



Cite this: *New J. Chem.*, 2016,
40, 1297

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

Morgane Rivoal, Elena Zaborova, Gabriel Canard, Anthony D'Aléo* and
Frédéric Fages

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 ($B^1 = 87\,000\text{ M}^{-1}\text{ cm}^{-1}$ 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.

Received (in Montpellier, France)
14th April 2015,
Accepted 3rd August 2015

DOI: 10.1039/c5nj00925a

www.rsc.org/njc

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 acetylacetonate ligand have shown

interesting properties such as high two-photon absorption cross sections,^{13,14} mechano-fluorochromic behaviors^{15,16} and efficient NIR emissions^{17,18} that led to their use for cells imaging¹⁹ or as sensors of volatile acid/base,^{20,21} fluorescent reporters for amyloid,^{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 BF_2 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 (Φ_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)s 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

Aix Marseille Université, CNRS, CINaM UMR 7325, Campus de Luminy, Case 913,
13288 Marseille, France. E-mail: daleo@cinam.univ-mrs.fr

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c5nj00925a

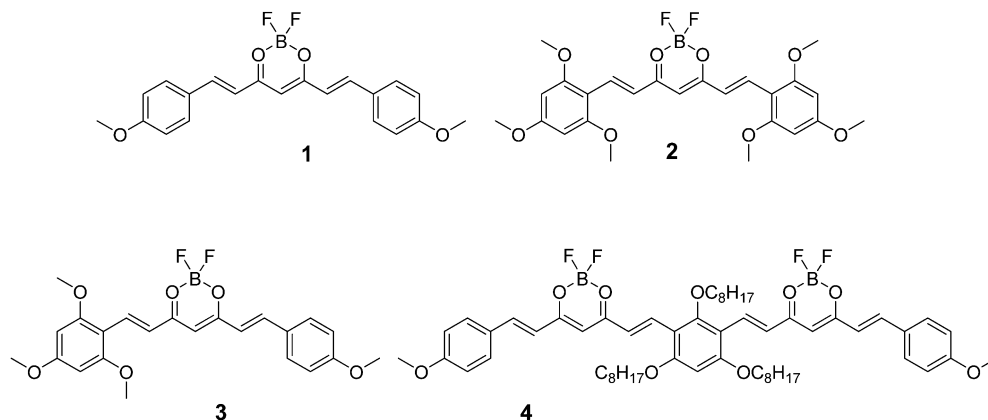


Chart 1 Molecular structures of the borondifluoride curcuminoid derivatives.

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.

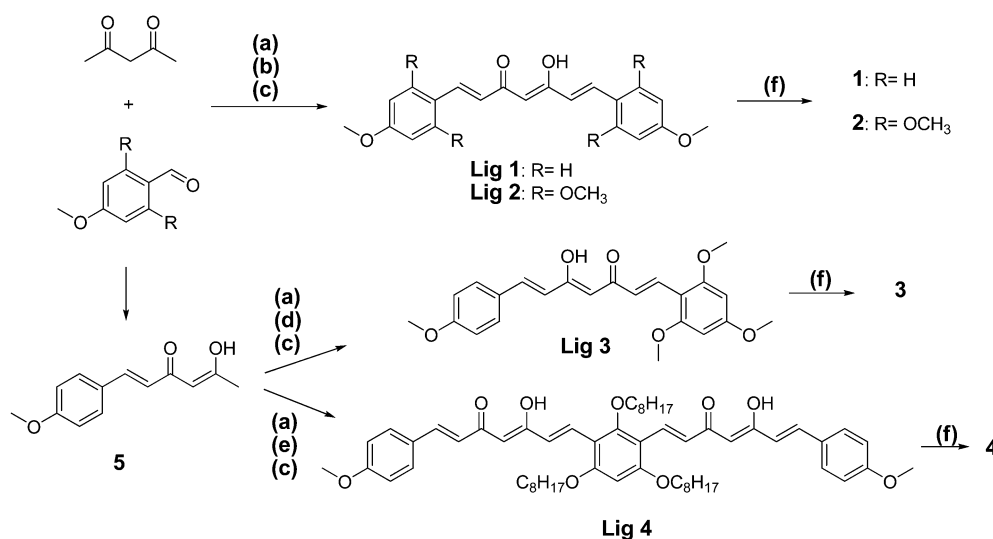
Results and discussion

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.²⁷ 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,²⁸ this intermediate was prepared by a Knoevenagel reaction using an excess of acetylacetone (acac/aldehyde 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).



