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To cite this version:
Sebastien Thiery, Céline Declairieux, Denis Tondelier, Gijun Seo, Bernard Geffroy, et al.. 2-Substituted vs 4-substituted-9,9’-spirobifluorene host materials for green and blue phosphorescent OLEDs: a structure-property relationship study. Tetrahedron, Elsevier, 2014, 70 (36), pp.6337 - 6351. 10.1016/j.tet.2014.05.081. hal-01072178

HAL Id: hal-01072178
https://hal-univ-rennes1.archives-ouvertes.fr/hal-01072178
Submitted on 7 Oct 2014

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2-substituted vs 4-substituted-9,9'-spirobisfluorene host materials for green and blue phosphorescent OLEDs: A Structure-Property Relationship Study

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Abstract
We report a structure-property relationship study of four 9,9'-spirobisfluorene (SBF) derivatives (4-5Pm-SBF, 2-5Pm-SBF, 4-Ph-SBF and 2-Ph-SBF), substituted with either phenyl or pyrimidine at the C2 or C4 position of the SBF core. Structural, thermal, electrochemical and photophysical properties have been examined and correlated to theoretical calculations in order to study the influence of the nature and the position of the substituent. The emission properties of 4- vs 2-substituted SBFs are noticeably different highlighting, in the excited state, the remarkable effect of substitution in ortho position of SBF. All compounds have been used as host material for green dopant in PhOLEDs with very high performances (2-5Pm-SBF: CE > 58 cd/A, PE > 35 lm/W, EQE > 14 %). More importantly, the two 4-substituted SBFs have been used as host materials in blue PhOLEDs, displaying high performance and a decrease of \(V_{TH}\) for 4-5Pm-SBF due to the incorporation of the electron-withdrawing pyrimidine.

Graphical Abstract

Introduction
In the past twenty years, the use of organic semiconducting materials for applications in “plastic electronics”, such as field effect transistors (FETs), organic light emitting diodes (OLEDs), and organic photovoltaic (OPVs), has gained fantastic interest. Particularly, OLEDs have attracted much scientific and commercial interest because of their potential applications in organic flat-panel displays and solid-state lighting. However, as full colour displays and white lighting require materials that emit over a broad range of colour, the weak link of the technology remains the lack of efficient and stable blue light emitters. Moreover, due to the triplet exciton/singlet exciton ratio of 3/1, the maximum achievable emission quantum yield is limited to 25% for fluorescent emitters. Among the significant progress recently made, phosphorescent organic emitting-diodes (PHOLEDs) which use heavy-metal complexes as emitters to harvest both singlet and triplet excitons are considered as the most promising technology because, theoretically the internal quantum efficiencies of these devices may reach 100%. The fabrication of blue phosphorescent devices is however still challenging because the prerequisites for designing organic host materials are (i) a high triplet energy level (higher than \( E_g \) of the blue dopant), (ii) a high glass-transition temperature (Tg) and (iii) matching HOMO/LUMO levels for hole and electron injection/transport. In this context, most of host materials for blue PHOLEDs are based on carbazole, triphenylamines, phosphine oxides or fluorene derivatives. However, novel host materials with good thermal and morphological stabilities to afford blue PHOLEDs with high efficiencies, stability and colour purity are still desirable. In this context, 9,9'-spirobifluorene (SBF) based semiconductors have experienced a fantastic development during the last thirty years, due to their use as chiral ligand, as electropolymerizable building block, as solid state laser, in third-order nonlinear optics, in Metal Organic Frameworks but more importantly in organic electronics, leading to fantastic breakthroughs in the field. Thus, numbers of electron donating and /or withdrawing functionalities have been introduced on the SBF scaffold in order to make the electronic properties of the resulting dyes fitting with an application. For that purpose, electron-deficient heterocycles such as 1,3,4-oxadiazoles, pyridine, quinoline, quinoxaline and pyridopyrazine connected at C2 and C7 positions of SBF were reported in literature. Among those electron-deficient heterocycles, only few examples describing the fluorene substitution by pyrimidine units (see chart 1) have been reported to date. However, pyrimidine possesses many striking characteristics, as it is notably more electron-deficient than pyridine, which has been widely studied. Thus Bryce’s group has for example demonstrated in 2003, that fluorene fluorophore F3 emits a pure blue emission in solution and thin films. However the electroluminescent spectrum of F3-based OLED mainly presents the known parasite green emission of dialkyfluoranes. A similar green emission is observed on PL spectrum of the liquid crystal F4. The introduction of a 9,9’ Spirobifluorene unit (SBF) in TBPSF leads to improve the thermal, optical and electronic properties and this new blue emitter has been used as host for fluorescent blue, red or green organic dopants leading to efficient devices.

![Chart 1](image)

**Chart 1.** Literature report of fluorene substitution by pyrimidine units
However, if 2-substituted SBFs have been impressively investigated for the last decades, it is far to be the case of the three other positional isomers, that are 1-, 3- or 4-substituted SBFs. Thus, for example, 4-substituted SBFs have only been reported in a few instances but appears as a very appealing scaffold to design rigid SBF-based organic host materials for PhOLEDs. In our quest for robust derivatives with high $E_I$, which can be used as host materials for green and blue phosphorescent dyes, we wish to report herein the synthesis and the detailed study of the properties of two pyrimidine-substituted SBFs (4-5Pm-SBF and 2-5Pm-SBF, see chart 2). Moreover, in order to give a detailed description of the influence of the pyrimidine unit, the 2- and 4-phenyl-substituted SBFs (2-Ph-SBF and 4-Ph-SBF) have been also studied in detail (4-Ph-SBF has been previously reported). Thus, after synthetic considerations, the electrochemical, thermal and photo-physical properties will be discussed and coupled to a theoretical approach. The influence of the nature of the substituent (pyrimidine vs phenyl) and of its position (2- vs 4-) will be notably carefully investigated. Finally, the incorporation of these four semiconductors as host material for green and blue PhOLEDs will be presented.

![Chart 2. Molecules studied in this work](image)

Results and discussion

The first step towards the synthesis of 4- and 2-substituted SBF was the access of the two key platforms 4- and 2-bromo-9,9'-spirobifluorene (see structures of 4-Br-SBF and 2-Br-SBF in scheme 1). As reported previously, 4-Br-SBF was synthesized by the mono lithium-halogen exchange of 2,2'-dibromobiphenyl followed by the trapping of the corresponding lithiated intermediate with 9-fluorenone. The intramolecular electrophilic ring closure of the corresponding fluorenol (not isolated), in acidic media (AcOH/HCl), further provides 4-Br-SBF with 75 % yield (2 steps). Finally, a subsequent Suzuki cross-coupling between 4-Br-SBF and either 5-pyrimidine boronic acid or phenylboronic acid was performed (Pd(dppf)Cl$_2$/K$_2$CO$_3$/DMF) providing 4-5Pm-SBF or 4-Ph-SBF with significant good yields (94% and 91% respectively). Similarly, access to 2-Br-SBF was performed by the lithium-halogen exchange of 2-bromobiphenyl followed by the trapping of the corresponding lithiated intermediate with 2-bromo-9-fluorenone. The intramolecular electrophilic ring closure of the corresponding fluorenol, in acidic media (AcOH/HCl), further provides 2-Br-SBF with 67% yield (2 steps), involved in a final Suzuki cross-coupling with either 5-pyrimidine boronic acid or phenylboronic acid to lead the corresponding 2-5Pm-SBF or 2-Ph-SBF with high yields (77% and 78% respectively).
Scheme 1. Synthesis of the four dyes, 4-Ph-SBF, 4-5Pm-SBF, 2-Ph-SBF and 2-5Pm-SBF

Structural Properties

Molecular structure of the four dyes was further confirmed by X-ray crystallography. The X-ray diffraction data of single crystals of 2-Ph-SBF (Figure 1, left) and 4-Ph-SBF (Figure 1, right) reveal an asymmetric unit containing two independent molecules 1 and 2. As previously reported, in 4-Ph-SBF, the angle between the mean plane of the pendant phenyl ring and that of its attached phenyl ring of the fluorene is as large as 51.2° in molecule 1 and 56.6° in molecule 2. In 2-Ph-SBF, the same angle appears to be noticeably smaller, i.e., 37.4° for molecule 1. Similar values have been reported in literature for 2-substituted SBF (ca 30°)46,47 and seems hence to be a characteristic of such family of 2-substituted SBFs. This difference between the angle observed in 2-Ph-SBF and 4-Ph-SBF may be at the origin of the restricted π-conjugation of the latter and will have remarkable consequences on the electronic properties (see below). In addition, we note that the fluorene unit bearing the pendant phenyl ring in the molecule 1 of 2-Ph-SBF is folded with a deformation of ca 7.1° between the mean planes of each phenyl unit. However, in the case of molecule 2 of 2-Ph-SBF, we surprisingly note that the pendant phenyl unit is almost coplanar with the fluorene unit with an angle between the mean plane of the pendant phenyl ring and that of its attached phenyl ring of the fluorene unit as low as 4.6°. In addition and oppositely to what was observed for molecule 1, the fluorene bearing the phenyl unit is now almost perfectly flat with only a small deformation of 1.5° (7.1° measured in molecule 1, see above). Such structural feature appears highly surprising and is assigned to the packing of the molecule. Indeed, numbers of short CH and CC distances are detected in the packing diagram of 2-Ph-SBF and especially two short intermolecular CH distances of ca 2.8 Å between the pendant phenyl ring and its close neighbours (see SI).
However, in the case pyrimidine based SBFs, ie 4-5Pm-SBF and 2-5Pm-SBF, we only managed to obtain single crystals of 2-5Pm-SBF good enough for X-Ray diffraction and hence the direct comparison is not possible. However, we are convinced that similar structural features structures occur in both 4-5Pm-SBF and 2-5Pm-SBF. Indeed, in 2-5Pm-SBF, the angle between the mean plane of the pyrimidine plane and that of its attached phenyl ring is recorded at 23.7° (See SI), lower than that recorded for its phenyl analogue 2-Ph-SBF (37.4°).

**Electrochemical Properties**

Electrochemical properties have been investigated by cyclic voltammetry (CV) in CH₂Cl₂ in oxidation and reduction and are summarized in table 1. Pyrimidine-substituted SBF were studied in absence of Al₂O₃ as their rapid adsorption on Al₂O₃ leads to a strong decrease of the intensity of the oxidation and reduction waves.

All compounds present at least two successive oxidation waves (with maxima E₁ and E₂) before 2.2 V (figure 2). Except for 2-5Pm-SBF, the oxidation is accompanied by an electropolymerization process signed on the present CVs by a reduction wave at the back sweep. This polymerization process is more clearly observed along recurrent sweeps (see CVs in SI). Compared one to another, polymerization process is highly efficient for 2-Ph-SBF and 4-Ph-SBF and largely less efficient for 4-5Pm-SBF. Moreover, the electroactivity of the corresponding polymers, ie Poly(2-Ph-SBF) and Poly(4-Ph-SBF) are similar with onset oxidation potentials at ca 1.1 V vs SCE (polymers HOMO: -5.5 eV). In addition, the p-doping process reversibility of the two polymers occurs in a large potential range between 1 and 2 V. Oppositely, Poly(4-5Pm-SBF) presents a p-
doping process only occurring since 1.35 V (polymer HOMO : -5.75 eV) and appearing less intense and reversible.

