Observation of Contrary Thermo-responsive Trend for Single Crystal and Powder Samples in Mechano-, Thermo- and Solvato-responsive Luminescent Cubane [Ag₄I₄L₄] Cluster

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A new silver(I) iodide cluster [Ag₄I₄(TMP)₄] (TMP = tris(3-methylphenyl)-phosphine) shows triply stimuli-responsive luminescent chromism, namely, mechano-, thermo- and solvent-sensitive chromism, which is isostructural to our previously reported [Cu₄I₄(TMP)₄] but shows quite different luminescence in response to the external stimuli. Especially, during the mechanical grinding, the relative intensities of HE and LE of 1 varied with a concomitant hypsochromic shift, and when the temperature was decreased from 300 to 5 K, unprecedented contrary thermo-responsive trend for single crystal and powdered samples (blueshift of single crystals and redshift of powdered samples) was observed. These distinct characters of 1 should be due to the different molecular packing modes, metallic interactions and the unique character of Ag(I) ion.

Phosphorescent coordination materials have received increasing attention due to their fascinating functionalities and widespread applications in lighting, sensing, memories and display devices, which provides a great motivation for research on coinage metal emissive complexes. As an important member of these materials, coinage metal emissive complexes, such as coinage metal-halide aggregates, exhibit modification of their luminescent color upon exposure to the external stimuli (for example, temperature, mechanical grinding, light, pH, solvent vapor, etc). Such functional compounds reported so far are mainly focused on copper-halide-based complexes due to their chromic luminescent properties which are generally related to the flexible geometries. The majority of known copper-halide-based complexes display stimuli-responsive chromic luminescent behaviors; however, the emission shifts of these compounds are unpredictable, and the origin of stimuli-responsive chromisms, particularly mechanochromism, remains ambiguous. Stimuli-responsive chromisms, except thermochromism, of copper halide complexes have not been well investigated yet because structural changes in response to the external stimuli cannot be comprehensively determined through single-crystal X-ray crystallography or other experimental methods. Especially, during mechanical grinding, the crystal state of copper cluster complexes is generally converted into the amorphous state, which is difficult to characterize through single-crystal X-ray diffraction analysis. Scholars have proposed different hypothetical interpretation of the origin of various chromisms, but no general rule has been established. Studies on the number of stimuli-sensitive luminescent complexes still remain in the infancy stage; thus, novel stimuli-responsive chromic luminescent complexes must be developed to comprehensively investigate chromic properties and provide new insights into their origins.

Similar to the Cu(I) ions of the stimuli-responsive copper(I) halide clusters, Ag(I) ions possess the tetrahedral coordination geometry and exhibit metallophilic interactions; hence, emissive silver(I) halide compounds are potential stimuli-sensitive materials. In addition, the coordinated geometry of Ag(I) ion can change from a...
tetrahedral geometry to a square-planar geometry under an external stimulus; this property is similar to that of copper-halide clusters and can induce the striking tunable luminescence. Nevertheless, some differences exist between the copper(I) halide and silver(I) halide compounds. (1) Ag(I) ion is more inert to oxidation than Cu(I), often resulting in an absence of the low-energy MLCT state and reduced complexity of the origin of stimuli-responsive luminescence; (2) In contrast to Cu(I) complexes, the geometrical arrangements and local environment around Ag(I) ions drastically affect luminescence. Considering these similarities and differences, scholars have focused on stimuli-responsive luminescence of emissive silver(I)-halide materials. Nevertheless, research on emissive silver(I) halides is extremely limited because of their potential photosensitivity and low luminescent efficiency. In particular, the luminescent properties of tetranuclear Ag(I) clusters formulated [AgI,L] (L = organic ligand) have been rarely investigated; however, a number of isostructural tetracopper(I) complexes have been intensively explored for their luminescent stimuli-responsive properties. Further studies on multinuclear silver(I)-halide materials must be conducted because of their interesting emission properties and distinct emissive mechanisms. Based on above considerations, we aim to design and synthesize new silver(I) cluster-based compounds with stimuli-responsive luminescence by using hydro/solvothermal synthesis method. Herein, we develop novel cubane [Ag4I4(TMP)4] complex (1) with weak Ag-Ag interaction; this complex is isostructural to the compound [Cu4I4(TMP)4] 2 in the absence of Cu-Cu interaction, as previously reported by our team. However, compound 1 exhibits quite different luminescence in response to external stimuli. In particular, an unprecedented contrary thermo-responsive trend was observed for single crystal and powder samples (blueshift of single crystals and redshift of powdered samples with decreasing temperature). Compound 1 displayed distinct chromic luminescence probably because of its closed packing mode, presence of metallic interactions, and unique characteristic of Ag(I) ions. These results contribute to understanding the effects of the metallicophilic interactions, molecular structure, and packing mode for stimuli-responsive luminescence.

