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Intramolecular excimer emission as a blue light source in fluorescent organic light emitting diodes: a promising molecular design

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Intramolecular excimer emission arising from organic molecules as a blue light source in fluorescent Small Molecule Organic Light Emitting Diodes (SMOLEDs) is almost absent from the literature. In this work, three aryl-substituted DiSpiroFluorene–IndenoFluorenes (DSF–IFs 1–3) possessing different fluorescent properties due to their different main emitters have been investigated through a structure–property relationship study. Due to its particular geometry, the rigid DSF–IF platform 2 allows an ‘aryl/fluorene/aryl’ dimer to be preformed in the ground state leading, in the excited state, to a deep blue fluorescent emission through strong π–π intramolecular interactions between the two ‘aryl/fluorene/aryl’ arms. 2 has been successfully used as an emitting layer in a SMOLED with electroluminescence arising from electrogenerated intramolecular excimers and the properties of these excimer-based OLEDs have been compared to those of two model compounds (1 and 3). The simple and non-optimized double-layer device displays a deep blue colour (CIE coordinates: 0.19; 0.18) exhibiting a luminance of 510 Cd m\(^{-2}\) with a luminous efficiency of ca. 0.1 Cd A\(^{-1}\). This work is, to the best of our knowledge, the first rational and comparative study describing an intramolecular excimer-based-SMOLED.

Introduction

Intramolecular π–π interactions play a key role in the field of Materials Science and especially in Organic Electronics. Indeed, in an Organic Field Effect Transistor (OFET), intramolecular π–π interactions between molecules in the thin solid film must be very strong to ensure high mobility of charge carriers.\(^1\)\(^4\) Conversely, in an Organic Light Emitting Diode (OLED), intramolecular π–π interactions should be usually suppressed to avoid any parasite and uncontrolled emission colour due to intermolecular excimer formation (dimer in the excited state). However, in white OLEDs the use of a single fluorescent emitter which can form an excimer through intramolecular π–π interactions has been designed in order to cover the whole visible range from 400 to 700 nm.\(^5\)\(^7\) Thus, due to their importance in OLED technology, intramolecular π–π interactions have been hence extensively studied for the last twenty years, but intramolecular π–π interactions have not been deeply investigated. Indeed, if one can control the formation of intramolecular π–π interactions within a fluorescent dye and in the meantime avoid intermolecular π–π interactions, then intramolecular excimer emission could be used as a light source in an OLED. Indeed, the strategy to generate blue light (and light in general) in an OLED is most of the time always the same and consists of designing a fluorophore with emission properties directly arising from its π-conjugated backbone modulated (or not) by the electronic effects of different substituents (for example, donor–acceptor π-conjugated dyes).\(^8\)\(^\text{12}\) In this work, we wish to report a unique and promising strategy to generate light in an OLED, that is using intramolecular excimer emission. Recently, our group has designed an original family of aryl-substituted DiSpiroFluorene–IndenoFluorene (DSF–IF) derivatives, which possess two ‘aryl-fluorene-aryl’ moieties in a rigid face-to-face arrangement. This particular face-to-face geometry predominantly leads for these ‘3π–2 spiro’\(^\ddagger\) compounds\(^9\)\(^\text{13}\) to conformationally controllable intramolecular excimer fluorescence emission.\(^14\)\(^\text{16}\) However, these systems have been mainly studied in solution and never incorporated in an optoelectronic device. In the present work, we wish to report preliminary results on intramolecular excimer emission as a deep blue light source in an OLED. Three structurally related DSF–IFs 1–3 (Scheme 1) possessing different emission properties have been investigated through a structure–property relationship study. Their optical properties have first been investigated in detail before successfully incorporating them as an emitting layer (EML) in blue SMOLEDs. The performances and electro-optic properties have been discussed. If the

\(\ddagger\) ‘3π–2 spiro’ is a geometric concept meaning three π-conjugated systems linked by two spiro bridges. See ref. 13.
The use of intermolecular excimer emission as a light source in a SMOLED is known for organic5–7 and organometallic compounds,15–20 the use of intramolecular excimer emission arising from pure organic molecules has been reported, to the best of our knowledge, only once.21 In the light of the marked difference observed between the excimer-based SMOLEDs (with \( \text{EML} \)) and the “classical” SMOLEDs (with \( \text{EML} \)) or \( \text{EML} \)), we are convinced that this strategy may open new avenues in the design of efficient blue fluorophores for optoelectronic applications.

**Results and discussion**

The molecular design adopted is the following: the \((2,1-a)-\text{DSF–IF}\) platform found in 2 has been chosen in order to impose a rigid arrangement of the two cofacial “aryl/fluorene/aryl” arms and to allow these arms to interact with one another in the ground state. This will then favour the facile and easily controllable formation of excimers in the excited state. In addition, the 3D geometry of 2 should suppress the non-desired intermolecular \( \pi-\pi \) interactions in thin film leading to a stable blue color only arising from intramolecular \( \pi-\pi \) interactions.

