

EFFECT OF DEFECTS ORIGINATING UNDER THE PROTON IRRADIATION ON THE ELECTROPHYSICAL AND DETECTOR PROPERTIES OF CdTe:Cl AND CdZnTe

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The work is dedicated to studying by computer modeling the mechanisms of the influence of radiation defects, originating under high energy proton irradiation, on the resistivity ρ , lifetime of nonequilibrium electrons τ_n and holes τ_p in CdTe:Cl and Cd_{0.9}Zn_{0.1}Te, and charge collection efficiency η of room temperature ionizing radiation detectors based on these materials. The effect of recombination at deep levels of radiation defects on the degradation of τ_n , τ_p , and η of detectors based on CdTe:Cl and Cd_{0.9}Zn_{0.1}Te was studied. Energy levels of radiation defects also substantially effect on compensation degree of semiconductor decreasing ρ . The main factors affecting the abrupt or gradual decrease in the resistivity and charge collection efficiency of these detectors during their bombardment by high-energy protons, leading to complete degradation of their recording ability, were found. The important role of purity and deep donor concentration in initial state of the detector material was indicated.

PASC: 61.80.-x; 61.82.Fk; 87.66.Pm

INTRODUCTION

CdTe and CdZnTe are traditionally considered as semiconductor materials for X-ray and gamma-ray high energy resolution detectors operating at room temperature. At present the detectors have been developed for industrial tomography and astrophysics, as well as for medical imaging equipment. Ions irradiation can generate electron-hole pairs in semiconductors, and the above-mentioned materials are also considered as very promising those for the manufacture of charged particle detectors [1–3]. Even high-energy charged particles in semiconductors have short mean free paths, not exceeding several tens of microns, therefore the preferential collection of single charge carrier, which is necessary to get high energy resolution [4, 5], can be easily achieved.

Semiconductor charged particle detectors are intended to operate under the influence of a harsh radiation environment, for example, under proton irradiation inducing radiation defects in the matrix, which affect the electrophysical properties and detector performance. Elucidation of the mechanisms of such influence is of decisive importance to apply the detectors in various fields, as well as to improve the methods of radiation monitoring. Earlier studies of radiation damage caused by charged particles such as protons in CdTe and CdZnTe detectors were mainly devoted not to studying the mechanisms of the influence of radiation defects energy levels on the detector properties, but to investigations of the effect of the radiation doses on the device characteristics [2, 6–14], the most important are the charge collection efficiency, resistivity, electron mobility, lifetimes of nonequilibrium electrons, and holes. For example, Yaxu Gu et al. [13] found that a high density of nonradiative recombination centers is generated in CdZnTe crystals after 2 MeV proton fluence of $6 \cdot 10^{13}$ p/cm², which causes a significant deterioration in the detector performance. Similar results were reported in paper

[14], where the charge collection of CdZnTe-based detector significantly degraded after irradiation of 2 MeV proton fluences of above $3.3 \cdot 10^{11}$ p/cm². Zanarini et al. [10] investigated damages caused by 2 MeV protons in CdZnTe and CdTe radiation detectors, and found that CdZnTe detectors have decreased performance after a fluence of $2.6 \cdot 10^{11}$ p/cm² which about an order of magnitude higher than limiting dose for CdTe-based detectors. The distribution of the charge collection efficiency η over the cathode surface of a CdZnTe detector bombarded with 2 MeV protons was estimated in paper [2], using ion beam induced charge microscopy. It was found that the bombardment of selected areas of the surface with a fluence of 10^5 ion/cm² causes structural damage in the irradiated areas and a noticeable deterioration of the charge collection efficiency.

The authors of paper [12] found that the fluence of 700 keV protons, which caused a complete degradation of the characteristics of CdTe and CdZnTe detectors, was identified to be the same for investigated detectors i.e. $2 \cdot 10^{12}$ p/cm², even if batches of detectors characterized by different initial transport properties had been chosen. The transport properties of the detectors were studied by measuring the product of the mobility by the lifetime of nonequilibrium charge carriers. The photoinduced current transient spectroscopy (PICTS) analysis was used to determine the activation energy of deep traps present in the materials. It was also suggested in [12] that some deep levels, which serve as traps for nonequilibrium electrons and holes, play an important role in changing the detector performance during irradiation. The CdTe and CdZnTe detector materials have a high resistivity of $10^{10} \dots 10^{11}$ Ω ·cm, which greatly complicates or makes it impossible, basing only on experimental techniques, to elucidate the mechanisms of change and degradation of the electrophysical and detector properties under the influence of radiation defects. In this regard, it is very important to use the computer modeling as an additional

research tool, based on known experimental results and proven physical models.

The aim of this work is to study by the simulation method the effect of radiation defects arising under the proton irradiation on the mechanisms of degradation of the electrophysical and detector properties of CdTe:Cl and CdZnTe. The calculations were made for the electron mobility μ_n , the resistivity ρ , the lifetime of nonequilibrium electrons τ_n , and holes τ_p , the charge collection efficiency η depending on the concentration of doping shallow donors N_d for different defect compositions of the samples before and after proton irradiation.

