Top–down fabrication of horizontally-aligned gallium nitride nanowire arrays for sensor development

Guannan Liu a,b, Baomei Wen b,c, Ting Xie a,b, Audie Castillo b,c, Jong-Yong Ha a,b, Nichole Sullivan c, Ratan Debnath b,c, Albert Davydov b, Martin Peckerar b, Abhishek Motayed a,b,c,⁎

a Department of Electrical and Computer Engineering, University of Maryland, College Park, MD 20742, USA
b Material Measurement Laboratory, National Institute of Standards and Technology, Gaithersburg, MD 20899, USA
c NS Sensors, Inc., 9610 Medical Center Drive, Suite 200, Rockville, MD 20850, USA

1. Introduction

Gallium nitride (GaN) is a versatile semiconductor used in optical devices (light-emitting diodes, laser diodes, UV sensors) and in power electronics (power switches, RF devices, and high power transistors) [1]. Due to its direct band gap and chemical and thermal stability, GaN nanowires (NWs) are gaining significant importance as chemical sensor elements [2–4].

To date, most of GaN NWs devices are fabricated using chemical vapor deposition growth with a follow-up detachment from the substrate and individual on-chip alignment. Due to the variation in morphology, dimensions, doping, and crystal quality, it is hard to control the quality of NWs grown via the bottom–up methods [5]. Besides, variations from a multi-step pick-and-place fabrication process result in a low yield of functioning devices. Thus, such a process has low large-area integration capabilities, rendering mass-manufacturing challenging. An alternative approach to overcome these drawbacks is the top–down fabrication, in which NWs are patterned on a uniform thin-film using standard lithography and etching. Vertically-aligned semiconductor NWs (nanorods, nanopillars or nanocolumns) have been demonstrated by such top–down methods [6–10]. The heights of these nanostructures are limited by the thickness of the starting thin-film material, and they often exhibit tapering as a result of etching. On the contrary, horizontally-aligned NWs can be fabricated without any length limitations. For photodetectors and photovoltaic devices, vertically-aligned nanowires represent a high efficiency platform, whereas for chemical sensors horizontally-aligned nanowires are the optimal choice [11–13]. Due to its inherent inertness to most wet chemical etches, GaN nanostructures are commonly produced using dry etch techniques [8]. However, high-aspect ratio GaN structures achieved by plasma etching are often associated with extensive sidewall damage, result in lower performance of the device [14]. Therefore, producing damage-free nitride structures with a precisely-defined geometry over large area remains a challenge.

By combining deep-UV projection lithography and inductively-coupled plasma (ICP) etching, this paper demonstrates horizontally-aligned NWs etched from a GaN thin film grown on sapphire. Since the rough and tapered NW sidewalls due to dry etching create leakage current and limit NW performance [14], a post ICP potassium hydroxide (KOH) wet etching procedure was developed to smooth the NW walls.
The KOH etching also allows us to control the final shape and dimensions of the NWs. The PL and current–voltage (I–V) results verify the improved performance resulting from the KOH treatment.

The top–down approach described here can potentially enable GaN NW and SMW to be integrated with LED, micro-pumps, and micro heaters for system-on-chip development [15–20]. Different types of sensors such as chemical, gas and bio sensors can be fabricated using the top–down method reported here.

2. Fabrication

Two different thicknesses of commercial intrinsic GaN epitaxial layers (NTT Advanced Technology) grown on c-plane sapphire were used in this study: 3 μm thick GaN films were used for characterization of the etch rate of ICP dry etching, optimization of wet etching and measurements of PL spectra; 1 μm thick GaN films were used in the optimization of the wet etching process as well as in I–V characterization. AlGaN is a buffer layer widely used to compensate lattice mismatch and reduce stress in the growing film, and thereby improves the crystalline quality of GaN on sapphire. In our experiment, a 500 nm AlGaN layer was used between the GaN and sapphire substrate. A silicon nitride (SiN) passivation layer with thickness of 50 nm was deposited by plasma enhanced chemical vapor deposition (PECVD). This layer prevents metal penetration into the bulk GaN. A bi-layer stack of photo-resist (MicroChem LOR3A and Ultra-i) was spun onto the wafer for photolithography patterning. The 570 mJ/cm² stepper lithography dose was optimized for the fabrication process.

