

## RESEARCH ARTICLE

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Cite this: *Inorg. Chem. Front.*, 2023, **10**, 6909Luminescence detection of CH<sub>2</sub>Cl<sub>2</sub> by varying Cu...Cu interactions in a flexible porous coordination polymer†Wei-Jie Zhang,<sup>‡a</sup> Wan-Tao Chen,<sup>‡a</sup> Chen-Hui Li,<sup>a</sup> Wen-Zhu Sun,<sup>a</sup> Jia-Wen Ye,<sup>ID \*a</sup> Ling Chen,<sup>ID \*a</sup> Hai-Ping Wang<sup>\*a</sup> and Xiao-Ming Chen<sup>ID b</sup>

Owing to the weakness of the interaction between chlorohydrocarbon and the host framework, the development of reversible and vapochromic coordination polymer (CP)-based luminescence sensors for the detection of CH<sub>2</sub>Cl<sub>2</sub> with a fast response is still challenging. In this study, a flexible Cu(I)-CP, [Cu<sub>2</sub>l<sub>2</sub>(PPh<sub>3</sub>)<sub>2</sub>(PYZ)] (**CIPP**, PPh<sub>3</sub> = triphenylphosphine, PYZ = pyrazine) is reported. The intra-cluster Cu...Cu distance in **CIPP** is quite sensitive to the external stimuli, resulting in the corresponding luminescence color and intensity change. Based on this feature, the fast (11 s), distinguishing (wavelength shift of 45 nm), and reversible luminescence response of **CIPP** for CH<sub>2</sub>Cl<sub>2</sub> is realized. Crystallographic analysis suggests that the presence/removal of CH<sub>2</sub>Cl<sub>2</sub> can affect the Cu...Cu distance, which is the origin of the responsive luminescence. In addition, the multiple weak interactions between CH<sub>2</sub>Cl<sub>2</sub> and the framework afford the strong adsorption of CH<sub>2</sub>Cl<sub>2</sub> into **CIPP**, which can be maintained for at least 5 minutes when exposed to air, thereby ensuring accuracy in the sensing process.

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## Introduction

Volatile organic compounds (VOCs) refer to organic compounds with high vapor pressure and low boiling point under ambient conditions, therefore, they have high volatility.<sup>1,2</sup> Long-term exposures to VOCs possibly cause damage to human health, such as damage to the liver, kidneys, and nervous system; thus the detection of VOCs is of critical significance.<sup>3–5</sup> Emitting units in porous coordination polymers (CPs) or metal–organic frameworks (MOFs) can interact with VOCs, leading to changes in the intensity or the wavelength of the luminescence. For example, by generating C–H...O or Cl...H interactions between the host and guest molecules, the luminescence detection of tetrahydrofuran (THF), *N,N*-dimethylformamide (DMF), dimethyl sulfoxide (DMSO), and CH<sub>2</sub>Cl<sub>2</sub> can be realized.<sup>6–8</sup> In addition, the process of guest-adsorption/desorption in a porous CP is usually

accompanied by the structural transformation, which further exhibits changes in the conformation or the charge transfer of the luminescent chromophore.<sup>9,10</sup>

Among VOCs, CH<sub>2</sub>Cl<sub>2</sub> is widely used as a solvent or reagent not only in laboratories but also in a wide range of industrial applications, such as the production of paint removers, herbicides, and pesticides.<sup>11</sup> However, CH<sub>2</sub>Cl<sub>2</sub> may cause damage to the liver and nervous system, and even potential carcinogenic effects.<sup>12,13</sup> Several methods have been developed for the detection of CH<sub>2</sub>Cl<sub>2</sub>, such as gas chromatography (GC)-mass spectrometry (GC-MS, 5 ng L<sup>–1</sup>), gas chromatography-photo-ionization detectors (GC-PID, 1 ppm), and screen-printed electrodes (SPE, 17.3 μmol L<sup>–1</sup>).<sup>14–17</sup> However, they display some shortcomings, such as the use of expensive instruments, tedious sample pretreatment processes, and the inability for *in situ* analysis. Compared to the above traditional ways, luminescence detection for VOCs based on porous CPs has unique advantages, such as fast response, visualization, and simplicity.<sup>18–23</sup> However, in comparison with usual VOCs, CH<sub>2</sub>Cl<sub>2</sub> has difficulty forming strong interactions with the luminescent probes because of its low polarity and rather low boiling points. Therefore, reports about the luminescence response for CH<sub>2</sub>Cl<sub>2</sub> in porous CPs and MOFs are rare.<sup>8,24</sup> Moreover, because of the weak interaction between CH<sub>2</sub>Cl<sub>2</sub> and CPs, it is difficult for CH<sub>2</sub>Cl<sub>2</sub> to directly affect the luminescent center in CPs and cause the solvatochromism/vapochromism. Generally, the reported luminescent CP sensors for

<sup>a</sup>School of Biotechnology and Health Sciences, Wuyi University, Jiangmen, Guangdong 529000, PR China. E-mail: wyuchemyjw@126.com, wyuchemcling@126.com, wanghp@wuyi.edu.cn

<sup>b</sup>MOE Key Laboratory of Bioinorganic and Synthetic Chemistry, School of Chemistry, Sun Yat-Sen University, Guangzhou 510275, PR China

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‡These authors contributed equally to this study.

