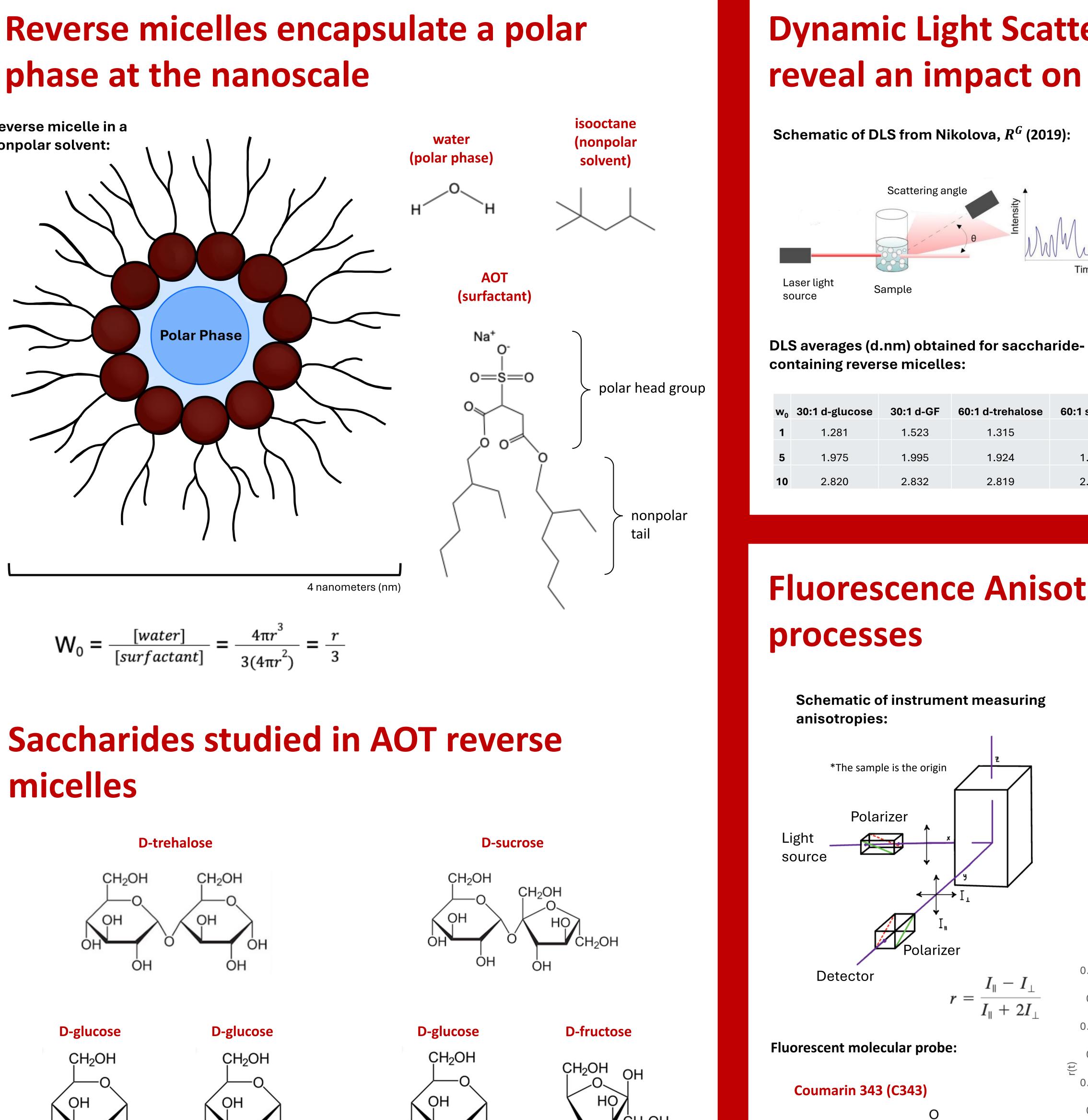
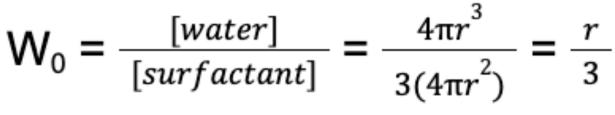
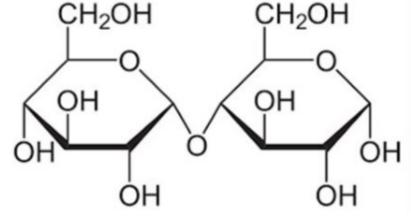