<table>
<thead>
<tr>
<th>Electrochemical studies</th>
<th>Theoretical calculations</th>
<th>Optical studies</th>
</tr>
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<tbody>
<tr>
<td>**E&lt;sup&gt;1&lt;/sup&gt;, E&lt;sup&gt;2&lt;/sup&gt; oxidation peaks/ E&lt;sub&gt; onset&lt;/sub&gt;&lt;sup&gt;red&lt;/sup&gt; [V]&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Reduction peak/ E&lt;sub&gt; onset&lt;/sub&gt;&lt;sup&gt;red&lt;/sup&gt; [V]&lt;sup&gt;a&lt;/sup&gt;</td>
<td>HOMO&lt;sup&gt;b&lt;/sup&gt; (eV)</td>
</tr>
<tr>
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<td>-2.38/ -2.17</td>
</tr>
<tr>
<td>4-Ph-SBF</td>
<td>1.68, 1.88/ 1.55</td>
<td>-2.75/ -2.45</td>
</tr>
<tr>
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<td>1.75, 2.00/ 1.56</td>
<td>-2.33/ -2.10</td>
</tr>
<tr>
<td>2-Ph-SBF</td>
<td>1.60, 1.93/ 1.48</td>
<td>No peak in CH&lt;sub&gt;2&lt;/sub&gt;Cl&lt;sub&gt;2&lt;/sub&gt;</td>
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</table>

- Measured in CH<sub>2</sub>Cl<sub>2</sub> + Bu<sub>4</sub>NPF<sub>6</sub> 0.2 M at 100 mV/s. b. Calculated from the onset oxidation potential E<sub> onset</sub><sup>red</sup>. c. Calculated from the onset reduction potential E<sub> onset</sub><sup>red</sup>. d. ΔE<sup>d</sup> = |HOMO-LUMO|. e. Calculated from the onset of the absorption spectrum in solution in cyclohexane (see spectra figure 5 left) by using ΔE<sup>d</sup> = hν/λ (ΔE<sup>d</sup> (eV)=1237.5/λ (nm)).

Figure 2. Cyclic voltammetry of 4-5Pm-SBF (5 x 10<sup>-3</sup> M) and 2-5Pm-SBF (2 x 10<sup>-3</sup> M) (top) and of 4-Ph-SBF (5 x 10<sup>-3</sup> M) and 2-Ph-SBF (5 x 10<sup>-3</sup> M) (bottom) solutions in CH<sub>2</sub>Cl<sub>2</sub> + Bu<sub>4</sub>NPF<sub>6</sub> 0.2 M, sweep-rate 100 mV.s<sup>-1</sup>. Platinum disk (∅: 1mm) working electrode.

From the onset oxidation of the CVs, we determined the HOMO levels<sup>58</sup> of 4-5Pm-SBF at -6.0 V close to that of 4-Ph-SBF (HOMO: -5.95 eV) and of its constituting building block SBF (HOMO: -5.94 eV)<sup>49</sup> indicating that the substitution in ortho position leads to a π-conjugation interruption surely due to the large angle observed in the X-ray structure between the pendant unit and the fluorene core. In addition, we note that the substitution by a 5-pyrimidine or a phenyl group have both very weak effects on the HOMO level of the 4-substituted-SBFs. Indeed, as seen in theoretical calculations, the pendant substituent is not involved in the HOMO character of 4-substituted SBFs (Figure 3) being hence in accordance with our electrochemical conclusions. On the other hand, HOMO levels of 2-5Pm-SBF and 2-Ph-SBF calculated from their respective onset oxidation potentials are estimated at -5.96 eV and -5.88 eV. The highest HOMO of 2-
Ph-SBF compared to that of SBF signs an extension of the π-conjugation between the fluorene and the phenyl group in 2-Ph-SBF. This effect is however neutralized in 2-5Pm-SBF by the electron withdrawing effect of the pyrimidine group leading to an HOMO level identical to that of SBF. Thus, in the case of 2-Ph-SBF, the substituent is involved in the character of the HOMO (Figure 3), which is not the case for the HOMO of 2-5Pm-SBF. Cathodonic exploration in dichloromethane, reveal that all compounds are reduced at negative potential in a non-reversible process close to the electrolytic medium reduction (see CVs in SI). However, when studied in THF, allowing reaching a more cathodic potential range, one or two reduction waves are observed before the reduction of the electrolytic solution (see CVs in SI). LUMO levels were determined from the onset reduction potential in CH₂Cl₂ at -2.23 ; -1.95 ; -2.30 and -2.10 eV for 4-5Pm-SBF ; 4-Ph-SBF : 2-5Pm-SBF and 2-Ph-SBF respectively (these values are close to that measured in THF, see in SI).

The lowest LUMO level is recorded for 2-5Pm-SBF as low as -2.3 eV, 0.41 eV lower than the LUMO of SBF (-1.89 eV)⁴⁴ signing the double effect of the extension of conjugation due to the substitution at the C2 of SBF and to the electron-withdrawing effect of the electron-deficient pyrimidinium group. Thus, we note that the pyrimidine unit is strongly involved in the LUMO character of 2-5Pm-SBF and even more strongly involved in the case of 4-5Pm-SBF (Figure 3). Thus, the LUMO level of 4-5Pm-SBF (-2.23 eV) is shifted by 0.07 eV compared to that of 2-5Pm-SBF. In addition, the LUMO level of 4-5Pm-SBF is shifted by 0.27 eV compared to that of SBF, (-1.96 eV) due to the effect of the incorporation of the electron-withdrawing pyrimidine fragment. Finally, due to the absence of π-conjugation between the phenyl and the C4-fluorene,⁴⁴ LUMO levels of 4-Ph-SBF is the highest of the present series, ie -1.96 eV, and very close to that of SBF.

Finally, HOMO-LUMO gaps (ΔE⁹) extend between 3.66 eV for 2-5Pm-SBF to 4.00 eV for 4-Ph-SBF. This trend in HOMO-LUMO gap changes is well reproduced by theoretical calculations (figure 3) with nevertheless a deviation of ca 0.7 eV.

![Figure 3. Calculated frontier molecular orbitals by DFT of 4-5Pm-SBF ; 4-Ph-SBF ; 2-5Pm-SBF and 2-Ph-SBF, after geometry optimization at the B3LYP/6-311G+(d,p) level of theory, shown with an isovalue of 0.04](image-url)

Geometry optimization of the four substituted dyes in the singlet and triplet states was performed using Density Functional Theory (DFT) at the Gaussian09 B3LYP/6-311G+G(d,p) level of theory. There is a clear resemblance between the HOMO values determined by CVs and by theoretical calculations. 2-Ph-SBF is the easiest oxidizable molecule due to the extension of conjugation between the phenyl and the 2-substituted fluorene unit and its HOMO presents a phenyl-fluorene character, perfectly translating this extension of conjugation. 4-5Pm-SBF is the more difficult oxidizable molecule due to the withdrawing effect of the pyrimidine group on the fluorenyl core. However, it should be noted that 2-5Pm-SBF and 4-5Pm-SBF presents almost identical HOMO character and HOMO energy level (-6.13 eV vs -6.16 eV resp) clearly meaning that the position of the pyrimidine group only has a weak influence on the HOMO energy levels and distribution. As exposed above, the more easily reduced molecule is 2-5Pm-SBF due to the double effect of the extension of conjugation and of the electron-accepting properties of the pyrimidine unit, its LUMO has hence a clear pyrimidine-fluorene character. In 4-5Pm-SBF, the LUMO exclusively has a pyrimidine character and its energy level is 0.2 eV higher than that of 2-5Pm-SBF due to the absence of
conjugation between the fluorene and the pyrimidine unit. This is a very important feature of the present chemical design and clearly highlights its efficiency. Thus, in 4-substituted SBFs, the HOMO character is almost identical with no (or weak) contribution of the substituent in accordance with our electrochemical conclusions (see above). In addition, it must be pointed out that the LUMO+1 of 4-5Pm-SBF is very close to its LUMO (0.01 eV) with a 4-5Pm-fluorenyl and 4,5Pm character respectively. Thus, both 2-5Pm-SBF and 4-5Pm-SBF have a clear fluorene/pyrimidine character in their LUMO and LUMO+1 (almost degenerated with the LUMO which exclusively possess a pyrimidine character) respectively but present different energy levels. This highlights the importance of the torsion angle between the fluorene and the pyrimidine, which is surely at the origin of these different energy values. The computed triplet adiabatic S0 to T1 excitation energies, defined as the energy difference between total energy of the molecule in their respective singlet and triplet states, is of 2.80 and 2.82 eV for 4-5Pm-SBF and 4-Ph-SBF and of 2.63 and 2.62 eV for 2-5Pm-SBF and 2-Ph-SBF respectively. These calculations predict that 4-5Pm-SBF will be, as 4-Ph-SBF, a good candidate as host for blue and green phosphorescent dyes and both 2-5Pm-SBF and 2-Ph-SBF will be both only adapted to green dopants.

The thermal properties of all SBF derivatives were investigated by thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC) and the results are summarized in table 2. The Td is defined as the temperature at which the sample has lost the first 5% of mass during heating.

<table>
<thead>
<tr>
<th></th>
<th>4-5Pm-SBF</th>
<th>4-Ph-SBF</th>
<th>2-5Pm-SBF</th>
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<tr>
<td>m.p. (°C)</td>
<td>173</td>
<td>213b</td>
<td>197-198</td>
<td>100-108a</td>
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<td>Tc (°C)</td>
<td>-</td>
<td>115</td>
<td>-</td>
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a: This transition appears to be not very clear, occurring on a large range. b: from reference 44

Figure 4. DSC curves from second heating cycle (Left) and TGA curves (right) of 4-Ph-SBF (black line), 4-5Pm-SBF (red line), 2-Ph-SBF (blue line), 2-5Pm-SBF (green line)

5% decomposition (Td) of 4-Ph-SBF (Figure 4 right, black line) and 4-5Pm-SBF (Figure 4 right, red line) respectively occurs at 254°C and 277°C, higher than the Td of SBF (234°C) showing the interest of the incorporation of a pendant phenyl or pyrimidine unit at the C4 of SBF. Decomposition temperature of the 2-substituted-SBF occurs at 242°C (2-5Pm-SBF) and 238°C (2-Ph-SBF) close to the Td of SBF, clearly highlighting that the substitution in position 2 only has a weak effect on the thermal stability, whereas the substitution in position 4 leads to a more stiff structure being hence more thermally stable. Thus, we note that the position of the substitution has a clear influence on the Td. On the other hand, 4-5Pm-SBF and 4-Ph-SBF respectively present melting transitions at 173 and 206°C, whereas that of 2-5Pm-SBF is detected at ca 197/198° (see SI). It should be noted that the fusion of 2-Ph-SBF was very surprising appearing on a wide
range of temperature. The substituent also has a clear influence on the whole thermal properties as the replacement of phenyl by pyrimidine leads to different glass transitions ($T_g$). Thus, we note that lower $T_g$ (Figure 4, left) are detected (recorded on the second heating cycle of DSC, Figure 4 left) for phenyl-substituted SBF (76 and 78°C for 4-Ph-SBF and 2-Ph-SBF respectively) compared to pyrimidine-substituted SBF (85 and 94°C for 4-5Pm-SBF and 2-5Pm-SBF respectively). The slightly higher $T_g$ of pyrimidine-substituted compounds compared to phenyl-substituted compounds may be the consequence of stronger intermolecular interactions due to CH–N hydrogens bonding interactions. It is important to stress that all compounds present $T_g$ higher than those of classical host materials for PhOLEDs such as CBP (62°C) or m-CP (55°C). Finally, it should be mentioned that 4-5Pm-SBF, 2-5Pm-SBF and 2-Ph-SBF do not present any crystallization phenomena, $T_c$, being hence highly promising for devices incorporation. A crystallization transition was only recorded for 4-Ph-SBF but at high temperature, 115°C, during the second heating cycle in DSC.

**Photophysical studies**

The UV-Vis absorption and emission spectra recorded in cyclohexane are presented figure 5. All compounds present similar absorption bands in the range 260-315 with the same maxima at 308 nm similar to the main absorption band of SBF attributed to $\pi-\pi^*$ transition of the fluorene unit in SBF.

![Absorbance and PL spectra](image)

Figure 5. Normalized absorption (left) and emission (right, $\lambda_{exc}=300$ nm for 4-Ph-SBF, 4-5Pm-SBF and 2-Ph-SBF and 305 nm for 2-5Pm-SBF) spectra of the four dyes recorded in cyclohexane, $C=10^{-6}$ M.