Scheme 1 Syntheses of the borondifluoride curcuminoid derivatives **1**, **2**, **3** and **4**. (a) B₂O₃, ethylacetate, 60 °C; (b) anisaldehyde, tri-*n*-butylborane, ethylacetate, 60 °C; (c) *n*-butylamine, ethylacetate, 80 °C, then HCl, 60 °C; (d) 2,4,6-trimethoxybenzaldehyde, tri-*n*-butylborane, ethylacetate, 60 °C; (e) 2,4,6-tris(octyloxy)isophthalaldehyde, tri-*n*-butylborane, ethylacetate, 60 °C; (f) BF₃·Et₂O, DCM, reflux.



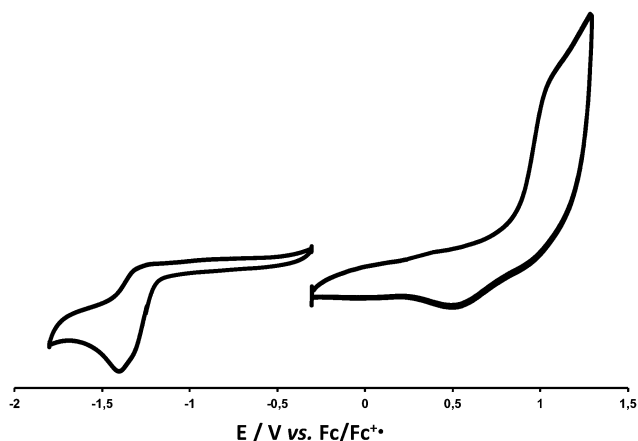


Fig. 1 Cyclic voltammogram of the bis borondifluoride complex **4** in DCM solution containing 0.1 M [$(n\text{-Bu}_4\text{N})\text{PF}_6$] (scan rate of 100 mV s^{-1}).

Table 1 Oxidation and reduction half-wave potential values for the studied dioxaborine derivatives (10 μM) in dichloromethane solution vs. $\text{Fc}/\text{Fc}^{+\bullet}$ using tetrabutylammonium hexafluorophosphate as electrolyte (100 μM), the values are given in volt

	$E_{1/2}^{\text{red}}$	$(E_{1/2}^{\text{ox}})_1$	$(E_{1/2}^{\text{ox}})_2$	ΔE_g^a
1	−1.27	1.10	— ^b	2.37
2	−1.44	0.72	1.09	2.16
3	−1.37	0.89	1.31	2.26
4	−1.30	1.05	— ^b	2.35

^a Electrochemical HOMO–LUMO gap $\Delta E_g = (E_{1/2}^{\text{ox}})_1 - E_{1/2}^{\text{red}}$. ^b 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 of $(n\text{-Bu})_4\text{NPF}_6$. 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 $\pi\text{--}\pi^*$ transition. A second electronic transition band of much lower intensity appears as a shoulder at higher energy ($<400\text{ nm}$, *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 three other dyes. The more complex shape and the larger Stoke shift of the absorption band is likely to stem 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_f (61%) is obtained for complex **4**, giving a high brightness value of ca. $87\,000\text{ M}^{-1}\text{ cm}^{-1}$.

