

Electronic Properties of (B) GaAs/GaAs Heterostructure Capped by Ga₂O₃Dalanda Slimi¹, Faouzi Saidi^{1,2*} and Hassen Maaref¹¹Laboratory of Micro-Optoelectronics and Nanostructures (LR99ES29), University of Monastir, Faculty of Sciences Monastir, Tunisia²Institut Supérieur des Sciences Appliquées et Technologie de Sousse, Université de Sousse, Tunisia***Corresponding Author**

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Submitted: 2024, Feb 03; **Accepted:** 2024, Jun 25; **Published:** 2024, Jul 04**Citation:** Slimi, D., Saidi, F., Maaref, H. (2024). Electronic Properties of (B)GaAs/GaAs Heterostructure Capped by Ga₂O₃. *Curr Res Env Sci Eco Letters*, 1(2), 01-10.**Abstract**

In this work we performed optical studies of Ga₂O₃/GaAs, BGaAs/GaAs and Ga₂O₃/BGaAs/GaAs epilayers obtained by chemical vapor deposition (MOCVD) at different growth temperatures (580°C and 670°C). This enables the development of heterostructures based on BGaAs/GaAs thin films covered by Ga₂O₃. We focused our research on the influence of power excitation and temperature on the photoluminescence (PL) properties. It has shown the presence of emission energy around 1.2 eV and a red shift after the deposition of Ga₂O₃ on BGaAs/GaAs at low temperature (10K). The power study highlights the dominance of the Ga₂O₃ layer during the emission via the donor-acceptor transitions. The evolution of the emission energy temperature for the BGaAs/GaAs thin film allowed us to explain the origin of the excitonic recombination. The luminescence within the new Ga₂O₃/BGaAs/GaAs heterostructure is controlled by the localization of carriers in energy fluctuation potentials created by inhomogeneous distributions of VGa families. We have shown that this proposed heterostructure based on III-V semiconductors could broaden the emission spectrum and hence increase the conversion efficiency as the absorption increases.

Keywords: GaAs, BGaAs, Ga₂O₃, MOCVD, PL**1. Introduction**

Right now, III-V semiconductor materials based on GaAs, InAs (BAs) and their ternary alloys BxGa1-xAs (InxGa1-xAs) are favored for their potential use in photonic, electronic and optoelectronic devices such as blue light emitting diodes and green high electron mobility transistors (HEMT) and metal oxide semiconductor capacitors (MOSCAP) diode lasers thin-film solar cells Electrooptic waveguide modulators metal-insulator-semiconductor photodetectors [1-10].

In fast electronics and optoelectronics, as silicon has been well developed and commercialized in the photovoltaic field due to its high absorbance, abundance and low cost, it remains insufficient in the optoelectronic field and more precisely for the design of transmitters due to the low mobility of the carriers and its indirect gap which reinforces the non-radiative transitions. On the other hand, III-V semiconductors are favored by their direct gap, their high mobility and their high electrical conductivity [11]. The main efforts have been undertaken to develop new photovoltaic conversion routes based on III-V semiconductors such as GaAs,

In ... etc. [12].

Moreover, they are materials of choice for the development of lasers or detectors operating in the telecommunication domain at wavelengths of 1.3μm [13]. In this context, our mastered heterostructure is based on the III-V semiconductors Ga₂O₃/BGaAs/GaAs. The latter generates a new structure applicable in optoelectronics emitting in the 1.3-1.55 micrometer range [12]. The choice of GaAs as substrate due to its high carrier confinement in the active region, its infrared emission and its low cost [14]. The introduction of Boron in GaAs has allowed to decrease the emission energy compared to GaAs and reach the wavelength 1.33 micrometer [15]. Indeed, it acts on the state of strain but it is limited by the incorporation of boron we cannot reach concentrations < 8% this is accompanied by the presence of energy levels in the band gap generally related to boron aggregates (clusters) [16,17]. The advantage of adding Ga₂O₃ is that it broadens the emission range since it emits in the UV range and is a transparent conductive material [18]. From an energy point of view, these materials have a different energy diagram. In fact,

these three semiconductors are distinguished by their band gaps which are 1.42 eV for GaAs, 1.34 eV for BGaAs and 4.6–4.9 eV for Ga₂O₃ [19,20]. In this heterostructure we find energy levels below the BC of BGaAs. Gallium oxide also has deeper energy levels in its optical gap at energies of 2.4, 2.75, 3 and 3.15 eV [21]. These different gap energies make a richness for us in terms of optoelectronic applications.

1.1. Experimental Details

The heterostructure we studied consists of a GaAs substrate, a buffer layer on which a thin layer of BGaAs is deposited, a gallium arsenide barrier followed by a gallium oxide Ga₂O₃ layer. The samples are grown by MOCVD in a horizontal T-shaped reactor at atmospheric pressure [22–24]. GaAs exhibits a 1° disorientation from the (001) plane in the [110] direction. Triethylgallium (TEG) and diborane (B₂H₆) are used as group III precursors for gallium (Ga) and boron (B), respectively. Arsine (AsH₃) is used as group V precursor for arsenic (As). All the growths of these structures took place under hydrogen, which is the carrier gas used in EPVOM.

