Exam. Code : 107406 Subject Code : 1882

B.Sc. (Biotechnology) 6th Semester **BT-7 : PHYSICAL, ORGANIC AND INORGANIC** ASPECTS OF SPECTROSCOPY-B

Time Allowed—3 Hours] [Maximum Marks—40

Note :- Attempt all questions of Section A and it is compulsory. Do any five questions from Section B and do any two questions from Section C.

SECTION-A

(Compulsory, do all questions)

- ¹H NMR spectrum of compound A was recorded in 1. CDCl, (99%) deuterated. It showed a signal at 7.26 ppm which is not due to compound A. What is the origin of the signal?
- 2. Suppose 'H NMR spectrum of methanol (CH₂OH) is recorded at a concentration of 10⁻³ M and OH proton showed a guartet at 4.5 ppm. In which direction this quartet shall move : to low field or high field, if concentration is changed to 10⁻⁴ M ? Give suitable explanation.
- Suppose you are recording ¹H NMR spectrum of 3. compound B using 100 MHz NMR probe. What is Larmor frequency of the spinning proton for resonance? Give explanation.

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- What type of magnets are used in FT NMR spectrometer?
 Is FT NMR a single scan technique or a multiscan technique ?
- 5. From the mass spectrum of benzene discuss major ions you can identify. 1
- 6. What is Nitrogen rule as used in mass spectrometry?
- Methyl bromide showed two molecular ions which nearly same intensity. Write the molecular ions : (⁷⁹Br 51% abundance; ⁸¹Br 49% abundance).
- What is the most important information which we can get from the mass spectrum of a compound ? Explain with a suitable example.

SECTION-B

(Do any five questions)

9. The ¹H NMR spectrum of benzyl acetate (A) showed three signals at 2.3, 5.4 and 7.5 ppm. Assign these signals to the respective type of protons of compound. 4



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10. Predict ¹H NMR spectrum of nitropropane. Draw the spectrum and explain occurrence of various peaks. Mark chemical shift and spin-spin coupling constants.

O₂N-CH₂-CH₂-CH₃

R

- How protons of benzene appear at low magnetic field in its proton NMR spectrum ? Explain the role of the ring current in this observation.
- 12. Suppose ¹H NMR spectrum of compound A (formula C_2H_6O) showed three signals : one triplet at $\delta = 2.5$ ppm; one quartet of doublets at $\delta = 3.5$ ppm and another triplet at $\delta = 5.5$ ppm. Suggest structure of compound with suitable justification.
- 13. Both n-butane and isobutane showed most intense peak at m/z = 43. Identify the species and also explain its high intensity.
- Illustrate with an example, Diels-Alder fragmentation encountered in mass spectrometry of organic compounds.
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- 15. What is the importance of metastable ions in mass spectrometry ? Illustrate using a suitable example. 4
- 16. Describe mass spectrum of butyrophenone (C₁₀H₁₂O; MW 148). Illustrate Mc Lafferty rearrangement shown, if any, by this compound.

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SECTION-C

(Do any two questions)

- 17. How the fragments of mass spectra of aliphatic and aromatic compounds differ ? Explain using suitable examples. 6
- 18. Explain the major mass spectral fragments and their intensity of the following compounds : cyclopentane (C_5H_{10}) ; toluene $(C_6H_5-CH_3)$ and ortho-xylene $\{C_6H_4(CH_3)_2\}$. 6
- The C-H protons of aliphatic compounds show proton NMR signals in the high field region, while those of aromatic rings show signals in the low field region. Explain the phenomenon/factors which govern this difference. Explain using suitable examples.
- Describe in detail the basic principles of NMR spectroscopy with special reference to : NMR transition, solvents needed, chemical shift, nuclear spin-nuclear spin coupling constant, radiations used and advantages in respect of structure elucidation.

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