

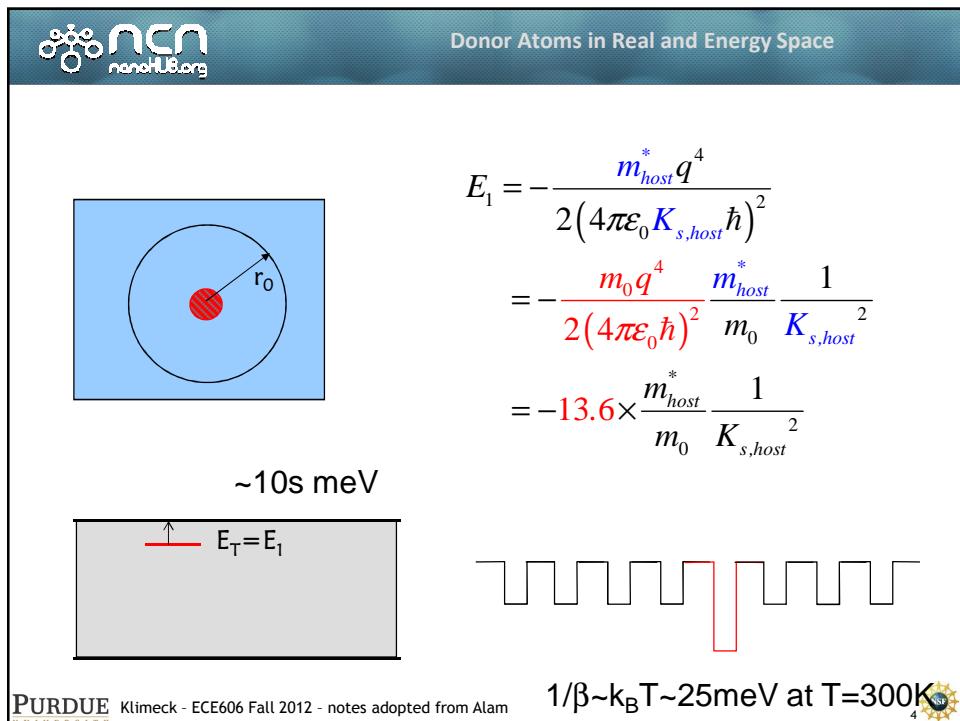
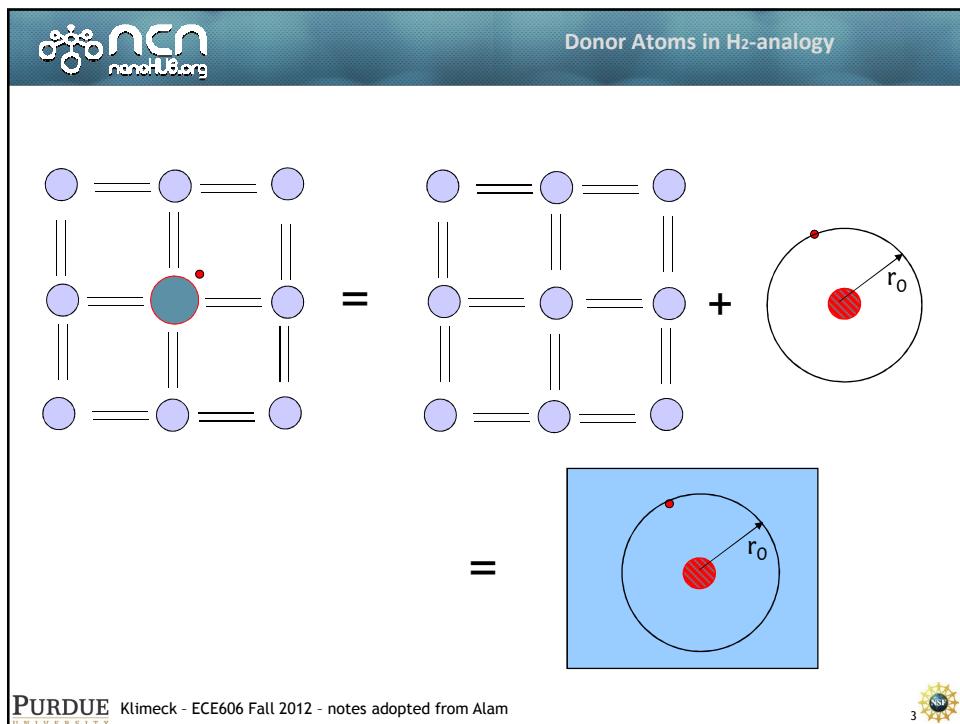
ECE606: Solid State Devices

Lecture 8

Gerhard Klimeck
gekco@purdue.edu

- Reminder:
 - »Basic concepts of donors and acceptors
 - »Statistics of donors and acceptor levels
 - »Intrinsic carrier concentration
- Temperature dependence of carrier concentration
- Multiple doping, co-doping, and heavy-doping
- Conclusion

Reference: Vol. 6, Ch. 4



How to Read the Table ...

	Li	Sb	P	As	S	Se	Tc	Cu	Ag	Ag
Ge	.0093	.0096	.012	.013	.18	.14	.11	.12	.04	.09
	GAP CENTER	—	—	—	—	—	—	.1	.2	.2
	.01	.01	.01	.011	.011	.06	.095	.21	.1	.13
B	Li	Al	Tl	Ga	In	Be	Zn	Cr	Cd	Hg
Si	.033	.019	.045	.054	.065	.14	.11	.14	.21	.22
	GAP CENTER	—	—	—	—	—	—	.25	.25	.17
	.3	.34	.35	.42	.35	.42	.32	.26	.26	.16
B	Al	Ga	In	Tl	Pd	Na	Be	Zn	Au	Co
Si	.045	.067	.072	.16	.045	.067	.072	.16	.28	.27
	GAP CENTER	—	—	—	—	—	—	.35	.33	.31
	.33	.33	.33	.36	.33	.36	.36	.36	.36	.36
Ge	.0058	.006	.006	.006	.006	.006	.006	.03	.0039	.4
	GAP CENTER	—	—	—	—	—	—	.67	.63	.63
	.44	.52	.53	.37	.44	.52	.53	.37	.37	.37
C	.026	.028	.028	.031	.035	.035	.023	.04	.023	.023
B	Mg	Zn	Si	Cd	Li	Ge	Au	Mn	Ag	Pb
Sn	.05	.07	.07	.04	.05	.05	.09	.11	.12	.16
	Ni	Cu	Fe	Cr						

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Summary ...

A bulk material must be charge neutral over all ...	
$\int [p - n + N_D^+ + N_A^-] dV = 0$	
Further if the material is <i>spatially homogenous</i>	
$p - n + N_D^+ + N_A^- = 0$ $N_V e^{-(E_F - E_V)/k_B T} - N_A e^{-(E_C - E_F)/k_B T} + \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} - \frac{N_A}{1 + 4e^{(E_A - E_F)/k_B T}} = 0$	

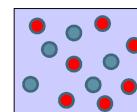
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A bulk material must be charge neutral over all ...

$$\int [p - n + N_D^+ + N_A^-] dV = 0$$



Further if the doping is spatially homogenous

$$p - n + N_D^+ + N_A^- = 0$$

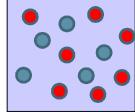
FD integral vs. FD function ?

$$N_V \frac{2}{\sqrt{\pi}} F_{1/2} [\beta(E_F - E_V)] - N_A \frac{2}{\sqrt{\pi}} F_{1/2} [\beta(E_C - E_F)] + \frac{N_D}{1 + 2e^{\beta(E_F - E_D)}} - \frac{N_A}{1 + 4e^{\beta(E_A - E_F)}} = 0$$

$$N_V e^{-(E_F - E_V)/k_B T} - N_A e^{-(E_C - E_F)/k_B T} + \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} - \frac{N_A}{1 + 4e^{(E_A - E_F)/k_B T}} = 0 \quad (\text{approx.})$$

Once you know E_F , you can calculate n , p , N_D^+ , N_A^- .

