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Internal Assessment Test 1 – Sept. 2017

Sub:	internal Assessment Test 1 – Sept. 2017			1		
	: Engineering Physics Sub Code: 15	SPHY12	Branch:	All		
Date:	: 21/09/2017 Duration: 90 mins Max Marks: 50 Sem / Sec:	,L,M,N,O	Л,N,O		BE	
	Note: Value of Constants: $h = 6.625 \times 10^{-34} Js$ $k = 1.38 \times 10^{-23} J/K$ $m = 9.11 \times 1$ $e = 1.6 \times 10^{-19} C$ , $c = 3 \times 10^{8} m/s$	10 <sup>-31</sup> kg.	MA	ARKS	СО	RBT
1 (a)	State and explain Planck's law of black body radiation. Show that it reduces to Rayleigh-Jeans law under suitable wavelength conditions.	Wien's law	and [	[07]	CO1	L3
(b)				03]	CO1	L2
2 (a)	Define phase velocity and group velocity. Derive an expression for group velocity superposition of two travelling waves.	y on the bas	is of [	06]	CO1	L2
(b)	If the kinetic energy of the electron is 3 eV, calculate de-Broglie wavelength and ph the de-Broglie wave associated with the electron.	hase velocity	of [	04]	CO1	L2
3 (a)	Using Heisenberg's uncertainty principle, show that an electron cannot exist inside	e the nucleus	s. [	07]	CO1	L3
(b)	An electron has a speed of 500 m/s measured with an accuracy of 0.003%. With what can locate the position of the electron?	at accuracy	one [	03]	CO1	L2
. —						
4 (a)	a) Derive the time independent Schrodinger wave equation for a free particle in one d	limension.		[06]	CO1	<u> </u>
4 (a) (b)			value	[06] [04]	CO1	
	b) An electron is bound in a one dimensional potential well of width 0.15 nm. Find (in eV) and the de-Broglie wavelength of the electron in the second excited state.	I the energy				l L
(b) 5 (a)	<ul> <li>b) An electron is bound in a one dimensional potential well of width 0.15 nm. Find (in eV) and the de-Broglie wavelength of the electron in the second excited state.</li> <li>a) Derive the expression for energy Eigen value and Eigen function for a particle in a</li> </ul>	I the energy	ional	[04]	CO1	l L
(b) 5 (a) (b)	<ul> <li>b) An electron is bound in a one dimensional potential well of width 0.15 nm. Find (in eV) and the de-Broglie wavelength of the electron in the second excited state.</li> <li>a) Derive the expression for energy Eigen value and Eigen function for a particle in a potential well of infinite height.</li> <li>b) Calculate the wavelength of the X-ray photon scattered by an electron at an</li> </ul>	I the energy	ional	[04] [07]	CO1	L L
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(b) 5 (a) (b) 6 (a)	<ul> <li>b) An electron is bound in a one dimensional potential well of width 0.15 nm. Find (in eV) and the de-Broglie wavelength of the electron in the second excited state.</li> <li>a) Derive the expression for energy Eigen value and Eigen function for a particle in a potential well of infinite height.</li> <li>b) Calculate the wavelength of the X-ray photon scattered by an electron at an incident photon has a wavelength of 1Å.</li> <li>a) Explain the failures of classical free electron theory.</li> <li>b) Discuss the dependence of Fermi factor on temperature at T = 0 K and T &gt; 0 K.</li> </ul>	the energy one dimens angle 90°,	ional if the	[04] [07] [03] [06]	CO1 CO1	l L L L L L L L L L L L L L L L L L L L

# SCHEME OF EVALUATION 2017-18 ODDSEMESTER IAT-1

## 1.a.(7)

Max Planck developed a structural model for black body radiation that leads to a theoretical equation for the wavelength distribution that is in complete agreement with the experimental results at all wavelengths.

