Formation of ohmic contacts to MOCVD grown p-GaN by controlled activation of Mg

E. Kamińska^{*1}, A. Piotrowska¹, A. Barcz^{1,2}, D. Bour³, M. Zielinski²,
J. Jasinski^{4,5}

¹Institute of Electron Technology, Warsaw, Poland

²Institute of Physics, PAS, Warsaw, Poland

³Agilent Laboratories, Palo Alto, USA

⁴Lawrence Berkeley Laboratory, Berkeley, CA, USA

⁵Institute of Experimental Physics, Warsaw University, Poland

Abstract

We report on the formation of low resistivity ohmic contacts to p-GaN, $r_c < 10^{-4} \Omega \text{cm}^2$, by increasing the concentration of the active Mg in the subcontact zone, via Zr-mediated release of hydrogen.

Keywords: ohmic contact, p-GaN, hydrogen, ZrN, ZrB₂.

1. Introduction

The development of procedures for the fabrication of ohmic contacts to p type GaN is one of the most stimulating goals in the technology of GaN-based devices. Recently, evidence was given that the subsurface region of MOCVD-grown p-GaN contains a substantial amount of hydrogen, which due to the passivation of magnesium, seems to be responsible for the difficulties in making ohmic contacts. To resolve this problem we have used Zr-based metallization and demonstrated its usefulness as a drain enabling the evolution of hydrogen from the semiconductor subcontact region under heat treatment [1]. This work presents a detailed study of the process of zirconium-mediated removal of hydrogen. We report the results of experiments designed to analyse the effect of temperature and gas ambient on the concentration profiles of hydrogen in the subcontact region of p-GaN as well as on the electrical properties and the microstructure of contacts.

^{*}Corresponding author: tel: +48 2254 87942; fax: +48 2284 70631; e-mail: eliana@ite.waw.pl

2. Experimental procedure

The GaN samples used in this study were grown by MOCVD on sapphire substrate with an undoped GaN buffer layer, followed by 1.5 μm thick p-GaN doped with Mg. The experiments were performed on both as grown and post growth-activated material. After the activation of Mg, the hole concentration was $1 \times 10^{18} \text{ cm}^{-3}$. Zr-based metallization, consisting of a ZrN(100nm)/ZrB₂(100nm) bilayer was deposited by sequential DC magnetron sputtering from ZrN and ZrB₂ targets, respectively. Heat treatments were conducted in a RTP system at 750-1150°C, for 30 s. in N₂ flow. Compositional depth profiling was performed with secondary ion mass spectrometry (SIMS) using a Cameca 6F instrument, with either cesium or oxygen as primary beam. The microstructure of ZrN/ZrB₂ metallization on GaN was investigated by cross-sectional transmission electron microscopy (XTEM) and selected area diffraction (SAED) analysis in JEOL 2000EX microscope. Contact resistivity was evaluated by circular TLM method.

3. Results and discussion

In order to understand the behaviour of hydrogen in p-GaN under annealing, the first question to be addressed concerns the effect of the post growth annealing, routinely performed to activate Mg as acceptor dopant, on hydrogen concentration. Our results show that, despite of a significant change in the free barrier concentration, the annealing of bare p-GaN at 800°C does not affect noticeably the amount and the distribution of hydrogen. The level of H remains stable upon annealing up to 800°C of ZrN/ZrB₂-capped semiconductor as well. Such a processing in N₂ ambient, however, enables the desorption of hydrogen trapped during the sputter-deposition of ZrB₂ and ZrN films, which was found to enhance the thermal stability of the contact. Fig.1 shows depth profiles of H and Mg in p-GaN after annealing at temperatures 800°C-1100°C in N₂, and at 800°C in O₂ ambient.

Fig.1. Depth profiles in p-GaN annealed with ZrN/ZrB₂ metallization of a) hydrogen and b) oxygen.

Concentrations of H and Mg in GaN were evaluated using relative sensitivity factors from ref. [2].

Both Mg and H are present throughout the thickness of the p-GaN film. The level of Mg is uniform and remains stable under annealing. In contrast, a substantial accumulation of hydrogen in the subsurface region is clearly seen. The process of hydrogen removal depends on the gas ambient during heat treatment. In N_2 atmosphere, it commences at 900 - 950°C. After annealing at ~1050°C almost all hydrogen from the superficial region and from the bulk of the p-type layer is released. Thermal processing in O_2 atmosphere enables to create a hydrogen-depleted subcontact zone at 800°C.

