Synthesis, electrochemical and photophysical studies of the borondifluoride complex of a meta-linked biscurcuminoid†

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The synthesis of the borondifluoride complex of a biscurcuminoid system is described, and its electrochemical and photophysical properties are compared to those of a series of monochromophoric models. The data show that the meta-linked biscurcuminoid exhibits enhanced optical properties in solution, such as high optical brightnesses obtained at one- and two-photon excitation ($B^1 = 87000 \text{M}^{-1} \text{cm}^{-1}$ and $B^2 = 313 \text{GM}$, respectively). UV/visible absorption of solid-state particles formed in water solution revealed that the four investigated dyes are strongly aggregated and fluorescence spectroscopy showed that they are emissive in the NIR with fluorescence quantum yields spanning from 1.5 to 12.5%. In the case of the meta-linked biscurcuminoid, a high brightness is obtained using one- and two-photon excitation ($B^1 = 6952 \text{M}^{-1} \text{cm}^{-1}$ and $B^2 = 70 \text{GM}$, respectively). Such red-shifted emission combined with a yet significant brightness in the solid-state is believed to arise from the choice of the meta-linkage which limits strong intermolecular packing.

Introduction

The design of organic dyes with tailored optical properties is of utmost importance for applications in advanced fields such as bioimaging,1 photodynamic therapy,2 theranostics, display3 and telecommunication technologies,4 and photovoltaics.5–7 In that context, compounds displaying high molar absorption coefficients, large two-photon absorption cross sections, high luminescence quantum yields in solution and in the solid state, large Stokes shifts, high thermal and photochemical stability represent attractive candidates as fluorescent reporters. Especially, dyes that emit in the near infrared (NIR, wavelengths longer than 700 nm according to The International Commission on Illumination) using two-photon excitation in the NIR region are of great interest for use in cell imaging because both excitation and detection operate in the biological transparency window.

Among the many classes of organic dyes, boron complexes, such as borondipyromethene (BODIPY) compounds,8–11 are particularly attractive. In addition of the interesting optical properties in solution, many borondifluoride complexes (other than BODIPY) have also been shown to yield rather efficient photoluminescent behavior in the solid state.12 In particular, compounds deriving from acetylacetone ligand have shown interesting properties such as high two-photon absorption cross sections,13,14 mechno-fluorochromic behaviors15,16 and efficient NIR emissions17,18 that led to their use for cells imaging19 or as sensors of volatile acid/base,20,21 fluorescent reporters for amylold,22,23 optical sensors for anaerobic environment24 and electron donors in solar cells.25

Recently, we and others have shown that curcuminoid structures containing the borondifluoride unit represent efficient emitters that fulfill many of the previous requirements. However, our work also led to the conclusion that solid-state fluorescence of BF₂ complexes of curcuminoids arose from highly stacked chromophores. Such interactions could explain the emission occurring in the NIR but inherently induce efficient face-to-face quenching, which limit the fluorescence quantum yield ($\Phi_f$) to ca. 5%.17,18 These results led us to consider dye 4 (Chart 1) in which two curcuminoid subunits are covalently connected via a meta-phenylene linker. Indeed, this structural design has proven to be successful in the case of electroluminescent conjugated polymers such as poly(phenylenevinylene) because the meta-linkage was observed to inhibit interchain interactions and excimer formation in the condensed phase, thereby ensuring brighter emission than in the para-linked counterparts.26 The introduction of three aliphatic long chains at the meta-phenylene bridge in 4 was anticipated to further limit solid-state interchromophoric interactions and to provide a higher solubility in organic solvents for the extended structure. As a result, compound 4 features a non symmetrical curcuminoid system, which required the investigation of the model compound 3 that
possesses a non symmetrical structure as well. Compounds 1 and 2 served as models to estimate the electronic influence of the electron donor (D) peralkylated phloroglucinol ring in the photophysical investigation.

