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III-Nitride semiconductors GaN, InN, and AlN and their heterostructures have triggered a rapid expansion of photonic and electronic device applications into new wavelength, voltage, and frequency regimes. Bringing new and unique physical properties into this semiconductor family by new, epitaxially compatible nitride materials holds the promise to significantly expand what is possible today.

Transition metal Sc is stable in a hexagonal crystal structure in its elemental form. The Sc ion, as a group III element, has a stable oxidation state of +3. This allows the formation of rock salt scandium nitride (ScN) with N in the −3 oxidation state and isovalent alloying when replacing the group III elements In, Ga, Al, or B for the III-nitride materials. This feature, combined with predictions of a metastable nonpolar h-BN-like, and wurtzite crystal structures, has led to significant interest in ScN and its alloys. These alloys promise to bring to the established GaN and AlN based electronics and photonics family missing properties such as plasmonic, thermoelectric, extremely high piezoelectric, and also ferroelectric behavior in ScAlN.

As Sc-based alloys with AlN and GaN (such as ferroelectric ScAlN) are beginning to be explored by epitaxy, it is essential that the epitaxial growth of the limiting binary ScN thin films and their physical properties are understood. The binary compound scandium nitride (ScN) is part of the family of the transition metal nitride semiconductors, with desirable physical properties such as high hardness, mechanical strength, and high temperature stability. Its equilibrium phase has a face-centered cubic (FCC) rock salt (NaCl) structure with a lattice constant of 4.505 Å. The (111) lattice constant of cubic ScN is nearly lattice-matched (only ~0.1% difference) to the c-plane lattice constant of wurtzite GaN. This has led to avenues for the use of ScN as a dislocation reduction buffer layer and for in situ Ohmic contacts for GaN based devices. Earlier studies have reported the growth of binary ScN thin films by reactive magnetron sputtering, hydride vapor phase epitaxy (HVPE), and gas-source and plasma MBE on Si, Al2O3, SiC, and MgO surfaces. Very few reports exist on ScN epitaxy on GaN, and no results have yet been shown for growth on AlN. One literature report mentions mixed (110) and (111) orientations for ScN grown on AlN, but the data are not presented.

Here, we present a comparative study of the plasma-MBE growth, structure, surface morphology, and electrical transport properties of ScN thin films (~30 nm) on c-plane GaN, AlN, and SiC surfaces. We find that extremely smooth epitaxial thin films of high crystalline quality of rock salt ScN grow with their 111 axes aligned with the polar 0001 c-axis of the wurtzite substrates of GaN, AlN, and SiC and are unintentionally n-type doped with degenerate electron concentrations. Because the cubic crystal of ScN grows on a hexagonal lattice, cubic twins in the ScN layer are observed.
in the MBE environment. An electron beam is steered into a W crucible with a magnetic coil to create the Sc flux. Flux stability was achieved using an Inficon® RF UNI-Bulb plasma source, with a growth pressure of approximately 10⁻³ Torr. In situ monitoring of film growth was performed using a Veeco® RF UNI-Bulb plasma source, with a growth pressure of approximately 10⁻⁴ Torr. Kikuchi patterns were indexed to face centered cubic ScN (Fm-3m space group, 225). Temperature-dependent Hall-effect measurements were taken using indium contacts in a Van der Pauw geometry in a Lakeshore® system with a 1 T magnet from approximately 250 °C to 350 °C.

ScN films were grown in nitrogen-rich conditions at a thermocouple temperature of 750 °C, a Sc flux of 0.16 A/s, a nitrogen plasma condition of 1.95 standard cubic centimeter per minute (sccm) and 200 W, and a chamber pressure of 1.5 × 10⁻⁵ Torr. Nitrogen rich growth conditions were utilized to suppress nitrogen vacancy formation and maintain a 1:1 Sc to N stoichiometry in the epitaxial film. The ScN layer thickness was 30 nm with a growth rate of 30 nm/h.