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[†]).²⁹

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



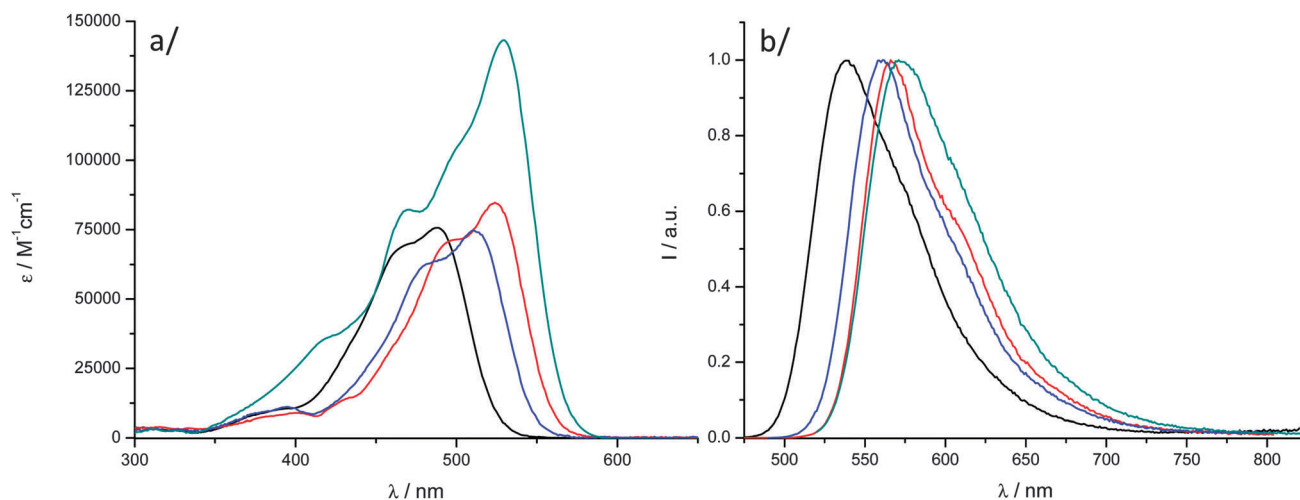


Fig. 2 (a) Overlaid electronic absorption spectra and (b) overlaid corrected normalized fluorescence emission spectra (conc. $\approx 10^{-6}$ M. λ_{exc} at the absorption maximum) of **1** (—), **2** (—), **3** (—) and **4** (—) recorded in DCM at room temperature.

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 solvato-kinetics) then decreases (positive solvato-kinetics), a maximum being observed. The positive solvato-kinetic 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.³⁰ 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 photo-degradation 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,¹⁴ **1** has a TPA cross section (σ^{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 σ^{TPA} value of ca. 248 GM. It is interesting to note that **3** presents an intermediate σ^{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 curcumin³¹ and borondifluoride complexes of curcuminoids,¹⁴ the TPA band of the four complexes

Table 2 Spectroscopic data and photophysical properties of all borondifluoride compounds solvated in DCM and as particles in water at room temperature^a

Compound	UV-vis		Fluorescence							TPEF		
	λ_{abs}	ϵ_{max}	λ_{em}	$\Delta\nu_{\text{ST}}$	Φ_f	B^1	τ_f	k_f	k_{nr}	λ_{max}^2	σ^{TPA}	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	2.3	800	513	313
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	—	—	— ^d	— ^d	— ^d
							2.91 ^c					
3 (water)	407	29 500	711	10 505 ^b	0.015	443	2.13	—	—	— ^d	— ^d	— ^d
							6.01 ^c					
4 (water)	468	55 620	692	6917 ^b	0.125	6952	1.89	—	—	850	560	70
							6.64 ^c					

^a Absorption maximum wavelengths λ_{abs} (nm); molar absorption coefficients at maximum ϵ_{max} ($\text{M}^{-1} \text{cm}^{-1}$); fluorescence maximum wavelengths λ_{em} (nm); Stokes shifts $\Delta\nu_{\text{ST}}$ (cm^{-1}); fluorescence quantum yields Φ_f ; brightness $B = \Phi_f \times \epsilon$ ($\text{M}^{-1} \text{cm}^{-1}$); fluorescence lifetimes τ_f (ns); radiative k_f (10^8 s^{-1}) and nonradiative $k_{nr} = (1 - \Phi_f)/\tau_f$ (10^8 s^{-1}) rate constants; two-photon absorption maximum λ_{max}^2 (nm); two-photon cross section σ^{TPA} (GM); two-photon brightness $B^2 = \Phi_f \times \sigma^{\text{TPA}}$ (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.