At higher wavelength, each compound possesses its own signature with a clear supplemental absorption peak at 320 or 319 nm for 2-5Pm-SBF or 2-Ph-SBF signing an extension of the conjugation between the fluorene and the pyrimidine or the phenyl rings when the fluorene is substituted at C2 position. Literature reports a $\lambda_{max}$ of 328 nm (CH$_2$Cl$_2$) for the disubstituted analogue 2,7-di-5Pm-9,9-dihexyfluorene (compound F2 in chart 1) clearly showing an even more important extension of $\pi$-conjugation due to the presence of two pyrimidine units. It means that efficient $\pi$-conjugation occurs between fluorene and pyrimidine unit leading to a red shift of ca 8/10 nm per pyrimidine unit. In the case of 4-substituted SBF, we note the presence of wavelength tail between 315 and 340 nm, which appears to be larger in the case of 4-5Pm-SBF than in the case of 4-Ph-SBF. This tail may be due to a more limited $\pi$-conjugation between the pyrimidine or the phenyl ring and the fluorene when substituted at the C4 position. It could also reflect a minor proportion of conformers of the molecules with a more planar structure, allowing a certain degree of $\pi$-conjugation between the pyrimidine or phenyl rings and the fluorene moiety. As these bands at low energy seem to be more intense for pyrimidine-based dyes than for phenyl-based dyes, one may conclude that more efficient $\pi$-conjugation seems to occur between fluorene and pyrimidine units.
From the onset absorption wavelength, optical energy gap ($\Delta E_{opt}$) varying from 3.66 eV for 2-5Pm-SBF to 3.82 eV for 4-Ph-SBF were determined (see Table 1) with a variation following the same trend as that obtained from electrochemical measurement and theoretical calculations.

The experimental UV-vis spectra of the four dyes have been compared with predicted electronic absorption spectrum from TD-DFT calculations (see Figures in SI) and compared to that of its constituting building block SBF. From those calculations one may observe that except 4-Ph-SBF, all compounds present an additional absorption band at a lower energy than that the main band observed in SBF (see figures in SI). This additional band presents an oscillator strength value as high as 0.355 for 2-Ph-SBF and 0.1677 for 2-5Pm-SBF, similar to that calculated for the neighboring band (0.3346 for 2-Ph-SBF and 0.1553 for 2-5Pm-SBF), itself calculated close to the lowest energy band of SBF. This is in accordance with the experimental conclusions from absorption spectroscopy presented above. Looking at the detailed transitions, for 2-Ph-SBF and 2-5Pm-SBF, the lowest energy absorption band corresponds to an HOMO-LUMO transition involving the 2-substituted-fluorene unit (see figure 3). For 4-5Pm-SBF, the two calculated lowest energy bands present strongly less intense oscillator strength (0.0198 and 0.0303 for excited state 1 and 2 respectively) and the more important transitions involved in these two bands correspond to (i) HOMO vs LUMO transition (65 % in excited state 1) involving two different molecular fragments of the fluorophore (HOMO possessing a spirobifluorenyl character and LUMO possessing a pyrimidine character) presenting very low orbital overlap and hence a very strong charge transfer character and (ii) HOMO vs LUMO+1 transition (58.7% in excited excited state 2) involving two fluorenyl units and hence an orbital overlap between them (HOMO possessing a spirobifluorenyl character and LUMO+1 possessing a pyrimidine/fluorenyl character). Finally, calculated UV-spectrum of 4-Ph-SBF is very similar to that calculated for SBF (see SI) and the lowest energy absorption bands with very weak oscillator strength (0.06 and 0.059 for calculated excited states 1 and 2) correspond to the HOMO vs LUMO and HOMO vs LUMO+1 transitions, corresponding mainly to fluorene-fluorene transitions.

Fluorescence spectra of the four dyes (figure 5, right) clearly show drastically different emission spectra depending on the substitution. The first type of emission spectra recorded for the 2-substituted-SBFs presents a well resolved emission spectra with maxima at 338/355 and 335/351 nm for 2-5Pm-SBF and 2-Ph-SBF respectively, signing the extension of $\pi$-conjugation compared to SBF ($\lambda_{max} = 311/323$ nm).49 Structured emission spectrum was also reported for 2,7-di-5Pm-9,9-dihexyfluorene (compound F2 in chart 1) with maxima at 361 and 378 nm due to an even more important extension of conjugation in the disubstituted-fluorenyl core.40 4-substituted SBFs presents structureless and large emission spectra centered at 365 nm for 4-5Pm-SBF and 358 nm for 4-Ph-SBF. This surprising loss of resolution was also observed for the rare examples of 4-substituted SBF found in literature, ie 4-diphenylphosphine oxide-SBF ($\lambda_{max}: 346$ nm)49, 4-dibenzo furan-SBF ($\lambda_{max}: 369$ nm)41 and 4-dibenzo thiophene-SBF ($\lambda_{max}: 355$ nm).42 This feature appears hence to be a unique characteristic of 4-substituted SBF derivatives as 2-substituted SBF analogues (and 9,9'dialkylfluorene as well) always present well resolved emission bands assigned to the double bond character of the C-C bond linking the pendant substituent and the fluorene core in the excited state.52

In addition, there is also an intriguing red shift observed between 4-substituted-SBF and 2-substituted-SBF derivatives (shift of 27 nm for 5Pm-SBFs and of 23 nm for Ph-SBFs). It is clear that the position of the substitution (C2 vs C4) is at the origin of this spectacular effect. We could assume that the C4 substitution of phenyl or pyrimidine fragments on the fluorene do not affect dramatically the $\pi$-conjugation in the ground state (absorption spectra are only slightly altered with a low energy tail, as described above), but contribute to a strong planarization in the excited state leading to an impressive Stokes shift. More detailed theoretical and spectroscopic investigations need to be conducted to unravel this critical issue.

The emission quantum yields ($\Phi_{sol}$) were determined in cyclohexane by using standard procedures with quinine sulphate as reference. On one hand, phenyl-substituted SBFs appear to be efficiently fluorescent with $\Phi_{sol}$ of 0.42 and 0.87 and lifetimes of 4.20 ns and 1.56 ns for 4-Ph-SBF and 2-Ph-SBF, respectively. On the other hand, a quenching of fluorescence is observed for pyrimidine-substituted SBFs with $\Phi_{sol}$ of 0.04 and 0.02 and lifetimes of 0.61 ns and 0.44 ns for 4-5Pm-SBF and 2-5Pm-SBF, respectively. It should be
noted that in other solvents, such as THF, the quantum yields of 4-5Pm-SBF and 2-5Pm-SBF remain very low and the solvent can hence not be responsible of these low values. It must be noted that such a decrease of the fluorescence was not observed for previously reported compounds with pyrimidine units such as TBPFS (Φsol=1 in solution)\(^\text{35}\) and F3 (Φsol=0.85 in solution)\(^\text{30}\) (see structures in chart 1) bearing respectively either a 5 pyrimidine or a 2 pyrimidine unit, both being also substituted with a phenyl ring. The presence of this phenyl ring on the pyrimidine unit appears hence to be very important and we believe that, in our case, the absence of this phenyl ring and hence the presence of the aromatic NCHN fragment in the 5-pyrimidine unit has a strong impact on the deactivation rates of the excited state. Indeed, pyrimidine substituted derivatives 4-5Pm-SBF and 2-5Pm-SBF show a lower radiative rate constant \(k_r\) and a much higher non-radiative rate constant \(k_{nr}\) compared to phenyl-substituted derivatives 4-Ph-SBF and 2-Ph-SBF, as mentioned in Table 3.

Table 3. Optical properties of the four SBF dyes.

<table>
<thead>
<tr>
<th></th>
<th>4-5Pm-SBF</th>
<th>4-Ph-SBF</th>
<th>2-5Pm-SBF</th>
<th>2-Ph-SBF</th>
</tr>
</thead>
<tbody>
<tr>
<td>(\lambda_{abs, solution} (\text{nm}))</td>
<td>297 (0.5), 308</td>
<td>297 (0.7), 308 (1.4), 296 (2.5), 308 (2.5), 297 (2.4), 308 (2.2),</td>
<td>301, 312, 326</td>
<td>301, 312, 329</td>
</tr>
<tr>
<td>(\varepsilon (10^4, \text{L.mol}^{-1}.\text{m}^{-2}))</td>
<td>(0.6), 318 (sh)</td>
<td>318 (sh)</td>
<td>320 (1.8)</td>
<td>319 (1.6)</td>
</tr>
<tr>
<td>(\lambda_{abs, thin-film} (\text{nm}))</td>
<td>298, 311</td>
<td>298, 311</td>
<td>301, 312, 326</td>
<td>301, 312, 329</td>
</tr>
<tr>
<td>(\lambda_{em, solution} (\text{nm}))</td>
<td>365</td>
<td>358</td>
<td>338, 355</td>
<td>335, 351</td>
</tr>
<tr>
<td>(\lambda_{em, thin-film} (\text{nm}))</td>
<td>372</td>
<td>363</td>
<td>355, 366</td>
<td>343, 359</td>
</tr>
<tr>
<td>(\Phi_{sol})</td>
<td>0.04</td>
<td>0.42</td>
<td>0.02</td>
<td>0.87</td>
</tr>
<tr>
<td>(\tau (\text{ns}))</td>
<td>0.61</td>
<td>4.20</td>
<td>0.44</td>
<td>1.56</td>
</tr>
<tr>
<td>(k_r (s^{-1}))</td>
<td>6.6×10^7</td>
<td>1.0×10^8</td>
<td>4.6×10^7</td>
<td>5.6×10^8</td>
</tr>
<tr>
<td>(k_{nr} (s^{-1}))</td>
<td>1.6×10^9</td>
<td>1.4×10^8</td>
<td>2.2×10^9</td>
<td>8.3×10^7</td>
</tr>
</tbody>
</table>

a. in cyclohexane, b. Spin-coated thin film from a solution in THF, [solution] = 10 mg/mL, c. Quantum yields in solution (Φsol) were calculated relative to quinine sulfate (Φsol = 0.546 in H_2SO_4 1N)

From a careful inspection of the dependence of the absorption behaviour of the four dyes on solvent polarity, we note that the absorption maximum is almost insensitive to the dielectric constant of the environment (see Figures in SI). This indicates that the electronic and structural characteristics of the ground and Franck Condon excited states do not differ much with a change in solvent polarity.\(^\text{56}\)

While phenyl-substituted SBFs showed minor differences of the PL spectra in various solvents (Figure 6, top), distinct fluorescence solvatochromism was observed in pyrimidine-substituted SBF when the solvent polarity increases from cyclohexane to ethanol (Figure 6, bottom). Indeed, both dyes are highly sensitive to the solvent polarity with fluorescence maxima ranging from 366 and 338 nm in cyclohexane to 390 and 375 nm in ethanol for 4-5Pm-SBF and 2-5Pm-SBF respectively. This dependence of the emission wavelength on the solvent polarity is indicative of dipole-dipole interactions between Pm-substituted SBFs and polar solvent molecules and hence of a photoinduced intramolecular charge transfer between the donor (fluorene) and the acceptor part (pyrimidine) of the dye.\(^\text{57}\)
Figure 6. Normalized PL spectra of solutions of 4-Ph-SBF (top left) and 2-Ph-SBF (top right) and of 4-5Pm-SBF (bottom left) and 2-5Pm-SBF (bottom right) in cyclohexane, toluene, THF, CH₂Cl₂, ethanol and CH₃CN.

The solid state absorption spectra of all dyes (figure 7, left) appear very similar to those in solution (figure 5 left) with only a small red shift of less than 8 nm detected and a slight increase of the lowest energy band intensity (compared to the band at 310 nm) for the 2-substituted-SBFs. Such similarity between absorption spectra in solution and in solid state clearly signs that there are minimal intermolecular interactions in the ground state in thin films.

Figure 7. Absorption (left) and Emission (right) spectra of the four dyes recorded in solid state (spin-coated thin film from a solution in THF, [solution] = 10 mg/mL).

