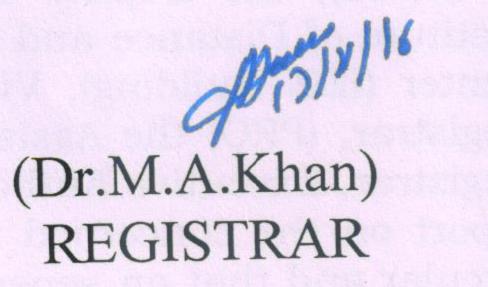
UNIVERSITY OF MUMBAI No. UG/ 168 of 2016-17

CIRCULAR:-

A reference is invited to the Syllabi relating to the B.Sc. degree course, vide this office Circular No. UG/179 of 2010, dated 13th July, 2010 and the Principals of affiliated Colleges in Science are hereby informed that the recommendation made by the Ad-hoc Board of Studies in Chemistry at its meeting held on 7th July, 2016 has been accepted by the Academic Council meeting held on 14th July, 2016 vide item No. 4.14 and that in accordance therewith, the revised syllabus as per the Choice Based Credit System for T.Y. B.Sc. programme in Chemistry (Three Units)(Sem. V & VI), which are available on the University's web site (www.mu.ac.in) and that the same has been brought into force with effect from the academic year 2016-17.



MUMBAI - 400 032 1 9 November, 2016

To,

The Principals of the affiliated Colleges in Science.

A.C/4.14/14.07.2016

MUMBAI-400 032 No. UG//68-A of 2016

November, 2016

Copy forwarded with Compliments for information to:-

1) The Co-ordinator, Faculties of Science, 2) The Chairman, Board of Studies in Chemistry,

- 3) The Professor-cum-Director, Institute of Distance & Open Learning (IDOL)
- The Director, Board of College and University Development, 4)
- 5) The Co-Ordinator, University Computerization Centre,
- 6) The Controller of Examinations.

(Dr.M.A.Khan)

REGISTRAR

PTO..

T. Y. B. Sc. CHEMISTRY (Three Units) Credit Based and Grading System To be implemented from the Academic year 2016-2017

SEMESTER V Theory Paper-I

Course	Un	Contents	Credit	L/Week
USCHP501	it	 1.1 Colligative Properties of Dilute Solutions (8L) 1.1.1 Dilute solution, colligate properties, Raoult's law, relative lowering of vapour pressure 1.1.2 Elevation in boiling point of a solution, thermodynamic derivation relating elevation in the boiling point of a solution and the molar mass of the non-volatile solute 1.1.3 Depression in freezing point of a solution, thermodynamic derivation relating the depression in the freezing point of a solution as solution and the molar mass of the non-volatile solute 1.1.4 Osmotic pressure, van't Hoff's equation for osmotic pressure, equation for osmotic pressure, determination of molar mass of the solute. Abnormal molar masses of solutes and van't Hoff factor (calculation of Degree of Association and Degree of Dissociation.) 1.2. Phase Rule 1.2.1. Gibb's phase rule and terms involved in the equation 1.2.2. Application of phase rule to ONE component systems (i) water system, (ii) sulphur system 1.2.3. Application of phase rule to TWO component systems, condensed systems, condensed phase rule, eutectic systems (Lead-Silver system), desilverisation of lead 1.2.4. Introduction to three component system, explanation of phase diagram for three liquids forming one immiscible pair. 	<u>s</u> 2.5	1
	2	 2.1.Electrochemistry –Electrochemical cells (15L) 2.1.1. Lewis concept of Activity and Activity coefficient, Mean ionic activity and mean ionic activity coefficient γ₊₋ of an electrolyte, expression for activities of electrolytes of different valence type, ionic strength 2.1.2. Classification of cells: 1.chemical cells without transference 2.Concentration cells with and without Transference (derivations of concentration cell EMF expected) Origin of liquid-liquid junction potential and its elimination using a salt bridge 2.1.3. Applications of EMF measurements in the determination of 1. pH of a solution using quin hydrone electrode and glass electrode. 2. Solubility and solubility product of sparingly soluble salt using chemical cell and concentration cell method. 3. Determination of liquid-liquid junction potential 3. Chemical Bonding And Solid State Chemistry (15L) 		

