

IMPROVING A BASED SEMICONDUCTOR III-V LASER STRUCTURE EMITTING AT A WAVELENGTH OF 400 nm

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ABSTRACT

This work focuses on the modeling of a ternary structure based on III-V semiconductor with strained quantum well, InGa_xN/GaN. We studied the effect of indium concentration (x) on different parameters of the In_xGa_{1-x}N/GaN material. Indeed, the incorporation of indium induces a variation of the bandgap energy. Increasing the indium concentration x decreases the bandgap, an important property to obtain the wavelength in the visible region. We also studied the strain effect on the band structure. We have considered the effect of the active zone thickness on the emission wavelength taking into account the effect of the strain $\epsilon(x)$ and the indium content x. Then, we modeled the optical gain as a function of the wavelength by varying the well width L_w , the injected electron-carrier density N and x at T = 300K. Finally, we optimized the structure to make a Laser component emitting around 400nm.

Keywords: Semiconductors III-V, Nanostructures, Laser, Optoelectronics.

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1. INTRODUCTION

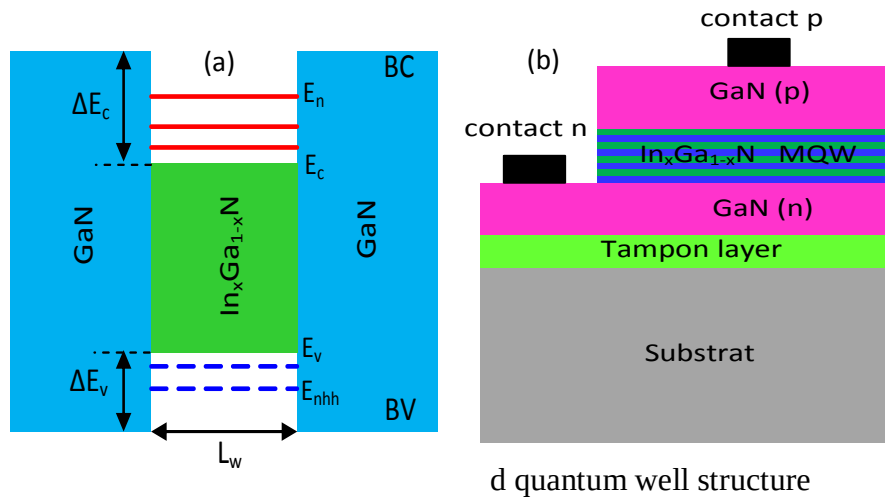
Currently, III-V semiconductors are frequently mentioned for their importance as part of advanced and ultra-fast optoelectronic components. The operational characteristics of these devices depend predominantly on the physical properties of the constituent materials, which

are often combined in quantum heterostructures based on carrier confinement to nanoscale dimensions such as quantum wells, nonwires, and quantum dots. Most of these materials are obtained by standard alloy on substrates. They could in principle cover a wide range of electronic, optical and optoelectronic applications. Most importantly, III-V semiconductor lasers that have played a crucial role in the development of emitters and detectors [1,2]. Semiconductor lasers are lasers based on III-V semiconductor gain media. In these types of lasers, the conduction band (BC) and valence band (BV) electronic structure is used to increase the radiative interband and intraband recombinations. However, the accessible spectral ranges depend on the choice of semiconductors [3]. The InGaN/GaN materials are of great potential interest for light emitting devices operating in the blue (0.4 μ m)–ultraviolet spectral range [4]. In addition, a basic technique for an extremely wide range of applications, harmonic generation or frequency conversion (doubling, tripling, etc.) allows for new features are often used with semiconductor lasers; and are of great potential interest for light emitting devices operating in the visible and the blue–ultraviolet spectral range [5,6]. Since the first experimental demonstration of the laser effect [7], the development of new laser sources, based on a variety of technologies and new material systems, operating in the spectral regions from deep UV to Terahertz has made a great progress [7]. It distinguishes some semiconductor materials available for the manufacture of lasers. Generally, the alloys of semiconductor materials are type III-V, II-VI or IV-VI. These semiconductors are matched lattice layer ($\epsilon=0$) and have a direct band gap, examples of such devices are the VCSELs structures, each interface allows to create a very high reflectivity Bragg mirrors exceeding 99% and a matched active zone on the same substrate which gives a very high crystalline proper. The improvement of the exploits of the optoelectronic components was the consequence of the appearance of the strained quantum wells which contributed to the design of new optoelectronic devices more efficient and more reliable by studying the strain effects on the electronic and optical properties of these III-V semiconductors [8]. The strain has significant impact on the characteristics of intrinsic semiconductors such as bandgap energy, deformation potentials as well as band structure parameters such as effective masses, which are due to the lattice mismatched epitaxial layers. These changes have been exploited to make more suitable components. The strain-degeneration, for example, in the valence band makes it possible to realize optical amplifiers insensitive to the polarization of light and several other devices [9].