Photoluminescence measures were performed between 10 and 300K by holding the samples in an unrestricted cycle helium cryostat. The sample was excited by the green line (wavelength 514.5 nm) of an argon-ion laser (Ar⁺) with a maximum power density of 80 W/cm². Photons emitted by the sample are scattered by a high-resolution spectrometer (Jobin-Yvon monochromator HDR1: focal length 0.6). Detection is via a silicon detector (spectral range between 800 and 1200 nm) with integrated amplifier.

2. Results and Discussion

Figure 1 shows that the low temperature PL spectrum of GaAs has a sharp emission peak at 1.516 eV [24,25]. When depositing Ga₂O₃/GaAs, we observe that the emission energy at 1.43 eV red shifted by 80 meV compared to GaAs dominates the GaAs PL emission peak. This shift may be a consequence of the excess of carbon concentration since the deposition was done by MOCVD [26].

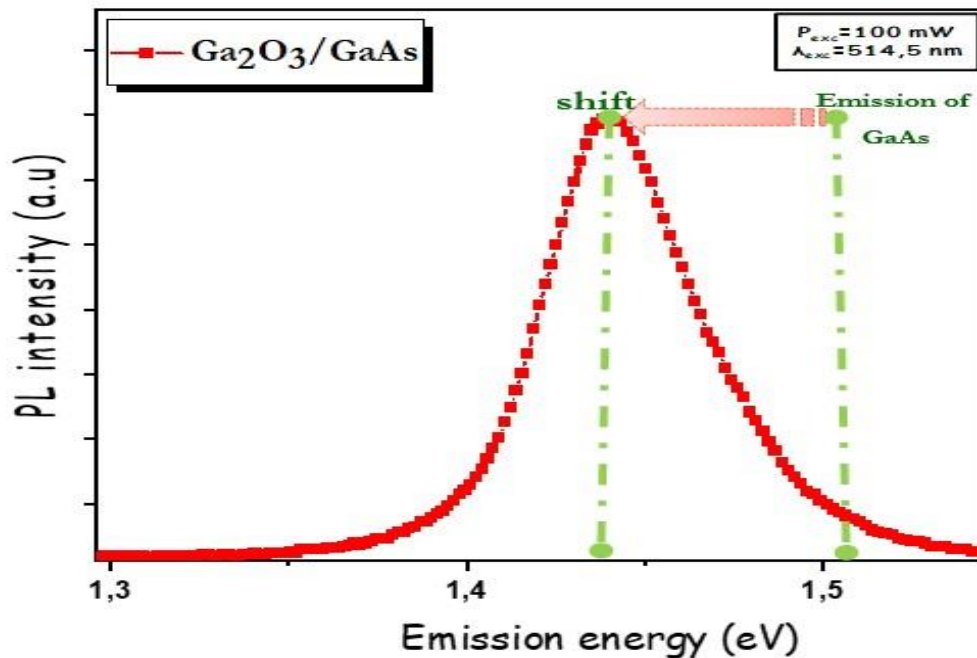


Figure 1: Room Temperature PL Spectrum for Ga₂O₃/GaAs.

To confirm the nature of this heterostructure, PL measurements are performed at 10K and under different excitation power densities on Ga₂O₃/GaAs (see Figure 2).