$\text{CH}_2\text{Cl}_2$  with fast response are based on the emission intensity change. It is rather difficult to design and prepare luminescent CPs with both chromatic and fast responses to  $\text{CH}_2\text{Cl}_2$ . Compared with the color-changeable luminescence detection, the intensity-depending method is more likely to cause errors due to test conditions and methods.<sup>25–27</sup>

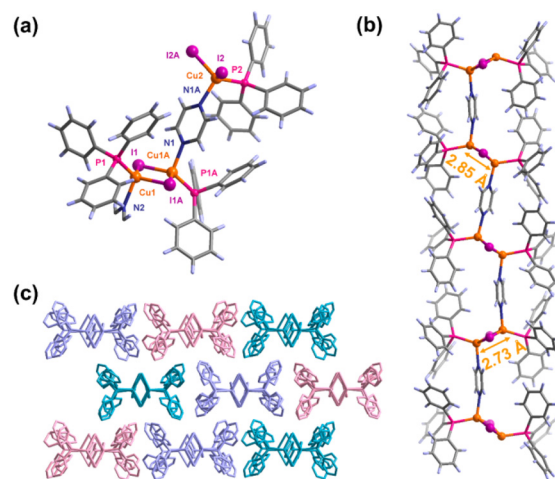
Cu(I)-CPs have attracted much attention because of their low toxicity, low cost, and broad photophysical behaviors.<sup>28–34</sup> In particular, the intra-cluster  $\text{Cu}\cdots\text{Cu}$  distance in copper clusters could be varied with external stimulus, resulting in changes in  $\text{Cu}\cdots\text{Cu}$  interaction and corresponding emissions.<sup>35,36</sup> Based on this principle, we recently reported a series of Cu(I)-CPs with attractive optical properties, *i.e.*, long afterglow and multi-stimuli-responsive, properties, including thermochromism, mechanochromism, and solvatochromism/vapochromism.<sup>37–40</sup>

Herein, we report a Cu(I)-CP,  $[\text{Cu}_2\text{I}_2(\text{PPh}_3)_2(\text{PYZ})]$  (**CIPP**,  $\text{PPh}_3$  = triphenylphosphine,  $\text{PYZ}$  = pyrazine), which has a flexible one-dimensional (1D) chain structure. **CIPP** displays different blue shifts in the luminescence after the adsorption of  $\text{CH}_2\text{Cl}_2$ ,  $\text{CHCl}_3$ , and  $\text{CH}_3\text{CN}$  vapors. In order to investigate the vapochromic mechanism, **CIPP** with corresponding guests was also prepared. Detailed structural analyses show that compared to **CIPP**, the adsorption of guest molecules makes the  $\text{Cu}\cdots\text{Cu}$  distances longer, resulting in luminescent blue shifts and higher emission energy. Meanwhile, **CIPP** exhibits fast response speed (11 s) and satisfactory reversibility. Furthermore, **CIPP** can retain  $\text{CH}_2\text{Cl}_2$  for a long time, which is helpful for applications in the detection processes.

## Results and discussion

### Syntheses and characteristics

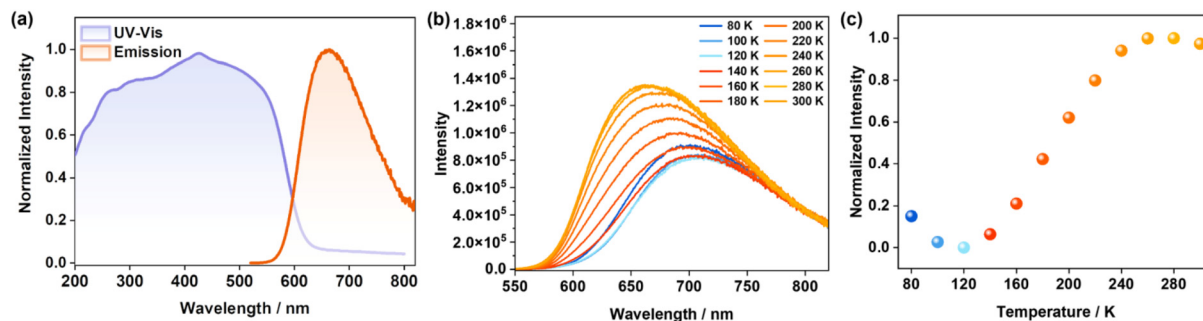
**CIPP**· $\text{CH}_2\text{Cl}_2$  (**CIPP-D**) was synthesized by the reaction of  $\text{CuI}$ ,  $\text{PPh}_3$ , and  $\text{PYZ}$  in a  $\text{CH}_2\text{Cl}_2$  solution. After drying under vacuum, the  $\text{CH}_2\text{Cl}_2$  guest in **CIPP-D** was removed and the red **CIPP** crystals were obtained, referring to the reported method of a similar structure.<sup>41</sup> Single-crystal X-ray diffraction (SCXRD) demonstrates that **CIPP** belongs to the monoclinic  $C2/c$  space group. The crystallographic asymmetric unit of **CIPP** contains one and a half  $[\text{Cu}_2(\mu\text{-I})_2]$  clusters as nodes, a bridging bidentate  $\text{PYZ}$ , and three monodentate  $\text{PPh}_3$  ligands, which directly coordinate with the copper ions (Fig. 1a). In contrast to the similar structure that has been reported,<sup>42</sup> there are two different clusters in **CIPP** with intra-cluster  $\text{Cu}\cdots\text{Cu}$  distances of 2.7338(13) Å ( $\text{Cu1}\cdots\text{Cu1A}$ ) and 2.8557(16) Å ( $\text{Cu2}\cdots\text{Cu2}$ ), respectively (Fig. S1†). These two types of copper clusters are arranged at the ABAB intervals along the CP chain. The  $[\text{Cu}_2(\mu\text{-I})_2]$  clusters are interconnected by  $\text{PYZ}$  as a bridging ligand into the 1D chains (Fig. 1b). These 1D chains are stacked with each other to form the 3D structure *via* aromatic  $\text{C-H}\cdots\pi$  interaction (Fig. 1c and S2†). Thermogravimetric (TG) analysis indicated that the **CIPP** was stable up to 115 °C, and no significant weight loss was observed before the collapse, proving that there was no guest molecule in the **CIPP** (Fig. S3†).