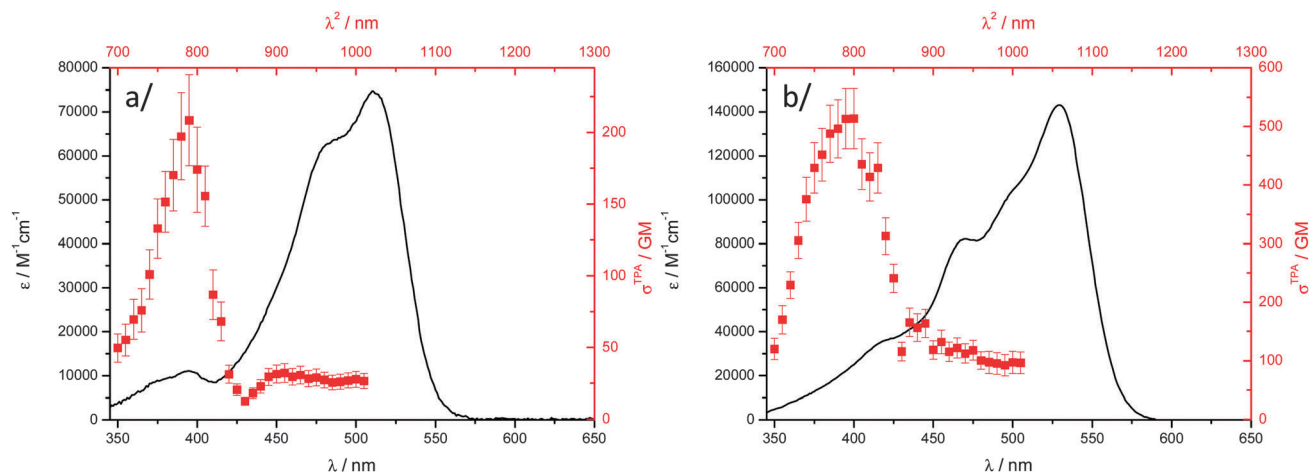


Fig. 3 Two-photon excitation (■, higher x-coordinate and —, right y-coordinate) with their error bars, OPA spectra (—, lower x-coordinate and —, left y-coordinate) in DCM: (a) dye **3** and (b) dye **4**.

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: σ^{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_1 transition was still observed.³¹