 Intrinsic Concentration

$$p - n + N_D^+ + N_A^- = 0$$


$$N_V e^{-(E_F - E_V)/k_B T} - N_C e^{-(E_C - E_F)/k_B T} + \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} - \frac{N_A}{1 + 4e^{(E_A - E_F)/k_B T}} = 0$$

$$n - p = 0 \Rightarrow N_C e^{-\beta(E_c - E_F)} = N_V e^{+\beta(E_v - E_F)}$$

$$E_F \equiv E_i = \frac{E_G}{2} + \frac{1}{2\beta} \ln \frac{N_V}{N_C}$$

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 Presentation Outline

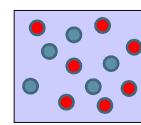
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In spatially homogenous field-free region ...

$$p - n + N_D^+ + N_A^- = 0$$

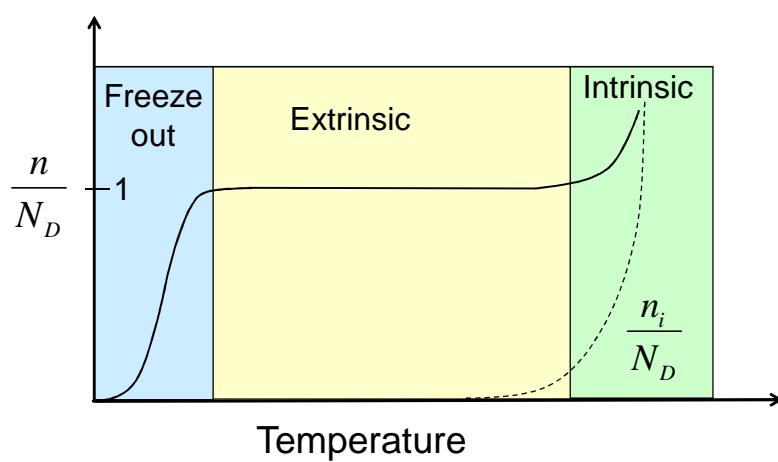


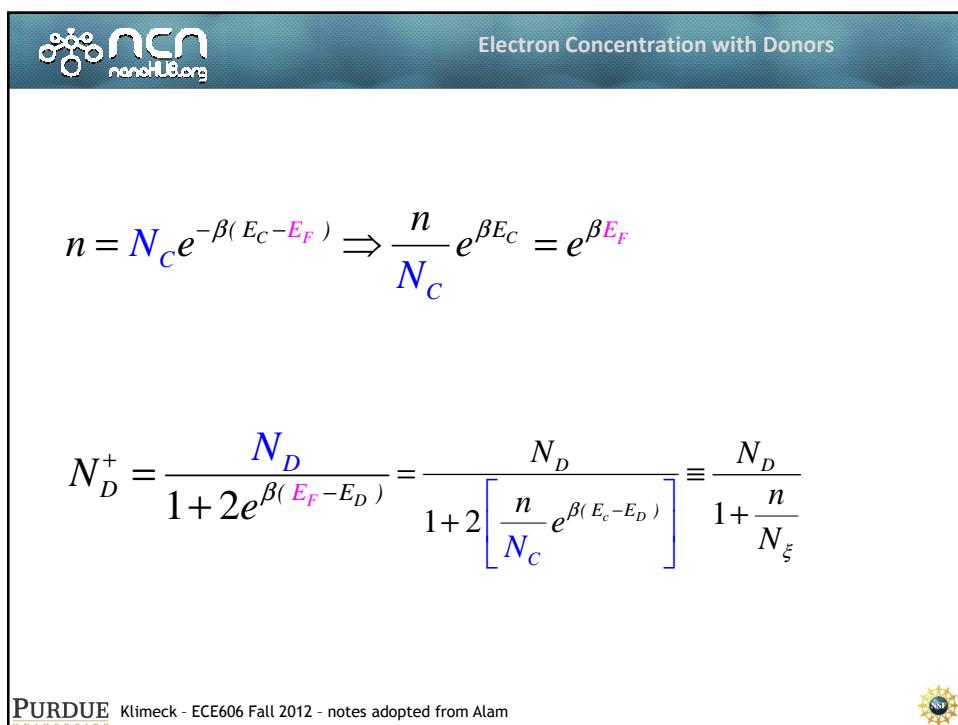
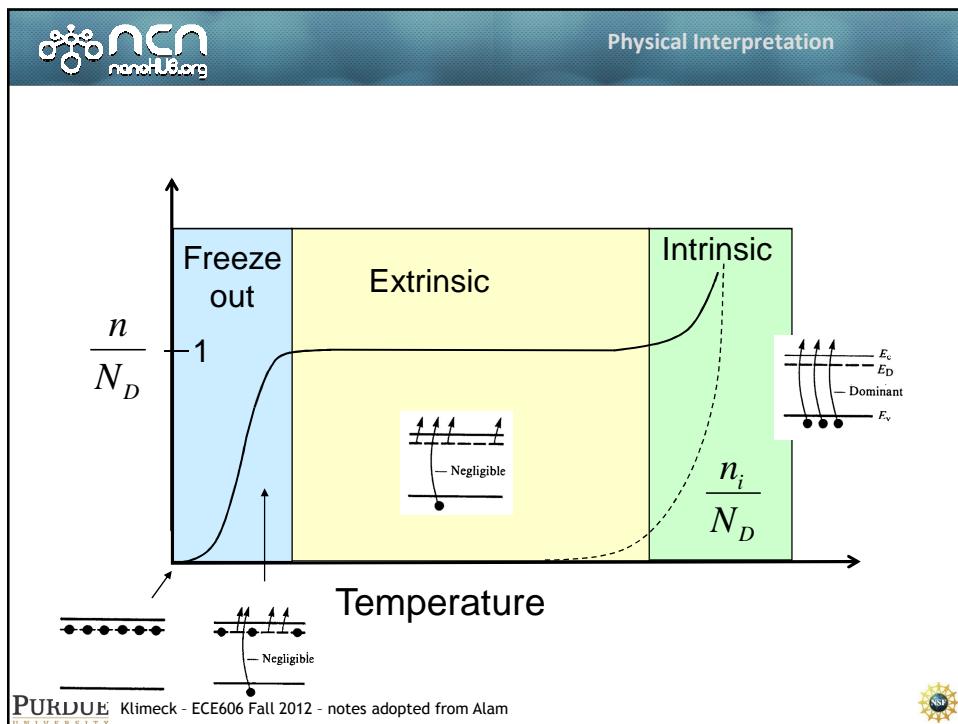
Assume
N-type doping ...

$$N_V e^{-(E_F - E_V)/k_B T} - \cancel{N_C} e^{-(E_C - E_F)/k_B T} + \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} - \cancel{\frac{N_A}{1 + 4e^{(E_A - E_F)/k_B T}}} = 0$$

n

(will plot in next slide)





$$\begin{array}{c}
 p - n + N_D^+ = 0 \\
 \downarrow \quad \downarrow \quad \searrow \\
 N_V e^{-(E_F - E_V)/k_B T} - N_C e^{-(E_C - E_F)/k_B T} + \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} = 0 \\
 \swarrow \quad \downarrow \quad \nearrow \\
 p \times n = n_i^2 \\
 \frac{n_i^2}{n} - \cancel{n} + \frac{N_D}{1 + \frac{n}{N_\xi}} = 0
 \end{array}$$

No approximation so far

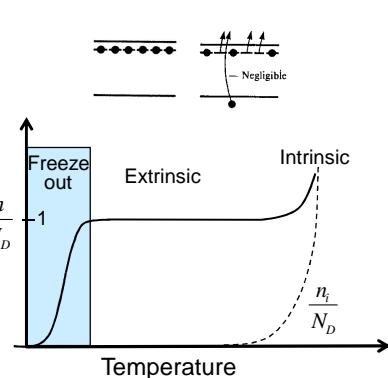
$$\frac{n_i^2}{n} - \cancel{n} + \frac{N_D}{1 + \frac{n}{N_\xi}} = 0$$

$$N_D \gg n_i$$

$$\Rightarrow -\cancel{n} + \frac{N_D}{1 + \frac{n}{N_\xi}} \approx 0$$

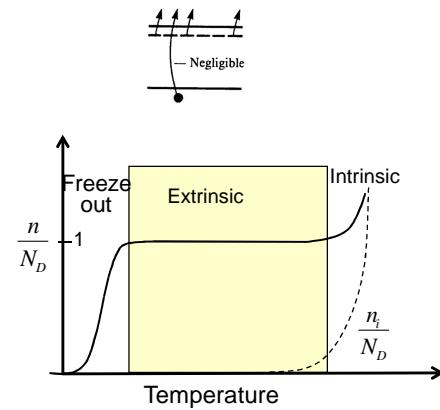
$$\Rightarrow \cancel{n}^2 + N_\xi \cancel{n} - N_\xi N_D = 0$$

$$N_\xi \equiv \frac{N_C}{2} e^{-\beta(E_C - E_D)} \quad \cancel{n} = \frac{N_\xi}{2} \left[\left(1 + \frac{4N_D}{N_\xi} \right)^{1/2} - 1 \right]$$



$$N_\xi \equiv \frac{N_C}{2} e^{-(E_C - E_D)/kT} \gg N_D$$

$$\begin{aligned} n &= \frac{N_\xi}{2} \left[\left(1 + \frac{4N_D}{N_\xi} \right)^{1/2} - 1 \right] \\ &\approx \frac{N_\xi}{2} \left[\left(1 + \frac{1}{2} \frac{4N_D}{N_\xi} \right) - 1 \right] \\ &\approx N_D \end{aligned}$$

