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A pillar[5]arene-based planar chiral charge-transfer dye with enhanced circularly polarized luminescence and multiple responsive chiroptical changes†

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The fabrication of circularly polarized luminescent (CPL) organic dyes based on macrocyclic architecture has become an importantly studied topic in recent years because it is of great importance to both chiral science and supramolecular chemistry, where pillar[*n*]arenes are emerging as a promising class of planar chiral macrocyclic hosts for CPL. We herein synthesized an unusual planar chiral charge-transfer dye (P5BB) by covalent coupling of triarylborane (Ar₃B) as an electron acceptor to parent pillar[5]arene as an electron donor. The intramolecular charge transfer (ICT) nature of P5BB not only caused a thermally responsive emission but also boosted the luminescence dissymmetry factor (*g*_{lum}). Interestingly, the specific binding of fluoride ions changed the photophysical properties of P5BB, including absorption, fluorescence, circular dichroism (CD), and CPL, which could be exploited as an optical probe for multi-channel detection of fluoride ions. Furthermore, the chiroptical changes were observed upon addition of 1,4-dibromobutane as an achiral guest.

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Introduction

Chirality, as one of the most significant phenomena, is ubiquitous in life and the environment, and determines the daily physiological activities and metabolism of life.¹ Chiral science unarguably promotes the development of life, medicine and materials science.² Chiroptical functional materials with circularly polarized luminescence (CPL) have drawn great attention in the last decade,³ not only for the understanding of the inherent principles of chirality but also owing to their wide potential applications in chiral sensing,⁴ photoelectric devices,⁵ 3D displays,⁶ asymmetric catalysis⁷ and so forth. Generally, CPL is generated due to the molecules or supramolecular aggregates having both chiral features and luminescent properties.⁸ Therefore, it has become an effective strategy to achieve CPL activity by connecting luminophores with chiral fragments (*e.g.*, binaphthyls, helicenes, and [2.2]paracyclophane).⁹ Recently, CPL-active

systems based on planar chiral analogues have attracted increasing attention,¹⁰ because their inherent macrocyclic skeletons have more important potential in the field of chiral supramolecular chemistry (*e.g.*, chiral network gelation and chiral host–guest recognition).

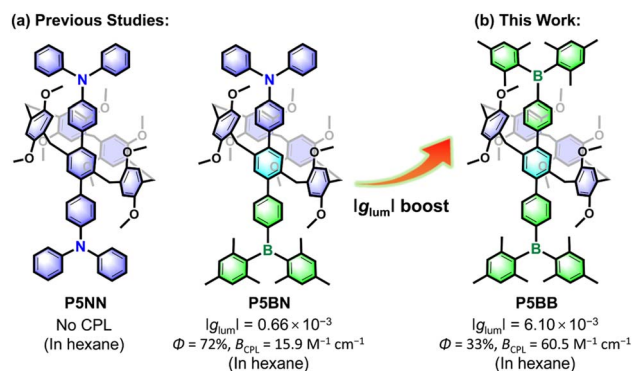
Pillararenes,¹¹ as an important type of macrocyclic arene, have attracted extensive studies and made significant contributions in host–guest recognition and self-assembly because of their unique structure and easy synthesis.¹² Pillararenes possess planar chirality, which comes from the different orientations of 1,4-alkoxyphenyl units.¹³ Although the enantiomers (*pS* and *pR*) are easy to interconvert due to the dynamic rotations of phenyl units, stable chiral configurations can still be achieved by reasonable molecular functionalization of the parent pillararenes. For example, introducing 10 cyclohexylmethyl groups at both rims of pillar[5]arene could prevent the rotations of benzene rings.¹⁴ Stoddart *et al.* also developed an effective strategy to obtain separable enantiomers by introducing bulky π -conjugated units at the A1/A2 positions of pillar[5]arene.¹⁵ Recently, pillararenes have been used to prepare CPL-active molecules by integrating with appropriate fluorophores.¹⁶ For example, Chen *et al.* reported two π -conjugated CPL-active systems (P5NN and P5BN) through axial functionalization of pillar[5]arene with sterically bulky triarylamine (Ar₃N) and triarylborane (Ar₃B) (Scheme 1); however, the *g*_{lum} values were only a 10^{−4} order because the luminescence largely depended on axial π -conjugated fluorophores and hence limited the transfer of chirality.¹⁷ In 2022, Ogoshi *et al.* reported a series of rim-differentiated C₅-symmetric pillar[5]arenes with