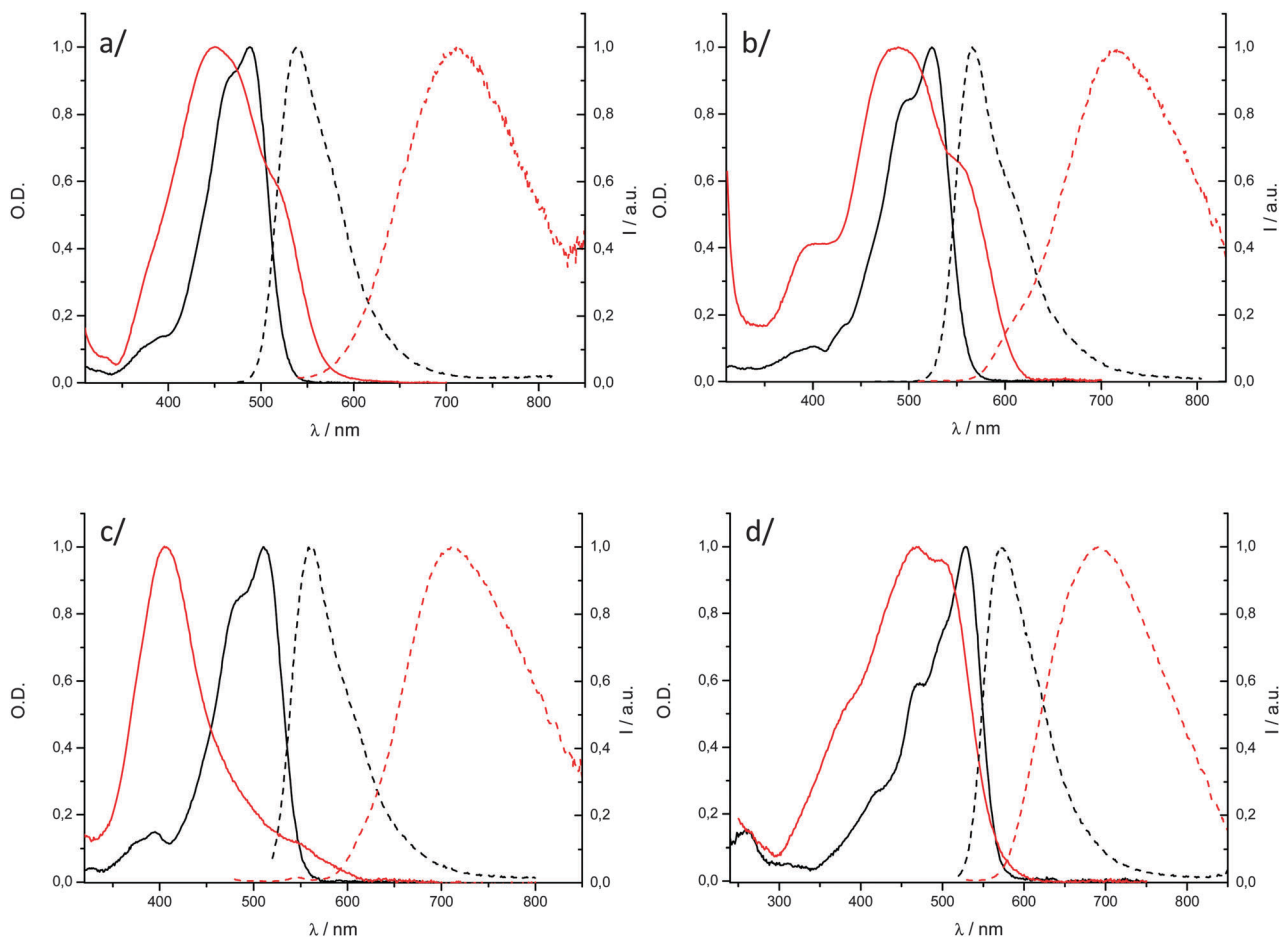


Fig. 4 Overlaid absorption (solid lines) and emission (dashed lines) spectra of DCM solutions (black) and particles in water (red) for (a) **1**; (b) **2**; (c) **3** and (d) **4**.



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.³⁶ 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 publication¹⁴ 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 Φ_f 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 **P4** 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

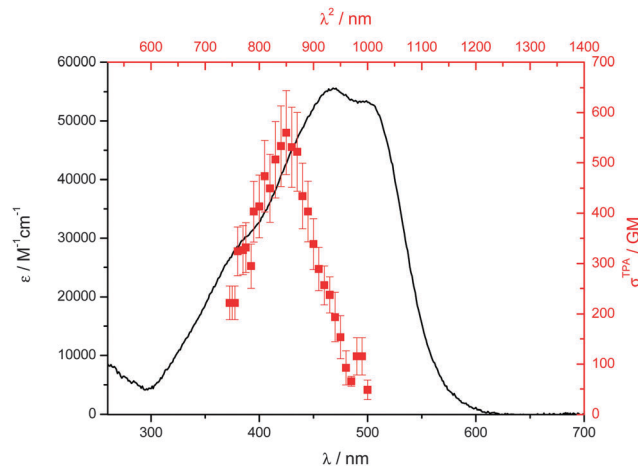


Fig. 5 Two-photon excitation (■, higher x-coordinate and —, right y-coordinate) with their error bars, OPA spectra (—, lower x-coordinate and —, left y-coordinate) of **P4** in water.

transition (Fig. 5) but it better matches the S₀–S₂ 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 , ^{13}C , ^{19}F) were recorded at room temperature on a BRUKER AC 250 operating at 400, 100, and 425 MHz for ^1H , ^{13}C , and ^{19}F , respectively. Data are listed in parts per million (ppm) and are reported relative to tetramethylsilane (^1H and ^{13}C); 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-JobinYvon 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^{-1} ($1200 \text{ grooves mm}^{-1}$). 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_1 was recorded at a wavelength λ , which corresponds to the apparent emission spectrum. A_1 is given by [eqn (1)], where F_1 and S_1 are the corrected emission spectrum and the spectroscopic sensitivity factor of the monochromator-photomultiplier setup, respectively.

$$A_1 = (F_1)(S_1)/\lambda^2 \quad (1)$$

To calculate S_1 , 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 (Φ_f) were measured in dilute DCM solutions with an absorbance below 0.1 by using [eqn (2)], where $\text{OD}(\lambda)$ is the absorbance at the excitation wavelength (λ), n the refractive index, and I the integrated luminescence intensity.

$$\Phi_{\text{fx}}/\Phi_{\text{fr}} = [\text{OD}_r(\lambda)/\text{OD}_x(\lambda)][I_x/I_r][n_x/n_r]^2 \quad (2)$$

Subscripts "r" 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 ($\Phi_{\text{fr}} = 0.021$) as a reference for compounds that absorbed in the 450 nm region, while rhodamine B ($\Phi_{\text{fr}} = 0.49$) 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 τ 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. $[(n\text{Bu})_4\text{N}]\text{PF}_6$ (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)isophthalaldehyde⁴¹ and (1*E*,4*Z*)-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 acetylacetonate derivative (1 mol eq.) and B_2O_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 ((1*E*,4*Z*)-5-hydroxy-1-(4-methoxyphenyl)hexa-1,4-dien-3-one). In a 50 mL round bottom flask, a solution of acetylacetone (3.00 g, 30 mmol) and B_2O_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_2O_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 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: ((1*E*,4*Z*,6*E*)-5-hydroxy-7-(4-methoxyphenyl)-1-(2,4,6-trimethoxyphenyl)hepta-1,4,6-trien-3-one). Orange solid; yield: 65%; ¹H NMR (400 MHz, CDCl₃, ppm): δ = 8.07 (d, ³*J* = 16.1 Hz, 1H), 7.57 (d, ³*J* = 15.8 Hz, 1H), 7.48 (d, ³*J* = 8.7 Hz, 2H), 6.98 (d, ³*J* = 16.1 Hz, 1H), 6.89 (d, ³*J* = 8.7 Hz, 2H), 6.48 (d, ³*J* = 15.8 Hz, 1H), 6.10 (s, 2H), 5.77 (s, 1H), 3.87 (s, 6H), 3.83 (s, 3H), 3.82 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ = 186.01, 182.18, 162.97, 161.47, 161.13, 139.27, 132.17, 129.69, 128.13, 123.98, 122.24, 114.41, 106.57, 101.50, 90.57, 55.85, 55.49, 55.47. HRMS (ESI⁺) [M + H]⁺ calcd for C₂₃H₂₅O₆⁺ *m/z* = 397.1651, found *m/z* = 397.1650.