PHYSICAL MODELS AND MATERIALS COMPOSITION

The applied models as well as their testing were described in detail in [15], and the spectral characteristics of the initial and irradiated CdTe:Cl and Cd_{0.9}Zn_{0.1}Te were taken from [10, 12, 13]. To determine the Fermi level F and the concentrations of free electrons n and holes p in the parabolic band approximation, we used a multilevel compensation model for an arbitrary number of impurities and defects with construction and numerical solution of the corresponding electroneutrality equation taking into account all impurities and defects experimentally recorded in [12, 13]. The electron mobility μ_n was calculated using the approximation of the pulse relaxation time and taking into account the mechanisms of scattering at ionized and neutral centers, acoustic, piezoelectric, and optical phonons. The hole mobility μ_p was considered unchanged and equal to 70 cm²/(V·s). Specific conductivity was determined as a quantity

consisting of electron and hole components, and it was calculated by the formula: $e \cdot n \cdot \mu_n + e \cdot p \cdot \mu_p$, where e is the electron charge. Resistivity was calculated as the reciprocal of conductivity. The lifetimes of nonequilibrium electrons and holes were calculated using the Shockley-Reed-Hall recombination model, and the charge collection efficiency of the detector was determined according to the Hecht equation [16, p. 489]. The experimentally measured and published values of the i th energy levels of defects E_i , their concentration N_i , and the capture cross section σ_i of nonequilibrium charge carrier traps were used as input parameters. As in [12], the distance between the detector electrodes was assumed to be 1 mm, and the electric field strength for CdTe:Cl-based detector was 500 V/cm, and for CdZnTe-based that did 1000 V/cm.

The typical initial composition of the unirradiated CdTe:Cl and Cd_{0.9}Zn_{0.1}Te detector materials, registered in [12], is presented in the Table. Since the PICTS technique used in that paper to determine the compositions of the samples allows one to measure accurately only the energy of defects, the concentrations of impurities and defects were determined by us in such a way as to provide hole conductivity as well as the known maximum values of resistivity $\rho = 10^{10}$ and 10^{11} Ω·cm for CdTe:Cl and Cd_{0.9}Zn_{0.1}Te respectively. In addition to [12], a similar composition was described, for example, in [17–20], where, as a rule, A centers were present, for which a doping donor was introduced to compensate. Aluminum, indium, yttrium can be used as a shallow dopant donor in CdZnTe, while chlorine is often doped in CdTe.

Composition of defects in CdTe:Cl and Cd_{0.9}Zn_{0.1}Te before irradiation

Defect	Energy levels E_i , eV		Concentrations N_i , cm ⁻³		Capture cross-sections σ_i , cm ²	
	CdTe:Cl	Cd _{0.9} Zn _{0.1} Te	CdTe:Cl	Cd _{0.9} Zn _{0.1} Te	CdTe:Cl	CdZnTe
AX ⁻	0.11	–	$3.3 \cdot 10^{14}$	–	$1.7 \cdot 10^{-21}$	–
A0 ⁻	0.12	–	$2 \cdot 10^{15}$	–	$3.6 \cdot 10^{-21}$	–
A ⁻	0.14		$2 \cdot 10^{15}$	$2 \cdot 10^{15}$	$1.8 \cdot 10^{-19}$	
A1 ⁻	0.16		$2 \cdot 10^{15}$	$2 \cdot 10^{15}$	$4 \cdot 10^{-19}$	
K ⁻	–	0.23	–	$6 \cdot 10^{15}$	–	$6 \cdot 10^{-19}$
X ⁻	0.29		$2 \cdot 10^{15}$	$6 \cdot 10^{15}$	$3.6 \cdot 10^{-19}$	$4.0 \cdot 10^{-19}$
D ²⁻	0.41		$2 \cdot 10^{15}$	$2 \cdot 10^{15}$	$7.0 \cdot 10^{-17}$	
Z ⁺	1.05 (E _c -0.47)	1.05 (E _c -0.52)	$1 \cdot 10^{14}$	$1 \cdot 10^{15}$	$5.0 \cdot 10^{-17}$	
J ⁻	0.53		$2 \cdot 10^{15}$	$5.0 \cdot 10^{14}$	$5.0 \cdot 10^{-18}$	
Y ⁻	0.63		$1.8 \cdot 10^{15}$	$8.0 \cdot 10^{15}$	$3.2 \cdot 10^{-17}$	
W ⁻	0.70		$1.5 \cdot 10^{15}$	$9.0 \cdot 10^{15}$	$1.3 \cdot 10^{-17}$	$1.3 \cdot 10^{-18}$
H ⁻	0.75	–	$2.0 \cdot 10^{15}$	–	$1.0 \cdot 10^{-18}$	–
H1 ⁺	0.73 (E _c -0.79)	0.75 (E _c -0.82)	$4.0 \cdot 10^{15}$	$7.0 \cdot 10^{15}$	$2.0 \cdot 10^{-17}$	$4.0 \cdot 10^{-18}$
I ⁺	0.52 (E _c -1.0)	0.57 (E _c -1.0)	$5 \cdot 10^{15}$	$6.0 \cdot 10^{15}$	$1.0 \cdot 10^{-18}$	

There are also deep acceptors – cadmium vacancies V_{Cd} and deep donors in the form of tellurium at site of cadmium Te_{Cd} . The presence of deep donors in the matrix is necessary to achieve the high resistance state required for the operation of detectors.