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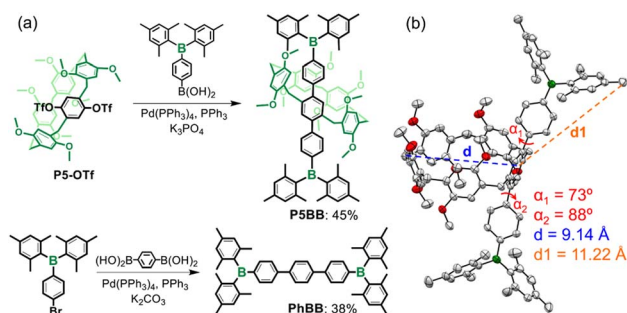
Scheme 1 Research foundation and design strategy of pillararene-based planar chiral charge transfer dye with amplified g_{lum} values.

improved g_{lum} values, but it was difficult to obtain proper fluorescence efficiency while improving the asymmetry factor.¹⁸ Therefore, the design and synthesis of CPL-active pillararenes at the molecular level with a good balance between g_{lum} factors and luminescence efficiency are highly anticipated.

Theoretically, the g_{lum} factor is simply approximated using $4|m|\cos\theta/|\mu|$, where m , μ and θ are the magnetic transition dipole moment, electric transition dipole moment and the angle between m and μ , respectively.¹⁹ If organic systems are appropriately designed to have weaker electric dipole transition and stronger magnetic dipole transition, it could lead to CPL-active materials with high g_{lum} . In fact, a large number of chiral organic molecules show relatively low g_{lum} values because of their electric dipole-allowed but magnetic dipole-forbidden transitions.²⁰ In contrast, because charge transfer (CT) systems possess relatively small $|\mu|$ and large $|m|$, the larger g_{lum} values obtained from the CT state are hypothesized.²¹ As a typical electron acceptor, Ar_3B has extensive application prospects in organic optoelectronic materials and stimulus-responsive materials.²² Notably, Ar_3B are enabled to show distinctive CT emission once they are electronically coupled with electron donors.²³ Pillararenes possess electron rich macrocyclic structures and can be used as a kind of electron donor. As one of our continuous pursuits of functionalized pillar[5]arenes,²⁴ we herein propose a facile methodology to amplify g_{lum} values through functionalizing the pillar[5]arene parent to construct planar chiral CT dye (**P5BB**) with Ar_3B . Sterically bulky Ar_3B fluorophores not only allow the enantiomeric resolution but also promote intramolecular charge transfer (ICT) from pillar[5]arene to Ar_3B . Based on the inherent host-guest nature of pillar[5]arene and the stimulus-responses of Ar_3B , the chiroptical response behaviors of this chiral system were further studied. The details are presented herein.

Results and discussion

The key synthetic process of **P5BB** is shown in Scheme 2a and the ESI.† The core planar chiral block **P5-OTf** was directly obtained *via* the previous report,²⁵ and then Pd-catalyzed Suzuki coupling by the reaction of **P5-OTf** with 2.0 equiv. Mes_2B -containing phenylboronic acid led to the formation of **P5BB** in 45% yield. Similarly, **PhBB** was also obtained by standard Suzuki coupling of (4-



Scheme 2 (a) Synthetic approach of **P5BB** and **PhBB**. (b) X-ray crystal structure of **P5BB** in elemental colors (C: grey, O: red, B: green; ellipsoid probability = 50%). All the hydrogen atoms and solvent molecules are omitted for clarity.

