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Acousto-defect interaction in irradiated and non-irradiated silicon n^+-p structure



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The influence of ultrasound on current-voltage characteristics of non-irradiated silicon n^+ -p structures as well as silicon structures exposed to reactor neutrons or ⁶⁰Co gamma radiation has been investigated experimentally. It has been found that the ultrasound loading of the n^+ -p structure leads to the reversible change of shunt resistance, carrier lifetime, and ideality factor. Specifically, considerable acoustically induced alteration of the ideality factor and the space charge region lifetime was observed in the irradiated samples. The experimental results were described by using the models of coupled defect level recombination, Shockley-Read-Hall recombination, and dislocation-induced impedance. The experimentally observed phenomena are associated with the increase in the distance between coupled defects as well as the extension of the carrier capture coefficient of complex point defects and dislocations. It has been shown that divacancies and vacancy-interstitial oxygen pairs are effectively modified by ultrasound in contrast to interstitial carbon-interstitial oxygen complexes. Published by AIP Publishing.

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I. INTRODUCTION

It is well known that ultrasound (US) can effectively interact with defects. As a defect engineering tool, US has the following advantages: (i) locality of action due to predominant absorption in the regions of lattice periodicity deviation; (ii) selectivity of influence, which depends on acoustic wave (AW) polarization and AW type; (iii) possibility to transform the defect system by applying resonance frequency; and (iv) reversibility of the effect of low intensity AW.

In piezoelectric semiconductors, the acousto-defect interaction (ADI) is mainly determined by the electric field that accompanies the vibration wave propagation. However, the ADI is also observed in such non-piezoelectric crystals as silicon, the basic material in microelectronics. It was experimentally observed that US can cause atomic diffusion, 1,2 transformation of native and impurity defects, 3-7 modification of interior surface states, 8-10 and appearance of new defects 11,12 in Si structures. Defects are known to determine most of the semiconductor device characteristics. In particular, the ADI governs the variation of tunneling, 13,14 generation–recombination, ^{15–17} and thermionic emission ^{18,19} currents in silicon barrier structures.

The change of population of impurity oscillator levels,²⁰ the displacement of impurity atoms with respect to their surroundings, 4,21,22 the decrease in the diffusion activation energy,²³ the local temperature increase caused by point defect clusters, 24 and the US absorption by dislocations 16,25,26 are believed to be the main mechanisms of elastic vibration-defect interactions in non-piezoelectric crystals. However, to the best of our knowledge, there is no comprehensive ADI theory for silicon suggested so far, the lack of experimental research focused on acoustically induced (AI) effects being one of the main reasons.

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The defects in silicon structures are not all acoustically active (AA) and can remain unmodified under the action of ultrasound. The ADI efficiency depends on the defect type and structure.⁸ For example, the force acting on the point defect in the crystal under US loading (USL) is determined by the relaxation of the defect volume. ^{21,22} The alterations of semiconductor defects are most widely produced by using the well-studied irradiation method. On the one hand, the high-power US treatment of irradiated silicon structures has been shown^{27–30} to result in residual changes in structure properties. This effect deals with AI annealing of radiation defects (RDs). On the other hand, irradiation can be the reason of reversible AI phenomenon initiation, ^{31,32} which is caused by the formation of acoustically active RDs. Unfortunately, there are but a few reports on the acoustically driven phenomenon in irradiated silicon structures.

The aim of our work is to investigate experimentally the AI electrical characteristic variation that takes place in nonirradiated and irradiated n^+ –p-Si structures. For this purpose, the samples were irradiated by reactor neutrons and ⁶⁰Co–gamma source rays. It is supposed that γ -rays introduce predominantly the VO_i complex, 33-35 whereas neutrons mainly create vacancy clusters, ^{36,37} disordered regions, ³⁸ and C_iO_i complexes. 35,39 Our work presents distinctions between AI effects in silicon structures with different RDs. The intensity of US applied was insufficient for a new defect formation, RD annealing, or long distance (a many interatomic distance) diffusion. As a result, the complete recovery of characteristics was observed after AW propagation had stopped. To describe the processes in the space charge region 81 (SCR) and in the diode base as well as to study shunt resistance, we used the models of coupled defect level recombination, 40,41 Shockley-Read-Hall (SRH) recombination, and

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TABLE I. The sample irradiation parameters.

Sample	Irradiation Type	D (rad)	Ψ (cm ⁻²)	NIEL ^a (MeV cm ² /g)	$\frac{\Psi \times \text{NIEL}}{(\text{MeV/g})}$
iSC	Non	0	0		0
nSC	neutron	4.5×10^{3}	4×10^{11}	2.04×10^{-3}	8.2×10^{8}
g6SC	γ - 60 Co	1×10^{6}	1.6×10^{15}	1.07×10^{-7}	1.7×10^{8}
g7SC	γ - 60 Co	1×10^7	1.6×10^{16}	1.07×10^{-7}	1.7×10^{9}

^aReference 44.

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dislocation-induced impedance, ^{42,43} respectively. The observed AI phenomena are accounted for in terms of the defect interaction with the AW strain field. ^{21,22} Our research not only provides a better understanding of ADI but could also facilitate the development of acoustically controlled devices or radiation sensors.

II. EXPERIMENTAL AND CALCULATION DETAILS

The n^+ -p-Si structure was fabricated from a 2 in. (300 µm thick) p-type boron doped Czochralski silicon wafer with $\langle 111 \rangle$ orientation and a resistivity of 10Ω cm. The n⁺ emitter with a carrier concentration of about 10¹⁹ cm⁻³ and a thickness of $0.5 \mu m$ was formed by phosphorus implantation. The front and rear aluminium electrodes were deposited by screen printing before rapid annealing. The samples used in the experiment were cut from the central part of the wafer and had the area of 2 cm². The samples were irradiated by reactor neutrons or by 60 Co γ -rays. The doses D, fluences Ψ , and sample labels are listed in Table I. To determine D and Ψ correlation, the data from Refs. 44 and 45 were used. The non-ionizing energy losses (NIEL) for neutron and γ -60Co are also shown in Table I. Since the displacement damage effect is characterized by $(\Psi \cdot NIEL)$, a similar damage was expected in the investigated samples as well. To avoid the impact of long-term annealing, which is typical for the neutron damaged structure, 35,36 the irradiated samples were stored for five years at room temperature before the measurements.

The dark forward current–voltage (I–V) characteristics of the samples both with and without USL were measured over a temperature range of 290–340 K. The temperature was controlled by a differential copper–constantan thermocouple. Some of the obtained curves are shown in Fig. 1.

The double-diode model of n^+ –p structure I–V characteristics is expressed in the following form:

$$I(V,T) = I_{SCR} + I_{base} + I_{sh}, \tag{1}$$

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$$I_{SCR} = \frac{qAn_i d}{2\tau_g} \left\{ \exp\left[\frac{q(V - IR_s)}{n_{id}kT}\right] - 1 \right\},\tag{2}$$

$$I_{base} = \frac{qAn_i^2}{p_p} \sqrt{\frac{\mu_n kT}{\tau_n}} \left\{ \exp\left[\frac{q(V - IR_s)}{kT}\right] - 1 \right\}, \quad (3)$$

$$I_{sh} = (V - IR_s)/R_{sh}, (4)$$

where I_{SCR} describes the overall SCR recombination, I_{base} is 118 closely related to the recombination in the quasi-neutral 119 region, I_{sh} is the shunt current, A is the sample area, n_i is the 120 intrinsic carrier concentration, τ_g is the SCR carrier lifetime, 121 d is the SCR thickness

$$d = \sqrt{\frac{2\varepsilon\varepsilon_0}{qp_p} \left[\frac{E_g}{q} - \frac{kT}{q} \ln\left(\frac{N_v N_c}{p_p n_n}\right) - \frac{2kT}{q} - V \right]}, \quad (5)$$

 ε is the permittivity (11.7 for Si), p_p and n_n are the majority 123 carrier concentration in p- and n-type regions, E_g is the semiconductor bandgap, N_c and N_v are the effective densities of 125 states in the conduction and valence bands; n_{id} is the ideality 126 factor, R_s and R_{sh} are the series and shunt resistances, and μ_n 127 and τ_n are the mobility and lifetime of the electron (minority 128 carrier) in the diode base.

We used Eqs. (1)–(5) to fit the experimental data taking 130 τ_g , τ_n , $n_{\rm id}$, R_{sh} , and R_s as the fitting parameters. Also, we 131 used the known 46–48 temperature dependences of n_i , E_g , 132 and μ_n . As a result, we obtained extremely good fit to the 133 experimental data—see Fig. 1. In particular, for all the samples, the value of R_s was found to be about 1 Ω . The broken 135 lines in Fig. 1(a) show an example of the calculated contributions of I_{SCR} , I_{base} , and I_{sh} to the total current.

In the case of USL, the transverse AWs with the fre- 138 quency of 4.2 MHz, which were excited by using a piezo- 139 electric transducer, were applied to the samples at the base 140 side in the [111] direction. The US intensities $W_{\rm US}$, ampli- 141 tudes of lattice deformation $\xi_{\rm US}$, and lattice atom displace- 142 ments $u_{\rm US}$ are listed in Table II. It was reported 143 previously 6,7,19 that the characteristic time of change in 144

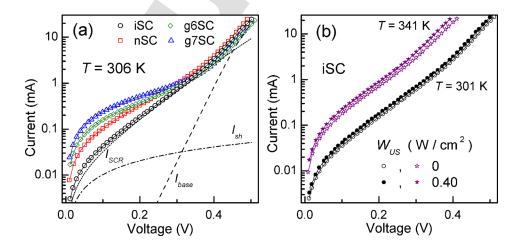


FIG. 1. Dark *I–V* characteristics measured (a) at 306 K for non-irradiated (circles), neutron-irradiated (squares), and gamma-irradiated (diamonds and triangles) structures without USL; (b) at 301 K (circles) and 341 K (asterisks) with (filled marks, Ui-2) and without (open marks) USL for the iSC. The marks are the experimental results, and the solid lines are the curves fitted by Eqs. (1)–(5). The dashed, dotted, and dotted–dashed lines in (a) represent the calculated base, SCR, and shunt components of total iSC current (black solid line).

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TABLE II. The ultrasound loading parameters.

