

## MULTIPLE CHOICE QUESTIONS (MCQ)

1. Which of the following has zero 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. Which of th following is not the product of dehydration of


(a)

(b)

(c)

(d)
5. In the hydrocarbon :


The state of hybridization of carbon 1,3 and 5 are in the order
(a) $\mathrm{sp}, \mathrm{sp}^{2}, \mathrm{sp}^{3}$
(b) $\mathrm{sp}^{3}, \mathrm{sp}^{2}, \mathrm{sp}$
(c) $\mathrm{sp}^{2}, \mathrm{sp}, \mathrm{sp}^{3}$
(d) $\mathrm{sp}, \mathrm{sp}^{3}, \mathrm{sp}^{2}$
6. Action of acetylene on dilute $\mathrm{H}_{2} \mathrm{SO}_{4}$ gives:
(a) Acetic acid
(b) Acetone
(c) Acetaldehyde
(d) Ethyl alcohol
7. Which of the following compounds will exhibit cis-trans (geometrical) isomerism?
(a) Butanol
(b) 2-Butyne
(c) 2-Butenol
(d) 2-Butene
8. Basic strength of :
$\mathrm{H}_{3} \mathrm{C} \mathrm{CH}, \mathrm{H}_{2} \mathrm{C}=\mathrm{CH}$ and $\mathrm{HC} \equiv \mathrm{C}$ is in the order of
(a) I $>$ III $>$ II
(b) I $>$ II $>$ III
(c) II $>$ I $>$ III
(d) III $>$ II $>$ I
9. Reaction of hydrogen bromide with propene in the absence of a peroxide is $\mathrm{a} / \mathrm{an}$
(a) free radical reaction
(b) nucleophilic substitution
(c) electrophilic addition
(d) nucleophilic substitution
10. Among the following compounds, the one which is most reactive towards electrophilic nitration is:
(a) Benzoic acid
(b) Nitrobenzene
(c) Tofuene
(d) Benzene

ANSWERS: 1.b 2.c $3 . \mathrm{d}$ 4.a $5 . \mathrm{d}$ 6.c $\quad 7 . \mathrm{d}$ 8.c $\quad 9 . \mathrm{c} \quad$ 10.c

## FILL IN THE BLANKS

1. The addition of HBr to an unsymmetrical alkene takes place in accordance with $\qquad$ rule.
2. Benzene favours $\qquad$ substitution reaction.
3. The Dipole moment of Benzene is $\qquad$ -
4. The nitro group in the benzene nucleus is $\qquad$ directing. It $\qquad$ the reactivity of the benzene ring.
5. Melting point and boiling point increase as the molar masses $\qquad$ .
ANSWERS : 1. Markownikov's
6. electrophilic
7. Zero
8. Meta, decreases
9. Increase

## 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 $>\mathrm{n}$.
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 $(4 n+2) \pi$ 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, $\mathrm{AlCl}_{3}$ is an electrophile.
10. An electron-donating substituent in benzene ring gives a meta product.
ANSWERS: 1. T
11. F
12. F
13. T
14. F
15. T 7.T
16. F
17. F
18. 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. Column I

a. $\mathrm{CH}_{4}$
b. $\mathrm{CH}_{2}=\mathrm{CH}_{2}$
c. $\mathrm{CH} \equiv \mathrm{CH}$
d. $\mathrm{PCl}_{5}$

Column II
i. $\mathrm{sp}^{2}$
ii. $\mathrm{sp}^{3}$
iii. $\quad s p^{3} d$
iv. sp

## Column III

e. Ozonalysis
f. Oxidising agent
g. Saturated nature
h. Un Saturated nature

Column I
a. Alkanes
b. Alkenes
c. Alkynes
d. Arenes

Column II
i. Saturated nature
ii. Ozonolysis
iii. Geometrical isomerism
iv. Aromatic character

ANSWERS:

1. a. $\rightarrow$ ii. -g .
b. $\rightarrow$ i.-h.-e.
c. $\rightarrow$ i.-h.-e.
d. $\rightarrow$ iii.-f.
2. a. $\rightarrow \mathrm{i}$.
b. $\rightarrow$ 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.

1. Assertion: The IUPAC name of $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}-\mathrm{C} \equiv \mathrm{CH}$ is pent-3-en1 -yne and not pent-2-en-4-yne.
Reason: While deciding the locants of double and triple Bonds, lowest sum rule is always followed.
2. 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 $\mathrm{AlCl}_{3}$ 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: Cyclopentadienyl anion is aromatic in nature.

Reason: Cyclopentaclienyl anion has six $\pi$-electrons.
6. 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 $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{CH}_{2}$ lie in one plane. Reason: All the carbon atoms in it are $\mathrm{sp}^{2}$ 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.
9. Assertion: Benzene does not decolourise bromine water.

Reason: Benzene is stabilised by resonance due to delocalisation of $\pi$ electrons.
10. Assertion: Acidity of $\mathrm{C}-\mathrm{H}$ bond decreases in the order:
$\mathrm{HC} \equiv \mathrm{CH}>\mathrm{H}_{2} \mathrm{C}=\mathrm{CH}_{2}>\mathrm{H}_{3} \mathrm{C}-\mathrm{CH}_{3}$
Reason: Greater the percentage s-character, more is the acidity of $\mathrm{C}-\mathrm{H}$ bond.
ANSWERS: 1.a $2 . \mathrm{c}$ 3.c $\begin{aligned} & \text { 4.c } \\ & \text { 5.a }\end{aligned}$ 6.b $\quad$ 7.c $\quad$ 8.d $\quad 9 . \mathrm{a} \quad$ 10.a

## ONE WORD TYPE QUESTIONS

1. What is the state of hybridisation of Carbon atoms in alkanes?
2. What is the number of bonds in But-3-en-1-yne?
3. Name the product formed when Propyne is treated with aqueous $\mathrm{H}_{2} \mathrm{SO}_{4}$ in the presence of dil. $\mathrm{HgSO}_{4}$.
4. Name the product formed when Benzene reacts with $\mathrm{CH}_{3} \mathrm{Cl}$ in the presence of anhydrous aluminium chloride.
5. -COOH is ortho, para directing or Meta directing group?

## 1-MARK QUESTIONS

1. Write the reaction involved in Kolbe's electrolytic method to prepare ethane.
2. Define term decarboxylation.
3. Why dry ether and not water is used as a solvent in the preparation of alkane by Wurtz reaction?
4. Sodium salt of which carboxylic acid will be needed for the preparation of propane by decarboxylation method?
5. Complete the following reaction:
$\mathrm{CH}_{3} \mathrm{Cl}+\mathrm{Na} \xrightarrow{\text { dry ether }}$
6. Amongst the following which one has the maximum boiling point? $n$-Pentane, iso-pentane, neo-pentane.
7. Write IUPAC name of $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CHCH}_{2} \mathrm{CH}=\mathrm{CCH}_{2} \mathrm{CH}=\mathrm{CH}_{2}$ $\mathrm{CH}_{2} \mathrm{CH}_{3}$
8. Draw the cis and trans isomers of $\mathrm{CHCl}=\mathrm{CHCl}$.
9. What happens when 2-Bromobutane is being treated with alc. KOH ?
10. Name the reagents used to carry out the following conversions:

$$
\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2} \rightarrow \mathrm{CH}_{3} \mathrm{CH}(\mathrm{OH}) \mathrm{CH}_{2} \mathrm{OH}
$$

11. Complete the following reaction :

12. An alkene A on ozonolysis gives a mixture of ethanol and pentan-3-one. Write IUPAC name of element.
13. When alkyne is treated with bromine water then what will be the colour of the product?
14. Why alkynes do not exhibit geometrical isomerism?
15. Complete the following reaction:
(i) $\mathrm{CH}_{3} \mathrm{C} \equiv \mathrm{CH} \xrightarrow{\mathrm{H}_{2} \mathrm{O}, \mathrm{Hg}^{2+} / \mathrm{H}^{+}}$?
(ii) $\mathrm{CaC}_{2}+2 \mathrm{H}_{2} \mathrm{O} \longrightarrow \longrightarrow+$ $\qquad$
16. How will you convert ethyne to benzene?
17. Write IUPAC name of $\mathrm{C}_{6} \mathrm{H}_{5}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}=\mathrm{CH}_{2}$.
18. Why is benzene extraordinarily stable although it contains three double bonds?
19. Write chemical reaction to exemplify Friedel-Crafts alkylation of benzene.
20. Why is not aromatic?
21. $\mathrm{C}-\mathrm{C}$ bond length in benzene is intermediate between $\mathrm{C}-\mathrm{C}$ and $\mathrm{C}=\mathrm{C}$. Why?
22. Starting from benzene, how would you synthesize $m$ - Bromonitrobenzene.
23. Give one example each of o , p -directing group and m -directing group.
24. Complete the reaction:

$$
\mathrm{C}_{6} \mathrm{H}_{6}+\mathrm{CH}_{3} \mathrm{COCl} \xrightarrow{\text { Anhy. } \mathrm{AlCl}_{3}}
$$

## 2-MARKS QUESTIONS

1. What effect does branching have on the boiling point of an alkane and why?
2. An alkene A Canteins three $\mathrm{C}-\mathrm{C}$ eight $\mathrm{C}-\mathrm{H}$ and one $\mathrm{C}-\mathrm{C}(\pi)$ bond upon ozonlysis A gives two moles of an aldehyde of molar mass 44 u . Write the IUPAC name of A.
3. Write the structures of all the alkenes which upon hydrogenation give 2-methyl butane.
4. Wurtz reaction cannot be used for the preparation of unsymmetrical alkanes? Give reason.
5. How can ethene be prepared from (i) Ethanol (ii) Ethyl bromide?
6. Melting point of cis-But-2-ene is lower than that of trans-But-2-ene. Give reason.
7. Draw the structures of cis and trans Hex-2-ene.
8. Explain with the help of equation : Ozonolysis of propene.
9. What do you understand by peroxide effect (Kharasch effect)?
10. What factor determines the stability of alkene?
11. Out of benzene, $m$-dinitro benzene and toluene, which will undergo nitration most easily and why.
12. Complete the reaction:

(ii)

13. An alkene on treatment with HBr in presence of peroxide can generate two types of free radicals $\mathrm{CH}_{3}-\mathrm{C} \cdot-\mathrm{CH}_{2}-\mathrm{Br}$ and $\mathrm{CH}_{3}-\mathrm{C}_{\mid}\left(\mathrm{CH}_{3}\right)-\mathrm{CH}_{2} \cdot$ $\mathrm{CH}_{3} \quad \mathrm{Br}$

Predict the final product of the reaction and give reason.
(Hint: Stability of free radicals)
14. What happens when But-2-ene reacts with acidified potassium permanganate solution?
15. You are provided with But-2-yne, how will you convert it into:
(i) cis-But-2-ene
(ii) trans-But-2-ene
16. An alkene $\mathrm{C}_{4} \mathrm{H}_{8}$ reacts with HBr both in the presence and absence of peroxide to give the same product. Identify the alkene.
17. Arrange ethane, ethene and ethyne in the order of increasing acidity.
18. Identify A and B in the following reaction:

19. Write the structures of the products A and B of the following reactions:

(ii) $\mathrm{BrH}_{2} \mathrm{C}-\mathrm{CH}_{2} \mathrm{Br} \xrightarrow{\text { Alc. } \mathrm{KOH}} \mathrm{A} \xrightarrow{\mathrm{NaNH}_{2}} \mathrm{~B}$

## 3-MARKS QUESTIONS

1. Write the structures and name of products obtained in the reaction of sodium 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) $\mathrm{HCl} \&$ (ii) $\mathrm{H}_{2} \mathrm{O}$.
3. Complete the following reactions:
(ii) $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}+\mathrm{O}_{3} \xrightarrow{\mathrm{Zn} / \mathrm{H}_{2} \mathrm{O}}$
(iii) $\mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{H}_{2} \mathrm{O}+[\mathrm{O}] \xrightarrow{\text { Dil. } \mathrm{KMnO}_{4}}$
4. (i) Write the structure of 3, 4-Dimethylhept-3-ene.
(ii) Name the compounds obtained by ozonolysis of 3-Methylpent-2-ene.
5. Complete the following reactions:
(i) $\mathrm{CH} \equiv \mathrm{CH} \xrightarrow{\mathrm{NaNH}_{2}, \mathrm{CH}_{3} \mathrm{Br}}$ ?
(ii) $\mathrm{CH} \equiv \mathrm{CH} \xrightarrow{\mathrm{H}_{2} \mathrm{O}, \mathrm{HgSO}_{4} / \mathrm{H}_{2} \mathrm{SO}_{4}}$ ?
(iii) $\mathrm{CH}_{3} \mathrm{C} \equiv \mathrm{CH}+\mathrm{H}_{2} \xrightarrow{\mathrm{Pt}}$ ? $\xrightarrow{\mathrm{H}_{2}}$ ?
6. Write the mechanism of nitration of benzene.
7. Arrange in the order of decreasing relative reactivity with an electrophile and explain:
Toluene, p-Nitrotoluene, 1, 4-Dinitrobenzene
8. What is meant by delocalization of $\pi$ electrons? How does it affect stability of benzene?
9. What are the conditions for a compound/species to be aromatic according to Huckel's rule?
10. How will you convert benzene into
(i) Acetophenone
(ii) m-Chloronitrobenzene?

## 5-MARKS QUESTIONS

1. Define isomerism. Write all the structural isomers of hexane $\left(\mathrm{C}_{6} \mathrm{H}_{14}\right)$ and arrange them in increasing order of boiling points.
2. Write short note on (i) Wurtz reaction (ii) Kolbe's electrolysis (iii) Ozonolysis
3. An alkene 'A' of molecular mass ' 28 u ' 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.
4. An organic compound ' A ' with formula $\mathrm{C}_{4} \mathrm{H}_{9} \mathrm{Br}$ on treatment with KOH (alc.) gave two isomeric compounds ' B ' and ' C ' with formula $\mathrm{C}_{4} \mathrm{H}_{8}$. Ozonolysis of ' B ' gave only one product CH 3 CHO while ' C ' gave two different products. Identify $\mathrm{A}, \mathrm{B}$ and C .
5. How will you convert Ethyne into (i) 1, 1, 2, 2-Tetrachloroethane (ii) Ethene (iii) Ethanal (iv) Benzene (v) Sodium ethynide

## UNIT TEST-I

Time Allowed: $1 \mathbf{H r}$.
Maximum Marks : 20

## General Instructions:

(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. Amongst the following which one has the maximum boiling point and why? n-Pentane, iso-pentane, neo-pentane
2. What is the number of $\sigma$ and $\pi$ bonds in But-3-en-1-yne?
3. Action of acetylene on dilute $\mathrm{H}_{2} \mathrm{SO}_{4} /$ dil. $\mathrm{HgSO}_{4}$ gives:
(a) Acetic acid
(b) Acetone
(c) Acetaldehyde
(d) Ethyl alcohol

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 but 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.
4. Assertion : Benzone reacts with chlorine in the form of light to from BHC.

