Multiple photoluminescence from pyrene-fused hexaarylbenzenes with aggregation enhanced emission features

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Multiple Photoluminescence from Pyrene-Fused Hexaarylbenzenes with Aggregation Enhanced Emission Features


Abstract: Multiple photoluminescence, involved in monomer emission, excimer emission and charge transfer emission origin from new pyrene-fused hexaarylbenzenes (HAB) compounds were observed, which were designed and synthesized (in high yield) via the Diels-Alder reaction of bis(2-tert-butyl)pyren-6-yl)acetylene and tetraphenylenecyclopentadiene. Although the distinction of between two molecules arises only from the geometrical position of one of the pyrenes, the NMR spectra, the crystal packing and the physicochemical properties of these pyrene-based HAB hybrids are distinctly different both in their solution state and in aggregation-state. The X-ray diffraction analysis clearly indicated that the pyrene moieties in this system would form different crystal packing in crystal state that can induce a fantastic multiple photoluminescence phenomenon.

Introduction

The construction of intriguing molecular structures is both an art and a science and has attracted considerable interest in recent years. Highlights include reports on cage-carborane-based dyads with aryl-substitution,1 propeller-shaped tetraphenylenethene-based aggregation-induced emission (AIE) luminogens with bulky pendant moieties,2 chiral helicenes with fused planar and nonplanar systems,3 and π-expanded fullerene-based dyads.4 Such systems have played a crucial role in developing the field of organic optoelectronics. Among the many promising hybrid candidates, hexaarylbenezene (HAB) appears to be an ideal scaffold given that it possesses a rigid core and adjustable aryl units, allowing for the formation of unusual and fascinating structures. It has also proved possible to extend the range of applications possible for HAB derivatives by the introduction of suitable radial substitution reactions.5

Pyrenes as typical polymeric aromatic hydrocarbons (PAHs) possess unique optoelectronic properties and features that can be readily modified which have allowed them to be extensively explored as fluorophores, particularly in view of their additional favorable stability and high fluorescence efficiency.6 Generally, the availability of efficient monomeric emission in the range of 370–430 nm and the excimer emission at around 480 nm, combined with the charge-transfer (CT) emission possible by means of the introduction of donor/acceptor moieties at the pyrene has established their position/reputation in comparison with other polyaromatic fluorophores (see examples in Scheme 1).6 However, the mechanisms associated with multiple photoluminescence from a single molecule in pyrene chemistry remains relatively unexplored. The minimal mechanistic work to-date reflects the limiting harsh prerequisites, such as the combination of multidimensional intermolecular interactions required, not to mention the pre-determined spatial arrangements combined with appropriate donor/acceptor constituents.7 Thus, the design and synthesis of organic molecules with multiple photoluminescence remains a challenging topic.

Scheme 1 Diagrammatic sketch illustrating the typical structures giving rise to emissions in the pyrene system.

Building on the individual emission mechanisms mentioned above, a new strategy to construct high-performance single molecules with multiple photoluminescence was conceived. Our target was to attempt to integrate a pyrene fluorophore with a propeller-shaped HAB scaffold into a single molecule, which would exhibit multiple photoluminescence, due to its multidimensional geometrical parameters, conformations and intra/inter-molecular interactions. In particular, we wished to construct this “propeller” by replacing two of the “propeller blades”. Herein, the synthesis of two conformers of the pyrene based HAB derivatives anti-6 and syn-6 was achieved by employing this molecular design. Fortunately, via diffusion of methanol vapor into a chloroform solution of each compound, single crystals were cultivated, and the intriguing structures of anti-6 and syn-6 are disclosed by X-ray crystallography (Fig. 1).