We report herein the synthesis of compounds 1–4 and a study of their electrochemical and optical properties in organic solvents. The two-photon excited fluorescence (TPEF) properties of the four dyes were characterized in solution. We describe the preparation and fluorescence emission of organic nanoparticles of 1–4. Our approach gives a strategy toward the design of extended biscurcuminoids with improved emission ability in both the solution and the solid states.

Results and discussion

Synthesis

The synthetic routes toward 1–4 are outlined in Scheme 1. The borondifluoride complexes 1 and 2 were prepared using a previously reported procedure. Syntheses of the unsymmetrical BF₂ complexes 3 and 4 required first of all the preparation of the hemicurcuminoid 5. Similarly to what has been published, this intermediate was prepared by a Knoevenagel reaction using an excess of acetylacetone (acac/alddehyde 3 : 1), which provided the compound 5 in a reasonable yield of 60%. Then, the reaction of the intermediate 5 with one equivalent of 2,4,6-trimethoxybenzaldehyde afforded the ligand Lig 3. Lig 4 was prepared in a yield of 34% by reacting two equivalents of 5 with 1,3,5-tris(n-octyloxy)benzaldehyde.

Complexation to boron difluoride was performed by reacting the ligands with a slight excess of the boron trifluoride etherate in dichloromethane solution (DCM). The dyes were purified by crystallization or, when necessary, by column chromatography. Dyes 1–4 were obtained as highly colored solids and characterized by ¹H- and ¹⁹F-NMR spectroscopies and by high resolution mass spectrometry (HRMS).
the oxidation and reduction half-wave potential values for the studied dioxaborine derivatives (10 μM) in dichloromethane solution vs. ferrocene/ferrocenium using tetrabutylammonium hexafluorophosphate as electrolyte (100 μM), the values are given in volt

| n  | E_{1/2}^{ox} | E_{1/2}^{red} | E_{1/2}^{ox} – E_{1/2}^{red} | ΔE_\text{g} \text{a} |
|----|-------------|---------------|-----------------------------|----------------|
| 1  | -1.27       | 1.10          | n                           | 2.37           |
| 2  | -1.44       | 0.72          | 1.19                        | 2.66           |
| 3  | -1.37       | 0.89          | 2.26                        | 2.26           |
| 4  | -1.30       | 1.05          | 2.35                        | 2.35           |

Electrochemical HOMO–LUMO gap ΔE_\text{g} = (E_{1/2}^{ox} – E_{1/2}^{red}). The second oxidation process occurs at potentials out of the solvent electrochemical window.

Electrochemistry

The electrochemical properties of dyes 1–4 were investigated in DCM solution containing 0.1 M [t[Bu_4N]PF_6] (scan rate of 100 mV s^-1). The cyclic voltammograms (CV) are given in Fig. 1 and Fig. S1 (ESI†) and the oxidation and reduction half-wave potential values (E_{1/2} vs. ferrocene/ferrocenium) are collected in Table 1.

The four complexes exhibit one-electron oxidation and reduction waves that can be attributed to the donor alkoxy-substituted phenyl (D) end-groups and the acceptor (A) dioxaborine ring, respectively. The first oxidation potential decreases with the strength of the D units, depending on the number of appended methoxy groups (Table 1). In contrast, increasing the donor strength induces a cathodic shift of the reduction potential but overall, the electrochemical gap is decreased by ca. 0.2 V.

The CV of 2 (Fig. S1, ESI†) clearly shows two successive and well separated one-electron oxidation waves that reflect the electronic communication between trimethoxyphenyl end-groups through the curcuminoid backbone. It can be noted that the first oxidation and reduction potential values of the unsymmetrical complex 3 are intermediate between those of the symmetrical complexes 1 and 2.

The first oxidation and reduction potential values obtained for 4 are closer to those recorded for 1 than for 3. Furthermore, the HOMO–LUMO electrochemical gap is also very close for both compounds 1 and 4. These observations suggest that the first oxidation in 4 is likely centered on the p-methoxy aryl groups. The increase of the oxidation potential in 4 relative to 3 could be due to presence the second curcuminoid moiety. In addition, since both reduction and oxidation processes of 4 involve the same number of electrons, we assume that methoxy-phenyl end-groups and dioxaborine rings in 4 are simultaneously oxidized and reduced, respectively, at the same potential. This effect is in agreement with the reduced conjugation between meta-linked chromophores.

