Heavily Si-doped InAs photoluminescence measurements

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In this paper, we present experimental results of photoluminescence for series of InAs:Si heavily doped samples, with doping level varying from 1.6×10^{16} cm⁻³ to 2.93×10^{18} cm⁻³. All samples were grown using MBE system equipped with a valved arsenic cracker. The measurements were performed in the temperature range of 20 K to 100 K. Although the Mott transition in InAs appears for electron concentrations above 10^{14} cm⁻³, Burstein-Moss broadening of photoluminescence spectra presented in this article was observed only for samples with concentration higher than 2×10^{17} cm⁻³. For the samples with lower concentrations two peaks were observed, arising from the band gap and defect states. The intensity of the defect peak was found to be decreasing with increasing temperature as well as increasing concentration, up to the point of disappearance when the Burstein-Moss broadening was visible.

Keywords: InAs; photoluminescence; Burstein-Moss; heavy doping

1. Introduction

InAs is a proper material for production of infrared photodetectors and infrared light emitting diodes working in 3 µm to 5 µm wavelength range [1-3]. Contrary to the other III-V materials, such as antimony compounds, InAs devices may operate at room temperature [4], which in turn, leads to the reduction of photodetector cost. Most of InAs photodiode heterostructures consist of undoped absorber and contacts made of p-type and strong n-type $(1 \times 10^{18} \text{ cm}^{-3})$ materials [5– 8]. InAs of both p-type and n-type have been already studied in many works [9–11]. Therefore, the great number of InAs parameters, such as band gap energy as well as defect, donor, acceptor energies, have been determined [11, 12]. However, the characterization of heavily n-doped InAs is still not sufficient. Theoretical calculations showed that for doping higher than 10^{17} cm⁻³, Burstein-Moss shift and band gap peak broadening appear [13]. These phenomena may be described as an increased probability of indirect transitions, which in turn, leads to their appearance, contributing

to the measured spectra [13]. This effect was also visible in experimental results presented by Gladkov et al. [14].

In this work, we present experimental results that confirm broadening of the band gap peak only for highly doped samples. Additionally observed effects are: the decrease of defect-induced peak at 35 meV [11, 15] and the decrease of defect peak intensity with increasing temperature. They may all be explained by the analysis of electron energy and momentum in highly doped InAs.

2. Experimental

The samples studied in this paper were grown on GaAs (0 0 1) substrate with 2° offcut towards $\langle 1 1 0 \rangle$ by a solid source RIBER COMPACT 21 DZ molecular beam epitaxy (MBE) system equipped with a valved arsenic cracker. The growth details have been reported in the literature [16]. The InAs layers were grown at 400 °C, with an As/In flux ratio of 8.5. The growth rate was 0.26 µm/h. The n-type doping was introduced by opening the shutter of Si effusion cell while growing InAs epilayers. The level of doping was calibrated by the temperature of Si effusion cell. Hall effect

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measurements using van der Pauw method were performed by ECOPIA Hall Measurements System, in order to assess the electrical parameters of the samples between 80 K and room temperature.

A high-resolution X-ray diffractometer of PANalytical X'Pert was utilized to evaluate the crystallographic properties of the samples. The CuK α_1 radiation ($\lambda \sim 1.5406$ Å) originating from a line focus was used. The X-ray beam was monochromatized by four bounce, Ge (0 0 4) hybrid monochromator. The measurements were made in both ω and 2θ - ω directions.

Basic parameters of the samples are presented in Table 1.

Table	1.	Concentratio	on	and	XRD	Full	Width	at	Half
Maximum (FWHMs) for measured samples.								les.	

Sample	Concentration	FWHM of XRD			
	$[cm^{-3}]$	peak [arc sec]			
A	$1.60 imes 10^{16}$	37			
В	$1.80 imes10^{16}$	27			
С	$2.00 imes 10^{16}$	32			
D	$2.13 imes 10^{17}$	39			
Е	$2.93 imes 10^{18}$	22			

Photoluminescence was measured using Vertex v70 FT-IR spectrometer in a step-scan mode, equipped with lock-in [17]. As an excitation source a 630 nm diode laser with mechanical chopper was used. During measurements, the samples were placed in a cryostat equipped with piezo table.