The first column of the Table contains the symbols and charge states of defects: ‘plus’ for electron traps, and ‘minus’ for hole traps. The designations of defects in this column are the same as in [12]. Energy levels are measured from the valence band. Additionally, the

energy levels relative to the conduction band for donor centers are indicated in parentheses. The Table also lists the values of the level capture cross sections σ_i , the order of magnitude of which was determined in [21], where the contribution of all levels to the currents emitted into the corresponding bands was taken into account. The designations of defects and their probable nature are given in papers [10–12, 18–21]. Dopants of a shallow donor introduced into the matrix of the studied materials are intended to compensate shallow acceptors (AX, A0, A, A1) and to achieve a high-resistance state necessary for the operation of detectors. It is generally accepted today that the levels AX, A0, A, A1, K, X are the complexes of cadmium vacancies with dopants and background impurities. The cadmium vacancy apparently corresponds to defect D [22]. The donor defects Z ($E_c-0.47$ eV in CdTe, $E_c-0.52$ eV in CdZnTe) and H1 ($E_c-0.79$ eV in CdTe, $E_c-0.82$ eV in CdZnTe) can be Te(I) and Te_{Cd} respectively. Donor defect I corresponds to a tellurium vacancy. The nature of the remaining defect levels has not yet been fully elucidated.

RESULTS AND DISCUSSION

The authors of paper [12] irradiated CdTe:Cl and $Cd_{0.9}Zn_{0.1}Te$ detectors with 700 keV protons and fluences from $2.5 \cdot 10^{11}$ to $2 \cdot 10^{12}$ p/cm². After irradiation of cadmium telluride detector with a fluence of $2 \cdot 10^{12}$ p/cm², the charge collection and the value of the product of the electron mobility μ_n by the lifetime of nonequilibrium electrons τ_n decreased by 13.5 times. To investigate this behavior, an analysis was performed using PICTS technique. The PICTS spectra, given in [12] for the initial and irradiated materials, were used in present work for a model study of changes in the electrophysical and detector properties of CdTe:Cl and $Cd_{0.9}Zn_{0.1}Te$ during proton bombardment.

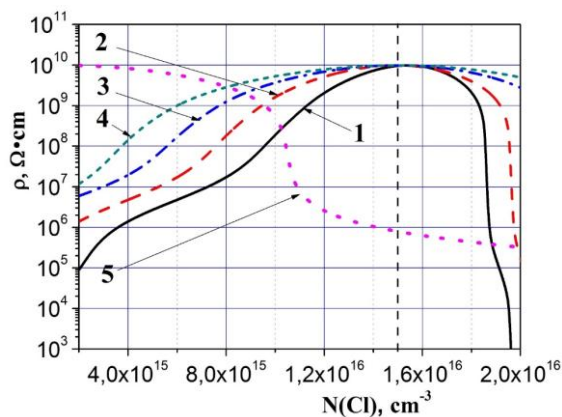


Fig. 1. Dependences of the specific resistance of CdTe:Cl on the chlorine concentration for the different defect concentrations N , cm⁻³, and after various fluences of protons J_p , p/cm²: 1 – the initial sample with the composition given in the table; 2 – $J_p=2.5 \cdot 10^{11}$, $N(Y)=1.8 \cdot 10^{15}$, $N(H)=3 \cdot 10^{15}$, $N(H1)=6 \cdot 10^{15}$; 3 – $J_p=6 \cdot 10^{11}$, $N(Y)=1.9 \cdot 10^{15}$, $N(H)=5 \cdot 10^{15}$, $N(H1)=8 \cdot 10^{15}$; 4 – $J_p=1.2 \cdot 10^{12}$, $N(Y)=2.0 \cdot 10^{15}$, $N(H)=7 \cdot 10^{15}$, $N(H1)=1 \cdot 10^{16}$; 5 – $J_p=2 \cdot 10^{12}$, $N(Y)=1.9 \cdot 10^{15}$, $N(H)=5 \cdot 10^{15}$, $N(H1)=8 \cdot 10^{15}$, $N(Z)=1 \cdot 10^{16}$. The vertical dashed line corresponds to the studied sample

