

Studies on n-type silicon after electron irradiation

Roxana Radu^{a),b)}, Eckhart Fretwurst^{a)}, Robert Klanner^{a)}, Gunnar Lindström^{a)}, Ioana Pintilie^{b)}

a) Institute for Experimental Physics, University of Hamburg, Germany

b) National Institute of Material Physics, Magurele, Bucharest, Romania

Outline

- ◆ **Motivation**
- ◆ **Radiation-induced bulk damage**
- ◆ **Results on electron irradiation from 1.5 to 15 MeV**
- ◆ **Conclusions**

- I. Joint project between NIMP Bucharest and University of Hamburg –” Electron irradiation of Si-diodes at $E = 1 - 15$ MeV’** presented by Prof. G. Lindström at the last *WODEAN Workshop in Bucharest, 13/14 May 2010*

- II. ”Comprehensive investigation on bulk radiation damage in defect engineered silicon - from point defects to clusters”** –
Project Director Ioana Pintilie, 2011

Aim: Identification of both the structure of the electrically active defects responsible for the electrical properties of irradiated silicon diodes and the possible reactions with different impurities in the material.

1) Irradiation with 1.5, 3.5, 6 and 15 MeV electrons – studies performed in Hamburg

- Electrical characterization (CV/IV) before/after irradiation - V_{dep} and I_{dep} space charge distribution, recombination-generation current also after annealing (isothermal and isochronal)
- Analysis of electrically active defects by means of DLTS and TSC methods before/after irradiation - correlation with results from diode characteristics

2) Electron induced damage in Si implanted with ^{17}O and ^{13}C – investigations performed in Bucharest

- Studies for defect identification by Electron Paramagnetic Resonance (EPR, ENDOR) methods - in defect engineered silicon (O enriched, O lean, C rich, C lean)
- Microstructural investigation of the extended and clustered defects by High Resolution-Transmission Electron Microscopy (HRTEM) - Identify the structure of the radiation-induced electrically active defects and establish the role of the impurities in their generation and kinetics.

Particle type	Damage created
$^{60}\text{Co-}\gamma$ irradiation (1.1 and 1.3 MeV) $E_{\text{rec.max}} = 200 \text{ eV}$	<u>only point defects</u>
Reactor neutron irradiation (1 MeV) $E_{\text{rec.max}} \sim 50 \text{ keV}$	<u>dominant cluster defects</u>
High energy protons (23 GeV)	<u>both point and cluster defects</u>
Threshold for formation of point defects : $E_{\text{rec.}} \sim 21 \text{ eV}$ Threshold for formation of cluster defects : $E_{\text{rec.}} \sim 1.2 \text{ keV}$	

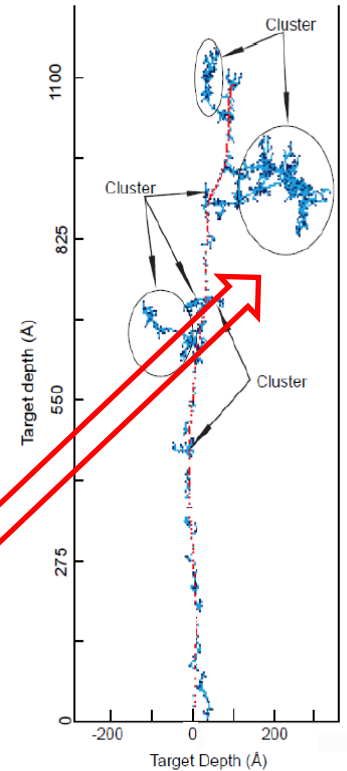
Irradiation with electrons, from low ($E_e=1,5 \text{ MeV}$) to higher energies ($E_e=15 \text{ MeV}$), in order to study the difference between radiation-induced point and cluster defects

$$T_{max} = \frac{2T_e(T_e + 2m_e c^2)}{M_0 c^2 A}$$

T_{max} = maximal energy transfer
 $T_e = E_{e,kin.}$ = electron kinetic energy

A. Akkerman et al. / Radiation Physics and Chemistry 62 (2001) 301–310

$E_{e,kin.}$	=	1 MeV ,	T_{max}	=	154 eV
$E_{e,kin.}$	=	3.5 MeV ,	T_{max}	=	1.2 keV
$E_{e,kin.}$	=	6 MeV ,	T_{max}	=	3.2 keV
$E_{e,kin.}$	=	15 MeV ,	T_{max}	=	18.3 keV
$E_{e,kin.}$	=	30 MeV ,	T_{max}	=	71 keV



