

Synthesis of nitric oxide probes with fluorescence lifetime sensitivity

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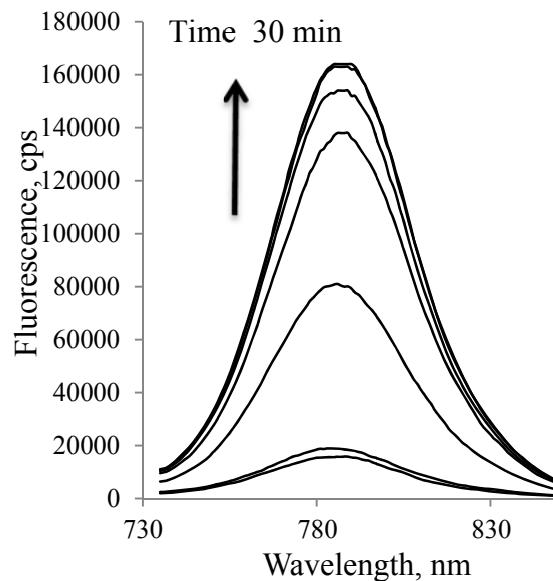
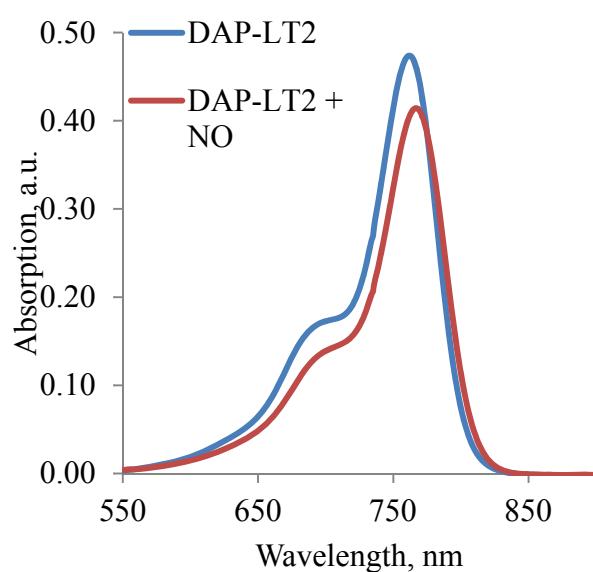


Figure S1 Absorption and emission spectra of DAP-LT2 before and after treatment with DEA NONOate