1. THEORETICAL MODEL

The quantum wells are thin layered semiconductor structures in which one semiconductor well material is surrounded by two "barrier" layers with a wider band gap ($E_{g1} < E_{g2}$). Since, the well is so thin, typically about 100 Å, both electrons and holes are confined. The energy states allowed for the carriers are quantified. The direct advantage of this configuration is to reduce the active volume, which makes it easier to reach a population inversion, the threshold for oscillation and increase the probability of transitions.

The quantum physic equations are valid when the active layer thickness is of the order of few nanometers and becomes comparable to the De Broglie wavelength $\lambda_F = 2\pi\hbar(2mE_F)^{-1/2}$, m is the electron effective mass, E_F refers to the Fermi level [10,11]. In the type I quantum well, the electrons and holes are confined within the same spatial region as shown in figure 1(a,b). [12].



By varying the well width and the alloy composition, the quantification energy levels can be tuned to modify the emission wavelength of the structure. Indeed, the shift between two subbands of the same quantum number n can be approximated by the equation given in reference [13]. The carrier confinement energies (electrons and holes) in quantum wells are obtained when solving the corresponding Schrödinger equation [14]. The resolution of the Schrödinger equation requires the knowledge of the structural potential $V(z)$ seen by the particle along the direction of interest z [15]. The solution of Schrödinger equation gives an improved estimate of the energy eigenvalue with variable E_n the quantification energy. In order to determine the quantization energies, the numerical method of dichotomy is used. When the Electron confinement energies are not negligible in front of the height of the

potential barrier, either because of ΔE_c is not very important or L_w is small. By asking $V_0 = \Delta E_c$, the potential may be written as mentioned in [16]. Schrödinger's equation in the three regions is given by [17]. The epitaxial growth techniques of lattice mismatched materials became well controlled. Thus, the lattice matching is not an imperative; moreover the strain effects are exploited in many high performance electronic and optoelectronic components. Indeed, the use of strained heterostructures changes the energy position of both conduction band and valence band and gives us the possibility to adjust the emission wavelength [17]. In the pseudomorphic growth along the (001) direction, the difference of lattice constants for the quantum well and barrier layers, results in a parallel ϵ_{xx} and a perpendicular strain ϵ_z [17]. The energy offset of gravity centers of the conduction band and the valence band in $k=0$ vary proportionally with the strain [18]. The energy offset induced by the shear strain for the valence band gap [18-20]. The material gain per unit propagation length $G(h\omega)$, is defined as the fractional increase of the photon-flux density propagating along the direction of interest in the crystal [7]. The gain is calculated using [21,22].

2. RESULTS AND DISCUSSION

Figure 2 shows the variation of the unstrained bandgap energy as a function of the indium concentration. The bandgap is equal to 3.05 eV for an indium concentration of $x=0.10$. When it reaches the value of 0.3, the bandgap energy is 2.6 eV. So, we note a decrease in the bandgap energy with about $\Delta E_g = 0.45$ eV. The effect of the strain on the bandgap energy is also distinguished. The strain induced a decrease in the bandgap energy and makes the valence bands ΔE_{ghh} strongly anisotropic. For an indium value of $x = 0.30$, the strain bandgap energy is equal to 2.43 eV with a deformation of $\epsilon = 3\%$. So, the strain induces a decrease of the bandgap of about $\Delta E_{ghh} = 0.17$ eV. As seen from figure 2, when the indium concentration becomes important $x=0.40$, we will have a compressive strain $\epsilon=4\%$, which makes the structure unstable. The results of simulations are very according to the experimental one.