**Fig. 1** (a) The asymmetric unit and (b) 1D chain of **CIPP**. Color codes: Cu, orange; I, purple; N, indigo; P, pink; C, gray; H, light purple. (c) The stacking mode between the chains of **CIPP**. Different colors are employed to distinguish separate chains. Hydrogen atoms are omitted for clarity.

### Photophysical properties

UV-vis absorption and emission spectra of **CIPP** are shown in Fig. 2a. It shows the strong absorption from 200 to 600 nm, and the maximum absorption wavelength ( $\lambda_{\text{abs}}$ ) was centered at approximately 427 nm, which can be assigned to the mixed metal-to-ligand charge transfer (MLCT) and halogen-to-ligand charge transfer (XLCT) transitions.<sup>43</sup> Because the  $\text{Cu}\cdots\text{Cu}$  distance in **CIPP** is about 2.8 Å, the strong  $\text{Cu(I)}\cdots\text{Cu(I)}$  interaction should be considered, which will affect the mixed MLCT/XLCT transition, and is usually accompanied by the longer wavelength emission. As observed under 365 nm excitation, **CIPP** shows orange-red luminescence with the maximum emission wavelength ( $\lambda_{\text{em}}$ ) at 663 nm (Fig. S4†). This  $\lambda_{\text{em}}$  with rather low energy indicates the fairly large MLCT composition and strong  $\text{Cu(I)}\cdots\text{Cu(I)}$  interaction. Meanwhile, the emissions of **CIPP** and **CIPP-D** do not change when varying the excited wavelength (260–460 nm), demonstrating the same emission center (Fig. S5†). Because the luminescence of Cu(I) clusters is reported to be sensitive to temperature, the emission spectra of **CIPP** at different temperatures were measured. As shown in Fig. 2b, from 80 K to 120 K, the emission of **CIPP** displayed both red-shifting and intensity-decreasing tendencies with increasing temperature, which agreed with the common thermal quenching (TQ) phenomenon.<sup>44</sup> However, when the temperature was further increased, contrastingly, the emissions of **CIPP** showed blue shifts with the gradually enhanced intensity (Fig. 2c). Detailedly, the emission of **CIPP** shifted from 710 nm to 663 nm (120–300 K), which showed a significant wavelength shift of 47 nm (0.12 eV). Although some studies have promoted that such temperature-dependent luminescence in Cu(I) complexes may be caused by the “thermally activated delayed fluorescence” mechanism,<sup>45,46</sup> in this work, we tend to attribute this phenomenon to the slight

