Volume I & II

CHEMISTRY

12th Standard

Strictly as per the Reduced (Prioritised) Syllabus released on 13th August, 2021 (G.O.(Ms).No126)

Sura's Model question paper is given based on the reduced syllabus, with answers.

Salient Features

- Complete Solutions to Textbook Exercises.
- Model Question Papers 1 to 6 (PTA) : Questions are incorporated in the appropriate sections.
- Govt. Model Question Paper 2019, Quarterly Exam 2019, Half Yearly Exam 2019, March Public Exam 2020 and Govt. Supplementary Exam September 2020 questions are incorporated in the appropriate sections.
- NEET based questions with answers are also given.



orders@surabooks.com

Public Exam Edition 2021-22

2021-22 Edition

All rights reserved © SURA Publications.

No part of this book may be reproduced, stored in a retrieval system, or transmitted in any form or by any means, digitally, electronically, mechanically, photocopying, recorded or otherwise, without the written permission of the publishers. Strict action will be taken.

ISBN : 978-93-5330-416-4

Code No : RPS_013



DIRECT DEPOSIT

A/c Name : Sura Publications	A/c Name : Sura Publications
Our A/c No. : 36550290536	Our A/c No. : 21000210001240
Bank Name : STATE BANK OF INDIA	Bank Name : UCO BANK
Bank Branch : PADI	Bank Branch : Anna Nagar West
IFSC : SBIN0005083	IFSC : UCBA0002100
A/c Name : Sura Publications	A/c Name : Sura Publications
A/c Name : Sura Publications Our A/c No. : 6502699356	A/c Name : Sura Publications Our A/c No. : 1154135000017684
Our A/c No. : 6502699356	Our A/c No. : 1154135000017684

For More Information - Contact

Doubts in Our Guides	:	enquiry@surabooks.com
For Order	:	orders@surabooks.com
Contact	:	80562 94222 / 80562 15222
Whatsapp	:	8124201000 / 9840926027
Online Site	:	www.surabooks.com
For Free Study Materials	Visi	t http://tnkalvi.in

Strictly as per the Reduced (Prioritised) Syllabus released on 13th August, 2021 (G.O.(Ms).No126)

Class: 12th

Subject: Chemistry

UNIT	CONTENT						
1. METALLURGY	Introduction	1.3.1	Conversion of ores into oxides				
	1.1 Occurrence of metals		Roasting				
	1.1.1 Mineral and ore		Calcination				
	1.2 Concentration of ores	1.3.2	Reduction of metal oxides				
	1.2.1 Gravity separation or Hydraulic wash		Smelting				
	1.2.2 Froth flotation		Reduction by carbon; Reduction by hydroger				
	1.2.3 Leaching	1.0	Reduction by metal; Auto-reduction:				
	Cyanide leaching	1.6	Refining process				
	Recovery of metal of interest from the complex		Distillation				
	by reduction	1.6.2	Liquation				
	Ammonia leaching	1.6.3	Electrolytic refining				
	Alkali leaching	1.6.4	Zone Refining				
	Acid leaching	1.6.5	Vapour phase method				
	1.2.4 Magnetic separation	1.0.0	Mond process for refining nickel; Van-Arkel metho				
	1.3 Extraction of crude metal		for refining zirconium / titanium				
0		0.05					
2. p-block	Introduction	2.2.5	Boric acid $[H_3BO_3 \text{ or } B(OH)_3]$				
elements - I	2.1 General trends in properties of p-block element	5	Preparation, Properties; Structure of Boric acid				
	2.1.1 Electronic configuration and oxidation state		Uses of boric acid				
	2.1.2 Metallic nature:	2.2.9	Alums				
	2.1.3 Ionisation Enthalpy		Examples				
	2.1.4 Electronegativity		Preparation; Properties of Alum; Uses of Alum				
	2.1.5 Anomalous properties of the first elements	2.3	Group 14 (Carbon group) elements:				
		2.3.1					
	2.1.6 Inert pair effect		Occurrence				
	2.1.7 Allotropism in p-block elements	2.3.2	Physical properties				
	2.2 Group 13 (Boron group) elements	2.3.3	Tendency for catenation				
	2.2.1 Occurrence	2.3.4	Allotropes of carbon				
	2.2.2 Physical properties		Structure of Graphite				
	2.2.3 Chemical properties of boron		Structure of Diamond				
	Uses of boron		Structure of Fullerenes				
	2.2.4 Borax $[Na_2B_4O_7.10H_2O]$		Structure of carbon nanotubes				
	Preparation		Structure of Graphene				
	Properties	2.3.8	Silicones				
	Uses of Borax		Preparation				
			Types of silicones				
			Properties, Uses				
3. p-block elements - II	Introduction	3.2.1	Allotropic forms of sulphur				
	3.1 Group 15 (Nitrogen group) elements	3.2.2	Sulphur dioxide				
	3.1.1 Occurrence		Preparation Properties				
	3.1.2 Physical properties		Chemical properites				
	3.1.3 Nitrogen		Uses of sulphur dioxide				
	Preparation		Structure of sulphur dioxide				
	Properties of Nitrogen		Structure of oxoacids of sulphur				
	Uses of nitrogen	3.3	Group 17 (Halogen group) elements:				
	3.1.4 Ammonia (NH ₃)	3.3.1	Chlorine				
	Preparation		Occurrence:				
	Properties of Ammonia		Physical properties of Chlorine				
	Chemical Properties	3.3.1	Manufacture of chlorine				
	Structure of ammonia		Physical properties				
	3.1.7 Allotropic forms of phosphorus		Chemical properties				
	3.1.8 Properties of phosphorus		Uses of chlorine				
	Uses of phosphorus	3.3.4	Inter halogen compounds:				
		0.0.7					
	Oxoacids of Phosphorus-Structure		Properties of inter halogen compounds				
	Group 16 (Oxygen group) elements		Structure of inter halogen compounds				
	Occurrence	3.4	Group 18 (Inert gases) elements:				
	Physical properties	3.4.1	Occurrence:				
	3.2 Oxygen		Physical properties				
	Preparation: Properties		Physical properties-Inert Gases				
	Chemical properties	1	Properties of inert gases				
	Uses of Oxygen		Chemical Properties				
			Chemical Properties Structures of compounds of Xenon				

