INORGANIC CHEMISTRY







FRONTIERS

RESEARCH ARTICLE

View Article Online
View Journal | View Issue



Cite this: *Inorg. Chem. Front.*, 2024, **11**, 5536

Reversible "on–off" conversion and ultra-high temperature sensitivity of a zero-dimensional lead-free $Cs_2InBr_5(H_2O):Sb^{3+}$ perovskite†

Maohao Yang,^a Wanyin Ge, (1) *a Kenshi Matsumoto,^b Masaki Saruyama, (1) b Ryota Sato, (1) h Haruka Takekuma, (1) b Ryo Takahata (1) and Toshiharu Teranishi (1) b

Zero-dimensional (0D) lead-free perovskites have garnered significant attention due to their unique optoelectronic properties and non-toxicity. However, the single response to stimuli in lead-free perovskites limits the versatility of multifunctional compounds. In this study, we abandoned the toxic bromic acid and utilized water as the reaction medium, to achieve an environmentally friendly green chemical route. We successfully prepared a zero-dimensional lead-free halide $Cs_2In_{1-x}Sb_xBr_5(H_2O)$ perovskite with a fractal structure using an *in situ* solution crystallization method. Herein, we discovered that the recrystallization occurred beneath the smooth-grown rhombic crystal surface, forming fractal branched crystals. Interestingly, reversible luminescence color transitions from orange-red to yellow and return to the initial state were achieved in response to the individual stimuli of temperature and humidity. We found that H_2O molecules played a crucial role in the color tuning, enabling reversible "on–off" switching modes with rapid modulation rates. Moreover, the OD lead-free halide perovskite $Cs_2InBr_5(H_2O):Sb^{3+}$ also exhibited excellent temperature sensitivity, with the relative sensitivity (S_R) reaching up to 9.39% K^{-1} . This study provides valuable insights for the further development of halide perovskites in design and application fields, laying a foundation for the development of multifunctional smart materials and temperature sensing applications.

Received 10th May 2024, Accepted 1st July 2024 DOI: 10.1039/d4qi01164c rsc.li/frontiers-inorganic

1. Introduction

Recently, lead halide perovskites have shown excellent optoelectronic properties, with growing applications in various fields. 1-8 However, the toxicity of the lead element severely limits the practical application. 9-15 Consequently, the global academic community is deeply engaged in the development of lead-free perovskite compounds. Various synthetic methods have been developed to obtain a diverse array of morphologies (e.g., nanocrystals, single-crystals, thin films, and powders) of lead-free halide perovskites. 16-22 Lead-free perovskite materials have gained widespread attention due to their superior optical properties and environmentally friendly nature. They are widely used in areas such as solar cells, photodetectors, lightemitting diodes, and lasers, bringing new development opportunities for modern optoelectronic technology. 23-26 However, the innovative synthesis route and optical characteristics of the zero-dimensional (0D) indium-based halide perovskite Cs₂InBr₅(H₂O), as well as its intrinsic physicochemical properties, still require further investigation.²⁷

Currently, the incorporation of dopants is recognized as an effective strategy for improving the luminescence efficiency of perovskites.²⁸ Substantial advancements have been achieved in the study of the photoluminescence (PL) properties of indiumbased halide perovskites doped with antimony (Sb3+). For instance, Sb3+-doping in 0D Cs2SnCl6 exhibits yellow-red emission.^{29,30} Compared to the undoped counterparts, the photoluminescence quantum yield (PLQY) of Sb3+-doped $A_2InCl_5(H_2O)$ (A = Rb, Cs) has significantly improved, increasing from below 2% to a range of 85-95%.31 In the family of $Cs_2InX_5(H_2O)$ (X = Cl⁻, Br⁻ and I⁻), the octahedral units exhibit pronounced electron-phonon coupling, resulting in a substantial Stokes shift observed in these 0D perovskite compounds.³² Substitution of the halide from Cl⁻ to Br⁻ and I⁻ also leads to redshifts in the photoluminescence (PL) and photoluminescence excitation (PLE) spectra of Sb3+-doped Cs₂InX₅(H₂O) compounds.³³ Additionally, applying high pressure enables precise modulation of the PL properties of Cs₂InBr₅(H₂O), culminating in the observation of anomalous anti-Stokes emission.34 In Cs2InBr5(H2O), photogenerated charge carriers are spatially confined within isolated octahedra, enhancing the radiative recombination probability.35-38 A-site cations have a slight effect on PL, 39 while water mole-