Regarding the solid-state fluorescent properties (figure 7, right), except for 2-5Pm-SBF, they are almost identical to those recorded in solution (figure 5, right) with only a small red shift ranging from 3 nm for 4-Ph-SBF to 8 nm for 4-5Pm-SBF and 2-Ph-SBF. Solid-state fluorescent spectrum of 2-5Pm-SBF (figure 7, right, pink line) nevertheless appears poorly resolved with two maxima recorded at 355/366 nm, red shifted by 17/11 nm compared to that in solution. Finally, to examine the suitability of the four dyes as host materials for phosphorescent green and/or blue dopant, the phosphorescence of the molecules was recorded at 77 K in a methylcyclohexane/2-methylpentane mixture (1:1), Figure 8. From those spectra, Eᵣ values of the molecules were calculated from the lowest phosphorescent peak of 2.75 eV for 4-5Pm-SBF and 2.56 eV for
**2-Ph-SBF.** Compared to the $E_T$ value of **4-Ph-SBF** (2.77 eV) and of **SBF** (2.87 eV) reported by our group recently, we note that the $E_T$ level of **4-5-Pm-SBF** is sufficiently high for its use as host material for both green and blue phosphorescent dyes. The $E_T$ of **2-Ph-SBF** (2.56 eV) and that of **2-5-Pm-SBF** (2.58 eV) is lower than that of the blue dopant Flrpic ($E_T$: 2.62 eV) and these molecules may be used as host material for green dopant Irppy$_3$ ($E_T$: 2.42 eV).

![Emission spectra of 2-Ph-SBF and 4-5-Pm-SBF](image)

Figure 8. Emission spectra at 77 K in a methycyclohexane/2-methylpentane mixture (1:1) of **4-Ph-SBF** ($\lambda_{exc}=300$ nm), **4-5Pm-SBF** ($\lambda_{exc}=300$ nm) and **2-Ph-SBF** ($\lambda_{exc}=250$ nm) and **2-5PmSBF** ($\lambda_{exc}=250$ nm). The determination of the $E_T$ value has been done at from the lowest phosphorescent peak (black arrow).

In conclusion of the above presented photophysical and physico-chemical analysis, the position (C2 vs C4) and the nature of the substituent (phenyl vs pyrimidine) on the SBF core leads to drastic alterations of the resulting properties. Substitution at the C2 leads to an increase of the conjugation length and therefore to a decrease of the singlet and triplet state energy level, while the substitution at the C4 only has a weak influence on these two key parameters, keeping them almost identical to those of their constituting building block SBF. This has been assigned to π-conjugation breaking induced by the large torsion angle made by the substituents in C4. This chemical design allows hence obtaining organic semiconductors based on a C4-substituted SBF with high singlet and triplet state energy level with excellent thermal/morphological properties. On the other hand, the substitution by the pyrimidine group leads to a decrease of the LUMO level due to the pyrimidine electron affinity, which should lead in principle to a decrease of the threshold voltage of the resulting devices. Thus, the four dyes possess the prerequisites for their uses as host in PhOLEDs that is, a high $T_g$ and $T_d$, HOMO/LUMO levels adapted for hole and electron injection/transport in the devices and $E_T$ higher that of the green phosphorescent emitter Irppy$_3$ (2.42 eV) and in the case of **4-Ph-SBF** and **4-5Pm-SBF** for the blue phosphorescent emitter Flrpic (2.62 eV).
Scheme 2. Energy level diagram of the devices.

In order to explore the potential of the present semi-conductors as host for triplet emitters, green (Ir(ppy)$_3$) and blue (Firpic) PhOLEDs were fabricated. The device configuration was ITO/CuPc(10 nm)/NPB (40 nm)/TCTA (10 nm)/**EML**: dopant (20 nm)/TPBI (40 nm)/LiF (1.2 nm)/Al (100 nm). ITO is used as the anode, CuPc (copper phthalocyanine) is the hole injecting layer, NPB (N,N’-di(1-naphthyl)-N,N’-diphenyl-[1,1’-biphenyl]-4,4’-diamine) is the hole-transporting layer, TCTA (4,4’,4”-Tris(carbazol-9-yl)-triphenylamine) is the electron/exciton blocking layer, TPBI (1,3,5-Tris(1-phenyl-1H-benzimidazol-2-yl)benzene) is both the electron transporting layer and the hole blocking layer and a thin film of lithium fluoride covered with aluminum is the cathode. Ir(ppy)$_3$ and Firpic are used as dopant for green and blue OLEDs respectively. The relative energy levels of the successive layers of the devices are reported in scheme 2. Table 4 gathers selected data obtained from the average of two experiments.

Due to their $E_{T}$ values higher than that of Ir(ppy)$_3$, the four substituted-SBFs were used as host for Ir(ppy)$_3$, green dopant. Performance of the best green PhOLEDs is presented in figure 9 (left) and average parameters are gathered in table 4. The best green PhOLED performance was obtained with **2-5Pm-SBF** and **4-5Pm-SBF** host materials doped with 10% Ir(ppy)$_3$. As shown in figure 9 (left), maximum current and power efficiencies of 58 cd/A and 35 lm/W are obtained respectively for the pyrimidine substituted SBF. Maximum luminance as high as 30 000 cd/m$^2$ is reached for these devices as shown in table 4. Phenyl substituted dyes, **4-Ph-SBF** and **2-Ph-SBF** presents lower performance with EQE of 10.4% and 11.3 %, highlighting the great interest of the incorporation of the pyrimidine unit in SBF scaffold. The threshold voltage for all devices appears to be low and as low as 2.5 eV for **2-5Pm-SBF**.

The EL spectra of the four devices based on the different host materials are identical, exclusively showing the emission of the green dopant at 516/540 nm (figure 9 right) close to the PL emission of pure Ir(ppy)$_3$ film (509/540 nm)$^{57}$ with no parasite emission in the range of the undoped device. This result demonstrates an efficient energy transfer from the four SBF-derivatives to Ir(ppy)$_3$, due to their high triplet energy levels.

The device performance obtained with small pure hydrocarbon host molecules **4-Ph-SBF** and **2-Ph-SBF** are comparable to those obtained with more complicated pure-hydrocarbon-based host materials recently reported.$^{38,37,58}$ Performances obtained with pyrimidine-substituted SBF are even more interesting with lowest turn-on voltages and highest EQE (close to 14%) and clearly demonstrate the interest of the pyrimidine substitution in notably favouring the injection of electron.
Table 4. Selected EL data of green and blue devices (average values calculated from two experiments).

<table>
<thead>
<tr>
<th>Host /Guest (%)</th>
<th>V_on (V)</th>
<th>EQE (%)</th>
<th>Current efficiency (cd.A^{-1})</th>
<th>Power efficiency (lm/W)</th>
<th>L_{max}^\text{at current density} (cd.m^{-2}) at 10 mA/cm²</th>
<th>CIE (x ; y) at 10 mA/cm²</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Green Phosphorescent OLEDs</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2-Ph-SBF /Ir(ppy)_3 (10)</td>
<td>2.6</td>
<td>11.3</td>
<td>42.7</td>
<td>17.8</td>
<td>27 990 (210)</td>
<td>0.33 ; 0.62</td>
</tr>
<tr>
<td>2-5Pm-SBF /Ir(ppy)_3 (9)</td>
<td>2.5</td>
<td>13.3</td>
<td>49.2</td>
<td>21.5</td>
<td>30 000 (160)</td>
<td>0.33 ; 0.61</td>
</tr>
<tr>
<td>4-Ph-SBF (9) /Ir(ppy)_3 (10)</td>
<td>3.5</td>
<td>10.4</td>
<td>40.3</td>
<td>16.7</td>
<td>25 700 (230)</td>
<td>0.32 ; 0.62</td>
</tr>
<tr>
<td>4-5Pm-SBF /Ir(ppy)_3 (10)</td>
<td>3.3</td>
<td>13.7</td>
<td>49.9</td>
<td>17.6</td>
<td>28 000 (150)</td>
<td>0.34 ; 0.61</td>
</tr>
<tr>
<td><strong>Blue Phosphorescent OLEDs</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4-Ph-SBF (9) /FirPic (19)</td>
<td>4.0</td>
<td>5.7</td>
<td>16.2</td>
<td>6.0</td>
<td>3800 (70)</td>
<td>0.20 ; 0.44</td>
</tr>
<tr>
<td>4-5Pm-SBF /FirPic (19)</td>
<td>3.3</td>
<td>4.8</td>
<td>14.1</td>
<td>5.5</td>
<td>2030 (50)</td>
<td>0.20 ; 0.45</td>
</tr>
</tbody>
</table>

Figure 9. Current (empty symbol) and power efficiencies (filled symbol) versus current density of the green devices using 4-Ph-SBF, 4-5Pm-SBF, 2-Ph-SBF and 2-5Pm-SBF doped with Ir(ppy)_3 (10% in mass) as emitting layer (left). Corresponding EL spectra recorded at 10 mA/cm² (right).

Due to their high E_T values (2.77 eV for 4-Ph-SBF and 2.75 eV for 4-5Pm-SBF), higher than that of Firpic (E_T: 2.62 eV), the two 4-substituted-SBFs were used as host for this blue dopant. Characteristics of the two blue devices are presented in figure 10. The best performance is obtained with 4-Ph-SBF with an EQE of 5.7 %, a LE of 16.2 cd/A and a threshold voltage of 4 V while blue PhOLEDs with 4-5Pm-SBF as host reach an EQE of 4.8 %, a LE of 14.1 cd/A and a threshold voltage of 3.3 V reported at a current density value of 10 mA/cm² (Figure 10 left and Table 4). The threshold voltage of 4-5Pm-SBF is hence decreased by ca 0.7 V compared to that of 4-Ph-SBF due to the incorporation of the pyrimidinium group in the former (which leads to a decrease of the LUMO level and hence an easier injection of the electron in the device). This highlights the efficiency of the chemical design investigated in this work. Both devices, reach brightness of 3800 / 2030 cd/m² for 4-Ph-SBF / 4-5Pm-SBF based devices respectively.

EL spectra of the devices (Figure 10 right) point a main emission of the blue dopant with maxima at 473 and 500 nm close to the PL emission of pure Firpic film (475/500 nm) but also, in the case of 4-PhSBF of an additional tiny emission in the range of the undoped device (360/440 nm). This emission may be due to a small amount of recombination in the other organic layers and especially in TCTA. CIE coordinates of 4-Ph-SBF-based device (0.20 ; 0.44) and of 4-5Pm-SBF-based device (0.20 ; 0.45) appear nevertheless almost identical as shown in Table 4.
Figure 10. Current (empty symbol) and power efficiencies (filled symbol) versus current density of the blue devices using 4-Ph-SBF and 4-5Pm-SBF doped with FIrPic (19% in mass) as emitting layer (left). Corresponding EL spectra recorded at a current density of 10 mA/cm² (right).

Conclusions

In summary, four substituted Sbf dyes have been synthesized and their thermal, electrochemical and photophysical properties have been scrutinized and compared to the properties of their constituting SBF unit. From these studies it has been demonstrated that, compare to SBF, the substitution by a phenyl unit at the C4 position of SBF leads to weaker modification of the HOMO and LUMO levels compared to the phenyl substitution at the C2 position, which leads to a conjugation extension and hence a decrease of the HOMO-LUMO gap. On the other hand, the substitution by the electro-deficient pyrimidine unit compared to the substitution by the phenyl unit leads to a significant lowering of the LUMO level. Overall, the combination of the two effects (lowering of the HOMO through the extension of conjugation, lowering of the LUMO through pyrimidine electro deficient effect) in 2-5Pm-SBF leads to the highest modification of the optical energy gap (from 4.05 eV in SBF to 3.66 eV in 2-5Pm-SBF). Thus, the optical energy gap of the four compounds is finely tuned from 4 eV for 4-Ph-SBF to 3.66 eV for 2-5Pm-SBF. In addition, the substitution in C2 or in C4 has an impressive influence on the Eg. Indeed, the C2-substituted dyes have Eg values at ca 2.42 eV, whereas the C4 substituted dyes have an Eg as high as 2.7 eV due to the π-conjugation disruption. This structure properties relationship study clearly highlights how a different substitution may lead to very different properties keeping others intact. As all compounds possess a triplet energy level higher than 2.42 eV, they have been used as host for green dopant in PhOLEDs. Remarkable performance has been obtained with 2-5Pm-SBF as host (CE > 58 cd/A, PE > 35 lm/W, EQE > 14 % and threshold voltage of 2.5 V). Furthermore, the two 4-substituted SBF with triplet energy levels as high as 2.8 eV have also been successfully used as host materials for blue PhOLEDs with high performance. The effect of the pyrimidine substituent leads to a lowering of the threshold working voltage of the device from 4 V for 4-Ph-SBF based blue device to 3.3 V for 4-5Pm-SBF based blue device.