Reason: BHC is also called gammexane or 666.
5. Assertion: Tropylium cation

Reason: The only property which decides the aromatic character is its planar nature.
6. Arrange the following alkenes in decreasing order of stability and give reason.

$$
\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{3}, \quad \mathrm{CH}_{2}=\mathrm{CH}_{2}, \quad \mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2}
$$

7. (i) Give a chemical test to distinguish between ethyne and ethene.
(ii) Melting point of cis-But-2-ene is lower than that of trans-But-2-ene. Give reason.
8. Complete the following reactions:
(i) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br} \xrightarrow{\text { Alc. } \mathrm{KOH}}$
(ii) $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}+\mathrm{O}_{3} \xrightarrow{\mathrm{Zn} / \mathrm{H}_{2} \mathrm{O}}$
(iii) $\mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{H}_{2} \mathrm{O}+[\mathrm{O}] \xrightarrow{\text { Dil. } \mathrm{KMnO}_{4}}$
9. (i) What are the conditions for a compound/species to be aromatic according to Huckel's rule?
(ii) How will you convert Benzene to acetophenone?
10. (i) An alkene 'A' of molecular mass ' 28 u ' 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.
(ii) Why is benzene extraordinarily stable although it contains three double bonds?
(iii) How can we convert ethyne into benzene?

## UNIT TEST-II

Time Allowed: $1 \mathbf{H r}$.
Maximum Marks : 20

## General Instructions:

(i) All questions are compulsory.
(ii) Maximum marks carried by each question are indicated against it.

1. The methyl group in Benzene ling is
(a) Ortho directing
(b) Ortho and meta directing
(c) Para directing
(d) Ortho and para directing
2. Action of acetylene on dilute $\mathrm{H}_{2} \mathrm{SO}_{4}$ gives :
(a) Acetic acid
(b) Acetone
(c) Acetaldehyde
(d) Ethyl alcohol
3. What is the other name for Geometrical isomerism?

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 but 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.
4. Assertion: Benzene does not decolourise bromine water.

Reason: Benzene is stabilised by resonance due to delocalisation of $\pi$ electrons.
5. Assertion: Acidity of $\mathrm{C}-\mathrm{H}$ bond decreases in the order:
$\mathrm{HC} \equiv \mathrm{CH}>\mathrm{H}_{2} \mathrm{C}=\mathrm{CH}_{2}>\mathrm{H}_{3} \mathrm{C}-\mathrm{CH}_{3}$
Reason: Greater the percentage s-character, more is the acidity of $\mathrm{C}-\mathrm{H}$ bond.
6. How will you convert methyl bromide to ethane?
7. Explain with the help of equation ozonolysis of propene. [2]
8. Write the mechanism of nitration of benzene.
9. How will you convert benzene into
(i) Acetophenone
(ii) Toluene
10. Write short note on the following :
(i) Wurtz reaction.
(ii) Kolbe's electrolysis
(iii) Ozonolysis

## QUESTION FOR PRACTICES-I

1. What are the oxygen moles in 0.5 mol of $\mathrm{CaCO}_{3}$ ?
(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) $(\mathrm{n}-1) \mathrm{f}^{1-14} \mathrm{nd}^{0-1} \mathrm{~ns} 2$
(b) $(\mathrm{n}-1) \mathrm{f}^{0-1} \mathrm{nd}^{2} \mathrm{~ns}^{2}$
(c) $(\mathrm{n}-2) \mathrm{f}^{1-14}(\mathrm{n}-1) \mathrm{d}^{0-1} \mathrm{~ns}^{2}$
(d) $(\mathrm{n}-2) \mathrm{f}^{1-14}(\mathrm{n}-1) \mathrm{d}^{0-2} \mathrm{~ns}^{0-1}$
4. The shape of $\mathrm{IBr}_{2}^{-}$is
(a) Tetrahedral
(b) Planar
(c) Linear
(d) V-shape
5. The body temperature of a normal healthy person is $37^{\circ} \mathrm{C}$. Calculate its value in ${ }^{\circ} \mathrm{F}$.
6. Cs shows maximum photoelectric effect. Why?
7. Which of the following orbitals are possible?
$1 \mathrm{~S}, 2 \mathrm{~S}, 2 \mathrm{P}$ and 3 f
8. $\Delta_{\mathrm{f}} \mathrm{H}^{0}$ for Graphite is $\qquad$ .

## 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 $\mathrm{A}+\mathrm{B} \rightleftharpoons \mathrm{C}, \mathrm{K}=4$ on addition of catalyst K becomes more than 4.
Statement-2 Catalyst only helps to attain the equilibrium faster from either end of reaction.
10. Statement- $1 \mathrm{Cl}_{2}+2 \mathrm{OH}^{-} \longrightarrow \mathrm{ClO}^{-}+\mathrm{Cl}^{-}$is a disproportion reaction.

Statement-2 In disproportionation, the same element get oxidised as well as reduce.
11. -COOH is ortho, Para directing or Meta directing group? Give reason.
12. Draw Cis and trans isomers of $\mathrm{CHCl}=\mathrm{CHCl}$
13. At what temperature the entropy of a perfectly Crystalline solid is zero?
14. Write the IUPAC name of following


OR

15. In what period and group will an element with $\mathrm{Z}=118$ will be present.
16. Calculate number of atoms in 52 u 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 $\mathrm{ClF}_{3}$.

OR, Draw the shape of $\mathrm{SF}_{6}$.
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. List the possible values for all the quantum numbers for the following sub shell
(i) 2 p
(ii) $4 f$
23. (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.
24. Balance the following reaction by (ion-electron or oxidation number method)
$\mathrm{Cl}_{2} \mathrm{O}_{7}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}_{2}$ (aq.) $\longrightarrow \mathrm{ClO}_{2}^{-}($aq. $)+\mathrm{O}_{2}(\mathrm{~g})+\mathrm{H}^{+}$(Acidic medium)
25. Calculate oxidation state of Fe in $\mathrm{Fe}_{3} \mathrm{O}_{4}$.
26. A sample of 0.5 g of an organic compound was treated according to Kjeldahl's method. The ammonia evolved was absorbed in 50 ml of $0.5 \mathrm{M} \mathrm{H}_{2} \mathrm{SO}_{4}$. The residual acid requirede 60 mL of 0.5 solution of NaOH for neutralisation. Find the percentage composition of nitrogen in the compound.
27. (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?
28. Give IUPAC name the following:
(i) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{OH}$
(ii) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CHO}$
29. Inductive effect is of permanent nature while electromeric effect is only temporary. Explain.
30. A crystalline salt on being rendered anhydrous loses $45.6 \%$ of its weight. The percentage composition of the anhydrous salt is
$\mathrm{Al}=10.50 \%, \mathrm{~K}=15.1 \%, \mathrm{~S}=24.96 \%, \mathrm{O}=49.92 \%$
Find the simplest formula of the anhydrous and crystalline salt.
(Atomic Mass : $\mathrm{K}=39, \mathrm{Al}=27, \mathrm{~S}=32, \mathrm{O}=16$ ).
31. Explain following with example :
(i) Aufbau's Rule
(ii) Hund's Rule
(iii) Pauli's Exclusion Principle
32. An ion with mass number 81 contains $31.7 \%$ more neutrons as compared to protons. Assign the atomic symbol.
33. Explain the bonding in $\mathrm{SF}_{6}$ using hybridisation concept and define what is hybridisation.
34. On the basis of molecular orbital theory find the bond order, molecular orbital configuration and magnetic nature of $\mathrm{O}_{2}{ }^{+}$.
35. Calculate pH of a solution obtained by mixing 50 mL of 0.2 MHCl with 49.9 mL of 0.2 m NaOH solution.
36. Explain Born Haber cycle with by considering example of formation of $\mathrm{MgCl}_{2}$ as given in the chemical reaction

$$
\mathrm{Mg}(\mathrm{~s})+\mathrm{Cl}_{2}(\mathrm{~g}) \longrightarrow \mathrm{MgCl}_{2}(\mathrm{~s})
$$

37. Calculate the standard enthalphy of formation of $\mathrm{CH}_{3} \mathrm{OH}(l)$ from the following data:

$$
\begin{array}{ll}
\mathrm{CH}_{2} \mathrm{OH}(l)+\frac{3}{2} \mathrm{O}_{2}(g) \longrightarrow \mathrm{CO}_{2}(g)+2 \mathrm{H}_{2} \mathrm{O}(l) & \Delta_{\mathrm{C}} \mathrm{H}^{\mathrm{o}}=-726 \mathrm{~kJ} \mathrm{~mol}^{-1} \\
\mathrm{C}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g}) & \Delta_{\mathrm{f}} \mathrm{H}^{\mathrm{o}}=-393 \mathrm{~kJ} \mathrm{~mol}^{-1} \\
\mathrm{H}_{2}(\mathrm{~g})+\frac{1}{2} \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{H}_{2} \mathrm{O}(l) & \Delta_{\mathrm{f}} \mathrm{H}^{\mathrm{o}}=-286 \mathrm{~kJ} \mathrm{~mol}^{-1}
\end{array}
$$

38. Arrange following in order of increasing stability
(a) $\stackrel{+}{\mathrm{CH}}_{3}$

$\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}^{+}$

(b)




(c) $\mathrm{CH}_{3} \mathrm{CH}_{2}, \quad \mathrm{CH}_{3}, \quad \mathrm{CH}_{2}-\mathrm{CH}=\mathrm{CH}_{2}, \quad \mathrm{CH}=\mathrm{CH}_{2}$
39. (i) What is ambident nucleophile? Mention one example.
(ii) Distinguish between homolytic and hetrolytic bond cleavage.
(iii) Which one is stronger acid and why

40. (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
41. (a) Find out $\mathrm{K}_{\mathrm{c}}$ for following reaction

$$
2 \mathrm{NOCl}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) ; \mathrm{K}_{\mathrm{p}}=1.8 \times 10^{-4} \text { at } 500 \mathrm{~K}
$$

(b) $\mathrm{K}_{\mathrm{p}}=0.04 \mathrm{~atm}$ at 899 K . What is the equilibrium concentration of $\mathrm{C}_{2} \mathrm{H}_{6}$ where it is placed in a flask at 4.0 atm pressure and allow to come to equilibrium

$$
\mathrm{C}_{2} \mathrm{H}_{6} \rightleftharpoons \mathrm{C}_{2} \mathrm{H}_{4}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g})
$$

(c) What is the unit of $\mathrm{K}_{\mathrm{p}}$ for the following chemical reaction?

$$
2 \mathrm{NH}_{3}(\mathrm{~g}) \rightleftharpoons \mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g})
$$

42. A compound contains $4.07 \% \mathrm{H}, 24.27 \% \mathrm{C}$ and $71.65 \% \mathrm{Cl}$. Its molar mass is 98.96 g . What are its empirical and molecular formula.
43 Calculate the uncertainty in the position of a dust particle with mass equal to 1 mg if the uncertainity in its velocity is $5.5 \times 10^{-20} \mathrm{~ms}^{-1}$.
43. (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 $\sigma$ and $\pi$ bond(s) in

44. (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


$$
* * * *
$$

## QUESTION FOR PRACTICES-II

1. The number of nodal planes in $p_{x}$ orbitals is
(a) 1
(b) 2
(c) 3
(d) 0
2. Which of the following has smallest bond angle?
(a) $\mathrm{H}_{2} \mathrm{O}$
(b) $\mathrm{H}_{2} \mathrm{~S}$
(c) $\mathrm{NH}_{3}$
(d)
$\mathrm{SO}_{2}$
3. For a reaction to be spontaneous at all the temperature:
(a) $\Delta \mathrm{G}-\mathrm{ve}, \Delta \mathrm{H}+\mathrm{ve}, \Delta \mathrm{S}+\mathrm{ve}$
(b) $\Delta \mathrm{G}+\mathrm{ve}, \Delta \mathrm{H}$-ve, $\Delta \mathrm{S}+\mathrm{ve}$
(c) $\Delta \mathrm{G}-\mathrm{ve}, \Delta \mathrm{H}-\mathrm{ve}, \Delta \mathrm{S}$ - ve
(d) $\Delta \mathrm{G}-\mathrm{ve}, \Delta \mathrm{H}$-ve, $\Delta \mathrm{S}+\mathrm{ve}$
4. The enthalpy of all elements in their standard states are:
(a) unity
(b) zero
(c) less than zero
(d) different for each element
5. Which of the following has largest ionic radii?
(a) $\mathrm{Na}^{+}$
(b) $\mathrm{Mg}^{2+}$
(c) $\mathrm{F}^{-}$
(d) $\mathrm{O}^{2-}$
6. When carbon is bonded to four other atoms or groups it uses $\qquad$ hybrid orbitals.
7. Surface tension $\qquad$ with increase in temperature.
8. The second electron gain enthalpy is $\qquad$ .
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.
10 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 $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{K}_{2} \mathrm{CO}_{3}$
12. Define mole fraction.
13. Mention the quantum number which determines the energy of electron in the H -atom.
14. How many unpaired electrons are there in $\mathrm{Ni}^{2+}$ ion? (Given : $\mathrm{Z}=28$ )
15. State the condition for the formation of precipitate.
16. Write the conjugate acid and conjugate base of $\mathrm{HSO}_{4}{ }^{-}$.
17. Using VSEPR theory draw the shape of $\mathrm{XeF}_{4}$ molecule.
18. Write IUPAC name of $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CH}(\mathrm{OH}) \mathrm{C} \equiv \mathrm{CH}$.
19. In a reaction between an oxidant and a reductant which will give up electrons and which will accept electrons?
20. Calculate oxidation number of Cr in $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$ and Mn in $\mathrm{KMnO}_{4}$.
21. Write the conjugate acids for the following Bransted bases:

$$
\mathrm{NH}_{2}^{-}, \mathrm{NH}_{3}, \mathrm{HCOO}^{-}
$$

22. What will be the mass of one ${ }^{12} \mathrm{C}$ atom in grane?
23. How are frequency and wave number related to each other.
24. Why the I.E. of Ga is greater than Al ?
25. What effect does branching of an alkane chain has on its boiling point?
26. Cis But-2-ene has lower melting point that trans But-2-ene. Give reason.
27. Balance the following redox reaction in acidic medium by ion electron method.
Zn (aq.) $+\mathrm{NO}_{3}^{-}$(aq.) $\longrightarrow \mathrm{Zn}^{2+}($ aq. $)+\mathrm{N}_{2} \mathrm{O}(\mathrm{g})+\mathrm{H}_{2} \mathrm{O}$
28. (i) $\mathrm{Mg}^{2+}$ ion is smaller than O 2 -ion, although both are isoelectronic. Give reason.
(ii) Write IUPAC name and symbol for the element with atomic no. 120.
29. (i) Mention the number of radial nodes in 6 s orbitals.
(ii) Write electronic configuration of $\mathrm{Fe}^{2+}$ ion. (Given, $\mathrm{Z}=26$ )
30. Calculate the wavelength of a ball of mass 0.1 kg moving with a velocity of $10 \mathrm{~ms}^{-1}$. (Given, $\mathrm{h}=6.626 \times 10^{-34} \mathrm{Js}$ )
31. Arrange Benzene, $n$-Hexane and Ethyne in decreasing order of acidic behaviour.
32. Which of the two $\mathrm{O}_{2} \mathrm{NCH}_{2} \mathrm{CH}_{2} \mathrm{O}^{-}$or $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{O}^{-}$is expected to be more stable and why?
33. Write the difference between precision and accuracy.
34. Out of molarity and molarity which changes with temperature.
35. (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?
36. (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?
37. The density of 3 M solution of NaCl is $1.25 \mathrm{~g} / \mathrm{mL}$. Calculate the molality of the solution. (Given: Atomic masses: $\mathrm{Na}=23 \mathrm{u}, \mathrm{Cl}=35.5 \mathrm{u}$ )
38. Calculate the molarity of nitric acid $\left(\mathrm{HNO}_{3}\right)$ in a sample having a density $1.41 \mathrm{~g} / \mathrm{mL}$ and mass percent of nitric acid in it being $69 \%$. (Atomic mass: $\mathrm{N}=14 \mathrm{u}, \mathrm{H}=1 \mathrm{u}, \mathrm{O}=16 \mathrm{u}$ )
39. (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 $\mathrm{l}=2$.
(iii) Chromium has configuration $3 d^{5} 4 s^{1}$ and not $3 d^{4} 4 s^{2}$. Explain.
40. (i) Explain non linear shape $\mathrm{H}_{2} \mathrm{~S}$ and non-planar shape of $\mathrm{PCl}_{3}$ using VSEPR theory.
(ii) Can we have a diatomic molecule with its ground state molecular orbitals full with electrons. Give reason for your answer.
41. Calculate enthalpy change for the reaction:
$\mathrm{CH}_{4}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
The enthalpy of formation of $\mathrm{CH}_{4}(\mathrm{~g}), \mathrm{CO}_{2}(\mathrm{~g})$ and $\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ are $-74.8 \mathrm{kJmol}^{-1}$, $-393.5 \mathrm{kJmol}^{-1}$ and $285.8 \mathrm{kJmol}^{-1}$ respectively.
42. Calculate the bond enthalpy of $\mathrm{Cl}-\mathrm{Cl}$ bond from the following data:
$\mathrm{CH}_{4}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightarrow \mathrm{CH}_{3} \mathrm{Cl}(\mathrm{l})+\mathrm{HCl} \quad \Delta \mathrm{H}=-100.3 \mathrm{kJmol}^{-1}$
Given: bond enthalpies of $\mathrm{C}-\mathrm{H}, \mathrm{C}-\mathrm{Cl}$ and $\mathrm{H}-\mathrm{Cl}$ bonds are 413,326 and $431 \mathrm{kJmol}^{-1}$ respectively.
43. 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: $\mathrm{O}=16 \mathrm{u}$, $\mathrm{Ne}=20 \mathrm{u}$ )
44. An organic compound contains $69 \%$ carbon and $4.8 \%$ Hydrogen, the remainder being oxygen. Calculate the masses of carbon dioxide and water produced when 0.20 g of this substance is subjected to complete combustion.
45. (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 $\pi$ system.
46. (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.
47. (i) Explain common ion effect with example.
(ii) The concentration of hydrogen ion in a sample of soft drink is $4 \times 10-3$ M. Calculate its pH .
(iii) What is the effect of removal of $\mathrm{CH}_{3} \mathrm{OH}$ on the equilibrium of the reaction, $2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{CO}(\mathrm{g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{g})$.
48. (i) Calculate $\mathrm{H}_{3} \mathrm{O}^{+}$ion concentration of a water sample having $\mathrm{pH}=6.78$.
(ii) Define buffer solutions.
(iii) State Lewis definition of acids and bases. Give one example of each.
49. (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) $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}+\mathrm{HBr} \xrightarrow{\text { organic peroxide }}$
(b) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br}+\mathrm{Na} \xrightarrow{\text { Dry Ether }}$
(iii) Write a short note on Friedel Crafts alkylation.
50. (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)?

## QUESTION FOR PRACTICES-III

1. Which of the following d-orbital has double disc (body shooter shape)
(a) $\mathrm{d}_{\mathrm{xy}}$
(b) $\mathrm{d}_{\mathrm{z}^{2}}$
(c) $d_{x^{2}-y^{2}}$
(d) $\mathrm{d}_{\mathrm{yz}}$
2. Among the following, the compound that contains, ionic, covalent and Coordinate linkage is
(a) $\mathrm{NH}_{4} \mathrm{Cl}$
(b) $\mathrm{NH}_{3}$
(c) NaCl
(d) $\mathrm{CO}_{2}$
3. Which of the following is a state function
(a) q (b) w
(c) $q+w$
(d) None of these
4. Which one of the following thermodynamic quantities is not a state function
(a) Gibb free energy
(b) Enthalpy
(c) Entropy
(d) Work
5. The correct order of electron gain enthalpy with negative sign of $\mathrm{F}, \mathrm{Cl}, \mathrm{Br}$ and I having atomic number $9,17,35$ and 53 respectively is
(a) $\mathrm{I}>\mathrm{Br}>\mathrm{Ci}>\mathrm{F}$
(b) $\mathrm{F}>\mathrm{Cl}>\mathrm{Br}>$ I
(c) $\mathrm{Cl}>\mathrm{F}>\mathrm{Br}>$ I
(d) $\mathrm{Br}>$ I $>\mathrm{Cl}>\mathrm{F}$
6. Arrange the following carbonations in order of their increasing stability $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}^{+}, \mathrm{CH}_{3} \mathrm{CH}_{2}^{+},\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CH}^{+}, \mathrm{CH}_{3}^{+}$
7. The IUPAC name and symbol of element with $\mathrm{Z}=120$ is $\qquad$ .
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.
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 be liqufied.

Reason (R) : Critical temperature is the temperature above which a gas cannot be liquefied 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?
12. How is pH scale affected by increasing temperature.
13. Why are alkali metals used in photoelectric cells?
14. Write electronic configuration of $\mathrm{Cr}^{3+}$ ion. (Atomic No . of $\mathrm{Cr}=24$ )
15. Draw the shape of $\mathrm{ClF}_{3}$ molecule according to VSEPR theory?
16. Write the conjugate acid and conjugate base of $\mathrm{H}_{2} \mathrm{O}$.
17. Write the relation between solubility and solubility product of $\mathrm{Cr}_{2}\left(\mathrm{SO}_{4}\right)_{3}$.
18. Consider the given standard reduction potentials of following elements A , $\mathrm{B}, \mathrm{C} \& \mathrm{D}$ and arrange them in order of their increasing reducing power.
$\mathrm{A}=-3.71 \mathrm{~V}, \mathrm{~B}=-0.76 \mathrm{~V}, \mathrm{C}=+0.34 \mathrm{~V}, \mathrm{D}=+0.80 \mathrm{~V}$
19. For the reaction $\mathrm{Co}(\mathrm{g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CoCl}_{2}(\mathrm{~g})$ the value of $\mathrm{kc} / \mathrm{kp}$ is equal to.
20. Chemical species having identical bond order have same bond dissociation enthalpies.
21. Write IUPAC name of $\mathrm{CH}_{3}-\mathrm{CO}-\mathrm{CH}_{2}-\mathrm{CHO}$.
22. Out of Cis-But-2-ene and Trans-but-2-ene which has greater boiling point and why?
23. Neither ' $q$ ' nor ' $w$ ' is state functions but $q+w$ is a state function. Explain.
24. (i) What is the lowest value of n that allows ' g ' orbitals to exist?
(ii) Why 4 s orbital is filled before 3 d ?
25. Calculate wave number for the longest wavelength transition in the Balmer Series of hydrogen atom.
26. Explain why?
(i) $\Delta_{\mathrm{i}} \mathrm{H}_{1}$ of ' N ' is more than that of ' O '.
(ii) A cation is smaller than parent atom.
27. Balance the following redox reaction in acidic medium by ion electron method.

$$
\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\text { aq. })+\mathrm{SO}_{2}(\mathrm{~g}) \longrightarrow \mathrm{Cr}^{3+} \text { (aq.) }+\mathrm{SO}_{4}{ }^{2-} \text { (aq.) }
$$

28. What is chromatography? Name different types chromatographic processes.
29. Write reaction is suitable for the preparation of both symmetrical and unsymmetrical alkanes.
30. In sulphur estimate, 0.157 g of an organic compound gave 0.4813 g of barium sulphate. What is the percentage of sulphur in the compound?
31. Write chemical equations to covert:
(i) Ethyne to Ethanal
(ii) Benzene to m-Nitrotoluene
32. The density of 3 M solution of NaCl is $1.25 \mathrm{~g} \mathrm{ml}^{-1}$ calculate molarity of the solution.
33. 3 g of $\mathrm{H}_{2}$ react with 29 g of $\mathrm{O}_{2}$ yield $\mathrm{H}_{2} \mathrm{O}$
(a) Which is the limiting reagent?
(b) Calculate the maximum amount of $\mathrm{H}_{2} \mathrm{O}$ that can be formed.
34. Calcium carbonate reacts with aqueous HCl according to the reaction:
$\mathrm{CaCO}_{3}(\mathrm{~s})+2 \mathrm{HCl}($ aq. $) \longrightarrow \mathrm{CaCl}_{2}($ aq. $)+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
What mass of CaCO 3 is required to react completely with 25 mL of 0.75 M HCl ?
35. (i) State Hund's rule of maximum multiplicity of electrons.
(ii) The mass of an electron is $9.1 \times 10^{-31} \mathrm{~kg}$. If its K.E. $3.0 \times 10^{-25} \mathrm{~J}$, calculate its wavelength.
36. $\mathrm{N}_{2}$ is diamagnetic while $\mathrm{O}_{2}$ is paramagnetic. Explain on the basis of molecular orbital theory.
37. Explain the structure of $\mathrm{PCl}_{5}$ according to hybridization. Why all $\mathrm{P}-\mathrm{Cl}$ bonds.
38. What will be the pressure exerted by a mixture of 3.2 g of methane and
4.4 g of carbon dioxide contained in $9 \mathrm{dm}^{2}$ flask at $27^{\circ} \mathrm{C}$.
39. Calculate the compressibility factor for $\mathrm{CO}_{2}$, if one mole of it occupies 0.4 litre at 300 K and 40 atm . Comment on the result.
40. The combustion of 1 mol of benzene takes place at 298 K . After combustion $\mathrm{CO}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ are formed and $3267 \mathrm{~kJ} \mathrm{~mol}^{-1}, \Delta_{\mathrm{f}} \mathrm{H}^{\ominus}\left(\mathrm{H}_{2} \mathrm{O}\right)=-393 \mathrm{~kJ} \mathrm{~mol}^{-1}$.
41. For the reaction : $2 \mathrm{~A}(\mathrm{~g})+\mathrm{B}(\mathrm{g}) \longrightarrow 2 \mathrm{D}(\mathrm{g}), \Delta \mathrm{U}^{\ominus}=-10.5 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{\circ}=-44.1$ $J \mathrm{~K}^{-1}$. Calculate $\Delta \mathrm{G}^{\circ}$ for the reaction, and predict whether the reaction will occur spontaneously.
42. Account for the following:
(i) Halogen have very high negative electron gain enthalpy
(ii) Ionisation enthalpy of nitrogen $(z=7)$ is more than oxygen $(z=8)$
43. (i) Out of $\mathrm{NO}_{2}-\mathrm{CH}_{2}-\mathrm{O}$ and $\mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{O}^{-}$which is more stable and why?
(ii) Why is it necessary to prepare Lassaigne extract for detection of $\mathrm{N}, \mathrm{S}$ and halogens?
(iii) Define the term hyperconjugation.
44. (i) State Le Chatelier's principle.
(ii) Calculate the pH of $10^{-8} \mathrm{M} \mathrm{HCl}$.
(iii) A sample of $\mathrm{HI}(\mathrm{g})$ is placed in flask at a pressure of 0.2 atm . At equalibrium the partial pressure of $\mathrm{HI}(\mathrm{g})$ is 0.04 atm . What is the Kp for given equilibrium?
$2 \mathrm{HI}(\mathrm{g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g})$
45. (i) Define common ion effect.
(ii) Write the relationship between $\mathrm{K}_{\mathrm{p}}$ and $\mathrm{K}_{\mathrm{c}}$ for the reaction:
$\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})$
(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 $\mathrm{K}_{\mathrm{sp}}=7.4 \times 10^{-8}$ )
46. Compare relative stability of following species and predict their magnetic
properties:
$\mathrm{O}_{2}, \mathrm{O}_{2}^{+}, \mathrm{O}_{2}^{-}, \mathrm{O}_{2}^{2-}$
47. Give reason for the following:
(i) $\mathrm{NH}_{3}$ has higher boiling point than $\mathrm{pH}_{3}$
(ii) Ionic compound do not conduct electricity in solid state
(iii) $\mathrm{H}_{2} \mathrm{O}$ has bent structure.
48. (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) $\mathrm{C}_{6} \mathrm{H}_{6}+\mathrm{CH}_{3} \mathrm{Cl} \xrightarrow{\text { Anhydrous } \mathbf{A l C l}_{3}}$
(b) $\mathrm{CH}_{3} \mathrm{Cl}+\mathrm{Na} \xrightarrow{\text { Dry ether }}$
(c) $\mathrm{CH}_{3}-\mathrm{CHCl}-\mathrm{CH}_{2}-\mathrm{CH}_{3}+\mathrm{KOH}($ alc. $) \longrightarrow$
49. (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.

## QUESTION FOR PRACTICES-IV

1. Number of matamers possible for molecular formula $\mathrm{C}_{4} \mathrm{H}_{10}$ are $\qquad$ .
2. How is the mass of an element related the number of atoms present in it.
3. Haemoglobincontain $0.25 \%$ iron by mass. The molecular mass of haemoglobinis 89600 . Calculate the number of iron atoms per molecule of haemoglobin.
4. Orbital which is not possible
(a) 2 p
(b) 3 d
(c) 3 s
(d) 3 f
5. Which of the following is paramagnetic
(a) CO
(b) $\mathrm{O}_{2}^{-}$
(c) $\mathrm{N}_{2}$
(d) $\mathrm{NO}^{+}$
6. Which of the following is not correct?
(a) $\Delta \mathrm{G}$ is zero for a reversible reaction.
(b) $\Delta \mathrm{G}$ is positive for a spontaneous reaction.
(c) $\Delta \mathrm{G}$ is negative for a spontaneous reaction.
(d) $\Delta \mathrm{G}$ is positive for a non-spontaneous reaction.

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 $n=n^{2}$.
8. Assertion : Ionic radius of $\mathrm{Na}^{+}$is smaller than Na .