^aSchool of Materials Science and Engineering, Shaanxi University of Science and Technology, Xi'an, Shaanxi 710021, P. R. China. E-mail: gewanyin@sust.edu.cn ^bInstitute for Chemical Research, Kyoto University, Gokasho, Uji, Kyoto 611-0011, Japan

[†]Electronic supplementary information (ESI) available. See DOI: https://doi.org/ 10.1039/d4qi01164c

cules, through their induced distortion of In3+ octahedra, result in a broad emission and exert a more pronounced influence on the self-trapped excitons (STEs).40

Halide perovskites are well recognized for their high sensitivity to environmental conditions, including air composition (humidity, oxygen), temperature, light exposure, and electric fields, which can lead to rapid degradation and limit their practical applications. 41-46 Fortunately, the novel lead-free 0D halide perovskite Cs2InBr5(H2O) stands out for its exceptional stability and tunable crystal structure. Interestingly, its isolated octahedral structure incorporates a water molecule (H2O). As the temperature increases, the water molecule is released, modulating its luminescence from red to yellow. 47,48 Furthermore, when the dehydrated octahedral structure is subjected to an atmosphere with 80% relative humidity (RH), the PL wavelength and intensity undergo gradual alterations, with the color transitioning reversibly from yellow back to red.49

High-performance functional compounds that are reversible and responsive to stimuli such as temperature or humidity are highly desirable for sophisticated anti-counterfeiting technologies. However, many existing anti-counterfeiting functional compounds exhibit limited temperature responsiveness and necessitate further development to enhance their reversibility. In this study, we report the successful synthesis of a 0D halide perovskite, Cs₂In_{1-x}Sb_xBr₅(H₂O), utilizing water as the reaction medium, thus abandoning the common chemical route involving bromic acid reported in the literature and demonstrating an environmentally friendly and green chemical approach. Furthermore, the diverse luminescence properties of Cs₂InBr₅(H₂O):Sb³⁺ are of significant importance. We observed that the water molecule in the Cs₂InBr₅(H₂O): Sb³⁺ molecular structure acts as a switch, facilitating an "onoff-on" cycle that can be repeated multiple times. We conducted in-depth analysis of the reversible structural and photoluminescence transitions triggered by water molecules or thermal effects. This halide perovskite, with its reversible photoluminescence properties, offers a fresh perspective, overcoming the limitations of traditional materials that exhibit singular responses. Additionally, we assessed the absolute sensitivity (S_A) and relative sensitivity (S_R) during temperature variations, with S_A achieving a maximum value of 3.55% K⁻¹ and $S_{\rm R}$ reaching a peak of 9.39% K⁻¹, indicating extremely high sensitivity. These research findings demonstrate the high sensitivity and responsiveness of Cs₂InBr₅(H₂O):Sb³⁺ to temperature changes within a certain temperature range, providing strong support for its applications in temperature- and humidity-sensitive fields. The reversibility of water molecules as switches and their multiple cycling capabilities presented in this study bring new opportunities for the development of lead-free 0D halide perovskites. This material, with its reversible photoluminescence characteristics, holds promise as a crucial component in anti-counterfeiting technology and multifunctional smart materials, while also exhibiting significant potential in high-performance temperature sensing applications.

Results and discussion

During the synthesis process, as depicted in Fig. 1a, distilled water was utilized as the solvent to formulate a transparent precursor solution by reacting InBr₃ (1 mmol, 99%, Sigma-Aldrich), CsBr (2 mmol, 99.9%, Sigma-Aldrich), and an appropriate amount of SbBr3 (99.9%, Wako Pure Chemical Corporation), without using bromic acid, to achieve a green synthetic route. The scanning electron microscopy (SEM) image in Fig. 1b illustrates the morphological features of the synthesis reaction at 150 °C for 2 min, clearly displaying granular grains with sizes around 20 µm. An individual microcrystal in Fig. 1c exhibits its energy dispersive X-ray spectroscopy (EDX) features. Hydrogen (H), the lightest element, is typically undetectable by this method. The EDX spectrum confirms the homogeneous distribution of Cs, In, Br, and O elements and indicates elemental molar ratios approximating the stoichiometry of Cs2InBr5(H2O), thereby validating the successful synthesis of the Cs₂InBr₅(H₂O) compound.