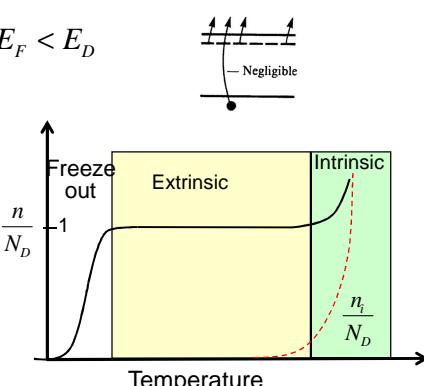


Electron concentration equals donor density
hole concentration by $n \times p = n_i^2$

$$N_D^+ = \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} \approx N_D \quad \text{for } E_F < E_D$$

$$\begin{aligned} \frac{n_i^2}{n} - n + \frac{N_D}{1 + \frac{n}{N_D}} &= 0 \\ \frac{n_i^2}{n} - n + N_D &\approx 0 \end{aligned}$$

$$\Rightarrow -n_i^2 + n^2 - N_D n = 0$$



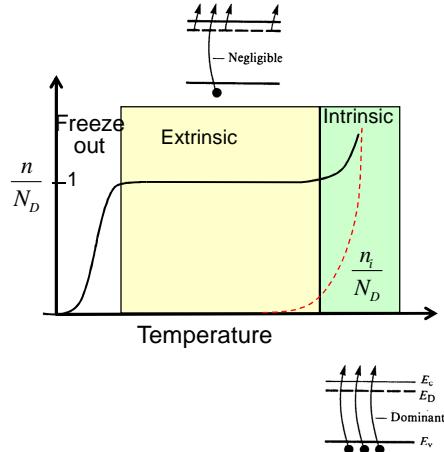
$$n = \frac{N_D}{2} + \left[\frac{N_D^2}{4} + n_i^2 \right]^{1/2}$$

For $N_D \gg n_i$

$$n = \frac{N_D}{2} + \left[\frac{N_D^2}{4} + n_i^2 \right]^{1/2} \approx N_D$$

For $n_i \gg N_D$

$$n = \frac{N_D}{2} + \left[\frac{N_D^2}{4} + n_i^2 \right]^{1/2} \approx n_i$$



What will happen if you use silicon circuits at very high temperatures ?
Bandgap determines the intrinsic carrier density.

$$n = N_c e^{-\beta(E_c - E_F)} \Rightarrow E_F = E_c + \frac{1}{\beta} \ln \left(\frac{n}{N_c} \right)$$

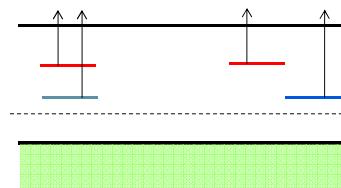
$$p - n + N_D^+ = 0$$

\swarrow \searrow \nearrow

$$N_V e^{-(E_F - E_V)/k_B T} - N_c e^{-(E_c - E_F)/k_B T} + \frac{N_D}{1 + 2e^{(E_F - E_D)/k_B T}} = 0$$

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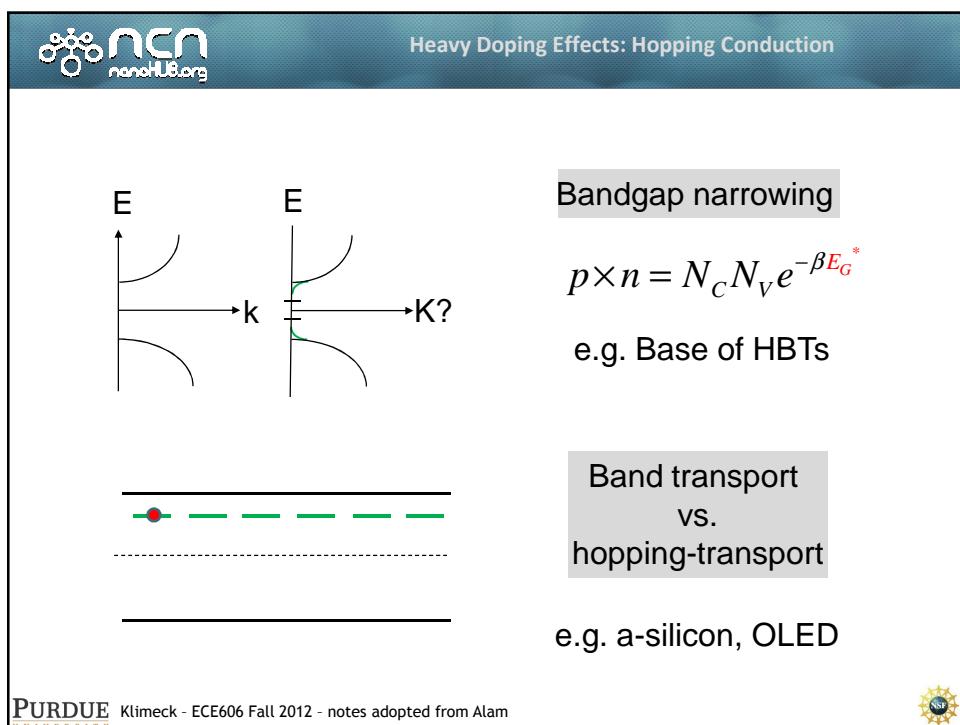
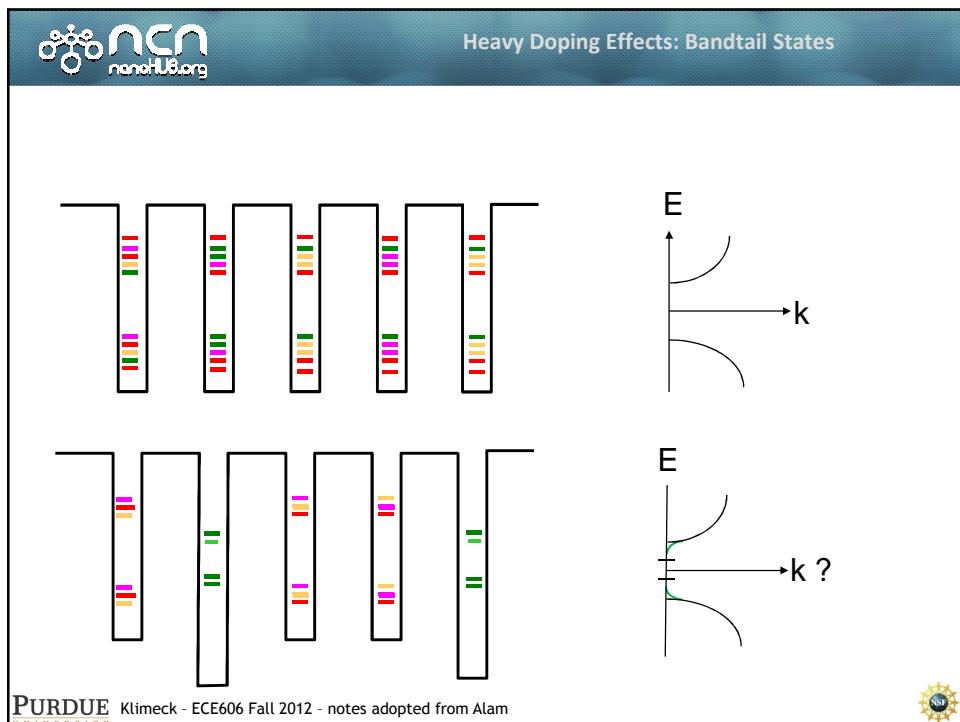