New way to study the change from purely point to cluster-dominated effects

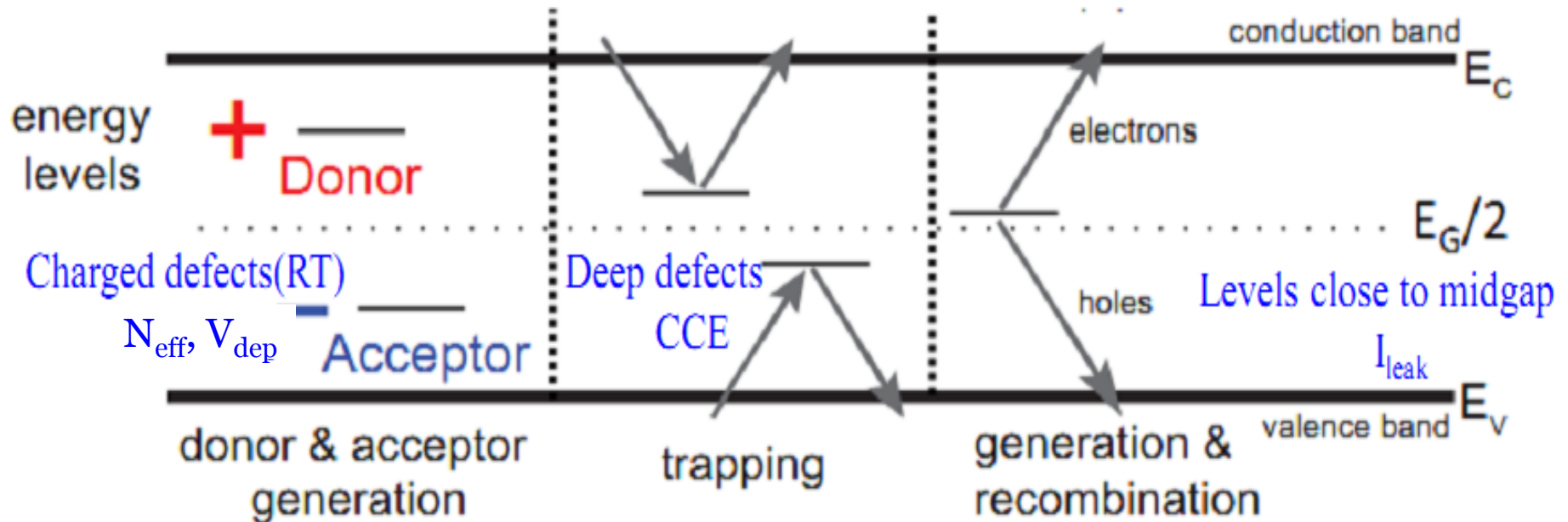
Materials and irradiation

Type	Orientation	d[μm]	$N_D[\text{cm}^{-3}]$	$\langle[O]\rangle[\text{cm}^{-3}]$	$\rho [\Omega\text{cm}]$	Diffusion oxygenated
EPI	$\langle 111 \rangle$	50	6×10^{13}	1×10^{17}	50	-
	$\langle 100 \rangle$	100	1×10^{13}	2.8×10^{17}	300	24 h /1100°C
ST-FZ	$\langle 100 \rangle$	280	8×10^{11}	1×10^{16}	5×10^3	-
DO-FZ	$\langle 100 \rangle$	280	8×10^{11}	1.2×10^{17}	5×10^3	72 h /1150°C

Electron fluences: $1 \times 10^{12} \rightarrow 1.5 \times 10^{15} \text{ cm}^{-2}$, $E_e = 1.5 - 15 \text{ MeV}$
 (for $E_e = 15 \text{ MeV}$: $\Phi_{\text{eq}} = 2.7 \times 10^{10} \rightarrow 4.1 \times 10^{13}$)

Hardness factor (k)	
1.5 MeV	1.88×10^{-3}
3.5 MeV	6.50×10^{-3}
6 MeV	1.24×10^{-2}
15 MeV	2.78×10^{-2}

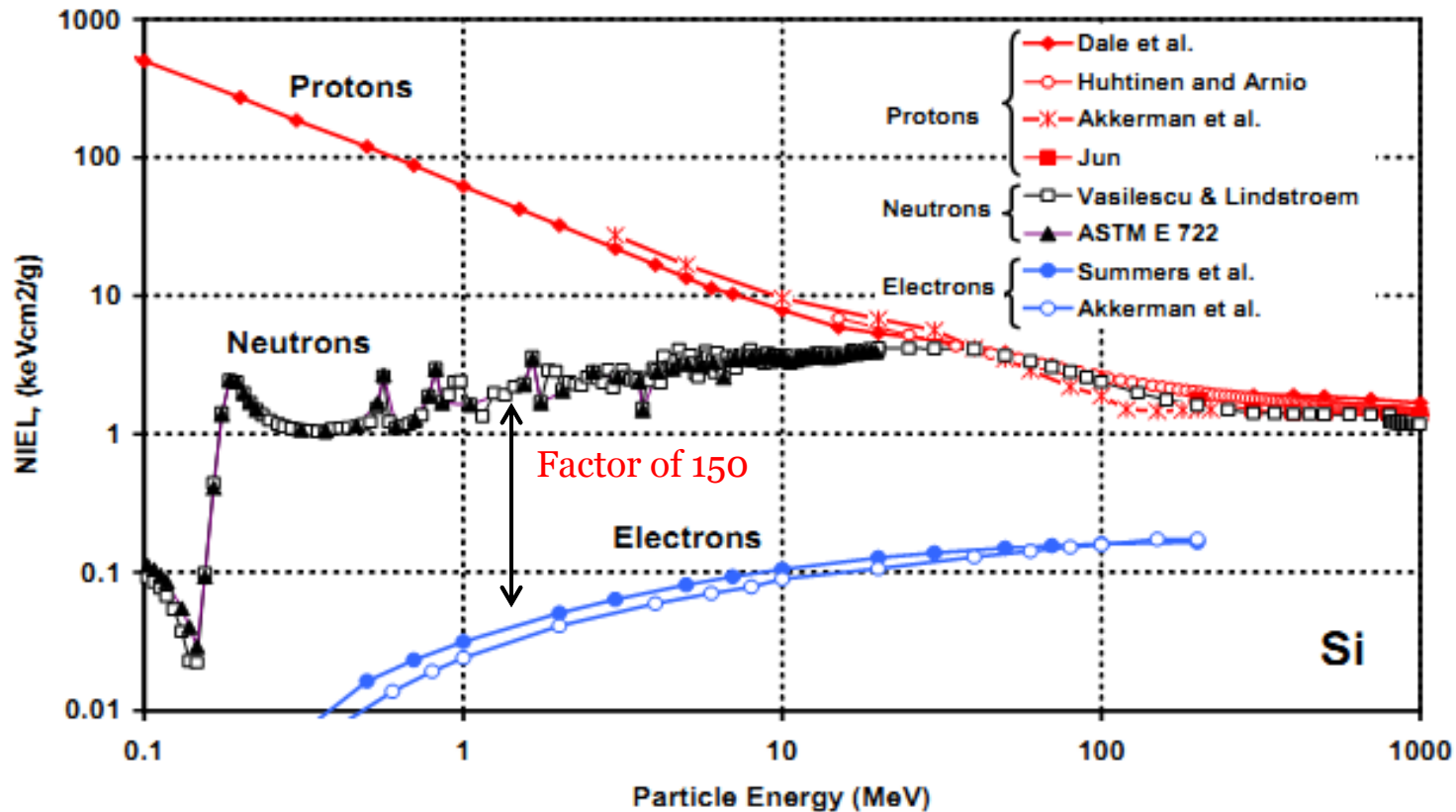
Radiation induced bulk damage



- Levels shown introduced by irradiation or by initial impurities
- Energy levels in the band gap with impact on electrical sensor properties