bromophenyl)dimesitylborane with 1,4-phenylenediboric acid in 38% yield. The chemical structures of **P5BB** and **PhBB** were fully characterized by 1H , ^{13}C , and ^{11}B NMR and high-resolution mass spectrometry (HRMS). A single crystal of *rac*-**P5BB** for X-ray diffraction analysis was collected by slowly evaporating the solution of acetone/MeOH ($v/v = 1:1$). In the crystalline form, a highly twisted π -conjugated skeleton was observed, as confirmed by dihedral angles (α_1 and α_2) that are measured to be 73° and 88° , respectively (Scheme 2b). Molecular size measurement revealed that the Ar_3B -substituents ($d_1 = 11.22 \text{ \AA}$) are greater than the cavity diameter of pillar[5]arene ($d = 9.14 \text{ \AA}$), so the racemization of enantiomers is sufficient to be inhibited through Ar_3B -substituents. The equimolar enantiomers of *pS*-**P5BB** and *pR*-**P5BB** are packed in a unit cell *via* C-H \cdots O, C-H \cdots π and C-H \cdots C interactions (Fig. S8†).

The photophysical properties of **P5BB** and **PhBB** were investigated in THF solution and the solid state (Fig. 1 and Table 1). **PhBB** shows a strong absorption band of π - π^* transition at 345 nm. However, **P5BB** exhibits two absorption peaks at 306 nm and 328 nm, corresponding to the local π - π^* transitions in the pillar[5]arene motif and in the axial conjugated skeleton, respectively (Fig. S9†).²²ⁱ In comparison to **PhBB**, **P5BB** showed significantly red-shifted emission in both the solid state and

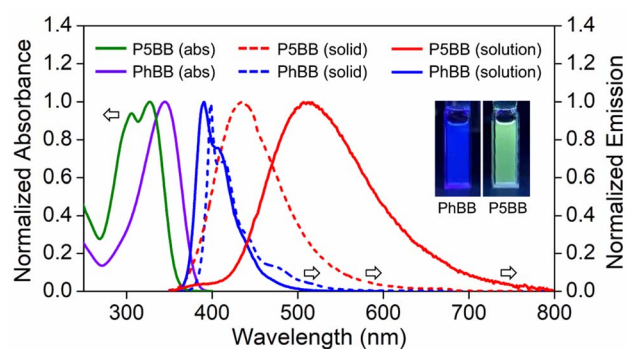


Fig. 1 UV-vis absorption and emission spectra of *rac*-**P5BB** ($\lambda_{ex} = 306 \text{ nm}$) and **PhBB** ($\lambda_{ex} = 345 \text{ nm}$): solid line, in THF ($c = 1.0 \times 10^{-5} \text{ mol L}^{-1}$); dashed line and solid state. Inset: photographs showing the emission colors of *rac*-**P5BB** and **PhBB** in THF under 365 nm UV irradiation.



Table 1 Photophysical, electrochemical and computational data of **P5NN**, **P5BN**, **P5BB** and **PhBB**

| | λ_{abs}^a (nm) | λ_{em}^a (nm) | Φ_L^b (%) | Φ_S^c (%) | τ_{ave}^d (ns) | E_{HOMO}^e (eV) | E_{LUMO}^e (eV) | $E_{\text{gap(DFT)}}^f$ (eV) | E_{ox}^g (V) | E_{red}^g (V) |
|--------------------------|-------------------------------|------------------------------|----------------|----------------|----------------------------|--------------------------|--------------------------|------------------------------|-----------------------|------------------------|
| P5NN ^h | 306 | 393 | 61 | 19 | 1.2 | −4.88 | −0.64 | 4.24 | +0.52 | — |
| P5BN ^h | 303, 330 | 493 | 99 | 57 | 6.4 | −4.89 | −1.67 | 3.22 | +0.59 | −2.49 |
| P5BB | 306, 328 | 510 | 12 | 16 | 20.5 | −4.91 | −1.77 | 3.14 | +0.61 | −2.46 |
| PhBB | 345 | 390 | 99 | 70 | 1.8 | −5.83 | −1.94 | 3.89 | — | — |

^a Measured in THF (1.0×10^{-5} mol L^{−1}) at room temperature. ^b Fluorescence quantum yield (Φ_L) measured in THF. ^c Φ_S measured in the solid state.