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Results and Discussion

As depicted in Scheme 2, the bis(2-tert-butylpyren-6-ylylacetylene 5 was synthesized via the Sonogashira coupling reaction of 1-bromo-7-tert-butylpyrene 2 with 7-tert-butyl-1-ethynylpyrene 4 in 75% yield. 1,2-Bis(7-tert-butyl-pyren-1-yly)-3,4,5,6-(tetraphenyl)benzenes (anti-6 and syn-6) were prepared by the Diels-Alder reaction of bis(2-tert-butylpyren-6-ylylacetylene 5 with 2,3,4,5-tetr phenylcyclopenta-2,4-dienone in one step (see ESI† for full synthetic procedures and characterisation data). These two stereoisomers could be cleanly separated by normal silica gel chromatography and recrystallization, affording (see ESI† for full synthetic procedures and characterisation data). These two stereoisomers could be cleanly separated by normal silica gel chromatography and recrystallization, affording anti-6 as a pale green solid and syn-6 as a yellow solid in 53% and 22% yields, respectively. The \(^1\)H / \(^13\)C NMR spectra, X-ray crystallography and HRMS clearly indicate that they are different compounds. Both compounds exhibited excellent thermal stability with decomposition temperatures \(T_D\) of 411 °C for anti-6, and 403 °C for syn-6 under a nitrogen atmosphere at a heating rate of 10 °C min\(^{-1}\), as shown in Table 1 and Supporting Information Fig. S11. Moreover, their fundamental photophysical properties and mechanism were investigated, which demonstrates the validity for the construction of multiple photoluminescence molecules utilizing this strategy.

Analysis of the \(^1\)H NMR spectra revealed well-resolved proton signals which could clearly distinguish these two isomers (see Fig. 2). As expected, for anti-6, six doublets (due to ortho coupling) (12H) and two singlets (4H) were observed for the pyrene moieties, though there is an overlap between one singlet \((c', \delta_H = 8.02\) ppm\) and one doublet \((d, \delta_H = 8.01\) ppm\). In comparison, the proton signals of syn-6 exhibited a compact distribution, especially for the hydrogen atoms at the K-region of the pyrene moieties \((d, e, f, g, 8H, \delta_H = 7.58-7.70)\) ppm. In addition, the protons at the 6, 8-positions of pyrene completely overlap into one singlet \((b, 4H, \delta_H = 7.89)\) ppm, reflecting the shielding effects induced by the spatial overlap of the aromatic rings. In addition, the thermostability and photochemical stability were further studied by \(^1\)H NMR (Fig. S12), the interconversions of anti-6 and syn-6 was not observed after heating (lasted 30 min at 180 °C) and irradiation by UV light under air. The NMR spectrum indicated the isomer of anti-6 and syn-6 are remarkable conformational stability with of unique optical properties.

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In contrast to their similar absorption spectra, the stereoisomers anti-6 and syn-6 exhibited distinct emission bands in solution and in the solid state due to the different molecular conformations. Specifically, anti-6 emitted blue light with two well-separated emission bands at 394 nm and 447 nm in THF solution. Based on our knowledge, the shoulder peak at 394 nm can be assigned to the monomeric emission of the pyrene-like chromophore,7 and the maximum peak at 447 nm can be attributed to the intermolecular interactions in this propeller-shaped multiple conjugated system. Similarly, syn-6 also presents a blue emission band centred at 412 nm in THF solution (Fig. 3 right). Interestingly, new subsidiary peaks in both stereoisomers appeared in the solid state, which may indicate different molecular conformations are present leading to different emission behavior when compared with those observed in solution. As shown in Fig. 3, anti-6 displays a red-shifted maximum emission peak at 482 nm with a shoulder peak at 463 nm, but no obvious emission band at 394 nm. It is reasonable to speculate that the new peak at 482 nm arises from the excimer emission and the shoulder peak at 463 nm may originate from a significant intermolecular interaction. We speculate that there is an intramolecular charge transfer (IntramCT) state resulting in the red-shift from the primary peak at 447 nm in solution to the subsidiary peak at 463 nm in the solid state. Also, similar optical characteristics were investigated for syn-6. In this system, it is easy to assign the monomeric emission and excimer emission in solution and in the solid state from the multiple photoluminescence spectra. Taking anti-6 as an example, we assumed that the emission band at 447 nm in solution and 463 nm in the solid state can be attributed to the IntermCT emission. Given the polarity of the solution can exert slight effects on the CT emission, a solvatochromism experiment was carried out to provide further evidence. As shown in Fig. S13, anti-6 exhibited a slight red-shift from the less polar cyclohexane (442 nm) to the more polar DMF (453 nm), while there is no obvious regularity with the increase of the polarity for the low wavelength emission band at around 394 nm. It further demonstrated that there is no intramolecular charge transfer (ICT) but IntermCT for the anti-6 system. Unsurprisingly, there is barely a change in the solvatochromism for syn-6. This is consistent with our conjecture and with the emission properties both in solution and in the solid state. Thus, the spectroscopic data strongly support the presence of multiple photoluminescence from the monomeric emission, excimer emission and CT emission in this system.

