

Module :- 3

- Optical transition in Bulk Semiconductors
- Absorption, Spontaneous Emission & Stimulated emission
- Einstein Coefficients
- Population Inversion
- Applications in Semiconductor Laser
- Joint density of States
- Density of States of Phonons
- Transition rates (Fermi Golden Rule)
- Optical loss & Gain
- Photovoltaic Effect
- Exciton
- Drude Model

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LASER

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Light Amplification by Stimulated Emission
of Radiation

→ Different Mechanisms

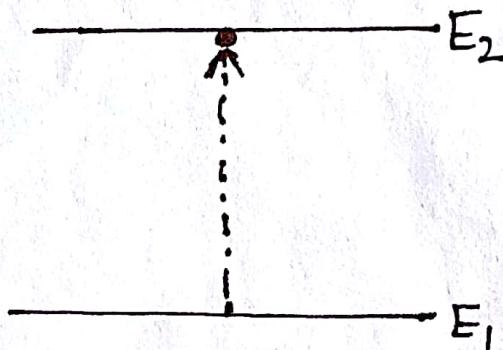
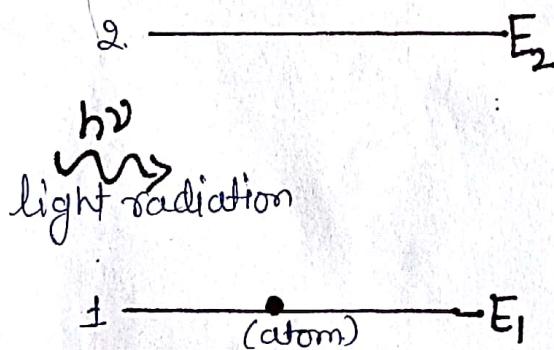
* Absorption

* Spontaneous emission

* Stimulated emission or Induced emission.

→ ABSORPTION

Let us consider two energy levels E_1 & E_2 . Let the atom is initially in lower energy state E_1 . Let atom is exposed to light radiation (Photon) with energy $h\nu$. Atom get the energy from photon & jump to higher energy level E_2 .



Before

$$\text{Now, } E_2 - E_1 = h\nu$$

$$\gamma = \frac{E_2 - E_1}{h}$$

After

where h = Planck's const.

Now; probable rate of transition from 1 to 2 depends on properties of states 1 & 2 & energy density $\nu(r)$ i.e.

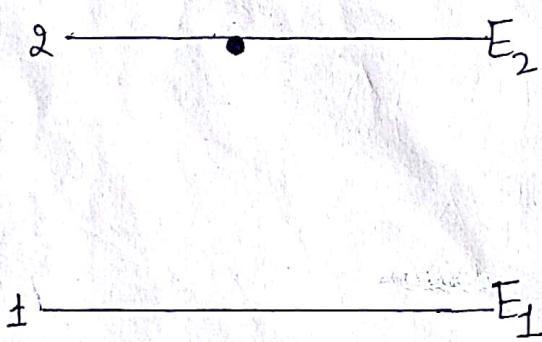
$$P_{12} = B_{12} \nu(r)$$

where B_{12} = constant of proportionality.
Called Einstein coefficient of absorption of radiation

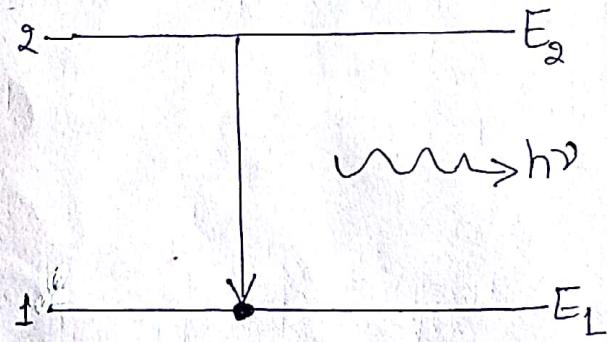
SPONTANEOUS EMISSION

Let us consider two energy levels E_1 & E_2 ($E_1 < E_2$)
Let initially atom in higher energy level E_2 .

An atom remains in excited state only for 10^{-8} sec. after this, its own jump to lower energy state E_1 by emitting a photon with frequency ν . This type of emission is called spontaneous emission of radiation.



Initial State



Final State

$$\nu = \frac{E_2 - E_1}{h}$$

Now, probable rate of transition from $2 \rightarrow 1$ depends on properties of states of 1 & 2.