Sample	W _{US} (W/cm ²)	$\xi_{\rm US} (10^{-6})$	u _{US} (nm)	USL label
iSC	0.22	3.1	0.67	Ui-1
	0.40	4.2	0.91	Ui-2
nSC	0.24	3.2	0.70	Un-1
	0.40	4.2	0.91	Un-2
g6SC	0.38	4.1	0.89	Ug6-2
g7SC	0.19	2.9	0.63	Ug7-1
	0.37	4.0	0.87	Ug7-2

silicon structure parameters under the US action did not exceed 2×10^3 s. In order to wait until the AI transitional period is completed, the following experimental procedure was used. When USL started, the sample was first exposed to room temperature for 60 min and then the I-V measurement and the sample heating were started. In order to avoid the effect of the piezoelectric field on I-V characteristics, the piezoelectric transducer was shielded.

Figure 2 illustrates the reversibility of AI effects. The time interval between USL initiation and "during" measurement was longer than 60 min, and the time interval between USL termination and "after" measurement was about 24 h. The data for nSC and g6SC are similar to those presented for iSC and g7SC.

The non-linear fittings were performed by using the differential evolution method. 49

III. RESULTS AND DISCUSSION

A. Space charge region

The parameters of I-V characteristics associated with 163 SCR phenomena are $n_{\rm id}$ and τ_g . The temperature dependences of the ideality factor and SCR carrier lifetime are shown 165 in Figs. 3 and 4, respectively.

As shown in Fig. 3, the ideality factor decreases with the increase in temperature, and the dependence of $n_{\rm id}$ on 1/T is close to linear. Thus, dependence $n_{\rm id}(T)$ can be expressed as 169

$$n_{\rm id}(T) = n_{\rm id,\infty} + T_{\rm id}/T. \tag{6}$$

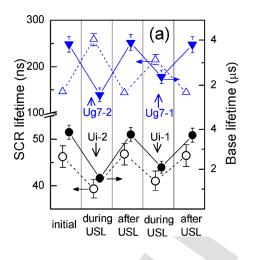
The thermoactivated growth of SCR lifetime is observed 170 over the explored temperature range—see Fig. 4. The temperature dependence of τ_g is well described by the following 172 equation: 173

$$\tau_g(T) = \tau_{g0} \exp\left(-\frac{E_{\tau g}}{kT}\right). \tag{7}$$

The values of $T_{\rm id}$ and $E_{\tau g}$ found for both non-irradiated and 174 irradiated samples under USL as well as without USL are 175 listed in Table III.

We would like to stress that

irradiation leads to changes in T_{id} and $E_{\tau g}$, and 178 g6SC's characteristic temperature of the ideality factor and SCR lifetime characteristic energy are closely 180 related to those of g7SC under similar conditions; 181



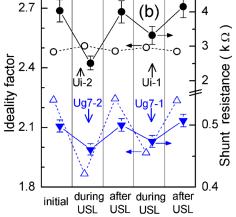
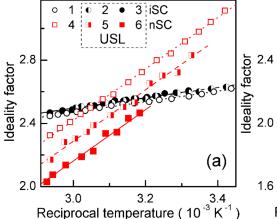


FIG. 2. SCR lifetime (a, left axis, open marks), base lifetime (a, right axis, filled marks), ideality factor (b, left axis, open marks), and shunt resistance (b, right axis, filled marks) obtained before, during, and after USL at 330 K. Data for iSC (circles) and g7SC (triangles) are presented.



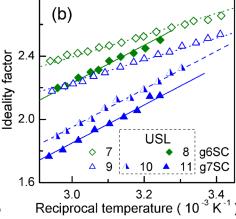


FIG. 3. Temperature dependences of the ideality factor for non-irradiated (curves 1–3, circles), neutron-irradiated (4–6, squares), and γ -irradiated (7–11, diamonds and triangles) samples. The curves 1, 4, 7, and 9 (open marks) are obtained without USL, and curves 2, 3, 5, 6, 8, 10, and 11 correspond to Ui-1, Ui-2, Un-1, Un-2, Ug6-2, Ug7-1, and Ug7-2, respectively. The marks are the experimental results, and the lines are the fitted curves using Eq. (6).

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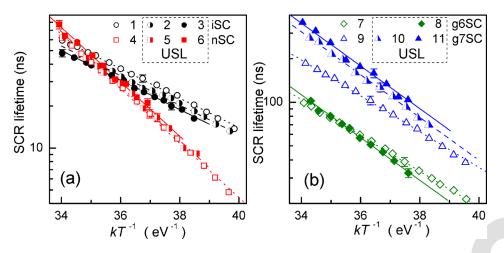


FIG. 4. Temperature dependences of SCR lifetime for non-irradiated (curves 1-3, circles), neutron-irradiated (4-6, squares), and γ-irradiated (7-11, diamonds and triangles) samples. The curves 1, 4, 7, and 9 (open marks) are obtained without USL, and curves 2, 3, 5, 6, 8, 10, and 11 correspond to Ui-1, Ui-2, Un-1, Un-2, Ug6-2, Ug7-1, and Ug7-2, respectively. The marks are the experimental results, and the lines are the fitted curves using Eq. (7).

- 182 USL affects n_{id} and τ_{ϱ} values; the absolute AI changes of the ideality factor $\Delta n_{\rm id} = n_{\rm id,US} - n_{\rm id,in}$ and the rela-183 tive AI changes of SCR lifetime $\varepsilon_{\tau g} = (\tau_{g,US} - \tau_{g,in})/$ 184 $\tau_{g,in}$ (where subscripts "US" and "in" indicate the val-185 ues obtained at the same temperature with and without 186 USL, respectively) are listed in Table IV; 187
- $\Delta n_{\rm id}$ and $\varepsilon_{\tau g}$ vary with $W_{\rm US}$ enhancement, whereas $T_{\rm id}$ 188 (iii) and $E_{\tau g}$ values practically do not depend on US 189 intensity; 190
- 191 (iv) USL leads to the increase in both T_{id} and $E_{\tau g}$ in γ -irradiated samples [see Figs. 3(b) and 4(b)], but this 192 effect is not observed in non-irradiated and neutron-193 irradiated samples [see Figs. 3(a) and 4(a)]; 194
- $\Delta n_{\rm id}$ and $\varepsilon_{\tau g}$ have opposite signs for non-irradiated and irradiated samples (for SCg6 not in the whole 196 temperature range);
- (vi) ideality factor is varied by USL more effectively in 198 the irradiated samples. 199

For the purpose of our analysis, it is important to discuss the recombination mechanism in SCR of the investigated samples. According to classical SRH theory, the ideality factor must be smaller than 2, and τ_g temperature dependence is expected^{50,51} to be described by the relation $\tau_g \simeq 2 \tau_n$ $\sqrt{\sigma_n/\sigma_p} \cosh[(E_t - E_i)/kT]$ (where σ_n , σ_p , and E_t are the electron and hole capture cross sections (CCSs) and the energy level of the recombination center, and E_i is the intrinsic energy level). In our case, n_{id} is greater than 2, and τ_{o}

TABLE III. Characteristics of temperature dependences of n^+ –p-Si structure parameters.

Sample	USL	$T_{\mathrm{id}}\left(\mathbf{K}\right)$	$E_{\tau g}$ (eV)	$R_{293,\mathrm{Al}}$ (k Ω)	$\sigma_{\rm dis}~(10^4~{ m K/\Omega})$
iSC	non	330 ± 30	0.24 ± 0.01	27 ± 3	41 ± 4
	Ui-1	310 ± 30	0.24 ± 0.01	27 ± 3	50 ± 4
	Ui-2	360 ± 30	0.24 ± 0.01	26 ± 3	58 ± 4
nSC	non	1610 ± 70	0.45 ± 0.02	2.2 ± 0.4	65 ± 7
	Un-1	1600 ± 70	0.44 ± 0.02	2.3 ± 0.4	95 ± 10
	Un-2	1680 ± 70	0.44 ± 0.02	2.2 ± 0.4	130 ± 10
g6SC	non	610 ± 40	0.28 ± 0.01	0.7 ± 0.1	19 ± 2
	Ug6-2	1080 ± 50	0.33 ± 0.02	0.8 ± 0.1	24 ± 2
g7SC	non	770 ± 50	0.29 ± 0.01	0.41 ± 0.06	26 ± 3
	Ug7-1	1260 ± 60	0.34 ± 0.02	0.39 ± 0.06	45 ± 4
	Ug7-2	1270 ± 60	0.35 ± 0.02	0.38 ± 0.06	55 ± 4

increases with temperature. Therefore, SRH theory cannot 209 be applied in our case. Several attempts to account for large 210 $n_{\rm id}$ values have been made by using different models.^{52–55} 211 However, all the observed features of SCR recombination 212 (large ideality factor, independence of light intensity, depen- 213 dence on temperature as well as short carrier lifetime) can be 214 explained by the model of coupled defect level recombina- 215 tion (CDLR)^{40,41} only. This mechanism provides a rapid 216 direct charge transfer between defect levels. The phenome- 217 non was first observed experimentally, 56,57 after which it 218 was recruited to explain the process in semiconductor 219 diodes. 40,41,58

According to the CDLR model, the recombination is the 221 result of carrier exchange between two defect levels and 222 crystal bands. In particular, it is supposed⁴¹ that the recombi- 223 nation rate is dominant at the sites where the acceptor-like 224 defect is coupled with the donor-like defect. In a simplified 225 case, when there is no carrier exchange between the donor 226 level $E_t^{\rm D}$ and the valence band, as well as between the accep- 227 tor level E_t^A and the conduction band, the recombination rate 228 R can be expressed⁴⁰ as

$$R = \frac{R_{12} - \sqrt{R_{12}^2 - 4\tau_n^D \tau_p^A (np - n_i^2)(1 - \epsilon)}}{2\tau_p^D \tau_p^A (1 - \epsilon)},$$
 (8)

$$R_{12} = \frac{(n+n_{\rm D})(p+p_{\rm A})}{R_{\rm DA}} + \tau_n^{\rm D}(p+p_{\rm D}) + \tau_p^{\rm A}(n+n_{\rm A}), \quad (9)$$

$$\tau_n^{\rm D} = (N_{\rm D} \, \sigma_n^{\rm D} \, v_{\text{th},n})^{-1}, \quad \tau_n^{\rm A} = (N_{\rm A} \, \sigma_n^{\rm A} \, v_{\text{th},p})^{-1},$$
(10)

TABLE IV. Acoustically induced change of n^+ –p-Si structure parameters (at 330 K).

