### Improving Fluorescence Emission of Graphene Quantum Dots through Surfactant-Assisted Stabilization The Michael and Sally McCracken Annual Student Research Symposium Mason McClure, Judah Crawford, Anton V. Naumov

## ABSIRACT

Graphene Quantum Dots (GQDs) are nanoscale carbon based graphene sheets that exhibit unique fluorescent properties throughout a wide range wavelengths. Given their uniquely small size, low toxicity, biocompatil ity, and fluorescent capabilities, GQDs have many unique and important roles. To name a few, GQDs are used in drug delivery, fluorescent image and biosensing thanks to their unique ability to fluoresce under different wavelengths of light. Furthermore, there are different types of GQDs w their own unique properties. Knowing this, five amphipathic molecules called surfactants, were added to two different types of GQDs to test if would impact the resulting fluorescence. Furthermore, concentrations of these added surfactants were varied to test how different concentrations given surfactant might affect the fluorescence for a given GQD. We ob served that some of these surfactants provided a beneficial boost to GQ fluorescence, while others slightly inhibited the fluorescence. Moreove saw that the increase in fluorescence varied based on the concentration surfactant added yielding lower fluorescence for extremely low and high concentrations, while increasing the fluorescence at a more moderate c centration.

# INTRODUCTION

Graphene Quantum Dots (GQDs) fluoresce (glow) under specific way lengths of light.

This study focuses on two types of GQDs:

•NGQDs (nitrogen-doped): graphene sheets with nitrogen and oxyge containing groups.

•rGQDs (reduced): similar structure, but without nitrogen—only gra phene and oxygen.

A third GQD, Nd-NGQDs (neodymium-doped), will be tested in futur work; it glows in the near-infrared (NIR). **Goal:** Understand how different surfactants affect the fluorescence of

NGQDs and rGQDs.

**Hypothesis**: Surfactants influence fluorescence by interacting with G hydrophobic ends face the GQDs, and hydrophilic ends face water, en hancing light emission. Surfactants tested:

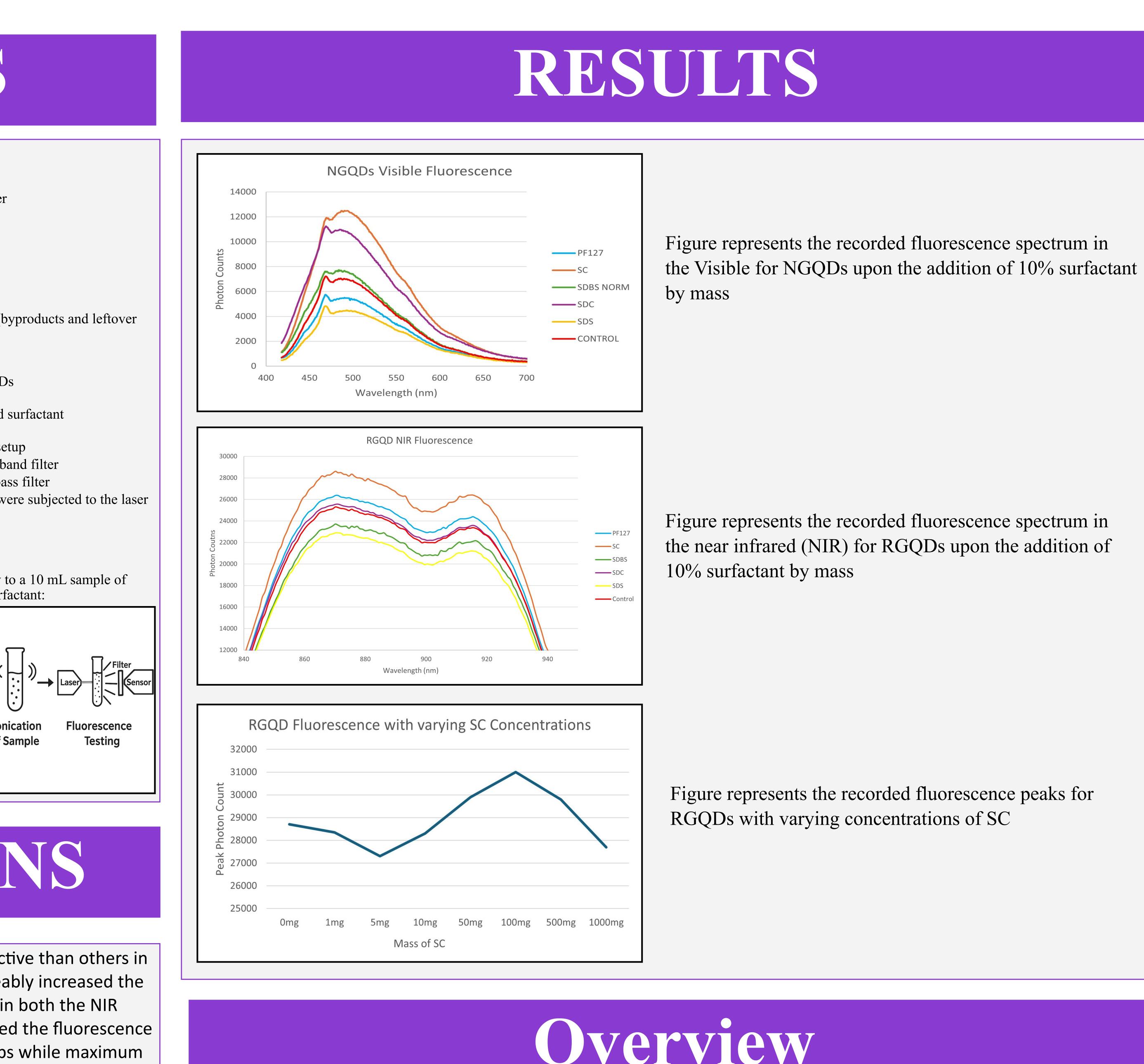
- •Sodium dodecyl sulfate (SDS)
- •Sodium deoxycholate (SDC)
- •Sodium dodecylbenzene sulfonate (SDBS)
- •Pluronic F127 (P-F127)
- •Sodium cholate (SC)

