

TOPIC :- GENERATION - RECOMBINATION

Let 'n' and 'p' denote the concentrations of electrons and holes in a semiconductor sample. At equilibrium we have

$$np = n_i^2 \quad (\text{law of mass action}).$$

where  $n_i \propto e^{-E_g/2kT}$ ,  
 $E_g$  being the band gap.

Under such conditions, there is a detailed balance of ~~re~~ carrier recombination due to various processes and the corresponding generation mechanisms.

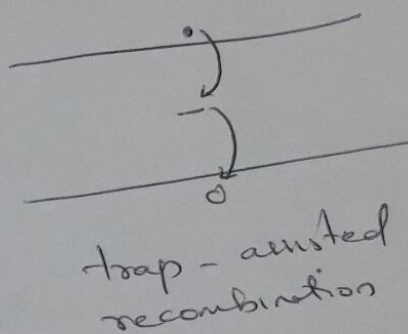
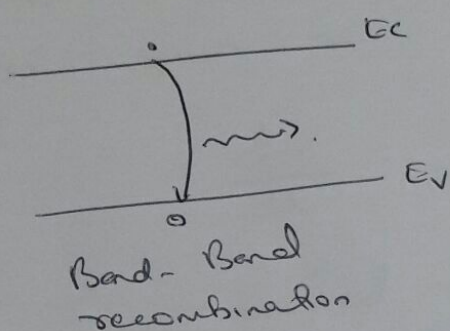
Once driven out of equilibrium, if the excitation or perturbation is held constant, the system will reach "steady state" conditions in which the time rate of change of all parameters will be zero.

The detailed recombination process will determine how the system reaches steady state or equilibrium condition.



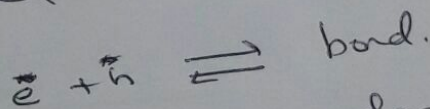
# A. Common RECOMBINATION MECHANISMS

In this course we will consider two recombination mechanisms, viz. Band-Band (radiative) recombination and trap-assisted SRH (non-radiative) recombination mechanism.



## A.1

Band-Band recombination can be represented as



Hence, the rate equation is given as

$$\frac{dp}{dt} = \frac{dn}{dt} = - \underset{\substack{\downarrow \\ \text{recombination} \\ \text{(forward)} \\ \text{reaction}}}{k n \cdot p} + \underset{\substack{\downarrow \\ \text{generation} \\ \text{(reverse)} \\ \text{reaction}}}{G}$$

$k$  - radiative recombination coefficient

The above equation, at equilibrium condition, should result in

$$\frac{dn}{dt} \bigg|_{\text{equilibrium}} = 0.$$



or rather

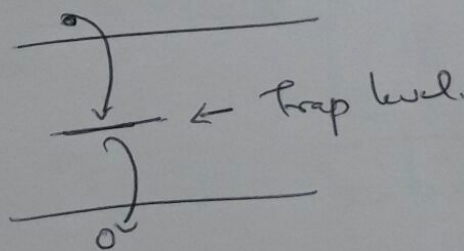
$$G_{\text{equilibrium}} = k n_i^2$$

then We further assume that the above generation rate remains the same for out of equilibrium condition as well. This indicates that the rate equation is given as

$$\frac{dn}{dt} = -k(np - n_i^2)$$

## A.2 Trap-assisted recombination.

For a proper derivation, see the text book AOSF. Here we provide an approximate analysis.

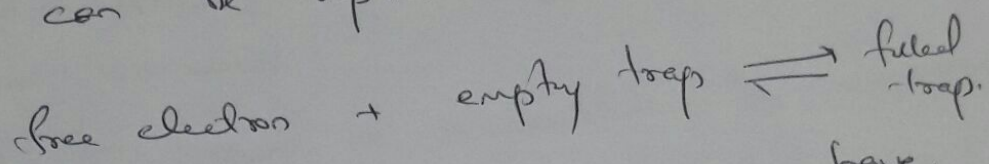


\* We consider near midgap trap states. As all recombination are pair-wise process we have

$$\frac{dn}{dt} = \frac{dp}{dt}$$



Let  $N_T$  be the density of trap states and  $n_T$  of them be filled with electrons at any time. The electron capture can be represented as.