Sample	USL	$\Delta n_{\rm id} \\ (\pm 0.01)$	$rac{arepsilon_{ au g}}{(\pm 5\%)}$	$\epsilon_{\tau n}$ (±0.2)	$\varepsilon_{\sigma \mathrm{dis}}$ $(\pm 10\%)$
iSC	Ui-1	0.02	-14	0.7	20
	Ui-2	0.03	-17	1.4	40
nSC	Un-1	-0.13	5	1.5	50
	Un-2	-0.26	13	3.0	100
g6SC	Ug6-2	-0.15	2	2.3	30
g7SC	Ug7-1	-0.26	49	0.9	70
	Ug7-2	-0.36	70	1.9	110

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where R_{DA} is the coupling parameter, N_{D} and N_{A} are the densities of donor and acceptor-like defects, $\sigma_n^{\rm D}$ and $\sigma_p^{\rm A}$ are the electron CCS of the donor and hole CCS of the acceptor, 232 $v_{\text{th},n}$ and $v_{\text{th},p}$ are the thermal electron and hole velocities, 233 $n_{\rm D,A}, p_{\rm D,A},$ and ϵ depend on $E_t^{\rm D}, E_t^{\rm A},$ and level degeneracy factors. Since $\tau_g \propto R^{-1}$, the last three values are expected to 235 provide a thermoactivated behavior of SCR lifetime. Unfortunately, the equation does not account for the func-237 tional relation between I-V characteristics parameters and 238 239 attributes of defects taking part in CDLR.

According to Steingrube *et al.*, 41 CCS for the defect in a pair differs from that for an isolated defect and depends on the distance r between the donor and the acceptor

$$\sigma_{n,p}^{D,A}(r) = C_{n,p}^{D,A} r^2, \tag{11}$$

where $C_n^{\rm D}$ and $C_p^{\rm A}$ are the constant values. $R_{\rm DA}$ is proportional to the overlap integral of the defect wave functions as well. If both defects are characterized by the H-like radial–symmetric wave function and equal Bohr radius a_0 , the following expression can be used:⁴¹

$$R_{\rm DA}(r) \propto N_{\rm D}N_{\rm A} \left[1 + \frac{r}{a_0} + \frac{1}{3} \left(\frac{r}{a_0} \right)^2 \right] e^{-r/a_0}.$$
 (12)

In our opinion, the observed reversible AI modifications of $n_{\rm id}$ and τ_g are induced by donor–acceptor distance alteration in the samples under USL. In fact, according to the data, ^{21,22} the force acting on a point defect during USL can be expressed as

$$F_d = \chi \, \Delta \Omega_d \, \frac{\partial \xi(z, t)}{\partial z},\tag{13}$$

where χ is the bulk elasticity modulus, $\Delta\Omega_d$ is the crystal volume change per defect, ξ is the crystal lattice deformation, and AW propagates along the z axis, $\partial \xi(z,t)/\partial z \propto \xi_{\rm US}$. For the interstitial atoms and substitutional impurities with ionic radius exceeding the ionic radius of matrix atoms, $\Delta\Omega_d>0$, whereas for the vacancies and substitutional impurities whose ionic radius is smaller than that of matrix atoms, $\Delta\Omega_d<0$. Therefore, a point defect vibrates under USL, so oscillation amplitude and phase are determined by both the defect character and AW intensity.

The simplest model, which is shown in Fig. 5, gives the following qualitative conclusion. Initially, the donor and the acceptor are separated by the distance r_{in} , and the X axis is drawn through the point defect initial positions. Under USL, the defects would vibrate with amplitudes $u_{\rm D}$ and $u_{\rm A}$. The vibration axis coincides with the AW displacement direction and forms angle φ with the X-axis. Depending on ξ_{US} , defect elastic strain $(\Delta\Omega_d^{\rm D})$ and $\Delta\Omega_d^{\rm A}$, and defect coupling, the defect vibration amplitudes can have different values. The donoracceptor distance in the sample under USL $r_{\rm US}$, according to the model, depends on time t

$$r_{\text{US}}(t) = \left\{ \left[r_{in} + u_{\text{A}} \cos \left(\omega_{\text{US}} t + \delta \right) - u_{\text{D}} \cos \left(\omega_{\text{US}} t \right) \right]^{2} \cos^{2} \varphi \right.$$
$$+ \left[u_{\text{A}} \cos \left(\omega_{\text{US}} t + \delta \right) - u_{\text{D}} \cos \left(\omega_{\text{US}} t \right) \right]^{2} \sin^{2} \varphi \right\}^{0.5},$$

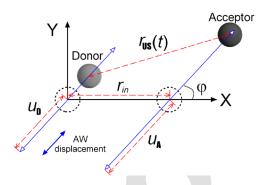


FIG. 5. Model of CDLR center behavior under US action.

where $\omega_{\rm US}$ is the US cyclic frequency and δ is the phase shift 274 between donor and acceptor vibration. 275

We use Eqs. (11) and (12) to estimate AI relative changes 276 of CCS $\varepsilon_{\sigma} = [\sigma_{\text{US}} - \sigma(r_{in})]/\sigma(r_{in})$ and coupling parameters 277 $\varepsilon_{\text{RDA}} = [R_{\text{DA,US}} - R_{\text{DA}}(r_{in})]/R_{\text{DA}}(r_{in})$, where σ_{US} and $R_{\text{DA,US}}$ 278 are averaged over the AW period T_{US} 279

$$\sigma_{ ext{US}} = rac{1}{T_{ ext{US}}} \int_0^{T_{ ext{US}}} \sigma(r_{ ext{US}}(t)) dt,
onumber \ R_{ ext{DA,US}} = rac{1}{T_{ ext{US}}} \int_0^{T_{ ext{US}}} R_{ ext{DA}}(r_{ ext{US}}(t)) dt.$$

In this estimation, the relaxation time in the CDLR subsystem is assumed to be considerably shorter than $T_{\rm US}$, and 281 we apply the previously used⁴¹ value $a_0=3.23$ nm. In addition, the chosen $u_{\rm D}$ and $u_{\rm A}$ values are commensurate with 283 $u_{\rm US}$. However, it should be taken into account that the displacement of the point defect without the covalent bond 285 could exceed a matrix atom displacement. Finally, no US 286 absorption by the defect is assumed. In this simple case, δ is 287 equal to 0° if $(\Delta\Omega_d^D \cdot \Delta\Omega_d^A) > 0$ or to 180° if $(\Delta\Omega_d^D \cdot \Delta\Omega_d^A)$ 288 < 0. In addition, $\varepsilon_{\rm RDA}$ exclusively depends on $|u_{\rm D} - u_{\rm A}|$ (109 the $\delta = 0^{\circ}$ case) or $|u_{\rm D} + u_{\rm A}|$ (in the $\delta = 180^{\circ}$ case). 290 Moreover, these dependences are identical in both cases. The 291 typical results of simulation of coupling parameter changes 292 are shown in Fig. 6.

Relative changes of CCS depend on oscillation amplitudes with similar features and do not depend on φ 295

$$\varepsilon_{\sigma} = (u_{\rm D} \pm u_{\rm A})^2 / 2 r_{in}^2 = K_{\rm US}^{\rm DA} W_{\rm US},$$
 (15)

where "+" and "-" correspond to $\delta=180^\circ$ and $\delta=0^\circ$, 296 respectively, $K_{\rm US}^{\rm DA}$ characterizes the defect couple–ultrasound 297 interaction and depends on property defects as well as the 298 crystal matrix. Equation (15) takes into account that $u_{\rm D}, u_{\rm A}$ 299 $\propto \xi_{\rm US} \propto \sqrt{W_{\rm US}}$.

It is worth keeping in mind that CDLR current flows 301 locally in the locations of extended defects. At the same 302 time, the dislocations are often located perpendicularly to the 303 p-n junction plane in the SCR region, and the investigated 304 samples are not exception (see Sec. III C). If coupled defects 305 and dislocations are close to each other, then the dislocations 306 with the edge component should affect the pair spatial orientation. Thus, the axis of donor-acceptor pair with $(\Delta\Omega_d^D)$ 308 $\Delta\Omega_d^A>0$) should be predominantly parallel to the dislocation line, whereas the axis of a pair of coupled defects with 310

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> irradiated samples. In fact, in the case of 344 $(\Delta\Omega_d^{\rm D} \cdot \Delta\Omega_d^{\rm A} < 0)$, the US efficiency is determined by 345 the sum of pair component displacements, whereas in 346 the contrary case, it is determined by their difference. 347 In our opinion, both the donor and the acceptor are 348 defects of interstitial-type in the non-irradiated sam- 349

ple, and one of the pair components is a defect of 350 vacancy-type in irradiated samples. The defect config- 351

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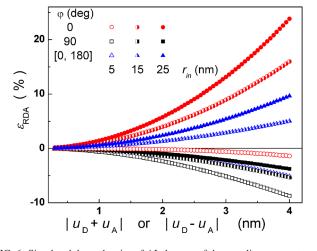


FIG. 6. Simulated dependencies of AI changes of the coupling parameter on the vibration amplitudes. Axis $|u_D - u_A|$ corresponds to the $\delta = 0^\circ$ case, whereas axis $|u_D + u_A|$ corresponds to the $\delta = 180^{\circ}$ case. The parameters are set to $a_0 = 3.23$ nm, $r_{in} = 5$ nm (open marks), 15 nm (semi-filled marks), and 25 nm (filled marks), and $\varphi = 0^{\circ}$ (circles) 90° (squares). The triangles correspond to the mean ε_{RDA} value for the $[0^{\circ} \div 180^{\circ}] \varphi$ range.

 $(\Delta\Omega_d^{\rm D} \cdot \Delta\Omega_d^{\rm A} < 0)$ should make a right angle with the dislocation line. As AW displacement is parallel to the p-n junction plane, the cases of most exciting interest are the 313 following:



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$$\begin{split} \delta &= 0\,^\circ, \;\; \phi = 90^\circ (\Delta\Omega_d^{\rm D} \cdot \Delta\Omega_d^{\rm A} > 0 \; {\rm case}); \;\; \delta = 180^\circ, \\ \phi &\in [0^\circ - 180^\circ] \big(\Delta\Omega_d^{\rm D} \cdot \Delta\Omega_d^{\rm A} < 0 \, {\rm case}\big). \end{split}$$

In other words, all curves in Fig. 6 can be realized if defect volume relaxation of the donor-like defect has the sign opposite to that of acceptor-like defect. Moreover, only squares should to be under consideration in the case of $\Delta\Omega_d^{\rm D} \cdot \Delta\Omega_d^{\rm A} > 0$.