		3.1 Molecular Symmetry (10L)		
		3.1.1 Introduction and Importance.		
		3.1.2 Symmetry elements and symmetry operations.		
		3.1.3 Concept of a Point Group with illustrations using the		
		following point groups: (i) Cav (HCl), (ii) $D_{\alpha h}$ (H ₂),		
		(iii) C_{2v} (H ₂ O), (iv) C_{3v} (NH ₃), (v) C_{2h} (trans-trichloroethylene),		
		and (vi) D_{3h} (BCl ₃).		
		3.2 Molecular Orbital Theory for Polyatomic Species (5L)		
		3.2.1 Simple triatomic species: H_3^+ and H_3 (correlation between		
		bond angle and Molecular orbitals).		
		3.2.2 Other molecules (considering only σ -bonding): i) BeH ₂ , ii)		
		H ₂ O.		
	4	Solution Chemistry		
	-	4.1 Acid-base Chemistry in Aqueous Medium (8L)		
		4.1.1 Acidity of mono- and polyatomic cations.		
		4.1.2 Basicity of mono- and polyatomic		
		anions (discussion for 4.1.1 as well as 4.1.2 to Include Latimer		
		equation and predominance diagrams).		
		4.2 Chemistry in Non-aqueous Solvents (7L)		
		4.2.1 Classification of solvents and importance of non-aqueous		
		solvents.		
		4.2.2 Characteristics and study of liquid ammonia, dinitrogen		
		tetraoxide and acetic acid as non-aqueous solvents with respect		
	1	to (i) acid-base reactions and (ii) redox reactions.	2.5	1
USCHP502	1	Mechanism of Organic Reactions (15L)	2.5	1
		1.1. Thermodynamic and Kinetic control of organic reactions:		
		Concept with mechanisms of the following reactions: addition of		
		HX to butadiene; sulfonation of naphthalene. Nucleophilicity/		
		electrophilicity vs Basicity/acidity.		
		1.2 Mechanism of elimination reactions, with stereochemistry:		
		E1 and E2 reactions: regioselectivity (Saytzeff and Hofmann		
		rules).		
		1.3 Mechanism of reactions of carbonyl compounds with		
		nucleophiles:		
		1.3.1 Formation of acetals/ketals from aldehydes and ketones.		
		1.3.2 Reaction of aldehydes and ketones with primary and		
		secondary amines.		
		1.3.3 Acyl nucleophilic substitution (tetrahedral		
		mechanism): Acid catalysed esterification of Carboxylic acids		
		and base promoted hydrolysis of esters.		
		1.4 Mechanism of rearrangements with examples and		
		stereochemistry wherever applicable.		
		1.4.1 Migration to electron deficient carbon: Pinacol,		
		Benzylic acid.		
		1.4.2 Migration to deficient nitrogen: Beckmann, Hofmann.		
		1.5 Mechanism of the following reactons with synthetic		
		application: Claisen condensation, Michael addition.		
	2	Heterocyclic Chemistry (8L)		
		2.1.1 Introduction: Electronic structure and aromaticity of furan,		
		pyrrole, thiophene and pyridine.		
	1		1	1