Reason : Effective nuclear charge of $\mathrm{Na}^{+}$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 $\mathrm{SO}_{2}$ are present in 11.2 L at STP?
12. Calculate the number of atoms in 35 g of Li (Atomic mass of $\mathrm{Li}=7 \mathrm{u}$ ).
13. For an isolated system, $\Delta \mathrm{U}=0$, what will be $\Delta \mathrm{S}$ ?
14. Write the statement of third law of thermodynamics.
15. Calculate the number of sigma and pie bonds in $\mathrm{C}_{2} \mathrm{H}_{2}$ ?
16. Give the relation between $\mathrm{K}_{\mathrm{a}}$ and $\mathrm{K}_{\mathrm{b}}$.
17. Write the conjugate base of $\mathrm{H}_{2} \mathrm{CO}_{3}$.
18. Write the oxidation number of Cr in $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$.
19. Write the stock notation $\mathrm{MnO}_{2}$.
20. Suggest angular and spherical node on (a) 4 s (b) 2 p (c) 3 d
21. Name the series of the line spectrum of H -spectrum which lies in visible region.
22. Assign the oxidation number of underline element

23. Identify oxidizing and reducing agent in:

$$
\mathrm{CH}_{4}+4 \mathrm{Cl}_{2} \longrightarrow \mathrm{CCl}_{4}+4 \mathrm{HCl} .
$$

24. Out of stagged and eclipsed form which is were stable and why?
25. What are carbocations? Give the stability order of different carbocations.
26. Explain why :
(a) The three electron present in 2 p subshell of nitrogen remain upaired.
(b) Cr has configuration $3 d^{5} 4 s^{1}$ and not $3 d^{4} 4 s^{2}$.
27. Calculate the radius of Bohr's fifth orbit for hydrogen atom.
28. (a) Why electron gain enthalpy of Cl is more -ve than F .
(b) Give group number, period and block of an element having atomic number :
(i) 86
(ii) 55
(iii) 74
(iv) 63
29. Balance the following redox reaction in acidic medium

$$
\mathrm{MnO}_{4}^{-}+\mathrm{H}^{+}+\mathrm{Fe}^{2+} \longrightarrow \mathrm{Mn}^{2+}+\mathrm{H}_{2} \mathrm{O}+\mathrm{Fe}^{3+} .
$$

30. Write the resonance structure of (a) $\mathrm{CO}_{3}{ }^{2-}$ (b) $\mathrm{NO}_{3}{ }^{-}$
31. Explain why $\mathrm{BeCl}_{2}$ molecule have zero dipole moment.
32. 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 $\mathrm{Ag} \& \mathrm{Br}$ are 105 and 80 u$]$
33. Propanal and pentan-3-one ate the ozonolysis product of an alkene. What is the structural formula of alkene?
34. Convert the following :
(a) 1-Bromopropane to 2-Bromopropane
(b) Ethanol to Glycol
35. Explain the following terms (a) Tautomerism (b) metamerism.
36. $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}+\mathrm{O}_{3} \longrightarrow \mathrm{X} \xrightarrow{\mathrm{Zn} / \mathrm{H}_{2} \mathrm{O}} \mathrm{A}+\mathrm{B}$.

Identify A and B . Also give reactions.
(a) Green house effect
(b) Green chemistry
37. In a compound $\mathrm{C}_{x} \mathrm{H}_{\mathrm{y}} \mathrm{O}_{z}$, 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 $\mathrm{C}_{\mathrm{x}} \mathrm{H}_{\mathrm{y}}$. Find the empirical formula of the compound.
38. When light with a wavelength of 400 nm falls on the surface of sodium, electrons with a kinetic energy of $1.05 \times 10^{5} \mathrm{~J} \mathrm{~mol}^{-1}$ 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?
39. (a) Describe the hybridisation in case of $\mathrm{C}_{2} \mathrm{H}_{2}$.
(b) Which out of $\mathrm{NH}_{3}$ and $\mathrm{NF}_{3}$ has higher dipole moment and why?
40. (a) Use molecular orbital theory to predict why the $\mathrm{He}_{2}$ molecule does not exist?
(b) Compare the stability of $\mathrm{O}_{2}$ and $\mathrm{O}_{2}^{+}$on the basis of molecular theory.
41. Calculate the maximum work of expansion of two moles of an ideal gas when it expand isothermally and reversibly from 20 L to 40 L at $27^{\circ} \mathrm{C}(\mathrm{R}=$ 1.987 cal per kelvin per mol.
42. Heat produced during the combustion of benzoic acid $(\mathrm{C} 6 \mathrm{H} 5 \mathrm{COOH})$ at 298 K and 1 atm is -2500 KJ per mole. What is $\Delta \mathrm{E}$ for the reaction.
43. The standard enthalpy of combustion of Surcose $\left(\mathrm{C}_{12} \mathrm{H}_{32} \mathrm{O}_{11}\right)$ at 298 K producing $\mathrm{CO}_{2}(\mathrm{~g})$ and $\mathrm{H}_{2} \mathrm{O}(l)$ is $-5200.7 \mathrm{~kJ} \mathrm{~mol}^{-1}$. If $\Delta_{\mathrm{f}} \mathrm{H}^{\circ}$ of $\mathrm{CO}_{2}(\mathrm{~g})$ and $\mathrm{H}_{2} \mathrm{O}(l)$ are $-393.0 \mathrm{~kJ} \mathrm{~mol}^{-1}$ and $-286 \mathrm{~kJ} \mathrm{~mol}^{-1}$.
44. Explain the First law of thermodynamics and how can you prove the following relation :
$\Delta \mathrm{H}=\Delta \mathrm{U}+\mathrm{P} \Delta \mathrm{V}$ from the first law of the thermodynamics $(\Delta \mathrm{U}=\mathrm{q}+\mathrm{w})$ ?
45. Give structural formula of following compound:
(a) But-2-enoic acid
(b) 4-Hydroxypent-3-enoic acid
(c) 3-Cyclopropyl-3-methyulbutanal.
46. Explain the resonance effect and draw the resonance structures of $\mathrm{CH}_{2}=$ $\mathrm{CH}-\mathrm{CHO}$. Indicate the relative stability of the contributing structures.
47. (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.
48. (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
49. (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 $=7.4 \times 10^{-8}$ ).
50. (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 $10^{-10} \mathrm{M} \mathrm{NaOH}$ solution.
51. If 20 mL of $0.1 \mathrm{M} \mathrm{BaCl}_{2}$ is mixed with 15 mL of $0.01 \mathrm{M} \mathrm{AgNO}_{3}$, Predict the precipitation of AgCl if its solubility product is $1.47 \times 10^{-10}$.
52. Describe the effect of
(a) addition of $\mathrm{Cl}_{2}$
(b) addition of $\mathrm{NO}_{2} \mathrm{Cl}$
(c) removal of $\mathrm{NO}_{2}$
(d) removal of $\mathrm{NO}_{2} \mathrm{Cl}$

On the equilibrium of reaction:
$\mathrm{Cl}_{2}(\mathrm{~g})+2 \mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}_{2} \mathrm{Cl}(\mathrm{g})$

# PRACTICE PAPER-I <br> CLASS - XI <br> CHEMISTRY THEORY (043) 

Time: 3 Hours
Max. Marks : 70

## General instructions: read the following instructions carefully.

1. There are 33 questions in this question paper with internal choice.
2. Section A consists of 16 multiple choice questions carrying 1 mark each.
3. Section B consists of 5 short answer questions carrying 2 marks each
4. Section C consists of 7 short answer questions carrying 3 marks each.
5. Section D consists of 2 case-based questions carrying 4 marks each.
6. Section E consists of 3 long answer questions carrying 5 marks each.
7. All questions are compulsory.
8. Use of $\log$ table and calculator is not allowed.

## SECTION-A

The following questions are multiple choice questions with one correct answer. Each question carries 1 mark. There is no internal choice in this section.

1. Among the following group, the group having highest priority in IUPAC nomenclature scheme is
(a) F
(b) -OCH
(c) $-\mathrm{CH}_{3}$
(d) -COOH
2. Which of the following is heavier in weight
(a) 2 g He
(b) 22.4 L at STP He
(c) 20 moles of $\mathrm{H}_{2}$
(d) 10 moles of $\mathrm{N}_{2}$
3. The molarity of a solution obtained by mixing 750 ml of 0.5 M HCI with 250 ml of 2 M HCI will be
(a) 0.975 M
(b) 0.875 M
(c) 1.00 M
(d) 1.175 M
4. What transition in H -spectrum would have the same wavelength as the Balmer transition $\mathrm{n}=4$ to $\mathrm{n}=2$ in the $\mathrm{He}^{+}$spectrum
(a) $n=4$ to $n=1$
(b) $\mathrm{n}=3$ to $\mathrm{n}=2$
(c) $\mathrm{n}=3$ to $\mathrm{n}=1$
(d) $n=2$ to $n=1$
5. The total number of orbitals associated with third shell will be
(a) 2
(b) 4
(c) 9
(d) 3
6. The correct order of ionic radii of the species $\mathrm{N}^{3-}, \mathrm{O}^{2-}, \mathrm{Na}^{+}$and $\mathrm{F}^{-}$is
(a) $\mathrm{Na}^{+}<\mathrm{F}^{-}<\mathrm{O}^{2-}>\mathrm{N}^{3-}$
(b) $\mathrm{F}<\mathrm{O}^{2-}<\mathrm{N}^{3-}>\mathrm{Na}^{+}$
(c) $\mathrm{O}^{2-}<\mathrm{N}^{3-}<\mathrm{F}^{-}>\mathrm{Na}^{+}$
(d) $\mathrm{N}^{3-}<\mathrm{Na}^{+}<\mathrm{F}^{-}>\mathrm{O}^{2-}$
7. Which molecule have zero dipole moment
(a) Ammonia
(b) water
(c) Carbon dioxide
(d) $\mathrm{CHCI}_{3}$
8. Which of the following molécule has coordinate bond
(a) $\mathrm{CH}_{4}$
(b) $\mathrm{H}_{2} \mathrm{O}$
(c) $\mathrm{CO}_{2}$
(d) $\mathrm{NH}_{4}^{+}$
9. The correct order of the bond order in the following species are
(a) $\mathrm{O}_{2}{ }^{2+}>\mathrm{O}_{2}^{+}>\mathrm{O}_{2}^{-}$
(b) $\mathrm{O}_{2}{ }^{2+}<\mathrm{O}_{2}^{-}<\mathrm{O}_{2}^{+}$
(c) $\mathrm{O}_{2}^{+}>\mathrm{O}_{2}^{-}>\mathrm{O}_{2}^{2-}$
(d) $\mathrm{O}_{2}^{-}<\mathrm{O}_{2}^{+}<\mathrm{O}_{2}^{2+}$
10. A process will be spontaneous at all temperature if
(a) $\Delta \mathrm{H}>0$ and $\Delta \mathrm{S}<0$
(b) $\Delta \mathrm{H}<0$ and $\Delta \mathrm{S}>0$
(c) $\Delta \mathrm{H}<0$ and $\Delta \mathrm{S}<0$
(d) $\Delta \mathrm{H}<0$ and $\Delta \mathrm{S}>0$
11. According to second law of thermodynamics
(a) $\Delta \mathrm{S}_{\text {total }}=+\mathrm{ve}$
(b) $\Delta \mathrm{S}_{\text {total }}=-\mathrm{ve}$
(c) $\Delta \mathrm{S}_{\text {system }}=+\mathrm{ve}$
(d) $\Delta \mathrm{S}_{\text {system }}=-\mathrm{ve}$
12. In what manner increase in pressure affect the following equilibrium?
$\mathrm{C}(\mathrm{s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightleftharpoons \mathrm{CO}(\mathrm{g})+\mathrm{Hg}$
(a) Shift in forward direction
(b) Shift in reverse direction
(c) Increase in yield of hydrogen
(d) No effect

In the following questions ( $\mathbf{Q} .13$ to 16) two statements are given labelled as Assertion (A) and Reason (R), choose the correct answer out of the following choices.
(a) Both A and R are true and R is correct explanation of A .
(b) Both A and R are true and R is not the correct explanation of A .
(c) A is true but R is false
(d) A is false but R is true
13. Assertion: ammonia is a lewis base

Reason: $\mathrm{NH}_{3}$ molecule is pyramidal in shape.
14. Assertion: a liquid crystallises in solid and is accompanied by decrease in entropy

Reason: In crystals molecules organise in an ordered manner
15. Assertion: Oxidation state of hydrogen in HCI is +1 but in NaH is -1

Reason: NaH is a metal hydride and for hydrides hydrogen is assigned an oxidation state of -1 .
16. Assertion: Benzene is non-aromatic compound

Reason: compounds which follow the Huckle's rule are aromatic in nature

## SECTION B

17. The uncertainty in momentum of a particle is $2.5 \times 10^{-14} \mathrm{gcms}^{-1}$ with what accuracy can its position be determined. $\left(\mathrm{h}=6.25 \times 10^{-27} \mathrm{gcms}^{-1}\right)$
18. Account for the following
(i) Nitrogen has more first ionization energy than oxygen.
(ii) The electron gain enthalpy of Cl is more negative than that of fluorine.
19. Calculate the pH of 0.02 M acetic acid solution. $\mathrm{K}_{\mathrm{a}}$ for acetic acid solution is $1.8 \times 10^{-5}$ at 298 K
20. (A) Give IUPAC name of $\mathrm{CH}_{2}=\mathrm{CH}-\mathrm{CH}_{2}-\mathrm{CHOH}-\mathrm{CH}_{3}$
(B) give metamer of N -Ethylethanamine $\left(\mathrm{C}_{2} \mathrm{H}_{5}-\mathrm{NH}-\mathrm{C}_{2} \mathrm{H}_{5}\right)$

OR
Explain (a) $\mathrm{ClCH}_{2}-\mathrm{COOH}$ is more acidic than $\mathrm{CH}_{3}-\mathrm{COOH}$
(b) arrange in order of increasing stability $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}^{+}, \mathrm{CH}_{3} \mathrm{CH}_{2}^{+}, \mathrm{CH}_{3} \mathrm{C}^{+} \mathrm{HCH}_{3}, \mathrm{CH}_{3}^{+}$
21. Explain the following (a) kolbe's electrolysis
(b) wurtz reaction

## SECTION-C

22. Answer the following
(a) Why cation is always smaller than parent atom
(b) Why lonisation energy of B is less then Be
(c) Give group number and period mumber of an element having atomic number 75
23. (a) molar heats of combustion of $\mathrm{C}_{2} \mathrm{H}_{2}(\mathrm{~g}), \mathrm{C}$ (graphite) and $\mathrm{H}_{2}(\mathrm{~g})$ are $310.62 \mathrm{kcal}, 94.05 \mathrm{kcal}$ and 68.12 kcal respectively. Calculate the standard heat of formation of $\mathrm{C}_{2} \mathrm{H}_{2}(\mathrm{~g})$.
(b) Define intensive property
24. (a) for a reaction $2 \mathrm{~A}+\mathrm{B} \longrightarrow \mathrm{C}$

$$
\Delta \mathrm{H}=400 \mathrm{KJ} \mathrm{~mol}^{-1} \text { and } \Delta \mathrm{S}=0.2 \mathrm{~kJ} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}
$$

At what temperature will the reaction become spontaneous considering $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ to be combatant over the temperature range.
(c) Define enthalpy of formation
25. At a certain temperature equilibrium constant $\left(\mathrm{K}_{\mathrm{c}}\right)$ is 16 for the reaction $\mathrm{SO}_{2}(\mathrm{~g})+\mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{SO}_{3}(\mathrm{~g})+\mathrm{NO}(\mathrm{g})$
If the container contain 1 M concentration of each component initially then what is concentration of SO, at equilibrium.
26. Balance the following equation by in oxidation number method
$\mathrm{P}_{4}(\mathrm{~s})+\mathrm{OH}^{-}(\mathrm{aq}) \longrightarrow \mathrm{PH}_{3}+\mathrm{H}_{2} \mathrm{PO}_{2}^{-}$
(Basic medium)
27. (a) identify the electrophiles among the following

$$
: \mathrm{CCl}_{2}, \mathrm{NH}_{3}, \mathrm{Br}^{+}, \mathrm{OH}^{-}
$$

(b) Give any two difference between inductive and electromeric effect.
28. One mole of a hydrocarbon A reacts with 1 mole of bromine giving a dibromo compound $\mathrm{B}, \mathrm{C}_{5} \mathrm{H}_{10} \mathrm{Br}_{2}$. Compound A on treatment with cold dilute $\mathrm{KMnO}_{4}$ solution form a compound $\mathrm{C}, \mathrm{C}_{5} \mathrm{H}_{12} \mathrm{O}$. On ozonolysis A gives equimolar quantities of propanone and ethanal. Deduce the structure of A and give reaction of A with $\mathrm{KMnO}_{4}$ and ozonolysis.