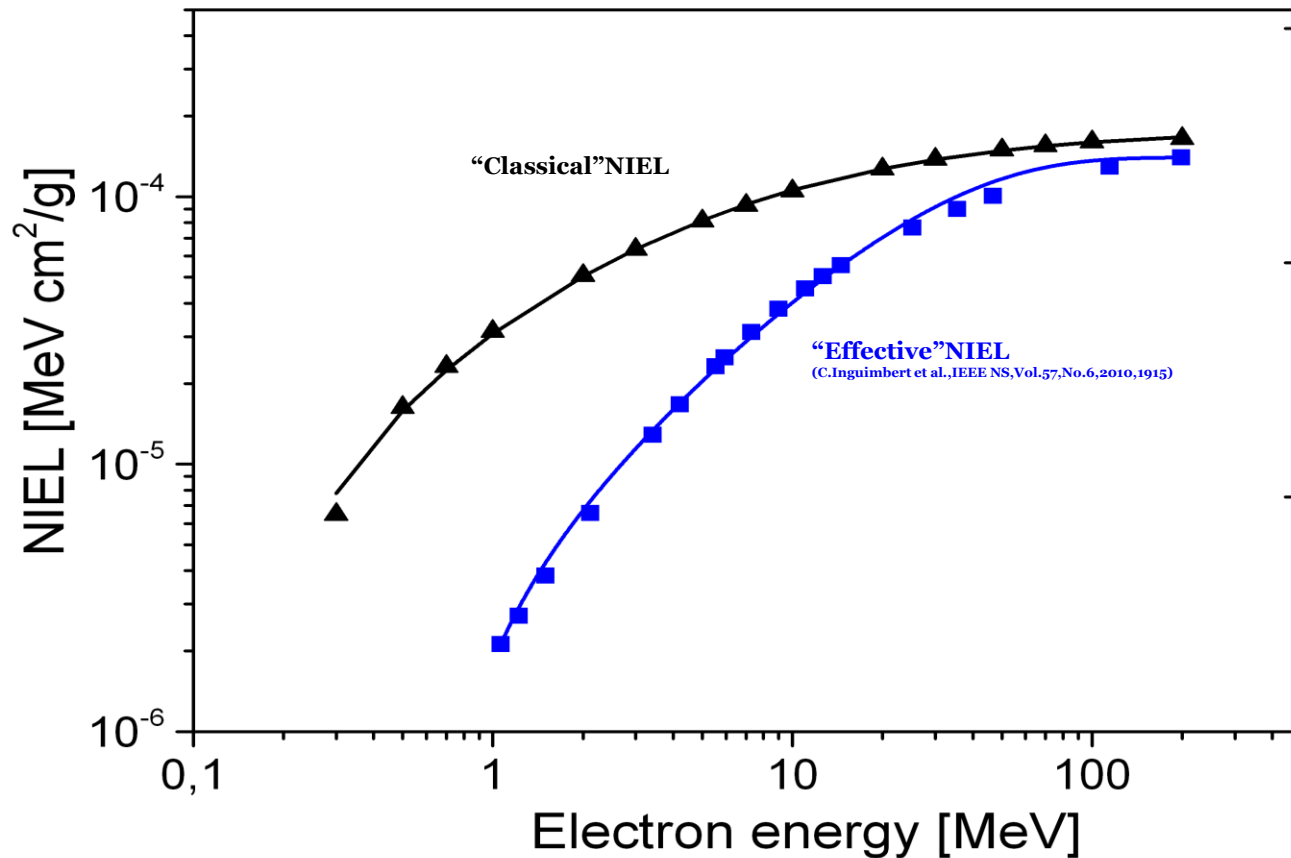
Defects	σ_n [cm ²]	σ_p [cm ²]	E_A [eV]	Assignment/References	Impact on the diodes electrical characteristics at room temperature
E(30K)	2.3×10^{-14}		$E_C - 0.1$	Electron trap with a donor level in the upper half of the Si bandgap / [11]	On the N_{eff} by introducing positive space charge
H(40K)		1.7×10^{-15}	$E_V + 0.09$	Hole trap/[11]	
VO_i^{-0}	1.44×10^{-14}		$E_C - 0.176$	VO_i^{-0} / [40]	
$C_i C_s^{-0}$	1.4×10^{-14}		$E_C - 0.171$	$C_i C_s^{A-0}$ / [41, 42]	
I_p^{+0}	1.7×10^{-15}		$E_V + 0.23$	Donor level of V_2O or of a still unknown C related defect/[11 ,30]	
I_p^{0-}	1.7×10^{-15}	9×10^{-14}	$E_C - 0.55$	Acceptor level of V_2O or of a still unknown C related defect/[11 ,30]	On the N_{eff} by introducing negative space charge and on LC
C_i^{+0}	1.11×10^{-15}	4.28×10^{-15}	$E_V + 0.284$	C_i^{+0} / [21]	
V_2^{-0}	2.1×10^{-15}		$E_C - 0.424$	V_2^{-0} / [21]	
E_4	1×10^{-15}		$E_C - 0.38$	V_3^{\pm} / [38]	On LC
E_5	7.8×10^{-15}		$E_C - 0.46$	V_3^{-0} / [38]	On LC
H(116K)		4×10^{-14}	$E_V + 0.33$	Hole trap with an acceptor level in the lower part of the Si bandgap - Extended defect (cluster of vacancies and/or interstitials) / [10,11]	On the N_{eff} by introducing negative space charge
H(140K)		2.5×10^{-15}	$E_V + 0.36$	Hole trap with an acceptor level in the lower part of the Si bandgap - Extended defects (clusters of vacancies and/or interstitials)/[10,11]	On the N_{eff} by introducing negative space charge
H(152K)		2.3×10^{-14}	$E_V + 0.42$	Hole trap with an acceptor level in the lower part of the Si bandgap - Extended defects (clusters of vacancies and/or interstitials)/[10,11]	On the N_{eff} by introducing negative space charge
H(87K)		0.3×10^{-15}	$E_V + 0.193$	V_3^{0+} / [37]	
H(98K)		1.2×10^{-15}	$E_V + 0.234$	$V_2O^{0+} + V_3O^{0+}$ / [37]	

LC = Leakage current



“Classical” NIEL – Non Ionizing Energy Loss is a quantity that describes the rate of energy loss due to atomic displacements as a particle traverses a material

- Final concentration of defects depends only on NIEL (total energy that goes into displacements) and not on the type of initial energy of the particle
- Number of displacement is proportional to PKA energy → nature of damage independent of PKA energy