Two other molecules (1 and 3), with a very similar molecular structure but with ‘‘classical’’ fluorescence properties arising from their \( \pi \)-conjugated backbone (without any excimer emission), will be also investigated, for comparison purpose, in order to evaluate the efficiency of using intramolecular excimer emission in an OLED. The molecule 1, a regioisomer of 2, presents a \((2,1-b)-\text{DSF–IF}\) core and two “3,4,5-trimethoxyphenyl/fluorene/3,4,5-trimethoxyphenyl” arms but the latter do not interact with one another (as it is the case in 2). The molecule 3 is also constituted of a \((2,1-b)-\text{DSF–IF}\) core, but this time the indenofluorenyl unit is substituted by two 3,4,5-trimethoxyphenyl moieties.

**Synthesis**

The synthesis of 1 and 2 has been previously reported.22,23,24,25,26,27 This has been synthesized through a Pd-catalyzed Suzuki–Miyaura cross-coupling reaction between the related dibromo-indenofluorene substituted compound \((1,2-b)-\text{DSF}-(\text{t-Bu})_2\text{IF}(\text{Br})_2\) ref. 12 and 3,4,5-trimethoxyphenylboronic acid using \( \text{Pd}_2(\text{dba})_2/\text{P}(\text{t-Bu})_3 \) as the catalytic system, potassium carbonate as the base in a mixture of toluene and water at 100 °C (yield: 55%, Scheme 2).

**Scheme 2** Synthesis of 3.

**Optical properties**

The absorption spectrum of 1, in solution in THF (Fig. 1A), is quite well defined with one shoulder at 301 nm and four maxima at \( \lambda = 314, 331, 337, 345 \) nm.14 As previously reported, the three main bands have been ascribed to the “aryl–fluorene–aryl” moieties as these bands are also found in other spirolinked oligoaromatic derivatives.22–24 The band at 345 nm has been assigned to the \( \pi-\pi^* \) electronic transition of the \((2,1-b)-\text{indenofluorenyl}\) core in perfect accordance with that of its non-substituted (without aryl rings) congener \((1,2-b)-\text{DSF–IF}\) \((\lambda_{\text{max}} = 345 \text{ nm})\).25,26 It should be stressed that this band is red-shifted by 11 nm compared to that of \((1,2-b)-\text{indenofluorenyl}\) \((1,2-b)-\text{IF}, \lambda_{\text{max}} = 334 \text{ nm})\), due to the electronic effects of the two spirolinked diarylfluorene units.25–27 The absorption spectrum of 2, in solution in THF (Fig. 1B), is less defined compared to that of 1 with nevertheless one shoulder at 314 nm and two maxima at 328 and 340 nm. As mentioned above, the bands at 314 and 328 nm have been ascribed to the “aryl–fluorene–aryl” moieties. The transition recorded at 340 nm has been assigned to the \((2,1-a)-\text{indenofluorenyl}\) core in accordance with that of its non-substituted congener \((2,1-a)-\text{DSF–IF}, \lambda_{\text{max}} = 339 \text{ nm})\).28 The slight blue-shift observed between the maxima of 2 \((\lambda_{\text{max}} = 340 \text{ nm})\) and 1 \((\lambda_{\text{max}} = 345 \text{ nm})\) has been assigned to a better delocalization of \( \pi \)-electrons in 1 compared to 2 due to the different indenoarylfluorene cores \((1,2-b\ vs. \ 2,1-a)\).25,26 In addition, a red shift of 18 nm is detected between the maximum of 2 and that of \((2,1-a)-\text{indenofluorenyl}\) \((2,1-a)-\text{IF}, \lambda_{\text{max}} = 322 \text{ nm})\), due to the electronic effects of the two spirolinked diarylfluorene units. The electronic influence on the indenoarylfluorenyl core of a cofacial diaryl-fluorene \( \pi \)-dimer found in 2, leading to a 18 nm red shift, is different from that of two non-directly interacting diaryl-fluorene units as found in 1 (red shift: 11 nm, vide supra). Such a difference was also observed in the \((2,1-a)-\text{IF}/(2,1-a)-\text{DSF–IF}\) (red shift: 17 nm) and \((1,2-b)-\text{IF}/(1,2-b)-\text{DSF–IF}\) (red shift: 11 nm) series. Thus, isomers 1 and 2 present absorption properties governed by both their indenoarylfluorenyl and their “aryl–fluorene–aryl” moieties.

