SCIENCES (52)

CHEMISTRY

SCIENCE Paper - 2

Aims:

1) To acquire the knowledge of terms, concepts, processes, techniques and principles related to the subject.

2) To develop the ability to apply the knowledge of contents and principles of chemistry in unfamiliar situations.

3) To acquire skills in proper handling of apparatus and chemicals.

4) To develop scientific temper, attitude and problem solving skills.

5) To recognize Chemical Science as having an important impact on the environment relating to cycles in nature; natural resources, pollution.

CLASS IX

There will be one paper of two hours duration of 80 marks and Internal Assessment of practical work carrying 20 marks.

The paper will be divided into two sections, Section I (40 marks) and Section II (40 marks).

Section I (compulsory) will contain short answer questions on the entire syllabus.

Section II will contain six questions. Candidates will be required to answer any four of these six questions.

Note: All chemical reactions should be studied with reference to the reactants, products, conditions, observations and the (balanced) equations.

1. The Language of Chemistry

   (i) Symbol of an element; valency; formulae of radicals and formulae of compounds. Balancing of simple chemical equations.
   - Symbol – definition; symbols of the elements used often.
   - Valency - definition; hydrogen combination and number of valence electrons of the metals and non-metals; mono, di, tri and tetra valent elements.
   - Radicals – definition; formulae and valencies
   - Compounds – name and formulae.
   - Chemical equation – definition and examples of chemical equations with one reactant and two or three products, two reactants and one product, two reactants and two products and two reactants and three or four products; balancing of equations. (by hit and trial method).

(ii) Relative Atomic Masses (atomic weights) and Relative Molecular Masses (molecular weights): either - standard H atom or 1/12th of carbon 12 atom.
   - Definitions
   - Calculation of Relative Molecular Mass and percentage composition of a compound.

2. Chemical changes and reactions

   (i) Types of chemical changes.

   - Direct combination
   - Decomposition
   - Displacement;
   - Double decomposition

   (The above to be taught with suitable chemical equations as examples).

(ii) Energy changes in a chemical change.

   Exothermic and endothermic reactions with examples – evolution/absorption of heat, light and electricity.

3. Water

   (i) Water as a universal solvent.

   - Solutions as 'mixtures' of solids in water; saturated solutions.
• Qualitative effect of temperature on solubility (e.g., solutions of calcium sulphate, potassium nitrate and sodium chloride in water).

(ii) Hydrated and anhydrous substances.

(a) Hydrated substances:

Water of Crystallisation – meaning and examples

(b) Anhydrous substances:

Meaning and examples only

(c) Properties:

• Efflorescence
• Deliquescence
• Hygroscopy
• Removal of hardness
  (i) By boiling
  (ii) By addition of washing soda

(Definition and examples of each of the above).

(iii) Drying and Dehydrating Agents

Meaning and examples only.

(iv) Soft water and Hard water

• Meaning, (in terms of action of soap)
• Advantages and disadvantages of soft water and hard water.
• Types and causes of hardness.

4. Atomic Structure and Chemical bonding

(i) Structure of an Atom, mass number and atomic number, Isotopes and Octet Rule.

• Definition of an atom
• Constituents of an atom - nucleus (protons, neutrons) with associated electrons; mass number, atomic number.
• Electron distribution in the orbits - \(2n^2\) rule, Octet rule. Reason for chemical activity of an atom.
• Definition and examples of isotopes (hydrogen, carbon, chlorine).

(ii) Electrovalent and covalent bonding, structures of various compounds – orbit structure

(a) Electrovalent Bond

• Definition
• Atomic orbit structure for the formation of Electrovalent compounds (e.g. NaCl, MgCl\(_2\), CaO);

(b) Covalent Bond

• Definition
• Atomic orbit structure for the formation of Covalent molecules on the basis of duplet and octet of electrons (examples: hydrogen, chlorine, oxygen, nitrogen, hydrogen chloride, water, ammonia, carbon tetrachloride, methane.)

5. The Periodic Table

Dobereiner’s Triads, Newland’s law of Octaves, Mendeleev’s contributions; Modern Periodic Law, the Modern Periodic Table. (Groups and periods)

• General idea of Dobereiner’s triads, Newland’s law of Octaves, Mendeleev’s periodic law.
• Discovery of Atomic Number and its use as a basis for Modern Periodic law.
• Modern Periodic Table (Groups 1 to 18 and periods 1 to 7).
• Special reference to Alkali metals (Group 1), Alkaline Earth metals (Group 2) Halogens (Group 17) and Zero Group (Group 18).