According to his theory

- a black body is imagined to be consisting of large number of electrical oscillators.
- an oscillator emits or absorbs energy in discrete units. It can emit or absorb
  energy by making a transition from one quantum state to another in the
  form of discrete energy packets known as photons whose energy is an
  integral multiple of hv where h is the planks constant and v is
  the frequency.
- 3. Energy emitted per unit volume per unit energy range is given by the product of number of modes of vibration in the given energy range and the energy per mode. The Energy density per unit wavelength range per unit volume is given by

$$E_{\lambda}d\lambda = \frac{8\pi\hbar c}{\lambda^{5}} \left[ \frac{1}{e^{\left[\frac{\hbar\gamma}{kT}\right]} - 1} \right] d\lambda$$
 (3)

Where h is Planck's constant, c is velocity of light, T is absolute temperature,  $\lambda$  is the wavelength and k is Boltzmann constant

#### Deduction of Weins law: (2)

It is applicable at smaller wavelengths.

For smaller wavelengths  $e^{rac{h\gamma}{kT}} >> 1$ 

$$\therefore e^{\frac{h\gamma}{kT}} >> 1 = e^{\frac{h\gamma}{kT}}$$

So Planck's radiation law becomes

$$E_{\lambda}d\lambda = \frac{8\pi hc}{\lambda^{5}} \left[ \frac{1}{e^{\left[\frac{h\gamma}{kT}\right]}} \right]$$

# Deduction of Rayleigh Jeans Law: (2)

It is applicable at longer wavelengths.

For longer wavelengths  $\ \, \frac{h\gamma}{kT} \ \, << 1$ 

$$\therefore e^{\frac{h\gamma}{kT}} = 1 + \frac{h\gamma}{kT} + \left(\frac{h\gamma}{kT}\right)^2 \frac{1}{2|} + \dots = 1 + \frac{h\gamma}{kT}$$

$$E_{\lambda}d\lambda=\frac{8\pi hc}{\lambda^{5}}.\frac{1}{1+\frac{h\gamma}{kT}-1}d\lambda=\frac{8\pi kT}{\lambda^{4}}d\lambda$$

## 1. b. (3)

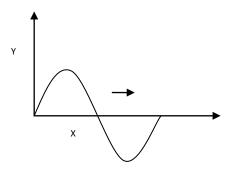
## From Weins displacement law

$$\lambda_{max} . T = 2.89 \times 10^{-3} \, \text{mK}$$

Put T=311K, 
$$\lambda_{\rm max}$$
. =  $9318.3nm$ 

## 2.a. (6)

<u>Phase velocity( $V_p$ ):</u> It is the speed with which an isolated pulse / constant phase propagates in a medium. (1)



A single pulse is shown in this diagram .It is represented as

$$Y = A \sin [wt - kx]$$

where Y is the displacement of a particle at a distance 'x' from the origin at a time 't', A is the amplitude , w is the angular velocity and k is the wave number.

Imagine two points A and B at X<sub>1</sub> and X<sub>2</sub> at same phase on the wave. Then

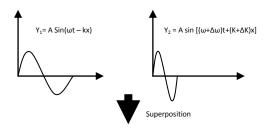
$$(\mathsf{wt}_1\text{-}\mathsf{kX}_1)=(\mathsf{wt}_2-\mathsf{kX}_2)$$

$$\frac{\mathbf{w}}{\mathbf{k}} = \frac{\mathbf{x}_1 - \mathbf{x}_2}{\mathbf{t}_1 - \mathbf{t}_2}$$

$$\therefore V_p = \frac{W}{k}$$

<u>Group Velocity( $V_g$ ):</u> It is the velocity with which the resultant envelope of varying amplitude formed by the superposition of two or more waves propagates. (1)

$$V_g = Lt_{dk \to 0} \frac{dw}{dk}$$





Resultant Envelope with varying amplitude

Let the waves be represented as

$$Y_1 = A \sin(wt - kx)$$
 ..... (1

$$Y_2 = A \sin [(w+\Delta w) t - (k + \Delta k) x]$$

The susultant superposition is  $Y = Y_1 + Y_2$ 

$$= 2A \cos\left[\left(\frac{\Delta w}{2}\right) t - \left(\frac{\Delta k}{2}\right) x\right] \sin\left(\frac{2w + \Delta w}{2}\right) t - \left(\frac{2k + \Delta k}{2}\right) \quad .... (2)$$