Photophysical study in solution

The electronic absorption spectra of 1–4 were recorded in DCM solutions (Fig. 2a) and the spectroscopic data are reported in Table 2. The spectra consist mainly of one intense transition band at low energy (450–550 nm) attributed to a strongly allowed π-π* transition. A second electronic transition band of much lower intensity appears as a shoulder at higher energy (<400 nm, vide infra). The spectra of compounds 1–3 display identical shape of the absorption profiles, they only differ by the position of the bands. The increase of donor strength from 1 to 2 induces a red-shift of the electronic transitions. As expected, the unsymmetrical dye 3 absorbs at a wavelength intermediate between those recorded for 1 and 2. In contrast, compound 4 exhibits a low-energy absorption band with very different shape, full width at half maximum, and intensity. The molar absorption coefficient determined for 4 is twice as high as that of the other three dyes. The more complex shape and the larger Stoke shift of the absorption band is likely from intramolecular excitonic coupling between the two curcuminoid chromophores. This excitonic coupling explains the low value of the optical gap determined for 4.

The complexes 1–4 are fluorescent in the visible region (540–575 nm) upon excitation into the low-energy transition band and exhibit fluorescence quantum yields ranging from 44 to 61% in DCM. In agreement with electronic absorption data, an increase of the donor strength causes a red-shift of the fluorescence emission from 538 nm (1) to 574 nm (4). It is worth noting that the highest value of Φ_\text{f} (61%) is obtained for complex 4, giving a high brightness value of ca. 87 000 M^-1 cm^-1.

The solvent dependence of absorption and emission properties was studied for the four borondifluoride complexes (Tables S1 and S2, ESI†). Both electronic UV/visible absorption and fluorescence emission spectra undergo a bathochromic shift and lose their vibronic structures when solvent polarity increases (Fig. S3–S6, ESI†). These features show that the Franck–Condon excited state S_1 is more polar in nature than the ground state. Furthermore, the positive slope of the Lippert–Mataga plots reveals that, in those systems, the relaxation occurs toward a solvent-equilibrated charge transfer (CT) singlet excited state along with an increase of dipole moment with respect to the ground state (Fig. S2, ESI†).

The fluorescence intensity of compounds 1–3 continuously increases with increasing solvent polarity. Both solvatochromism and the negative solvatokinetic behavior suggest the formation of an emissive CT state. The fact that the fluorescence lifetimes also

![Fig. 1. Cyclic voltammogram of the bis borondifluoride complex 4 in DCM solution containing 0.1 M [t[Bu_4N]PF_6] (scan rate of 100 mV s^-1).](image-url)
increase with solvent polarity leads to constant values of the radiative rate constants $k_f$ for the three dyes (1–3). Concomitantly, the nonradiative rate constants $k_{nr}$ decreases in polar solvents. The case of 4 is different. Upon increasing solvent polarity, the fluorescence quantum yield first increases (negative solvatochemical effects) then decreases (positive solvatochemicals), a maximum being observed. The positive solvatochemical behavior of 4 in polar solvent suggests an enhanced population of a highly polar CT state prone to a strong nonradiative deactivation. At this stage, the peculiar photophysical behavior of 4 remains unclear.