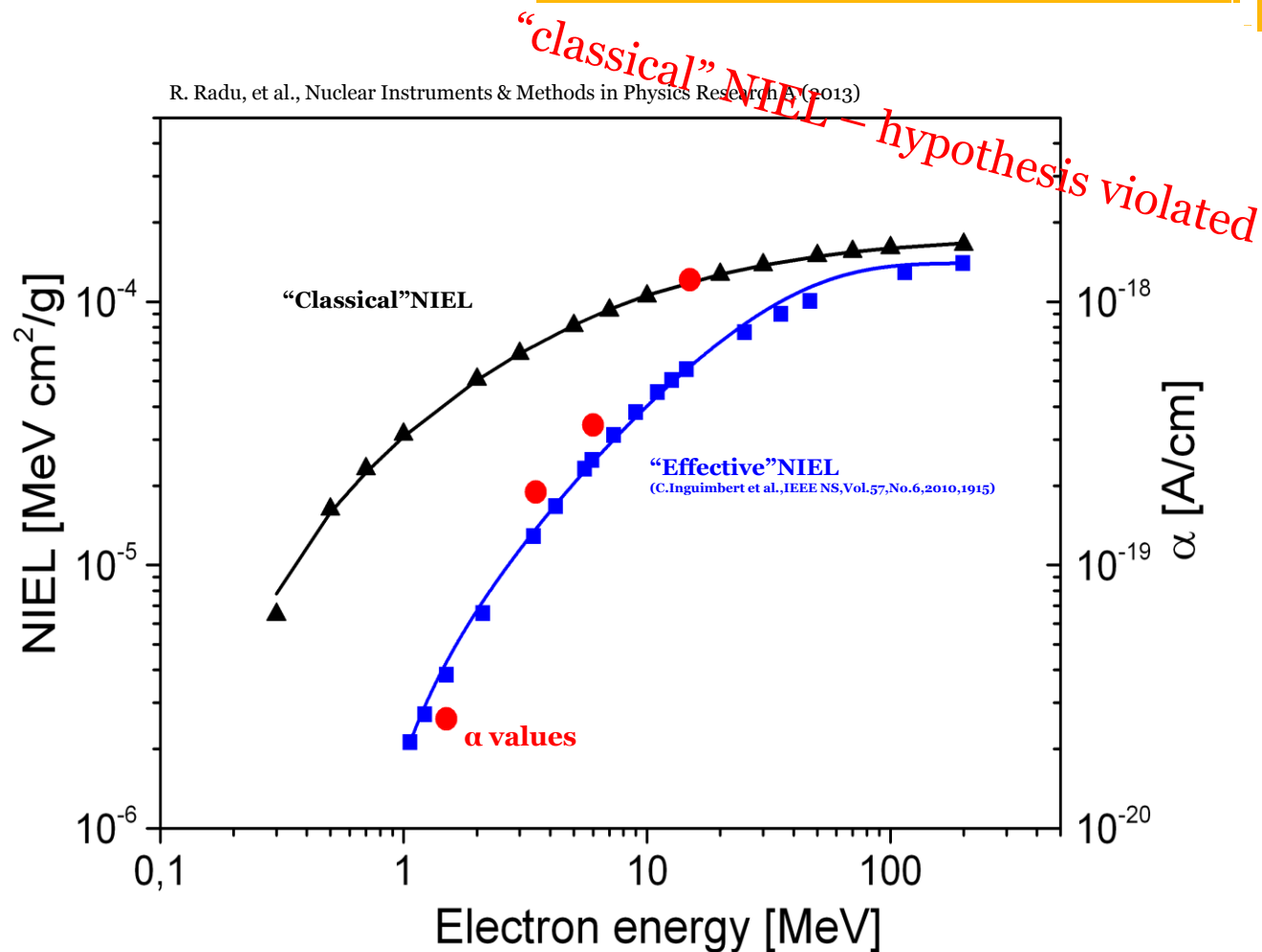


“**effective**” NIEL - based on molecular dynamics (MD) simulation - recombination of displacements in disordered regions is taken into account

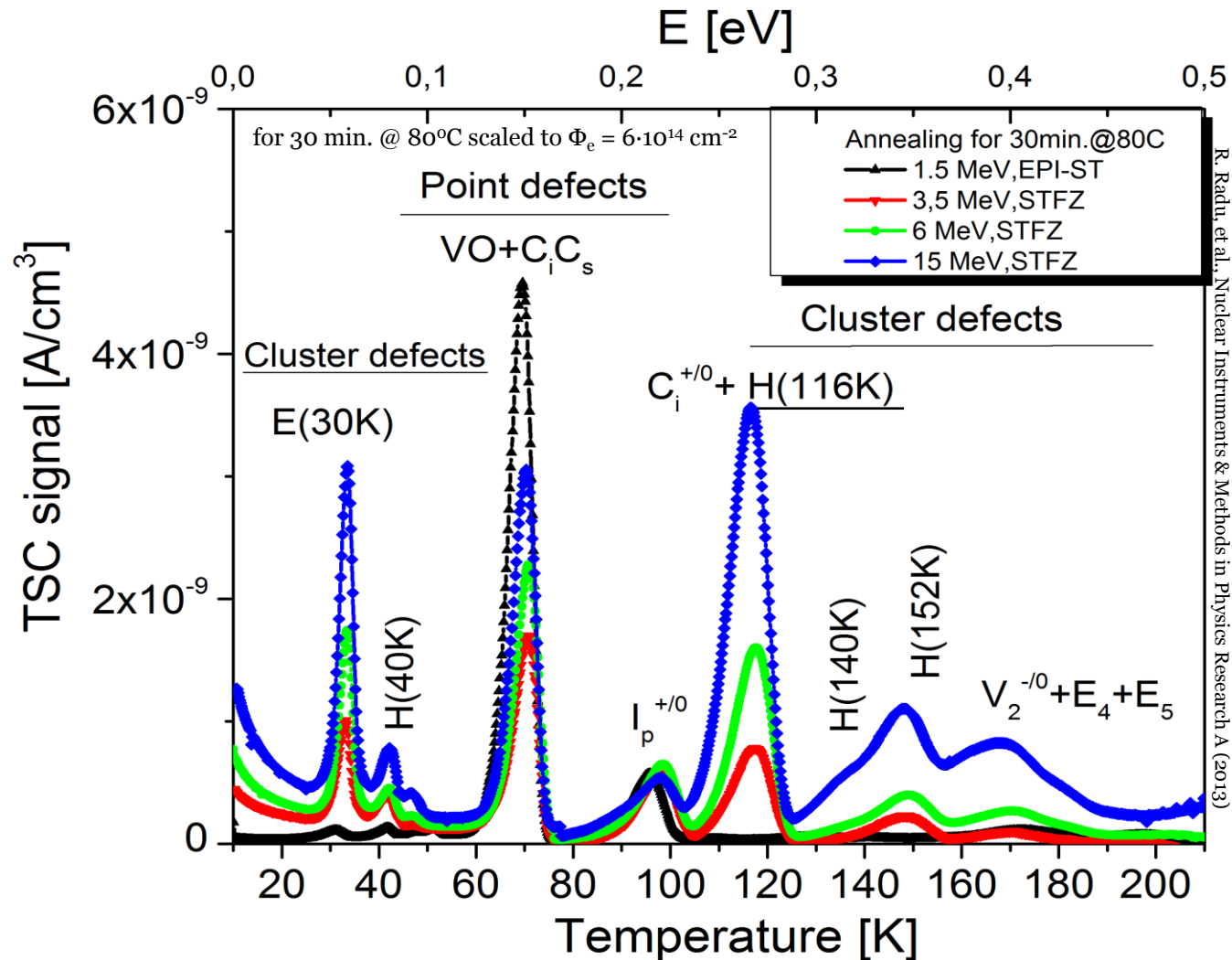
$$\alpha = \Delta I / (V \cdot \Phi_e)$$