During growth, RHEED was utilized to continuously monitor the surface crystal structure of ScN. During growth on hexagonal substrates, the (110) azimuth evolved from first order 1 × 1 streaks for GaN to pairs of symmetric spots on either side of the original first order streaks for ScN. These spots can be viewed as rotated variants of (110) (002) families of planes, as shown in Fig. 1. This RHEED pattern and indexing have not been reported before for ScN (111) films but were seen in Cu (111) thin films grown on Al₂O₃.²³

The observed RHEED pattern implies the existence of twin domains that are expected to result from the symmetry constraints encountered upon growing a threefold symmetric cubic crystal on a sixfold symmetric hexagonal substrate. The (110) zone axis for a cubic single crystal would only have two 111 and two 002 diffraction spots, one each on each side of the zone center (000). Here, four 111 and four 002 diffraction spots are seen, two on each side of the zone center, indicating a (111) cubic twinned crystal.

To further test for the existence of twin domains in the epitaxial ScN layers, electron backscatter diffraction (EBSD) measurements were performed on ScN films grown on 6H-SiC and GaN/SiC template substrates (Fig. 2). In EBSD, the incident beam-sample-detector geometry is such that backscattered electrons escape the sample through the Bragg angle and diffract and form Kikuchi bands. If the chemical composition is known, the orientation sensitive Kikuchi bands can be indexed to determine the crystal orientation in different regions. Accordingly, Kikuchi patterns were indexed to face centered cubic (FCC) ScN to verify the symmetry of the ScN film. EBSD topography images showed striped patterns with alternating orientations. Two different colors indicate two different orientations, as expected for domains on either side of a twin boundary. Pole figures with the (111) direction out of plane indicated sixfold symmetry, as shown in Fig. 2 for samples grown on 6H-SiC and GaN/SiC. The corresponding AFM images indicate highly smooth surface morphologies, with rms roughness below 1 nm for 10 × 10 µm scans for ScN layers that are 30 nm thick. Grain misorientation statistics (not shown) indicated that the ScN epitaxial film was entirely (111) oriented, with 60° being the dominant in-plane misorientation angle.

Hence, we find that the ScN samples are not polycrystalline with random grain orientations, but rather highly oriented crystals with grain misorientation. EBSD determined twin domain sizes differing from approximately 800 nm for samples grown on SiC and 60 nm when grown on GaN/SiC. It is currently unclear if step-terrace or other surface features from the substrate such as threading dislocations influence the nucleation of cubic-twin domains and subsequent size of these domains. Growth on bulk substrates combined with transmission electron microscopy (TEM) studies can answer this question definitively in the future.

The orientation of the ScN crystal planes was further assessed using X-ray diffraction (XRD) measurements, as seen in Fig. 3. Whereas the ScN peak is clearly resolved for growth on SiC and AlN, the films grown on GaN/SiC template substrates show no separate ScN peak. This is because the ScN peak lies within the background of the GaN (002) peak. This is expected given the lattice constant of ScN (111) is only 0.18% different from the in-plane lattice constant of wurtzite GaN, 3.189 Å. For films grown on AlN/Al₂O₃ and SiC, separate ScN (111) peaks are observed; yet, none of the samples showed cubic ScN (002) reflections, indicating that ScN forms a highly oriented film with the (111) direction aligned along the 0001 c-axis of the substrate and perpendicular to the growth plane. The peak positions of ScN (111) in the films grown on SiC and are consistent with those reported in the literature occurring near 2θ = 34.5°.²⁵ Scans of the ScN (224) peak for a sample grown on SiC indicate sixfold symmetry, with peaks separated by sixty degrees. This is further support for the nature of a cubic-twinned ScN (111) film.

