Abstract
This work reports on an experimental investigation of the influence of vertical stacking of quantum dots, the thickness of GaAs potential barriers, and their isovalent doping with bismuth on the photoluminescence properties of InAs/GaAs heterostructures. The experimental samples were grown by ion-beam deposition. We showed that using three vertically stacked layers of InAs quantum dots separated by thin GaAs barrier layers was accompanied by a red-shift of the photoluminescence peak of InAs/GaAs heterostructures. An increase in the thickness of the GaAs barrier layers was accompanied by a blue shift of the photoluminescence peak. The effect of isovalent Bi doping of the GaAs barrier layers on the structural and optical properties of the InAs/GaAs heterostructures was investigated. It was found that the Bi content up to 4.96 atom % in GaAs decreases the density of InAs quantum dots from $1.53 \times 10^{10}$ to $0.93 \times 10^{10}$ cm$^{-2}$. In addition, the average lateral size of the InAs quantum dots increased from 14 to 20 nm, due to an increase in the surface diffusion of In. It is shown that isovalent doping of GaAs potential barriers by bismuth was accompanied by a red-shift of the photoluminescence peak of InAs quantum dots of 121 meV.

Introduction
The formation of III–V nanoheterostructures with quantum dots (QDs) raises the possibility of developing a new generation of photodetectors in the infrared range [1-3]. The significant problems of existing HgCdTe detectors are low yield and high cost in comparison with quantum-well infrared photodetectors (QWIPs) [4]. The QWIPs, in turn, have a simpler technology but a low quantum efficiency and require cooling. One way to solve these problems is to grow semiconductor heterostructures in which QDs are embedded. The localization of photogenerated charge carriers in a quantum dot along three directions
leads to a low thermionic emission and decreases the dark current [5,6]. InAs/GaAs, Ge/Si, and others are examples of such heterostructures [7-14]. The structural and optical properties of InAs QDs rely heavily on their surrounding matrix, which is separated into two parts: the top with a GaAs strain-reducing layer; the bottom with a strained buffer layer (wetting layer) [15-19]. Thus, the simplified energy band diagram of the active region of an InAs/GaAs heterostructure is an InAs quantum dot built into a GaAs matrix in the form of a quantum well. It follows from the above that it is possible to manipulate the photosensitivity spectrum of heterostructures through three methods: 1) the size, shape and density of the quantum dot; 2) the vertical stacking of quantum dot arrays; 3) the material of the barrier layers surrounding the quantum dots. We and other researchers described the first method in [20-23]. Mechanical strains are a technological problem in the development of effective IR devices based on InAs/GaAs nanoheterostructures with vertically stacked QD layers [24,25]. The surface density and geometric sizes of InAs quantum dots varies depending on the mechanism and the sign of the stresses, and consequently, the photoluminescence properties of InAs/GaAs heterostructures will vary too. The distinctive feature of the first and second methods is that the optical transitions are performed inside the potential well formed by the quantum dot. In the case of using the third method, other mechanisms of optical transitions are possible.

The growth of QDs occurs in the Stranski–Krastanov mode, when the substrate wetting layer is elastically stressed. An InAs/GaAs heteropair is ideal for creating conditions for a transition from 2D to 3D growth but has some limitations for a number of functional applications such as manipulating the photosensitivity spectrum in the infrared region. The solution to the problem can be found by the selection of a potential-barrier material that can be found by the selection of a potential-barrier material for QDs based on III–V multicomponent solid solutions. GaAsSb [26], InGaAs [27], InGaAsN [28,29] and GaAsNSb [30,31] have been used for these purposes. In this work, it is proposed to use GaAsBi. Bismuth is an isovalent doping and forms ternary, quaternary and quinary solid solutions with III–V compounds [18,32-38]. Bi incorporation into GaAs makes it possible to change the lattice mismatches. The GaAs energy band diagram varies greatly due to the large size of Bi atoms. Emerging strains in the GaAsBi layer should have an influence on the surface density and sizes of InAs QDs and also on the photoluminescence properties of InAs/GaAs heterostructures [17,18,20,32,33]. In this regard, a study of the influence of Bi on the optical properties of InAs/GaAs heterostructures is an actual requirement.

In this work, ion-beam deposition (IBD) is used to grow the samples. IBD has significant advantages over many methods of growth from the gas phase, since it allows for a control of the energy of sputtered atomic fluxes and their interaction with the growth surface [39]. IBD plays an important role in the semiconductor materials technology [40-48].

In this work, we study the influence of QD vertical stacking, the thickness of GaAs potential barriers and their doping by bismuth on the structural and photoluminescence properties of InAs/GaAs heterostructures.