But  $\Delta w$  and  $\Delta k$  are small

 $2w + \Delta w \approx 2w$ ,  $2k + \Delta k \approx 2k$ 

$$\therefore Y = 2A \cos \left[ \left( \frac{\Delta w}{2} \right) t - \left( \frac{\Delta k}{2} \right) x \right] \sin (wt-kx) \dots (2)$$

Comparing equations (1) and (2), the coefficient of sin (wt-kx) in equation (2) can be considered as the amplitude of the wave.

Amplitude of the resultant wave =2A cos  $[(\Delta w/2) t - (\Delta k/2) x]$ 

This amplitude varies as a wave .The velocity with which the variation in amplitude is propagated is referred as group velocity

$$V_g = (\Delta w/2) / (\Delta k/2)$$

$$V_{g} = \Delta w / \Delta k = Lt \frac{dw}{dk}$$
 (4)

# 2.b. (4)

Debroglie wavelength 
$$\lambda = \frac{h}{\sqrt{2mE}}$$

 $E=3eV = 3x1.6x10^{-19}J$ 

 $\lambda = 7x10^{-10}\,\text{m} \quad \text{ particle velocity = 1.02x10}^{\text{6}}\text{m/s}$ 

Phase velocity =  $c^2/v_g = 8.76 \times 10^{10} \text{m/s}$ 

# 3.a. (7)

The position and momentum of a particle cannot be determined accurately and simultaneously. The product of uncertainty in the measurement of position  $(\Delta x)$  and momentum  $(\Delta p)$  is always greater than or equal

to 
$$\frac{h}{2\Pi}$$
.(2)

$$(\Delta x) \cdot (\Delta p) \ge \frac{h}{4\Pi}$$

This uncertainty is not due to discrepancy with the apparatus or with the method of measurement, but because of the very wave nature of the object. This uncertainty persists as long as matter possesses wave nature.

Different forms of Heisenberg's Principle:

$$(\Delta x) \cdot (\Delta p) \ge \frac{h}{4\Pi}$$

$$(\Delta L).(\Delta \theta) \ge \frac{h}{4\pi}$$

$$(\Delta E).(\Delta t) \ge \frac{h}{4\pi}$$

Here  $\Delta L$  is the uncertainty in angular momentum

 $\Delta\theta$  is the uncertainty in angular displacement

ΔE is the uncertainty in the energy

 $\Delta t \;\; \text{is the uncertainty in the time interval during which the particle exists}$  in the state E

#### To Show that electron does not exist inside the nucleus:

We know that the diameter of the nucleus is of the order of  $10^{-15}$ m.If the electron is to exist inside the nucleus, then the uncertainty in its position  $\Delta x$  cannot exceed the size of the nucleus

$$\Delta x \le 10^{-14} m$$

Now the uncertainty in momentum is

$$\Delta p \ge \frac{h}{4\pi \cdot \Delta x}$$

$$\Delta p \ge \frac{6.62 \times 10^{-34}}{4\pi \times 10^{-15}}$$

$$\Delta p \ge 0.5 \times 10^{-20} Ns$$

Then the momentum of the electron can at least be equal to the uncertainty in momentum.

$$p \ge 0.5 \times 10^{-20} \, Ns$$

Now the energy of the electron with this momentum supposed to be present in the nucleus is given by - for small velocities (non-relativistic) case-For high velocities,

$$\mathbf{E} = \left(\frac{1}{\sqrt{1 - \left(\frac{\mathbf{v}^2}{\mathbf{c}^2}\right)}}\right) \mathbf{m}_o \mathbf{c}^2$$

$$\mathbf{E}^2 = \left(\frac{1}{1 - \left(\frac{\mathbf{v}^2}{\mathbf{c}^2}\right)}\right) \mathbf{m}^2 \circ \mathbf{c}^4$$

$$\mathbf{p} = \left(\frac{1}{\sqrt{1 - \frac{\mathbf{v}^2}{c^2}}}\right) \mathbf{m}_o \mathbf{v}$$

$$p^2 = \left(\frac{1}{1 - \left(\frac{v^2}{c^2}\right)}\right) m^2 \circ v^2$$

$$p^{2}c^{2} = \left(\frac{1}{1 - \left(\frac{v^{2}}{c^{2}}\right)}\right) m^{2} v^{2}c^{2} \dots (2)$$