FIG. 1. RHEED pattern during evolution along the [110] azimuth from GaN (0001) to ScN (111). 111 and 002 pairs of kinematically allowed diffraction spots are symmetrically rotated about the zone center, illustrating that ScN grows as a cubic-twinned crystal.
The ScN films grown on all three hexagonal substrates (SiC, AlN, and GaN) adopted a twinned face centered cubic crystal structure instead of a hexagonal structure. Reasons for the stability of the cubic phase of ScN can be qualitatively found in the principles of molecular orbital energies and quantitatively verified by first-principles energy calculations. For binary nitrides, the three 2p orbitals of the nitrogen anion and the three lower t<sub>2g</sub> states (of the crystal-field split five 3d orbitals) of the scandium transition metal cation hybridize to form bonding and antibonding orbitals. The remaining two 3d orbitals remain as nonbonding energy states, with energy levels close to the transition metal 3d orbitals. A schematic of this pd coupling dominated bonding is shown in Fig. 4, and a similar bonding diagram has been illustrated previously in the literature. 15

**FIG. 2.** (a) EBSD topography image of ScN grown on 6H-SiC (left). EBSD pole figure of ScN with the [111] direction out of plane (center). 10 × 10 µm AFM image (right). (b) EBSD topography of ScN grown on GaN/SiC (left). EBSD pole figure of ScN with the [111] direction out of plane (center). 2 × 2 µm AFM image (right). Alternating stripe patterns in the EBSD topography images indicate domains with a 180° orientation device, with six spots in the EBSD pole figures indicating sixfold symmetry in the [111] direction.

**FIG. 3.** (a) and (b) XRD 2theta-omega scans of ScN grown on 6H-SiC, AlN/Al<sub>2</sub>O<sub>3</sub>. (c) and (d) XRD Phi scan of the ScN (224) peak grown on 6H-SiC and XRD 2theta-omega of ScN grown on GaN/SiC. The XRD results indicate that ScN grows epitaxially in an (111) orientation on the respective substrates, with sixfold in-plane rotational symmetry as seen from the Phi scans.
The three outermost electrons of Sc \([\text{Ar}4s^23d^1]\) bond with three N 2p electrons in a crystal to make Sc\(^{3+}\). Adding more electrons (e.g., using Ti, V, Cr, Mn... instead of Sc) will populate the antibonding and then nonbonding energy states, making the rock salt crystal structure less stable and the crystal metallic. On the other hand, ScN is an extremely stable rock salt semiconductor because of the filled bonding states, with a completely empty band in the Sc\(^{3+}\) configuration. This is in accordance with first principles calculations that show that the rock salt crystal structure is the ground state for ScN and that ScN has one of the lowest formation energies for any binary nitride.\(^{34,35}\) ScN is a semiconductor with an indirect bandgap of 0.9 eV and a direct bandgap of \(\sim 2.1\) eV.\(^{36}\) XPS results (not shown) verify the presence of Sc-N bonding, and optical absorption measurements (not shown) show a weak band edge absorption near 2.1 eV.