**Experimental**

**Heterostructures with vertically stacked InAs QDs**

Several samples were grown to investigate the influence of QD vertical stacking on the photoluminescence spectrum of the heterostructures. The heterostructures were synthesized by using an IBD system [21]. The deposition experiments were performed on GaAs(100) substrates. The deposition of the material was carried out by sputtering the target with an Ar⁺ ion beam. Calibration dependencies of InAs and GaAs sputtering yields on the beam energy, the slope angle and the flux density were obtained earlier [21,49,50]. Two types of samples were grown for the research of vertically stacked QDs. The first type of samples (ST#1) contained one to three QD layers separated by 15 nm GaAs layers. The second type of samples (ST#2) contained three QD layers that were separated by GaAs barriers of 15, 20 and 30 nm.

During the growth process, pressure in the vacuum chamber was $3.7 \times 10^{-7}$ Pa. The 500 nm thick n⁺-GaAs buffer layer at $T = 610$ °C was grown first. The accelerating voltage of the ion beam was 450 V, and the current density was $3.2 \times 10^{-4}$ A/cm². Furthermore, the i-GaAs barrier layer was deposited under similar conditions, then a 15 second pause was made, and the temperature was reduced to 535 °C. InAs QDs were grown at an accelerating ion voltage of 250 V and an ion current density of $4.5 \times 10^{-6}$ A/cm². The covering of QDs by the i-GaAs layer was accompanied by an increase of the temperature to 550 °C.

**InAs/GaAs heterostructures with potential barriers of GaAsBi**

Growth of GaAs$_{1-x}$Bi$_x$ layers was carried out using polycrystalline targets, which were made as follows. The batch composition was calculated in accordance with the solid phase composition of the finished GaAs$_{1-x}$Bi$_x$ target. The total mass of the melt was calculated taking into account the ingot volume of the polycrystalline target. After chemical treatment, the components of the batch were placed in a graphite container. Heating of the container with the batch was carried out by a resistive heater to a temperature above the melting point of GaAs (1250 °C), which is the highest-melting compound. The melt...
was kept for 10 min under such conditions, then rapid shut-
down of the heating elements was followed by crystallization of
the initial melt. The resulting polycrystalline ingot was cut to
2 mm thick plates.

For investigation of the influence of the potential barrier materi-
al on the photoluminescence properties of InAs/GaAs hetero-
structures, experimental samples with GaAs\(_{1-x}\)Bi\(_x\) potential
barriers with bismuth fractions of 1, 3 and 5 atom % were
grown. One experimental sample contained GaAs barriers for
comparison. The Bi-containing layers were grown at 360 °C.
The samples contained one layer of InAs QDs.

Characterization methods
Photoluminescence (PL) measurements of InAs/GaAs hetero-
structures were carried out at temperatures of 90 and 300 K in
the spectral range from 0.9 to 1.4 eV. An injection laser with
402 nm wavelength and power of 12.5 mW was used as a
source of excitation optical radiation. The PL signal was re-
corded by an MDR-23 monochromator with a cooled germa-
nium photodiode PDG-3600. Protection of the monochromator
entrance slit from the reflected harmonics of the exciting laser
radiation was carried out by means a Y-1.4 optical filter (light
yellow color).

A study of the surface morphology of the grown heterostruc-
tures was carried out using a Solver HV atomic force micro-
scope. The structural properties of the GaAsBi/GaAs, InAs/
GaAs, and InAs/GaAsBi heterointerfaces were investigated by
methods of Raman spectroscopy and X-ray diffraction (XRD).
X-ray diffraction reflection curves were investigated on a high-
resolution TRS-1 X-ray diffractometer with a third-crystal ge-
ometry using the Cu K\(_\alpha\) emission line (\(\lambda = 0.154 \text{ nm}\)). A

Results and Discussion
Photoluminescence properties of
InAs/GaAs heterointerfaces

The photoluminescence properties of vertically stacked QD
arrays grown by using molecular beam epitaxy (MBE) are well
studied in [24,25]. But the IBD method differs from MBE.
Therefore, it remains relevant to research the vertical stacking
of QDs grown by using IBD. In the beginning, ST#1 samples
were studied. The statistical analysis by the threshold tool in the
Image Analysis 2.1.2 program showed that the density of QDs
in the bottom layer reached 1.42 × 10\(^{10}\) cm\(^{-2}\), the average
lateral size of the quantum dots was 17 nm, and their height was
6.3 nm.

