Polymorphic Copper Iodide Anions: Luminescence Thermochromism and Mechanochromism of (PPh₄)₂[Cu₂I₄]

*Yaouen Thefioux,^a Marie Cordier,^b Florian Massuyeau,^a Camille Latouche,^a Charlotte
Martineau-Corcos,^{c,d} and Sandrine Perruchas^{a*}*

^a Université de Nantes, CNRS, Institut des Matériaux Jean Rouxel, IMN, F-44000 Nantes, France.
Phone: (+33) (0)2 40 37 63 35. E-mail: sandrine.perruchas@cnrs-imn.fr

^b Université de Rennes, CNRS, ISCR (Institut des Sciences Chimiques de Rennes), UMR 6226, F-35000
Rennes, France.

^c MIM, Institut Lavoisier de Versailles (ILV), UMR CNRS 8180, Université de Versailles St-Quentin
en Yvelines (UVSQ), 45, avenue des Etats-Unis, 78035 Versailles Cedex, France.

^d CNRS, CEMHTI UPR 3079, Université d'Orléans, F-45071 Orléans, France.

Abstract. The photoluminescent stimuli-responsive properties of two crystalline polymorphs of (PPh₄)₂[Cu₂I₄] formula, are reported. Distinct luminescence properties are exhibited by these ionic copper iodide compounds with blue or yellow emission and original luminescence thermochromism and mechanochromism are demonstrated. While one polymorph displays contrasted temperature-dependent emission properties, the other shows great modification of its emission upon mechanical solicitation. Establishment of structure-properties relationships supported by theoretical approach, permit to get insights into the origin of the photoluminescence properties and the mechanisms at play. According to DFT calculations, the different emission bands originate either from the (PPh₄)⁺ organic cation or from the [Cu₂I₄]²⁻ anion. Activation of these two emissive centers appears to be dependent on the crystalline packing of the polymorph. The thermochromism displayed by one polymorph can be attributed to variation in temperature of the relative intensities of two emission bands of two different excited states. The origin is different for the other polymorph with emission bands coming from two independent emissive centers namely, (PPh₄)⁺ and [Cu₂I₄]²⁻. The luminescence mechanochromism is attributed to a polymorphic transition. The mechanical solicitation induces a partial transformation of one polymorph into the other within a disordered phase. The mechanochromic mechanism can be related to mechanical modifications of inter-molecular interactions between the (PPh₄)⁺ cations. By displaying luminescence properties that dependent on crystalline structure, excitation wavelength, temperature and mechanical solicitation, the studied copper iodides offer a great possibility of emissive color control and switch, a clear demonstration of great potentialities of this family of compounds for developing photoactive materials.

Introduction.

Luminescent materials exhibiting multicolor emission switching are attracting attention due to their potential applications in data storage devices and security systems, for instance.^{1,2} Because of their outstanding photophysical properties and economic advantages, copper complexes are particularly appealing for developing light-emitting functional materials.^{3,4,5,6,7,8} Amongst Cu-based compounds, copper(I) iodide derivatives have recently emerged as attractive emissive materials due to their rich optical properties.^{9,10,11,12,13}

The d¹⁰ electronic configuration of Cu(I) gives rise to diverse coordination number/geometries and associated to the multiple connectivity modes of halides, the family of copper iodides present a great structural diversity, leading consequently to various photophysical properties. Actually, their potential applications as sensors,^{14,15,16,17} photocatalytic systems,¹⁸ and emitters in organic light-emitting diodes (OLEDs),^{19,20,21,22,23} have been recently reported. Copper(I) iodide compounds can also display diverse stimuli-responsive properties such as luminescence thermochromism,^{24,25} mechanochromism,^{26,27} rigidochromism,²⁸ or vapo/solvatochromism.²⁹ Combining multistimuli-responsive properties is actually particularly appealing to access original multifunctional materials. The molecular cubane clusters formulated [Cu₄I₄L₄] (L = ligand) is indeed well known for exhibiting multistimuli-responsive properties that are closely related to metallophilic interactions. Within the family of neutral copper iodides, another iconic complex is the rhomboid one formulated [Cu₂I₂L₂].^{19,30,31} While its emissive properties have been well described, its stimuli-responsive properties remain rare.^{27a} Although fewer compared with neutral derivatives, charged copper iodide complexes are documented whose structural geometry vary with the counter-ion.¹³ Common anions are the binuclear [Cu₂I₆]⁴⁻ and the tetranuclear [Cu₄I₈]⁴⁻³² ones for which iodides replace the organic ligands, present in the neutral complexes. Note that zwitterionic compounds which covalently bind the anion and cation, have been recently reported.³³ Another binuclear anion is [Cu₂I₄]²⁻ which has been encountered in several associations with different organic and inorganic cations.^{34,35,36,37,38,39,40,41,42,43,44,45,46,47,48,49,50,51,52,53,54,55,56,57,58,59,60,61,62,63,64,65,66} The photophysical properties of these compounds have been rarely characterized, [Cu₂I₄]²⁻ being simply treated as an inorganic counter-anion and not as an emissive species by itself. In particular, with the tetraphenylphosphonium cation, the polymorphic structure of (PPh₄)₂[Cu₂I₄] has been reported but no photophysical characterization has been investigated.^{67,68} Occurrence of crystalline polymorphism provides a great tool to explore structure-properties relationships of molecular solids and to get insights into their physical properties.⁶⁹ In particular, crystalline polymorphs of luminescent compounds usually exhibit different photophysical properties allowing determining the mechanisms at the origin of their light-emitting properties and even stimuli-responsive ones. This opportunity motivated our investigations for exploring the photophysical properties of the (PPh₄)₂[Cu₂I₄] compounds.

Here, we report on the stimuli-responsive properties of two crystalline polymorphs of (PPh₄)₂[Cu₂I₄] resulting from the association of the [Cu₂I₄]²⁻ anion and the (PPh₄)⁺ cation (Figure 1). Original luminescence thermochromism and mechanochromism are demonstrated for these compounds. Comparative structural and photophysical characterisations of the two polymorphs are presented in order to elucidate the mechanochromism mechanism at play. Theoretical studies have been also conducted aiming at rationalizing the experimental light-emitting properties.

Figure 1. Schematic drawing of the $[\text{Cu}_2\text{I}_4]^{2-}$ anion and $(\text{PPh}_4)^+$ cation.

Results and Discussions.

The two crystalline polymorphs of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$, namely **A** and **B**, were synthesized in solution by reacting copper iodide (CuI) and tetraphenylphosphonium iodide ($\text{P}(\text{C}_6\text{H}_5)_4\text{I}$) in dichloromethane, acetone or acetonitrile, at room temperature (details in experimental section). By slowly adding an anti-solvent (diethyl-ether or cyclohexane) to the reaction solution, the products precipitated as colorless crystalline powders. Both polymorphs are formed concomitantly but their relative proportion vary with the nature of the used solvents. For the dichloromethane/cyclohexane solvents, polymorph **A** is largely predominant but for the acetonitrile/diethyl-ether combination this is the polymorph **B**. Rapid concentration of an acetone solution leads exclusively to polymorph **B**. The $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ formula is in agreement with elemental analysis (experimental section). The reported synthesis for these two polymorphs (**A** and **B**), uses Cu , I_2 and PPh_4I as reactants in acetone. Concomitant formation was also reported in this previous work.⁶⁷ The synthesis of two other known polymorphs namely $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ -**C** and **D** was also described with acetone as solvent with Cu , I_2 and PPh_4I as reactants.⁶⁸ None of these two latter were obtained in our synthesis conditions.

Figure 2. Photos of the crystalline polymorphs $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ under ambient light and under UV irradiation (lamp at 365 nm) with the yellow-emitting **A** and blue-emitting **B**, at room temperature.

The colorless crystals of the two polymorphs exhibit distinctive emission colors under UV irradiation, with a yellow emission for **A** while that of **B** is blue (Figure 2). Real crystalline polymorphs displaying such different emission properties in the visible spectra are quite rare. Their respective crystal structures were determined by SCXRD (single crystal X-Ray diffraction) analysis at different temperatures (100, 150, 200 and 293 K, Table S1). Their structures were already solved⁶⁷ at room temperature but here are reported low temperature measurements to correlate with the photoluminescence data. The purity of the

samples was confirmed by powder X-ray diffraction analysis (Figure S3) and elemental analysis (experimental part in SI). **A** and **B** crystallize in the monoclinic $P2_1/c$ space group. The unit cell contents are shown in Figure 3. Both structures can be described as an alternation of organic layers formed by the PPh_4^+ cations and inorganic layers formed by the $[\text{Cu}_2\text{I}_4]^{2-}$ anions. No solvent of crystallization is included in the structures giving to these two compounds the important characteristic of real crystalline polymorphs. Indeed, the comparative study is in this case straightforward and not hampered by the potential effect of solvent of crystallization.

Figure 3. View of the unit cell of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ -**A** and **B** polymorphs at 293 K.