Paal-Knor synthesis. Pyridines by Hantzsch synthesis and from 1.5-diketones. 21.3 Reactivity: Reactivity towards electrophilic substitution reactions- of furan, pyrrole and thiophene on basis of stability of intermediate; and of pyridine on the basis of electron distribution.Nucleophilic substitution reaction of pyridine on the basis of electron distribution. 21.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation, Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole and pyrrolidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 22.2 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi-component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 22.2.1 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione). Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 3 Treatment of analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, ave	_		
 1,5-diketones. 2.1.3 Reactivity: Reactivity towards electrophilic substitution reactions of furan, pyrrole and thiophene on basis of stability of intermediate; and of pyridine on the basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation, Nitration, Sulphonation, Vilsmeir formylation reaction. Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole-comparison of basicity of pyrrole. Pyrrole: Acidity and basicity. Comparison of basicity of pyrrole and pperidine. Sulphonation of pyrridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyrridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.3.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione). Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 3 Treatment of analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7 L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, a		2.1.2 Synthesis: Synthesis of furans, pyrroles, and thiophenes by	
2.1.3 Reactivity: Reactivity towards electrophilic substitution reactions- of furan, pyrrole and thiophene on basis of stability of intermediate; and of pyridine on the basis of electron distribution.Nucleophilic substitution reaction of pyridine on the basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation , Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and piperidine. Acid catalysed polymerization of pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.1.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-1, pigment red 2			
 reactions- of furan, pyrrole and thiophene on basis of stability of intermediate; and of pyridine on the basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation, Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole. Acidity and basicity of pyrrole-Comparison of basicity of pyrrole. Portidine: Basicity. Comparison of basicity of pyrrole. Pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyrdidine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione). Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Introduction to the use of the following in organic synthesis: Classond, microwaves, PTC. 3 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, coefficient of variation, [Numerical problems expected] 3.2 Sampling (8L) Treatment of Analytical Data (7L) Types of enrors, ensures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficie		1,5-diketones.	
 reactions- of furan, pyrrole and thiophene on basis of stability of intermediate; and of pyridine on the basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation, Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole. Acidity and basicity of pyrrole-Comparison of basicity of pyrrole. Portidine: Basicity. Comparison of basicity of pyrrole. Pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyrdidine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione). Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Introduction to the use of the following in organic synthesis: Classond, microwaves, PTC. 3 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, coefficient of variation, [Numerical problems expected] 3.2 Sampling (8L) Treatment of Analytical Data (7L) Types of enrors, ensures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficie		2.1.3 Reactivity: Reactivity towards electrophilic substitution	
antermediate; and of pyridine on the basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation , Nitration, Sulphonation, Vilsmeir formylalion reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole: Apyrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyrrole. Pyridine: Basicity. Comparison of basicity of pyrrole and piperidine. Sulphonation of pyrdine, pyrrole and piperidine. Sulphonation of pyrdines and action of sodamide (Chichibabin reaction).Nmethylation of pyrdine. Quaternization of piperdine, pyrrolidine, addyl pyrdines and action of sodamide (Chichibabin reaction).Nmethylation of pyrdine. Quaternization of piperdine, pyrrolidine, addyl pyrdines and action f sodamide (Chichibabin reaction).Nmethylation of pyrdine: Cuaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Mult: component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanilin, methyl dihydrojasmonate (Hedinoe), Bifenox -1, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 3 Treatment of analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersi			
distribution.Nucleophilic substitution reaction of pyridine on the basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation , Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Dels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and piperidine. Sulphonation of puscility. Gropparison of basicity of pyrrole and piperidine. Sulphonation of puscility of pyridine, pyrrole and piperidine. Sulphonation of puscility, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine, and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.3.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-1, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 3 Treatment of analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average devia			
 basis of electron distribution. 2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation , Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole: Acidity and basicity. Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyrrole and piperidine. Sulphonation of pyrdine, with and without catalyst. Reduction. Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction). Nmethylation of pyrdine. Quaternization of piperdine, pyrroleine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione). Bifenox-1, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, stanace, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
2.1.4 Reactions of heterocycles: The following reactions of furan, pyrrole and thiophene: Halogenation, Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole. Pyridine: Basicity. Comparison of basicity of pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrroleidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi-component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 3 Treatment of analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, rainace, coefficient of variation. [Numerical problems expected] 3.2 Sampling (8L) Treatment of Sampling, sampling techniques, sampling of gases, ambient and stack			
furan, pyrrole and thiophene: Halogenation , Nitration, Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis - with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.1 Ilustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. <			
Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.3.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-L pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.3. Newer methods of organic synthesis: Iltrasound, microwaves, PTC. 3 3 T			
reaction. Furan: Diels-Alder reaction. Ring opening of furan. Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.1. Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.1 Ilutroduction: Criteria for ideal organic synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanilin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, det		furan, pyrrole and thiophene: Halogenation, Nitration,	
Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole and piperidine: Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.3.3 Newer methods of organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of analytical pota (7L) Types of erro		Sulphonation, Vilsmeir formylation reaction, Friedel-Crafts	
Pyrrole: Acidity and basicity of pyrrole-Comparison of basicity of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole and piperidine: Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.3.3 Newer methods of organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of analytical pota (7L) Types of erro		reaction. Furan: Diels-Alder reaction. Ring opening of furan.	
of pyrrole and pyrrolidine, Acid catalysed polymerization of pyrrole, Pyridine: Basicity. Comparison of basicity of pyrdine, pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.1.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-1, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.3.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of Analytical data-1 and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors			
 pyrrole. Pyridine: Basicity. Comparison of basicity of pyridine, pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction). Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi-component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2. 21 Ilustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione). Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
signal pyrrole and piperidine. Sulphonation of pyridine, with and without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average <td></td> <td></td> <td></td>			
 without catalyst. Reduction.Oxidation of alkyl pyridines and action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of Analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
action of sodamide (Chichibabin reaction).Nmethylation of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of Analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Num			
of pyridine. Quaternization of piperdine, pyrrolidine and Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (TL) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gas			
 Hofmann elimination of the quaternary salts. 2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. Treatment of Analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
2.2. Organic Synthesis (7L) 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
 2.2.1 Introduction: Criteria for ideal organic synthesis. Yield and selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of Analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (<i>TL</i>) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 		Hofmann elimination of the quaternary salts.	
 selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 		2.2. Organic Synthesis (7L)	
 selectivity. Multi- component synthesis – with examples, Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
Mannich reaction, Hanztsch synthesis of pyridines (without mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
mechanism). 2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
2.2.2 Illustrative synthesis of industrially important compounds: Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
Ibuprofen (chiral synthesis), paracetamol (green synthesis), Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid.2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC.3Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
Lascorbic acid (from D-glucose), norfloxacin, thyroxine, vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid.2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC.3Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
vanillin, methyl dihydrojasmonate (Hedione), Bifenox-I, pigment red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid.2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC.3Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic acid. 2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
acid.2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC.3Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
2.2.3 Newer methods of organic synthesis: Introduction to the use of the following in organic synthesis: Ultrasound, microwaves, PTC. 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack		red 242, indigo, 2- hydroxy-3-amino-5-nitrobenzene sulphonic	
use of the following in organic synthesis: Ultrasound, microwaves, PTC.3Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack		acid.	
use of the following in organic synthesis: Ultrasound, microwaves, PTC.3Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack		2.2.3 Newer methods of organic synthesis: Introduction to the	
microwaves, PTC. 3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
3 Treatment of analytical data-I and sampling (15 L) 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
 3.1 Treatment of Analytical Data (7L) Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
Types of errors, determinate and indeterminate errors, minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack		$\mathbf{I} = \mathbf{I} + $	
 minimization of errors, constant and proportionate errors, accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
 accuracy and precision, measures of dispersion and central tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
 tendency: mean, median, average deviation, relative average deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 			
 deviation, standard deviation, variance, coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack 		• •	
coefficient of variation.[Numerical problems expected] 3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack		deviation, standard deviation, variance,	
3.2 Sampling (8L) Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack		coefficient of variation.[Numerical problems expected]	
Terms involved, importance of sampling, sampling techniques, sampling of gases, ambient and stack			
sampling of gases, ambient and stack			
sampling equipment used sampling of homogeneous and			
sampling, equipment used, sampling of homogeneous and			
heterogeneous liquids, sampling of static and flowing liquids,			
methods and equipments used, sampling of solids, importance of			
particle size and sample size, samples used, need for the		particle size and sample size, samples used, need for the	