DSF–IF 3 displays a different behaviour due to its different substitution pattern compared to 1 and 2. Indeed, 3 possesses a \((2,1-b)-\text{indenofluorenyl}\) core substituted by two 3,4,5-trimethoxyphenyl moieties. Thus, the optical properties of 3 should be almost fully governed by the “aryl/indenofluorenyl/aryl” moiety. Indeed, 3 presents an absorption spectrum with a maximum recorded at 362 nm, red shifted by 17 nm compared to 1 due to the extension of the conjugation length of the 2,6-diaryl-(1,2-b)-indenofluorenyl core. This wavelength (362 nm) has been assigned to a \( \pi-\pi^* \) transition of the “aryl/indenofluorenyl/aryl” moiety as it fits well with the maximum reported for a structurally related compound, that is 2,6-diphenyldibenzoindolylfluorenyl \((\lambda_{\text{max}} = 350 \text{ nm})\) in DMF \(28 \) with nevertheless a 12 nm red shift caused by the electron donating behaviour of the six methoxy groups borne by 3.28–30

In thin film, 1–3 display the same behaviour that is a broader absorption spectrum and a slight red shift, usually assigned to the different environment surrounding the molecules (solid vs. liquid) and hence the different dielectric constants.22,31

The fluorescence spectrum of 1, in solution in THF (Fig. 2A), which mainly arises from its “aryl/fluorene/aryl” moieties, presents two maxima at 381 and 393 nm.44 These maxima are red
shifted compared to those of 2,2′,7,7′-tetraphenyl-9,9′-spirobi-
fluorene (λ = 359, 378 nm), which can be considered as two
spirolinked “phenyl/fluorene/phenyl” units.22 This red shift,
ascribed to the electron-donating effect of the methoxy groups
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and has been ascribed to the electron-donating effect of the methoxy groups
This behaviour highlights the remarkable effect of the cofacial
fluorenes arrangement found in 2 and has been ascribed to the
emission of intramolecular excimers, due to the interactions
between “aryl–fluorene–aryl” moieties in the excited state.5 Several
Several groups have also reported similar behaviour for various
molecular systems with π–π interactions.32–40 An important
feature in the fluorescent spectrum of 2 is related to the large
Stokes shift i.e. 116 nm which leads to an emission in the deep
blue region. This feature is of key importance as no self-
absorption may hence occur in this blue emitter. The well-
resolved emission spectrum of 3 (Fig. 2C, λmax = 391/412 nm)
compared to its absorption spectrum (Fig. 1C) suggests a more
rigid/planar structure of 3 in the excited state since the bonds
joining the indenofluorene unit and the aryl rings acquire some
double bond characters.16,41,42 Indeed, for species that comprise
torsional degrees of freedom in their π-conjugated core, varying
degrees of deviation from the classical mirror image behaviour
can be detected. Such differences between absorption and emission
bandshapes may be related to the flexible character of the
molecules, as already studied for p-terphenyl and indenofluorene
derivatives by ab initio quantum chemical methods.43 Compared
to 1, there is a red shift of the two main maxima (11 and 19 nm)
for 3 due to the extension of the conjugation length from diaryl-
fluorene in 1 to diaryl-indenofluorene units in 3. However and
despite this extension of the conjugation length, the emission
maxima of 3 (λmax = 391/412 nm) are remarkably blue-shifted
compared to that of 2 (λmax = 457 nm, vide infra). This feature
highlights the efficiency of the molecular design found in 2, which
allows reaching a deep blue emission with relatively short π-
conjugated molecular fragments (i.e. “aryl/fluorene/aryl”),
whereas this deep blue emission wavelength is almost impossible
to reach with structurally related “aryl–fluorene–aryl” deriva-
tives.44 For example, in the series of comparable spiro–aryl-
fluorene–aryl compounds, described by Salbeck and co-workers,
possessing two, four, six, or eight phenyl units connected to the
fluorene core, the fluorescence maxima in solution only gradually
increase in the order 359, 385, 395, and 402 nm.45 Thus, reaching
an emission wavelength of 460 nm, as it is the case for 2, seems to
be almost impossible (even with the introduction of appropriate
electron-donor substituents) as a limiting value for long chains
exists.24 Indeed, even bridged-polyphenylene derivatives such as
polyfluorene, polyindenofluorene or polypentaphenylene deriva-
tives, known to possess high wavelengths in the blue region,
only present an emission maximum at around 420, 430 nm and
445 nm respectively.44 The use of intramolecular excimers emis-
sion resulting from ‘aryl–fluorene–aryl’ fragments to obtain deep
blue fluorescent emission appears hence as a possible solution to
this issue.

