

## PAPER

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2, 4859Asymmetric D–A–D' Scaffold inducing distinct  
mechanochromic luminescence†Huifang Yang,<sup>a</sup> Zhiyuan Fu,<sup>b</sup> Sai Wang,<sup>a</sup> Haichao Liu,<sup>c</sup> Jianwei Zhou,<sup>a</sup>  
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Mechanochromic luminescence (MCL) materials have a high requirement in high-tech applications such as pressure sensors and data storage. Nevertheless, the diversity of these emitters is rather restricted owing to the lack of clear molecular structure design guidelines, and it is imperative to develop creative multi-stimulus responsive compounds with polychromatic emission under different types of mechanical forces. In order to overcome difficulties of a single molecule exhibiting obvious MCL responses towards mechanical forces, in this study, a powerful approach of distinct multi-stimulus responsive MCL materials is to integrate the distinguished ability of donor units (D'), i.e. stronger phenothiazine (PTZ) and weaker bis(4-(*tert*-butyl)phenyl)amine (tDPA) compared with 9H-carbazole, directly into the same donor–acceptor (D–A) scaffold of 2-(4-(carbazol-9-yl)phenyl)-isoindoline-1,3-dione (CzPL) and hence obtaining **CzPL-PTZ** and **CzPL-tDPA**. Distinct MCL were primarily found for the parent **CzPL-PTZ** and **CzPL-tDPA** crystals under tensile force, and anisotropic and isotropic pressures. Interestingly, both crystals show an abnormal blue-shifted emission under crushing force. When under anisotropic pressure, the **CzPL-PTZ** ground sample further displays blue-shifted emission, but the **CzPL-tDPA** ground sample displays red-shifted emission. In contrast, both parent crystals show a similarly red-shifted and weakened emission under increasing isotropic pressure. Furthermore, these emissions exhibit a linear relationship between wavelength and pressure. More importantly, crystal **CzPL-PTZ** can completely return to the initial state once decompressed due to the reversible 3D short-range intermolecular interaction network; in contrast, the crystal **CzPL-tDPA** cannot fully recover to its original state after removing pressure due to the lattice vibration under a high pressure. The interesting and distinct MCL behaviors of **CzPL-PTZ** and **CzPL-tDPA** indicate that the D–A–D' molecular design should provide pragmatic insights towards MCL materials.

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## 1. Introduction

Mechanochromic luminescence (MCL) materials with polychromatic switches can exhibit distinctly variational emission when subjected to external mechanical stimuli (e.g., smashing, grinding, stretching, or hydrostatic pressure).<sup>1–6</sup> Such materials have been widely studied due to their potential applications in

optical data storage, pressure sensors, anti-counterfeiting inks, and optoelectronic devices.<sup>7–24</sup> In general, MCL can be realized by chemical structure change (e.g., chemical bond destruction and recombination)<sup>25,26</sup> or physical structure change (e.g., phase transition<sup>27–29</sup> and conformational transition<sup>30–32</sup>). The emission properties of MCL materials largely depend on molecular stacking modes, which are closely related to the molecular conformation and noncovalent interactions.<sup>33–35</sup> External forces can change the molecular stacking mode, subsequently managing the emission colors and intensities.<sup>10–17,36,37</sup>

Most MCL materials show reversible conversion luminescence under only one mechanical force.<sup>38–42</sup> However, materials with distinct MCL responses towards different mechanical stimuli are very rare. For instance, Wang certificated that boron-diketonate crystals exhibited unique emissions under different mechanical stimuli, with a blue-shift after crushing and a red-shift under grinding and compressing.<sup>43</sup> Yamaguchi reported that a tetra-thiazolylthiophene fluorophore displayed distinctly emissive responses to mechanical grinding and hydrostatic pressures.<sup>44</sup>

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Liu found that a donor–acceptor charge-transfer cocrystal CT-R also exhibited differently emissive responses to anisotropic grinding and isotropic compressing.<sup>23</sup> From the viewpoint of applications, materials with different MCL responses seem to be more attractive. Nevertheless, the diversity of these emitters was still rather restricted owing to the lack of clear molecular structure design guidelines, and it is imperative to develop creative multi-stimulus responsive MCL materials with polychromatic emission under different types of mechanical forces.