4. TRANSITION AND INNER	Introduction	4.3.10	Formation of complexes
TRANSITION ELEMENTS	 4.1 Position of d- block elements in the periodic table 4.2 Electronic configuration 4.3 General trend in properties 4.3.1 Metallic behavior 4.3.2 Variation of atomic and ionic size 4.3.3 Ionization enthalpy 4.3.4 Oxidation state 4.3.5 Standard electrode potentials of transition metals 4.3.6 Magnetic properties 4.3.7 Catalytic properties 4.3.8 Alloy formation 4.3.9 Formation of interstitial compounds 	4.4	Important compound of Transition elements f-block elements – Inner transition elements The position of Lanthanoids in the periodic table Electronic configuration of Lanthanoids Oxidation state of lanthanoids Atomic and ionic radii Causes of lanthanoid contraction Consequences of lanthanoid contraction Actinoids Electronic configuration of actinoids Oxidation state of actinoids Differences between lanthanoids and actinoids
5. COORDINATION	Introduction		Classification based on the net charge on the
CHEMISTRY	 5.1 Coordination compounds and double salts 5.2 Werner's theory of coordination compounds Postulates Werner's theory 5.2.1 Limitations of Werner's theory 5.3 Definition of important terms pertaining to co-ordination compounds 5.3.1 Coordination entity 5.3.2 Central atom/ion 5.3.3 Ligands Coordination sphere, Coordination polyhedron, Coordination number, Oxidation state (number) Types of complexes 	5.4 5.6 5.6.1	complex Classification based on kind of ligands Nomenclature of coordination compounds a. Naming the ligands b.Naming the central metal More examples with names – IUPAC Nomenclature Theories of coordination compound Valence Bond Theory Main assumptions of VBT Illustration(1-4) Limitations of VBT
6. Solid state	 6. Introduction 6.1 General characteristics of solids 6.2. Classification of Solids 6.3. Classification of Crystalline Solids 6.3.1. Ionic solids 6.3.2. Covalent Solids 6.3.3. Molecular Solids 6.3.4. Metallic Solids 6.4. Crystal lattice and unit cell 6.5 Primitive and Non Primitive unit 6.5.1 Primitive (or) Simple Cube unit cell 6.5.2 Body Centered cubic unit cell 6.5.3 Face centered cubic unit cell 	6.6.2 6.6.3 6.6.4. 6.7. 6.7.1 6.7.2. 6.7.3. 6.7.4.	Calculations involving unit cell Dimensions Calculation of density Packing in Crystals Linear arrangement of spheres in one direction Two dimensional Close Packing Simple Cubic arrangement Body Centered Cubic arrangement Imperfections in solids Schottky defect Frenkel defect Metal Excess defect Metal Deficiency defect Impurity defect
7. CHEMICAL KINETICS	Introduction	7.5.1	Integrated rate law for First order, Pseudo first order
	 7.1 Rate of Chemical reaction 7.1.1 Stoichiometry and rate of a reaction 7.1.2 Average and instantaneous rate 7.3 Rate law and Rate Constant 7.4 Molecularity 7.5 Integrated Rate Equation 	7.5.2 7.6 7.8	reaction Integrated rate law for a Zero order reaction Half life period of a reaction Arrhenius Equation-The effect of temperature on reaction rate
8. IONIC EQUILIBRIUM	Introduction8.1.Acids and bases8.1.1Arrhenius concept8.1.2Lowry - Bronsted Theory8.1.3Lewis Concept8.2Strength Of Acids and Bases8.3Ionisation of water8.4The pH Scale8.4.1Relation between pH and pOH	8.5 8.5.1 8.6. 8.7 8.7.1 8.7.3 8.9 8.9.1	Ionistion of Weak Acids Ostwalds Dilution Law Common ion effect Buffer Solution Buffer Action Henderson Hasselbalch Equation Solubility Product Determination of solubility Product from Molar Solubility
9. Electro chemistry	Introduction9.1Conductivity of electrolytic solution9.1.1Molar conductivity9.1.2Equivalent conductance9.1.3Factors affecting Electrolytic conductance9.1.4Measurement of conductivity of ionic solutions9.2Variation of molar conductivity with concentration9.2.2Kohlrausch's law and Applications	9.3.2 9.3.4 9.4 9.4.1	Galvanic cell notation Measurement of electrode potential Thermodynamics of cell reactions Nernst equation Electrolytic cell and Electrolysis Faraday's law of electrolysis First law, Second law Electrochemical series

10. SURFACE CHEMISTRY	Introduc	ction	10.5.2	Preparation of Colloids
	10.1	Adsorption and Absorption	101012	(1) Dispersion methods [mechanical dispersion,
		Characteristics of adsorption		electro dispersion, ultrasonic dispersion,
	10.1.1	Types of Adsorption		peptisation]
		Distinction between Physical and		(2) Condensation method [oxidation, reduction,
		Chemical Adsorption		hydrolysis, double decomposition,
	10.1.2	Factors affecting Adsorption		Decomposition]
	10.1.3	Adsorption isotherms and isobars		(3) By exchange of solvent
	10.1.3.1		10.5.3	Purification of colloids
	10.1.3.1	Catalysis Positive and Negative Catalysis	10.5.5	(i) Dialysis
	10.2	Characteristics of Catalysis		(ii) Electrodialysis
	10.2.1	Promoters and Catalytic poison		(ii) Ultrafiltration
			10.5.4	
	10.0.0	Auto Catalysis, Negative Catalysis	10.5.4	Properties of colloids 14 points [colour, size,
	10.2.2	Theories of Catalysis		Heterogeneous nature, Filtrability, Non-Setting
		The Intermediate compound formation theory,		nature, Concentration & density, Diffusability,
	10 5	Adsorption Theory & Active Centers		Colligative Properties, Shape of Colloidal
	10.5	Colloid, dispersion Phase and dispersion		Particles, Optical, Kinetic and Electrical properties,
		medium		Coagulation, Protective action]
	10.5.1	Classification of colloidal solution		
11. Hydroxy	Introduc			Acidity of phenols
COMPOUNDS AND		Classification of Alcohols		Preparation of phenol
ETHERS		JPAC Nomenclature		Physical Properties of Phenol
		tructure of functional group of alcohols		Chemical properties of phenols
		Physical Properties of Alcohols		Test to differentiate Alcohols & Phenols
		Preparation of Alcohols		Uses of phenol
		lethods to differentiate primary,		ETHERS
		econdary,Tertiary		Ethers Classification
		hysical properties of alcohols		Structure of funtional group
	C	Chemical Properties of Alchols (without		IUPAC system
	m	nechanism)		Preparation of Ethers except mechanism
	U	Jses of Alcohols		Physical properties
	A	acidity of alcohols		Chemical Properties of Ethers (except mechanism)
19 Curroury	Indua dara	-41	CADDO	uses DXYLIC ACIDS
12. CARBONYL COMPOUNDS AND	Introduce 12.1 N	Cuon Iomenclature of Aldehyde and Ketones	12.8	
CARBOXYLIC ACIDS		-		IUPAC Nomenclature of carboxylic acids
CAMBOATLIC ACIDS		structure of carbonyl group	12.9	Structure of carboxyl group
		Seneral methods of preparation of	12.10	Methods of preparation of carboxylic acids except
		ldehydes and Ketones	10.11	Sno 5
		Physical properties of Aldehydes and Ketones		Physical properties of carboxylic acids
		Chemical properties of Aldehydes and Ketones	12.12	Chemical properties of carboxylic acids (except
	(I) (I	Mechanism only for aldol and cannizaro reaction)		mechanism of esterification)
	12.6 Te	est for Aldehydes (First two test only)		Test for carboxylic acid
				Acidity of carboxylic acids
	Introduc	ction		Acidity of carboxylic acids Chemical properties of Nitro Alkanes
13. Organic nitrogen compounds	Introduc 13.1 N	ction Jitro Compounds	13.1.8	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes
	Introduc 13.1 N 13.1.1 C	c tion litro Compounds Classification of Nitro compounds	13.1.8 13.2	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification
	Introduc 13.1 N 13.1.1 C 13.1.2 N	ction litro Compounds Classification of Nitro compounds Iomenclature of nitro alkanes	13.1.8 13.2 13.2.1	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines
	Introduce 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is	ction litro Compounds Classification of Nitro compounds Iomenclature of nitro alkanes somerism	13.1.8 13.2 13.2.1 13.2.2	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines
	Introduce 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.4 A	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes	13.1.8 13.2 13.2.1 13.2.2 13.2.3	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines
	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.4 A 13.1.5 P	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only	13.1.8 13.2 13.2.1 13.2.2 13.2.3	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines
	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.4 A 13.1.5 P	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines
	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines
COMPOUNDS	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P	ction litro Compounds Classification of Nitro compounds lomenclature of nitro alkanes somerism acidic Nature of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only Preparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties
COMPOUNDS	Introduct 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P 13.1.7 P 13.1.7 P Introduct Introduct	ction litro Compounds Classification of Nitro compounds lomenclature of nitro alkanes somerism acidic Nature of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only Preparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6 14.2.1	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties Chemical properties of Amines
13. Organic nitrogen compounds 14. Bio molecules	Introduct 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P 13.1.7 P Introduct 14.1	ction litro Compounds Classification of Nitro compounds lomenclature of nitro alkanes somerism acidic Nature of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only Preparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes ction	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6 14.2.1 14.2.3	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties Chemical properties of Amines Amino acids
COMPOUNDS	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P 13.1.7 P 13.1.7 P 14.1 C 14.1.2 C	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only Preparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes Ction Carbohydrate Classification of carbohydrate	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6 14.2.1 14.2.3 14.2.4	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties Chemical properties of Amines Amino acids Properties of Amino acids
COMPOUNDS	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.3 Is 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P 13.1.7 P 13.1.7 P 14.1 C 14.1.2 C 14.1.3 G	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes Preparation of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only Preparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes Ction Carbohydrate Classification of carbohydrate Slucose (except cyclic structure of glucose)	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6 14.2.1 14.2.3 14.2.4 14.5	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties Chemical properties Chemical properties of Amines Amino acids Properties of Amino acids Peptide bond formation Nucleic acids
COMPOUNDS	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.3 Is 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P 13.1.7 P 13.1.7 P 13.1.7 C 14.1 C 14.1.2 C 14.1.3 G 14.1.4 F	ction litro Compounds Classification of Nitro compounds lomenclature of nitro alkanes somerism acidic Nature of Nitro Alkanes reparation of Nitro Alkanes reparation of Nitro Alkane first 3 methods only reparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes ction Carbohydrate Classification of carbohydrate Slucose (except cyclic structure of glucose) ructose (except cyclic structure of fructose)	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6 14.2.1 14.2.3 14.2.4 14.5 14.5.1	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties Chemical properties Chemical properties of Amines Amino acids Properties of Amino acids Peptide bond formation Nucleic acids Composition and structure of nucleic acid
COMPOUNDS	Introduc 13.1 N 13.1.1 C 13.1.2 N 13.1.3 Is 13.1.3 Is 13.1.3 Is 13.1.4 A 13.1.5 P 13.1.6 P 13.1.7 P Introduc 14.1 14.1.2 C 14.1.3 G 14.1.4 F 14.1.5 D	ction Nitro Compounds Classification of Nitro compounds Nomenclature of nitro alkanes somerism Acidic Nature of Nitro Alkanes Preparation of Nitro Alkanes Preparation of Nitro Alkane first 3 methods only Preparation of Nitro Arenes first method only Physical Properties of Nitro Alkanes Ction Carbohydrate Classification of carbohydrate Slucose (except cyclic structure of glucose)	13.1.8 13.2 13.2.1 13.2.2 13.2.3 13.2.4 13.2.5 13.2.6 14.2.1 14.2.3 14.2.4 14.5 14.5.1	Acidity of carboxylic acids Chemical properties of Nitro Alkanes Chemical properties of Nitro benzenes Amines - Classification Nomenclature IUPAC system of Amines Structure of Amines General Methods of Preparation of Amines Properties of amines Chemical properties Chemical properties Chemical properties of Amines Amino acids Properties of Amino acids Peptide bond formation Nucleic acids

