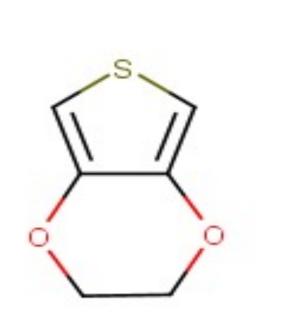


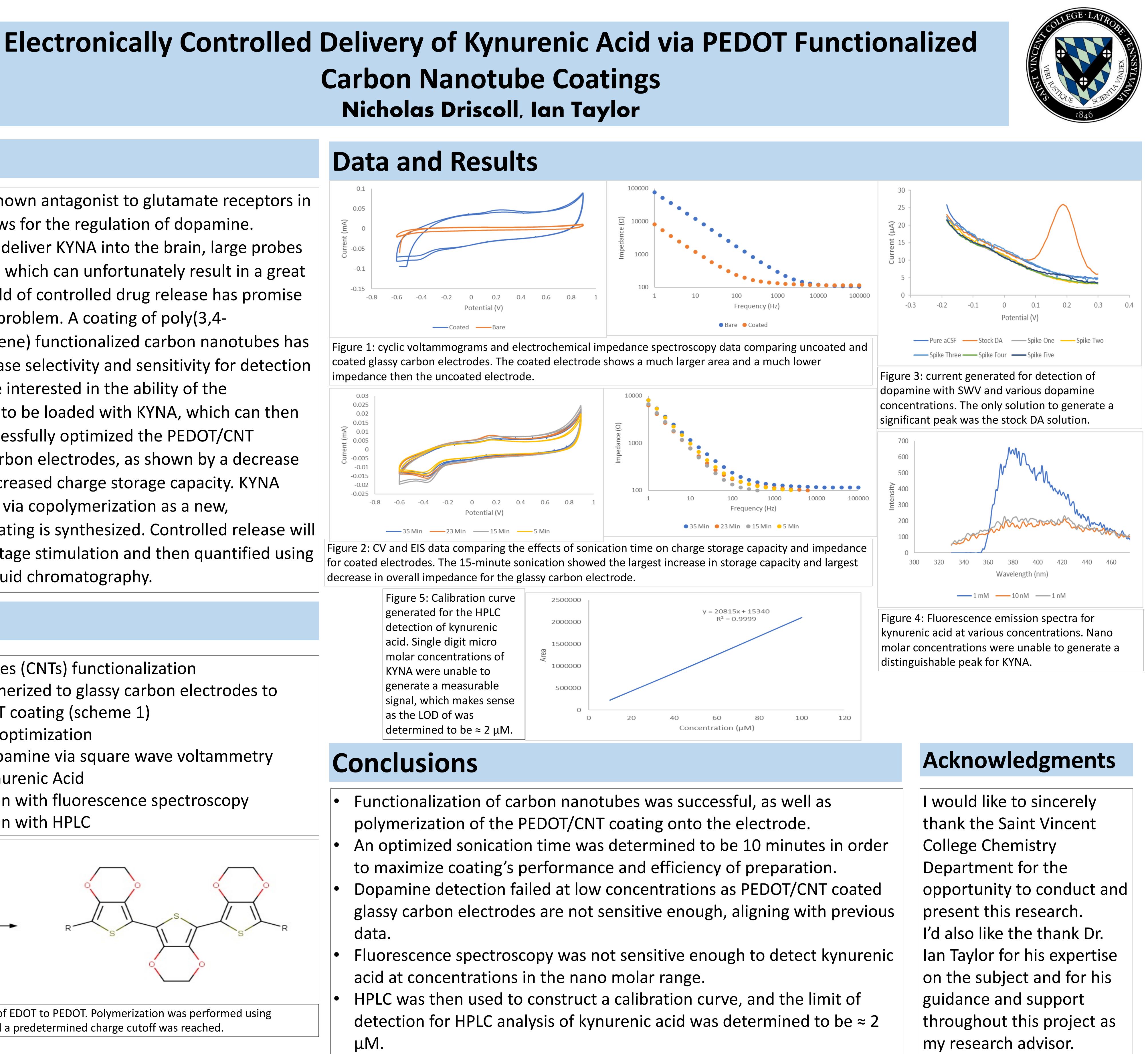
## Abstract

Kynurenic Acid is a known antagonist to glutamate receptors in the brain, which allows for the regulation of dopamine. However, in order to deliver KYNA into the brain, large probes have to be employed which can unfortunately result in a great deal of harm. The field of controlled drug release has promise as an answer to this problem. A coating of poly(3,4ethylenedioxythiophene) functionalized carbon nanotubes has been shown to increase selectivity and sensitivity for detection of resting DA. We are interested in the ability of the PEDOT/CNT coatings to be loaded with KYNA, which can then be released. We successfully optimized the PEDOT/CNT coatings on glassy carbon electrodes, as shown by a decrease in impedance and increased charge storage capacity. KYNA loading is performed via copolymerization as a new, PEDOT/CNT-KYNA coating is synthesized. Controlled release will be performed via voltage stimulation and then quantified using high performance liquid chromatography.

# Methods

- Carbon nanotubes (CNTs) functionalization EDOT/CNT polymerized to glassy carbon electrodes to form PEDOT/CNT coating (scheme 1)
- Sonication time optimization
- Detection of dopamine via square wave voltammetry Detection of Kynurenic Acid
  - Detection with fluorescence spectroscopy Detection with HPLC





Scheme 1: The polymerization of EDOT to PEDOT. Polymerization was performed using chronocoulometry at 0.9 V until a predetermined charge cutoff was reached.

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