

# UNIVERSITY OF MUMBAI

## S.Y.B.Sc. CHEMISTRY Syllabus Credit Based and Grading System To be implemented from the Academic year 2012-2013

<b>Course Code</b>	UNIT	TOPICS	Credits	L / Week
	I	Chemical Thermodynamics		1
USCH301	Π	Solution of Electrolytes	2	1
	III	Introduction to Analytical Chemistry and Visible Spectroscopy		1
	I	Chemical bonding and Molecular structure		1
USCH302	п	Chemistry of transition Elements	2	1
	ш	Industrial Inorganic chemistry		1
	I	Nomenclature Mechanism of organic Reactions Aromatic Electrophilic Substitution Reactions		1
USCH303	II	Aromatic Hydrocarbons Haloarenes Phenols Ether and Epoxides		1
	ш	Sources of Organic compounds Chemical Industry Environmental aspects of chemical Industry		1
USCHP3	Practical	s based on both courses in theory	3	9

## SEMESTER III

## SEMESTER IV

Course Code	UNIT	TOPICS	Credits	L / Week
	Ι	Phases in Equilibria		1
USCH401	П	Electrochemical cells & Ionic equilibria	2	1
	III	Introduction to Titrimetric Analysis & use of Instrumental methods in titrimetric analysis		1
	Ι	Acids and Bases,Chemistry of elements and organometallic		1
USCH402	п	Chemistry of Group-17 & Group- 15 elements and Bio-inorganic chemistry		1
	III	Industrial inorganic chemistry		1
	Ι	Aromatic Nitrogen compounds Aromatic Aldehydes and Ketones		1
	II Aromatic a	Aromatic acids		1
USCH403	III	Stereochemistry Structure determination and Multistep synthesis Green Chemistry Manufacture of some bulk Chemicals including flow diagrams		1
USCHP4	Practical	s based on both courses in theory	3	9

Semester I		SEMESTER I	I		
CREDITS 3		CREDITS 3			
Type of Experiment	Expts	Type of Experiment	Expts		
Physical-		Physical-			
Instrumental	03	Instrumental	03		
Non Instrumental	02	Non Instrumental	02		
Organic		Organic			
Derivatives	07	Spotting	12		
Estimations	04				
Inorganic		Inorganic			
SMQA	08	Gravimetric	03		
		Volumetric	03		
	24		23		

In each semester, total marks for Practical Examination are 150.PRACTICAL EXAMINATION IS TO BE CONDUCTED FOR TWO HRS PER PAPER.

Instrumental: Semester I: 1. To verify Ostwald's dilution law conductometrically. 2. To determine amount of strong acid present in the given solution by titrating against stong base pHmetrically. 3. To determine standard emf of the cell potentiometrically.

Non-Instrumental: 1. To determine dissolved oxygen in the given waste water sample. 2. To carry out assay of commercial sample of Aspirin.

Semester II: Instrumental:Semester I: 1. To titrate strong acid against strong base conductometrically. 2. To determine dissociation constant of weak acid using Henderson's equation by the method of incomplete titration pHmetrically. 3. To determine  $\lambda_{max}$  and molar extinction coefficient for KMNO4 solution using photometer.

Non-Instrumental: 1. To investigate the reaction between equimolar concentration of  $K_2S_2O_8$  and KI. 2. To determine partition coefficient of oxalic acid/succinic acid between water and CCl<sub>4</sub>.

Semester III----- CREDITS 3

Session	External Component		Internal Component		
	Practical Examination				
	Experiment	Marks	Details	Marks	
1	Physical- Instrumental/	30	2 Best	10	
	Non Instrumental		Practicals	05	
	Non Instrumental		Journal	05	
			Journal	05	
			Viva-		
			Voce		
	Organic	30	2 Best	10	
	Derivatives/Estimations		Practicals	05	
			Journal	05	
			Journal	05	
			Viva-		
			Voce		
111	Inorganic SMQA	30	2 Best	10	
			Practicals	05	
			Journal	05	
				05	
			Viva-		
			Voce		
		90		60	