measured at 20°C

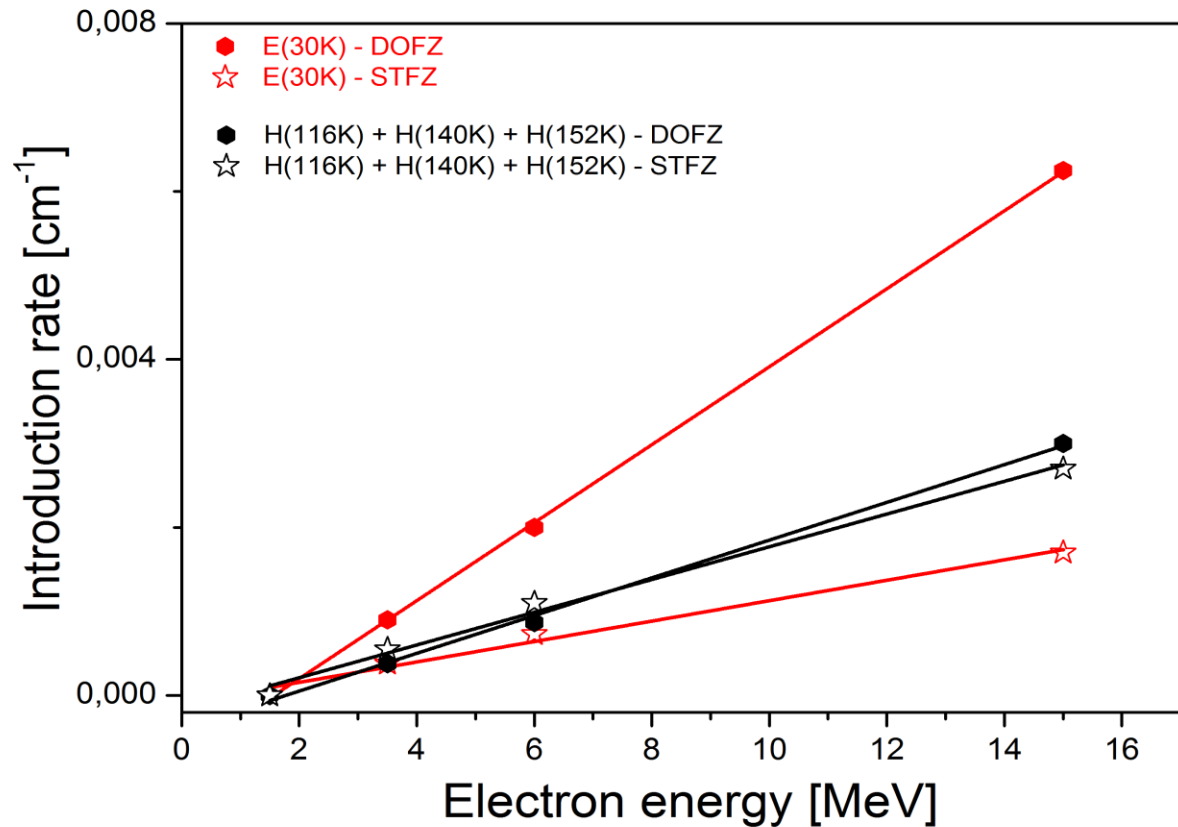
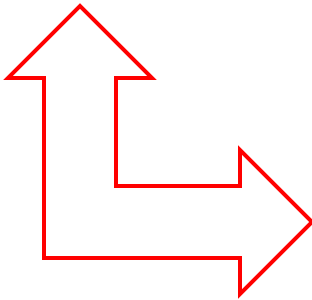
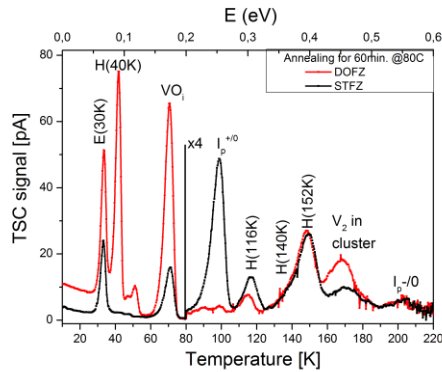
E_e [MeV]	α_e [A/cm]	α_{eq} [A/cm]
1.5	2.6×10^{-20}	1.38×10^{-17}
3.5	1.89×10^{-19}	2.91×10^{-17}
6	3.4×10^{-19}	2.74×10^{-17}
15	1.21×10^{-18}	4.35×10^{-17}