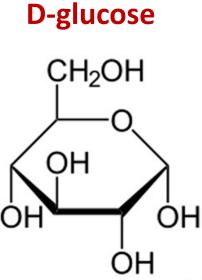
Summer 2024

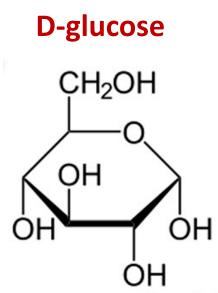
Reverse micelle in a nonpolar solvent:

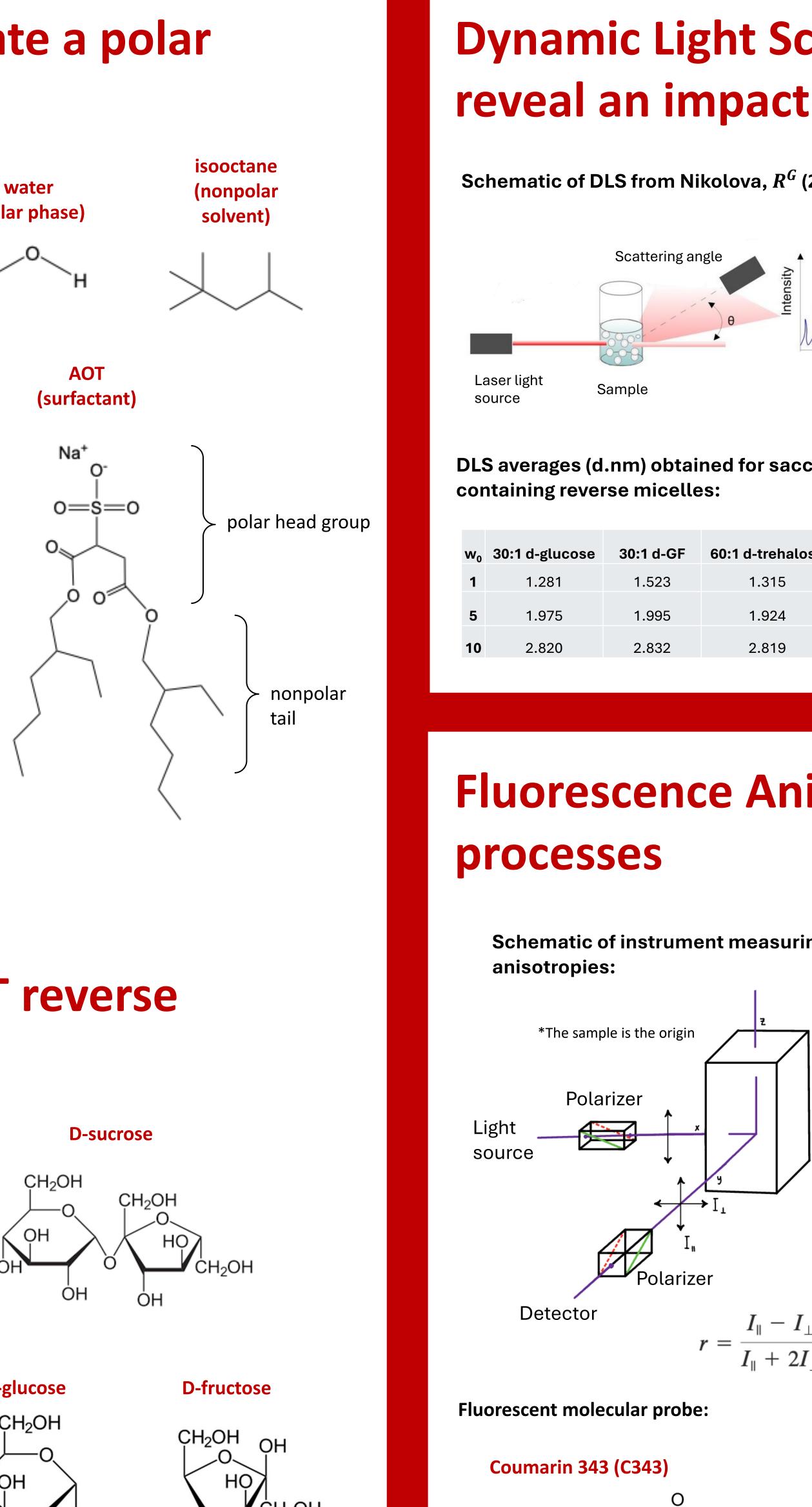


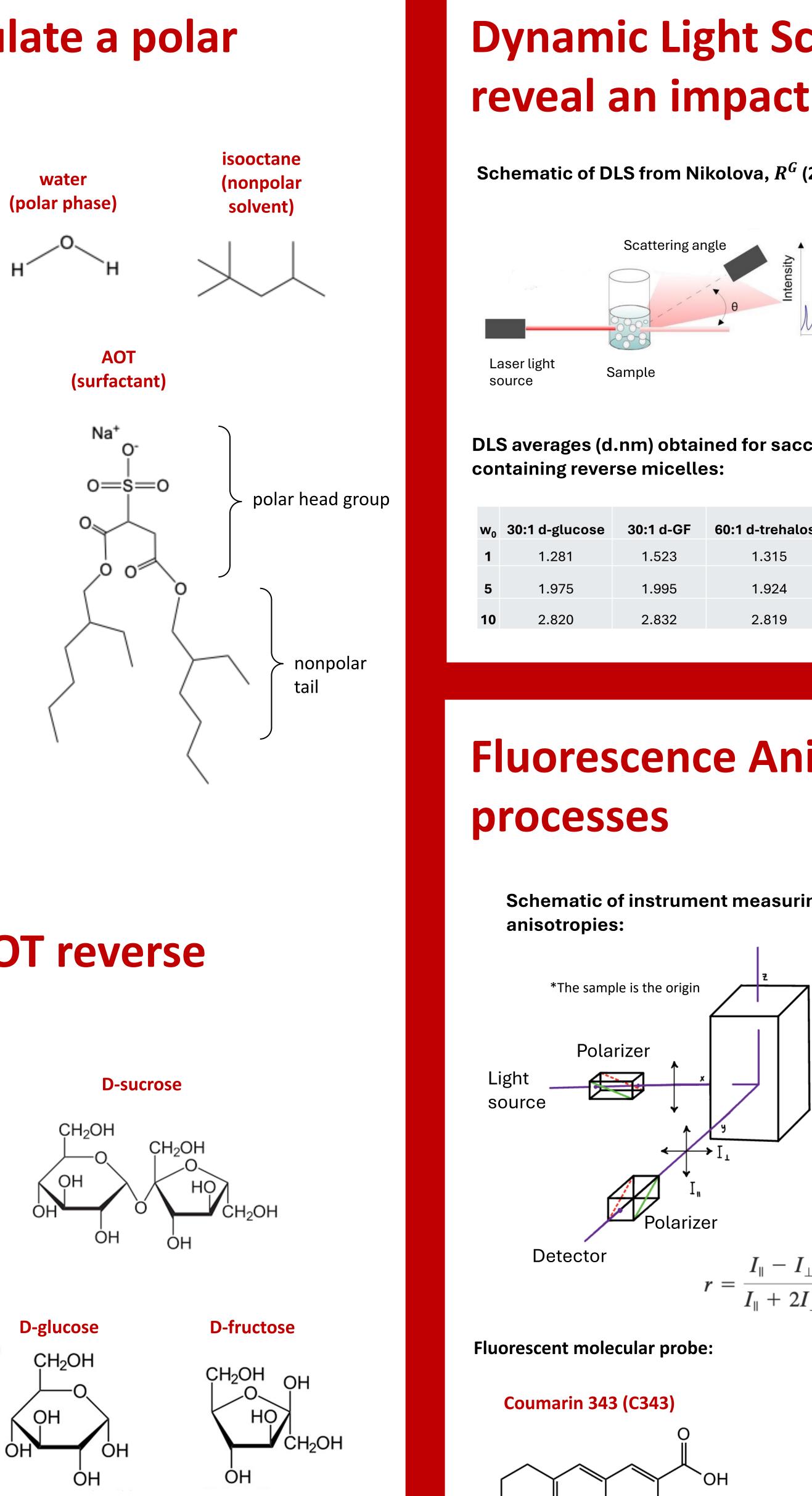












Denoted as "D-GF" in tables and figures

Comparison of two disaccharides and their saccharide components in AOT reverse micelles: Impacts of D-trehalose and D-sucrose

Delaney Collier ¹, Nancy E. Levinger ², Ph.D., and Bridget L. Gourley ¹, Ph.D.