Table 1 The photophysical properties of HAB derivatives anti-6 and syn-6.

<table>
<thead>
<tr>
<th>Py-based HAB transition</th>
<th>( \lambda_{\text{max}} ) (nm)</th>
<th>( \lambda_{\text{PL}} ) (nm)</th>
<th>( \phi_{\text{PL}} )</th>
<th>HOMO (eV)</th>
<th>LUMO (eV)</th>
<th>( E_{\text{g}} ) (eV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>S(_1) ( \rightarrow ) S(_0)</td>
<td>403</td>
<td>394, 447</td>
<td>463, 482</td>
<td>0.60 (0%)</td>
<td>0.95 (60%)</td>
<td>5.01</td>
</tr>
<tr>
<td>S(_2) ( \rightarrow ) S(_0)</td>
<td>353</td>
<td>472</td>
<td>394, 447</td>
<td>463, 482</td>
<td>0.60 (0%)</td>
<td>0.95 (60%)</td>
</tr>
<tr>
<td>S(_3) ( \rightarrow ) S(_0)</td>
<td>283</td>
<td>394</td>
<td>447</td>
<td>0.46 (0%)</td>
<td>0.80 (60%)</td>
<td>4.98</td>
</tr>
<tr>
<td>S(_2) ( \rightarrow ) S(_0)</td>
<td>404</td>
<td>412</td>
<td>420, 484</td>
<td>0.46 (0%)</td>
<td>0.80 (60%)</td>
<td>4.98</td>
</tr>
<tr>
<td>S(_2) ( \rightarrow ) S(_0)</td>
<td>334</td>
<td>468</td>
<td>412</td>
<td>420, 484</td>
<td>0.46 (0%)</td>
<td>0.80 (60%)</td>
</tr>
<tr>
<td>S(_2) ( \rightarrow ) S(_0)</td>
<td>272</td>
<td>472</td>
<td>394</td>
<td>447</td>
<td>0.46 (0%)</td>
<td>0.80 (60%)</td>
</tr>
</tbody>
</table>

[a] Measured in THF at room temperature. [b] Measured as powder. [c] the different fraction water in mixed solvents of THF and water are shown in bracket. [d] DFT (B3LYP/6-31G*) calculations were carried out with the use of structures optimized at the B3LYP/6-31G* level of theory. [e] Measured from the oxidation potential in THF solution by cyclic voltammetry. [f] LUMO = E\(_g\) + HOMO. [g] Estimated from the absorption edge of UV-Vis spectra.
As is well known, the propeller-shaped HAB scaffold has proved to be a natural attraction due to its rigid hub and adjustable peripheries. The peripheral groups of the HAB scaffold generally undergo interlocking to exhibit interesting structural variations with fascinating photophysical properties. Hence, we further attempted to interpret the mechanism on the basis of the X-ray crystallographic results. Close inspection reveals that the HAB derivatives anti-6 and syn-6 present distinctively different spatial arrangements to each other. As shown in Fig. 4, a shell-like intermolecular donor-acceptor system was observed for anti-6, which was achieved by utilizing the electron-rich pyrene molecules as the donor “shell”, and the electron-deficient phenyl moieties as the acceptor “pearl”. The fragments comprising pyrene-phenyl-pyrene were present with displaced face-to-face patterns linked via C-H-π interactions with distances ranging from 2.79 to 2.87 Å. The pyrene moieties are almost mutually perpendicular. The formation of a donor-acceptor system is prefered to the formation of a charge transfer system. More interestingly, an obvious InterCT was observed and demonstrated via experimental and crystallographic studies, and the maximum InterCT emission was observed at 447nm in THF solution. In contrast to anti-6, a clear pyrene…pyrene π-π stacking interaction with a distance of 3.15 Å was observed in the packing of syn-6. Obviously, the π-π stacking interaction plays a dominant role to affect the emission. The most direct evidence is that the excimer emission peak located at 484 nm appeared in the solid state, but was not observed in solution. These results are explicit and provide powerful evidence for the emission phenomenon and mechanism.