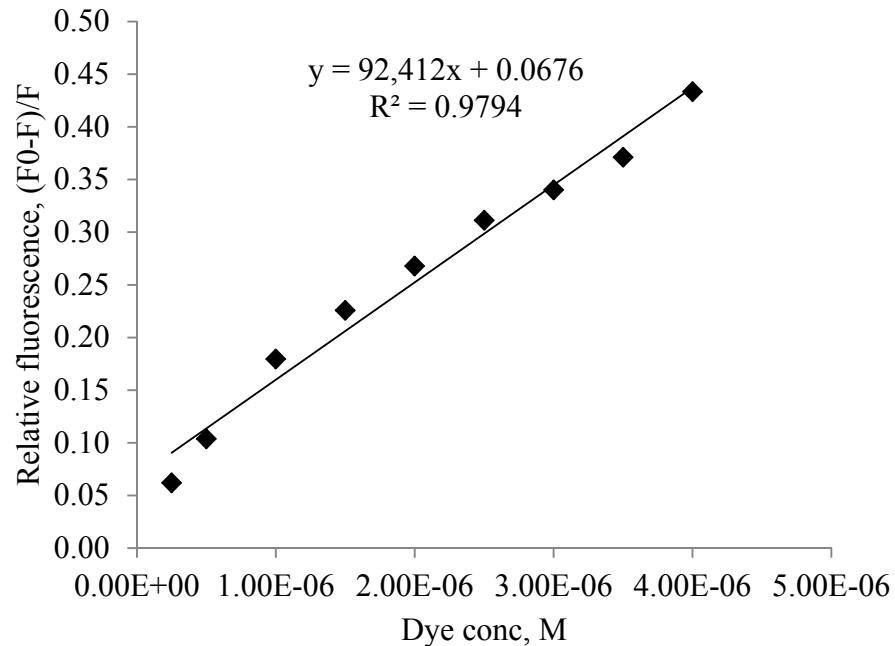


Figure S2 Binding constant of DAP-LT1 to albumin

Ex/em: 279/345 nm, BSA concentration 1.4×10^{-6} M in PBS. The slope of the trend lines corresponds to the binding constants for DAP-LT1 $\sim 92,400 \text{ M}^{-1}$.

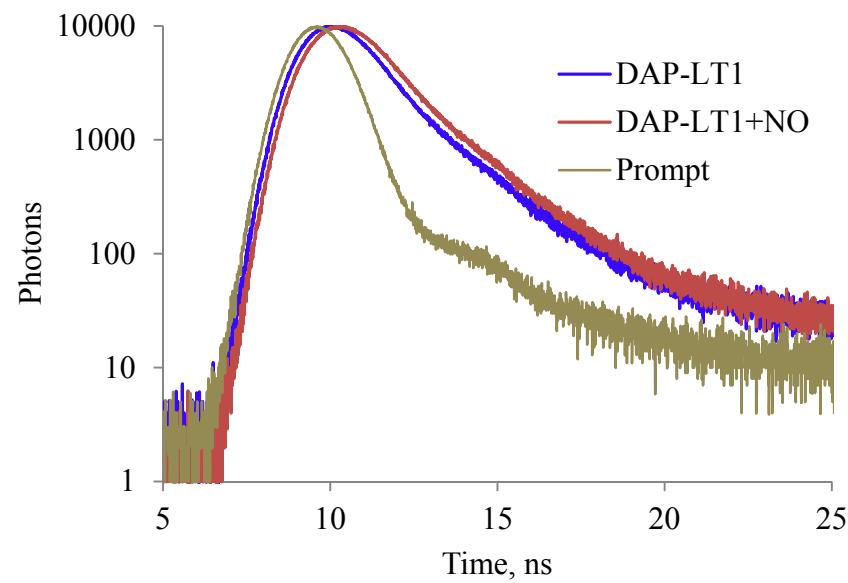


Figure S3 Fluorescence lifetime decays of DAP-LT1 before and after treatment with DEA NONOate

Ex/em 740/790 nm. 10% serum/PBS buffer

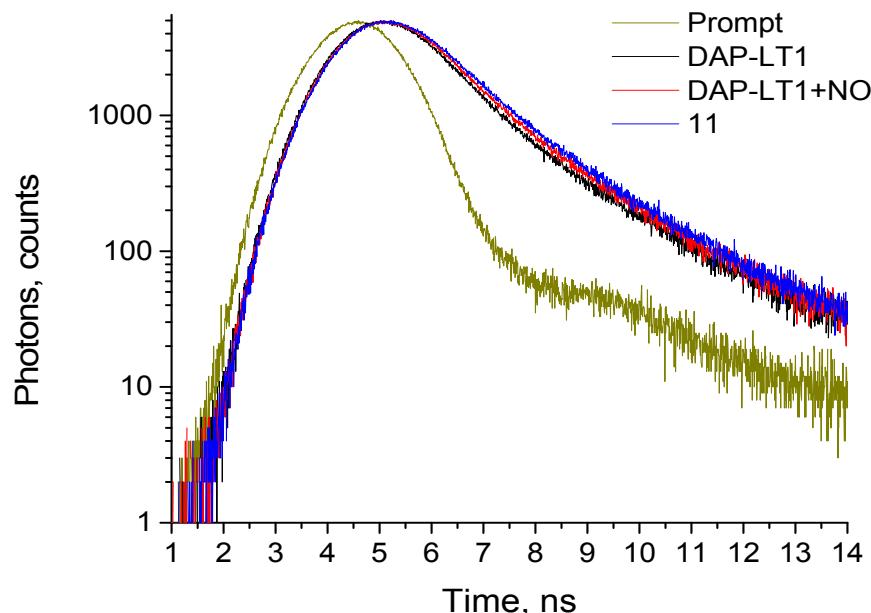


Figure S4 Fluorescence lifetime decays of DAP-LT1 before and 5 min after treatment with DEA NONOate and the decays of entry 11. All at 37 °C