¹ DePauw University, Department of Chemistry and Biochemistry, Greencastle, IN ² Colorado State University, Department of Chemistry, Fort Collins, CO

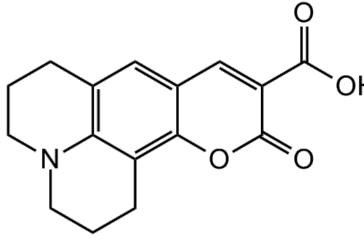
Dynamic Light Scattering (DLS) measurements reveal an impact on reverse micelle size

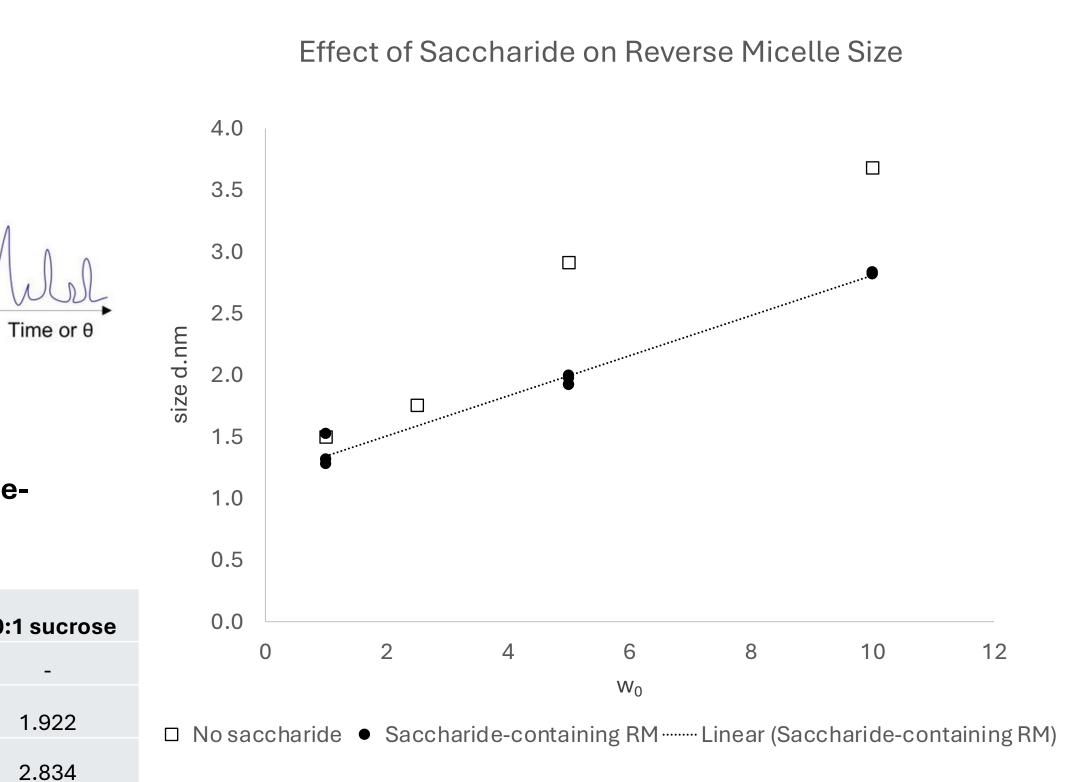
Schematic of DLS from Nikolova, R^G (2019):