Position of the non-metal (Hydrogen) in the periodic table and general group characteristics with reference to valency electrons, burning, ion formation applied to the above mentioned element.

(i) Hydrogen from: water, dilute acids and alkalis.

(a) Hydrogen from water:

• The action of cold water on sodium potassium and calcium.
• The action of hot water on magnesium.
• The action of steam on aluminium, zinc, and iron; (reversibility of reaction between iron and steam).
• The action of steam on non-metal (carbon).

Students can be shown the action of sodium and calcium on water in the laboratory. They must be asked to make observations and write equations for the above reactions.

Application of activity series for the above mentioned reactions.

(b) Displacement of hydrogen from dilute acids:

The action of dilute sulphuric acid or hydrochloric acid on metals: Mg, Al, Zn and Fe

(To understand reasons for not using other metals and dilute nitric acid)

(c) Displacement of hydrogen from alkalis:

The action of Alkalis ((NaOH, KOH) on Al, Zn and Pb – unique nature of these elements.

(ii) The preparation and collection of hydrogen by a standard laboratory method other than electrolysis.

In the laboratory preparation, the reason for using zinc, the impurities in the gas, their removal and the precautions in the collection of the gas must be mentioned.

(iii) Industrial manufacture of hydrogen by Bosch process:

• Main reactions and conditions.
• Separation of CO₂ and CO from hydrogen.

(iv) Oxidation and reduction reactions

Differences in terms of addition and removal of oxygen / hydrogen.

7. Study of Gas Laws

(i) The behaviour of gases under changes of temperature and pressure; explanation in terms of molecular motion (particles, atoms, molecules); Boyle’s Law and Charles’ Law; absolute zero; gas equation; simple relevant calculations.

• The behaviour of gases under changes of temperature and pressure; explanation in terms of molecular motion (particles, atoms, molecules).
• Boyle’s Law: statement, mathematical form, simple calculations.
• Absolute zero Kelvin scale of temperature.
• Gas equation \( P_1 V_1 / T_1 = P_2 V_2 / T_2 \); simple relevant calculations based on gas equation.

(ii) Relationship between Kelvin scale and Celsius Scale of temperature; Standard temperature and pressure. Conversion of temperature from Celsius Scale to Kelvin scale and vice versa. Standard temperature and pressure. (Simple calculations).

8. Atmospheric pollution

(a) Acid rain – composition, cause and its impact.

Sulphur in fossil fuels giving oxides of sulphur when burnt. High temperatures in furnaces and internal combustion engines produce oxides of nitrogen. (Equations to be included). Acid rain affects soil chemistry and water bodies.

(b) Global warming:

Greenhouse gases – their sources and ways of reducing their presence in the atmosphere.

(Water vapour, carbon dioxide, methane and oxides of nitrogen)

(c) Ozone depletion

• Formation of ozone – relevant equations
• Function in the atmosphere.
• Destruction of the ozone layer – chemicals responsible for this to be named but reactions not required.
INTERNAL ASSESSMENT OF PRACTICAL WORK

Candidates will be asked to observe the effect of reagents and/or of heat on substances supplied to them. The exercises will be simple and may include the recognition and identification of certain gases listed below.

**Gases:** Hydrogen, Oxygen, Carbon dioxide, Chlorine, Hydrogen chloride, Sulphur dioxide, Hydrogen sulphide, Ammonia, Water vapour, Nitrogen dioxide.

Candidates are expected to have completed the following minimum practical work.

**Simple experiments on:**

1. Action of heat on the following compounds:
   (a) copper carbonate, zinc carbonate
   (b) washing soda, copper sulphate crystals
   (c) zinc nitrate, copper nitrate, lead nitrate
   (d) ammonium chloride, iodine, ammonium dichromate
   Make observations, identify the products and make deductions where possible.

2. Action of dilute sulphuric acid on the following substances. (warm if necessary)
   (a) a metal
   (b) a carbonate
   (c) a sulphide
   (d) a sulphite
   Make observations, identify the gas evolved and make deductions

3. Apply the flame test to identify the metal in the unknown substance.
   (a) a sodium salt
   (b) a potassium salt
   (c) a calcium compound


5. Find out the sources of pollution of water bodies in the locality. Suggest preventive steps to control it.
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Section I (compulsory) will contain short answer questions on the entire syllabus.