During PL excitation, certain steady-state conditions are estab-
lished in the sample, in which the radiation rate depends on the
lifetime of the photogenerated charge carriers in the QD or in
the wetting layer. Therefore, the photoluminescence in the
grown samples can go through the ground (PL\(_{\text{GS}}\)) and excited
(PL\(_{\text{ES}}\)) states of the electrons at the QDs and also through the
energy levels in the wetting layer (PL\(_{\text{WL}}\)). The measured PL
spectra of the vertically stacked QD arrays are shown in
Figure 1.

The appearance of PL\(_{\text{WL}}\) peaks indicates the Stranški–Kras-
tanov growth mode of QDs. The photoluminescence of a sam-
ple with one QD layer is characterized by the absence of a peak
of the excited states, and the PL\(_{\text{GS}}\) peak has a full width at half
maximum (FWHM) of ca. 118 meV with a maximum at an
energy of 1.22 eV (Figure 1a). The spectra of samples with two

![Figure 1: a) Photoluminescence spectra of samples with vertically stacked InAs QD arrays; b) full width at half maximum of ground state peaks as a function of the number of InAs QD layers.](image-url)
and three QD layers differ by a decrease of FWHM of the PL_{GS} peaks (Figure 1b) and also by an appearance of PL_{ES} peaks. In this case, the PL_{GS} peaks in the samples with QD vertical stacking are shifted to lower energies (red-shift). The shift value between PL_{WL} peaks and PL_{GS} peaks in the samples increases by 70 meV. It is seen that the FWHM of PL_{GS} peaks decreases with increasing number of InAs QD layers (Figure 1b). As a result, the luminescence spectrum of vertically stacked QD arrays narrows and becomes similar to the large single-QD spectrum [51-53].

The red-shift is also described in [24] for heterostructures with five QD layers of InAs and in [25] for heterostructures with 1–20 QD layers of In_{0.5}Ga_{0.5}As. In both cases, the heterostructures were grown by using MBE. As a comparison of the possibilities of methods for the formation of heterostructures, we note that the average lateral sizes of quantum dots are the same, and their height, in the case of IBD, is higher. The reasons for this may be related to the difference in the formation of growth fluxes. In IBD, much depends on the calibration of the sputtering yield of the target, which affects the kinetics of the surface processes on the substrate.

The influence of the thickness of the i-GaAs barrier layer (ST#2 samples) on the photoluminescence properties of the InAs/GaAs heterostructures is shown in Figure 2. It is seen that a decrease in the i-GaAs layer thickness from 30 to 15 nm is accompanied by a red-shift of the PL peaks with an increase in the intensity (Figure 2a) and decrease of FWHM (Figure 2b).

The energy difference between PL_{GS} and PL_{WL} peaks decreased from 205 to 158 meV when the barrier layer thickness was changed from 30 to 15 nm. Obviously, the thicker the i-GaAs layer, the less it is strained. The coating of QDs with a thin i-GaAs layer creates the conditions for the inheritance of properties from the bottom QD layer by a new layer of InAs QDs. Otherwise, if the thickness of the i-GaAs layer is much higher than the height of the QDs, their sizes and density will be determined by the self-organization on the new growth surface, which can cause a fluctuation in QD sizes during vertical stacking and a degradation of the structural and optical properties of the InAs/GaAs heterostructures. The results of the investigations showed that in the case of three QD layers of InAs separated by 15 nm i-GaAs barriers, the density of the array decreased insignificantly from $1.42 \times 10^{10}$ to $1.33 \times 10^{10}$ cm$^{-2}$. In addition, the average lateral size of the QDs increased by 2 nm. For 30 nm layers of i-GaAs, the average lateral size of the QDs is increased by 2.6 nm, while the density of the top array of quantum dots is decreased to $1.27 \times 10^{10}$ cm$^{-2}$. A similar situation was observed in [24,54]. A significant increase in the characteristic QD sizes during vertical stacking was described in [25] only after the deposition of ten QD layers through thin barriers. The observed effect of red-shift and simultaneous increase of PL intensity with increasing number of QD layers was also described in [24] and mainly is explained by a suppression of relative oscillations of QD heights accompanied by high homogeneity of lateral QD sizes. Thus, a decrease in the i-GaAs layer thickness, on the one hand, limits the fluctuation of QD heights in vertically stacked arrays and stabilizes them and, on the other hand, improves the charge transfer due to the resonant tunneling between the vertical InAs QD layers.