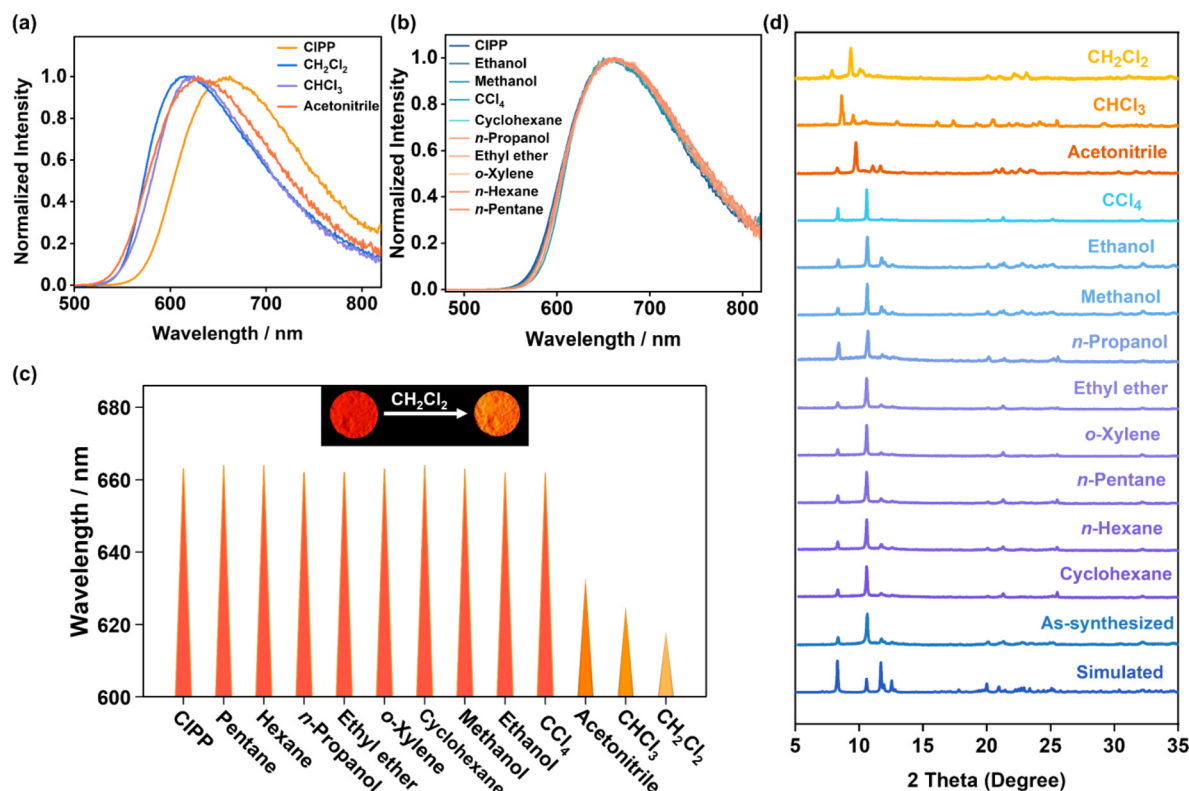


**Fig. 2** (a) UV-vis absorption and emission spectra (excited at 365 nm) of **CIPP** in air at room temperature. (b) Temperature-dependent emission spectra of **CIPP**, from 80 K to 300 K, excited at 365 nm. (c) A scatter plot of the integral emission intensity of **CIPP** at the corresponding temperature, based on (b).

lengthening of the Cu(I)⋯Cu(I) distance,<sup>47,48</sup> as the SCXRD measurements at variable temperatures further confirmed it (Fig. S6 and Table S1†). The increase in the Cu(I)⋯Cu(I) distance weakens the Cu⋯Cu interaction, resulting in the luminescent blue shift. Because the luminescence blue shifts from the near-infrared to the visible region, its non-radiative transition is reduced. Thus, besides the blue shift, it also exhibits luminescence intensity enhancement at the same time.

Interestingly, the obvious vapochromism of **CIPP** was observed (Fig. 3). Especially, when exposed to CH<sub>2</sub>Cl<sub>2</sub> vapor,

the luminescence of **CIPP** exhibited a distinct blueshift (45 nm), with the emission color change from orange-red (663 nm) to orange-yellow (618 nm). Moreover, other VOC vapors, such as CH<sub>3</sub>CN and CHCl<sub>3</sub>, can also result in similar blue shifts but with smaller changes (30 and 38 nm, respectively, Fig. 3a and c, respectively). When **CIPP** was exposed to other types of halohydrocarbon vapors, for example, CH<sub>2</sub>Br<sub>2</sub>, CH<sub>2</sub>I<sub>2</sub>, and CHBr<sub>3</sub>, it exhibited similar luminescent vapochromism behavior, indicating that **CIPP** can detect the molecules with a similar size to CH<sub>2</sub>Cl<sub>2</sub> (Fig. S7†). On the contrary,



**Fig. 3** Emission spectra of **CIPP** before and after exposure in (a) acetonitrile, CH<sub>2</sub>Cl<sub>2</sub>, CHCl<sub>3</sub> vapors, and (b) other VOCs. (c) The λ<sub>em</sub> that was obtained from (a) and (b), excited at 365 nm. Insert: a photograph of **CIPP** before and after exposure in CH<sub>2</sub>Cl<sub>2</sub> under 365 nm UV light. (d) PXRD patterns of **CIPP** before and after exposure in different VOCs.

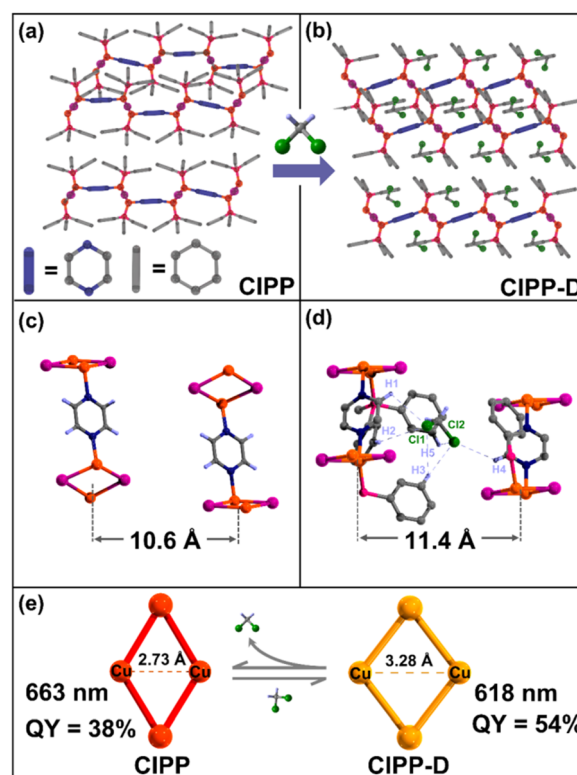
exposure to other common VOC vapors did not lead to the obvious luminescence variation for **CIPP** (Fig. 3b and c), which suggested the selective luminescence sensing and the potential application of halohydrocarbon and  $\text{CH}_3\text{CN}$  detection.

