

Supporting Information

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***N*-Methyl-Substituted Fluorescent DAG–Indololactone Isomers Exhibit Dramatic Differences in Membrane Interactions and Biological Activity**

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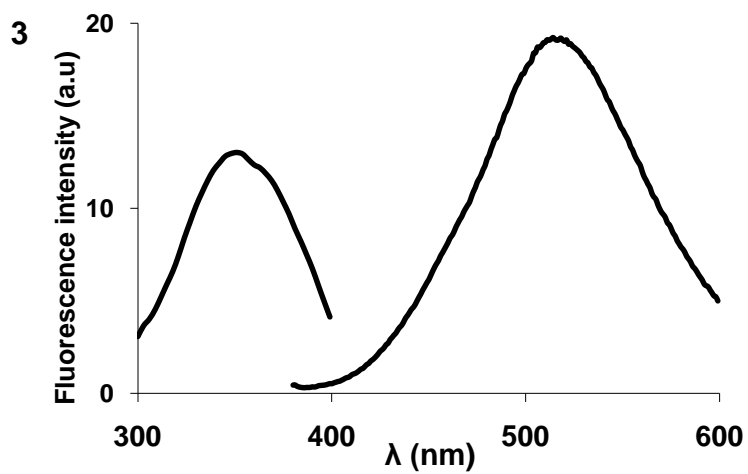
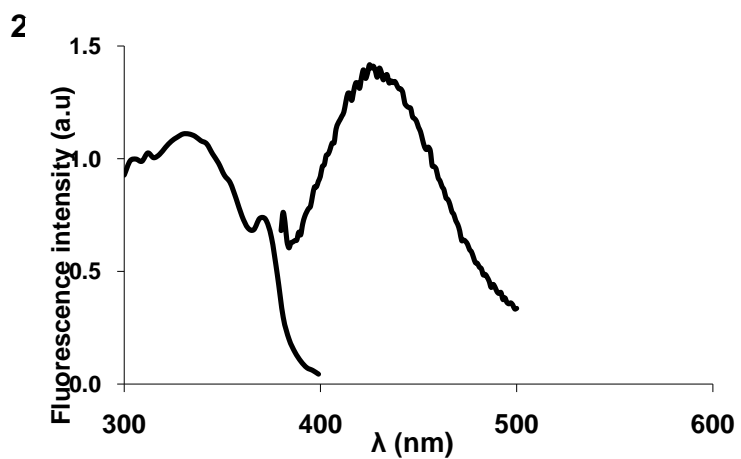
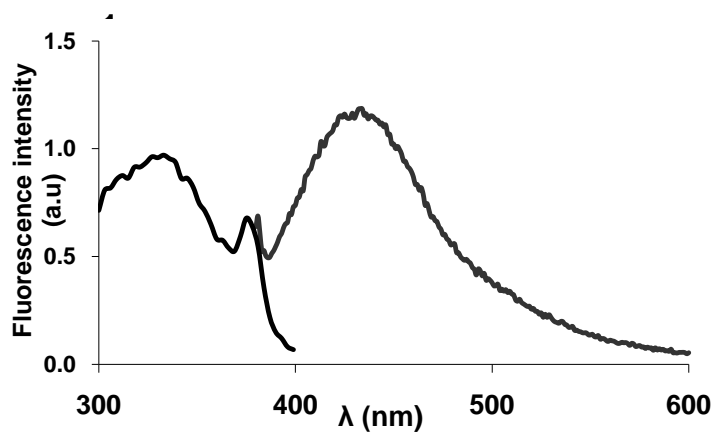


Figure 1,SI: Excitation spectra (left) and emission spectra (right) of the DAG-indololactones (dissolved in water).