Two-photon excited fluorescence emission and excitation spectra of 1–4 were recorded in the 700–1000 nm wavelength range using a femtosecond Ti-Sapphire pulsed laser source, according to the experimental protocol described by Webb et al. using coumarin-307 and rhodamine B as references. The observation of a quadratic dependence of the fluorescence intensity versus incident laser power at several wavelengths unambiguously confirmed that the origin of the fluorescence emission can be assigned to a TPA process in DCM solution (Fig. S7, ESI†). In the experimental laser power range used for these measurements, we checked that no saturation or photodegradation occurred. The two-photon excitation spectra of 1, 2, 3 and 4 in DCM are shown in Fig. 3 and Fig. S9 (ESI†) and the corresponding data are reported in Table 2. As reported elsewhere, 1 has a TPA cross section ($\sigma_{TPA}$) of 155 GM at 770 nm while dye 2, containing stronger donating groups, has its maximum red-shifted to 810 nm with a larger $\sigma_{TPA}$ value of ca. 248 GM. It is interesting to note that 3 presents an intermediate $\sigma_{TPA}$ value of 208 GM at 790 nm. For dye 4, as already observed for the molar absorption coefficient, the two-photon cross section is slightly more than twice (i.e. 513 GM) the value determined for 3. This gives a two-photon brightness of 313 GM, that is much higher than those obtained with the model borondifluoride complexes 1–3.

As noticed previously for curcumin and borondifluoride complexes of curcuminoids, 14 the TPA band of the four complexes

| Compound | UV-vis | Fluorescence | TPEF |
|----------|--------|--------------|------|
|          | $\lambda_{abs}$ (nm) | $\lambda_{em}$ (nm) | $\Delta\mu_{CT}$ (cm$^{-1}$) | $\Phi_f$ | $B^1$ | $\tau_f$ | $k_f$ | $k_{nr}$ | $\lambda_{max}$ (nm) | $\sigma_{TPA}$ (GM) | $B^2$ |
| 1 (DCM)  | 488 | 75 480 | 538 | 1904 | 0.44 | 33 211 | 1.30 | 3.4 | 4.3 | 770 | 155 | 68 |
| 2 (DCM)  | 524 | 84 860 | 566 | 1416 | 0.52 | 44 127 | 1.72 | 3.0 | 2.8 | 810 | 248 | 129 |
| 3 (DCM)  | 511 | 74 510 | 561 | 1957 | 0.46 | 34 275 | 1.74 | 2.6 | 3.1 | 790 | 208 | 96 |
| 4 (DCM)  | 529 | 143 140 | 574 | 1421 | 0.61 | 87 315 | 1.69 | 3.6 | 3.1 | 800 | 248 | 129 |
| 1 (water)| 451 | 34 170 | 710 | 8088$^b$ | 0.06 | 2050 | 6.44 | 0.09 | 1.5 | 780 | 210 | 13 |
| 2 (water)| 489 | 27 380 | 717 | 6503$^b$ | 0.035 | 783 | 1.32 | — | — | — | — | — |
| 3 (water)| 407 | 25 950 | 711 | 10 505$^b$ | 0.015 | 443 | 2.13 | — | — | — | — | — |
| 4 (water)| 468 | 55 620 | 692 | 6917$^b$ | 0.125 | 6952 | 1.89 | 6.01$^c$ | 6.64$^c$ | 850 | 560 | 70 |

$^a$Absorption maximum wavelengths $\lambda_{abs}$ (nm); molar absorption coefficients at maximum $c_{max}$ (M$^{-1}$ cm$^{-1}$); fluorescence maximum wavelengths $\lambda_{em}$ (nm); Stokes shifts $\Delta\mu_{CT}$ (cm$^{-1}$); fluorescence quantum yields $\Phi_f$; brightness $B = \Phi_f \times \varepsilon$ (M$^{-1}$ cm$^{-1}$); fluorescence lifetimes $\tau_f$ (ns); radiative $k_f$ (10$^5$ s$^{-1}$) and nonradiative $k_{nr} = (1 - \Phi_f)\varepsilon$ (10$^5$ s$^{-1}$) rate constants; two-photon absorption maximum $\lambda_{max}$ (nm); two-photon cross section $\sigma_{TPA}$ (GM); two-photon brightness $B^2 = \Phi_f \times \sigma_{TPA}$ (GM). $^b$Pseudo-Stokes shift determined using the maximum absorption. $^c$A biexponential decay was found. $^d$Not determined due to high scattering of light with those particles.
does not match the one-photon transition as it is strongly blue-shifted, corresponding to the two-photon allowed transition at higher energy mentioned previously. The lowest-energy transitions of 1–4 are not completely TPA forbidden: $\sigma^{\text{TPA}}$ values of ca. 10–15, 30–35, 25–30 and 100 GM were measured for 1, 2, 3 and 4, respectively (Fig. 3 and Fig. S8, ESI†). Such situation is often encountered for D–A–D chromophores$^{32-35}$ and, in the case of free curcumin, Hernández et al. have shown that the $S_0$–$S_2$ transition was more TPA-allowed than the $S_0$–$S_1$ transition. However, a much weaker contribution from the $S_0$–$S_2$ transition was still observed.$^{31}$

