

# **ELECTRONIC SUPPLEMENTARY INFORMATION**

## **(ESI)**

### **Amplification of Light Collection in Solid-State Dye-Sensitized Solar Cells via Antenna Effect through Supramolecular Assembly**

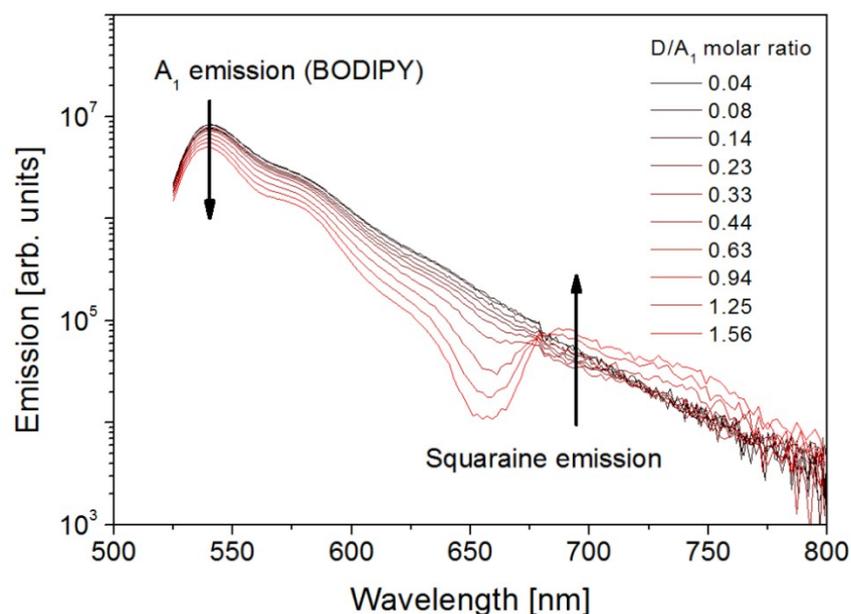
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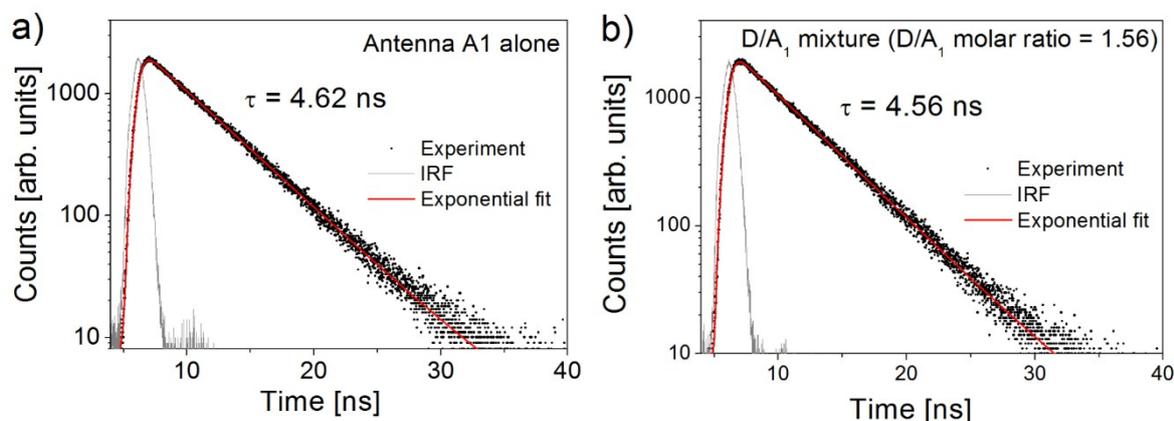
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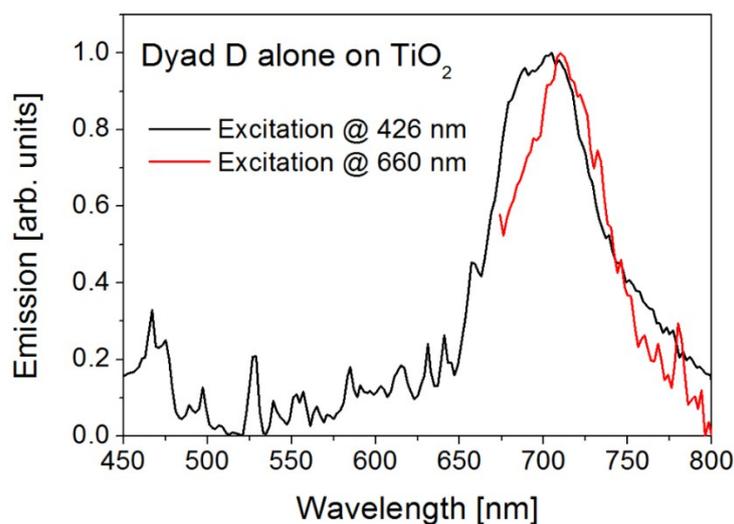
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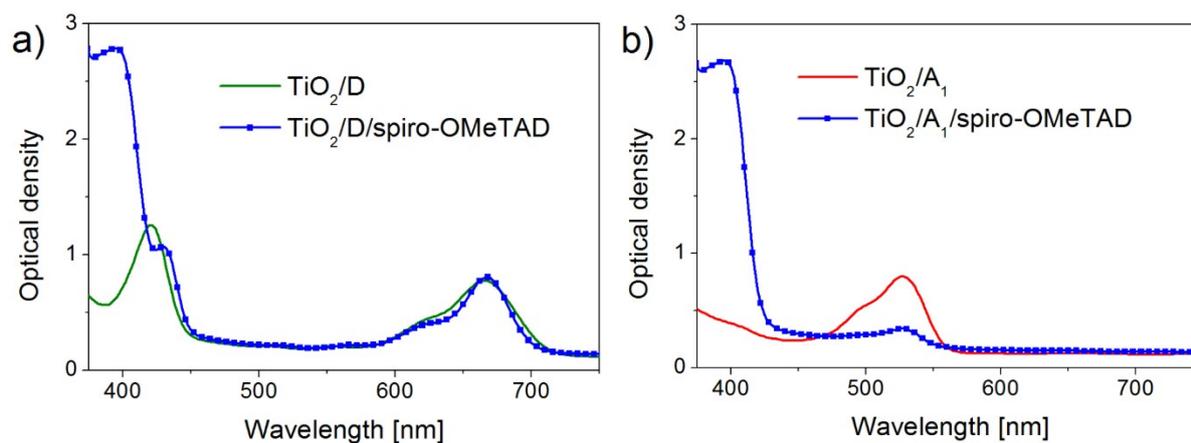
**Figure S1.** Photoluminescence spectra of a dyad D/Antenna A1 mixture as a function of the molar ratio. The solvent is a mixture of ethanol:dichloromethane. The excitation is at 520 nm (initial concentration of antenna A1 dyad D is 0.0125 mM).



**Figure S2.** Photoluminescence decays of (a) antenna A<sub>1</sub> alone in solution (acetonitrile, 0.0125mM) and of (b) a mixture of dyad D and antenna A<sub>1</sub> (mixture ethanol:dichloromethane, 1:1 in volume) corresponding to a D/A<sub>1</sub> molar ratio of 1.56). The deconvolution of the experimental data by a mono-exponential function, taking into account the instrument response function (IRF), are also presented, as well as the corresponding decay time.



**Figure S3.** Photoluminescence spectra of dyad D alone on porous  $\text{TiO}_2$  electrode ( $2 \mu\text{m}$  thick films) for an excitation at 426 and 660 nm. The signal to noise ratio is reduced due to the poor emission intensity of the squaraine unit centered at approximately 706 nm.



**Figure S4.** Optical absorption spectra of sensitized  $\text{TiO}_2$  electrodes ( $2 \mu\text{m}$  thick) sensitized with (a) dyad D alone and (b) antenna  $A_1$  alone, before and after infiltration by the molecular HTM (spiro-OMeTAD), and used as porous electrodes for the ssDSSC devices. In both cases, a significant removal of chromophores following the HTM infiltration is evidenced, especially in the case of antenna  $A_1$ .