



Fatima Mata National College **(Autonomous)** **Kollam**

Scheme & Syllabus of
Post Graduate Degree in Chemistry
2019 Admission Onwards

PREAMBLE

The syllabus of M.Sc programme in Chemistry offered under Semester system has been revised and the revised syllabus is to be effective from 2019 admission. M.Sc programme shall extend over a period of two academic years comprising of four semesters, each of 450 hours in 18 weeks duration. The syllabus and scheme of examinations of this programmes are detailed below.

M.Sc. PROGRAMME IN CHEMISTRY (Revised syllabus under semester system with effect from 2019 admission) SYLLABUS AND SCHEME OF EXAMINATION

| Course No and Title | Hours per week | | Duration for ESA in hours | Marks for CA | Marks for ESA | Total marks |
|---|----------------|---|------------------------------|--------------------------------|----------------------------|-------------|
| | L | P | | | | |
| Semester I | | | | | | |
| 19PCH11 Inorganic Chemistry I | 5 | | 3 | 25 | 75 | 100 |
| 19PCH12 Organic Chemistry I | 5 | | 3 | 25 | 75 | 100 |
| 19PCH13 Physical Chemistry I | 5 | | 3 | 25 | 75 | 100 |
| 19PCH14 Inorganic Practicals I | | 3 | | To be continued in semester II | | |
| 19PCH15 Organic Practicals I | | 3 | | To be continued in semester II | | |
| 19PCH16 Physical Practicals I | | 4 | | To be continued in semester II | | |
| | | | | | Total marks for semester I | 300 |
| Distribution of teaching hours/week: Theory-15 hours, practicals-10 hours | | | | | | |

| Course No and Title | Hours per week | | Duration for ESA in hours | Marks for CA | Marks for ESA | Total marks |
|---|----------------|---|------------------------------|-----------------|-----------------------------|-------------|
| | L | P | | | | |
| Semester II | | | | | | |
| 19PCH21 Inorganic Chemistry II | 5 | | 3 | 25 | 75 | 100 |
| 19PCH22 Organic Chemistry II | 5 | | 3 | 25 | 75 | 100 |
| 19PCH23 Physical Chemistry II | 5 | | 3 | 25 | 75 | 100 |
| 19PCH14 Inorganic Practicals I | | 3 | 6 | 25 | 75 | 100 |
| 19PCH15 Organic Practicals I | | 3 | 6 | 25 | 75 | 100 |
| 19PCH16 Physical Practicals I | | 4 | 6 | 25 | 75 | 100 |
| | | | | | Total marks for semester II | 600 |
| Distribution of teaching hours/week: Theory-15 hours, practicals-10 hours | | | | | | |

| Course No and Title | Hours per week | | Duration for ESA in hours | Marks for CA | Marks for ESA | Total marks |
|---|----------------|---|------------------------------|--------------------------------|------------------------------|-------------|
| | L | P | | | | |
| Semester III | | | | | | |
| 19PCH31 Inorganic Chemistry III | 5 | | 3 | 25 | 75 | 100 |
| 19PCH32 Organic Chemistry III | 5 | | 3 | 25 | 75 | 100 |
| 19PCH33 Physical Chemistry III | 5 | | 3 | 25 | 75 | 100 |
| 19PCH34 Inorganic Practicals II | | 3 | | To be continued in semester IV | | |
| 19PCH35 Organic Practicals II | | 3 | | To be continued in semester IV | | |
| 19PCH36 Physical Practicals II | | 4 | | To be continued in semester IV | | |
| | | | | | Total marks for semester III | 300 |
| Distribution of teaching hours/week: Theory-15 hours, practicals-10 hours | | | | | | |

| Course No and Title | Hours per week | | Duration for ESA in hours | Marks for CA | Marks for ESA | Total marks |
|---|----------------|---|------------------------------|--|------------------|-------------|
| | L | P | | | | |
| Semester IV | | | | | | |
| 19PCH41 Chemistry of Advanced Materials | 5 | | 3 | 25 | 75 | 100 |
| *19PCH42 (a) Inorganic Chemistry IV | 5 | | 3 | 25 | 75 | 100 |
| *19PCH42(b) Organic Chemistry IV | | | | | | |
| *19PCH42(c) Physical Chemistry IV | | | | | | |
| 19PCH34 Inorganic Practicals II | | 3 | 6 | 25 | 75 | 100 |
| 19PCH35 Organic Practicals II | | 3 | 6 | 25 | 75 | 100 |
| 19PCH36 Physical Practicals II | | 4 | 6 | 25 | 75 | 100 |
| 19PCH43(a) Dissertation | 5 | | | | 70 | 70 |
| 19PCH43(b) Visit to R&D centre | | | | | 5 | 5 |
| Internship (2 weeks to 4 weeks) in collaboration with research institutions | No weightage | | | | | |
| 19PCH44 Comprehensive Viva-Voce | | | | | 25 | 25 |
| | | | | Total marks for semester IV | | 600 |
| | | | | Grand total (semester I +II+III+IV) | | 1800 |
| Distribution of teaching hours/week: Theory-15 hours, Practicals-10 hours | | | | | | |
| *Each student has to choose either (a), (b) or (c) as elective in accordance with the dissertation chosen | | | | | | |

PROGRAMME OUTCOME OF MSc CHEMISTRY

The outcome of the program are as follows.

- To give students a comprehensive understanding of the principles of Chemistry.
- To gain the skill to design and carry out scientific experiments and interpret the data.
- To understand the interdisciplinary nature of Chemistry and to be aware of the emerging fields in Chemistry.
- To build a scientific temper and to learn the necessary skills to succeed in research or industrial field.
- To be able to define and resolve new problems in Chemistry and participate in the future development of chemistry.

SEMESTER I

19PCH11: Inorganic Chemistry-I

Total 90 h

Course Outcome:

CO 1. To get an insight into the various theories of metal complexes.

CO 2. Identify the principles, structure and reactivity of selected coordination complexes.

CO 3. To understand the applications of various techniques to evaluate analytic data and Solve problems based on various analytical concepts.

CO 4. Quantify analytes with proper data handling and analysis

CO 5. To be able understand the structure of various metal-metal bonds and metal clusters

CO 6. To gain a comprehensive idea of the preparation, properties and use of isopoly, heteropoly acids, noble gases and interhalogen compounds

CO 7. To understand the chemistry of processes in atmosphere, and impacts of various pollutants on destruction of ozone, hydrosphere, lithosphere.

Unit I Coordination chemistry-I: Theories of metal complexes 18 h

Types of ligands and complexes. Isomerism: Structural, geometrical and optical isomerism. Crystal field theory: Splitting of d orbitals in octahedral, tetragonal, square planar, tetrahedral, trigonal bipyramidal and square pyramidal fields. Jahn-Teller theorem, evidence for JT effect, static and dynamic JT effect. Crystal field stabilization energy (CFSE) and its calculations. Octahedral Site Stabilization Energy. Factors affecting the splitting parameter. Spectrochemical series. Evidence of covalency in Metal-Ligand bond, introduction to Ligand field theory. Molecular orbital theory. Sigma and pi bondings in complexes. MO diagrams of octahedral and tetrahedral complexes with and without pi bonds. Experimental evidence of pi bond on the stability of sigma bond. Nephelauxetic effect.

Unit II Analytical principles 18 h

Evaluation of analytical data: Accuracy and precision. Standard deviation, variance and coefficient of variation. Student 't' test, 'Q' test, and 'F' test. Confidence limits. Errors: Classification, distribution, propagation, causes and minimization of errors. Significant figures and computation rules. Correlation analysis: Scatter diagram. Correlation coefficient, r. Calculation of r by the method of least squares. Volumetric methods: Classification of reactions in volumetry. Theories of indicators. Acid-base, redox, adsorption, metallochromic indicators. Complexometric titrations: Titration using EDTA-direct and back titration methods. Precipitation titrations. Redox titrations. Titrations in non-aqueous solvents. Organic reagents used in gravimetry: Oxine, dimethylglyoxime and cupferron. Principle and instrumentation of TG, DTA and DSC. Factors affecting TG and DTA curves. Applications of TG, DTA and DSC in the study of metal complexes.

Unit III Metal-metal bonds and metal clusters 18 h

Metal-metal bonds: Factors affecting the formation of metal-metal bond. Dinuclear compounds of Re, Cu and Cr, metal-metal multiple bonding in $(Re_2X_8)^{2-}$, Trinuclear clusters, tetranuclear clusters, hexanuclear clusters. Polyatomic zintl anion and cations. Infinite metal chains. Metal carbonyl clusters. Anionic and hydrido clusters. LNCCs and HNCCs. Isoelectronic and isolobal relationships. Hetero atoms in metal clusters: Carbide and nitride containing clusters. Electron counting schemes for HNCCs. Capping rule. Chalcogenide clusters. Chevrel phases.

Unit IV Isopoly and heteropoly acids, Noble gases, interhalogens 18 h

Preparation, properties and structure of isopoly acids of Mo, W and V and Heteropoly acids of Mo and W. Preparation and properties of Xenon fluorides and Krypton compounds ($KrCl_4$, KrF_4 , KrF_2 , $KrBr_6$, $Kr_2Cr_2O_7$, $KrCrO_4$ & KrO_2), structure of XeF_2 (MO theory only). Preparation, bonding and uses of inter halogen compounds. Properties and structure of aluminosilicates and zeolites, shape selectivity. Preparation, properties and applications of silicones.

Unit V Chemistry of Natural Environmental Processes 18 h

The chemistry of processes in atmosphere; Composition of the atmosphere. Automobile pollutants and the catalytic converter. Photochemical smog. Chemistry of the stratosphere. Catalytic destruction of ozone. Depletion of the ozone layer. Hazards of common air pollutants on the human health. The Chemistry of

processes in hydrosphere; The hydrologic cycle. Cycling and purification. The unique properties of water. Acid base properties. CO₂ in water. Alkalinity. O₂ consuming waste. DO, BOD and COD. The chemistry of processes in Lithosphere; Redox status in soil. pE, pH predominance diagrams for redox sensitive elements. Acidity in soil materials. Acid neutralization capacity and the quantification of the soil acidity. Ion speciation in soil solution. Cation exchange capacity and exchange phase composition.

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10. A.S. Kunju and G. Krishnan, Group Theory and its Applications in Chemistry, PHI Learning,
11. R.L.Carter, Molecular Symmetry and Group Theory, John Wiley & Sons, 1998
12. E. James Girard, Principles of Environmental Chemistry, Jones and Bartlett Publishers, 3rd Edition,
13. H.V. Jadhav, Elements of Environmental Chemistry, Himalya Publication House, 2010
14. E. Michael Essington, Soil and Water Chemistry, CRC Press, 2nd edition, 2015.

19PCH12: ORGANIC CHEMISTRY-I

Total 90 h

Course Outcome:

CO 1. To understand the basic concepts and mechanism in organic chemistry.

CO 2. To be familiarize with the mechanism of organic reactions and different factors which affect the reaction rate.

CO 3. To understand the role of various reaction intermediates like carbanion, carbocation, carbenes, radicals etc. in organic reactions. Comprehend the structure-reactivity pattern of reactive intermediates involved in organic reactions.

CO4. To get insight into the chemistry of substitution reactions, elimination and addition reactions.

CO5. To able to apply the various organic synthetic reagents in the field organic synthesis.

Unit I Stereochemistry of organic compounds

18h

Nomenclature of organic compounds - Cyclic, fused polycyclic and bridged polycyclic hydrocarbons, Bridged and fused hydrocarbon systems, Spirocyclic hydrocarbon systems, Heterocyclic systems containing Nitrogen and Oxygen.

Introduction to molecular symmetry and chirality, Axial Chirality, Planar Chirality and Helicity, Relative configuration, Stereochemical nomenclature, *R* and *S*, *E* and *Z*. Prostereoisomerism, stereotopicity and stereoprojections. Non-carbon chiral centres –Nitrogen, phosphorus and sulfur as chiral centres. Axial stereochemistry: Atropisomerism and its designation, *M* and *P* configurations. Stereoselectivity: enantioselectivity, diastereoselectivity and stereoconvergence. Basic introduction to chiral separation methods and estimation of enantiomeric excess.

Conformational analysis of alkanes and cycloalkanes, Biased systems. Effect of conformation on reactivity of cyclohexanes – conformation of decalin.

Introduction to ORD, CD- their application in assigning configuration. Sector rules such as octant and axial haloketone rules. Cotton effect.

Chiral drugs: Ibuprofen, Methyldopa, and Thalidomide – Structure, chirality and activity (Basic concepts only)

Unit II Structure, reactivity and intermediates

18 h

Reaction coordinates- difference between transition state and intermediates, Homolytic and heterolytic bond fissions. Formation and structure of carbocations, carbanion and free radicals, Stability of intermediates, influence of field effect, inductive, mesomeric and steric effects on controlling stability of carbocations, carbanions and carbon-centered radicals. Influence of structural features on acidity, basicity and reactivity of organic compounds. Alkyl, aralkyl and allylic cations –influence of substituents. General reactions of carbocations, carbanions and free radicals. Introduction to radical ions. Formation, structure, stability and chemical reactions of carbenes, nitrenes and arynes.

Unit III Substitution reactions

18 h

Nucleophilic substitution at sp^3 carbon - S_N1 and S_N2 mechanisms. Walden inversion, stereochemistry. Effect of solvent, leaving group and substrate structure on rates of S_N1 and S_N2 substitutions. Neighbouring group participation, Non-classical carbocations, Competition between S_N1 and S_N2 reactions. S_N1' , S_N2' , S_Ni mechanism.

Mechanism of esterification and ester hydrolysis-acid catalysed and base catalysed reactions.

Aromatic Substitution reactions - Electrophilic substitution: mechanism and evidence- Reactions involving nitrogen, sulphur, carbon, halogen and oxygen electrophiles. Directive and rate controlling factors in aromatics with one or more substituents.

Aromatic Nucleophilic Substitution reactions - S_N1 , S_NAr , benzyne and $S_{RN}1$ mechanism and evidence with examples.

Unit IV Elimination and addition reactions

18h

Elimination reactions leading to C=C bond formation and their mechanisms. E1, E2 and E1CB mechanisms. Stereoaspects of C=C bond formation. Effect of leaving group and substrate structure. Hoffmann and Saytzeff elimination. Cis elimination. Stereoaspects of the addition of H_2O , X_2 , HX , and boranes to C=C systems. Effect of substituents on the rate of additions. Cis and trans hydroxylation of cycloalkenes. Nucleophilic addition to activated C=C systems. Structure of the transition state in the addition reactions. Michael addition: mechanism. Addition to Carbon-Heteroatom multiple bonds: Aldol condensation (normal, crossed and directed), Perkin, Stobbe, Knoevenagel, Darzen, Reformatsky and benzoin condensations. Grignard, Cannizzaro, Wittig and Wittig-Horner reactions. Application of Cram's rule, Felkin- Ann model.