The $(\text{PPh}_4)^+$ cation in **A** and the two independent ones in **B**, present a classical geometry with a tetrahedral environment around the phosphorus atom. Free rotation around the P-C bond leads to different orientation of the phenyl groups (Figure S1). The molecular structure of the $[\text{Cu}_2\text{I}_4]^{2-}$ anions are depicted in Figure 4. All Cu atoms present a pseudo-trigonal CuI_3 environment with iodide atoms in μ_1 and μ_2 coordination modes. In polymorph **A**, $[\text{Cu}_2\text{I}_4]^{2-}$ has an inversion center whereas in **B** it is in general position. Selected bond lengths and angles in the $[\text{Cu}_2\text{I}_4]^{2-}$ anions at 293 K are listed in Table 1. The Cu-I bond distances and I-Cu-I angle values are within the range of reported values for such copper iodide compounds.³⁴⁻⁶⁶ The Cu-I μ_1 -bond distances are shorter than the Cu-I μ_2 ones in agreement with the stronger ionic interaction. The intramolecular Cu-Cu bond distances in **A** ($\text{Cu}_1\text{-Cu}_1 = 2.955(1)$ Å) is much longer compared with that in **B** ($\text{Cu}_1\text{-Cu}_2 = 2.662(1)$ Å at 293 K). These Cu-Cu distances lie in the range of values encountered in copper iodide compounds.^{9,13,26} Nevertheless comparison with the value of the sum of the van der Waals radii of Cu of 2.80 Å, suggests that strong cuprophilic bonding interaction ($d^{10}\text{-}d^{10}$) occurs in **B**, whereas it is much weaker in the case of **A**.^{70,71} Values of the torsional angles in the $[\text{Cu}_2\text{I}_4]^{2-}$ anions are significantly different in the two polymorphs (mean values of 173.86(4) in **A** and 162.07(4)° in **B**), as reported in Table 1. As shown in Figure 4, these geometric differences lead in **A** to an almost planar form of the $[\text{Cu}_2\text{I}_4]^{2-}$ anion and to a strongly folded one in **B**. Bond lengths and angles in the $[\text{Cu}_2\text{I}_4]^{2-}$ anions at lower temperatures are reported in Table 1 (100 and 293 K) and S3

(100, 150, 200 and 293 K). Similar temperature dependent behavior is observed for both polymorphs with crystal lattice contraction (from 293 to 100 K) of 2.3 and 3.0 % in **A** and **B**, respectively. The Cu-Cu distances present a shortening upon temperature decreasing of 3.0 and 2.1 % in **A** and **B**, respectively. Concerning the Cu-I bond distances (mean values), slight increases of 0.2 % for **A** and 0.1 % for **B** are observed by lowering the temperature. These observations mean that upon decreasing the temperature, the Cu-Cu distances contract while Cu-I ones remain almost constant. This can be explained by the Cu-Cu interactions of d^{10} - d^{10} type which are weaker compared with the ionic Cu-I bonds and their modification minimizes the crystal packing energy. This phenomena has been already observed in copper iodide clusters by applying isostatic pressure.⁷² Even if the thermal lattice contraction is larger in **B** compared with that of **A**, this is the $[\text{Cu}_2\text{I}_4]^{2-}$ anion in **A** that presents the more important structural changes. This can be due to the longer Cu-Cu distances in **A** that is far from the van der Waals value, being therefore more easily compressible. The density can be a criteria to evaluate the relative stability of crystalline polymorphs, with more stable compounds having denser phase.⁶⁹ However, the very close density values of **A** and **B** (1.863 and 1.851 respectively at 293 K) does not give useful information in the present case.

Figure 4. Molecular structure of $[\text{Cu}_2\text{I}_4]^{2-}$ anion in $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ -**A** and **B** polymorphs at 293 and 100 K.

Table 1. Selected intramolecular bonds lengths and angles of (PPh₄)₂[Cu₂I₄] at 100 and 293 K from SCXRD.

Compound	A		B	
	100 K	293 K	100 K	293 K
Cu⁺...Cu	2.8652(9)	2.9549(13)	2.6072(5)	2.6621(9)
Cu-I_{μ2}	2.5833(6)*2	2.5806(7)*2	2.5711(5)	2.5720(8)
	2.5907(6)*2	2.5932(7)*2	2.5740(4)	2.5780(8)
Cu-I_{μ1}	2.5126(7)*2	2.4954(7)*2	2.6033(4)	2.6029(8)
			2.6076(5)	2.6086(7)
			2.4969(4)	2.4858(7)
<i>mean</i>	2.5622(7)	2.5564(7)	2.4976(4)	2.4904(7)
I_{μ1}-Cu-I_{μ2}	125.87(2)*2	127.09(3)*2	125.19(2)	124.71(3)
	120.98(2)*2	122.26(3)*2	119.92(2)	121.27(3)
I_{μ2}-Cu-I_{μ2}	112.75(2)*2	110.34(2)*2	124.57(2)	124.58(3)
			119.99(2)	121.12(3)
			114.80(1)	113.94(3)
I-Cu-I-Cu	172.76(3)*2	173.68(4)*2	114.85(2)	113.93(3)
	173.16(3)*2	174.04(4)*2	159.14(3)	160.15(4)
			159.27(3)	160.22(4)
			164.21(3)	163.89(4)
			164.51(3)	164.00(4)

The great majority of the reported compounds incorporating the [Cu₂I₄]²⁻ anion present a planar geometry. In addition to the case of (PPh₄)₂[Cu₂I₄],^{67,68} only three examples of [Cu₂I₄]²⁻ anion with a folded geometry are known, with the As(Ph₄)⁺,⁴⁰ (CH₃PPh₃)⁺,⁴⁵ and Ba(tetraglyme)₂²⁺,³⁹ cations. The spherical geometry of these three cations may account for structural constraints leading to the folded geometry. This reflects the flexibility of copper(I) halide complexes that can adapt their molecular geometry to the crystalline packing. Regarding the Cu-Cu distances, the values with the As(Ph₄)⁺,⁴⁰ (CH₃PPh₃)⁺,⁴⁵ and Ba(tetraglyme)₂²⁺,³⁹ cations are 2.663(4), 2.678(3) and 2.636(1) Å, respectively, which are similar to that of **B** (2.662(1) Å). They indeed all lie within the range of 2.6-2.8 Å of the literature values. The relatively long value of 2.955(1) Å found in **A**, can be associated to its planar geometry. Indeed, the planar [Cu₂I₄]²⁻ anion with comparable Cu-Cu distance of 2.938(1) Å has been reported with a phosphonium anion (*i*-Pr(Ph₃P))⁺.³⁴

Short intermolecular contacts (< 3 Å) are present in the crystalline structure of **A** and **B**. Several C-H...C contacts occur between the phenyl groups of the phosphonium cations (2.897(1), 2.911(1) and 2.914(1) Å in **A** and 2.928(1), 2.949(1), 2.922(1) and 2.854(1) Å in **B**). Cu...H contacts also exist in **A** of 2.631(1) Å whereas the shortest one in **B** is of 2.938(1) Å (Figure S2). As shown in Figure 5, π-π stacking can be also observed in **A** with C...C contacts of 3.398(1) and 3.950(1) Å. Such interaction is not found in **B**. The different networks of interactions between the phenyl groups in **A** and **B** are confirmed by the different FTIR signatures around 3050 cm⁻¹ (Figure S4).

Figure 5. Intermolecular π - π interactions in $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]\text{-A}$.

Structural comparison with the two other known polymorphs, **C** and **D**,⁶⁸ indicates similarities between **A** and **C** and, **B** and **D**. $[\text{Cu}_2\text{I}_4]^{2-}$ anion presents a planar geometry in **C** as observed in **A** and a similar close packing of $(\text{PPh}_4)^+$ cations is found in these two polymorphs. In **D**, the geometry of the $[\text{Cu}_2\text{I}_4]^{2-}$ anion is folded as in **B**. These polymorphs however present different arrangements of the $(\text{PPh}_4)^+$ cations that are less dense compared with that of **A** and **C**.

Luminescent mechanochromism is exhibited by polymorph **B** whereas no emission modification was observed for **A** under mechanical sollicitation. As shown in Figure 6, the manual grinding of crystalline powder of **B** results in a great emission change by shifting from blue to yellow light. Meanwhile, the ground phase, noted **Bg** (*g* stands for ground), maintains the slightly yellowish body-color of the initial crystalline sample. To our knowledge, **B** is the only example of CuI-based mechanochromic compounds of ionic nature. Note that mechanosynthesis was tested to synthesize $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ by grinding the reactants with drops of solvent. This was unsuccessful to obtain polymorph **A** but from PXRD analysis, polymorph **B** was detected. After careful examination, polymorph **B** was not obtained but the ground form **Bg** instead. The mechanosynthesis has therefore to be employed carefully when mechanochromic compounds are involved.

Figure 6. Photos of crystalline powder of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]\text{-B}$ partially ground (right side) under visible light and UV irradiation (365 nm UV lamp), at room temperature.

Results of elemental analysis of **Bg** are similar to those of **B** corresponding to the $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ formula (experimental section), indicating that no chemical reaction related to adsorption or removal of atmospheric molecules are induced by the grinding process. Solid state ^{31}P NMR analysis (*vide infra*) also demonstrates that the molecular structure of the complex is preserved in **Bg**. The powder X-Ray

diffraction (PXRD) analysis of **B** revealed numerous sharp and intense diffraction peaks which indicate good crystallinity (Figure 7). After grinding, intensity decrease and broadening of the peaks is observed that indicates partial alteration of the crystalline packing and possible amorphization.

FTIR spectra were recorded for the three compounds along with that of PPh₄I for comparison. DFT calculations performed on (PPh₄)⁺ and [Cu₂I₄]²⁻ moieties (anharmonic computations, detailed in SI), permit to simulate their IR signatures and to assign the vibration bands. Post-treatments were performed using the VMS package.⁷³ Experimental FTIR spectra (4000-500 cm⁻¹) are similar for all the compounds and clearly correspond to the tetraphenylphosphonium cation's signature. Simulated spectrum is in very good agreement with that of PPh₄I (Figures S4-5). Main peaks observed at 1433, 1103, 722, 686 and 526 cm⁻¹ can thus be attributed to combination of C-C and C-H vibrations. More specifically, the three highest wavenumbers comprise C-H bending, rocking and wagging motions. The two lowest wavenumbers correspond to C-H scissoring together with C-C breathing. Furthermore, the anharmonic treatment demonstrates that all peaks between 1800 and 2800 cm⁻¹ correspond to overtones and combination bands. Spectra have been also recorded in the far-IR region (650-50 cm⁻¹) and present differences between the two polymorphs but with a common peak at 177 cm⁻¹ (Figure S6). The spectra of **Bg** has different peaks compared with that of **B** (203, 71 cm⁻¹) which are observed in **A** suggesting structural similarities between them. The peak at 177 cm⁻¹ in **A** and **B** is found at 179 cm⁻¹ in **Bg**. This peak, simulated at 168 cm⁻¹, corresponds to Cu-I bond stretching vibrations of [Cu₂I₄]²⁻ (Figure S7). The experimental peak at 152 cm⁻¹ can also be attributed to Cu-I bond vibrations calculated at 127 cm⁻¹. At lower frequencies, two peaks are observed which are different for the two polymorphs (**A**: 71, 53 cm⁻¹ and **B**: 68 and 48 cm⁻¹). Corresponding values for **Bg** (71 and 51 cm⁻¹) are closer to those of **A** indicating again structural similarities. From simulations, these two vibrations (73 and 31 cm⁻¹) may correspond to an out of plane bending and a wagging modes of the [Cu₂I₄]²⁻ anion. Differences observed can be attributed to the different planar and folded geometries of the anion. Values reported for (PNP)₂[Cu₂I₄] (71 and 54 cm⁻¹)⁶⁶ whose geometry is planar, are indeed close to those of **A**.