Results

Description of Structures. X-ray diffraction data reveals that compound 1 contains cubane [Ag4I4] clusters with four silver ions and four iodide ions alternatively occupying the corners of a distorted cube. Compound 1 with chiral C3-symmetric cubane structure crystallizes in cubic space group I-43d. The asymmetric unit contains one third of a cubic [Ag4I4] core located on the crystallographic threefold axis; in the axis, two crystallographically independent Ag atoms adopt a distorted AgIP4 tetrahedral geometry and are coordinated three μ3-iodide ions and one phosphorus donors from the TMP ligand (Fig. 1). The Ag-P bond lengths of 2.460(3) and 2.450(4) Å at 293 K are comparable with those of the related clusters. The Ag-I bond distances are within the range of 2.8782(11) and 2.9311(9) Å, and the I-Ag-I bond angles range from 96.83(3) and 107.74(3) °. The Ag-Ag distances within the range of 3.3762(15) and 3.3764(15) Å at 293 K are shorter than the sum of the van der Waals radii (3.44 Å), indicating the presence of argentophilic interaction in compound 1. The intermolecular Ag···Ag distance is absent at room temperature. Intermolecular offset π−π interactions exist between two phenyl rings of neighboring [Ag4I4(TMP)4] clusters. The centroid-to-centroid distance between two closest benzene rings is 3.900(8) Å and the corresponding dihedral angle is 2.82 ° (Table S2 and Figure S1).

Photoluminescence Properties. The photoluminescent spectra of 1 were measured in depth and reveal triply 3-symmetric cubane structure crystallizes in cubic space group I-43d. The asymmetric unit contains one third of a cubic [Ag4I4] core located on the crystallographic threefold axis; in the axis, two crystallographically independent Ag atoms adopt a distorted AgIP4 tetrahedral geometry and are coordinated three μ3-iodide ions and one phosphorus donors from the TMP ligand (Fig. 1). The Ag-P bond lengths of 2.460(3) and 2.450(4) Å at 293 K are comparable with those of the related clusters. The Ag-I bond distances are within the range of 2.8782(11) and 2.9311(9) Å, and the I-Ag-I bond angles range from 96.83(3) and 107.74(3) °. The Ag-Ag distances within the range of 3.3762(15) and 3.3764(15) Å at 293 K are shorter than the sum of the van der Waals radii (3.44 Å), indicating the presence of argentophilic interaction in compound 1. The intermolecular Ag···Ag distance is absent at room temperature. Intermolecular offset π−π interactions exist between two phenyl rings of neighboring [Ag4I4(TMP)4] clusters. The centroid-to-centroid distance between two closest benzene rings is 3.900(8) Å and the corresponding dihedral angle is 2.82 ° (Table S2 and Figure S1).

A detailed photophysical study of the single-crystal 1 shows a weak high-energy emission (HE) centered at 448 nm (τ = 0.8 μs) and a strong low-energy emission (LE) centered at 476 nm (τ = 1.3 μs) under excitation at 337 nm. With carefully reviewing the relative literatures, the HE and LE emission bands can be generally clarified by our team. However, compound 1 exhibits quite different luminescence in response to external stimuli. In particular, an unprecedented contrary thermo-responsive trend was observed for single crystal and powder samples (blueshift of single crystals and redshift of powdered samples with decreasing temperature). Compound 1 displayed distinct chromic luminescence probably because of its closed packing mode, presence of metallic interactions, and unique characteristic of Ag(I) ions. These results contribute to understanding the effects of the metallicophilic interactions, molecular structure, and packing mode for stimuli-responsive luminescence.
grinding. Upon dropping ethanol to the powder sample and subsequent evaporating of ethanol, emission color slowly returned to the original blue-green.

