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Progress in periodically oriented III-nitride materials

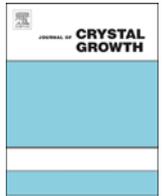
Jennifer Hite

U.S. Naval Research Laboratory, jennifer.hite@nrl.navy.mil

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Progress in periodically oriented III-nitride materials



Jennifer Hite

U.S. Naval Research Laboratory, Washington, DC, USA

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ABSTRACT

The ability to grow III-nitride structures with alternating c-plane orientation has garnered interest in using these materials for new application spaces, such as frequency conversion. An overview of recent progress in growing periodically oriented (PO) III-nitrides is discussed, including AlN, AlGa_N, and GaN. Successes in fabricating thick PO GaN structures (> 500 nm) for uses in frequency conversion are highlighted.

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1. Introduction

The family of III-nitride materials is well known for applications in electrical and optoelectronic devices. Some of the more common applications include transistors and visible LEDs, along with a push towards using Al-containing alloys for DUV LEDs [1–7]. However, all of these structures are based on using a single crystal orientation. Engineering structures with opposing polar faces (III- or N-polar) abutting one another on a single substrate opens a promising avenue to using these materials for frequency conversion via quasi-phase matching [8–10]. The interest in III-Ns in frequency conversion over more conventional materials stems from their wide band gaps, high thermal conductivity, and wide transparency windows [11].

On sapphire substrates, creating these engineered structures takes advantage of the ability to control the polar orientation of III-nitrides by manipulation of the buffer layer. Growth of GaN, AlN, and alloys of the two on sapphire substrates are generally III-polar when grown on low temperature AlN buffer layers of sufficient thickness (> 15 nm), while those on sapphire substrates without a buffer layer are N-polar [12–14]. The process to create periodically oriented (PO) structures, which consist of alternating regions of opposing crystal orientations in the c-direction consists of several steps. First, the buffer layer, or buffer layer plus III-polar nitride layer, is grown. It is then patterned and etched back to the sapphire substrate. Afterwards, III-N layers are grown, with N-polar material grown over the regions of revealed substrate while simultaneously growing III-polar material over the areas with the buffer layer. This succession is shown in Fig. 1. This has successfully been demonstrated with GaN, AlGa_N, and AlN [15–17].

Although this process on sapphire substrates is useful in most

thin film nitride device structures and could be employed in waveguide structures for frequency conversion, to achieve bulk frequency conversion equivalent to commercially available materials, thick layers of III-nitrides, on the order of millimeter thicknesses, are required. This is because thin films on sapphire are strained and have a large density of dislocations. Thick layers also reduce internal reflection and scattering. To reach these thicknesses, native substrates are required to allow for thick growth with high crystal quality. The most successful method of creating thick PO GaN structures on N-polar substrates has subtle differences from the methods on sapphire, namely in the use of a deposition of patterned oxide to selectively grow the inversion layer. Although other inversion layers, such as Mg-doping have been demonstrated, they have issues in interface quality or require large concentrations of Mg, which can lead to uncontrolled inversion domains and clustering [18,19]. The lift-off oxide approach results in a patterned inversion layer, but maintains an epi-ready surface without plasma etch-induced damage [20–22] prior to regrowth [23–25].

2. Al-containing nitrides

Recently, PO AlN structures have been grown on sapphire substrates via metal organic chemical vapor deposition (MOCVD) [17,26]. These structures showed a well-defined boundary between the Al- and N- polar regions with very little difference in growth rate between the two. The latter is very different than what has been observed with PO GaN growth [13,27]. In this case, the N-polar regions also showed columnar growth in the initial nucleation on sapphire and is associated with rougher growth.

Some of the first PO AlGa_N structures reported were thin GaN/AlGa_N/GaN structures grown on sapphire by a hybrid molecular beam epitaxy (MBE) and MOCVD approach, and used Al

E-mail address: jennifer.hite@nrl.navy.mil

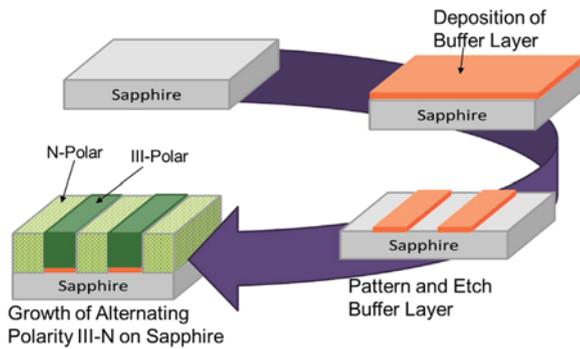


Fig. 1. Process schematic used to create periodically oriented III-nitride materials. First, a buffer, or inversion, layer is grown or deposited. This layer is patterned and etched back to the sapphire substrate. A III-N layer is deposited, resulting in III-polar material over the buffer layer and N-polar over the bare sapphire.