Unit V Separation techniques

18 h

Classification of chromatographic methods. Theory of chromatography. Applications of chromatographic methods. Adsorption and partition chromatography. Paper, thin layer and column chromatographic methods. Centrifugal TLC, LC, Pressure column chromatography, HPLC and GC. Column matrices. Detectors. Affinity and chiral separations using HPLC. Normal and ultra-centrifugation. Gel and Capillary electrophoresis and their applications. Solvent extraction. Extraction using supercritical liquid CO_2 , Craig's technique of liquid-liquid extraction.

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19PCH13 PHYSICAL CHEMISTRY –I

90 h

Course Outcome:

- CO1. At the end of the course, the learners should be able to: Use mathematical techniques in linear algebra for eigenvalues and eigenvectors and first and second order differential equations not only in quantum chemistry but in other areas of physical and theoretical chemistry that will be offered during the whole programme.
- CO2. Solve all the model problems in quantum mechanics for which exact analytical methods and solutions are available and will apply them to analyze the basis behind the postulatory method of quantum mechanics and which forms the foundations for advanced study of the subject.
- CO3. Relate concepts that were originally introduced purely as modern atomic physics to molecular systems through harmonic oscillator, spin and rigid rotator.
- CO4. Utilize the principles of group theory to predict the hybridization of molecules and find out the selection rule of various spectral techniques.
- CO5. Understand the various theories of reaction rate.
- CO6. Solve problems on rate/rate constants/efficiency for (i) complex reactions (ii) unimolecular and bimolecular reactions, and (iii) electronically excited state dynamics
- CO7. To know the basics principle of different techniques employed in molecular spectroscopy.
- CO8. To study the origin, instrumentation and important applications of Microwave, IR, Raman, UV techniques.

Unit 1- Quantum Chemistry I

18 h

Classical mechanics and its limitations, need of quantum mechanics, concept of matter wave, de Broglie relation and its experimental proof, uncertainty principle and its consequences.

Postulates of Quantum Mechanics

State function postulate: Born interpretation of the wavefunction, well behaved functions, orthonormality of wave functions. Operator postulate: operator algebra, linear and nonlinear operators, Laplacian operator, commuting and non-commuting operators, Hermitian operators and their properties, eigen functions and eigen values of an operator. Eigen value postulate: eigen value equation, eigen functions of commuting operators. Expectation value postulate. Postulate of time-dependent Schrödinger equation, Quantization of angular momentum, quantum mechanical operators corresponding to angular momenta (L_x , L_y , L_z and L^2)-commutation relations between these operators.

Application of Quantum mechanics to Exactly Solvable Model Problems

Translational motion: free particle in one-dimension, particle in a box with infinite potential barrier- one dimensional box, three dimensional box and cubical box - degeneracy -particle with finite potential barriers-one potential barrier, two finite barriers, potential barriers of definite thickness-Quantum mechanical tunneling (Qualitative concept only)

Vibrational motion: one-dimensional harmonic oscillator (complete treatment), Hermite equation (solving by method of power series), Hermite polynomials, recursion relation, wave functions and energies-important features, Harmonic oscillator model and molecular vibrations.

Symmetric and antisymmetric wave functions, Pauli's antisymmetry principle, the postulate of spin.

Spin orbitals. Spin-orbit coupling.

Vector atom model- Term symbols, explanation of spectral lines (for sand pblocks only)

Unit II Molecular symmetry and Applications of Group Theory

18 h

Symmetry and Character table: Symmetry elements and symmetry operation. Matrix representation of symmetry operations. Character of a matrix. Conditions for a set of elements to form a group. Point groups. Multiplication of operations. Group multiplication table. Similarity transformation and classification of

symmetry operation, Matrix representation of point group. Reducible and Irreducible representations. The Great Orthogonality theorem. Rules derived from GOT (proof not required). Setting up of character table of C_{2v} , C_{3v} and C_{2h} groups. Direct product representations. Reduction formula, reduction of reducible representation to IRs. Transformation properties of atomic orbitals. Hybridisation: identification of atomic orbitals taking part in hybridisation of triangular planar, square planar, trigonal bipyramidal, square pyramidal and tetrahedral molecules.

Spectroscopic applications: Transition moment integral transition moment operator. Vanishing matrix element. Symmetry selection rule for IR, Raman and electronic spectra.

Unit III Spectroscopy –I

18 h

Microwave spectroscopy: Rotational spectrum, intensity of spectral lines, calculation of internuclear distance. Non-rigid rotors and centrifugal distortion. Rotational spectra of polyatomic molecules-linear and symmetric top molecules. Introduction to instrumentation.

Vibrational Spectroscopy: Vibrational spectra of harmonic and anharmonic oscillator. Selection rules. Morse curve, fundamentals and overtones. Determination of force constant. Rotational fine structure, P,Q,R branches of spectra. Vibrational spectra of polyatomic molecules: Normal modes, classification of vibrational modes into stretching (asymmetric, symmetric), bending, parallel and perpendicular vibrations. Finger print region and group frequencies. Introduction to FTIR and instrumentation.

Raman spectroscopy: Raman scattering, polarisability and classical theory of Raman spectrum. Rotational and vibrational Raman spectrum. Raman spectra of poly atomic molecules. Complementarity of IR and Raman spectra. Mutual exclusion principle. Introduction to instrumentation. Laser Raman spectrum.

Electronic spectra. Electronic spectra of diatomic molecules. Vibrational coarse structure and rotational fine structure of electronic spectrum. Franck-Condon principle. Types of electronic transitions. Forrat diagram. Predissociation. Calculation of heat of dissociation. Electronic spectra of polyatomic molecules: Electronic transition among molecular orbitals and absorption frequencies. Effect of conjugation. Introduction to instrumentation. Simultaneous determination of two components.

Unit IV Chemical kinetics

18 h

Theories of reaction rates: Collision theory and its failure,. Transition state theory-Eyring equation. Comparison of the two theories. Thermodynamic formulation of the reaction rates. potential energy surfaces Theories of unimolecular reactions- Lindemann theory, Lindemann-Hinshelwood mechanism, qualitative idea of RRKM theory,

Kinetics of complex reactions-parallel reactions, opposing reactions, consecutive reactions and chain reactions, steady state treatment, kinetics of H_2-Cl_2 and H_2-Br_2 reactions, decompositions of ethane, acetaldehyde and N_2O_5 . Rice-Herzfeld mechanism, branching chain reactions Hinshelwood mechanism of chain reactions and explosion.

Fast reactions: Relaxation method, relaxation spectrometry, flow method, shock method, fast mixing method, field jump method, pulse method, Flash photolysis and NMR method.

Reactions in solution: Factors affecting reaction rates in solutions, effect of dielectric constant and ionic strength, cage effect, Bronsted-Bjerrum equation, primary and secondary kinetic salt effect, influence of solvent on reaction rates, significance of volume of activation, linear free energy relationship. Hammett equation and Taft equation.

Unit V Gaseous and liquid state

18 h

Maxwell's distribution of molecular velocities, influence of temperature on molecular velocities, types of molecular velocities-average velocity and most probable velocity and its determination from Maxwell's equation Transport phenomena in gases-viscosity of gases, Chapman equation, determination of viscosity of gases, calculation of mean free path, Thermal conductivity, diffusion, Degrees of freedom of gaseous molecules Translational, Rotational and vibrational. Equation of state of real gases-van der Waal's equation, Other equation of states—Redlich-Kwong equation, Clausius equation, Virial equation, second virial coefficient and determination of diameter of a molecule.

Inter molecular forces—dipole-dipole interaction, induced dipole- dipole, induced dipole-induced dipole interactions

Liquid state Liquid vapour equilibria, vapour pressure- methods of measuring vapour pressure - barometric method and dynamic method - equation of state for liquids, structure of liquids, X-ray diffraction of liquids-

vacancy model for a liquid, pair correlation function, surface tension, determination of surface tension, drop weight method and drop number method, viscosity, determination of coefficient of viscosity using Ostwald viscometer.

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19PCH14: INORGANIC CHEMISTRY PRACTICALS -I

Total 125 h

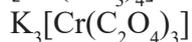
Course Outcome:

CO1. To be able to identify and separate rare/less familiar ions such as Ti, W, Se, Mo, Ce, Th, Li, Zr, V, U etc.

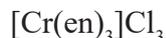
CO2. to estimate colorimetrically ions such as Fe, Cu, Ni, Mn, Cr and Mn etc.

CO3. Developing the skill to prepare various metal complexes thereby developing research aptitude.

1. Separation and identification of rare/less familiar cations such as Ti, W, Mo, Th, Zr, V, U and Li
2. Volumetric estimation using EDTA, ammonium vanadate, ceric sulphate etc.
3. Colorimetric estimation of Cr, Fe, Mn, Ni, Cu etc.
4. Preparation of metal complexes: selection can be made from the following or any other from the existing literature.



Cis and trans isomers of $[\text{Co}(\text{en})_2\text{Cl}_2]\text{Cl}$



References

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2. A. I. Vogel, A Text Book of Qualitative Inorganic Analysis, Longman 5th edition, 1979.

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19PCH15: ORGANIC PRACTICALS-I

Total 125 h

Course Outcome:

- CO1. To learn the separation and purification of an organic mixture by chemical/solvent separation methods.
 CO2. To gain the skill to separate the mixture of organic compound using TLC plate.
 CO3. To identify the organic compounds based on their R_f values.
 CO4. To be able to conduct two stage organic preparations.
 CO5. To draw the structure of compounds using Chemdraw software

A. Separation and identification of organic compounds

1. Quantitative wet chemistry separation of a mixture of two components by solvent extraction
2. TLC of the purified samples along with the mixture in same TLC **plates (component 1 with mixture and component 2 with mixture on separate TLC plate)** and calculation of R_f values-Reporting and recording TLC in standard formats- preparation of sample solution, adsorbent, dimensions of the plate, saturation time, developing time, visualization and detection, R_f Value, Drawing - in the form of a table.

B. Separation of a mixture of by column chromatography (not for End semester evaluation)

- 1) Malachite green and methylene blue
- 2) *o*-nitroaniline and *p*-nitroaniline.

C. Preparation of compounds by two stages.

Recording/downloading UV, IR, ^1H NMR and ^{13}C NMR and EI mass spectra of synthesized compounds.
 TLC analysis- (stage 1 reactants and products on TLC plate 1 and stage 2 reactants and products on plate 2)-
 Record TLC in standard format as in separation

All preparations must be restricted to 1 g level

Nitration

- 1) Acetanilide \rightarrow *p*-nitroacetanilide \rightarrow *p*-nitroaniline
- 2) Methylbenzoate \rightarrow methyl *m*-nitrobenzoate \rightarrow *m*-nitrobenzoic acid

Bromination

- 3) Acetanilide \rightarrow *p*-bromoacetanilide \rightarrow *p*-bromoaniline

Aldol condensation- Synthesis of heterocycles

- 4) benzaldehyde \rightarrow Dibenzylideneacetone \rightarrow 1,5-Diphenyl-3-styryl-2-pyrazoline

Diazocoupling

- 5) Aniline \rightarrow Diazoaminobenzene \rightarrow *p*-aminoazobenzene

Rerrangement

- 6) Pthalic anhydride \rightarrow Pthalimide \rightarrow anthranilic acid

Synthesis of Dyes

- 7) N,N-Dimethylaniline \rightarrow N,N-dimethyl-4-nitrosoaniline \rightarrow methylene blue

The board of examiners have to select either TLC of separated components *OR* TLC of preparation for an examination. But both TLC examinations are to be practiced and entered in the record of experiments

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7. P. F Shalz, *Journal of Chemical Education*, **1996**, 173: 267.
8. Monograph on green laboratory experiments, DST, Government of India, pp 1-79.
9. For spectral data of organic compounds, see: <http://sdfs.riondb.aist.go.jp/sdfs/cgi-bin/direct frame top.cgi>.

19PCH16 PHYSICAL PRACTICALS –I

125 h

Course Outcome:

CO1. To verify the some important principles in physical chemistry and to determine various physical properties.

CO2. To learn to carry out some simple computational chemistry calculations

Adsorption

Freundlich and Langmuir isotherms for adsorption of acetic/oxalic acid on active charcoal. Determination of concentration of acetic/ oxalic acid.

Kinetics

Determination of rate constant of acid hydrolysis of methyl acetate.

Determination of Arrhenius parameters.

Determination of concentration of given acid.

Determination of rate constant of the saponification of ethyl acetate and evaluation of Arrhenius parameters.

Determination of rate constant of reaction between $K_2S_2O_8$ and KI.

Study the kinetics of iodination of acetone in acid medium.

Phase rule

Solid-liquid equilibria

Construction of phase diagram and determination of the composition of unknown mixture (naphthalene/biphenyl, naphthalene/benzophenone, naphthalene/diphenyl amine)

Construction of phase diagram with congruent melting point- naphthalene/metadinitrobenzene

Partially miscible liquid pairs- CST of phenol-water system.

Effect of impurities (KCl/ NaCl/ succinic acid) on the miscibility temperature of phenol-water system and hence the determination of concentration of given unknown solution.

Three component system- Construction of ternary phase diagram of acetic acid chloroform-water system and hence the composition of given homogeneous mixture. Construction of tie-line.

Distribution law

Distribution coefficient of ammonia between chloroform and water. Determination of equilibrium constant of copper- ammonia complex by partition method or coordination number of Cu^{2+} in copper-ammonia complex.

Distribution coefficient of benzoic acid between toluene and water.

Distribution coefficient of iodine between hexane and water/ $CHCl_3$ and water/ CCl_4 and water

Determination of the equilibrium constant of the reaction $KI + I_2 \rightleftharpoons KI_3$ and hence the concentration of given KI in hexane and water/ $CHCl_3$ and water/ CCl_4 and water.

Determination of hydrolysis constant of anilinium hydrochloride.

Dilute Solutions

Determination of K_f of solid solvent, molar mass of non-volatile solute, mass of solvent and composition of given solution (Solvent- Naphthalene/Biphenyl/ Benzophenone etc.

Solute- Naphthalene/ Biphenyl/ Diphenylamine etc)

Determination of vant Hoff's factor for benzoic acid in Naphthalene. Determination of atomicity of sulphur.

Transition temperature

Determination of K_T of salt hydrate, molar mass of solute, mass of salt hydrate and composition of given solution (Solvent- $Na_2S_2O_3 \cdot 5H_2O/CH_3COONa \cdot 3H_2O$, Solutes glucose, sucrose, urea)

Thermochemistry

Determination of the concentration of given strong acid/alkali.

Thermometric titration of NaOH Vs standard HCl.

Heat of displacement of Cu^{2+} by Zn.

Determination of the heat of ionisation of acetic acid.