The thermo-responsive luminescent chromism of the solid-state compound 1 was also studied in detail. A unique contrary thermo-responsive trend was found for both single and powdered sample. After immersion in liquid nitrogen for a few minutes, both unground and ground samples were collected and exposed to the UV light. The reversible thermochromic luminescence for both samples can be easily distinguished by the naked eye (Figs 3 and 4). For single-crystal sample, the LE emission band progressively blue-shifted to 466 nm and the HE emission band slightly shifted to 440 nm with decreasing the temperature to 5 K. Such blue-shifted emission observed with decreasing temperature is very rare in coinage-metal thermochromic luminescent coordination compounds. This unusual phenomenon is very complicated and possibly related to variations in synergistic π···π and Ag···Ag interactions with decreasing the temperature. The intensities of both LE and HE emissions gradually increased with decreasing temperature, but the intensity of HE band increased faster than that of LE band. When the unground single-crystal sample was cooled down to 5 K, the intensity of the HE band is almost equal to that of the LE band (Fig. 3). The measured lifetimes were increased drastically by approximately 100 times from 0.8 μs at 300 K to 97.4 μs at 5 K for HE band and from 1.3 μs at 300 K to 90.5 μs at 5 K for LE band, respectively (Table S3 and Figures S5–13). The emission lifetimes significantly increased with decreasing the temperature. This finding could be due to the relation of decreased non-radiative rate to the high spin-orbit coupling of Ag(I) ions.
Figure 2. The luminescence spectra of compound 1 at room temperature in different states.

Figure 3. Temperature-dependent emission spectra of single crystal 1 from 5 to 300 K. Inset: Images for single crystals under ambient light at room temperature (left), under 365 nm UV lamp irradiation at room temperature (middle) and in liquid nitrogen (77K) (right).
and fast intersystem crossing. This behavior is also observed in the emissive copper(I) and silver(I) complexes with the thermally activated delayed fluorescence.

The thermochromic luminescence properties of compound 1 were also affected by the mechanical grinding. After crushing, the luminescence of the powdered sample 1 exhibited unprecedented contrasting thermochromic luminescence trend from that of single-crystal 1. With decreasing temperature, the HE emission band of the powder sample progressively increased and red shifted to 435 nm with a concomitant increase and slight red shift of the LE band, which is contrary to the blue-shifted emission observed in the single crystal sample. To be noted, two different changing steps in the thermochromic luminescent process of powder sample were observed in the solid-state emission spectra (Figure S14 and S15). Initially, the intensity of LE emission band increased faster than that of HE emission band from room temperature down to 150 K until the intensities of the LE and HE bands are almost equal. While with continuous decrease in temperature to 5 K, the intensity of the HE emission band increased faster; at 5 K, the HE intensity is much higher than LE intensity (Fig. 4). When the temperature was lowered to 5 K, the lifetimes largely increased by hundreds of times from 0.4 μs at 300 K to 168.5 μs at 5 K for HE band and from 0.6 μs at 300 K to 175.6 μs at 5 K for LE band, respectively (Table S4 and Figure S16–24). It can be seen that the luminescent lifetimes of powder samples increased much faster than that of single crystal sample.

Solvato-stimuli experiments were also carried out. The crushed powder sample of 1 was immersed into six different organic solvents, ultrasonicated, and then aged for 48 hours to form the stable emulsion solutions. The luminescent spectra in most solvents such as EtOH, BuOH, DMF, acetone, and cyclohexane displayed obvious changes. The luminescent emission bands became nearly the same as that of unground sample, indicating that these solvents have effect to mediate the recovery of powder sample to original structural arrangement. While in CH₃CN solvent, the emission bands showed no changes (Figure S25). In addition, we carefully performed vapor-stimuli experiments. When the colorless block crystals of 1 were exposed to CH₃CN, CH₂Cl₂, C₂H₅OH, BuOH and DMF vapors respectively, the luminescent color and emission bands remained unchanged, indicating none vapochromism.