Taking into account the experimental results and the estimation obtained from our model:

- $E_{\tau g}$ and $T_{\rm id}$ are mainly determined by couple compo-(i) nent energy levels. The alteration of $E_{\tau g}$ and $T_{\rm id}$ for nSC, g6SC, and g7SC in comparison with iSC testifies to the change of defect (donor, acceptor, or both) which take part in CDLR after irradiation. g6SC defects are coincident to g7SC defects and differ from the neutron-irradiated sample defect.
- USL causes the donor-acceptor distance change and 328 results in ε_{σ} and $\varepsilon_{\rm RDA}$, which increase with $W_{\rm US}$. 329
- Acoustically induced $E_{\tau g}$ (and T_{id}) modification, 330 (iii) 331 which is observed in g6SC and g7SC only, testifies to the rebuilding of γ -induced RD, i.e., γ -induced RD is 332 configurationally bistable (or metastable) and trans-333 forms from the ground state under US action. Similar 334 335 defect variations were previously.3,5,32,59 336
- ε_{σ} sign is immutable—see Eq. (15), whereas the ε_{RDA} 337 sign can vary for the pair with the opposite relaxation 338 339 volume component (see Fig. 6). Therefore, the change of $\Delta n_{\rm id}$ and $\varepsilon_{\tau g}$ sign is the evidence of transformation 340 from $(\Delta\Omega_d^{\mathrm{D}}\cdot \overset{\circ}{\Delta}\Omega_d^{\mathrm{A}}>0)$ to $(\Delta\Omega_d^{\mathrm{D}}\cdot \Delta\Omega_d^{\mathrm{A}}<0)$ after 341 irradiation. The transformation is confirmed by the 342 enhanced efficiency of US action on defects in 343

B. Quasi-neutral region

Base lifetime describes the processes which occur in the 354 quasi-neutral region of the p-n structure. Figure 7 shows τ_n 355 behaviour in the explored temperature range. As expected, 356 minority carrier lifetime increases as the temperature 357 increases, and at 320 K, τ_n values comprise 2–5 μ s for different samples, which correspond to the 80–130 µm range of 359 diffusion lengths. In our opinion, the observed τ_n dispersion 360 is caused not by irradiation but rather deals with 361 sample-ancestor wafer inhomogeneity, which is often the 362 case. 60,61

urations are discussed below, in Sec. III D.

In fact, the irradiation induced lifetime reduction is 364 described by the Messenger-Spratt equation⁴⁸ 365

$$\tau_n^{-1} = \tau_{n0}^{-1} + K_\tau \Psi \,, \tag{16}$$

where τ_{n0} is the minority carrier lifetime in the non- 366 irradiated sample and K_{τ} is a lifetime damage constant. The 367 known K_{τ} values and estimated changes of reciprocal base 368 lifetime $K_{\tau}\Psi$ are shown in Table V. As seen from the table, 369 the estimated value of the radiation-induced τ_n^{-1} change 370 comprises (8-17), 4, and 29% of the values measured for 371 samples nSC, g6SC, and g7SC, respectively, so this cannot 372 explain the dispersion observed experimentally. At the same 373 time, the calculated lifetime changes $K_{\tau}\Psi$ are in quite good 374 agreement with the changes expected from RDs produc- 375 tion—see Sec. III D. 376

Base lifetime can be expressed as follows:⁶⁴

$$\tau_n^{-1} = \tau_{\text{bb}}^{-1} + \tau_{\text{CEAuger}}^{-1} + \tau_{\text{SRH}}^{-1},$$
 (17)

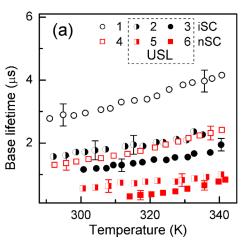
where $\tau_{bb},\,\tau_{CEAuger},\,\tau_{SRH}$ are the lifetimes of band-to-band 378 recombination, Coloumb-enhanced Auger recombination, 379 and SRH recombination, respectively. The calculation shows 380 that $\tau_{bb}^{-1} = 14 \text{ s}^{-1}$ and $\tau_{CEAuger}^{-1} = 6 \text{ s}^{-1}$. Therefore, band-381 to-band recombination and Auger recombination can be 382 neglected. In the case of the low injection level and single 383 recombination centre, SRH lifetime is described by Eq. (10). 384 If there are several centers of recombination, the following 385 equation should be applied:

$$\tau_n^{-1} = \sum_{i}^{M_d} \tau_{n,i}^{-1} = \sum_{i}^{M_d} N_{d,i\tau} \, \sigma_{n,i} \, v_{\text{th},n} \,, \quad \boxed{ (18)}$$

where M_d is the total number of centers, $\tau_{n,i}$ characterizes 387 lifetime due to recombination by the *i*-th defect, and $N_{d,i}$ and 388 $\sigma_{n,i}$ are the concentration and electron CCS of the *i*-th defect, 389 respectively.

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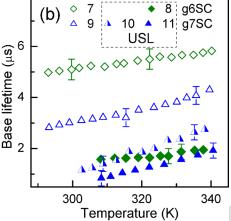


FIG. 7. Temperature dependences of base lifetime for non-irradiated (curves 1–3, circles), neutron-irradiated (4–6, squares), and γ -irradiated (7–11, diamonds and triangles) samples. The curves 1, 4, 7, and 9 (open marks) are obtained without USL, and curves 2, 3, 5, 6, 8, 10, and 11 correspond to Ui-1, Ui-2, Un-1, Un-2, Ug6-2, Ug7-1, and Ug7-2, respectively.

Figure 7 shows that USL results in a decrease in τ_n . Relative AI changes of piprocal base lifetime $\varepsilon_{\tau n} = (\tau_{n,in} - \tau_{n,US})/\tau_{n,US}$ are listed in Table IV. As AI changes are reversible, the lifetime alteration, in our opinion, deals with the increase in σ_n under US action. Following the empirical relation proposed by Ref. 65, we assume that Eq. (11) is valid for a complex point defect as well. In this case, however, r is the distance which separates the components of a complex. According to the model suggested in Sec. III A, USL leads to r variation and σ_n change in line with Eq. (15). In the case of CDLR, the AI change of the capture cross section of donor (or/and acceptor) is supplemental to the variation of both the coupling parameter and the couple distance, but only the CCS change determines the AI variation of base lifetime.

However, not every defect effectively takes part in AID. If $M_d^{\rm AA}$ and $M_d^{\rm nonAA}$ are the total numbers of acoustically active (AA) and non-acoustically active (non-AA) centers, Eq. (18) for τ_n^{-1} under USL and without it takes the following shape:



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$$\begin{split} \tau_{n,in}^{-1} &= \sum_{j}^{M_d^{\text{AA}}} N_{d,j\overline{\imath}} \, \sigma_{n,j}^{in} \, v_{\text{th},n} + \sum_{l}^{M_d^{\text{nonAA}}} N_{d,l\overline{\imath}} \, \sigma_{n,l} \, v_{\text{th},n} \,, \\ \tau_{n,\text{US}}^{-1} &= \sum_{j}^{M_d^{\text{AA}}} N_{d,j\overline{\imath}} \, \sigma_{n,j}^{\text{US}} \, v_{\text{th},n} + \sum_{l}^{M_d^{\text{nonAA}}} N_{d,l\overline{\imath}} \, \sigma_{n,l} \, v_{\text{th},n} \,. \end{split}$$

By using Eq. (15), $\varepsilon_{\tau n}$ is transformed as follows:

$$\varepsilon_{\tau n} = K_{\rm US}^{\rm eff} W_{\rm US} \,, \tag{19}$$

where $K_{\text{US}}^{\text{eff}}$ characterizes ADI in the sample and depends on the concentration of both AA and non-AA centers

TABLE V. Measured and estimated base lifetime parameters.

Sample	$\tau_{n,in}^{-1} (320 \mathrm{K})$	$K_{ au}$	$K_{\tau} \times \Psi$	$K_{ m US}^{ m eff}$
	$(10^5 \mathrm{s}^{-1})$	(cm ² /s)	$(10^4 \mathrm{s}^{-1})$	(cm ² /W)
iSC	2.9			3.5
nSC	4.7	$10^{-7}(\text{Ref. }33)$	4–8	7.1
		$2 \times 10^{-7} (\text{Ref. } 62)$		
g6SC	1.8	5×10^{-12}	0.8	6.0
g7SC	2.8	(Refs. 33 and 63)	8	5.2

$$K_{\rm US}^{\rm eff} = \sum_{j}^{M_d^{\rm AA}} \frac{\tau_{n,in}}{\tau_{n,j,in}} K_{{\rm US},j}$$
 (20)

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Here, $K_{\mathrm{US},j}$ deals with the j-th defect—ultrasound interaction. The obtained dependences of $\varepsilon_{\tau n}$ vs W_{US} are shown in 415 Fig. 8. The linearity of these dependences proves the correctness of our assumptions. The obtained $K_{\mathrm{US}}^{\mathrm{eff}}$ values are listed 417 in Table V. The non-monotonic $K_{\mathrm{US}}^{\mathrm{eff}}$ alteration with γ dose is 418 discussed in Sec. III D. 419

C. Shunt resistance

Figure 9 shows the shunt resistance over the explored 421 temperature range. As seen from the figure, the irradiation 422 results in a decrease in R_{sh} . Also, the R_{sh} temperature dependence changes after γ irradiation. In particular, the shunt 424 resistance decreases with the temperature growth in iSC and 425 nSC, whereas in g6SC and g7SC, the increase in R_{sh} vs T 426 is close to linear in the vicinity of 293 K. It should be 427 noted that the R_{sh} axis is logarithmic in Fig. 9(a) and linear 428 in Fig. 9(b).

The shunt resistance is known⁶⁶ to occur in the p-n 430 structure due to several non-mechanical reasons. It can be 431 caused by aluminum particles, macroscopic Si_3N_4 inclusions, 432

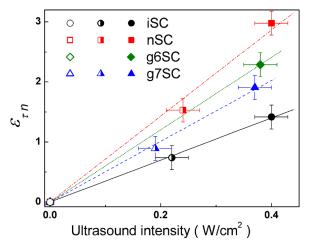


FIG. 8. Dependences of base lifetime relative change on US intensity for non-irradiated (circles), neutron-irradiated (squares), and γ -irradiated (triangles and diamonds) samples. Lines are the fitted curves using Eq. (19).