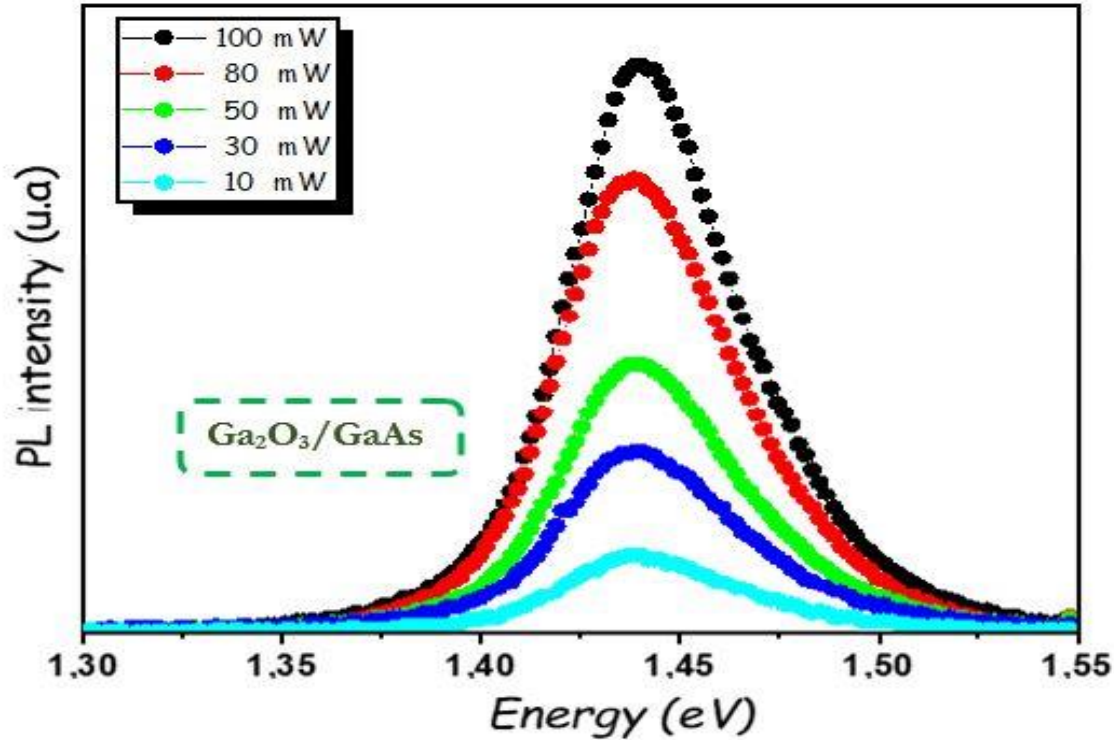


Figure 2: Shows the Low Temperature Photoluminescence PL Spectra of Ga_2O_3 Grown on GaAs

Indeed, as the excitation power increases, the intensity increases as well as the full width at half maximum. Moreover, a small blue energy shift is observed in the 10 - 100 mW excitation power range. From this power study, we can see that this heterostructure is of type I [27]. In order to clarify the PL transition medium, the

analysis of PL intensity as a function of the excitation density is performed. A power law fitting allowed determining the nature of this transition as shown in Figure 3. This law is given by the following expression:

$$I_{PL} \propto P_{ex}^n$$

Where I_{pl} is the PL intensity, P_{ex} is the excitation power, and n is the dimensionless exponent which can take the following forms [27].

- $n \leq 1$: Recombination from the free state to the bound state or from the bound state to another bound state.
- $n = 1$: Band to band recombination.
- $0 < n < 1$: Excitonic recombination.

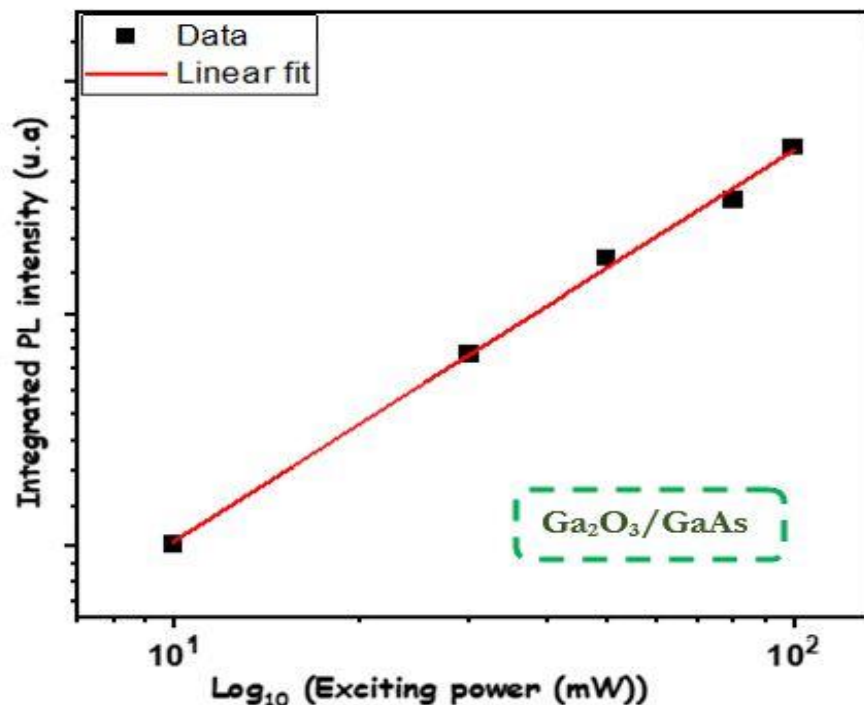


Figure 3: PL Intensity as a Functions of Exciter Power for Ga₂O₃/Gagas Heterostructure

In our structure n= 0, 7: the type of recombination is a transition from the donor state to the acceptor state (D-A), which has already been proven by the work of R. Fornari in which the excitation is done by cathodoluminescence with an energy varying between 5 and 30 KeV [21].

Figure 4 shows the different spectra of the Ga₂O₃/GaAs sample by

varying the temperature from 10 to 300 k. An increase in PL intensity and width at half maximum is detected in the range of 10 to 60 K. For T<60 K, a drop in PL intensity while increase in LWHM is marked this and justified by the PL intensity ratio $I_{10} / I_{300} \sim 10$ which shows that this reflects a strongly present non-radiative character due to defects and impurities.