Figure 7. PXRD of **B** and **Bg** recorded at room temperature.

The reversibility of the mechanochromic effect was tested by solvent treatment but without success. With the different tested solvents, the initial blue emission of **B** is not recovered and instead either the yellow emission of **Bg** is maintained (cyclohexane, diethyl-ether) or a white emission appears

(acetonitrile, acetone). This irreversibility can be explained by the formation of these copper iodide compounds which is highly dependent to the experimental conditions such as the solvent. This is illustrated by the rich polymorphism and isomerism displayed by this family of compounds.

To get insight into the local structure of the ground phase in light of that of the two polymorphs, solid-state NMR analysis has been conducted. In addition to the classical ^1H and ^{13}C nuclei, ^{31}P and $^{63,65}\text{Cu}$ ones have been studied. Those two latter are particularly relevant to probe the copper iodide structural geometries.^{74,75} ^1H MAS NMR spectra (Figure S8) present broad signatures centered at 7.5 ppm which are in agreement with the phosphonium cation ($\text{P}(\text{C}_6\text{H}_5)_4^+$) with phenyl groups characteristics. ^{13}C MAS NMR spectra present two broad features around 135 and 120 ppm, the latter corresponding to carbon atoms bonded to the phosphorus atom (Figure S8). No significant difference is observed between the samples. The ^{31}P signatures shown in Figure 8, are also in accordance with the phosphonium cation. The singlet observed at 22 ppm for **A** results from the crystallographically independent $(\text{PPh}_4)^+$ and for **B** the two peaks at 21 and 23 ppm correspond to the two inequivalent cations. The different structural conformation of the phosphonium within the structure of the two polymorphs are thus revealed thanks to the sensitivity of the ^{31}P nucleus. The two peaks are also present in the ^{31}P spectrum of **Bg** but with a broadened feature. This indicates the preservation of the structure of **B** upon grinding but with creation of some local structural disorder in accordance with the PXRD analysis.

Figure 8. ^{31}P CPMAS solid state NMR spectra of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ -**A**, **B** and **Bg**.

The ^{63}Cu static NMR spectra are shown in Figure 9. They present particularly broad signals extending over 1.5 MHz. For **A**, the quadrupolar coupling constant (C_Q) is about 45.6 MHz, with asymmetry parameter (η_Q) of 0.34. This large C_Q value is typical of copper atoms in three-coordinated CuI_3 geometry and is clearly distinct from the four-coordinate PCuI_3 , P_2CuI_2 and CuI_4 copper environment signatures (usually 20-25 MHz C_Q values).⁷⁶ The ^{63}Cu NMR spectrum of **A** can be simulated using a single component (Figure S9), in agreement with the single Cu site in the crystal structure. In **B**, the two individual components are still strongly overlapped (a deconvolution is suggested in Figure S10), despite the use of high magnetic field (17.6 T). Both C_Q values are in the 45-47 MHz range, again, in agreement with copper atoms in three-coordinated CuI_3 geometry. The two polymorphs present distinct ^{65}Cu NMR signatures but the correlation with their structure is difficult to establish at this stage. The spectra of **Bg** is very similar to that of **B**, indicating the preservation of the $[\text{Cu}_2\text{I}_4]^{2-}$ anion upon grinding. Partial

amorphization is however not obvious in this case, probably due to predominance of the signal of intact crystalline part.

Figure 9. ^{63}Cu static WURST-QCPMG NMR spectra of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ -**A**, **B** and **Bg**. The span of the patterns is about 9000 ppm, corresponding to ca 1.7 MHz. The stars indicate signals due to the copper coil of the NMR probehead.

As shown in Figure 10, all studied compounds are slightly yellowish crystalline powders under daily light at room temperature whereas under UV illumination they present distinct luminescence colors with bright yellow, blue and yellowish emissions for **A**, **B** and **Bg**, respectively. By lowering the temperature down to 77 K, significant change of the emissions colors is observed with a yellow to orange shift for **A**, a yellow to blue shift for **Bg** while for **B** the pristine blue emission turns into deeper blue. This phenomenon is reversible upon warming up with recovery of the initial emissions. **A** and **Bg** thus present contrasted luminescence thermochromism. Because the temperature dependent emission color change is lesser for **B** compared with that of **Bg**, the grinding process induces in some way luminescence thermochromism to the compound.

Figure 10. Photos of crystalline powder of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ -**A**, **B** and **Bg** under visible light and UV irradiation (365 nm UV lamp) at room temperature (298 K) and in liquid nitrogen (77 K).

Solid-state emission and excitation spectra were recorded from 77 to 293 K. Corresponding temperature-dependent spectra are shown in Figure S12 and related data are reported in Table 2. As shown in Figure 11, at 293 K the emission spectra of **A** displays an unstructured emission band centered at 545 nm ($\lambda_{\text{ex}} = 380$ nm), in accordance with the observed yellow emission. The emission band of **B** centered at 465 nm presents a shoulder at 545 nm whereas for **Bg** this is the opposite, with the main emission band at 545 nm and the shoulder at 465 nm ($\lambda_{\text{ex}} = 380$ nm). The grinding thus induces a significant red-shift of the main emission of 80 nm that is among the largest values reported for mechanochromic compounds.⁷⁷ This agrees with the contrasted color emission change from blue to yellow observed upon grinding (Figure 6). Because the emission spectra of **Bg** is clearly a combination of the emission of the **A** and **B** polymorphs, the mechanical sollicitation seems to induce a transition between the two polymorphs.

The associated emission lifetimes present single exponential decays for **A** and **B** with $\tau = 1.01$ and 0.07 μs respectively, whereas **Bg** displays a biexponential decay with $\tau_1 = 0.14$ and $\tau_2 = 0.93$ μs values (Figure S13). These measurements reveal the formation upon grinding of new emissive species with longer lifetime values, close to that of **A**. This confirms the mechanical transformation of the **B** polymorph into the **A** one. The absolute internal quantum yields determined at room temperature are $\text{QY} = 10$, 2 and 1 % for **A**, **B** and **Bg**, respectively (Table 2).

Figure 11. Solid-state luminescence spectra of **A**, **B** and **Bg** at 293 and 77 K with normalized emission in solid lines and excitation spectra in dotted lines.

The emission band of **A** does not present large change upon lowering the temperature down to 175 K with a moderate intensity increase ($\lambda_{\text{ex}} = 380$ nm). Below 175 K, a new band centered at 650 nm appears whose intensity increases down to 77 K, thus becoming eventually predominant compared with the 545 nm band (Figure 11 and S12). This agrees with the orange emission displayed at 77 K (Figure 10). Relatively narrow excitation bands are observed around 400 nm for the higher energy band ($\lambda_{\text{em}} = 540$ nm) at 293 and 77 K compared to the broad feature of the low energy one ($\lambda_{\text{em}} = 650$ nm). For **B**, the emission at 465 nm presents a moderate blue-shift (from 465 to 455 nm) by lowering the temperature down to 77 K but with a great intensity increase ($\times 50$). In opposite, the intensity of the secondary band at 545 nm decreases rapidly upon cooling and is not observed anymore at 77 K (Figure 11). No significant change is observed for the broad excitation bands ($\lambda_{\text{em}} = 460$ nm). **Bg** presents similar temperature-dependent behavior compared with **B** by displaying a great increase of the intensity of the 460 nm emission band upon cooling. However, intensity of the emission band at 545 nm progressively increases from 293 to 200 K. At 225 K, the two emission bands have similar intensity and at 77 K the high energy one is largely predominant (Figure 11 and S12). The excitation bands present more structured profiles at low temperature ($\lambda_{\text{em}} = 450$ nm).

Figure 12. Solid-state photoluminescence maps (excitation vs. emission) of **A**, **B** and **Bg** at 293 and 77 K.

In addition to temperature-dependent emission properties, the three compounds also display excitation-dependent emissions with important color tuning possibility. This is illustrated by the photoluminescence maps recorded at 293 and 77 K and reported in Figure 12. With a narrow excitation

range around 415 nm, the emission properties of polymorph **A** are not excitation-dependent at 293 K (temperature dependent luminescence spectra at $\lambda_{\text{ex}} = 415$ nm are reported in Figure S12). In contrast, at 77 K, the additional emission band at 650 nm is excitable at wavelength below 390 nm. At 77 K, the two emission bands (545 and 650 nm) can thus be either simultaneously ($\lambda_{\text{ex}} = 385$ nm) or independently excited ($\lambda_{\text{ex}} = 415$ and 350 nm respectively, Figure S12). In the case of **B**, there is no excitation dependent properties at 77 K with a wide range of possible excitations (280-400 nm) for the 465 nm emission band. This is different at 293 K with the second emission band at 550 nm that is excitable for wavelengths above 390 nm (Figure 12). This band cannot be solely excited but is, at $\lambda_{\text{ex}} = 415$ nm, of comparable intensity compared with the blue one (Figure S12). For **Bg** at 293 K, the two emission bands at 560 and 465 nm can be both excited around 380 nm. However, at $\lambda_{\text{ex}} = 415$ nm, the emission band at 560 can be solely excited (Figure S12). This dual emissions system is also present at 77 K but with a more contrasted excitation dependency. Indeed, the more intense emission band is this time the blue one (460 nm) accompanied with the yellow one (560 nm). The latter can be excited individually at wavelengths above $\lambda_{\text{ex}} = 410$ nm leading to a contrasted emission color change.

Table 2. Photoluminescence data of **(PPh₄)₂[Cu₂I₄]-A**, **B** and **Bg**. Lifetimes and quantum yields have been measured at 293 K.

