# 2. Answer *any five* questions :

 $2\frac{1}{2} \times 5$ 

- a) What do you mean by ORD?
- b) Discuss the nature of CD spectra of (-) Menthone in different solvents.
- c) How we can isolate different isomers of tris(s-alaninato) cobalt(III) complex ?
- d) Considering one specific example explain how position of a functional group of a molecule can be determined using CD.
- e) What is LASER Raman spectroscopy?
- f) What are the advantages of Raman spectroscopy over Infrared spectroscopy?

# M. Sc. CHEMISTRY EXAMINATION, 2017

(3rd Semester)

#### INORGANIC CHEMISTRY SPECIAL

## PAPER - XII - I

Time: Two hours

Full Marks: 50

(25 marks for each unit)

Use a separate answerscript for each group.

### UNIT - I - 3121 - a

- 1. Starting from Bloch equation derive the required relation between FID signal and  $T_1$  that can be applied in determining longitudinal relaxation time  $(T_1)$ . Describe in detail the Inversion Recovery (IR) method. 2+2
- 2. a) How could you identify all the possible isomers of the compound  $SnF_4(base)_2$  from their NMR spectra?
  - b) Describe in detail the <sup>19</sup>F NMR spectrum of XeF<sub>4</sub>.

3+2

- 3. Explain why  $J_{13C-H}$  coupling constants of  $-CH_2$  moiety in  $(C_6H_5)_2CH_2$  and  $(CH_3)_2CH_2$  differ from their corresponding carbonium ions.
- 4. Comment on *any one* of the followings:
  - a)  $^{19}$ F NMR of HPF<sub>2</sub>.
  - b)  $^{31}$ P NMR of  $P_4S_3$ .

 $1\frac{1}{2}$ 

[ Turn over

#### [3]

#### UNIT - I - 3121b

- 5. "While PFCl<sub>4</sub> retains its molecular symmetry in crystalline state PPhCl<sub>4</sub> does not." Explain on the basis of NQR spectra.  $2\frac{1}{2}$
- 6. Complexes of composition FeX<sub>2</sub> (pyridine)<sub>2</sub>, may be monomeric with four coordination or polymeric with six coordination in case of Fe. From the data given below for X = Cl and I, giving proper reason, deduce which one is polymeric.

<u>Complex</u>	<u>IS/mms<sup>-1</sup></u>	QS/mms <sup>-1</sup>	
$FeCl_2(Py)_2$	1.21	1.25	
$FeI_2(py)_2$	0.86	1.33	3

- 7. Explain why the complex  $[Co(NH_3)_6]^{3+}$  shows NQR resonance for <sup>14</sup>N but not for <sup>59</sup>Co. While both isotopes have  $I \ge 1$ .
- Deduce the required relation between quadrupole coupling constant and radiofrequency applied to observe NQR spectrum of CH<sub>3</sub>D molecule.
- 9. Why Mössbauer spectrum of <sup>57</sup>Fe often appears as doublet?

### **UNIT - 3122**

## Answer the following questions

- a) What makes a molecule Raman active? Using classical theory, explain the occurrence of Stokes and anti-stokes Raman scattering.
  - b) The equilibrium vibration frequency of the iodine molecule (I<sub>2</sub>) is 215 cm<sup>-1</sup>, and the anharmonicity constant (x) is 0.003; what, at 300K, is the intensity of the 'hot band' (v = 1  $\rightarrow$  v = 2 transition) relative to that of the fundamental (v = 0  $\rightarrow$  v = 1)?
  - c) How will you prove the occurrence of linkage isomerism in  $[Ru(dmso)_6]^{2+}$  (dmso = dimethylsulfoxide) with the help of IR spectroscopy?
  - d) Taking  $v_{CO}$  as a probe, how will you monitor the oxidative addition reaction in Vaska compound?
  - e) The symmetrical stretching mode of CO<sub>2</sub> is Infrared inactive but Raman active. Explain.
  - f) Justify the infrared stretching frequencies observed for the following isolectronic species:

 $[Mo(CO)_6]^+$  : 2090 cm<sup>-1</sup>

 $[Cr(CO)_6] : 2000 \text{ cm}^{-1}$ 

 $[V(CO)_6]^-$  :  $1858 \text{ cm}^{-1}$   $[3+2+2+2+\frac{1}{2}+2]$ 

[ Turn over