





#### Preparation and Study of Cross-Sectioned GaN HEMT Devices

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# Outline



- Motivation: Understanding Device Physics of Failure by Looking at Features Underneath the Gate
- How Cross-Sectional Samples are Prepared and How They Perform
- Brief Discussion of Scanning Probe Methods
- Results From Scanning Probe and Optical Studies



#### What Happens Beneath the Gate?







#### What Happens Beneath the Gate?





Delayered AlGaN/GaN surface after chemical removal of device structures and passivating layers.

# **Device Layout:**





Custom devices from several foundries have been designed with gate pads on one side so as to be operable after sectioning. Devices are produced at a commercial foundry.

Devices with and without source-connected field plates are available for test.

Gate is nominally ~1 $\mu$ m wide, channel is a few  $\mu$ m wide.

GaN thickness is a few  $\mu$ m.





### **Cross-Sectioning:**



- Devices are prepared with an initial mechanical polish followed by Ar ion milling.
- Resulting surfaces need to be as smooth as possible for scanning probe microscope experiments.
- Cross-sectioned devices need to perform similarly to intact devices.



#### **Cross-Sectioning: Mechanical Polishing**





Sample is polished with a series of SiC, diamond, and colloidal alumina polishing media.

Mechanical polishing with fine colloidal polishing media still leaves some polishing damage at the nanoscale.



# **Cross-Sectioning: Ion Milling**





Leica TIC (Triple Ion Cutter) three-beam wide area milling tool.

Sample is placed behind a mask and milled with 8keV Ar ion guns for several hours.



#### **Cross-Sectioning:**





Ion milling tends to redeposit material on the active region of the device, which can be protected using an easily-removable photoresist layer.



### **Cross-Sectioning:**





Cross-sectioned devices perform in a reasonably similar manner to intact devices



### **Cross-Section Mount:**



Will hold the device on edge for SPM or optical microscopy experiments, and allows for electrical operation and temperature control of the device.







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#### Photoemission Study: Predicted Hot-Spot Migration





Modeled channel heat generation and temperature under equivalent 10W power conditions.

E. Heller, S. Choi, D. Dorsey, R. Ventury, S. Graham, *Microelectronics Reliability* 53 (2013) 872-877.



### **Photoemission Study**





#### Photoemission map at 0.5mW power, 10V drain



### **Photoemission Study**





0.5mW power, 20V Vd, 25  $\mu$ A Id 0.5mW power, 80V Vd, 6.25  $\mu$ A Id Shift and change in size in photoemission "hot spot" with increasing drain bias







# Scanning Probe Microscopy (SPM)





In an atomic force microscopy (AFM) experiment, a sharp probe is scanned across a sample surface.

Probe deflection alters the path of the reflected laser beam, which is measured by a position-sensitive diode.

Topographic images with nanometer-scale resolution can be recorded.

From: http://en.wikipedia.org/wiki/File:Atomic\_force\_microscope\_block\_diagram.svg





With metal-coated probes, an electric potential can be applied during imaging, and local electric properties can be mapped along with topography.

A wealth of SPM techniques are available, including

- Current-Sensing AFM
- Electric Force Microscopy
- Kelvin Probe Microscopy
- Magnetic Force Microscopy
- Scanning Tunneling Microscopy
- Scanning Capacitance Microscopy
- Scanning Microwave Microscopy
- Scanning Thermal Microscopy







For a conducting tip having a DC potential separated a short distance from the surface, the electrostatic force on the tip is

$$F_{electrostatic} = \frac{1}{2} \frac{\partial C}{\partial z} \Delta V^2$$

Which is sensitive to local surface charge.



# Electrostatic Force Microscopy (EFM) and Kelvin Probe Force Microscopy (KPFM or KFM)



For a conducting tip having a DC potential *and* an AC potential positioned above a surface with surface potential  $V_{CPD}$ ,

$$V = (V_{DC} - V_{CPD}) + V_{AC} \cdot \sin(\omega t)$$

As before, the electrostatic force on the tip is  $F = \frac{1}{2} \frac{dC}{dz} V^2$ , so....

$$F = \frac{dC}{dz} \left[\frac{1}{2}(V_{DC} - V_{CPD})^2 + \frac{1}{4}V_{AC}^2\right] + \frac{dC}{dz}[V_{DC} - V_{CPD}]V_{AC}\sin(\omega t) - \frac{1}{4}\frac{dC}{dz}V_{AC}^2\cos(2\omega t)$$

With a lock-in detector at  $\omega$  and a little math this simplifies to,

$$F_{\omega} = \frac{dC}{dz} [V_{DC} - V_{CPD}] V_{AC} \sin(\omega t)$$

In KPFM, a feedback mechanism adjusts  $V_{DC}$  such that  $F_{\omega} = 0$ .

When this happens,  $V_{DC}$  = a measurement the local surface potential.



#### Surface Topography

#### Kelvin Probe Surface Potential Map (Unbiased)





Surface Topography



Surface Potential (unbiased)



#### Surface Potential -2 Vg, 0 Vd

#### Surface Potential -4 Vg, 5 Vd







Trapped charges beneath the gate are believed to give rise to a "virtual gate," temporarily altering device behavior.



Reduced drain current after semi-on stress, which recovers after exposure to light.



### **Time-Varying Surface Potential Maps**





Charge buildup dissipates over time, and device is restored to initial conditions after ~1 day Similar results observed for hot-carrier stress conditions. DISTRIBUTION A. Approved for public release; distribution unlimited.



# **Raman Spectroscopy**



Vibrational spectroscopy to assess molecular movement, assess structure, temperature, uniformity, mechanical stress, thickness

- Inelastic scattering of a monochromatic excitation source
- Incident radiation interacts with phonons (lattice vibrations) resulting in an energy shift of the incident light.
- Incident photons excite vibrational modes in the sample, yielding scattered photons with diminished energy (Stokes scattering)
- Or, incident light interacts with phonons at a raised energy level, scattering photons at frequencies above the incident frequency (anti-Stokes scattering)







- Quantify surface charge
- Measure the kinetics and distribution of charge traps
- Study devices that have been degraded through electrical/thermal stress
- Use cross-sectional measurements to validate modeling results

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