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Citation: Journal of Vacuum Science & Technology B 23, 2514 (2005); doi: 10.1116/1.2131872

View online: http://dx.doi.org/10.1116/1.2131872

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Effects of doping type and concentration on precipitation of nanometer arsenic clusters in low-temperature-grown GaAs

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(Received 27 April 2005; accepted 10 October 2005; published 11 November 2005)

In this study, the effects of doping type and concentration on arsenic precipitation in low-temperature-grown GaAs upon postgrowth annealing at 600, 700, and 800 °C were investigated. Three undoped/Si-doped/undoped (i-n-i) regions and three undoped/Be-doped/undoped (i-p-i) regions were grown by low-temperature molecular beam epitaxy. The results show that arsenic precipitation is dependent on doping type and doping concentration. Arsenic depletion was observed in all Be-doped layers for all annealing temperatures. However, a “dual” arsenic precipitation behavior was observed in Si-doped layers: As accumulates in [Si]=2×10^{18} \text{cm}^{-3} doped layers, while it depletes in [Si]=2×10^{16} and 2×10^{17} \text{cm}^{-3} doped layers. We attribute this “dual” As precipitation phenomenon in Si-doped layers to the different depletion depths. © 2005 American Vacuum Society. [DOI: 10.1116/1.2131872]

I. INTRODUCTION

GaAs and related compounds grown by low-temperature molecular-beam epitaxy (LT-MBE) have attracted much attention due to their unique electronic and optical properties.\textsuperscript{1–3} When grown at 200–300 °C, the LT-GaAs layers are very nonstoichiometric, containing about 1 at. \% excess As and a high concentration of As related defects.\textsuperscript{4} Upon postgrowth annealing above 500 °C, the excess As precipitate into clusters\textsuperscript{5} and the LT layer’s resistance changes to an extremely high-resistivity state (>10^6 \Omega \text{cm}).\textsuperscript{6} This property offers the benefits of excellent device isolation in integrated circuits for the elimination of sidegating or backgating effects.\textsuperscript{3} The reduced recombination time of about 400 fs\textsuperscript{7,8} makes LT GaAs very suitable for ultrafast optoelectronic applications.

Although the annealed LT GaAs has demonstrated successful device applications, the out diffusion of As precipitates during annealing remains a major constraint in device design. It is, therefore, very important to control the density and distribution of As precipitates for potential device applications. Previous studies showed that As precipitation in annealed LT materials can be controlled by doping effects.\textsuperscript{5,8–15} It has been found that As precipitates preferentially form in Si-doped GaAs then intrinsic and least favorably in Be-doped GaAs for moderately doped GaAs. However, when the Si and Be doping concentrations reach certain levels (≥5×10^{18} \text{cm}^{-3}), an opposite precipitation behavior occurs.\textsuperscript{9,16} Chang et al.\textsuperscript{13} examined the As precipitation in an n-p_{1}-n-p_{2}-n-p_{3} or an i-n_{1}-i-n_{2}-i-n_{3}-i structure, where “i,” “n,” and “p” denotes undoped, Si-doped, and Be-doped GaAs, respectively, with doping levels ranging from 2×10^{16} (n_{1} or p_{1}) to 2×10^{18} \text{cm}^{-3} (n_{3} or p_{3}). Upon postgrowth annealing at 800 °C, distinctive depletion zones could be observed along the i-n_{1} interface, as generally expected. However, the depletion zone was less well defined or even unrecognizable at the i-n_{2} and i-n_{3} interfaces, and depletion zones along the p_{1}-n_{1} interface were narrower than those at the p_{2}-n_{3} and p_{3}-n_{3} interfaces; these precipitation behaviors cannot be fully explained by the so-called “doping” or “Fermi-level” effect. Both the inter-diffusion of As related defects and doping effects during annealing among the LT layers make this a more complicated system for understanding the As precipitation behavior in lightly-doped LT GaAs. Thereby, in this study we designed a LT-GaAs structure consisting of three i-n-i and three i-p-i active regions, with doping levels ranging from 10^{16} to 10^{18} \text{cm}^{-3}, in which each active region is separated with a 10 nm AlAs marker layer. Owing to the high activation energy but small diffusion constant for Al and Ga inter-diffusion,\textsuperscript{17} the thin AlAs layer also acts as an inter-diffusion barrier between adjacent active regions. By this way, the dependence of doping type and concentration on As precipitation behavior can be clearly distinguished. The As precipitation behavior was carefully characterized by transmission electron microscopy (TEM) and discussed.