Ex/em 740/790 nm. 10% serum/PBS buffer. Mean lifetime: DAP-LT1 ~ 0.94 ns, DAP-LT1+NO donor (5 min) ~ 1.02 ns and entry 11 ~ 1.09 ns.

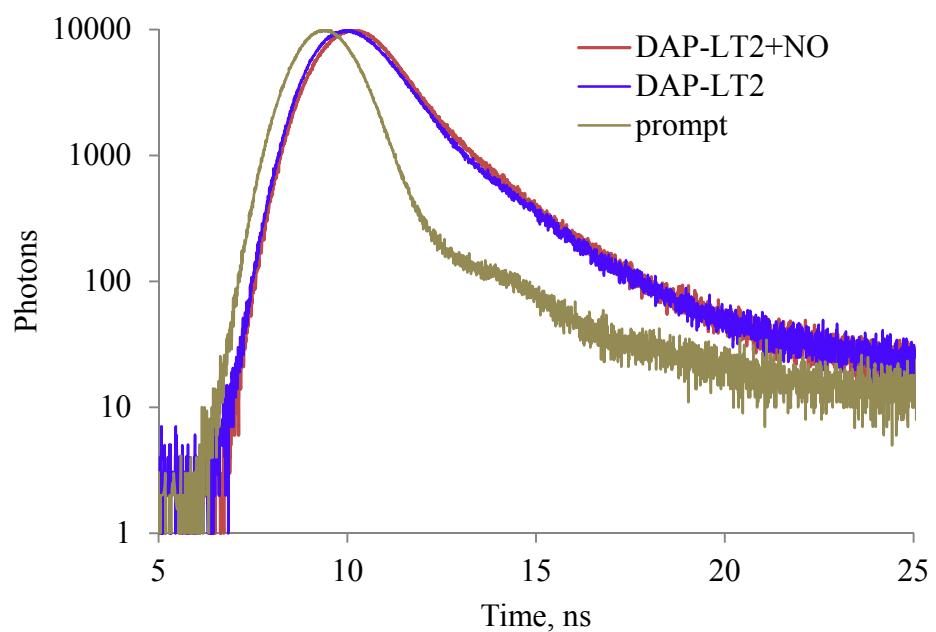
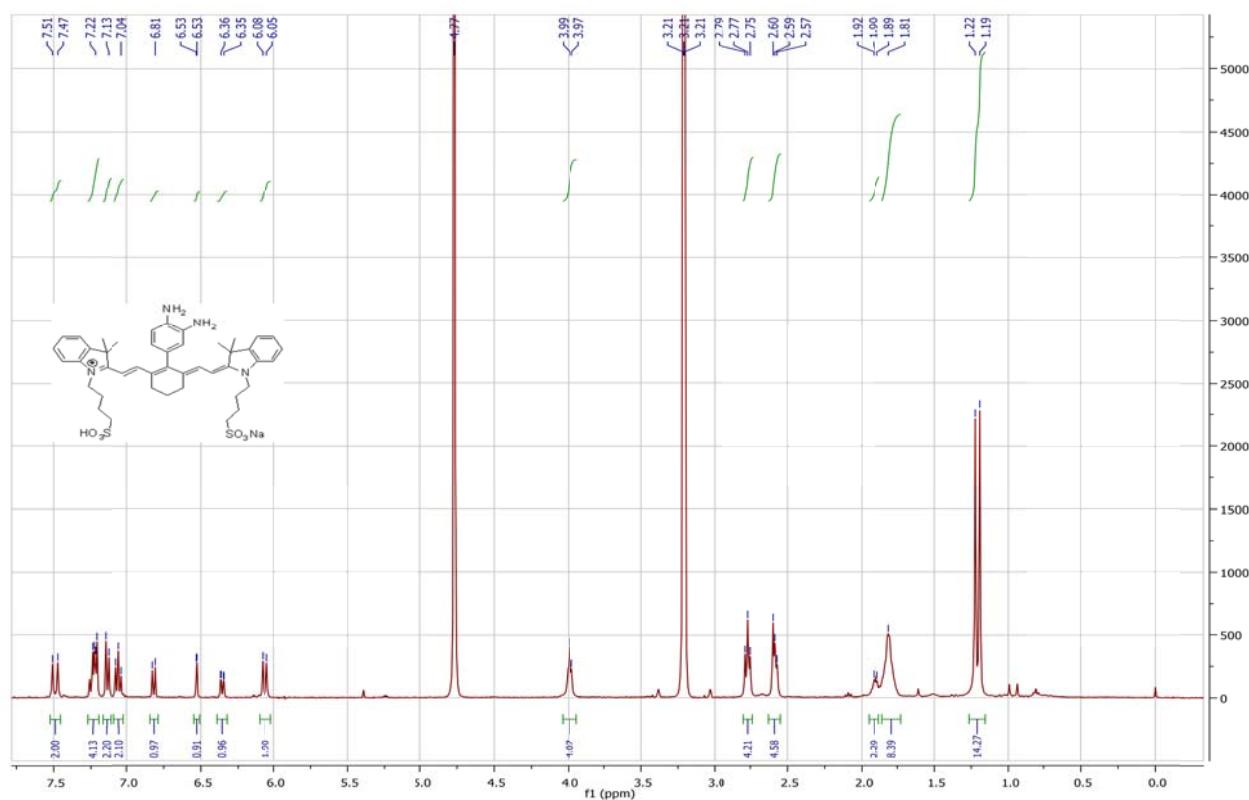