So,

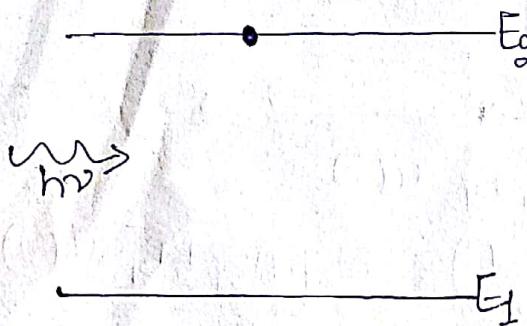
$$(P_{21})_{\text{spont}} = A_{21}$$

where A_{21} = Einstein coefficient of spontaneous emission

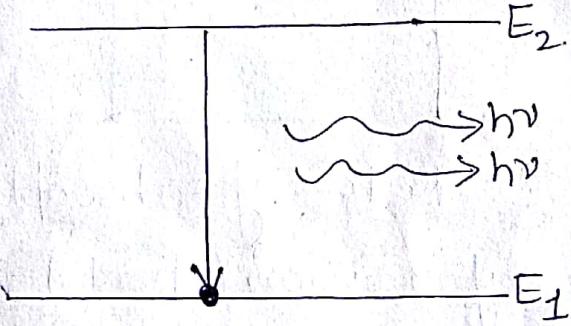
STIMULATED OR INDUCED EMISSION

2

Let us consider two energy levels E_1 & E_2 ($E_1 < E_2$). Initially atom is in higher energy level E_2 . If a photon of frequency ν is incident on it & cause the atom to jump to lower energy state E_1 by emitting an additional photon of frequency ν . This process is called stimulated emission of radiation.



Initial State



final State

Now, probable rate of transition from $2 \rightarrow 1$ depends on energy density of as well as properties of states of 1 & 2.

$$(P_{g1})_{\text{stimulated}} = B_{21} u(\nu)$$

→ Difference b/w spontaneous & stimulated emission

SPONTANEOUS EMISSION

- Emission has broad spectrum i.e. many wavelengths
- It gives incoherent radiation
- Less intense
- This emission gives radiations with less directionality & more angular spread.

e.g.- Ordinary source of light

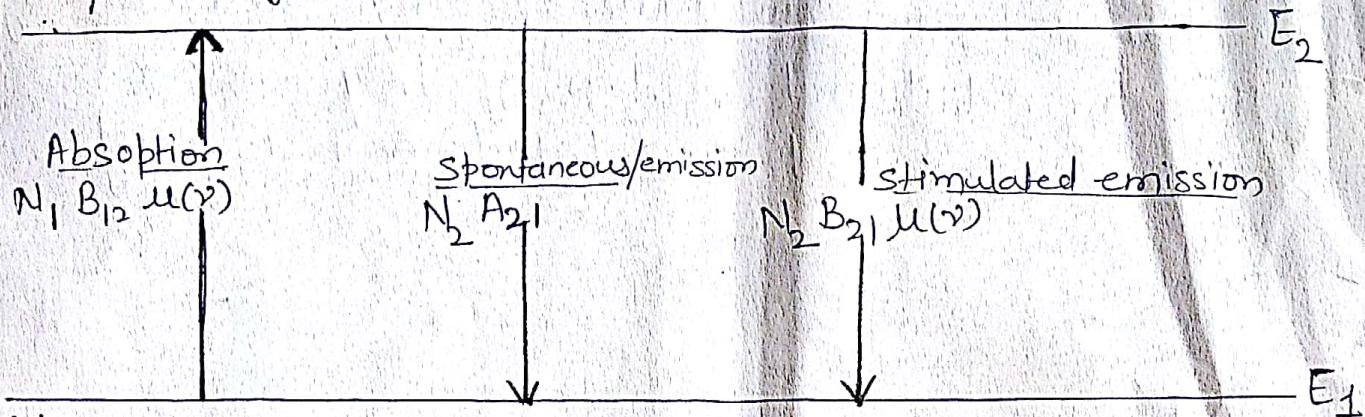
STIMULATED EMISSION

- Emission has monochromatic radiation
- If gives coherent radiations highly intense
- This emission gives radiations with high directionality & less angular spread

e.g:- Laser

→ Einstein Coefficients of Relations.

Let us consider two energy level E_1 & E_2 ($E_1 < E_2$)
 Let N_1 & N_2 number of atoms in energy states 1 &
 2 respectively.