^d Average fluorescence lifetime in THF at room temperature. ^e HOMO and LUMO energy levels obtained using DFT calculations (B3LYP, 6-31G(d,p)). ^f $E_{\text{gap(DFT)}} = E_{\text{LUMO}} - E_{\text{HOMO}}$ (B3LYP, 6-31G(d,p)). ^g The first half-wave potentials of oxidation and reduction processes. ^h See ref. 17.

solution, which was ascribed to the ICT between the pillar[5]arene donor and Ar₃B acceptor. The ICT nature could be verified by a visible solvatochromic emission in various polar solvents (Fig. S11†). Owing to the temperature dependence of the equilibrium between the local excited (LE) state and the ICT excited state,²⁶ we explored the thermally responsive emission of **P5BB** in 2-methyltetrahydrofuran as a low melting point solvent (Fig. S12†). At low temperature (150 K), **P5BB** showed a significantly dual emission band with a main ICT emission at 552 nm ($\tau = 33.9$ ns) slightly overlapped with a minor LE emission at 405 nm ($\tau = 2.3$ ns). With temperature increasing from 150 to 330 K, the main emission band of **P5BB** experienced an apparent hypsochromic shift, and the emission color change from yellow to blue was monitored using CIE coordinates. There is a good linear relationship of the maximum emission wavelength *versus* temperature with a correlation coefficient of 0.976. It is noteworthy that the above thermochromic response is completely reversible. The good accuracy and reversibility suggested that the system is an ideal candidate for high-performance fluorescent thermometers.

In order to further understand the correlation between the molecular structures and the photophysical properties of **P5BB** and **PhBB**, the electronic structure calculations were performed using DFT (B3LYP, 6-31G(d,p)) and TD-DFT (B3LYP, 6-31G(d)). TD-DFT computations revealed that the absorption of **PhBB** is mainly attributed to the π - π^* transition to the S₁ state (HOMO \rightarrow LUMO, $f = 1.2994$) (Fig. S14 and Table S5†). In **P5BB**, the HOMO is fully located on the electron-donor pillar[5]arene backbone; however, the LUMO is delocalized over the B-conjugated π -extension (Fig. S15†). By means of TD-DFT calculations, it was found that the first three CT transitions are the results of vertical excitations from the pillar[5]arene-localized HOMO, HOMO−1 and HOMO−2, to the LUMO (Fig. S16†). The higher excited states (S₄ and S₅) are excitations to the LUMO+1 level from the HOMO and HOMO−1. Cyclic voltammetry (CV) of **P5BB** showed a reversible reduction potential at −2.46 V (*vs.* Fc+/Fc, in THF), representing the reduction of the electron-deficient Ar₃B segments (Fig. S17†). Three reversible oxidation curves with the first oxidation potentials at +0.61 V for **P5BB** (*vs.* Fc+/Fc, in CH₂Cl₂) were identified, corresponding to the oxidation of the pillar[5]arene skeleton. The electrochemical gap (3.07 eV) is almost consistent with the HOMO–LUMO gap (3.14 eV) *via* the DFT calculations. As expected, the first oxidation potential of **P5BB** is slightly higher than that of compounds with the N donor (**P5NN**,

+0.52 V; **P5BN**, +0.59 V), which is completely consistent with the slightly lower HOMO energy level (**P5BB**, −4.91 eV; **P5NN**, −4.88 eV; **P5BN**, −4.89 V).

The definite enantiomeric configuration inspired us to prepare their optically pure forms for studying the chiroptical properties. Two separated peaks with a 1 : 1 area were observed by the initial injection of *rac*-**P5BB** into a chiral HPLC with a Daicel Chiralpak IB N-5 column (hexane/2-propanol = 96/4, v/v). After each fraction was well isolated with an enantiomeric excess (>99% ee, Fig. S18†), the CD spectra of the enantiomers exhibited mirror-image relationships with strong CD absorption peaks in various organic solvents (Fig. 2a). The CD signals belonging to pillar[5]arene cores are clearly observed at 310 nm,²⁷ and the absorption dissymmetry factor $|g_{\text{abs}}|$ was calculated to be 1.53×10^{-3} in hexane. On the basis of a comparison of the experimental CD absorption signals with the results of previous reports,²⁷ the *pS* configurations correspond to the first peak and the second peak correspond to *pR* configurations in HPLC traces. In the CPL spectra, the enantiomers exhibited almost mirror-imaged signals in various solvents (Fig. 2b). The $|g_{\text{lum}}|$ reached 10^{-3} in solution as well as in the solid state (Table 2), which is significantly higher than that of **P5NN** and **P5BN**. The CPL spectra of **P5BB** gradually redshifted with the increase in solvent polarity, and the positions of CPL signals were almost consistent with that of fluorescence. In fact, the enantiomers of **P5NN** and **P5BN** did not show apparent CPL signals in various solvents (except for hexane), suggesting that the chiral transfer did not occur