Fig. 1 shows the dynamics of the change in the resistivity of CdTe:Cl depending on the concentration of the chlorine dopant after various proton irradiation fluxes J_p . The range of variation of the Cl dopant concentration was chosen so that for an unirradiated material the resistivity dependence passed through a maximum. The vertical dashed line intersects this ρ dependence near the maximum, marks the chlorine concentration on the horizontal axis, at which hole conductivity is observed in cadmium telluride, and therefore can denote the material under study. According to the spectra given in [12], during irradiation the concentrations of defects Z (0.47 eV), Y (0.63 eV), H (0.75 eV), H1 (0.79 eV) increase, and the authors of this paper supposed that defects Z and H1 are the only candidates which cause degradation of electron transport properties. Moreover, the appearance and increase in the content of the donor defect Z always causes a complete degradation of the detector properties of CdTe:Cl. All these circumstances were taken into account when simulating changes in ρ during irradiation, shown in Fig. 1. It can be seen from the figure that with an increase in the content of radiation defects, the resistivity ρ first stabilizes together with an expansion of the Cl concentration range where a high-resistance state is reached, and then there is a sharp drop in the resistivity by about four orders of magnitude, which leads to a complete degradation of the detector properties.

The fact of an abrupt deterioration of the detector properties was established experimentally in [12] (see Fig. 1 [12]). There it was shown that after the fluence of $2 \cdot 10^{12}$ p/cm² the gamma spectrum of CdTe:Cl-based detector from the ⁵⁷Co demonstrates drastically degradation of collecting properties. According to the simulation performed in present work, the reason for such a jump in properties is the appearance in large quantities of a radiation defect Z which pulls the Fermi level closer to the conduction band, thereby increasing the concentration of free electrons and decreasing ρ . The behavior of the Fermi level F is shown in Fig. 2.

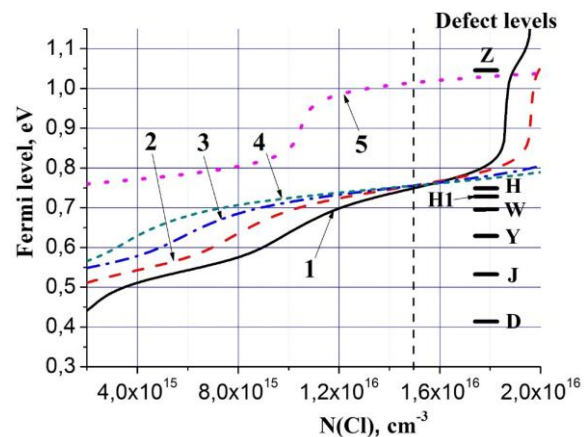


Fig. 2. Behavior of the Fermi level, measured from the valence band, as a function of the concentration of the doping chlorine after various proton fluences. The fluences and defect concentrations are the same as for the numbered curves in Fig. 1. The vertical dashed line corresponds to the studied sample

We see for curves 1–4 the pinning of Fermi level near the middle of the band gap and, hence, the stabilization of ρ , which occurs due to a large increase in the concentration of acceptor defects H, especially donor H1 (see Fig. 2 [12]), the energy levels of which attract F . A gradual increase in the concentrations of donor radiation defects Z and H1 increases the number of electrons n in the conduction band. Finally, at certain concentrations of Z and n , a sharp displacement of F to the vicinity of level Z occurs (see Fig. 2, curve 5).

After an abrupt displacement of the Fermi level to the vicinity of Z, the energy level of which becomes partially ionized, a recombination of nonequilibrium electrons at this level occurs, which, in turn, reduces the charge collection efficiency η and further deteriorates the detector properties.

Fig. 3 shows the dynamics of changes in charge collection efficiency during the proton bombardment. Here for dependencies 1–4, the concentration of Z (10^{13} cm^{-3}) is much lower than the content of any impurities and defects. For curves 5–7, the concentration of Z gradually increases, which causes a noticeable decrease in charge collection due to recombination at the Z level of nonequilibrium electrons, which, together with the influence of Z on a sharp decrease in ρ , clearly demonstrates the well-known experimental fact about the decisive role of this defect in degradation detector properties of CdTe:Cl. Based on the character of dependences 1–4 in Fig. 3, we see that the charge collection for the investigated detector material changes little. The reason of this effect is the small variability of the position of the Fermi level (see Fig. 2) before the appearance of a defect Z in an appreciable amount and, therefore, the little variability of the ionization degree of energy levels close to F , which capture nonequilibrium charge carriers.

Simulation also showed that the main contribution to the degradation of charge collection is made by a decrease in the lifetime of nonequilibrium electrons τ_n by more than an order of magnitude. At the same time, the lifetime of nonequilibrium holes τ_p is reduced by less than two times.