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II. EXPERIMENT

The samples used in this study were grown by a Varian Modular GEN-II MBE system. The growth rates of 0.8 μm/h for GaAs and 0.2 μm/h for AlAs and the V/III beam equivalent pressure (BEP) ratio of 25 were used. Following native oxide desorption, a 100 nm GaAs buffer layer was first grown at 580 °C to smooth the surface, followed by a 10 nm AlAs diffusion barrier at the same temperature. Growth was then interrupted by closing the Al effusion furnace shutter, the substrate temperature was ramped down to 250 °C and the As shutter was closed when the substrate temperature was below 400 °C to maintain a clear 2 × 4 surface reconstruction, as observed by reflection high-energy electron diffraction (RHEED). It took about 15 min to stabilize the substrate temperature. Subsequently, the low-temperature active layers and a 35 nm GaAs cap layer were grown. As shown in Fig. 1, the LT-GaAs sample consisted of six active regions, i.e., three regions of undoped/Be-doped/undoped (i-p-i) multilayers and another three regions of undoped/Si-doped/undoped (i-n-i) multilayers. Each i-n-i or i-p-i region was separated by a thin AlAs layer, which acted as both a diffusion barrier and a marker layer. The thickness of each doped or undoped GaAs layer was 35 nm. The doping levels in the p1, p2, and p3 layers (and n1, n2, and n3 layers) were 2 × 10^16, 2 × 10^17, and 2 × 10^18 cm^-3, respectively.

Postgrowth annealing was carried out in a rapid thermal annealing (RTA) system in forming gas ambient with a GaAs proximity cap at 600, 700 and 800 °C for 30 s. Arsenic precipitation in annealed samples was examined using a JEOL JEM-2010 transmission electron microscope. Cross-sectional samples parallel to (110) planes were prepared conventionally by mechanical thinning and Ar-ion milling. To further characterize the concentration of excess As incorporated into the LT-GaAs layers, three 1 μm thick LT GaAs control samples [Si]=10^18 cm^-3 doped, [Be]=10^18 cm^-3 doped, and undoped were also grown under the same growth conditions as aforementioned. Double-crystal x-ray diffraction (DXRD) rocking curves were examined using a Philip DCD-3 double-crystal diffractometer.

III. RESULTS AND DISCUSSION

Figures 2(a)–2(c) show the bright-field TEM images of six active regions after 30 s anneals at 600, 700, and 800 °C, respectively. The bright stripes seen in each picture are the AlAs layers. It is noted that AlAs layers provide good markers among the (i-n-i) or (i-p-i) regions. As precipitation depletion was observed in all Be-doped layers for all annealing temperatures; moreover, the As clusters accumulate toward the undoped layers. The DXRD rocking curves of GaAs (400) reflection for the three control samples are shown in Fig. 3. Clearly, peak separations of 108 and 72 arc s caused by excess As incorporated in LT GaAs, were observed in Be-doped and undoped LT-GaAs epilayers, respectively; however, no obvious peak separation was observed in the Si-doped LT GaAs. This suggests that the Be doping enhanced the incorporation of excess As into the lattice of LT GaAs while the Si doping suppressed this effect. According to a previous report,11 the concentration of As antisites, [AsGa], is directly proportional to the lattice expansion of the LT-GaAs epilayer. Therefore, we can conclude that [AsGa] = [AsGa]_o/[AsGa]_i, where [AsGa]_o denotes the As antisite concentration in the x-doped control sample. Moreover, during postgrowth annealing, the reduction of strain caused by the AsGa defects is also a driving force for precipitation. For the i-p3-i region, the AsGa defects diffused from the p3 to i layers, and for the i-n3-i region from i to n3 layer, which is opposite to the VGa defects. Our experimental result about i-n3-i region and all i-p-i regions is therefore consistent with the vacancy-assisted As antisite diffusion mechanism.8,10,12,13