## SECTION D

29. Head the case study given below and answer the following questions:

The valence Shell Electron Pair Repulsion theory by Sidgwick and powell in 1940, proposed a simple theory based on the repulsive interaction of the electron pairs in the valence shell of the atoms. It was further developed and redefined by NYholm and Gillespie (1957) and they put forward the concept of important difference of lone pair and the bonding pairs of electrons. While the lone pairs are localized on the central atom, each bonded pair is shared between two atoms. In the molecules the bond pair, Lone pairs (if any) will occupy such position around the central atom to obtained a min-
imum repulsion energy states. These repulsion effects result in deviation from idealized shapes and alteration in bond angles in molecules.
(A) Which have higher bond angle NH , or $\mathrm{H}, \mathrm{O}$. Give reason

OR
Give shape of following compounds on Basis of VSPER theory
(a) $\mathrm{PCl}_{5}$ and (b) $\mathrm{XeF}_{2}$
(B) Give an example of molecule having tetrahedral geometry
(C) Arrange in order of increasing repulsion

Lone pair-bond pair, bond pair-bond pair, lone pair-lone pair
30. Alkenes undergoes addition reaction and oxidation reactions due to presence of unsaturation Addition reactions are those in which the attacking reagent adds to the substrate molecule without elimination. ie. two molecules combine to form a single molecule. Such reactions are given by those molecules which possess at least one multiple bond. In such reactions for each $\pi$ bond of the molecules, two sigma bond is formed and hybridization state of the C -atom changes from $\mathrm{sp}^{2}$ to $\mathrm{sp}^{3}$. This addition may be electrophilic addition, nucleophilic addition or free radical addition. Addition takes place either according to markownkov's rule or according to antimarkownikov's rule depending on molecule to be added or conditions of additions
(A) Complete the following reaction


An organic compound X of molecular formula $\mathrm{C}_{6} \mathrm{H}_{12}$ absorb one mole of bromine to give $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{Br}_{2}$. On ozonolysis it gives two molecules of propanal: Give structure of X and its reaction with ozone.
(B) How will you convert 2-bromopropane to 1-bromopropane.
(C) Name the monomer of polyethene.

## SECTION-E

31. (a) calculate the ratio of energy of electron in the ground state of hydrogen to electron in the first excited state of $\mathrm{Be}^{+3}$.
(b) Give electronic configuration of i) Cr ii) $\mathrm{Fe}^{+2}$
(c) an electron is moving with velocity $2 \times 10^{3} \mathrm{mS}^{-1}$ find the wavelength of the electron ( $\mathrm{h}=6.62 \times 10^{-34} \mathrm{j}$ s)
32. Give answer to the following questions:
(a) State markownikov's rule. Explain with example.
(b) Draw sawhorse projection of conformation of ethane.
(c) Halogens are ortho and para directing but deactivating. Explain.

OR
(a) What is peroxide effect. Explain with example.
(b) State homolytic fission.
(c) In sulphur estimation, 0.157 g of an organic compound gave 0.4813 g of BaSO4. What is percentage of sulphur in organic compound.
33. (A) a compound containing sodium, sulphur, hydrogen and oxygen gave the following results on analysis; $\mathrm{Na} 14.28 \% \mathrm{~S}=9.92 \% \mathrm{H}=6.2 \%$. Calculate the molecular formula of the anhydrous compound. If H is present in the form of water of crystallization only, what is the structure of the crystalline salt? (molecular weight of crystalline salt=322)
(B) why do we prefer molality over molarity for expressing the concentration of a solution?
(C) State limiting reagent

OR
(a) Calculate mole fraction of benzene in a solution containing $30 \%$ by mass in $\mathrm{CC}_{4}$.
(b) What volume of oxygen at NTP is needed to cause the complete combustion of 200 ml of acetylene? Also calculate the volume of $\mathrm{CO}_{2}$ formed,

## Solution

1. (d)
2. (d)
3. (b)
4. (d)
5. (c)
6. (b)
7. (c)
8. (d)
9. (a)
10.(b)
10. (a)
11. (b)
12. (b)
13. (a)
15.(a)
14. (d)
15. $\Delta \mathrm{x}=\mathrm{h} / 4 \times 3.14 \Delta \mathrm{p}=2.11 \times 10^{-14}$
16. (i) due to stable exactly half filled $\mathrm{p}^{3}$ configuration in nitrogen
(ii) because of interelectronic repulsions in F due to small sire some of energy is used to overcome the repulsion
17. $\left[\mathrm{H}^{+}\right]=\sqrt{ } \mathrm{K}_{\mathrm{a}} \cdot \mathrm{C}=\sqrt{ } 1.8 \times 10^{-5} \times 0.02=6 \times 10^{-4}$ $\mathrm{pH}=-\log \left[\mathrm{H}^{+}\right]=-\log \left(6 \times 10^{-4}\right)=3.22$
18. (a) Pent-4-en-2-ol.
(b) $\mathrm{CH}_{3}-\mathrm{NH}-\mathrm{C}_{3} \mathrm{H}_{7}$

OR
(a) $\mathrm{ClCH}_{2} \mathrm{COOH}$ is more acidic than $\mathrm{CH}_{3}-\mathrm{COOH}$ due to -1 effect of Cl
(b) $\mathrm{CH}_{3}^{+}<\mathrm{CH}_{3} \mathrm{CH}_{2}^{+}<\mathrm{CH}_{3} \mathrm{C}^{+} \mathrm{HCH}_{3}<\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}^{+}$,
21. (a) $\mathrm{CH}_{3} \mathrm{COONa} \rightarrow \mathrm{CH}_{3} \mathrm{COO}^{-}+\mathrm{Na}^{+}$

At anode: $\mathrm{CH}_{3} \mathrm{COO}^{-} \rightarrow \mathrm{CH}_{3} \mathrm{COO}+\mathrm{e}^{-}$
$\mathrm{CH}_{3} \mathrm{COO} \rightarrow \stackrel{\dot{\mathrm{C}} \mathrm{H}_{3}+\mathrm{CO}_{2}}{ }$
$\dot{\mathrm{C}} \mathrm{H}_{3}+\dot{\mathrm{C}} \mathrm{H}_{3} \rightarrow \mathrm{CH}_{3}-\mathrm{CH}_{3}$
At cathode: $2 \mathrm{H}_{2} \mathrm{O}+\mathrm{e}^{-} \rightarrow \mathrm{H}_{2}+2 \mathrm{OH}^{-}$
(b) $\mathrm{CH}_{3} \mathrm{Cl}+2 \mathrm{Na}+\mathrm{CH}_{3} \mathrm{Cl} \rightarrow \mathrm{CH}_{3}-\mathrm{CH}_{3}+2 \mathrm{NaCl}$
22. (a) due to increase of effective nuclear charge with lose of electron.
(b) due to more penetration of s electron to be removed from Be than B in which p electron is removed.
23. (a) $\Delta \mathrm{H}=2 \times(-94.05)+(-68.32)-(-310.62)=54.20 \mathrm{kcal}$
(b) the properties which depend on nature of substance and not on quantity of substance.
24. $\Delta \mathrm{G}=\Delta \mathrm{H}-\mathrm{T} \Delta \mathrm{S}$ for spontaneous process $\Delta \mathrm{G}<0$
i.e. $\Delta \mathrm{H}-\mathrm{T} \Delta \mathrm{S}<0 \Rightarrow \Delta \mathrm{H}<\mathrm{T} \Delta \mathrm{S} \Rightarrow \mathrm{T}>\Delta \mathrm{H} / \Delta \mathrm{S}$
$\Delta \mathrm{T}>400 / 0.2 \Rightarrow>2000 \mathrm{~K}$
25.
$\mathrm{T}=0 \quad 1$
$\mathrm{T}=\mathrm{t}_{\text {eq }} \quad 1-\mathrm{X}$
Therefore
$\mathrm{SO}_{2}(\mathrm{~g})+\mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{SO}_{3}(\mathrm{~g}) \quad+\quad \mathrm{NO}(\mathrm{g})$

Hence $\left[\mathrm{SO}_{2}\right]=1-0.6=0.4 \mathrm{moles} /$ Litre
26. $\mathrm{P}_{4}(\mathrm{~s})+3 \mathrm{OH}^{-}(\mathrm{aq})+3 \mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{PH}_{3}+3 \mathrm{H}_{2} \mathrm{PO}_{2}^{-}$
27. (a) : $\mathrm{CC}_{2}, \mathrm{Br}^{+}$
(b) Inductive effect 1. It is permanent displacement of electron of C-chain due to presence of an atom or group of atom attached with C chain having electronegativity different from C.

## 2. Presence of multiple bond is not required

Electromeric effect: 1 it is temporary shifting of electrons of rt bond towards one of the bonded atom
2. Presence of multiple bond is required. Or any two.
28. $\mathrm{A}=\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}=\mathrm{CHCH}_{3}$
$\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}=\mathrm{CHCH}_{3}+\mathrm{H}_{2} \mathrm{O}+[\mathrm{O}] \xrightarrow{\mathrm{KMnO}_{4} / \mathrm{OH}-}\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}(\mathrm{OH})-\mathrm{CH}(\mathrm{OH}) \mathrm{CH}_{3}$
$\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}=\mathrm{CHCH}_{3}, \xrightarrow{\text { (i) } \mathrm{O}_{3} \text { - (ii) } \mathrm{Zn} / \text { water }}\left(\mathrm{CH}_{3}\right)_{2} \mathrm{C}=\mathrm{O}+\mathrm{O}=\mathrm{CHCH}_{3}$
29. (A) $\mathrm{NH}_{3}$

Reason: NH, Have only one lone pair while water have two lone pair of electron. Due to higher lone pair-lone pair repulsion in water bond angle decrease.

OR
(a) PC, trigonal bipyramidal
(b) Xef, linear
(B) Methane
(C) bond pair-bond pair < Lone pair-bond pair < lone pair-lone pair
30. (a) $\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{HBr} \longrightarrow \mathrm{CH}_{3}-\mathrm{CHBr}-\mathrm{CH}_{3}$
$\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{HBr} \xrightarrow{\text { peroxide }} \mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{CH}_{2} \mathrm{Br}$
$\mathrm{X}=\mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{2}-\mathrm{CH}_{3}$
$\mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{2} \mathrm{CH}_{3} \xrightarrow{\text { (i) } \mathrm{O}_{3} \text { - (ii) } \mathrm{Zn} / \text { water }} \mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{CHO}+$ $\mathrm{OCH}-\mathrm{CH}_{2}-\mathrm{CH}_{3}$
(b) $\mathrm{CH}_{3}-\mathrm{CHBr}-\mathrm{CH}_{3} \xrightarrow{\mathrm{KOH} / \text { alcohol }} \mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2} \xrightarrow{\mathrm{HBr} / \text { peroxide }}$ $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{Br}$
(c) ethene
31. (a) $\mathrm{E}_{1} / \mathrm{E}_{2}=\mathrm{Z}_{1}{ }^{2} \mathrm{n}_{2}{ }^{2} / \mathrm{Z}_{2}^{2} \mathrm{n}_{1}^{2}=1 / 4$
(b) $\mathrm{Cr}=[\mathrm{Ar}] 3 \mathrm{~d}^{5} 4 \mathrm{~s}^{1} \mathrm{Fe}^{+2}=[\mathrm{Ar}] 3 \mathrm{~d}^{6}$
(c) $\lambda=\mathrm{h} / \mathrm{mv}=6.62 \times 10^{-34} / 9.1 \times 10^{-31} \times 2 \times 10^{3}=0.364 \times 10^{-6} \mathrm{~m}=364 \mathrm{~nm}$
32. (a) when an unsymmetrical molecule is added to unsymmetrical alkene Then negative part of molecule goes to the C -atom which has less number of H -atom
e.g. $\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{HBr} \longrightarrow \mathrm{CH}_{3}-\mathrm{CHBr}-\mathrm{CH}_{3}$
(b)


Eclipse


Staggered
(c) Halogens are ortho directing due to +R effect but deactivating due to effect.

## OR

(a) When HBr is added to an unsymmetrical alkene in presence of peroxide then negative part $(\mathrm{Br})$ goes to the C -atom which has more number of H -atom.
eg $\mathrm{CH}-\mathrm{CH}=\mathrm{CH}_{2}+\mathrm{HBr} \xrightarrow{\text { peroxide }} \mathrm{CH}_{3}-\mathrm{CH}_{2}-\mathrm{CH}_{2} \mathrm{Br}$
(b) when a covalent bond is cleaved in such a way that each atom retain one electron of the shared pair, the fission of called homolytic fission.
(c) $\%$ of Sulphur $=32 \times$ weight of $\mathrm{BaSO}_{4} \times 100 / 233 \times$ weight of organic compound

$$
\begin{aligned}
& =32 \times 0.4813 \times 100 / 233 \times 0.157 \\
& =42.1 \%
\end{aligned}
$$

33. (A)

| Element | \%age | atomic mass | relative no. of <br> moles | whole no.ratio of <br> atoms |
| :--- | :--- | :--- | :--- | :--- |
| Na | 14.28 | 23 | $14.28 / 23=0.62$ | $0.62 / 0.31=2$ |
| S | 9.92 | 32 | $9.92 / 23=0.31$ | $0.31 / 0.31=1$ |
| H | 6.2 | 1 | $6.2 / 1=6.2$ | $6.2 / 0.31=20$ |
| O | 69.6 | 16 | $69.6 / 16=4.35$ | $4.35 / 0.31=14$ |

Empirical formula $=\mathrm{Na}_{2} \mathrm{SH}_{20} \mathrm{O}_{14}$
Empirical formula mass $23 \times 2+32+1 \times 20+16 \times 14-322$
$\mathrm{n}=$ molecular formula mass/ empirical formula mass $=322 / 322=1$
molecular formula $=($ Empirical formula $) \mathrm{n}=\left(\mathrm{Na}_{2} \mathrm{SH}_{20} \mathrm{O}_{14}\right) 1=\mathrm{Na}_{2} \mathrm{SH}_{20} \mathrm{O}_{14}$
since all hydrogen is present in form of water hence molecular formula is $\mathrm{Na}_{2} \mathrm{SO}_{4} \cdot 10 \mathrm{H}_{2} \mathrm{O}$
(b) Because molality is independent of temperature while molarity decrease with increase of temperature.
(c) The reactant which is completely consumed in reaction is called limiting reagent.