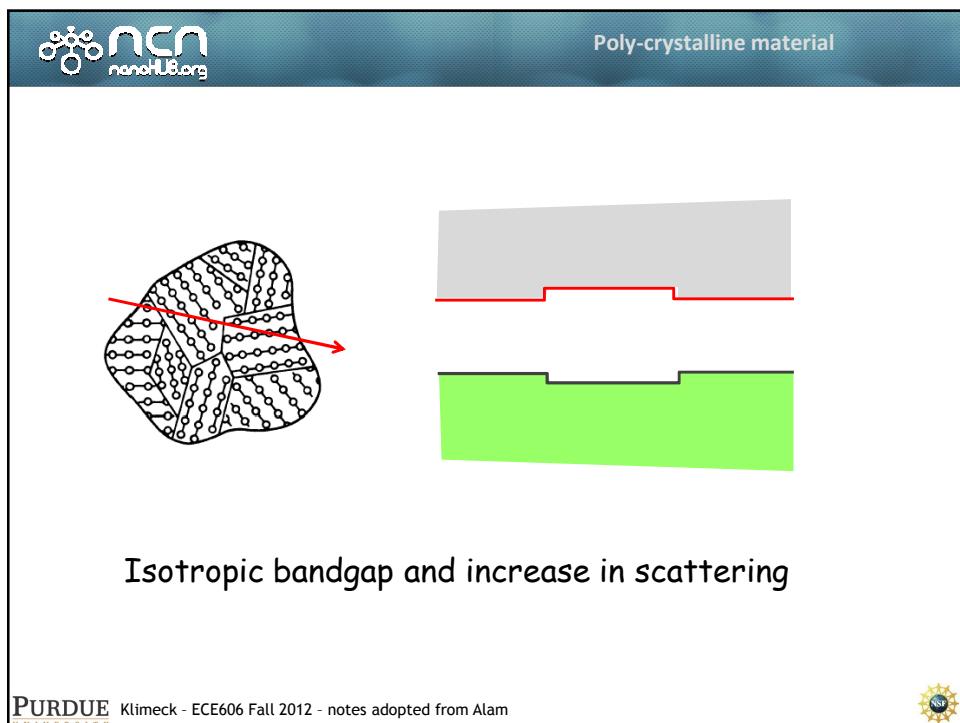
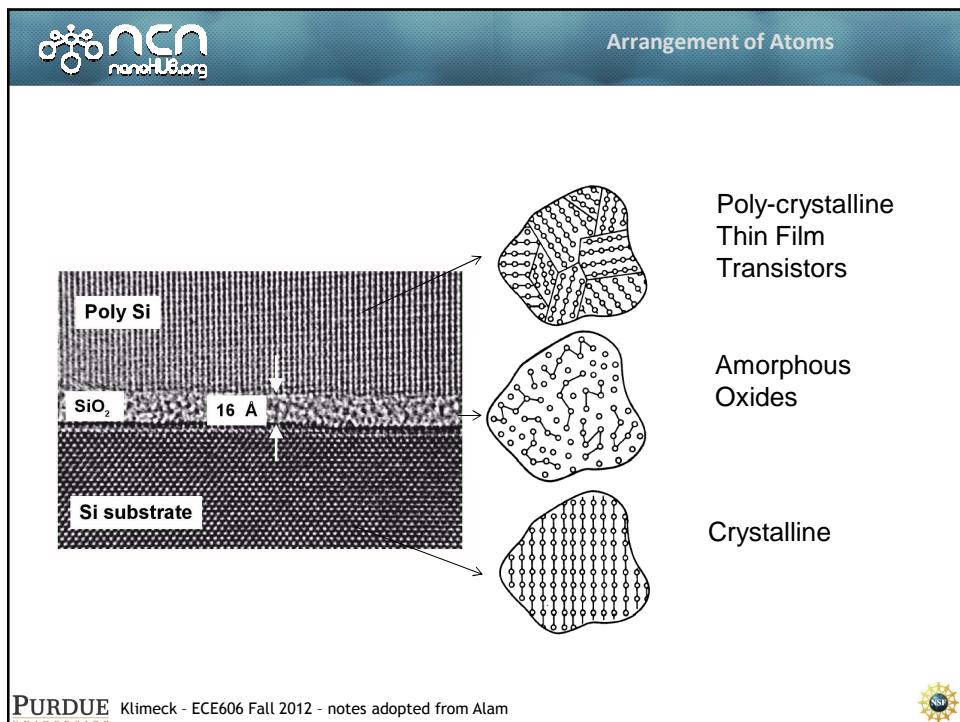
Multiple levels of same donor ...

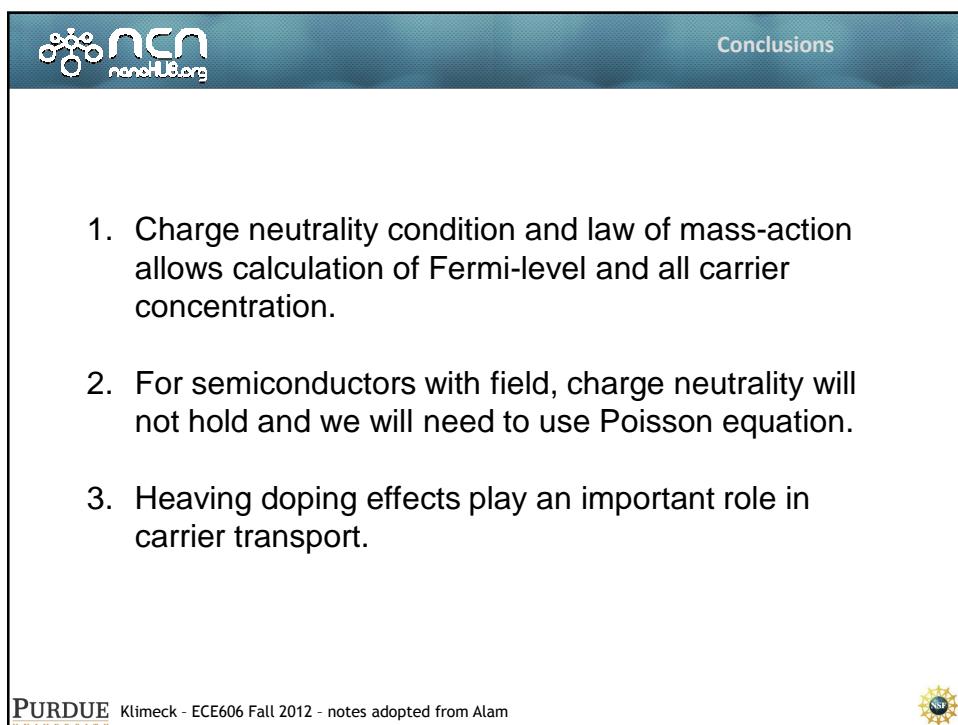
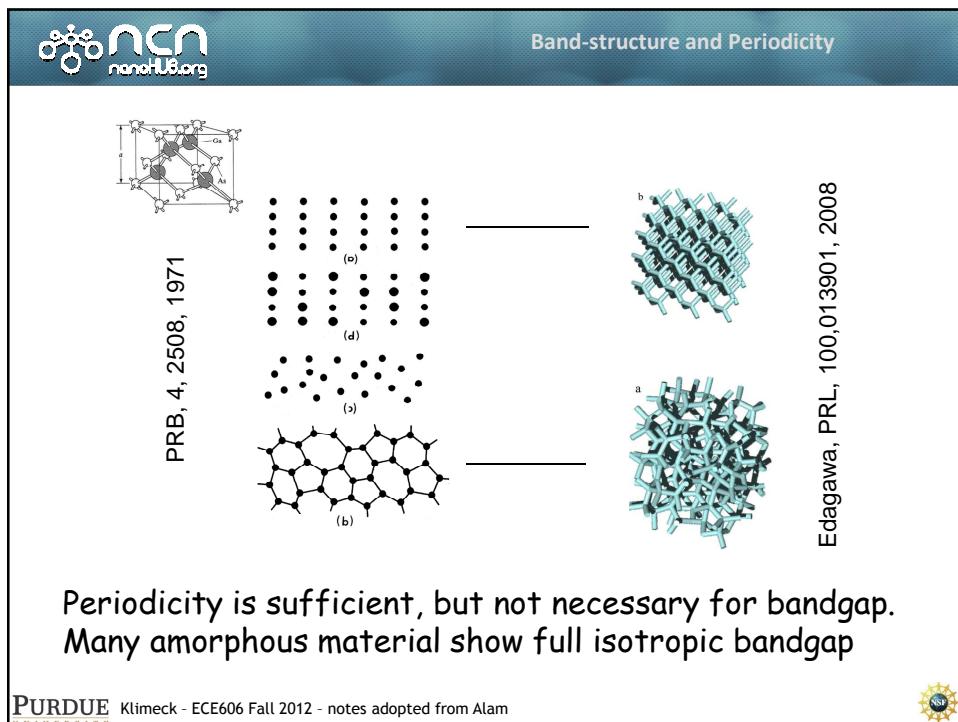
$$p - n + \frac{N_D}{1 + 2e^{(E_F - E_{D1})/k_B T}} + \frac{N_D}{1 + 2e^{(E_F - E_{D2})/k_B T}} - \frac{N_A}{1 + 4e^{(E_A - E_F)/k_B T}} = 0$$

Codoping...

$$p - n + \frac{N_{D1}}{1 + 2e^{(E_F - E_{D1})/k_B T}} + \frac{N_{D2}}{1 + 2e^{(E_F - E_{D2})/k_B T}} - \frac{N_A}{1 + 4e^{(E_A - E_F)/k_B T}} = 0$$



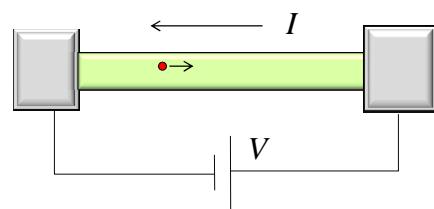




1) Non-equilibrium systems

- 2) Recombination generation events
- 3) Steady-state and transient response
- 4) Derivation of R-G formula
- 5) Conclusion

Ref. Chapter 5, pp. 134-146



$$I = G \times V$$

$$= q \times n \times v \times A$$

Carrier Density

velocity

Depends on chemical composition, crystal structure, temperature, doping, etc.

