Electrical and optical properties of p-type InGaN

B. N. Pantha, A. Sedhain, J. Li, J. Y. Lin, and H. X. Jiang^{a)}

Department of Electrical and Computer Engineering, Texas Tech University, Lubbock, Texas 79409, USA

(Received 14 November 2009; accepted 4 December 2009; published online 29 December 2009)

Mg-doped $In_xGa_{1-x}N$ alloys were grown by metal organic chemical vapor deposition on semi-insulating c-GaN/sapphire templates. Hall effect measurements showed that Mg-doped $In_xGa_{1-x}N$ epilayers are p-type for x up to 0.35. Mg-acceptor levels (E_A) as a function of x, (x up to 0.35), were experimentally evaluated from the temperature dependent hole concentration. The observed E_A in Mg-doped $In_{0.35}Ga_{0.65}N$ alloys was about 43 meV, which is roughly four times smaller than that in Mg doped GaN. A room temperature resistivity as low as 0.4 Ω cm (with a hole concentration $\sim 5 \times 10^{18}$ cm⁻³ and hole mobility ~ 3 cm²/V s) was obtained in Mg-doped $In_{0.22}Ga_{0.78}N$. It was observed that the photoluminescence (PL) intensity associated with the Mg related emission line decreases exponentially with x. The Mg energy levels in InGaN alloys obtained from PL measurements are consistent with those obtained from Hall-effect measurements. © 2009 American Institute of Physics. [doi:10.1063/1.3279149]

Achieving highly conductive p-type GaN and AlGaN is very difficult due to the high activation energies (E_A) of the Mg-acceptor, which is a universally accepted p-type dopant for GaN and related alloys.^{1–3} Since E_A decreases with a decrease in band gap energy,³ Mg-doped InGaN (InGaN:Mg) is expected to have a higher hole concentration (p) than Mgdoped GaN. Additionally, p-type InGaN is synthesized at a much lower temperature compared to p-type GaN. Thus, the use of p-type InGaN instead of p-type GaN in device structures is beneficial, particularly in devices such as green laser diodes, long wavelength emitters, and solar cells, where the active region has to be grown at temperatures much lower than that of the top p-type GaN layer. Low etching damage and low contact resistance are other superior characteristics of p-type InGaN over p-type GaN.¹ Improvements in the performance of GaN based devices using p-type InGaN either as a contact layer or as the p-layer itself has already been demonstrated.⁴⁻⁶ In recent years, applications of InGaN alloys have expanded into areas such as optoelectronics/ photonics, solar cells, photoelectrochemical cells for hydrogen generation, and thermoelectric materials for converting heat to electricity.⁷⁻¹² For all these applications, high optical and electrical quality p-type InGaN is highly sought after. For hydrogen generation, p-type InGaN alloys are expected to be more stable in aqueous solutions than n-type materials.

P-type doping in relatively high In content InGaN alloys is highly challenging due to the presence of high background electron concentrations, which is believed to originate from defects such as oxygen and hydrogen impurities or nitrogen vacancies.^{13,14} Nitrogen vacancies could be the consequence of an insufficiency of nitrogen atoms which results from the low decomposition rate of ammonia, as high In-content In-GaN has to be grown at temperatures lower than <800 °C (while the growth temperature of GaN is generally >1000 °C). Although the synthesis of p-type InGaN has been reported as early as in 1995,¹⁵ there are only a few reports on the properties of p-type InGaN.^{1,16–20} However, the use of high In-content p-InGaN is inevitable in the near future as nitride based devices are rapidly expanding toward longer wavelength emitters.⁷ We report on the synthesis and systematic studies on the electrical and optical properties of Mg doped p-type $In_xGa_{1-x}N$ alloys, for x up to 0.35.