Neglecting the reverse reaction, ~~the~~ we have.

$$\frac{dn}{dt} = -c_n (N_T - n_T) n$$

$c_n$  - electron capture coefficient

Similarly  $\frac{dp}{dt} = -c_p (n_T) p$

Since  $\frac{dn}{dt} = \frac{dp}{dt}$ , we find.

$$n_T = \frac{c_n N_T n}{c_n n + c_p p}$$

Accordingly  $\frac{dn}{dt} = \frac{dp}{dt} = - \frac{c_n c_p N_T n p}{c_n n + c_p p}$

the above equation needs further modification

- as
- ① The emission process is neglected.
  - ② The influence of trap location within



⑤

the band gap should be incorporated.

Item ① can be easily accounted for by replacing  $n_p$  with  $(n_p - n_i^2)$ . This will ensure that the rate will be zero at equilibrium conditions as well.

Item ② can be accounted for by approximating the rate equation as follows.

$$\frac{dn}{dt} = \frac{dp}{dt} = \frac{-(n_p - n_i^2)}{Z_p(n + n_1) + Z_n(p + p_1)}$$

where

$$Z_p = \frac{1}{C_p N_T}$$

$$Z_n = \frac{1}{C_n N_T} \exp\left(\frac{E_T' - E_i}{kT}\right)$$

$$n_1 = n_i \exp\left(\frac{E_i - E_T'}{kT}\right)$$

$$p_1 = p_i$$

Note that  $n_1 \neq p_1$  ensure that the rate  $\rightarrow 0$  as the trap levels are close to the conduction and valence band edges.



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B. Effective lifetime

Under certain conditions, the rate equation could be reduced to the form

$$\frac{dn'}{dt} = -\frac{n'}{\tau}, \text{ where } \tau \text{ could be}$$

the effective lifetime of excess carriers.

B1. Effective lifetime in the presence of Band-Band recombination (only).

We have

$$\frac{dn}{dt} = -k(np - n_i^2)$$

As any generation process will lead to pair-wise creation of e-h, we have  
(this assumes no spatial variation)

$$n = n_0 + \Delta n$$

$$p = p_0 + \Delta p$$

$$\left. \begin{array}{l} n_0 - \text{equilibrium} \\ p_0 - \text{conc.} \\ n_0 p_0 = n_i^2 \end{array} \right\}$$

$$\Delta n = \Delta p$$

Substituting in the rate equation, we have

$$\frac{d(n_0 + \Delta n)}{dt} = -k((n_0 + \Delta n)(p_0 + \Delta n) - n_i^2)$$

$$\Rightarrow \frac{d(\Delta n)}{dt} = -k(\Delta n(n_0 + p_0) + \Delta n^2)$$



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Assuming low-level injection (i.e.  $\Delta n < n_0$  or  $p_0$  the majority carrier concentration), we have

$$\frac{d(\Delta n)}{dt} = -k \Delta n (n_0 + p_0)$$

$$\text{or } \tau_{\text{radiative}} = \frac{1}{k (n_0 + p_0)}$$

B2. Effective lifetime for SRH process (only)

Here we have

$$\frac{dn}{dt} = - \frac{(np - n_i^2)}{\tau_n(p + p_1) + \tau_p(n + n_1)}$$

Assume the traps are near midlevel.

(i.e.  $n_1 = p_1 = n_i$ ) and the sample is

P-type doped, (i.e.  $p_0 \gg n_0$ ).

Following the analysis in previous section, we have

$$\frac{d(n_0 + \Delta n)}{dt} = - \frac{(n_0 + \Delta n)(p_0 + \Delta n) - n_i^2}{\tau_n(p_0 + \Delta n + n_i) + \tau_p(n_0 + \Delta n + n_i)}$$

Assuming low level injection ( $\Delta n = \Delta p \ll p_0$ ), the term that dominates the denominator is  $\tau_n p_0$ .



$$\Rightarrow \frac{dp}{dt} = \frac{dn}{dt} = -\frac{\Delta n}{\tau_n} = -\frac{\Delta p}{\tau_n}$$

or rather the minority carrier effective lifetime  $\tau_{SRH} = \tau_n$ .

\* Note that the above analysis could lead to different expressions depending on the specific values of  $n_0$  &  $p_0$ . For example for intrinsic semiconductor, low level injection assumption will not hold, and the  $\tau_{SRH}$  will be different.

B.3: Effective life time in the presence of both recombination and SRH processes (low level injection)

Here we have.

$$\begin{aligned} \frac{dn}{dt} = \frac{dp}{dt} &= -k N_A \Delta n - \frac{\Delta n}{\tau_n} \\ &= -\frac{\Delta n}{\tau_{eff}} \end{aligned}$$

$$\Rightarrow \frac{1}{\tau_{eff}} = k N_A + \frac{1}{\tau_n}$$



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### C. Carrier Concentrations in steady state.