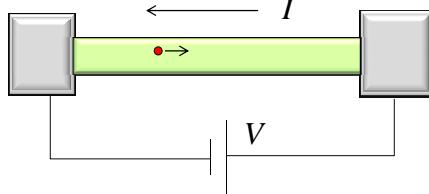
Quantum Mechanics + Equilibrium Statistical Mechanics

⇒ Encapsulated into concepts of effective masses and occupation factors (Ch. 1-4)

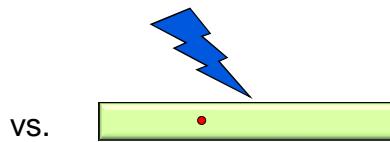
Transport with scattering, non-equilibrium Statistical Mechanics

⇒ Encapsulated into drift-diffusion equation with recombination-generation (Ch. 5 & 6)

Chapter 6



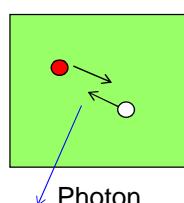
Chapter 5



vs.

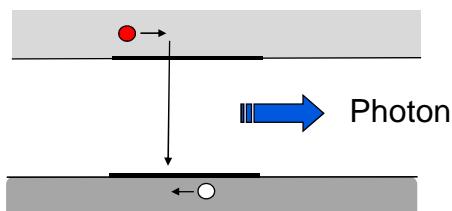
How does the system
go BACK to
equilibrium?

In real space ...



e and h must
have same wavelength
1 in 1,000,000 encounters

In energy space ...



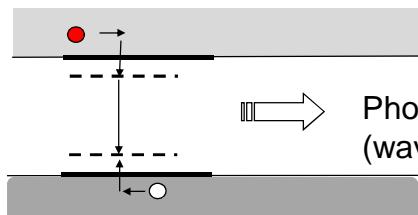
Direct transition –
direct gap material

GaAs, InP, InSb (3D)

Lasers, LEDs, etc.

Direct Excitonic Recombination

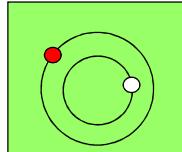
In energy space ...



Mostly in 1D systems
Requires strong coulomb interactions

Photon
(wavelength reduced from bulk)

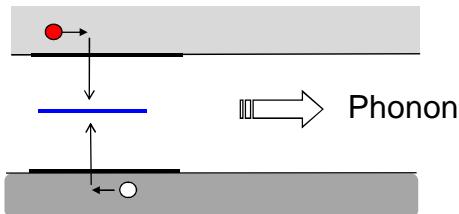
In real space ...



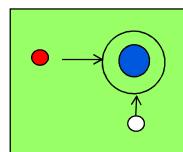
CNT, InP, 1D-systems

Transistors, Lasers, Solar cells, etc.

Indirect Recombination (Trap-assisted)



Trap needs to be mid-gap to be effective.
Cu or Au in Si



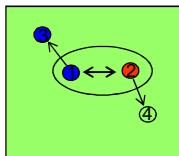
Ge, Si,

Transistors, Solar cells, etc.

Auger Recombination



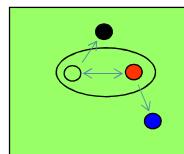
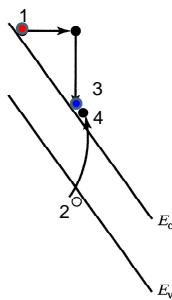
Requires very high electron density



InP, GaAs, ...

Lasers, etc.

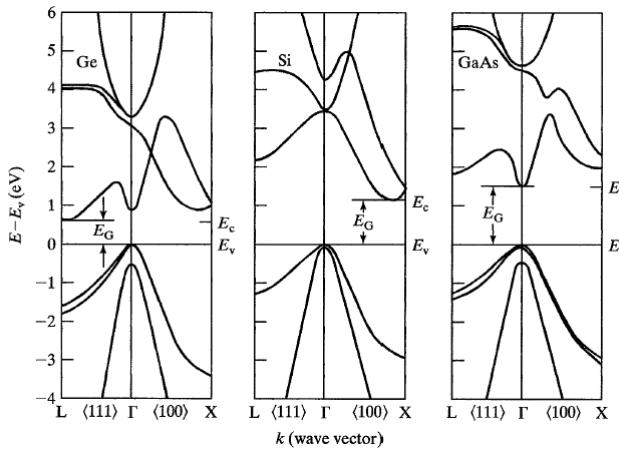
Impact Ionization - A Generation Mechanism



Si, Ge, InP

Lasers, Transistors, etc.

Indirect vs. Direct Bandgap

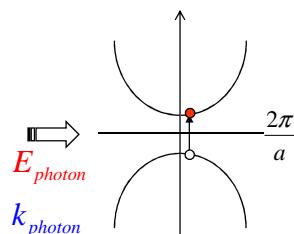


The top & bottom of bands do not align at same wavevector k for indirect bandgap material

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Photon Energy and Wavevector



$$E_V + E_{\text{photon}} = E_C$$

$$\hbar k_V + \hbar k_{\text{photon}} = \hbar k_C$$

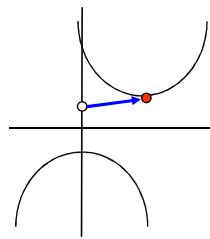
$$k_{\text{photon}} = \frac{2\pi}{\lambda \text{ in } \mu\text{m}} = \frac{2\pi}{1.21/E_{\text{photon}} \text{ in eV}}$$

$$<< \frac{2\pi}{a} = \frac{2\pi}{5 \times 10^{-4} \mu\text{m}}$$

Photon has large energy for excitation through bandgap, but its wavevector is negligible compared to size of BZ

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$$E_V + E_{phonon} = E_C$$

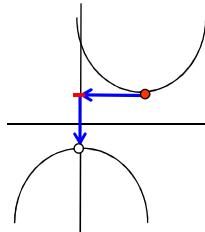
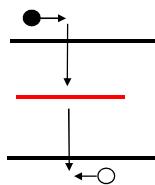
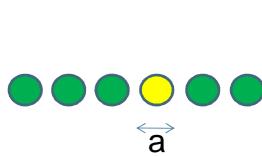
$$\hbar k_V + \hbar k_{phonon} = \hbar k_C$$

$$v_{sound} \sim 10^3 \text{ m/s} \ll v_{light} = c \sim 10^6 \text{ m/s}$$

$$\lambda^{sound} \gg \lambda^{light}$$

$$k_{phonon} = \frac{2\pi}{\lambda} = \frac{2\pi}{\hbar v_{sound} / E_{phonon}} \approx \frac{2\pi}{a} = \frac{2\pi}{5 \times 10^{-4} \mu\text{m}}$$

Phonon has large wavevector comparable to BZ,
but negligible energy compared to bandgap

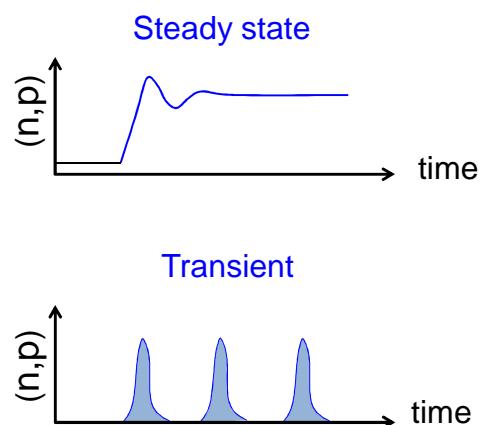
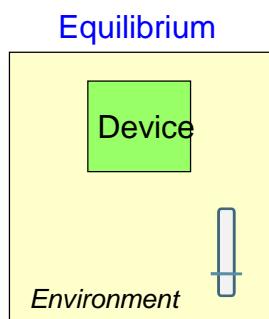


$$k_{trap} \approx \frac{2\pi}{a} \sim \frac{2\pi}{5 \times 10^{-4} \mu\text{m}}$$

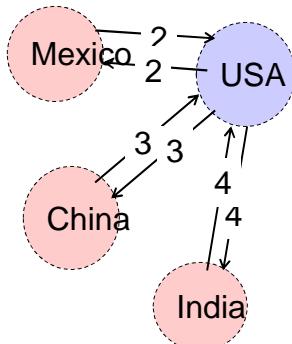
Trap provides the wavevector
necessary for indirect transition

- 1) Non-equilibrium systems
- 2) Recombination generation events
- 3) Steady-state and transient response**
- 4) Derivation of R-G formula
- 5) Conclusion

Ref. Chapter 5, pp. 134-146



Detailed Balance: Simple Explanation



The rates of exchange of people (particles) between every pair of countries (energy levels) is balanced. Hence the name “Detailed Balance”.

Detailed balance is the property of equilibrium

The population of each of the countries (energy levels) remains constant under detailed balance.

The concept of detailed balance is powerful, because it can be used for many things (e.g. reduce the number of unknown rate constants by half, and derive particle distributions like Fermi-Dirac, Bose-Einstein distributions, etc.)

Equilibrium is a very active place

- 9 in & 9 out
- All numbers are people/unit time.

Fermi-Dirac distribution demands exploration of allowed states

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Steady-state Response

Disturbing the detailed balance requires non-equilibrium conditions (needs energy). Unidirectional forces (red lines) can create such Non-equilibrium conditions.