#### Semester I V----- CREDITS 3

Session	External Component	External Component		
	Practical Examination			
	Experiment	Marks	Details	Marks
I	Physical- Instrumental/	30	2 Best	10
	Non Instrumental		Practicals	05
			Journal	05
			Viva-Voce	
II	Organic Spotting	30	2 Best	10
			Practicals	05
			Journal	05
			Viva-Voce	
	Inorganic	30	2 Best	10
	Gravimetric/Volumetric		Practicals	05
			Journal	05
			Viva-Voce	
		90		60

## **SEMESTER III**

Course Code	UN IT	TOPICS	Cr edi ts	L / Wee k
	Ι	<ol> <li>Chemical Thermodynamics: 15L</li> <li>Gibbs free energy and He,mboltz's free energy, variation of Gibbs's free energy with temperature and pressure, Gibb's-Helmholtz equation.</li> <li>Physical equilibbria involving pure substances, Clayperon's equation and variation of vapour pressure with temperature, Clausius-Clayperon equation and its application. Partial molal properties, partial molal valume and chemical potential, Gibbs-Duhem equation.</li> <li>Variation of chemical potential potential with pressure and temperature, fugacity, activity and their relationship with chemical potential, activity and activity coefficient.</li> <li>Thermodynamic derivation of Law of mass action, Kp, Kc and their inter-relation, van't Hoff's reaction isotherm and reaction isochore.</li> </ol>		1
USCH 301	Π	<ul> <li>Solutions of Electrolyte: (15L)</li> <li>1) Electronic and electrolytic conductors, conductance, specific conductance, equivalent conductance, molar conductance, determination of molar conductance, variation of molar conductance with concentration for strong and weak electrolytes, concept of limiting molar conductance,</li> <li>2) Debye-Huckel theory of conductance of strong electrolytes, ionic atmosphere, relaxation effect, electrophoretic effect.</li> <li>3) Kohlrausch's law of independent migration of ions.</li> <li>Applications of Kohlrausch's law</li> <li>a) Determination of limiting molar conductance of a weak electrolyte.</li> <li>b) Determination of dissociation constant of a weak acid.</li> <li>c) Determination of solubility product of a sparingly soluble salt.</li> <li>4) Migration of ions, transport number, dependence</li> </ul>	2	1

<ul> <li>of transport number on the velocity of the ion, Hittorf's rule for unattackable electrodes illustrate with ions having equal and unequal speeds, moving boundary method for the determination of the transport number, factors affecting the transport number of an ion.</li> <li>5) Relationship between transport number ionic mobility, and equivalent conductance.</li> <li>Introduction to Analytical Chemistry and Visble</li> </ul>	
<ul> <li>Introduction to Ynaryteer Chemistry and Visite Spectroscopy (15L)</li> <li>(A) Introduction to analytical chemistry</li> <li>1) Analytical chemistry, qualitative and quantitative analysis, analytical chemistry and chemical analytical methods with emphasis on detection limit and sensitivity.</li> <li>2) Steps involved in chemical analysis: sampling, obtaining a sample, processing the sample, the sample, selection of a method for analysis, calibration and actual analysis, data collection, data processing, presentation of results and interpretation.</li> <li>3) Performance characteristics of an analytical method: accuracy, precision, detection limit, dynamic range, sensitivity, selectivity.</li> <li>4) Quantitative analysis with calibration curve and standard addition method.</li> <li>5) Applications of analytical methods in various fields such as organic, pharmaceuticals, electronic and environmental analysis.</li> <li>(B)Visible Spectroscopy</li> <li>1) U. V. and Visible spectroscopy, absorption spectroscopy, terms involved; radiant power, absorbance, transmittance, percentage transmittance, wavelength of maximum absorption.</li> <li>2) Statement of the Beer's law and the Lambert's law, (derivation expected) combined expression, molar extinction coefficient deviations from the Beer-Lambert's Law, limitations.</li> <li>3) Components of an optical instrument, photometers and spectrophotometers, construction of a single beam photometer.</li> </ul>	1