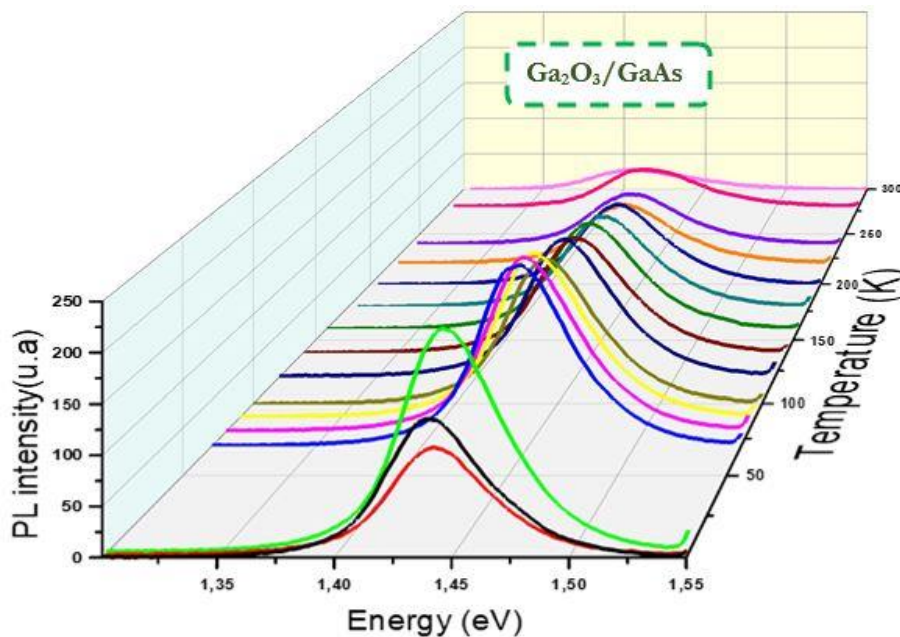


Figure 4: PL Spectra at Different Temperatures of the Ga₂O₃/GaAs Layer

In order to confirm and validate the presence of the localization phenomenon, we studied the variation of the energy position of the photoluminescence peak as a function of the temperature of

the Ga₂O₃/GaAs epilayers (see Figure 5). These spectra could be adjusted by the empirical law proposed by Varshni using Equation (1) [28,29].

$$E_g(T) = E_g(0) - \frac{\alpha T^2}{\beta + T} \quad (1)$$

Where $E_g(0)$ is the energy of the gap at T=0 K, α is an empirical parameter related to the joint density of states and β an effective temperature. By fitting the experimental results with Equation (1), we have deduced the different values of α and β shown in Table.

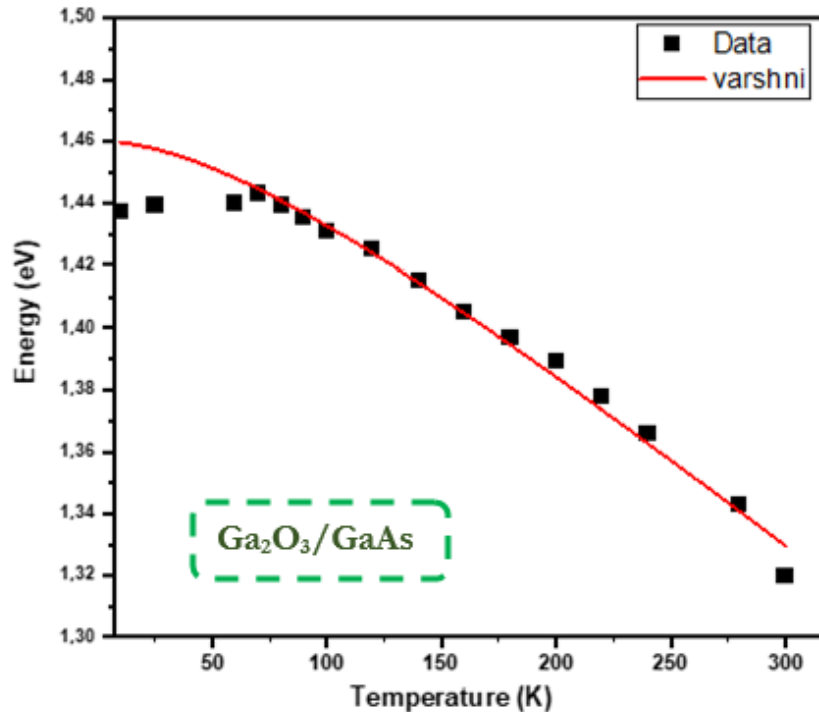


Figure 5: The Evolution of PL Energy as a Function of Temperature

Sample Name	α (ev/K) 10^{-4}	β (K)	$E_g(0)$ (eV)
Ga ₂ O ₃ /GaAs	6,159	125	1,46
BGaAs/GaAs	5,969	168	1,343