From (1) and (2) 
$$E^2 = p^2c^2 + m^2_oc^4$$
 (5)

On substitution

$$E = (9.1x10^{-31})^2 (3x10^8)^2 + (0.5x10^{-19})^2 (3x10^8)^4 = 9.4 MeV$$

## 3.b. (3)

Given v=500m/s accurate to 0.003%

$$\Delta v = 500x \frac{0.003}{100} = 0.015 \text{m/s}$$

From Heisenberg's Principle,

$$\Delta x.\Delta p \ge \frac{h}{4\pi}$$

$$\Delta x.m\Delta v = \frac{h}{4\pi}$$

$$\Delta x = \frac{h}{4\pi . \Delta v.m} = 3.86 x 10^{-3} m$$

# 4.a. (6)

#### Time independent Schrödinger equation

A matter wave can be represented in complex form as

 $\Psi = A \sin kx (\cos wt + i \sin wt)$ 

$$\Psi = A \sin kx e^{iwt}$$

Differentiating wrt x

$$\frac{d\Psi}{dx} = kA\cos kxe^{iwt}$$

$$\frac{d^2\Psi}{dx^2} = -k^2 A \sin kx e^{iwt} = -k^2 \Psi \dots (1)$$

From debroglie's relation (2)

$$\frac{1}{\lambda} = \frac{h}{mv} = \frac{h}{p}$$

$$k = \frac{2\pi}{\lambda} = \frac{2\Pi p}{h}$$

$$k^2 = 4\Pi^2 \frac{p^2}{h^2} \dots (2)$$

Total energy of a particle E = Kinetic energy + Potential Energy

$$E = \frac{1}{2} \text{ m } v^2 + V$$

$$E = \frac{p^2}{2m} + V$$

$$p^2 = (E - V)2m$$
(2)

$$k^{2} = \frac{4\Pi^{2}(E-V)2m}{h^{2}}$$

$$\therefore \text{ From (1)}$$

$$\frac{d^2\Psi}{dx^2} + \frac{8\Pi^2 m(E-V)\Psi}{h^2} = 0$$

(For one dimension) (2)

## 4.b. (4)

Energy of a particle in an infinite potential well  $E = \frac{n^2 h^2}{8ma^2}$ 

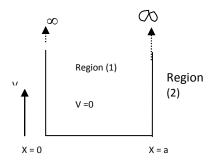
Second excited state corresponds to n= 3

$$E = \frac{3^2 h^2}{8.(9.1x10^{-31}).(0.15x10^{-9})^2} = 2.42x10^{-17} J = 151eV$$
$$\lambda = \frac{h}{\sqrt{2mE}} = 1x10^{-10} m$$

## 5.a. (7)

#### Particle in an infinite potential well problem:

Consider a particle of mass m moving along X-axis in the region from X=0 to X=a in a one dimensional potential well as shown in the diagram. The potential energy is zero inside the region and infinite outside the region.