In the presence of external perturbation like generation due to an optical source the rate equation becomes. (low level injection)

$$\frac{dn}{dt} = \frac{dp}{dt} = -\frac{\Delta n}{\tau_{eff}} + G_{ex}$$

$G_{ex}$  - Generation due to external source

At steady state, we have

$$\frac{dn}{dt} = \frac{dp}{dt} = 0 \Rightarrow$$

$$\Delta n = G_{ex} \tau_{eff}$$

Hence  $n = n_0 + \Delta n$   
 $p = p_0 + \Delta n$

Note  $np \neq n_i^2$ .

So, it is convenient to use quasi-Fermi levels as defined below.

$$n = n_i e^{(E_{Fn} - E_i) / kT}$$

$$p = n_i e^{(E_i - E_{Fp}) / kT}$$



## Sample Problems

1\* Assuming low level injection, find the effective carrier life. (Sample doped with  $N_A = 10^{17} \text{ cm}^{-3}$ )

0.01s

(a) In the presence of Band-Band recombination only ( $k = 10^{16} \text{ cm}^{-3} \text{ s}^{-1}$ ).

10<sup>-4</sup>

(b) In the presence of SRH recombination only ( $\tau_n = \tau_p = 100 \text{ ns}$ ).

(c) In the presence of both SRH & band-band recombination

2\* The above sample is (in condition is specified in part c) is subjected to optical illumination which leads to  $G_{ex} = 10^{18} \text{ cm}^{-3} \text{ s}^{-1}$ .

10<sup>-17</sup>

(\*) Assume low level injection, what would be the steady state carrier density.

(\*) Is the assumption of low level injection valid?

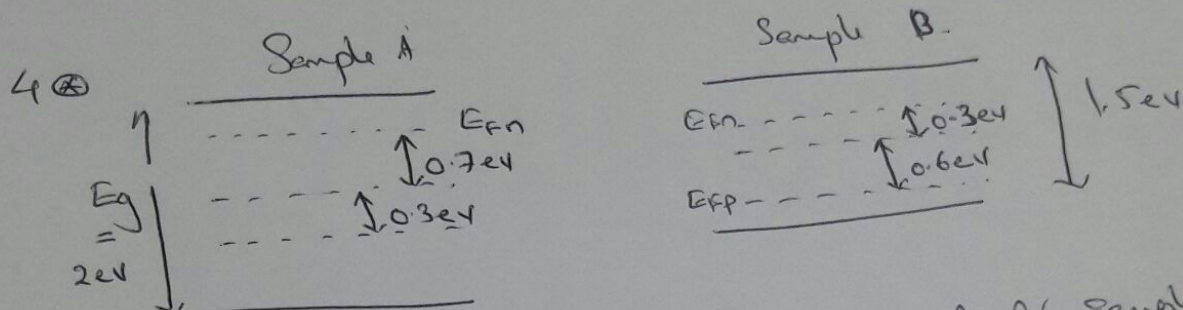
(\*) If  $G_{ex} = 10^{21} \text{ cm}^{-3} \text{ s}^{-1}$ , comment on the assumption of low level injection. If not valid, what would be the carrier density?

(Hint → You might have to start from the basic rate equation of B-B & SRH processes).

yes as  $\Delta n \ll n$ 2.6\*10<sup>-17</sup>



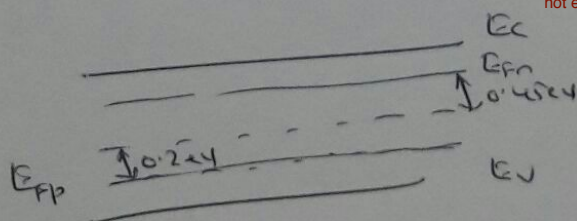
3\* Find  $E_{Fn}$  &  $E_{Fp}$  for the two cases discussed in Question 2.



Consider the above two semiconductor samples (A) & (B). Assume ~~the~~  $N_c$ ,  $N_v$  for them to be the same as that of Si and (at  $T = 300 \text{ K}$ ) and that they are temperature independent.

- \* Find the carrier densities in both the samples.
- \* Find the doping densities in both the samples.
- \* Which of them is under higher intensity of illumination?

5\* Without using calculator or any calculations, figure out whether the below sample is under low level illumination?



low level as  $E_{Fp}$  and  $E_{Fn}$  are not equally separated from  $E_i$