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1. V. D. Athawal, "Experimental Physical Chemistry", New Age International, 1st edn., 2001.
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MODEL QUESTION PAPER
19PCH11: INORGANIC CHEMISTRY- I

Time : 3 Hrs

Max. Marks: 75

SECTION A

Answer **two** among (a), (b) and (c) from **each** question carries **2** marks

- What is meant by Nephelauxetic effect?
 - Explain linkage isomerism with suitable example.
 - What is meant by crystal field stabilization energy?
- What do you mean by significant figure? How many significant figures are in the following?
 - 0.0026
 - 6.023×10^{23}
 - What are metallochromic indicators? Give an example.
 - In a volumetric experiment the volumes of the titrant used are 9.98, 9.99, 9.98, 9.95, 10.00 and 10.02 mL. Calculate the standard deviation.
- The electronic spectrum of $[\text{Ti}(\text{H}_2\text{O})_6]^{3+}$ exhibits a double humped curve. Why?
 - What are Pascals constant? Where are they used?
 - Chromium acetate dihydrate crystals are diamagnetic at room temperature. Why?
- Explain the term 'shape selectivity'.
 - Give the preparation of KrCl_4 and KrO_2 .
 - What are zeolite? Explain their use as water softeners.
- Brief the role of catalytic converters in automobiles.
 - Explain the formation of photochemical smog.
 - Mention the different regions of atmosphere.

(2x10= 20 marks)

SECTION B

Answer either among (a) or (b) from **each** question carries **5** marks

- State and illustrate Jahn Teller effect.
 - Explain the crystal field theory of octahedral complexes.
- Give a brief note on scatter diagram and its significance.
 - Explain the titrations in non-aqueous solvents.
- Two complexes (A and B) of Ni (II) are believed to be octahedral and tetrahedral. Each has three absorption bands, but 'A' has $\epsilon = 10$ and 'B' has $\epsilon = 150$. Which one of them is probably a tetrahedral complex? Explain measurement of which physical property would exclude the probability of either complex being square planar. Explain.
 - Compare the electronic spectra of lanthanides with that of the first row transition metals. Why do the lanthanides ions give rise to very sharp bands in their electronic spectra?
- Give an account of inter halogen compounds.
 - Write a short note on silicones.
- List out the major pollutants. Outline how they affect human health?
 - Describe how we can quantify soil acidity.

(5x5= 25 marks)

SECTION C

Answer **any three** questions. **Each** question carries **10** marks

11. Explain the bonding in octahedral complexes with and without pi bonds using MO Theory.
12. Briefly explain the principle, instrumentation and applications of TG and DTA.
13. Explain, what is meant by spin-only magnetic moment. Illustrate with suitable examples the application of the values of spin-only magnetic moments in the structural elucidation of transition metal complexes.
14. Write a short note on the preparation and properties of heteropoly acids of Mo and W.
15. What are Pourbaix diagrams? Outline its role in explaining the chemistry of processes in lithosphere.

(10x3= 30 marks)

MODEL QUESTION PAPER 19PCH12: Organic Chemistry-I

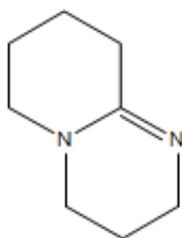
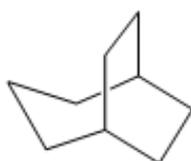
Time: 3 hours

Maximum marks:75

Section A

Answer **any two** among (a), (b) and (c) from each question. Each sub question carries 2 marks

1. a) Write IUPAC names of the following.



- b) Indicate the element of symmetry present in each of the following molecules.

- i) *trans*-1,4-dichlorocyclohexane-



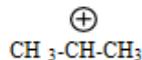
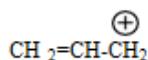
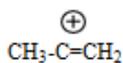
i)

- c) What is atropisomerism?

2. a) How arynes are formed?

- b) *p*-Nitroaniline is less basic than *m*-nitroaniline, explain.

- c) Arrange the following carbocations in order of increasing stability and give reasons.



3. a) What is S_Ni reaction?

- b) Alkaline hydrolysis of $\text{Et}_2\text{NCH}(\text{Cl})\text{CH}_2\text{CH}_3$ produces $\text{Et}_2\text{NCH}(\text{Et})\text{CH}_2\text{OH}$. Account for this observation

- c) Write two examples of non classical carbocations.

4. a) How would you prepare *trans*-1,2-dihydroxycyclohexane from cyclohexene?

- b) Write the mechanism of benzoin condensation.

- c) How can the E1CB pathway be distinguished from the kinetically indistinguishable E2 pathway?

5. a) What is the principle of chromatography

- b) What is paper chromatography? How is it helpful in identifying various alpha amino acids?

- c) Outline the applications of Gel electrophoresis.

(2x10= 20 marks)

Section B

Answer **either (a) or (b)** from each question. Each sub question carries 5 marks

6. a) Compare ORD and CD and explain their relationship

- b) Write a note on stereochemistry of nitrogen compounds

7. a) Explain the Felkin-Ann model with an example

- b) How carbenes are generated? Explain its structure and properties

8. a) Explain why anti Markonikoff's addition is not exhibited by HCl or HI when reacted with 1-butene?
b) After standing in aqueous acid *R*-2-butanol is found to have lost its optical activity. Account for this observation.
9. a) Write a note on cis and trans hydroxylation of alkenes
b) State Cram's rule. Explain it with suitable example
10. a) Explain the principle of gas chromatography and ion exchange chromatography. What type of substances are analysed using the above?
b) Describe the principle and instrumentation of HPLC

(5x5= 25 marks)

Section C

Answer **any three** questions. **Each** question carries **10** marks

11. Give a brief account on stereoselectivity, enantiomeric excess and chiral separation.
12. Explain the following
a) S_NAr mechanism, b) Orientation effect in aromatic electrophilic substitution
13. Discuss the following
a) competition between S_N1 and S_N2 b) Stereochemistry of nucleophilic substitution
14. Describe the following
a) Mechanism with evidences of aldol condensation
b) Wittig reactions and applications
15. With a schematic diagram explain the principle, instrumentation, and applications of GC

(10x3= 30 marks)

MODEL QUESTION PAPER 19PCH13: Physical Chemistry- I

Time: 3 h

Max.Marks:75

Section A

Answer any two from **a,b,c**, of each question. Each sub question carries **2** marks.

1. (a) Calculate de Broglie wave length of mass 1 mg moving with a velocity of 10 m s^{-1}
(b) What is an operator? Give example.
(c) Write spectroscopic term symbol for the ground state of O atom
2. (a) Identify the symmetry elements present in the following and assign the point group i) H_2O ii) HCl
(b) Explain improper axis of symmetry.
(c) What is meant by character table.
3. (a) What are overtones? Why are they weak?
(b) State and explain the rule of mutual exclusion with one example.
(c) State Franck-Condon principle.
4. (a) How is nmr spectroscopy made use of in the study of fast reactions?
(b) What is steady state approximation?
(c) Define quantum yield
5. (a) What is the effect of temperature on the distribution of molecular velocities of a gas? Explain.
(b) Calculate the root mean square velocity of nitrogen at 27°C
(c) Write the virial equation of state. Explain the terms.

(10 x 2 = 20 marks)

Section B

Answer either **a** or **b** of each question. Each question carries **5** marks.

6. (a) Explain Quantum mechanical Tunneling
(b) Write kinetic energy operator. Show that it is a Hermitian operator
7. (a) Construct the multiplication table for the symmetry operations of NH_3 molecule.
(b) State and explain orthogonality theorem.
8. (a) Explain the principle and application of Laser Raman Spectrum.
(b) Explain the origin of P and R branches in rotational-vibrational spectrum.

9. (a) Explain Jabalonski diagram
(b) Derive the rate law for the decomposition of N_2O_5
10. (a) Calculate the viscosity of O_2 at 25 °C. The molecular diameter is 3.6 Å
(b) Write any one method for the determination of surface tension of liquid

(5 x 5 = 25 marks)

Section C

Answer any **three** questions. Each question carries **10** marks.

11. Apply Schrodinger Wave equation for a simple harmonic oscillator. Find eigen values and eigen functions
12. Construct the character table for C_{2v} and explain.
13. (i) Give an account of rotation spectra of diatomic molecules. Explain the effect of non rigidity of the bond on the spectra.
(ii) How is the rotational spectrum of a diatomic molecule affected by isotopic substitution?
14. Explain chain reactions. Discuss Semionoff Henshelwood theory of branching chain reactions
15. Discuss viscosity of a gas and Chapman equation. How can we calculate mean free path and collision diameter from viscosity determination

(3 x 10 = 30 marks)

SEMESTER II

19PCH21: INORGANIC CHEMISTRY –II

Total 90 h

Course Outcome:

CO1. To get an insight into the preparation and properties of sulphur, nitrogen, phosphorus and boron compounds.

CO2. To know the structure and bonding of important coordination compounds.

CO3. To understand the magnetic properties of complexes and to know how magnetic moments can be employed for the interpretation of their structure.

CO4. To learn the extraction and separation of lanthanides and actinides.

CO5. To be able to apply the extraction techniques to explore the mineral wealth of beach sands of Kerala.

CO6. Arrive at the chemical compositions based on unit cell contents and fractional coordinates.

CO8. Identify and apply a suitable strategies for synthesizing inorganic crystalline solids in polycrystalline and single crystal forms.

CO9. Correlate and Predict structure-composition-properties (magnetic, electrical and optical) in inorganic crystalline solids.

Unit I Sulphur, nitrogen, phosphorus and boron compounds

18 h

Sulphur-nitrogen compounds: Tetrasulphur tetranitride, disulphur dinitride and polythiazyl S_xN_y compounds. S-N cations and anions. Sulphur-phosphorus compounds: Molecular sulphides such as P_4S_3 , P_4S_7 , P_4S_9 and P_4S_{10} . Phosphorous-nitrogen compounds: Phosphazines. Cyclo and linear phosphazines. Boron-nitrogen compounds: Borazine, substituted borazines and boron nitride. Boron hydrides: Reactions of diborane. Structure and bonding. Polyhedral boranes: Preparation, properties, structure and bonding. The topological approach to boron hydride structure. *Styx* numbers. Importance of icosahedral framework of boron atoms in boron chemistry. Closo, nido and arachno structures. Wade's rules. Carboranes and metallocarboranes.

Unit II Coordination chemistry-II: Spectral and magnetic properties of transition metal complexes

18 h

Electronic spectra of metal complexes- Term symbols of d^n system, Racah parameters, splitting of terms in weak and strong octahedral and tetrahedral fields. Correlation diagrams for d^n and d^{10-n} ions in octahedral and tetrahedral fields (qualitative approach), $d-d$ transition, selection rules for electronic transition-effect of spin orbit coupling and vibronic coupling Interpretation of electronic spectra of complexes- Orgel diagrams, Tanabe-Sugano diagrams, Calculation of Dq , B and $\hat{\alpha}$ (Nephelauxetic ratio) values, charge transfer spectra. Magnetic properties of complexes-paramagnetic and diamagnetic complexes, molar susceptibility, Gouy's method for the determination of magnetic moment of complexes, spin only magnetic moment. Temperature dependence of magnetism. Temperature Independent Paramagnetism (TIP). Spin state crossover, Antiferromagnetism-inter and intra molecular interaction. Application of magnetic measurements in the determination of structure of transition metal complexes.

Unit III Crystalline state

18 h

Crystal systems and lattice types. Bravais lattices. Crystal symmetry- Introduction to point groups and space groups. Miller indices. Reciprocal lattice concept. Close packed structures: BCC, FCC and HCP. Voids. Coordination number. Crystal binding: Molecular, covalent, metallic and hydrogen bonded crystals. X- Ray diffraction by crystals: Function of crystals. Transmission grating and reflection grating. Braggs equation. Diffraction methods: Powder and rotating crystal. Indexing and determination of lattice type and unit cell dimensions of cubic crystals. Crystal defects: Perfect and imperfect crystals. Point, line and plane defects. Thermodynamics of Schottky and Frenkel defects. Colour centers in alkali halide crystals. Defect clusters. Extended defects: Crystallographic shear structure and stacking faults. Dislocations and crystal structure. Structure of compounds of AX (Zinc blende, Wurtzite), AX_2 (Rutile, fluorite, antiferrofluorite), A_mX_2 (Nickel arsenide), ABX_3 (Perovskite, Ilmenite). Spinel. Inverse spinel structures.

Unit IV Lanthanides and actinides

18 h

Lanthanides: Characteristic properties. Electronic configurations and term symbols. Occurrence and extraction. Separation techniques. Oxidation states. Spectral and magnetic properties. Shapes of f orbital and their splitting in cubic ligand field. Lanthanide complexes as shift reagents. Actinides: Occurrence and general properties.

Extraction of thorium and uranium. Electronic configuration and term symbol. Oxidation states. Spectral and magnetic properties. Comparative properties of lanthanides and actinides. Trans-uranium elements and their stabilities. Applications of lanthanide and actinide compounds. Comprehensive study of the beach sands of Kerala and their important components such as monazite, ilmenite, zircon and sillimanite.

Unit V Solid state chemistry

18 h

Kronig-Penney model, Free electron theory, Zone theory and MO theory of solids. Energy bands-conductors and non-conductors, intrinsic and extrinsic semiconductors. Electrons and holes. Mobility of charge carriers. Hall Effect. Pyroelectricity, piezo electricity and ferro electricity. Conductivity of pure metals.

Solid state reactions-diffusion coefficient, mechanisms, vacancy diffusion, thermal decomposition of solid-Type I reactions, Type II reactions.

Phase transition in solids: classification of phase transitions-first and second order phase transitions, Martensitic transformations, order-disorder transitions and spinodal decomposition. Kinetics of phase transitions, sintering. Growing single crystals-crystal growth from solution, growth from melt and vapor deposition technique.

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6. L. V. Azaroff, Introduction to Solids, McGraw-Hill, 1960.
7. S. Cotton, Lanthanides and Actinides, Macmillan, 1991.
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12. A. Earnshaw, Introduction to Magnetochemistry, Academic Press, 1968.

19PCH22: ORGANIC CHEMISTRY- II

Total 90 h

Course Outcome:

CO1. Comprehend and Predict the role of temperature, solvents, and catalysts in organic reactions.

CO2. Elucidate reaction mechanisms using isotope effects.

CO3. Comprehend the orbital interactions and orbital symmetry correlations of various pericyclic reactions.

CO4. Write mechanism of organic reactions involving reactive intermediates and concerted processes

CO5. Apply these reactions in organic synthesis.

CO6. To familiarize with the important photochemical reactions in Organic Chemistry.

CO7. Understand the bio molecules and their role in metabolic activities.

Unit I Physical organic chemistry

18 h

Reactivity in relation to molecular structure and conformation. Steric effects. *F* strain. Ortho effect, Bond angle strain. Linear free energy relationships. The Hammett equation and its applications. Taft equation. Solvent polarity and parameters. *Y*, *Z* and *E* parameters and their applications. Primary and secondary kinetic isotope effects. Salt effects and special salt effects in S_N reactions. Kinetic and thermodynamic control of reactions. Energy profiles, Hammond postulate. Principle of microscopic reversibility. Marcus theory. Methods of determining reaction mechanisms. Phase transfer catalysis and its applications.