Mg-doped p-type $In_rGa_{1-r}N$ alloys $(0 \le x \le 0.35)$ were grown on semi-insulating c-GaN (SI-GaN)/sapphire templates by metal organic chemical vapor deposition (MOCVD). We have chosen SI-GaN templates to minimize electrical measurement errors of top InGaN:Mg alloys. Since the p-type $In_rGa_{1-r}N$ layer is relatively thin (~200 nm), resistivity of the layer underneath has to be high in order to assure accuracy of the measurement results. Ammonia (NH₃), trimethylgallium (TMGa), trimethylindium (TMIn), and biscyclopentadienyl-magnesium (Cp2Mg), were used as N, Ga, In, and Mg sources. N₂ gas was used as a carrier gas. Growth temperatures were varied from 1050 to 740 °C to increase x from 0 to 0.35. A variable temperature Hall-effect experiment was performed to measure the hole concentration (p), hole mobility $(\mu_{\rm h})$, and resistivity (p) of the samples. Photoluminescence (PL) spectra were measured using a Tisapphire laser spectroscopy system coupled with a tripler. This system gives an average output power of about 40 mW at 4.7 eV and a spectral resolution of about 0.2 meV.²¹ Indium contents were determined from the peak angles of (002) ω -2 θ x-ray diffraction (XRD) curves and using Vegard law.

The room temperature electrical properties of p-type $In_rGa_{1-r}N$:Mg alloys as functions of x are plotted in Fig. 1. It was found that p continuously increases from 2×10^{17} for x=0 (GaN) to 5×10^{18} cm⁻³ when x=0.22. $\mu_{\rm h}$ was found to decrease from 15 to 1.8 cm²/V s as x increases from 0 to 0.35. The variation in ρ with x shows that ρ decreases as x increases and reaches a minimum value of 0.4 Ω cm at x =0.22 (In_{0.22}Ga_{0.78}N). This value of ρ is among the lowest reported for p-type InGaN. Though an even higher ρ value has been reported for similar In-content InGaN:Mg¹⁸ and GaN:Mg grown by molecular beam epitaxy,²² μ_h values of our samples are higher, which results in lower values of ρ . Higher $\mu_{\rm h}$ values are a result of enhanced material quality, while reduced ρ values will significantly improve device performance for many practical applications. The reduction in pvalues observed in p-type $In_xGa_{1-x}N$ with x > 0.22 are due to

0003-6951/2009/95(26)/261904/3/\$25.00

95, 261904-1

Downloaded 12 Jul 2010 to 129.118.86.45. Redistribution subject to AIP license or copyright; see http://apl.aip.org/apl/copyright.jsp

^{a)}Electronic mail: hx.jiang@ttu.edu.



FIG. 1. (Color online) Room temperature (a) resistivity ρ , (b) hole mobility μ_h , and (c) hole concentration p as functions of In content (x) in Mg doped In_xGa_{1-x}N alloys.

the effect of hole compensation by background electrons. The presence of high background electron concentration is the main hindrance for obtaining p-type conductivity and p-type $In_xGa_{1-x}N$ alloys with In content x > 0.35.

Temperature dependent p of p-type $In_xGa_{1-x}N$ alloys was also measured. Figure 2 shows the Arrhenius plot of free hole concentration p for x=0.05, 0.15, 0.22, and 0.35. Straight lines are linear fits of the experimental data of p by the following equation:

$$p = p_o e^{-E_{\rm A}/k_{\rm B}T},\tag{1}$$

where E_A is Mg energy level and k_B is the Boltzmann constant. The plot of E_A as a function of x for $In_xGa_{1-x}N:Mg$ alloys is shown in Fig. 3 along with other reported values.^{1,17,19} It was found that E_A continuously decreases with an increase in x. Lower values of E_A are the main physical reason for higher values of p in $In_xGa_{1-x}N:Mg$ alloys of higher x. An E_A value as low as 43 meV was measured in $In_{0.35}Ga_{0.65}N:Mg$. This value is approximately four times



FIG. 3. (Color online) Energy level (E_A) of Mg acceptors in Mg doped p-In_xGa_{1-x}N alloys as a function of In content *x*. Data from references available up to x=0.17 are also included.

smaller than that of Mg-doped GaN. $In_xGa_{1-x}N$ materials with x > 0.35 are generally highly n-type and conversion of these materials to p-type by Mg doping is still very difficult. Our results indicate that p-type conductivity in InGaN:Mg could be further improved if a better control of the background electron concentration could be achieved. The issues of high background electron concentration in InGaN alloys with relatively high In content is currently under intensive investigation^{11,12} and a significant improvement in the p-type conductivity of relatively high In content alloys is anticipated.