### Mechanism of sensing for VOCs

Powder X-ray diffraction (PXRD) patterns were measured to reveal the structural transformation during the vapochromism. As shown in Fig. 3d, The PXRD results show that the structure of **CIPP** does not change after exposure to the most common VOC vapors, including  $\text{CCl}_4$ , ethanol, methanol, and cyclohexane, consistent with the corresponding irresponsive emission spectra shown in Fig. 3b and c. It is worth mentioning that the structure of **CIPP** in the gate-closing state is close-packed, making it difficult to adsorb these VOC molecules. The PXRD patterns in other VOC vapors, including  $\text{CH}_2\text{Cl}_2$ ,  $\text{CHCl}_3$ , and  $\text{CH}_3\text{CN}$ , show remarkable changes (Fig. 3d). This means that these organic molecules are able to interact with **CIPP** by vapor diffusion, which opens the channels, illustrating the structural flexibility of **CIPP**. Such flexibility enables it to adsorb different molecules to form the structures in the opening state. The luminescence lifetimes of **CIPP** before and after exposure to VOC vapors are listed in Fig. S8,<sup>†</sup> which are all  $\mu\text{s}$ -scale and assigned to the phosphorescence.

To further clarify the structural transformations of **CIPP** in the vapochromic processes, the crystallographic structures of **CIPP** with different guest molecules were analyzed in detail, including **CIPP-D** (with  $\text{CH}_2\text{Cl}_2$  guest), **CIPP-C** (with  $\text{CHCl}_3$  guest), and **CIPP-A** (with acetonitrile guest), which crystallize from the corresponding organic solvents (Fig. S9<sup>†</sup>). The  $\lambda_{\text{em}}$  of the as-synthesized **CIPP-D** is almost the same as that of the fumigated **CIPP** powder in  $\text{CH}_2\text{Cl}_2$  vapor. The comparison of the PXRD patterns of  $\text{CH}_2\text{Cl}_2$ -fumigated **CIPP** and **CIPP-D** indicates that the transformation from **CIPP** to **CIPP-D** is mostly complete (Fig. S10<sup>†</sup>). Meanwhile, the luminescence sensing is still sufficient by using only the surface of the compound, as the excitation light is difficult to deeply penetrate the crystals (Fig. S11<sup>†</sup>). The completed structural transformation was observed by directly immersing **CIPP** in  $\text{CH}_2\text{Cl}_2$  solvent (Fig. S12<sup>†</sup>). Similar transformations for **CIPP** after exposure to  $\text{CHCl}_3$  and  $\text{CH}_3\text{CN}$  can be also observed (Fig. S13 and S14<sup>†</sup>).

As shown in Fig. 4a and b, the molecular chains in **CIPP-D** are similar to those in guest-free **CIPP**, suggesting that they are not reconstructed after absorbing  $\text{CH}_2\text{Cl}_2$ . For **CIPP-D**, the gaps between molecular chains are filled with  $\text{CH}_2\text{Cl}_2$  molecules, resulting in the expansion of the structure, which exhibits the flexibility of the crystal (Fig. 4b). Each  $\text{CH}_2\text{Cl}_2$  molecule in **CIPP-D** is enclosed by four phenyl rings of  $\text{PPh}_3$  and two PYZ ligands from the two adjacent chains (Fig. S15<sup>†</sup>). The chlorine atoms of the  $\text{CH}_2\text{Cl}_2$  molecule point toward the pyrazine ligands of two adjacent chains. The dihedral angle between the pyrazine ligands of the two adjacent chains is *ca.*  $25^\circ$  and creates a void to accommodate a  $\text{CH}_2\text{Cl}_2$  molecule (Fig. 4d). The chains of **CIPP-D** are also stacked by weak intermolecular interactions, such as those of **CIPP** (Fig. S16<sup>†</sup>). The structures in **CIPP-C** with  $\text{CHCl}_3$  and **CIPP-A** with  $\text{CH}_3\text{CN}$  are



**Fig. 4** Molecular chains stacking in (a) **CIPP** and (b) **CIPP-D** viewed along the *b*-axis, the purple cylinders represent the PYZ ligands and the gray ones are for the phenyl rings of  $\text{PPh}_3$ . The distance between adjacent molecular chains in (c) **CIPP** and (d) **CIPP-D**. Interactions in **CIPP-D**: Cl1...H1, 3.33 Å; Cl1...H2, 3.26 Å; Cl1...H3, 3.15 Å; Cl2...H3, 3.35 Å; Cl2...H4, 3.33 Å; Cl2...H5, 3.40 Å. (e) Cu...Cu distances of the  $[\text{Cu}_2(\mu\text{-l})_2]$  clusters in **CIPP** and **CIPP-D**. Color codes: Cu, orange; l, purple; Cl, green; N, indigo; P, pink; C, gray; H, light purple.

