

White-Light Electroluminescence from a Layer Incorporating a Single Fully-Organic Spiro Compound with Phosphine Oxide Substituents

P. Tourneur^{a,†}, F. Lucas^{b,†}, C. Quinton^b, Y. Olivier^c, R. Lazzaroni^{a,d}, P. Viville^d, J. Cornil^a, and C. Poriel^b

^a *Laboratory for Chemistry of Novel Materials, Materials Research Institute, University of Mons, Mons, Belgium. E-mail: jerome.cornil@umons.ac.be.*

^b *Université de Rennes, CNRS, ISCR-UMR 6226, F-35000 Rennes, France. E-mail: cyril.poriel@univ-rennes1.fr.*

^c *Unité de Chimie Physique Théorique et Structurale (UCPTS) & Laboratoire de Physique du Solide (LPS), Namur Institute of Structured Matter (NISM), University of Namur, Namur, Belgium. E-mail: yoann.olivier@unamur.be.*

^d *Materia Nova, Materials R&D center, Mons, Belgium. E-mail: pascal.viville@materianova.be.*

[†] First authorship shared.

Supporting information

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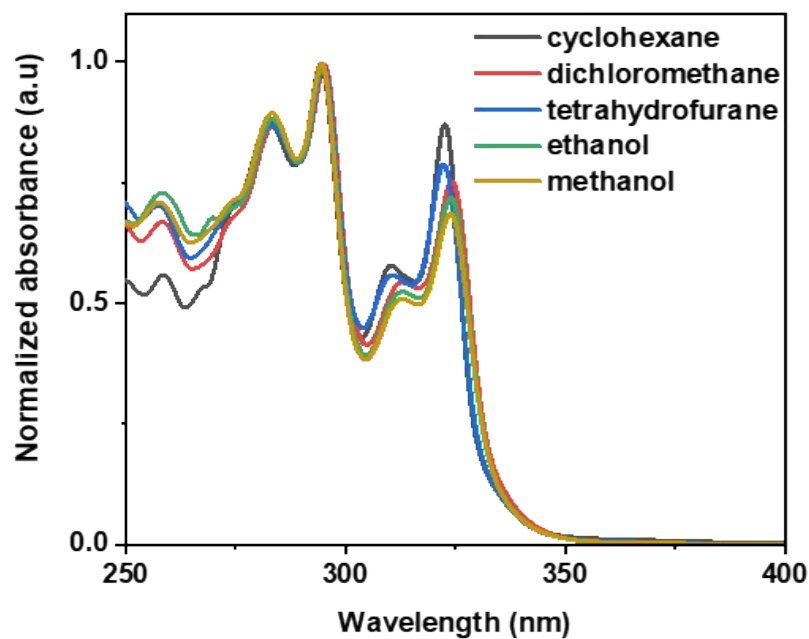


Figure S1 - Normalized absorption spectrum of SPA-F-(POPh₂)₂ in different solvents recorded at room temperature without heating.

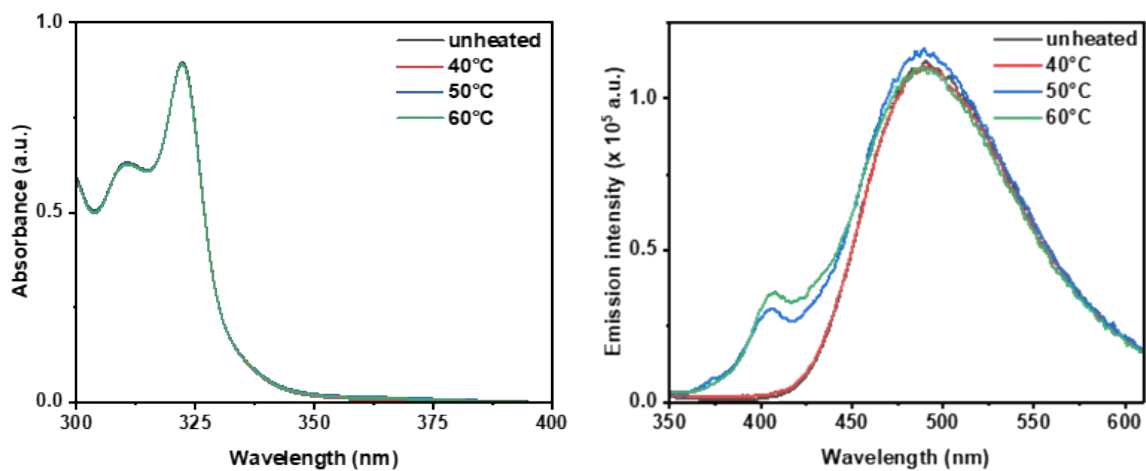


Figure S2 - Evolution of the absorption (left) and emission (right) spectra of SPA-F-(POPh₂)₂ in THF upon heating at a concentration of 4.4×10^{-5} mol/L. The solution is heated during 30 minutes for each temperature and the spectra are recorded once the solution has cooled down to room temperature.