**Photoluminescence properties of InAs/GaAsBi heterointerfaces**

To study the photoluminescence properties of InAs/GaAsBi heterointerfaces, it was necessary to investigate the morphological and structural properties. At first it was necessary to verify the conservation of the stoichiometric composition of
GaAs$_{1-x}$Bi$_x$ during ion-beam deposition. For this purpose, GaAs$_{1-x}$Bi$_x$ films were grown on GaAs(100) substrates. The thickness of the GaAs$_{1-x}$Bi$_x$ layers was 20 nm. The results of XRD studies of the epitaxial layer of GaAsBi are shown in Figure 3a.

Bi is an isovalent dopant material for III–V compounds and replaces group-V element. Consequently, the GaAs$_{1-x}$Bi$_x$ crystal lattice should be a zinc blende lattice. Therefore, a diffraction maximum corresponding to the GaAs(004) lattice planes was measured. This is due to its high structure factor, which governs the maximum diffraction intensity. In Figure 3a, two main peaks are identified: a sharp peak of high intensity at 0 radians ($\theta_s = 33^\circ$) from the GaAs substrate, and low-intensity peaks with negative $\Delta \theta$ due to the strained GaAs$_{1-x}$Bi$_x$ epitaxial lattice. That kind of peak splitting (with respect to the substrate) is usually observed in epitaxial layers with lattice relaxation. The Bi contents were calculated from obtained X-ray diffraction patterns over the linear dependence $x = 6.77 \Delta \theta$ under the condition of a completely strained GaAs$_{1-x}$Bi$_x$ layer [55]. The calculated values of bismuth contents were $x_{Bi} = 0.99\%, 2.98\%$ and 4.96%. The composition of GaAs$_{1-x}$Bi$_x$ epitaxial layers differs from the composition of the sputtering targets. In our opinion, this difference is due to the desorption of Bi from the surface, despite the low deposition temperature of 360 °C. The influence of bismuth on the structural properties of InAs/GaAsBi and InAs/GaAs heterointerfaces was investigated by using Raman spectroscopy.

In accordance with the selection rule, modes of longitudinal optical phonons (LO) and longitudinal acoustic phonons (LA) are allowed in the heterointerfaces. The Raman spectroscopy results are shown in Figure 3b. In addition to the allowed modes, forbidden transverse optical (TO) modes for optical phonons of bulk-like GaAs (268 cm$^{-1}$) and GaBi (182 cm$^{-1}$) are also observed. This indicates a change in the selection rule under Raman scattering in the InAs/GaAs system caused by lattice strain during the incorporation of Bi into the GaAs matrix. In the spectra we can clearly see the longitudinal modes of optical phonons of bulk GaAs (LO) at 283 cm$^{-1}$, InAs QDs (LO) at 248 cm$^{-1}$ and GaBi at 235 cm$^{-1}$. Effects not detectable in the Raman spectrum of the InAs/GaAs heterostructure appear in the spectra of InAs/GaAs$_{1-x}$Bi$_x$ samples in the interval of 150–250 cm$^{-1}$. An increase in $x_{Bi}$ is accompanied by the Raman shift of the GaAs-like and GaBi-like peaks to smaller wavenumbers. Due to decrease of lattice mismatch and, thus, less strain in the InAs/GaAs$_{1-x}$Bi$_x$ heterointerface compared to InAs/GaAs, there is a Raman shift of the peaks of InAs/GaAs$_{1-x}$Bi$_x$ heterostructures relative to the spectrum of the InAs/GaAs heterostructure. The results of Raman spectroscopy are in good agreement with XRD.

Features of PL in the InAs/GaAs$_{1-x}$Bi$_x$ heterostructures are shown in Figure 4a.

The difference between the energies of the PL$_{GS}$ peaks of a sample without Bi and a sample with $x_{Bi} = 4.96$ atom % is 121 meV. This result differs from the red-shift of 23 meV obtained in [56] at room temperature. The reason for this is that InAs QDs in our work have larger sizes and lower density. The results of the investigation of the surface morphology with InAs QDs in samples with GaAs and GaAs$_{1-x}$Bi$_x$ potential barriers are shown in Figure 4b,c. It is seen that there is a decrease of InAs QD density at growth on the GaAs$_{0.95}$Bi$_{0.05}$ sur-
The AFM analysis showed that the density of QDs in InAs/GaAs$_{0.95}$Bi$_{0.05}$ and InAs/GaAs heterosystems was $0.91 \times 10^{10}$ cm$^{-2}$ and $1.53 \times 10^{10}$ cm$^{-2}$, respectively. Doping GaAs with bismuth is accompanied by an increase in the average size of QDs from 14 to 20 nm. The results can be explained by an increasing rate of In surface diffusion on GaAs$_{1-x}$Bi$_x$ during the ion-beam deposition due to the Bi surfactant effect. Note that scientists have different points of view about it. Some argue that doping of Bi during the QD growth process by MBE [17,18,20] and metal-organic vapor-phase epitaxy (MOVPE) [32,33] reduces the In adatom diffusion length and prevents the coalescence of InAs QDs. Then, the QD array density increases. Others argue that the usage of Bi during MOVPE and MBE of InAs QDs [57,58] enhances In adatom diffusion length. In this case, QD sizes increase, and the QD density decreases. Detailed analysis of these works elucidated that in the first case, doping with bismuth was being done during InAs QD growth on GaAs.