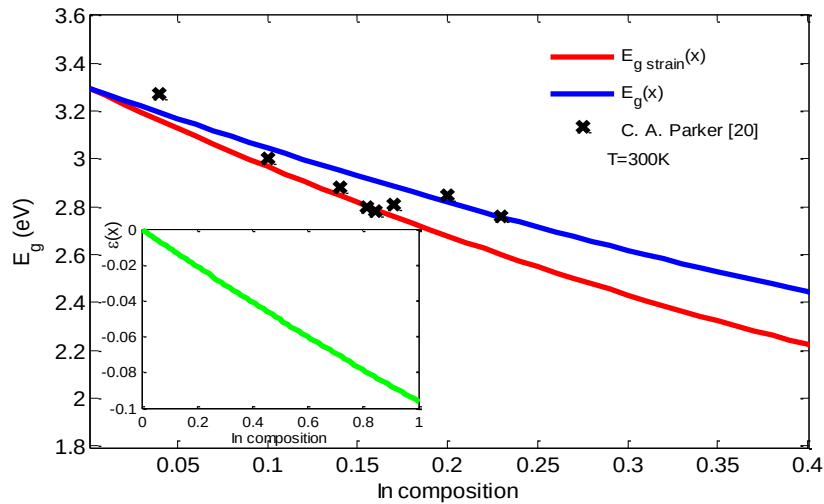


Fig.2. Variation of the bandgap energy as a function of the indium concentration.

Figures 3 (a,b) illustrate the variation of electron confinement energies for the $\text{In}_x\text{Ga}_{1-x}\text{N}/\text{GaN}$ quantum well structure as a function of the well width (L_w) for different indium concentrations. It is noted that the increase in the well width L_w induces a decrease in the quantized energy. On the other hand, the increase of the indium concentration x increases the number of the confinement energy levels. This leads to an increase of the transition probability of the direct electron-hole transitions. Figures 4 (a,b) present the heavy hole confinement energies of the $\text{In}_x\text{Ga}_{1-x}\text{N}/\text{GaN}$ quantum well structure as a function of the well width for different indium contents. The heavy hole confinement energies increase with reducing the well width. Figures 5 (a, b) show the calculated transition energies occurred between the conduction band and the heavy holes band in the $\text{In}_x\text{Ga}_{1-x}\text{N}/\text{GaN}$ quantum well structure versus the well width for different concentrations of indium. It is observed that the increase of the well width, leads to a decrease of the transition energy.

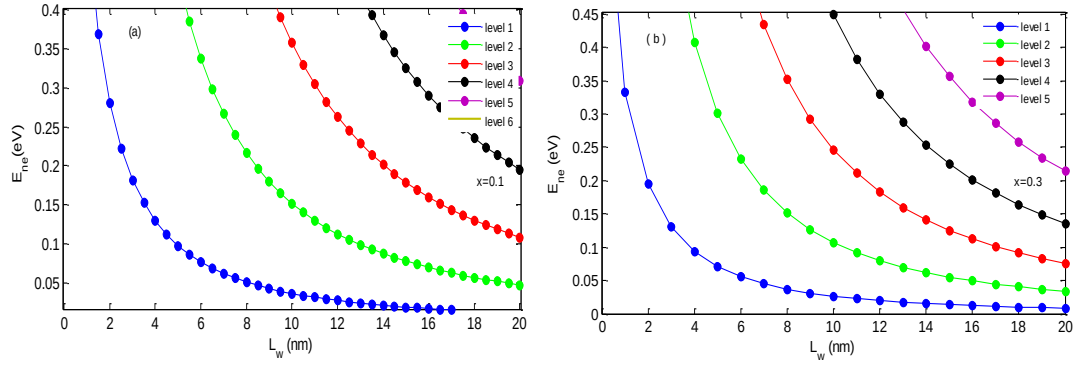


Fig.3. Variation of electron confinement energies levels according to the well width L_w a) $x=0.10$ b) $x=0.30$

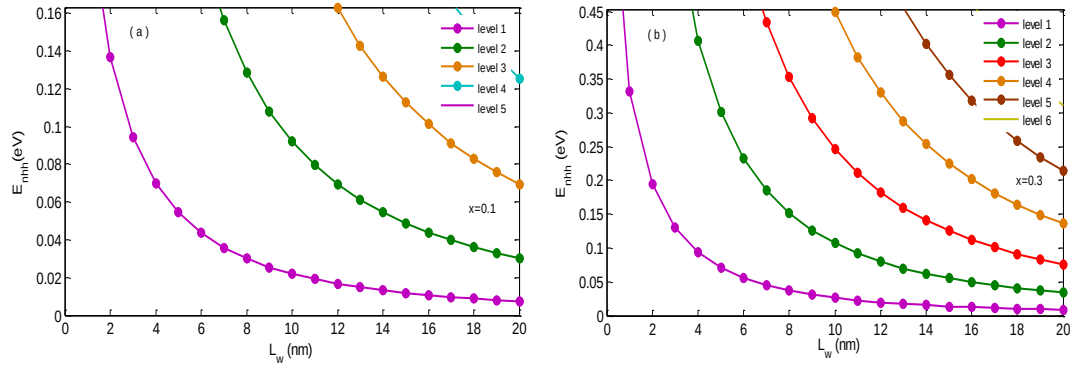


Fig.4. Variation of heavy holes confinement energies levels according to the well width L_w a) $x=0.10$ b) $x=0.30$

