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Carbon doping of non-polar cubic GaN by CBr₄

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ABSTRACT

Non-polar carbon-doped cubic GaN (c-GaN:C) films were grown by the plasma-assisted molecular beam epitaxy (MBE) using carbon tetra-bromide (CBr₄) as a carbon source. The growth was in-situ monitored by the reflection high-energy electron diffraction (RHEED) and for the atomic carbon detection quadrupole mass spectrometry (QMS) was applied. Time-of-flight secondary ion mass spectroscopy (ToF-SIMS) was used to quantify the carbon incorporation behavior. The structural, morphological and optical properties of the epilayers were studied by high-resolution X-ray diffraction (HRXRD), atomic force microscopy (AFM) and photoluminescence (PL) measurements at room temperature and at 4 K. The electrical properties of c-GaN:C samples were determined by measuring the current–voltage (I-V) characteristics at room temperature.

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1. Introduction

Due to the absence of polarization fields in non-polar cubic group III-nitrides phase pure cubic GaN/AlGaN heterostructures may offer the realization of normally-on and normally-off operation of hetero field-effect transistors (HFETs) with the large technological allowance and the same flexible device design ability as the GaAs/AlGaAs HFET technology [1]. For such devices a highly insulating GaN buffer layer has a pivotal role in HFET fabrication since it provides a good electrical isolation from the substrate interface for accurate determination of electrical properties and considerably improves the rf-performance of the devices by suppressing the parasitic capacitance. In polar hexagonal nitrides carbon [2], beryllium [3] and iron [4] have been used to achieve high-quality semi-insulating GaN buffer layers. In this contribution we show that C-doping by carbon tetra-bromide (CBr_4) is suitable to produce highly resistive nonpolar cubic GaN (c-GaN) epilayers.

2. Experiment

c-GaN epilayers were grown by rf-plasma-assisted molecular beam epitaxy (MBE) on conducting free-standing 3C-SiC (001) substrates [5]. Carbon doping via CBr_4 is investigated as a means to produce semi-insulating GaN buffer layers for device applications [6]. The growth was in situ monitored by reflection highenergy electron diffraction (RHEED) [7] and quadrupole mass spectrometry (QMS) was applied to detect the atomic carbon. Time-of-flight secondary ion mass spectrometry (ToF-SIMS) was used to quantify the C incorporation behavior. The C-concentration was calibrated via reference samples, in which a defined amount of C was built in by the ion implantation. The electrical and optical properties were investigated by low-temperature photoluminescence (PL) and current–voltage (*I–V*) characteristics, respectively.

3. Results and discussion

For ToF-SIMS measurements a 500-nm-thick GaN sample containing three different amounts of C-concentrations was prepared on free-standing 3C-SiC substrate. In Fig. 1 the measured depth profiles of C, Br and GaN is depicted. Whereas the GaN concentration is constant all over the c-GaN epilayer thickness, three clear regions with different C-concentrations can be identified in the C-profile. In the inset of Fig. 1 the incorporated C-concentration is plotted vs. the CBr₄ flux. This measurement shows that the amount of incorporated C is linearly related to the incident CBr₄ flux and varies between 2×10^{19} and 6×10^{19} cm⁻³. The Br concentration is at the SIMS detection limit (~ 10^{15} cm⁻³) and indicates that no Br is built in. This behavior is similar to that observed in hexagonal GaN [2].

The reciprocal space map of the (002) reflection of a c-GaN epilayer homogeneously doped with a C-concentration of $\sim 4 \times 10^{19} \, \text{cm}^{-3}$ is shown in Fig. 2. Besides the 3C-SiC and the c-GaN reflection at (0Å⁻¹, 2.83Å⁻¹) and (0Å⁻¹, 2.785Å⁻¹),

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Fig. 1. SIMS depth profiles of GaN, C and Br of a C-doped cubic GaN epilayer which contains three different C-doping levels. The inset shows the incorporated C-concentration measured by SIMS vs. CBr_4 flux.



Fig. 2. Reciprocal space map of the (002) reflection of C-doped cubic GaN/3C-SiC epilayer ([C] \sim 4 × 10¹⁹ cm⁻³).

respectively, no additional (0002) or (1011) reflections of the hexagonal GaN is detectable at (0Å⁻¹, 2.4Å⁻¹) or (-0.4Å⁻¹, 2.5Å⁻¹). Therefore, no degradation of the phase purity and structural properties of the c-GaN epilayers is observed up to a C-concentration of ~10²⁰ cm⁻³. Atomic force microscopic (AFM) measurements further show that also the surface roughness is not deteriorated due to C-doping and a typical rms roughness (5 × 5 µm²) of about 5 nm is measured for both undoped and C-doped samples.