The release of hydrogen from p-GaN has a direct impact on the electrical properties of p-GaN/ZrN/ZrB₂ contacts. As for contacts processed in N_2 ambient, they become low resistive after annealing at 1000°C with $r_c = 6 - 8 \times 10^{-5} \Omega \text{cm}^2$. Contacts heat treated at 800°C in O_2 flow were ohmic as well, and exhibited somewhat higher resistivity $r_c = 1 - 3 \times 10^{-4} \Omega \text{cm}^2$.

As the thermal integrity of the metal/semiconductor system is a key issue in the fabrication of reliable devices, the evolution of p-GaN/ZrN/ZrB₂ contact microstructure under annealing was of primary concern. As-deposited metallization is amorphous. Annealing at 800°C in N_2 causes the recrystallization of both metallic compounds. Fig.2 shows XTEM micrographs and corresponding diffraction patterns of contacts heat treated in N_2 and O_2 atmospheres, at temperatures required to obtain the ohmic behaviour.

Fig.2. XTEM micrographs and corresponding diffraction patterns of p-GaN/ZrN/ZrB₂ contact
a) annealed at 1000°C in N_2 flow, b) annealed at 800°C in O_2 flow.

While the metal/semiconductor interface in both contacts is sharp and abrupt, the microstructure of metallization films considerably differs. The final morphology of the contact annealed in N_2 is polycrystalline. The metallization is composed of ZrN and ZrB_2 layers, as indicated by SAED ring patterns (Fig.2.a). The processing in O_2 significantly alters the contact microstructure. The upper part of metallization becomes amorphous. While, SAED pattern of the lower, polycrystalline part revealed mainly the presence of ZrN (Fig.2.b), SIMS analysis revealed an important intermixing between ZrN and ZrB_2 compounds [3]. To get further insight into the interaction of O_2 with the contact, the profile of oxygen in the metallization and in the semiconductor subcontact region has been measured and compared with contact processed in N_2 . SIMS data evidence an important incorporation of oxygen in the two parts of metallization during annealing. An increase of the oxygen level in the subsurface film of p-GaN has been observed, Fig.1.b. The annealing behaviour of Zr-based metallization in O_2 atmosphere indicates that such a metallization is not oxidation resistant and does not prevent the diffusion of oxygen into the semiconductor. Taking into account that oxygen is a shallow donor in GaN [4], when incorporated into the superficial region of p-GaN it could compensate freshly activated Mg acceptor centres. This could be the cause of higher resistivity of contacts processed in O_2 as compared to N_2 , in spite of a comparable level of hydrogen in both contacts.

4. Conclusions

We have investigated the process of evolution of hydrogen from MOCVD grown p-GaN via Zr-based metallization, and determined the optimum processing conditions (temperature and gas ambient) for fabrication of low resistance ohmic contacts. When the process is conducted in N_2 flow, the metallization remains stable at temperatures required to achieve the ohmic behaviour, and the morphology of the metal/semiconductor interface is unaltered by such a heat treatment. The processing in O_2 , on the contrary, causes the interdiffusion of metallization constituents and the incorporation of oxygen into the semiconductor subcontact region, which could be responsible for increased resistivity of these contacts.

Acknowledgements

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References

- [1] E. Kaminska, A. Piotrowska, A. Barcz, J.Jasinski, M. Zielinski, K.Golaszewska, R.F.Davis, E. Goldys, K. Tomsia, MRS Internet J. Nitride Semicond. Res. 5S1 (2000) W10.9.
- [2] J.W. Erikson, Y. Gao, R.G. Wilson, Mat. Res. Soc. Symp. Proc. 395 (1996) 369.
- [3] J. Jasinski, E.Kaminska, A. Piotrowska, A. Barcz, M. Zielinski, accepted for publication in MRS Internet J. Nitride Semicond. Res.
- [4] C. G. Van de Walle, C. Stampfl, J. Neugebauer, M. D. McCluskey, N.M. Johnson, MRS Internet J. Nitride Semicond. Res. 4S1 (1999) G10.4.