Now; Rate of absorption = $N_1 B_{12} u(v)$

Rate of emission (spont + stimulated) = $N_2 [A_{21} + B_{12} u(v)]$
 emission emission

AT equilibrium Rate of absorption = Rate of emission

So;

$$N_1 B_{12} u(v) = N_2 [A_{21} + B_{12} u(v)]$$

$$N_1 B_{12} u(v) - N_2 B_{12} u(v) = N_2 A_{21}$$

$$u(v) = \frac{N_2 A_{21}}{N_1 B_{12} - N_2 B_{12}}$$

$$u(v) = \frac{A_{21}}{\frac{N_1 B_{12} - B_{12}}{N_2}}$$

÷ dividing Numerator
 & Denominator by N_2

$$u(v) = \frac{A_{21}}{B_{21} \left[\frac{N_1}{N_2} \frac{B_{12}}{B_{21}} - 1 \right]} \quad (1)$$

According to Boltzmann distribution law, number
 of atoms N_1 & N_2 in Energy state E_1 & E_2 at temp T

$$N_2 = N_1 e^{-hv/kT} \quad \text{where } h = \text{Planck's Const.} \quad (3)$$

$$\frac{N_2}{N_1} = e^{-hv/kT}$$

$$\text{So, } \frac{N_1}{N_2} = e^{hv/kT}$$

Put the value of $\frac{N_1}{N_2}$ in Eq (1.)

$$u(v) = \frac{A_{21}}{B_{21}} \frac{1}{e^{hv/kT} B_{12} - 1}$$

Now, comparing it by Planck's radiation law

$$u(v) = \frac{8\pi hv^3}{c^3} \frac{1}{e^{hv/kT} + 1}$$

We get,

$$\boxed{\frac{A_{21}}{B_{21}} = \frac{8\pi hv^3}{c^3}} \quad \neq \quad \boxed{B_{12} = B_{21}}$$

$\therefore B_{21} \propto \frac{1}{v^3}$, So rate of stimulated emission is higher at lower frequencies

$B_{12} = B_{21}$ means probability of stimulated emission is same as that of absorption

Ques What are the units of Einstein Coefficients?

Answer:- Energy density $u(v) = \frac{\text{Energy}}{\text{Volume} \times \text{freq.}}$

$$\text{Energy density } u(v) = \frac{\text{Energy} \times \text{Time}}{\text{Volume}} = \frac{\text{JS}}{\text{m}^3}$$

Units of $(P_{12})_{\text{spot emission}}, (P_{21})_{\text{stimulated emission}}$] = $\frac{\text{Number}}{\text{Time}} = \text{s}^{-1}$

Now for absorption

$$P_{12} = B_{12} N_1 u(\gamma)$$

$$\Rightarrow B_{12} = \frac{P_{12}}{N_1 u(\gamma)} = \frac{\text{s}^{-1} \text{m}^3}{\text{J s}} = \text{m}^3 \text{J}^{-1} \text{s}^{-2}$$

Similarly, for Stimulated Emission

$$\Rightarrow B_{21} = \text{m}^3 \text{J}^{-1} \text{s}^{-2}$$

for Spontaneous Emission

$$(P_{21})_{\text{spot emission}} = A_{21}$$

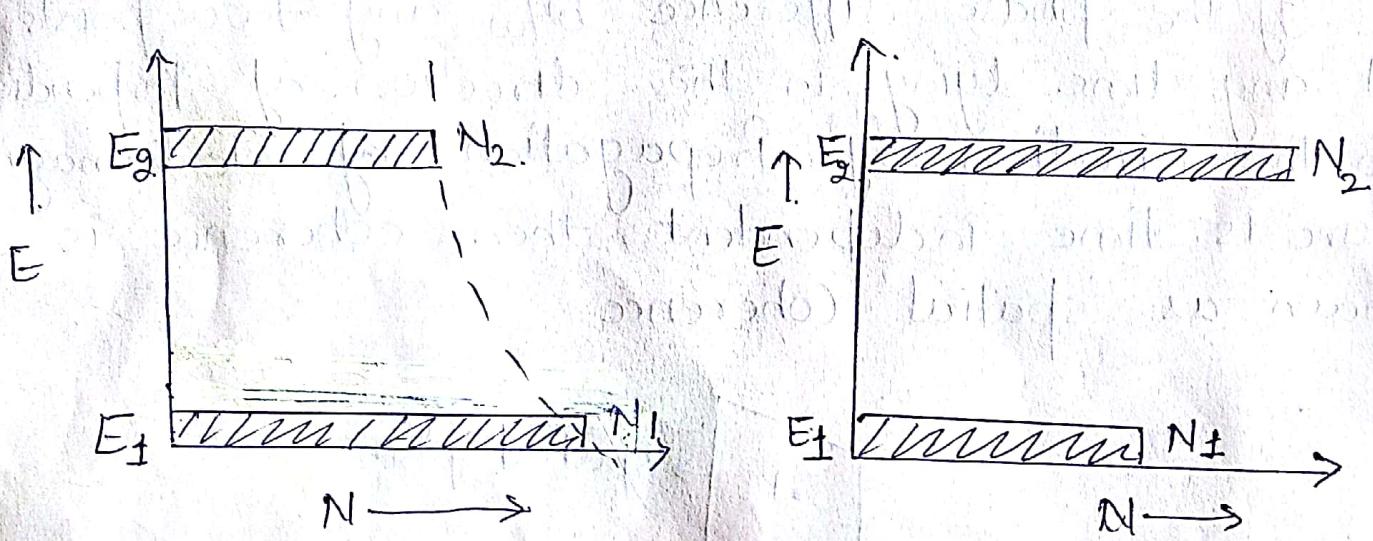
$$A_{21} = \text{s}^{-1}$$

Population Inversion

Let us consider two Energy levels (E_1 & E_2) where ($E_1 < E_2$). Let N_1 & N_2 be the number of atoms in energy levels E_1 & E_2 respectively.