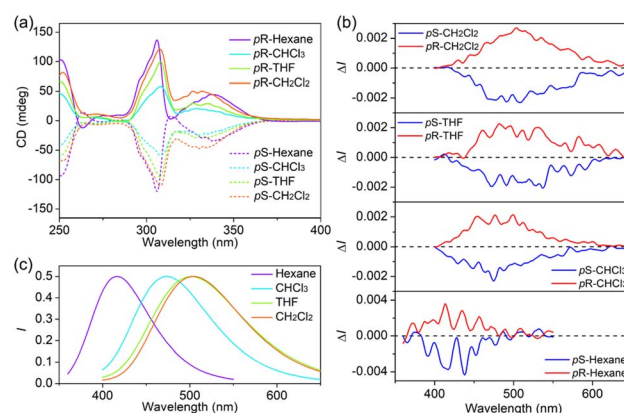


Fig. 2 (a) CD, (b) CPL and (c) fluorescence spectra of enantiomers of **P5BB** in different solvents ($c = 5.0 \times 10^{-5}$ mol L^{−1}, $\lambda_{\text{ex}} = 306$ nm).



Table 2 The chiroptical property data of **P5NN**, **P5BN** and **P5BB**

| | | λ_{abs} (nm) | ϵ^a ($\text{M}^{-1} \text{cm}^{-1}$) | λ_{em} (nm) | Φ^b (%) | g_{lum}^c (10^{-3}) | B_{CPL}^d ($\text{M}^{-1} \text{cm}^{-1}$) |
|------------------------------|---------------------------------|-----------------------------|---|----------------------------|--------------|----------------------------------|---|
| pS- P5NN ^e | Hexane | 306 | 7.31×10^4 (306) | 391 | 52 | — | — |
| | Solid | — | — | 398 | 19 | +0.22 | — |
| pS- P5BN ^e | Hexane | 303, 330 | 6.71×10^4 (303) | 403 | 72 | +0.66 | 15.9 |
| | Solid | — | — | 455 | 57 | +0.88 | — |
| pS- P5BB | Hexane | 305, 329 | 6.01×10^4 (305) | 422 | 33 | −6.10 | 60.5 |
| | CHCl ₃ | 308, 328 | 5.98×10^4 (308) | 473 | 23 | −3.64 | 25.1 |
| | THF | 306, 328 | 6.08×10^4 (306) | 510 | 12 | −3.34 | 12.2 |
| | CH ₂ Cl ₂ | 306, 327 | 6.41×10^4 (306) | 512 | 12 | −4.62 | 17.8 |
| | Solid | — | — | 435 | 16 | −5.13 | — |
| | CHCl ₃ | 308, 328 | 6.00×10^4 (308) | 460 | 18 | −4.16 | 22.4 |
| pS- P5BB + G | THF | 295 | 3.35×10^4 (295) | 365 | 56 | −5.08 | 47.6 |

^a Molar absorption coefficient (ϵ) at a given wavelength. ^b Fluorescence quantum yield (Φ). ^c Luminescence dissymmetry factor (g_{lum}). ^d The CPL brightness (B_{CPL}). ^e See ref. 17.

effectively in this case. These phenomena indicated that the ICT character of **P5BB** not only amplifies the g_{lum} factors but also adjusts the color of CPL *via* the selection of different solvents. With all the necessary photophysical and chiral optical data in hand, the CPL brightness (B_{CPL} : defined as $B_{\text{CPL}} = \epsilon \times \Phi \times |g_{\text{lum}}|/2$) in solutions was further calculated to evaluate the overall performance of the CPL dyes. The B_{CPL} of pS-**P5BB** was calculated to be $60.5 \text{ M}^{-1} \text{cm}^{-1}$ in hexane, which is significantly higher than that of pS-**P5BN** ($15.9 \text{ M}^{-1} \text{cm}^{-1}$), indicating that **P5BB** possesses excellent chiroptical properties for future CPL applications.

