# **PHOTOELECTRIC CHARACTERISTICS OF THE HETEROJUNCTION**   $n$ **-GaAs-** $p$ **-(GaAs)<sub>1-x-y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub>**

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The photoelectric properties of *n*-GaAs –  $p$ -(GaAs)<sub>1–x–y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> heterostructures have been investigated both in photodiode and photovoltaic modes. It has been revealed that the spectral dependence of the photocurrent covers a wide range of energy intervals, ranging from 1.07 eV to 3 eV. It has been demonstrated that as the temperature of the crystallization onset  $(T<sub>oc</sub>)$  increases, the peaks of the spectral dependencies of the photoelectromotive force (photo-EMF) shift towards shorter wavelengths. It has been observed that as the crystallization onset temperature (T<sub>oc</sub>) of the solid solution layer (GaAs)<sub>1–x–y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> increases, the lifetime of photo carriers increases from  $10^{-7}$  s at T<sub>oc</sub>=650°C to 5 $\cdot 10^{-5}$  s at T<sub>oc</sub>=730°C. It is demonstrated that the peaks of the intrinsic photoluminescence band shift towards shorter wavelengths with an increase in the temperature of the crystallization onset. Additionally, the study of the intrinsic spectral region of photoluminescence in samples across the thickness of the epitaxial layer confirms the variability of the obtained structures.

**Keywords***: Solid solution; Heterostructure; Temperature; Photoelectric property; Photo-EM; Photoluminescence; Photo carriers; Varizon structure* 

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#### **INTRODUCTION**

Semiconductors  $A<sup>3</sup>B<sup>5</sup>$  and  $A<sup>2</sup>B<sup>6</sup>$  are well known as promising materials for creating optoelectronic devices in the infrared and visible spectrums. Intensive research is being carried out to create various electronic devices based on GaP, GaAs, CdTe, ZnSe and their solid solutions [1-4]. Among these materials, GaAs-based compounds are of particular interest, as their electron and hole mobilities are much higher [5-9], thus enabling them to be used in high-speed optoelectronic devices. In addition, solid substitution solutions  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  make it possible to expand the spectral range of functioning of structures based on them to 400 nm [10, 11]. However, the mass use of device structures made on the basis of the A<sup>3</sup>B<sup>5</sup> compound is limited, on the one hand, by the economic inexpediency of large-scale use of such device structures, and on the other hand, by insufficient knowledge of the photoelectric properties of such materials and heterojunctions based on them.

In this regard, this work reports the results of studies of some photoelectric properties of the  $n$ -GaAs - *p*-(GaAs)<sub>1-x-v</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> heterojunction, which was synthesized by growing epitaxial layers from solid solutions  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  on an n-GaAs substrate oriented in the (100) direction.

### **MATERIALS AND METHODS**

The growth processes of epitaxial layers were carried out from a limited volume of tin solution-melt on a GaAs substrate using forced cooling method. The thicknesses of the resulting films varied within 10-15 μm depending on the size of the gap between the substrates and the growth mode. X-ray diffraction patterns of layers  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$ , taken on a thirdgeneration diffractometer of the Empyrean Malvern PANalytical L.T.D type, confirmed that the grown layers have a sphalerite crystal lattice and are characterized by the lowest mechanical stresses, both in volume and near the interface of the layers.

The lux-volt characteristics were measured on a setup that can simulate sunlight. A 150 W tungsten incandescent lamp was used as the radiation source. The illuminance measurements were conducted using a digital illuminance meter, MASTKCH LUXMETER MS6610.

An investigation was conducted to analyze the spectral sensitivity of the samples by employing an optical spectrometer that featured a CARLZEISJENA mirror monochromator with quartz optics. This sophisticated setup allowed for the examination of the samples within the photon energy range of 1 to 3 eV.

The photoluminescence spectrum of the samples was studied at the KSVU-23 installation. Optical pumping of the surface of the epitaxial layer at a temperature of 300 K was carried out by a mercury lamp, the signal was recorded on the SDL-2 installation.

#### **RESULTS AND DISCUSSION**

The surfaces of the epitaxial layers of  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  solid solutions had p- type conductivity with Hall carrier concentrations  $p=5.10^{17}$  sm<sup>-3</sup>. The literature highlights that the ZnSe epitaxial layer typically exhibits n-type

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conductivity as a result of the self-compensation effect under standard conditions [12]. We successfully produced epitaxial films containing the chemical composition  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  that exhibited p-type conductivity. Based on quantum chemical concepts, the possibility of increasing the binding energy of selenium in the composition of the Se-Ge dumbbell structure located at the nodes of the crystal lattice of the  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  solid solution compared with the nodal position in the crystalline ZnSe lattice. This can lead to a significant reduction in the number of  $V_{S_6}$  vacancies, which usually lead to the appearance of electronic conductivity in ZnSe. In addition, part of the vacancies in the selenium  $V_{\text{Se}}$  sublattice in solid solutions can be occupied by Ge atoms, which also leads to a decrease in the number of  $V_{S_e}$  vacancies. It is assumed that the appearance of epitaxy layers of a solid solution of the  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  type during the formation of the *n*-GaAs –  $p$ -(GaAs)<sub>1–x–y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> heterostructure is associated with partial dissolution GaAs substrates.