concentrations between 15% and 30% [15]. This showed the feasibility of creating such structures. A more recent study shows PO AlGaN stripes with much higher Al content (60–100%) [16]. Both PO AlN and PO AlGaN could be useful for DUV applications.

3. Progress in GaN

Although PO $\text{Al}_x\text{Ga}_{1-x}\text{N}$ structures are beginning to be investigated, PO GaN structures on sapphire have been more rigorously examined, with an eye towards both frequency conversion as well as more conventional electrical device structures.

3.1. PO GaN on sapphire substrates

Using the method described in the introduction, several groups have reported PO GaN structures on sapphire substrates using MBE and MOCVD [12,13,28–31]. As in the case for PO AlGaN structures, the N-polar regions are generally rougher than the Ga-polar regions [32]. From shifts in the $A_1(\text{LO})$ peak, micro-Raman spectroscopy showed an elevated free-carrier concentration in the N-polar region, which indicates higher impurity incorporation of the N-polar face over the Ga-polar face even when grown simultaneously [33]. This is a trend that has been shown in blanket film growth as well [34]. This study also showed a slight shift in the Raman E_2 peak with polarity, indicative of slightly larger compressive in-plane stress for the Ga-polar material [33]. Work function studies have shown a difference of 0.2–0.25 eV between the two polar regions [35,36].

One interesting aspect of these PO structures is the inversion domain boundary (IDB) that occurs at the junction of the two polar regions. In examining these structures using photoluminescence (PL) spectroscopy, an intense recombination was observed at the IDB [13,32,37]. In PL, it has been shown to be an order of magnitude higher in intensity and 30–40 meV lower in energy than bulk GaN, as well as being polarized parallel to the boundary [37]. Temperature dependent PL studies correlate with calculations by Fiorentini [38] indicating the emission arises from localized potential maximum in the IDB coupled with minima in the surrounding polar regions [32]. This intense emission was also observed in UV photoelectron emission microscopy (PEEM) [35]. In transmission electron microscopy (TEM) images, scanning electron microscopy (SEM) images, etching studies, and work function investigations, the IDB has been shown to range laterally in thickness from 50 nm to 5 μm and have mixed polarity [12,36,39]. This mixed polarity region has been attributed to imprecise etching of the AlN buffer layer and could contribute to a non-vertical IDB [32,39]. However, the PEEM study indicated that

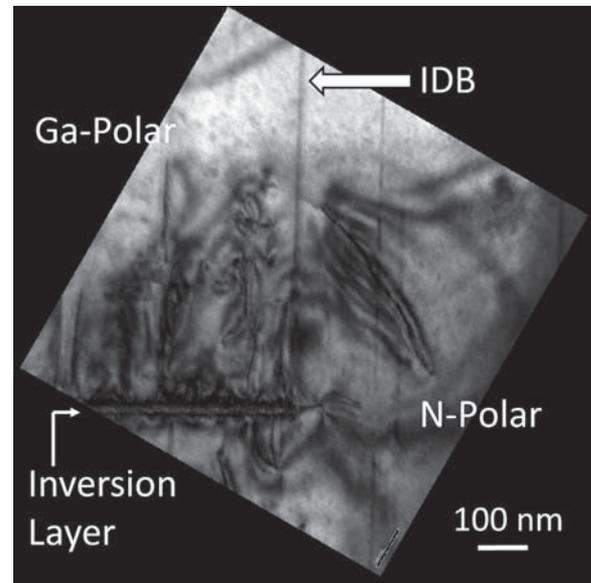


Fig. 2. TEM cross-section of PO GaN structure on an N-polar GaN template. New dislocations initiating at inversion layer interface in Ga-polar region can be observed.

the IDB could exhibit rectifying behavior for electronic transport across the IDB [35].