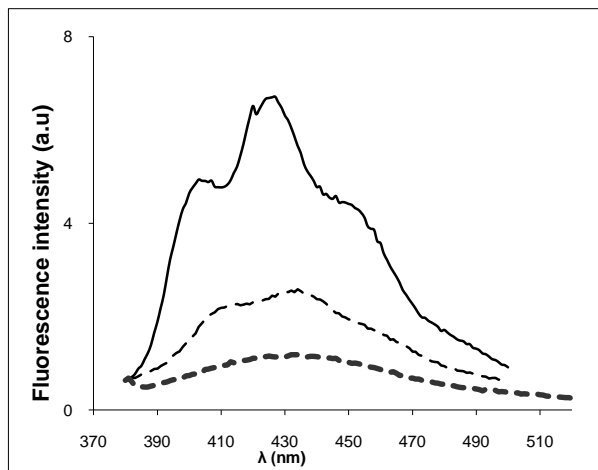
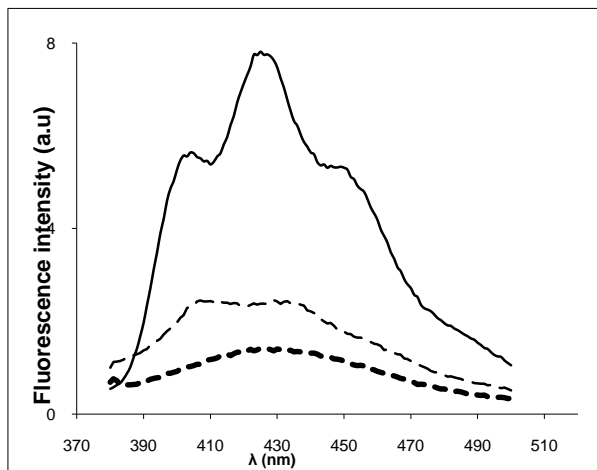
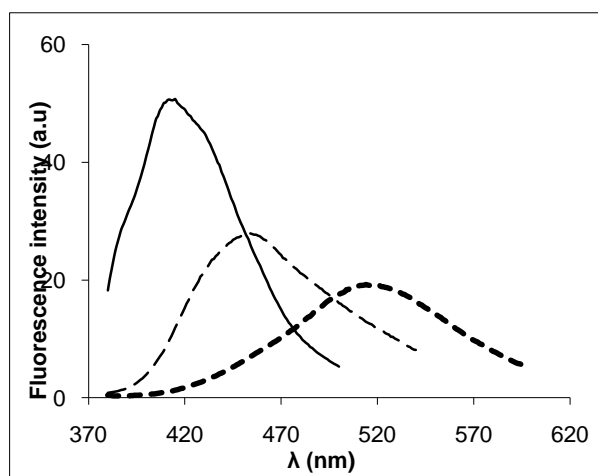
1**2****3**

Figure 2,SI: Dependence of DAG-indololactone fluorescence emission upon solvent polarity. Short dash: water; long dash: ethanol; solid line: n-hexane.

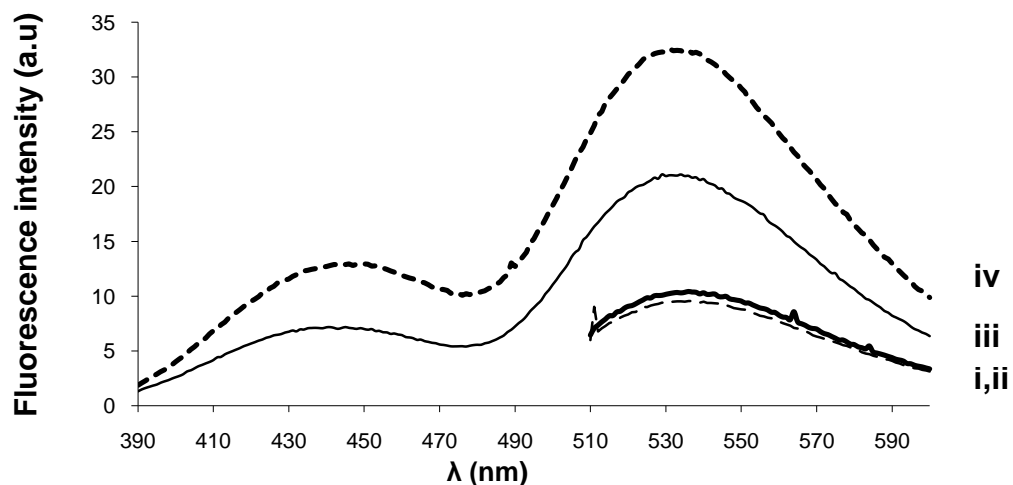


Figure 3,SI: Fluorescence energy transfer from 3 to NBD-PE. Fluorescence emission spectra of NBD-PE/PC (1:100 mole ratio) vesicles incubated with **3**. The spectra were acquired using two different excitation wavelengths: 330 nm (excitation of **3**), and 469 nm (excitation of NBD-PE). **i** – excitation at 469 nm, low concentration of **3** (40 μm); **ii** - excitation at 469 nm, high concentration of **3** (190 μm); **iii** - excitation at 330 nm, low concentration of **3** (40 μm); **iv** - excitation at 330 nm, high concentration of **3** (190 μm).