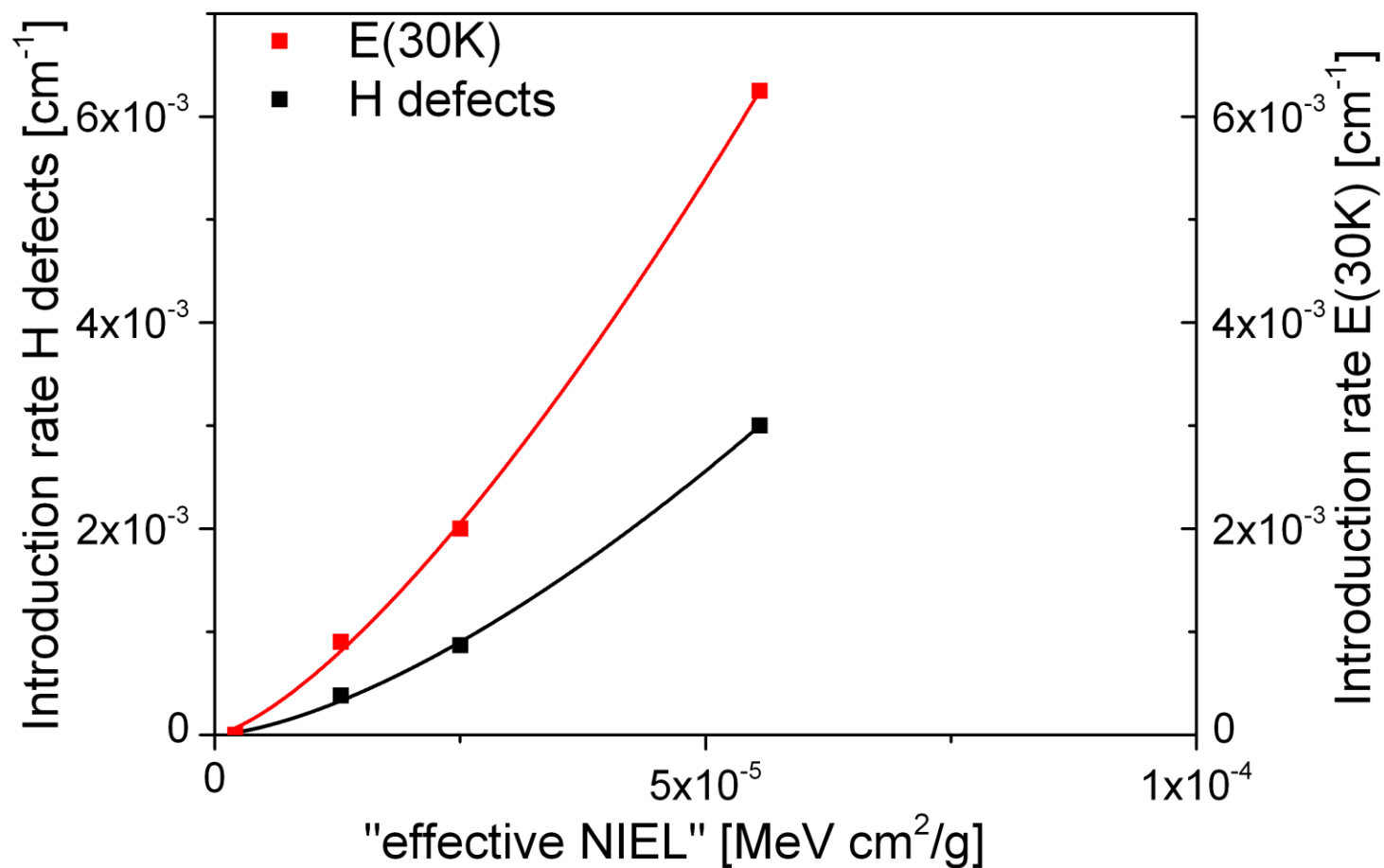
- α for electrons compared with “effective” and “classical” NIEL
- “effective” NIEL describes the energy dependence of α much better



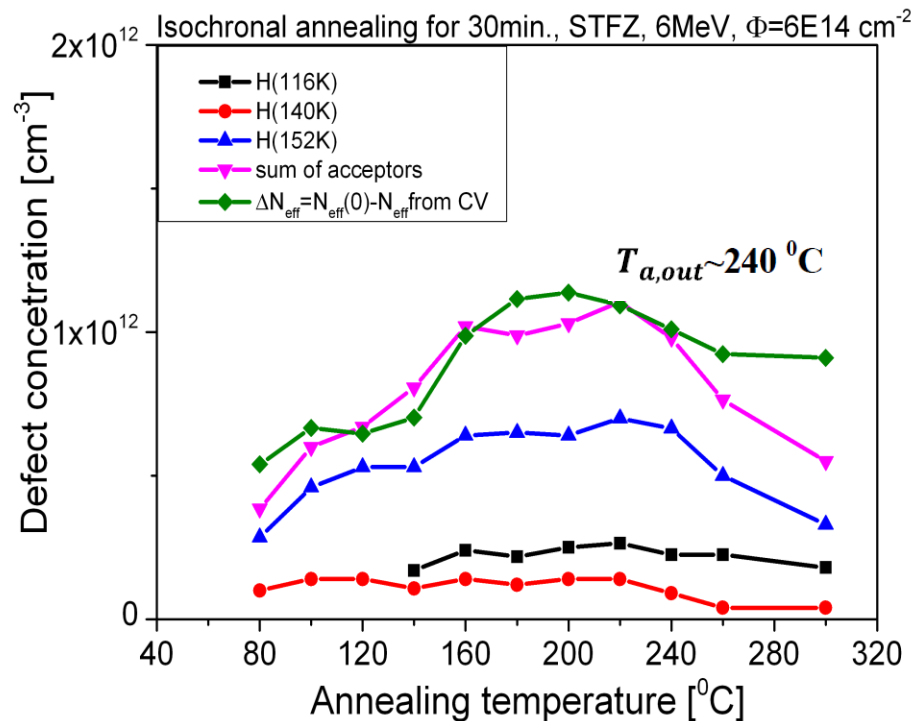
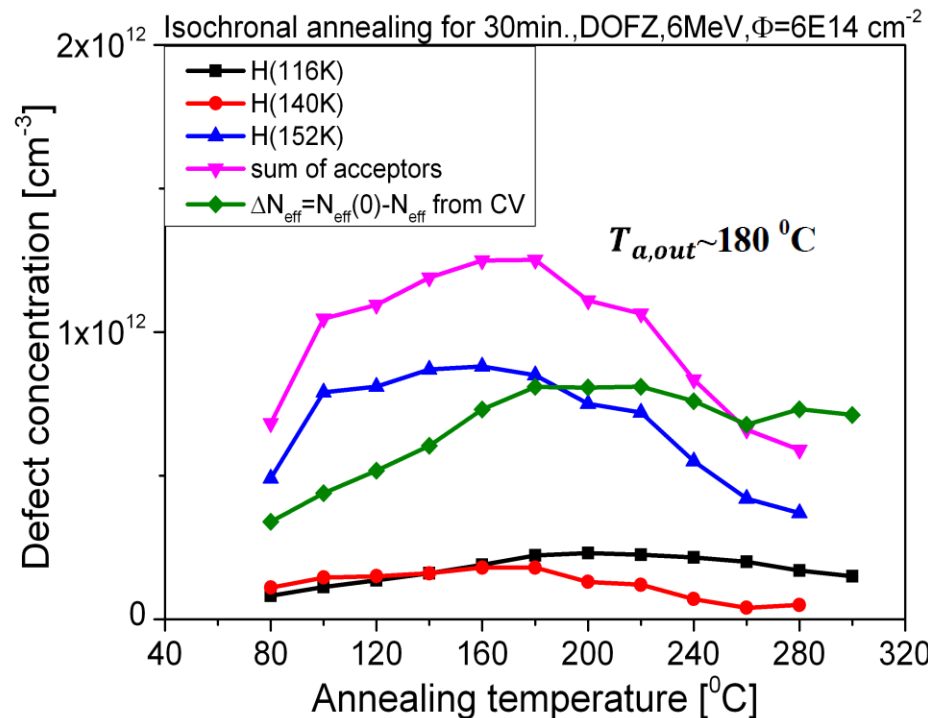
Increasing electron energy \rightarrow increase of local density of vacancies and interstitials \rightarrow cluster defects