Unit II Molecular rearrangement and transformation reactions

18h

Types of organic rearrangements: Anionic, cationotropic, prototropic, free radical, carbene and nitrene intermediates. Mechanism with evidence of Wagner-Meerwein, Pinacol, Demjanov, Hofmann, Curtius, Schmidt, Lossen, Beckmann, Wolf, Fries, Fischer-Hepp, Hofmann-Martius, von-Richter, Orton, Bamberger,

Smiles, Dienone-phenol, Benzilic acid, Benzidine, Favorskii, Stevens, Wittig, Sommelet-Hauser, Bayer-Villiger rearrangements.

Unit III Aromaticity and symmetry controlled reactions

18 h

Aromaticity and antiaromaticity. Homo, hetero and non-benzenoid aromatic systems. Aromaticity of annulenes, mesoionic compounds, metallocenes, cyclic carbocations, carbanions.

Symmetry properties of MOs. Classification of pericyclic reactions. Mechanism and stereochemistry of electrocyclic, cycloaddition and sigmatropic reactions. Woodward-Hoffmann rules. FO, CD and Huckel-Mobius analysis of electrocyclic, cycloaddition and sigmatropic reactions. FO analysis of [1, j] and [3, 3] migrations. 1,3-dipolar cycloaddition. Stereo aspects of Diels-Alder reaction and Cope rearrangement. Intramolecular Diels-Alder, Retro Diels-Alder, Alder-ene, retro-ene and cheletropic reactions. Synthetic applications of Diels-Alder reactions. Fluxional molecules.

Unit IV Organic photochemistry

18 h

Photochemical processes. Energy transfer, sensitization and quenching. Singlet and triplet states and their reactivity. Photoreactions of carbonyl compounds, enes, dienes and arenes. Norrish Type I and Type II reactions of acyclic ketones. Free radical reactions: Paterno-Buchi and Barton reactions, photo-Fries and Di- δ methane rearrangements. Photoreactions of Vitamin D. Photosynthesis, photochemistry of vision. Singlet oxygen generation and their reactions. Introduction to chemiluminescence. Applications of photochemistry.

Unit V Chemistry of natural products and biomolecules

18 h

Introduction to primary and secondary metabolites in plants. Extraction methods of chemical constituents from plants, such as fractionation using solvents, specific extraction of alkaloids and supercritical fluid extraction. Characterizations of isolated compounds (terpenes, sterols, alkaloids, carbohydrates, flavonoids and polyphenols) by colour reactions and spray reagents. Biosynthesis of terpenes from mevalonic acid and sterols from squalene. Structure elucidation of ocimene, classification of pigments, structure elucidation of β -carotene. Structural differences between a triterpene and a sterol. Synthesis of quercetin, synthesis of testosterone, androsterone, estrone and progesterone. Determination of carbon skeleton of alkaloids (Hofmann, Emde and Von Braun degradation methods). Structural elucidation of ephedrine, nicotine, atropine.

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19PCH23: PHYSICAL CHEMISTRY –II

Total 90 h

Course Outcome:

CO1. Understand the theory and instrumentation of NMR, ESR, NQR, PES spectroscopic techniques.

CO2. Interpret the above spectroscopic data of unknown compounds. Use these spectroscopic techniques in their research.

CO3. To know the basic concepts in classical thermodynamics and to learn the thermodynamic aspects of various processes and reactions.

CO4. To understand the different aspects of statistical thermodynamics and its applications.

CO5. Write equations representing electrochemical cell, explain various overpotential involved during the operation of the cell.

CO6. Calculate electrochemical cell parameters, electrochemical active surface area, current and over potential under given condition, amount of corrosion and its rate.

CO7. Solve problems based on Debye-Huckel limiting law.

Unit I

Quantum Chemistry II

18 h

Rotational motion: Cartesian and spherical polar coordinates. The wave equation in spherical polar coordinates - particle on a ring, the ϕ equation and its solution, wave functions in the real form. Non-planar rigid rotor (or particle on a sphere)-separation of variables, the ϕ and the θ equations and their solutions, Legendre and associated Legendre equations, Legendre and associated Legendre polynomials. Spherical harmonics (imaginary and real forms)-polar diagrams of spherical harmonics.

Quantum Mechanics of Hydrogen-like systems-Potential energy of hydrogen-like systems. The wave equation in spherical polar coordinates: separation of variables-R, θ and ϕ equations and their solutions, wave functions and energies of hydrogen-like systems. Orbitals-radial functions, radial distribution functions, angular functions and their plots.

Unit II: Classical Thermodynamics

18 h

Entropy- dependence of entropy on variables of a system (S, T and V; S, T and P). Thermodynamic equations of state. Criteria for equilibrium and spontaneity, Euler's relation, Gibbs and Helmholtz free energy, Maxwell relations and significance, temperature dependence of free energy, Gibbs Helmholtz equation and its applications.

Partial molar quantities: chemical potential, Gibbs-Duhem equations, determination of partial molar properties—partial molar volume and partial molar enthalpy.

Fugacity- relation between fugacity and pressure, determination of fugacity of a real gas, variation of fugacity with temperature and pressure. Fugacity of liquid mixtures, fugacity of mixture of gases, Lewis Randall rule. Activity, activity coefficients, dependence of activity on temperature and pressure. Determination of activity and activity coefficients of electrolytes and non electrolytes

Thermodynamics of mixing, Duhem-Margules equation, Konowaloff's rule, Henry's law, excess thermodynamic functions- Determination of excess enthalpy and volume.

Chemical affinity and thermodynamic functions, effect of temperature and pressure on chemical equilibrium-van't Hoff reaction isochore and isotherm.

Unit III Spectroscopy II

18 h

Resonance spectroscopy: Nuclear Magnetic resonance Spectroscopy, Nuclear spin. Interaction between nuclear spin and applied magnetic field. Proton NMR . Population of energy levels. Nuclear resonance. Chemical shift. Relaxation methods. Spin-spin coupling. Fine structure. Elementary idea of 2D and 3D NMR. Introduction to instrumentation.

ESR spectroscopy: Electron spin. Interaction with magnetic field. Kramer's rule. The g factor. Determination of g values. Fine structure and hyperfine structure. Elementary idea of ENDOR and ELDOR.

Mossbauer spectroscopy: Basic principles. Doppler effect, chemical shift, recording of spectrum, application. Quadrupole effect.

NQR spectroscopy - Principle and application

Photoelectron spectroscopy. Introduction to UV photoelectron and X-ray photoelectron spectroscopy.

Unit IV Statistical Mechanics –I

18 h

Microstates. Concept of ensembles Canonical and Grand canonical ensemble. Classical distribution of particles-Maxwell Boltzmann distribution.

Bose-Einstein statistics, Bose-Einstein distribution. Thermodynamic probability, Bose Einstein distribution function. Examples of particles. Theory of Para magnetism. Bose Einstein condensation, Liquid Helium. Super cooled liquid.

Fermi- Dirac Statistics. Fermi- Dirac distribution, examples of particles Fermi-Dirac distribution function. Thermionic emission. Relation between Maxwell Boltzmann, Bose Einstein and Fermi -Dirac Statistics.

Unit V Electrochemistry I

18 h

Ionics- Ions in solution. Deviation from ideal behaviour. Ionic activity. Ion-solvent interaction. Born equation. Ion-ion interaction. Strong electrolytes Debye-Huckel theory of strong electrolytes, Onsager equation. Limitation of the model Conductance at high frequencies and high potentials –Wein effect—Activity coefficient and its determination.

Ionic strength, Debye-Huckel limiting law. Equation for appreciable concentration. Osmotic coefficient. Activities in concentrated solutions. Ion associations. Ion transport.

Electrodics: Different type of electrodes. Origin of electrode potential, Electrochemical cells, Concentration cells and activity coefficient determination. Liquid junction potential. evaluation of thermodynamic properties, the electrode double layer, Electrode-electrode interface. Theory of multilayer capacity. Electrocapillarity. Lippmann potential and membrane potential.

Electro kinetic phenomena. Mechanism of charge transfer at electrode- electrolyte interface. Electrolysis. Current- potential curve. Dissolution, deposition and decomposition potentials. Energy barriers at metal – electrolyte interface. Different types of overpotentials. Butler-Volmer equation. Tafel and Nernst equation. Rate determining step in electrode kinetics. The hydrogen and oxygen over voltage. Theories of overvoltage.

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MODEL QUESTION PAPER
19PCH21: INORGANIC CHEMISTRY-II

Time:3 h

Max marks:75

SECTION-A

Answer any two among (a), (b), and (c) from each question. Each sub-question carries 2 marks.

1. a) Classify the following as closo, nido, arachno or hypho.
1) B_5H_9 2) B_5H_{11} 3) B_6H_{12} 4) B_9H_{15} .
b) Which sulphur –nitrogen compound is known as ‘one-dimensional’ metal? Why is it called so?
c) Why are $P_4N_4C_{18}$ puckered and $P_4N_4F_8$ planar?
2. a) How would you distinguish between ferro magnetic and anti ferromagnetic material?
b) Explain non-crossing rule.
c) The effective magnetic moment of a complex is 4.90 BM. Calculate the no: of unpaired electron per unit complex.
3. a) What is the type of defect observed in AgBr crystals? Why?
b) Give one example each for molecular, covalent, metallic and H-bonded crystals.
c) Differentiate between H-centre and V-centre in NaCl crystals.
4. a) Write any two differences between 4f and 5f orbitals.
b) Give the term symbols for Eu^{3+} and Lu^{3+} .
c) Explain why Actinides have greater tendency for complex formation than lanthanides?
5. a) What is band gap?
b) What is meant by order-disorder transitions ?
c) Write a short note on sintering.

[2 x 10 = 20 marks]

SECTION-B

Answer either (a) or (b) of each question carries 5 marks.

6. a) How is diborane prepared? Discuss the structure and bonding in diborane.
b) Write a note on metallocarboranes.
7. a) Describe the Guoy’s method to determine magnetic susceptibility . How are these measurements used to calculate effective magnetic moments?
b) Even though d-d transitions are forbidden, why such transitions occur in many transition metal complexes? Illustrate with examples.
8. a) Distinguish between spinels and inverse spinels with suitable examples.
b) Write briefly on line and plane defects in solids.
9. a) Briefly discuss the basis of the ion-exchange method for the separation of Lanthanides.
b) Compare the spectral and magnetic properties of Lanthanides and Actinides.
10. a) With suitable examples explain the phenomenon of metal conductivity.
b) Write a short note on the applications of ferro, piezo and pyroelectrics.

[5x 5 = 25 marks]

SECTION-C

Answer any three questions and each question carries 10 marks.

11. How is Borazine prepared? Discuss its structure and compare the bonding with phosphazene molecule.
12. Write an account on the selection rules and characteristics of d-d transition and application of each electronic spectra in elucidating the structure of metal complexes.
13. Discuss briefly on the packing of atoms and ions in solids.
14. a) Correlate the oxidation states and ionic radii with electronic configuration of lanthanides.
b) Write a short note on the beach sands of kerala.
15. Discuss the salient features of band theory of solids and compare it with the free electron theory of solids.

[10x3 = 30 marks]

MODEL QUESTION PAPER
19PCH22: Organic Chemistry-II

Time-3 hours

Maximum marks :75

Section A

Answer any two among a), b) and c) from each question. Each sub-question carries, 2 marks.

1. a) Give the mechanism of rearrangement of aryl hydroxylamines to aminophenols.
b) Describe Stevens rearrangement.
c) Show the mechanism involved in the rearrangement of an unsubstituted amide to a primary amine.
2. a) Explain why cyclodecapentaene with 10 π electrons is not aromatic.
b) Write briefly on Homoaromaticity.
c) State Woodward Hoffmann rules.
3. a) Explain Phosphorescence.
b) Write a short note on Norrish type I reaction.
c) Describe anyone method of generation of singlet oxygen.
4. a) Explain von Braun reaction.
b) Show the products formed when Quercetin is treated with dimethyl sulphate followed by boiling with ethanolic KOH.
c) Draw the structure of Cholesterol.
5. a) Explain Taft equation
b) Describe salt effect in substitution reaction
c) State Marcus theory

[2x10=20 marks]

Section B

Answer either a) or b) of each question, and each question carries 5 marks.

6. a) Discuss the mechanism and applications of Baeyer Villiger reaction.
b) Discuss the mechanism of Benzidine rearrangement. Also write proof to support the mechanism.
7. a) Explain the acidity of Cyclopentadiene and Cycloheptatriene.
b) Explain briefly Claisen rearrangement.
8. a) Briefly explain Barton reaction.
b) Explain the photochemistry of olefins.
9. a) Discuss the structure elucidation of Carotene.
b) Briefly describe the biosynthesis of terpenes.
10. a) Explain kinetic and thermodynamic control in reactions involving ketones.
b) Explain the reason for the difficulty in the hydrolysis of 2, 6 - disubstituted benzoic acid esters.

[5x5=25 marks]

Section C

Answer any three questions and each question carries 10 marks.

11. i) Discuss the mechanism and applications of Beckmann rearrangement.
ii) Discuss the similarity in the intermediates of Curtius, Schmidt and Lossen rearrangements.
12. Explain briefly on
i) Sigmatropic reactions
ii) 1, 3 - Dipolar and Ene reactions.
13. Discuss the following:
i) Photochemistry of vision
ii) Photoreaction of Vitamin D.
14. Explain the following:
i) Structure of Estrone.
ii) Chemical, spectroscopic and chiroptical methods for establishing carbon skeleton. .
15. Discuss the following:
i) Principles and applications of phase transfer catalysis.
ii) Methods of determination of reaction mechanism.

[10x3=30 marks]

MODEL QUESTION PAPER
19PCH23: Physical Chemistry- II

Time : 3 Hours

Max mark : 75

Section A

Answer any two among (a), (b) and (c) from each question. Each sub-division carries 2 marks .

1. a) Set up the Schrodinger equation for a rigid rotator.
b) Give plots of (a) radial probability distribution functions of 2S orbital and (b) angular plot of $2P_x$ orbital.
c) Write the expression for Fock operator and explain the terms
2. a) Define chemical potential
b) State Lewis –Randall rule of fugacity
c) Write Konowaloff's rule
3. a) Write the expression for chemical shift in Mossbauer spectroscopy and explain the terms.
b) Calculate the ESR frequency of an unpaired electron in a magnetic field 0.33 Tesla. Given for free electron $g=2$, $\beta=9.273 \times 10^{-27}$ J/T
c) Explain the basic principle of X-ray photoelectron spectroscopy.
4. a) Derive ideal gas law from translational partition function.
b) Explain the term canonical ensemble.
c) Electron would never follow Maxwell Boltzmann statistics. Why?
5. a) What is Lippmann potential? How does it arise?
b) Calculate the mean activity coefficient of 0.01M $BaCl_2$ in water at 25 °C.
c) Explain the origin of concentration overpotential.

