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Reg. No.:

St Aloysius College (Autonomous) Mangaluru

Semester IV - P.G. Examination - M.Sc. Analytical Chemistry
May - 2024

ORGANIC SYNTHETIC METHODS

Time: 3 Hours

Max. Marks: 70

PART - A

1. Answer any FIVE questions of the following:

(5x2=10)

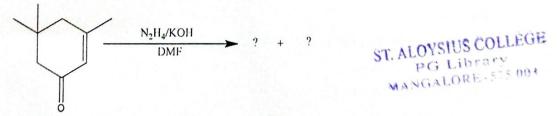
- a) How do transition metal catalysts participate in homogenous catalysis reactions?
- b) What is acyloin condensation? Give an example.
- c) Give Cutrius rearrangement.
- d) Explain the mechanism of the reaction of peracid with keto compound.
- e) Describe the structure of Penicillin -V.
- f) Write any two carbon-carbon single bond forming reactions.
- g) Perform the retrosynthetic analysis of phenacetin.
- h) Discuss any two basic principles in disconnection approach.

PART - B

Answer any <u>FIVE</u> of the following choosing at least one (5x12=60) full question from each unit:

UNIT - I

- 2.a) Explain the role of sodium borohydride as a reducing agent in organic (4) chemistry.
 - b) What is heterogenous catalysis? Discuss the catalysts used in it. (4)
 - c) Outline the bimolecular reduction of esters.
- 3.a) Predict the product with the mechanism for the following and also mention the name of the reaction:



- b) Compare and contrast the reactivity of sodium borohydride with lithium (4) aluminum hydride. Provide examples.
- c) Write a note on reduction using diimide.

UNIT - II

- 4.a) Describe the importance of Pinacol-Pinacolone rearrangement. (4)
 - b) Explain the mechanism of oxidation of aldehydes by SeO₂. (4)
 - c) Discuss the mechanism of Neber rearrangement. (4)

Contd...2

(4)

(4)

- 5.a) Explain Demaynov rearrangement. (4)
 - b) Propose a suitable mechanism for the oxidation of aromatic side chain by KMnO₄.
 - c) Outline the application of HIO₄ in organic synthesis. (4)

UNIT - III

- 6.a) Explain the mechanism of 6-Methoxy-1-tetralone. (4)
 - b) Write a note on Thorpe condensation. (4)
 - c) Explain the mechanism of Robinson annulations. (4)
- 7.a) Describe 1,3-dipolar cycloaddition reaction with mechanism. (4)
 - b) Write a note on Suzuki coupling reaction. (4)
 - c) Explain the mechanism of Arndt-Eistert reaction with mechanism. (4)

UNIT - IV

8.a) Provide all the possible disconnections and suggest a synthetic scheme (4) for the following molecule. Describe the rationale of your chosen synthetic method.

- b) Suggest any two reagents for the protection of carbonyl compounds.
 (4)
 Write the protection and deprotection reactions.
- c) What is FGI? What is its importance in synthesis? Explain with examples. (4)
- 9.a) Give one group and two group C-C disconnections with suitable examples. (4)
 - b) Suggest retrosynthetic schemes for chlorobenzide and 2,4-dichloro phenoxy acetic acid.
 - c) Explain the conversion:

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Semester IV - P.G. Examination - M.Sc. Analytical Chemistry

May - 2004

SPECTROSCOPIC METHODS OF ANALYSIS

Time: 3 Hours

Max. Marks: 70

PART - A

Answer any <u>FIVE</u> questions of the following:

(5x2=10)

- a) What is Auger effect?
- b) What are the factors affecting g-values in ESR spectroscopy?
- c) What are two key factors that influence the precision of AAS measurements for trace element analysis?
- d) Discuss the significance of the detector in atomic absorption spectroscopy.
- e) Differentiate between steady-state and time-resolved fluorimetry techniques in one sentence.
- f) What is a typical application of phosphorimetric analysis in forensic science?
- g) How are nephelometry and fluorimetry different different?
- Explain the application optical rotation method in the determination of rate constant.

PART - B

Answer any <u>FIVE</u> of the following choosing at least one (5x12=60) full question from each unit:

UNIT - I

2.a) Explain the principle behind Auger Spectroscopy.