Figure 4 shows low temperature (10 K) PL spectra of $In_xGa_{1-x}N:Mg$ for x=0, 0.05, 0.11, and 0.22. We observed that the spectra for GaN:Mg is dominated by a broad emission band centered around 2.90 eV, which is a donor-acceptor-pair type transition involving a deep donor, D^o, and an Mg-acceptor, Mg^o. Deep donors appeared in Mg doped GaN but disappeared in all InGaN:Mg alloys. One speculation is that the lower growth temperatures employed for In-GaN alloys somehow suppresses the formation of these deep donors. A relatively weak feature in the higher energy side



FIG. 2. (Color online) The Arrhenius plot of free hole concentration in Mg doped $In_rGa_{1-x}N$ alloys.



FIG. 4. (Color online) PL spectra of Mg-doped p-type $In_xGa_{1-x}N$ alloys measured at temperature 10 K.

Downloaded 12 Jul 2010 to 129.118.86.45. Redistribution subject to AIP license or copyright; see http://apl.aip.org/apl/copyright.jsp



FIG. 5. (Color online) Mg acceptor energy levels E_A , peak position of Mg related emission E_p , and integrated PL intensity I_{int} as functions of In content *x*. (a) E_A , (b) E_p , and (c) I_{int} .

(with a peak at 3.29 eV) is a band-to-impurity type transition involving the conduction band (e⁻) and acceptor (Mg^o) which is believed to dominate the spectra of GaN with light or moderate Mg doping. All p-type $In_rGa_{1-r}N:Mg$ spectra except the GaN:Mg sample show the dominant peak with the same mechanism as that of the 3.29 eV line in GaN:Mg \Box namely, a band-to-impurity transition involving the conduction band (e⁻) and acceptor (Mg^o). An emission line at 3.29 eV also appeared in the spectra for samples with x=0.11 and 0.22. This emission line is related to the SI-GaN template, which was lightly doped with Mg to achieve semi-insulation. From the PL peak positions observed here and the band-gap values already reported in the literature,¹¹ we have estimated the acceptor level of Mg in our $In_xGa_{1-x}N$ epilayers, which is plotted in Fig. 5. As expected, Mg levels in $In_rGa_{1-r}N$ alloys decrease with In-content and are consistent with those obtained from the electrical measurement results shown in Fig. 3. The Mg impurity related PL emission intensity is found to decrease exponentially with In-content, as shown in Fig. 5. The PL emission intensity of Mg doped In_{0.22}Ga_{0.78}N is almost three orders of magnitude lower than that of Mg doped GaN. The reduction in PL intensity may be related with the incorporation of impurities, which are also responsible for the high background electron concentrations in high In content InGaN alloys.

In summary, we have synthesized Mg-doped p-type $In_xGa_{1-x}N$ alloys by MOCVD for *x* up to 0.35 and analyzed their electrical and optical properties. P-type resistivity in Mg-doped InGaN alloys was found to be lower than that of Mg-doped GaN. Resistivity as low as 0.4 Ω cm with a free hole concentration as high as 5×10^{18} cm⁻³ was measured in

Mg doped $In_{0.22}Ga_{0.78}N$. We have measured the Mg acceptor energy levels (E_A) in p- $In_xGa_{1-x}N$ alloys for x up to 0.35. E_A as low as 43 meV was obtained in Mg doped $In_{0.35}Ga_{0.65}N$. From low temperature PL measurements, we found Mgacceptor levels for x up to 0.22 and the results are in close agreement with those obtained from electrical measurements. The difficulties of obtaining p-type InGaN in relatively high In-content are due to the high background election concentrations caused by the lower growth temperature of InGaN alloys, which seems to promote the incorporation of donor like defects and impurities.