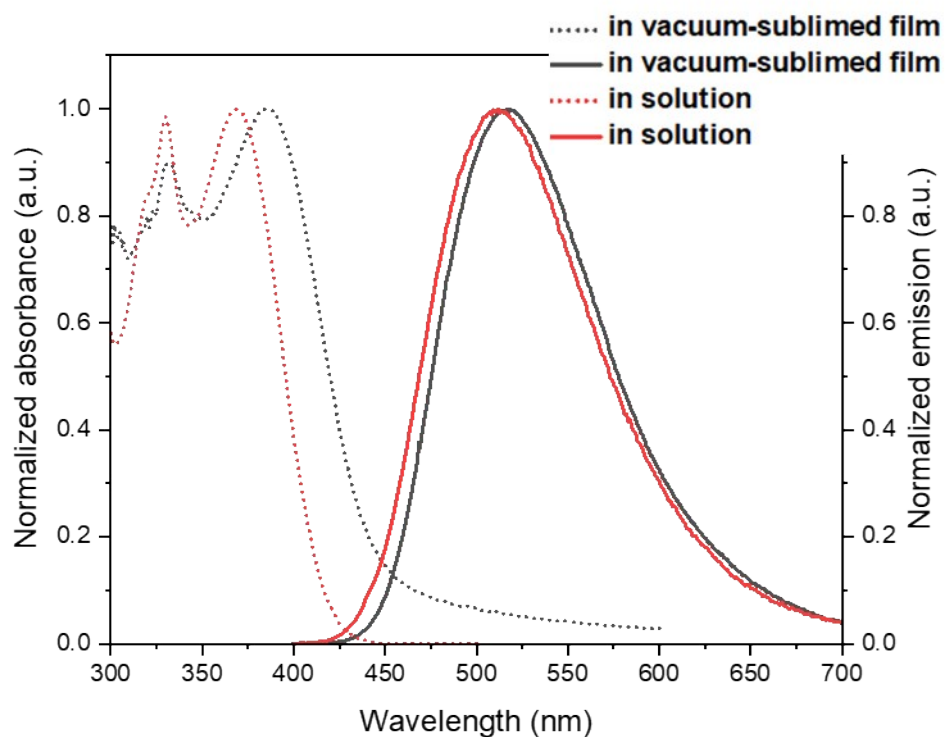


Figure S3 - Absorption and emission spectra of 2CZPN in THF and in vacuum-sublimed films.

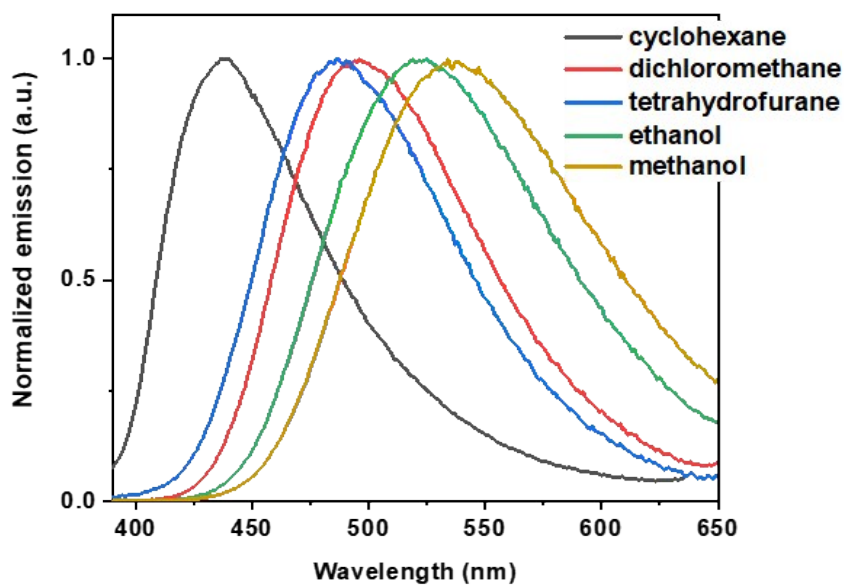


Figure S4 - Normalized emission spectrum of SPA-F-(POPh₂)₂ in different solvents upon excitation at 330 nm, as recorded at room temperature without heating ($A_{(330\text{ nm})} < 0.1$). The concentration used is $> 6 \times 10^{-5}$ mol/L.