Thus, the reason for the experimentally observed in a number of cases a sharp degradation of the detector properties of CdTe:Cl after a certain dose of proton irradiation should be considered the appearance in a large amount of donor radiation defect, presumably tellurium interstitial Te (I), the energy level of which located in the upper half of the band gap. The emergence of Te (I) and a noticeable increase of its concentration is accompanied by a jump in the Fermi level F to the upper half of the band gap with a change in conductivity from hole to electronic and a sharp drop in the resistivity, which is the main reason of detector degradation. To increase the radiation resistance of CdTe:Cl, it is necessary to stabilize the position of the Fermi level during irradiation, which can be achieved by the formation of a sufficient amount of deep acceptor with energy level located below the middle of band gap. Such deep acceptor is cadmium vacancy V_{Cd} not bound in complexes with background and dopant impurities. The formation of such complexes is characterized by an increase in the charge number of cadmium vacancies

from $^{-2}V_{\text{Cd}}$ to $^{-1}V_{\text{Cd}}$ or $^0V_{\text{Cd}}$, and, as a consequence, a decrease in their effect on the Fermi level. To restrict the formation of these complexes, the purity of the detector material should be increased, and the concentration of doping shallow donors should be reduced to some value.

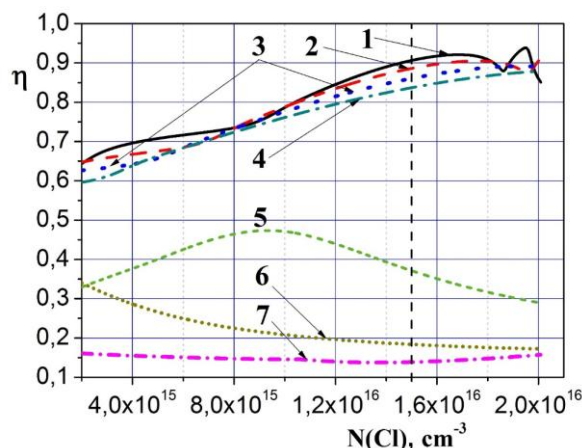


Fig. 3. Dependences of the charge collection efficiency on the concentration of doping chlorine after bombardment by protons with different fluences. The sample characteristics for 1–5 curves are the same that for numbered curves in Fig. 1; 6 – $N(Z)=3 \cdot 10^{16} \text{ cm}^{-3}$; 7 – $N(Z)=4 \cdot 10^{16} \text{ cm}^{-3}$. The vertical dashed line corresponds to the studied sample

Now we consider the results of modeling changes in the electrophysical and detector properties of CdZnTe under the influence of proton irradiation, experimentally investigated in [12], where two batches of samples no. 1 and 2 were irradiated with the same proton fluence as CdTe:Cl discussed above. Samples of batch no. 1 at first did not change their properties during irradiation, and then at a final stage suddenly deteriorated the detector properties after a fluence of $2 \cdot 10^{12} \text{ p/cm}^2$. We have already considered above a similar effect by the example of CdTe:Cl in which almost the same defects and energy levels were registered, as shown in the Table. On the contrary, detectors of batch no.2 gradually lost their registering properties during irradiation, so we will first consider this specific case.

It follows from the PICTS spectra, shown in Fig. 5 of paper [12], that in the process of proton bombardment, only the relative content of radiation defects K (0.23 eV) and X (0.29 eV) increases noticeably, and a some concentration of these defects already occurs in the initial $\text{Cd}_{0.9}\text{Zn}_{0.1}\text{Te}$ (Fig. 4 [12]). For this reason, the calculations were carried out taking into account the gradual increase in the content of K and X during irradiation. Fig. 4 shows the dependences of the change in the resistivity (a) and the charge collection efficiency (b) of CdZnTe-based detectors on the concentration of doping shallow donor for unirradiated sample and with different concentrations of radiation defects K, X. Curves 5 in both plots approximately correspond to the state of the detector after the proton fluence of $2 \cdot 10^{12} \text{ p/cm}^2$ with complete degradation of registering properties. It can be seen that, as in the experiment of paper [12], there is a gradual degradation of the detector properties of CdZnTe. We will try to

understand the reasons for this phenomenon. Let us consider the behavior of the Fermi level shown in Fig. 5, which also shows the defects levels with the exception of the donor defect I (tellurium vacancy), the level of which practically coincides with the level of the acceptor defect J.