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	METHODS
e ge of bil- nt ging, nt vith s,	<ul> <li>Synthesis</li> <li>Synthesis of GQDs</li> <li>Batches of NGQDs were synthesized through a microwave synthesis process</li> <li>2.156g of glucosamine was added to a 500mL flask containing 250mL of DI water</li> <li>Flask was placed in microwave at power level three for one hour</li> <li>Synthesis of rGQDs</li> <li>Batches of rGQDs were synthesized through a microwave synthesis process</li> <li>5g of reduced graphene oxide dissolved in a 20mL aliquot of DI water</li> <li>1 mL of Sodium Hypochlorite is added to the solution</li> <li>Solution is stored in darkness and allowed to sit for 48 hours</li> </ul>
f they of is of a o- QDs er, we	<ul> <li>Both NGQD and rGQD batches were dialyzed and filtered to remove impurities (by reactants)</li> <li>Surfactant Test         Addition of surfactant         10% surfactant by weight was added to a 10mL aliquot of both NGQDs and rGQD         Sonication         Individual aliquots were sonicated to ensure normalized distribution of GQDs and so Testing     </li> </ul>
n of gh con-	<ul> <li>4mL samples were pipetted into cuvettes which were placed in our fluorescence set</li> <li>NGQD samples were configured with a 400 nm laser with a corresponding broadba</li> <li>rGQD samples were configured with an 808 nm laser with a corresponding longpas</li> <li>NGQD samples were subjected to the laser for one second while rGQD samples were for 15 seconds</li> <li>Data was recorded via an Avantes spectrophotometer</li> </ul>
	<ul> <li>Addition of surfactant</li> <li>Starting with the control with no surfactant, surfactants were added incrementally t rGQDs which were then tested for peak fluorescence at the following masses of surfactant = 1 mg, 5 mg, 10 mg, 50 mg, 100 mg, 500 mg, and 1000 mg</li> <li>Sonication</li> <li>Sonicated in between each individual test in the same manner as the surfactant test</li> <li>Testing</li> </ul>
ve- en-	<ul> <li>4mL samples were pipetted into cuvettes which were placed in our fluorescence setup with an 808 nm laser and corresponding longpass filter</li> <li>Samples were subjected to the laser for 15 seconds</li> <li>Data was recorded via an Avantes spectrophotometer</li> </ul>
a- re	CONCLUSIO
QDs 1-	Our experiment showed that some surfactants were more effect enhancing fluorescence. Specifically, SC, SDC, and PF127 noticea fluorescence of rGQDs in the NIR, with SC performing the best in rGQD and visible NGQD tests. At low concentrations, SC inhibite of rGQDs, likely due to interactions with RGQD functional groups fluorescence was observed at ~10% by mass. We theorize these to the hydrophobic interactions between the GQDs and the surface

e results were due factant's nonpolar regions while hydrophilic parts interacted with water, reducing inhibition. Similar trends were observed with SDC and PF127, which we also theorize enhanced fluorescence through hydrophilic interactions. However, less effective surfactants appeared to suppress fluorescence, likely due to unfavorable interactions with GQDs that diminished their inherent fluorescence, similar to what we saw with SC at low and very high concentrations. Future work will expand surfactant testing to Nd-GQDs and investigate concentration effects on both NGQDs and Nd-GQDs.





We studied how soap-like molecules (surfactants) affect the glow (fluorescence) of tiny carbon particles called graphene quantum dots (GQDs). These dots can be used in things like medical imaging or sensors. We found that different surfactants can either make the dots glow more or less depending on how they interact. Understanding this helps us control the glow, which is important for future uses in science and technology.