A bi-layer metal stack (50 nm Ti and 120 nm Ni) was deposited by e-beam evaporator (Denton Infinity 22) and lifted-off. This was followed by ICP etching (Oxford Plasmalab 100) with a combination gases (Cl/N₂/Ar 25/5/1 sccm) at 40 °C temperature, 300 W RF power and 800 W ICP power. After removal of Ti/Ni metal shadow mask and SiN, potassium hydroxide (KOH) wet etching was studied at various temperatures (40 °C, 60 °C, and 80 °C) and in different solvents (deionized water, isopropanol, and ethylene glycol), and for different etching times from 10 min to 5 h. The fabrication steps for NW formation are shown in Fig. 1. PL measurements and SEM imaging were conducted before and after KOH etching as well as on the control sample (pristine GaN film). The optimized condition of KOH treatment (10% KOH in ethylene glycol etching for 2 h at 80 °C) was used on NWs. An ohmic contact metal stack (Ti/Al/Ti/Au) was deposited by e-beam evaporation for I–V characterization of the samples. The electrical properties of the no KOH treatment control group and a 2 h KOH etched group with different length NWs were studied. Due to the highly resistive property of intrinsic GaN, I–V measurements were conducted under UV (355 nm wavelength, 48 μW/cm² intensity) illumination. The width of both
Fig. 4. SEM images of a nanowire section after ICP etch. (a) plan-view, (b) 45° tilted. The top surface appears smooth, however sidewall damage is evident.

3. Results and discussion

Fig. 2(a) shows SEM image of the longest length (5000 μm) wire in the meander-shape. The upper inset of Fig. 2(a) shows a low magnification image of 6 identical NWs. The lower inset of Fig. 2(a) shows a high magnification of a NW. The wire bonded NWs in a chip carrier for IV and photoconductance measurement are shown in Fig. 2(b). On a single chip, the total 9 groups of NW arrays with length range from 10 μm to 5000 μm were fabricated. And within each group, there are 9 identical NWs arranged in a 3 × 3 matrix.

Several GaN layers (3 μm thickness) were ICP etched for different times to calibrate the etch rate. The etched thickness was measured by a profilometer and the data summary is shown in Fig. 3. Black dots show the original data, and the regression linearization analysis (green line) gives the dry etching rate of 428.9 nm/min. Based on this rate, an 3 min etching was performed on the 1 μm GaN/500 nm AlGaN films to fabricate the nanowire devices.

Both top view and tilted SEM images of NWs after ICP etching are shown in Fig. 4. The vertical texture on the sidewalls formed by plasma bombardment is clearly seen in this figure. To remove the surface defects, KOH wet etching methods are explored with different etching temperatures, solvent compositions and times.

3.1. Wet etching temperatures

The samples were etched for 10 min in a 10% KOH in deionized water (DIW) solution at 40 °C, 60 °C and 80 °C. PL measurements were conducted before and after KOH etching as well as on the control sample (pristine GaN film). In Fig. 5, intensities are normalized with respect to peaks at near-band edge (NBE) of 355 nm. Surface defects are created during the ICP etch, which are non-radiative recombination centers, results in the decreasing of the NBE emission compared to the impurity-related PL peaks at 420 nm and 600 nm. The subsequent KOH etch removes the ICP-damaged layers from the sidewall surfaces and remarkably reduces the surface non-radiative recombination centers density. After etching at 80 °C, the overall shape of the PL spectrum of GaN NW surface resembles that of the pristine GaN film surface. Similar results also observed and reported on by Q. Li et al. [9].

3.2. Wet etching solvents

The 10% KOH in DIW etchant used above shows the improvement of PL intensity at 80 °C. However, the KOH in DIW shows aggressive etching of the a-plane of GaN, leading to an anisotropic etching profile as shown in the first column of Fig. 6. To address this issue, three different types of etch baths, namely 10% KOH in DIW, isopropanol (IPA) and ethylene glycol (EG), were studied in order to find a truly isotropic etch, one that is more controllable when used in batch fabrication.

All three groups were studied under the same conditions: 10 min etching at 80 °C. PL measurements were conducted after treatment. These results are similar to those shown in Fig. 5 (purple line). SEM images (Fig. 6) reveal different etching rates of different facets: for the top surfaces (c-plane) of GaN NWs, etching was not observable for all three cases. The sidewalls of the NWs (a-plane and m-plane) were found to be more easily attacked by KOH.

Generally speaking, the a-plane is less stable than m-plane due to a higher surface energy [21,22], so the etch rate is faster on a-plane than on m-plane. SEM images (Fig. 6) show NWs disconnected at several points after etching with KOH in DIW, while m-plane NWs remain the same. Thus, we conclude KOH in DIW etches a-plane much faster than the other solutions. A high reaction rate is not favorable in achieving the smooth etched surfaces [23,24]. An organic solvent addition, such as IPA or EG, slows down the a-plane etching process and making it comparable in rate to m-plane etching in order to achieve “isotropic” etching. The organic solvent contained in the solution plays an important role in the etching process. The organic molecules of will selectively adhere to some crystallographic planes (a-plane in our case), thereby hindering the access of etching agent (KOH) and slowing down the corresponding plane's etching rate. KOH in IPA etching shows an improvement in the isotropy of the etch process. IPA can slow down the etch rate of a-plane. However, it still etches faster than the m-plane. KOH in EG etching shows the most isotropic result since both a-plane and m-plane end up with the same NW width. Both the first and second
columns of Fig. 6 highlight the original NW sidewalls before treatment by blue dashed lines. It is obvious that the a-plane etches faster than m-plane NWs in DIW and IPA. The high magnification image of the 90° bend parts proves this fact. The inner side of the NWs is etched to m-planes with 120° inter-planar angles between them. This is highlighted by the yellow dashed line in these figures. These results