In thin-solid film, 1 and 3 present a broader and red-shifted
spectrum compared to their solution ones due to the different
dielectric constants of the environments (liquid vs. solid) as

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\[ \text{It should be noted that the fluorescence spectra of 2 are independent of the concentration (10}^{-7} \text{ to 10}^{-3} \text{ M).} \]
discussed above. However, such behaviour can also translate some \( \pi-\pi \) intermolecular interactions which can further disturb the purity of the colour. We also noted a marked difference between solution and thin-film fluorescence spectra of 3 in the relative intensity of the emission bands, with the second fluorescence band at 422 nm being more intense compared to the first at 402 nm (Fig. 2C). This might be ascribed to self-absorption phenomena due to the overlap of the 0–0 transition emission band and the absorption band which lead to a relative decrease of the intensity of the 0–0 transition.\(^\text{46} \) However, the shift observed for 3 is without any comparison with that described in the case of the previously described 2,6-diphenylindenofluorene for which the thin-film emission is detected at ca. 500–530 nm clearly signing a stacking of the molecules in the solid state due to strong intermolecular \( \pi-\pi \) interactions.\(^\text{28} \) Thus, the absence of strong \( \pi-\pi \) interactions in the case of 3 is hence due to the presence of the two spiro-(r-Bu-fluorene) units on each side of the indenofluorene core and highlights the efficiency of the present molecular design to avoid any intermolecular \( \pi-\pi \) interaction.\(^\text{12} \)

Interestingly, the fluorescence spectrum of 2 in solution is almost identical to that in the solid state (Fig. 2B) with only a slight red shift of around 6 nm. Thus, the intramolecular "aryl–fluorene–aryl" dimer formed in the excited state seems to lead to a stable deep blue fluorescence emission even in the solid state. This feature shows that no intermolecular \( \pi-\pi \)-interactions occur in the thin-film between the molecules of 2, due to the steric protection induced by the geometry of the molecule. The molecular design of 2 allows hence gathering in a single molecule, a wide HOMO/LUMO gap (ca. 3.4 eV, Table 1) and its associated properties (high stability toward oxidation for example) and a deep blue emission in the solid state. Finally, the quantum yield of 1–3 has been evaluated. 1 possesses a high quantum yield, ca. 75% relative to quinine sulfate, as already observed for similar compounds with the "aryl–fluorene–aryl" frameworks.\(^\text{24} \) 3 possesses a higher quantum yield, ca. 90% than that of 1. This result is in accordance with the fact that the indenofluorene molecule possesses a higher quantum yield than that of the fluorene or spirofluorene molecule.\(^\text{26} \) 2 presents again a drastically different behaviour compared to its analogues 1 and 3 as its quantum yield is of only 35%, fully consistent with the fact that the fluorescence of 2 arises from intramolecular excimers.\(^\text{32,46,47} \) However, despite being smaller than those of 1 and 3, the quantum yield of 2 remains relatively high for an excimer-based emission. Thus, in the case of 2, due to the rigidity of the backbone, the "aryl–fluorene–aryl" chromophores are located closely, in a favourable conformation, and require only a very slight spatial reorganization to form excimers. This close vicinity between the two "aryl–fluorene–aryl" chromophores probably leads to a decrease of any non-radiative decay processes and hence keeps the quantum yield at a reasonable value (ca. 35%), hence promising for further OLED applications.

To conclude on the optical properties of these molecular systems, one can say that their fluorences are drastically different and are mainly driven: in 1, by the "aryl/fluorene/aryl" moieties, in 2 by the excited dimer constituted of two face-to-face "aryl/fluorene/aryl" moieties and in 3 by the "aryl/indenofluorene/aryl" moiety. In addition, we noted for 2 very appealing properties for a fluorophore in which the fluorescence arises from excimer emission. Indeed, its quantum yield is in an acceptable range and its thin-film fluorescence spectrum is almost unchanged compared to its solution spectrum.

An important feature in OLED technology and particularly for blue light is related to the stability of the colour as a function of the temperature.\(^\text{46} \) Indeed, it is known that an OLED device can reach a temperature of ca. 86 °C and the stability of the emitted colour upon heating in air is hence of key importance.\(^\text{49} \)

