

The Effect of the Ion Beam Energy on M-plane InGaN Layer Preparation for STEM

Trang Nguyen¹, Brandon Dzuba¹, Yang Cao¹, Alexander Senichev², Rosa Diaz², Geoffrey Gardner², Michael Manfra^{1,2,3,4}, Oana Malis¹

¹. Department of Physics and Astronomy, Purdue University, West Lafayette, IN, USA.

². Birck Nanotechnology Center, West Lafayette, IN, USA.

³. School of Electrical and Computer Engineering, Purdue University, West Lafayette, IN, USA.

⁴. School of Materials Engineering, Purdue University, West Lafayette, IN, USA.

Nonpolar AlGaIn/InGaIn superlattices are investigated as a strain balanced potential alternative for nonpolar high Al composition AlGaIn/GaIn for near infrared intersubband absorption [5]. Inhomogeneity and defects appearing during InGaIn growth are still challenges to the realization of high performance nitride infrared optoelectronic devices. These defects can be imaged with various TEM imaging modes [1-4]. Nonetheless, damage caused by the sample preparation process reduces the quality of images and raises difficulties to the study of defects in InGaIn thin films. We discuss in detail our attempts to prepare cross section TEM samples of 30nm m-plane $\text{In}_x\text{Ga}_{1-x}\text{N}$ ($x = 0.1-0.15$) using the traditional polishing method followed by ion polishing with various beam energies and the resultant quality of the specimens.

Our InGaIn thin films are grown on m-plane free standing GaIn substrates by plasma assisted molecular beam epitaxy. These samples are prepared for TEM using traditional methods followed by an ion beam milling process with a Gatan Precision Ion Polishing System II (PIPSII). Prior to the mechanical polishing process, samples are glued together and cured at 160°C for 1.5 hours with the M-bond 610. Then, specimens are pre-thinned using an Allied Optiprep Polishing System with diamond lapping paper over a wide range of roughnesses (0.1-15 μm), before being attached onto a TEM grid. Subsequently, additional material is removed using a Gatan Dimple Grinder II with 0-4 μm diamond pastes and 10gr of weight. At the end of the dimpling process, each sample is around 10 μm thick. A series of PIPSII beam energies (1-5keV) is used on different specimens to create an electron-thin region for TEM imaging. Table 1 summarized these different PIPSII procedures.

Finally, the samples are imaged with an FEI Talos 200kV Microscope with a Super X-EDS detector. Figure 1 shows the STEM images for samples A, B, and C. Figure 2 includes EDS elemental 1D maps for In and Ga of sample B. The bright lines from the top to the bottom of images in Fig. 1 correspond to damage caused by PIPSII process. While sample A shows not only damage but also contamination from the sample preparation process, sample B only shows damages from the ion beam of PIPSII. Moreover, EDS line scans show that the nanostructures in samples B are not related to contamination. However, the origin on these unique structures is still not well understood. Sample C with milder ion polishing procedure shows cleaner images with neither damage nor contamination. The red circles in Figure 1c are electron beam damage from TEM during alignment process. Figure 1d is a zoom-in section of 1c. It confirms that the procedure used to make sample C is optimal for cross section InGaIn. No evidence for Indium segregation was found in this sample.

Despite the useful information from TEM including the growth dynamics and structural properties of the material, sample preparation is as crucial as TEM imaging. A good sample preparation process produces a precise and certain TEM characterization of the material. Due to the sensitivity of In atoms in the InGaIn thin film, a complex sample preparation process is needed to provide suitable samples for TEM. Using the traditional polishing process, we introduce a systematic way of providing the optimal ion polishing procedure. Our STEM images confirm the reduction of the ion beam damage through a series of samples, and insure we have an optimal recipe for cross section InGaIn sample preparation [6].