USCH 302	Ι	<ul> <li>Chemical Bonding and Molecular Structure 15L</li> <li>1. Valence Bond Theory (VBT) 5L</li> <li>(i) Hydridization involving the use of 'd' orbitals as in BaCl2 (sd), MnO4 (sd3) and PCI5 (sp3d)</li> <li>(ii) Concept of Resonance and Resonance Energy, Formal Charge, with examples.</li> <li>(iii) Limitations of VBT.</li> <li>2. Molecular Orbital Theory (MOT): 10L</li> <li>(i) Conditions for the formation of molecular orbitals.</li> <li>(ii) Linear combination of atomic orbitals to obtain molecular orbitals (LCAO-MO approach)</li> <li>(iii) Application of LCAO-M) approach to the formation of: <ul> <li>a. Homo-nuclear diatomic molecules: H<sub>2</sub>, He<sub>2</sub>, Li<sub>2</sub>, Be<sub>2</sub>, C<sub>2</sub>, N<sub>2</sub>, O<sub>2</sub>, F<sub>2</sub> and Ne<sub>2</sub>. (Discussion should include 2s-2p interaction; stabilization of p(2p<sub>x</sub>, 2p<sub>y</sub> orbitals) Mos in B<sub>2</sub>, C<sub>2</sub> and N<sub>2</sub> with experimental evidences; bond order and correlation with stability, bond length, bond energy and magnetic properties of all the homo-nuclear diatomic molecular ions in (O<sub>2</sub>) + (PtF<sub>6</sub>)', KO<sub>2</sub> and Na<sub>2</sub>O<sub>2</sub>.</li> <li>c. Hetero-nuclear diatomic molecules or molecular ions: CO, No, CN-, and HCI. (Discussion should include comparison with the approach for homo-nuclear diatomic molecules or molecular ions of bond order. Stability, magnetic and polarity)</li> </ul></li></ul>	2	1
	II	<ol> <li>Transition Elements: 3L         <ol> <li>Definition and general characteristics of transition elements.</li> <li>Chemistry of 3d-transition elements with reference to:</li> </ol> </li> </ol>		1

			(a) Electronic Configuration	
			(b) Oxidation States	
			(c) Colour and Magnetic property	
			(d) Formation of Coordination compounds	
			(e) Application in general.	
	2.	Bondi	ing in Coordination Compounds: Valence	
			Theory 4L	
		(i)	Application to 4,5,6-Coordinate compounds	
		(ii)	Electro-neutrality principle and back-	
			bonding.	
	3.	Gravi	metric Analysis:	
			<b>8L</b>	
		(i)	Definition and Types of gravimetric analysis	
		(ii)	Precipitation Gravimetry with respect to Theory and Practice:	
			(a) Solubility considerations: Common ion effect; diverse ion effect, pH, temperature and nature of solubility	
		(iii)	<ul><li>(b) Controlling particle size.</li><li>Treatment of precipitates in gravimetry:</li><li>(a) Digestion</li></ul>	
		(iv)	<ul><li>(b) Filtration and Washing</li><li>(c) Drying and Ignition</li><li>Use of Organia Paragenta in gravimatric</li></ul>	
		(iv)	Use of Organic Reagents in gravimetric analysis	
	(Num	erical p	problems on this topic expected.	
	1.	Physic	co-Chemical Principles:	
		(i)	Criteria for spontaneity of Chemical	
			Reactions	
		. ,	Electrolysis	
		. ,	Effect of catalysts	
	-	(iv)	General Principles of Metallurgy	
	2.		facture of Bulk Chemicals:	
		(i)	Sulfuric acid (by Contact process)	
	2	(ii) Extra	Ammonia (by Haber's process)	
	5.	Extra	ction and Purification of:	
III		(i)	5L Coppor (from pyrites) by pyrometallyrgy	1
		(i)	Copper (from pyrites) by pyrometallurgy and electrolysis	
		(ii)	Silver by hydrometallurgy	
		(iii)	Aluminium by electrometallurgy.	
		(111)	mannun oy electionicunutgy.	