	reduction in the sample size, methods of reduction in sample size, collection, preservation and dissolution of the sample.		
	 Titrimetric analysis-I and UV-Visible Spectroscopy. (15L) 4.1 Acid-base Titrations (5L) Construction of titration curves and choice of indicators in the titration of [1] strong acid and strong base, [2] strong acid and weak base, [3] weak acid and strong base, [4] weak acid and weak base. 4.2 Precipitation titrations (4L) Argentimetric titrations, construction of the titration curve, Volhard's method, Mohr's method, adsorption indicators, theory and applications. 4.3 U.V. Visible Spectroscopy (4L) Photometers and spectrophotometers, Instrumentation in the case of single and double beam spectrophotometers, Qualitative and quantitative analysis, calibration cure method. 		
US3CHP05	 Physical Chemistry Practicals Chemical Kinetics – To determine the order between K2S2O8 & KI by fractional change method. Potentiometry – To determine the solubility product and solubility of AgCl potentiometrically using chemical cell. OR To determine the solubility product and solubility of AgCl potentiometrically using concentration cell. Colorimetry – To determine the amount of Fe(III) present in the given solution by using salicylic acid by colorimetric titration.(static method) (=525 nm) Inorganic Chemistry Practicals	1.5	
US3CHP06	 1. We compleximite the final cator Organic Chemistry Practicals Binary Mixture Separation, drying, weighing & Melting Point (No identification) (Solid + Solid) (4 Expts) Analytical Chemistry Practicals 1. Estimation of persulphate in the given sample by the method of back titration. 2. Determination of Vitamin C content of a given tablet by titration with sodium hydroxide pH metrically 	1.5	

