

SOLUTION 6

The aim of this exercise was to understand the following equations:

$$dr_a = c(E_t) \cdot g(E) f(E) \cdot N_t [1 - f(E_t)] dE \quad (1)$$

$$dr_b = e(E_t) \cdot N_t f(E_t) \cdot g(E) [1 - f(E)] dE \quad \text{with} \quad (2)$$

$$f(E) = \frac{1}{1 + e^{\frac{E - E_F}{kT}}} \quad (3)$$

Please be aware that we consider the system to be in thermal equilibrium here!

- a)
- E_t is the energy level of the traps – in the SRH-model all traps are considered to be at the same energy level.
 - E_F is the energy of the Fermi level.
 - E is the energy of electron states in the conduction band, i.e. it can lie in the range between the conduction band edge E_c and infinity.
 - g is the density of electron states of the conduction band, i.e. the number of electron states per volume and energy.
 - f is the occupation function which describes the probability that an electron state is occupied.
 - N_t is the constant trap- or recombination center density.
 - c and e are capture and emission coefficients, respectively: The whole physics of the capture and emission processes are considered therein. Multiplied with a density they give a capture and emission rate. The relationship between the capture cross section σ_c and c may also help for the understanding: $c = v_{th} \cdot \sigma_c$ with v_{th} the thermal velocity of electrons.
 - r_a / r_b are the rates of electrons that are captured in / emitted from traps of the conduction band tail and contribute therefore to the Shockley-Read-Hall recombination. They depend on the (fixed) energy of the traps. To get the net SRH-recombination rate one has to integrate $dr_a - dr_b$ over the whole energy range of the conduction band.
- b) The units of the parameters of the equations (1) to (3) are the following (not in SI system, but as commonly used in photovoltaics):

$$[r_a] = [r_b] = \text{cm}^{-3} \text{s}^{-1} \quad (4)$$

$$[E] = [E_t] = [E_F] = \text{eV} \quad (5)$$

$$[c] = [e] = \text{cm}^3 \text{s}^{-1} \quad (6)$$

$$[g] = \text{cm}^{-3} \text{eV}^{-1} \quad (7)$$

$$[f] = 1 \quad (8)$$

$$[N_t] = \text{cm}^{-3} \quad (9)$$

c) As the net electron capture rate in equilibrium must vanish, it is

$$0 \stackrel{!}{=} dr_a - dr_b \quad (10)$$

$$= g(E)N_t \cdot \{c(E_t)f(E)[1 - f(E_t)] - \quad (11)$$

$$e(E_t)f(E_t)[1 - f(E)]\} dE \quad (12)$$

$$\Rightarrow c(E_t)f(E)[1 - f(E_t)] = e(E_t)f(E_t)[1 - f(E)] \quad \forall E \quad (13)$$

$$\Rightarrow \frac{e(E_t)}{c(E_t)} = \frac{f(E) \cdot [1 - f(E_t)]}{f(E_t) \cdot [1 - f(E)]} \quad (14)$$

$$\stackrel{(*)}{=} \frac{f(E) \cdot e^{\frac{E_t - E_F}{kT}} \cdot f(E_t)}{f(E_t) \cdot e^{\frac{E - E_F}{kT}} \cdot f(E)} \quad (15)$$

$$= e^{\frac{E_t - E_F - E + E_F}{kT}} \quad (16)$$

$$= e^{\frac{E_t - E}{kT}}. \quad (17)$$

where in (*) has been used that

$$f(E) = \frac{1}{1 + e^{\frac{E - E_F}{kT}}} \Rightarrow 1 - f(E) = \frac{1 + e^{\frac{E - E_F}{kT}} - 1}{1 + e^{\frac{E - E_F}{kT}}} = e^{\frac{E - E_F}{kT}} \cdot f(E). \quad (18)$$