Lig 4: ((1*E*,1'*E*,4*Z*,4'*Z*,6*E*,6'*E*)-1,1'-(2,4,6-tris(octyloxy)-1,3-phenylene)bis(5-hydroxy-7-(4-methoxyphenyl)hepta-1,4,6-trien-3-one)). Dark orange solid; yield: 34%; ¹H NMR (400 MHz, CDCl₃, ppm): δ = 7.92 (d, ³*J* = 16.1 Hz, 2H), 7.60 (d, ³*J* = 15.8 Hz, 2H), 7.49 (d, ³*J* = 8.7 Hz, 4H), 7.04 (d, ³*J* = 16.1 Hz, 2H), 6.89 (d, ³*J* = 8.7 Hz, 4H), 6.48 (d, ³*J* = 15.8 Hz, 2H), 6.23 (s, 1H), 5.69 (s, 2H), 4.05 (t, ³*J* = 6.2 Hz, 4H), 3.83 (m, 6H), 3.77 (t, ³*J* = 6.3 Hz, 4H), 1.88 (m, 6H), 1.53 (m, 6H), 1.30 (m, 24H), 0.87 (m, 9H); ¹³C NMR (100 MHz, CDCl₃): δ = 184.09, 183.78, 161.66, 161.23, 139.80, 131.86, 129.78, 128.00, 125.68, 122.29, 114.41, 111.44, 101.76, 92.63, 69.04, 55.47, 31.97, 29.68, 29.53, 29.44, 29.17, 26.42, 26.36, 22.81, 14.23. HRMS (ESI⁺) [M + 2H]²⁺ calcd for C₅₈H₈₀O₈⁺ *m/z* = 460.2901, found *m/z* = 460.2903.

General procedure for borondifluoride complexes

In a 50 mL round bottom flask, the ligand (1 mol eq.) was solubilized in dichloromethane (20 mL) before boron trifluoride etherate (1.1 mol eq.) was added. The reaction mixture was refluxed overnight. After cooling to room temperature, the solvent was evaporated and the resulting solid was suspended in diethyl ether. The precipitate was filtered off yielding the pure complex except for **2**. For the latter dye, the crude complex was purified by column chromatography using silica gel with a mixture of cyclohexane and dichloromethane (1/4).

2: ((1*E*,4*Z*,6*E*)-5-(difluoroboryloxy)-1,7-bis(2,4,6-trimethoxyphenyl)hepta-1,4,6-trien-3-one). Purple solid; yield: 52%; ¹H-NMR (400 MHz, (CD₃)₂CO, ppm): δ = 8.41 (d, ³*J* = 15.7 Hz, 2H), 7.24 (d, ³*J* = 15.7 Hz, 2H), 6.34 (s, 4H), 6.20 (s, 1H), 3.98 (s, 12H), 3.92 (s, 6H); Satisfactory ¹³C NMR spectra could not be obtained; ¹⁹F-NMR ((CD₃)₂CO, 235 MHz): δ = -141.77 (¹⁰B, 0.2F), -142.87 (¹¹B, 0.8F). HRMS (ESI⁺) [M + Na]⁺ calcd for C₂₅H₂₇O₈BF₂Na⁺ *m/z* = 527.1664, found *m/z* = 527.1661.