In order to further evaluate the chiroptical changes induced through host-guest chemistry, we selected the neutral small molecule 1,4-dibromobutane (**G**) as the representative guest to bind **P5BB** by 1 : 1 complexation.²⁸ As shown in Fig. S19,† upon the addition of excess 1,4-dibromobutane to **P5BB** in CDCl₃ solution, the proton signals of H₁₋₃ and H₅₋₁₁ on **P5BB** were shifted downfield, and meanwhile, all signal peaks on the guest were found to shift upfield in ¹H NMR spectra. Additionally, the proton peaks of the guest were substantially reduced after complexation due to inclusion-induced shielding effects. The association constant (K_a) was determined to be 187.8 M^{-1} using the nonlinear data fitting of ¹H NMR titrations (Fig. S20 and S21†). This implies that host-guest recognition between **P5BB** and 1,4-dibromobutane has occurred. Notably, the addition of a guest results in dramatic enhancement of CD signals at 310 nm and 328 nm in CHCl₃, while the absorption does not change significantly, implying that the host-guest complexation decreases the configuration rotation of the pillar[5]arene skeleton of **P5BB** (Fig. 3). The addition of excessive 1,4-dibromobutane led to a decrease in fluorescence intensity, and meanwhile, the CPL signals showed a mild enhancement, indicating the host-guest recognition further affected the excited state chiral conformation of **P5BB**.

Considering the Lewis acidity of Ar₃B, another interesting thing is the response behavior of **P5BB** toward small Lewis bases.²⁹ Herein, the optical responses of four common halogen anions (including F[−], Cl[−], Br[−] and I[−]) as Lewis bases to **P5BB** were primarily investigated in THF solution. In the UV-vis absorption spectra (Fig. 4a), the absorption bands at around 306 nm and

328 nm of **P5BB** gradually decrease with the addition of F[−] anions, which is due to the fact that the conjugation of the Ar₃B moiety was broken by the formation of tetra-coordinated boron complexing with F[−] anions (Scheme S1 and Fig. S24†). The limit of detection (LOD) was further calculated to be 43.3 nM based on the $3\sigma/S$ values (Fig. S22†). Remarkably, other halogen ions (Cl[−], Br[−] and I[−]) could not cause any significant changes in absorption spectra, which was likely ascribed to the smaller steric size of the F[−] anion. Furthermore, the strong affinity between fluoride and boron also played a crucial role. As displayed in Fig. 4b, as the concentration of F[−] increased, the emission peak at 510 nm of **P5BB** slowly declined, while the emission peak at 365 nm rapidly enhanced. Quite evidently, the emission spectrum exhibited a blue shift ($\sim 145 \text{ nm}$) with the emission color changing from green to purple, and the fluorescence intensity increased dramatically with Φ up to 56%. These phenomena indicated that the binding of F[−] anions with the Ar₃B unit prevented the ICT process in the excited state, leading to the enhancement of LE-state emission.²⁹ Moreover, in the presence of an excess amount of F[−] anions, the CD signals of the enantiomers at 328 nm disappeared, while the signals at 310 nm were effectively enhanced. Even after complexation with F[−] anions, significant CPL signals with a blue-shift were also detected. The $|g_{\text{lum}}|$ values were calculated to reach $5.08 \times$