Under Ordinary Conditions of.

thermal equilibrium no. of atoms in Energy level E_1 is greater than E_2 . i.e. ($N_1 > N_2$) But the condition in which no. of atoms in energy level E_2 is greater than E_1 i.e. ($N_2 > N_1$) is known as population Inversion. It is necessary & essential Condition for Laser action to takes place.



Pumping
The process through which state of Population Inversion is achieved is known as Pumping. There are various types of pumping like.

- (1) Optical Pumping (used in Ruby Laser)
- (2) Electric discharge (used in He-Ne Laser)
- (3) Inelastic atom-atom collision
- (4) Direct Converstion (used in Semiconductor Laser)
- (5) chemical reaction (used in CO_2 Laser)

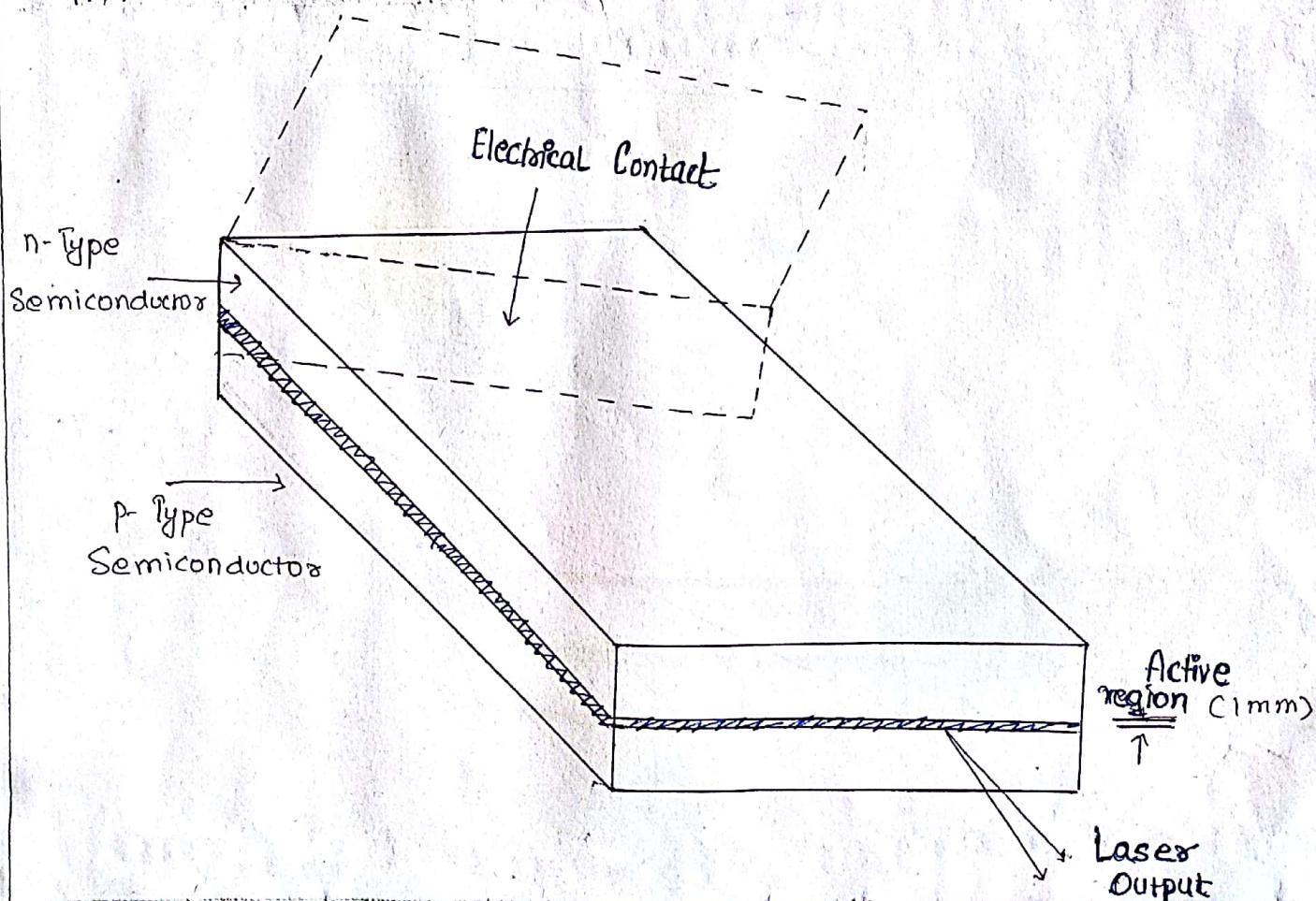
Semiconductor Laser

Semiconductor Lasers are also known as laser diode. It is pn junction device which emit coherent light in forward bias condition.