- Introduction rates for H defects for DOFZ & STFZ are similar → **no** [O] dependent
- Introduction rate for E (30K) is **3 times** larger in DOFZ material → [O] dependent
- Chemical structure of these defects unknown → **next step: isochronal annealing to get an overview of defect kinetics**

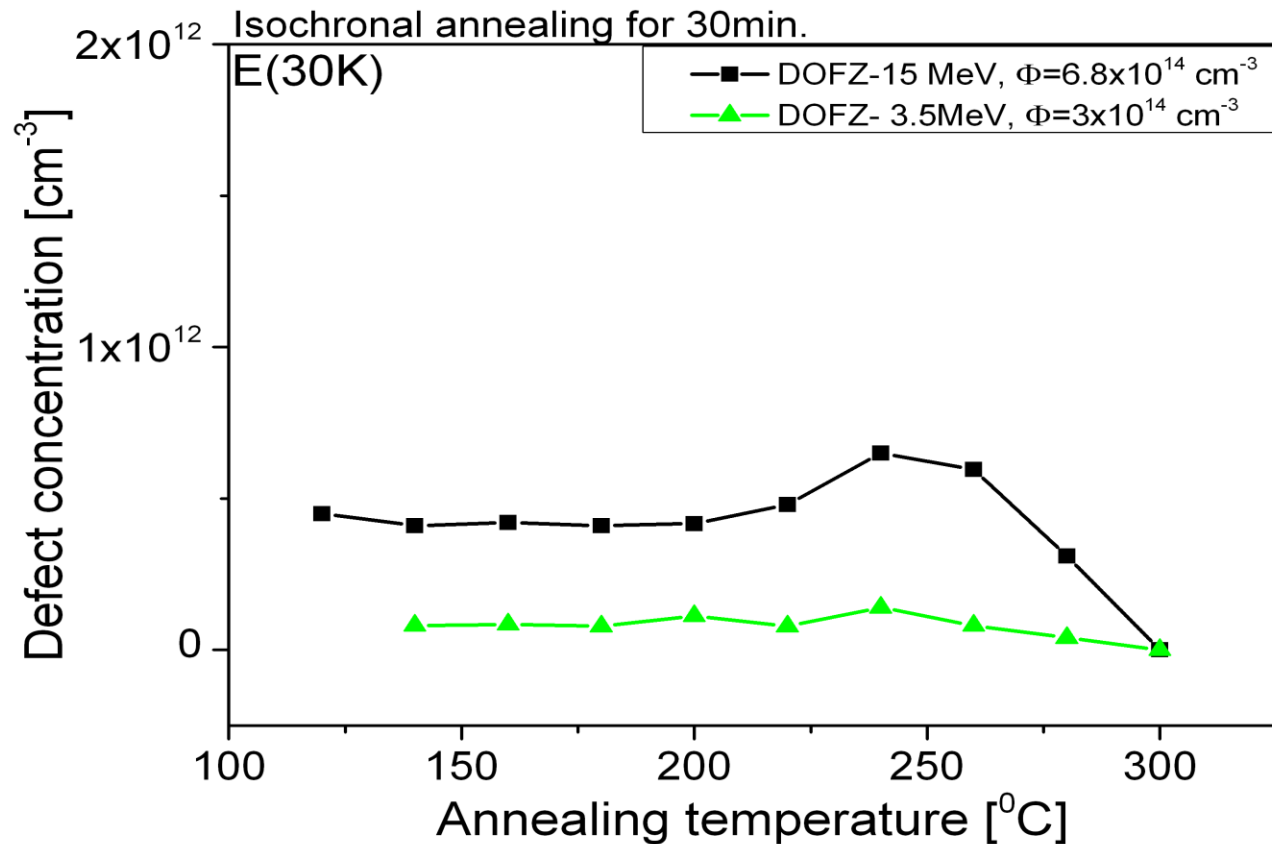


Introduction rates for E(30K) and H defects versus “effective” NIEL follows a power law function



What is the origin of H defects?

- After irradiation – small concentration, than start to increase
- **Puzzle:** If the introduction rates for H defects is not [O] dependent why the annealing in/out temperature is different for the two materials?
- Is there a relation between H defects and higher order vacancies V_n ($n > 3$)?