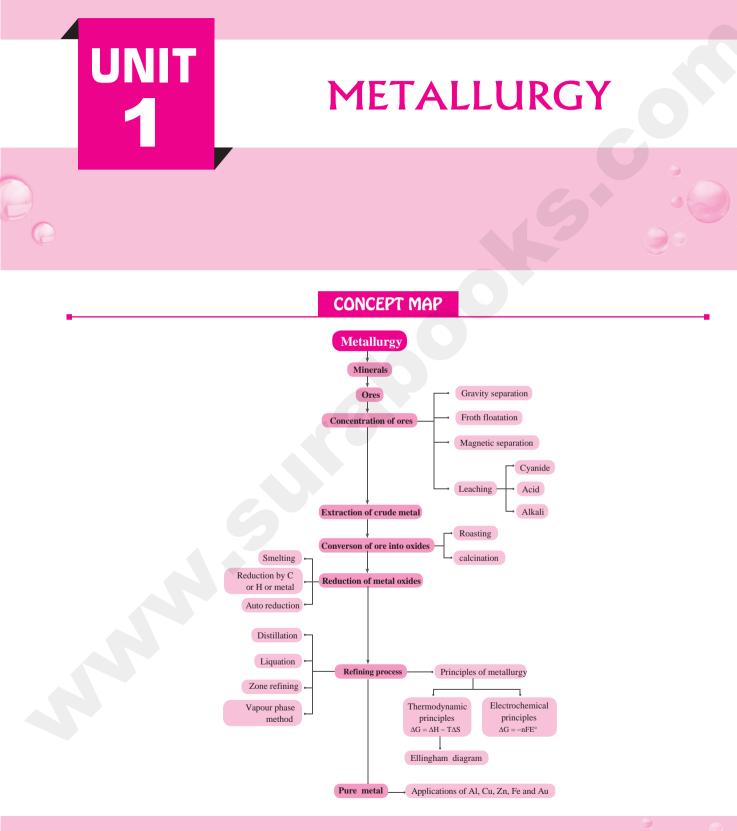
orders@surabooks.com



VOLUME - I	
Unit 1 Metallurgy	1 - 8
Unit 2 p-Block Elements-I	9 - 16
Unit 3 p-Block Elements - II	17 - 24
Unit 4 Transition and Inner Transition Elements	25 - 36
Unit 5 Coordination Chemistry	37 - 48
Unit 6 Solid State	49 - 64
Unit 7 Chemical Kinetics	65 - 80
VOLUME - II	
Unit 8 Ionic Equilibrium	81 - 96
Unit 9 Electro Chemistry	97 - 111
Unit 10 Surface Chemistry	112 - 123
Unit 11 Hydroxy Compounds and Ethers	124 - 139
Unit 12 Carbonyl Compounds and Carboxylic Acids	140 - 154
Unit 13 Organic Nitrogen Compounds	155 - 173
Unit 14 Biomolecules	174 - 182
Unit 15*	

LAB MANUAL	183 - 192
NEET based questions and answers	193 - 200
Sura's Model Question Paper, based on Reduced Syllabus, with answers.	201 - 210
*Removed as per Reduced Syllabus for 2021-22.	

VOLUME - I



orders@surabooks.com

🖞 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

FORMULAE TO REMEMBER

Metal	Ore	Composition	Metal	Ore	Composition
	Bauxite	Al ₂ O ₃ .nH ₂ O		Zinc blende or Sphalerite	ZnS
Aluminium	Diaspore	Al ₂ Si ₂ O ₅ (OH) ₄	Zinc	Calamine	ZnCO ₃
	Kaolinite	Al ₂ O ₃		Zincite	ZnO
	Haematite	Fe ₂ O ₃		Galena	PbS
	Magnetite	Fe ₃ O ₄	Lead	Anglesite	PbSO ₄
_	Siderite	FeCO ₃		Cerrusite	PbCO ₃
Iron	Iron pyrite	FeS ₂	Tin	Cassiterite (Tin stone)	SnO ₂
	Limonite	Fe ₂ O ₃ .3H ₂ O		Silver glance (Argentite)	Ag ₂ S
	Copper pyrite	CuFeS ₂		Pyrargyrite (Ruby silver)	Ag ₃ SbS ₃
Copper	Copper glance	Cu ₂ S	Silver	Chlorargyrite (Horn Silver)	AgCl
Copper	Cuprite	Cu ₂ O		Stefinite	Ag ₅ SbS ₄
	Malachite	CuCO ₃ .Cu(OH) ₂		Proustite	Ag ₃ AsS ₃
	Azurite	2CuCO ₃ .Cu(OH) ₂			

MUST KNOW DEFINITIONS

Mineral	:	A naturally occurring substance obtained by mining which contains the metal in free state or in the form of compounds like oxides, sulphides etc is called a mineral .
Ores	:	Minerals that contains a high percentage of metal, from which it can be extracted conveniently and economically are called ores .
Concentration of Ore	:	The preliminary step in metallurgical process is removal of impurities. This removal process is known as concentration of ore .
Roasting	:	Roasting is the method, usually applied for the conversion of sulphide ores into their oxides. The concentrated ore is oxidised by heating it with excess of oxygen in a suitable furnace below the melting point of the metal.
Calcination	:	Calcination is the process in which the concentrated ore is strongly heated in the absence of air.
Refining process	:	Removal of unreacted oxide ore, other metals, non-metals associated with the isolated crude metal is called refining process .

2

orders@surabooks.com

	3	urd S 🗰 XII Sta - Chemistry 🗰 Volume - I
Ellingham diagram	:	The graphical representation of variation of the standard Gibbs free energy of reaction for the formation of various metal oxides with temperature is called Ellingham diagram.
Application of Ellingham diagram	:	Ellingham diagram helps us to select a suitable reducing agent and appropriate temperature range for reduction.
Electrolytic refining	:	In electrolytic refining of the metal : Cathode : Pure metal Anode : Impure metal Electrolyte : Acidified aqueous solution of salt of the metal

EVALUATION

7.

3



Currela

- 1. Bauxite has the composition [HY. 2019]
 - b) Al₂O₃.nH₂O d) None of these c) $Fe_2O_3.2H_2O$ $[Ans. (b) Al_2O_3.nH_2O]$
- 2. Roasting of sulphide ore gives the gas (A). (A) is a colourless gas. Aqueous solution of (A) is acidic. The gas (A) is
 - a) CO_2 b) SO_3 d) H₂S c) SO₂ $[Ans. (c) SO_2]$
- 3. Which one of the following reaction represents calcination?
 - a) $2Zn + O_2 \longrightarrow 2ZnO$

a) Al_2O_2

- b) $2ZnS + 3O_2 \longrightarrow 2ZnO + 2SO_2$
- c) MgCO₃ \longrightarrow MgO + CO₂
- d) Both (a) and (c) $[Ans. (c) MgCO_3 \longrightarrow MgO + CO_7]$
- 4. The metal oxide which cannot be reduced to metal by carbon is

a) PbO b) Al_2O_3 c) ZnO d) FeO $[Ans. (b) Al_2O_2]$

Which of the metal is extracted by Hall-5. Heroult process?