Section II will contain six questions. Candidates will be required to answer any four of these six questions.

Note: All chemical process/reactions should be studied with reference to the reactants, products, conditions, observation, the (balanced) equations and diagrams.

   (i) Periodic properties and their variations in groups and periods.

   Definitions and trends of the following periodic properties in groups and periods should be studied:
   • atomic size
   • metallic character
   • non-metallic character
   • ionisation potential
   • electron affinity
   • electronegativity

   (ii) Periodicity on the basis of atomic number for elements.

   • The study of modern periodic table up to period 3 (students to be exposed to the complete modern periodic table but no questions will be asked on elements beyond period 3 – Argon);
   • Periodicity and other related properties to be explained on the basis of nuclear charge and shells (not orbitals).

   (Special reference to the alkali metals and halogen groups).

2. Chemical Bonding

   Electrovalent, covalent and co-ordinate bonding, structures of various compounds, Electron dot structure.

   (a) Electrovalent bonding:

   • Electron dot structure of Electrovalent compounds NaCl, MgCl₂, CaO.

   • Characteristic properties of electrovalent compounds – state of existence, melting and boiling points, conductivity (heat and electricity), dissociation in solution and in molten state to be linked with electrolysis.

   (b) Covalent Bonding:

   • Electron dot structure of covalent molecules on the basis of duplet and octet of electrons (example: hydrogen, chlorine, nitrogen, ammonia, carbon tetrachloride, methane.

   • Polar Covalent compounds – based on difference in electronegativity:

     Examples – HCl and H₂O including structures.

   • Characteristic properties of Covalent compounds – state of existence, melting and boiling points, conductivity (heat and electricity), ionisation in solution.

   Comparison of Electrovalent and Covalent compounds.

   (c) Coordinate Bonding:

   • Definition

   • The lone pair effect of the oxygen atom of the water molecule and the nitrogen atom of the ammonia molecule to explain the formation of H₃O⁺ and OH⁻ ions in water and NH₄⁺ ion.
The meaning of lone pair; the formation of hydronium ion and ammonium ion must be explained with help of electron dot diagrams.

3. Study of Acids, Bases and Salts
   (i) Simple definitions in terms of the molecules and their characteristic properties.
   (ii) Ions present in mineral acids, alkalis and salts and their solutions; use of litmus and pH paper to test for acidity and alkalinity.
   • Examples with equation for the ionisation/dissociation of ions of acids, bases and salts.
   • Acids form hydronium ions (only positive ions) which turn blue litmus red, alkalis form hydroxyl ions (only negative ions) with water which turns red litmus blue.
   • Salts are formed by partial or complete replacement of the hydrogen ion of an acid by a metal. (To be explained with suitable examples).
   • Introduction to pH scale to test for acidity, neutrality and alkalinity by using pH paper or Universal indicator.

(iii) Definition of salt; types of salts.
   Types of salts: normal salts, acid salt, basic salt, definition and examples.

(iv) Action of dilute acids on salts.
   Decomposition of hydrogen carbonates, carbonates, sulphites and sulphides by appropriate acids with heating if necessary. (Relevant laboratory work must be done).

(v) Methods of preparation of Normal salts with relevant equations. (Details of apparatus or procedures not required).
Methods included are:
• Direct combination
• Displacement
• Precipitation (double decomposition)
• Neutralization of insoluble base
• Neutralisation of an alkali (titration)

4. Analytical Chemistry
   (i) Action of Ammonium Hydroxide and Sodium Hydroxide on solution of salts: colour of salt and its solution; formation and colour of hydroxide precipitated for solutions of salts of Ca, Fe, Cu, Zn and Pb; special action of ammonium hydroxide on solutions of copper salt and sodium hydroxide on ammonium salts.

On solution of salts:
• Colour of salt and its solution.
• Action on addition of Sodium Hydroxide to solution of Ca, Fe, Cu, Zn, and Pb salts drop by drop in excess. Formation and colour of hydroxide precipitated to be highlighted with the help of equations.
• Action on addition of Ammonium Hydroxide to solution of Ca, Fe, Cu, Zn, and Pb salts drop by drop in excess. Formation and colour of hydroxide precipitated to be highlighted with the help of equations.
• Special action of Ammonium Hydroxide on solutions of copper salts and sodium hydroxide on ammonium salts.

(ii) Action of alkalis (NaOH, KOH) on certain metals, their oxides and hydroxides.
   The metals must include aluminium, zinc and lead, their oxides and hydroxides, which react with caustic alkalis (NaOH, KOH), showing the amphoteric nature of these substances.