T. Y. B. Sc. CHEMISTRY (Three Units) Credit Based and Grading System To be implemented from the Academic year 2016-2017

SEMESTER VI

Theory Paper-I

Course	Unit	Content	Credi ts	L/Week
USC3CH601	1	Molecular Spectroscopy (15 L)	2.5	1
0.00001001	-	1.1 Dipole moment: Dipole moment, polarization of a		-
		bond, bond moment, dipole moment and molecular		
		structure.		
		1.2 Rotational Spectrum: Rotational spectrum of a		
		diatomic molecule, rigid rotor, moment of inertia,		
		energy levels, conditions for obtaining pure rotational		
		spectrum, selection rule, nature of		
		pectrum, determination of inter nuclear distance and		
		isotopic shift.		
		1.3 Vibration (IR) spectrum:		
		Vibrational motion, degrees of freedom, modes of		
		vibration, vibrational spectrum of a diatomic molecule,		
		simple harmonic oscillator, energy levels, zero point		
		energy, conditions for obtaining vibrational spectrum,		
		selection rule, nature of spectrum.		
		1.4 Vibration-Rotation spectrum of		
		diatomic molecule		
		vibrating rotor, energy levels, selection rule, nature of		
		spectrum, R and P branches, anharmonic oscillator :		
		energy levels, selection rule, fundamental band,		
		overtones. Application of vibrationrotation spectrum in		
		determining Force constant, determination and		
		significance. Introduction to infrared spectra of simple		
		molecules like H2O and CO2		
		1.5 Raman Spectroscopy : Scattering of		
		electromagnetic radiation, Rayleigh scattering, Raman		
		scattering, nature of Raman spectrum, Stoke's		
		lines, anti- Stoke's lines, Raman shift, quantum theory of		
		Raman spectrum, comparative study of IR and Raman		
		spectra, rule of mutual exclusion.(example of		
		CO2molecule).		
	2	2.1 Renewable Energy Sources (5L)		
		2.1.1. Lithium ion cell.		
		2.1.2 . Fuel cells; Choice of fuel and oxidant, Bacon's		
		H2 and O2 fuel cell.		
		2.1.3 . Solar cells, solar energy, photovoltaicneffect,		
		semiconductors as solar energy converters, silicon solar		
		cell		

		3.3 Electronic States and Terms for Poly electronic		
		Atoms (4L)		
		3.3.1 Introduction: electronic configuration and		
		electronic states, Term symbols, coupling of spin		
		momenta (Ms), orbital momenta (Ml)and spin- orbit		
		coupling or Russell-Saunders coupling.		
		3.3.2 Determination of Terms for <i>p</i> 2 electronic		
		configuration (as in a carbon atom).		
		3.3.3 Terms and micro-states for transition metal		
		atoms/ions.		
	4.	Some Selected Topics (15L)		
		4.1 Inorganic Polymers (3L)		
		4.1.1 Various methods of classification with		
		examples.		
		4.1.2 Chemistry of borazine with reference to		
		preparation, properties, structures, bonding and		
		applications.		
		4.2 Characteristics and Treatment of		
		Liquid Effluent (06L)		
		4.2.1 Characterization of waste: biochemical oxygen		
		demand (BOD), chemical oxygen demand (COD), total		
		organic carbon (TOC), aerobic and anaerobic processes.		
		4.2.2 Removing of solid contaminants, physical and		
		chemical principles such as coagulation, flocculation		
		and sedimentation.		
		4.2.3 Primary, secondary and tertiary of liquid effluents.		
		4.3 Nanomaterials (04L)		
		4.3.1 Introduction and importance of nanomaterials.		
		4.3.2 Properties (Comparison between bulk and		
		nanomaterials): (i) Optical properties, (ii) Electrical		
		conductivity, and (iii) Mechanical properties.		
		4.3.3 Forms of nanomaterials: nanofilms, nanolayers,		
		nanotubes, nanowires, and nanoparticles.		
		4.3.4 Chemical methods of preparation: (i) Colloidal		
		route, and (ii) Sol-gel method.		
		4.4 Inorganic Pharmaceuticals (2L)		
		4.4.1Gastrointestinal agents viz., (i) antacids		
		(aluminium hydroxide, milk of magnesia, sodium		
		bicarbonate and (ii) cathartics (magnesium sulphate and		
		sodium phosphate). Topical agents viz., (i) protectives		
		and adsorbents (talc, calamine), (ii) antimicrobial agents		
		(potassium permanganate, tincture iodine, boric acid)		
		and astringents (alum).		
USC3CH602				
	1.	Spectroscopy (15L)	2.5	1
		1.1 Introduction : Electromagnetic spectrum, units of		
		wavelength and frequency.		
		1.2 UV- Visible Spectroscopy: Basic theory, solvents,		
		nature of UV-VIS spectrum, concept of Chromophore,		
		auxochrome, bathochromic shift, Hypsochromic shift		