Herein, two novel molecules, namely 2-(4-(9H-carbazole-9-yl)-phenyl)-5-(10H-phenothiazine-10-yl)isoindoline-1,3-dione (**CzPL-PTZ**) and 2-(4-(9H-carbazole-9-yl) phenyl)-5-(bis(4-*tert*-butyl)phenyl)-amino)isoindoline-1,3-dione (**CzPL-tDPA**), were designed and synthesized by integrating distinguishing ability of electron-donating units (D') into the same donor–acceptor (D–A) scaffold of 2-(4-(carbazol-9-yl)phenyl)-isoindoline-1,3-dione (CzPL). In both molecules, phthalimide is chosen as an electron-withdrawing acceptor (A) owing to its excellent stability, mechanical toughness, and rigid structure.<sup>45</sup> 9H-carbazole (Cz) is employed as an end-capping electron-donating group due to its planar  $\pi$  conjugation and good charge transport ability. Pseudo-planar phenothiazine (PTZ) and propeller-like bis(4-*tert*-butyl)phenylamine (tDPA) are used as the D' units make the molecular packing can be adjusted, eventually resulting in exhibit distinct MCL of **CzPL-PTZ** and **CzPL-tDPA**. Overall, the sturdy intramolecular hydrogen bonds in phthalimide should rigidify the molecular conformation and suppress the nonradiative decay.<sup>46–48</sup> Moreover, the abundant hydrogen bonds between the CzPL unit and distinguishing stereochemical structures are available for the resulting molecules to form different three-dimensional (3D) networks, and different multi-stimuli responsive MCL materials can be obtained by adjusting the electron donating ability of D' units. Our study provides a new design strategy of asymmetric D–A–D' to develop novel MCL materials.

## 2. Results and discussion

### 2.1. Synthesis and characterization

Fig. 1 shows the chemical structures of **CzPL-PTZ** and **CzPL-tDPA**. Their synthetic routes are shown in Scheme S1 ESI†. They were synthesized by the Buchwald–Hartwig C–N cross-coupling reaction between CzPL-Br and PTZ or tDPA in moderate yields. Subsequently, they were purified *via* vacuum sublimation. Their molecular structures were fully characterized by <sup>1</sup>H, <sup>13</sup>C NMR spectroscopy and TOF-MS (see ESI†). Thermogravimetric analysis (TGA) measurements indicate that both **CzPL-PTZ** and **CzPL-tDPA** have good thermal stability with high decomposition temperatures

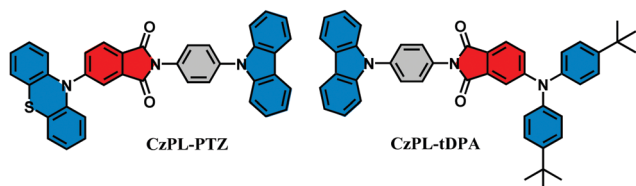


Fig. 1 Chemical structures of **CzPL-PTZ** and **CzPL-tDPA**.

( $T_d$ , corresponding to 5% weight loss) of 418 °C and 428 °C, respectively (Fig. S1 in ESI†).