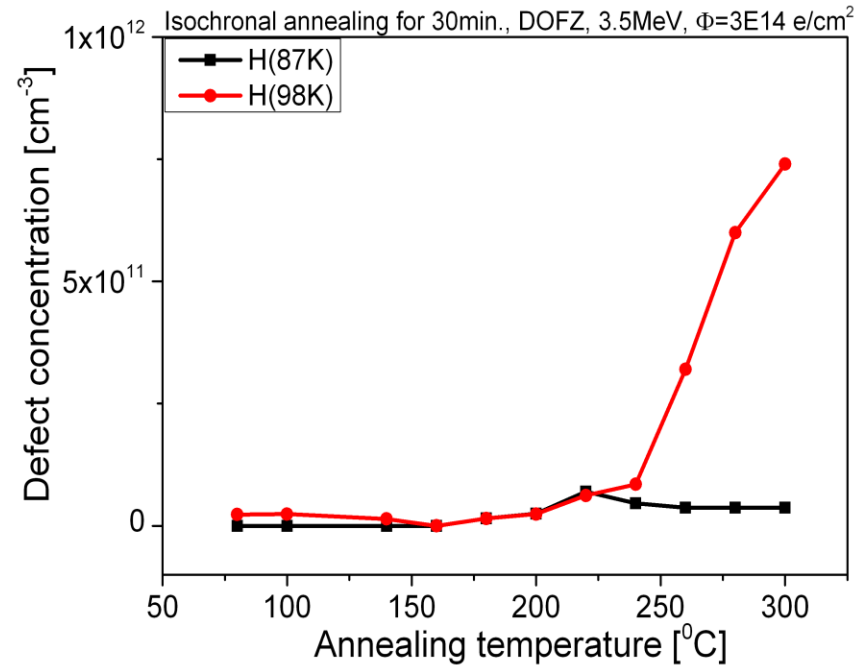
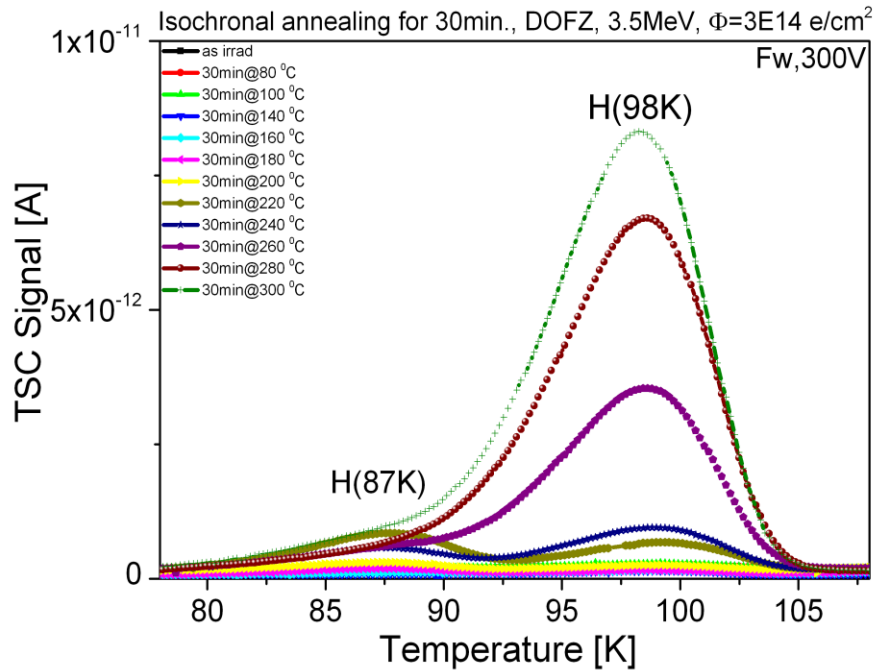


Unknown kinetic mechanism

What is the origin of E(30) defect?

- The E (30K) defect, a shallow donor, increases the positive space charge
- The E(30K) maximum concentration $\sim 240^\circ\text{C}$ and anneals out at 300°C

Next step: isothermal annealing at high temperature to identify its formation kinetics



- V_2 and V_3 become mobile at $T > 200^\circ\text{C}$, they are trapped by oxygen
- Two donor levels have been detected in TSC only after injection of holes
- 1) Hole trap (87K): $V_2^{+/0}$ - stable up to 220°C
 $V_3^{+/0}$ - changed to the ffc. – configuration
 - 2) Hole trap (98K): overlap of $V_2\text{O} + V_3\text{O}$ donor levels

→ consistent with *V.P.Markevich et al: Phys.Status Solidi A 208,No3, 568-571,2011*

Next step: isothermal annealing at high temperature to identify their formation kinetics

Energy dependence of current related damage parameter α :

- α proportional “Effective” NIEL; “Classical” NIEL scaling violated

Energy dependence of cluster-related defects E(30K) and H-defects :

- Increasing electron energy \rightarrow introduction rates increase (linear with E_e , power law with “effective” NIEL)
- Rates for H-defects \rightarrow **no [O] dependence**
- Rate for E (30K) \rightarrow **[O] dependence**

Isochronal annealing (80°C \rightarrow 300°C)

- Concentration of the H defects increase with T_{an} up to $\sim 180^\circ\text{C}$ for DOFZ and $\sim 240^\circ\text{C}$ for STFZ materials followed by a decrease at higher T_{an}
- Annealing of E(30K) shows a maximum at $\sim 240^\circ\text{C}$ for DOFZ

Next steps:

- Isothermal annealing at high temperatures (already started)
- Aim: get more information of defect kinetics of H-defects and E(30K) (activation energies for the formation and decay, frequency factors) – for comparison with results from other methods like EPR \rightarrow identify defect structures

Thank you for your attention!

- [10] I. Pintilie, E. Fretwurst, and G. Lindstroem, Appl. Phys. Lett. 92 (2008) 024101.
- [11] I. Pintilie, et al., Nucl. Instr. and Meth. in Phys. Res. A 611 (2009) 52-68.
- [21] M. Moll, PhD Thesis, University of Hamburg, DESY-THESIS-1999-040, December 1999.
- [30] I. Pintilie, et al., Appl. Phys. Lett. 81 (2002) 165.
- [37] V.P. Markevich, et al., Phys. Status Solidi A 208 (2011) 568-571.
- [38] R.M. Fleming, et al., J. Appl. Phys. 111 (2012) 023715.
- [40] G.D. Watkins, Materials Science in Semiconductor Processing 3 (2000) 227-235.
- [41] L.W. Song, et al., Phys. Rev. Lett. 60 (1988) 460-463.
- [42] L.W. Song, et al. Phys. Rev. B42 (1990) 5765.