3: ((1*E*,4*Z*,6*E*)-5-(difluoroboryloxy)-7-(4-methoxyphenyl)-1-(2,4,6-trimethoxyphenyl)hepta-1,4,6-trien-3-one). Red solid; yield: 85%; ¹H NMR (400 MHz, DMSO-*d*₆, ppm): δ = 8.30 (d, ³*J* = 15.7 Hz, 1H), 7.90 (d, ³*J* = 15.7 Hz, 1H), 7.80 (d, ³*J* = 8.8 Hz, 2H),

7.16 (d, ³*J* = 15.7 Hz, 1H), 7.06 (d, ³*J* = 8.8 Hz, 2H), 6.99 (d, ³*J* = 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]⁺ calcd for C₂₃H₂₃O₆BF₂Na⁺ *m/z* = 467.1452, found *m/z* = 467.1453.

4: ((1*E*,1'*E*,4*Z*,4'*Z*,6*E*,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, ppm): δ = 8.25 (d, ³*J* = 16 Hz, 2H), 7.97 (d, ³*J* = 15.5 Hz, 2H), 7.55 (d, ³*J* = 8.5 Hz, 4H), 7.14 (d, ³*J* = 16 Hz, 2H), 6.93 (d, ³*J* = 8.5 Hz, 4H), 6.54 (d, ³*J* = 15.5 Hz, 2H), 6.26 (s, 1H), 5.56 (s, 1H), 4.14 (t, ³*J* = 6.5 Hz, 4H), 3.85 (s, 6H), 3.80 (t, ³*J* = 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]²⁺ calcd for C₅₈H₇₆O₉B₂F₄Na₂²⁺ *m/z* = 530.2707, found *m/z* = 530.2705.

Acknowledgements

The authors want to thank Dr Wladimir Marine for his help in setting up the two-photon excitation set-up.

References

- 1 Z. Guo, S. Park, J. Yoon and I. Shin, *Chem. Soc. Rev.*, 2014, **43**, 16–29.
- 2 A. B. Ormond and H. S. Freeman, *Materials*, 2013, **6**, 817–840.
- 3 S. S. Kelkar and T. M. Reineke, *Bioconjugate Chem.*, 2011, **22**, 1879–1903.
- 4 P.-A. Bouit, G. Wetzel, G. R. Berginc, B. Loiseaux, L. C. Toupet, P. Feneyrou, Y. Bretonnière, K. Kamada, O. Maury and C. Andraud, *Chem. Mater.*, 2007, **19**, 5325–5335.
- 5 M. Liang and J. Chen, *Chem. Soc. Rev.*, 2013, **42**, 3453–3488.
- 6 A. Zitzler-Kunkel, M. R. Lenze, N. M. Kronenberg, A.-M. Krause, M. Stolte, K. Meerholz and F. Würthner, *Chem. Mater.*, 2014, **26**, 4856–4866.
- 7 A. Mishra and P. Bäuerle, *Angew. Chem., Int. Ed.*, 2012, **51**, 2020–2067.
- 8 A. Loudet and K. Burgess, *Chem. Rev.*, 2007, **107**, 4891–4932.
- 9 G. Ulrich, R. Ziessel and A. Harriman, *Angew. Chem., Int. Ed.*, 2008, **47**, 1184–1201.
- 10 R. Ziessel, G. Ulrich and A. Harriman, *New J. Chem.*, 2007, **31**, 496–501.
- 11 N. Boens, V. Leen and W. Dehaen, *Chem. Soc. Rev.*, 2012, **41**, 1130–1172.
- 12 A. D'Aléo and F. Fages, *Disp. Imaging*, 2014, **2**, 149–174.
- 13 E. Cogné-Laage, J.-F. Allemand, O. Ruel, J.-B. Baudin, V. Croquette, M. Blanchard-Desce and L. Jullien, *Chem. – Eur. J.*, 2004, **10**, 1445–1455.
- 14 A. D'Aleo, A. Felouat, V. Heresanu, A. Ranguis, D. Chaudanson, A. Karapetyan, M. Giorgi and F. Fages, *J. Mater. Chem. C*, 2014, **2**, 5208–5215.