### 2.2. Crystal structures

Since molecular packing plays a key role in MCL, the crystal structures of **CzPL-PTZ** and **CzPL-tDPA** were deeply investigated, which might give important insights into the MCL mechanism. The yellow–green (533 nm) and blue–green (498 nm) photoluminescent (PL) peaks are observed for the **CzPL-PTZ** and **CzPL-tDPA** parent crystals made *via* vacuum sublimation (Fig. S2 in ESI†), respectively. Both parent crystals possess quantum yield ( $\Phi_f$ ) values of 39% and 66% correspond to fluorescence lifetimes ( $\tau$ ) of 12.07 ns and 16.56 ns for **CzPL-PTZ** and **CzPL-tDPA** (Fig. S3 and Table S1 in ESI†). Fig. 2 shows the crystal structures and intermolecular interactions of the **CzPL-PTZ** and **CzPL-tDPA** crystals measured by single crystal X-ray diffraction (XRD) experiments. Table S2 (ESI†) summarizes the detailed data of these crystal structures. The results indicate that **CzPL-PTZ** adopts a quasi-axial conformation and a regular head-to-tail arrangement throughout some C–H $\cdots\pi$ , C–H $\cdots$ N, C–H $\cdots$ O, C–H $\cdots$ S, and weak  $\pi\cdots\pi$  interactions (Fig. 2a). In contrast, **CzPL-tDPA** features a head-to-tail arrangement but Z-shaped packing between adjacent molecules (Fig. 2b). These multiple intermolecular interactions can restrict the intramolecular motions and vibrations for both crystals, which is beneficial to the increase the  $\Phi_f$  values. Moreover, the twisty molecular configurations with large dihedral angles between phenylene and phthalimide/carbazolyl (Fig. S4 in ESI†) and the  $\pi\cdots\pi$  distance over 3.518 Å between anti-parallel molecules make both crystals present a weak  $\pi\cdots\pi$  stacking interaction, in which the aggregation caused quenching (ACQ) can be efficiently suppressed. Their densities of 1.386 g cm<sup>−3</sup> and 1.224 g cm<sup>−3</sup> for **CzPL-PTZ** and **CzPL-tDPA** crystals indicate that the triclinic **CzPL-PTZ** crystal has much tighter stacking than the monoclinic **CzPL-tDPA** crystal. Interestingly, both crystals exhibit multiple intermolecular interactions observed from the *a*, *b*, *c* axes, and form a huge 3D short-range intermolecular interaction network (Fig. S5 in ESI†). Such short-range 3D networks with weak intermolecular interactions should be easy to be deformed and get distinct emissive responses under different mechanical forces (*e.g.*, crushing, grinding, or hydrostatic pressure).

### 2.3. Photophysical properties

The UV-Visible and PL spectra of **CzPL-PTZ** and **CzPL-tDPA** were obtained in their solutions with different solvents and neat films. Two typical absorption bands were observed in Fig. S6 (ESI†), which contain a high-lying band of absorption in the range of 300–350 nm and a low-lying band of absorption in the range of 350–450 nm. The low-lying bands of absorption are ascribed to the intramolecular charge transfer (ICT) absorption.<sup>49–51</sup> Fig. S6 (ESI†) also shows the PL spectra of **CzPL-PTZ** and **CzPL-tDPA** in different solvents ( $1.0 \times 10^{-5}$  M). Dual emissions were observed for **CzPL-PTZ** in toluene (Tol), chloroform (CF), and tetrahydrofuran (THF), which were assigned to the dual conformers of PTZ (Fig. S6a in ESI†).<sup>52</sup> The excitation spectra of **CzPL-PTZ** at 480 nm and 560 nm in the CF solvent ( $1.0 \times 10^{-5}$  M) with different spectra indicate that the two emission bands at