 1		
	hyperchromic effect and chromophore- auxochrome	
	interactions.	
	1.3 IR Spectrocopy: Basic theory, nature of IR	
	spectrum, selection rule, fingerprint region.	
	1.4 PMR Spectroscopy: Basic theory of NMR, nature of	
	PMR spectrum, chemical shift (∂ unit), standard for	
	PMR, solvents used. Factors affecting chemical shift:	
	(1) inductive effect (2) anisotropic effect (with reference	
	to C=C, C \equiv C, C=O and benzene ring). Spin- spin	
	coupling and coupling constant. Proton exchange	
	application of deuterium exchange Application of PMR	
	in structure determination.	
	1.5 Spectral characteristics of following classes of	
	1 0	
	organic compounds, including benzene and	
	monosubstituted benzenes, with respect to UV-VIS,	
	IR,PMR: (1)alkanes (2)alkenes and polyenes (3) alkynes	
	(4) haloalkanes (5) alcohols (6) carbonyl compounds (7)	
	ethers (8) carboxylic acids (9) esters (10)amines (11)	
	amides (broad regions characteristic of different groups	
	are expected).	
	1.6 Problems of structure elucidation of simple organic	
	compounds using individual or combined use of the	
	above spectroscopic technique are expected.(index of	
	hydrogen deficiency should be the first step in solving	
	the problems).	
2.	2.1 Organometallic Chemistry (5L)	
	2.1.1 Introduction : Carbon-metal bond- Nature, types	
	reactivity.	
	2.1.2 Organo magnesium Compounds: Grignard	
	reagent : Preparation, structure, and stability, Reaction	
	with compounds containing acidic hydrogen, carbonyl	
	compounds, cyanides and CO2.	
	2.1.3 Organolithium Compounds : Preparation using	
	alkyl/aryl halides. Reactions with compounds containing	
	acidic hydrogen, alkyl halides, carbonyl compounds,	
	cyanides and CO2. Lithium dialkyl cuprates:	
	Preparation and reactions with aliphatic	
	/aromatic/vinylic halides.	
	2.1.4 Organozinc compounds : Preparation of dialkyl	
	zinc. Reaction with water, acid chlorides and alkyl	
	halides. Reformatsky reaction (with mechanism).	
	2.2 Chemistry of some Important Biomolecules:	
	(10L)	
	2.2.1 α-Amino acids: Structure, configuration, Essential	
	amino acids and their abbreviations, classification,	
	Properties: pH dependency of ionic structure and	
1		
	isoelectric point. Methods of preparations. Strecker	
	isoelectric point. Methods of preparations: Strecker	
	isoelectric point. Methods of preparations: Strecker synthesis, amidomalonate synthesis, Erlenmeyer azalactone synthesis.	

2.2.2 Polypeptides and Proteins: Polypeptides: Pept	ide	
bond. Nomenclature and representation of polypep Merrifields solid phase peptide synthesis (example di- and tri- peptides for nomenclature and synthesis	tides. of	
Proteins: Sources, types,functions,colloidal nature, separation based on isoelectric point, denaturation a	and	
functions. Partial and total hydrolysis. General idea primary, secondary, tertiary and quartenary structur		
2.2.3 Nucleic acids: Selective hydrolysis of nucleic		
acids.Sugars and bases in nucleic acids. Stuctures o		
nucleosides an nucleotides in DNA and RNA. Struc of nucleic acids (DNA and RNA): Base pairing in	cture	
nucleic acids. Importance of nucleic acids-self		
 duplication, protein synthesis.	of 25	1
 3.1 Treatment of Analytical Data (6L): Distribution random errors, Gaussian curve, students' t, confider limits and confidence interval, criteria for rejection result: 2.5d rule,4.0 rule and Q test, F teset, testing t significance, null hypothesis, method of averages, le squares method. [Numerical problems expected] 3.2 Complexometric Titrations (5L): General introduction, EDTA titrations, advantages and limitations of EDTA as the titrant, absolute and conditional formation constants of metal EDTA complexes, construction of titration curves, types of EDTA titrations, methods of increasing the selectivi EDTA as a titrant, metallochromic indicators, theor and applications. 3.3 Redox Titrations (4L): General introduction, the of redox indicators, criterion for choosing an indicator curves in the case of (1) Fe (II) Vs. Ce(IV) (2) Fe (Vs. dichromate, use of diphenyl amine and ferroin a redox indicators. 	nce of for east f ty of y eory tor II)	1
 redox indicators. Concepts in Quality and miscellaneous methods (15L)	2.5	1
4.1 Total quality management (5L) : concept of qua quality control, quality assurance total quality	llity,	
management, ISO series, Good laboratory practices		
4.2 Mass Spectrometry (2L): Basic principles, introduction of components only		
4.3 Thermal Methods (5L): Classification of therma		
methods, thermogravimetric analysis, basic principle instrumentation factors affecting the TG curve,	es,	
applications		
4.4 Introduction to Radio Analytical Techniques (3) Classification of the techniques, introduction to neu activation analysis and its applications.		
· · · · · · · · · · · · · · · · · · ·		
 Practicals		