> a) Al b) Ni c) Cu d) Zn [Ans. (a) Al]

Which of the following statements, about the advantage of roasting of sulphide ore before reduction is not true?

a) ΔG_{f}° of sulphide is greater than those for CS₂ and H₂S.

Metallurgy

- b) ΔG_r° is negative for roasting of sulphide ore to oxide
- c) Roasting of the sulphide to its oxide is thermodynamically feasible.
- d) Carbon and hydrogen are suitable reducing agents for metal sulphides.

[Ans. (d) Carbon and hydrogen are suitable reducing agents for metal sulphides.]

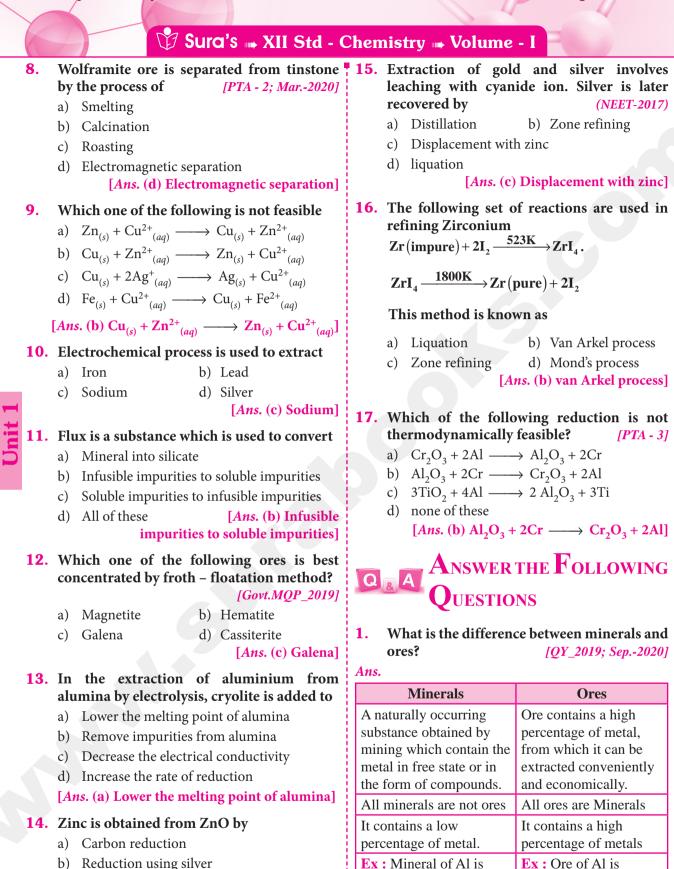
Match items in column - I with the items of column - II and assign the correct code.

	Column - I	Column - II		
Α	Cyanide process	(i)	Ultrapure Ge	
B	Froth floatation process	(ii)	Dressing of ZnS	
С	Electrolytic reduction	(iii)	Extraction of Al	
D	Zone refining	(iv)	Extraction of Au	
		(v)	Purification of Ni	
	A B C D)		

(a) (i) (ii) (iii) (iv)

- **(b)** (iii) (iv) (v) (i)
- (c) (iv) (ii) (iii) (i)
- (d) (ii) (iii) (i) (v) [Ans. (c) (iv) (ii) (iii) (i)]

orders@surabooks.com



c) Electrochemical process

orders@surabooks.com

d) Acid leaching [Ans. (a) Carbon reduction]

Ph:9600175757 / 8124201000

bauxite

bauxite and china clay

4

🖞 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

- 2. What are the various steps involved in extraction of pure metals from their ores?
- *Ans.* The various steps involved in the extraction of pure metals from their ores are
 - (*i*) concentration of the ore
 - (ii) extraction of crude metal
 - (iii) refining of crude metal

3. What is the role of Limestone in the extraction of Iron from its oxide Fe₂O₃?

- *Ans.* (*i*) Lime stone acts as a Flux.
 - (*ii*) It combine with silica and get converted into Calcium silicate called as slag.

$$\begin{array}{c} \text{CaO}_{(s)} + \text{SiO}_{2(s)} \longrightarrow \text{CaSiO}_{3(s)} \\ \text{Flux} \quad \text{Gangue} \quad \text{Slag} \end{array}$$

- 4. Which type of ores can be concentrated by froth floatation method? Give two examples for such ores.
- *Ans. (i)* Sulphide ores can be concentrated by froth floatation method.
 - (*ii*) **Ex** : Lead sulphide **galena** (PbS) and zinc blende (ZnS).

5. Describe a method for refining nickel.

Ans. Mond process for refining nickel : [PTA - 3]

- (*i*) The impure metal is heated in a stream of carbon monoxide at around 350K.
- *(ii)* The nickel reacts with the CO to form a highly volatile nickel tetra carbonyl.
- (*iii*) The solid impurities are left behind.

$$\operatorname{Ni}_{(s)} + 4\operatorname{CO}_{(g)} \xrightarrow{350 \text{ K}} [\operatorname{Ni}(\operatorname{CO})_4]_{(g)}$$

(iv) On heating the nickel tetracarbonyl around 460K, the complex decomposes to give pure metal.

 $[\operatorname{Ni}(\operatorname{CO}_4)]_{(g)} \xrightarrow{460 \text{ K}} \operatorname{Ni}_{(s)} + 4\operatorname{CO}_{(g)}$

- 6. Explain zone refining process with an example. [PTA 6; Mar.-2020]
- *Ans. (i)* Zone refining method is based on principles of Fractional Crystallisation. When an impure metal is melted and allowed to solidify, the impurities will prefer to remain in the molten region.
 - (*ii*) The impure metal is taken in the form of a rod.
 - *(iii)* When the metal rod is heated with a heater the metal melts.

- *(iv)* The heater is slowly moved from one end to the other end.
- (*v*) The impurity dissolves in the molten zone.
- (*vi*) When the heater moves, the molten zone also moves.
- (vii) This process is repeated again and again to get pure metal.
- (*viii*) The process is carried in an inert gas atmosphere to prevent oxidation of metal.

Example :

(*i*) Elements such as Germanium (Ge), Silicon (Si) and Galium (Ga) are refined using this process.

7. Give the uses of zinc.

[PTA - 4]

- *Ans. (i)* Metallic zinc is used in **galvanising** metals such as iron and steel structures to protect them from rusting and corrosion.
 - *(ii)* Zinc is also used to produce die-castings in the automobile, electrical and hardware industries
 - (iii) Zinc oxide is used in the manufacture of many products such as paints, rubber, cosmetics, pharmaceuticals, plastics, inks, batteries, textiles and electrical equipment.
 - *(iv)* Zinc sulphide is used in making luminous paints, fluorescent lights and x-ray screens.
 - (v) Brass an alloy of zinc is used in water valves and communication equipment as it is highly resistant to corrosion.

8. Explain the following terms with suitable examples. [PTA - 2; Sep.-2020]

- (i) Gangue (ii) Slag
- Ans. (i) Gangue : The non-metallic impurities, rocky materials and siliceous matter, associated with the ore is called gangue.
 Example : SiO₂ is the gangue present in the iron ore.
 - *(ii)* **Slag :** Slag is the fusible product formed when flux reacts with gangue during the extraction of metal.

Example :

$$\begin{array}{ccc} \text{CaO}_{(s)} + \text{SiO}_2 & \longrightarrow & \text{CaSiO}_3 \\ \text{Flux} & \text{Gangue} & (\text{slag}) \end{array}$$

orders@surabooks.com

5

🗘 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

- 9. Give the basic requirement for vapour phase refining.
- *Ans.* (*i*) The metal should form a volatile compound when treated with a suitable reagent.
 - *(ii)* Then the volatile compound is decomposed to give the pure metal.
- **10.** Describe the role of the following in the process mentioned.
 - (i) Silica in the extraction of copper.
 - (ii) Cryolite in the extraction of a luminium. $[QY_2019]$
 - (iii) Iodine in the refining of Zirconium. $[QY_2019]$

(iv) Sodium cyanide in froth floatation.