The photovoltaic properties of the obtained structures were studied by both photovoltaic and photodiode methods at a temperature of 300K. The surface was illuminated from the side of the solid solution. The lux-ampere characteristics of the structures, obtained under integral light illumination, have shown that the dependence of the short-circuit current  $(I_{sc})$ on the illuminance exhibits a nonlinear relationship in the initial region. In other words, an increase in the incident light intensity leads to a wave-like increase in Isc. This phenomenon may be attributed to changes in the parameters of the contacting semiconductors under the influence of the light flux, such as the lifetime of non-majority charge carriers and the diffusion length (Fig. 1). With increasing intensity of the incident light, the open-circuit EMF tended to saturate, reaching a value of 0.6 V, which may be due to the impossibility of reducing the thickness of the transition layer between the epitaxial film and the substrate under the influence of incident light.



**Figure 1.** The dependence of the short-circuit current (1-curve) and the no-load EMF (2-curve) on the illumination by integral light

The lifetime of photocarriers in structures grown under various conditions is determined by the relaxation decay of the photo-EMF. The investigation revealed that as the temperature of the onset of crystallization  $(T<sub>oc</sub>)$  of the  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  solid solution layer rose, the lifetime of photo carriers extended from  $10^{-7}$  seconds at  $T_{\text{oc}} = 650^{\circ}\text{C}$ to  $5.10^{-5}$  seconds at T<sub>oc</sub>=730 °C. The spectral characteristics of I<sub>s.c</sub>.(short-circuit current) and U<sub>o.c</sub>. (open-circuit voltage), as well as the photocurrent were measured in the photodiode mode. Monochromatic radiation was carried out from the solid solution  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$ . Measurement of the spectral dependence of photocurrent in photodiode mode (Fig. 2, curves *a* and *b*) showed that it covered a wide range of incident photon energies from 1.07 to 3.0 V.

From Fig. 2, it can be seen that the spectrum of photodetectors in the energy range of 1-3 eV is characterized by six peaks with maximums at photon energies (eV): 1-at 1.37, 2-1.47, 3-1.65, 4-1.88, 5-2.3, 6-2.62 (Figure 2a).



**Figure 2.** Spectral dependence of the photocurrent of the *n*-GaAs –  $p$ -(GaAs)<sub>1-x-y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> structure obtained in photodiode mode at various crystallization onset temperatures:  $a - T_{oc} = 730^{\circ}C$ ,  $b - T_{oc} = 650^{\circ}C$ .

In curve b, Figure 2. the spectrum begins at a photon energy of 1.07 eV, which may be due to the bonding energy of paired Ge atoms, which partially substitute some of the gallium arsenide molecules and create the corresponding energy

level [13]. Additionally, impurity germanium atoms self-organize into nanocrystals in defectable regions of the gallium arsenide lattice [10], forming their own energy level with an acceptor-like characteristics [10]. At a photon energy of 1.48 eV, a photopic corresponding to recombination from the conduction band to acceptor states in p-GaAs is observed [11]. The maximum of the spectral sensitivity dependence is observed at 1.64 eV, which may be due to the structural bandgap of the valence zone and isovalent impurities of Ge-Se compounds in the GaAs layer [14]. The following photopic peaks, curve *b*, Fig. 2, corresponding to photon energies of 1.89, 2.18, and 2.67 eV, may be attributed to compounds GaSe (h $v_{\text{max}} = 1.88 \text{ eV}$ ), ZnAs (hv<sub>max</sub> = 2.15 eV), and ZnSe (hv<sub>max</sub> = 2.69 eV) with deep levels in the valence zone of gallium arsenide [12].

The spectral dependencies of the photo-EMF of heterostructures *n*-GaAs –  $p$ -(GaAs)<sub>1–x–v</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> are shown in Figure 3. It can be seen that the peaks of the spectral dependencies, characterized by the crystallization onset temperatures, shift towards the short-wavelength region with increasing displacement. Since carrier separation occurs in the solid solution  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  at the *p-n* junction, such mixing is likely associated with a decrease in the germanium content relative to ZnSe at the heterojunction boundary in the solid solution and an increase in the epitaxial onset temperature. Although the radiation falls on the side of the solid solution, some carriers generated from the surface of the solid solution at high energies of incident photons likely recombine without reaching the p-n junction boundary, where their separation occurs. Therefore, in almost all investigated cases of photo-EMF, a decrease in photosensitivity is observed in the shortwavelength region of the spectral dependence of photo-EMF. Some difference in the values of photosensitivity observed in the short-wavelength region for different samples can be explained by differences in film thickness.



**Figure 3.** Spectral dependence of the open-circuit photo-EMF of the structures *n*-GaAs – *p*-(GaAs)<sub>1–x–y</sub>(Ge2)<sub>x</sub>(ZnSe)<sub>y</sub> obtained at different crystallization onset temperatures:  $a - T_{oc} = 730^{\circ}C$ ,  $b - T_{oc} = 650^{\circ}C$ .