![Fig. 6. SEM images of NWs after KOH in different solvents. The first row indicates the a-plane and m-plane wires under study. The second row shows the conjoined parts. This is a short a-plane NW connects two m-plane NWs. For DIW and IPA solvents, a-plane NWs were over-etched and disconnected after treatment. Blue dash lines indicate the original NW sidewall positions before the treatment. Yellow dash lines indicate that the final sidewall position after treatment. The m-planes replaced the original a-plane at the inner loop of the part because of a higher etch rate on the a-plane (in contrast to the m-plane.) For the EG solvent, NWs were not severely etched after the treatment. This result shows that the KOH in EG solvent is the best candidate for isotropic etching among the three explored. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)](image)

![Fig. 7. SEM images of NWs after KOH in EG treatments. (a)–(d) correspond to different etching times from 10 min to 3 h under condition of 10% KOH in EG at 80 °C. Blue dash lines in (c) and (d) indicate the original NW sidewall position before the treatment. Yellow dash lines in (c) and (d) indicate the interface between GaN and AlGaN layers after treatment. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)](image)
show that KOH in EG solvent is the best candidate for an isotropic etching for GaN.

3.3. Wet etching time

A series of studies of etching times from 10 min to 5 h based on the optimized condition of 10% KOH in EG at 80 °C were conducted. SEM images were taken to observe the NW sidewall surface roughness changes with the increasing etching time. Fig. 4 shows the control group without wet etching treatment, and sidewalls were rough created by plasma bombardment can be clearly seen. Fig. 7 shows a set of SEM images of NW sidewalls after wet etching of 10 min, 20 min, 2 h, and 3 h. The NWs had similar surface morphology with etching times greater than 3 h etching. Both 2 h and 3 h of etching show smooth sidewalls. Comparing Figs. 4 and 7(a), we conclude KOH goes into the deep trenches to widening them and causing the vertical textures to become more pronounced at the beginning of the wet etching. After that, the horizontal textures and steps on the tapered sidewall become visible. The horizontal and vertical texture “cut” the sidewalls into “brick-like” blocks (Fig. 7(b)). These convex block textures are attacked by KOH with further wet etching and decreases the size into smaller convex spots (Fig. 7(c)). After 3 h of etching, the convex structures largely disappeared, ending up with smooth sidewalls (Fig. 7(d)). In Fig. 7(c), (d), the interface layer, highlighted with yellow dash lines, of GaN and AlGaN are identified easily on the smooth sidewall. The original sidewall’s position before wet etching can be determined by a KOH resistive nucleation layer as highlighted with blue dash lines.

3.4. Electrical characterization

The electrical properties of the no KOH treatment control group NWs and a 2 h KOH etched NWs with different lengths were studied. I–V measurements were conducted under UV (355 nm wavelength, 48 μW/cm² intensity) illumination. Due to the high resistivity of intrinsic GaN, dark currents measured in the pA range (from 10 pA to 100 pA). With UV excitation, resistances tend to increase with increasing NW length, shown in Fig. 8. The KOH treated NWs show lower resistance compared to those no KOH treated NWs at the same length under UV excitation. This enhancement becomes more significant as the NW length decreases. This phenomenon can be explained either by the suppression of the non-irradiative recombination process due to surface improvement or the surface treatment may lead to a lower barrier resistance and form the ideal ohmic contacts.

UV photocurrent measurements were conducted with 200 μm and 1000 μm NWs. For both lengths, NWs with and without KOH wet etching samples compared. The same UV source intensity and voltage biases were used for 5 cycles, 2 min for each cycle (1 min UV on and 1 min off). Fig. 9 shows the response of the NWs’ conductance to this cycle. Note: the NW is almost insulating without UV exposure. For the same dimension, surface-treated NWs show almost 3 times higher current under UV exposure. Furthermore, the UV response is proven to be repeatable and reliable over multiple cycles.

4. Conclusions

In this study, we were able to reliably and reproducibly fabricate different lengths of alignment-free, horizontal nanowire arrays for various applications. Stepper lithography, ICP etching and KOH wet etching techniques were used to ensure mass production capability and competitive performance to conventional bottom-up nanowires. Different wet etching temperatures, solvents, and times were explored for surface improvement of NWs. PL, SEM and I–V characterization were used to show wet etching is necessary for optimal GaN NW devices. On the basis of our experimental results, this NW fabrication method shows promise, potentially enabling mass production of sensors, optical and power devices based on single NW structures. The top–down approach described here enables GaN NW and SMW to be integrated with LEDs, micro-pumps, micro heaters to provide useful sensor structures. Many sensors (chemical, gas and biology) using bottom-up GaN NWs can be benefit from the top–down approach.

References