References:

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	A (5° angle)	B (5° angle)	C (4° angle)
Eye opens	5keV	3keV	3keV
Fringes	4keV	3keV	2keV
Hole opens	3keV	3keV	1keV
Final polish	1keV, 0.5keV, 0.25keV	1keV, 0.5keV, 0.25keV	0.5keV, 0.25keV

Table 1. Ion beam milling process using a Gatan PIPSII with different beam energy between samples A, B, and C.

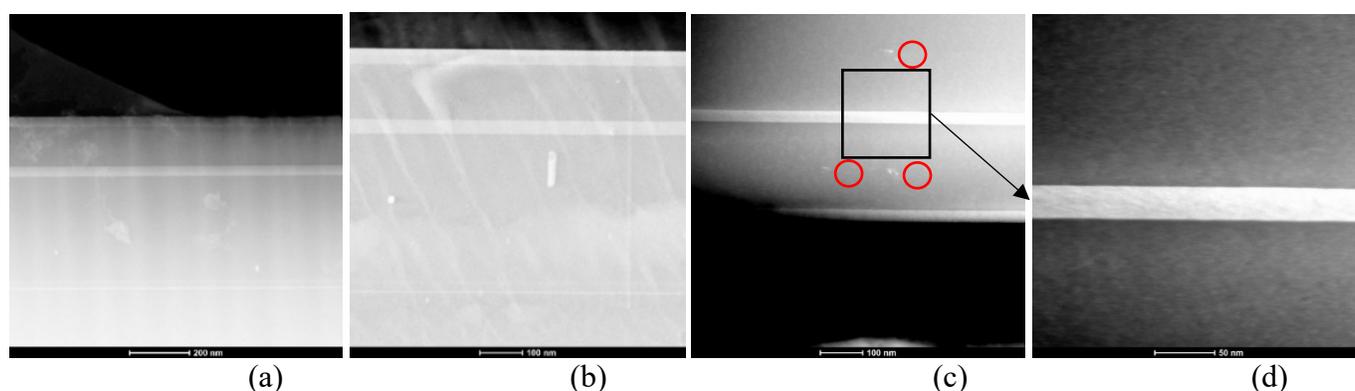


Figure 1. STEM images of sample A, (a), sample B, (b), and sample C, (c) and (d). The bright vertical/tilted lines are due to damage from the PIPSII process

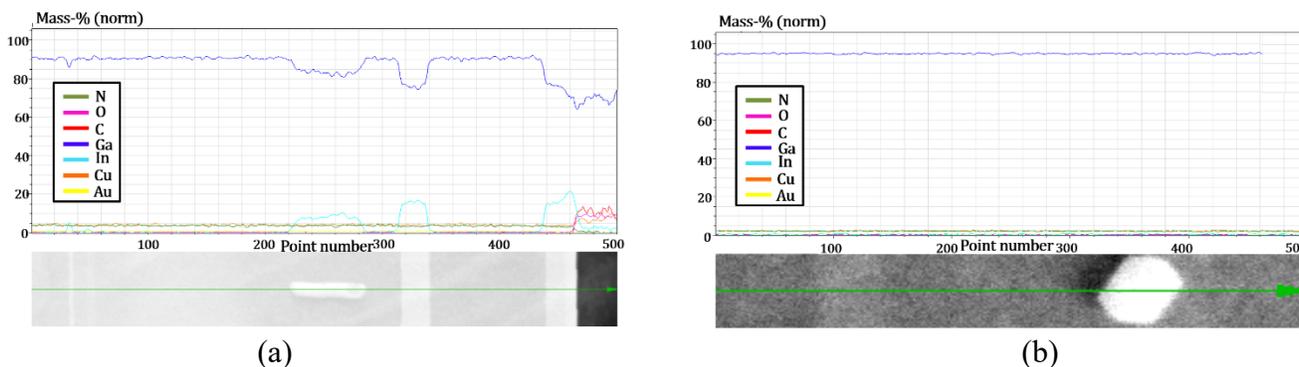


Figure 2. EDS elemental line scans for unique nanostructures in the GaN layer of sample B.