similar with those of **CIPP-D** (Fig. S17<sup>†</sup>). The  $\text{CH}_2\text{Cl}_2$  molecules are ordered in the crystal structure and stabilized by multi-weak interactions with PYZ and  $\text{PPh}_3$  ligands in the lattice (Fig. 4d and S18<sup>†</sup>). The closest distance between Cl atoms of  $\text{CH}_2\text{Cl}_2$  and H atoms of triphenylphosphine is 3.08 Å, which shows the large tendency of Cl...H hydrophobic interaction. In addition, the closest C-H... $\pi$  distance between  $\text{CH}_2\text{Cl}_2$  and the aromatic ring is only 3.56 Å. In order to analyze the proportion of multi-weak interactions, Hirshfeld surfaces analysis was used (Fig. S19<sup>†</sup>).<sup>49,50</sup> The 2D fingerprint plots illustrated that the weak host-guest Cl...H interactions occupied a higher fraction (58.5%). The theoretical calculation demonstrated that the total binding energy between  $\text{CH}_2\text{Cl}_2$  and **CIPP** framework is  $85 \text{ kJ mol}^{-1}$  (Table S2<sup>†</sup>). Meanwhile, the adjacent inter-chain distance in **CIPP-D** (11.44 Å) is markedly longer compared with that in **CIPP** (10.60 Å), due to the insertion of  $\text{CH}_2\text{Cl}_2$  (Fig. 4c and d). Furthermore, the average occupancy area per chain for **CIPP-D** ( $31.2 \text{ \AA}^2$ ) is also larger than that for **CIPP** ( $27.9 \text{ \AA}^2$ ). Similarly, there is also a difference in the average occupancy areas of the molecular chains for **CIPP-C** ( $35.8 \text{ \AA}^2$  per chain) and **CIPP-A** ( $30.3 \text{ \AA}^2$  per chain), which are mainly dependent on the sizes of the guest molecules (Fig. S20<sup>†</sup>).



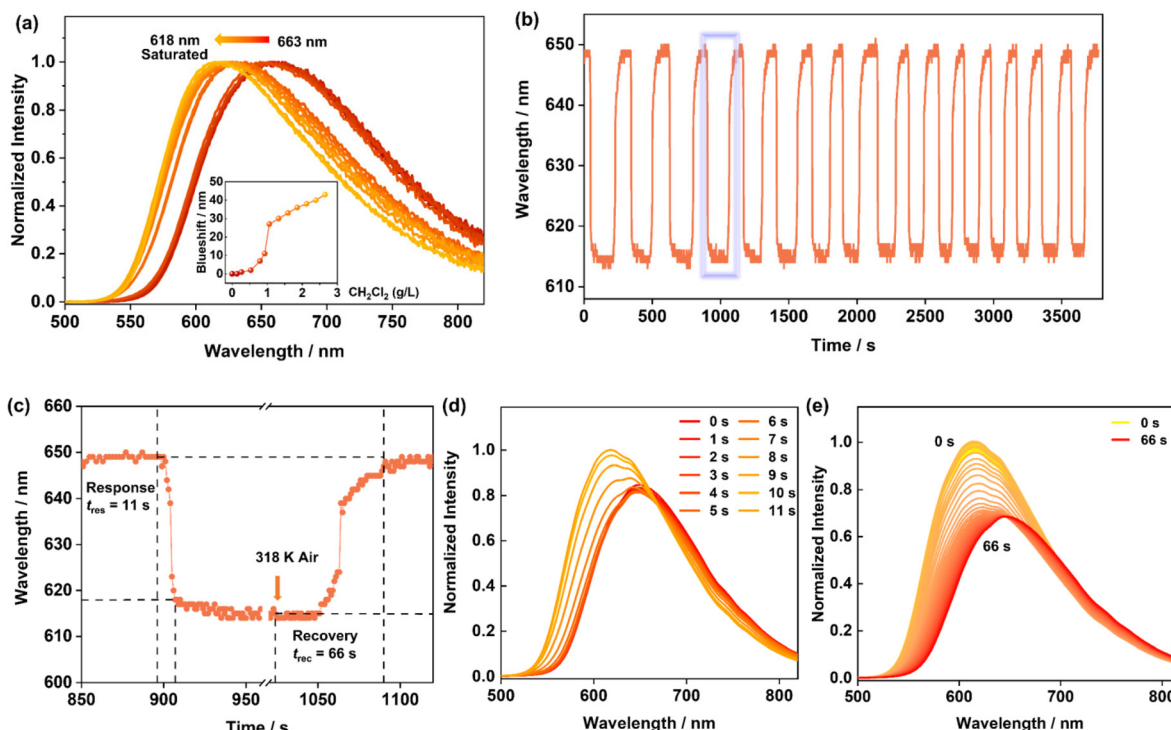
The intra-cluster Cu...Cu distance in **CIPP** is 2.73 Å, indicating a strong Cu...Cu interaction that leads to greatly reduced emission energy (1.87 eV). While for **CIPP-D**, with the lengthening of the inter-chain distance, [Cu<sub>2</sub>(μ-I)<sub>2</sub>] cluster stretches and the Cu...Cu distances increased to 3.28 Å, and 3.2378(8) Å for **CIPP-A** and 3.1479(7) Å for **CIPP-C** (Fig. S21†). For this reason, the intra-cluster Cu...Cu interaction in **CIPP-D** almost disappeared, thus it showed an obvious blueshift emission compared to that of **CIPP** (Fig. 4e). In addition, the increase of the Cu...Cu distance may enhance luminescence efficiency, as the photoluminescence quantum yields (PLQYs) are 38% for **CIPP**, and 54% for **CIPP-D** (Fig. 4e). This conclusion also explains why **CIPP** exhibits thermal luminescence enhancing phenomenon mentioned above, as the Cu...Cu distance in **CIPP** can slightly increase the warming process (120–298 K, Fig. S6†).