US3CHP07	Physical Chemistry Practicals	1.5	4
	Potentiometry –		-
	1. To determine the strength of the given strong acid		
	(HCl) by potentiometric titration using quinhydrone		
	electrode (Calculation of pH from Ecell and the plot of		
	(a) against V (b) pH against V graphs are expected).		
	OR		
	2. To determine pKa value of the given weak monobasic		
	acid (CH3COOH) by e.m.f. measurements.		
	Conductometry –		
	3.To determine the amount of dibasic acid (Oxalic acid)		
	by conductometric titration against strong base.		
	OR		
	4. To determine the relative strength of		
	monochloroacetic acid and acetic acid nductometrically.		
	Inorganic Chemistry Practicals		
	Inorganic Chemistry Practicals Inorganic preparations		
	1. Tris-(acetylacetonato) iron (III)		
	2. Bis-(Dimethylglyoximato) nickel (II)		
	Inorganic estimations/ Analysis		
	1. Acidity of a water sample.		
	1. Acturty of a water sample.		
US3CHP08	Organic Chemistry Practicals	1.5	4
	Organic Preparations: Drying, Weighing		
	& Melting Point (No Purification)		
	1. Aniline/p-toluidine \rightarrow N-Acetyl derivative		
	2. Salicylic acid/nitrobenzene/ Acetanilide \rightarrow		
	Nitro derivative		
	3. β - naphthol \rightarrow Methyl Ether derivative (Using		
	dimethyl sulphate)		
	4. Methyl salicylate/ethyl benzoate \rightarrow Acid derivative		
	(Hydrolysis)		
	(Hydrolysis) Analytical Chemistry Practicals		
	Analytical Chemistry Practicals		
	Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water		
	Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample.		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of 		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of common salt using a cation 		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of common salt using a cation exchanger. 		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of common salt using a cation exchanger. 3. Determination of acetic acid content of a vinegar 		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of common salt using a cation exchanger. 3. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium 		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of common salt using a cation exchanger. 3. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium hydroxide using quinhydrone. 		
	 Analytical Chemistry Practicals 1. Determination of chemical oxygen demand of a water sample. 2. Determination of percentage purity of a sample of common salt using a cation exchanger. 3. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium 		
	 Analytical Chemistry Practicals Determination of chemical oxygen demand of a water sample. Determination of percentage purity of a sample of common salt using a cation exchanger. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium hydroxide using quinhydrone. Suggested References 		
	 Analytical Chemistry Practicals Determination of chemical oxygen demand of a water sample. Determination of percentage purity of a sample of common salt using a cation exchanger. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium hydroxide using quinhydrone. Suggested References Physical Chemistry 		
	 Analytical Chemistry Practicals Determination of chemical oxygen demand of a water sample. Determination of percentage purity of a sample of common salt using a cation exchanger. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium hydroxide using quinhydrone. Suggested References Physical Chemistry Physical Chemistry, Ira Levine, 5th Edition, 		
	 Analytical Chemistry Practicals Determination of chemical oxygen demand of a water sample. Determination of percentage purity of a sample of common salt using a cation exchanger. Determination of acetic acid content of a vinegar sample by potentiometric titration with sodium hydroxide using quinhydrone. Suggested References Physical Chemistry Physical Chemistry, Ira Levine, 5th Edition, 2002 Tata McGraw Hill Publishing Co.Ltd. 		