InGaN growth and optical characterization work is supported by NSF (Grant No. DMR0906879). The effort of achieving p-type conduction in InGaN is supported by DARPA and monitored by Dr. Michael Haney and Dr. John Zavada. H.X.J. and J.Y.L. would like to acknowledge the support of Edward Whitacre and Linda Whitacre endowed chairs through the AT&T foundation.

- ¹K. Kumakura, T. Makimoto, and N. Kobayashi, J. Appl. Phys. **93**, 3370 (2003).
- ²W. Götz, N. M. Johnson, J. Walker, D. P. Bour, and R. A. Street, Appl. Phys. Lett. **68**, 667 (1996).
- ³J. Li, T. N. Oder, M. L. Nakarmi, J. Y. Lin, and H. X. Jiang, Appl. Phys. Lett. **80**, 1210 (2002).
- ⁴T. Makimoto, K. Kumakura, and N. Kobayashi, Appl. Phys. Lett. **79**, 380 (2001).
- ⁵K. Kumakura, T. Makimoto, and N. Kobayashi, Appl. Phys. Lett. **79**, 2588 (2001).
- ⁶P. C. Chen, C. H. Chen, S. J. Chang, Y. K. Su, P. C. Chang, and B. R. Huang, Thin Solid Films **498**, 113 (2006).
- ⁷J. Wu, J. Appl. Phys. **106**, 011101 (2009).
- ⁸R. Dahal, B. Pantha, J. Li, J. Y. Lin, and H. X. Jiang, Appl. Phys. Lett. **94**, 063505 (2009).
- ⁹K. Fujii and K. Ohkawa, Jpn. J. Appl. Phys. 44, L909 (2005).
- ¹⁰J. Li, J. Y. Lin, and H. X. Jiang, Appl. Phys. Lett. **93**, 162107 (2008).
- ¹¹B. N. Pantha, R. Dahal, J. Li, J. Y. Lin, H. X. Jiang, and G. Pomrenke, Appl. Phys. Lett. **92**, 042112 (2008); J. Electron. Materials **38**, 1132 (2009).
- ¹²N. Kobayashi, T. Narumi, and R. Morita, Jpn. J. Appl. Phys., Part 2 44, L784 (2005).
- ¹³C. G. Van de Walle and D. Segev, J. Appl. Phys. **101**, 081704 (2007).
- ¹⁴J. W. L. Yim, R. E. Jones, K. M. Yu, J. W. Ager III, W. Walukiewicz, W. J. Schaff, and J. Wu, Phys. Rev. B 76, 041303(R) (2007).
- ¹⁵S. Yamasaki, S. Asami, N. Shibata, M. Koike, K. Manabe, T. Tanaka, H. Amano, and I. Akasaki, Appl. Phys. Lett. 66, 1112 (1995).
- ¹⁶K. Kumakura, T. Makimoto, and N. Kobayashi, Jpn. J. Appl. Phys., Part 2 39, L337 (2000).
- ¹⁷K. Kumakura, T. Makimoto, and N. Kobayashi, J. Cryst. Growth 221, 267 (2000).
- ¹⁸T. C. Wen, W. I. Lee, J. K. Sheu, and G. C. Chi, Solid-State Electron. 45, 427 (2001).
- ¹⁹D. Iida, M. Iwaya, S. Kamiyama, H. Amano, and I. Akasaki, Appl. Phys. Lett. **93**, 182108 (2008).
- ²⁰C. A. Chang, T. Y. Tang, P. H. Chang, N. C. Chen, and C. T. Liang, Jpn. J. Appl. Phys., Part 1 46, 2840 (2007).
- ²¹Research page, nanophotonics research at TTU (http://www2.ece.ttu.edu/ nanophotonics/).
- ²²G. Namkoong, E. Trybus, K. K. Lee, M. Moseley, W. A. Doolittle, and D. C. Look, Appl. Phys. Lett. **93**, 172112 (2008).