In general, HAB homologues exhibit a pronounced AIE property. To further investigate the AIE properties of the title compounds, the PL spectra of anti/syn-6 (5 × 10^{-7} M) were recorded in a water/THF mixture with differing water content (Fig. 5a). Taking anti-6 as an example, interesting AEE behavior was observed compared with previous reports.47 In particular, on increasing the water fraction in the water/THF mixture from 0% to 60%, the emission intensity was enhanced by about 2-fold (solid line). This is due to the restricted intramolecular rotation (RIR) and led to enhanced emission in this propeller-shaped system.48,49 However, when the content of water was increased up to 99%, the emission gradually decreased and a new peak was observed compared with the original emission in THF solution (dashed line), which resulted from the formation of excimer and solvent polarity dependent InterCT emission.10 It is worth mentioning that the emission profiles present a coherent transition from the organic solution to the solvent mixtures, aggregates and solid state. In the pure organic phase, two well-separated emission bands were observed as monomeric emission (394 nm) and CT emission (447 nm), and this type of emission band remains unaltered until the water percentages were increased up to 60%, inducing the formation of aggregation. Gradually, as the water fraction increased to more than 60%, multiple emission spectra gradually appeared, which are due to the monomeric emission (416 nm), CT emission (453 nm), and excimer formation (481), respectively. The PL spectra exhibited during the ultimate stage, suggested that excimer emission plays a dominant role and CT emission is secondary in the solid state. Similar AEE properties and transitions for the emission can also be observed for syn-6 (Fig. S14). In case of syn-6, there are two main emissions, including monomeric (412 nm) and excimer emission (484 nm). In particular, it is worth mentioning that the concentration-dependent luminescence observed for the two compounds are in excellent agreement with the AEE properties (Figures S15–16). Increasing emission intensity is displayed on increasing the concentration until the luminescence reaches its maximum intensity, following which the luminescence decreases in intensity when the concentration is further increased; similar red-shifts and multiple photoluminescence were also observed in these results. Furthermore, the fluorescence quantum yields were measured in solution and in the solid state, and the quantum yields (ΦE) are listed in Table 1. Specifically, a noticeable enhancement of quantum yield from 60% (in THF solution) to 95% (at 60% water fraction) was observed for anti-6, and a subdued quantum yield (45%) was obtained in the solid state, which is due to the formation of excimer. Similar optical behavior was observed in molecule syn-6.

To provide further evidence for the energy states, cyclic voltammetry (CV) and DFT calculations (B3LYP/6-31G*) were performed on anti-6 and syn-6 to interpret the electronic states combined with the photophysical properties. As shown in Fig. S18, both the compounds 6 displayed irreversible redox processes with distinct positive potentials in the vicinity of 1.4 V in solution. As shown in Figures S19–20, the electronic distributions of 6 in the ground state were further studied by frontier orbital analysis. As expected, both optimized geometries are consistent with their single crystal X-ray structures (Table S1). In order to derive explicit evidence, Time-dependent DFT (TD-DFT) calculations were carried out with the program.

Fig. 4 The packing structures of compounds anti-6 (a) and syn-6 (b) with principal intermolecular packing interactions (most hydrogen atoms are omitted for clarity). Distances are in Å.

Fig. 5 a) PL spectra of anti-6 in THF/H2O mixtures (5 × 10^{-7} M). b) Plots of I/I0 values versus the compositions of aqueous mixtures of anti-6.