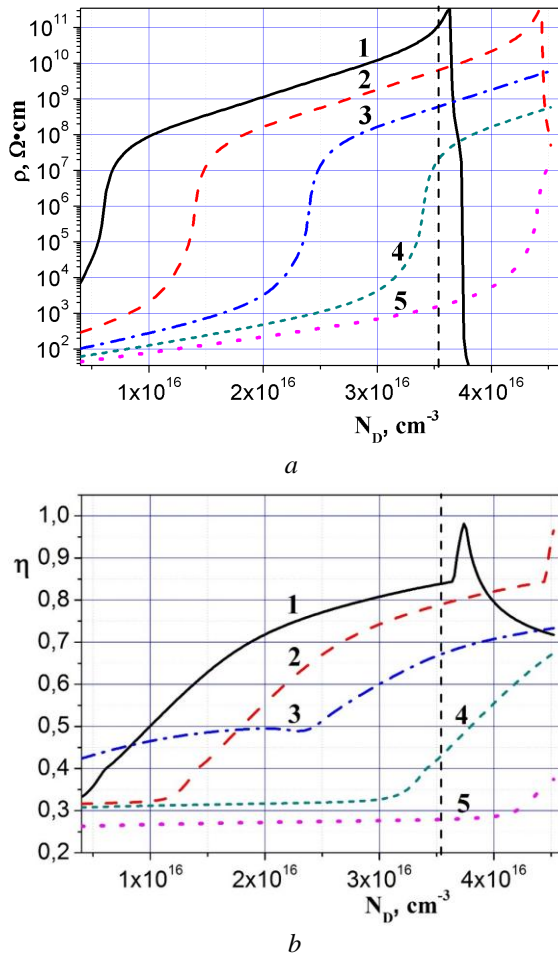


Fig. 4. Dependences of the resistivity (a) and the charge collection efficiency (b) in a $\text{Cd}_{0.9}\text{Zn}_{0.1}\text{Te}$ detector on the content of doping shallow donors for different concentrations of K and X defects $N(K, X), \text{cm}^{-3}$. 1 – initial, unirradiated sample, $6 \cdot 10^{15}$; 2 – $1 \cdot 10^{16}$; 3 – $1.5 \cdot 10^{16}$; 4 – $2 \cdot 10^{16}$; 5 – $2.5 \cdot 10^{13}$.

The vertical dashed lines denote the sample under study

For the material under study, the maximum values of the resistivity and charge collection, required for the operation of the detectors, are observed at the Fermi level position $F \sim 0.8 \text{ eV}$, i.e. in the vicinity of the deep donor level of H1, which, by the opinion of the most investigators, is the substitutional defect Te_{Cd} [20, 23, 24]. During the proton bombardment, carried out by the authors of paper [12], the interstitial tellurium $\text{Te}(\text{I})^{2+}$, Te_{Cd}^+ , and also cadmium vacancies V_{Cd}^{2-} are formed; the last form complexes K and X with various background impurities. Apparently, the latter mechanism turned out to be predominant, which led to the prevailing formation of radiation defects K and X of the acceptor type. As the concentrations of K and X increase, they begin to shift gradually the Fermi level F towards the valence band with a gradual decrease in the resistivity. In this case, a jump in the position of F , and hence ρ , as

it was for $\text{CdTe}:\text{Cl}$, does not occur, since the Fermi level in its drift gradually passes by the levels W, Y, J, D, I which prevent abrupt changes of F position. In the vicinity of each of these levels, within a certain range of concentrations of radiation defects, the Fermi level changes small, i.e. some pinning effect takes place, and it has a stabilizing effect.

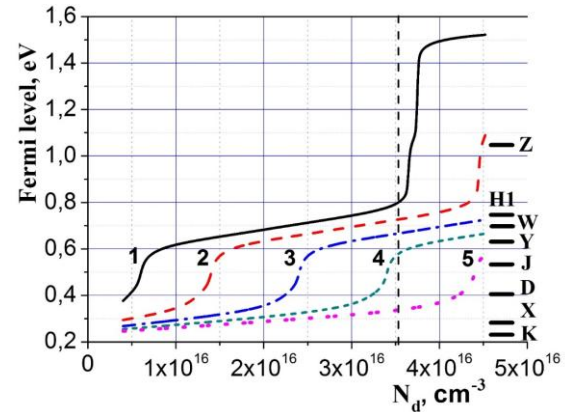


Fig. 5. Dependences of the Fermi level on the concentration of a doping shallow donor for different contents of radiation defects K and X, the concentrations of which are the same as for numbered curves 1–5 in Fig. 4. The vertical dashed line denotes the sample under study

In addition, partially ionized levels (W, Y, J, D, I) serve as centers of recombination of nonequilibrium charge carriers. As the Fermi level drifts during irradiation, the concentration of the radiation centers of recombination increases. Ultimately, the Fermi level is fixed near the K and X levels, which is accompanied by a decrease in the resistivity by several orders of magnitude, as well as a decrease in charge collection by a factor of two to three compared to the initial state due to the recombination at these levels. K and X levels are of the acceptor type; therefore, they are traps for holes, and it was just experimentally confirmed in [12]. The behavior of the Fermi level in $\text{CdTe}:\text{Cl}$ (see Fig. 2) is completely different from that just described. In $\text{CdTe}:\text{Cl}$, with an increase in the concentration of the radiation defect Z during irradiation, there are no other defect levels near which the level F could be pinned along the path of its displacement towards the level Z in the band gap, therefore, precisely a jump of the Fermi level took place, and a sharp degradation of detector properties was experimentally observed.