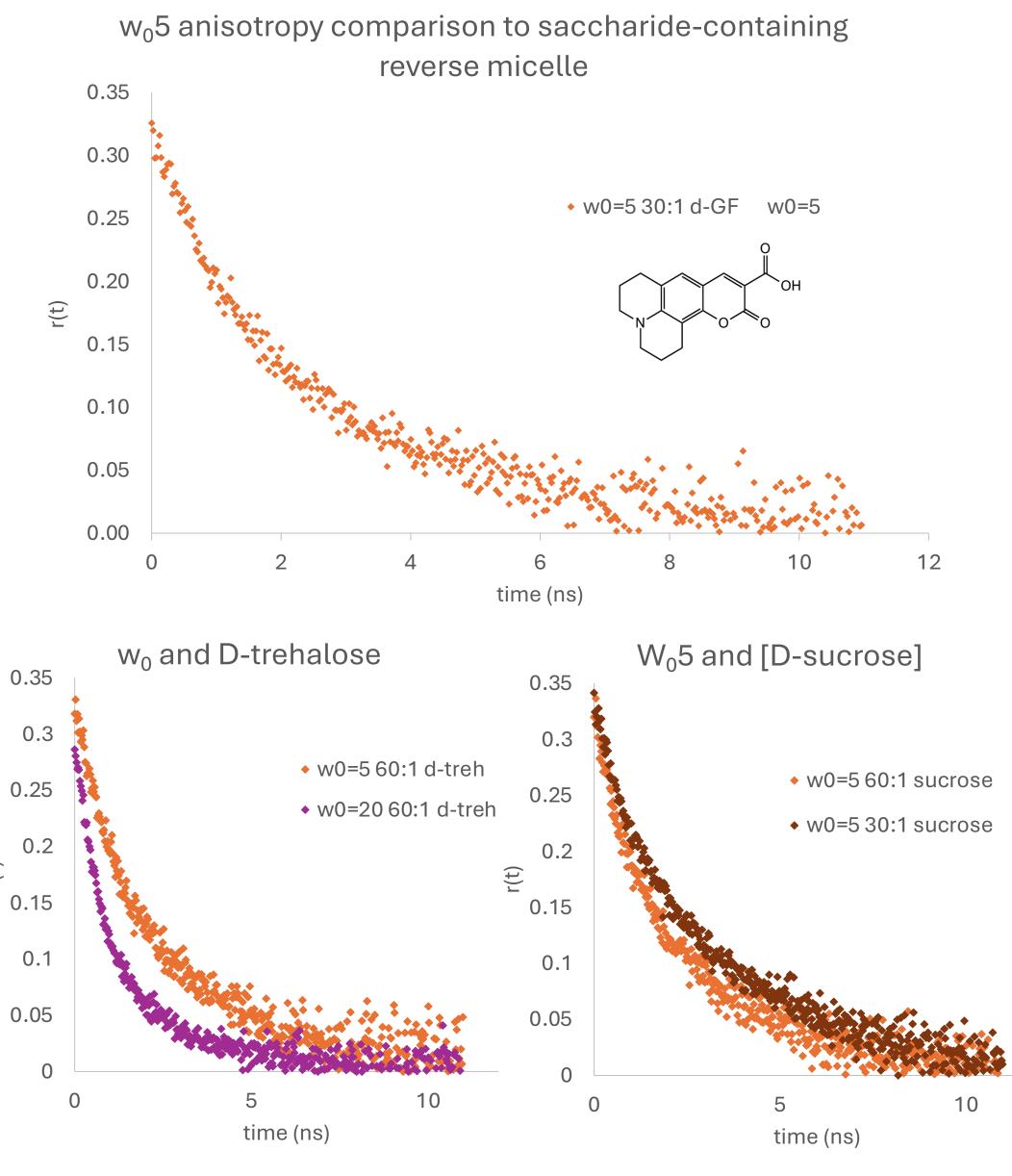
w _o	30:1 d-glucose	30:1 d-GF	60:1 d-trehalose	60:
1	1.281	1.523	1.315	
5	1.975	1.995	1.924	
10	2.820	2.832	2.819	

Fluorescence Anisotropy reveals slower relaxation w₀5 anisotropy comparison to saccharide-containing

Schematic of instrument measuring









Reverse Micelles encapsulate a polar phase at the nanoscale

- In a nonpolar solvent, the use of a surfactant (AOT) allows for water to be encapsulated at the nanoscale
- w₀ represents a ratio of water to surfactant, and we use this parameter in relation to size

Saccharides studied in AOT reverse micelles

- Prepared by sonication with 0.1 M AOT, 30:1 monosaccharide solutions, and 60:1 disaccharide solutions for at least 10 minutes
- Stirring and heating at 60 degrees C was applied for no more than 3 minutes for select samples in addition to sonication for 10 minutes

Dynamic Light Scattering measurements reveal an impact on reverse micelle size

- DLS determines the size of the reverse micelles for each ratio of water to surfactant (w_0)
- Plotted is a comparison of averages of newly reported saccharide RM size to RM with no saccharide. The trendline for saccharide RMs reveals that saccharides do not only make reverse micelles smaller, but they also impact reverse micelles differently as w0 increases

Fluorescence Anisotropy reveals slower relaxation processes

- Fluorescence anisotropy uses polarized excitation and emission to measure the rotational dynamics of a fluorescent molecule
- Plotted are 3 modes of analysis: saccharide RMs vs water only RMs, varying w₀, and varying saccharide concentration
- The top plot shows that rotational dynamics are slower in saccharide RMs
- The lower plots show that smaller reverse micelles and higher concentrations of reverse micelles result in slower rotational dynamics

Acknowledgements

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