Table: Varshni parameter of Ga₂O₃/GaAs and BGaAs grown at 670 °C and 580 °C for Boron Composition X_b = 1.5%, Respectively

At low temperature, Varshni's law does not correspond to the experiment given for the Ga₂O₃/GaAs epilayer. It exhibits an anomalous behavior, the so-called S-shape. From 10 to 70 K the emission energy increases, but a decrease is observed between 70 and 100 K. This shape is a signature of the localization phenomenon, which can be related to localized states in the Ga₂O₃ layer. At high temperature, no distinction can be observed between the expected Varshni energy band gap and the PL energy peak because the carriers are thermally delocalized and one joins the evolution guided by band-to-band recombination; this behavior can be attributed to

the emission of GaAs.

After studying the Ga₂O₃/GaAs reference structure, we now proceed to the characterization of the Ga₂O₃/BGaAs/GaAs heterostructure to compare the effect of boron and gallium oxide on GaAs. To compare the optical properties of BGaAs/GaAs layer and Ga₂O₃/BGaAs/GaAs heterostructures, we first demonstrate low-temperature photoluminescence spectra. For the BGaAs/GaAs layer (Figure 6a), PL emission from GaAs substrates is dominated by the carbon impurity transition (e-CAs) and its phonon-optical

replica (LO) [30]. In addition, an asymmetric PL band named (HE) centered at 1.334 eV with a half-value width of 89 meV has been reported. This emission band is associated with exciton recombination of BGaAs [13]. When the gallium oxide layer is deposited on BGaAs/GaAs, a broad peak centered around 1.20 eV with a full width at half maximum of 185 meV is observed (see Figure 6b). Thus, we notice that after the deposition a broader emission

peak is observed with an increase in the full width at half maximum. Figure 7 shows PL spectra at different powers of the BGaAs/GaAs structure and of the Ga₂O₃/BGaAs/GaAs heterostructure performed at low temperature and with an excitation wavelength equal to 514.5 nm ($X_b=1.5\%$). For both structures, we observe that when we increase the excitation power, the PL intensity increases with increasing full width at half maximum.

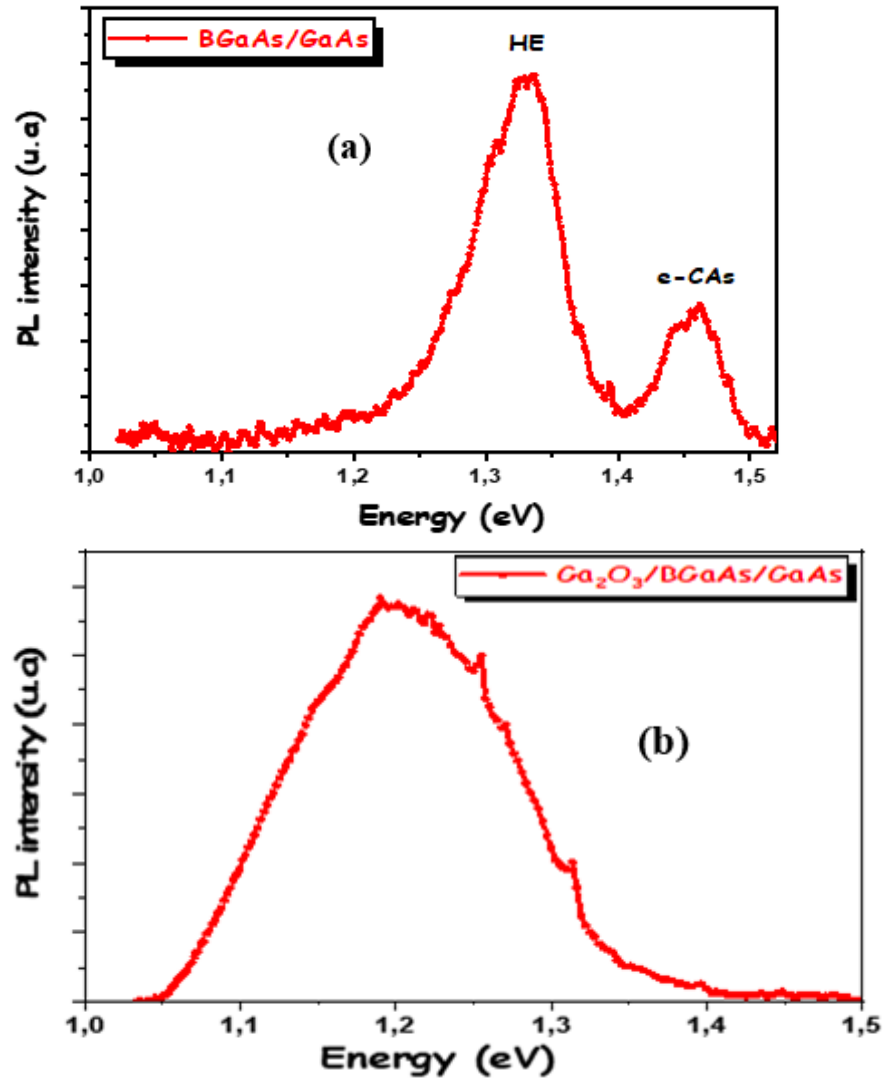


Figure 6: PL Spectra at 10K: of the BGaAs/GaAs Layer (A) and of the Ga₂O₃/BGaAs/GaAs Heterostructure (B).