(10 X 2 = 20 Marks)

Section- B

Answer either (a) or (b) of each question and each question carries 5 marks

6. a) Explain self-consistent field method to solve many electron systems.
b) Write the Schrodinger equation for hydrogen atom in polar coordinates and separate the variables.
7. a) Define fugacity. Write the method for the determination of fugacity of a gas
b) Derive Gibbs Duhem equation
8. a) Write a brief account of 2D-NMR spectroscopy.
b) What is Kramer's degeneracy? Discuss.
9. a) Apply Fermi-Dirac statistics to understand paramagnetism in solids.
b) Derive the expression for partition function for particle executing (i) free linear motion and (ii) free linear harmonic vibration.
10. a) Derive Debye-Huckel limiting law.
b) Discuss the various models for electrical double layer.

(5 X 5 = 25 Marks)

Section-C

Answer any three questions and each question carries 10 marks

11. (i) Apply Schrodinger equation for particle in a ring. Find eigen values and eigen functions.
(ii) Show that any two associated Legendre functions satisfy orthonormality condition.
12. Write a brief account of the methods for the determination of activity coefficient of electrolytes and non electrolytes
13. Explain the principle and applications of NQR spectroscopy.
14. Derive the expression for the distribution function of a Boson and Bose-Einstein condensation.
15. (i) Derive Butler-Volmer equation. Deduce the expression for the low and high field limits of this equation.
(ii) Discuss the application of Debye-Huckel Onsagar equation as applied to strong electrolytes and point out its limitations.

(10 x 3 = 30Marks)

SEMESTER III

19PCH31: INORGANIC CHEMISTRY- III

Total 90 h

Course Outcome:

CO1. Identify the structure and bonding aspects of simple organometallic compounds.

CO2. Apply different electron counting rules to predict the shape/geometry of low and high nuclearity metal carbonyl clusters.

CO3. Identify the different types of organometallic reactions and apply the above concepts to explain different catalytic reactions.

CO4. To know the important reactions of coordination compounds.

CO5. To get an insight into the role of essential and trace elements in biological systems and understand the mechanism of photosynthesis, respiration etc.

CO6. Use various spectroscopic principles to characterize inorganic and organometallic compounds.

CO7. To impart essential theoretical knowledge on nuclear chemistry.

Unit I Organometallic compounds

18h

Nomenclature of organometallic compounds. Hapto nomenclature. 18 and 16 electron rule, isoelectronic and isolobal analogy. Types of metal complexes. Metal carbonyls, bonding in metal carbonyls. Bonding in metal nitrosyls and cyanides. Synthesis, structure and bonding of polynuclear carbonyls with and without bridging. Complexes with linear π donor ligands: Olefins, acetylenes, dienes and allyl complexes. Complexes with cyclic π donors: Cyclopentadiene, benzene complexes, structure and bonding of ferrocene and dibenzenechromium complexes (MO treatment). Oxidative addition and reductive elimination, insertion and elimination reactions
Catalysis by organometallic compounds:

Alkene hydrogenation using Wilkinson's catalyst, hydroformylation of olefins using cobalt catalyst and polymerization reaction by Ziegler-Natta catalyst. Fluxional molecules.

Unit II Coordination chemistry-III: Reactions of metal complexes

18 h

Energy profile of a reaction - Thermodynamic and kinetic stability, Stability of complex ions in aqueous solutions: Formation constants. Stepwise and overall formation constants. Factors affecting stability of complexes. Determination of stability constants: spectrophotometric, polarographic and potentiometric methods. Stability of chelates. Thermodynamic explanation, macrocyclic effects. Classification of ligand substitution reactions -kinetics and mechanism of ligand substitution reactions in square planar complexes, trans effect- theory and synthetic applications. Kinetics and mechanism of octahedral substitution- water exchange, dissociative mechanism, associative mechanism- Eigen-Wilkins mechanism, Eigen-Fuoss equation, base hydrolysis, racemisation and isomerisation reactions. Electron transfer reactions: Outer sphere mechanism- Marcus theory, inner sphere mechanism- Taube mechanism. Photochemical reactions- substitution and redox reactions of Cr(III), Ru(II), and Ru(III) complexes. Photo-isomerisation and photo-aquation reactions of metal complexes.

Unit III Bioinorganic chemistry

18 h

Essential and trace elements in biological systems, structure and functions of biological membranes, mechanism of ion transport across membranes, sodium-potassium pump. Photosynthesis, porphyrin ring system, chlorophyll, PS I and PS II. Synthetic model for photosynthesis. Role of calcium in biological systems. Oxygen carriers and oxygen transport proteins- haemoglobin and myoglobin. Non-haeme iron-sulphur proteins involved in electron transfer-ferredoxin and rubredoxin. Iron storage and transport in biological systems- ferritin and transferrin. Redox metalloenzymes-cytochromes, peroxidases and superoxide dismutase and catalases. Nonredox metalloenzymes- Carboxypeptidase A- structure and functions. Nitrogenases, biological nitrogen fixation. Vitamin B₁₂ and coenzymes. Toxic effects of metals (Cd, Hg, Cr and Pb).

Unit IV Spectroscopic Methods in Inorganic Chemistry

18 h

Infrared spectra of coordination compounds. Structural elucidation of coordination compounds containing the following molecules/ ions as ligands- NH₃, H₂O, CO, NO, OH⁻, SO₄²⁻, CN⁻, SCN⁻, NO₃⁻, NO₂⁻, CH₃COO⁻ and X⁻ (X= halogen). Changes in ligand vibration on coordination with metal ions.

Vibrational spectra of metal carbonyls- CD and ORD spectra of metal complexes.

ESR spectra : Application to Cu(II) complexes and inorganic free radicals such as PH_4 , F^- and $[\text{BH}_3]^-$. Nuclear Magnetic Resonance Spectroscopy: The contact and pseudocontact shifts, some applications including biological systems, an overview of NMR of metal nuclides with emphasis on P-31 and F-19 NMR. Mossbauer Spectroscopy : Application of the technique to the studies of iron and tin complexes.

Unit IV Nuclear chemistry

18 h

Fission products and fission yield. Neutron capture cross section and critical size. Nuclear fusion reactions and their applications. Chemical effects of nuclear transformations. Positron annihilation and autoradiography. Principles of counting technique such as G.M. counter, proportional, ionization and scintillation counters. Cloud chamber. Synthesis of transuranic elements such as Neptunium, Plutonium, Curium, Berkelium, Einsteinium, Mendeleevium, Nobelium, Lawrencium and elements with atomic numbers 104 to 109. Analytical applications of radioisotopes-radiometric titrations, kinetics of exchange reactions, measurement of physical constants including diffusion constants, Radioanalysis, Neutron Activation Analysis, Prompt Gama Neutron Activation Analysis and Neutron Absorptiometry. Applications of radio isotopes in industry, medicine, autoradiography, radiopharmacology, radiation safety precaution, nuclear waste disposal. Radiation chemistry of water and aqueous solutions. Measurement of radiation doses. Relevance of radiation chemistry in biology, organic compounds and radiation polymerization.

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19PCH32: ORGANIC CHEMISTRY-III

Total 90 h

Course Outcome:

- CO1. Apply NMR, IR, MS, UV-Vis spectroscopic techniques in solving structure of organic molecules and in determination of their stereochemistry.
- CO2. Interpret the above spectroscopic data of unknown compounds.
- CO3. Use these spectroscopic techniques in their research.
- CO4. To understand the concept of various organic synthetic reactions and different separation techniques and its use in research.

Unit I UV-Vis and IR Spectroscopy and Mass spectrometry

18h

Electronic transitions and analysis of UV spectra of enes, enones and arenes. Woodward-Fieser rules. Effect of solvent polarity on UV absorption. Principle of characteristic group frequency in IR. Identification of functional groups and other structural features by IR, Hydrogen bonding and IR bands. Sampling techniques. Mass spectrometry- EI, CI, FAB, Electrospray and MALDI ion sources. Magnetic, High resolution (Double focusing), TOF and quadrupole mass analysers. Characteristic EIMS fragmentation modes and MS rearrangements. Mass spectral fragmentation patterns of long chain alkanes, alkenes, alkynes, alcohols, ethers, thiols, aromatic compounds, aldehydes, ketones, acids, amides, nitro, amino and halo compounds.

Unit II NMR spectroscopy and structural elucidation

18h

Theory of NMR spectroscopy, chemical shifts, anisotropic effects and coupling constant. Spin-spin interactions in typical systems. First order and higher order spectra. Simplification methods of complex spectra by high field NMR, shift reagents, chemical exchange and double resonance. ^{13}C NMR chemical shifts. Applications of NOE, DEPT, and 2D techniques such as COSY, HSQC, HMQC and HMBC. Spectral interpretation and structural elucidation. Solving of structural problems on the basis of numerical and spectrum based data.

Unit III Organic synthesis

18 h

C-C and C=C bond forming reactions. Mannich, Reimer-Tiemann, Synthesis of small rings. Simmons-Smith, Vilsmeier-Haack, Ullmann and Chichibabin reactions. Ring formation by Dieckmann, Kostanecki, Thorpe, Pschorr and acyloin condensations. Stork enamine, Shapiro, Peterson, Heck, Stille, Ritter and Prilezhaev reactions. Reduction and oxidation in synthesis. Catalytic hydrogenation. Alkali metal reduction, Birch reduction, Clemmensen reduction and Wolff-Kishner reduction, Huang-Minlon modification. Boranes, LAH and sodiumborohydride as reductants. Hydrogenation, Oppenauer oxidation, Jones oxidation. Applications of HIO_4 , OsO_4 and mCPBA. Organo palladium catalysts -Heck, Negishi, Sonagashira and Suzuki coupling

Unit IV Methods in organic synthesis

18 h

Retrosynthetic analysis and disconnection approach. Synthetic strategy and synthons. Regioselectivity in enol and enamine alkylation. Stereospecific and stereoselective synthesis, Sharpless asymmetric epoxidation, Chiral pool, chiral auxiliary, Chiral reagents, BINAP, Mitsunobu reaction. 1,3-dipolar cycloaddition in the construction of rings. Olefin synthesis by extrusion reactions. Olefin metathesis – first and second generation Grubbs' catalysts. Umpolung concept, functional group equivalents. Reductive coupling reactions. Epoxide to alkene. Introduction to combinatorial synthesis. Electrochemical reduction of organohalogen, nitro and carbonyl compounds. Electrochemical Kolbe oxidation. Protecting group strategy: Tetrahydropyranyl, silyl, *t*-butyl, trichloroethyl, acetal and thioacetal as hydroxyl, thiol, carboxyl and carbonyl protecting groups in synthesis.

Unit V Reagents in organic synthesis

18h

Applications of hydrogenation catalysts, hindered boranes, bulky metal hydrides. NaCNBH_3 , DIBAL, Li trialkylborohydrides, tri-*n*-butyltin hydride, diimide, Lindlar catalysts and aluminium alkoxide. Rosenmund reduction and McFadayan-Stevens reaction. Oxidation using SeO_2 , lead tetraacetate, ozone, peracids, DDQ, manganese (IV) oxide, silver carbonate and Cr(VI) reagents. Swern oxidation, Moffatt oxidation, allylic and benzylic oxidation. Sommelet reaction. Elbs reaction. Oxidative coupling of phenols. Chemo and regioselectivity in reductions and oxidations.

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19PCH33: Physical Chemistry -III

90 h

Course Outcome:

- CO1. To understand the requirement of approximation methods in quantum mechanics.
- CO2. To gain the knowledge to apply important approximation methods to problems in quantum mechanics.
- CO3. To gain insight in to valance bond theory molecular orbital theory and the concept of hybridisation
- CO4. To get an exposure to the emerging world of computational chemistry
- CO5. To have a basic idea about computational chemistry calculations.
- CO6. Calculate change in thermodynamic properties, equilibrium constants, partial molar quantities, chemical potential.
- CO7. Identify factors affecting equilibrium constant.
- CO8. Apply phase rule and, draw phase diagrams for one, and two component systems, identify the dependency of temperature and pressure on phase transitions, and identify first/second order phase transitions.
- CO9. Solve problems based on partition functions.
- CO10. Calculate excess thermodynamic properties.
- CO11. To understand the principle and applications various electroanalytical and spectrophotometric methods.

Unit I Chemical Bonding

18 h

Approximate methods: method of Variation - Variation theorem and its proof. Linear variation functions. Secular equations and secular determinants. Perturbation method- Successive correction to an unperturbed problem. Detailed treatment of first order non-degenerate case only. Wave equation for multielectron systems. Hartree-Fock Self- Consistent Field (HF-SCF) method for atoms, Hartree-Fock equations (derivation not required) & the Fock operator.

Treatment of molecules- The Born- Oppenheimer approximation- LCAO-MO Theory- MO theory of H_2 and H_2^+ . MO treatment of other homo diatomic molecules Li_2 , Be_2 , B_2 , C_2 , O_2 and F_2 . MO treatment of hetero diatomic molecules LiH , CO , NO and HF . Spectroscopic term symbols for homo diatomic molecules.

Valance bond theory of diatomic molecules H_2 , O_2 and F_2 . Comparison of MO and VB theories, Quantum mechanical treatment of sp , sp^2 and sp^3 Hybridisation. HMO theory of conjugated systems. Bond order and charge density calculations, Free valance. Application of HMO method to ethylene, allyl system, butadiene and benzene.

Unit II Computational Chemistry

18 h

Introduction to computational chemistry: as a tool and its scope. Potential energy surface- stationary point, saddle point or transition state, local and global minima. Slater and Gaussian functions and its properties. **Basis sets** : minimal, double zeta, triple zeta basis sets, contracted basis sets, Pople's style basis sets and their nomenclature. Basis functions- Roothan's concept, Slater type orbitals (STO) and Gaussian type orbitals (GTO). Slater determinants

Quantum mechanical computational methods- Ab initio methods: Introduction to SCF. Wave functions for open shell state, RHF, ROHF and URHF. (no calculation). Electron correlation and introduction to post HF methods.

Semiempirical methods. Huckels and extended Huckel methods. Strengths and weaknesses. PPP, ZDO and CNDO approach. (Mentioning only).

Density functional theory- Hohenberg-Kohn theorems, Exchange correlational functional. (Only the basic principles and terms to be introduced).

Non-quantum mechanical computational methods-Molecular mechanics: Force fields, bond stretching, angle bending, torsional terms, non-bonded interactions, electrostatic interactions and the corresponding mathematical expressions. Names of some commonly used force fields.

Construction of Z-matrix for simple molecules. H_2O , H_2O_2 , H_2CO , CH_3CHO , NH_3 and CO_2 .

Unit III Applications of Thermodynamics

18 h

Thermodynamics of irreversible processes: Simple examples of irreversible processes. General theory of non equilibrium processes The phenomenological relations. Onsager reciprocal relation. Generalized equation for entropy production, Entropy production from heat flow, matter flow and current flow. Application of irreversible thermodynamics to diffusion. Thermal diffusion, Thermo osmosis and thermomolecular pressure difference. Electro kinetic effects, the Glansdorf- Pregogine equation. Far from equilibrium region. Principle of minimum entropy production, Le-Chatelier Bra(u)wn Principle.

Three component systems: Graphical representation. Three component liquid systems with one pair of partially miscible liquids. Influence of temperature. Systems with two pairs and three pairs of partially miscible liquids.

