

Abstract Submitted  
for the MAR12 Meeting of  
The American Physical Society

**Controlled electrochemical functionalization of epitaxial graphene**<sup>1</sup> CALVIN CHAN, THOMAS BEECHEM, TAISUKE OHTA, DAVID WHEELER, Sandia National Laboratories, KEITH STEVENSON, University of Texas, Austin — Chemical functionalization is a promising means of modifying graphene for applications ranging from nanoelectronics to transparent electrodes. Various schemes have been demonstrated, but control over functionalization density with well-specified molecules is still a challenge. We report on the controlled electrochemical functionalization of epitaxial graphene with trifluoromethylphenylene ( $\text{CF}_3\text{Ph}$ ), where the functionalization density was controlled by the electron injection rate.  $\text{CF}_3\text{Ph}$  peaks were observed in x-ray photoemission spectroscopy, along with binding energy shifts consistent with bonding between  $\text{CF}_3\text{Ph}$  and graphene. A maximum functionalization density of one molecule per six graphene carbons was inferred from the peak intensities. Spectroscopic Raman mapping revealed increasing graphene D:G peak intensity ratios that scaled with increasing functionalization-induced localized defects. While a minimal shift in the  $\pi$  orbital structure and the emergence of  $\text{CF}_3\text{Ph}$  related features were observed in ultraviolet photoemission spectroscopy, a work function increase by 0.5 eV in  $\text{CF}_3\text{Ph}$ -graphene suggests a shift of the electron distribution towards the  $\text{CF}_3$  moieties on the surface. This work has positive implications for transparent electrode applications.

<sup>1</sup>This work was supported by the Early Career LDRD program at SNL, and the DOE EFRC in Charge Separation and Transfer (DE-Calvin Chan SC0001091). SNL is operated by Sandia Corp., a Lockheed Martin Co. Laboratory for the US DOE-NNSA (DE-AC04-94AL85000).

Date submitted: 11 Nov 2011

Electronic form version 1.4