In [12], the batch of CdZnTe samples no.1 was also investigated, which after a proton flux $J_p = 2 \cdot 10^{12} \text{ p/cm}^2$ demonstrated an abrupt degradation of the detector properties. However, the PICTS spectra of these samples after irradiation are not presented in the article; therefore, it is difficult for them to indicate the degradation mechanism. Nevertheless, in paper [10] where the damage caused by 2 MeV protons in CdTe and CdZnTe radiation detectors was investigated, the gamma spectra of $\text{Cd}_{0.9}\text{Zn}_{0.1}\text{Te}$ detector from the ^{57}Co source (see Fig. 4 [10]) are presented, and one can see sharp deterioration of the detector properties after the fluence $(2.1+2.6) \cdot 10^{11} \text{ p/cm}^2$. In this case, the PICTS spectra for $\text{Cd}_{0.9}\text{Zn}_{0.1}\text{Te}$ are presented (see Fig. 5 in

[10]), in which it can be seen that a defect Z appears in a completely degraded material, at that in a large amount. Thus, it can be assumed that in the batch samples of $Cd_{0.9}Zn_{0.1}Te$ no.1 of paper [12], a sharp degradation of the detector properties occurred due to a jump-like transition of the Fermi level to the vicinity of the donor level Z located in the upper half of the band gap.

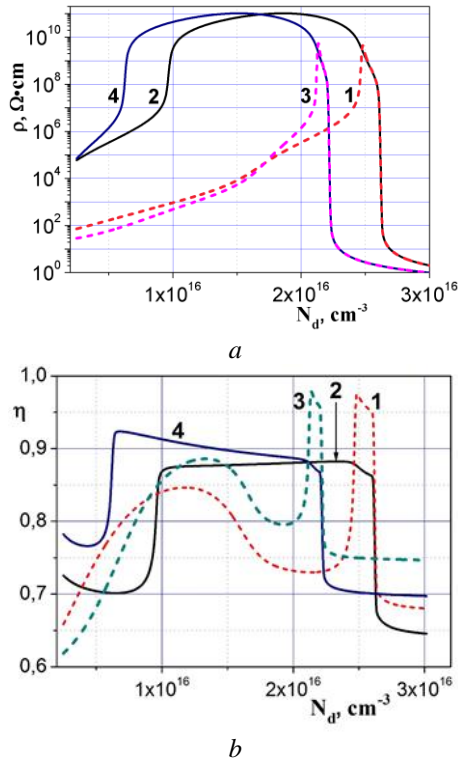


Fig. 6. Dependences of resistivity (a) and charge collection efficiency (b) on the concentration of the dopant donor impurity in $Cd_{0.9}Zn_{0.1}Te$ -based detector.

Dashed dependences are for the composition experimentally determined in [13]; solid curves are for the same composition, but with introducing of a deep donor with an energy $E_C-0.79$ eV and concentration of $1.5 \cdot 10^{16} \text{ cm}^{-3}$; 1, 2 – initial samples; 3, 4 – after irradiation by 2 MeV proton fluence of $6 \cdot 10^{13} \text{ p/cm}^2$

In conclusion, we consider one of the results obtained in [13], the authors of which investigated radiation damage by the photoluminescence method, and the concentration of defects, their level energies and capture cross sections were determined using the thermally stimulated current (TSC) technique. The values of resistivity and gamma spectra of detectors based on the material under study was not presented in that paper. Modeling the electrophysical properties, with taking into account the parameters experimentally determined in [13] and presented there in Table, showed that it is problematically to create an acceptable gamma radiation detector based on such a material, since the maximum possible resistivity at room temperature does not exceed $10^9 \text{ } \Omega \cdot \text{cm}$, which is clearly insufficient for $Cd_{0.9}Zn_{0.1}Te$ -based detector. In addition, this value of ρ can be achieved only in a very narrow range of concentrations of the doping shallow donor from $2.47 \cdot 10^{16}$ to $2.51 \cdot 10^{16} \text{ cm}^{-3}$, which requires samples with very uniform properties and the need for very precise control of dopant and defect composition. Finally, in the

indicated narrow high-resistance region, the conductivity of the detector material turns out to be electronic with $n/p \sim 10^3$ at room temperature, although the materials for gamma and X-rays detectors have to be of hole conductivity. Modeling has shown that to eliminate the indicated drawbacks it is sufficient to introduce into the matrix a structural defect or impurity, with a deep donor level located in the band gap within the range from 0.75 to 0.80 eV. This is demonstrated in Fig. 6 which shows the dependences of the resistivity ρ and the charge collection efficiency η on the concentration of the shallow donor for the initial and irradiated samples from paper [13] (curves 1, 3), as well as with introduced deep donor having energy level at 0.79 eV measured from the bottom of the conduction band and a concentration of $1.5 \cdot 10^{16} \text{ cm}^{-3}$ (curves 2, 4). The antisite defect Te_{Cd} can act as such a donor.