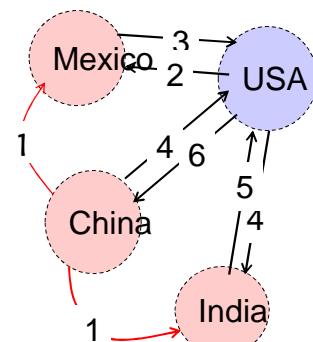
The rates of exchange of people (particles) between every pair of countries (energy levels) is NOT balanced, but the sum of all arrival and departures to all countries is zero.

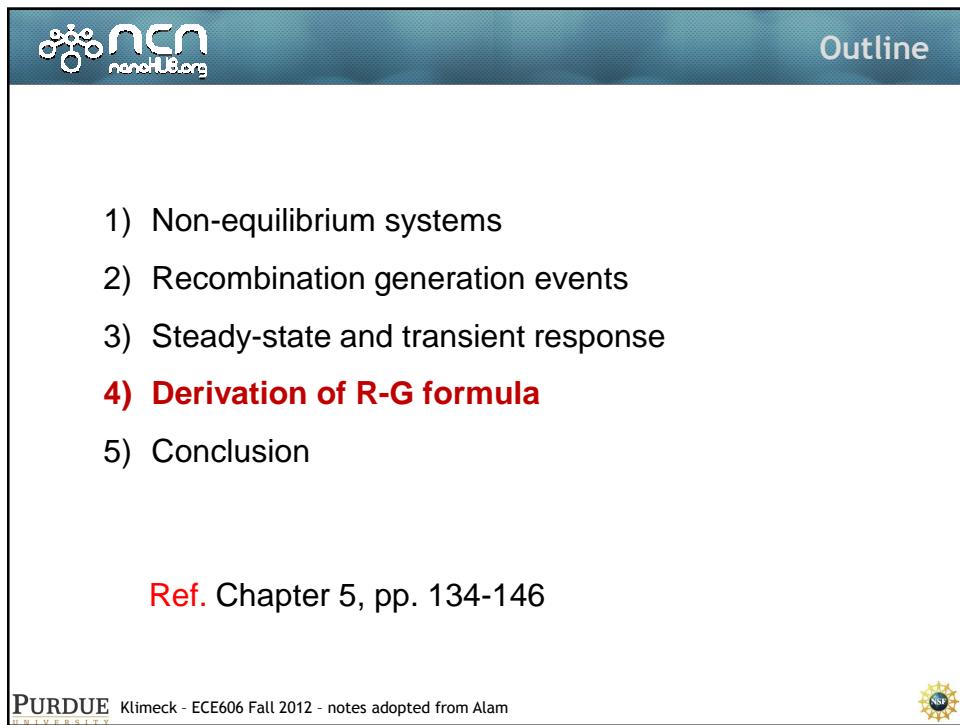
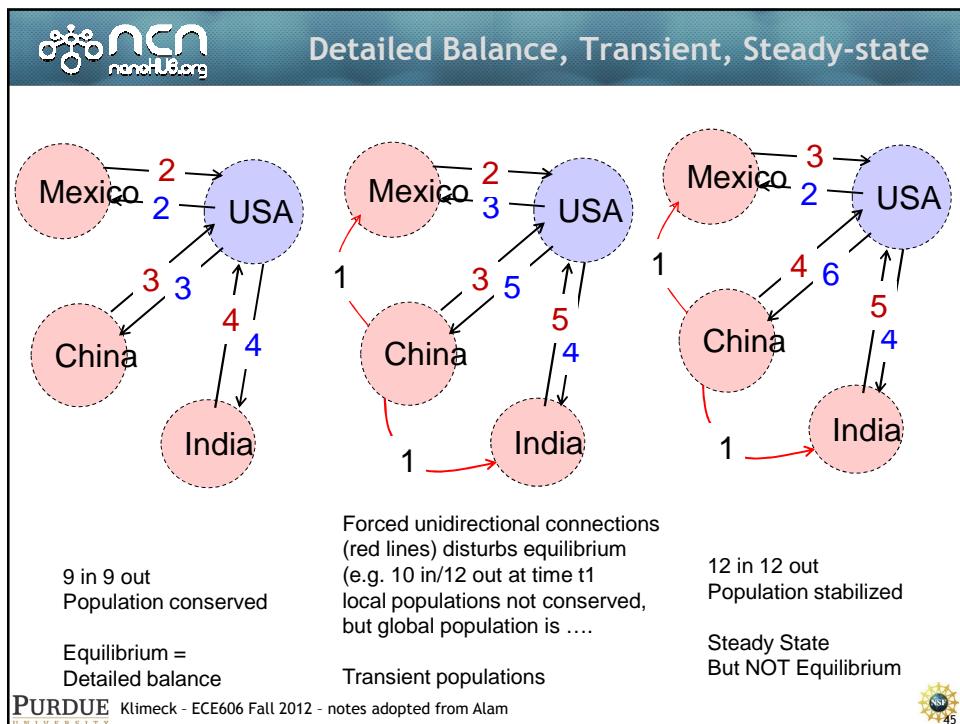
The flux at steady state is balanced overall, but the flux is NOT the same as in detailed balance (e.g. 12 in and 12 out in SS vs. 9 in and 9 out for Detailed Balance, for example).

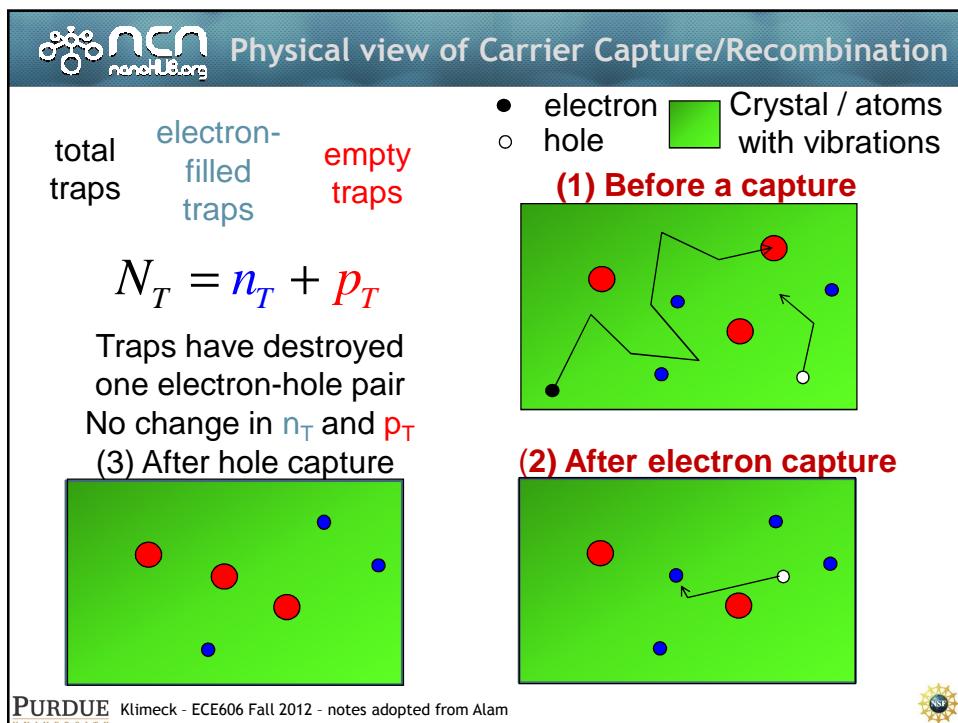
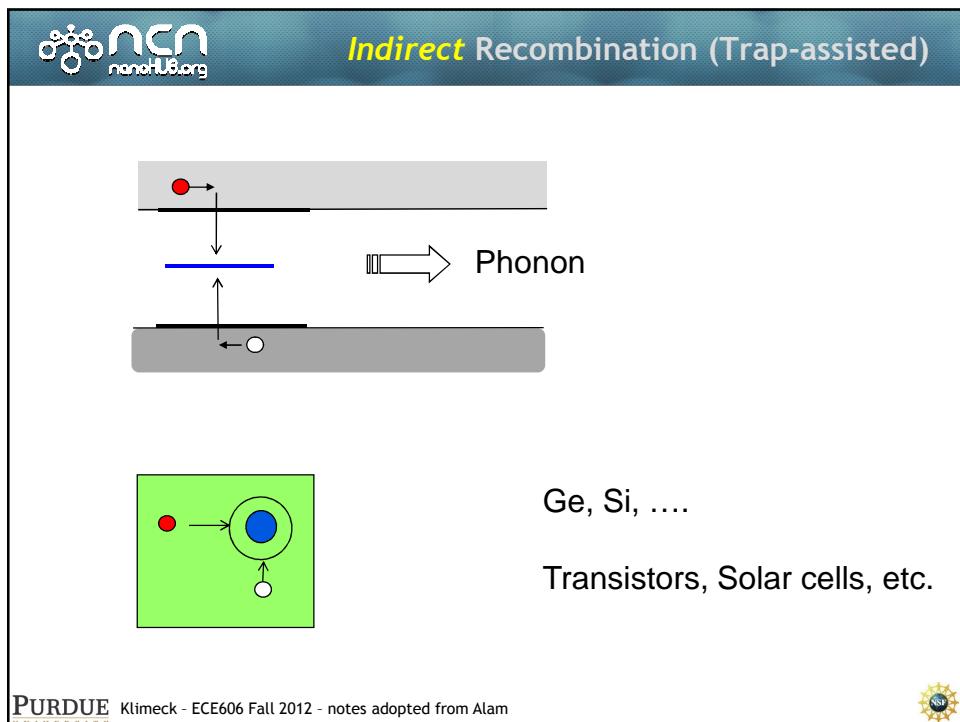
The population of a country (energy level) remains constant with time after steady state is reached.