- *Ans.* (*i*) Silica, is used as an acidic flux is used to remove slag during the process of roasting.
 - *(ii)* Lowers the melting point to 1173K and improves the electrical conductivity of the aluminium.
 - (iii) To form a volatile compound which on further heating decomposes to give pure Zn.
 - (iv) Sodium cyanide is used as an depressing agent in froth flotation. It prevents other metal sulphides coming to the froth.
 For Example, when impurities such as ZnS is present in galena (PbS), sodium

cyanide (NaCN) is added to depresses the flotation property of ZnS by forming a layer of zinc complex $Na_2[Zn(CN)_4]$ on the surface of zinc sulphide.

- **11.** Explain the principle of electrolytic refining with an example. [HY_2019]
- Ans. (i) Electrolytic refining is carried out in an electrolytic cell.
 - (ii) Anode : Impure metal
 Cathode : Thin strips of pure metal
 Electrolyte : Aqueous solution of the salts of the metal.
 - (*iii*) The metal of interest dissolves from the anode, pass into the solution while the same amount of metal ions from the solution will be deposited at the cathode.
 - *(iv)* During electrolysis, the less electropositive impurities in the anode, settle down at the bottom and are removed as anode mud.

- (v) Electrolytic refining of silver as an example.
 Cathode : Pure silver
 Anode : Impure silver rods
 Electrolyte : Acidified aqueous solution of silver nitrate.
- (vi) When a current is passed through the electrodes the following reactions will take place

Reaction at anode

$$Ag_{(s)} \longrightarrow Ag^{+}_{(aa)} + 1e^{-}$$

Reaction at cathode

$$Ag^+_{(aq)} + 1e^- \longrightarrow Ag_{(s)}$$

- (*vii*) During electrolysis, at the anode the silver atoms lose electrons and enter the solution.
- (*viii*) The positively charged silver cations migrate towards the cathode and get discharged by gaining electrons and deposited on the cathode.

Evaluate Yourself

1. Write the equation for the extraction of silver by leaching with sodium cyanide and show that the leaching process is a redox reaction.

Ans. $Ag_2S + 4NaCN \longrightarrow 2Na [Ag(CN)_2] + Na_2S$

 $2Na[Ag(CN)_2] + Zn \longrightarrow Na_2[Zn(CN)_4] + 2Ag$

- (*i*) Silver is leached with an aqueous solution of NaCN to form soluble sodium dicyano argentite.
- *(ii)* Silver is recovered from dicyano argentite. Silver is reduced to its elemental state.
- 2. Magnesite (Magnesium carbonate) is calcined to obtain magnesia, which is used to make refractory bricks. Write the decomposition reaction.

Ans. MgCO₃
$$\longrightarrow$$
 MgO + CO_{2(g)} \uparrow

6

orders@surabooks.com

Ph:9600175757 / 8124201000

Unit]

🖞 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

PTA MODEL QUESTIONS AND ANSWERS

1.

CHOOSE THE CORRECT ANSWER

1 MARK Answer The Questions

Define roasting.

3 MARKS

1. Sulphide ores of metals are usually concentrated by froath floatation process. Which one of the following sulphide ore offers an exception and is concentrated by chemical leaching. [PTA - 4]

- a) Argentite b) galena
- c) Copper pyrites d) Sphalerite [Ans. (a) Argentite]
- 2. Which method of purification represented by the equation? [PTA 5]

 $Ti(impure) + 2l_2 \xrightarrow{550K} Til_4$

 $\underbrace{1800K}{} Ti(pure) + 2I_2$

- a) Cupellation
- b) Zone refining
- c) Van Arkel method
- d) Mond's process

[Ans. (c) Van - Arkel method]

- **3**. The process of converting hydrated aluminia into anhydrous alumina is called. [PTA - 6]
 - a) Roasting
- b) Smelting
- c) Auto-reduction d) Calcination [*Ans.* (d) Calcination]

Answer The Questions

- 2 MARKS
- 1. What is the role of depressing agent in froath floatation process? [PTA 1]
- *Ans.* When impurities such as ZnS is present in galena (PbS), sodium cyanide (NaCN) is added to depresses the flotation property of ZnS by forming a layer of zinc complex $Na_2[Zn(CN)_4]$ on the surface of zinc sulphide.
- 2. Describe the underlying principle of froth floatation process. [PTA 3]
- *Ans.* Froth floatation process is based on the principle that the ore particles are wetted by oil and the gangue particles by water. This is used for the concentration of sulphide ores.

[PTA - 4]

Ans. In roasting, the concentrated ore is oxidised by heating it with excess of oxygen in a suitable furnace below the melting point of the metal.

 $2Pbs + 3O_2 \xrightarrow{\Delta} 2PbO + 2SO_2 \uparrow$

2. Explain calcination with an example.

- *Ans. (i)* Calcination is the process in which the concentrated ore is strongly heated in the absence of air.
 - (*ii*) During this process, the water of crystallisation present in the hydrated oxide escapes as moisture.
 - *(iii)* Any organic matter (if present) also get expelled leaving behind a porous ore.
 - *(iv)* This method can also be carried out with a limited supply of air.
 - (v) During calcination of carbonate ore, carbon dioxide is expelled

$$PbCO_3 \xrightarrow{\Delta} PbO + CO_2\uparrow$$

Answer The Questions

5 MARKS

1. Explain electrolytic refining of silver.

[PTA - 5]

Metallurgy

- Ans. (i) Electrolytic refining of silver as an example.
 Cathode : Pure silver
 Anode : Impure silver rods
 Electrolyte : Acidified aqueous solution of silver nitrate.
 - *(ii)* When a current is passed through the electrodes the following reactions will take place

Reaction at anode

 $Ag_{(s)} \longrightarrow Ag^{+}_{(aq)} + 1e^{-}$

Reaction at cathode

$$\operatorname{Ag}^{+}_{(aq)} + 1e^{-} \longrightarrow \operatorname{Ag}_{(s)}$$

orders@surabooks.com

7

🖞 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

- 2. Explain extraction of copper from copper pyrites. [PTA 5]
- *Ans. (i)* In this method, a flux (a chemical substance that forms an easily fusible slag with gangue) and a reducing agent such as carbon, carbon monoxide (or) aluminium is added to the concentrated ore and the mixture is melted by heating at an elevated temperature (above the melting point of the metal) in a smelting furnace.
 - (*ii*) For example the oxide of iron can be reduced by carbon monoxide as follows. $Fe_2O_{3(s)} + 3CO_{(g)} \longrightarrow 2Fe_{(s)} + 3CO_{2(g)\uparrow}$
 - (iii) In this extraction, a basic flux, limestone (CaO) is used. Since the silica gangue present in the ore is acidic in nature, the limestone combines with it to form calcium silicate (slag).

$$CaO_{(s)} + SiO_{2(s)} \longrightarrow CaSiO_{3(s)}$$

Flux Gangue Slag

- *(iv)* In the extraction of copper from copper pyrites, the concentrated ore is heated in a reverberatory furnace after mixing with silica, an acidic flux.
- (v) The ferrous oxide formed due to melting is basic in nature and it combines with silica to form ferrous silicate (slag).

(vi) The remaining metal sulphides Cu_2S and FeS are mutually soluble and form a copper matte.

$$2\text{CuFeS}_{2(s)} + \text{O}_{2(g)} \xrightarrow{2\text{FeS}_{(l)}} + \text{Cu}_{2}\text{S}_{(l)} + \text{SO}_{2(g)}$$

$$2\text{FeS}_{(l)} + 3\text{O}_{2(g)} \longrightarrow 2\text{FeO}_{(l)} + 2\text{SO}_{2(g)}$$

$$\text{FeO}_{(s)} + \text{SiO}_{2(s)} \longrightarrow \text{FeSiO}_{3(s)}$$

$$\text{Flux Gangue Slag}$$

- (*vii*) The matte is separated from the slag and fed to the converting furnace.
- (*viii*) During conversion, the FeS present in the matte is first oxidised to FeO.
- *(ix)* This is removed by slag formation with silica.
- (x) The remaining copper sulphide is further oxidised to its oxide which is subsequently converted to metallic copper.

$$2\operatorname{Cu}_{2}\operatorname{S}_{(l,s)} + 3\operatorname{O}_{2(g)} \longrightarrow 2\operatorname{Cu}_{2}\operatorname{O}_{(l,s)} + 2\operatorname{SO}_{2(g)}$$
$$2\operatorname{Cu}_{2}\operatorname{O}_{(l)} + \operatorname{Cu}_{2}\operatorname{S}_{(l)} \longrightarrow 6\operatorname{Cu}_{(l)} + \operatorname{SO}_{2(g)}$$

(xi) The metallic copper is solidified and it has blistered appearance due to evolution of SO_2 gas formed in this process. This copper is called blistered copper.