5. Mole Concept and Stoichiometry
   (i) Gay Lussac’s Law of Combining Volumes; Avogadro’s Law.
   • Idea of mole – a number just as a dozen, a gross (Avogadro’s number).
• Avogadro’s Law - statement and explanation.
• Gay Lussac’s Law of Combining Volumes. – Statement and explanation.
• Understanding molar volume- “the mass of 22.4 litres of any gas at S.T.P. is equal to its molar mass”. (Questions will not be set on formal proof but may be taught for clear understanding).
• Simple calculations based on the molar volume and Gay Lussac’s law.

(ii) Refer to the atomicity of hydrogen, oxygen, nitrogen and chlorine (proof not required).

The explanation can be given using equations for the formation of HCl, NH₃, and NO.

(iii) Vapour Density and its relation to relative molecular mass:
• Molecular mass = 2 × vapour density (formal proof not required)
• Deduction of simple (empirical) and molecular formula from:
  (a) the percentage composition of a compound.
  (b) the masses of combining elements.

(iv) Mole and its relation to mass.
• Relating mole and atomic mass; arriving at gram atomic mass and then gram atom; atomic mass is a number dealing with one atom; gram atomic mass is the mass of one mole of atoms.
• Relating mole and molecular mass; arriving at gram molecular mass and gram molecule – molecular mass is a number dealing with a molecule, gram molecular mass is the mass of one mole of molecules.
• Simple calculations based on relation of mole to mass, volume and Avogadro’s number.

(v) Simple calculations based on chemical equations Related to weight and/or volumes of both reactants and products.

6. Electrolysis

(i) Electrolytes and non-electrolytes.

Definitions and examples.

(ii) Substances containing molecules only, ions only, both molecules and ions.
• Substances containing molecules only ions only, both molecules and ions.
• Examples; relating their composition with their behaviour as strong and weak electrolytes as well as non-electrolytes.

(iii) Definition and explanation of electrolysis, electrolyte, electrode, anode, cathode, anion, cation, oxidation and reduction (on the basis of loss and gain of electrons).

(iv) An elementary study of the migration of ions, with reference to the factors influencing selective discharge of ions (reference should be made to the activity series as indicating the tendency of metals, e.g. Na, Mg, Fe, Cu, to form ions) illustrated by the electrolysis of:
• Molten lead bromide
• Acidified water with platinum electrodes
• Aqueous copper (II) sulphate with copper electrodes; electron transfer at the electrodes.

The above electrolytic processes can be studied in terms of electrolyte used, electrodes used, ionization reaction, anode reaction, cathode reaction, use of selective discharge theory, wherever applicable.

(v) Applications of electrolysis:
• Electroplating with nickel and silver, choice of electrolyte for electroplating.
• Electro refining of copper;

Reasons and conditions for electroplating; names of the electrolytes and the electrodes used should be given. Equations for the reactions at the electrodes should be given for electroplating, refining of copper.
7. Metallurgy

(i) Occurrence of metals in nature:
- Mineral and ore - Meaning only.
- Common ores of iron, aluminium and zinc.

(ii) Stages involved in the extraction of metals:
(a) Dressing of the ore – hydrolytic method, magnetic separation, froth flotation method.
(b) Conversion of concentrated ore to its oxide - roasting and calcination (definition, examples with equations).
(c) Reduction of metallic oxides - some can be reduced by hydrogen, carbon and carbon monoxide (e.g. copper oxide, lead (II) oxide, iron (III) oxide and zinc oxide) and some cannot (e.g. Al₂O₃, MgO - refer to activity series). Active metals by electrolysis e.g. sodium, potassium and calcium. (reference only).

Equations with conditions should be given.

(d) Electro refining – reference only

(iii) Extraction of Aluminium.
(a) Chemical method for purifying bauxite by using NaOH – Baeyer’s Process.
(b) Electrolytic extraction – Hall Heroult’s process:
Structure of electrolytic cell - the various components as part of the electrolyte, electrodes and electrode reactions.
Description of the changes occurring, purpose of the substances used and the main reactions with their equations.

(iv) Alloys – composition and uses
Stainless steel, duralumin, brass, bronze, fuse metal / solder.