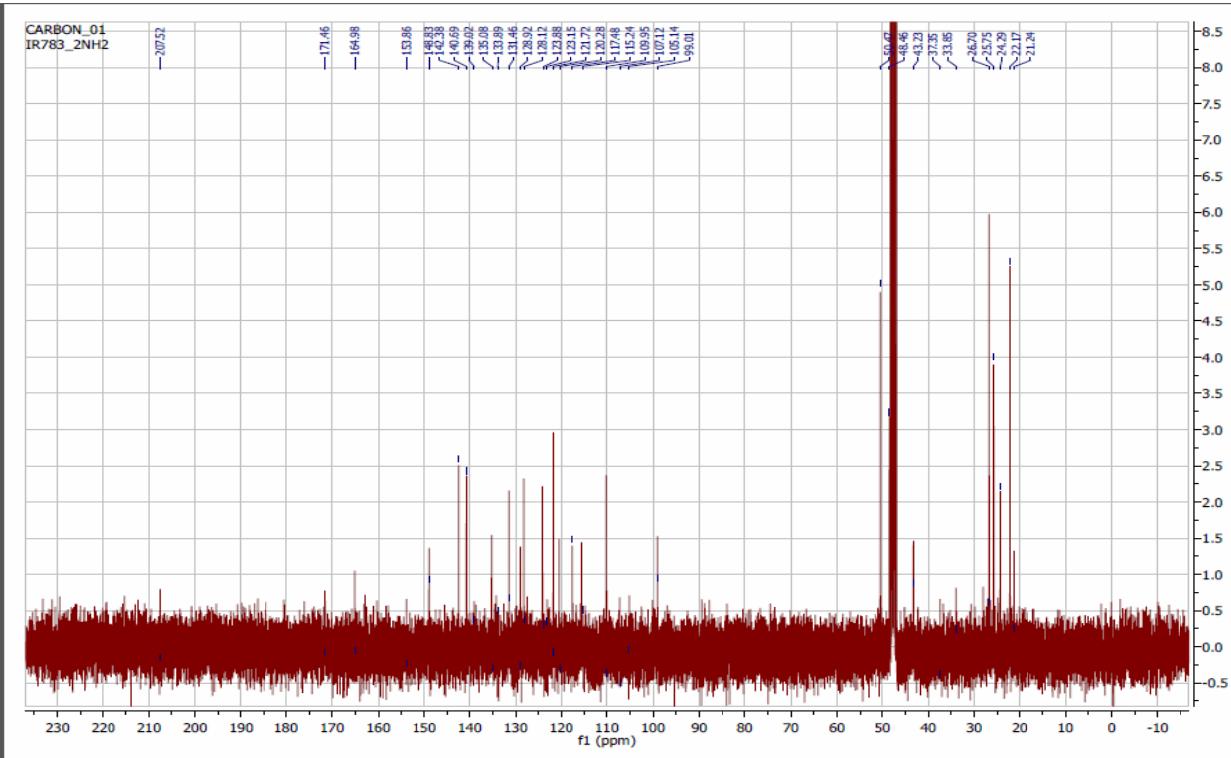
Figure S5 Fluorescence decays of DAP-LT2 before and after reaction with NO