In order to determine the stability of the colour, spin-coated films of DSF–IFs 1–3 on quartz substrates were exposed to thermal stress conditions under ambient atmosphere (Fig. 3). Gradual heating of a spin-coated film of 1, in air, from room temperature to 200 °C (1 h for each stage and finally one day at 200 °C), leads to a red-shift of around 6 nm and a broader spectrum. However, even after 24 h at 200 °C in the presence of air, no sign of low-energy emission band usually called Green Emission Band (GEB) was observed as it is usually the case for fluorene derivatives.\(^\text{48,50} \) For example, Lahti and co-workers have recently investigated the behaviour of fluorescent spectra as a function of the temperature for \( E,E-(2,7\text{-bis}(3,4,5\text{-trimethoxyphenylethyl}-9\text{'-diethylfluorene})) \), a structurally related compound to 1. The authors have notably reported the existence of an important GEB (\( \lambda = 540 \text{ nm} \)) after heating a thin film at 200 °C in the presence of air.\(^\text{51} \) This GEB emission has been assigned to the presence of fluorenone moieties at the fluorene bridge due to the oxidation process. However, in the case of 1 possessing spiro aryl bridges (and not alkyl bridges), no oxidation degradation processes leading to a GEB is observed. The emission colour of 1 appears then to be highly stable upon heating in the presence of air. In the case of 3, gradual heating of a spin-coated film, in air, from room temperature to 160 °C does not lead to any modification of the shape of the fluorescent spectra, highlighting the high colour stability. This feature has been assigned to the presence of two \( \text{r-butyl-fluorene} \) units on each side of the indenofluorenyl core leading to a steric protection and then avoiding any \( \pi-\pi \) intermolecular interaction.\(^\text{12} \)

The most striking feature here is again related to the compound 2. Thus, from room temperature to 160 °C, the fluorescence spectrum of 2 remains perfectly stable with a main emission centred at 463 nm (no additional GEB and no broadening of the spectrum compared to the solution spectrum). This feature is remarkable since the fluorescence emission of 2 results from excimer emission. Interestingly, the heating at 200 °C is accompanied by a slight modification of the fluorescence spectrum which now presents a maximum at 453 nm, surprisingly in accordance with that of its solution fluorescence spectrum.

Table 1 Electronic properties of compounds 1, 2 and 3

<table>
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<th>HOMO/( \text{eV} )</th>
<th>LUMO/( \text{eV} )</th>
<th>( \Delta E^{\text{opt}}/\text{eV} )</th>
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<td>-2.26</td>
<td>3.17</td>
<td>This work</td>
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\(^a\) Calculated from the onset oxidation potential \( E_{\text{onset}}^{a} \) (HOMO (eV) = \( -[E_{\text{onset}}^{a} \text{(vs. SCE)} + 4.4] \), based on an SCE energy level of 4.4 eV relative to the vacuum).\(^\text{6} \) Calculated from the HOMO energy level and the optical band gap.\(^\text{7} \) Optical band gap \( \Delta E^{\text{opt}} = h\nu - \Delta E^{\text{opt}}(\text{eV}) = 1237.5\lambda/\text{nm} \) has been estimated from the liquid UV-vis spectra in THF (\( \lambda \) is the edge of the absorption spectrum).
Similar changes of the solid-state fluorescence spectrum with temperature are described in the literature for butadiene derivatives, but in these examples, the monomer-like or excimer-like fluorescence comes from specific “intramolecular packing” which may be modified with the temperature. In a “herringbone” arrangement, minimal electronic coupling between molecules leads to a monomer-like emission, whereas in a “brickstone” arrangement, molecular packing leads to excimer-like fluorescence. In the case of 2, as previously observed for spiroterfluorene face-to-face fluorophores, heating of the films leads to a slight reorganization of the face-to-face “aryl-fluorene-aryl” moieties in the solid with less π–π interactions and hence a blue shift of the emission band.

These studies showed the high colour stability upon heating of these three DSF–IFs derivatives. As the fluorescent emission of 2 arises from intramolecular excimer, the high stability of the deep blue colour appears even remarkable and to the best of our knowledge without any precedent.

**Small Molecule Organic Light-Emitting Diodes (SMOLEDs) using 1, 2 and 3 as the emitting layer**

Several single-layer and double-layer OLEDs using 1, 2 and 3 as the emitting layer were fabricated and characterized (Fig. 4). The HOMO/LUMO levels of the three compounds have been then evaluated from electrochemical data (see figures in the ESI†) and optical data (Table 1).