GOVERNMENT EXAM QUESTIONS AND ANSWERS

8

CHOOSE THE CORRECT ANSWER

1. The metal which is used in packing metarial for food items : [Sep.-2020] (a) Zn (b) Zr (c) Al (d) A

Answer The Questions

2 MARKS

[Ans. (c) Al]

1 MARK

- 1. Oxides like Ag₂O and HgO undergo self reduction. Why? [QY_2019]
- Ans. Decomposition temperature of Ag_2O and HgOare 600 and 700 K respectively.

These oxides are unstable at moderate temperature so undergo self reduction.

- 2. Name the collector and depressing agent used in froth flotation process. [HY_2019]
- *Ans. (i)* Sodium ethyl xanthate acts as a collector.
 - (*ii*) Sodium cyanide, Sodium carbonate are used as depressing agents in froth flotation process.

Answer The Questions

3 MARKS

- 1. Explain how gold ore is leached by cyanide process [Govt.MQP_2019]
- *Ans.* (*i*) Gold is usually found in native state.
 - *(ii)* The leaching process is intended to concentrate the gold metal.

$$4\operatorname{Au}_{(s)} + 8 \operatorname{NaCN}_{(aq)}^{-} + 2\operatorname{H}_{2}\operatorname{O}_{(aq)} + \operatorname{O}_{2(g)} \longrightarrow$$
$$4 \operatorname{Na} \left[\operatorname{Au}(\operatorname{CN})_{2}\right]_{(aq)}^{-} + 4\operatorname{NaOH}_{(aq)}^{-}$$

 $2Na [Au(CN)_2] + Zn \longrightarrow$

 $Na_2[Zn(CN)_4] + 2Au \downarrow$

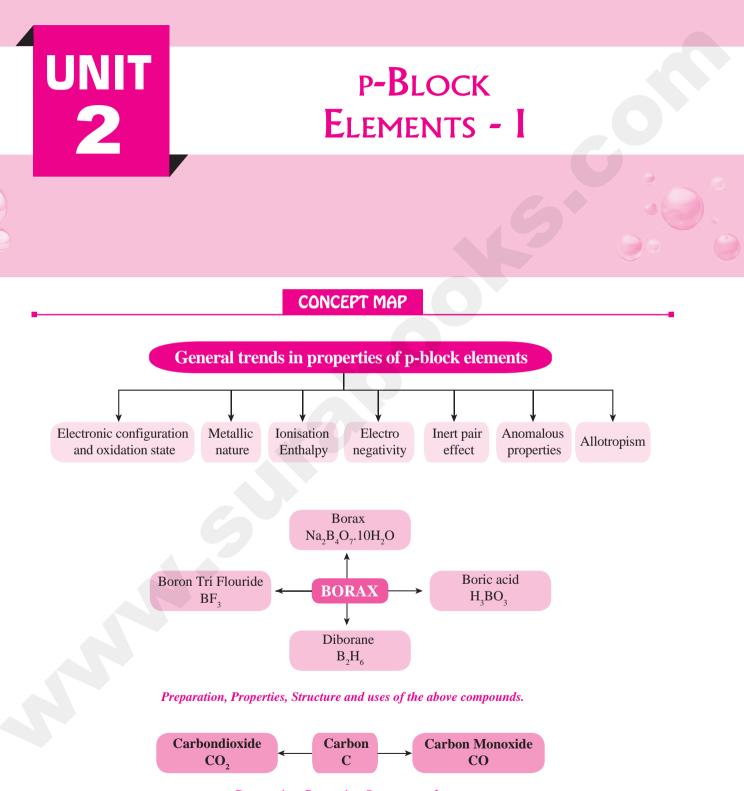
(*iii*) In this reaction, gold is reduced to its elemental state and the process is called cementation.

 $\diamond \diamond \diamond$

orders@surabooks.com

Ph:9600175757 / 8124201000

Unit



Preparation, Properties, Structure and uses.

orders@surabooks.com

🖞 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

FORMULAE TO REMEMBER

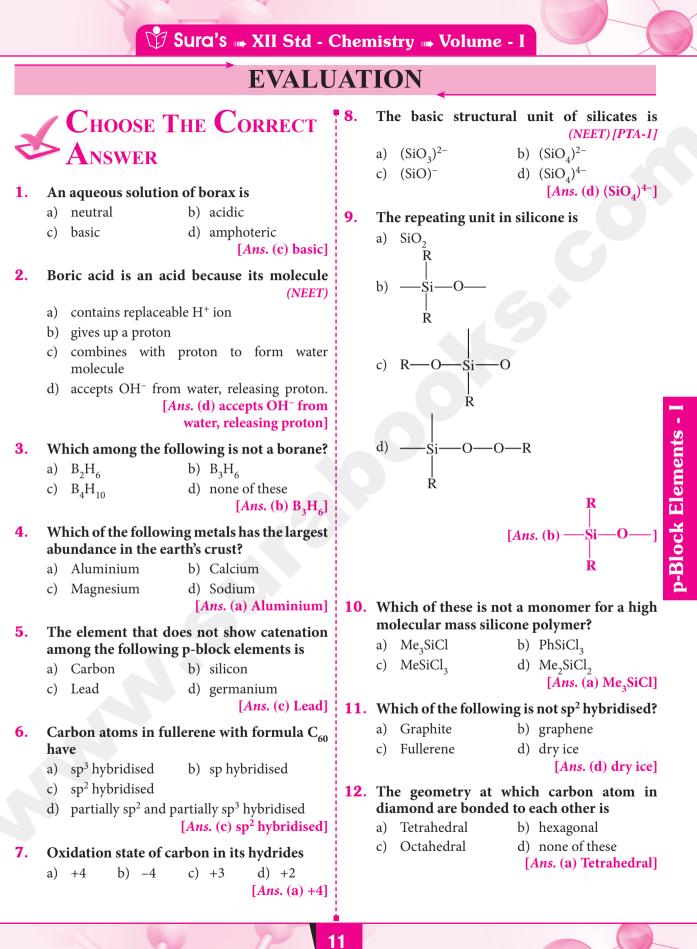
Borax	: $Na_2B_4O_7.10H_2O$
Boric acid	: H_3BO_3 or $B(OH)_3$
Di Borane	: B ₂ H ₆
Boron Fluoride	: BF ₃
Aluminium Chloride	: AlCl ₃
Silicon tetrachloride	: SiCl ₄
Silicones	: R ₂ SiO
Potash Alum	: $K_2SO_4.Al_2(SO_4)_3.24H_2O$
Sodium Alum	: $Na_2SO_4.Al_2(SO_4)_3.24H_2O$
Ammonium Alum	: $(NH_4)_2SO_4.Al_2(SO_4)_3.24H_2O$
Chrome Alum	: $K_2SO_4.Cr_2(SO_4)_3.24H_2O$
Nickel Tetra Carbonyl	: [Ni(CO) ₄]
Iron Penta Carbonyl	: $[Fe(CO)_5]$
Chromium Hexa Carbonyl	: $[Cr(CO)_6]$
Thortveitite	: $Sc_2Si_2O_7$
Spodumene	: $LiAl(SiO_3)_2$

MUST KNOW DEFINITIONS

Unit 2

Metallic Character	:	The tendency of an element to form a cation by loosing electrons is known as electropositive or metallic character.	
Allotropism		Some elements exist in more than one crystalline or molecular forms in the same physical state. This property is called allotropism .	
Hydroboration		Diborane adds on to alkenes and alkynes in ether solvent at room temperature. This reaction is called hydroboration .	
Catenation	:	Catenation is an ability of an element to form chain of atoms.	
Silcones	:	Silicones or poly siloxanes are organo silicon polymers with general empirical formula (R_2 SiO).	
Silicates	:	The mineral which contains silicon and oxygen in tetrahedral $[SiO_4]^{4-}$ units linked together in different patterns are called silicates.	
Zeolites	:	Zeolites are three dimensional crystalline solids containing aluminium, silicon, and oxygen in their regular three dimensional framework.	
Inert pair effect	:	In heavier post-transition metals, the outer s electrons (ns) have a tendency to remain inert and show reluctance to take part in the bonding, which is known as inert pair effect.	