The electrical transport properties of the 30 nm thick MBE grown ScN on GaN/SiC and AlN/Al\(_2\)O\(_3\) template substrates were assessed using temperature-dependent Hall-effect measurements from room temperature to 20 K, as shown in Fig. 5. A high n-type carrier concentration independent of temperature was observed. The lack of carrier freeze out at low temperatures indicates that the nominally undoped ScN obtained in this work is a degenerately doped semiconductor. As shown in Fig. 5, the carrier concentrations and mobilities were 1.55 \(\times 10^{20}/\text{cm}^3\) and 23 cm\(^2/\text{V}\cdot\text{s}\) and 1.05 \(\times 10^{20}/\text{cm}^3\) and 11 cm\(^2/\text{V}\cdot\text{s}\) for samples grown on GaN/SiC and AlN/Al\(_2\)O\(_3\), respectively. The high carrier concentration is an order of magnitude higher than the reported values of the conduction band density of states for many semiconductors, giving support for their degenerate, metallic electrical behavior. At this stage, it is unclear if ScN is strained or relaxed to AlN and if the crystal quality and carrier mobility are affected by the approximately 2.4% lattice mismatch. The obtained mobility values are lower than those reported previously.\(^{26,27,29,31,34,42}\) for similar carrier concentrations, and this may be due to increased impurity scattering from domain boundaries in (111) oriented films. The (111) surface is a higher energy surface than the (100) surface for a face centered cubic crystal. Higher surface energy planes are more likely to trap impurities like oxygen, potentially due to a decrease in adatom mobility. This has been shown in ScN, where the oxygen concentration was higher for cubic-twinned (111) growth on c-plane sapphire compared to untwinned cubic (111) growth on MgO (111).\(^{31}\) This has also been reported in the case of Si and Ti segregation to twin domain boundaries in Mg\(_2\)Al\(_2\)O\(_4\).\(^{26}\) A high electron carrier concentration has previously been reported in nominally undoped ScN, with possible causes being linked to nitrogen vacancies, Sc-N antisite defects, and atomic level concentrations of oxygen and fluorine originating from source and crucible material. Density functional theory (DFT) calculations have shown that oxygen substitutional defects are lower in formation energy than the other defect mechanisms mentioned above.\(^{38}\) Support for oxygen incorporation into the film and donating electrons comes from scandium metal’s high affinity for oxygen, as evidenced in its large negative enthalpy of formation for Sc\(_2\)O\(_3\) from Ellingham diagrams.\(^{39}\) Carrier mobilities up to 100 cm\(^2/\text{V}\cdot\text{s}\) at carrier concentrations of \(10^{21}/\text{cm}^3\) have been previously obtained in ScN on MgO (001),\(^{26}\) and mobilities up to 284 cm\(^2/\text{V}\cdot\text{s}\) at a carrier concentration of \(3.7 \times 10^{18}/\text{cm}^3\) have been obtained in ScN grown using hydride vapor phase epitaxy (HVPE) on m and r-plane sapphire substrates. The lower carrier concentration and higher mobility in HVPE growth were due to a reduction in impurities in the film, notably oxygen concentration, through utilization of 6N (99.9999%) pure ScCl\(_3\) and NH\(_3\) as the source materials instead of Sc metal.\(^{41}\) This points toward an important role that impurities play in determining the carrier concentration and limiting the mobility of ScN films. Using ionized impurity scattering models,\(^{42}\)
assuming that all donors are ionized and a density of states effective mass of \( \sim 0.35 m_t \), where \( m_t \) is the free electron mass, a close agreement with the obtained mobility values is found. However, we point out that the interface between rock salt ScN and hexagonal GaN or AlN is a non-polar/polar interface, and the polar discontinuity across the interface, assisted by the conduction and valence band offsets, can give rise to mobile carrier concentrations even in the absence of defects and impurities. \(^{42}\) Similar polar/nonpolar interfaces have been found in several oxides, but not in the nitride crystals yet. Future work involving the measurement of the electronic structure and electron mobility will give further insight into the mobility limiting mechanisms in ScN and, more importantly, the origin of the mobile charges in the bulk and at interfaces.

In this work, we have reported the MBE growth of highly crystalline ScN thin films on hexagonal GaN, AlN, and SiC substrates. ScN films exhibited solely cubic twinned (111) orientation on all three hexagonal substrates and did not adopt a hexagonal crystal structure. ScN films exhibited large n-type carrier concentrations of approximately \( 10^{10}/\text{cm}^2 \) with mobilities of approximately \( 20 \text{ cm}^2/\text{V s} \). SEM-EBSD and Mike Hawkridge for discussions related to XRD. to acknowledge Don Werder for discussions related to EBSD and Air Force Office of Scientific Research, Grant No. AFSOR FA9550-15-C-0001 (N00014-15-1-2521). The authors would like to acknowledge the National Nanotechnology Coordinated Infrastructure (NNCI), performed in part at the Cornell NanoScale Facility, a member of MRSEC program (No. DMR-1719875). This work was also supported by the Cornell Center for Materials Research Shared Facilities which are supported through the NSF/C24.

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The origin of the mobile charges in the bulk and at interfaces.

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18. See https://icsd.fiz-karlsruhe.de/search/basic.xhtml for “Inorganic Crystal Structure Database.”

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