We first decided to focus on single-layer devices to study the intrinsic properties of 1–3 when used as EML. Single-layer devices, ITO/PEDOT/EML/Ca (devices A), have been fabricated and characterized. In these devices, ITO/PEDOT is referred to as the anode, in which PEDOT (poly(3,4-ethylenedioxythiophene)) doped with PSS (poly(styrene sulfonate)) is the hole injecting layer and calcium is the cathode. These devices are considered as single-layer ones as PEDOT is not taken into account in the number of layers. The choice of the low work-function calcium cathode ( 2.9 eV) was evident in the light of the very high LUMO levels of all the compounds investigated (Table 1). The luminance of the device A using 1 as EML (50 nm) reaches around 52 Cd m⁻² with luminous and energetic efficiencies of 3.1 × 10⁻³ Cd A⁻¹ and 1.1 × 10⁻² Lm W⁻¹ respectively (see figures in the ESI†). The turn-on voltage of the device, defined as the voltage for a luminance of 1 Cd m⁻², is recorded around 7.2 V. For 3, the performance of the device A appears to be very similar to that discussed above for 1. Thus, the luminance of the device A using 3 as EML (40 nm) reaches around 50 Cd m⁻² (see figure in the ESI†) with luminous and energetic efficiencies of 1.6 × 10⁻² Cd A⁻¹ and 4 × 10⁻³ Lm W⁻¹ respectively (turn on voltage: 10.8 V). The device A with 2 as EML (45 nm) was finally investigated and a maximum luminance of 100 Cd m⁻² with luminous and energetic efficiencies of 3.7 × 10⁻² Cd A⁻¹ and 1.8 × 10⁻² Lm W⁻¹ were obtained. This device not only exhibits the best performances among the investigated single-layer devices but also presents the lowest turn on voltage (5 V).

Three main information may be extracted from this dataset: (i) the three single-layer devices A display modest performances translating the difficulty in injecting charges in the three compounds. This feature was assigned to the wide HOMO/LUMO gap of these materials coupled to their very high LUMO energy level (around −2.0/−2.2 eV, Table 1) and low HOMO energy level (around −5.3/−5.5 eV, Table 1), leading to a very high injection barrier for both electrons and holes. Even if the performances are modest, single-layer devices allow a perfect comparison of the intrinsic properties of each molecule and a clear definition of their potential for OLED applications. (ii) The molecules 1 and 3 which possess a similar HOMO/LUMO
gap present nearly identical performances. (iii) The best results were obtained with 2 as EML. Indeed, despite 2 possessing a similar HOMO/LUMO gap compared to that of 1, the emission of 2 results from an “aryl–fluorene–aryl” dimer and then the electrogenerated exciton is delocalized over this dimer, leading to a red-shifted emission band (vide infra) and hence to enhanced performances.

In order to increase the performances, a thin layer of NPB (N,N’-di(1-naphthyl)-N,N’-diphenyl-[1,1’-biphenyl]-4,4’-diamine) was deposited between the anode and the EML. The turn-on voltage of the ITO/PEDOT/NPB (40 nm)/Ca (device B) appears at ca. 6.8 V with a luminance magnified by around 5 times compared to device A, reaching ca. 230 Cd m⁻² with efficiencies also magnified, 6.2 × 10⁻² Cd A⁻¹ and 2 × 10⁻³ Lm W⁻¹ (Fig. 4, top). Similarly, the ITO/PEDOT/NPB (40 nm)/Ca device B with 2 as EML also leads to a strong enhancement of the performances (Fig. 4, bottom). Indeed, the maximum luminance reaches around 510 Cd m⁻² with a luminous efficiency of ca. 0.1 Cd A⁻¹. The turn-on voltage of the double-layer device B (5.8 V) is slightly higher than that of the single-layer device A due to the increase thickness of the organic layers. It is obvious that the performances of this excimer-based SMOLED are modest but: (i) the devices were not optimized, (ii) such intramolecular excimer based devices were without any precedent, (iii) the performances are magnified compared to a classical “aryl–fluorene–aryl” emitter such as 1, (iv) with a more accurate molecular design of the molecules (hole/electron transporting moieties) and/or of the devices (“band engineering” with hole/electron transporting/blocking layers, modification of the electrodes) the performances of the SMOLEDs will be surely and easily enhanced.

The last and crucial feature is finally related to the emission colour of these devices. The electro-optic properties of the devices A and B are presented below (Fig. 5). For 1, two important features were noted. First, the Electroluminescent (EL) spectra seem to be dependent on the nature of the device. Indeed, the device A presents an EL spectrum with a blue emission recorded at 432 nm (Fig. 5A, red line) red shifted by 18 nm compared to its thermally evaporated thin film, emission recorded at 448 nm (Fig. 5A, blue line). The second important feature is related to the shape of the EL spectra which appears to be not only larger than the PL spectrum but also with an additional GEB recorded at ca. 540 nm. The origin of such contribution at low energy has been extensively studied for the last decade as it remains one important problem in blue OLED technology. 48,50,54,55 It is widely agreed nowadays that the GEBs appearing in poly or oligo(dialkyl)fluorene arise from keto-defects in the form of fluorenone moieties.56 Similarly, GEBs have also been observed in structurally related poly-p-phenylenes such as poly(alkyldiindenofluorene) or ladder-type poly or oligo(alkylpentaphenylene).48,55–59 However, in the case of 1, the bridges are protected by a spirofluorene unit, not sensitive to oxidation degradation processes55,60 and hence the oxidation of the bridges leading to keto-defects appears to be impossible. In addition, we have also shown above that even annealing in the presence of oxygen, the fluorescent emission spectra do not possess any GEB (Fig. 3A). Thus, it is difficult to assign the GEB observed in the EL spectra of 1 to intermolecular π–π interactions leading to excimer emissions. We believe that this GEB may arise from interactions with the electrodes48,55 and/or exciplex emissions at the organic/organic interface,61,62 as previously described. Indeed, Ca-EML and/or PEDOT/PSS-EML interactions at the interface between the EML and the electrode have been previously suggested for similar extended π-conjugated backbones.54,55 The chromatic coordinates of the devices A and B with 1 as EML are (0.24; 0.24)/(0.25; 0.27) respectively and correspond to a light blue colour (see the CIE diagram in the ESIF).