In the second case [57,58], the QD growth was being done on GaAs$_{1-x}$Bi$_x$ surfaces. Among other things, temperature conditions also differed from each other. In [20,32,33], doping with Bi on InAs QDs was carried out at 510–530 °C. In our case, as in [52,53], GaAs$_{1-x}$Bi$_x$ layers were grown at temperatures below 500 °C. We also want to emphasize the fact that the covalent radii [59] of Bi = 148 pm and In = 142 pm atoms are very close. Therefore, in the InAs QD growth process on the GaAs$_{1-x}$Bi$_x$ surface, In adatom surface diffusion can be realized both through substituting Ga or Bi vacancies. Increase of QD heights in the InAs/GaAs$_{0.95}$Bi$_{0.05}$ heterosystem is an indirect demonstration of this. Bi has a large atomic size, so it creates elastic strain in the GaAs lattice and thus influences the QD self-organization. Consequently, using Bi as a surfactant on the GaAs growth surface changes the kinetics of In surface diffusion during ion-beam deposition. Unchanged is the circumstance that with an increase in Bi content in GaAs, an increase in the intensity of photoluminescence is observed in spite of a decrease in InAs QD density (Figure 4b,c). The effect is due to the fact that Bi incorporation leads to valence-band splitting of GaAs, shifting the energy maximum of heavy holes deep into the energy gap. This induces a change in the profiles of the valence and the conduction bands of the InAs/GaAs heterostructure. The InAs/GaAs$_{1-x}$Bi$_x$ heterointerface forms a type-II misaligned heterojunction [60]. In accordance with the band diagram in Figure 5, the profile of the valence band of the GaAs$_{1-x}$Bi$_x$ layer shifts in such a way that it forms a well for holes on either side of the InAs QD. In this case, the optical transitions in the direction “quantum dot–barrier layer” begin to
appear. The energy of optical transitions is less than the energy of transitions through the ground states of QDs, which explains the red-shift of the PL peaks (Figure 5b). In addition, we suppose that an increase in the lateral QD sizes also contributes to the red-shift of the photoluminescence peak.

Note that in this case density reduction of the InAs QDs is not a negative factor. QDs being at a more remote distance from each other are able to form a narrower energy sub-band in the conduction band. This effect can be used for the fabrication of multicolor photodetectors.

**Conclusion**

The properties of InAs/GaAs and InAs/GaAsBi heterointerfaces grown by ion-beam deposition are analyzed. Methods of controlling the photoluminescence spectrum of InAs/GaAs heterostructures are considered. It is demonstrated that the vertical stacking of three QD layers decreases the FWHM of the photoluminescence peaks. According to the studies of photoluminescence, it is established that the variation of the photoluminescence spectrum of InAs/GaAs heterostructures can be carried out by isovalent doping of potential barriers for QDs. It is shown that a content of 4.96 atom % Bi in GaAs leads to a red-shift of the PL peaks (Figure 5b). In addition, we suppose that an increase in the lateral QD sizes also contributes to the red-shift of the PL peaks.

**Acknowledgements**

This work was carried out within the framework of the state assignment of the SSC RAS for 2018, (the project registration No. 01201354240), and also with financial support by the Russian Foundation for Basic Research [grant numbers 16–38–60127 mol_a_dk, 17-08-01206].