		1. Nomenchlature	
		<ol> <li>Nomenclature</li> <li>Nomenclature of polysbstituted aromatic compounds containing different functional Groups : Polysubstituted benzenes, tri substituted napgthalenes, disubstituted anthracences. Nomenclature of ethers, expoxides, and sulphur compounds.</li> </ol>	
		<ol> <li>Mechanism of organic reaction</li> <li>7L</li> </ol>	
		2.1 The mechanism of reactions involving the following reactive intermediates:	
		<ul> <li>2.1.1 Carbocations : Different types of carbocations, such as alkyl, allyl, benzyl. SN<sup>1</sup> reaction. Electrophilic addition across an olefinic double bond. Elimination – El mechanism. Reareeagement Wagner Meerwein rearrangement.</li> </ul>	
USCH 303	Ι	2.1.2 Carbanions : concept of cabon acid. Alkylation of carbon acids (active methelene compounds and terminal alkynes) using alkyl halides, and synthetic applications of these reactions. Reactions of Grignard reagents at carbonyl group. Aldol addition.	1
		<ul> <li>2.1.3 Carbon radicals : General reactions of radicals – abstraction, addition to C = C, combination, disproportination .</li> <li>Addition of HBr to alkenes in presence of perozide. Polymerization, Redox reaction – Kolbe electrolytic method.</li> </ul>	
		2.1.4 Carbenes : Generation of cabenes : through alpha elimination, from diazolakanes, from ketenes. Sturcture and stability. Reactions : insertion into C-H bond and addition to olefin.	
		2.2 Tautomerism : Keto-enol tautomerism in adlehudes and keyones, acid and base catalysed	

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enolisation, enol content and stability enols B- ketoester, B-diketones, phenols.		
<ol> <li>Aromatic Electrophilic Substitution Reaction</li> <li>6L</li> </ol>		
3.1 Huckel rule of aromaticity and its application to carbocyclic benzenoid and nonbenxenoid compounds and ions. Concept of antiaromaticity and homoaromaticity.		
3.2 General mechamism of aromatic electrophilic substation reaction with energy profile diagram.		
3.3 Drawing resonance structure of monosubstituted benzens – Activited and deactivated aromatic rings.		
<ul> <li>3.4 Effect of electron withdrawing and election donating substituents on the orientation of an incoming electrophile on the basis of - (i) electron density distribution (ii) stability of intermediate.</li> </ul>		
Cases of studied : Monosubstituted benezenes containing groups – Alkyl, amino, hedroxyl, alkoxy, halo, formyl, acyl, nitril, nitro, carboxy.		
1. Aromatic Hydrocabon		
3L		
1.1 Stucture of benzene, naphthalene, linear and angular acenes.		
<ul> <li>1.2 Alky arenes : Preparation of alkyl arences through reforming, Fridedel – Crafts Alkyation, with mechanism, using – Olefins, alcohols, alkylgalides.</li> </ul>		1
1.3 Reactions of alkyl arenes – side chian oxidation, ring vs side chain halogenations (mechanism)		
1.4		
	<ul> <li>ketoester, B-diketones, phenols.</li> <li>Aromatic Electrophilic Substitution Reaction 6L</li> <li>1.1 Huckel rule of aromaticity and its application to carbocyclic benzenoid and nonbenxenoid compounds and ions. Concept of antiaromaticity and homoaromaticity.</li> <li>2.2 General mechamism of aromatic electrophilic substation reaction with energy profile diagram.</li> <li>3.3 Drawing resonance structure of monosubstituted benzens – Activited and deactivated aromatic rings.</li> <li>3.4 Effect of electron withdrawing and election donating substituents on the orientation of an incoming electrophile on the basis of – (i) electron density distribution (ii) stability of intermediate.</li> <li>Cases of studied : Monosubstituted benzenes containing groups – Alkyl, amino, hedroxyl, alkoxy, halo, formyl, acyl, nitril, nitro, carboxy.</li> <li>1.1 Stucture of benzene, naphthalene, linear and angular acenes.</li> <li>1.2 Alky arenes : Preparation of alkyl arenees through reforming, Fridedel – Crafts Alkyation, with mechanism, using – Olefins, alcohols, alkylgalides.</li> <li>1.3 Reactions of alkyl arenes – side chian oxidation, ring vs side chain halogenations (mechanism)</li> </ul>	<ul> <li>ketoester, B-diketones, phenols.</li> <li>Aromatic Electrophilic Substitution Reaction 6L</li> <li>1.1 Huckel rule of aromaticity and its application to carbocyclic benzenoid and nonbenxenoid compounds and ions. Concept of antiaromaticity and homoaromaticity.</li> <li>2.2 General mechamism of aromatic electrophilic substation reaction with energy profile diagram.</li> <li>3.3 Drawing resonance structure of monosubstituted benzens – Activited and deactivated aromatic rings.</li> <li>3.4 Effect of electron withdrawing and election donating substituents on the orientation of an incoming electrophile on the basis of – (i) electron density distribution (ii) stability of intermediate.</li> <li>Cases of studied : Monosubstituted benezenes containing groups – Alkyl, amino, hedroxyl, alkoxy, halo, formyl, acyl, nitril, nitro, carboxy.</li> <li>1.1 Stucture of benzene, naphthalene, linear and angular acenes.</li> <li>1.2 Alky arenes : Preparation of alkyl arences through reforming, Fridedel – Crafts Alkyation, with mechanism, using – Olefins, alcohols, alkylgalides.</li> <li>1.3 Reactions of alkyl arenes – side chian oxidation, ring vs side chain halogenations (mechanism)</li> </ul>