Acknowledgements

S.T. and G. S. thank ANR (HOME-OLED project n°ANR-11-BS07-020-01) for a studentship. C. D thanks ANR (HOME-OLED project n°ANR-11-BS07-020-01) for financial support as postdoctoral position. We wish to thank the C.R.M.P.O (Centre Régional de Mesures Physiques de l’Ouest, Rennes) for high resolution mass measurements, Service de Microanalyse-CNRS (Gif sur Yvette) for CHN analyses, CINES (Montpellier) for computing time, the CDIFX (Rennes) for X-Ray data collection, the ”Institut des Sciences Analytiques” (UMR CNRS 5280, Villeurbanne) for TGA. Arnaud Brousseau (Cachan) is thanked for his precious assistance in photophysical measurements. Dr Franck Camerel (Rennes) is warmly acknowledged for his help and advice.
in the understanding of thermal properties. Prof. Abdou Boucekkine is also warmly thanked for his help in DFT and TD-DFT calculations.

**Experimental Section**

**Synthesis:**

All manipulations of oxygen- and moisture-sensitive materials were conducted with a standard Schlenk technique. Commercially available reagents and solvents were used without further purification other than those detailed below. THF was distilled from sodium/benzophenone prior to use. Light petroleum refers to the fraction with bp 40-60°C. 2.5 M solutions of n-BuLi in hexanes or THF were purchased from Sigma Aldrich. 2,2’-dibromobiphenyl was purchased from Fluorochem. Reactions were stirred magnetically, unless otherwise indicated. Analytical thin layer chromatography was carried out using aluminum backed plates coated with Merck Kieselgel 60 GF254 and visualized under UV light (at 254 and 360 nm). Chromatography was carried out using Teledyne Isco CombiFlash® RF 400 (UV detection 200-360 nm), over standard silica cartridges (Redisper® Isco, GraceResolv™ Grace or Puriflash® columns Interchim). 1H and 13C NMR spectra were recorded using Bruker 300 MHz instruments (1H frequency, corresponding 13C frequency: 75 MHz); chemical shifts were recorded in ppm and J values in Hz. In the 13C NMR spectra, signals corresponding to C, CH, CH2 or Me groups, assigned from DEPT, are noted. The residual signals for the NMR solvents are: CDCl3: 7.26 ppm for the proton and 77.00 ppm for the carbon, CD2Cl2: 5.32 ppm for the proton and 53.80 ppm for the carbon. The following abbreviations have been used for the NMR assignment: s for singlet, d for doublet, t for triplet and m for multiplet. High resolution mass spectra were recorded at the Centre Régional de Mesures Physiques de l’Ouest (Rennes) on (i) Bruker MicroTof-Q II (Source: Atmospheric Pressure Chemical Ionization (APCI - direct introduction) (ASAP–Atmospheric Solids Analysis Probe) at a temperature of 30°C - positive mode) or on (ii) Waters Q-Tof II. The synthesis of 4-bromo-9,9’-spirob[fluorene] (1) and of 4-Ph-SBF have been previously described.

2-bromo-9,9’-spirob[fluorene] (2)33 2-bromo-1,1’-biphenyl (2.16 g, 9.27 mmol, 1.15 eq) was dissolved in dry THF (40 mL) under argon atmosphere, cooled at -78°C, and stirred during 10 minutes at this temperature. A 2.5 M n-BuLi solution in THF (4.31 mL, 10.78 mmol, 1.34 eq) was then slowly injected via a seringue, at -78°C. The resulting yellow mixture was stirred at the same temperature for 30 min. 2-bromo-9H-fluoren-9-one (2.08 g, 8.03 mmol) in dry THF (35 mL) was then added dropwise, and the mixture was stirred for another 30 min at -78°C, and allows warming up to room temperature gradually and stirred overnight. Saturated brine solution (10 mL) was added, and the mixture was extracted twice with ethyl acetate (2x30 mL). The combined organic extracts were dried over anhydrous magnesium sulfate, filtered, and concentrated under reduced pressure. Without other purification, the crude was dissolved into a mixture of acetic acid/hydrochloric acid (50 mL/5 mL) and warmed at 70°C during 2 h under stirring. Then, the mixture was poured into water/ice (200 mL), and the solution was neutralized with solid sodium hydroxide until pH 7. Then the organic layer was extracted three times with dichloromethane (3x50 mL). The combined organic extracts were dried over magnesium sulfate, filtered, and concentrated under reduced pressure. The residue was purified by flash chromatography on silica gel (light petroleum/dichloromethane 95/5) to afford a colorless solid (2.14 g). [column conditions: Silica cartridge 40 g (Serlabo); solid deposit on Celite®; λ detection: (254 nm, 280 nm); light petroleum/dichloromethane (95/5) at 25 mL/min; collected fraction: 18-42 min]. Yield: 67%. mp: 168-170°C (lit. 160 °C);64 1H NMR (300 MHz, CD2Cl2): δ 7.89 (m, 3H, ArH), 7.76 (d, J = 8.1 Hz, 1H, ArH), 7.52 (dd, J = 8.1, 1.8 Hz; 1H, ArH), 7.42 (m, 3H, ArH), 7.15 (m, 3H, ArH), 6.87 (m, 1H, ArH), 6.71 (m, 3H, ArH); 13C NMR (75 MHz, CD2Cl2): δ 151.5 (C), 149.1 (C), 148.5 (C), 142.4 (C), 141.4 (C), 141.3 (C), 131.4 (CH), 128.8 (CH), 128.6 (CH), 128.54 (CH), 128.48 (CH), 127.6 (CH), 124.37 (CH), 124.35 (CH), 122.1 (CH), 121.8 (C), 120.8 (2xCH seen from HSQC), 66.4 (Siro C).

**General procedure for Suzuki coupling**

9,9’-spirob[fluorene] brominated compound, boronic acid compound, K₂CO₃ and Pd(dppf)Cl₂ were dissolved in dry DMF or xylenes. The mixture was warmed to 150°C (DMF) and 135°C (xylenes) stirred overnight. After cooling to room temperature, saturated solution of ammonium chloride (50 mL) was ad-

---

Pd(dppf)Cl₂ : [1,1’-Bis(diphenylphosphino)ferrocene]palladium(II) dichloride complex with dichloromethane
ded, and organic layer was extracted three times with dichloromethane (3x30 mL) and washed with brine (3x30 mL). The combined organic extracts were dried over magnesium sulfate, filtered, and concentrated under reduced pressure. The residue was purified by flash chromatography on silica gel using Teledyne Isco CombiFlash® Rf 400. Column conditions can be found in each compounds description below.

5-(9,9'-spirobi[fluoren]-4-yl)pyrimidine (4-SPM-SBF). 5-(9,9'-spirobi[fluoren]-4-yl)pyrimidine was synthesized according to the general coupling procedure between 4-bromo-9,9'-spirobi[fluorene] (1) (0.808 g, 2.045 mmol), pyrimidin-5-yl-boronic acid (0.333 g, 2.688 mmol, 1.31 eq), K$_2$CO$_3$ (2.800 g, 20.260 mmol, 9.91 eq) and Pd(dpdf)Cl$_2$ (0.088 g, 0.108 mmol, 0.05 eq) in dry DMF (30 mL). The residue was purified by flash chromatography on silica gel (light petroleum/ethyl acetate 85/15) to afford a colorless solid (0.759 mg, 1.924 mmol) [column conditions: Silica cartridge 12 g (Serlabo); solid deposit on Celite®; λ$_{detection}$: (254 nm, 280 nm); ethyl acetate in light petroleum (85/15) at 20 mL/min; collected fraction: 8-17 min]. Yield: 94%. mp: 167°C; IR (ATR, cm$^{-1}$) ν = 593, 646, 723, 752, 908, 1030, 1157, 1282 1405, 1444, 1545, 3020, 3072; $^1$H NMR (300MHz, CD$_2$Cl$_2$): δ 9.36 (s, 1H, ArH), 9.04 (s, 2H, ArH), 7.89 (m, 2H, ArH), 7.41 (td, J = 7.5, 1.2 Hz, 2H, ArH), 7.22-7.01 (m, 7H, ArH), 6.76 (m, 3H, ArH), 6.68 (s, 1H, ArH); $^{13}$C NMR (75MHz, CD$_2$Cl$_2$): δ 158.6 (CH), 157.4 (CH), 150.7 (C), 149.9 (C), 149.0 (C), 142.5 (C), 141.3 (C), 139.7 (C), 135.1 (C), 130.8 (C), 130.7 (CH), 128.7 (CH), 128.53 (CH), 128.48 (CH), 128.3 (CH), 128.2 (CH), 124.9 (CH), 124.7 (CH), 124.3 (CH), 123.0 (CH), 120.8 (CH), 66.24 (Spío C); HRMS calculated for C$_{29}$H$_{39}$N$_2$: 395.1542, found 395.1542 [M+H]$^+$; Elemental analysis calculated for C$_{29}$H$_{39}$N$_2$: C, 88.30%; H, 4.60%; N, 7.10%. Found: C, 87.81%; H, 4.81%; N, 6.81%; λ$_{abs}$ [nm] ([ε$^{10^4}$ L.mol$^{-1}$.cm$^{-1}$]) = 297 (0.5), 308 (0.6).

2-phenyl-9,9'-spirobi[fluorene] (2-Ph-SBF). 2-phenyl-9,9'-spirobi[fluorene] was synthesized according to the general procedure between 2-bromo-9,9'-spirobi[fluorene] (2) (0.751 g, 1.901 mmol), phenylboronic acid (0.304 g, 2.488 mmol, 1.31 eq), K$_2$CO$_3$ (1.227 g, 8.883 mmol, 4.67 eq) and Pd(dpfd)Cl$_2$ (0.061 g, 0.076 mmol, 0.04 eq) in xylene (20 mL). The residue was purified by flash chromatography on silica gel (light petroleum/dichloromethane 9/1) to afford a colorless solid (0.581 g, 1.482 mmol), [column conditions: Silica cartridge 24 g (Serlabo); solid deposit on Celite®; λ$_{detection}$: (254 nm, 280 nm); dichloromethane in light petroleum (9/1) at 20 mL/min; collected fraction: 35-45 min]. Yield: 78%. mp: 100-108°C IR (ATR, cm$^{-1}$) ν = 476, 494, 617, 634, 650, 692, 727, 748, 829, 1020, 1155, 1280, 1444, 1597, 3014, 3059; $^1$H NMR (300 MHz, CD$_2$Cl$_2$): δ 7.96 (dd, J = 8.0, 0.6 Hz, 1H, ArH), 7.93 – 7.88 (m, 3H, ArH), 7.66 (dd, J = 8.0, 1.7 Hz, 1H, ArH), 7.45 – 7.36 (m, 5H, ArH), 7.35 – 7.20 (m, 3H, ArH), 7.13 (td, J = 7.5, 1.1 Hz, 3H, ArH), 6.94 (dd, J = 1.7, 0.6 Hz, 1H, ArH), 6.78 – 6.72 (m, 2H, ArH), 6.72 – 6.67 (m, 1H, ArH); $^{13}$C NMR (75 MHz, CD$_2$Cl$_2$): δ 150.1 (C), 149.7 (C), 149.2 (C), 142.4 (C), 142.0 (C), 141.7 (C), 141.4 (C), 141.3 (C), 129.1 (CH), 128.40 (2xCH$_3$), 128.38 (CH), 128.37 (CH), 128.78 (CH), 127.43 (CH), 127.38 (CH), 124.4 (CH), 124.3 (CH), 123.0 (CH), 121.0 (CH), 120.8 (CH), 120.7 (CH), 66.6 (Spío C); HRMS calculated for C$_{30}$H$_{32}$: 393.1643, found 393.1646 [M+H]$^+$; Elemental analysis calculated for C$_{30}$H$_{32}$: C, 94.86%; H, 5.14%. Found: C, 94.93%; H, 5.24%; λ$_{abs}$ [nm] ([ε$^{10^4}$ L.mol$^{-1}$.cm$^{-1}$]) = 297 (2.4), 308 (2.2), 319 (1.6).