**References**

1. Martynyuk, P.; Rogalski, A. Prog. Quantum Electron. 2008, 32, 89–120. doi:10.1016/j.pquantelec.2008.07.001
2. Phillips, J.; Kamath, K.; Bhattacharya, P. Appl. Phys. Lett. 1998, 72, 2020–2022. doi:10.1063/1.121252
3. Berryman, K. W.; Lyon, S. A.; Segev, M. Appl. Phys. Lett. 1997, 70, 1861–1863. doi:10.1063/1.118714
4. Abedin, M. N.; Refaat, T. F.; Zawodny, J. M.; Sandford, S. P.; Singh, U. N.; Bandara, S. V.; Gunapala, S. D.; Bhat, I.; Barnes, N. P. Proc. SPIE 2003, 5152, 10. doi:10.1117/12.505887
5. Ryzhii, V. J. Appl. Phys. 2001, 89, 5117. doi:10.1063/1.1359759
6. Phillips, J. J. Appl. Phys. 2002, 91, 4590–4594. doi:10.1063/1.1455130
7. Kudo, M.; Mishima, T.; Ishimoto, S.; Nakaoka, T.; Arakawa, Y. Phys. E (Amsterdam, Neth.) 2004, 21, 275–278. doi:10.1016/j.physe.2003.11.030
8. Shuhui, Z.; Lu, W.; Zhenwu, S.; Yanxiang, C.; Hailao, T.; Huaiju, G.; Haiqiang, J.; Wexin, W.; Hong, C.; Liangch, Z. Nanoscale Res. Lett. 2012, 7, 87. doi:10.1186/1556-276x-7-87
9. Böhrer, J.; Kroost, A.; Bimberg, D. Appl. Phys. Lett. 1994, 64, 1992–1994. doi:10.1063/1.111716
10. Hayne, M.; Provooost, R.; Zundel, M. K.; Manz, Y. M.; Eberl, K.; Moshchalkov, V. V. Phys. Rev. B 2000, 62, 10324–10328. doi:10.1103/physrevb.62.10324
11. Sugisaki, M.; Ren, H.-W.; Nair, S. V.; Nishi, K.; Masumoto, Y. Phys. Rev. B 2002, 66, 235309. doi:10.1103/physrevb.66.235309
12. Ribeiro, E.; Maitlez, R. L.; Carvalho, W.; Jr.; Ugarte, D.; Medeiros-Ribeiro, G. Appl. Phys. Lett. 2002, 81, 2935–2935. doi:10.1063/1.1513215
13. Liu, G.; Chuang, S.-L.; Park, S.-H. J. Appl. Phys. 2000, 88, 5554–5561. doi:10.1063/1.1319328
14. Liang, B.; Lin, A.; Pavarelli, N.; Reyner, C.; Tatebayashi, J.; Nunna, K.; He, J.; Ochalski, T. J.; Huyet, G.; Huffer, D. L. Nanotechnology 2009, 20, 455604. doi:10.1088/0957-4484/20/45/455604
15. Souafi, M.; Baira, M.; Nasr, O.; Alounae, M. H. H.; Maaref, H.; Sfaxi, L.; Ilahi, B. Materials 2015, 8, 4699–4709. doi:10.3390/ma8064899
16. Nabataani, Y.; Matsumoto, T.; Sasikala, G.; Suemune, J. I. Appl. Phys. 2005, 98, 063502. doi:10.1063/1.2041846
17. Wang, L.; Pan, W.; Chen, X.; Wu, X.; Shao, J.; Wang, S. Opt. Mater. Express 2017, 7, 4249–4257. doi:10.1364/ome.7.004249
18. Wang, L.; Zhang, L.; Yue, L.; Liang, D.; Chen, X.; Li, Y.; Lu, P.; Shao, J.; Wang, S. Crystals 2017, 7, 63. doi:10.3390/cryst7030063
19. Maia, A. D. B.; da Silva, E. C. F.; Quivy, A. A.; Bindilatti, V.; de Aquino, V. M.; Dias, I. F. L. J. Phys. Appl. Phys. 2013, 114, 083708. doi:10.1063/1.4818610
20. Dasika, V. D.; Kriyov, E. M.; Nair, H. P.; Maddox, S. J.; Park, K. W.; Jung, D.; Lee, M. L.; Yu, E. T.; Bank, S. R. Appl. Phys. Lett. 2014, 105, 253104. doi:10.1063/1.4904825
21. Chebotarev, N. S.; Pashchenko, A. S.; Libin, L. S.; Zivotova, E. N.; Erimeev, G. A.; Libin, M. L. Beilstein J. Nanotechnol. 2017, 8, 12–20. doi:10.3765/bjnano.8.2
22. Chakraborti, S.; Stiff-Roberts, A. D.; Su, X. H.; Bhattacharya, P.; Aryawanisa, G.; Perera, A. G. J. U. J. Phys. D: Appl. Phys. 2005, 38, 2135–2141. doi:10.1088/0022-3727/38/13/0309