Laser diode is very popular for their compactness & operational efficiency.

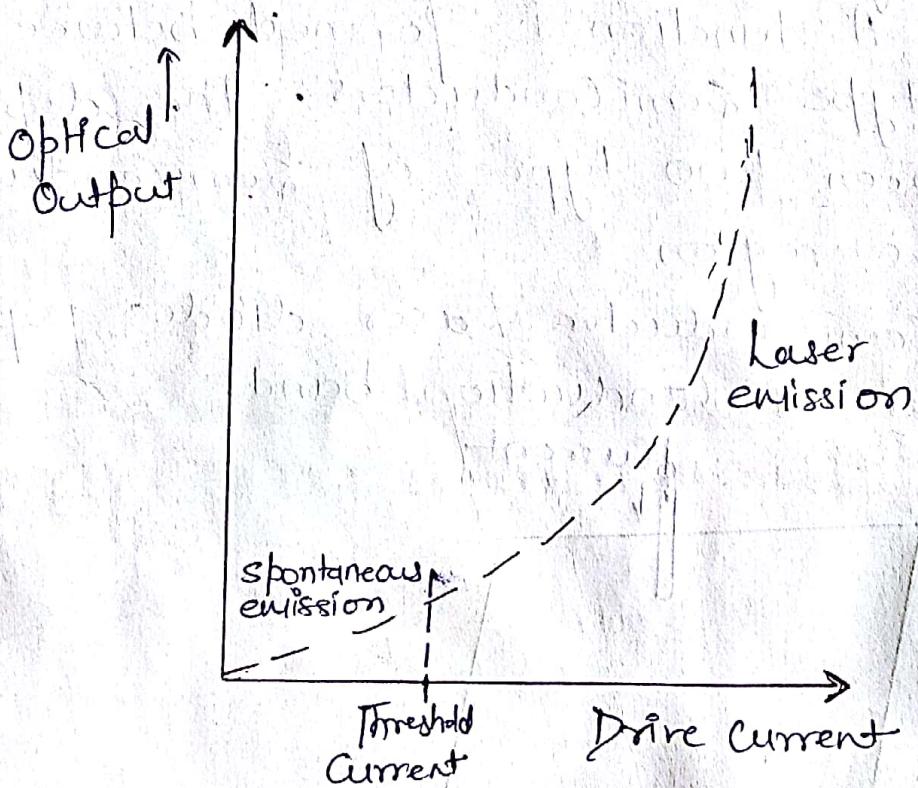
The Semiconductor laser diode are made up of direct band Semiconductors like GaAs. A Junction is formed between P-type & N-type Semiconductors. The depletion region between two types of Semiconductors act as active region.

In Semiconductor Laser diodes population inversion occur in conduction band.



When laser diode is in forward bias electrons from n-side & holes from p-side diffuse into depletion region of both carriers combine & energy is released in form of light photons.

Under forward biased condition when drive current through electrical connections reaches threshold level, population inversion results of laser emission will takes place.



Semiconductor Laser is different from Solid Gas & liquid laser as

- (1) Junction laser is quite small (approx $0.1 \times 0.1 \times 0.3$ mm)
- (2) High efficiency.
- (3) Laser output is easily modulated by.

Controlling the Junction Current.

- (4) Semiconductor Lasers operate at low power compared to ruby or CO_2 lasers.
- (5) Output power is higher than He-Ne Laser.
- (6) These lasers are portable also

Optical Transitions in Bulk Semiconductors

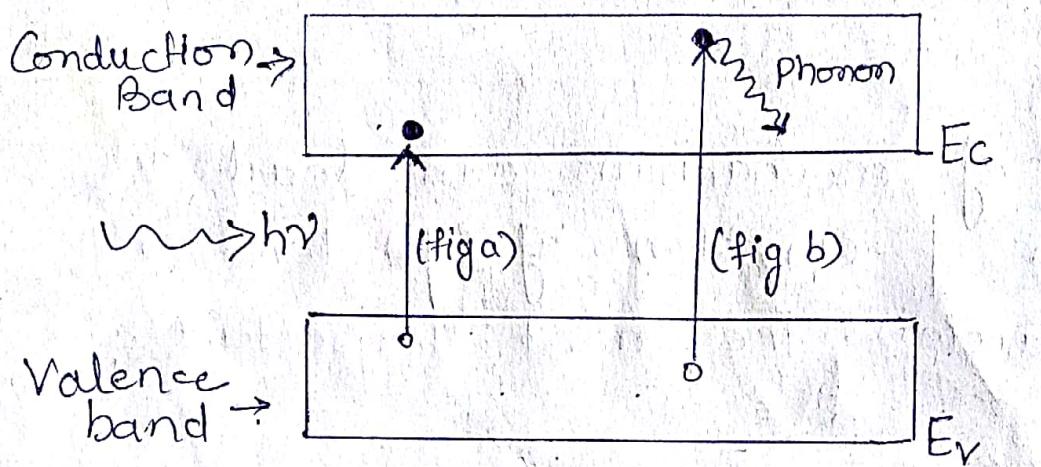
A number of mechanisms can lead to the absorption & emission of photons in bulk Semiconductors. The most important of these are