- 15 N. D. Nguyen, G. Zhang, J. Lu, A. E. Sherman and C. L. Fraser, *J. Mater. Chem.*, 2011, **21**, 8409–8415.
- 16 G. Zhang, J. P. Singer, S. E. Kooi, R. E. Evans, E. L. Thomas and C. L. Fraser, *J. Mater. Chem.*, 2011, **21**, 8295–8299.
- 17 A. D'Aléo, D. Gachet, V. Heresanu, M. Giorgi and F. Fages, *Chem. – Eur. J.*, 2012, **18**, 12764–12772.
- 18 A. D'Aléo, V. Heresanu, M. Giorgi, B. Le Guennic, D. Jacquemin and F. Fages, *J. Phys. Chem. C*, 2014, **118**, 11906–11918.
- 19 G. Bai, C. Yu, C. Cheng, E. Hao, Y. Wei, X. Mu and L. Jiao, *Org. Biomol. Chem.*, 2014, **12**, 1618–1626.
- 20 X. Cheng, D. Li, Z. Zhang, H. Zhang and Y. Wang, *Org. Lett.*, 2014, **16**, 880–883.
- 21 J. Hu, Z. He, Z. Wang, X. Li, J. You and G. Gao, *Tetrahedron Lett.*, 2013, **54**, 4167–4170.
- 22 C. Ran, X. Xu, S. B. Raymond, B. J. Ferrara, K. Neal, B. J. Bacskaï, Z. Medarova and A. Moore, *J. Am. Chem. Soc.*, 2009, **131**, 15257–15261.
- 23 X. Zhang, Y. Tian, Z. Li, X. Tian, H. Sun, H. Liu, A. Moore and C. Ran, *J. Am. Chem. Soc.*, 2013, **135**, 16397–16409.
- 24 A. Pfister, G. Zhang, J. Zareno, A. F. Horwitz and C. L. Fraser, *ACS Nano*, 2008, **2**, 1252–1258.
- 25 S. Chambon, A. D'Aléo, C. Baffert, G. Wantz and F. Fages, *Chem. Commun.*, 2013, **49**, 3555–3557.
- 26 A. M. Sarker, E. E. Gürel, M. Zheng, P. M. Lahti and F. E. Karasz, *Macromolecules*, 2001, **34**, 5897–5901.
- 27 A. Felouat, A. D'Aléo and F. Fages, *J. Org. Chem.*, 2013, **78**, 4446–4455.
- 28 G. Mann, L. Beyer and A. Arrieta, *Z. Chem.*, 1987, **27**, 172–173.
- 29 Z. R. Grabowski, K. Rotkiewicz and W. Rettig, *Chem. Rev.*, 2003, **103**, 3899–4031.
- 30 C. Xu and W. W. Webb, *J. Opt. Soc. Am. B*, 1996, **13**, 481–491.
- 31 J. A. Tiburcio-Moreno, J. J. Alvarado-Gil, C. Diaz, L. Echevarria and F. E. Hernández, *Chem. Phys. Lett.*, 2013, **583**, 160–164.
- 32 H. Hu, O. V. Przhonska, F. Terenziani, A. Painelli, D. Fishman, T. R. Ensley, M. Reichert, S. Webster, J. L. Bricks, A. D. Kachkovski, D. J. Hagan and E. W. Van Stryland, *Phys. Chem. Chem. Phys.*, 2013, **15**, 7666–7678.
- 33 W. M. McClain, *Acc. Chem. Res.*, 1974, **7**, 129–135.
- 34 F. Terenziani, A. Painelli, C. Katan, M. Charlot and M. Blanchard-Desce, *J. Am. Chem. Soc.*, 2006, **128**, 15742–15755.
- 35 Q. Bellier, N. S. Makarov, P.-A. Bouit, S. Rigaut, K. Kamada, P. Feneyrou, G. Berginc, O. Maury, J. W. Perry and C. Andraud, *Phys. Chem. Chem. Phys.*, 2012, **14**, 15299–15307.
- 36 H. Kawai, H. S. Nalwa, H. Oikawa, S. Okada, H. Matsuda, N. Minami, A. Kakuda, K. Ono, A. Mukoh and H. Nakanishi, *Jpn. J. Appl. Phys.*, 1992, **31**, L1132–L1134.
- 37 C. A. Parker, *Photoluminescence of Solutions*, Elsevier Publishing, Amsterdam, 1969.
- 38 J. R. Lakowicz, *Principles of Fluorescence Spectroscopy*, Kluwer, NewYork (NY), 2006.
- 39 N. G. Connelly and W. E. Geiger, *Chem. Rev.*, 1996, **96**, 877–910.
- 40 Y. Zuo, J. Huang, B. Zhou, S. Wang, W. Shao, C. Zhu, L. Lin, G. Wen, H. Wang, J. Du and X. Bu, *Eur. J. Med. Chem.*, 2012, **55**, 346–357.
- 41 S. B. Bharate, R. Mudududdla, R. Sharma and R. A. Vishwakarma, *Tetrahedron Lett.*, 2013, **54**, 2913–2915.