	-
3. Physical Chemistry, R.J. Silbey, & R.A.	
Alberty, 3rd edition, John Wiley & Sons, Inc [part 1]	
4. Physical Chemistry, G. Castellan, 3rd	
edition, 5th Reprint, 1995 Narosa Publishing	
House.	
5. Modern Electrochemistry, J.O.M Bockris &	
A.K.N. Reddy, Maria Gamboa – Aldeco 2nd	
Edition, 1st Indian reprint,2006 Springer	
6. Visible & U.V. Spectroscopy, Analytical	
Chemsitry by Open Learning R. Demny and	
R. Sinclair M 1991 John Wiley & Sons	
7. Classical Methods, Vol 1 Analytical	
Chemistry by Open Learning D. Cooper & C. Devan,1991 John Wiley & Sons	
8. Physical Chemistry, G.M. Barrow, 6th	
Edition, Tata McGraw Hill Publishing Co.	
Ltd. New Delhi.	
9. The Elements of Physical Chemistry, P.W.	
Atkins, 2nd Edition, Oxford University	
Press Oxford	
10. Physical Chemistry, G.K. Vemullapallie,	
1997, Prentice Hall of India, Pvt.Ltd. New Delhi.	
Denn.	
Inorganic Chemistry.	
1. D. Banerjea, <i>Coordination chemistry</i> , Tata	
McGraw Hill, New Delhi, (1993).	
2. D. F. Shriver and P. W. Atkins, Inorganic	
chemistry, 3 rd Ed., Oxford University Press,	
(1999).	
3. K. F. Purcell and J. C. Kotz, <i>Inorganic chemistry</i> ,	
Saunders, Hongkong, (1977).	
Saunders, Hongkong, (1977).4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of</i>	
 Saunders, Hongkong, (1977). 4. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 	
Saunders, Hongkong, (1977).4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of</i>	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and</i> 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of</i> 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of</i> <i>elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and</i> <i>models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of</i> <i>bioinorganic chemistry</i>, Dhuri and Sons, 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of bioinorganic chemistry</i>, Dhuri and Sons, Calcutta, (1988). 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of bioinorganic chemistry</i>, Dhuri and Sons, Calcutta, (1988). 8. R. W. Hay, <i>Bioinorganic chemistry</i>, Ellis 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of bioinorganic chemistry</i>, Dhuri and Sons, Calcutta, (1988). 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of bioinorganic chemistry</i>, Dhuri and Sons, Calcutta, (1988). 8. R. W. Hay, <i>Bioinorganic chemistry</i>, Ellis Harwood, England, (1984). 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of bioinorganic chemistry</i>, Dhuri and Sons, Calcutta, (1988). 8. R. W. Hay, <i>Bioinorganic chemistry</i>, Ellis Harwood, England, (1984). 9. R. C. Mehrotra and A. Singh, <i>Organometallic chemistry: A unified approach</i>, Wiley Eastern, New Delhi, (1991). 	
 Saunders, Hongkong, (1977). 4. N. N. Greenwood and E. Earnshaw, <i>Chemistry of elements</i>, Pergamon Press, Singapore, (1989). 5. W. L. Jolly, <i>Modern inorganic chemistry</i>, 2nd Ed. McGraw Hill Book Co., (1991). 6. B. E. Douglas and H. McDaniel, <i>Concepts and models in inorganic chemistry</i>, 3rd Ed., John Wiley & Sons, Inc., New York, (1994). 7. G. N. Mukherjee and A. Das, <i>Elements of bioinorganic chemistry</i>, Dhuri and Sons, Calcutta, (1988). 8. R. W. Hay, <i>Bioinorganic chemistry</i>, Ellis Harwood, England, (1984). 9. R. C. Mehrotra and A. Singh, <i>Organometallic chemistry: A unified approach</i>, Wiley Eastern, 	

r	
	and B. W. Rockett,Van Nostrand Reinhold Company London1972. P 34. 11. For preparation of CuCl ₂ .2DMSO Refer Microscale Inorganic Chemistry by Z. Szafran, Ronald M. Pike and Mono M. Singh. Pub. John Wiley and Sons1991.p.218.
	 Organic Chemistry 1. Organic Chemistry, Francis A Carey, Pearson Education, 6th Edition, Special Indian Edition 2008 2. Organic Chemistry, R.T. Morrison and R.N. Boyd, 6th Edition, Pearson Edition 3. Organic Chemistry, T.W.G. Solomon and C.B. Fryhle, 8th Edition, John Wiley & Sons, 2004 4. A guide to mechanism in Organic Chemistry, 6th Edition, Peter Sykes, Pearson Education 5. Fundamentals of Organic Chemistry, G. Marc Loudon, 4th Edition Oxford 6. Organic Chemistry, L.G. Wade Jr and M.S. Singh, 6th Edition,2008 7. Organic Chemistry, J.G. Smith, 2nd Editionm Special Indian Edition, Tata McGraw Hill 9. Organic Chemistry, S.H. Pine, McGraw Hill Kogakusha Ltd. 10. Stereochemistry, P.S. Kalsi, New Age International Ltd. 4th Edition,2006
	 Analytical Chemistry 1. D. Harvey, Modern Analytical Chemistry, The McGraw-Hill Pub. 1st Edition (2000) 2. H.S. Ray, R Sridhar and K.P. Abraham, Extraction of Nonferrous Metals, AffiliatedEast- West Press Pvt. Ltd. New Delhi (1985) reprint 2007. 3. G.H. Jeffery, J. Bassett, J. Mendham and R.C. Denney , Vogel's Textbook of Qunatitative Chemical Analysis, Fifth edition, ELBS Publication (1996) 4. D.A. Skoog D.M. West and F.J. Holler, Fundametals of Analytical Chemistry, 7thEdition (printed in India in 2001) ISBN Publication. 5. Analytical Chemistry, J.G. Dick,1973 Tata McGraw Hill Publishing Co. Ltd. New Delhi. 6. Quantitative analysis, Dey & Underwood,

		7. Fundamentals of Analytical Chemistry, Skoog etal 8th edition, Saunders college publishing.		
--	--	---	--	--