2.1	Ialoarenes	4L	
	1.5 Preparation of haloarenes arenes – Halogenation substituted benezenes with (mechanism), limitations.	of benzene and	
	1.6 Reactions of haloarenes : L arylhaides under $S_nl$ and $S_N$ mechanism (addition – aromatic nucleophilic sur reaction – hydrolysis a haloarenes. Benzyne intern (elimination – addition nucleophilic substitution substitution) Grignard r Ullmann reaction.1.7 Applications of aromatic haloarenes	2 reactions. General elimination) of bstitutents on the and animation of nediate mechamism n) of aromatic reaction (cine reagent formation.	
2.	Phenols	3L	
	2.1 Preparation of phenols : P halobenzenes, (ii) from a acids (benezene and nag acids) (iii) isopropyl and hydroperoxide method.	aromatic sulphonic phthalene sulhonic	
	2.2 Reaction of phenols : Ac effect of substituents on acid formation, Etherification – alcohol, Williamson Sym Halogenation, Nitration, Fri aryl caboxylates, Claisen allyloxyarence	dity of phenols. Salt direct reaction with thesis O-acylation, es rearrangement of	
	2.3 Applications of phenols		
3.	Ether and Epoxides	5L	
	3.1 Ethers :		
	3.1.1 Preparation : Dehyo (mechanism), Wil (mechanism)	dration of alcohols lamson synthesis	

r			 
		3.1.2 Reactions : Acid catalyzed cleavage reaction with HX (mechanism)	
		3.1.3 Applications : Application of ethers, Crown ethers : Structure; 12-crown-4 and 18 crown-6 and their uses.	
		3.2 Epoxides :	
		3.2.1 Preparation : Oxidation of olefins – ethylene oxide; Reactions of per acids with olefins; from vicinal halohydrins	
		<ul> <li>3.2.2 Reactions : Reactivity. Ring opening reactions by nucleophiles (a) In acid conditions: hydrolysis, reaction with – HX, alcohol, HCN (b) In neutral or basic conditions ; ammonia, amines metal cyanides, Grignard reagents, alkoxides</li> </ul>	
		3.2.3 Applications of epoxides.	
		1. Sources of Organic Compounds	
		1. Sources of Organic Compounds 8L	
		-	
	III	8L 1.1 Sources : (a) Non-renewable : coalm petroleum (crude oil) and natural gas. (b) Renewable :	1
	III	<ul> <li>8L</li> <li>1.1 Sources : (a) Non-renewable : coalm petroleum (crude oil) and natural gas. (b) Renewable : biomass</li> <li>1.2 Coal : Stuctuture and types of coal, origin of coal. Distructive distillation of coal, coal tar refining, coal liquefaction (coal to liquid), coal gasification – synthesis gas (syn gas),</li> </ul>	1
	ш	<ul> <li>8L</li> <li>1.1 Sources : (a) Non-renewable : coalm petroleum (crude oil) and natural gas. (b) Renewable : biomass</li> <li>1.2 Coal : Stuctuture and types of coal, origin of coal. Distructive distillation of coal, coal tar refining, coal liquefaction (coal to liquid), coal gasification – synthesis gas (syn gas), hydropyrolysis</li> <li>1.3 Petroleum : Charactrisitics, composition and origin of petroleum Refining of petroleum. Catalytic cracking and reforming, hydrocracking, thermal cracking, steam</li> </ul>	1