	λ_{max} (nm) [λ_{ex} (nm)]		Lifetime τ (μs) [λ_{em} (nm)]	Quantum Yield QY (%) [λ_{ex} (nm)]
	293 K	77 K		
A	545 [380]	650 [380]	1.01 [550]	10 [415]
	545 [415]	533 [415]		
B	465 [380]	455 [380]	0.07 [460]	2 [380]
Bg	545 [380]	455 [380]	0.14 (41 %) 0.93 (59 %) [560]	1 [380]

Three different emission bands are thus observed for **(PPh₄)₂[Cu₂I₄]** (whatever the polymorph and the temperature) at roughly $\lambda_{\text{max}} = 460$, 550, and 650 nm. These transitions can originate from the two different emissive centers which are the organic cation **(PPh₄)⁺** and the inorganic anion **[Cu₂I₄)²⁻**. Luminescence of tetraphenylphosphonium halides have been indeed reported and in particular **PPh₄I** exhibits an emission at $\lambda_{\text{max}} = 430$ nm which has been attributed to phosphorescence.⁷⁸ On the anionic side, the emission properties of ionic copper halide compounds (no ligand in this case) can be attributed to ³XMCT (X stands for halide) and ³MC (metal centered) transitions with for the latter, possible influence of Cu-Cu interactions. Only few photophysical characterizations have been reported in the literature for **[Cu₂I₄)²⁻** based compounds. For **(btx)[Cu₂I₄]** (btx = 1,6-bis(triethylmethylammonium)xylol), an emission band was observed at $\lambda_{\text{max}} = 463$ nm which was surprisingly attributed to a MLCT transition.³⁵ Single emission bands were also reported for **[Ba(tetraglyme)₂][Cu₂I₄]**³⁹ and **[Cu(Me-PPy₃)I]₂[Cu₂I₄]**⁶³ with $\lambda_{\text{max}} = 680$ and 660 nm respectively, and were assigned to mixed XMCT and MC transitions. In the cases of **(PPh₃iPr)₂[Cu₂I₄]**³⁷ and **{[K(benzo-18-crown-6)]₂[Cu₂I₄]}_n**,³⁸ dual emissions were recorded at $\lambda_{\text{max}} = 447$, 558 nm and at $\lambda_{\text{max}} = 465$, 632 nm. These emissions were also attributed to XMCT and MC excited states, excepted the high energy band of **(PPh₃iPr)₂[Cu₂I₄]**³⁷ that was assigned to a cation centered transition. From these previous studies,

the assignment of the emission bands is not straightforward and were usually not supported by theoretical study. The only one theoretical approach was reported for $[\text{Cu}(\text{Me-PPy}_3)\text{I}]_2[\text{Cu}_2\text{I}_4]$ ⁶³ and despite large cation-anion distances, transitions from the MOs of the anion to those of the cation were suggested.

The diverse and nebulous emission origins, prompted us to perform DFT (Density Functional Theory) computations, in order to get a clear vision of the electronic structure of $(\text{PPh}_4)_2[\text{Cu}_2\text{I}_4]$ and have an accurate assignment of the emission properties. These calculations were carried on with the Gaussian16 package⁷⁹ using the PBE0 functional together with the LANL2DZ basis augmented with polarization functions for H, C and P atoms, and the triple ζ basis set DEF2TZVP for Cu and I atoms (details in SI).^{80,81} The two potential emissive centers, $(\text{PPh}_4)^+$ and $[\text{Cu}_2\text{I}_4]^{2-}$, were treated independently because of the large intermolecular distances in the crystalline structures that precludes electronic transfer.

The $(\text{PPh}_4)^+$ moiety was computed without symmetry constrain but the relaxed structure can be approximated to the S_4 symmetry point group. The four P-C bond lengths have a mean value of 1.811 Å and the C-P-C angles lie between 108 and 110°, leading to a slightly distorted tetrahedron. These values are in agreement with the corresponding experimental ones of 1.795(5) Å and 109.50(2)° (mean values). As shown in the orbital diagram in Figure 13, the last occupied orbitals are π -bonding ones and centered on phenyl moieties whereas the unoccupied ones are localized on the phosphorus atom together with phenyl moieties with a π^* -antibonding character. The optimized structure exhibits a gap of *ca* 6 eV, it is therefore a very stable structure from thermodynamics viewpoint. The computed triplet excited state of $(\text{PPh}_4)^+$ keeps the pseudo S_4 symmetry point group. In that case the P-C bonds are ranging from 1.792 to 1.813 Å and angles from 107 to 110°. Therefore, the structural distortion appears to be stronger at the excited state than at the ground state. The corresponding simulated electronic phosphorescence is centered at 462 nm. Vibrational contributions to electronic transitions were then taken into account to simulate the emission spectrum using the so-called Adiabatic Shift ansatz (vibrational modes detailed in SI).^{82,83} For this compound, temperature effects were also considered during the vibronic calculation.⁸⁴ The simulated emission curve (Figure S14), is in a very good agreement with the emission spectra of PPh_4I recorded at room temperature. These results demonstrate that PPh_4^+ can be responsible for the emission at 460 nm observed for **B**. Furthermore, the simulated IR spectra and experimental FTIR spectra of PPh_4I are in good agreement (Figure S5), giving confidence in this analysis.

Figure 13. Kohn-Sham orbital diagrams of PPh_4^+ and $[\text{Cu}_2\text{I}_4]^{2-}$ at their ground states.

For $[\text{Cu}_2\text{I}_4]^{2-}$, the two different geometries found in the polymorphs, planar and folded, were used as starting points to optimize the GS. It turned out that both structures converge to the same planar optimized geometry with the D_{2h} symmetry. This could indicate that the folded geometry results from constraints of the crystalline packing. In this planar relaxed geometry, the computed Cu-I μ_1 and Cu-I μ_2 bond distances values of 2.571 and 2.658 Å respectively, are in agreement with the experimental data of 2.495(1) and 2.587(1) Å (polymorph A). This is similar for the angles with computed values of I μ_1 -Cu-I μ_2 = 128.2° and I μ_2 -Cu-I μ_2 = 103.6° and corresponding experimental ones of 124.68(3)° and 110.34(3)°. Calculated Cu-Cu bond distance of 3.288 Å is in relative good accordance with the experimental one of 2.955(1) Å. As shown in the orbital diagram in Figure 13, the HOMO and HOMO-1,2 are mainly localized on iodide atoms with smaller contribution of Cu atoms (3d). These latter present a strong Cu-I μ_2 bonding and Cu-I μ_1 antibonding characters. The LUMO possesses a strong Cu-Cu bonding character which will lead to a strong shortening of the bond length at the excited state. The computed gap of 5.2 eV is also large. The most intense absorption transitions calculated by TD-DFT mainly correspond to, HOMO $\rightarrow$ LUMO, HOMO-1 $\rightarrow$ LUMO transitions with wavelengths around 290 nm. These calculated wavelengths are in reasonable agreement with the experimental broad absorption band centered at 325 nm (Figure S11). The absorption transitions have thus mixed $^1\text{XMCT}$ and ^1MC character. Two triplet excited states were calculated for $[\text{Cu}_2\text{I}_4]^{2-}$ and, as expected, their optimized geometries present a strong shortening of the Cu-Cu distance (2.463 and 2.468 Å) relative to that of the fundamental state. In both cases, the I μ_2 atoms leave their positions and become bonded to a single copper atom, leading to two CuI_2 entities linked by the short Cu-Cu bond. The flexibility of copper iodides may account for this coordination change. The first excited state computed in TD-DFT, has for origin a transition from the HOMO-1 into the LUMO with Cu-Cu bonding characteristic. Therefore, the corresponding transition is of $^3\text{XMCT}$ and ^3MC character. The calculated emission wavelength is around 620 nm which could represent the experimental emission centered at 550 nm. In this excited state, the structure is indeed strongly distorted, both I μ_2 become I μ_1 and two sets of Cu-I appear (2 $\times$ 2.649 and

2×2.602 Å) with I-Cu-I angle value of 108°. The second excited state was calculated using the unrestricted approach. In that case, the C_s symmetry is found upon optimization. The corresponding mean Cu-I distances are 2.598 Å and the I-Cu-I angle is of 109°. It should be pointed out that both triplets have a similar geometrical shape, the Cu-I being shorter and the structure less distorted for this second excited state comparatively to the first one. The spin density (Figure S15) clearly demonstrates that the unpaired electrons are shared by the Cu atoms. This excited state gives rise to an emission wavelength computed at 840 nm which is comparable to the experimental one at 650 nm. Significant energetic deviations between the computed emissions and the experimental ones are here obtained (0.25 eV and 0.43 eV). The computed redshift between the two emissions around 0.52 eV is however relatively close to the experimental one (0.35 eV).

According to the DFT calculations, the different emission bands observed for $(PPh_4)_2[Cu_2I_4]$ originate either from the $(PPh_4)^+$ organic cation ($\lambda_{max} = 460$ nm) or from the $[Cu_2I_4]^{2-}$ anion ($\lambda_{max} = 550$, and 650 nm) (Figure S16). Activation of these two emissive centers appears to be dependent on the molecular arrangement in the solid-state. The lack of the blue band for polymorph **A** can be related to the π -orbital overlap present in its crystalline structure that are deleterious for the emission of phenyl-based system. This effect can be referred to ACQ ‘aggregation-caused quenching’ phenomenon commonly encountered for π -conjugated motifs.⁸⁵ The fact that the red band ($\lambda_{max} = 650$ nm) is not observed for polymorph **B** can be attributed to the structural geometry of the $[Cu_2I_4]^{2-}$ anion. This low energy band is indeed theoretically predicted for the planar geometry but could be non-radiative for the folded form as in **B**.

The difference between the two polymorphs is also evidenced by their temperature-dependent emission properties. Indeed, a contrasted luminescence thermochromism is observed for **A** whereas for polymorph **B** its temperature dependent emission is less pronounced. Luminescence thermochromism of **B** is however induced by mechanical solicitation, **Bg** displaying a yellow emission at room temperature that becomes blue upon decreasing the temperature. This blue shift contrasts with the observed redshift of **A** upon cooling. The thermochromism displayed by **A** can be related to that of $[Cu_4I_4L_4]$ cubane clusters with variation in temperature of the relative intensities of two emission bands corresponding to two different excited states (Figure S16).^{24,25} For **Bg**, the origin is different because the emission bands do not originate from the same molecule but come from two independent emissive centers namely $(PPh_4)^+$ and $[Cu_2I_4]^{2-}$.