8. Study of Compounds

A. Hydrogen Chloride
Hydrogen chloride: preparation of hydrogen chloride from sodium chloride; refer to the density and solubility of hydrogen chloride (fountain experiment); reaction with ammonia; acidic properties of its solution.
- Preparation of hydrogen chloride from sodium chloride; the laboratory method of preparation can be learnt in terms of reactants, product, condition, equation, diagram or setting of the apparatus, procedure, observation, precaution, collection of the gas and identification.
- Simple experiment to show the density of the gas (Hydrogen Chloride) – heavier than air.
- Solubility of hydrogen chloride (fountain experiment); setting of the apparatus, procedure, observation, inference.
- Method of preparation of hydrochloric acid by dissolving the gas in water - the special arrangement and the mechanism by which the back suction is avoided should be learnt.
- Reaction with ammonia
- Acidic properties of its solution - reaction with metals, their oxides, hydroxides and carbonates to give their chlorides; decomposition of carbonates, hydrogen carbonates, sulphides, sulphites.
- Precipitation reactions with silver nitrate solution and lead nitrate solution.

B. Ammonia
Ammonia: its laboratory preparation from ammonium chloride and collection; ammonia from nitrides like Mg₃N₂ and AlN and ammonium salts. Manufacture by Haber’s Process; density and solubility of ammonia (fountain experiment); aqueous solution of ammonia; its reactions with hydrogen chloride and with hot copper (II) oxide
and chlorine; the burning of ammonia in oxygen; uses of ammonia.

- Laboratory preparation from ammonium chloride and collection; (the preparation to be studied in terms of; setting of the apparatus and diagram, procedure, observation, collection and identification)
- Ammonia from nitrides like Mg$_3$N$_2$ and AlN using warm water.
- Ammonia from ammonium salts using alkalis.

*The reactions to be studied in terms of reactants, products, conditions and equations.*

- Manufacture by Haber’s Process.
- Density and solubility of ammonia (fountain experiment).
- The burning of ammonia in oxygen.
- The catalytic oxidation of ammonia (with conditions and reaction)
- Its reactions with hydrogen chloride and with hot copper (II) oxide and chlorine (both chlorine in excess and ammonia in excess).

*All these reactions may be studied in terms of reactants, products, conditions, equations and observations.*

- Aqueous solution of ammonia - reaction with sulphuric acid, nitric acid, hydrochloric acid and solutions of iron(III) chloride, iron(II) sulphate, lead nitrate, zinc nitrate and copper sulphate.
- Uses of ammonia - manufacture of fertilizers, explosives, nitric acid, refrigerant gas (Chlorofluoro carbon – and its suitable alternatives which are non-ozone depleting), and cleansing agents.

C. Nitric Acid

Nitric Acid: one laboratory method of preparation of nitric acid from potassium nitrate or sodium nitrate. Large scale preparation. Nitric acid as an oxidizing agent.

- Laboratory preparation of nitric acid from potassium nitrate or sodium nitrate; the laboratory method to be studied in terms of reactants, products, conditions, equations, setting up of apparatus, diagram, precautions, collection and identification.
- Manufacture of Nitric acid by Ostwald’s process (Only equations with conditions where applicable).
- As an oxidising agent: its reaction with copper, carbon, sulphur.

D. Sulphuric Acid

Large scale preparation, its behaviour as an acid when dilute, as an oxidizing agent when concentrated - oxidation of carbon and sulphur; as a dehydrating agent - dehydration of sugar and copper (II) sulphate crystals; its non-volatile nature.

- Manufacture by Contact Process Equations with conditions where applicable).
- Its behaviour as an acid when dilute - reaction with metal, metal oxide, metal hydroxide, metal carbonate, metal bicarbonate, metal sulphite, metal sulphide.
- Concentrated sulphuric acid as an oxidizing agent - the oxidation of carbon and sulphur.
- Concentrated sulphuric acid as a dehydrating agent- (a) the dehydration of sugar (b) Copper (II) sulphate crystals.
- Non-volatile nature of sulphuric acid - reaction with sodium or potassium chloride and sodium or potassium nitrate.

9. Organic Chemistry

(i) Introduction to Organic compounds.

- Unique nature of Carbon atom – tetra valency, catenation.
• Formation of single, double and triple bonds, straight chain, branched chain, cyclic compounds (only benzene).

(ii) Structure and Isomerism.
• Structure of compounds with single, double and triple bonds.
• Structural formulae of hydrocarbons. Structural formula must be given for: alkanes, alkenes, alkynes up to 5 carbon atoms.
• Isomerism – structural (chain, position)

(iii) Homologous series – characteristics with examples.
Alkane, alkene, alkyne series and their gradation in properties and the relationship with the molecular mass or molecular formula.