One can use the requirement that net flux at steady state be zero to calculate steady state population of a country (Eq. 5.21)

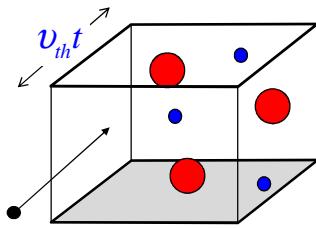
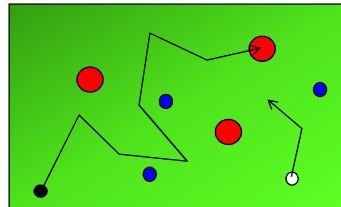
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Carrier Capture Coefficients



$$\frac{dn}{dt} = -n \times \left[\frac{\text{Volume} \times p_T \times \text{RelArea}}{\text{TotalArea} \times t} \right]$$

$$\frac{1}{2} m^* v_{th}^2 = \frac{3}{2} kT$$

$$v_{th} \approx 10^7 \frac{cm}{s}$$

$$\frac{dn}{dt} = -n \times \left[\frac{A \times v_{th} t \times p_T \times \sigma_n}{A \times t} \right]$$

$$\equiv -c_n p_T n \quad c_n \equiv \sigma_n v_{th}$$

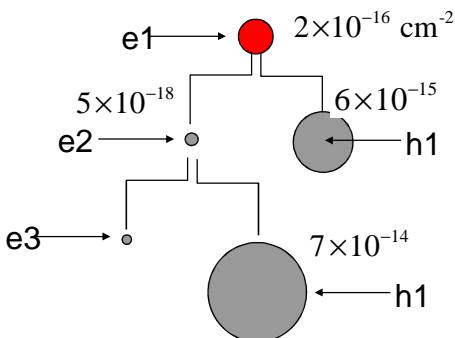
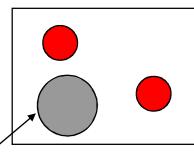
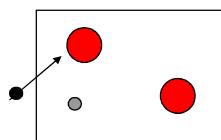
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Capture Cross-section

$$\sigma_n = \pi r_0^2$$

Zn capture model ...



Cascade model for capture

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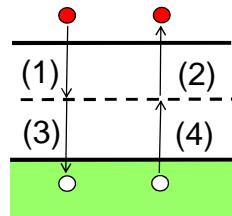


- 1) There are wide variety of generation-recombination events that allow restoration of equilibrium once the stimulus is removed.
- 2) Direct recombination is photon-assisted, indirect recombination phonon assisted.
- 3) Concepts of equilibrium, steady state, and transient dynamics should be clearly understood.

- 1) **Derivation of SRH formula**
- 2) Application of SRH formula for special cases
- 3) Direct and Auger recombination
- 4) Conclusion

Ref. ADF, Chapter 5, pp. 141-154

Sub-processes of SRH Recombination

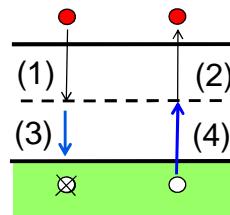


(1)+(3): one electron reduced from Conduction-band & one-hole reduced from valence-band

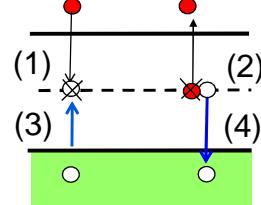
(2)+(4): one hole created in valence band and one electron created in conduction band

SRH Recombination

Physical picture



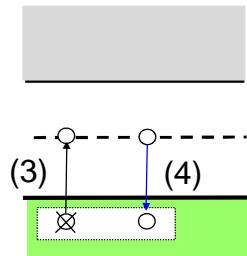
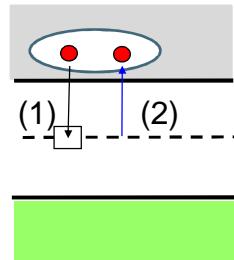
Equivalent picture



(1)+(3): **one electron reduced from C-band & one-hole reduced from valence-band**

(2)+(4): **one hole created in valence band & one electron created in conduction band**

Changes in electron and hole Densities



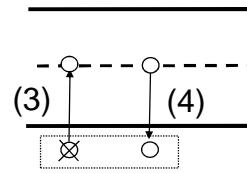
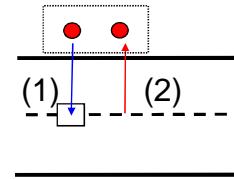
$$\frac{\partial n}{\partial t} \Big|_{1,2} = -c_n n p_T + e_n n_T (1 - f_c)$$

$$\frac{\partial p}{\partial t} \Big|_{3,4} = -c_p p n_T + e_p p_T f_v$$

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Detailed Balance in *Equilibrium*



$(1 - f_c) \approx 1$ Assume non-degenerate

$$\frac{\partial n}{\partial t} \Big|_{1,2} = -c_n n_0 p_{T0} + e_n n_{T0}$$

$$0 = -c_n n_0 p_{T0} + e_n n_{T0}$$

$$e_n = c_n \frac{n_0 p_{T0}}{n_{T0}} \equiv c_n n_1$$

$$0 = -c_n (n_0 p_{T0} - n_{T0} n_1)$$

$$\frac{\partial p}{\partial t} \Big|_{3,4} = -c_p p n_T + e_p p_T$$

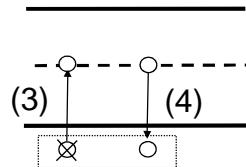
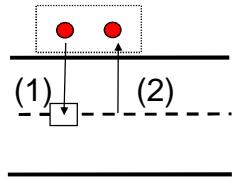
$$0 = -c_p p_0 n_{T0} + p_{T0} e_p$$

$$e_p \equiv \frac{c_p p_0 n_{T0}}{p_{T0}} = c_p p_1$$

$$0 = -c_p (p_0 n_{T0} - p_{T0} n_1)$$

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$$n_1 = \frac{n_0 p_{T0}}{n_{T0}}$$

$$p_1 = \frac{p_0 n_{T0}}{p_{T0}}$$

$$n_1 p_1 = \frac{n_0 p_{T0}}{n_{T0}} \times \frac{p_0 n_{T0}}{p_{T0}} = n_0 p_0 = n_i^2$$

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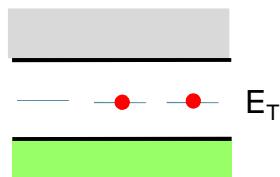


Trap is like a donor!

$$n_{T0} = N_T (1 - f_{00}) = \frac{N_T}{1 + g_D e^{\beta(E_T - E_F)}}$$

$$(1 - f_{00}) = \frac{1}{1 + g \exp} \quad f_{00} = 1 - \frac{1}{1 + g \exp}$$

$$f_{00} = \frac{g \exp}{1 + g \exp} \quad \cancel{f_{00}/(1 - f_{00})} = \frac{g \exp}{1 + g \exp} / \frac{1}{1 + g \exp} = g \exp$$



$$n_1 = \frac{n_0 p_{T0}}{n_{T0}} = n_0 \frac{(N_T f_{00})}{N_T (1 - f_{00})}$$

$$p_1 n_1 = n_i^2$$

$$n_1 = n_i e^{\beta(E_F - E_i)} g_D e^{\beta(E_T - E_F)}$$

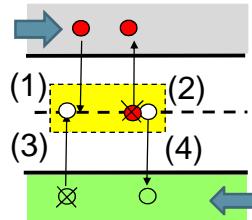
$$= n_i g_D e^{\beta(E_T - E_i)}$$

$$p_1 = n_i^2 / n_1$$

$$= n_i g_D^{-1} e^{\beta(E_i - E_T)}$$

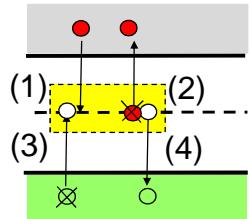
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$$\begin{aligned}
 \frac{\partial n_T}{\partial t} &= -\left. \frac{\partial n}{\partial t} \right|_{1,2} + \left. \frac{\partial p}{\partial t} \right|_{3,4} \\
 &= c_n n p_T - e_n n_T - c_p p n_T + e_p p_T \\
 &= c_n (n p_T - n_T n_1) - c_p (p n_T - p_T p_1)
 \end{aligned}$$