![Figure 3](image1.png)

**Fig. 3** Two-photon excitation (higher $x$-coordinate and right $y$-coordinate) with their error bars, OPA spectra (lower $x$-coordinate and left $y$-coordinate) in DCM: (a) dye 3 and (b) dye 4.

![Figure 4](image2.png)

**Fig. 4** Overlaid absorption (solid lines) and emission (dashed lines) spectra of DCM solutions (black) and particles in water (red) for (a) 1; (b) 2; (c) 3 and (d) 4.
Solid-state optical properties

We prepared solid-state particles, P-2–P-4, by quickly adding a concentrated THF solution of the dyes 2–4 into water according to the classical fast precipitation method. The so-obtained suspensions enabled the measurement of the UV/visible absorption and fluorescence spectra of the aggregated molecules. We reported the preparation of P-1 in a previous publication and the resulting spectroscopic data are recalled here for the sake of comparison. The shape of the electronic absorption spectra of P-1–P-4 is strongly affected relative to the solution spectra showing that Davydov splitting is occurring within such dyes. The absorption profiles of P-1, P-2 and P-4 (Fig. 4a, b, and d, respectively) are complex, broad, and their maximum presents a hypsochromic shift (ca. 30–60 nm). These observations show that excitonic interactions prevail in the condensed phase, which could be correlated with single-crystal X-ray diffraction data in the case of 1. Such complex structure is in line with Davydov splitting into high- and low-energy transitions. The absorption spectrum of P-3 displays a very different effect (Fig. 4c). The low-energy transition is considerably blue-shifted (of ca. 96 nm) and has a narrower shape as compared to those of the above mentioned particles P-1, P-2 and P-4. These spectroscopic features are the signature of H-aggregation in the solid state for compound 3. Noticeably, P-4, also containing unsymmetrical curcuminoid chromophores, does not adopt such arrangement.

Particles P-2–P-4 fluoresce in the NIR, from 692 to 717 nm (Fig. 4, Table 2). The spectra are considerably red-shifted (superior to 95 nm) as compared to those obtained in solution in the highly polar acetonitrile solvent (Fig. S3–S6, ESI†). Such long emission wavelengths can be attributed to chromophore packing in the solid state, as shown by X-ray diffraction for P-1 and by electronic absorption spectroscopy for the others (Fig. 4). The fluorescence quantum yields of P-1–P-3 are found in the range of 1.5–6.0% which are significant values for aggregated organic dyes emitting in the NIR. Not surprisingly, the H-aggregated compound 3 affords the lowest value of 1.5%. Remarkably, the Φf of dye 4 reaches a substantial value of 12.5% in the solid state, which makes P-4 a very bright solid-state NIR fluorophore (brightness at 700 nm of 6952 M⁻¹ cm⁻¹). Actually, a rather large part of emitted photons by the four compounds have energies below 700 nm because the particle emission spectra are broad. However, it can be highlighted that, when only photons at longer wavelength than 700 nm are integrated, dye 4 remains the most luminescent borondifluoride complex of the series with a NIR luminescence quantum yield of ca. 6.5% while the other three dyes have quantum yields of 4.0, 1.5 and 1.0%, respectively.