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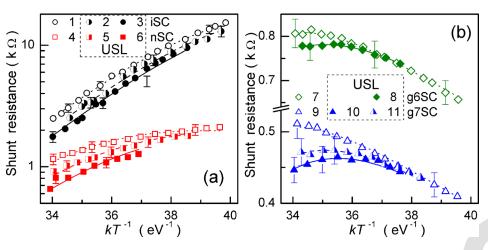


FIG. 9. Temperature dependences of shunt resistance for non-irradiated (curves 1-3, circles), neutron-irradiated (4–6, squares), and γ -irradiated (7–11, diamonds and triangles) samples. The curves 1, 4, 7, and 9 (open marks) are obtained without USL, and curves 2, 3, 5, 6, 8, 10, and 11 correspond to Ui-1, Ui-2, Un-1, Un-2, Ug6-2, Ug7-1, and Ug7-2, respectively. The marks are the experimental results, and the lines are the fitted curves using Eqs. (21)-(23).

or inversion layers at precipitates. In the course of firing, the All particle can penetrate into the sample creating the p^+ doped region around it, which compensates the emitter and remains in ohmic contact with the base. Inversion layers and Si₃N₄ inclusions occur mainly in multicrystalline silicon cells⁶⁶ and cannot cause shunt resistance in the investigated samples. Dislocations, however, which intersect the junction, are generally held responsible as a possible source of ohmic current.66-68 In our opinion, both aluminum particles and dislocations are present in the investigated structures, so the overall shunt resistance can be expressed as

$$R_{sh}^{-1} = R_{sh,Al}^{-1} + R_{sh,dis}^{-1}, (21)$$

where $R_{sh,Al}$ and $R_{sh,dis}$ deal with the aluminum particles and dislocations, respectively. The linear temperature depen-445 dence of metal particles $R_{sh,Al}$ is suggested by 446

$$R_{sh,A1} = R_{293,A1}[1 + \alpha(T - 293)],$$
 (22)

where $R_{293,\text{Al}}$ is the shunt resistance at 293 K and α is the 447 resistance temperature coefficient.

According to the model of dislocation-induced imped-449 ance of the photovoltaic detector suggested by Gopal and Gupta, ${}^{42,43}R_{sh,dis}$ can be given by

$$R_{sh,\text{dis}} = \frac{T}{\sigma_{\text{dis}}} \left[\cosh\left(\frac{E_{\text{dis}} - E_i}{kT}\right) + \cosh\left(\frac{U_s}{kT}\right) \right], \quad (23)$$

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$$\sigma_{\rm dis} = \rho_{\rm dis} A q^2 A_{\rm dis} \sqrt{K_n K_p} N_{\rm dis} (n_p + p_p) / k , \qquad (24)$$

where E_{dis} is the energy level which significantly contributes to the dislocation recombination current, U_s is the potential at the surface of the dislocation core, $\rho_{\rm dis}$ and $A_{\rm dis}$ are the dislocation density and surface area, respectively, K_n and K_p are the probabilities for electrons and holoptured by the dislocation states, and $N_{\rm dis}$ is the density of surface states at each dislocation. Equation (23) is true for the simplified case of $K_p = K_n$.

The resistance temperature coefficient was estimated from data on g7SC. The obtained value $8.3 \times 10^{-3} \text{ K}^{-1}$ is not far from the resistance temperature coefficient of bulk Al $(4.3 \times 10^{-3} \text{ K}^{-1})$. To fit the experimental data for R_{sh} , we

used Eqs. (21)–(23). As the fitting parameters, $R_{293,A1}$, 465 $(E_{\rm dis}-E_i)$, U_s , and $\sigma_{\rm dis}$ were taken. It has been found that 466 the experimental data are in good agreement with the fitting 467 curves (see Fig. 9) for values $(E_{\text{dis}} - E_i) = (0.46 \pm 0.02) \text{ eV}$ 468 and $U_s = (5\pm4) \times 10^{-8}$ eV, which were independent of irra-469 diation and USL. The obtained value of $(E_{\rm dis}-E_i)$ corre- 470 sponds to the carrier activation energy $0.10 \pm 0.02 \,\mathrm{eV}$ and is 471 comparable with the activation energy of dislocation levels 472 $0.08 \,\mathrm{eV}$, which was earlier reported $^{69-73}$ in Cz–Si:B too. $^{69-71}$ 473

The obtained values of $R_{293, \mathrm{Al}}$ and σ_{dis} are given in 474 Table III. $R_{293,A1}$ does not depend on USL and increases with 475 the irradiation level. In our opinion, $R_{sh,dis}$ is smaller than 476 $R_{sh,Al}$ in iSC. The irradiation facilitates the formation of 477 vacancies as well as Al diffusion out of the electrodes. As a 478 consequence, the number of Al particles increases, and $R_{sh,Al}$ 479 decreases and becomes the key factor contributing to the 480 overall shunt resistance. Al diffuses more effectively in the 481 samples exposed to γ radiation due to a more uniform distribution of irradiation-induced single vacancies.

Dispersion of $\sigma_{\rm dis}$ correlates with dispersion of τ_n over 484 the sample set. Hence, $\sigma_{\rm dis}$ dispersion deals with wafer inhomogeneity too. USL causes the $\sigma_{\rm dis}$ increase, and relative AI 486 changes $\varepsilon_{\sigma \rm dis} = (\sigma_{\rm dis,US} - \sigma_{\rm dis,in})/\sigma_{\rm dis,in}$ are shown in Table 487 IV. In our opinion, this is caused by an $A_{\rm dis}$ augmentation. 488 Namely, the dislocation core atom displacement is normal to 489 the current direction. As a result, the carriers are captured by 490 dislocation levels from enlarged volume. Therefore, the 491 effective surface area increases and $R_{sh,dis}$ decreases due to 492 US action.

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D. Defect type speculation

Lifetime killers in boron-doped Czochralski-grown Si 495 are boron-oxygen related (BO) defects, 74,75 iron-boron 496 pairs 64,76,77 (or another Fe-related trap in the n^+p junctions^{78,79}), and oxide precipitates. ^{60,61,64,80-82} The first two 498 defects are sensitive to intensive illumination at room tem- 499 perature. To determine the major recombination center in the 500 investigated samples, the following experimental procedure 501 was used. The non-irradiated sample was light soaked under 502 illumination using a halogen lamp (2 Suns) at approximately 503 305 K. The illumination time varied from 1 h to 8 h. After 504 illumination was terminated, the sample was exposed to 505 room temperature in the darkness. Over 5h period, I-V 506

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characteristics were measured with the interval of the parameters at room temperature. To estimate the permanent light-induced change, the measurements of I-V characteristics were performed in 48 h after illumination. After the total time under illumination ran up to 15 h, the iSC was annealed at 200 °C for 10 min in darkness, after which the measurements were carried out at room temperature. Then, the illumination and measurements were repeated.

Intensive light is known^{74,75} to cause permanent transformation of BO defects and considerable decrease in minority–carrier lifetime (as low as 10% of initial value at long term illumination). Annealing at 200 °C for 10 min in the darkness results in both the recovery of state and readiness to light-induced degradation of BO defects. Figure 10 shows the changes of structure parameters in comparison with those prior to illumination. As seen from the figure, illumination does not result in a considerable permanent change of τ_g , τ_n , and n_{id} before as well as after annealing. Therefore, the BO influence on recombination can be neglected in both the SCR and the base.

At the same time, the vast majority of impurity iron exists in iron–boron pairs. Fe_iB_s can be readily dissociated under intense illumination to release interstitial iron, which results in lifetime changes. In the darkness, Fe_iB_s is repaired and Fe_i concentration decreases according to 64,83

$$N_{Fe}(t) = (N_{Fe,0} - N_{Fe,eq}) \exp\left[-\frac{t}{\tau_{\text{rep}}}\right] + N_{Fe,eq},$$
 (25)

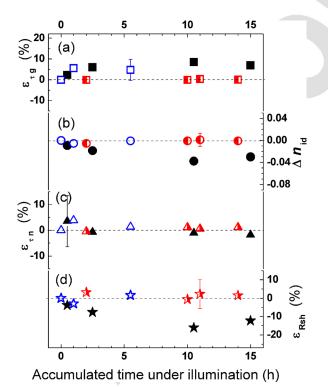


FIG. 10. Permanent changes of SCR lifetime [(a), squares], ideality factor [(b), circles], base lifetime [(c), triangles], and shunt resistance [(d), asterisks] versus accumulated illumination time. Sample iSC, $T\!=\!295\,\mathrm{K}$. Filled, semi-filled, and open marks correspond to the sample before annealing, after first 10 min 200 °C annealing, and after second 10 min 200 °C annealing, respectively.

where $N_{Fe,0}$ is the concentration immediately after illumination, and $N_{Fe,eq}$ is the equilibrium concentration which 534 remains for a long time after dissociation; the characteristic 535 time of repairing τ_{rep} depends on the doping level 536

$$\tau_{\text{rep}} = 770 \cdot p_p^{-2/3} \exp\left(\frac{E_{\text{D,Fe}}}{kT}\right),\tag{26}$$

where $E_{\rm D,Fe}=0.68~{\rm eV}$ is the activation energy of Fe $_i$ 537 diffusion.

It was found that $n_{\rm id}$ increased (by about 0.03) and τ_g 539 decreased (by about 10%) immediately after illumination— 540 see Fig. 11(a). These changes vanished gradually. We supposed that τ_g and $n_{\rm id}$ evolutions could be described by 542 expressions similar to Eq. (25). The obtained Eq. (26) was used to calculate characteristic time, and the fitting lines are 544 presented in Fig. 11(a). The fittings with $E_{\rm D,Fe}=0.68$ eV are 545 in good agreement with the experimental data. Hence, it is 546 evident that iron–boron pairs take part in SCR recombination. At the same time, electron and hole CCS of Fe $_i$ are 1.7 548 and 0.04 times 64 as much as those of Fe $_i$ B $_s$. A small τ_g alteration (by about 10%) caused by light is the evidence of the 550 supporting role of iron–boron pair in SCR recombination. 551 Furthermore, since τ_n does not depend on illumination [see 552 Fig. 11(b)], Fe $_i$ B $_s$ does not influence the base lifetime. 553

Thus, a conclusion can be made that oxide precipitates 554 are number one agents in SCR and base recombinations. 555 According to Murphy *et al.*^{80,81} there exist at least two independent oxide precipitate related defects. These defects have 557 $\sigma_n/\sigma_p=157$ and $\sigma_p/\sigma_n=1200$, respectively, 81 which is 558 suitable for CDLR. These facts allow us to conclude that the 559 defect responsible for AI phenomena in iSC is mainly oxide 560 precipitate.

