



Abstract

Cell visualization through cell labeling has previously been an effective way to learn about the cell and cell biology. Small-molecule fluorescent dyes are used for cell imaging, especially the RNA. This method of binding fluorescent dyes to the cell is less expensive compared to other methods such as oligonucleotide-based probes or protein-based reporters. Although using RNA dyes are less expensive, it it still not practical due to qualities such as prone to scattering and having low photostability. Styrene dyes however, will be more stable and give more fluorescence, increasing the application of using RNA dyes. With this improved fluorescent dye, visualizing RNA will be easier and can lead to further research and understanding of RNA biology

Objectives

By using fluorogenic styrene dyes and binding them to RNA and DNA, it makes it easier for cell imaging and visualization of the cell. Our objective is to create 2 new styrene dyes that will give off the most amount of light emission under fluorescent wavelengths (250-700 nm), and then binding it to the cell for cell imaging.

Purification Methods

There are different methods of purifying and extracting the products after the reaction:

• If there are no waste chemicals then filtration is used. Filtration uses a funnel with filter paper to separate the liquid and solid. • If there are unwanted chemical waste in the product, then Thin Layer Chromatography (TLC) and Column Chromatography should be used. Column chromatography uses the polarity of the chemicals to separate the product and waste.

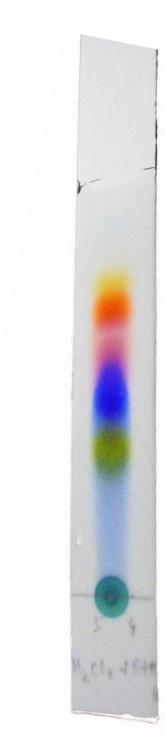


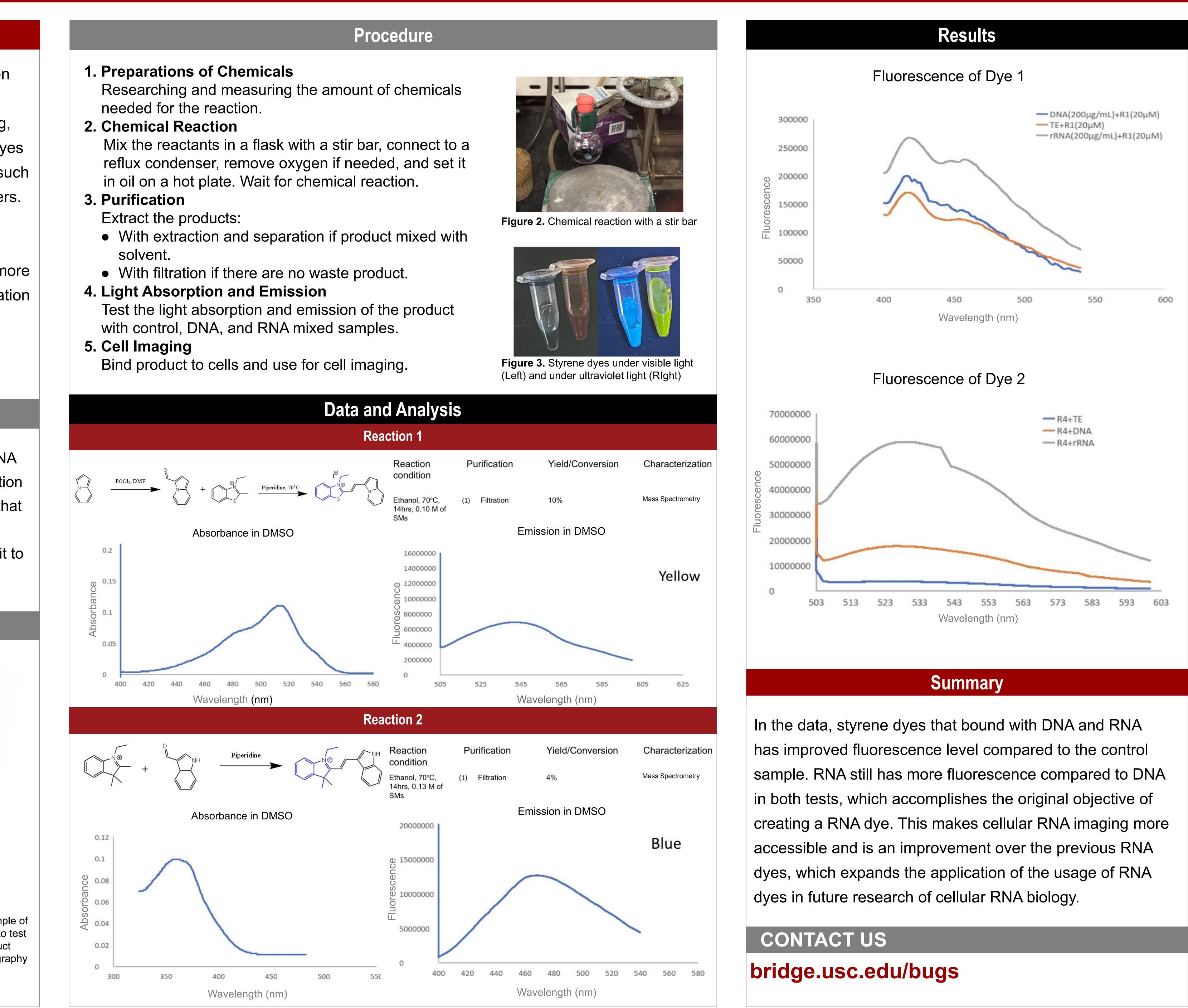
Figure 1. This is an example of TLC, which can be used to test the formation of the product before column chromatography

Development of Fluorogenic Styrene Probes for Cell Imaging

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