The luminescence mechanochromism displayed by **B** appears to be the result of a polymorphic transition. The mechanical solicitation induces a partial transformation of the **B** polymorph into the **A** one. From DFT geometrical optimization, **A** is actually the most thermodynamically stable polymorph. Structural characterizations indicate that this created **A** polymorph is in a disordered (amorphous) phase in **Bg**. The mechanochromic mechanism can be thus related to modification of the structural packing and in particular of the π - π interactions between the $(PPh_4)^+$ cations. Structural disorder may favors the formation of such intermolecular interactions that quench the emission band of $(PPh_4)^+$ and consequently enhance that of the low-energy exhibited by $[Cu_2I_4]^{2-}$. The present mechanochromism thus implies intermolecular mechanical transformation that contrasts to intramolecular mechanism reported for $[Cu_4I_4L_4]$

cubane clusters for which cuprophilic interactions are modified.^{26f} By displaying mechanochromic properties in close relation with polymorphism, the flexible adaptation of copper iodide compounds to the structural environment is highlighted in the present study.

Conclusion.

The two studied copper iodide crystalline polymorphs displaying distinct luminescence properties constitute a neat example of molecular packing-dependent emission. Structural and photophysical characterizations of these real polymorphs permit to reveal their original stimuli-responsive properties. Theoretical investigations allowed to get insight into the mechanisms at play by rationalizing the emission properties and showing that the cationic and anionic counterparts are both emissive centers.

Combined luminescence thermochromism and mechanochromism are demonstrated for these ionic copper iodide compounds. Their emission properties are additionally highly dependent on the excitation wavelength. By displaying luminescence properties that dependent on crystalline structure, excitation wavelength, temperature and mechanical solicitation, these compounds offer a great possibility of color control and switch. By reporting original photoactive properties of ionic copper iodide compounds, this study demonstrates the potentialities of these family of compounds for developing optical stimuli-responsive materials.

Acknowledgments. CMC is grateful for financial support from the Institut Universitaire de France (IUF). Financial support from the IR-RMN-THC Fr3050 CNRS for conducting the research is gratefully acknowledged. H. Brault and R. Utrera-Melero are thanked for their assistance. CL thanks access to the HPC resources of TGCC under allocation 2019-A0070911052.

ASSOCIATED CONTENT

Supporting Information Available. The X-ray crystallographic files (CIF), the experimental section, the DFT and other characterization data. This information is available free of charge via the Internet at <http://pubs.acs.org>.

References

- (1) Naren, G.; Hsu, C.-W.; Li, S.; Morimoto, M.; Tang, S.; Hernando, J.; Guirado, G.; Irie, M.; Raymo, F. M.; Sundén, H.; Andréasson, J.; An all-photonic full color RGB system based on molecular photoswitches. *Nature Com.* **2019**, 3996-4003.
- (2) Huang, G.; Xia, Q.; Huang, W.; Tian, J.; He, Z.; Li, B. S.; Tang, B. Z. Multiple Anti-Counterfeiting Guarantees from a Simple Tetraphenylethylene Derivative - High-Contrasted and Multi-State Mechanochromism and Photochromism. *Angew. Chem. Int. Ed.* **2019**, 58, 17814-17819.
- (3) Wenger, O. S. Photoactive Complexes with Earth-Abundant Metals. *J. Am. Chem. Soc.* **2018**, 140, 13522-13533.
- (4) Hamze, R.; Peltier, J. L.; Sylvinson, D.; Jung, M.; Cardenas, J.; Haiges, R.; Soleilhavoup, M.; Jazzar, R.; Djurovich, P. I.; Bertrand, G.; Thompson, M. E. Eliminating nonradiative decay in Cu(I) emitters: >99% quantum efficiency and microsecond lifetime. *Science* **2019**, 363, 601-606.

- (5) Yam, V. W.-W.; Au, V. K.-M.; Leung, S. Y.-L. Light-Emitting Self-Assembled Materials based on d^8 and d^{10} Transition Metal Complexes. *Chem. Rev.* **2015**, *115*, 7589-7728.
- (6) Cariati, E.; Lucenti, E.; Botta, C.; Giovannella, U.; Marinotto, D.; Righetto, S. Cu(I) hybrid inorganic-organic materials with intriguing stimuli responsive and optoelectronic properties. *Coord. Chem. Rev.* **2016**, *306*, 566-614.
- (7) Hu, Z.; Deibert, B. J.; Li, J. Luminescent metal-organic frameworks for chemical sensing and explosive detection. *Chem. Soc. Rev.* **2014**, *43*, 5815-5840.
- (8) Wenger, O. S. Vapochromism in Organometallic and Coordination Complexes: Chemical Sensors for Volatile Organic Compounds. *Chem. Rev.* **2013**, *113*, 3686-3733.
- (9) Peng, R.; Li, M.; Li, D. Copper(I) halides: A versatile family in coordination chemistry and crystal engineering. *Coord. Chem. Rev.* **2010**, *254*, 1-18.
- (10) Kobayashi, A.; Kato, M. Stimuli-responsive Luminescent Copper(I) Complexes for Intelligent Emissive Devices. *Chem. Lett.* **2017**, *46*, 154-162.
- (11) Wallesch, M.; Volz, D.; Zink, D. M.; Schepers, U.; Nieger, M.; Baumann, T.; Bräse, S. Bright copper opportunities: Multinuclear CuI complexes with N-P ligands and their applications. *Chem. Eur. J.* **2014**, *20*, 6578-6590.
- (12) Fang, Y.; Liu, W.; Teat, S. J.; Dey, G.; Shen, Z.; An, L.; Yu, D.; Wang, L.; O'Carroll, D. M.; Li, J. A Systematic Approach to Achieving High Performance Hybrid Lighting Phosphors with Excellent Thermal- and Photostability. *Adv. Funct. Mater.* **2017**, *27*, 1603444.
- (13) Yu, T.-L.; Guo, Y.-M.; Wu, G.-X.; Yang, X.-F.; Xue, M.; Fu, Y.-L.; Wang, M.-S. Recent progress of d^{10} iodoargentate(I)/iodocuprate(I) hybrids: Structural diversity, directed synthesis, and photochromic/thermochromic properties. *Coord. Chem. Rev.* **2019**, *397*, 91-111.
- (14) Yu, Y.; Zhang, X.-M.; Ma, J.-P.; Liu, Q.-K.; Wang, P.; Dong, Y.-B. Cu(I)-MOF: naked-eye colorimetric sensor for humidity and formaldehyde in single-crystal-to-single-crystal fashion. *Chem. Commun.*, **2014**, *50*, 1444-1446.
- (15) Wang, J.-H.; Li, M.; Li, D. A dynamic, luminescent and entangled MOF as a qualitative sensor for volatile organic solvents and a quantitative monitor for acetonitrile vapour. *Chem. Sci.* **2013**, *4*, 1793-1801.
- (16) Conesa-Egea, J.; Gallardo-Martínez, J.; Delgado, S.; Martínez, J. I.; Gonzalez-Platas, J.; Fernández-Moreira, V.; Rodríguez-Mendoza, U. R.; Ocón, P.; Zamora, F.; Amo-Ochoa, P.; Multistimuli Response Micro- and Nanolayers of a Coordination Polymer Based on Cu_2I_2 Chains Linked by 2-Aminopyrazine. *Small* **2017**, *13*, 1700965.
- (17) Kitagawa, H.; Ohtsu, H.; Kawano, M. Kinetic Assembly of a Thermally Stable Porous Coordination Network Based on Labile CuI Units and the Visualization of I_2 Sorption. *Angew. Chem. Int. Ed.* **2013**, *52*, 12395-12399.
- (18) Shi, D.; Zheng, R.; Sun, M.-J.; Cao, X.; Sun, C.-X.; Cui, C.-J.; Liu, C.-S.; Zhao, J.; Du, M. Semiconductive Copper(I)-Organic Frameworks for Efficient Light-Driven Hydrogen Generation Without Additional Photosensitizers and Cocatalysts. *Angew. Chem. Int. Ed.* **2017**, *56*, 14637-14641.
- (19) Zink, D. M.; Volz, D.; Baumann, T.; Mydlak, M.; Flüge, H.; Friedrichs, J.; Nieger, M.; Bräse, S. Heteroleptic, Dinuclear Copper(I) Complexes for Application in Organic Light-Emitting Diodes. *Chem. Mater.* **2013**, *25*, 4471-4486.
- (20) Liu, Z.; Qiu, J.; Wei, F.; Wang, J.; Liu, X.; Helander, M. G.; Rodney, S.; Wang, Z.; Bian, Z.; Lu, Z.; Thompson, M. E.; Huang, C. Simple and High Efficiency Phosphorescence Organic Light-Emitting Diodes with Codeposited Copper(I) Emitter. *Chem. Mater.* **2014**, *26*, 2368-2373.
- (21) You, Z.; Li, H.; Zhang, L.; Yu, B.; Zhang, J.; Wu, X. A Host Material for Deep-Blue Electrophosphorescence Based on a Cuprous Metal-Organic Framework. *J. Phys. Chem. C* **2017**, *121*, 23072-23079.
- (22) Liu, W.; Fang, Y.; Li, J. Copper Iodide Based Hybrid Phosphors for Energy-Efficient General Lighting Technologies. *Adv. Funct. Mater.* **2018**, *28*, 1705593.
- (23) Xie, M.; Han, C.; Zhang, J.; Xie, G.; Xu, H. White Electroluminescent Phosphine-Chelated Copper Iodide Nanoclusters. *Chem. Mater.* **2017**, *29*, 6606-6610.
- (24) (a) Ford, P. C.; Cariati, E.; Bourassa, J. Photoluminescence Properties of Multinuclear Copper(I) Compounds. *Chem. Rev.* **1999**, *99*, 3625-3647. (b) Kim, T. H.; Shin, Y. W.; Jung, J. H.; Kim, J. S.; Kim, J. Crystal-to-crystal transformation between three CuI coordination polymers and structural evidence for luminescence thermochromism. *Angew. Chem., Int. Ed.* **2008**, *47*, 685-688. (c) Lapprand, A.; Dutartre, M.; Khiri, N.; Levert, E.; Fortin, D.; Rousselin, Y.; Soldera, A.; Jugé, S.; Harvey, P. D. Luminescent P-chirogenic copper clusters. *Inorg. Chem.* **2013**, *52*, 7958-7967. (d) Fu, Z.; Lin, J.; Wang, L.; Li, C.; Yan, W.; Wu, T. Cuprous Iodide Pseudopolymorphs Based on Imidazole Ligand and Their Luminescence Thermochromism. *Cryst. Growth Des.*

2016, *16*, 2322-2327. (e) Zhan, S.-Z.; Li, M.; Zhou, X.P.; Wang, J.-H.; Yang J.-R.; Li, D. When Cu₄I₄ cubane meets Cu₃(pyrazolate)₃ triangle: dynamic interplay between two classical luminophores functioning in a reversibly thermochromic coordination polymer. *Chem. Commun.*, **2011**, *47*, 12441-12443. (f) Zhan, S.-Z.; Li, M.; Ng, S. W.; Li, D. Luminescent Metal-Organic Frameworks (MOFs) as a Chemopalette: Tuning the Thermochromic Behavior of Dual-Emissive Phosphorescence by Adjusting the Supramolecular Microenvironments. *Chem. Eur. J.* **2013**, *19*, 10217-10225.