orders@surabooks.com



orders@surabooks.com

								4 4
		/		Ŵ	Sura's 🛶 XII Std - C	Chen	nistr	ry 🖦 Volume - I
		a) b) c) d) Ma colu	rect? Beryl is a cy Mg_2SiO_4 is a SiO_4^{4-} is the Feldspar is n [<i>Ans.</i> (d) F tch items in	follo clic si un ort basic tot alu celdsp	wing statements is not			
		D	Borax	4	SiO ₂		(iii)	Absence of d-orbitals in their valance shell.
Unit 2		ABCD(a) 2 1 4 3 (b) 1 2 4 3 (c) 1 2 3 4 (d)None of these[Ans. (a) 2, 1, 4, 3] 5. Duralumin is an alloy ofa)Cu, Mnb)Cu, Al, Mgc)Al, Mnd)Al, Cu, Mn, Mg[Ans. (d) Al, Cu, Mn, Mg] 6. The compound that is used in nuclear reactors as protective shields and control rods isa)Metal boridesb)Metal oxidesc)Metal carbonates				 Describe briefly allotropism in p- block elements with specific reference to carbon. Ans. Allotropism in p- block elements : (i) Some elements exist in more than one crystalline or molecular forms in the same physical state. (ii) Carbon exists as diamond and graphite. This phenomenon is known as allotropism. (iii) Other important allotropes of carbon are graphite, fullerenes, carbon nanotubes. Give the uses of Borax. [HY_2019] Ans. (i) Borax is used for the identification of coloured metal ions. (ii) In the manufacture optical and borosilicate glass, enamels and glazes for pottery. (iii) It is also used as a flux in metallurgy and also acts as a preservative. What is catenation? Describe briefly the 		
	17.	The in t a) b) c)	Metal carbic e stability of he sequence Al < Ga < In Tl < In < Ga In < Tl < Ga Ga< In < Al	f +1 c c < Tl c < Al c < Al c < Tl	[<i>Ans.</i> (a) Metal borides] oxidation state increases [QY_2019] Ans. (a) Al < Ga < In < Tl]	Ans.		 nation property of carbon. [Sep2020] Catenation is an ability of an element to form chain of atoms. The following conditions are necessary for catenation. Mar2020] (a) The valency of element is greater than or equal to two, (b) Element should have an ability to bond with itself (c) The self bond must be as strong as its bond with other elements (d) Kinetic inertness of catenated compound towards other molecules.

12

orders@surabooks.com

This i ores

is Only for Sample For Full Book Order Online or Available at All Leading Booksto						
	🗘 Sura's 🛶 XII Std - O	Chemis	try 🖦 Volume - I			
(iii)	Carbon possesses all the above properties	8 . Co	omplete the following reactions			
	and forms a wide range of compounds with itself and with other elements such as H, O,	a.	$B(OH)_3 + NH_3 \longrightarrow$			
	N, S and halogens.	b.	$Na_2B_4O_7 + H_2SO_4 + H_2O \longrightarrow$			
Give	the uses of silicones.	c.	$B_2H_6 + 2NaOH + 2H_2O \longrightarrow$			
. (i)	Silicones are used for low temperature	d.	$B_2H_6 + CH_3OH \longrightarrow$			
	lubrication and in vacuum pumps, high	e.	$BF_3 + 9H_2O \longrightarrow$			
(ii)	temperature oil baths etc. They are used for making water proofing	f.	$\text{HCOOH} + \text{H}_2\text{SO}_4 \longrightarrow$			
()	clothes.	g	$SiCl_4 + NH_3 \longrightarrow [QY_2019]$			
(iii)	They are used as insulting material in electrical	h.	$SiCl_4 + C_2H_5OH \longrightarrow$			
(iv)	motor and other appliances. They are mixed with paints and enamels to	i.	$B + NaOH \longrightarrow [QY_2019]$			
()	make them resistant towards high temperature,	j.	$H_2B_4O_7 \xrightarrow{\text{Red hot}}$			
	sunlight, dampness and chemicals.	Ans. a.	$B(OH)_3 + NH_3 \longrightarrow BN + 3H_2O$			
	one example for each of the following					
(i)	icosogens (ii) tetragen	b.	$Na_2B_4O_7 + H_2SO_4 + 5H_2O \longrightarrow Na_2SO_4 + 4H_3BO_3$			
(iii)	prictogen (iv) chalcogen					
. (i) (ii)	icosogens - Boron tetragen - Carbon	C.	$B_2H_6 + 2NaOH + 2H_2O \longrightarrow$ $2NaBO_2 + 6H_2$			
(iii)	prictogen - Nitrogen	d.				
(iv)	chalcogen - Oxygen	u.	$B_2H_6 + 6CH_3OH \longrightarrow 2B(OCH_3)_3 + 6H_2$			
	e a note on metallic nature of p-block	e.	$4BF_3 + 3H_2O \longrightarrow H_3BO_3 + 3HBF_4$			
	The ten denses of an element to form a set in	f.	$\text{HCOOH} + \text{H}_2\text{SO}_4 \longrightarrow$			
. (i)	The tendency of an element to form a cation by losing electrons is known as electropositive		$CO + H_2O + H_2SO_4$			
	or metallic character.	g	$2\text{SiCl}_4 + \text{NH}_3 \xrightarrow{335\text{K}} \text{Cl}_3\text{Si} - \text{NH} - \text{SiCl}_3$			
(ii)	This character depends on the ionisation		ether			
(iii)	energy. Generally on moving down a group ionisation	h.	$SiCl_4 + 4C_2H_5OH \longrightarrow$			
()	energy decreases and hence the metallic		$Si(OC_2H_5)_4 + 4HCl$			
(\cdot, \cdot)	character increases.	i.	$2B + 6NaOH \longrightarrow 2Na_3BO_3 + 3H_2$			
(iv)	In p-block, the elements present in lower left part are metals while the elements in	j.	$H_2B_4O_7 \xrightarrow{\text{Red hot}} 2B_2O_3 + H_2O_3$			
	the upper right part are non metals.	ישי ו ו	2 + 1 2-3 2-			

How will you identify borate radical?

[PTA - 5; Govt.MQP_2019; QY_2019]

- Ans. (i) When boric acid or borate salt is heated with ethyl alcohol in presence of conc. sulphuric acid, an ester, triethylborate is formed.
 - (ii) The vapour of this ester burns with a green edged flame and this reaction is used to identify the presence of borate.

$$H_3BO_3 + 3C_2H_5OH \xrightarrow{Conc.} B(OC_2H_5)_3 + 3H_2O$$

orders@surabooks.com

Non-metals

В

С

O, S, Se

F, Cl, Br, I

He, Ne, Ar, Kr, Xe

5. Ans.

6.

Ans.

7.

Ans.

Group

13

14

15

16

17

Metals

Al, Ga, In, Tl

Sn, pb

Bi

-

_

13

9.

Metalloids

-

Si, Ge

Te, Po

_

_

Ph:9600175757 / 8124201000

p-Block Elements -

🗘 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

10. Write a note on zeolites. *[PTA - 2; QY_2019]* **12.** A

- *Ans. (i)* Zeolites are three-dimensional crystalline solids containing aluminium, silicon, and oxygen in their regular three dimensional framework.
 - (ii) They are hydrated sodium alumino silicates with general formula $Na_2O.(Al_2O_3).x(SiO_2).yH_2O$ (x = 2 to 10; y = 2 to 6).
 - (*iii*) Zeolites have porous structure in which the monovalent sodium ions and water molecules are loosely held.
 - *(iv)* The Si and Al atoms are tetrahedrally coordinated with each other through shared oxygen atoms.
 - (*v*) Zeolites are similar to clay minerals but they differ in their crystalline structure.
 - (*vi*) Zeolites have a three dimensional crystalline structure looks like a honeycomb consisting of a network of interconnected tunnels and cages.
 - (vii) Water molecules moves freely in and out of these pores but the zeolite framework remains rigid.
 - (viii) Another special aspect of this structure is that the pore/channel sizes are nearly uniform, allowing the crystal to act as a molecular sieve.
 - *(ix)* The removal of permanent hardness of water using zeolites.
- 11. How will you convert boric acid to boron nitride? [PTA 3]

Ans. Fusion of urea with B(OH)₃, in an atmosphere of ammonia at 800 - 1200 K gives boron nitride.

$$B(OH)_3 + NH_3 \xrightarrow{\Delta} BN + 3H_2O$$

[*PTA - 2; QY_2019*] **12.** A double salt which contains fourth period alkali metal (A) on heating at 500K gives (B). aqueous solution of (B) gives white precipitate with BaCl₂ and gives a red colour compound with alizarin. Identify A and B.