In addition, two-photon properties of P-4 could be measured and compared to the previously reported P1 particles. Noticeably, the P2 and P3 particles could not be measured due to the much higher light scattering in those samples which precluded the obtention of reliable data. As observed in DCM solution and for P1 in water, the two-photon maximum of P4 does not overlap the maximum of the one photon absorption S₀–S₁ transition (Fig. 5) but it better matches the S₀–S₁ one (vide supra). This maximum is located at 850 nm with a two-photon cross section of ca. 560 GM. Such two-photon cross section value is 2.5 times higher than that of P1 which, associated to the higher fluorescence quantum yield of P4, results in a much higher two-photon brightness for P4 (more than 5 times greater than the one of P1).

Conclusion

We have described the synthesis of three borondifluoride complexes of curcuminoid derivatives, 2–4. Compound 4 contains two curcuminoid subunits connected via a meta-position of an aryl linkage, and as such it represents a first example of a new class of extended biscurcuminoids. The electrochemical investigations show the interruption of conjugation in 4, the two chromophoric subunits behaving independently. However, the absorption spectrum reveals that an intramolecular excitonic interaction exists in solution, showing that the two chromophoric units are not optically independent. Compound 4 displays a high fluorescence quantum yield, and a good value of the two-photon absorption cross section, which makes it an attractive fluorophore. Compounds 2–4 are fluorescent in the solid state with wavelengths reaching the NIR. Like 1–3, compound 4 experience intermolecular interactions in the condensed phase. However, it exhibits the higher value of fluorescence quantum yield within the series investigated, which may result from the meta-linkage that limits strong π–π stacking with respect to the monochromophoric analogues. Indeed, the unsymmetrical model compound 3 forms H-aggregates, which illustrates the strong propensity of monochromophoric curcuminoids to form tightly packed solid-state arrangements. These findings should help in the design of new fluorophores with improved NIR emitting properties in solution and in the solid state for imaging applications.
Experimental

Material and steady state spectroscopy

Materials and Instrumentation. All solvents for synthesis were of analytic grade. Spectroscopy measurements were carried out with spectroscopic grade solvents. NMR spectra (1H, 13C, 19F) were recorded at room temperature on a BRUKER AC 250 operating at 400, 100, and 425 MHz for 1H, 13C, and 19F, respectively. Data are listed in parts per million (ppm) and are reported relative to tetramethylsilane (1H and 13C); residual solvent peaks of the deuterated solvents were used as internal standards. Mass spectra were realized in Spectropole de Marseille (http://www.spectropole.fr/). Solid state spectra and luminescence quantum yield were measured using an integrating sphere. UV/Vis-absorption spectra were measured on a Varian Cary 50. Emission spectra were measured on a Horiba-Jobin Yvon Fluorolog-3 spectrofluorimeter that was equipped with a three-slit double-grating excitation and a spectrograph emission monochromator with dispersions of 2.1 nm/mm (1200 grooves mm⁻¹). Steady-state luminescence excitation was done using unpolarized light from a 450W xenon CW lamp and detected at an angle of 90° for dilute-solution measurements (10 mm quartz cell) and with a red-sensitive Hamamatsu R928 photomultiplier tube. Special care was taken to correct NIR-emission spectra that were obtained with the latter device. The detector was corrected according to the procedure described by Parker et al.37 The observed photomultiplier output $A_i$ was recorded at a wavelength $\lambda_i$, which corresponds to the apparent emission spectrum. $A_i$ is given by [eqn (1)], where $F_i$ and $S_i$ are the corrected emission spectrum and the spectroscopic sensitivity factor of the monochromator-photomultiplier setup, respectively.

$$A_i = (F_i)(S_i)l^{-2}$$  \hspace{1cm} (1)

To calculate $S_i$, we used 4-N,N-dimethylamino-4’-nitrostilbene (DMANS) as a standard NIR fluorophore for which its corrected emission spectrum has been precisely determined.38 Luminescence quantum yields ($\Phi_0$) were measured in dilute DCM solutions with an absorbance below 0.1 by using [eqn (2)], where OD($\lambda$) is the absorbance at the excitation wavelength ($\lambda$), $n$ the refractive index, and $I$ the integrated luminescence intensity.