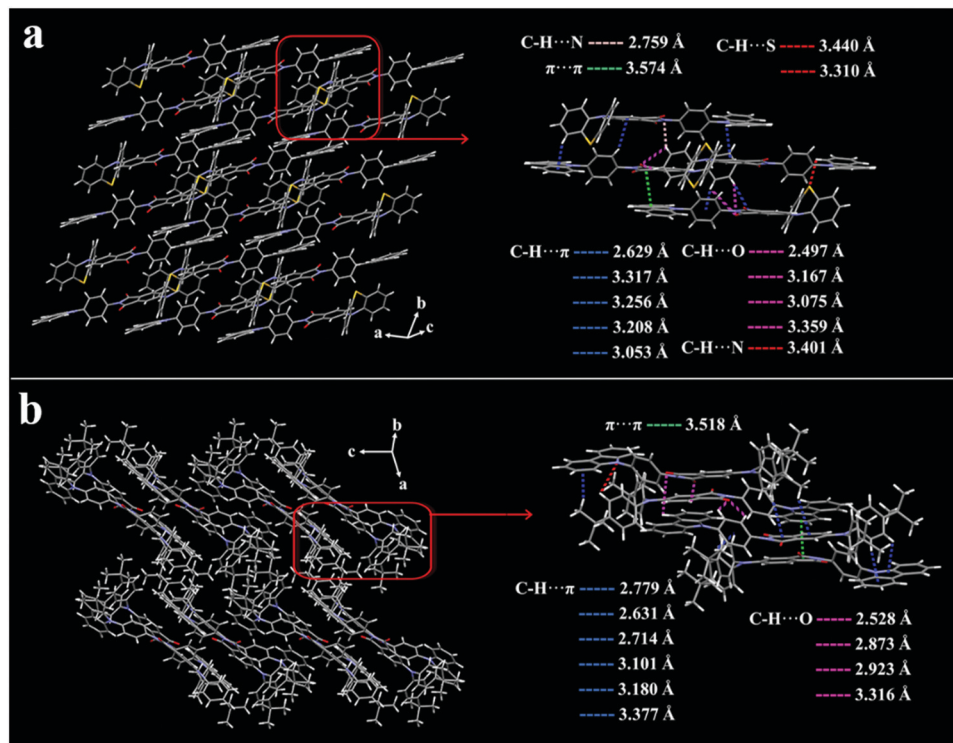


Fig. 2 Crystal structures and intermolecular interactions of the (a) **CzPL-PTZ** and (b) **CzPL-tDPA** crystals.

480 nm and 560 nm come from different excited states, *i.e.* locally excited (LE) and CT states (Fig. S7 in ESI<sup>†</sup>).<sup>52–58</sup> In contrast, there is only single emission peak for **CzPL-tDPA** in different solutions (Fig. S6b in ESI<sup>†</sup>). Both molecules show obvious solvent dependence in different polar solvents owing to their ICT effects. In addition, different PL spectra are observed for both molecules under two solid states. Due to the special quasi-axial conformation of the PTZ unit (bending to one side, small steric hindrance), the linear **CzPL-PTZ** molecules are easy to pile up in the long-range order and exhibit relatively strong intermolecular interactions. In contrast, **CzPL-tDPA** can easily form a large twisty type between **tDPA** and **CzPL**, which results in a relatively weak intermolecular interaction owing to the large steric hindrance of *tert*-butyl. Therefore, **CzPL-PTZ** displayed more red-shifted luminescence than **CzPL-tDPA** in crystal and film states (Fig. S2 and S8 in ESI<sup>†</sup>).<sup>56</sup>

#### 2.4. Theoretical calculations

Density functional theory (DFT) discloses that the asymmetric D–A–D' strategy has an impact on the lowest-lying electronic transition. The electronic structures of **CzPL-PTZ** and **CzPL-tDPA** are optimized at the ground state ( $S_0$ ) in the gas state using Gaussian 09W program at the B3LYP/6-31G (d)/LanL2DZ level.<sup>59,60</sup> As shown in Fig. S9 (ESI<sup>†</sup>), the lowest unoccupied molecular orbitals (LUMOs) of **CzPL-PTZ** and **CzPL-tDPA** are mainly located on electron deficient phthalimide moieties, and the highest occupied molecular orbital (HOMO) of **CzPL-PTZ** is dominantly distributed over the electron donor phenothiazine moieties. Since carbazole has stronger electron donating ability than bis (4-(*tert*-butyl) phenyl) amine, the HOMO of **CzPL-tDPA** is mainly distributed in the

carbazole unit.<sup>61</sup> Clearly, electron-donating D' units with distinguishing ability led to distinction in the molecular level and hence influence the MCL properties in the aggregation state. In addition, the calculated HOMO and LUMO energy levels are  $-5.28/-2.63$  eV and  $-5.18/-2.08$  eV for **CzPL-PTZ** and **CzPL-tDPA**, respectively. The more effective spatial separation of HOMO and LUMO in the **CzPL-PTZ** compound compared with **CzPL-tDPA** should demonstrate stronger ICT nature.<sup>62</sup>

#### 2.5. Electrochemical characterization

Electrochemical properties of **CzPL-PTZ** and **CzPL-tDPA** were investigated by cyclic voltammetry (CV) adopting ferrocene as a reference standard in anhydrous  $\text{CH}_3\text{CN}$  solutions. The recorded CV curves are shown in Fig. S10 (ESI<sup>†</sup>) and their electrochemical data are listed in Table S4 (see ESI<sup>†</sup>); **CzPL-PTZ** and **CzPL-tDPA** present reversible oxidation process in CV curves. According to the onset voltages of the oxidation curves, their HOMO levels are determined to be  $-5.16$  and  $-5.41$  eV for **CzPL-PTZ** and **CzPL-tDPA**, respectively. The lowest unoccupied molecular orbital (LUMO) energy level of **CzPL-PTZ** and **CzPL-tDPA** can be calculated using  $E_{\text{HOMO}}$  and optical band gap ( $E_g$ ), which are  $-2.04$  and  $-2.48$  eV, respectively.<sup>63,64</sup> In addition, the calculated HOMO and LUMO energy levels for **CzPL-PTZ** and **CzPL-tDPA** are  $-5.28/-2.63$  eV and  $-5.18/-1.18$  eV, respectively, which are reasonable deviations from the experimental data.