In Fig. 6 there is a noticeable shift in the dependences of $\rho(N_d)$ and $\eta(N_d)$ after irradiation towards lower concentrations of a doping shallow donor, i.e. in the course of proton bombardment, the relative content of donor-type radiation defects increases. In other words, during proton irradiation the total content of ionized radiation defects of the donor type increases relative to overall concentration of electrically active acceptors. Such phenomenon can occur in materials with an increased content of background impurities, which interact with radiation defects – cadmium vacancies, and form complexes, neutral or with an increase in the charge number by +1 relative to the initial state of V_{Cd}^{2-} . If this assumption is correct, then the purity of the initial $Cd_{0.9}Zn_{0.1}Te$ should be improved.

Fig. 6 also shows a clear correlation between the region of high resistance state and high values of the charge collection efficiency η , which makes it possible to obtain a high-quality detector material, provided that a deep donor is introduced into its matrix.

CONCLUSION

The simulation results are in qualitative agreement with the experimental data of [10, 12].

It was found that the abrupt degradation of the detector properties after proton irradiation of the $CdTe:Cl$ and $CdZnTe$ materials is caused by the formation of such radiation defect as interstitial tellurium $Te(I)$, accompanied by a jump of the Fermi level F to the vicinity of the level of this defect and with a decrease in the resistivity by several orders of magnitude, and change in the conductivity from hole to n-type with noticeable reduced charge collection efficiency.

It was found for the first time that the condition for the jump in properties at the final stage of irradiation is the absence of energy levels of defects in the band gap within the range between the Fermi level positions in the initial and degraded states of detector material. It was determined for the first time that in the case of the presence of the mentioned levels, the degradation of the detector properties occurs gradually as F pass by these levels with some pinning near each of them in a certain range of radiation defect concentrations. Understanding the established mechanisms of degradation of $CdTe$ and

CdZnTe under the influence of proton irradiation will contribute to the development of the technology for producing radiation-resistant detectors based on these materials.

The high purity of CdTe and Cd_{0.9}Zn_{0.1}Te materials and the presence of a deep donor in their matrix, which can be an antisite defect Te_{Cd} or a donor impurity, with an energy level close to the middle of the band gap, are of particular importance for producing high-resistance, radiation-resistant detectors.

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Статья поступила в редакцию 25.01.2021 г.

ВЛИЯНИЕ ДЕФЕКТОВ, ВОЗНИКАЮЩИХ ПРИ ОБЛУЧЕНИИ ПРОТОНАМИ, НА ЭЛЕКТРОФИЗИЧЕСКИЕ И ДЕТЕКТОРНЫЕ СВОЙСТВА CdTe:Cl И CdZnTe

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Методом компьютерного моделирования изучены механизмы влияния радиационных дефектов, возникающих при облучении протонами высоких энергий, на удельное сопротивление ρ , время жизни неравновесных электронов τ_n и дырок τ_p в CdTe:Cl и Cd_{0,9}Zn_{0,1}Te и на эффективность сбора заряда η работающих при комнатной температуре детекторов ионизирующего излучения на основе этих материалов. Исследовано влияние рекомбинации на глубоких уровнях радиационных дефектов на деградацию τ_n , τ_p и η детекторов на основе CdTe:Cl и Cd_{0,9}Zn_{0,1}Te. Установлены основные факторы, влияющие на резкое или постепенное снижение удельного сопротивления и эффективности сбора заряда этих детекторов при их бомбардировке протонами высоких энергий, приводящих к полной деградации их регистрирующей способности. Отмечена важная роль чистоты и концентрации глубоких доноров в исходном состоянии необлученного материала детектора.

ВПЛИВ ДЕФЕКТІВ, ЩО ВИНІКАЮТЬ ПРИ ОПРОМІНЕННІ ПРОТОНАМИ, НА ЕЛЕКТРОФІЗИЧНІ ТА ДЕТЕКТОРНІ ВЛАСТИВОСТІ CdTe:Cl І CdZnTe

О.І. Кондрик

Методом комп'ютерного моделювання вивчено механізми впливу радіаційних дефектів, що виникають при опроміненні протонами високих енергій, на питомий опір ρ , час життя нерівноважних електронів τ_n і дірок τ_p у CdTe:Cl і Cd_{0,9}Zn_{0,1}Te та ефективність збору заряду η детекторів іонізуючого випромінювання на основі цих матеріалів, що працюють при кімнатній температурі. Досліджено вплив рекомбінації на глибоких рівнях радіаційних дефектів на деградацію τ_n , τ_p і η детекторів на основі CdTe:Cl і Cd_{0,9}Zn_{0,1}Te. Встановлені основні фактори, які впливають на різке або поступове зниження питомого опору та ефективності збору заряду цих детекторів при їх бомбардуванні протонами високих енергій, що призводить до повної деградації їх реєструючої здатності. Відзначено важливу роль чистоти і концентрації глибоких донорів у початковому стані неопроміненого детекторного матеріалу.