

# **2015 MRS Fall Meeting & Exhibit, November 29-December 4, 2015, Boston Massachusetts USA**

## **L6.14**

### **The Extended Core Coax: A Novel Nanoarchitecture for Electrochemical Detection of Infectious Disease Biomarkers**

Amy Valera<sup>1</sup>, Michelle Archibald<sup>1</sup>, Jeff Naughton<sup>1</sup>, Michael Burns<sup>1</sup>, Michael J Naughton<sup>1</sup>, Thomas C Chiles<sup>1</sup>

<sup>1</sup>Boston College, Chestnut Hill, Massachusetts, United States.

Highly specific and sensitive platforms for detection of clinically relevant biomarkers are critical for accurate disease diagnosis. Pathogens such as *Vibrio cholerae* continue to cause significant mortality in resource-limited areas, where low cost, point-of-care (POC) diagnosis is ideal. While standard tools such as an enzyme linked immunosorbant assay (ELISA) meet diagnostic specificity and sensitivity needs, they cannot be utilized outside a clinical setting, at the site of the patient. To fill this unmet need for specific and sensitive disease detection with POC accessibility, we propose to use a novel nanoarchitecture for electrochemical sensing, the extended core coax (ECC). Each ECC is a vertically oriented nanocoax comprised of an extended inner metal core and an outer metal shield, separated by a dielectric annulus. The inner core, comprised of gold, acts as a working electrode which extends ~200 nm above the chrome counter electrode. Arrays with a base area of ~2000  $\mu\text{m}^2$  each contain ~2000 individual ECCs connected in parallel. The extended gold core provides a potential substrate for molecular imprinting of proteins, making the ECC an attractive candidate for development as a biosensor for electrochemical detection of infectious disease biomarkers such as cholera toxin..