In foreseeing the RD type, it is worth keeping in mind 562 the doping level, oxygen concentration, and irradiation dose. 563 In our case (Czochralski, oxygen-rich, $\sim 7 \times 10^{17}$ cm⁻³, p-Si 564 with boron concentration $\sim 10^{15}$ cm⁻³, and low dose), it is 565 expected that C_iO_i , vacancy clusters V_n (divacancy V_2 , trivacancy V_3 ,...) and VO_i are produced mainly by neutron irradiation V_i and V_i are produced by V_i -rays. V_i The RD concentration V_i and V_i are produced by V_i -rays. V_i irradiation in V_i -rays. V_i -rays irradiation in V_i -rays shown in Table VI. The expected 571 alues of V_i -ray for the investigated samples are listed in 572 Table VI as well.

The other defects that can be created by irradiation in 574 silicon are the I_p center, bistable donor (BD), B_iO_i , and C_iC_s . 575 At the same time, the I_p center and BD are characterized by 576 a small introduction rate. For example, the expected 85,93 concentration of BD is only $(1-2) \times 10^{10}$ cm $^{-3}$ in nSC and 578 g7SC. The lack of B_iO_i in the investigated samples deals 579 with the low boron concentration. 94 The formation of C_iC_s is 580 suppressed in the oxygen-rich crystal 84,87,88 and, what is 581 more, C_iC_s is not an active recombination center. 95

The influence of RD on base lifetime could be estimated 583 by Eq. (18) taking into account the fact that VO_i is a recombination center which is not active in p-Si. $^{63,96-99}$ The estimated $\tau_{n,\text{RD}}$ for C_iO_i , V_2 , and V_3 are shown in Table VI. As 586 seen from the table, τ_n is effected mainly by C_iO_i in 587

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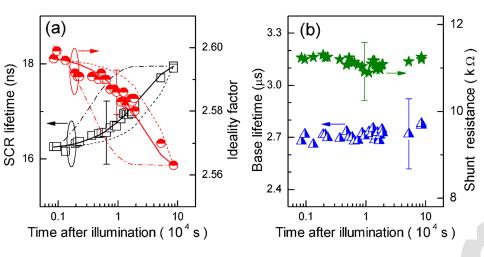


FIG. 11. SCR lifetime [(a), squares, left axis], ideality factor [(a), circles, right axis], base lifetime [(b), triangles, left axis], and shunt resistance [(b), asterisks, right axis] as a function of time since illumination has stopped. Sample iSC, T = 295 K. Lines are calculated by using Eqs. (25) and (26) and $E_{\rm D,Fe} = 0.63$ eV (dashed–dotted lines), 0.68 eV (solid lines), and 0.73 eV (dashed lines).

TABLE VI. Cited and calculated defect parameters.

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	_	$\eta_n (\text{cm}^{-1})$		Λ	$N_{t,\text{RD}}(10^{11} \text{ cm}^{-3})$			$\tau_{n,RD}^{-1}(10^4 \text{ s}^{-1})$	
Defect	$(10^{-15} \mathrm{cm}^2)$	Reference 86	$oldsymbol{\eta}_{\gamma}$	nSC	g6SC	g7SC	nSC	g6SC	g7SC
C_iO_i	0.7 (Ref. 87) 0.9 (Ref. 88)	1.38	$6 \times 10^5 \text{ rad}^{-1} \text{ cm}^{-3} \text{ (Ref. 87)}$ $4 \times 10^{-4} \text{ cm}^{-1} \text{ (Ref. 88)}$	5.5	6	60	0.8–1	0.9–1.1	9–11
V_2	3 (Ref. 87) 2 (Ref. 90)	1.21	$3 \times 10^4 \text{rad}^{-1} \text{cm}^{-3} (\text{Ref. 87})$	4.8	0.3	3	2.2–3.3	0.1-0.2	1–2
V_3	2.4 (Ref. 91)	0.37		1.5			0.7		
VO _i	2.4 (Ref. 89) 4 (Ref. 92)	0.52	$7 \times 10^{5} \text{ rad}^{-1} \text{ cm}^{-3} \text{ (Ref. 87)}$ $4 \times 10^{-4} \text{ cm}^{-1} \text{ (Ref. 88)}$	2	6–7	60–70			

 γ -irradiated samples and by vacancy clusters in nSC. It should be noted that nSC, g6SC, and g7SC sums of $\tau_{n,\text{RD}}^{-1}$ are in quite good agreement with $(K_{\tau} \cdot \Psi)$ values.

We shall now consider $K_{\text{US}}^{\text{eff}}$ for the non-irradiated sample assuming $M_d^{\text{AA}} = 1$ and $M_d^{\text{nonAA}} = 1$. We shall also assume that US interactions with C_iO_i and V_n are described by $K_{\text{US}}^{\text{CO}}$ and K_{US}^{V} , respectively. Then, Eq. (20) gives the following expression for $K_{\text{US}}^{\text{eff}}$ in iSC and irradiated samples:

$$\begin{split} K_{\mathrm{US}}^{\mathrm{eff}} &= K_{\mathrm{US}}^{\mathrm{AA}} \, \tau_{n,in} / \tau_{n,in}^{\mathrm{AA}}, \\ K_{\mathrm{US}}^{\mathrm{eff}} &= K_{\mathrm{US}}^{\mathrm{AA}} \tau_{n,in} / \tau_{n,in}^{\mathrm{AA}} + K_{\mathrm{US}}^{\mathrm{CO}} \tau_{n,in} / \tau_{n,\mathrm{RD}}^{\mathrm{CO}} + K_{\mathrm{US}}^{\mathrm{V}} \tau_{n,in} / \tau_{n,\mathrm{RD}}^{\mathrm{V}}. \end{split}$$

Here, $au_{n,in}^{AA}$ is the base lifetime in the sample with the sole non-radiative AA defect and K_{US}^{AA} describes ADI.

For the analysis, the following two limit cases are appropriate. In the first one, non-AA defects are distributed uniformly across the wafer, and AA defects define $(\tau_{n,in}^{-1} - K_{\tau} \cdot \Psi)$ values in different samples. In the second one, a non-AA defect distribution is not uniform, and $\tau_{n,in}^{AA}$ is identical for iSC, nSC, g6SC, and g7SC. However, in the first case (as well as in the case of $M_d^{\text{nonAA}} = 0$), the experimental values of $K_{\text{US},j}^{\text{eff}}$ lead to unreal (negative) values of $K_{\text{US},j}$. In the second case, Eq. (20) and the data from Tables V and VI give the following array equations:

iSC:
$$3.5 = K_{\text{US}}^{\text{AA}} \cdot (\tau_{n,in}^{\text{AA}})^{-1} / 2.9$$
,
nSC: $7.1 = K_{\text{US}}^{\text{AA}} \cdot (\tau_{n,in}^{\text{AA}})^{-1} / 4.7 + 0.09 K_{\text{US}}^{\text{V}} + 0.02 K_{\text{US}}^{\text{CO}}$,
g6SC: $6.0 = K_{\text{US}}^{\text{AA}} \cdot (\tau_{n,in}^{\text{AA}})^{-1} / 1.8 + 0.01 K_{\text{US}}^{\text{V}} + 0.05 K_{\text{US}}^{\text{CO}}$,
g7SC: $5.2 = K_{\text{US}}^{\text{AA}} \cdot (\tau_{n,in}^{\text{AA}})^{-1} / 2.8 + 0.05 K_{\text{US}}^{\text{V}} + 0.35 K_{\text{US}}^{\text{CO}}$,

where $(\tau_{n,in}^{AA})^{-1}$ in 10^4 /s. These equations are valid for K_{US}^{AA} · 608 $(\tau_{n,in}^{AA})^{-1} = (10\pm3)~{\rm cm}^2/{\rm W},~K_{US}^{\rm V} = (42\pm15)~{\rm cm}^2/{\rm W},~{\rm and}~{\rm 609}$ $K_{US}^{\rm CO} = 0$. Since $(\tau_{n,in}^{AA})^{-1} < 1.83,~K_{US}^{AA} > 5~{\rm cm}^2/{\rm W}$. Thus, the 610 observed change in base lifetime is caused by AI modification of the same defect (most likely oxide precipitates) in 612 both non-irradiated and γ -irradiated samples. This effect is 613 enhanced by AI alteration of the divacancy in neutron-614 irradiated samples. In other words, C_iO_i is the non-AA 615 defect, whereas V_2 is the AA defect.

In our opinion, under US action, τ_g and $n_{\rm id}$ in the non- 617 irradiated sample depend on the modification of coupled 618 oxide precipitate related defects. As assumed in Sec. III A, in 619 irradiated samples, the AA radiation defects with $\Delta\Omega_d < 0$ 620 take part in CDLR. The divacancy is a quite suitable expla- 621 nation for the AI influence on τ_g and $n_{\rm id}$ in nSC, but in γ -irra-622 diated samples, a bistable (or metastable) defect is expected. 623 A few similar defects with $\Delta\Omega_d < 0$ are known in Si, viz, 624 VO_2 , ¹⁰⁰ V_3 , ⁹¹ and VO_i . ¹⁰¹ VO_2 is formed after 300 °C 625 annealing of the irradiated crystal, V₃ is not the typical 626 defect for γ -60Co exposed silicon, while VO_i is largo 627 manum produced and can take part in CDLR around n^+ -p 628 interface in g6SC and g7SC. The metastable state commonly 629 observed at low temperature is remarkable for the large oxy- 630 gen-vacancy distance and deeper energy level. 101 The vol- 631 ume change of the entire complex is negative, whereas for 632 the complex component, $\Delta\Omega_d(V) < 0$ and $\Delta\Omega_d(O_i) > 0$. 633 Hence, under the assumption made, VO; is a favorable pair 634 for AI alteration of the distance between components and 635 therefore can be transformed into the metastable configura- 636 tion by USL, which causes changes in both T_{id} and $E_{\tau g}$.