Ans. (i) A double salt which contains 4th period alkali metal (A) is

$$K_2SO_4$$
· $Al_2(SO_4)_3$ ·24 H_2O

(ii)
$$K_2SO_4Al_2(SO_4)_3.24H_2O \xrightarrow{500K}$$

 $K_{2}SO_{4}.Al_{2}(SO_{4})_{3} + 24H_{2}O_{4}$

А	K ₂ SO ₄ .Al ₂ (SO ₄) ₃ 24H ₂ O	Potash alum
В	$K_2SO_4.Al_2(SO_4)_3.$	Burnt alum

Evaluate Yourself

- 1. Why group 18 elements are called inert gases? Write the general electronic configuration of group 18 elements.
- Ans. (i) Group 18 consists of 6 elements, helium, neon, argon, krypton, xenon and radon. All these are gases and almost chemically inert. Since they have completely filled valence shell. (least reactive elements). Hence they are called inert gases.
 - (ii) The general electronic configuration of group 18 elements is ns², np⁶.

PTA MODEL QUESTIONS AND ANSWERS

Сн	IOOSE THE COP	RRECT ANSWER			1 MARK
1.	Elements like	silicon and Germ	anium to be used	as a semiconduct	tor is purified by [PTA - 1]
	a) heating un	ider vaccum	b) Van - Arl	kel method	
	c) Zone refin	ing	d) Electroly	sis	[Ans. (c) Zone refining]
2.	The basicity	of hypophosphoru	is acid is		[PTA - 2]
	a) 1	b) 2	c) 3	d) 4	[Ans. (a) 1]

14

orders@surabooks.com

Ph:9600175757 / 8124201000

Unit 2

🗊 Sura's 🛶 XII Std - Chemistry 🛶 Volume - I

- Ortho boric acid on dehydration at 373 K
 Answer The QUESTIONS produces mainly
 [PTA 3]
 AlCl₂ is more stable w
 - a) metaboric acid b) boric anhydride
 - c) Boron metal and Oxygen
 - d) tetraboric acid. [Ans. (a) metaboric acid]
- **4.** On hydrolysis BF₃ gives Boric acid and converted to fluroboric acid. The fluoroboric acid contains the species. [*PTA* 6]

a)
$$H^+$$
, F^- & BF_3 b) H^+ & $[BF_4]^-$

c) $[H BF_3]^+ \& F^-$ d) $H^+, B^{3+} \& F^-$

 $[Ans. (b) H^+ \& [BF_4]^-]$

2 MARK

2.

Answer The Ouestions

- 1. Nitrogen does not form any penta halides like phosphorus. why? [PTA 4]
- *Ans.* Nitrogen does not form penta halides because it cannot extend its valency upto 5 due to unavailability of d-orbitals.
- 2. Why formic acid act as strong reducing agent? Give one equation to show its reducing property. [PTA - 6]
- *Ans. (i)* Formic acid (HCOOH) is unique because it contains both an aldehyde group and carboxyl group also.
 - *(ii)* Hence it can act as a **reducing agent**. It reduces Fehling's solution Tollen's reagent and decolourises pink coloured KMnO₄ solution.
 - (*iii*) Whereas in acetic acid, there is no aldehyde group and it cannot act as reducing agent.
 - (*iv*) Formic acid reduces ammoniacal silver nitrate solution (Tollen's reagent) to metallic silver. $HCOOH + Ag_2O \longrightarrow H_2O + CO_2 + 2Ag \downarrow$ (metallic silver)
 - (v) Formic acid reduces Fehling's solution. It reduces blue coloured cupric ions to red coloured cuprous ions.

. AlCl₃ is more stable where as TlCl₃ is highly unstable. Why? [PTA - 2]

3 MARKS

- Ans. (i) Aluminium(III)chloride is stable whereas thallium(III)chloride is highly unstable and disproportionates to thallium(I) chloride and chlorine gas.
 - (ii) This shows that in thallium the stable lower oxidation state corresponds to the loss of np electrons only and not ns electrons.
 - (iii) Thus in heavier posttransition metals, the outer s electrons (ns) have a tendency to remain inert and show reluctance to take part in the bonding, which is known as inert pair effect.

How is potash Alum prepared? [PTA - 4]

Ans. The alunite the alum stone is the naturally occurring form and it is K_2SO_4 . $Al_2(SO_4)_3$. $4Al(OH)_3$. When alum stone is treated with excess of sulphuric acid, the aluminium hydroxide is converted to aluminium sulphate. A calculated quality of potassium sulphate is added and the solution is crystallised to generate potash alum. It is purified by recrystallisation.

 $K_2SO_4.Al_2(SO_4)_3.4Al(OH)_3 + 6H_2SO_4 \longrightarrow$

$$K_2SO_4 + 3Al_2(SO_4)_3 + 12H_2O_4$$

 $K_2SO_4 + Al_2(SO_4)_3 + 24H_2O \longrightarrow$ $K_2SO_4.Al_2(SO_4)_3.24H_2O$

3. What are amphiboles? Give example.[*PTA*-5]

Ans. Double chain silicates (or amphiboles): These silicates contains $[Si_4O_{11}]_n^{6n-}$ ions. In these silicates there are two different types of tetrahedra :

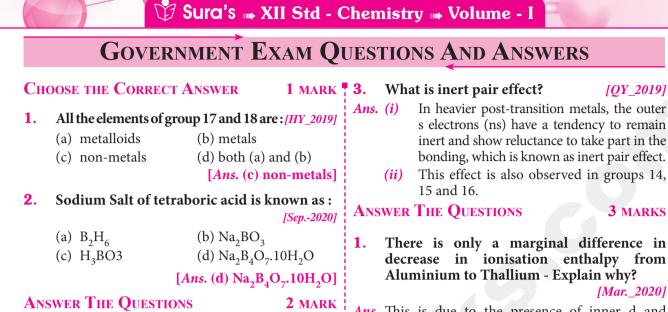
- (a) Those sharing 3 vertices
- (*b*) those sharing only 2 vertices. **Examples :**

Asbestos : These are fibrous and non-combustible silicates.

p-Block Elements

orders@surabooks.com

15



Ans. This is due to the presence of inner d and f-electrons which has poor shielding effect compared to s and p-electrons.

Answer The Ouestions 5 MARKS

Write the preparation of potash alum. 1.

[HY 2019] Ans. $K_2SO_4.Al_2(SO_4)_3.4Al(OH)_3 + 6H_2SO_4 \longrightarrow$ $K_{2}SO_{4} + 3Al_{2}(SO_{4})_{3} + 12H_{2}O$

$$K_{2}SO_{4} + Al_{2}(SO_{4})_{3} + 24H_{2}O \longrightarrow$$
$$K_{2}SO_{4}.Al_{2}(SO_{4})_{3}.24H_{2}O$$

- 1. Although Graphite and Diamond are allotropes of carbon, graphite is soft whereas diamond is hard. why? [QY_2019]

Ans. Both diamond and graphite are made of carbon. In diamond, each carbon atom forms 4 covalent bonds in a tetrahedral structure. Whereas, in graphite the carbon atoms only form 3 covalent bonds, creating hexagonally packed sheets of carbon. The sheets of carbon become bonded by weaker intermolecular forces. Which makes layers of graphite slide over each other, making the overall substance softer than diamond.

Give the uses of Potash alum. 2. [QY_2019]

- Ans. (i) It is used for purification of water.
 - It is also used for water proofing and *(ii)* textiles. It is used in dyeing, paper and leather tanning industries.
 - (iii) It is employed as a styptic agent to arrest bleeding.

 $\diamond \diamond \diamond$

16

3 MARKS

Ph:9600175757 / 8124201000

orders@surabooks.com