### CH<sub>2</sub>Cl<sub>2</sub> sensing properties

As **CIPP** shows the best sensing performance to CH<sub>2</sub>Cl<sub>2</sub> with the greatest change of the luminescence among the VOCs, its responding performance to CH<sub>2</sub>Cl<sub>2</sub> was analyzed in detail. First, when **CIPP** was exposed to the atmosphere of CH<sub>2</sub>Cl<sub>2</sub> at different concentrations, it displayed a regularly changing tendency. As shown in Fig. 5, with the increase in CH<sub>2</sub>Cl<sub>2</sub> concentration, the emission of **CIPP** gradually shifted from 663 nm to 618 nm. Moreover, a significant blueshift occurred at the

CH<sub>2</sub>Cl<sub>2</sub> concentration of 1 g L<sup>-1</sup> (Fig. 5a), suggesting that the channel begins to open at this concentration.

The sensing speed and reversibility of **CIPP** were also tested by *in situ* experiments. The real-time continuous emission spectra were recorded using a charge-coupled device (CCD). CH<sub>2</sub>Cl<sub>2</sub> molecules absorbed in **CIPP** could be removed by heating at 318 K, which was proved by both PXRD patterns and emission spectra (Fig. S22 and S23†). Therefore, to test the reversibility, **CIPP** powder was alternately placed in the atmosphere of saturated CH<sub>2</sub>Cl<sub>2</sub> vapors at room temperature, and warm air without CH<sub>2</sub>Cl<sub>2</sub> at 318 K. It should be noted that **CIPP** still showed a similar response to CH<sub>2</sub>Cl<sub>2</sub> vapors after 16 cycles and the structure was maintained even after cyclic responses (Fig. 5b and S22†). As shown in Fig. 5d, in the saturated CH<sub>2</sub>Cl<sub>2</sub> vapor, the luminescence of **CIPP** rapidly blue shifts within 11 s, suggesting a rather fast response. After the stabilization of the luminescence spectra, which suggested the saturation of CH<sub>2</sub>Cl<sub>2</sub> absorption, CH<sub>2</sub>Cl<sub>2</sub> from the sample was removed by blowing heated air (318 K), and the recovery time (back to the luminescence of **CIPP** phase) was about 66 s (Fig. 5e). The regenerated **CIPP-D** was able to maintain the emission without significant shift for about 5 min at room temperature (Fig. S24†), indicating that **CIPP** showed a relatively strong absorption of CH<sub>2</sub>Cl<sub>2</sub>, thus reducing the luminescence change in the measurement and improving the accuracy. In addition to CH<sub>2</sub>Cl<sub>2</sub>, the response time for CHCl<sub>3</sub> and



**Fig. 5** (a) Emission spectra of **CIPP** in the atmosphere with various CH<sub>2</sub>Cl<sub>2</sub> concentrations (0, 0.133, 0.265, 0.53, 0.795, 0.928, 1.06, 1.33, 1.59, 1.86, 2.12, 2.39, 2.65 g L<sup>-1</sup>, from right to left) at room temperature. (b) The cyclic luminescence responses of **CIPP** under the alternation between saturated CH<sub>2</sub>Cl<sub>2</sub> vapor and hot air (318 K). The vertical coordinate is the  $\lambda_{em}$  of the spectra per second. (c) Response curves in the range of purple square in (b) with a time range from 896 s to 907 s. (d) Time-dependent emission spectra of **CIPP** in saturated CH<sub>2</sub>Cl<sub>2</sub> vapor, from 0 s to 11 s. (e) Time-dependent emission spectra of CH<sub>2</sub>Cl<sub>2</sub>-absorbed **CIPP** sample at 318 K, recorded once per second.