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IV. CONCLUSION 638

639 The influence of ultrasound on I-V characteristics of non-irradiated silicon n^+ -p structures as well as silicon 640 structures exposed to reactor neutrons or ⁶⁰Co gamma radia-641 tion has been investigated experimentally. The investigation 642 has revealed an acoustically driven reversible decrease in 643 both the minority carrier lifetime and shunt resistance in the structure base. The effect is intensified in the irradiated struc-645 tures. The analysis has shown that these effects are caused by the acoustically induced increase in the carrier capture 647 coefficient for point or extended defects. It has also been found that ultrasound loading leads to the reversible modifi-649 cation of the SCR carrier lifetime and ideality factor. The 650 changes are opposite in non-irradiated and irradiated struc-651 tures. The qualitative model of the observed phenomenon, which is based on the increase in the distance between cou-653 pled defects or between complex defect components due to 654 ultrasound action, has been suggested. It has been shown that 655 interstitial carbon-interstitial oxygen complexes practically do not take part in acousto-defect interactions, whereas the 657 divacancy in neutron-exposed structures and vacancy-inter-658 stitial oxygen pairs in γ -exposed structures can be effectively 659 modified by applying ultrasound. Thus, ultrasound can be an 660 effective tool for controlling silicon structure characteristics. 661 662

- ¹M. Jivanescu, A. Romanyuk, and A. Stesmans, J. Appl. Phys. 107, 664 114307 (2010).
- ²A. Romanyuk, P. Oelhafen, R. Kurps, and V. Melnik, Appl. Phys. Lett. 665 90, 013118 (2007). 666
- 667 ³I. A. Buyanova, S. S. Ostapenko, M. K. Sheinkman, and M. Murrikov, 668 Semicond. Sci. Technol. 9, 158 (1994).
- 669 ⁴O. Korotchenkov and H. Grimmliss, Phys. Rev. B **52**, 14598 (1995).
- 670 ⁵O. Y. Olikh, Semiconductors **43**, 745 (2009).
- 671 ⁶S. S. Ostapenko and R. E. Bell, J. Appl. Phys. 77, 5458 (1995).
- 672 ⁷I. Ostrovskii, O. Korotchenkov, O. Olikh, A. Podolyan, R. Chupryna, and 673 M. T. Cisneros, J. Opt. A: Pure Appl. Opt. 3, S82 (2001).
- 674 ⁸D. Kropman, V. Seeman, S. Dolgov, and A. Medvids, *Phys. Status Solidi* 675 C 13, 793 (2016).
- 676 ⁹N. Zaveryukhina, E. Zaveryukhina, S. Vlasov, and B. Zaveryukhin, Tech. 677 Phys. Lett. 34, 241 (2008).
- ¹⁰S. A. Mirsagatov, I. B. Sapaeva, and Z. Nazarov, Inorg. Mater. 51, 1 678 679 (2015).
- 680 ¹¹R. Savkina, A. Smirnov, T. Kryshtab, and A. Kryvko, Mater. Sci. 681 Semicond. Process. 37, 179 (2015). ¹²M. Virot, R. Pflieger, E. V. Skorb, J. Ravaux, T. Zemb, and H. Mohwald, 682
- 683 J. Phys. Chem. C 116, 15493 (2012). 684
- ¹³O. Y. Olikh, K. V. Voytenko, R. M. Burbelo, and J. M. Olikh, 685 J. Semicond. 37, 122002 (2016).
- 686 ¹⁴O. Olikh, Semiconductors **45**, 798 (2011).
- 687 ¹⁵A. Davletova and S. Z. Karazhanov, J. Phys. Chem. Solids 70, 989 688 (2009)
- ¹⁶A. Davletova and S. Z. Karazhanov, J. Phys. D: Appl. Phys. 41, 165107 689 690 (2008).
- ¹⁷V. Melnik, Y. Olikh, V. Popov, B. Romanyuk, Y. Goltvyanskii, and A. 691 692 Evtukh, Mater. Sci. Eng., B 124-125, 327 (2005).
- 693 ¹⁸O. Y. Olikh, K. V. Voytenko, and R. M. Burbelo, J. Appl. Phys. 117, 694 044505 (2015).
- 695 ¹⁹O. Olikh, Ultrasonics **56**, 545 (2015).
- ²⁰V. N. Pavlovich, Phys. Status Solidi B **180**, 97 (1993). 696
- ²¹F. Mirzade, J. Appl. Phys. **110**, 064906 (2011).
- ²²R. Peleshchak, O. Kuzyk, and O. Dan'kiv, Ukr. J. Phys. **61**, 741 (2016). 698
- 699 ²³V. D. Krevchik, R. A. Muminov, and A. Y. Yafasov, Phys. Status Solidi A 700 63, K159 (1981).
- 701 ²⁴F. Mirzade, J. Appl. Phys. **97**, 084911 (2005).
- ²⁵I. Ostrovskii and O. Korotchenkov, Solid State Commun. **82**, 267 (1992). 702
- 703 ²⁶O. Olikh and K. Voytenko, Ultrasonics **66**, 1 (2016).
- ²⁷N. Guseynov, Y. Olikh, and S. Askerov, Tech. Phys. Lett. 33, 18 (2007). 704

²⁸P. Parchinskii, S. Vlasov, and L. Ligai, Semiconductors **40**, 808 (2006).

Total Pages: 13

- ²⁹A. Gorb, O. Korotchenkov, O. Olikh, and A. Podolian, IEEE Trans. Nucl. Sci. 57, 1632 (2010).
- ³⁰A. O. Podolian, A. B. Nadtochiy, and O. A. Korotchenkov, Tech. Phys. Lett. 38, 405 (2012).
- ³¹Y. Olikh, M. Tymochko, and A. Dolgolenko, Tech. Phys. Lett. **32**, 586
- ³²Y. Olikh and M. Tymochko, Tech. Phys. Lett. **37**, 37 (2011).
- ³³H. Jafari and S. Feghhi, Nucl. Instrum. Methods Phys. Res., Sect. A 816, 713 62 (2016).
- ³⁴Y. P. Rao, K. Praveen, Y. R. Rani, A. Tripathi, and A. G. Prakash, Nucl. Instrum. Methods Phys. Res., Sect. B 316, 205 (2013).
- ³⁵M. Moll, H. Feick, E. Fretwurst, G. Lindström, and C. Schütze, Nucl. Instrum. Methods Phys. Res., Sect. A 388, 335 (1997)
- ³⁶J. Srour, C. Marshall, and P. Marshall, IEEE Trans. Nucl. Sci. **50**, 653 (2003).
- ³⁷I. Pintilie, G. Lindstroem, A. Junkes, and E. Fretwurst, Nucl. Instrum. Methods Phys. Res., Sect. A 611, 52 (2009).
- ³⁸N. Arutyunov, N. Bennett, N. Wight, R. Krause-Rehberg, V. Emtsev, N. Abrosimov, and V. Kozlovski, Phys. Status Solidi B 253, 2175 (2016).
- ³⁹C. A. Londos, G. Antonaras, and A. Chroneos, J. Appl. Phys. 114, 193513 (2013).
- ⁴⁰A. Schenka and U. Krumbein, J. Appl. Phys. **78**, 3185 (1995).
- ⁴¹S. Steingrube, O. Breitenstein, K. Ramspeck, S. Glunz, A. Schenk, and P. P. Altermatt, J. Appl. Phys. 110, 014515 (2011).
- ⁴²V. Gopal and S. Gupta, IEEE Trans. Electron Devices **50**, 1220 (2003).
- ⁴³V. Gopal and S. Gupta, IEEE Trans. Electron Devices **51**, 1078 (2004). ⁴⁴A. Akkerman, J. Barak, M. Chadwick, J. Levinson, M. Murat, and Y.
- Lifshitz, Radiat. Phys. Chem. 62, 301 (2001). ⁴⁵D. Bräunig and F. Wulf, Radiat. Phys. Chem. **43**, 105 (1994).
- ⁴⁶A. B. Sproul and M. A. Green, J. Appl. Phys. **73**, 1214 (1993).
- ⁴⁷D. K. Schroder, Semiconductor Material and Device Characterization,
- 3rd ed. (John Wiley & Sons, New Jersey, 2006).
- ⁴⁸Solar Cells Materials, Manufacture and Operation, 2nd ed., edited by A. McEvoy, T. Markvart, and L. Castaner (Academic Press, Oxford, 2013).
- ⁴⁹K. Wang and M. Ye, Solid-State Electron. **53**, 234 (2009).
- ⁵⁰D. Schroder, IEEE Trans. Electron Devices **29**, 1336 (1982).
- ⁵¹H. Aharoni, T. Ohmi, M. M. Oka, A. Nakada, and Y. Tamai, J. Appl. Phys. 81, 1270 (1997).
- ⁵²A. S. H. van der Heide, A. Schonecker, J. H. Bultman, and W. C. Sinke, Prog. Photovoltaics: Res. Appl. 13, 3 (2005).
- ⁵³J. Beier and B. Voss, in *Proceedings of the 23rd IEEE Photovoltaic* Specialists Conference, Louisville, KY, USA (1993), pp. 321–326.
- ⁵⁴J. M. Shah, Y.-L. Li, T. Gessmann, and E. F. Schubert, J. Appl. Phys. 94, 2627 (2003).
- ⁵⁵A. Kaminski, J. J. Marchand, H. E. Omari, A. Laugier, Q. N. Le, and D. Sarti, in Proceedings of the 25th IEEE Photovoltaic Specialists Conference, Washington, DC, USA (1996), pp. 573-576.
- ⁵⁶W. M. Chen, B. Monemar, E. Janzén, and J. L. Lindström, Phys. Rev. Lett. 67, 1914 (1991).
- ⁵⁷A. M. Frens, M. T. Bennebroek, A. Zakrzewski, J. Schmidt, W. M. Chen, E. Janzén, J. L. Lindström, and B. Monemar, Phys. Rev. Lett. 72, 2939 (1994).
- ⁵⁸O. Breitenstein, J. Bauer, P. P. Altermatt, and K. Ramspeck, Solid State Phenom. 156-158, 1 (2010).
- ⁵⁹T. Wosinski, A. Makosa, and Z. Witczak, Semicond. Sci. Technol. 9, 2047 (1994).
- ⁶⁰L. Chen, X. Yu, P. Chen, P. Wang, X. Gu, J. Lu, and D. Yang, Sol. Energy Mater. Sol. Cells 95, 3148 (2011).
- ⁶¹J. Schön, A. Youssef, S. Park, L. E. Mundt, T. Niewelt, S. Mack, K. Nakajima, K. Morishita, R. Murai, M. A. Jensen, T. Buonassisi, and M. C. Schubert, J. Appl. Phys. **120**, 105703 (2016).
- ⁶²E. Gaubas, A. Uleckas, and J. Vaitkus, Nucl. Instrum. Methods Phys. Res., Sect. A 607, 92 (2009).
- ⁶³I. I. Kolkovskii, P. F. Lugakov, and V. V. Shusha, Phys. Status Solidi A 83, 299 (1984).
- ⁶⁴J. D. Murphy, K. Bothe, M. Olmo, V. V. Voronkov, and R. J. Falster, J. Appl. Phys. 110, 053713 (2011).
- ⁶⁵D. G. Thomas, J. Hopfield, and W. M. Augistyniak, Phys. Rev. 140, A202 (1965).
- ⁶⁶O. Breitenstein, J. P. Rakotoniaina, M. H. Al Rifai, and M. Werner, Prog. Photovoltaics: Res. Appl. 12, 529 (2004).
- ⁶⁷V. Gopal, J. Appl. Phys. **116**, 084502 (2014).
- ⁶⁸I. Baker and C. Maxey, J. Electron. Mater. **30**, 682 (2001).