(4)

- b) Examine the importance of hyperfine couplings in electron paramagnetic (4) resonance (EPR) spectroscopy, and illustrate their significance through relevant examples.
- c) Explain the shifts in energies due to chemical forces in photoelectron (4) spectroscopy.
- Describe the theory behind the NQR spectroscopy and mention any two (4) applications of NQR spectroscopy.
 - b) Describe the methodology employed by Mössbauer spectroscopy to (4) differentiate between Fe⁺² and Fe⁺³ compounds. Discuss the characteristic features of their Mössbauer spectra and how they relate to oxidation states.
 - c) Discuss the following aspects of Mössbauer Spectroscopy:

(4)

- i) Chemical Isomer Shift
- ii) Mössbauer Nuclide.

UNIT - II

| 4.a) | Analyze the influence of flame background in atomic absorption spectroscopy, considering its sources, effects on measurement accuracy, and methods to mitigate interference. | (4) | | | | |
|------|--|-----|--|--|--|--|
| b) | | | | | | |
| | operation and applications. | | | | | |
| c) | Write a note on ICP spectroscopy. | | | | | |
| 5.a) | What are the advantages and limitations of using premix burners in atomic | (4) | | | | |
| | absorption spectroscopy compared to total consumption burners? | | | | | |
| b) | Write a note on applications of AAS. | | | | | |
| c) | Briefly describe the instrumentation of flame photometry. | | | | | |
| | UNIT - III | | | | | |
| 6.a) | Evaluate the role of concentration on fluorescence and phosphorescence | (4) | | | | |
| | intensity. Discuss how changes in concentration can lead to quenching. | | | | | |
| b) | Compare and contrast filter fluorimeters and spectrofluorimeters, detailing | (4) | | | | |
| | their instrumentation, operational principles, and applications. | | | | | |
| c) | Write a note on instruments used in phosphorimetry. | (4) | | | | |
| 7.a) | Describe the types of condensed systems studied using photoacoustic | (4) | | | | |
| | spectroscopy (PAS), including liquids, solids, and colloidal suspensions. | | | | | |
| b) | Explain the concept of intersystem crossing and its role in determining the | (4) | | | | |
| | quantum yield of phosphorescence. Discuss how the rate of intersystem | | | | | |
| | crossing influences the population of triplet states and subsequent | | | | | |
| | phosphorescent emission. | | | | | |
| c) | Explain the principle of chemiluminescence. | (4) | | | | |
| | UNIT - IV | | | | | |
| 8.a) | Explain the instrumentation of single crystal XRD? | (4) | | | | |
| b) | Give an account of interaction of x-ray with matter. | (4) | | | | |
| c) | Explain the effects of concentration and particle size of suspension on | (4) | | | | |
| | nephelometry and turbidemitry. | | | | | |
| 9.a) | Describe the principle and applications of X-ray absorption. | (4) | | | | |
| b) | How do you relate scattering of light with turbidimetry | | | | | |
| c) | Write a note on Octant rule by taking a suitable example. | (4) | | | | |
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Semester IV - P.G. Examination - M.Sc. Analytical Chemistry

May - 2024

CHEMISTRY OF POLYMERS AND NATURAL PRODUCTS

Max. Marks: 70 Time: 3 Hours

PART - A

- (5x2=10)1. Answer any FIVE questions of the following:
- a) What is the tensile strength of a polymer?
- b) What are linear and branched polymers?
- c) Write any two differences between glass transition temperature and melting point temperature.
- d) Write the principle of end group analysis.
- e) How is glycolysis important in the biogenesis of natural products?
- f) What are the various groups of opium derived alkaloids? Give any examples to each group
- g) Define pericyclic reactions.
- ST. ALOYSIUS COLLEGE h) What is an ene reaction? How is it similar to Diels-Alder reaction?

