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Please use this identifier to cite or link to this item: http://hdl.handle.net/123456789/2153 Title: How to study picosecond solvation dynamics using fluorescent probes with small Stokes shifts Authors: Silori, Y. (/jspui/browse?type=author&value=Silori%2C+Y.) Dey, Shivalee (/jspui/browse?type=author&value=Dey%2C+Shivalee) De, A.K. (/jspui/browse?type=author&value=De%2C+A.K.) Solvation dynamics Keywords: Time-correlated single photon counting Xanthene dyes Issue Date: 2018 Publisher: Elsevier B.V. Citation: Chemical Physics Letters, 693, pp. 222-226 Xanthene dyes have wide ranging applications as fluorescent probes in analytical, biochemical and Abstract: medical contexts. Being cationic/anionic in nature, the solvation dynamics of xanthene dyes confined within a negatively/positively charged interface are very interesting. Unfortunately, the floppy structure and small Stokes shift render any xanthene dye unsuitable for use as a solvation probe. Using di-sodium fluorescein, we present our work on the picosecond solvation dynamics of bulk and confined water (at pH = 9.2). We also propose a new methodology for studying picosecond solvation dynamics using any fluorescent dye with a small Stokes shift. We discuss how scattering contributions can be effectively removed, and propose an alternative way of

defining zero time of solvation. Finally, we demonstrate the tuning location of the probe within confinement.

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