AQ3

705

706

707

708

709

710

711

712

714

715

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717

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720

721 722

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771

772

773 774

775

776

777

778

000000-12 Olikh et al.

820

J. Appl. Phys. 123, 000000 (2018)

779	⁶⁹ A. Castaldini, D. Cavalcoli, A. Cavallini, and S. Pizzini, Phys. Rev. Lett.
780	95 , 076401 (2005).
781	⁷⁰ I. Isakova, A. Bondarenko, O. Vyvenko, V. Vdovin, E. Ubyivovk, and O.
782	Kononchuk, J. Phys.: Conf. Ser. 281, 012010 (2011).
783	⁷¹ X. Yu, O. Vyvenko, M. Kittler, W. Seifert, T. Mtchedlidze, T. Arguirov,
784	and M. Reiche, Semiconductors 41, 458 (2007).
785	⁷² V. Kveder, M. Kittler, and W. Schröter, Phys. Rev. B 63 , 115208 (2001).
786	⁷³ M. Trushin, O. Vyvenko, T. Mchedlidze, O. Kononchuk, and M. Kittler,
787	Solid State Phenom. 156–158 , 283 (2010).
788	⁷⁴ J. Lindroos and H. Savin, Sol. Energy Mater. Sol. Cells 147 , 115 (2016).
789	⁷⁵ T. Niewelt, J. Schöon, W. Warta, S. W. Glunz, and M. C. Schubert, IEEE
790	J. Photovoltaics 7, 383 (2017).
791	⁷⁶ V. Vahanissi, A. Haarahiltunen, H. Talvitie, M. Yli-Koski, and H. Savin,
792	Prog. Photovoltaics: Res. Appl. 21 , 1127 (2012).
793	⁷⁷ J. Schmidt, Prog. Photovoltaics: Res. Appl. 13 , 325 (2005).
) 94	⁷⁸ T. Mchedlidze and J. Weber, Phys. Status Solidi B 251 , 1608 (2014).
95	⁷⁹ T. Mchedlidze, L. Scheffler, J. Weber, M. Herms, J. Neusel, V. Osinniy,
796	C. Moller, and K. Lauer, J. Appl. Phys. 103 , 013901 (2013).
797	⁸⁰ J. Murphy, J. McGuire, K. Bothe, V. Voronkov, and R. Falster, Sol.
798	Energy Mater. Sol. Cells 120 , 402 (2014).
799	⁸¹ J. D. Murphy, K. Bothe, R. Krain, V. V. Voronkov, and R. J. Falster,
800	J. Appl. Phys. 111 , 113709 (2012).
801	⁸² M. Porrini and P. Tessariol, Mater. Sci. Eng. B 73 , 244 (2000).
802	⁸³ W. Wijaranakula, J. Etectrochem. Soc. 140 , 275 (1993).
803	⁸⁴ G. Lindström, M. Ahmed, S. Albergo, P. Allport, D. Anderson, L.
804	Andricek, M. Angarano, V. Augelli, N. Bacchetta, P. Bartalini, R. Bates,
805	U. Biggeri, G. Bilei, D. Bisello, D. Boemi, E. Borchi, T. Botila, T.
806	Brodbeck, M. Bruzzi, T. Budzynski, P. Burger, F. Campabadal, G. Casse,
807	E. Catacchini, A. Chilingarov, P. Ciampolini, V. Cindro, M. Costa, D.
808	Creanza, P. Clauws, C. Da Via, G. Davies, W. De Boer, R. DellOrso, M.
809	D. Palma, B. Dezillie, V. Eremin, O. Evrard, G. Fallica, G. Fanouraki, H.
810	Feick, E. Focardi, L. Fonseca, E. Fretwurst, J. Fuster, K. Gabathuler, M.
811	Glaser, P. Grabiec, E. Grigorievt, G. Hall, M. Hanlond, F. Hauler, S.
812	Heising, A. Holmes-Siedle, R. Horisberger, G. Hughes, M. Huhtinen, I.
813	Ilyashenko, A. Ivanov, B. Jones, L. Jungermann, A. Kaminsky, Z.
814	Kohout, G. Kramberger, M. Kuhnke, S. Kwan, F. Lemeilleur, C. Leroy,
815	M. Letheren, Z. Li, T. Ligonzo, V. Linhart, P. Litovchenko, D. Loukas,
816	M. Lozano, Z. Luczynski, G. Lutz, G. MacEvoy, S. Manolopoulos, A.
817	Markou, C. Martinez, A. Messineo, M. Mikuž, M. Moll, E.
818	Nossarzewska, G. Ottaviani, V. Oshea, G. Parrini, D. Passeri, D. Petre, A.
819	Pickford, I. Pintilie, L. Pintilie, S. Pospisil, R. Potenza, C. Raine, J. Rafi,

P. Ratoff, H. Richter, P. Riedler, S. Roe, P. Roy, A. Ruzin, A. Ryazanov,

```
A. Santocchia, L. Schiavulli, P. Sicho, I. Siotis, T. Sloan, W. Slysz, K. 821
  Smith, M. Solanky, B. Sopko, S. K. B. Sundby Avset, S. B. C. Tivarus, 822
  G. Tonelli, A. Tricomi, S. Tzamarias, G. Valvo, A. Vasilescu, A. Vayaki, 823
  E. Verbitskaya, P. Verdini, V. Vrba, S. Watts, E. Weber, M. Wegrzecki, 824
  I. Wegrzecka, P. Weilhammer, R. Wheadon, C. Wilburn, I. Wilhelm, R. 825
   Wunstorf, J. Wüstenfeld, J. Wyss, K. Zankel, P. Zabierowski, and D. 826
                                                                                827
  Žontar, Nucl. Instrum. Methods Phys. Res., Sect. A 466, 308 (2001).
 <sup>85</sup>I. Pintilie, G. Lindstroem, A. Junkes, and E. Fretwurst, Nucl. Instrum. 828
   Methods Phys. Res., Sect. A 611, 52 (2009).
                                                                                829
 <sup>86</sup>M. Moll, "Radiation damage in silicon particle detectors: Microscopic
                                                                                830
  defects and macroscopic properties," Ph.D. thesis (Universität Hamburg,
                                                                                831
                                                                                832
 <sup>87</sup>J. Stahl, E. Fretwurst, G. Lindström, and I. Pintilie, Nucl. Instrum. 833
                                                                                834
  Methods Phys. Res., Sect. A 512, 111 (2003).
 <sup>88</sup>I. I. Kolkovskii and V. Lukyanitsa, Semiconductors 31, 340 (1997).
                                                                                835
<sup>89</sup>R. Siemieniec, W. Sudkamp, and J. Lutz, in Proceedings of the Fourth 836
  IEEE International Caracas Conference on Devices, Circuits and 837
                                                                                838
  Systems (Oranjestad, Aruba, Netherlands, 2002), pp. D029-1-D029-6.
<sup>90</sup>S. D. Brotherton and P. Bradley, J. Appl. Phys. 53, 5720 (1982).
                                                                                839
<sup>91</sup>V. P. Markevich, A. R. Peaker, S. B. Lastovskii, L. I. Murin, J. Coutinho, 840
  V. J. B. Torres, P. R. Briddon, L. Dobaczewski, E. V. Monakhov, and B. 841
                                                                                842
  G. Svensson, Phys. Rev. B 80, 235207 (2009).
 <sup>92</sup>H. Bleichner, P. Jonsson, N. Keskitalo, and E. Nordlander, J. Appl. Phys.
                                                                                843
                                                                                844
  79, 9142 (1996).
<sup>93</sup>E. Fretwurst, F. Hönniger, G. Kramberger, G. Lindström, I. Pintilie, and <sup>845</sup>
                                                                                846
  R. Röder, Nucl. Instrum. Methods Phys. Res., Sect. A 583, 58 (2007).
 <sup>94</sup>L. C. Kimerling, M. Asom, J. Benton, P. Drevinsky, and C. Caefer, <sup>847</sup>
                                                                                848
  Mater. Sci. Forum 38-41, 141 (1989).
<sup>95</sup>L. W. Song, X. D. Zhan, B. W. Benson, and G. D. Watkins, Phys. Rev. B 849
                                                                                850
  42, 5765 (1990).
 <sup>96</sup>J. L. Benton, S. Libertino, P. Kringhøj, D. J. Eaglesham, J. M. Poate, and 851
  S. Coffa, J. Appl. Phys. 82, 120 (1997).
                                                                                852
<sup>97</sup>S. Coffa, V. Privitera, F. Priolo, S. Libertino, and G. Mannino, J. Appl. 853
                                                                                854
  Phys. 81, 1639 (1997).
<sup>98</sup>N. Ganagona, B. Raeissi, L. Vines, E. V. Monakhov, and B. G. Svensson,
                                                                                855
   Phys. Status Solidi C 9, 2009 (2012).
                                                                                856
<sup>99</sup>L. Vines, E. V. Monakhov, A. Y. Kuznetsov, R. Kozłowski, P. Kaminski, 857
  and B. G. Svensson, Phys. Rev. B 78, 085205 (2008).
                                                                                858
100 L. I. Murin, V. P. Markevich, I. F. Medvedeva, and L. Dobaczewski, 859
  Semiconductors 40, 1282 (2006).
                                                                                860
```

¹⁰¹B. N. Mukashev, K. A. Abdullin, and Y. V. Gorelkinskii, Phys.–Usp. 43, 861

862

139 (2000).