From Figure 7a, we observe a broadening of the PL peak of BGaAs. This is due to the gradual filling of the energy levels introduced by boron in the GaAs gap. Moreover, Figure 7b shows that the broad luminescent band is shifted towards the low energy. Indeed, this

important shift towards high wavelengths is related to the gradual filling of energy levels introduced by the Ga₂O₃ oxide deposited on the BGaAs layer.

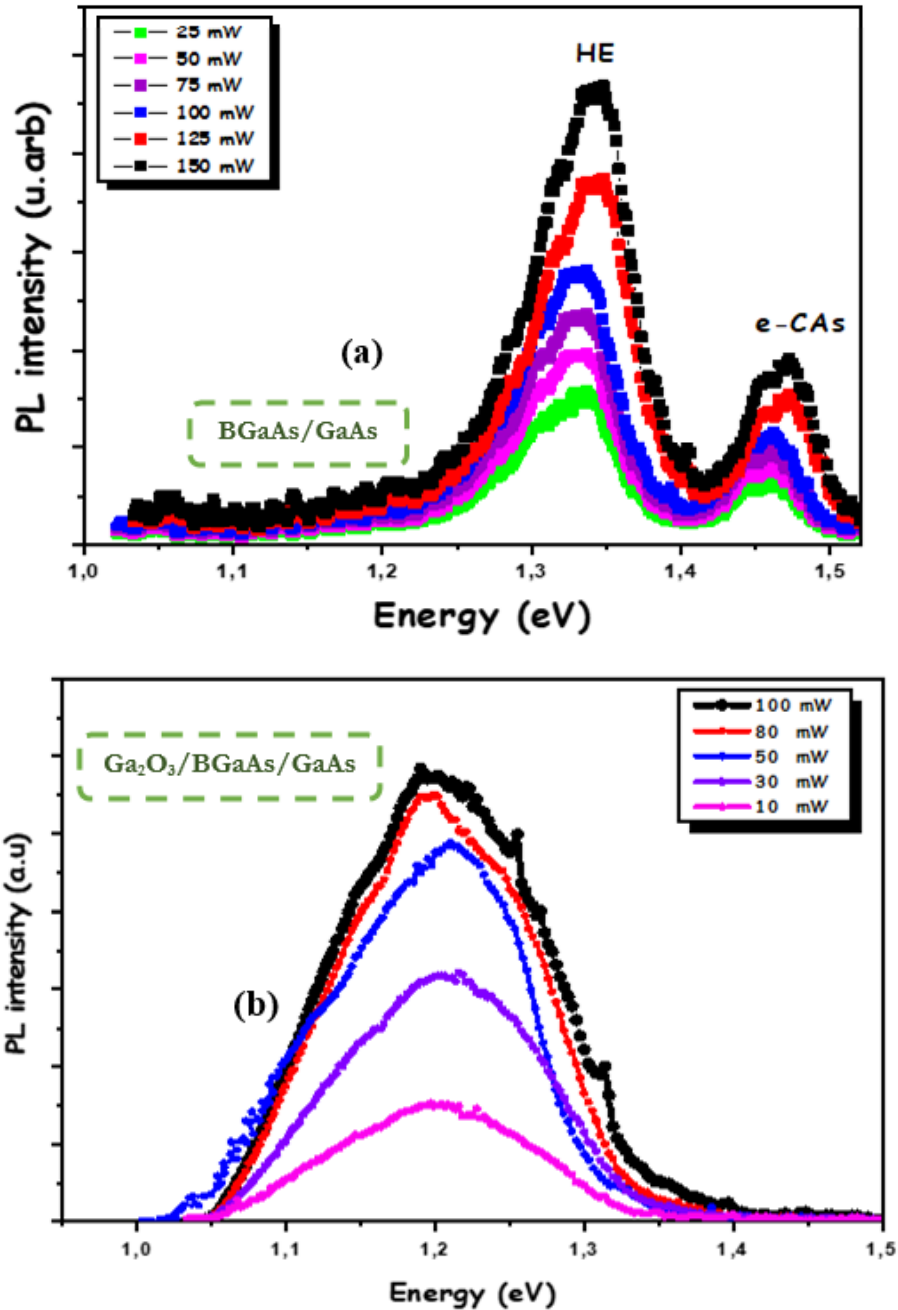


Figure 7: PL Spectra at Different Excitation Powers: of the BGaAs/GaAs Structure (A) and of the Ga₂O₃/BGaAs/GaAs Heterostructure (B)

In the same framework, the evolution of the PL intensity as a function of the exciter power is shown for the Ga₂O₃/BGaAs/GaAs heterostructure (see Figure 8). A power law fitting has allowed us to

determine the coefficient n which proves that the transition is of donor-acceptor type since $n < 1$ ($n = 0.5$). This n is comparable with the one found by Roberto and his collaborators [20].