5-(9,9'-spirobi[fluoren]-2-yl)pyrimidine (2-SPM-SBF). 5-(9,9'-spirobi[fluoren]-2-yl)pyrimidine was synthesized according to the general procedure between 2-bromo-9,9'-spirobi[fluorene] (2) (0.749 g, 1.897 mmol), pyrimidin-5-yl-boronic acid (0.312 g, 2.519 mmol, 1.33 eq), K$_2$CO$_3$ (1.226 g, 8.872 mmol, 4.68 eq) and Pd(dpfd)Cl$_2$ (0.059 g, 0.073 mmol, 0.04 eq) in dry DMF (20 mL). The residue was purified by flash chromatography on silica gel (light petroleum/ethyl acetate 8/2) to afford a colorless solid (0.5743 g, 1.456 mmol) [column conditions: Silica cartridge 24 g (Serlabo); solid deposit on Celite®; λ$_{detection}$: (254 nm, 280 nm); ethyl acetate in light petroleum (8/2) at 20 mL/min; collected fraction: 34-50 min]. Yield (77%). mp: 197-198°C; IR (ATR, cm$^{-1}$) ν = 491, 604, 619, 636, 721, 731, 750, 833, 862, 1018, 1105, 1153, 1186, 1267, 1400, 1414, 1446, 1554, 3016, 3039, 3053; $^1$H NMR (300 MHz, CD$_2$Cl$_2$): δ 8.94 (s, 1H, ArH), 8.67 (s, 2H, ArH), 7.93 (dd, J = 7.9, 0.6 Hz, 1H, ArH), 7.87 – 7.83 (m, 1H, ArH), 7.83 – 7.79 (m, 2H, ArH), 7.56 (dd, J = 7.9, 1.7 Hz, 1H, ArH), 7.37 – 7.26 (m, 3H, ArH), 7.06 (dd, J = 10.2, 7.5, 1.1 Hz, 3H, ArH), 6.86 (dd, J = 1.7, 0.6 Hz, 1H, ArH).

* This transition appears to be not very clear, occurring on a large range.

1 In the light of the protons assignment of 2-SPM-SBF, we believe that 2 carbons are present at 128.40 ppm.
ArH), 6.68 – 6.60 (m, 3H, ArH); 13C NMR (75 MHz, CD₂Cl₂): δ 157.8 (CH), 155.2 (CH), 150.7 (C), 149.7 (C), 148.8 (C), 143.2 (C), 142.4 (C), 141.4 (C), 134.53 (C), 134.45 (C), 129.0 (CH), 128.6 (CH), 128.5 (2xCH seen from HSQC), 127.3 (CH), 124.42 (CH), 124.35 (CH), 122.9 (CH), 121.6 (CH), 121.1 (CH), 120.8 (CH), 66.6 (Spiro C); HRMS calculated for C₂₉H₁₈N₂ 395.1548, found 395.1548 [M+H⁺]; λₘₚ [nm] (ε[10⁴ L·mol⁻¹·cm⁻¹]) = 296 (2.5), 308 (2.5), 320 (1.8).

X-Ray:
Crystal was picked up with a cryoloop and then frozen at 150 K under a stream of dry N₂ on a APEX II Brucker AXS diffractometer for X-ray data collection (Mo Kα radiation, λ = 0.71073 Å). 4-Ph-SBF structure was published previously.⁴⁴

2-Ph-SBF (C₁₀₂H₆₀); M = 784.94. APEXII, Brucker-AXS diffractometer, Mo-Kα radiation (λ = 0.71073 Å), T = 150(2) K; triclinic P ̅ 1 (I. T. 2), a = 10.9027(4), b = 13.7902(6), c = 15.7771(5) Å, α = 102.1400(10), β = 97.4710(10), γ = 111.5900(10)°, V = 2099.47(14) Å³, Z = 2, d = 1.242 g·cm⁻³, μ = 0.07 mm⁻¹. The structure was solved by direct methods using the SIR97 program,⁵⁵ and then refined with full-matrix least-square methods based on F² (SHELXL-97)⁶⁶ with the aid of the WINGX⁶⁷ program. All non-hydrogen atoms were refined with anisotropic atomic displacement parameters. H atoms were finally included in their calculated positions. A final refinement on F² with 9543 unique intensities and 559 parameters converged at ωR(F²) = 0.1128 (R(F) = 0.0447) for 7649 observed reflections with I > 2σ(I).

Spectroscopic studies:
Cyclohexane (analytical grade, VWR) was used without further purification. Standard 1N solution of sulfuric acid was purchased from Alfa Aesar. UV-visible spectra were recorded using an UV-Visible spectrophotometer SHIMADZU UV-1605. The energy gap was calculated from the absorption edge of the UV-vis absorption spectrum in cyclohexane, using the formula ΔEₕₒₒ (eV) = hc/λ, λ being the absorption edge (in meter). With h = 6.6x10⁻³⁴ J·s (1eV = 1.6x10⁻¹⁹ J) and c = 3.0x10⁸ m·s⁻¹, this equation may be simplified as: ΔEₕₒₒ (eV) = 1237.5/λ (in nm). Triplet energy level Eₜ was calculated from the maximum of the first phosphorescence emission peak, and conversion in electron-volt was obtained with the previous formula. Emission spectra were recorded with a PTI spectrophuorimeter (PTI-814 PDS, MD 5020, LPS 220B) using a Xenon lamp. Quantum yields in solution (φₒₒ) were calculated relative to quinine sulfate (φₒₒ = 0.546 in H₂SO₄ 1N). φₒₒ was determined according to the following equation,

φₙₑᵣ = φₘᵣₑ × 100 × \frac{(Iₙ × Aₙ)}{(Iₗ × Aₗ)} × \frac{Iₖ}{Iₙ}

where, subscripts s and r refer respectively to the sample and reference. The integrated area of the emission peak in arbitrary units is given as T, n is the refracting index of the solvent (n = 1.426 for cyclohexane) and A is the absorbance. Three solutions of different concentration of the substrate (A<0.1) and three solutions of the reference (quinine sulfate) were prepared. The quinine sulfate concentration was chosen so as the absorption of the reference and the substrate were the same at the excitation wavelength. Three quantum yields were then calculated at this wavelength and the average value is reported. IR spectra were recorded on a Bruker Vertex 70 using a diamond crystal MIRacle ATR (Pike). Thin films were prepared by spin-coating 300 μL of solution of THF (10 g/L) on a sapphire plate (10 mm x 10 mm) at 2500 tr/min on a Süss MicroTech Labspin 6.

Fluorescence intensity decays were obtained by the time-correlated single-photon counting (TCSPC) method with femtosecond laser excitation using a Spectra-Physics set-up composed of a Titanium Sapphire Laser pumped by a doubled YAG laser Millennia. Light pulses at 990 nm (resp. 900 nm) were selected by optoacoustic crystals at a repetition rate of 4 MHz, and the third harmonics at 330 nm
(resp. 300 nm) was obtained through non-linear crystals. Fluorescence photons were detected through a monochromator by means of a Hamamatsu MCP R3809U photomultiplier. The time-to-amplitude converter was purchased from Tennelec. The fluorescence data were analyzed by a nonlinear least-squares global method using the Globals software package developed at the Laboratory for Fluorescence Dynamics at the University of Illinois, Urbana-Champaign.

**Electrochemical studies.** Electrochemical experiments were performed under argon atmosphere using a Pt disk electrode (diameter 1 mm), the counter electrode was a vitreous carbon rod and the reference electrode was a silver wire in a 0.1M AgNO₃ solution in CH₂CN. Ferrocene was added to the solution at the end of a series of experiments. The ferrocene/ferrocnium (Fc/Fc⁺) couple served as internal standard. The three electrodes cell was connected to a PAR Model 273 potentiostat/galvanostat (PAR, EG&G, USA) monitored with the ECHEM Software. Activated Al₂O₃ was added in the electrolytic solution to remove excess moisture. For a further comparison of the electrochemical and optical properties, all potentials are referred to the SCE electrode that was calibrated at ~0.405 V vs.Fc/Fc⁺ system. Following the work of Jenekhe,⁴⁸ we estimated the electron affinity (EA) or lowest unoccupied molecular orbital (LUMO) and the ionisation potential (IP) or highest occupied molecular orbital (HOMO) from the redox data (tangent to the oxidation or reduction wave). The LUMO level was calculated from: LUMO (eV)= [E_{onset}^{red} (vs SCE) +4.4] and the HOMO level from: HOMO (eV) = [E_{onset}^{ox} (vs SCE) + 4.4], based on an SCE energy level of 4.4 eV relative to the vacuum. The electrochemical gap was calculated from: ΔE° = |HOMO-LUMO| (in eV).

**Theoretical modeling.** Full geometry optimization with Density Functional Theory (DFT)⁶⁸ ⁶⁹ and Time-Dependent Density Functional Theory (TD-DFT) calculations were performed with the hybrid Becke-3 parameter exchange⁷⁰-⁷² functional and the Lee-Yang-Parr non-local correlation functional⁷³ (B3LYP) implemented in the Gaussian 09 (Revision B.01) program suite⁷⁶ using the 6-311G+(d,p) basis set and the default convergence criterion implemented in the program. The figures were generated with GaussView 5.0. The computed triplet adiabatic S₀ to T₁ excitation energies were calculated from the difference between the total energy of the molecule in their respective singlet and triplet states.

**Thermal analysis:**

Thermal Gravimetric Analysis (TGA) was carried out by using TA SDT Q600 instrument, at the "Ecole Nationale Supérieure de Chimie de Rennes" or at the "Institut des Sciences Analytiques" (UMR CNRS 5280) of Villeurbanne. TGA curves were measured at 10°C/min from 0 to 600°C under an azote atmosphere. Differential Scanning Calorimetry (DSC) was carried out by using NETZSCH DSC 200 F3 instrument equipped with an intracooler. DSC traces were measured at 10°C/min. Two heating/cooling cycles were successively carried out and the glass transition Tg was determined from the 2nd heating cycle. Temperatures of the different transitions given are obtained with the tangent to the transition peak.

**Device fabrication and characterisation:**

OLEDs based on a multilayer structure have been fabricated onto patterned ITO coated glass substrates from XinYan Tech (thickness: 100 nm and sheet resistance: less of 20 W/m). The organic materials (from Aldrich and Lumtec) are deposited onto the ITO anode by sublimation under high vacuum (< 10⁻⁵ Torr) at a rate of 0.2 – 0.3 nm/s. The structure of the device is the following: ITO/CuPc (10 nm)/NPB (40 nm)/TCTA (10 nm)/Hosts:dopant (20 nm)/TPBi (40 nm)/LiF (1.2 nm)/Al (100 nm). In this device, ITO is used as the anode, CuPc (copper phtalocyanine) is the hole injecting layer, NPB (N,N’-di(1-naphthyl)-N,N’-diphenyl-[1,1’-biphenyl]-4,4’-diamine) is the hole-transporting layer, TCTA (4,4’,4”-Tris(carbazol-9-yl)-triphenylamine) is the electron/exciton blocking layer, TPBi (1,3,5-Tris(1-phenyl-1H-benzimidazol-2-yl)benzene) is both the electron transporting layer and the hole blocking layer and a thin film of lithium fluoride over aluminum is the cathode. The entire device is fabricated in the same run without breaking the vacuum. In this study, the thicknesses of the different organic layers were kept constant for all the devices. The active area of the devices defined by the overlap of the ITO anode and the metallic cathode was 0.3 cm². The current-voltage-luminance (I-V-L) characteristics of the devices were measured with a regulated power supply (Laboratory Power Supply EA-PS 3032-10B) combined with a multimeter and a cm² area silicon calibrated photodiode (Hamamatsu). The spectral emission was recorded with a SpectraScan PR650 spectrophotometer. All the measurements were performed at room temperature and at ambient atmosphere with no further encapsulation of devices.