OR
(a) Moles of benzene $=30 / 78=0.38$ moles

Moles of $\mathrm{CCl}_{4}=70 / 154=0.45$ moles
Total moles $0.38+0.45=0.83$
Mole fraction of benzene $=$ moles of benzene/ total moles $=0.38 / 0.83=0.45$.
(b) $\mathrm{C}_{2} \mathrm{H}_{2}+5 / 2 \mathrm{O}_{2} \rightarrow \quad 2 \mathrm{CO}_{2}+\mathrm{H}_{2} \mathrm{O}$
$22400 \mathrm{ml} 56000 \mathrm{ml} \quad 44800 \mathrm{ml}$ at NTP

200 ml ? ?
Volume of oxygen required $=56000 \times 200 / 22400=500 \mathrm{ml}$
Volume of $\mathrm{CO}_{2}$ produced $=44800 \times 200 / 22400=400 \mathrm{ml}$

# PRACTICE PAPER- II <br> CLASS - XI <br> CHEMISTRY THEORY (043) 

## Time: 3 Hours

Max. Marks : 70

## General Instructions: Read the following instructions creafully

1. There are 33 questions in the questions paper with internal choice.
2. Section A consists of 16 multiple-choice questions carrying 1 mark each.
3. Section $B$ consists of 5 short answer questions carrying 2 marks each.
4. Section C consists of 7 short answer questions carrying 3 marks each
5. Section D consists of 2 case based questions carrying 4 marks each.
6. Section E consists of 3 long answer questions carrying 5 marks each.
7. All questions are compulsory
8. Use of $\log$ tables and calculators is not allowed.

## SECTION-A

1. One mole of oxygenn gas at STP is equal to $\qquad$ .
(a) $6.022 \times 10^{23}$ molecules of oxygen
(b) $6022 \times 10^{23}$ atoms of oxygen
(c) 16 g of oxygen
(d) 32 g of oxygen
2. In which of the following pairs, the ions are iso electronic?
(a) $\mathrm{Na}^{+}, \mathrm{Mg}^{2+}$
(b) $\mathrm{Al}^{3+}, 0^{-}$
(c) $\mathrm{Na}^{2+}, \mathrm{O}^{2-}$
(d) $\mathrm{N}^{3-}, \mathrm{Cl}^{-}$
3. The radius of which of the following orbit is same as that of first obit of hydrogen atom?
(a) $\mathrm{He}^{+}(\mathrm{n}=2)$
(b) $\mathrm{Li}^{2+}(\mathrm{n}=2)$
(c) $\mathrm{Li}^{2+}(\mathrm{n}=3)$
(d) $\mathrm{Be}^{3+}(\mathrm{n}=2)$
4. Which of the following pairs of d-orbitals have electron density along the axis?
(a) $\mathrm{dz}^{2}, \mathrm{dxz}$
(b) dxz, dyz
(c) $\mathrm{dz}^{2}, \mathrm{dx}^{2} \mathrm{y}^{2}$
(d) dzy, $\mathrm{dx}^{2}-\mathrm{y}^{2}$
5. The electronic configuration of gadolinium (Atomic No.64) is
(a) $[\mathrm{Xe}] 4 \mathrm{f}^{3} 5 \mathrm{~d}^{5} 6 \mathrm{~s}^{2}$
(b) $[\mathrm{Xe}] 4 \mathrm{f}^{7} 5 \mathrm{~d}^{2} 6 \mathrm{~s}^{1}$
(c) $[\mathrm{Xe}] 4 \mathrm{f}^{7} 5 \mathrm{~d}^{1} 6 \mathrm{~s}^{2}$
(d) $[\mathrm{Xe}] 4 \mathrm{f}^{8} 5 \mathrm{~d}^{6} 6 \mathrm{~s}^{2}$
6. The types of hybrid orbitals of nitrogen in $\mathrm{NO}_{2}^{+}, \mathrm{NO}_{3}{ }^{-}$and $\mathrm{NH}_{4}^{+}$respecting are expected to be
(a) $\mathrm{sp}, \mathrm{sp}^{3}$ and $\mathrm{sp}^{2}$
(b) $\mathrm{sp}, \mathrm{sp}^{2}$ and $\mathrm{sp}^{3}$
(c) $\mathrm{sp}^{2}, \mathrm{sp}$ and $\mathrm{sp}^{3}$
(d) $\mathrm{sp}^{2}, \mathrm{sp}^{3}$ and sp
7. $\Delta_{f} \mathrm{U}$ of formation of $\mathrm{CH}_{4}(\mathrm{~g})$ at certain temperature is $-393 \mathrm{~kJ} \mathrm{~mol}^{-1}$. The value of $\Delta_{f} H$ is
(a) zero
(b) $<\Delta_{f} U$
(c) $>\Delta_{f} U$
(d) equal to $\Delta_{\mathrm{f}} U$
8. $0.1 \mathrm{M} \mathrm{CH}_{3} \mathrm{COOH}$ is $1.34 \%$, ionised, calculate its ka
(a) $1.8 \times 10^{-5}$
(b) $1.8 \times 10^{-4}$
(c) $5 \times 10^{-4}$
(d) $4 \times 10^{-5}$
9. Which of the following compound contain all the carbon atoms Same hybridisation state?
(a) $\mathrm{H}-\mathrm{C} \equiv \mathrm{C}-\mathrm{C} \equiv \mathrm{C}-\mathrm{H}$
(b) $\mathrm{CH}_{3}-\mathrm{C} \equiv \mathrm{C}-\mathrm{CH}_{3}$
(c) $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{CH}_{2}$
(d) $\mathrm{CH}_{2}=\mathrm{C}=\mathrm{C}=\mathrm{CH}_{2}$
10. The IUPAC name of the compound H
(a) 5-formyl hex-2-en-3-one
(b) 5-methyl-4- oxohex-2-en-5-al
(c) 3-keto-2 methyl hex-5-enal
(d) 3-keto-2-methyl hex-4-enal
11. Which of the following structure is atomatic?
(a)

(b)

(c)

(d)

12. Which of the following will not show geometrical isomenim 1
(a)

(b)

(c) $\mathrm{CH}_{3}>\mathrm{C}=\mathrm{C}<\mathrm{C}_{2}<\mathrm{C}_{2} \mathrm{H}_{5}$
(b)


In the following questions (Q.13-Q.16), a statement of assertion followed by a statement of reason is given. Choose the Correct answer out of the following choices
(a) Both Assertion \& Reason are true \& Reason is correct explanation of the Assertion
(b) Both Assertion and Reason are true but Reason is not correct explanation of Assertion.
(c) Assertion is a true but Reason in false
(d) Assertion is false but Reason is true
13. Assertion. It is impossible to determine the exact position and exact momentum of an electron simultaneously.
Reason: The path of an electron in an atom is clearly defined.
14. Assertion: Boron has a smaller 1st Ionisation enthalpy than Beryllium

Reason: The penetration of a 2 s electron to the nucleus is more than the $2 p$ electron hence 2 p electron is more shielded by the inner core of electron than the 2 s electron.
15. Assertion: Heat capacity is the amount of heat required to raise temperature of body by 1 K .
Reason: Heart capacity is an extensive property and it depends upon the size of the body.
16. Assertion: Black body is an ideal body that units and absorb all radiations. Reason: The frequency of radiation emitted by a body goes from a lower frequency to higher frequency with an increase in temperature.

## SECTION B

This section antains 5 questions with internal choice in one question. The following questions are very short answer type and carry 2 mark each.
17. (i) Write the significant figures in the following:
(a) 2.00
(b) 4.8046
(ii) Write the electronic configuration of $\mathrm{Cu}(29)$.
18. How will you convert following:
(i) Propene to 2, 3-Dimethyl butane
(ii) Ethyne to Ethanal
19. If 50 g of $\mathrm{CaCO}_{3}$ is treated with 50 g of HCl , how many grams of $\mathrm{CO}_{2}$ can be produced according to following equations:
$\mathrm{CaCO}_{3}(\mathrm{a})+2 \mathrm{HCl}(\mathrm{dil}) \longrightarrow \mathrm{CaCl}_{2}(\mathrm{aq})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
20. The wavelength of 1 st spectral line in the Balmer series is $6561 \mathrm{~A}^{\circ}$. Calculate the wavelength of the $2^{\text {nd }}$ spectral line in Balmer series.

Or
The radius 1st Bohr's orbit of hydrogen atom is $0.529 \mathrm{~A}^{\circ}$. Calculate the radius of-
(i) the $3^{\text {rd }}$ orbit of $\mathrm{He}^{+}$ion
(ii) the $2^{\text {nd }}$ orbit of $\mathrm{Li}^{2+}$ ion
21. What are values of ' $n$ ' \& 'I' for valence election of $\mathrm{Na}(11)$.

## SECTION C

This section contains 7 questions with internal choice in one question. The following questions are short answer type and carry 3 marks each.
22. (i) Define Markonikov's Rule. Illustrate with the help of an example.
(ii) Distinguish b/w 1-Butyne \& 1-butene
(iii) Write all possible isomers of $\mathrm{C}_{4} \mathrm{H}_{9} \mathrm{Br}$.
23. (i) State Heisenbergs uncertainty principle.
(ii) Calculate the uncertainity in the position of an election if uncertainty in its velocity is $0.001 \%$.

$$
\left(\mathrm{m}=9.1 \times 10^{-31} \mathrm{~kg}, \mathrm{~h}=6.63 \times 10^{-34} \mathrm{Js}\right)
$$

24. (i) Identify the oxidising and reducing agent int he following reaction

$$
\mathrm{H}^{+}+\mathrm{H}_{2} \mathrm{O} \longrightarrow \mathrm{OH}^{-}+\mathrm{H}_{2}
$$

(ii) Balance the following equation-
$\mathrm{MnO}_{4}^{-}+\mathrm{C}_{2} \mathrm{O}_{4}^{2-}+\mathrm{H}^{+} \longrightarrow \mathrm{Mn}^{2+}+\mathrm{CO}_{2}+\mathrm{H}_{2} \mathrm{O}$
(iii) What is the oxidation state of Fe in $\mathrm{K}_{4}\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]$ ?
25. (i) Define molar heat capacity
(ii) The enthalpy of combustion of benzoic acid at 298 K and 1 atm is $-2546 \mathrm{~kJ} \mathrm{~mol}^{-1}$. What is $\Delta \mathrm{U}$ of the reaction)
26. (i) State Hund's Rule
(ii) Calculate the wavelength associated with a Proton moving with $1.0 \times$ $10^{3} \mathrm{~ms}^{-1}$. (mass of proton $=1.67 \times 10^{-27} \mathrm{~kg}, \mathrm{~h}=6.63 \times 10^{-34} \mathrm{Js}$ ]

## Or

(i) Why are Bohr's orbits called stationary state?
(ii) Which series of hydrogen spectrum lies in UV region?
(ii) Which of the following have highest frequency.

X-rays, $\gamma$-rays, Microware
27. A compound is made up of 2 elements $A \& B$ has $A=70 \%$., $B=30 \%$. Their relative number of moles in the compound is $1.25 \& 1.88$. Calculate:
(i) Atomic Masses of the elements A \& B.
(ii) Molecular formula of the compound if molecular mass of compound is 160u.
28. (i) Define Le-chatelier's Principle
(ii) The Equilibrium constant for the reaction $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{Br}(\mathrm{g}) \rightleftharpoons 2 \mathrm{HBr}$ at 1024 k is $1.6 \times 10^{5}$. Find the equilibrium pressure of all gases if 10.0 bar of HBr is introduced into sealed container at 1024 k .

## SECTION D

The following questions are case based questions. Each question has an internal choice and carries 4 marks each. Read the passage carefully and answer the following questions.
29. To locate the address of a person, we require name, house number, city State, pin code. Similarly the electron in an atom can be located with the help of quantum numbers. These quantum numbers describe the energy level of an orbital and define the shape and orientation of the region in space where the election will be found. The principal, azimuthal and magnetic quantum numbers come from the solution of schro-dinger wave equation. The fourth quantum number called spin quantum number represents the spin of the election about its own axis. Answer the following questions.
(a) How many electrons in the ground state of neon have $1+\mathrm{m}_{1}=0$ (Atomic No. of $\mathrm{Ne}=10$ )
(b) Which quantum number distinguish between the 2 electrons present in the same obital
(c) Predict whether the following set of quantum numbers can exist or not for an electron.
(i) $\mathrm{n}=1, \mathrm{l}=0, \mathrm{~m}_{1}=0, \mathrm{~ms}=-1 / 2$
(ii) $\mathrm{n}=5, \mathrm{l}=3, \mathrm{~m}_{1}=-4, \mathrm{~ms}=+1 / 2$

OR
What will be the maximum possible, no of electrons having $\mathrm{ms}=-1 / 2$ for n $=5$ ?
30. The identity of a substance is defined not only by the types of atoms or ions it contains, but by the quantity of each type of atom or ion. The expermental approach required the introduction of new unit for amount of substances, the mole, which remains indispensable in modern chemical science. The mole is an amount unit similar to unit like pair, dozon. gross, etc. It provides a specific measure of the number of atoms of molecules in a bulk sample of matter. A mole is defined as the amount of substance containing the same number of discrete entities (atoms, molecules, ions etc.) as the number of atoms in a sample of pure 12 ( weighing exactly 12 g one latin connotation for the word "mole" is large mass") or bulk" which is consistent with its use as the name for this unit. The mole provides a link between an easily measured macroscopic propety, bulk mass and an extremely important fundamental property, number of atoms molecules and so forth. The number of entities composing a mole has been experimentally determined to be $6.022 \times 10^{23}$. $6.022 \times 10^{23}$, a fundamental constant named Avogaro's number (NA) or the Avogadro constant in honor of Italian scientist Amedco Avogactor. This constant is properly reported with an explicit unit of "per mole", a conveniently rounded version being $6.022 \times 10^{23} / \mathrm{mol}$. Consistent with its definition as an amount unit, 1 mole of any element contains the same number of atoms as 1 mole of any other element. The masses of 1 mole of different elements, however, are different since the masses of the individual atoms are drastically different, the molar mass of an element (or compound) is the mass in grams of 1 mole of that substance, a property expressed in units of grams per mole ( $\mathrm{g} / \mathrm{mol}$ ).
(a) When an antacid tablet is used, $\mathrm{Ca}(\mathrm{OH})_{2}$ reacts with HCl in the stomach to form inert $\mathrm{CaCl}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$. If the molar mass of $\mathrm{Ca}\left(\mathrm{OH}_{2}\right)$ is $75 \mathrm{~g} /$ mol, how many moles of HCl are required to fully react with 150 g of $\mathrm{Ca}(\mathrm{OH})_{2}$.
(b) Calculate no. of atoms of He present in 100u.
(c) How are $0.5 \mathrm{~mol} \mathrm{Na}_{2} \mathrm{CO}_{3}$ and $0.5 \mathrm{M} \mathrm{Na}_{2} \mathrm{CO}_{3}$ different from each other?