23. Brunkov, P. N.; Kovsh, A. R.; Ustinov, V. M.; Musikhin, Y. G.; Ledentsov, N. N.; Koninkov, S. G.; Polimeri, A.; Patane, A.; Main, P. C.; Eaves, L.; Kapteyn, C. M. A. J. Elecrhon. Mater. 1999, 28, 486–490. doi:10.1016/s1164-9989(99)00099-6
24. Nakata, Y.; Sugiyama, Y.; Futatsugi, T.; Yokoyama, N. J. Cryst. Growth 1997, 175–176, 719–719. doi:10.1016/s0022-2488(96)00814-7
25. Tsatsulnikov, A. F.; Egorov, A. Y.; Zhukov, A. E.; Kovsh, A. R.; Ustinov, V. M.; Ledentsov, N. N.; Maksimov, M. V.; Volovik, B. V.; Suvorova, A. A.; Bert, N. A.; Kop'ev, P. S. Semiconductors 1997, 31, 722–725. doi:10.1134/s1063782697050178
26. Liu, H. Y.; Steer, M. J.; Badcock, T. J.; Mowbray, D. J.; Skolnick, M. S.; Suarez, F.; Ng, J. S.; Hopkinson, M.; David, J. P. R. J. Appl. Phys. 2006, 99, 046104. doi:10.1063/1.2173188
27. Popescu, D. P.; Eliseev, P. G.; Stintz, A.; Malloy, K. J. Semicond. Sci. Technol. 2004, 19, 33–38. doi:10.1088/0268-1242/19/1/005
28. Mamutin, V. V.; Egorov, A. Y.; Kryzhanovskaya, N. V.; Mikhrin, V. S.; Nadtochy, A. M.; Pirogov, E. V. Semiconductors 2008, 42, 805–812. doi:10.1134/s1063782608070105
29. Wu, J.; Shan, W.; Walukiewicz, W. Semicond. Sci. Technol. 2002, 17, 860–869. doi:10.1088/0268-1242/17/8/315
30. Keizer, J. G.; Ullod, J. M.; Ulltra, A. D.; Koennaad, P. M. Appl. Phys. Lett. 2014, 104, 053116. doi:10.1063/1.4864159
31. Ulltra, A. D.; Reyes, D. F.; Ullod, J. M.; González, D.; Ben, T.; Guzman, A.; Hierro, A. Appl. Phys. Lett. 2014, 105, 043105. doi:10.1063/1.4891557
32. Zvonkov, B. N.; Karpovich, I. A.; Baidus, N. V.; Filatov, D. O.; Morozov, S. V. Semiconductors 2001, 35, 93–98. doi:10.1134/1.340297
33. Zvonkov, B. N.; Karpovich, I. A.; Baidus, N. V.; Filatov, D. O.; Morozov, S. V.; Gushina, Y. Y. Nanotechnology 2000, 11, 221–226. doi:10.1088/0957-4484/11/4/306
34. Lunin, L. S.; Sysoev, I. A.; Blagina, L. V.; Blagin, A. V.; Barannik, A. A. Inorg. Mater. 2009, 45, 841–845. doi:10.1134/s0020163809060020
35. Affimova, D. L.; Lunin, L. S.; Lunina, M. L. Inorg. Mater. 2014, 50, 113–119. doi:10.1134/s0020163814020010
36. Reyes, D. F.; Bestfitman, F.; Hunter, C. J.; Sales, D. L.; Sanchez, A. M.; David, J. P. R.; Gonzalez, D. Nanoscale Res. Lett. 2014, 9, 23. doi:10.1186/1556-276x-9-23
37. Plak, A. J.; France, R.; Beaton, D. A.; Alberi, K.; Simon, J.; Mascarenhas, A.; Jiang, C.-S. J. Cryst. Growth 2012, 338, 107–110. doi:10.1016/j.jcrysgro.2011.10.040
38. Janotti, A.; Wei, S.-H.; Zhang, S. B. Phys. Rev. B 2002, 65, 115203. doi:10.1103/physrevb.65.115203
39. Greene, J. E. Crit. Rev. Solid State Mater. Sci. 1983, 11, 189–227. doi:10.1080/01611598308244063
40. Seshan, K., Ed. Handbook of Thin Film Deposition, 3rd ed.; Elsevier: Amsterdam, Netherlands, 2012. doi:10.1016/c2009-0-64359-2
41. Reagor, D. W.; Bulko, V. Y. Nat. Mater. 2005, 4, 593–596. doi:10.1038/nmat1402
42. Razek, N.; Schindler, A.; Rauschenbach, B. Vacuum 2007, 81, 974–978. doi:10.1016/j.vacuum.2006.12.003
43. Kim, W. J.; Koo, W. H.; Jo, S. J.; Kim, C. S.; Baik, H. K.; Lee, J.; Im, S. J. Vac. Sci. Technol., B: Microelectron. Nanometer Struct.–Process., M eas., Phenom. 2005, 23, 2357–2362. doi:10.1116/1.2102969