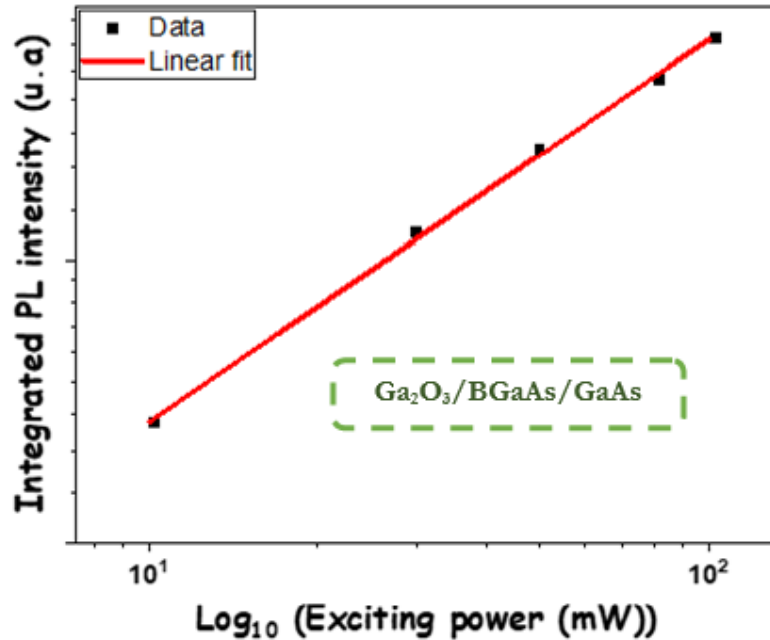


Figure 8: PL Intensity as a Function of Exciter Power for Ga₂O₃/BGaAs/GaAs Heterostructure

In order to deeply understand the recombination mechanisms of the Ga₂O₃/BGaAs/GaAs heterostructure we have performed a study of the PL intensity as a function of temperature. From this study, we observe that the PL intensity increases with temperature and then decreases. Moreover, we notice that the full width at half

maximum is also increasing with temperature. However, for low temperatures we have an energy shift towards the blue and a shift towards the red of the energy peak from the temperature 50 K (see Figure 9).