(25) Perruchas, S.; Tard, C.; Le Goff, X. F.; Fargues, A.; Garcia, A.; Kahlal, S.; Saillard, J.-Y.; Gacoin, T.; Boilot, J.-P. Thermochromic Luminescence of Copper Iodide Clusters: The Case of Phosphine Ligands. *Inorg. Chem.* **2011**, *50*, 10682-10692.

(26) (a) Perruchas, S.; Le Goff, X. F.; Maron, S.; Maurin, I.; Guillen, F.; Garcia, A.; Gacoin, T.; Boilot, J.-P. Mechanochromic and Thermochromic Luminescence of a Copper Iodide Cluster. *J. Am. Chem. Soc.* **2010**, *132*, 10967-10969. (b) Shan, X.-C.; Jiang, F.-L.; Zhang, H.-B.; Qian, X.-Y.; Chen, L.; Wu, M.-Y.; Al-Thabaiti, S. A.; Hong, M.-C. A solid AND logic stimuli-responsive material with bright nondestructive performance designed by sensitive cuprophilicity. *Chem. Commun.* **2013**, *49*, 10227-10229. (c) Shan, X.-C.; Jiang, F.-L.; Chen, L.; Wu, M.-Y.; Pan, J.; Wan, X.-Y.; Hong, M.-C. Using cuprophilicity as a multi-responsive chromophore switching color in response to temperature, mechanical force and solvent vapors. *J. Mater. Chem. C*, **2013**, 4339-4349. (d) Benito, Q.; Le Goff, X. F.; Maron, S.; Fargues, A.; Garcia, A.; Martineau, C.; Taulelle, F.; Kahlal, S.; Gacoin, T.; Boilot, J.-P.; Perruchas, S. Polymorphic Copper Iodide Clusters: Insights into the Mechanochromic Luminescence Properties. *J. Am. Chem. Soc.*, **2014**, *136*, 11311-11320. (e) Huitorel, B.; El Moll, H.; Cordier, M.; Fargues, A.; Garcia, A.; Massuyeau, F.; Martineau-Corcos, C.; Gacoin, T.; Perruchas, S.; Luminescence Mechanochromism induced by Cluster Isomerization. *Inorg. Chem.* **2017**, *56*, 12379-12388. (f) Huitorel, B.; Utrera-Melero, R.; Massuyeau, F.; Mevelec, J.-Y.; Baptiste, B.; Polian, A.; Gacoin, T.; Martineau-Corcos, C.; Perruchas, S. Luminescence mechanochromism of copper iodide clusters: a rational investigation. *Dalton Trans.* **2019**, *48*, 7899-7909.

(27) (a) Kobayashi, A.; Yoshida, Y.; Yoshida, M.; Kato, M. Mechanochromic Switching between Delayed Fluorescence and Phosphorescence of Luminescent Coordination Polymers Composed of Dinuclear Copper(I) Iodide Rhombic Cores. *Chem. Eur. J.* **2018**, *24*, 14750-14759. (b) Wen, T.; Zhang, D.-X.; Zhang, H.-X.; Zhang, H.-B.; Zhang, J.; Li, D.-S. Redox-active Cu(I) boron imidazolate framework for mechanochromic and catalytic applications. *Chem. Commun.* **2014**, *50*, 8754-8756. (c) Xiao, Q.; Zheng, J.; Li, M.; Zhan, S.-Z.; Wang, J.-H.; Li, D., Mechanically Triggered Fluorescence/Phosphorescence Switching in the Excimers of Planar Trinuclear Copper(I) Pyrazolate Complexes. *Inorg. Chem.* **2014**, *53*, 11604-11615. (d) Deshmukh, M. S.; Yadav, A.; Pant, R.; Boomishankar, R. Thermochromic and Mechanochromic Luminescence Umpolung in Isostructural Metal–Organic Frameworks Based on Cu₆I₆ Clusters. *Inorg. Chem.* **2015**, *54*, 1337-1345. (e) Chen, K.; Nenzel, M. M.; Brown, T. M.; Catalano, V. J. Subtle Modulation of Cu₄X₄L₂ Phosphine Cluster Cores Leads to Changes in Luminescence. *Inorg. Chem.* **2015**, *54*, 6900-6909. (f) Kwon, E.; Kim, J.; Lee, K. Y.; Kim, T. H. Non-Phase-Transition Luminescence Mechanochromism of a Copper(I) Coordination Polymer *Inorg. Chem.* **2017**, *56*, 943-949. (g) Hu, L.-X.; Gao, M.; Wen, T.; Kang, Y.; Chen, S. Synthesis of Halide-Modulated Cuprous(I) Coordination Polymers with Mechanochromic and Photocatalytic Properties. *Inorg. Chem.* **2017**, *56*, 6507-6511. (h) Zhang, D.-X.; Zhang, H.-X.; Wen, T.; Lia, D.-S.; Zhang, J. Mechanochromic Cu(I) boron imidazolate frameworks with low-dimensional structures and reducing function. *Inorg. Chem. Front.* **2016**, *3*, 263-267. (i) Lu, T.; Wang, J.-Y.; Tu, D.; Chen, Z.-N.; Chen, X.-T.; Xue, Z.-L. Luminescent Mechanochromic Dinuclear Cu(I) Complexes with Macrocyclic Diamine-Tetracarbene Ligands. *Inorg. Chem.* **2018**, *57*, 13618-13630. (j) Feng, N.; Gao, C.; Guo, C.-Y.; Chen, G. Copper-Phenylacetylide Nanobelt/Single-Walled Carbon Nanotube Composites: Mechanochromic Luminescence Phenomenon and Thermoelectric Performance. *ACS Appl. Mater. Interfaces* **2018**, *10*, 5603-5608. (k) Hupp, B.; Nitsch, J.; Schmitt, T.; Bertermann, R.; Edkins, K.; Hirsch, F.; Fischer, I.; Auth, M.; Sperlich, A.; Steffen, A. Stimulus-Triggered Formation of an Anion-Cation Exciplex in Copper(I) Complexes as a Mechanism for Mechanochromic Phosphorescence. *Angew. Chem. Int. Ed.* **2018**, *57*, 13671-13675.

(28) (a) Tran, D.; Bourassa, J. L.; Ford, P. C. Pressure-Induced Luminescence Rigidochromism in the Photophysics of the Cuprous Iodide Cluster Cu₄I₄py₄. *Inorg. Chem.* **1997**, *36*, 439-442. (b) Vogler, A.; Kunkely, H. Photoluminescence of Tetrameric Copper(I) Iodide Complexes in Solution. *J. Am. Chem. Soc.* **1986**, *108*, 7211-7212. (c) Roppolo, I.; Celasco, E.; Sangermano, M.; Garcia, A.; Gacoin, T.; Boilot, J.-P.; Perruchas S. Luminescence variation by rigidity control of acrylic composite materials. *J. Mater. Chem. C*, **2013**, *1* (36), 5725-5732.