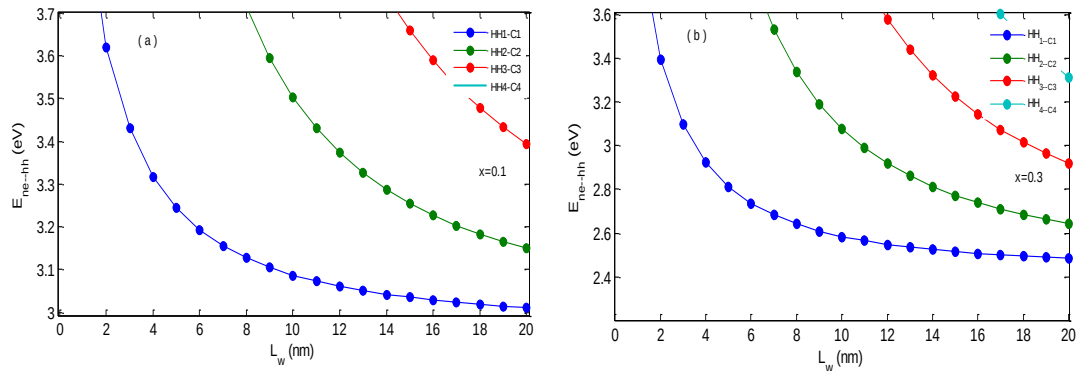


Fig.5. Variation of transition energy according to the well width L_w a) $x=0.10$ b) $x=0.30$

Figures 6 (a, b) illustrate the variation of emission wavelengths as a function of the quantum well width L_w for different indium mole fractions. It is noted that increasing the well width induces an increase of the wavelength. When the indium concentration increases, the emission wavelength varies from 0.1 to 0.5 μm . For example, for a thickness well of 8 nm and an indium concentration $x = 0.10$, the emission wavelength λ_{e1-hh1} is 0.396 μm . For the same transition e_1-hh_1 and an indium content of $x=0.30$, the emission wavelength becomes 0.469 μm .