Solid- Liquid systems: Two salts and water systems- no chemical combination, double salt formation, one salt forms a hydrate, double salt forms hydrate, Isothermal evaporation.

Unit IV Statistical Mechanics II

18 h

Molecular partition functions. Translational (1D, 2D and 3D), vibrational, rotational and electronic partition functions. Total partition functions Langevin function and its use for the determination of dipole moment and molecular energies

Relation ship between Partition functions and thermodynamic properties, The principle of equipartition of energy. Chemical equilibrium. Law of mass action. Transformation of the equilibrium expressions. Statistical derivation. Thermodynamic probability and entropy.

Equipartition principle Quantum theory of heat capacity. Calculation of heat capacity of gases, limitation of the method. Heat capacity of solids. Dulong and Petit's law, Kopp's law, Classical theory and its limitation. The vibrational properties of solids. Einstein theory of heat capacity. The spectrum of normal modes. Limitations of Einstein's theory. The Debye theory, the electronic specific heat.

Unit V Electro Analytical and Spectrophotometric methods.

18 h

Potentiometric methods: Reference electrodes and indicator electrodes. The hydrogen, calomel, Ag-AgCl electrode. The glass electrode- its structure, performance and limitations. Measurement of pH. Potentiometric titrations- redox and precipitation titrations.

Electrogravimetry: Principle and method. Determination of Copper. Separation of metal ions.

Conductometry: Principle and method. Conductometric titrations.

Coulometry: Principle and method. Coulometric titrations.

Voltametry: principle and method of polarography, cyclic voltammetry, stripping voltammetry and amperometry.

Flame emission and atomic absorption spectrometry. Instrumentation for AAS. The flame characteristics. Atomiser used in spectroscopy. Hollow cathode lamp. Interference in AAS. Application of AAS.

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19PCH34: Inorganic Chemistry Practicals -II

Total-125 h

Course Outcome:

CO1. Enable the students to estimate the binary mixtures of metallic ions by volumetric and gravimetric methods

CO2. To acquire the skill to analyse some common alloys and ores.

1. Estimation of simple mixture of ions (involving quantitative separation) by volumetric and Gravimetric methods.
2. Analysis of typical alloys and ores
3. Ion exchange separation of binary mixtures.
4. Spectral Interpretation of metal complexes using IR, UV-Vis. spectral data. Supplementary information like metal estimation, CHN analysis, conductivity measurements and magnetic measurements to be provided to the students. Assessment is based on arriving at the structure of the complex and assignment of IR spectral bands.
5. Interpretation of TG and DTA curves of metal oxalate hydrates. Assessment is based on the identification of various stages.

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19PCH35: ORGANIC PRACTICALS-II

Total 125 h

Course Outcome:

CO1. To gain the skill to prepare organic compounds using greener protocols

CO2. Enable the students to prepare organic compounds via two step synthetic sequences • To know about enzyme/coenzyme catalysed reactions

A. Volumetric estimation of

1) Aniline 2) Phenol 3) glucose 4) Ascorbic acid 5) Aspirin

B. Colorimetric estimation

6) paracetamol with potassium ferricyanide

7) protein by biuret method

8) Ascorbic acid by folin-phenol reagent or phosphor tungstic acid methods

C. Spectral identification

9) UV, IR, ¹H NMR, ¹³C NMR, EI mass spectral identification of Organic compounds from a library of organic compounds (Each students have to record the spectral analysis of a minimum of 40 compounds)

D. Separations of mixtures by Paper Chromatography

10) Identification of amino acids

E. Single stage preparation of organic compounds by green chemistry

11) Preparation of *p*-bromoacetanilide using CAN

12) Radical coupling – 1,1-Bis-2-naphthol

13) Synthesis of dihydropyrimidinone

14) Synthesis of dibenzalacetone- with lithium hydroxide

15) Photoreduction of benzophenone to benzopinacol (not for end semester evaluation)

The board of examiners have to choose the combination of a volumetric estimation, a colorimetric estimation, a green synthesis OR paper chromatography and spectral analysis. The choice of experiments should change every year.

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19PCH36: PHYSICAL PRACTICALS –II

125 h

Course Outcome:

Enable the students to determine the various physical properties using simple instrumental methods like polarimetry, refractometry etc.

Conductometry

Determination of strength of strong and weak acids in a mixture

Determination of strength of a weak acid.

Determination of solubility product of a sparingly soluble salt (PbSO₄, BaSO₄ etc.)

Hydrolysis of NH₄Cl or CH₃COONa or aniline hydrochloride

Determination of order of reaction, rate constant and energy of activation for saponification of ethyl acetate

Precipitation titrations.

Determination of critical micellar concentration (CMC) of sodium lauryl sulphate from measurement of conductivities at different concentrations.

Equivalent conductance at infinite dilutions and verification of Kohlrausch's law.

Determination of Onsager constants.

Potentiometry

Determination of emf of Daniel cell.

Determination of the emf of various ZnSO_4 solutions and hence the concentration of unknown ZnSO_4 solution.

Determination of valency of mercurous ion.

Determination of temperature dependence of EMF of a cell

Determination of stoichiometry and formation constant of silver-ammonia complex.

Determination of activity and activity constant of electrolytes.

Determination of thermodynamic constants of reactions.

pH metric titrations.

Acid alkali titrations using Quinhydrone electrode.

Titration(double) involving redox reactions – Fe^{2+} Vs KMnO_4 , $\text{K}_2\text{Cr}_2\text{O}_7$, $(\text{NH}_4)_4\text{Ce}(\text{SO}_4)_4$ and KI Vs KMnO_4

Determination of strengths of halides in a mixture.

Determination of pH of buffer solutions and hence to calculate the E° of quinhydrone electrode

Spectrophotometry

Verification of Beer-Lambert's law.

Absorption spectra of conjugated dyes.

Determination of concentration of potassium dichromate and potassium permanganate in a mixture.

To study the complex formation between Fe^{3+} and salicylic acid.

Determination of pKa of an indicator.

Polarimetry

Measurement specific rotation of glucose.

Determination of specific rotation of sucrose

Determination of unknown concentration of glucose solution. and rate constant of its hydrolysis in presence of HCl

Polarography :

Determination of half wave potential $E_{1/2}$ and unknown concentration of Cd^{2+} ion.

Determination of concentrations of metal ions in a mixture.

Surface tension

Determination of surface tension of various liquids (water-ethanol, water-glycerol, water-sorbitol, nitrobenzene-toluene) by Stalagmometric method (drop number/ drop weight)

Determination of parachors of molecules and various groups.

Determination of concentration of a mixture.

Determination of surface tension and parachor of liquids using double capillary method.

Variation of surface tension with concentration. Unknown concentration of a mixture. Interfacial tension.

Determination of surface excess and area per molecule.

Viscosity: Viscosity of liquids and mixtures of liquids. Verification of Kendall's equation. Composition of unknown mixtures. Determination of molecular masses polymers by viscosity measurements.

Refractometry

Determination of molar refraction of pure liquids

Determination of concentration of KCl solution/glycerol solution

Determination of solubility of KCl in water.

Determination of molar refraction of solid KCl

Study the stoichiometry of potassium iodide-mercuric iodide complex.

Determination of concentration of KI solution.

Part B Computational chemistry experiments

Experiments illustrating the capabilities of modern open source/free computational chemistry packages in computing single point energy, geometry optimization, vibrational frequencies, population analysis,

conformational studies, IR and Raman spectra, transition state search, molecular orbitals, dipole moments etc. Geometry input using Z-matrix for simple systems, obtaining Cartesian coordinates from structure drawing programs like Chems sketch,

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MODEL QUESTION PAPER 19PCH31: INORGANIC CHEMISTRY- III

Time : 3 Hrs

Max. Marks: 75

SECTION A

Answer **two** among (a), (b) and (c) from **each** question carries **2** marks

1. a) Draw the structure of $Rh_4(CO)_2$.
b) How do sigma allyl complexes differ from pi allyl complexes?
c) What are fluxional molecules?
2. a) What are the factors affecting the stability of complexes?
b) Explain Macrocyclic effect?
c) What is anation reaction? Give an example.
3. a) Why electron transfer process in photosynthesis is called an uphill process?
b) Explain the mechanism of oxygen binding in haemocyanin.
c) What is $Na^+ - K^+$ pump? How does it function?
4. a) What is group frequency concept? Illustrate with example.
b) Why are solid samples used for recording Mossbauer spectra?
c) What is Doppler broadening? Explain with an example.
5. a) Explain binding energy.
b) What is compound nucleus? How is it formed?
c) Give a note on breeder reactors.

(2x10= 20 marks)

SECTION B

Answer either among (a) or (b) from **each** question carries **5** marks

6. a) What are metal carbonyls? Explain the structure and bonding in $Ni(CO)_4$
b) Discuss the mechanism of polymerization of Zeigler- Natta catalyst.
7. a) Explain Trans effect with suitable examples.
b) Give an account of photochemical reactions of complexes.
8. a) Discuss the role of calcium in blood clotting process.
b) Explain the structural features of haemoglobin.
9. a) How does IR spectroscopy help for the structural elucidation of complexes containing ammonia and water as ligands.
b) Explain CD and ORD spectra of complexes.

10. a) Distinguish between Transient and Secular equilibria.
 b) What is meant by half life period? How is it related to decay constant? The $t_{1/2}$ of a radio nuclide is 20 years. If a sample of this nuclide has an initial activity of 8000 disintegrations per minute today, what will be its activity after 80 years?

(5x5= 25 marks)

SECTION C

Answer **any three** questions. **Each** question carries **10** marks

11. Explain the bonding of ferrocene by MO Theory.
 12. Briefly explain outer sphere and inner sphere mechanism of electron transfer reactions
 13. i) Discuss the function of PS-I and PS-II in photosynthetic activity.
 ii) Outline the probable mechanistic pathways Nitrogenase activity in nitrogen fixation.
 14. i) Discuss the application of ESR spectroscopy to Cu(II) complexes.
 ii) Explain how Mossbauer spectroscopy helps to the studies of iron and tin complexes
 15. Discuss about different types of nuclear reactions with suitable examples.

(10x3= 30 marks)

MODEL QUESTION PAPER 19PCH32: Organic Chemistry-III

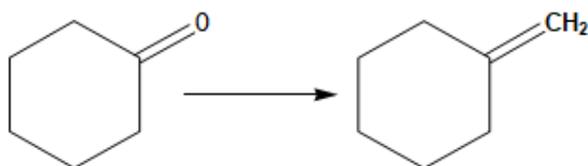
Time-3 hours

Maximum marks :75

Section A

Answer any two among (a), (b) and (c) from each question. Each sub question carries 2 marks

1. a) Explain how CH stretching vibrations of sp , sp^2 , sp^3 hybridised carbon varies.
 b) Polar solvents usually produces a red shift in the π to π^* transition. Explain.
 c) Explain how the presence of bromine atom in a molecule can be detected by mass spectrum.
 2. a) Explain why acetylenic hydrogens are more upfield than vinylic hydrogens.
 b) What multiplicities are observed for the signals of off resonance decoupled ^{13}C spectrum of 2-chloropropene?
 c) Account for the fact that splitting is observed between Hydrogens 'a' and 'b' in 2-methyl propene (CH_3^a $C=CH_2^b$) and not in neo pentylchloride ($(CH_3^a)_3CH_2^bCl$).
 3. a) Suggest a method for conversion



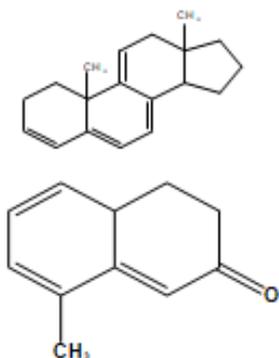
- b) What is Clemmenson reduction
 c) What are enamines? Write one synthetic application of enamines.
 4. a) Explain the use of silyl group as protecting group in organic synthesis.
 b) Explain the term combinatorial synthesis
 c) What is umpolung?
 5. a) Write two important reaction of $NaCNBH_3$
 b) Explain the importance of DDQ in organic synthesis.
 c) Explain Swern Oxidation

(2x10= 20 marks)

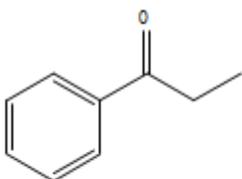
Section B

Answer either (a) or (b) from each question. Each sub question carries 5 marks

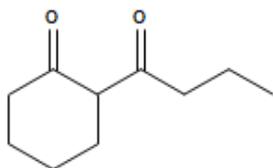
6. a) By using Woodward Fieser rules calculate the λ_{max} values



- b) How IR spectrum can be used to distinguish
 i) Primary amine from primary amide ii) Ethyl benzene from o-xylene
7. a) Deduce the identity of the compound $C_9H_{10}O$ that has an IR absorption at 1688 cm^{-1} and 1 H NMR signals at 1.22 (triplet, 3 H), 2.98 (quartet) and 7.28-7.95 (multiplet, 5H). Assign the data



- b) Explain the paramagnetic anisotropy of alkenic, aldehydic and aromatic protons.
8. a) Discuss Vilmeier-Hack reaction.
 b) Write short notes on important metal hydrides used as reducing agents in organic synthesis
9. a) Explain the regioselective synthesis of the following compound from cyclohexanone



- b) Discuss the synthesis of α -hydroxy ketones and aldehydes using umpolung
10. a) Write a note on oxidation using SeO_2
 b) Explain briefly the role of Lead tetra acetate in organic synthesis

(5x5= 25 marks)

Section C

Answer any three questions. Each question carries 10 marks

11. Discuss the functional group and finger print regions in the IR spectrum. How the IR spectrum is useful in distinguishing the inter and intra molecular hydrogen bonding
12. Write short notes on
 i) DEPT spectra, ii) Nuclear Overhauser effect, iii) Shift reagents in NMR, iv) Double resonance NMR
13. Explain the following named reactions with mechanism and example
 i) Mannich reactions ii) Robinson annulations reactions iii) Suzuki coupling
14. Explain olefin metathesis and Mitsunobu reaction with applications of each
15. Write a note on
 a) Sharpless asymmetric epoxidation
 b) Chemoselectivity in reduction reactions

(10x3= 30 marks)

MODEL QUESTION PAPER
19PCH33: Physical Chemistry III

Time : 3 Hours

Max. Marks : 75

Section A

Answer any two among (a), (b) and (c) from each question. Each sub question carries 2 marks

1. a) Arrange O_2 , O_2^+ , O_2^- in the increasing order of stability. Justify your answer
b) Write briefly about "Perturbation theory"
c) Explain the more dipole moment in ethyl chloride than in chlorobenzene.
2. a) Construct the z-matrix of CH_3CHO .
b) Name any two chemistry related software.
c) Write the determinantal wavefunction for the configuration $1s^2 2p_z^1$
3. a) Explain the terms 'Force' and 'Flux' with reference to irreversible thermodynamics.
b) Show the influence of temperature on the miscibility curve in a three component system forming a pair of partially miscible liquids.
c) What are the conditions under which linear relations are valid to understand irreversible processes?
4. a) Calculate the value of $\ln 6!$ with and without Stirling's theorem. Find the difference between the values if any. Comment on the result.
b) What is meant by the law of equipartition of energy?
c) State and explain Dulong Petit's law. Explain its limitations.
5. a) What are the requirements for choosing a reference electrode?
b) Define half wave potential. Explain its significance.
c) Why do we use three electrodes in cyclic voltametry.