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package Gaussian at the B3LYP/6-31G* level of theory. Simulated electronic spectra were produced based on the TD-DFT results, which are largely consistent with the experimental results. The TD-DFT calculations show that the pyrene moiety plays a dominant role on the orbital transition to the electronic excitation according to the respective contributions. In this system, the stronger S_{2}→S_{0} transition is described by a near 50:50 contribution from HOMO-1→LUMO+1 and HOMO-1→LUMO of the shoulder absorption band of anti-6, so these three key frontier molecular orbitals should be considered in any discussion of this system. Specifically, the shoulder absorption is due to the different combination of HOMO-1→LUMO+1 and HOMO-1→LUMO with a very high oscillator strength (f = 0.416, and f = 0.289), and the low oscillator strengths of 0.078 for S_{2}→S_{0} (n = 1.23), respectively. These theoretical results provided further insight into the intensity of the absorption spectra shown in Fig. 3. The detailed results of the TD-DFT calculations for the ground-state optimized geometries for anti-6 and syn-6 are presented in the Supporting Information, along with simulated absorption spectra.

Conclusions
In summary, by utilizing a propeller-shaped HAB scaffold, we have observed intriguing multiple photoluminescence. In particular, the thermally stable anti-6 and syn-6 pyrene-fused hexaarylbenezes synthesized herein exhibit unusual AEL characteristics. Multiple emission mechanisms, including monomer, excimer and CT emission were explored and deciphered in a single molecule. Single crystal X-ray structures of these two compounds 6 provided further insight into their differing properties. It is envisioned that the present work will widen our thinking in photophysics and stimulate work on differing properties. It is envisioned that the present work will widen our thinking in photophysics and stimulate work on differing properties. It is envisioned that the present work will widen our thinking in photophysics and stimulate work on differing properties.

Experimental Section
General procedures
Synthetic routes for the two compounds anti-6 and syn-6 are shown in Scheme 2. All reactions were carried out under a dry argon atmosphere. Solvents were Guaranteed reagent (GR) for cyclohexane, 1,4-dioxane, tetrahydrofuran (THF), dichloromethane (CH2Cl2), and dimethylformamide (DMF), and stored over molecular sieves. Other reagents were obtained commercially and used without further purification. Reactions were monitored using thin layer chromatography (TLC). Commercial TLC plates (Merck Co.) were developed and the purification. Reactions were monitored using thin layer chromatography (TLC). Commercial TLC plates (Merck Co.) were developed and the purification.

Synthesis of 7'-butyl-1-ethynylpyrene (4)
To a stirred solution of 7'-tert-butyl-1-[trimethylsilyl]ethynyl]pyrene (3) (1.00 g, 2.82 mmol) and THF (20 mL), was added MeOH (20 mL) and K2CO3 (585 mg, 4.23 mmol). The reaction was stirred at room temperature for 12 h. The mixture was quenched by addition of a large amount of water, extracted with dichloromethane (2 x 50 mL), washed with water and dried over anhydrous MgSO4 and concentrated. The residue was recrystallized in methanol to obtain 7'-tert-butyl-1-(trimethylsilyl)ethynylpyrene (4) (728 mg, 91%) as light green needles. M.p. 119–120°C; H NMR (300 MHz, CDCl3): δ = 1.59 (s, 9H, t-Bu), 3.61 (s, 1H, –C≡CH). 8.01 (d, J = 8.9 Hz, 1H, pyrene-δ), 8.05–8.17 (m, 4H, pyrene-γ), 8.24 (s, 1H, pyrene-γ), 8.26 (s, 1H, pyrene-γ), 8.56 ppm (d, J = 9.2 Hz, 1H, pyrene-γ). 13C NMR (100 MHz, CDCl3): δ = 29.11, 35.27, 82.41, 82.88, 116.24, 122.45, 122.99, 123.02, 124.17, 124.27, 125.17, 127.04, 128.62, 129.77, 129.82, 130.86, 131.03, 131.44, 132.34, 149.51 ppm; FAB-MS: m/z calc'd for C25H25Si: 354.18 [M]+; found 354.17 [M]+; anal. calcd (%) for C25H25Si: (354.56): C = 93.64, H = 6.36; found: C 93.57, H 6.43.