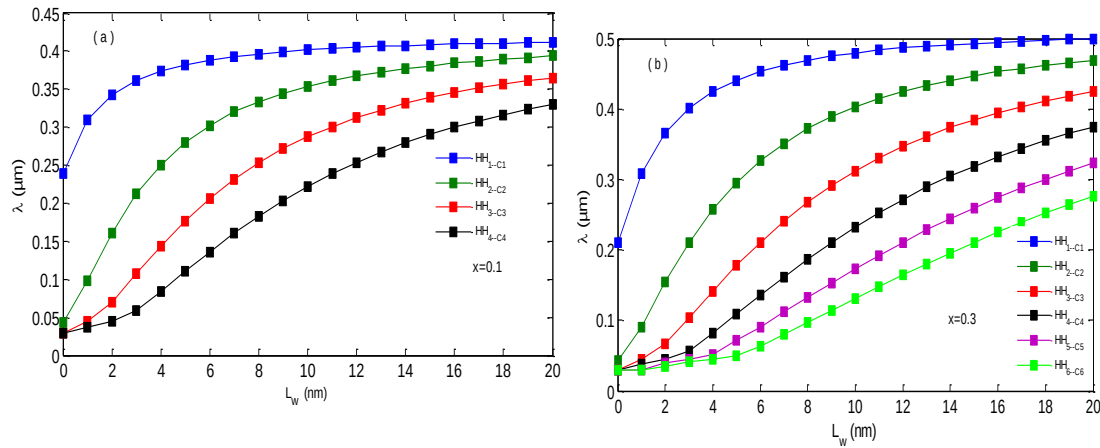


Fig.6. Variation of the wavelength according to well width L_w

a) $x=0.10$ b) $x=0.30$

Figure 7 shows the optical gain variation as a function of the emission wavelengths for a 6nm $\text{In}_{0.3}\text{Ga}_{0.7}\text{N}/\text{GaN}$ quantum well structure and different carrier densities N . It can be seen that the optical gain increases with increasing carrier density to achieve the saturation for $N=8 \times 10^{18} \text{cm}^{-3}$. It can be seen that more the carrier injection becomes larger, more the spectral width $\Delta\lambda$ increases significantly. Figure 8 shows the evolution of optical gain versus the wavelength. An increase of the well width leads to a decrease of the maximum optical gain. However, the spectral width becomes large and the emission wavelength achieves the value of 0.5 μm . For example, if the thickness of the quantum well is varied from 6 to 10 nm, the maximum optical gain decreases through 976 cm^{-1} .

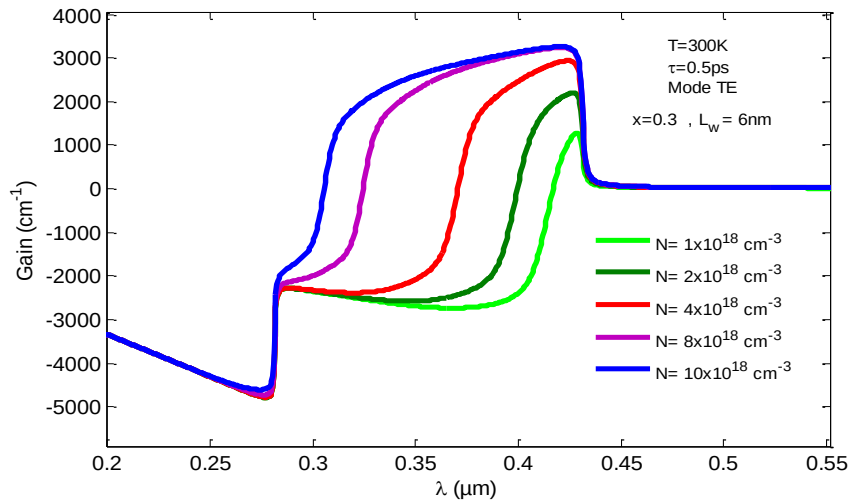


Fig.7. Variation of optical gain versus wavelength for different electron densities

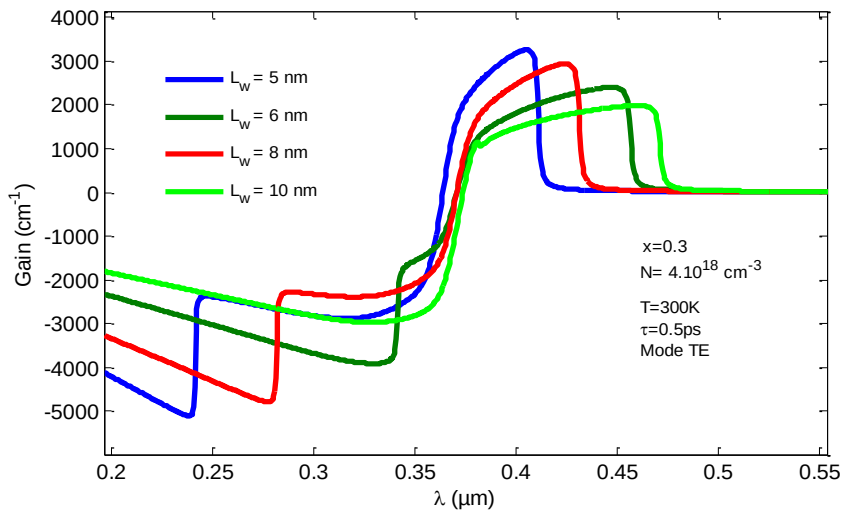


Fig.8. Variation of optical gain versus wavelength for different widths L_w

Figure 9 shows the optical gain variation as a function of the emission wavelength for several indium concentrations in the $\text{In}_x\text{Ga}_{1-x}\text{N}/\text{GaN}$ active zone with an injected carrier density $N=4.10^{18} \text{ cm}^{-3}$ at room temperature and a well width of 6 nm. It is noted that the increase in the indium concentration induces a shift of the gain spectra towards long wavelengths. For an increasing of the indium concentration from 0.10 to 0.30, the emission wavelength shifts from 0.365 to 0.425 μm .