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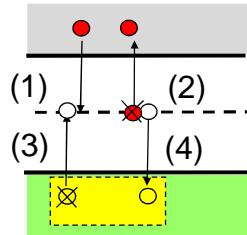


$$\frac{\partial n_T}{\partial t} = 0 = c_n (n p_T - n_T n_1) - c_p (p n_T - p_T p_1)$$

$$n_T = \frac{c_n N_T n + c_p N_T p_1}{c_n (n + n_1) + c_p (p + p_1)} = c_n (n p_T - n_T n_1)$$

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$$\begin{aligned}
 R = -\frac{dp}{dt} &= c_p (p n_T - p_T p_1) \\
 &= \frac{np - n_i^2}{\left(\frac{1}{c_p N_T}\right)(n + n_1) + \left(\frac{1}{c_n N_T}\right)(p + p_1)}
 \end{aligned}$$

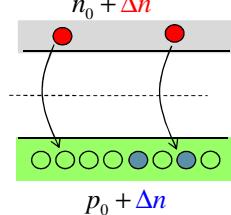
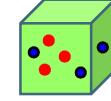
τ_n ↗ ↗ τ_p

- 1) Derivation of SRH formula
- 2) **Application of SRH formula for special cases**
- 3) Direct and Auger recombination
- 4) Conclusion

Case 1: Low-level Injection in p-type

$$\begin{aligned}
 R &= \frac{np - n_i^2}{\tau_p(n + n_1) + \tau_n(p + p_1)} \\
 &= \frac{(n_0 + \Delta n)(p_0 + \Delta p) - n_i^2}{\tau_p(n_0 + \Delta n + n_1) + \tau_n(p_0 + \Delta p + p_1)} \\
 &= \frac{\Delta n(n_0 + p_0) + \cancel{\Delta n^2}}{\tau_p(n_0 + \Delta n + n_1) + \tau_n(p_0 + \Delta p + p_1)} \\
 &= \frac{\Delta n(p_0)}{\tau_n(p_0)} = \frac{\Delta n}{\tau_n} \quad \Delta n^2 \approx 0 \\
 &\qquad\qquad\qquad p_0 \gg \Delta n \gg n_0
 \end{aligned}$$

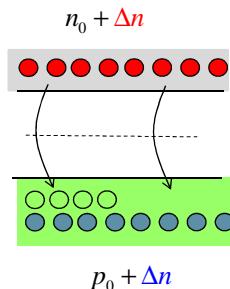
Lots of holes, few electrons => independent of holes



Case 2: High-level Injection

$$\begin{aligned}
 R &= \frac{np - n_i^2}{\tau_p(n + n_1) + \tau_n(p + p_1)} \quad \text{e.g. organic solar cells} \\
 &= \frac{(n_0 + \Delta n)(p_0 + \Delta p) - n_i^2}{\tau_p(n_0 + \Delta n + n_1) + \tau_n(p_0 + \Delta p + p_1)} \\
 &= \frac{\cancel{\Delta n}(n_0 + p_0) + \Delta n^2}{\tau_p(n_0 + \Delta n + n_1) + \tau_n(p_0 + \Delta p + p_1)} \\
 &= \frac{\Delta n^2}{(\tau_n + \tau_p)\Delta n} = \frac{\Delta n}{(\tau_n + \tau_p)} \quad \Delta n \gg p_0 \gg n_0
 \end{aligned}$$

Lots of holes, lots of electrons => dependent on both relaxations



$$R_{high} = \frac{\Delta n}{(\tau_n + \tau_p)} \quad \Delta n \gg p_0 \gg n_0$$

$$R_{low} = \frac{\Delta n}{\tau_p} \quad p_0 \gg \Delta n \gg n_0$$

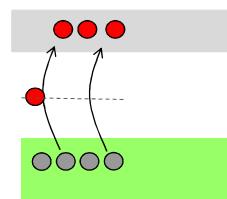
which one is larger and why?

Depletion region – in PN diode: $n=p=0$

$$n \ll n_1 \quad p \ll p_1$$

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

$$= \frac{-n_i^2}{\tau_p(n_1) + \tau_n(p_1)}$$



NEGATIVE Recombination => Generation

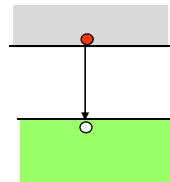
$n=p=0 \ll n_i \Rightarrow$ generation to create n,p
Equilibrium restoration!

- 1) Derivation of SRH formula
- 2) Application of SRH formula for special cases
- 3) Direct and Auger recombination**
- 4) Conclusion

$$R = B(np - n_i^2) \quad B \text{ is a material property}$$

Direct recombination at low-level injection

$$n_0 \ll (\Delta n = \Delta p) \ll p_0$$



$$R = B[(n_0 + \Delta n)(p_0 + \Delta p) - n_i^2] \approx Bp_0 \times \Delta n$$

Direct generation in depletion region

$$n, p \sim 0$$

$$R = B(np - n_i^2) \approx -Bn_i^2$$

Auger Recombination

2 electron & 1 hole

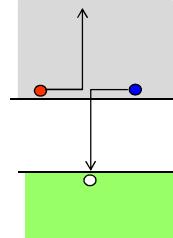
$$R = c_n (n^2 p - n_i^2 n) + c_p (np^2 - n_i^2 p)$$

$$c_n, c_p \sim 10^{-29} \text{ cm}^6/\text{sec}$$

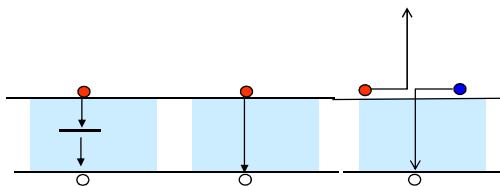
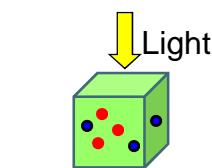
Auger recombination at low-level injection

$$n_0 \ll (\Delta n = \Delta p) \ll (p_0 = N_A)$$

$$R \approx c_p N_A^2 \Delta n = \frac{\Delta n}{\tau_{auger}} \quad \tau_{auger} = \frac{1}{c_p N_A^2}$$



Effective Carrier Lifetime

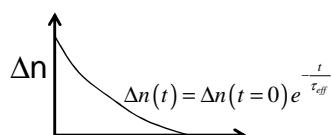


$$R = R_{SRH} + R_{direct} + R_{Auger}$$

$$= \Delta n \left(\frac{1}{\tau_{SRH}} + \frac{1}{\tau_{direct}} + \frac{1}{\tau_{Auger}} \right)$$

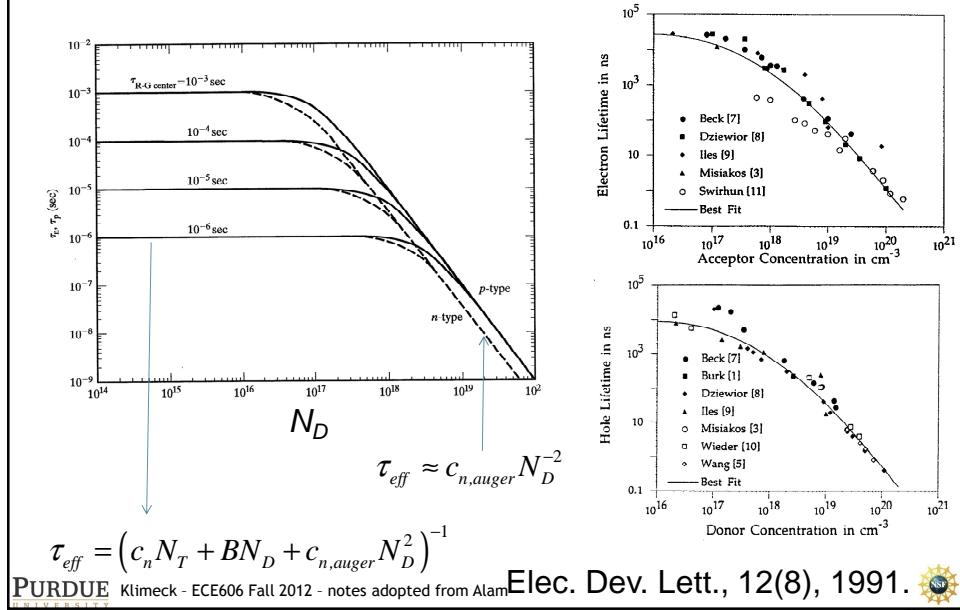
$$= \Delta n (c_n N_T + B N_D + c_{n,auger} N_D^2)$$

$$\tau_{eff} = (c_n N_T + B N_D + c_{n,auger} N_D^2)^{-1}$$



$$\Delta n(t) = \Delta n(t=0) e^{-\frac{t}{\tau_{eff}}}$$

Effective Carrier Lifetime with all Processes



Conclusion

SRH is an important recombination mechanism in important semiconductors like Si and Ge.

SRH formula is complicated, therefore simplification for special cases are often desired.

Direct band-to-band and Auger recombination can also be described with simple phenomenological formula.

These expressions for recombination events have been widely validated by measurements.

