

## MANIPAL UNIVERSITY

Reg No					

## DEPARTMENT OF SCIENCES I SEMESTER M.Sc (CHEMISTRY) END SEMESTER EXAMINATIONS, NOV/DEC 2016

SUBJECT: ORGANIC SPECTROSCOPY [CHM 607]

## REVISED CREDIT SYSTEM

Time: 3 Hours Date: 30/11/2016 MAX. MARKS: 50

## **Instructions to Candidates:**

Answer ANY FIVE FULL questions.

Write chemical equations wherever necessary.

- **1A.** Discuss the moment of inertia equations involved in spherical top and asymmetric top molecules in Microwave Spectroscopy. Comment on their microwave activity.
- **1B.** Distinguish symmetry element from symmetry operation. Describe all the symmetry operations that can be carried on BF<sub>3</sub> and NH<sub>3</sub> molecules.
- **1C.** Describe the basic principles of FTIR instrument. Write any four advantages of FTIR instrument over the dispersive instruments.

2+4+4

- **2A.** Give reason for the following:
  - i) The lifetime of a tungsten-halogen lamp is more than double that of an ordinary tungsten lamp.
  - ii) Stokes lines are more intense than anti-stokes lines in Raman spectrum.
- **2B.** i) Differentiate between the following;
  - a) IR and Raman Spectroscopy
  - **b)** Rigid and non-rigid Microwave Spectra of a diatomic molecule
  - **ii**) Pure rotational (microwave) spectrum of the gaseous molecule CO consist of a series of equally spaced lines separated by 3.7978 cm<sup>-1</sup>. Calculate the inter nuclear distance and the first excited rotational level energy of the molecule. The molar masses are  $^{12}$ C = 1.99 x  $10^{-26}$  kg and  $^{16}$ O = 2.66 x  $10^{-26}$  kg.
- **2C.** State and deduce Beer-Lambert's law. Discuss the instrumental deviations from Beer's law.

2+4+4

- **3A.** Discuss the various types of electronic transitions involved in organic molecules in UV Spectroscopy with suitable examples.
- **3B.** Compute the  $\lambda_{max}$  for the following compounds based on the Woodward Fieser rules for diene.

$$_{i)}$$
  $_{H_3}$ C  $_{OCH_3}$ 

ii)



- **3C.** i) Calculate the Raman shift in cm<sup>-1</sup> and the wavelength in which anti-Stokes line will appear in the Raman spectrum if a sample was excited by the 430 nm line of mercury and a Raman line was observed at 460 nm.
  - ii) Give reasons for the following;
  - a) Inter and intra hydrogen bonding can be distinguished by IR spectral data.
  - **b)** Photo cell is a better detector in UV region compare to photo voltaic cell

2+4+4

- **4A.** Describe the various energy levels possessed by organic molecules and write the interaction of Microwave and IR regions of the electromagnetic spectrum with these molecules.
- **4B.** Identify the point group of water molecule and prove that the set of symmetry operations of this molecule forms an Abelian group.
- **4C.** Explain the procedure for the quantitative determination of calcium present in a sample of water through Atomic Absorption Spectroscopy.

2+4+4

- **5A.** Calculate the approximate wave number of the fundamental absorption peak due to the stretching vibrations of a carbonyl group. The force constant for a double bond has an approximate value of 1 x  $10^6$  dynes/cm. The masses of carbon and oxygen atoms are  $1 \times 10^{-26}$  and  $2.6 \times 10^{-26}$  kg per atom, respectively.
- **5B.** Write a note on Nebulization of the liquid sample carried out in Atomic Absorption Spectroscopy. Describe the functions of hollow cathode lamp, chopper and monochromator.
- **5C.** i) What is a point group? Mention the special point groups and dihedral point groups.
  - ii) Identify the point groups for the following molecules.
  - a) CCl<sub>3</sub>F
- **b**) Ethylene
- c) Ni(CO)<sub>4</sub>

2+4+4

- **6A.** Write the advantages of Atomic Absorption Spectroscopy over Flame Photometry. Mention two applications of Atomic Absorption Spectroscopy.
- **6B** Discuss in detail the flame and non-flame atomizers used for atomizing the liquid sample in Atomic Absorption Spectroscopy.
- **6C.** Explain the factors responsible for the width and intensity of spectral lines.

2+4+4

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