Applying, Schrodingers equation for region (1) as particle is supposed to be present in region (1) (2)

$$\frac{d^2\Psi}{dx^2} + \frac{8\Pi^2 mE\psi}{h^2} = 0 \qquad \because V = 0$$
But  $k^2 = \frac{8\Pi^2 mE}{h^2}$ 

$$\therefore \frac{d^2 \Psi}{dx^2} + k^2 \Psi = 0$$

The general solution to this expression is given by

$$\Psi = C\cos kx + D\sin kx$$

At x=0, 
$$\Psi = 0$$
  $\therefore C = 0$ 

At x=a, 
$$\Psi=0$$
 D sin ka = 0  $\implies$  ka = n  $\Pi$  where n = 1, 2 3  $\therefore$ 

$$\Psi = D \sin \left( n \frac{\Pi}{a} \right) x$$

$$E = \frac{n^2 h^2}{8ma^2}$$
 (2)

#### To evaluate the constant D:

Normalisation: For one dimension

$$\int_{0}^{a} \Psi^{2} dx = 1$$

$$\int_{0}^{a} D^{2} \sin^{2}(\frac{n\Pi}{a})x dx = 1$$

But 
$$\cos 2\theta = 1 - 2\sin^2 \theta$$

$$\int_{0}^{a} D^{2} \frac{1}{2} (1 - \cos 2(\frac{n\Pi}{a})x) dx = 1$$

$$\int_{0}^{a} \frac{D^{2}}{2} dx - \int_{0}^{a} \frac{1}{2} \cos 2(\frac{n\Pi}{a})x dx = 1$$

$$\frac{D^{2}a}{2} - \left[\sin 2\left(\frac{n\Pi}{a}\right)\frac{x}{2}\right]_{0}^{a} = 1$$

$$D^2 \frac{a}{2} - 0 = 1$$

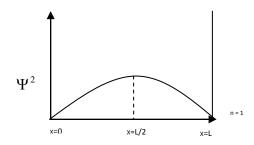
$$D = \sqrt{\frac{2}{a}}$$
 (2)

$$\therefore \Psi_n = \sqrt{\frac{2}{a}} \sin\left(n\frac{\Pi}{a}\right)x$$

For n = 1, First state

$$\therefore \Psi_1 = \sqrt{\frac{2}{a}} \sin\left(1.\frac{\Pi}{a}\right) x$$

The graph of  $\Psi^2$   $versus_x$  is shown below.

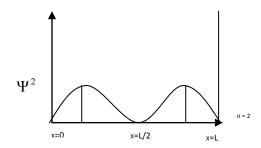


It is seen from the graph that probability density is maximum at the centre for the particle in the first state.

For n = 2, Second state

$$\therefore \Psi_2 = \sqrt{\frac{2}{a}} \sin\left(2.\frac{\Pi}{a}\right) x \text{ (1)}$$

The graph of  $\Psi^2$  versusx is shown below.



It is seen from the graph that probability density is maximum at x = L / 4 and x = 3L/4 for the particle in the second state.

# 5.b. (3)

Change in wavelength of x ray after Compton scattering is given by

$$d\lambda = \lambda^1 - \lambda = \frac{h}{m_0 c} (1 - \cos \theta)$$

Wavelength of Incident photon =  $\lambda = 1x10^{-10} \, m$ 

Here, when  $\theta=90^{\circ}$ ,

Wavelength of scattered photon

$$\lambda^{\!\scriptscriptstyle 1} = \lambda + \frac{h}{m_0 c} \big( 1 - cos 90 \big) = 1 x 10^{-\!10} + 0.0242 x 10^{-\!10} = 1.0242 x 10^{-\!10} \, m$$

## 6.a. (6)

## Failures of Classical free electron theory:

1. Prediction of low specific heats for metals: (2)

Classical free electron theory assumes that conduction electrons are classical particles similar to gas molecules. Hence, they are free to absorb energy in a continuous manner. Hence metals possessing more electrons must have higher heat content. This resulted in high specific heat given by the expression

$$c_v = c_v = \frac{3}{2}R.$$

This was contradicted by experimental results which showed low specific heat for metals given by  $c_v = 10^{-4} \, R$  .