#### 2.6. Mechanochromic luminescence properties

The MCL properties of the **CzPL-PTZ** and **CzPL-tDPA** crystals were studied under different forces, such as tensile force,



anisotropic, and isotropic pressures. Fig. 3a and b show the PL spectra of **CzPL-PTZ** and **CzPL-tDPA** in different solid states, respectively. Here, the crushed and ground samples were obtained under crushing using a stainless-steel spoon and ground for 30 min in a mortar. In addition, the PL spectra of **CzPL-PTZ** and **CzPL-tDPA** ground samples were recorded at room temperature after different retention time periods. It can be seen from the Fig. S12 (ESI†) that there is almost no change in their PL spectra, which indicates that the two ground samples are very stable at room temperature. The fumed samples were obtained under fuming dichloromethane (DCM) vapor for 10 min. It was found that both crushed crystals exhibited scarce blue-shifted emissions in comparison with their corresponding parent crystals. It implies that a proper splitting of the parent crystals can greatly disturb the ordered molecular arrangement on the surface, thus resulting in a depressed conjugated system and blue-shifted emission.<sup>42</sup> More surprisingly, the  $\Phi_f$  values of the crushed crystals of **CzPL-PTZ** (48%) and **CzPL-tDPA** (91%) greatly improved in comparison with those of their corresponding parent crystals of **CzPL-PTZ** (39%) and **CzPL-tDPA** (66%) (Table S6 in ESI†).<sup>65</sup>

When the crushed crystals are further ground by a mortar, the resulting ground sample of **CzPL-tDPA** shows bathochromic-shift emission like most of the reported MCL materials.<sup>13,14,27–31</sup> However, the ground sample of **CzPL-PTZ** shows a rare hypochromic-shift emission.<sup>21,23,24,66</sup> A broad emission band peak at 501 nm is observed in the **CzPL-PTZ** ground sample, which is blue-shifted by 18 nm in comparison with that of the **CzPL-PTZ** crushed crystal (519 nm). Under further fuming the ground samples with the polar solvent vapor of DCM for 10 min, the resulting **CzPL-PTZ** fumed

sample exhibited a red-shifted emission peaked at 520 nm (Fig. 3a). In contrast, the **CzPL-tDPA** ground sample presents a significantly red-shifted emission peak at 547 nm in comparison with that of the **CzPL-tDPA** crushed crystal (491 nm). The emission color was partially recovered after fumigation with DCM (Fig. 3b). Therefore, the alternative appendage of another electron-donating unit of **PTZ** and **tDPA** in the D–A unit of **CzPL** can make their crystals exhibit distinct MCL response in the above four states. Furthermore, the film emission of the two compounds was compared with that of crystals and ground samples (Fig. S13, ESI†). The results show that the crystal and ground sample emission of the two compounds blue-shifted compared with that of the films. The conformation planarization and the closer molecular packing of **CzPL-PTZ** and **CzPL-tDPA** molecules in the films should be the origins for their red-shifted emission.

The powder X-ray diffraction (PXRD) diagrams of **CzPL-PTZ** and **CzPL-tDPA** in different states indicate that these various MCL responses are related to the change in the molecular packing structure, which are recorded in Figs S14a and b (ESI†), respectively. The results show that **CzPL-PTZ** and **CzPL-tDPA** form different molecular stacking architectures over the crushing-grinding-fuming treatment. These observations indicate that the strong and few diffraction peaks of **CzPL-PTZ** and **CzPL-tDPA** in the parent crystal state indicate that their crystals are polycrystalline. After crushing, the stacked polycrystals are split, the crystal size becomes smaller and the diffraction peaks increase. With further grinding, the PXRD pattern of the ground sample of **CzPL-PTZ** and **CzPL-tDPA** are composed of several blunt “peaks”, indicating that they are mainly amorphous in the ground state or their crystallites are extremely small in size.