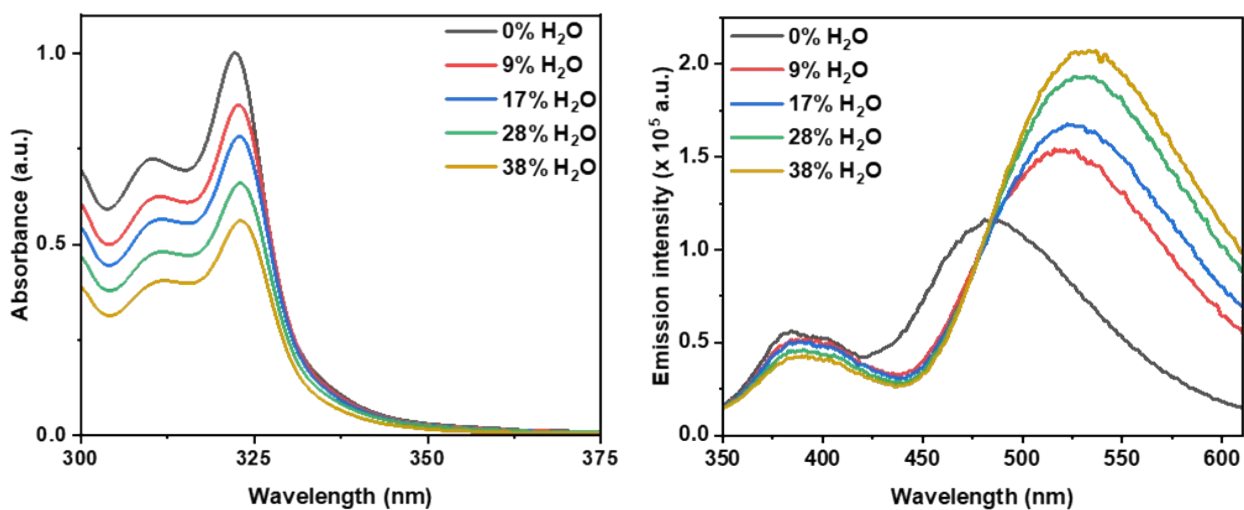


Figure S5 - Evolution of the absorption (left) and emission (right) spectrum in THF upon progressive addition of water. The initial concentration is 4.8×10^{-5} mol/L.

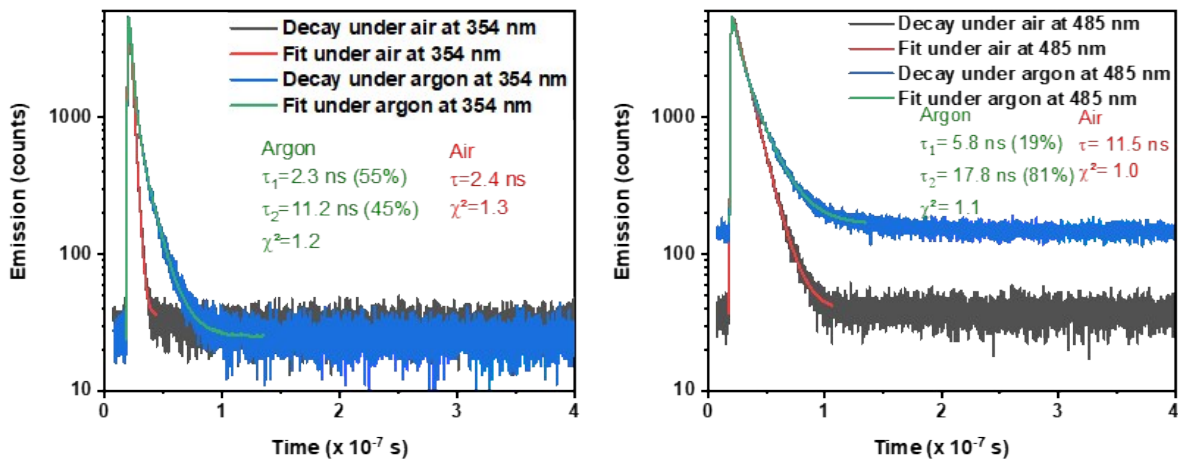


Figure S6 - Transient PL decay of SPA-F(POPh₂)₂ in THF under air (black) argon (blue) at (left) $\lambda_{em} = 354$ nm and (right) $\lambda_{em} = 485$ nm. The concentration is $< 6 \times 10^{-5}$ mol/L.