OR
At STP, what will be the volume of $6.022 \times 10^{23}$ molecules of $\mathrm{H}_{2}$.

## SECTION E

Following questions are long answer type and carry 5 marks each. All question have an internal choice.
31. (a) Equilibrium constant for a reaction is 100 . What will be equilibrium constant for reverse reaction?
(b) Write the conjugate acids for bronsted base $\mathrm{OH}^{-} \& \mathrm{CH}_{3} \mathrm{COO}^{-}$
(c) Determine the pH of $10^{-8} \mathrm{M} \mathrm{HCl}$ solution taking into account the $\mathrm{H}+$ produced by water also. $(\log 11=1.04 \mathrm{M})$

OR
(a) What is effect of temperature $m$ value of Ksp ?
(b) Calculate pH of buffer solution Containing 0.01 M solution of $\mathrm{NH}_{4} \mathrm{OH}$ and 0.1 M solution of $\mathrm{NH}_{4} \mathrm{Cl}$. pKb of $\mathrm{NH}_{4} \mathrm{OH}=4.75$.
(c) Classify the following into Acidic and Basic Salt. $\mathrm{CH}_{3} \mathrm{COONa}, \mathrm{NH}_{4} \mathrm{Cl}, \mathrm{Na}_{2} \mathrm{CO}_{3}, \mathrm{CuSO}_{4}$
32. (a) Propanal and Propan-2-one are the ozonolysis products of an alkene what is the structural formula of the alkene.
(b) Give the major products of the reactions
(i)

(ii)

(iii)

(a) An alkyl halide 'A' of the formula $\mathrm{C}_{6} \mathrm{H}_{13} \mathrm{Cl}$ on treatment alc. KOH give two isomeric alkene $(\mathrm{B}) \&(\mathrm{C})$ with formula $\mathrm{C}_{6} \mathrm{H}_{12}$. Both alkenes on hydrogenation give 2,3-Dimethylbutane Predict the structures of ' A ', ' B ' and ' C '.
(b) How is benzene obtained from ethyne! What is the name of the process?
(c) Convert 1-Propanol to 2-propanol.
33. (a) (i) Use Molecular orbital theory to predict why $\mathrm{Be}_{2}$ molecule does nto exist.
(ii) Compare the stability of $\mathrm{N}_{2}^{+}$and $\mathrm{N}_{2}$.
(b) (i) Why is bond length of all $\mathrm{C}=\mathrm{O}$ bonds is equal in carbonate ion)
(ii) Why dipole moment of $\mathrm{BF}_{3}$ is zero but $\mathrm{NF}_{3}$ is not?
(iii) Why does water have boiling point higher than HF?

## PRACTICE PAPER- III <br> CLASS - XI <br> CHEMISTRY THEORY (043)

Time: 3 Hours
Max. Marks : 70

## General Instructions: Read the following instructions carefully

1. There are 33 questions in the questions paper with internal choice.
2. Section A consists of 16 multiple-choice questions carrying 1 mark each.
3. Section B consists of 5 short answer questions carrying 2 marks each.
4. Section C consists of 7 short answer questions carrying 3 marks each
5. Section D consists of 2 case based questions carrying 4 marks each.
6. Section E consists of 3 long answer questions carrying 5 marks each.
7. All questions are compulsory
8. Use of $\log$ tables and calculators is not allowed.

## SECTION-A

The following questions are multiple choice questions with one correct answer. Each question carries 1 mark there is no internal choice in this section.

1. Which has maximum number of atoms
(a) 24 g of $\mathrm{C}-12$
(b) 56 g of $\mathrm{Fe}-56$
(c) 27 g of $\mathrm{Al}-27$
(d) 108 g of $\mathrm{Ag}-108$
2. The maximum number of electron in a sub-shell with $1=3$ and $\mathrm{n}=4$ is.
(a) 14
(b) 16
(c) 10
(d) 12
3. Which of the following elements has highest first ionisation energy.
(a) Rb
(b) Na
(c) k
(d) Sc
4. The hydration energy of $\mathrm{Mg}^{2+}$ ion is larger than that of
(a) $\mathrm{Al}^{3+}$
(b) $\mathrm{Na}^{+}$
(c) $\mathrm{Be}^{2+}$
(d) $\mathrm{Ca}^{2+}$
5. Which of the following species is not paramagnetic
(a) CO
(b) $\mathrm{O}_{2}$
(c) $\mathrm{B}_{2}$
(d) NO
6. The species in which the N atom is in a state of sp hybridisation is:
(a) $\mathrm{NO}^{2+}$
(b) $\mathrm{NO}_{2}^{-}$
(c) $\mathrm{NO}_{3}^{-}$
(d) $\mathrm{NO}_{2}$
7. The concentration of hydrogen in a sample of soft drink is $3.8 \times 10^{-3} \mathrm{M}$. The pH of the solution is:
(a) 3.84
(b) 2.42
(c) 4.44
(d) 1.42
8. The oxidation state of 's' in $\mathrm{KAl}\left(\mathrm{SO}_{4}\right)_{2} .12 \mathrm{H}_{2} \mathrm{O}$.
(a) -2
(b) -1
(c) +2
(d) +6
9. Which of the following can act as an electrophile.
(a) $\mathrm{CN}^{-}$
(b) $\mathrm{OH}^{-}$
(c) $\mathrm{H}_{2} \mathrm{O}$
(d) $\mathrm{BF}_{3}$
10. The most stable free radical among the following is:
(a) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{2} \dot{\mathrm{C}} \mathrm{H}_{2}$
(b) $\mathrm{C}_{6} \mathrm{H}_{5} \dot{\mathrm{C}} \mathrm{HCH}_{3}$
(c) $\mathrm{CH}_{3} \dot{\mathrm{C}} \mathrm{H}_{2}$
(d) $\mathrm{CH}_{3} \mathrm{CHCH}_{3}$
11. Action of acetylene on dilute $\mathrm{H}_{2} \mathrm{SO}_{4}$ gives
(a) Acetic acid
(b) Acetone
(c) Acetaldehyde
(d) Ethyl alcohol
12. $\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{3}+\mathrm{O}_{3} \xrightarrow[\mathrm{H}_{2} \mathrm{O}]{\mathrm{Zn}} \mathrm{A}$. A in the following reaction is:
(a) $\mathrm{CH}_{3} \mathrm{COCH}_{3}$
(b) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CHO}$
(c) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{COCH}_{3}$
(d) $\mathrm{CH}_{3} \mathrm{CHO}$

Question no. 13-16 are assertion reason based questions select the most appropriate answer from the options given below:
(a) Both A and R are true and R is the correct explanation of A .
(b) Both A and R are true but R is not the correct explanation of A .
(c) A is true but R is false.
(d) A is false but R is true.
13. Assertion: Empirical and molecular formulea of $\mathrm{NaHCO}_{3}$ are the same.

Reason: Upon heating $\mathrm{NaHCO}_{3}$ evolves $\mathrm{CO}_{2}$ gas.
14. Assertion: The spectrum of $\mathrm{He}^{+}$ion is expected to be similar to that of H atoms.

Reason: $\mathrm{He}^{+}$ion is also one electron species.
15. Assertion: $\mathrm{ClF}_{3}$ has T -shape structure

Reason: It has two lone pairs arranged at angle of 180.
16. Assertion: Addition of $q$ and $w$ gives $\Delta U$.

Reason: Addition of two path functions can not give state function.

## SECTION-B

This section contains 5 question with internal choice in one question. The following questions are very short answer types and carry 2 marks each.
17. Identify A and B in the following reaction
$\mathrm{A} \xrightarrow{\mathrm{Na}} \mathrm{CHC} \equiv \mathrm{CH} \xrightarrow[\text { Fe tube } 873 \mathrm{k}]{\text { Red not }} \mathrm{B}$.
OR
Write the structures of the products A and B of the following reaction:

18. Give IU PAC name of th following compound?
(i) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CHCH}_{2} \mathrm{CHO}$
(ii) $\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}-\mathrm{COOH}$
19. Write two functions of salt bridge in an eletrochemical cell.
20. Out of $\mathrm{NH}_{3}$ and $\mathrm{NF}_{3}$ which is more polar. Explain with the help of dipole moment.
21. Calculate the number of moles in the following masses:
(a) 7.85 g of Fe
(b) 7.9 g of Ca

## SECTION-C

This section contains 7 questions with internal choice in one question. The following questions are short Ans type and carry 3 marks each.
22. 3.0 g of $\mathrm{H}_{2}$ react with 29.0 g of $\mathrm{O}_{2}$ yield $\mathrm{H}_{2} \mathrm{O}$
(i) Which is the limiting reagent
(ii) Calculate the maximum amount of $\mathrm{H}_{2} \mathrm{O}$ that can be formed.
(iii) Calculate the amount of reactant left unreacted.
23. Electrons are emitted with zero velocity from a metal surface when it is exposed to radiation of wavelength 6800 A . Calculate threshold frequency $\left(v_{0}\right)$ and work function $\left(\mathrm{W}_{0}\right)$ of the metal.
24. Among the second period elements the actual ionisation enthalpies are in the order.
$\mathrm{Li}<\mathrm{B}<\mathrm{Be}<\mathrm{C}<\mathrm{O}<\mathrm{N}<\mathrm{F}<\mathrm{Ne}$. Explain why
(a) Be has higher $(\Delta \mathrm{iH})_{1}$ than B
(b) O has lower $(\Delta \mathrm{iH}) 1$ than N and F ?
25. Explain the structure of $\mathrm{PCl}_{5}$ according to hybridization. Why all $\mathrm{P}-\mathrm{Cl}$ bonds lengths are not equivalent in $\mathrm{PCl}_{5}$.
26. Calculate the pH of a solution obtained by mixing 50 ml of 0.2 M HCl with 49.9 ml of 0.2 M NaOH solution.
27. Account for the following:
(i) $\mathrm{HNO}_{3}$ acts as oxidizing agent while $\mathrm{HNO}_{2}$ can act both as reducing and oxidizing agent.
(ii) $\mathrm{AgF}_{2}$ is unstable compound and act as a strong oxidizing agent
(iii) Ozone act as an oxidising agent
28. State mark ownikov's rule. Using this rule write the reaction of propene with (i) HCl (ii) $\mathrm{H}_{2} \mathrm{O}$
Complete the following reactions:
(a) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br} \xrightarrow[\mathrm{KOH}]{\mathrm{Alc}}$
(b) $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}+\mathrm{O}_{3} \xrightarrow[\mathrm{H}_{2} \mathrm{O}]{\mathrm{Zn}}$
(c) $\mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{H}_{2} \mathrm{O}+(\mathrm{O}) \xrightarrow{\mathrm{KMnO}_{4}}$

## SECTION-D

The following questions are case based each question has an internal choice and carries $4(1+1+2)$ marks. Read the passage carefully and answer the questions that follow.
29. Heat of neutralization is defined as amount of heat released when one grams equivalent of a strong base reacts with one gram equivalent of strong acid the heat of neutralization is cause out to be as follows:
$\mathrm{H}^{+}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq}) \longrightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{aq})$
$\Delta \mathrm{H}(\mathrm{Neu})=\mathrm{X}$
(i) If enthalpy of neutialization of $\mathrm{CH}_{3} \mathrm{COOH}$ by HCl is y then enthalpy of ionization of $\mathrm{CH}_{3} \mathrm{COOH}$ acid will be.
(ii) What is the enthalypy change for a complete neutralization reaction of stong base $\mathrm{R}(\mathrm{OH})_{2}$ by HCl
(iii) When the conditions are identical how many ml of 0.1 M of strong base A OH and 0.05 M strong $\mathrm{H}_{2} \mathrm{~A}$ acid solution should be mixed for a total volume of 100 ml produce the highest rise in temperature.

OR
If heat of neutralization of $\mathrm{HA}, \mathrm{HB}, \mathrm{HC}, \mathrm{HD}$ acid follows the order $\mathrm{HA}>$ $\mathrm{HB}>\mathrm{HC}>\mathrm{HD}$ then among the given acid which one is strongest acid.
30. Presence of double bond in an alkene is tested by reacting with either water or with dilute solution of Bayer's reagent. The position of double bond is located with the help of ozonalysis it is done by Joining together the product of ozonalysis which are carbonyl compound at their carbonyl carbon atoms by double bond one mole of ozone is used in the ozonalysis reaction per mole of double bond in a particular alkene.
(i) The alkene $\mathrm{C}_{6} \mathrm{H}_{10}$ producing $\mathrm{OHC}\left(\mathrm{CH}_{2}\right)_{4} \mathrm{CHO}$ on Ozonalysis is:
(ii) Complete the following reaction

$$
\mathrm{C}_{2} \mathrm{H}_{5}-\mathrm{CH}=\mathrm{CH}-\mathrm{C}_{2} \mathrm{H}_{5} \xrightarrow\left[\left(\text { (i) } \mathrm{O}_{3}\right]{\mathrm{Zn}_{1} \mathrm{H}_{2} \mathrm{O}}\right.
$$

(iii) Alkene give two moles of $\mathrm{CH}_{3} \mathrm{CHO}$ on ozonalysis give the structure and IU PAC name of alkene.

OR
A alkene give $\mathrm{CH}_{3} \mathrm{CHO}$ and $\mathrm{CH}_{3} \mathrm{COCH}_{3}$ on ozonlyasis give the structure and IUPAC name of alkene.

## SECTION E

The following questions are long answer type are carry 5 marks each. All questions have an internal choice.
31. A welding fuel gas contains carbon and hydrogen only. Burning 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 in found to weight 11.6 g . Calculate.
(i) Empirical formula
(ii) Molar mass of the gas and
(iii) Molecular formaula.

OR
(i) What is difference between molarity and molality
(ii) The molarity of a solution of sulphuric acid is 1.5 M . Calculate its molality (given density of solution is $1.029 \mathrm{gml}^{-1}$ )
32. (a) Define photoelectric effect. Mention its one practical application in daily life.
(b) The work function for Cs atom is 1.9 eV . Find threshold wavelength $(\lambda 0)$ and threshold frequency $(\mathrm{vo})$ of this light radiation. If Cs metal is irradiated with radiation of wavelength 500 nm find kinetic energy and velocity of emitted electrons.

## OR

(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 $\mathrm{He}^{+}$and $\mathrm{Li}^{2+}$ ions.
33. (i) Calculate the solubility of silver chloride in water at room temperature if the $\mathrm{K}_{\text {sp }}$ of AgCl is $1.6 \times 10^{-10}$.
(ii) Calculate the molar solubility of $\mathrm{Ni}(\mathrm{OH})_{2}$ in 0.10 M NaOH . The ionic product of $\mathrm{Ni}(\mathrm{OH})_{2}$ is $2 \times 10^{-15}$.