Fig. 3 PL spectra of (a) **CzPL-PTZ** and (b) **CzPL-tDPA** in their parent crystals, crushed crystals, ground samples, and fumed samples; (c) the photographs of **CzPL-PTZ** and **CzPL-tDPA** during the crushing-grinding-fuming processes under UV light irradiation at 365 nm.





Similarly, the ground samples of **CzPL-PTZ** and **CzPL-tDPA** after fumigation in dichloromethane (DCM) vapor for 10 min exhibit almost same PXRD patterns as those of single crystal simulation, indicating that the molecular arrangement in fumigated samples return to the crystalline state. There is no doubt that the molecular arrangement of **CzPL-PTZ** and **CzPL-tDPA** changed during mechanical grinding. For the crystal **CzPL-PTZ**, grinding may cause the aggregated molecules to slip, disrupt the 3D short range intermolecular interaction network to ongoing single molecules, which will lead to the blue-shift emission.<sup>44</sup> The red-shift luminescence of **CzPL-tDPA** may be due to the different molecular arrangements such as  $\pi$ -stacked dimers have been formed after grinding.<sup>13,14,43</sup>

As mentioned above, the anisotropic pressure causes the **CzPL-PTZ** and **CzPL-tDPA** crystals to exhibit distinct MCL. To reveal the effects of isotropic pressure on emission, the PL spectra of both parent crystals under different hydrostatic pressures were further studied. As shown in Fig. 4, the **CzPL-PTZ** crystal exhibits a gradually red-shifted and weakened emission from 480 to 600 nm, with the hydrostatic pressure increasing from ambient pressure to 10.31 GPa. Furthermore, its absorption is also observed to maintain a red-shifted trend (Fig. S15 in ESI<sup>†</sup>). In contrast, when the hydrostatic pressure is returned to ambient pressure, its PL

spectrum turns back to the original PL profile (Fig. 4b). Identically, **CzPL-tDPA** shows similarly red-shifted emission and absorption with the increase in the hydrostatic pressure (Fig. 5 and Fig. S16 in ESI<sup>†</sup>). However, the PL spectrum of **CzPL-tDPA** just partly returns to the initial PL profile after the hydrostatic pressure is released to ambient pressure. More interestingly, whether it pressuring or decompressing the crystal, the emissive wavelength and energy have a direct good linear relationship to the hydrostatic pressure for both crystals (Fig. S17 and S18 in ESI<sup>†</sup>). This emissive uniqueness endows this type of MCL materials with potential application in pressure sensors.<sup>5</sup> Better emissive restorability of both crystals indicates that the molecular packing mode should not change substantially during a high-pressure process. Considering this point, the intense red-shift and weakened PL of both crystals may be attributed to the pressure-induced conformational planarization with increasing pressure, which leads to severe  $\pi$ - $\pi$  intermolecular interactions.<sup>67,68</sup>

According to the crystal structure analysis, there are weak  $\pi$ - $\pi$  interactions in both crystals. The  $\pi$ - $\pi$  distances of **CzPL-PTZ** and **CzPL-tDPA** are 3.574 Å and 3.518 Å, respectively, which are beyond the range of van der Waals interaction (Fig. 2b-d). The molecular distance between adjacent molecules should be shortened and  $\pi$ - $\pi$  interaction should be enhanced with increasing pressure.



Fig. 4 (a) PL spectra of crystal **CzPL-PTZ** under the pressures from 1 atm to 10.31 GPa, (b) PL spectra of crystal **CzPL-PTZ** during the decompression processes, (c) the fluorescence images of the **CzPL-PTZ** crystal under photo excitation at 350 nm and different pressures.