Also, investigations of photoluminescence were conducted on the solid solutions  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  obtained by us in their intrinsic spectral range. Samples obtained at different crystallization onset temperatures were subjected to measurement. Scanning of the samples was performed on the surface of epitaxial layers. Optical pumping of the surface layers at a temperature of 300 K was performed using a mercury lamp.

Meanwhile, upon visual observation, luminescent glow ranging from green to blue was detected on the surface of epitaxial layers, depending on the sample conditions. By scanning the surface of the films in various directions, it was also possible to approximately visually assess the condition of the sample surfaces based on color and the uniformity of luminescence. In the photoluminescence spectra obtained by continuous recording (Figure 4 - *a* and *b*), it was found that the peaks of the intrinsic photoluminescence bands shift towards the short-wavelength region with increasing crystallization onset temperature.



**Figure 4.** Intrinsic spectral range of photoluminescence spectra taken on the surface and upon sequential removal of the surface of epitaxial layers  $(a', a'', b')$ : a - T<sub>oc</sub> = 730°C, b - T<sub>oc</sub> = 650°C.

Investigating the photoluminescence spectrum of the solid solution  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$  it was found that with the removal of surface layers, the peaks of the intrinsic band shift towards the long-wavelength region (Figure 4 - *a*', *a*'', and *b*'). The research [15] demonstrated that in the solid solution, the zinc selenide content increases along the growth direction, consequently leading to an increase in the width of the bandgap.

Thus, the dependence of the bandgap width on the composition of germanium and zinc selenide in the volume of solid solution has a non-monotonic character, and the obtained solid solutions exhibit varizon structures. The bandgap widths, estimated from the photoluminescence spectra for the two specified samples, were 2.51 and 2.21 eV.

### **CONCLUSIONS**

Thus, the photoelectric properties of *n*-GaAs –  $p$ -(GaAs)<sub>1–x–y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub> heterostructures have been investigated both in photodiode and photovoltaic modes. It has been found that the spectral dependence of the photo-current covers a wide range of energy intervals from 1.07 eV to 3 eV. It has been demonstrated that with an increase in the temperature of the crystallization onset (Toc), the peaks of the spectral dependencies of the photo-electromotive force (photo-EMF) shift towards shorter wavelengths. It has been discovered that with an increase in the temperature of the crystallization onset (Tc) of the solid solution layer  $(GaAs)_{1-x-y}(Ge_2)_x(ZnSe)_y$ , the lifetime of photo-carriers increases from  $10^{-7}$  s at  $T_{\text{oc}} = 650^{\circ}\text{C}$  to  $5.10^{-5}$  s at T<sub>oc</sub> = 730°C. Investigation of the photoluminescence on the surface of epitaxial layers (GaAs)<sub>1-x-y</sub>(Ge<sub>2</sub>)<sub>x</sub>(ZnSe)<sub>y</sub>, it has been shown that the peaks of the intrinsic photoluminescence band shift towards shorter wavelengths with increasing crystallization onset temperature. Additionally, investigation of the intrinsic spectral range of photoluminescence in samples through the thickness of the epitaxial layer confirms the varizonality of the obtained structures.

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### **ФОТОЕЛЕКТРИЧНІ ХАРАКТЕРИСТИКИ ГЕТЕРОПЕРЕХОДУ n-GaAs-p-(GaAs)1-x-y(Ge2)x(ZnSe)y Акрамжон Ю. Бобоєвa,b**

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Досліджено фотоелектричні властивості гетероструктур *n*-GaAs – *p*-(GaAs)1–x–y(Ge2)x(ZnSe)y як у фотодіодному, так і у фотоелектричному режимах. Виявлено, що спектральна залежність фотоструму охоплює широкий діапазон енергетичних інтервалів від 1,07 еВ до 3 еВ. Показано, що з підвищенням температури початку кристалізації (Toc) піки спектральних залежностей фотоелектрорушійної сили (фото-ЕРС) зміщуються в бік коротших довжин хвиль. Було помічено, що зі збільшенням температури початку кристалізації (Toc) шару твердого розчину (GaAs)1–x–y(Ge2)x(ZnSe)y час життя фотоносіїв збільшується з  $10^{-7}$  с при  $T_{\text{oe}} = 650^{\circ}$ С до  $5 \cdot 10^{-5}$  с при  $T_{\text{oe}} = 730^{\circ}$ С. Показано, що піки власної смуги фотолюмінесценції зміщуються в бік коротших довжин хвиль зі збільшенням температури початку кристалізації. Крім того, дослідження власної спектральної області фотолюмінесценції зразків по товщині епітаксійного шару підтверджує варіабельність отриманих структур.

**Ключові слова:** *твердий розчин; гетероструктура; температура; фотоелектричні властивості; фото-ЕМ; фотолюмінесценція; фотоносії; варізонна структура*