- Band to Band (Interband Transition)
- Impurity to band transitions
- free Carriers (Intraband Transitions)
- Phonon Transitions
- Excitonic Transitions

I. Band to Band (Interband Transitions)

When a Semiconductor is illuminated, photons are absorbed to create e^- -hole pairs. as shown in fig (a) for a photon of energy $h\nu = E_g$.

If a 'incident' photon having Energy $(h\nu)$ is greater than E_g , again e^\ominus hole is generated of the excess Energy $(h\nu - E_g)$ is dissipated through phonons as heat, as shown in fig(b). Both process is called Intrinsic transitions or band to band transitions.



If it is true for reverse situation also. For eg., an e^\ominus at Conduction band edge (E_c), combining with hole at Valence band edge (E_v), will gives emission of photon with Energy $(h\nu)$ equal to E_g . i.e

$$h\nu = E_g$$

for Absorption & Emission of photons

$$h\nu > E_g$$

The minimum frequency ν_{\min} necessary for transition is

$$\nu_{\min} = \frac{E_g}{h}$$

f. Corresponding wavelength (λ_{\max}) called
Cut off wavelength is

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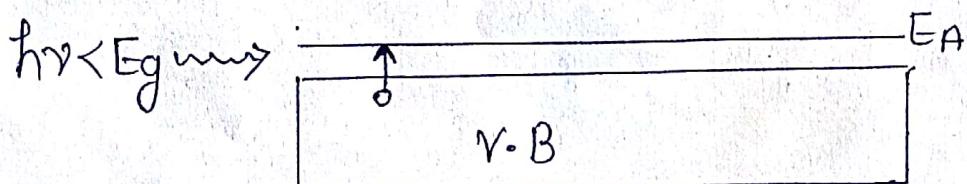
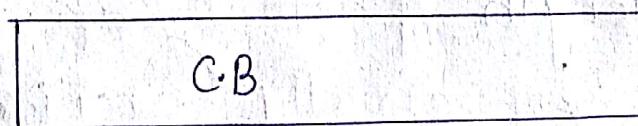
$$\lambda_{\max} = \frac{c}{\nu_{\min}} = \frac{c h}{E_g} \Rightarrow \frac{1.24 \text{ (nm)}}{E_g \text{ (eV)}}$$

2. Impurity to band Transition

If $h\nu < E_g$, a photon will be absorbed only if there are available energy states within forbidden gap (like donor level or acceptor level) due to impurities

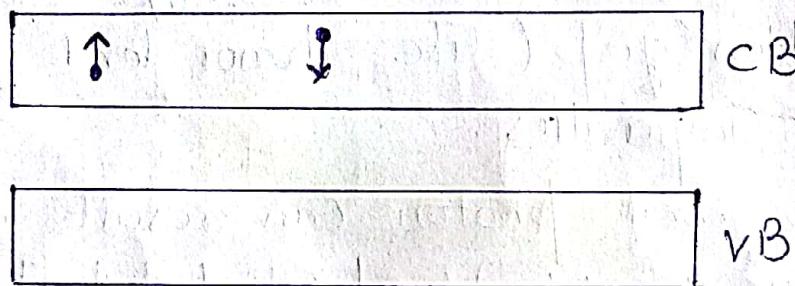
An absorbed photon can result in transition b/w donor (or acceptor) level called Impurity to band transition

e.g:- A low energy photon lift an electron from valence band of p-type Semiconductor to the acceptor level, where it get trapped inside the inside the acceptor atom. In this process hole is created in valence band & acceptor atom get ionized.



3. free Carrier (Intraband Transition)

An absorbed photon can impart its energy to the electron in a given band, causing it to move higher within that of band. for example An electron in Conduction band can absorb a photon & move to the higher Energy level within the Conduction band.



This will followed by the process of thermalization, where the electron relax down to the bottom of Conduction band while releasing its Energy in form of lattice vibrations.

4. Phonon Transition

long wavelength photons can release their Energy by directly exciting lattice vibrations i.e. by creating phonons.