Synthesis of bis(2-tert-butyl-6-phenylpyrene) (5)
To a stirred solution of 7'-tert-butyl-1-bromopyrene 2 (500 mg, 1.48 mmol), Et3N (20 mL) and DMF (20 mL), was added 7'-tert-butyl-1-ethynylpyrene 4 (503 mg, 1.76 mmol) and PPh3 (62.2 mg, 0.237 mmol). The mixture was stirred at room temperature under argon. PdCl2(PPh3)2 (72.8 mg, 0.1 mmol) and Cu (31.1 mg, 0.163 mmol) were then added, and the mixture was heated to 100°C with stirring for 24 h. After it was cooled, the mixture was diluted into CH2Cl2 (200 mL) and washed successively with saturated aqueous NH4Cl, H2O and brine. The organics were dried (MgSO4), and the solvents were evaporated. In order to obtain pure product, the crude product was purified twice by column chromatography, eluting with hexane/CH2Cl2 (9:1). Commercial TLC plates (Merck Co.) were developed and the synthesis of bis(2-tert-butyl-6-phenylpyrene) (5) was confirmed by FAB-MS: m/z calc'd for C25H25Si: 354.18 [M]+; found 354.17 [M]+; anal. calcd (%) for C25H25Si: (354.56): C = 93.64, H = 6.43; found: C 93.57, H 6.43.

Synthesis of 1,2-bis(7'-tert-butyl-pyren-1-yl)-3,4,5,6-(tetraphenyl)benzenes (6)
Compound 5 (100 mg, 0.279 mmol) and 2,3,5,4-tetraphenylcylooctene-2,4-dieneone (161 mg, 0.419 mmol) were dissolved in PhOH (2.0 mL) under argon. The mixture was refluxed for 24 h and then the solvent was removed in vacuo. The residue was subjected to column chromatography on silica gel (CH2Cl2/ hexane 4:6) to afford anti-6 (88.5 mg, 53%) as a pale-green solid and syn-6 (36.1 mg, 22%) as a yellow solid. 1H NMR (400 MHz, CDCl3): δ = 1.54 (s, 18H, t-Bu), 8.05 ppm (d, J = 8.9 Hz, 2H, pyrene-γ), 8.23–8.29 (m, 6H, pyrene-γ), 8.36 (d, J = 8.0 Hz, 2H, pyrene-γ), 8.86 ppm (d, J = 9.2 Hz, 2H, pyrene-γ). 13C NMR (100 MHz, CDCl3): δ = 31.93, 35.28, 94.46, 117.92, 122.66, 122.95, 124.46, 125.44, 125.59, 125.76, 128.40, 128.70, 129.51, 131.01, 131.17, 131.76, 149.48 ppm; FAB-MS: m/z calc'd for C22H17Si: 538.27 [M]+; found: 538.25 [M]+; anal. calcd (%) for C22H17Si: (538.27): C = 93.97, H 6.43; found: C 93.97, H 6.42.
141.96, 148.09 ppm; FAB-HRMS (MALDI-TOF): 130.71, 130.81, 131.49, 131.65, 135.69, 140.12, 140.25, 140.71, 140.87, 126.60, 126.63, 126.78, 127.01, 128.88, 129.14, 129.92, 130.48, 130.58, 121.56, 122.61, 122.70, 123.77, 125.10, 125.23, 125.97, 126.26, 126.54, 141.96, 148.09 ppm; FAB-HRMS (MALDI-TOF): m/z calcd for C_{70}H_{54} 894.4426 [M]; found 894.4418 [M].

Acknowledgements

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Keywords: Multiple Photoluminescence • Hexaarylbenzenes • Pyrene Chemical • Anti/Syn Conformations • Aggregation Enhanced Emission


Entry for the Table of Contents (Please choose one layout)

Layout 1:

FULL PAPER

Chuan-Zeng Wang, Yuki Noda, Chong Wu, Xing Feng,* Perumal Venkatesan
Hang Cong, Mark R.J. Elsegood,
Thomas G. Warwick, Simon J. Teat,
Carl Redshaw, and Takehiko Yamato*

Page xx – Page xx
Multiple Photoluminescence from
Pyrene-Fused Hexaarylbenzenes with
Aggregation Enhanced Emission
Features