In the case of 3, the device A presents an EL spectrum with a maximum recorded at ca. 452 nm and a long wavelength tail until 800 nm, whereas the device B presents a maximum red shifted by 38 nm (λ_max = 490 nm) and an even larger long wavelength tail with a shoulder at 700 nm, Fig. 5C. This large band (with its shoulder at 700 nm) found in device B can be tentatively assigned to exciplex emission due to the interactions at the organic/organic interface.61,62 This is confirmed by the fact that both devices possess EL spectra strongly red-shifted compared to the fluorescent spectrum of 3 (Fig. 5C, black line).

Finally for 2, the EL spectrum of the device A exhibits a nice structureless band with a maximum at 447 nm (Fig. 5B, red line), whereas the device B displays a maximum slightly red-shifted at 462 nm (Fig. 5B, blue line). These two EL spectra, opposite to those of 1 and 3, are in accordance with the thermally evaporated PL spectrum of 2 (Fig. 5B, black line) clearly signifying that the fluorescent emission of the device arises, without any doubt, from electrogenerated intramolecular excimer. A slight blue shift is nevertheless noted between the EL spectrum of device A (λ_max = 447 nm) and the PL spectrum of 2 (λ_max = 461 nm). Such

![Fig. 5](Image) Electroluminescent spectra using 1 (A), 2 (B) and 3 (C), as EML: device A (red), device B (blue) and thermally evaporated (~260 °C for 1 and 2) and spin-coated (3) thin-film PL spectra (black).
a feature has been also recently reported by Müllen and co-workers for dendrimer based blue OLEDs. This point needs to be nevertheless elucidated in further studies. In addition, no sign of intermolecular excimer leading to undesired GEB was detected, clearly signifying that the molecules of 2 in the thin-film are isolated enough to avoid any π–π intermolecular interaction. This site isolation is a key feature to ensure an efficient fluorescence and highlights the potential of the present molecular design for blue OLED applications. In addition, no band arising from keto-defects or from exciplex formation was detected as it was the case for I and 3. The chromatic coordinates of the devices A and B with 2 as EML are almost identical (0.19; 0.19) and (0.19; 0.18) respectively, and correspond to a deep blue colour (see the CIE diagram in the ESI†). In summary, the intramolecular excimer based-OLEDs not only display a stable deep blue colour with no trace of GEB due to “site isolation” of the molecules in the film but also present magnified performances compared to those of other devices with structurally related compounds (I and 3). Generating intramolecular excimers emission through the highly rigid (2,1-a)−DSF−IF molecular platform appears hence as an interesting new strategy to design electroluminescent devices. Indeed, this strategy allows obtaining a deep blue emission (PL and EL) centred at 460 nm with a short τ emissions compared to those of structurally related compounds. Furthermore, the intramolecular excimer based-OLEDs exhibit almost identical CIE coordinates, clearly signifying that the molecules of 2 in the film but also present magnified performances compared to those of structurally related compounds. Hence, this strategy allows obtaining a deep blue emission (PL and EL) centred at 460 nm with a short τ emission and 0.19; 0.18) exhibiting a luminance of 510 Cd m−2 with a luminous efficiency of ca. 0.1 Cd A−1. More sophisticated SMOLED designs, such as the use of carrier blocking layers, will easily allow the device performances to be significantly optimized. In addition, more experiments are nevertheless necessary in order to perfectly understand the physical and electronic properties of these intramolecular excimer based devices and to take full advantage of the favourable properties of this new molecular design. Regarding the importance of fluorene derivatives in organic electronics and the number of molecular platforms (pyrene, naphthalene, etc.) which have been used to induce intramolecular excimer emission, the present structure/property relationship study may open new avenues in the design of efficient blue OLEDs.