5. Excitonic Transitions

These transitions appear only in pure Semiconductors. The pairing of electron hole pair produced near the band gap is called exciton. (It is very much likely the hydrogen atom where electron f proton pair with a definite binding energy). The excitons appear in pure Semiconductors because of their low binding Energy f they are strongly temperature dependent.

Optical Joint Density of States

OR.

Density of States for photons

The quantity $D(\nu)$, which incorporates the density of states in both Conduction f Valence band is called Optical joint density of states because a photon is emitted only when an electron jump from one of the state of Conduction band to one of the state in Valence band

We know density of states in Conduction band is given by:

$$D(E) = N(E) = \frac{4\pi}{h^3} (2m_e^*)^{3/2} (E - E_c)^{1/2} \quad \text{from unit-2}$$

where $E > E_c$

E_c = energy at bottom of conduction band

$$D(E) = N(E) = 4\pi \left(\frac{2m_e^*}{h^2} \right)^{3/2} (E - E_c)^{1/2} \quad (\pm)$$

We know

$$E = E_c + \frac{m_r}{m_e^*} (hv - E_g)$$

$$\therefore E - E_c = \frac{m_r}{m_e^*} (hv - E_g) \quad (2)$$

Put Eqⁿ (2) in (1)

$$D(E) = N(E) = 4\pi \left(\frac{2m_e^*}{h^2} \right)^{3/2} \left(\frac{m_r}{m_e^*} \right)^{1/2} (hv - E_g)^{1/2} \quad (3)$$

Now; density of states for photons is

$$D(\gamma) = \frac{dE}{d\gamma} |D(E)|$$

$$D(\gamma) = \left(\frac{m_r}{m_e^*} h \right) 4\pi \left(\frac{2m_e^*}{h^2} \right)^{3/2} \left(\frac{m_r}{m_e^*} \right)^{1/2} (hv - E_g)^{1/2}$$

$$D(\gamma) = \frac{4\pi}{h^2} (2m_e^*)^{3/2} (hv - E_g)^{1/2}$$

Fermi Golden Rule

Fermi Golden rule is a formula that describes the transition rate from one energy state into another energy state.

In general, the transition rate (or the transition probability per unit time) depends on

- (1) Strength of coupling b/w initial & final state
- (2) the number of ways the transition can happen i.e. density of final state.

The transition probability per unit has the form

$$\lambda_{if} = \frac{2\pi}{\hbar} |M_{if}|^2 f_F$$

where $\lambda_{if} \Rightarrow$ Transition probability per unit time

$|M_{if}| \Rightarrow$ Coupling term b/w initial & final state

$f_F \Rightarrow$ Density of final state

The transition is high if the coupling b/w initial & final state is stronger. The coupling term is traditionally called "the matrix" element for the transition,

is given by

$$M_{if} = \int \Psi_f^* V \Psi_i dV$$

where Ψ_i = Wave function of initial state

V = operator for the physical interaction which couples the initial & final state.

Ψ_f^* = Wave function of final state

The transition rate b/tw two energy states is given by

$$A_{if} = \frac{2\pi}{\hbar} |H_f'|^2 S(E_f - E_i - \hbar\omega) + \frac{2\pi}{\hbar} |H_{if}'|^2 S(E_f - E_i + \hbar\omega)$$

where first term corresponds to the absorption of photon where second term corresponds to emission of photon.

Optical Gain & Loss in two level System

Let us consider a two level system. Suppose a coherent monochromatic radiation of unit cross-sectional area passing through the laser medium.

Inside the medium, absorption of radiation takes place so intensity varies as

$$I = I_0 e^{-\alpha x} \quad (1)$$

Now, differentiating. (where I_0 = intensity of incident radiation ($x=0$))

$$\frac{dI}{dx} = -\alpha I_0 e^{-\alpha x} \quad \alpha = \text{absorption coefficient}$$

$$\frac{dI}{dx} = -\alpha I \quad \text{[using Eq 1]}$$

If $N_1 + N_2$ no. of electrons in Energy level $E_1 + E_2$ respectively then Net rate of loss of photon density N_p is

$$-h\nu \frac{dN_p}{dt} = N_1 B_{12} \mu(\nu) - N_2 B_{21} \mu(\nu) \quad (2)$$

as $B_{12} = B_{21}$ {relation b/w Einstein coefficients}

as Eq (2) becomes

$$-h\nu \frac{dN_p}{dt} = (N_1 - N_2) B_{21} \mu(\nu) \quad (3)$$

Now; Energy density is

$$u(\nu) = N_p h \nu \quad (4)$$

(where N_p = number of photons per unit volume having freq ν)