Discussion

The developed new cubic silver(I)-iodide cluster exhibits mechano-, thermo- and solvato-responsive luminescent chromism. Although this compound is isostructural to our reported copper(I)-halide compound, their multistimuli-responsive luminescence differ upon exposure to the external stimuli. In particular, after mechanical grinding, the relative intensities of HE and LE of 1 varied with a concomitant hypsochromic shift, when the temperature was decreased from 300 to 5 K, an unprecedented contrary thermo-responsive trend was observed for single crystal and powder samples (blueshift of single crystals and redshift of powdered samples with decrease of temperature). The chromic luminescence of 1 is considerably different from those of 2 and other chromic luminescent cubic Cu₄I₄ clusters. These distinct characters of 1 might be due to the different molecular packing modes, metallic interactions and the unique character of Ag(I) ion.

Methods

Testing Methods. All reagents and solvents were obtained commercially and used without further purification. The FT-IR spectra were recorded from KBr pellets in range 400–4000 cm⁻¹ on a Perkin-Elmer Spectrum BX FT-IR spectrometer. Elemental analysis was performed on a Vario EL-II elemental analyzer. XRPD data were recorded in a Bruker D8 ADVANCE X-ray powder diffractometer (CuKα, λ = 1.5418 Å).
Steady-state photoluminescence spectra and lifetime measurements were measured from 300 K down to 5 K by a single-photon counting spectrometer using an Edinburgh FLS920 spectrometer equipped with a continuous Xe-900 xenon lamp, and a μF900 microsecond flash lamp. The corrections of excitation and emission for the detector response were performed ranging from 200 to 900 nm. Temperature-dependent measurements were carried out in a JANIS SHI-4S-1 cold head cooled with HC-4A compression engine. The data were analyzed by iterative convolution of the luminescence decay profile with the instrument response function using the software package provided by Edinburgh Instruments. Lifetime data were fitted with triple-exponential-decay functions. The quantum yields were measured by use of an integrating sphere with Edinburgh Instrument FLS920 spectrometer.

**Synthesis.** Synthesis of Ag$_4$I$_4$(TMP)$_4$(1). A mixture of AgI (0.5 mmol, 0.152 g), KI (1 mmol, 0.165 g), TMP (0.2 mmol, 0.059 g) and CH$_3$CN solvent (5 mL) was stirred and then sealed in a 25-ml Teflon-lined stainless container and heated to 140 °C for 7 days. With a cooling rate of 5 °C min$^{-1}$ to room temperature, clourless block crystals of 1 in 56% yield were recovered. Anal. Calc. for C$_{84}$H$_{87}$Ag$_4$I$_4$P$_4$: C, 46.72; H, 4.06; N, 9.47. Found: C, 46.65; H, 4.15; N, 9.53. IR(KBr, cm$^{-1}$): 3440 s, 3118 s, 2906w, 2361w, 1603m, 1481 m, 1403 m, 1280w, 1102 s, 980w, 890w, 780 m, 679 m, 612w, 545w, 456w.

**Single Crystal X-ray Crystallography.** X-ray single-crystal diffraction data for 1 at 293 K were performed with Mo-Kα radiation ($\lambda = 0.71073$ Å) on a Bruker Apex CCD diffractometer at 298(2) K. The program SAINt was used for integration of diffraction profiles, and the program SADABS was used for absorption correction. The structure was solved with XS structure solution program by direct method and refined by full-matrix least-squares technique using Olex2. All non-hydrogen atoms were refined with anisotropic thermal parameters. Hydrogen atoms of organic ligands were generated theoretically onto the specific carbon atoms, and refined isotropically with fixed thermal factors. Further details regarding the structural analysis are summarized in Table 1 and selected bond lengths and bond angles are shown in Table S1.

**Solvatochromism Experiment section.** Ground powder of 1 was immersed into various pure organic solvents such as DMF, acetone, cyclohexane, EtOH, MeCN, n-Butanol. After carried out 20 min ultrasound, the corresponding suspensions were obtained for measurements.

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Author Contributions
Shi-Li Li, Min Han, Bin Wu, Jie Wang carried out the experiment and data extraction. Shi-Li Li finished the manuscript. Fu-Qing Zhang and Xian-Ming Zhang were involved in data analysis. Xian-Ming Zhang designed the study and revised the manuscript. All authors reviewed the manuscript.

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