Supplementary Materials for

2-substituted vs 4-substituted-9,9'-spirobifluorene host materials
for green and blue phosphorescent OLEDs: A Structure-Property Relationship
Study

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Summary

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## X-ray Diffraction

### Table 1: Crystal data and structure refinement for 2-Ph-SBF

<table>
<thead>
<tr>
<th>Identification code</th>
<th>2-Ph-SBF</th>
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<tr>
<td>Empirical formula</td>
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<tr>
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<td>F(000)</td>
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<tr>
<td>Crystal size</td>
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<td>Completeness to theta = 27.50°</td>
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<td>Final R indices [I&gt;2sigma(I)]</td>
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<tr>
<td>R indices (all data)</td>
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<tr>
<td>Largest diff. peak and hole</td>
<td>0.261 and -0.269 e Å⁻³</td>
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**Table 2** Crystal data and structure refinement for 2-5Pm-SBF

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<th>Characteristic</th>
<th>2-5Pm-SBF</th>
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</tr>
<tr>
<td>Empirical formula</td>
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<tr>
<td>Formula weight</td>
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<td>Temperature</td>
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</tr>
<tr>
<td>Wavelength</td>
<td>0.71073 Å</td>
</tr>
<tr>
<td>Crystal system</td>
<td>orthorhombic</td>
</tr>
<tr>
<td>space group</td>
<td>P c a b</td>
</tr>
</tbody>
</table>
| Unit cell dimensions                        | a = 8.4342(3) Å, α = 90°  
|                                             | b = 13.6588(5) Å, β = 90°  
|                                             | c = 38.5506(13) Å, γ = 90° |
| Volume                                      | 4441.1(3) Å³ |
| Z                                           | 8         |
| Calculated density                          | 1.276 (g.cm⁻³) |
| Absorption coefficient                      | 0.078 mm⁻¹ |
| F(000)                                      | 1792      |
| Crystal size                                | 0.36 x 0.31 x 0.05 mm |
| Theta range for data collection             | 1.06 to 27.46° |
| h_min, h_max                                | -10, 9     |
| k_min, k_max                                | -17, 16    |
| l_min, l_max                                | -45, 50    |
| Reflections collected / unique              | 21000 / 5055 [R(int) = 0.0327] |
| Reflections [I>2σ]                          | 3820       |
| Completeness to theta_max                   | 0.994      |
| Absorption correction type                  | multi-scan |
| Refinement method                           | Full-matrix least-squares on R² |
| Data / restraints / parameters              | 5055 / 0 / 300 |
| Goodness-of-fit                             | 1.124      |
| Final R indices [I>2σ]                      | R1 = 0.0461, wR2 = 0.1246 |
| R indices (all data)                        | R1 = 0.0711, wR2 = 0.1511 |
| Largest diff. peak and hole                 | 0.36 and -0.315 e⁻.Å⁻³ |
2-Ph-SBF

2-Ph-SBF MOLECULE 1

S1 Top: Angle between the mean plane of the pendant phenyl ring and that of its attached phenyl ring of the fluorene: 37.37°; Bottom: Angle between the two mean planes of each fluorene: 7.08°
**2-Ph-SBF MOLECULE 2**

\[ \text{S. 2 Angle between the mean plane of the pendant phenyl ring and that of its attached phenyl ring of the fluorene: } 4.58^\circ \]
S.3 Short contacts (CH and C-C, distance < (sum of $r_{vdw}$) - 0.1) detected in crystal packing of 2-Ph-SBF.

S.4 Short contacts (CH and C-C, distance < (sum of $r_{vdw}$)) detected for the 2-Ph-SBF molecule 2 (in red above)
**S5** Short contacts (CH and C-C$_s$ distance < (sum of r$_{vdw}$)) detected for the pendant phenyl unit of 2-Ph-SBF molecule 2 (in red above)

**S6** Short contacts (CH and C-C$_s$ distance < (sum of r$_{vdw}$)) detected for the pendant phenyl unit of 2-Ph-SBF molecule 1 (in blue above)
$S\,7\quad 2\text{-}5\text{Pm-SBF}$: Angle between the mean plane of the pyrimidine plane and that of its attached phenyl ring: 23.69°

**Photophysical properties**

![Normalized absorption vs Wavelength graph]

- Cyclohexane
- Toluene
- THF
- DCM
- EtOH
- MeCN
S8 UV-Vis absorption spectrum of 4-Ph-SBF in various solvents

S9 UV-Vis absorption spectrum of 4-5Pm-SBF in various solvents

S10 UV-Vis absorption spectrum of 2-Ph-SBF in various solvents
**S 11** UV-Vis absorption spectrum of 2-5Pm-SBF in various solvents

**S 12** Quantum yield measurements: Absorption of solutions of 4-5Pm-SBF in cyclohexane and quinine sulfate (QS) in H₂SO₄ 1N
Table 2 Quantum yield calculation of 4-5Pm-SBF

<table>
<thead>
<tr>
<th>Solution</th>
<th>$\lambda$ (nm)</th>
<th>$\Lambda$</th>
<th>$T_S$</th>
<th>$T_{QS}$</th>
<th>$n_{D}^{25}$ (cyclohexane)</th>
<th>$n_{D}^{25}$ (H$_2$SO$_4$ 1N)</th>
<th>$\varphi$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>308.5</td>
<td>0.1011</td>
<td>14134.4</td>
<td>238032</td>
<td>1.42662</td>
<td>1.3325</td>
<td>4%</td>
</tr>
<tr>
<td></td>
<td>299</td>
<td>0.0721</td>
<td>7610.04</td>
<td>120670</td>
<td>1.42662</td>
<td>1.3325</td>
<td>4%</td>
</tr>
<tr>
<td>2</td>
<td>309.5</td>
<td>0.0569</td>
<td>10314.6</td>
<td>150249</td>
<td>1.42662</td>
<td>1.3325</td>
<td>4%</td>
</tr>
<tr>
<td></td>
<td>302</td>
<td>0.0442</td>
<td>6316.68</td>
<td>89861.1</td>
<td>1.42662</td>
<td>1.3325</td>
<td>4%</td>
</tr>
<tr>
<td>3</td>
<td>309</td>
<td>0.0435</td>
<td>8040.89</td>
<td>122287</td>
<td>1.42662</td>
<td>1.3325</td>
<td>4%</td>
</tr>
<tr>
<td></td>
<td>300.5</td>
<td>0.0328</td>
<td>4539.62</td>
<td>67376.1</td>
<td>1.42662</td>
<td>1.3325</td>
<td>4%</td>
</tr>
</tbody>
</table>

S 13 Quantum yield measurements: Absorption of solutions of 4-5Pm-SBF in tetrahydrofurane and quinine sulfate (QS) in H$_2$SO$_4$ 1N

Table 3 Quantum yield calculation of 4-5Pm-SBF

<table>
<thead>
<tr>
<th>Solution</th>
<th>$\lambda$ (nm)</th>
<th>$\Lambda$</th>
<th>$T_S$</th>
<th>$T_{QS}$</th>
<th>$n_{D}^{25}$ (cyclohexane)</th>
<th>$n_{D}^{25}$ (H$_2$SO$_4$ 1N)</th>
<th>$\varphi$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>299</td>
<td>0.073</td>
<td>5032.56</td>
<td>60164.3</td>
<td>1.42662</td>
<td>1.3325</td>
<td>5%</td>
</tr>
<tr>
<td>2</td>
<td>299</td>
<td>0.058</td>
<td>4262.56</td>
<td>52110.2</td>
<td>1.42662</td>
<td>1.3325</td>
<td>5%</td>
</tr>
<tr>
<td>3</td>
<td>298</td>
<td>0.049</td>
<td>5132.74</td>
<td>43404.5</td>
<td>1.42662</td>
<td>1.3325</td>
<td>7%</td>
</tr>
</tbody>
</table>
Quantum yield measurements: Absorption of solutions of 2-Ph-SBF in cyclohexane and quinine sulfate (QS) in H2SO4 1N

Table 4 Quantum yield calculation of 2-Ph-SBF

<table>
<thead>
<tr>
<th>Solution</th>
<th>λ (nm)</th>
<th>A</th>
<th>T_S</th>
<th>T_QS</th>
<th>n_DSI (cyclohexane)</th>
<th>n_DSI (H2SO4 1N)</th>
<th>φ</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>312</td>
<td>0.0815</td>
<td>65759.6</td>
<td>48449.5</td>
<td>1.42662</td>
<td>1.3325</td>
<td>85%</td>
</tr>
<tr>
<td>2</td>
<td>312</td>
<td>0.064</td>
<td>51316.1</td>
<td>37356.8</td>
<td>1.42662</td>
<td>1.3325</td>
<td>86%</td>
</tr>
<tr>
<td>3</td>
<td>312.5</td>
<td>0.0413</td>
<td>36521.4</td>
<td>25376.8</td>
<td>1.42662</td>
<td>1.3325</td>
<td>90%</td>
</tr>
</tbody>
</table>
Quantum yield measurements: Absorption of solutions of 2-Ph-SBF in cyclohexane and quinine sulfate (QS) in H₂SO₄ 1N

Table 5: Quantum yield calculation of 2-Ph-SBF

<table>
<thead>
<tr>
<th>Solution</th>
<th>( \lambda ) (nm)</th>
<th>A</th>
<th>( T_s )</th>
<th>( T_{QS} )</th>
<th>( n_{D}^{25} ) (cyclohexane)</th>
<th>( n_{D}^{25} ) (H₂SO₄ 1N)</th>
<th>( \varphi )</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>301</td>
<td>0.113</td>
<td>119399</td>
<td>95186.3</td>
<td>1.42662</td>
<td>1.3325</td>
<td>79%</td>
</tr>
<tr>
<td>2</td>
<td>299</td>
<td>0.100</td>
<td>96821.3</td>
<td>77070.8</td>
<td>1.42662</td>
<td>1.3325</td>
<td>79%</td>
</tr>
<tr>
<td>3</td>
<td>300</td>
<td>0.061</td>
<td>73910.6</td>
<td>57692.5</td>
<td>1.42662</td>
<td>1.3325</td>
<td>80%</td>
</tr>
</tbody>
</table>
**S 16** Quantum yield measurements: Absorption of solutions of 2-SPm-SBF in cyclohexane and quinine sulfate (QS) in H₂SO₄ 1N

**Table 6** Quantum yield calculation of 2-SPm-SBF

<table>
<thead>
<tr>
<th>Solution</th>
<th>λ (nm)</th>
<th>A</th>
<th>Tₛ</th>
<th>TQS</th>
<th>n₁D²⁵ (cyclohexane)</th>
<th>n₁D²⁵ (H₂SO₄ 1N)</th>
<th>Φ</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>309</td>
<td>0.129</td>
<td>10082.1</td>
<td>240980</td>
<td>1.42662</td>
<td>1.3325</td>
<td>3%</td>
</tr>
<tr>
<td>2</td>
<td>309.5</td>
<td>0.0881</td>
<td>6580.11</td>
<td>176643</td>
<td>1.42662</td>
<td>1.3325</td>
<td>2%</td>
</tr>
<tr>
<td>3</td>
<td>309.5</td>
<td>0.0593</td>
<td>5162.04</td>
<td>134942</td>
<td>1.42662</td>
<td>1.3325</td>
<td>2%</td>
</tr>
</tbody>
</table>
**S1** Quantum yield measurements: Absorption of solutions of 2-5Pm-SBF in cyclohexane and quinine sulfate (QS) in H$_2$SO$_4$ 1N