Interestingly, we observed a “dual” As precipitation phenomenon in Si-doped regions: As precipitates accumulate toward the center of the [Si]=2 × 10^18 cm^-3 doped layer for all annealed temperatures; however, As precipitates are depleted away the other two lightly Si-doped regions (n=2 × 10^16 and 2 × 10^17 cm^-3) toward the undoped regions. The latter depletion behavior contrasts with the earlier observation by Chang et al.13 In their work, the i-n3-i-n3-i-n3-i multilayer structure did not consist of any isolation layer to prevent the inter-diffusion of defects and doping effect from n3 to n2 or n3. As a result, As precipitation behaviors were disturbed and the correlation of doping concentration to As precipitation could not be clearly identified. The “dual” As precipitation phenomenon in Si-doped GaAs layers indicates that other effects must be considered in addition to the vacancy-assisted As antisite diffusion mechanism. According to the analysis of O’Hagan et al.16
As precipitation at the \( i/n \) interface can be influenced by the Debye length,\(^{18}\) \( \lambda_D = (\varepsilon \varepsilon_0 k T/[g^2(N_D-N_A)])^{1/2}. \) Following this argument, band bending\(^{14,18,19}\) and depletion of the “n” regions to a depth of up to a few Debye lengths would occur at each \( i/n \) interface. Moreover, according to the model proposed by Chandra \textit{et al.},\(^{18}\) the Debye lengths at the current annealing temperatures in our structure are estimated to be around 560, 180, and 50 Å for \( n_1, n_2, \) and \( n_3 \) layers, respectively. Therefore, the total depletion depth should be \( \sim 1120, \sim 360, \) and \( \sim 100 \) Å for \( n_1, n_2, \) and \( n_3 \) layers, respectively, due to the double \( n/i \) interfaces in each active region. Therefore, since each Si-doped layer is 350 Å thick, it is likely that complete depletion occurred in the \( n_1 \) and \( n_2 \) layers but an incomplete depletion in the \( n_3 \) layer, the latter was also noted as an “As accumulation” layer.

Figure 4 shows the average density of As clusters in each \( i-x-i \) region after annealing at 700 and 800 °C, respectively. The density of As clusters was determined by assuming that the sample thickness under TEM investigation was 100 nm. The analysis reveals that the cluster densities of the \( i-p-i \)
regions are higher than those of the i-n-i regions under the same doping concentration and annealing condition, which is consistent with the DXRD results.

IV. CONCLUSION

The effect of doping type and concentration on As precipitation in LT-GaAs upon post-growth annealing at 600, 700, and 800 °C was investigated. The results show that the As precipitation is dependent on doping type and doping concentration. As depletion were observed in all Be-doped regions. However, a “dual” arsenic precipitation phenomenon was observed in Si-doped regions: As precipitation accumulation was observed in $\text{Si}=2 \times 10^{18}$ cm$^{-3}$ doped layer for all annealing temperatures, while As precipitation depletion was observed in $\text{Si}=2 \times 10^{16}$ and $2 \times 10^{17}$ cm$^{-3}$ doped layers. The “dual” arsenic precipitation phenomenon in Si-doped layers can be attributed to the different depletion depths caused by various doping concentrations.

ACKNOWLEDGMENTS

This work was supported by the National Science Council, Republic of China, under Contract Nos. NSC 92-2112-M-007-032 and NSC 92-2120-M-007-006.