- (29) (a) Zhao, C.-W.; Ma, J.-P.; Liu, Q.-K.; Wang, X.-R.; Liu, Y.; Yang, J.; Yang, J.-S.; Dong Y.-B. An in situ self-assembled Cu₄I₄-MOF-based mixed matrix membrane: a highly sensitive and selective naked-eye sensor for gaseous HCl. *Chem. Commun.*, **2016**, 52, 5238-5241. (b) Cho, S.; Jeon, Y.; Lee, S.; Kim, J.; Kim, T. H. Reversible Transformation between Cubane and Stairstep Cu₄I₄ Clusters Using Heat or Solvent Vapor. *Chem. Eur. J.* **2015**, 21, 1439-1443. (c) Park, H.; Kwon, E.; Chiang, H.; Im, H.; Lee K. Y.; Kim, J.; Kim, T. H. Reversible Crystal Transformations and Luminescence Vapochromism by Fast Guest Exchange in Cu(I) Coordination Polymers. *Inorg. Chem.* **2017**, 56, 8287-8294. (d) Killarney, J. P.; McKinnon, M.; Murphy, C.; Henline, K. M.; Wang, C.; Pike, R. D.; Patterson, H. H. Amine- and sulfide-sensing copper(I) iodide films. *Inorg. Chem. Comm.* **2014**, 40, 18-21. (e) Benito, Q.; Balogh, C. M.; El Moll, H.; Gacoin, T.; Cordier, M.; Rakhmatullin, A.; Latouche, C.; Martineau-Corcus, C.; Perruchas, S.; Luminescence Vapochromism of a Dynamic Copper Iodide Mesocate. *Chem. Eur. J.* **2018**, 24, 18868-18872.
- (30) Liu, W.; Fang, Y.; Wei, J. Z.; Teat, S.J.; Xiong, K.; Hu, Z.; Lustig, W. P.; Li, J.; A Family of Highly Efficient CuI-Based Lighting Phosphors Prepared by a Systematic, Bottom-up Synthetic Approach. *J. Am. Chem. Soc.* **2015**, 137, 9400-9408.
- (31) Busch, J. M.; Zink, B. M.; Di Martino-Fumo, P.; Rehak, F. R.; Boden, P.; Steiger, S.; Fuhr, O.; Nieger, M.; Kloppe, W.; Gerhards, M.; Bräse, S.; Highly soluble fluorine containing Cu(I) AlkylPyrPhos TADF complexes. *Dalton Trans.*, **2019**, 48, 15687-15698.
- (32) Jalilian, E.; Liao, R.-Z.; Himo, F.; Brismar, H.; Laurell, F.; Lidin, S.; Luminescence properties of the Cu₄I₆²⁻ cluster. *Cryst. Eng. Comm.* 13, **2011**, 4729-4734.
- (33) Liu, W.; Zhu, K.; Teat, S. J.; Dey, G.; Shen, Z.; Wang, L.; O'Carroll, D. M.; Li, J. All-in-One: Achieving Robust, Strongly Luminescent and Highly Dispersible Hybrid Materials by Combining Ionic and Coordinate Bonds in Molecular Crystals. *J. Am. Chem. Soc.*, **2017**, 139, 9281-9290.
- (34) Jalilian, E.; Lidin, S. Bis(isopropyl-triphenyl-phospho-nium) di-μ-iodido-bis-[iodidocuprate(I)]. *Acta Cryst.* **2010**, E66, m432-m433.
- (35) Du, H. J.; Liang, Y.; Wang, L. F.; Niu, Y. Y. Syntheses, Crystal Structures, UV-Vis Absorption, and Fluorescent Properties of a Complex: [btx][Cu₂I₄]. *Synth. React. Inorg.* **2012**, 42, 364-368.
- (36) Liu, Y.-F.; Chen, J.-Z.; Huang, C.-C. Bis{4-[4-(dimethylamino)styryl]-1-ethylpyridinium} di-μ-iodido-bis[iodidocuprate(II)]. *Acta Cryst. E* **2007**, m2957-m2962.
- (37) Zhang, W.-T.; Liu, J.-Z.; Liu, J.-B.; Song, K.-Y.; Li, Y.; Chen, Z.-R.; Li, H.-H.; Jiang, R. Quaternary Phosphorus-Induced Iodocuprate(I)-Based Hybrids: Water Stabilities, Tunable Luminescence and Photocurrent Responses. *Eur. J. Inorg. Chem.* **2018**, 4234-4244.
- (38) Wu, F.; Tong, H.; Li, Z.; Lei, W.; Liu, L.; Wong, W.-Y.; Wong, W.-K.; Zhu, X. A white phosphorescent coordination polymer with Cu₂I₂ alternating units linked by benzo-18-crown-6. *Dalton Trans.*, **2014**, 43, 12463-12466.
- (39) Mishra, S.; Jeanneau, E.; Ledoux, G.; Daniele, S. Novel Barium–Organic Incorporated Iodometalates: Do They Have Template Properties for Constructing Rare Heterotrimetallic Hybrids? *Inorg. Chem.* 53 **2014** 11721-11731.
- (40) Asplund, M.; Jagner, S. Crystal Structure of Bis(tetrapropylammonium) Di-μ-iodo-diiodocuprate(I), [N(C₃H₇)₄]₂[Cu₂I₄]. *Acta Chem. Scand. A* 38, **1984**, 411-421. and references therein
- (41) Basu, A.; Bhaduri, S.; Sapre, N.Y.; Jones, P.G. Copper Complexes of 1,1'-Di(2-pyridylethanol): X-Ray Structures and Reaction with Oxygen. *J. Chem. Soc., Chem. Commun.* **1987** 1724-1725.
- (42) Cunningham, D.; Gallagher, J.F.; Higgins, T.; McArdle, P.; McGinley, J.; Sheerin, D. A New Co-ordinating Role for a Metal Salicylaldimine. *J. Chem. Soc., Chem. Commun.* **1990**, 959-961.
- (43) Bhaduri, S.; Sapre, N.Y.; Jones, P.G. Copper Complexes of 1,1'-Di(2 pyridyl)ethyl Methyl Ether: X-Ray Crystal Structures and Reactions with Dioxygen. *J. Chem. Soc., Dalton. Trans.* **1991**, 2539-2543.
- (44) Hoyer, M.; Hartl, H. Iodocuprate(I) mit solvatisierten Kationen: [Li(CH₃CN)₄]⁺[Cu₂I₃]⁻ und [Mg((CH₃)₂CO)₆]²⁺[Cu₂I₄]²⁻. *Z. Anorg. Allg. Chem.* 612 **1992** 45-50.
- (45) Ramaprabhu, S.; Ferretti, R.; Lucken, E.A.C.; Bernardinelli, G. NQR studies of polyhalocuprate(I) anions. *Inorg. Chim. Acta*, 227, **1994**, 153-157.
- (46) Allen, D. W.; Mifflin, J. P. L.; Coles, S. A copper(I)-catalysed template synthesis of solvatochromic aryl-arsonium and -stibonium systems and a synchrotron structural study of a tetraarylstibonium di-iodocuprate. *Chem. Commun.* **1998**, 2115-2116.

- (47) (a) Su, C.-Y. ; Cai, Y.-P. ; Chen, C.-L. ; Lissner, F. ; Kang, B.-S. ; Kaim, W. Self-Assembly of Trigonal-Prismatic Metallocages Encapsulating BF_4^- or CuI_3^{2-} as Anionic Guests: Structures and Mechanism of Formation. *Angew. Chem. Int. Ed.* 41, **2002**, 3371-3375. (b) Su, C.-Y.; Cai, Y.-P. ; Chen, C.-L. ; Smith, M.D.; Kaim, W.; zur Loye, H.-C. Ligand-Directed Molecular Architectures: Self-Assembly of Two-Dimensional Rectangular Metallacycles and Three-Dimensional Trigonal or Tetragonal Prisms. *J. Am. Chem. Soc.* 125, **2003**, 8595-8613.
- (48) Feng, H.; Zhou, X.-P.; Wu, T.; Li, D.; Yin, Y.-G.; Ng, S.W. Hydrothermal synthesis of copper complexes of 40-pyridyl terpyridine: From discrete monomer to zigzag chain polymer. *Inorg. Chim. Acta* 359, **2006**, 4027-4035.
- (49) Kia, R.; Mirkhani, V.; Harkema, S.; van Hummel, G.J. Synthesis and characterization of the 1:1 adducts of copper(I) halides with bidentate N,N'-bis(benzophenone)-1,2-diiminoethane Schiff base: Crystal structures of $[\text{Cu}(\text{bz}_2\text{en})_2][\text{CuX}_2]$ (X = Br, I) complexes. *Inorg. Chim. Acta* 360 **2007** 3369-3375.
- (50) Hu, G.; Holt, E. M. Tris(trimethylphenylammonium) hexaiodotricuprate(I). *Acta Cryst. C*, **1994**, 50, 1576-1578.
- (51) Herres-Pawlis, S.; Haase, R.; Akin, E.; Flörke, U.; Henkel, G. Syntheses and X-ray Structure Analyses of the First Bis(chelated) Copper and Iron Bisguanidine Complexes. *Z. Anorg. Allg. Chem.* **2008** 643 295-298.
- (52) Mishra, S.; Jeanneau, E.; Daniele, S. ; Hubert-Pfalzgraf, L.G. Rare example of a polynuclear heterometallic yttrium(III)-copper(I) iodide cluster with a $[\text{Y}_6(\mu_6\text{-O})(\mu_3\text{-OH})_8]^{8+}$ core structure showing single crystal-to-single crystal transformation. *Cryst. Eng. Comm.* 10 **2008** 814-816.
- (53) Liu, B. ; Pan, S. Liu, B. Chen, W. Di-, Tri-, and Tetranuclear Copper(I) Complexes of Phenanthroline-Linked Dicarbene Ligands. *Inorg. Chem.* 53 **2014** 10485-10497.
- (54) Zink, D. M. ; Grab, T. ; Baumann, T. ; Nieger, M. ; Barnes, E.C. ; Kloppe, W. ; Bräse, S. Experimental and Theoretical Study of Novel Luminescent Di-, Tri-, and Tetranuclear Copper Triazole Complexes. *Organometallics* 30 **2011** 3275-3283.
- (55) Haase, R.; Beschmitt, T.; Flörke, U.; Herres-Pawlis, S. Bidentate guanidine ligands with ethylene spacer in copper-dioxygen chemistry: Structural characterization of bis(μ -hydroxo) dicopper complexes. *Inorg. Chim. Acta* **2011** 374 546-557.
- (56) Chai, W.-X.; Lin, J.; Song, L.; Qin, L.-S.; Shi, H.-S.; Guo, J.-Y.; Shu, K.-Y. Three iodometalate organic-inorganic hybrid materials based on methylene blue cation: Syntheses, structures, properties and DFT calculations. *Solid State Sci.* **2012** 14 1226-1232.
- (57) Haldon, E.; Alvarez, E. ; Nicasio, M.C.; Perez, P.J. Copper(I) Complexes with Trispyrazolylmethane Ligands: Synthesis, Characterization, and Catalytic Activity in Cross-Coupling Reactions. *Inorg. Chem.* **2012** 51 8298-8306.
- (58) Ghorai, D.; Mani, G. Single-Step Substitution of all the α , β -Positions in Pyrrole: Choice of Binuclear versus Multinuclear Complex of the Novel Polydentate Ligand. *Inorg. Chem.* **2014** 53 4117-4129.
- (59) Williams, T.J. ; Bray, J.T.W. ; Lake, B.R.M. ; Willans, C.E. ; Rajabi, N.A. ; Ariafard, A. ; Manzini, C. ; Bellina, F. ; Whitwood, A.C. ; Fairlamb, I.J.S. Mechanistic Elucidation of the Arylation of Non-Spectator N-Heterocyclic Carbenes at Copper Using a Combined Experimental and Computational Approach. *Organometallics* **2015** 34 3497-3507.
- (60) Wang, H.; Zhang, J.; Lin, Z.; Xie, Z. Synthesis and Structural Characterization of Carbene-Stabilized Carborane-Fused Azaborolyl Radical Cation and Dicarbolyl-Fused Azaborole. *Organometallics* **2016** 35 2579-2582.
- (61) Begum, A.; Seewald, O.; Seshadri, T.; Flörke, U.; Henkel, G. Secondary Structures in Inorganic Helicates of an Octadentate Phenanthroline-Type Schiff Base Ligand. *Eur. J. Inorg. Chem.* **2016** 1157-1160.
- (62) Mendoza-Espinosa, D.; Alvarez-Hernández, A.; Angeles-Beltrán, D.; Negrón-Silva, G.E.; Suárez-Castillo, O.R.; Vásquez-Pérez, J.M. Bridged N-Heterocyclic/Mesoionic (NHC/MIC) Heterodicarbenes as Ligands for Transition Metal Complexes. *Inorg. Chem.* 56 **2017** 2092-2099.
- (63) Artem'ev, A. V.; Pritchina, E. A.; Rakhmanova, M. I.; Gritsan, N. P.; Bagryanskaya, I. Y.; Malysheva, S. F.; Belogorlova, N. A. Alkyl-dependent self-assembly of the first red-emitting zwitterionic $\{\text{Cu}_4\text{I}_6\}$ clusters from $[\text{alkyl-P}(2\text{-Py})_3]^+$ salts and CuI : when size matters. *Dalton Trans.* **2019**, 48, 2328.
- (64) Cariati, E. ; Macchi, R. ; Roberto, D. ; Ugo, R. ; Galli, S. ; Masciocchi, N. ; Sironi, A. Sequential Self-Organization of Silver(I) Layered Materials with Strong SHG by J Aggregation and Intercalation of Organic Nonlinear Optical Chromophores through Mechanochemical Synthesis. *Chem. Mater.* 19, **2007**, 3704-3711.