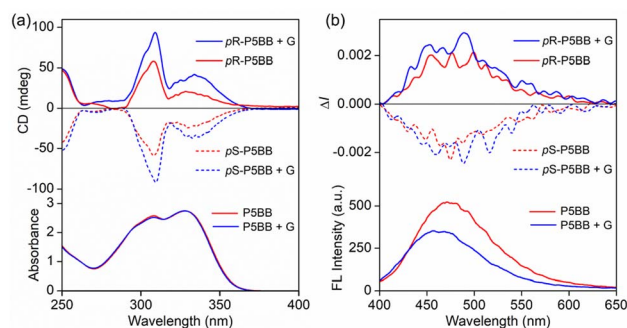


Fig. 3 (a) CD and UV-vis absorption spectra and (b) CPL and fluorescence spectral responses of enantiomers for **P5BB** in CHCl₃ ($c = 5.0 \times 10^{-5} \text{ mol L}^{-1}$, $\lambda_{\text{ex}} = 306 \text{ nm}$) upon addition of 50 equivalents of 1,4-dibromobutane **G** as a neutral guest.



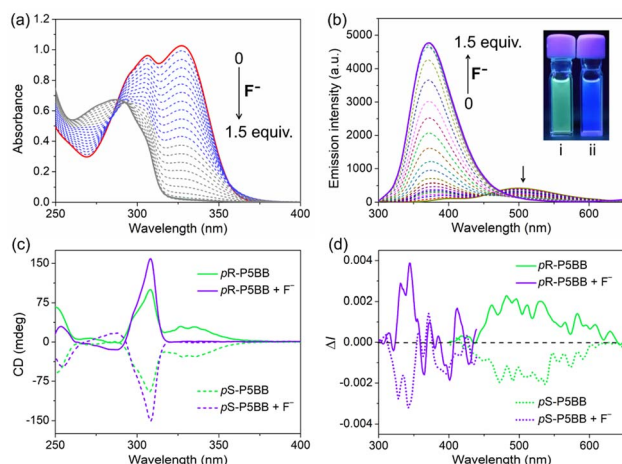


Fig. 4 (a) UV-vis absorption and (b) emission spectra with the addition of different concentrations of F^- anions to *rac*-P5BB in THF solution ($c = 2.0 \times 10^{-5} \text{ mol L}^{-1}$, $\lambda_{\text{ex}} = 306 \text{ nm}$). Inset: photographs of solutions before (i) and after (ii) the addition of F^- (1.5 equiv.) under 365 nm UV irradiation. (c) CD and (d) CPL spectral responses of enantiomers for P5BB ($c = 5.0 \times 10^{-5} \text{ mol L}^{-1}$, $\lambda_{\text{ex}} = 306 \text{ nm}$) in the presence of 3.0 equiv. of F^- anions.

10^{-3} with a B_{CPL} of 47.6 in this case and were much higher than those of chiral pillararene derivatives reported in ref. 16b (Scheme S2[†]). These investigations confirmed that the P5BB system could realize UV-vis absorption/fluorescence/CD/CPL quadruple-mode sensing of F^- anions.

Conclusions

In summary, we have developed an effective strategy to design and synthesize a π -conjugated planar chiral CT dye P5BB through integration of pillar[5]arene with organoborane. The ICT character in P5BB not only amplified the g_{lum} values but also could tune the CPL color by the selection of various solvents with different polarities. Approximately 10-fold enhancements in g_{lum} were observed from P5BB to P5BN in hexane, which was ascribed to the fact that the photo-responsive unit itself contains planar chiral pillar[5]arene in P5BB. Inclusion of the achiral guest 1,4-dibromobutane directly enhanced chiral optical signals, including CD and CPL. Furthermore, the coordination of F^- anions with boron leads to remarkable changes in absorption, fluorescence, CD and CPL signals of the P5BB system. Consequently, the P5BB platform could specifically detect F^- anions with high sensitivity and favorable selectivity. The current design strategy to fabricate a CPL-active ICT dye with the amplification of the g_{lum} factor is expected to promote the development of future planar chiral pillararene materials. We envision that this work will catalyze the future application of pillararene-based CPL-active systems in chiroptical sensing, chiral supramolecular chemistry and CPL-based photoelectric devices.

Data availability

All the data supporting this article have been included in the main text and the ESI.[†]

Author contributions

J.-F. C. initiated and coordinated the study. J.-F. C., Q.-X. G. and T.-B. W. designed all experiments, analyzed the data, and wrote the manuscript. L. L. carried out the computational work. P. C. and T.-B. W. supervised and administrated the project. All authors approved the final version.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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