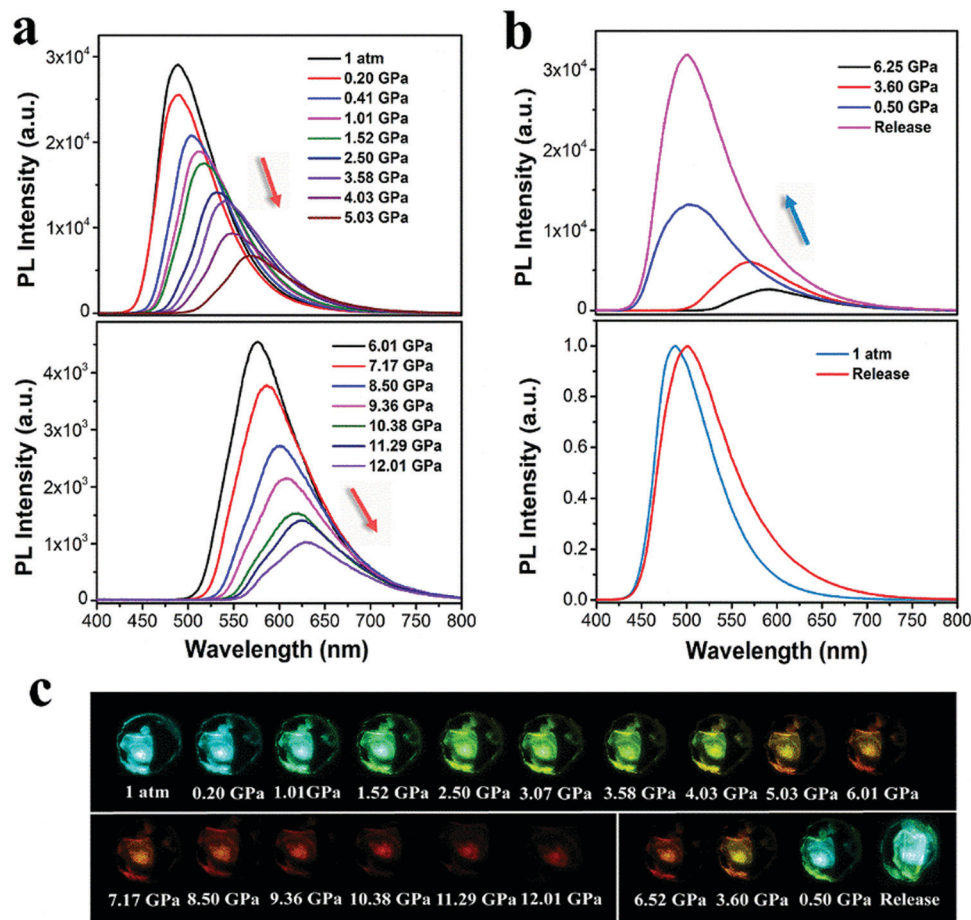


Fig. 5 (a) PL spectra of crystal **CzPL-tDPA** under the pressures from 1 atm to 12.01 GPa, (b) PL spectra of crystal **CzPL-tDPA** during the decompression processes, (c) the fluorescence images of the crystal **CzPL-tDPA** under photo excitation at 350 nm and different pressures.

As a result, red-shifted and weakened emissions are observed. The squashed **CzPL-PTZ** crystal can rebound to its original shape after decompressing. That is, its 3D intermolecular short-range interaction network undergoes a simple harmonic vibration within a certain pressure range during the compression-decompression process. Alternatively, the deformation of a 3D intermolecular short-range interaction network is reversible in this condition. Regarding the behavior of the **CzPL-tDPA** crystal under decompression, it might have to overcome some potential barrier to recover the 3D intermolecular short-range interaction network. Such an incomplete recovery is partially caused by the lattice vibration under high pressure.<sup>69</sup>

### 3. Conclusion

In summary, we have demonstrated a simple and effective molecular design strategy by integrating the distinguishing ability of electron-donating units (D') on the **CzPL** scaffold to obtain two novel MCL materials of **CzPL-PTZ** and **CzPL-tDPA**. The D' units of pseudo-planar PTZ and propeller-like tDPA play an important role in the MCL properties of **CzPL-PTZ** and **CzPL-tDPA** parent crystals. As a result, both crushed crystals show an abnormal blue-shifted emission under tensile force in comparison with

corresponding parent crystals; **CzPL-PTZ** crystals exhibit hypochromatic-shift, while **CzPL-tDPA** exhibit bathochromic shift when under anisotropic pressure. In contrast, both crystals show a red-shifted emission under increasing isotropic pressures. Significantly distinct MCL for both parent crystals under tensile force, anisotropic and isotropic pressures certify the effectiveness of the strategy D-A-D' scaffold. Interestingly, crystal **CzPL-PTZ** can completely return to the initial state after decompression, while crystal **CzPL-tDPA** cannot fully recover to its original state after removing pressure. More importantly, there is a linear relationship between emissive wavelength and pressure. The distinct and interesting MCL phenomena may have a potential application in pressure sensors and data storage.

### Conflicts of interest

There are no conflicts to declare.

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