The change in fluorescence lifetime was from 1.02 to 1.05 ns. Ex/em. 740/790 nm, 10% serum/PBS buffer, room temperature

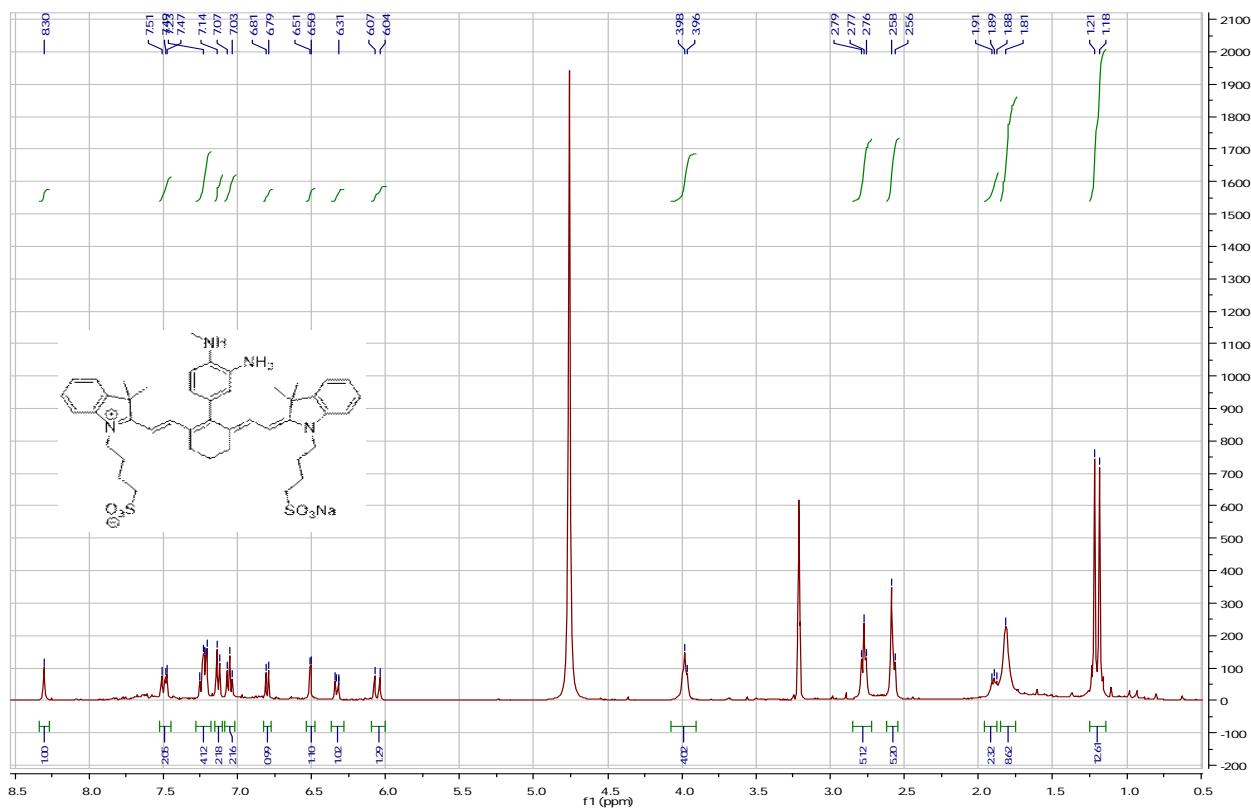
1H NMR DAP-LT1, CD₃OD



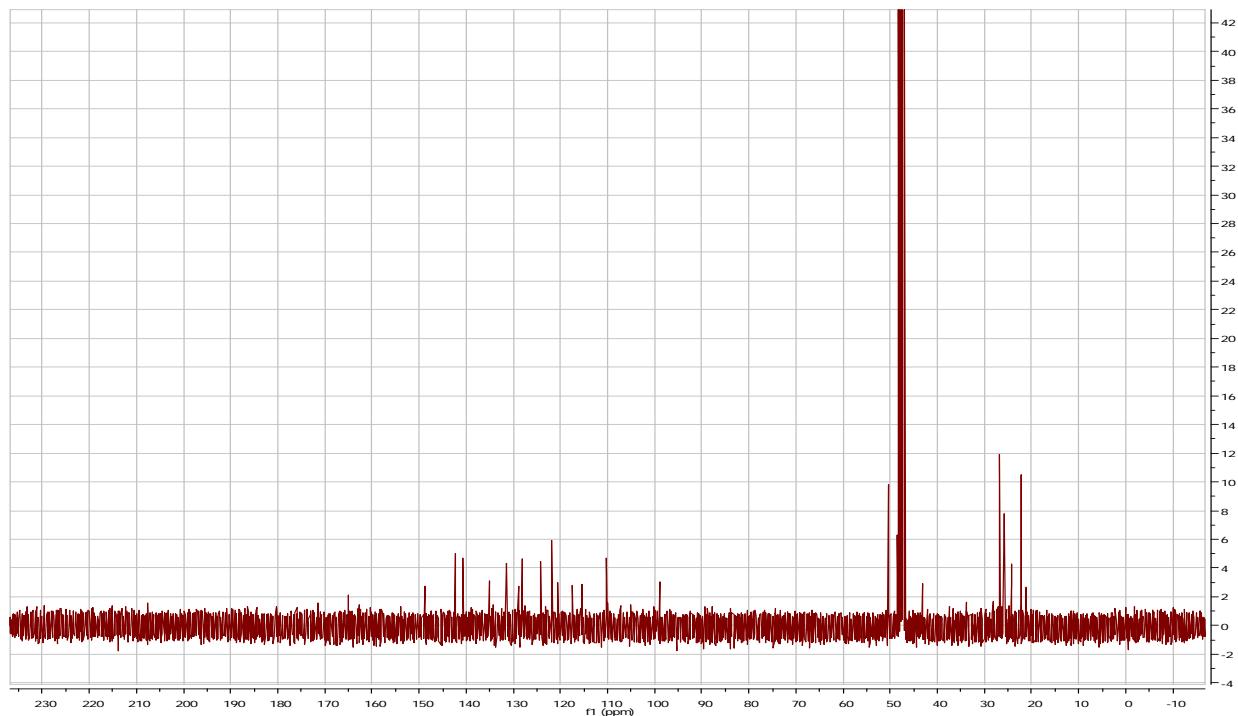
13C NMR DAP-LT1, CD₃OD



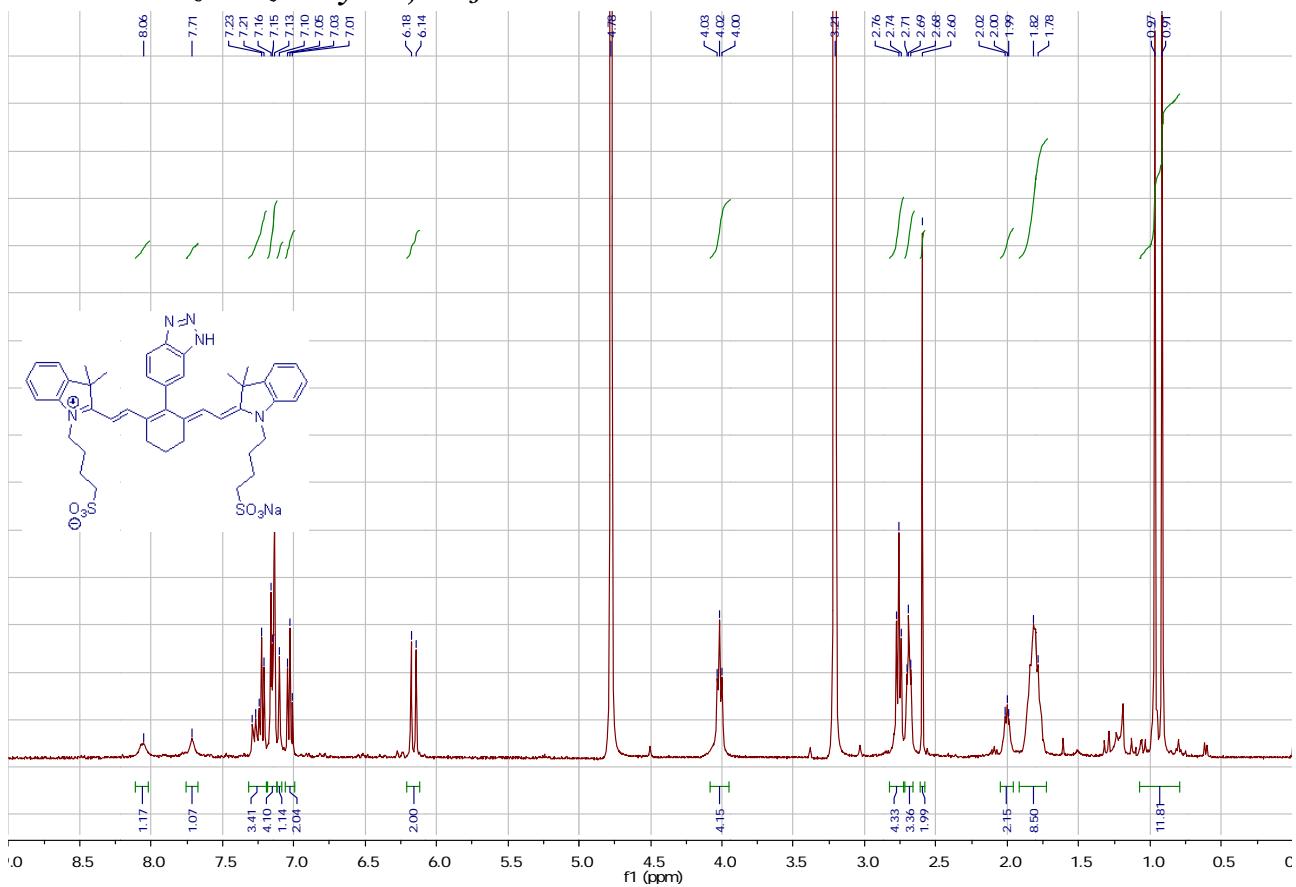
1H NMR DAP-LT2, CD₃OD



13C NMR DAP-LT2, CD₃OD



1H NMR Benzotriazole dye 11, CD₃OD



13C NMR Benzotriazole dye 11, CD₃OD