2. Temperature dependence of electrical conductivity: (2)

From the assumption of kinetic theory of gases

$$\frac{3}{2}kT = \frac{1}{2}mv^2$$
$$\therefore v \propto \sqrt{T}$$

Also mean collision time  $\tau$  is inversely proportional to velocity,

$$\tau \propto \frac{1}{v}$$

$$\tau \propto \frac{1}{\sqrt{T}}$$

$$\therefore \sigma = \frac{ne^2 \tau}{m} \Rightarrow \sigma \propto \frac{1}{\sqrt{T}}$$

However experimental studies show that  $\,\sigma \propto rac{1}{T}\,$ 

3. Dependence of electrical conductivity on electron concentration: (2)

As per free electron theory,  $\sigma \propto n$ 

The electrical conductivity of Zinc and Cadmium are  $1.09 \times 10^7$ /ohm m and  $.15 \times 10^7$ /ohm m respectively which are very much less than that for Copper and Silver for which the values are  $5.88 \times 10^7$  /ohm m and  $6.2 \times 10^7$ /ohm m. On the contrary, the electron concentration for zinc and cadmium are  $13.1 \times 10^{28}$  /m<sup>3</sup> and  $9.28 \times 10^{28}$  /m<sup>3</sup> which are much higher than that for Copper and Silver which are  $8.45 \times 10^{28}$  /m<sup>3</sup> and  $5.85 \times 10^{28}$  /m<sup>3</sup>.

These examples indicate that  $\sigma \propto n$  does not hold good.

4. Mean free path, mean collision time found from classical theory are incorrect

## 6.b. (4)

#### Case 1:

For  $E < E_F$ , at T = 0,

$$f(E) = \frac{1}{e^{(\frac{E-E_F}{kT})} + 1} = 1$$

This shows that energy levels below Fermi energy are completely occupied:

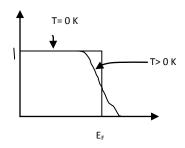
For E >  $E_{\scriptscriptstyle F}$  , at T=0

$$f(E) = \frac{1}{e^{(\frac{E-E_F}{kT})} + 1} = 0$$

This shows that energy levels above Fermi energy are empty:

At ordinary temperatures, for  $E = E_{F}$ .

$$f(E) = \frac{1}{2}$$



## 7.a. (6)

#### **Expression for Electrical conductivity:**

Imagine a conductor across which an electric field E is applied. The equation of motion for an electron moving under the influence of external field is given by F = dp/dt = eE

Let the wave number change from  $k_1 \, to \, k_2$  in time interval  $\, \tau_F$  in the presence of electric field.

$$k = \frac{2\pi}{\lambda} = \frac{2\pi}{h/p} = \frac{2\pi p}{h}$$

$$p = \frac{hk}{2\pi}$$

$$\frac{dp}{dt} = \frac{h}{2\pi} \left(\frac{dk}{dt}\right)$$

$$dk = \frac{2\pi}{h} eEdt$$
(3)

On integration

$$k_2 - k_1 = \Delta k = \frac{2\pi .eE.\tau_F}{h}$$
 .....(1)

From quantum theory, conductivity J =  $\Delta k.ne.\frac{h}{2\pi.m}$  .....(2)

Substituting (1) in (2)

We get J = 
$$\frac{ne^2 \tau_F}{m^*} E$$
 ...(3) (1)

Since from Ohm's,  $J = \sigma E$ , conductivity  $\sigma$  can be written as

$$\sigma = \frac{ne^2}{m^*} \frac{\lambda}{v_F}$$
 where  $\tau = \frac{\lambda}{v_F}$  (2)

# 7.b. (4)

### Fermi energy ( $E_{\scriptscriptstyle F}$ ):

It is the highest energy possessed by an electron at zero Kelvin.

$$E_F = \frac{h^2}{8m} \left(\frac{3n}{\Pi}\right)^{\frac{2}{3}} \tag{1}$$

<u>Fermi Temperature:</u> It is the temperature to which a metal is to be heated such that the free electrons acquire energy equal to Fermi energy.

$$T_F = \frac{E_F}{k}$$
 (1)

### Drift velocity:

The net displacement in the position of electrons per unit time caused by the application of electric field is known as drift velocity.

$$v_d = \frac{eE\tau}{m}$$
 (2)