PART - B

Answer any <u>FIVE</u> of the following choosing at least one (5x12=60)full question from each unit:

UNIT - I

- 2.a) Discuss Gradient elution technique with a schematic representation. (4)
 - b) Write the mechanism of free radical polymerization. (4)
 - c) Give the comparative account of step growth and chain growth (4)polymerisation.
- 3.a) Discuss Polydispersity and molecular weight distribution in polymers. (4)
 - b) With a neat diagram, compare crystalline and amorphous polymers. (4)
 - c) Describe emulsion polymerization technique. (4)

UNIT - II

- (4) 4.a) Explain calendaring and film casting techniques.
 - b) Describe sedimentation method of determination of molecular weight (4)of polymer
 - c) Elucidate the use of DSC technique in polymer characterisation. (4)
- (4) 5.a) Discuss the blow moulding method of polymer processing.
 - (4)b) Discuss the injection moulding method of polymer processing.
 - c) Discuss the technique of reinforcement in polymers with an example. (4)

Contd...2

UNIT - III

- 6.a) Illustrate the total synthesis of Menthol. What are the important reagents and reactions involved?
 - b) Explain the biogenesis and precursors of alkaloids. (4)
 - c) What are the steps involved in the total synthesis of Morphine? (4)
- 7.a) What are the various methods used in the structural elucidation of abietic acid?
 - b) Outline the total synthesis of a-Pinene. (4)
 - c) Illustrate the total synthesis of Papaverine. (4)

UNIT - IV

- 8.a) Describe the suprafacial and antarafacial sigmatropic rearrangement (4) of hydrogen and carbon moeity.
 - b) How FMO analysis can be used to study Diels-Alder reaction? (4)
 - c) With suitable rearrangement reaction, predict the most favourable product of allyl vinyl ether. (4)
- 9.a) Detail Cope rearrangement. Explain the regio and stereo specificity involved.
 - b) Predict the products and explain. (4)

c) Predict the products for these reactions and explain the salient (4) features of Ketene reactions.

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Semester IV - P.G. Examination - M.Sc. Analytical Chemistry

May - 2024

APPLIED ANALYSIS AND AUTOMATION

Time: 3 Hours Max. Marks: 70

PART - A

- Answer any <u>SEVEN</u> questions of the following: (7x2=14)
- Define reaction stoichiometry and provide an example illustrating its significance in chemical reactions.
- b) Explain the concept of a rate law and its significance in understanding the kinetics of chemical reactions.
- c) Mention two differences between automated and automatic systems.
- d) Write the significance of segmented flow analysis.
- e) Explain the difference between quality control and quality assurance in the context of pharmaceutical production.
- f) Explain why quality control is crucial at each stage of the manufacturing process, from raw materials to finished products.
- g) Describe the process of multicomponent analysis in kinetic methods.
- h) List the minerals present in milk.
- i) Describe the role of quality control in petrochemical industries.

PART - B

Answer any <u>FOUR</u> of the following choosing at least one (4x14=56) full question from each unit:

UNIT - I

- 2.a) Discuss in detail potentiometric and spectrophotometric methods to (4) determine rate of a reaction.
 - b) How do you determine sGPT in human samples? (5)
 - c) Derive Michaelis-Menten equation and give the rates at $K_M=[S]$ and (5) $K_M>>>[S]$.
- 3.a) Explain the half-life and isolation methods to determine order of a (4) reaction.
 - b) Discuss about enzyme specificity and effects of pH and Temperature (5) etc.
 - c) Explain Logarithmic Extra Polation method. (5)

UNIT - II

| | | 141 |
|------|---|-----|
| 4.a) | Discuss the principle of automated glucose analyser. | (4) |
| b) | at a of determination of moisture and ash content in food | (5) |
| c) | the working principle of centrifugal fast scan analyser. | (5) |
| 5.a) | Discuss the identification of arsenic in the biological sample. | (4) |
| b) | | (5) |
| c) | Write the source of cyanide poisoning and its symptoms. | (5) |
| | UNIT - III | |
| 6.a) | Describe the key elements of a product specification in the context of | (4) |
| b) | Design a quality control plan for a pharmaceuted beauty manufacturing tablets. Include specific tests and parameters to be monitored during different stages of production, from raw material | (5) |
| c) | evaluation to finished product testing Provide an overview of acts such as the Drugs and Cosmetics Act, 1940, and the Pharmacy Act, 1948, highlighting their key provisions and objectives. | (5) |
| 7.a) | Discuss various methods of sampling in case of pharma industries. | (4) |
| b) | Compare and contrast ISO 14001 and ISO/IEC 17025 with ISO 9001 in terms of their focus and application. | (5) |
| c) | Discuss quality control in raw materials, in process and finished product | (5) |
| | in pharma industries. | |
