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Reg. No.....

FOURTH SEMESTER M.Sc. DEGREE EXAMINATION, JUNE 2016

(CUCSS)



Physics

PHY 4C 12 - ATOMIC AND MOLECULAR SPECTROSCOPY

(2012 Admission onwards)

Time: Three Hours

Maximum: 36 Weightage

Section A

Answer all questions.

Each question carries 1 weightage.

- 1. Write a note on space quantisation.
- 2. Explain Hund's rule with examples.
- 3. What are the salient features of rotational spectra?
- 4. Outline the effect of isotopic substitution on the rotational spectra of a molecule.
- 5. Diatomic molecule do not show vibrational spectra. Justify the statement.
- 6. State the conditions to be satisfied for a vibration to be Raman active considering the normal vibrations of CO₂ molecules as example.
- 7. Explain inverse Raman scattering.
- 8. Write a note on Franck-Condon principle.
- 9. Explain Fortrat diagram.
- 10. What is chemical shift? Explain it with an example.
- 11. Deduce the condition for NMR.
- 12. With the help of a block diagram, explain Mossbauer spectrometer.

 $(12 \times 1 = 12 \text{ weightage})$

Section B

Answer any two questions.

Each question carries 6 weightage.

- 13. (a) Derive an expression for Lande's splitting factor and explain the anamalous Zeeman effect of sodium doublet lines D_1 and D_2 with its help.
 - (b) State rules of Zeeman components.
- 14. Discuss in detail the construction and working of a Microwave spectrometer.

Turn over

- 15. Describe with necessary theory how structure of diatomic and triatomic molecules can be determined by combining Raman and Infrared spectroscopy.
- 16. Explain the different relaxation processes from nuclei and derive Block equations.

 $(2 \times 6 = 12 \text{ weightage})$

Section C

Answer any four questions.

Each question carries 3 weightage.

- 17. In an atom obeying 2-S coupling the components of a normal triplet state have separations 20 cm⁻¹ and 40cm⁻¹ between adjacent components. There is a higher state for which the separations are 22 cm⁻¹ and 33 cm⁻¹ respectively. Determine the terms for the two states and show the allowed transitions on an energy level diagram.
- 18. In the intra-red spectrum of HCl molecule the first line falls at 20.8 cm⁻¹. Calculate the moment of inertia reduced mass and the bond length of molecule.
- 19. A Raman line is observed at 4768.5 Å when acetylene was irradiated by 4358.3 Å radiations. Calculate the equilibrium vibrational frequency that causes their shift.
- 20. The rotational lines of a band system of electronic vibration spectra is given by $v = (24,762 + 25 \text{ m} 2.1 \text{ m}^2) \text{ cm}^{-1} \text{ m} = \pm 1, \pm 2, \dots$ Deduce the values of B, B and the position of the band head. Comment on the internaction distance of the two states and on the degradation of the band system.
- 21. The band origin of a transition in C_2 is observed at 19378 cm⁻¹, while the rotational fine structure indicates that the rotational constants in excited and ground states are, respectively, $B^1 = 1.7527 \text{ cm}^{-1}$ and $B^{11} = 1.6326 \text{ cm}^{-1}$. Estimate the position of the band head. Which state has the larger internuclear distance?
- 22. Consider the ESR spectrum of the free radicle CH_3 observed in a magnetic field of 0.34 T (i) If $\mathbf{g} = 2.0023$ for free electron, find the frequency at which resonance is obtained; (ii) How many hyperfine components are observes? (iii) Represent the transitions in an energy level diagram. Given $\mu_B = 9.274 \times 10^{-24} \, \mathrm{JT}^{-1}$.

 $(4 \times 3 = 12 \text{ weightage})$