3. Results and discussion

Photoluminescence spectra for all measured samples are presented in Fig. 1. The spectrum for sample A consists of two spectrally separated peaks attributed to InAs band gap (0.41 eV) [14] and defect band (0.38 eV) [11]. On the other hand, the spectrum of sample E consists of solely one wide peak arising from band gap, broadened by Burstein-Moss effect. It is worth noting that we do not observe Mahan exciton [9], since we do not observe any peak higher than band gap energy, so it cannot interfere with Burstein-Moss broadening. Although the concentration for all samples is much higher than 1×10^{14} cm⁻³ (Mott transition in InAs), we observe Burstein-Moss broadening only for the two most heavily doped samples: D (2×10^{17} cm⁻³) and E (3×10^{18} cm⁻³). It is in agreement with previous calculations [13].

In the presented spectra we observe no correlation between intensity of defect peak and samples quality obtained from the XRD measurements. For sample C, Full Width at Half Maximum (FWHM) of the XRD peak is higher than for sample B (Table 1), while the intensity of photoluminescence defect peak is much lower. Moreover, for sample D, in spite of the highest FWHM value of the XRD peak (Table 1), there is no defect peak observed. We suspect the intensity of defect peak decreases rapidly with increasing concentration (above 1×10^{16} cm⁻³). Similar effect was observed for the low doped p-type GaSb samples [18]. However, in that case the defect peak was created by acceptor-donor interaction while Fermi level was inside band gap or even below valence band [18]. In our experiments Fermi level is above conduction band and the defect states are 35 meV above the valence band [11]. All those details suggest that the defect peak intensity decrease mechanisms for InAs:Si and GaSb samples are different.

One of the possible explanations may be the appearance of non-zero momentum electrons that take part in creation of band gap peak, whereas the defect peak is formed due to defect states which possess very low momentum. If defect states momentum is close to zero, then for low-doped sample in which almost all electrons have zero momentum there are many electrons that may recombine into defect states. If the concentration is increased, there are much fewer electrons with zero momentum and much more with higher momentum [13]. In this case, electrons have much lower probability to recombine into the defect states.

The disappearance of the defect peak is also observed with increasing temperature (Fig. 2). For samples A and B defect peak intensity is the highest at 20 K and disappears at 80 K and 100 K, respectively. In this case, there is no Burstein-Moss shift, but instead Fermi-Dirac distribution

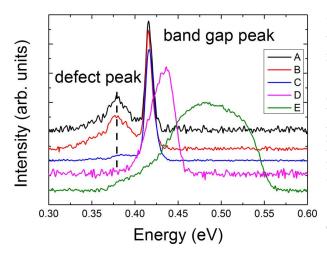


Fig. 1. Photoluminescence spectra for all samples, measured at 20 K.

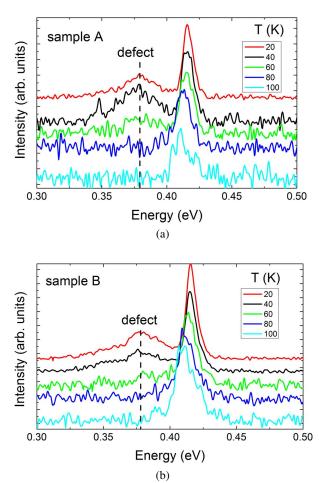


Fig. 2. Photoluminescence spectra for samples A (a) and B (b) obtained in the temperature range from 20 K to 100 K.

flattening takes place. It may cause the increase of the amount of electrons with higher energy and momentum. However, there are also other effects that shift the position of band gap peak to lower energies [12]. Thus, it is difficult to conclude whether the momentum of electrons is higher or not. Other possible explanation is that the energy dispersion at high temperature is stronger for defect states than for the bottom of conductive band and the top of valence band.

4. Conclusions

Presented results confirm previous calculations which predict the appearance of Burstein-Moss broadening only for InAs samples doped higher than 1×10^{17} cm⁻³. We have also shown the decrease of the defect peak intensity with both increasing concentration (taking place above Mott transition) and increasing temperature.

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