(iv) Simple nomenclature.
Simple nomenclature - of the hydrocarbons with simple functional groups – (double bond, triple bond, alcoholic, aldehydic, carboxylic group) longest chain rule and smallest number for functional groups rule – trivial and IUPAC names (compounds with only one functional group)

(v) Hydrocarbons: alkanes, alkenes, alkynes.
• Alkanes - general formula; methane (greenhouse gas) and ethane - methods of preparation from sodium ethanoate (sodium acetate), sodium propanoate (sodium propionate), from iodomethane (methyl iodide) and bromoethane (ethyl bromide). Complete combustion of methane and ethane, reaction of methane and ethane with chlorine through substitution.
• Alkenes – (unsaturated hydrocarbons with a double bond); ethene as an example. Methods of preparation of ethene by dehydro halogenation reaction and dehydration reactions.
• Alkynes -(unsaturated hydrocarbons with a triple bond); ethyne as an example of alkyne; Methods of preparation from calcium carbide and 1,2 dibromoethane ethylene dibromide).

Only main properties, particularly addition products with hydrogen and halogen namely Cl₂, Br₂ and I₂ pertaining to alkenes and alkynes.

• Uses of methane, ethane, ethene, ethyne.

(vi) Alcohols: ethanol – preparation, properties and uses.
• Preparation of ethanol by hydrolysis of alkyl halide.
• Denatured and spurious alcohol.
• Important uses of Ethanol.

(vii) Carboxylic acids (aliphatic - mono carboxylic acid): Acetic acid – properties and uses of acetic acid.
• Structure of acetic acid.
• Properties of Acetic Acid: Physical properties – odour (vinegar), glacial acetic acid (effect of sufficient cooling to produce ice like crystals). Chemical properties – action with litmus, alkalis and alcohol (idea of esterification).
• Uses of acetic acid.

INTERNAL ASSESSMENT OF PRACTICAL WORK
Candidates will be asked to observe the effect of reagents and/or of heat on substances supplied to them. The exercises will be simple and may include the recognition and identification of certain gases and ions listed below. The examiners will not, however, be restricted in their choice to substances containing the listed ions.


Ions: Calcium, Copper, Iron, Lead, Zinc and Ammonium, Carbonate, Chloride, Nitrate, Sulphide, Sulphite and Sulphate.
Knowledge of a formal scheme of analysis is not required. Semi-micro techniques are acceptable but candidates using such techniques may need to adapt the instructions given to suit the size of the apparatus being used.

Candidates are expected to have completed the following minimum practical work:

1. Action of heat on the following substances:
   (a) Copper carbonate, zinc carbonate
   (b) zinc nitrate, copper nitrate, lead nitrate
   Make observations, identify the products and make deductions where possible. (equations not required)

2. Make a solution of the unknown substance: add sodium hydroxide solution or ammonium hydroxide solution, make observations and give your deduction. Warming the mixture may be needed. Choose from substances containing Ca²⁺, Cu²⁺, Fe²⁺, Fe³⁺, Pb²⁺, Zn²⁺, NH₄⁺.

3. Supply a solution of a dilute acid and alkali. Determine which is acidic and which is basic, giving two tests for each.

4. Add concentrated hydrochloric acid to each of the given substances, warm, make observations, identify any product and make deductions: (a) copper oxide (b) manganese dioxide.

**EVALUATION**

The assignments/project work are to be evaluated by the subject teacher and by an External Examiner. (The External Examiner may be a teacher nominated by the Head of the school, who could be from the faculty, **but not teaching the subject in the section/class**. For example, a teacher of Chemistry of Class VIII may be deputed to be an External Examiner for Class X Chemistry projects.)

The Internal Examiner and the External Examiner will assess the assignments independently.

**Award of marks (20 Marks)**

Subject Teacher (Internal Examiner) 10 marks
External Examiner 10 marks

The total marks obtained out of 20 are to be sent to the Council by the Head of the school.

The Head of the school will be responsible for the entry of marks on the mark sheets provided by the Council.

**NOTE:** According to the recommendation of International Union of Pure and Applied Chemistry (IUPAC), the groups are numbered from 1 to 18 replacing the older notation of groups IA .... VIIA, VIII, IB ...... VIIB and 0. However, for the examination both notations will be accepted.

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