(2x10= 20 marks)

Section B

Answer either (a) or (b) from each question. Each sub question carries 5 marks

6. a) Derive the expression for the bond angle and wave function in sp^2 hybridisation.
b) Apply HMO theory to butadiene molecule and discuss the molecular orbitals and their corresponding energy levels.
7. a) Differentiate between Slater type orbitals and Gaussian type orbitals
b) What is potential energy surface? Explain its significance.
8. a) Derive generalized equation for entropy production from heat flow.
b) Give the Onsagar-reciprocal relations. What are its applications?
9. a) Give comparison between Bose-Einstein, Maxwell-Boltzmann and Fermi-Dirac statistics.
b) Explain briefly how heat capacity of gases can be calculated?
10. a) Explain the working of glass electrode.
b) Discuss the advantages and disadvantages of amperometric titrations.

(5x5= 25 marks)

Section C

Answer any three questions. Each question carries 10 marks

11. Write a note on the secular equations.
12. Discuss Density functional theory and give its advantages and limitations.
13. (i) Draw the phase diagram of a three component liquid system with three pairs of partially miscible liquids. Explain.
(ii) How would you understand (a) thermo osmosis and (b) thermal diffusion from irreversible thermodynamics?
14. Derive Einstein's heat capacity equation for solid.
15. Describe the theory and instrumentation of AAS.

(10x3= 30 marks)

SEMESTER IV

19PCH41: Chemistry of Advanced Materials

Course Outcome:

CO1. To get a brief idea about emerging branches in chemistry like nanoscience and nanotechnology.

CO2. Understand the applications of various smart materials and conducting polymers in the field of medicine, defence and communication technology.

Unit 1: Introduction to nanoscience and nanotechnology

Underlying physical principles of nanotechnology: Nanostructured Materials: fundamental physicochemical principles - size dependence of the properties of nanostructured matter - quantum confinement, single electron charging, importance of nanoscale morphology. Top down and bottom up approaches to building materials. Metal nanoparticles: Size control of metal nanoparticles and their characterization, study of their properties, optical, electronic, magnetic. Surface plasmon band and its applications, role in catalysis, alloy nanoparticles, stabilization in sol, glass, and other media, change of band gap, blue shift, colour change in sol, glass, and composites.

Unit 2: The basic tools of nanotechnology

Scanning electron microscopy (SEM), TEM and EDAX analysis and X-ray diffraction, A brief historical overview of atomic force microscopy (AFM) and an introduction to its basic principles & applications. Optical microscope and their description, operational principle and application for analysis of nanomaterials, UV-VIS, IR spectrophotometers, Principle of operation and application for band gap measurement.

Carbon nano structures: Introduction. Fullerenes, C60, C80 and C240 nanostructures. Properties & applications (mechanical, optical and electrical). Functionalization of carbon nanotubes, reactivity of carbon nanotubes. Nanosensors: Temperature sensors, smoke sensors, Accelerometer, pressure sensor, night vision system, nano tweezers, nano-cutting tools, integration of sensor with actuators and electronic circuitry biosensors

Unit 3: Polymerization processes

Free radical addition polymerization - kinetics and mechanism. Chain transfer. Molecular weight distribution and molecular weight control. Cationic and anionic polymerization: Kinetics and mechanism. Living polymers. Step growth polymerization - Linear Vs cyclic polymerization. Other methods of polymerization - bulk, solution, melt, suspension, emulsion and dispersion techniques. Polymer stereochemistry: Configuration and conformation. Tacticity. Chiral polymers. Polymer characterization. Molecular weights - Methods for determining molecular weights - static, dynamic, viscometry, light scattering and GPC. Crystalline and amorphous states. glassy and rubbery States. Glass transition temperature and crystalline melting of polymers. Degree of crystallinity - X-ray diffraction. Thermal stability of polymers. Polymer solutions: Flory-Huggins theory. Chain dimension - chain stiffness. End-to-end chain distance of polymers. Conformation - random coil, solvation and swelling. Determination of degree of cross linking and molecular weight between cross links. Industrial polymers - synthesis, structure and applications of industrially important polymers. Specialty polymers: Polymers as aids in organic synthesis. Polymeric reagents, catalysts, substrates.

Unit 4: Conducting polymers

Synthesis & applications of polyacetylenes, polyanilines, polypyrroles & polythiophenes. Photoresponsive and photorefractive polymers. Polymers in optical lithography - Drug delivery - Drug carriers-Polymer based nanoparticles. Polymer based LEDs, lithium-polymer batteries, Liquid crystalline polymers- Main chain and side chain liquid crystalline polymers. Phase morphology.

Unit 5: Smart materials

Piezoelectric, magnetostrictive, halochromic, electrochromic, thermochromic, magneto caloric and hermoelectric materials. Chemistry behind photochromism in spiropyrans, spirooxazines, diarylethenes, azobenzenes, quinones. Examples for Photochromic Coordination Compounds. Shape-memory polymers, pH-sensitive polymers, Temperature-responsive polymers, dielectric elastomers, self-healing polymers and concept of mechanophores, polymorphism in polycaprolactone, introduction to ferrofluids, concept of pseudoelasticity.

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19PCH42(a): INORGANIC CHEMISTRY IV

Total 90 h

Course Outcome:

- CO1. To apply the Principles of group theory in predicting the hybrid orbitals involved in inorganic complexes.
CO2. To understand the concepts of supramolecular chemistry and its applications in catalysis.
CO3. To understand the importance of metals in the biological systems.
CO4. To apply non aqueous solvents for synthetic reactions.

Unit I Applications of group theory

18 h

Hybrid orbitals and molecular orbitals for simple molecules. Transformation properties of atomic orbitals. Hybridisation schemes for σ and π bonding with examples. MO theory for AB_n type molecules. Molecular orbitals for regular octahedral, tetrahedral and metal sandwich compounds. Ligand field theory: Splitting of d orbitals in different environments using group theoretical considerations. Construction of energy level diagrams. Correlation diagram. Method of descending symmetry. Tanabe-Sugano diagrams. Selection rules for electronic spectra. Molecular orbitals in octahedral complexes. Formation of symmetry adapted group orbitals of ligands. MO diagram. Symmetry and selection rules: Symmetry properties of common orbitals. Application of character tables to infrared and Raman spectroscopy. Infrared and Raman active modes for C_{2v} , C_{3v} and D_{4h}

Unit II Supramolecular Chemistry

18 h

Concepts and language. Molecular recognition: Molecular receptors for different types of molecules, design and synthesis of co receptors and multiple recognition. Strong, weak and very weak Hydrogen bonds. Utilisation of H-bonds to create supramolecular structures. Use of H bonds in crystal engineering and molecular recognition. Supramolecular reactivity and catalysis. Transport processes and carrier design. Supramolecular devices. Supramolecular photochemistry, supramolecular electronic, ionic and switching devices Some examples of self- assembly in supramolecular chemistry.

Unit III Metal-metal bonds and metal clusters

18 h

Metal-metal bonds: Factors affecting the formation of metal-metal bond. Dinuclear compounds of Re, Cu and Cr, metal-metal multiple bonding in $(Re_2X_8)^{2-}$, Trinuclear clusters, tetranuclear clusters, hexanuclear clusters. Polyatomic zintl anion and cations. Infinite metal chains. Metal carbonyl clusters. Anionic and hydrido clusters. LNCCs and HNCCs. Isoelectronic and isolobal relationships. Hetero atoms in metal clusters: Carbide

and nitride containing clusters. Electron counting schemes for HNCCs. Capping rule. Chalcogenide clusters. Chevrel phases.

Unit IV Selected topics in Bioinorganic Chemistry

18 h

Copper on biochemical systems. Oxidase activity, super oxide dismutase activity. Electron transport in biology. Structure and function of copper proteins in electron transport process. Oxygen transport copper proteins. Hemocyanin- copper transport, copper enzymes-Azurin, plastocyanin. Inorganic medicinal chemistry. Metals in medicine. Metal deficiency and diseases. Toxic effects of metals. Effect of deficiency and excess of essential metal ions. Toxicity due to non essential elements and speciation. Detoxification mechanism. Role of lithium and aluminium in biological systems. Chelation therapy and chemotherapy. Anticancer drugs and vanadium based diabetics drugs.

Unit V Acids and Bases and Non-aqueous Solvents

18 h

Acid base concept in non aqueous media-HSAB concept, solvent effects, linear free energy relationship-mechanism and methods of determination, super acids, Reactions in non-aqueous solvents. Ammonia - solutions of metals in liquid ammonia. Protic solvents: anhydrous sulfuric acid, hydrogen halides. Aprotic solvents: non-polar solvents, non-ionizable polar solvents, polar solvents undergoing autoionization, liquid halogens, inter halogen compounds, oxy halides, dinitrogen tetroxide, sulphur dioxide.

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19PCH42(b): ORGANIC CHEMISTRY-IV

Course Outcome:

CO1. To get a brief idea about emerging branches in chemistry like supramolecular chemistry, nanochemistry, medicinal chemistry, polymer chemistry and its applications

CO2. To learn the principles of green chemistry and to know the various green protocols in organic synthesis.

CO3. To study the chemistry of biopolymers and polymers

Unit I Organometallic chemistry

18 h

Preparation of organo Mg, Al, Li, Cu, Zn, Cr, Fe, Ce and sulphur stabilised compounds. Grignard reagents in organic synthesis. Alkylation, oxirane addition, carbon dioxide addition, carbonyl addition, enone addition (1,2- and 1,4-additions), reduction, conjugate addition and enolisation reactions. Selectivity in Grignard reactions. Reactions of organo Li reagents, Li exchange reaction, its use in the preparation of RLi compounds, addition to C=O, COOH and CONR₂, Li dialkylcuprates (Gilman reagent)-preparation and reaction with alkyl

halides, aryl halides and enones. Alkynyl Cu(1) reagents, Glaser coupling. Dialkyl cadmium compounds- preparation and reaction with acyl halides. Benzene tricarbonyl chromium- preparation and reaction with carbanions. Tebbe's reagent, Silane carbanion and its reactions.

Unit II Molecular recognition and supramolecular chemistry

18 h

Introduction to supramolecular chemistry. One-pot reactions. The concepts of molecular recognition, host, guest and receptor systems. Forces involved in molecular recognition. Hydrogen bonding, ionic bonding, δ -stacking, Van der Waals and hydrophobic interactions. Introduction to molecular receptors. Tweezers, cryptands and carcerands. Cycophanes, cyclodextrins and calixarenes- typical examples. Non-covalent interactions in biopolymer structure organization. Role of self-organization and self-association in living systems. Importance of molecular recognition in DNA and protein structure, their function and protein biosynthesis. Supramolecular systems like Organic zeolite, Clathrate hydrates of gases, Helicates, Nanotubes, liquid crystals, nanotechnology and other industrial applications of supramolecular chemistry.

Unit III Medicinal chemistry

18 h

Combinatorial organic synthesis, introduction, methodology, automation, solid supported and solution phase synthesis, study of targeted or focused libraries and small molecule libraries, Application- drug discovery. Drug design and development-Discovery of a drug, a lead compound. Development of drug-Pharmacophore identification, modification of structure, structure-activity relationship, structure modification to increase potency. The Hammett equation, Taft equation and lipophilicity. Computer assisted drug design. Receptors and drug action. Natural products and drug development. Different classes of drugs with examples. Synthesis of paracetamol, phenobarbital, diazepam, sulphamethoxazole, benzylpenicillin, chloramphenicol.

Unit IV Chemistry of biopolymers and polymers

18 h

Peptide bond formation methods. SPPS, Mechanism, Amino and carboxy protection in SPPS. Synthesis of tripeptides, A, G, C, T, U, adenosine, ADP and ATP. Automated polypeptide and oligonucleotide synthesis. Structure organization of proteins and polynucleotides. Protein sequencing by Edman's method. Protein denaturation. Structure of polysaccharides including starch, cellulose, glycogen and chitin. Synthesis of stereoregular polymers. Ziegler-Natta catalyst. Polymers in organic synthesis- supports, reagents and catalysts. Biodegradable polymers.

Unit V Green chemistry

18 h

Twelve principles of green chemistry. Green chemical strategies for sustainable development- Reaction mass balance, atom economy evaluation for chemical reaction efficiency, green solvents, reaction media- Synthesis under water, solventless, fluorous and ionic liquid media. Synthesis using scavenger resins, catalysis and biocatalysis. Green computation. Green processes- Microwave synthesis- fundamentals of microwave synthesis- Two Principal Mechanisms for Interaction with matter- The Microwave Effect with examples - Single-Mode and Multimode Microwave cavities. Microwave technology- Techniques and applications. Sonochemical synthesis. Applications of sonication in the synthesis of organic compounds.

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19PCH42(c): PHYSICAL CHEMISTRY IV

Total 90 h

Course Outcome:

- CO1. To understand the requirement of approximation methods in quantum mechanics.
 CO2. To gain the knowledge to apply important approximation methods to problems in quantum mechanics.
 CO3. To gain insight in to valance bond theory molecular orbital theory and the concept of hybridisation
 CO4. To get an exposure to the emerging world of computational chemistry
 CO5. To have a basic idea about computational chemistry calculations.

Unit I Applications of group theory

18 h

Spectroscopic applications: Transition moment integral transition moment operator. Vanishing matrix element. Symmetry selection rule for IR, Raman and electronic spectra. Dipole and polarizability transition moment operator. Identification of IR and Raman active normal modes in molecules coming under various point groups such as C_{2v} , C_{3v} , C_{4v} , D_{3h} , T_d and O_h . Mutual exclusion and complementarity principle of IR and Raman spectra and their use in the identification of molecular structures. Probability of overtone and combination bands. Identification of allowed and forbidden electronic transitions in carbonyl groups. Vibronic transitions. Application to MO theory: Symmetry adapted LCAO-MO theory of σ -bonded hydrocarbons. Projection operator and its use in the construction of wave functions of σ -molecular orbitals, secular equations and use of symmetry for simplifying the calculations of energy and wave functions of ethylene, butadiene and carbocyclic systems such as benzene and naphthalene.

Unit II Exactly solvable systems.

18 h

Simple Harmonic Oscillator: Wave equation for 1D harmonic oscillator. Complete solution and their properties. Three dimensional Harmonic oscillator. Potential energy in three dimension and Schrodinger wave equation in Cartesian coordinate. Separation of variables and solution of the equation for energy and wave function. Degeneracy.

Rigid rotor: Schrodinger equation in polar Coordinate. Angular momentum operator for rigid rotor. Separation of variables and complete solution for ϕ and θ equations. Legendre polynomials and associated Legendre functions. Normalisation of associated Legendre functions and evolution of the values of orbital angular momentum quantum number. Recursion relations. Rigid rotor wave function and energy.