Conclusion

DSF−IF 2 has been successfully used as an emitting layer in a SMOLED leading to a stable deep blue emission arising from electrogenerated intramolecular excimers with better performances compared to those of structurally related compounds. This is to the best of our knowledge the first rational and comparative study describing the use of intramolecular excimers arising from organic molecules as blue light source in a SMOLED. Thus, due to the geometry of the rigid (2,1-a)−DSF−IF platform, the ‘aryl/fluorene/aryl’ dimer is preformed in the ground state favouring the facile and controllable formation of excimers in the excited state and leading to a deep blue fluorescent emission (λ ≈ 460 nm) through strong π–π intramolecular interactions. This emission wavelength is almost impossible to obtain with structurally related oligo ‘aryl–fluorene–aryl’ derivatives even with a large number of aryl substituents.

Experimental section

Synthesis

Commercially available reagents and solvents were used without further purification other than those detailed below. THF was distilled from sodium/benzophenone prior to use. Toluene was distilled from sodium prior to use. Light petroleum refers to the fraction with bp 40–60 °C. Reactions were stirred magnetically, unless otherwise indicated. Analytical thin layer chromatography was carried out using aluminium backed plates coated with Merck Kieselgel 60 GF254 and visualized under UV light (at 254 and 360 nm). Chromatography was carried out using silica 60A CC 40–63 μm (SDS). 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 CH, CH2 or Me groups, assigned from DEPT, are noted; all others are C. The residual signals for the NMR solvent CDCl3 are 7.26 ppm for the proton and 77.0 ppm for 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). The synthesis and the full characterization of 1, 2 and DSF(t-Bu)2−IF(Br)2 may be found in our previous works.

Spectroscopic studies

UV-visible spectra were recorded using a UV-Visible spectrophotometer Shimadzu UV-1605. The optical band gap was calculated from the absorption edge of the UV-vis absorption spectrum using the formula ΔEg = h/λ, λ being the absorption edge (in metres). With h = 6.6 × 10−34 J s (1 eV = 1.6 × 10−19 J) and c = 3.0 × 108 m s−1, this equation may be simplified as: ΔEg = (eV) = 1237.5/λ (in nm). Photoluminescence spectra were recorded with a PTI spectrofluorimeter (PTI-814 PDS, MD 5020, LPS 220B) using a xenon lamp. Quantum yields in solution (ϕsol) were calculated relative to quinine sulfate (ϕsol = 0.546 in H2SO4 1 N) using standard procedures. ϕsol was determined according to the following eqn (1):

\[
ϕ_{sol} = ϕ_{ref} \times 100 \times \left( \frac{T_s}{T_r} \times A_r \right) \left( \frac{n_s}{n_r} \right)^2
\]

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 (ns = 1.4072 for THF) and A is the absorbance (A < 0.1). IR spectra were recorded on a Bruker Vertex 70 using a diamond crystal MIRacle ATR (Pike).

Electrochemical studies

All 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 0.1 M AgNO₃ solution in CH₂CN. Ferrocene was added to the electrolyte solution at the end of a series of experiments. The ferrocene/ferrocenium (Fe/Fe⁺) couple served as the internal standard. The three electrode cell was connected to a PAR Model 273 potentiostat/galvanostat (PAR, EG&G, USA) monitored with the ECHEMA Software. Activated Al₂O₃ was added to the electrolytic solution to remove excess moisture. All potentials are referred to the SCE electrode that was calibrated at −0.405 V vs. Fe/Fe⁺ system.

OLED fabrication and testing

OLEDs were fabricated using the following procedure. Indium-tin oxide (ITO) substrates on glass from Merck underwent a solvent ultrasonic cleaning using acetone, ethanol and isopropanol followed by a 15 min UV-ozone treatment. A layer of poly(3,4-ethylene dioxythiophene) doped with poly(styrene sulfonate) (PEDOT/PSS from Aldrich) was then deposited onto ITO by spin-coating at 6000 rpm, from a 3 wt% water dispersion to form a 40 nm thick layer. PEDOT/PSS was subsequently annealed at 120 °C under vacuum for 40 minutes. This layer improves hole injection from the ITO to the HOMO level of the organic material and increases the performances and the lifetime of the device. Then a layer of 1, 2 or 3 or a NPB (N,N-di[1-naphthyl]-N,N′-diphenyl-[1,1′-biphenyl]-4,4′-diamine) layer followed by a layer of 1, 2 or 3 or 4 was thermally evaporated under vacuum (ca. 10⁻⁶ mbar). The layer thickness was monitored in situ during the evaporation via a piezoelectric quartz. Calcium cathodes (200 nm) were evaporated through a shadow mask. The OLEDs were then stored and characterized under inert atmosphere in a nitrogen glove box ([O₂] < 1 ppm). Current–voltage–luminance (I–V–L) curves were recorded using a Keithley 4200 SCS. Light emission was collected using a calibrated photodiode. Electroluminescence spectra were measured with a CCD spectrometer (Ocean Optics HR 2000).

References