**Table 7** Quantum yield calculation of 2-5Pm-SBF

<table>
<thead>
<tr>
<th>Solution</th>
<th>λ (nm)</th>
<th>Λ</th>
<th>$T_S$</th>
<th>$T_{QS}$</th>
<th>$n_D^{23}$ (cyclohexane)</th>
<th>$n_D^{23}$ (H$_2$SO$_4$ 1N)</th>
<th>φ</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>299</td>
<td>0.100</td>
<td>2965.74</td>
<td>77070.8</td>
<td>1.42662</td>
<td>1.3325</td>
<td>2%</td>
</tr>
<tr>
<td>2</td>
<td>299</td>
<td>0.074</td>
<td>2565.15</td>
<td>63262.2</td>
<td>1.42662</td>
<td>1.3325</td>
<td>3%</td>
</tr>
<tr>
<td>3</td>
<td>299</td>
<td>0.058</td>
<td>2084.54</td>
<td>52110.2</td>
<td>1.42662</td>
<td>1.3325</td>
<td>3%</td>
</tr>
</tbody>
</table>
THEORETICAL MODELING

Predicted UV-vis Spectra from TD-DFT energy calculations

SBF

4-Ph-SBF

4-5Pm-SBF

2-5Pm-SBF

2-Ph-SBF

S 2 Predicted UV-Vis spectra from TD-DFT energy calculations
SBF

LUMO+1: -1.25 eV

LUMO+2: -0.84 eV

LUMO+3: -0.84 eV

LUMO: -1.25 eV

HOMO: -5.99 eV

HOMO-2: -6.79 eV

HOMO-1: -6.26 eV

Excited state 1
λ: 297.6 nm
f: 0.064

Excited state 2
λ: 297.59 nm
f: 0.064

Excited states 3 and 4
f: 0

Excited state 5
λ: 271.95 nm
f: 0.1449

Excited state 6
λ: 271.95 nm
f: 0.1449

Excited state 7
λ: 261.09 nm
f: 0.0905

Excited state 8
λ: 261.09 nm
f: 0.0906

§3 Calculated frontier molecular orbitals by DFT and the 6th first calculated electronic transitions by TD-DFT of SBF, after geometry optimization with DFT B3LYP/6-311G+(d,p), show with an isovalue of 0.04.
4-Ph-SBF

LUMO+1: -1.24 eV
LUMO+2: -0.94 eV
LUMO+3: -0.84 eV

LUMO: -1.27 eV

HOMO: -5.97 eV

HOMO-1: -6.25 eV

Excited state 1
\( \lambda: 300.25 \text{ nm} \)
f: 0.06
Excited state 2
\( \lambda: 298.49 \text{ nm} \)
f: 0.059
Excited states 3 and 4
f: 0
Excited state 5
\( \lambda: 276.78 \text{ nm} \)
f: 0.1519
Excited state 6
\( \lambda: 274.41 \text{ nm} \)
f: 0.1249
Excited state 7
\( \lambda: 266.68 \text{ nm} \)
f: 0.06
Excited state 8
f: 0
Excited state 9
\( \lambda: 263.08 \text{ nm} \)
f: 0.01

\textbf{S4} Calculated frontier molecular orbitals by DFT and the first calculated electronic transitions by TD-DFT of 4-Ph-SBF, after geometry optimization with DFT B3LYP/6-311G+(d,p), show with an isovalue of 0.04.
4-5Pm-SBF

LUMO+1: -1.645 eV
LUMO+2: -1.38 eV
LUMO+3: -1.18 eV

Excited state 1
λ: 315.225 nm
f: 0.0198
Excited state 2
λ: 311.07 nm
f: 0.0303
Excited state 4
λ: 293.96 nm
f: 0.0744
Excited state 5
λ: 292.97 nm
f: 0.0937
Excited state 9
λ: 277.29 nm
f: 0.0513
Excited state 10
λ: 270.34 nm
f: 0.1271

HOMO: -6.16 eV
HOMO-1: -6.25 eV

S 5 Calculated frontier molecular orbitals by DFT and the 6th first calculated electronic transitions by TD-DFT of 4-5Pm-SBF, after geometry optimization with DFT B3LYP/6-311G+(d,p), show with an isovalue of 0.04
2-Ph-SBF

LUMO+1: -1.26 eV  LUMO+2: -0.88 eV  LUMO+3: -0.86 eV

LUMO: -1.48 eV

HOMO: -5.87 eV

Excited state 1
λ: 315.93 nm
f: 0.3552
Excited state 3
λ: 295.96 nm
f: 0.3346
Excited state 6
λ: 278.70 nm
f: 0.1324
Excited state 7
λ: 268.32 nm
f: 0.0941

HOMO-1: -6.17 eV

S 22 Calculated frontier molecular orbitals by DFT and the 6th first calculated electronic transitions by TD-DFT of 2-Ph-SBF, after geometry optimization with DFT B3LYP/6-311G+(d,p), show with an isovalue of 0.04
2-5Pm-SBF

LUMO+1: -1.50 eV  LUMO+2: -1.38 eV

LUMO: -1.85 eV

HOMO: -6.13 eV

HOMO-1: -6.37 eV

Excited state 1
\( \lambda: \) 327.18 nm
f: 0.1677

Excited state 2
\( \lambda: \) 304.91 nm
f: 0.1553

Excited state 3
\( \lambda: \) 304.41 nm
f: 0.239

Excited state 10
\( \lambda: \) 272.97 nm
f: 0.1497

\textbf{S 23} Calculated frontier molecular orbitals by DFT and the 6 first calculated electronic transitions by TD-DFT of 2-5Pm-SBF, after geometry optimization with DFT B3LYP/6-311G+(d,p), show with an isovalue of 0.04
THERMAL PROPERTIES

![Graph showing DSC curves of 4-5Pm-SBF](image)

\[ T_m = 164 \, ^\circ C \]

\[ T_g = 80 \, ^\circ C \]

*DSC curves of 4-5Pm-SBF*
$T_g = 92 \degree C$

$T_m = 194 \degree C$

S 26 DSC curves of 2-5Pm-SBF
ELECTROCHEMICAL PROPERTIES

4-5Pm-SBF in CH$_2$Cl$_2$ + 0.2 M Bu$_4$NPF$_6$

$E_{\text{onset}}$ = -2.17 V
LUMO: -2.23 eV

4-5Pm-SBF in THF + 0.2M Bu$_4$NPF$_6$

$E_{\text{onset}}$ = -2.17 V
LUMO: -2.23 eV

S 6 Cyclic voltammetry of 4-5Pm-SBF 5x10$^{-3}$ M in CH$_2$Cl$_2$ (left) or THF (right) + Bu$_4$NPF$_6$ 0.2 M, sweep-rate 100 mV.s$^{-1}$, platinum disk (Ø: 1mm) working electrode.

2-5Pm-SBF in CH$_2$Cl$_2$ + 0.2M Bu$_4$NPF$_6$

$E_{\text{onset}}$ = -2.10 V
LUMO: -2.3 eV

2-5Pm-SBF in THF + 0.2M Bu$_4$NPF$_6$

$E_{\text{onset}}$ = -2.1 V
LUMO: -2.3 eV

S 7 Cyclic voltammetry of 2-5Pm-SBF 2x10$^{-3}$ M in CH$_2$Cl$_2$ (left) or THF (right) + Bu$_4$NPF$_6$ 0.2 M, sweep-rate 100 mV.s$^{-1}$, platinum disk (Ø: 1mm) working electrode.
**S 8** Cyclic voltammetry of 4-Ph-SBF $5 \times 10^{-3}$ M in CH$_2$Cl$_2$ (left) or THF (right) + Bu$_4$NPF$_6$ 0.2 M, sweep-rate 100 mV.s$^{-1}$, platinum disk (Ø: 1mm) working electrode.

**S 30** Cyclic voltammetry of 2-Ph-SBF $5 \times 10^{-3}$ M in CH$_2$Cl$_2$ (left) or THF (right) + Bu$_4$NPF$_6$ 0.2 M, sweep-rate 100 mV.s$^{-1}$, platinum disk (Ø: 1mm) working electrode.
**S 31 Left:** Cyclic voltammetry of 2-Ph-SBF 5 $10^{-3}$ M in CH$_2$Cl$_2$ + Bu$_4$NPF$_6$ 0.2 M, 10 repeated cycles between 495 and 2045 mV, sweep-rate 100 mV.s$^{-1}$, platinum disk ($Ø$: 1mm) working electrode. **Right:** Cyclic voltammetry in CH$_2$Cl$_2$ + Bu$_4$NPF$_6$ 0.2 M, 3 repeated cycles between 580 and 1895 mV, sweep-rate 100 mV.s$^{-1}$, platinum disk ($Ø$: 1mm) covered previously in left with poly(2-Ph-SBF) working electrode.

**S 32 Left:** Cyclic voltammetry of 4-Ph-SBF 5 $10^{-3}$ M in CH$_2$Cl$_2$ + Bu$_4$NPF$_6$ 0.2 M, 12 repeated cycles between 0.28 and 2.13 V, sweep-rate 100 mV.s$^{-1}$, platinum disk ($Ø$: 1mm) working electrode. **Right:** Cyclic voltammetry in CH$_2$Cl$_2$ + Bu$_4$NPF$_6$ 0.2 M, 3 repeated cycles between 280 and 1530 mV, sweep-rate 100 mV.s$^{-1}$, platinum disk ($Ø$: 1mm) covered previously in left with poly(4-Ph-SBF) working electrode.
**S33 Left:** Cyclic voltammetry of 4-5Pm-SBF $5 \times 10^{-3}$ M in CH$_2$Cl$_2$ + Bu$_4$NPF$_6$ 0.2 M, 10 repeated cycles between 0.35 and 2.35 V, sweep-rate 100 mV.s$^{-1}$, platinum disk (Ø: 1mm) working electrode. **Right:** Cyclic voltammetry in CH$_2$Cl$_2$ + Bu$_4$NPF$_6$ 0.2 M, 3 repeated cycles between 350 and 1750 mV, sweep-rate 100 mV.s$^{-1}$, platinum disk (Ø: 1mm) covered previously in left with poly(4-5Pm-SBF) working electrode.
4-5Pm-SBF-HSQC-CD$_2$Cl$_2$
4-5Pm-SBF-HSQC (Zoom)-CD$_2$Cl$_2$
2-Ph-SBF-HSQC (Zoom)-CD$_2$Cl$_2$
2-5Pm-SBF-HSQC-CD$_2$Cl$_2$
2-5Pm-SBF-HSQC (Zoom)-CD$_2$Cl$_2$
$2\text{-}5\text{Pm-SBF}$-HMBC (C Spiro)-CD$_2$Cl$_2$
COPY OF NMR SPECTRA

$2^1\text{H-CD}_2\text{Cl}_2$

Solvent residual peak

Chemical shift (ppm)
$^2\text{H}^1\text{C}-\text{CD}_2\text{Cl}_2$

Solvent residual peak
4-5Pm-SBF-$^{13}$C-CD$_2$Cl$_2$
4-5Pm-SBF-DEPT-CD₂Cl₂
$2\text{-Ph-SBF}^{-1}\text{H-CD}_{2}\text{Cl}_2$
2-5Pm-SBF-^1^H-C^2^Cl^2^
2-5Pm-SBF-\textsuperscript{13}C-CD\textsubscript{2}Cl\textsubscript{2}
COPY OF MASS SPECTRA

Centre régional de mesures physiques de l'Ouest (CRMPO) - RAPPORT D'ANALYSE

Analysis Info
Analysis Name: D:\Data\CRMPO\ASAP_156_MS_01.d
Method: ASAP buoy wax 50_30K0_pos.m
Sample Name: 45 Pm SBF
Comment: G. THIERY 45 Pm SBF Température : 165°C

Acquisition Date: 3/11/2013 4:48:36 PM
Operator: Fabian LAMBERT
Instrument: microTOF-Q II

S.9 High Resolution Mass Spectrum of 4-5Pm-SBF