$$\Phi_{\text{ir}}/\Phi_{\text{x}} = [\text{OD}((\lambda))/\text{OD}(\lambda)]\text{[I]/[I]}\text{[n]/[n]}l^{-2}$$  \hspace{1cm} (2)

Subscripts “ir” and “x” stand for reference and sample, respectively. The luminescence quantum yields were not corrected by the refractive indices. We used ruthenium trisbipyridine bischloride in water ($\Phi_{\text{ir}} = 0.021$) as a reference for compounds that absorbed in the 450 nm region, while rhodamine B ($\Phi_{\text{ir}} = 0.49$) in EtOH was used for excitation between 540 and 560 nm.

Lifetime measurements were carried out on a HORIBA Jobin Yvon IBH FluoroLog-3 spectrofluorimeter that was adapted for time-correlated single-photon counting. For these measurements, pulsed LEDs with an appropriate wavelength were used. Emission was monitored perpendicular to the excitation pulse and spectroscopic selection was achieved by a passage through the spectrograph. A thermoelectrically cooled single-photon-detection module (HORIBA Jobin Yvon IBH, TBX-04-D) incorporating a fast-rise-time photomultiplier tube, a wide-bandwidth pre-amplifier, and a picosecond-constant fraction discriminator was used as the detector. Signals were acquired using an IBH DataStation Hub photon counting module and data analyses were performed by using the commercially available DAS 6 decay-analysis software package from HORIBA Jobin Yvon IBH; the reported $\tau$ values are given with an estimated uncertainty of about 10%.

Cyclic voltammetric (CV) data were acquired using a BAS 100 Potentiostat (Bioanalytical Systems) and a PC computer containing BAS100W software (v2.3). A three-electrode system with a Pt working electrode (diameter 1.6 mm), a Pt counter electrode and an Ag/AgCl (with 3 M NaCl filling solution) reference electrode was used. [nBu][N]PF$_6$ (0.1 M in dichloromethane) served as an inert electrolyte. Cyclic voltammograms were recorded at a scan rate of 100 mV s⁻¹. Ferrocene was used as internal standard.39

Synthetic procedure

Dye 1, its ligand (Lig 1)27 the ligand of 2 (Lig 2)40 2,4,6-tris(octyloxy)sophthalaldehyde41 and (1E,4Z)-5-hydroxy-1-[4-methoxyphenyl]hexa-1,4-dien-3-one (Lig 5)28 were prepared as previously reported.

General synthesis of Lig 1 and Lig 2. In a 50 mL round bottom flask, a solution of the acetylacetone derivative (1 mol eq.) and B$_2$O$_3$ (0.5 mol eq.) dissolved in ethyl acetate (15 mL) was stirred at 60 °C for 30 min. A solution of the appropriate aldehyde (1 mol eq.) and tri(n-butyl)borane (1 mol eq.) in ethyl acetate (10 mL) was added and the resulting mixture was stirred for 30 min at 60 °C. A catalytic amount of n-butylamine (0.4 mol eq.) was then added and the reaction mixture was refluxed overnight. After cooling to 60 °C, 30 mL of 0.4 M HCl were added and the mixture stirred for 30 min. After cooling, the precipitate was filtered off and dried in vacuo to yield pure ligands Lig 1 and Lig 2.