The incorporation of C has a profound influence on the optical properties as studied by the low-temperature PL. In Fig. 3 the spectra of a nominally undoped c-GaN sample (blue curve) and a C-doped c-GaN sample (red curve, $[C] \sim 4 \times 10^{19} \text{ cm}^{-3}$) are



Fig. 3. Low-temperature photoluminescence spectra of undoped (blue) and C-doped (red, $[C]\sim 4 \times 10^{19} \, \text{cm}^{-3}$) cubic GaN/3C-SiC epilayers. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

depicted. In the undoped GaN sample the PL-spectrum is dominated by the exitonic transition X at 3.28 eV and the omnipresent donor acceptor (D⁰A⁰) transition at 3.15 eV [8]. The PL-spectrum of the C-doped sample is dominated by a deep-red luminescence band, which appeared at 2.1 eV. In addition, the integral PL-intensity is severely reduced indicating the compensation effects. The PL peaks assignment is based on previous publications [9-11], where temperature- and intensitydependent PL measurements were performed on c-GaN epilayers deposited on GaAs substrates. In these c-GaN samples grown on GaAs substrates, where an e-beam evaporation source was used to incorporate C, a clear correlation between different PL transitions and the build in C was found [9–11]. It is well known that C can be incorporated at different lattice sites acting either as an acceptor, if it replaces nitrogen (C_N) or acting as a donor if it replaces gallium (C_{Ga}) [12]. Following the discussions in Refs. [10] and [13], we attribute the $(D_C^0 A_C^0)$ transition at 2.85 eV to the donor acceptor-transitions from C_{Ga} to C_N and the $(D^0A_C^0)$ transition at 3.085 eV to a donor acceptor-transition form an omnipresent residual donor in GaN to C_N. Both transitions are indicated in Fig. 3 by arrows. Up to now, the chemical identity of the residual donor is unknown, however O may be a good candidate for it.

Among substitutional insertion of C into the GaN films carbon may also be incorporated on an interstitial site or form diverse complexes. Such complexes may act as non-radiative centers, which reduce the overall PL-intensity or form deep defects, which may compensate the free carriers resulting in semi-insulating epilayers. The appearance of the deep-red luminescence band at 2.1 eV at highly C-doped GaN epilayers ($[C] \ge 1 \times 10^{18} \text{ cm}^{-3}$) is a sign for such compensation effects.

Concomitant to the reduction in the integral PL-intensity a strong increase in resistivity of the GaN:C epilayer is observed. Unfortunately, due to the low specific resistivity of the free-standing 3C-SiC substrates ($\rho \sim 6.6 \times 10^{-3} \Omega \,\mathrm{cm}$) Hall-effect measurements on the GaN epilayers itself could not be performed. Therefore, to get an idea of the influence of C-doping on the electrical properties of c-GaN *I–V* measurements on well-defined



Fig. 4. Room temperature current–voltage characteristics (log scale) measured between two ohmic Ti/Al contacts of an undoped (blue, upper curve) and a C-doped (red, lower curve, $[C] \sim 4 \times 10^{19} \text{ cm}^{-3}$) cubic GaN epilayer. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

Ti/Al contacts have been carried out. Two circular Ti/Al (30 nm/ 100 nm) ohmic contacts with a diameter of $300 \mu \text{m}$ and a center distance of 1.1 mm were produced by thermal evaporation. Fig. 4 shows the *I–V* characteristics of both the undoped and the C-doped samples in a semi-logarithmic scale. One clearly sees that the current flowing through the C-doped samples (red lower curve) is by more than four orders of magnitude lower than that for the undoped c-GaN layer (blue upper curve).

4. Conclusion

The growth of highly resistive non-polar c-GaN:C films via rfplasma-assisted MBE under Ga-rich growth conditions is demonstrated. SIMS measurements showed that the amount of incorporated C is linearly related to the incident CBr₄ flux and that no Br is built in. No degradation of the phase purity and structural properties of the c-GaN epilayers is observed up to a C-concentration of $\sim 10^{19}$ cm⁻³. At high C-doping levels the near band edge luminescence is strongly decreased and a deep-red emission band appeared. At the same time the resistivity of the GaN:C epilayer is increased by more than four orders of magnitude. These results demonstrate that C-doped GaN epilayers via CBr₄ can effectively be used to produce reliable and stable semi-insulating templates for future electronic devices.

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