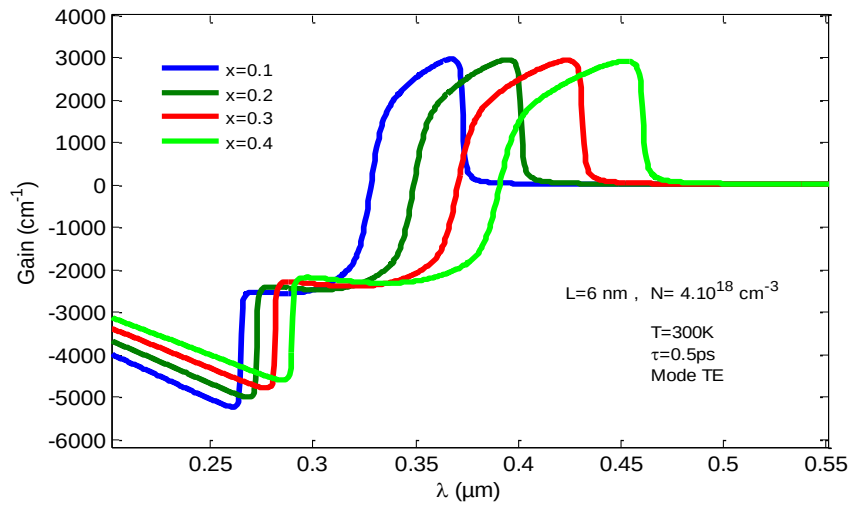


Fig.9. Variation of optical gain versus wavelength for different concentrations x

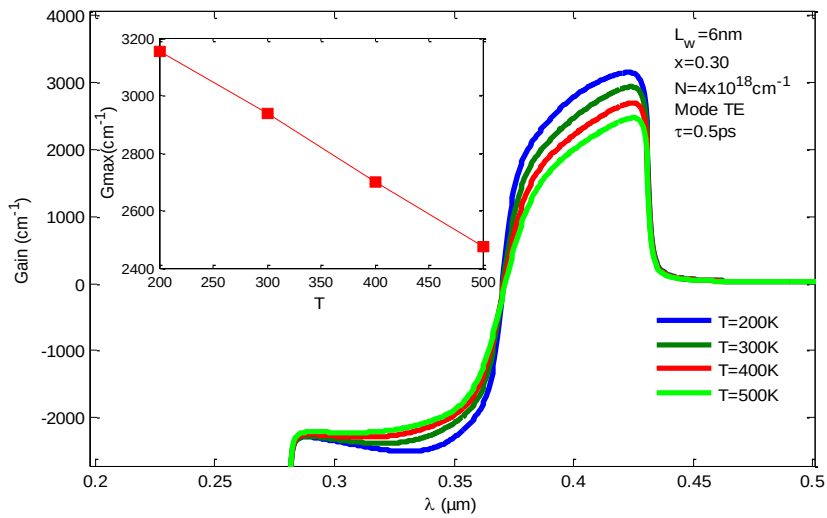


Fig.10. Variation of optical gain versus wavelength for different temperatures T

Figure 10 shows the optical gain variation as a function of the wavelength. It is noted that the increase in temperature leads to a significant decrease of the maximum optical gain. When the temperature increases from 300 to 400K, the maximum gain decrease from 2936 to 2698 cm^{-1} , that is to say, we have a loss of $G_{\text{max}} = 238 \text{ cm}^{-1}$. From this study, we can optimize the quantum well structure based on $\text{In}_x\text{Ga}_{1-x}\text{N}/\text{GaN}$ to achieve a reliable optoelectronic component emitting around 400nm. It suffices to find a compromise between the well width, the indium mole fraction, the carrier density and the temperature.

4. CONCLUSION

In this paper, a theoretical study has been carried out. Simulations have been performed for the design and optimization of an InGaN/GaN strained quantum well structures. In our study, the effects of $\varepsilon(x)$ and x on the bandgap energy, the quantum confinement energies, the transition energy and the wavelength emission are taken into account. Then, we have studied the material gain a based InGaN/GaN quantum well material system and present the factors that influence the material gain performance as well width, electron density, x concentration and temperature T . The optimized InGaN/GaN quantum well structures have better optical properties with a maximum optical gain of 2936 cm^{-1} and an emitting wavelength of 423 nm, at a well width of 6 nm, injected electron density $N=4 \times 10^{18} \text{ cm}^{-3}$, $x=0.30$, and a room temperature $T=300\text{K}$. This work allows us the possibility to realize a strained multiple quantum well VCSEL structure emitting in the blue region of the electromagnetic spectrum.

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