	<ul> <li>1.5 Synthesis gas (syn gas) : Production of syn gas from – coal, natural gas, biomass, Composition, Synthetic use of syn gas- Separation of H2 Production of – methanol, alkanes, hydroformylation of olefins, synthesis diesel (biomass to liquid)</li> <li>1.6 Oligomerisation and metathesis of olefins.</li> <li>1.7 Biomass : Trasforming biomass into chemicals (pyrolysis) and synthesis gas.</li> <li>1.8 Biofuels: Ethanol, biodiesel, synthetic diesel, methanol</li> </ul>		
	2. Chemical Industry 4L		
	Idea of chemical plant, different units, block digrams and flow digrams. Flow diagrams : Principle, importance, typical elements. Typical flow diagram of single unit process. Idea of flow diagrams of multiple process units. Continous vs batch operations. Raw materials, intermediates, end products, by- products, waste. Unit processes – nitration as an example.		
	3. Environmental aspects of Chemical Industry		
	Volatile Organic Compounds (VOC), Greehouse effect : Ozone depletion, important greenhouse gases and their sourses. Hydrocarbons as air pollutants, Harmful effects of industrial effluents. Carbon emission – carbon credit, carbon neutrality, carbon offsetting. Material safely Data Sheet (MSDS).		
USCH P3	Practicals based on both courses in theory	3	9

## SEMESTER IV

Course Code	UN IT	TOPICS	Cr edi ts	L / We ek
USCH401	Ι	<ul> <li>Phases in Equilibra: (15L)</li> <li>1) Introduction to the terms: phase, component, degrees of freedom, Gibbs phase rule.</li> <li>2) Two component systems:</li> <li>A) Liquid-liquid mixture, completely miscible liquids, phase diagrams, pressure composition and temperature diagrams, Raoult's law, ideal solutions, distillation of mixtures forming ideal solution, fractional distillation, distillation under reduced pressure, deviations from the Raoult's law, positive and negative deviations, nonideal solutions, azeotropes distillation of azeotropic mixtures, breaking of azeotropoes.</li> <li>B) Liquid-liquid mixtures, partially miscible liquids, partially miscible liquid pairs exhibiting lower critical solution temperature.</li> <li>C) Liquid-liquid mixtures, completely immiscible liquids, steam distillation.</li> <li>3) Nernst's distribution law, partition coefficient distribution ratio, solutes undergoing association and dissociation (Qualitative explanation only)</li> </ul>	2	1
	Π	<ul> <li>Electrochemical Cells and Ionic Equilibria (15L)</li> <li>Electrochemical Cells <ol> <li>Galvanic and electrolytic cells, types of electrodes, ion specific and ion selective, comparison, types of ion specific electrodes, (iv) redox electrodes.</li> <li>Cell representation and cell reaction, (Students are expected to write the cell reaction for a given cell and vice versa.</li> <li>Reversible and irreversible cells, chemical and concentration cells, primary and secondary cells.</li> <li>Thermodynamic parameters (G, H and S) for the cell reaction taking place ion a cell.</li> <li>Nernst's equation for cell emf and hence for a single electrode potential (Derivation expected.)</li> </ol> </li> </ul>		1

	6) Determination of equilibrium constant from emf		
	measurements.		
	Ionic equilibria:		
	1) Concept of PH, POH, calculations of PH of		
	solution of acids and bases.		
	2) Buffer, buffer capacity, Henderson's equation for		
	acidic and basic buffers (derivation expected).		
	Introduction to Titrimetric Analysis and Use of	-	
	e e		
	Instrumental Methods Ion (15L)Titrimetric Analysis		
	Inter destion to Tituins the Matheda of Anologie		
	Introduction to Titrimetic Methods of Analysis		
	1) Requirements for a reaction to be used in		
	titrimetric analysis, classification of titrimetric		
	analysis, terms: titration, titrand, titrant, titre value,		
	indicator, endpoint, equivalence point, titration		
	error.		
	2) Calibration of the volumetric glassware, burettes,		
	pipettes and volumetric flasks.		
	3) Theory of acid base indicators, choice of an		
	indicator for the titration dependence on the PH at		
	-		
	the equivalence point.		
	Use of Instrumental Methods in Titrimetric		
	Analysis		
	1) Conductometric Titrations: Basic principles,		
ттт	experimental set up titration curves in the titration		1
III	of (i) strong acid vs, strong base, (ii) weak acid vs,		1
	strong base, (iii) weak acid vs, weak base, (iv)		
	Mixture of strong and weak acid/strong weak base		
	vs, strong base/weak base or strong acid/weak		
	acid, (v) sodium chloride vs, silver nitrate (vi)		
	barium hydroxide vs, magnesium sulphate		
	advantages and limitations.		
	2) Potentiometric Titrations: Basic principles, concept		
	of indicator electrode, indicator electrodes for		
	different types of titration, determination of end		
	point from the graphs of E vs V, $E/E$ vs, mean		
	volume, $2E/V$ vs, mean volume, advantage and		
	•		
	limitations. (3) Photometric titrations: Basic		
	principles, titration curves, advantages and		
	limitations. (4) Use of computes, use of spread		
	sheets.		

