M 27248

Reg. No. :	
Name :	

II Semester M.A./M.Sc./M.Com. Degree (Reg./Sup./Imp.)
Examination, March 2015
(2014 Admn. Onwards)
PHYSICS

PHY2C06: Quantum Mechanics - I

Time: 3 Hours

Max. Marks: 60

SECTION - A

Answer both questions (Either a or b). Each question carries 12 marks.

 a) Discuss the problem of similarity transformation. Prove that the matrix representing a similarity transformation is a unitary matrix. Show that a Hermitian operator remains Hermitian under a unitary transformation.

OR

- b) Distinguish between Heisenberg and Schrodinger pictures. Show that the state vectors and operators are the same in both the pictures at t=0.
- a) Discuss the problem of conservation of angular momentum as a consequence of the rotational invariance of the system.

OR

b) Outline the Schrodinger perturbation theory for non degenerate levels and apply it to explain first order Stark effect in hydrogen. (2×12=24)

SECTION-B

Answer any four. Each question carries 9 marks.

(1 mark for Part a, 3 marks for Part b and five marks for Part c)

- 1. a) Define linear vector space.
 - (b) Explain the properties of linear vector space.
 - c) Show that commuting operators posses simultaneous eigen functions.

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- 2. a) Outline Dirac's bra and ket rotation.
 - b) Explain the properties of ket and bra space.
 - c) Prove that the two eigen vectors of a Hermitian operator belonging to different eigen values are orthogonal.
- 3. a) State the uncertainty principle.
 - b) Prove that the simultaneous measurement of potential and kinetic energies is not possible.
 - c) The wave function of a particle in a state is $\psi = N \exp \left[\frac{-x^2}{2\alpha} \right]$ where

$$N = \left(\frac{1}{\pi\alpha}\right)^{1/4}$$
. Evaluate $(\Delta P \Delta x)$.

- 4. a) Outline the interaction picture.
 - b) Obtain the equation of motion for the state vector in the interaction picture.
 - c) Derive the equation of motion for operator in the interaction picture.
- 5. a) What do you mean by spin of an electron?
 - b) Explain spin up and spin down states. What are spinors?
 - c) Using Pauli's spin matrix reduce each of the operators :
 - i) Sx2 Sy Sz2
 - ii) Sx² Sy² Sz².
- 6. a) What is symmetry transformation?
 - b) Prove that a symmetry transformation conserves probabilities.
 - c) Prove that the parity operator is Hermitian and unitary.

 $(4 \times 9 = 36)$

新加州 阿斯田安國 Reg. No. : Name :

II Semester M.A./M.Sc./M.Com. Degree (Reg./Sup./Imp.) Examination, March 2015 (2014 Admn. Onwards) PHYSICS

PHY2C09: Spectroscopy

Time: 3 Hours

Max Marks: 60

SECTION-A

Answer both questions (Either a or b).

1 a) Distinguish between Normal Zeeman effect and Anomalous Zeeman effect. Discuss the theory of Normal Zeeman effect and prove that the frequency shift is the same for all Normal Zeeman effect lines.

- b) What is Stark effect? Describe the importance of the study of quadrapole hyperfine interaction in microwave spectra. Explain the different types of information obtained from rotational spectra.
- 2. a) Describe the construction and working of a Raman spectrometer. Explain the importance of Raman effect for phase transition studies.

b) Explain Frank-Condon principle Explain band origin and band head in the (2×12=24) rotational fine structure of electronic vibration spectra.

SECTION - B

Answer any four. (1 mark for part a, 3 marks for part b, 5 marks for part c).

- 1. a) What is fine structure of spectral lines?
 - b) Explain singlet and triplet states with examples.
 - c) The wavelength of the first line of the Lyman series of hydrogen is 1215A°. Calculate the wavelength of the second line of the series and the series limit.

P.T.O.

- a) What are linear molecules ? Give example.
- b) Explain how rotational energy transitions take place
- c) Discuss the rotational spectra of rigid molecules. Draw the rotational energy levels and transitions for a rigid diatomic molecule.
- a) What is a diatomic vibrating rotator?
- b) In the vibration rotation spectrum of HBr, why is it that the rotational lines at the high frequency end of the R branch are closely spaced and those at the low frequency end of the P-branch widely spaced.
- c) The fundamental band for HCL is centered at 2886 cm⁻¹. Assuming that the internuclear distance is 1276 A*. Calculate the wave number of the first two lines of each of the P and R branches of HCL.
- a) What is NMR?
- b) Explain how NMR frequency is related to the external magnetic field applied
- c) Find the energy difference between the spin up and spin down states of a proton in a magnetic field of B = 1.00 Testa. Estimate the Larmour frequency of the proton in the field (g = 5.586 and μ_N = 5.051x 10⁻²⁷ J/T.
- a) Give the principle of ESA.
- b) What are the factors responsible for the hyperfine structure in ESR spectra? Explain.
- c) Calculate the resonance frequency of a free electron in a magnetic field of 1.69 Testa.
- a) What is a symmetric top molecule?
- b) Derive an expression for the rotational constant of a diatomic molecule.
- c) Calculate the rotational constant and bond length of carbon monoxide if the first line in the rotation spectrum of GO has a frequency of 3.8424 cm. (4x9)