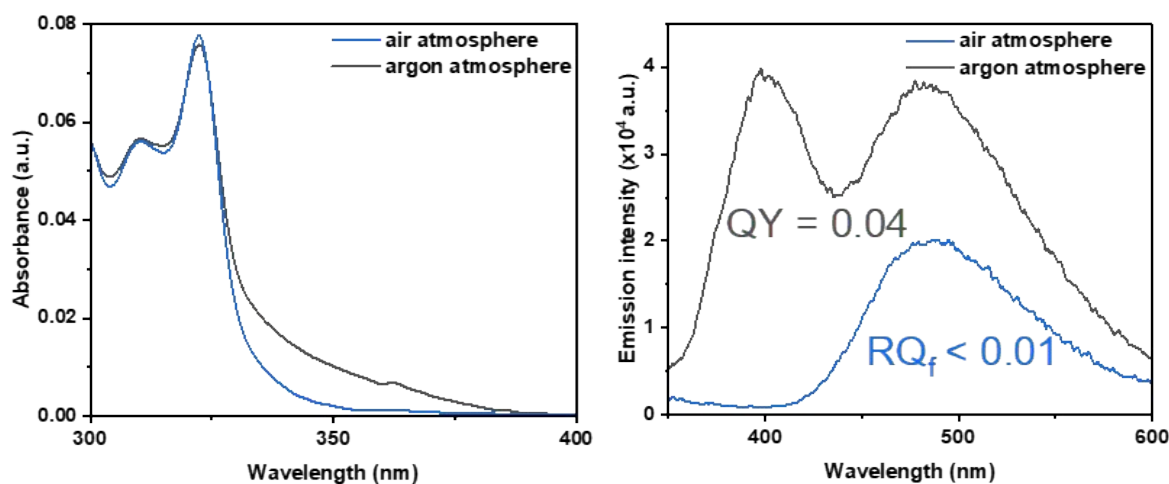


Figure S7 - Absorption (left) and emission (right) spectra of **SPA-F(POPh₂)₂** in THF under air (blue) and argon (black). Quantum yields (QY) are noted in the right graph. The concentration is $< 6 \times 10^{-5}$ mol/L.

Estimation of the fluorescence quantum yield

Quantum yields in solution (ϕ_{sol}) were calculated relative to quinine sulfate ($\phi_{ref} = 0.546$ in H₂SO₄ 1 N). ϕ_{sol} was determined according to the following equation:

$$\phi_{sol} = \phi_{ref} \times \frac{S_s \times A_r}{S_r \times A_s} \times \left(\frac{\eta_s}{\eta_r} \right)^2$$

where subscripts *s* and *r* refer respectively to the sample and reference, *S* is the integrated fluorescence intensity and *A* the absorbance at the excitation wavelength, η is the refractive index of the solvent ($\eta_s = 1.405$ for THF).

Film preparation

Thin films were prepared either by spin-coating ca 300 μ L of a THF solution of **SPA-F(POPh₂)₂** on sapphire plate (10 mm \times 10 mm) at 2500tr/min on a Süss+MicroTec Lab Spin6/8 or by vacuum-deposition using a Kurt J. Leskers "Spectros" thin film evaporator system in glovebox (JACOMEX) and on a glass substrate.

OLED devices

General fabrication procedures

Commercial ITO coated-glasses from Naranjo have been used as patterned substrates for device fabrication. Prior deposition, substrates were first cleaned in different ultrasonic baths during 10 min with

solutions of 10% RBS detergent in demineralized water, tap water, demineralized water and acetone, and finally with an UV/ozone cleaning system from UVOCS.

All devices have been fabricated in glovebox (JACOMEX) with a Kurt J. Leskers “Spectros” thin film evaporator system.

Device encapsulation with a dense layer of Al₂O₃ has been carried out using an Atomic Layer Deposition (ALD) SAVANNAH 100 system (from Annealsys) in order to protect the devices prior measurements of the EQE.

Characterization:

Curves of current density (mA/cm²) and luminance (Cd/m²) have been obtained using a luminance meter LS-110 system from MINOLTA equipped with a 135 Lens model and a Keithley 2400 Source measuring unit.

The EQE of the devices have been calculated using the equation:

$$EQE = \frac{P_{ext}}{n_e} = \frac{\int_{380}^{780} \frac{Radiance}{Photon\ energy}}{\frac{J \times A}{q}} = \frac{\int_{380}^{780} \frac{W(\lambda) d\lambda}{h \times \frac{c}{\lambda}}}{\frac{J \times A}{q}}$$

where A is the device area, J the measured current density and q the charge of an electron.

The radiance (λ) spectrum of the devices has been measured with a LED analyzer OmniLED-LMS10 system from Labsphere equipped with a 10” integrating sphere and a Keithley 2400 Source measuring unit. The use of an integrating sphere allows to collect all emitted photons in all directions and hence to avoid a correction in the emission spectrum with a Lambertian’s law.

Color Recording Index (CRI) coordinates, from CIE, have been recorded with the same integrating sphere.

The turn-on voltages (V_{ON}) have been estimated from the crossing of the two tangents of the current-voltage curves.