Synthesis of 5 ([1E,4Z]-5-hydroxy-1-[4-methoxyphenyl]hexa-1,4-dien-3-one). In a 50 mL round bottom flask, a solution of acetylacetonate (3.00 g, 30 mmol) and B$_2$O$_3$ (1.050 g, 15 mmol) dissolved in ethyl acetate (10 mL) was stirred at 60 °C for 30 min. A solution of anisaldehyde (1.36 g, 10 mmol) and tri(n-butyl)borane (2.30 g, 10 mmol) in ethyl acetate (8 mL) was added and the mixture was stirred for 30 min at 60 °C. A catalytic amount of n-butylamine (0.44 g, 6 mmol) was then added to the solution and the reaction mixture was refluxed overnight. After cooling to 60 °C, 30 mL of 0.4 M HCl were added and the mixture was stirred for 30 min. After cooling, the precipitate was filtered off (Lig 1). The residual solution was evaporated. The oily compound was dissolved in dichloromethane. The organic layer was washed with water, brine, dried over MgSO$_4$ and evaporated to dryness. The oily crude was purified by column chromatography on silica using a mixture of cyclohexane and dichloromethane (gradient from 1/1 to 3/1) yielding the pure 5 as a yellowish solid (1.20 g, 55%).

Synthesis of Lig 3 and Lig 4. In a 50 mL round bottom flask, a solution of 5 (1 mol eq.) and B$_2$O$_3$ (0.5 mol eq.) dissolved in
ethyl acetate (10 mL) was stirred at 60 °C for 30 min. A solution of the appropriate aldehyde (1 mol eq. of aldehyde function) and tri(n-butyl)borane (1 mol eq. of aldehyde function) in ethyl acetate (10 mL) was added and the mixture was stirred for 30 min at 60 °C. A catalytic amount of n-butylamine (0.5 mol eq.) was then added to the solution and the reaction mixture was refluxed overnight. After cooling to 60 °C, 30 mL of 0.4 M HCl were added and the mixture and stirred for 30 min. After cooling, the precipitate was filtered off and dried in vacuo to yield the pure ligands Lig 3 and Lig 4.

Lig 3: ([(E,1E,4Z,6E)-5-hydroxy-(4-methoxyphenyl)-1-(2,4,6-trimethoxyphenyl)hepta-1,4,6-trien-3-one]). Orange solid; yield: 85%; 1H NMR (400 MHz, CDCl3, ppm): δ = 8.72 (d, 3J = 15.7 Hz, 1H), 7.06 (d, 3J = 15.8 Hz, 2H), 6.99 (d, 3J = 15.8 Hz, 1H), 6.47 (s, 2H), 6.35 (s, 1H), 3.95 (s, 6H), 3.90 (s, 3H), 3.84 (s, 3H); satisfactory 13C NMR spectra could not be obtained; 19F NMR (235 MHz, CDCl3): δ = −141.59 (10B-F, 0.2), −141.65 ppm (11B-F, 0.8). HRMS [ESI] M + Na+ for C26H22OBF2Na+ m/z = 467.1452, found m/z = 467.1453.

4: ([(E,1E,4Z,6E,6′E)-1,1′-(2,4,6-tris(octyloxy)-1,3-phenylene)bis(5-(difluoroboryloxy)-7-(4-methoxyphenyl)hepta-1,4,6-trien-3-one]). Red solid; yield: 97%; 1H NMR (400 MHz, DMSO-d6, ppm): δ = 8.25 (d, 3J = 16 Hz, 2H), 7.97 (d, 3J = 15.5 Hz, 2H), 7.55 (d, 3J = 8.5 Hz, 4H), 7.14 (d, 3J = 16 Hz, 2H), 6.93 (d, 3J = 8.5 Hz, 4H), 6.54 (d, 3J = 15.5 Hz, 2H), 2.62 (s, 1H), 5.56 (s, 1H), 4.14 (t, 3J = 6.5 Hz, 4H), 3.85 (s, 6H), 3.80 (t, 3J = 6.5 Hz, 2H), 1.92 (m, 6H), 1.32 (m, 30H), 0.90 ppm (m, 9H); satisfactory 13C NMR spectra could not be obtained; 19F NMR (235 MHz, DMSO): δ = −140.91 (10B-F, 0.2), −140.97 ppm (11B-F, 0.8). HRMS [ESI] M + 2Na+ for C58H26O8BF2Na2 m/z = 530.2707, found m/z = 530.2705.

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