The Hydrogen atom: Schrodinger wave equation in polar coordinate. Separation of variables and complete solution of the radial part. The associated Laguerre polynomial. Normalisation. The evolution of the value of the principal quantum number. The spherical harmonics and the radial part of the wave function. The total wave function of H atom. The wave functions of Hydrogen like atomic orbitals and explanations for the shapes of various orbitals.

Angular momentum, angular momentum operators (L_x , L_y , L_z and L^2) and their commutation properties. Spherical harmonics as eigen functions of angular momentum operator L_z and L^2 . Ladder operator method for angular momentum. Space quantisation.

Unit III Approximate method I

18 h

Schrodinger wave equation for He atom and anharmonic oscillator and difficulty to get the exact solution. The Variation method: Variation theorem and its proof. The variation integral and its properties. Variational

parameters. Trial wave functions, Illustration of trial wave functions for calculation of H atom and particle in a 1D box as examples.

Trial functions as linear combination of orthonormal functions, linear combinations of functions containing variational parameters as trial functions. Setting up of secular determinants. Variation methods of normal state of He . The SCF method, SCF and variation method. Strength and limitation of the method.

Unit IV Approximate method II

18 h

The perturbation method. The generalised perturbation method. The idea of successive correction to unperturbed systems. First order perturbation. Correction of wave function and energy . Theory of non- degenerate level perturbation. The normal Helium atom. The first order perturbation of the degenerate level. The hydrogen atom. Second order perturbation theory. Correction for wave function and energy. Stark effect.

Time dependant wave equation: Variation in the state of a system with time. Emission and absorption of radiation. The Einstein s transition probability and its calculation. Selection rules and intensity of spectrum for harmonic oscillator , rigid rotor and hydrogen atom.

Unit V Computational methods

18 h

Computational methods as potential tools for practicing chemistry. Potential energy surface, saddle point, local minima and global minima. Geometry optimisation. Exchange and overlap integrals. Difficulty in evaluating them with H-like wave functions. Slater Type functions(STO), approximation of STO with Gaussian type functions. Contracted Gaussians.

Basis sets: minimal basis set, split valance basis set, polarised basis set and diffused basis set. Model chemistry and notations.

Geometry input- in terms of Cartesian coordinates and internal coordinates. Z-matrix, construction of z-matrices of simple molecules H_2 , H_2O , H_2O_2 , H_2CO , CH_3CHO , CH_4 , C_2H_6 and with dummy atom, CO_2 , NH_3 , C_6H_6 .

Molecular mechanics method: Force fields, potential energy expressions for bond stretching, bending, torsion, non-bonded interactions, electrostatic interaction and H-bonding. Setting up of force field expressions. Method of parameterisation. Use of molecular mechanics. Brief introduction to commonly using force fields(MM3, MMFF, AMBER and CHARMM) and Softwares.

Ab-initio method: Hartree-Fock Self Consistent Field method. Slater determinant. Post Hartree-Fock methods- Configuration Interaction(CI) and Moller Plesset(MP)methods.

Semiempirical method: Basic principle of the method. Its variants, ZDO, CNDO and INDO. Density Functional method: Functional. Hohenberg-Kohn theorems. Kohn-Sham orbitals. Basic idea of Local Density(LD) approximation, Generalised Gradient approximation and hybrid (BLYP, B3LYP) methods.

Comparative study of Molecular Mechanics, Ab-initio method, Semi-empirical method and DFT method of computations.

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19PCH43(a): DISSERTATION

Each of the students has to carry out original research in a topic in accordance with the Elective paper chosen for Semester IV under the guidance and supervision of a teacher in the concerned Department of the College.

Instructions to Question Papers Setters

The Syllabus of each theory has five units. While setting the question papers, equal weight is to be given to each of the Units for choosing the questions. Each question paper is of 3 hours duration and has three Sections, namely Section A, Section B and Section C constituting a total 75 marks as detailed.

Section A Five questions, one from each Unit containing three short answer questions marked (a), (b), and (c), each of which has 2 marks. One has to answer any two of (a), (b) or (c) from each of the five questions. (2x10=20 marks)

Section B Five questions, one from each unit containing two short essay questions marked (a) and (b), each of which has marks. One has to answer either (a) or (b) from each of the five questions. (5x5=25 marks)

Section C Five essay questions, one from each unit having 10 marks. One has to answer any three questions from the five questions asked. (10x3=30 marks)

Structures should be in chemdraw or ISIS draw

MODEL QUESTION PAPER 19PCH41: Chemistry of Advanced Materials

Time: 3 h

Max. Marks: 75

Section A

(Answer any two among (a), (b) and (c) from each question. Each sub question carries 2 marks)

1. a) Define surface plasmon resonance?
b) What is meant by quantum confinement?
c) What are alloy nanoparticles?
2. a) Write the principle of EDAX?
b) What are fullerenes?
c) Write the significance of nano tweezers?
3. a) What do you mean by chain transfer in polymerization process?
b) What is meant by tacticity of a polymer?
c) Give a short note on GPC?
4. a) What are conducting polymers?
b) Name any two polymeric reagents.
c) What are photo responsive polymers?
5. a) Write the important characteristics of piezo electric materials?
b) What are halochromic materials?
c) Write examples of any two photochromic coordination compounds.

[2 x 10 = 20 marks]

Section B

(Answer either (a) or (b) of each question Each question carries 5 marks)

6. a) Explain the relation between size and properties of nano-materials.
b) Write a note on the Sol-Gel method for preparing nano particles.
7. a) Describe the use of powder XRD in determination of particle size of nano materials.
b) Give an account of the synthesis of C60 and C80.
8. a) Give a brief account of the kinetics of free radical addition polymerization.
b) Explain DSC method for determination of Glass transition temperature.
9. a) How will you synthesis of polyacetylenes?
b) Write in detail the synthesis of polythiophenes.
10. a) Explain the concept of pseudo elasticity.
b) Write a note on shape-memory polymers.

[5x 5 = 25 marks]

Section C

(Answer any three question and each question carries 10 marks)

11. (a) Describe the principle of the following techniques (i) SEM (ii) TEM.
(b) Write the applications of SEM and TEM in the characterization of nanomaterials.
12. Explain in detail the synthesis of nano materials using sonochemical and Hydrothermal methods
13. Explain determination of molecular weights by viscometry and light scattering methods.
14. Explain the application of Polymers in catalysis.
15. Write a note on the chemistry behind photochromism in spiropyrans, spirooxazines, diarylethenes and azobenzenes.

[10x3 = 30 marks]

MODEL QUESTION PAPER 19PCH42(a): INORGANIC CHEMISTRY-IV

Time:3 hrs

Max marks:75

SECTION-A

Answer any two among (a), (b), and (c) from each question. Each sub-question carries 2 marks.

1. a) What is the point group of NO_3^- ? How does its symmetry change when complexed as unidentate ligand?
b) Give the splitting of d-orbitals in a square pyramidal geometry?
c) Demonstrate that C_3 and C_3^2 belong to the same class in C_{3v} point group.
2. a) What is meant by macrocyclic effect?
b) Give any two examples of self-assembly in supramolecular chemistry.
c) What are the 3 basic functions of supramolecular species?
3. a) What are anionic clusters? Give examples.
b) Explain Wade's rules to calculate the no: of frame work electrons in crystals.
c) Clusters of $\text{Co}(\text{CO})_{12}$ and $\text{Co}_3\text{FeH}(\text{CO})_{12}$ are isolobal to each other. Explain.
4. a) What is cisplatin? What is its importance?
b) What is the role of Lithium in biological systems?
c) Write any two diseases caused by the deficiency of copper in the body.
5. a) Give examples of room-temperature molten salts that could be used as non-aqueous solvents?
b) What is meant by leveling effect of solvent?
c) Which species would act as a base in (1) BrF_2 and (2) liq SO_2 . Why?

[2 x 10 = 20 marks]

SECTION-B

Answer either (a) or (b) of each question carries 5 marks.

6. a) Give the salient features of Tanabe-Sugano diagram taking d^5 ion as an example.
b) Discuss the application of group theory in arriving at the selection rules for the electronic spectrum of metal complexes.
7. a) Explain the importance of supramolecular chemistry in the field of catalysis.
b) What are the advantages and applications of supramolecular chemistry in Nanotechnology?
8. a) Explain the concept of Chevrel phases.
b) What are LNCC's? Give eg: Discuss their structures.
9. a) The toxicity of metals have been variously correlated with their
(1) electronegativity, (2) insolubility of sulphides and (3) stability of chelates. Discuss.
b) What are dismutation reactions? Give eg: What type of metal is in superoxide dismutase?
10. a) What are the advantages and disadvantages of using aprotic solvents as non-aqueous solvents?
b) Write a note on the reactions in liquid HF.

[5x 5 = 25 marks]

SECTION-C

Answer any three questions and each question carries 10 marks.

11. Deduce the normal modes of $\text{trans-N}_2\text{F}_2$ molecule and predict the IR and Raman activity of the modes.
Given the character table for C_{2h} .
12. Discuss with suitable eg: the different types of interaction in supramolecular compounds.

13. Discuss and draw the structure and bonding of $[\text{Re}_2\text{Cl}_8]^{2-}$.
14. Explain the structure and functions of hemocyanin and plastocyanin.
15. Write a detailed account of the reactions in the following non-aqueous solvents:
(1) NH_3 and (2) N_2O_4 .

[10x3 = 30 marks]

MODEL QUESTION PAPER
19PCH42(b): Organic Chemistry-IV

Time-3 hours

Maximum marks :75

Section A

Answer any two among (a), (b) and (c) from each question. Each sub question carries 2 marks

1. a) Describe any one method for preparation of organo zinc reagents.
b) Describe the preparation of benzenetricarbonyl chromium.
c) Write short note on Tebbe reagent.
2. a) Explain host-guest system with an example
b) Explain the importance of Hydrogen bonding in molecular recognition
c) What are cryptands?
3. a) Draw the structure of benzyl penicillin and diazepam
b) What is in vitro and in vivo analysis of drugs?
c) What is meant by lead in drug analysis?
4. a) Write the examples of amino protecting groups .How it can be deprotected?
b) What is the difference between nucleotide and nucleoside?
c) How cellulose is different from chitin?
5. a)What is biodiesel?
b) What is meant by sonochemical synthesis? Give an application.
c) Write few examples of green solvents.

[2 x 10 = 20 marks]

Section B

Answer either (a) or (b) from each question. Each sub question carries 5 marks

6. a) What is Tebbe reagent? How it is prepared? Mention any two applications
b) Illustrate the uses of Grignard reagent in organic synthesis.
7. a) Write short note on a) Molecular tweezers, b) Calixarene c) cyclodextrins
b) Explain the structure and importance of cyclodextrins
8. a) Describe the synthesis-of paracetamol
b) Explain a) pharmacophore b) lead compound with examples
9. a) Explain the role of polymers in organic synthesis.
b) Write any two techniques used in protein sequencing
10. a) Explain atom economy with aldol condensation as example.
b) Write a note on sonochemistry in organic synthesis.

[5x 5 = 25 marks]

Section C

Answer ant three questions.Each question carries 10 marks

11. Write notes on the preparation and application of the following:
a) Dialkyl Cd compounds. b) Benzenetricarbonyl chromium.
12. Describe the importance of molecular recognition in DNA and protein structure.
13. Explain the structure-activity relation in the development of drugs.
14. Write an SPPS method for synthesis of Phe-ala-gly, explaining each step and advantages of SPPS
15. Explain the twelve basic principles of green chemistry.

[10x3 = 30 marks]

MODEL QUESTION PAPER
19PCH42(c): Physical Chemistry IV

Time: 3hrs

Max. Marks: 75

Section – A

Answer any two among a, b and c from each question. Each sub question carries 2 marks.

1. a) What is transition moment integral?
b) Write projection operator P_A for C_{2v} point group
c) Find the symmetry species of the asymmetric mode of vibration of H_2O molecule. C_{2v} character table is given.
2. a) Write down the Hamiltonian for Li^{2+} ion.
b) What are Hermite polynomials? Write first three.
c) What are spherical harmonics. Write two examples.
3. a) State and explain variation theorem.
b) Briefly explain SCF method.
c) What are variational parameters? Explain.
4. a) What do you mean by Slater determinant? Illustrate using He atom
b) What is Stark effect?
c) The potential energy of an anharmonic oscillator is $V(x) = \frac{kx^2}{2} + \frac{kx^3}{6}$. Find the first order correction to the energy by perturbation method
5. a) Explain the terms local minima and global minima.
b) What does the notation 3-21 G stand for?
c) How is a double zeta set obtained?

(2x 10 = 20 marks)

Section – B

Answer either a or b from each question. Each question carries 5 marks.

6. a) Find the IR and Raman active vibrations of NH_3 . C_{3v} character table is given.
b) From symmetry consideration account for the low intensity of n- σ^* transition in HCHO
7. a) Find the commutator of \hat{L}_x and \hat{L}_y
b) Set up the Schrodinger equation for H atom in spherical polar coordinate, separate the variables and find the phi solution.
8. a) State and prove variation theorem.
b) Write a short note on the Self Consistent Field method. Give its strengths and limitations.
9. a) Find the ground state energy of He by first order perturbation method.
b) What is Stark effect? Rationalize using first order perturbation method.
10. a) Write the Z-matrices of CH_4 and H_2O .
b) Outline the procedure used for performing ab initio calculations

(5 x 5 = 25 marks)

Section – C

Answer any three. Each question carries 10 marks.

11. Using group theoretical principles find π -molecular orbitals of butadiene and arrive at their energy expressions. (use C_{2v} character table)
12. Obtain the complete solution of the simple harmonic oscillator.
13. a) Explain briefly the basis of the Self-consistent field theory
b) Find the ground state energy of H atom by variation method using the trial function $f = e^{-ar}$, where a is the variational parameter
14. Discuss the perturbation method and set up expression for the first and second order corrections.
15. What are the wave function based approaches in computational chemistry? What are its advantages and disadvantages

(3x10 = 30 marks)

| C_{2v} | E | C_2 | σ_{xz} | σ_{yz} | | |
|----------|---|-------|---------------|---------------|-------|-----------------|
| A_1 | 1 | 1 | 1 | 1 | z | x^2, y^2, z^2 |
| A_2 | 1 | 1 | -1 | -1 | Rz | xy |
| B_1 | 1 | -1 | 1 | -1 | X, Ry | xz |
| B_2 | 1 | -1 | -1 | 1 | Y, Rx | yz |

| C_{3v} | E | $2C_3$ | $3\sigma_v$ | | |
|----------|---|--------|-------------|---------------|------------------------|
| A_1 | 1 | 1 | 1 | z | x^2+y^2, z^2 |
| A_2 | 1 | 1 | -1 | Rz | |
| E | 2 | -1 | 0 | (x,y) (Rx,Ry) | $(x^2-y^2, xy)(xz,yz)$ |