So, for photons of frequency ν is

$$I = u(\nu) \frac{c}{n_r} \quad (4(a))$$

(where n_r = refractive index of medium)

$$I = \frac{N_p h \nu c}{n_r}$$

$$N_p = \frac{I n_r}{h \nu c}$$

If incident beam is passed through the medium b/w x & $x + \Delta x$, then change in photon density dN_p is

$$-dN_p = \frac{[I(x) - I(x + \Delta x)] n_r}{h \nu c}$$

$$-dN_p = -\frac{dI}{dx} \frac{n_r \Delta x}{h \nu c} \quad (5)$$

Now, photons travel the distance Δx in time dt is $dt = \frac{\Delta x}{c/n_r}$.

so Eq (5) becomes

$$-\frac{dN_p}{dt} = -\frac{dI}{dx} \cdot \frac{1}{h\nu} dt$$

$$-\frac{dN_p}{dt} = -\frac{dI}{dx} \cdot \frac{1}{h\nu} \quad \text{--- (6)}$$

Now, put the $\frac{dI}{dx}$ using Eq(1a) in above.

Eq.

$$-\frac{dN_p}{dt} = \frac{\alpha I}{h\nu}$$

$$\frac{dN_p}{dt} = -\frac{\alpha I}{h\nu}$$

Now put the value of I using 4(a)

$$\frac{dN_p}{dt} = -\alpha \mu(v) \frac{c}{n_g} \frac{1}{h\nu}$$

$$-h\nu \frac{dN_p}{dt} = -\alpha \mu(v) \frac{c}{n_g} \quad \text{--- (7)}$$

Now compare Eq (3) + (7) we get

$$(N_1 - N_2) B_{g1} \mu(v) = -\alpha \mu(v) \frac{c}{n_g}$$

$$\alpha = \frac{(N_1 - N_2) B_{g1} n_g}{c}$$

If $N_1 > N_2$ so α is positive which represent the medium with loss whereas if $N_2 > N_1$ then α become negative. It represent optical gain.

Photovoltaic Effect

In 1839, Becquerel discovered that when a pair of electrodes is dipped into electrolyte. If light is allowed to incident on one of the electrodes, a potential difference is created b/w these electrodes. The phenomenon is known as photovoltaic effect.

When photons are absorbed by photovoltaic cell, the energy from photon is transferred to an electron in an atom. The energized e^- is then escape its bond with in atom & generates an electric current.

Drude-Model

According to drude Model optical Conductivity & dielectric constant can be determined by considering the motion of free electrons. In presence of collisions, the equation of free e^- in electric field is

$$m \left[\frac{d^2x}{dt^2} \right] + \frac{m}{\tau} \left[\frac{dx}{dt} \right] = -eE \quad \text{--- (1)}$$

Collision term which is proportional
to velocity also called
friction term

where m = mass of e^-
 x = displacement
 τ = relaxation time

Now if

$$x = x_0 e^{-i\omega t} \quad \text{--- (2)}$$

then differentiate above Eqⁿ

$$\frac{dx}{dt} = -i\omega x_0 e^{-i\omega t}$$

$$\frac{dx}{dt} = -i\omega x \quad \{ \text{using Eq } 2 \}$$

again differentiate Eqⁿ (2)

$$\frac{d^2x}{dt^2} = -i\omega x - i\omega x_0 e^{-i\omega t}$$

$$\frac{d^2x}{dt^2} = -\omega^2 x \quad \text{using Eq 2}$$

Now put the value of $\frac{dx}{dt}$ & $\frac{d^2x}{dt^2}$ from above Eq in Eq (1)

$$-m\omega^2 x - i\frac{\omega m x}{c} = -eE$$

$$-x \left[m\omega^2 + i\frac{\omega m}{c} \right] = -eE$$

$$x = \frac{eE}{m\omega^2 + i\frac{\omega m}{c}} \quad (3)$$

To obtain optical conductivity, we consider $x = v/c$ — (4) where v = mean velocity of e^-
So current density

$$J = nev = \frac{ne x}{c}$$

Now put the value of x from Eq (3)

$$J = \frac{ne}{c} \left[\frac{eE}{m\omega^2 + i\frac{\omega m}{c}} \right] \quad (4)$$

$$\text{we know } J = \sigma E$$

$$\sigma = \frac{J}{E}$$

$$\sigma = \frac{ne^2}{c \left[m\omega^2 + i\frac{\omega m}{c} \right]} \quad \text{using Eq 4}$$

$$\sigma = \frac{ne^2}{mw\gamma \left[\omega + i\frac{\gamma}{c} \right]}$$

when collision frequency $\gamma \left(\frac{1}{c} \right)$ is small as compared with frequency ω of light i.e. in absence of collision above eq reduces to

$$\sigma = \frac{ne^2}{mw\gamma}$$

$$\gamma = \frac{eE}{mw^2}$$

This is known as Drude formula for optical conductivity.