CH<sub>3</sub>CN vapors were also measured. For CHCl<sub>3</sub>, the blue shift of  $\lambda_{em}$  was completed within 3 min (Fig. S25†). Such wavelength shift is smaller, and the response time is longer compared with that of the CH<sub>2</sub>Cl<sub>2</sub> vapor. While for CH<sub>3</sub>CN, the luminescence blueshift was completed within 60 s (Fig. S26†). The faster response speed for CH<sub>2</sub>Cl<sub>2</sub> may be attributed to the rather high saturation vapor pressure of CH<sub>2</sub>Cl<sub>2</sub> (46.5 kPa at 293 K), compared with those of CHCl<sub>3</sub> (21.28 kPa at 293 K) and CH<sub>3</sub>CN (9.33 kPa at 293 K).<sup>51,52</sup>

## Conclusions

A Cu(I)-CP named **CIPP** was synthesized and characterized extensively. **CIPP** displayed luminescence vapochromic response to CH<sub>2</sub>Cl<sub>2</sub>, CHCl<sub>3</sub>, and CH<sub>3</sub>CN vapors. Benefitting from the flexibility of **CIPP**, the intra-cluster Cu...Cu distance varies when it absorbs guest molecules and transforms into the gate-opening state. Especially, the interaction with CH<sub>2</sub>Cl<sub>2</sub> allows **CIPP** to display a remarkable luminescence wavelength for the blue shift (45 nm). Moreover, **CIPP** exhibits fast luminescence color variation (*ca.* 11 s) and excellent reversibility for sensing CH<sub>2</sub>Cl<sub>2</sub>. Due to the multi-weak interactions, **CIPP** can hold CH<sub>2</sub>Cl<sub>2</sub> for 5 minutes at room temperature in air.

## Experimental

All the chemicals were obtained from commercial sources and used as received without further purification unless otherwise noted. Powder X-ray diffraction (PXRD) patterns were recorded on a Rigaku Mini diffractometer with Cu-K $\alpha$  ( $\lambda$  = 1.54184 Å) radiation. UV-vis absorption spectra were recorded on a Shimadzu UV-3600i Plus UV-VIS-NIR absorption spectrometer. Thermogravimetric (TG) analyses were performed on a METTLER TOLEDO TGA/DSC 3+ instrument with a ramping rate of 10.0 °C min<sup>-1</sup> under a nitrogen atmosphere. Elemental analysis (C, H, and N) was performed on an Elementar Vario EL Cube.

### Preparation

For **CIPP-D** and **CIPP**: PYZ (16 mg, 0.2 mmol) and PPh<sub>3</sub> (105 mg, 0.40 mmol) were completely dissolved in dichloromethane (5 mL), followed by the addition of CuI (76 mg, 0.4 mmol) at room temperature. The orange **CIPP-D** crystals were formed from the transparent liquid after about 7 hours. The crystals were filtered and washed with CH<sub>2</sub>Cl<sub>2</sub>, then dried under vacuum to obtain orange-red **CIPP** crystals (yield = 60%). Elemental analysis calcd for C<sub>20</sub>H<sub>17</sub>CuINP (**CIPP**): C, 48.75; H, 3.48; N, 2.84. Found: C, 48.80; H, 3.83; N, 2.64.

For **CIPP-A**: **CIPP-A** was synthesized by a similar method to that of **CIPP-D** by replacing the CH<sub>2</sub>Cl<sub>2</sub> solvent with the CH<sub>3</sub>CN solvent (yield = 54%).

For **CIPP-C**: PYZ (32 mg, 0.4 mmol) and PPh<sub>3</sub> (105 mg, 0.40 mmol) were completely dissolved in CHCl<sub>3</sub> (5 mL) at

278 K, followed by the addition of CuI (76 mg, 0.4 mmol). Then, the solution was kept at 278 K for about one day, and orange **CIPP-C** crystals were observed. The crystals were filtered and washed with cold CHCl<sub>3</sub> (yield = 47%). Among these, **CIPP-A** was reported in the previous literature.<sup>36</sup>

### Single-crystal X-ray diffraction

Diffraction data were collected on a Rigaku XtaLAB single-crystal diffractometer by using Cu-K $\alpha$  radiation ( $\lambda$  = 1.54184 Å). The structures were solved using the direct methods and refined with the full-matrix least-squares method on  $F^2$  using the SHELXTL package.<sup>53</sup> Anisotropic thermal parameters were used to refine all non-hydrogen atoms. All hydrogen atoms were generated geometrically. Crystallographic data and details of data collection and refinements are summarized in Table S1.† CCDC 2258386 (**CIPP**), 2258387 (**CIPP-D**), 2258388 (**CIPP-C**), and 2258389 (**CIPP-150K**) contain supplementary crystallographic data for this work.†

### Photoluminescence measurement

Photoluminescence measurements were performed on an Edinburgh FLS1000 fluorescence spectrometer. The emission spectra were recorded on the spectrometer equipped with a continuous Xe lamp. The luminescence decay experiments were performed on the same spectrometer equipped with a variable pulsed laser (VPL) at 375 ± 10 nm as the excitation source. The temperature was controlled using an Oxford temperature controller. The photoluminescence quantum yields (PLQYs) were measured on the same spectrometer in the integrating sphere.

The *in situ* experiment was recorded using an Ocean QE Pro charge-coupled device (CCD) equipped with a 365 nm LED. Real-time sensing of CH<sub>2</sub>Cl<sub>2</sub> was realized using CH<sub>2</sub>Cl<sub>2</sub> liquid to provide the saturated vapor under ambient conditions. After removing CH<sub>2</sub>Cl<sub>2</sub> using hot air at 318 K, **CIPP** was cooled to room temperature by blowing normal air (detected by thermometer).

## Conflicts of interest

There are no conflicts to declare.

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