44. Ke, J. J.; Tsai, K. T.; Dai, Y. A.; He, J. H. Appl. Phys. Lett. 2012, 100, 053503. doi:10.1063/1.3680251
45. Szczytbowiski, J.; Czapla, A.; Jacobowski, J. Thin Solid Films 1996, 288, 191–194. doi:10.1016/0040-6090(96)00174-5
46. Greene, J. E.; Barnett, S. A.; Cadlen, K. C.; Ray, M. A. J. Cryst. Growth 1982, 56, 389–401. doi:10.1016/0022-2211(82)90458-4
47. Takeuchi, M.; Sakagawa, Y.; Nagaoka, H. Thin Solid Films 1986, 139–152. doi:10.1016/0040-6090(86)90421-6
48. Chebotarev, S. N.; Pashchenko, A. S.; Williamson, A.; Lunin, L. S.; Ikhva, V. A.; Gamidov, V. A. Tech. Phys. Lett. 2015, 41, 661–664. doi:10.1134/s1063785015070056
49. Chebotarev, S. N.; Pashchenko, A. S.; Ikhva, V. A.; Lunina, M. L. J. Nanotechnol. 2016, 5340218. doi:10.1155/2016/5340218
50. Dekel, E.; Regelman, D. V.; Gershoni, D.; Ehrenfreund, E.; Schoenfeld, W. V.; Petroff, P. M. Solid State Comm. 2001, 117, 395–400. doi:10.1016/s0038-1098(00)00483-x
51. Shtrichman, I.; Metzner, C.; Gerhardt, B. D.; Schoenfeld, W. V.; Petroff, P. M. Phys. Rev. B 2002, 65, 081303. doi:10.1103/physrevb.65.081303
52. Regelman, D. V.; Dekel, E.; Gershoni, D.; Ehrenfreund, E.; Williamson, A. J.; Shurinway, J.; Zunger, A.; Schoenfeld, W. V.; Petroff, P. M. Phys. Rev. B 2001, 64, 165301. doi:10.1103/physrevb.64.165301
53. Binberg, D.; Ledentsov, N. N.; Grundman, M.; Kirstaedter, N.; Schmidt, O. G.; Mao, M. H.; Ustinov, V. M.; Egorov, A. Y.; Zhukov, A. E.; Kopiev, P. S.; Allerov, Zh. I.; Ruvimov, S. S.; Göesele, U.; Heydenreich, J. Jpn. J. Appl. Phys., Part 1 1996, 35, 1311. doi:10.1143/jjap.35.1311
54. Steel, J. A.; Lewis, R. A.; Horvat, J.; Nancarrow, M. J. B.; Henini, M.; Fan, D.; Mazur, Y. I.; Schmidlbauer, M.; Ware, M. E.; Yu, S.-Q.; Salamo, G. J. Nat. Sci. Rep. 2016, 6, 28860. doi:10.1038/srep28860
55. Wang, P.; Pan, W.; Wu, X.; Liu, J.; Cao, C.; Wang, S.; Gong, Q. Nanoscale Res. Lett. 2016, 11, 280. doi:10.1186/s11671-016-1470-1
56. Okamoto, H.; Tawara, T.; Gotoh, H.; Kamada, H.; Sogawa, T. Jpn. J. Appl. Phys. 2010, 49, 056GJ01. doi:10.1143/jap.49.056GJ01
57. Fan, D.; Zeng, Z.; Dorogan, V. G.; Hirono, Y.; Li, C.; Mazur, Y. I.; Yu, S.-Q.; Johnson, S. R.; Wang, Z. M.; Salamo, G. J. Mater. Sci.: Mater. Electron. 2013, 24, 1635–1639. doi:10.1007/s10854-012-0987-z
58. Kressel, H.; Butler, J. K. Semiconductor lasers and heterojunction LEDs; Academic Press: New York City, NY, U.S.A., 2012. doi:10.1016/b978-0-12-426250-8.x5001-5
59. Nag, B. R. Physics of Quantum Well Devices; Kluwer Academic Publishers: Dordrecht, Netherlands, 2000. doi:10.1007/0-306-47127-2

License and Terms
This is an Open Access article under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0). Please note that the reuse, redistribution and reproduction in particular requires that the authors and source are credited.

The license is subject to the Beilstein Journal of Nanotechnology terms and conditions: (https://www.beilstein-journals.org/bjnano)

The definitive version of this article is the electronic one which can be found at: doi:10.3762/bjnano.9.261