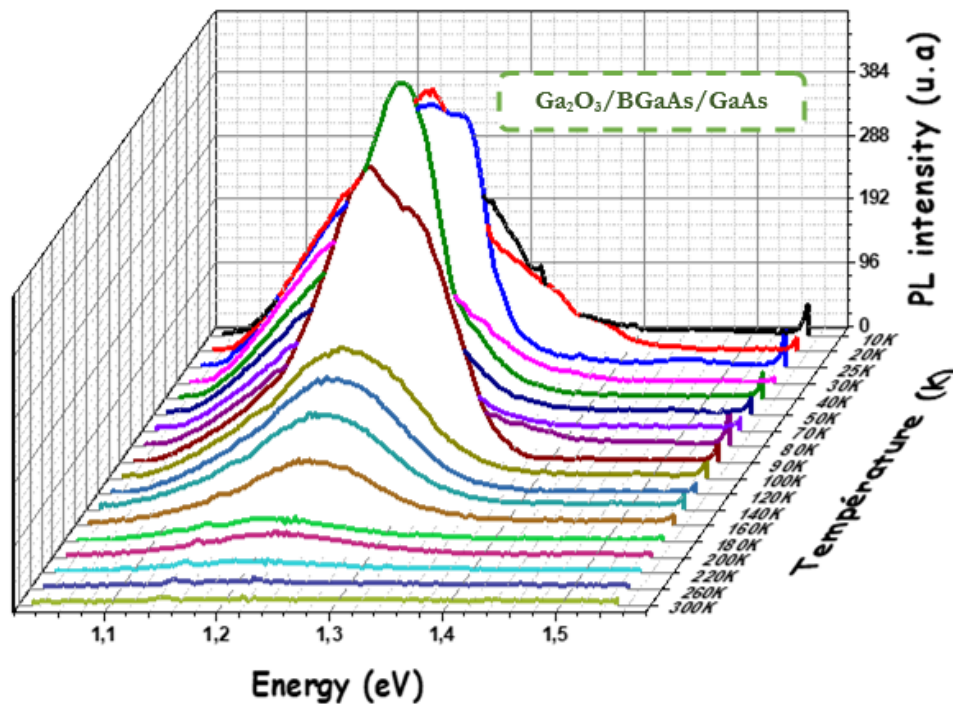


Figure 9: PL Spectra of the Ga₂O₃/BGaAs/GaAs Sample at Different Temperatures

For the Ga₂O₃/BGaAs/GaAs heterostructure, the peak is very broad and we could not follow the evolution of energy, intensity and full width at half maximum as a function of temperature. Whereas for the BGaAs/GaAs epilayer, Figure 10 shows the temperature dependence of the PL band energy which gives a good fit according to Varshni's laws (Equation (1)). We have a good correlation between the results of the experiment and the Varshni adjustment. Moreover, the difference ΔE (10 K) between the measured PL peak energies and the expected transition energies at 10 K (obtained by simulating the measured temperature dependence of the PL peak

positions by the well-known Varshni relation) are suggested to be related to Stokes shifts, is about 5 meV, may be due to the recombination of photogenerated carriers trapped by localized states in BGaAs [23,30]. Indeed, it exhibits anomalous behavior, the so-called S-shape between 10 and 90 K. This shape is the result of carriers jumping between localized states created by boron clusters [31]. At high temperature, the PL band is similar to the variation of the energy band gap. This indicates that the carriers acquire more thermal energy to reach the conduction band. They relax giving rise to band-to-band recombination [25-31].

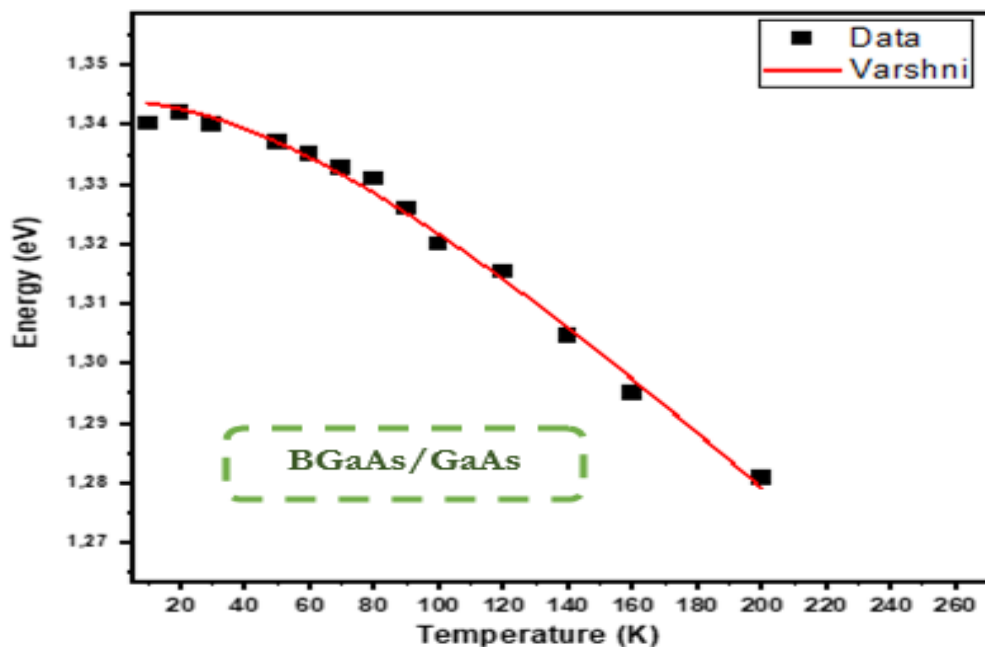


Figure 10: The Evolution of PL Energy as a Function of Temperature

3. Conclusion

In summary, the study of optical properties by steady-state photoluminescence of the active layer shows that there is an emission and red shift of about 1.2 eV after low temperature deposition of Ga₂O₃ on BGaAs/GaAs. However, performance studies highlight the dominance of the Ga₂O₃ layer in the emission process of the donor-acceptor transition. Boron segregates in the direction of the Ga₂O₃ layer, resulting in a reduction in boron content and a reduction in tensile stress, resulting in a reduction in the layer. The study of PL as a function of temperature on BGaAs/GaAs structures leads us to conclude that the observed luminescence is the origin of exciton recombination, which is associated with discrete states formed by the aggregation of boron atoms under band gap conditions. The luminescence in the new Ga₂O₃/BGaAs/GaAs structure is governed by the localization of charge carriers in the energy fluctuation potential created by the inhomogeneous distribution of the VGa family.

Data Availability Statements

We confirm that all data presented in the paper are original and

realized in our laboratory by the authors.

Authors Contribution Statement

We are pleased to submit an original research article entitled "Optical properties of Ga₂O₃ deposited on (B)GaAs/GaAs heterostructures for optoelectronic application". The results presented in this paper were carried out in our lab by the authors: Dalanda Slimi, Faouzi Saidi, and Hassen Maaref, submitted for possible publication in this journal. We confirm that all authors contributed equally to the paper.

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