Testing out this indication, several device structures have been fabricated, both electrical and optical. The electrical structures have included p/n junctions and a MESFET [31,40,41]. These demonstrate the feasibility of using the IDB as a lateral junction. For optical devices, there was a single study designing and growing PO GaN stripes for wavelength conversion [8]. Though the reported single harmonic generation was low in efficiency, it also demonstrated the possibilities for this type of structure.

3.2. PO GaN on bulk substrates

Although several methods to control GaN polarity on GaN have been reported [42–45], this section will focus on the effort that have resulted in thick PO GaN structures using a low temperature AlN inversion layer on N-polar GaN substrates [23–25]. TEM images show the IDB to be extremely sharp and vertical (Fig. 2). Along with these TEM results, etching studies and SEM images have shown a very distinct boundary between polar regions without indications of mixed polarity at the boundary. With careful optimization of growth conditions, the growth rates of the two polar materials can be nearly matched, with height differences under 80 nm [27]. PO GaN templates up to 2 μm in thickness have been grown by MOCVD on bulk GaN substrates. For these thin layers, a large difference in dislocation density has been observed in the two polarities. Low densities are observed in the homoepitaxially grown N-polar regions, matching that of the substrate. The Ga-polar regions show higher dislocation densities, as they are grown heteroepitaxially over the inversion layer [23]. Scanning Kelvin probe microscopy (SKPM) on these samples have shown a 0.3 eV difference in surface potential, similar to that seen on PO GaN structures on sapphire [46]. The study also indicated that the native oxide on the N-polar regions was better at charge trapping than that over the Ga-polar regions.

To produce thick PO GaN structures, 2 μm thick MOCVD-grown PO GaN structures have been used as templates for additional hydride vapor phase epitaxy (HVPE) growth. In order to produce thick PO GaN structures, it is imperative to reduce differences in the growth rates of the two polarities. If not equalized, these

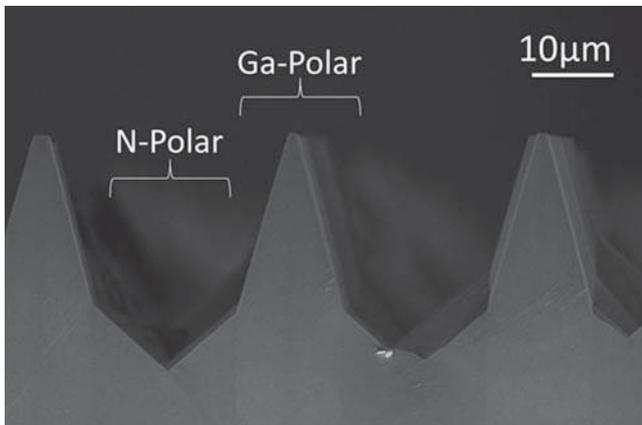


Fig. 3. Cross-sectional SEM image of the saw-tooth pattern at the surface of a thick (80 μm) PO GaN HVPE extension on a 2 μm thick MOCVD-grown PO GaN template on a bulk HVPE substrate. This is formed due to differences in growth rate between the two polar regions.

differences result in a saw-tooth or picket-fence type of morphology at the surface, as seen in Fig. 3. This can lead to nucleation of polycrystalline material, which rapidly dominates the growth, ruining the structures. The saw-tooth pattern can also affect impurity incorporation, with low impurity incorporation on +c-oriented facets, higher incorporation on -c-oriented facets, and even higher incorporation on semi-polar facets [34]. Even with differences in growth rate, TEM and SEM images show that the IDB remains vertical throughout the PO structure.

Overcoming these troubles, PO GaN structures consisting of 16 μm wide stripes have been produced with thicknesses reaching 1 mm. To date, structures with thicknesses of 100, 200, and 500 μm have been optically measured for frequency conversion [47]. These samples show second harmonic generation with a strong peak with a width of 10 nm, using a pump wavelength of 1.6 μm . The peak position variation between samples was $\pm 1\%$. Calculations show that the efficiency of these samples is 40% of the ideal structure. The efficiency was reduced by large variations in free carrier concentration as well as errors in the period of the structure. Further optimization in growth and processing should increase the efficiency.

4. Summary

In summary, PO III-nitride structures of AlN, AlGaIn, and GaN have been grown. The Al-containing PO structures are still being developed for device applications. Thick PO GaN structures have been produced that show single harmonic generation. This research offers the promise of engineered materials with custom lateral and vertical polarity variations for applications in novel electronic and optoelectronic devices, a subset of which are expected to be suitable for non-linear optics.

Acknowledgments

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