OR
(i) The molar solubility of $\mathrm{Cd}(\mathrm{OH})_{2}$ is $1.84 \times 10^{-5} \mathrm{M}$. Calculate the expected solubility of $\mathrm{Cd}(\mathrm{OH})_{2}$ in a buffer solution of $\mathrm{pH}=12$
(ii) Calculated the pH of a solution obtained by mixing 50 ml of 0.2 M HCl with 49.9 mL of 0.2 M NaOH solution.

## COMMON ANNUAL SCHOOL EXAMINATION

## General Instructions:

1. All questions are compulsory.
2. This question paper consists of 35 questions divided into five sections $\mathrm{A}, \mathrm{B}, \mathrm{C}, \mathrm{D}$ and E .
3. Section A: Q1 to Q18 are objective type questions. Carrying 1 mark each.
4. Section B: Q19 to Q25 are very short answer type questions carrying 2 marks each.
5. Section C: Q26 to Q30 are short answer type questions carrying 3 marks each.
6. Section D: Q31 to Q32 are of passage/case study based questions carrying 4 marks each.
7. Section E: Q33 to Q35 are long answer type questions carrying 5 marks each.
8. There is no overall choice in question paper, only internal choices are available.
9. Use of calculator and $\log$ tables is not permitted.

## SECTION-A

The following questions (Q. No.1-14) are multiple choice questions with one correct answer. Each question carries 1 mark. There is no internal choice in this section.

1. Which of the following terms are unitless?
(a) Molality
(b) Molarity
(c) Mole fraction
(d) Density
2. One of the statements of Dalton's atomic theory is given below:
"Compounds are formed when atoms of different elements combine in fixed ratio."

Which of the folowing law is related to this statement?
(a) Law of conservation of mass
(b) Law of definite proportion
(c) Gay Lucssa's law of gaseous volumes
(d) Avogadro law
3. The number of radial nodes for $3 p$ orbital is:
(a) 3
(b) 4
(c) 6
(d) 1
4. Amongst halogens, which one has highly negative electron gain enthapy?
(a) F
(b) Cl
(c) Br
(d) 1
5. Which of the following species has distorted tetrahedral geometry?
(a) $\mathrm{CO}_{2}$
(b) $\mathrm{CO}_{3}^{2-}$
(c) $\mathrm{NH}_{3}$
(d) $\mathrm{SF}_{4}$
6. A reaction $\mathrm{A}+\mathrm{B} \rightarrow \mathrm{C}+\mathrm{D}+\mathrm{q}$ is found to have a postive entropy change. The reaction will be:
(a) possible at the temperature
(b) possible only at low temperature
(c) not possible at any temperature
(d) possible of any temperature
7. $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2(\mathrm{~g})} \rightarrow 2 \mathrm{NH}_{3(\mathrm{~g})} ; \Delta \mathrm{rH}^{\circ}=-92.4 \mathrm{~kJ} \mathrm{~mol}^{-1}$

What is the standard enthalpy of formation of $\mathrm{NH}_{3}$ gas?
(a) $-46.2 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(b) $46.2 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(c) $-92.4 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(d) $92.4 \mathrm{~kJ} \mathrm{~mol}^{-1}$
8. Acidity of $\mathrm{BF}_{3}$ can be explained on the basis of which of the following concepts?
(a) Arrhenius concept
(b) Bronsted Lowry concept
(c) Lewis Concept
(d) Bronsted Lowry as well as Lewis concept
9. At 500 K , equlibrium constant, $\mathrm{K}_{\mathrm{c}}$, for the following reaction is 25:

$$
\mathrm{H}_{2(\mathrm{~g})}+\mathrm{I}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{HI}_{(\mathrm{g})}
$$

then what is the value of equilibrium constant $\mathrm{K}_{\mathrm{c}}$ for the reverse reaction of above process at same temperatrue?
(a) 0.4
(b) 0.04
(c) 0.25
(d) 625
10. What is the oxidation number of Mn in $\mathrm{KMnO}_{4}$ ?
(a) +2
(b) +3
(c) +5
(d) +7
11. The oxidation state/states exhibited by hydrogen in its various compounds are:
(a) -1 only
(b) +1 only
(c) 0 only
(d) $+1,-1$, and 0
12. Which of the following species is distorted tetrahedral in shape?
(a) Free radical
(b) Carbanion
(c) Carbocation
(d) None of these
13. The IUPAC name for $\mathrm{CH}_{3} \mathrm{CCH}_{2} \mathrm{CH}_{2} \mathrm{C}-\mathrm{OH}$ is $\qquad$ .
(a) 1-hydroxypentane-1, 4-dione
(b) 1,4-dioxopentanol
(c) 1-carboxybutan-3-one
(d) 4-oxopentanoic acid
14. Which of the following has zero dipole moment?
(a) But-1-ene
(b) Cis-But-2-ene
(c) trans-But-2-ene
(d) 2-Methylprop-1-ene

In the following questions, a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices:
(a) Assertion and reason both are correct and reason is correct explanation for assertion.
(b) Assertion and reason both are correct but reason is not correct explanation for assertion.
(c) Assertion is correct but reason is wrong statement.
(d) Assertion is wrong but reason is correct statement.
15. Assertion : The last electron in potassium atom enters into 4 s orbital, not 3 d orbital.

Reason: $(\mathrm{n}+l)$ rule is followed for determining the subshell lowest energy state.
16. Assertion : Assertion: Fluorine is the most electronegative element in the periodic table.
Reason: In the periodic table, electronegativity generally decreases on moving from left to right across a period.
17. Assertion : $\mathrm{NF}_{3}$ and ${\mathrm{C} / \mathrm{F}_{3}}$ do not have similar shapes.

Reason: The central atom in these two molecules have different electronegativities.
18. Assertion : Among isomeric pentanes, 2,2-dimethylpentane has lowest boiling point.
Reason: Branching does not affect the boiling point.

## SECTION-B

The following questions (Q. No.19-25) are very short answer type questions, carrying 2 marks each.
19. (a) State law of multiple proportions.
(b) How are $0.5 \mathrm{Na}_{2} \mathrm{CO}_{3}$ and $0.5 \mathrm{M} \mathrm{Na}_{2} \mathrm{CO}_{3}$ different?
20. (a) Draw the shape of that d-orbital which does not have four lobes?
(b) How many subshells are associated with $\mathrm{n}=3$ ?

OR
Write the electronic configuration of $\mathrm{Cr}(\mathrm{z}=24)$. Why is it different from the expected configuration?
21. (a) State Pauli's exclusion principle.
(b) Write relationship between energy of photon and wavelength of radiation.
22. Give reasons for the following:
(a) Cations are smaller in radii than their parent atoms.
(b) Boron has smaller first ionization enthalpy than beryllium.

OR
(a) What do you understand by isoelectronic species?
(b) Write a cation and anion which is isoelectronic with Ar?
23. Among the elements of third period from Na to Ar, pick out the element:
(a) With highest first ionization enthalpy.
(b) With biggest atomic radius.
24. Balance the following equation:

$$
\begin{aligned}
& \mathrm{Zn}_{\text {(s) }}+\mathrm{NO}_{3_{(\text {(aq) }}^{-}} \rightarrow \mathrm{Zn}^{2+}{ }_{\text {(aq) }}+\mathrm{NH}_{4(\mathrm{aq)}}^{+}+\mathrm{H}^{+}{ }_{\text {(aq) }} \\
& \text { (Acidic medium) }
\end{aligned}
$$

25. Draw sawhorse projections for the eclipsed and staggered conformations of ethane. Which of these conformation is more stable?
26. $\mathrm{CaCO}_{3}$ reacts with aqueous HCl to give $\mathrm{CaCl}_{2}$ and $\mathrm{CO}_{2}$ according to reaction: What mass of $\mathrm{CaCO}_{3}$ is required to react completely with 25 mL of 0.75 M HCl ?
(Atomic mass : $\mathrm{Ca}=40 \mathrm{u}, \mathrm{C}=12 \mathrm{U}, \mathrm{O}=16 \mathrm{U}, \mathrm{H}=1 \mathrm{u}, \mathrm{Cl}=35.5 \mathrm{u}$ )
27. (a) State Heisenberg's Uncertainty Principle.
(b) It the position of the electron is measured within an accuracy of $\pm 0.002$ nm , cacluate the uncertainty in the momentum of electron. (Planck's constant $=6.2626 \times 10^{-34} \mathrm{Js}$ ).
28. For the reaction $2 \mathrm{~A}_{(\mathrm{g})}+\mathrm{B}(\mathrm{g}) \rightarrow \mathrm{D}(\mathrm{g}) \Delta \mathrm{U}^{\circ}=-10.5 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{\circ}=-44.1 \mathrm{JK}^{-1}$. Calculate $\Delta \mathrm{G}^{\circ}$ for the reaction, and predict whether the reaction may occur spontaneously.
29. (a) 0.378 g of an organic compound gave 0.5740 g of silver chloride in carious estimation. Calculate the percentage of chlorine in the compound.
(b) Write the principle of chromatography.
30. (a) Sodium salt of which acid will be needed for the preparation of propane when heated with sodalime? Write chemical equation for the reaction.
(b) Write the mechanism of the following reaction:


OR
Explain the following :
(a) Trans-but-2-ene has higher melting point than cis-isomer.
(b) More substituted alkene is more stable.
(c) $\mathrm{C}-\mathrm{C}$ bond length in benzene ring is 139 pm which is in between $\mathrm{C}-\mathrm{C}$ single bond ( 154 pm ) and $\mathrm{C}=\mathrm{C}$ double bond (133 pm).

## SECTION-D

The following questions 31 and 32 case based questions.
31. Read the passage and answer the following questions:

Chemical energy stored by molecules can be released as heat during chemical reactions. When a fuel like methane, cooking gas or coal burns in air. The chemical energy may also be used to do mechanical work when a fuel burns in an engine or to provide electrical energy through a galvanic cell like dry cell. Thus, various forms of energy are interrelated and under certain conditions, these may be transformed from one form into another.
Thermodynamics deals with energy changes in chemical or physical processes and enables us to study these changes quantitatively and to make useful predictions. For these purposes, we divide the universe into the system and the surroundings. Chemical or physical processes lead to evolution or absorption of heat (q), part of which may be converted into work (w). These quantities are related through the first law of thermodynamics.
(a) State first law of thermodynamics.
(b) Heat (q) and work done (w) individually are not state functions but their sum is always a state function. Explain why?
(c) In a process, 701 J of heat is absorbed by a system and 394 J of work is done by the system. What is the change in internal energy for the process?

## OR

Give one point to differentiate the following thermodynamic terms:
(a) Extensive properties and internsive properties.
(b) Isothermal process and isobaric process.
31. Read the passage and answer the questions follow:

The general reaction is depicted as follows:


Substrate is that reactant which supplies carbon to the new bond and the other reactant is called reagent. If both the reactants supply carbon to the new bond then choice is arbitrary and in that case the molecule on which attention is focussed is called substrate.

In such a reaction a covalent bond between two carbon atoms or carbon and some other atom is broken and a new bond is formed. A sequential account of each step, describing details electron movement, energetics during bond cleavage and bond formation, and the rates of transformation of reactants into products (kinetics) is referred to as reaction mechanism.
(a) Identify the reagent shown underlined as electrophile or nucleophile:

$$
\mathrm{CH}_{3} \mathrm{COOH}+\underline{\mathrm{OH}}^{-} \rightarrow \mathrm{CH}_{3} \mathrm{COO}^{-}+\mathrm{H}_{2} \mathrm{O}
$$

(b) What is the state of hybridisation of underlined carbon atom in $\mathrm{CH}_{3} \mathrm{CH}=\mathrm{CH}_{2}$ ?
(c) Which of the two. $\mathrm{O}_{2} \mathrm{NCH}_{2} \mathrm{CH}_{2} \mathrm{O}^{-}$or $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{O}^{-}$is expected to be more stable and why?
Arrange the following:

## OR

(i) $-1,-\mathrm{Br},-\mathrm{C},-\mathrm{F}$ [Decreasing order of -I effect]
(ii) $\left(\mathrm{CH}_{3}\right)_{3} \dot{\mathrm{C}}, \mathrm{CH}_{3} \dot{\mathrm{C}} \mathrm{H}_{2}, \mathrm{CH}_{3} \dot{\mathrm{C}} \mathrm{HCH}_{3}, \dot{\mathrm{C}} \mathrm{H}_{3}$ [Increasing order of stability] (Section-E)

Questions 33 to 35 are long answer type questions carrying 5 marks each.
33. (a) Define Le-Chatelier principle. What is the effect of (i) addition of $\mathrm{H}_{2}$, (ii) removal of CO on the equilibrium:

$$
2 \mathrm{H}_{2(\mathrm{~g})}+\mathrm{CO}_{(\mathrm{g})} \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}_{(\mathrm{g})}
$$

(b) Equal volumes of 0.002 M solutions of sodium iodate $\left(\mathrm{NaO}_{3}\right)$ and 0.002 M cupric chlorate $\mathrm{Cu}\left(\mathrm{ClO}_{4}\right)_{2}$ are mixed together. Will it lead to precipitation of cupric iodate $\left[\mathrm{Cu}\left(\mathrm{IO}_{3}\right)_{2}\right]$ ?

$$
\begin{gathered}
\left(\mathrm{K}_{\mathrm{SP}}\left[\mathrm{Cu}\left(\mathrm{IO}_{3}\right)_{2}=7.4 \times 10^{-8}\right]\right. \\
\mathrm{OR}
\end{gathered}
$$

(a) Determine pH of $10^{-8} \mathrm{M} \mathrm{HCl}$ solution. $(\log 11=1.0414)$
(b) Explain common ion effect with example.
(c) Write the conjugate acids for the Bronsted base $\mathrm{OH}^{-}$and $\mathrm{CH}_{3} \mathrm{COO}^{-}$.
34. (a) Account for the following:
(i) $\mathrm{NH}_{3}$ has higher dipole moment than $\mathrm{NF}_{3}$ although $\mathrm{N}-\mathrm{F}$ bond is more polar than $\mathrm{N}-\mathrm{H}$ bond.
(ii) Bond angle of water is $104.5^{\circ}$.
(b) Compare the relative stability of the following species on the basis of molecular orbital theory and indicate their magnetic properties:

$$
\mathrm{O}_{2}^{+}, \mathrm{O}_{2}, \mathrm{O}_{2}^{2-}
$$

35. (a) Propanal and pentan-3-one are the ozonolysis product of an alkene. What is the structural formula of the alkene?
(b) How can you convert the following:
(i) Phenol to nitrobenzene
(ii) 2-bromopropane to 1-bromopropane
(c) Give a chemical test to distinguish between ethene and ethane.

OR
Identify $\mathrm{A}, \mathrm{B}, \mathrm{C}, \mathrm{D}$ and E in the following sequence of reactions :


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[^0]:    Note: The Article 51A containing Fundamental Duties was inserted by the Constitution (42nd Amendment) Act, 1976 (with effect from 3 January 1977).
    *(k) was inserted by the Constitution (86th Amendment) Act, 2002 (with effect from 1 April 2010).