USCH402	Ι	Acids and Bases, Chemistry of Elements and Organometallic Compounds       15L         1. Acids and Bases:       5L         (i) Appreciation of various concepts due to:       (a) Arrhenius         (b) Lowry-Bronstead       (c) Lux-Flood         (d) Lewis       (e) Solvent-System         (f) Usanovich       (ii) Pearson's classification of acids and bases and his principle of HSAB.         2. Chemistry of Group 15 and Group 16 elements:         SL         General discussion of trends in their physical and chemical properties, Physical properties of hydrides of Group 15 and Group 16 elements with respect to hydrogen bonding.         3. Organometallic Componds:         SL         (i) Introduction, definition, classification on the basis of:         (a) Hapticity, and         (b) Nature of metal-carbon (M-C) bond         (ii) Eighteen Electron Rule with respect to: applications and exceptions:         (iii) Metal carbonyls with respect to:         (a) Classification and bonding         (b) General methods of preparation, and (c) Properties.         (iv) Applications of organometallic compounds	1
	П	<ol> <li>Chemistry of Group 17 and Group 18 elements: 6L         (i) General trends in physical and chemical properties         (ii) Chemistry of pseudo-halogens with respect to: (a) Comparison with halogens (b) Preparation and uses.         </li> <li>Chemistry of Group – 18 elements with respect to: 4L i) History, and ii) Preparation and structures of xenon         </li> </ol>	1

fluorides and oxy fluorides		٦
3. Bio-Inorganic Chemistry: 5L (i) Metaloporphyrins (ii) Chlorophy II (iii) Cytochromes (iv) Hemoglobin and myoglobin: oxygen transport and storage.		
<ol> <li>Corrosion and Methods of Protection of Metals 7L         <ul> <li>(i) Introduction (to include economics and importance of corrosion)             <li>(ii) Types of corrosion                 <ul></ul></li></li></ul></li></ol>	1	

		1. Aromatic Nitrogen compounds8L	
		1.1 Aromatic nitrocompounds	
		1.1.1 Preparation : Nitration using mixed acid, Preparation of mononitro and dinitro- compounds through nitration of benzene (mechanism), nitrobenzene, toluene, chlorobenzene, naphthalene, anisole.	
		1.1.2 Reactions : Reduction of nitro- compounds under different conditions.	
		1.1.3 Applications of nitro compounds : In the preparion of amines and explosives.	
		11.2. Aromatic amino compounds	
USCH403	Ι	<ul> <li>11.2.1 Preparation : Reduction of aromatic nitro compounds using catalytic hydrogenation, dissolving metal reduction using – Fe-HCI, Sn-HCI, Zn-HOAc, NaHS, Animation of halobenzenes, Chemoselective reduction of dinitrobenzene, Hoffmann bromanide reaction.</li> <li>11.2.2. Reactions : Basicity of aromatic amines – effect of substituents on basicity of aniline, salt formation, N-alkylation, N-acylation, halogenations, reductive alkaylation, diazotization of aromatic primary amines (mechanism) Reactions of aryl diazonium salts-Sandmeyer and Gattermann reactions, Replacemet of diazo group by H – OH- CN Gomberg reaction. Azo-coupling reaction with phenols/naphthols and aromatic amines. Reducation of azo-and hydrazo-benzenes.</li> </ul>	1
		<ul> <li>4. Chromophore – auzochrome concept, azo group as a chromophore, azo dyes. Aromatic Aldehydes and Ketones 7L</li> </ul>	
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