- (65) Qin, X.; Zhang, G.; Gao, Y.; Liu, H.; Du, C.; Liu, Z. Metallo-supramolecular grid-type architectures for highly and selectively efficient adsorption of dyes in water. *RSC Advances*, **2015**, 5, 43334–43337.
- (66) Bowmaker, G. A.; Bruce, M. I.; Skelton, B. W.; Somers, N.; White, A. H.; Structural and Spectroscopic Studies of Halocuprate(I) and Haloargentate(I) Complexes $[M_2X_nX'_{4-n}]^{2-}$. *Z. Anorg. Allg. Chem.* **2007**, 633, 1024-1030.
- (67) Hartl, H.; Brüdgam, I.; Mahdjour-Hassan-Abadi, F. Synthese und Strukturuntersuchungen von Iodocupraten(I) VI. Iodocuprate(I) mit zweikernigen Anionen $[Cu_2I_4]^{2-}$ und $[Cu_2I_6]^{4-}$. *Z. Naturforsch.* **1985**, 40b, 1032-1039.
- (68) Pfitzner, A.; Schmitz, D. Two New Modifications of $[P(C_6H_5)_4]_2[Cu_2I_4]$. *Z. Anorg. Allg. Chem.* **1997**, 623, 1555-1560.
- (69) (a) Bernstein, J. *Polymorphism in Molecular Crystals*, Clarendon Press Oxford, **2002**. (b) Khorloo, M.; Cheng, Y.; Zhang, H.; Chen, M.; Sung, H. H. Y.; Williams, I. D.; Lam, J. W. Y.; Tang, B. Z. Polymorph selectivity of an AIE luminogen under nano-confinement to visualize polymer microstructures. *Chem. Sci.*, **2020**, 11, 997–1005. (c) Mutai, T.; Satou, H.; Araki, K. Reproducible on–off switching of solid-state luminescence by controlling molecular packing through heat-mode interconversion. *Nat. Mater.* **2005**, 4, 685-687. (d) Ito, H.; Muromoto, M.; Kurenuma, S.; Ishizaka, S.; Kitamura, N.; Sato, H.; Seki, T. Mechanical stimulation and solid seeding trigger single-crystal-to-single-crystal molecular domino transformations. *Nat. Commun.*, **2013**, 4, 2009-2014. (e) Malwitz, M. A.; Lim, S. H.; White-Morris, R. L.; Pham, D. M.; Olmstead, M. M.; Balch, A. L. Crystallization and Interconversions of Vapor-Sensitive, Luminescent Polymorphs of $[(C_6H_{11}NC)_2AuI](AsF_6)$ and $[(C_6H_{11}NC)_2AuI](PF_6)$. *J. Am. Chem. Soc.* **2012**, 134, 10885-10893. (f) Krishnan, B. P.; Sureshan, K. M. A Spontaneous Single-Crystal-to-Single-Crystal Polymorphic Transition Involving Major Packing Changes. *J. Am. Chem. Soc.* **2015**, 137, 1692–1696.
- (70) Bondi, A. van der Waals volumes and radii. *J. Phys. Chem.* **1964**, 68, 441-451.
- (71) Sculfort, S.; Braunstein, P. Intramolecular d^{10} - d^{10} interactions in heterometallic clusters of the transition metals. *Chem. Soc. Rev.* **2011**, 40, 2741-2760.
- (72) Benito, Q.; Baptiste, B.; Polian, A.; Delbes, L.; Martinelli, L.; Gacoin, T.; Boilot, J.-P.; Perruchas, S. Pressure Control of Cuprophilic Interactions in a Luminescent Mechanochromic Copper Cluster. *Inorg. Chem.* **2015**, 54, 9821-9825.
- (73) Licari, D.; Baiardi, A.; Biczysko, M.; Egidi, F.; Latouche, C.; Barone, V.; Implementation of a graphical user interface for the virtual multifrequency spectrometer: The VMS-Draw tool, *Journal of Computational Chemistry*, **2015**, 36, 321-334.
- (74) Tang, J. A.; Ellis, B. D.; Warren, T. H.; Hanna, J. V.; Macdonald, C. L. B.; Schurko, R. W. Solid-State ^{63}Cu and ^{65}Cu NMR Spectroscopy of Inorganic and Organometallic Copper(I) Complexes. *J. Am. Chem. Soc.* **2007**, 129, 13049-13065.
- (75) Benito, Q.; Maurin, I.; Poggi, M.; Martineau-Corcos, C.; Gacoin, T.; Boilot, J.-B.; Perruchas, S. Impact of crystalline packing on the mechanochromic luminescence properties of copper based compounds: towards functional coatings. *J. Mater. Chem. C*, **2016**, 4, 11231-11237.
- (76) Benito, Q.; Le Goff, X.; Nocton, G.; Fargues, A.; Garcia, A.; Berhault, A.; Kahlal, S.; Saillard, J.-Y.; Martineau, C.; Trebosc, J.; Gacoin, T.; Boilot, J.-P.; Perruchas, S. Geometry Flexibility of Copper Iodide Clusters: Variability in Luminescence Thermochromism. *Inorg. Chem.* **2015**, 54, 4483-4494.
- (77) (a) Balch, A. L. Dynamic Crystals: Visually Detected Mechanochemical Changes in the Luminescence of Gold and Other Transition-Metal Complexes. *Angew. Chem. Int. Ed.* **2009**, 48, 2641-2644. (b) Chi, Z.; Zhang, X.; Xu, B.; Zhou, X.; Ma, C.; Zhang, Y.; Liu, S.; Xu, J. Recent advances in organic mechanofluorochromic materials. *Chem. Soc. Rev.* **2012**, 41, 3878-3896. (c) Zhang, X.; Chi, Z.; Zhang, Y.; Liu, S.; Xu, J. Recent advances in mechanochromic luminescent metal complexes. *J. Mater. Chem. C*, **2013**, 1, 3376-3390.
- (78) Chen, G.; Zhou, J.; Feng, H.; Feng, F.; Xu, P.; Pan, S.; Xu, J.; Qian, Z. A simple and efficient phosphorescent probe for iodide-specific detection based on crystallization induced phosphorescence of organic ionic crystals. *J. Mater. Chem. C*, **2019**, 7, 43-47.
- (79) Gaussian 16, Revision A.03, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E.

Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman, and D. J. Fox, Gaussian, Inc., Wallingford CT, **2016**. Gaussian, Inc.

(80) (a) Perdew, J. P.; Ernzerhof, M.; Burke, K. M. Rationale for Mixing Exact Exchange with Density Functional Approximations. *J. Chem. Phys.* **1996**, *105*, 9982-9985. (b) Perdew, J. P.; Burke, K.; Ernzerhof, M. Generalized Gradient Approximation Made Simple. *Phys. Rev. Lett.* **1996**, *77*, 3865-3868. (c) Perdew, J. P.; Burke, K.; Ernzerhof, M. Erratum: Generalized Gradient Approximation Made Simple. *Phys. Rev. Lett.* **1997**, *78*, 1396-1396.

(81) (a) Dunning Jr., T. H.; Hay, P. J. *Methods of Electronic Structure Theory*, H. F. Schaeffer ed., Plenum Press, New York, **1977**. (b) Hay, P. J.; Wadt, W. R. Ab Initio Effective Core Potentials for Molecular Calculations. Potentials for the Transition Metal atoms Sc to Hg. *J. Chem. Phys.* **1985**, *82*, 270-283. (c) Hay, P. J.; Wadt, W. R. Ab Initio Effective Core Potentials for Molecular Calculations. Potentials for the Main Group Elements Na to Bi. *J. Chem. Phys.* **1985**, *82*, 284-298. (d) Hay, P. J.; Wadt, W. R. Ab Initio Effective Core Potentials for Molecular Calculations. Potentials for K to Au Including the Outermost Core Orbitals. *J. Chem. Phys.* **1985**, *82*, 299-310.

(82) Barone, V.; Baiardi, A.; Bloino, J.; New Developments of a Multifrequency Virtual Spectrometer: Stereo-Electronic, Dynamical, and Environmental Effects on Chiroptical Spectra, *Chirality*, **2014**, *26*, 588-600.

(83) Barone, V.; Baiardi, A.; Biczysko, M.; Bloino, J.; Cappellia, C.; Lipparini, P.; Implementation and validation of a multi-purpose virtual spectrometer for large systems in complex environments, *Phys. Chem. Chem. Phys.*, **2012**, *14*, 12404-12422.

(84) Massuyeau, F.; Faulques, E.; Latouche, C.; New Insights To Simulate the Luminescence Properties of Pt(II) Complexes Using Quantum Calculations, *J. Chem. Theory Comput.* **2017**, *13*, 4, 1748-1755.

(85) Li, J.; Wang, J.; Li, H.; Song, N.; Wang, D.; Tang, B. Z. Supramolecular materials based on AIE luminogens (AIEgens): construction and applications. DOI: 10.1039/c9cs00495e