Figure 1d demonstrates crystallization near room temperature (around 30 °C), where the evaporative concentration of the solution droplets on glass slides leads to the emergence of dendritic crystals at the solution-glass interface. Subsequently, diamond-shaped, rectangular, and blade-like crystals nucleate and grow on the solution surface, eventually aggregating and overlaying to form butterfly-petal-shaped crystals. Notably, as the solution evaporates, leaving a partial liquid beneath the petal-shaped crystals, the solute concentration increases, altering the crystal growth conditions and kinetics, resulting in the growth of fractal dendritic crystals beneath the crystal surface, consistent with the recent literature.⁵⁰ The resulting fractal structure consists of an upper-layer butterfly petal-shaped crystal and a lower-layer dendritic crystal (Fig. 1e). ESI Fig. S1b† exhibits fractal structures obtained via the in situ solution method at different temperatures. The increasing temperature correlates with a decrease in the grain size and a corresponding increase in the quantity. This trend is attributed to the accelerated evaporation rate of water molecules at elevated temperatures, which in turn shortens the duration of crystal growth. Fig. 1f displays the EDX mapping characterizing the elemental composition and distribution of petal-shaped fractal structures, with the expected elements Cs, In, Sb, Br, and O. In Fig. 1g, the elemental molar ratio Cs:In:Br = 2:1:6, implying bromine ions exceeding the stoichiometric value, is potentially attributable to defects within the fractal structure, which could account for the slight deviations or fluctuations in the chemical stoichiometry of the crystal structure.

To ascertain the 0D halide perovskite structure of the asprepared compound, we conducted X-ray diffraction (XRD) analysis. Subsequently, the high-precision XRD data were refined to validate the compound's phase and crystallographic characteristics. The refinement results, as presented in Fig. 2a, perfectly matched the synthesized sample's XRD pattern with the PDF card (01-087-4184) for Cs₂InBr₅(H₂O), with an R_{wp} value of 6.44% and a goodness-of-fit factor of 1.52, signifying a high degree of fitting accuracy. The crystallographic paraResearch Article

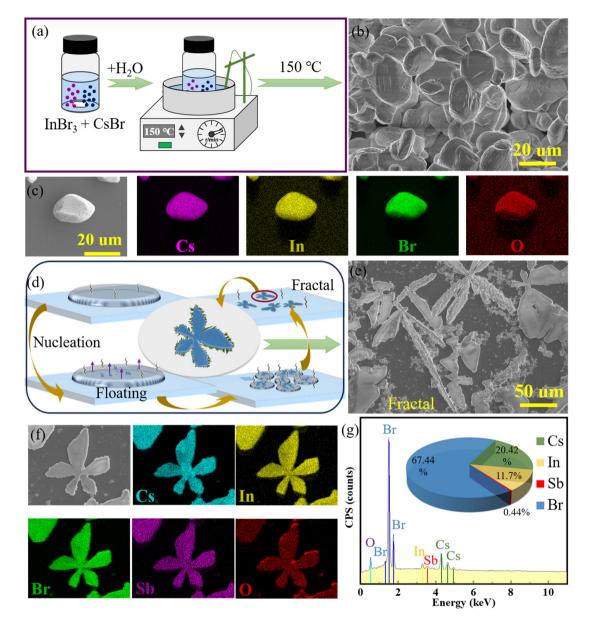


Fig. 1 (a) Preparation process of Cs₂InBr₅(H₂O); (b) SEM image of Cs₂InBr₅(H₂O) prepared at 150 °C; (c) EDX mapping of Cs, In, Br, and O elements; (d) fractal structure of Cs₂In_{1-x}Sb_xBr₅(H₂O) samples obtained via the in situ solution crystallization method; (e) SEM image of the fractal structure post-crystallization; the EDX mapping (f) and EDX spectrum (g) of Cs₂InBr₅(H₂O):Sb.

meters derived were a = 14.8763 Å, b = 10.8116 Å, and c =7.6437 Å, consistent with the pnma space group. Additionally, we observed a color change in the luminescence of Cs₂InBr₅(H₂O):Sb³⁺ under UV light after annealing at different temperatures. To further investigate this phenomenon, temperature-dependent XRD tests were conducted. As depicted in Fig. 2b, significant structural differences were evident in the samples annealed at different annealing temperatures. On increasing the temperature, the intensity of the diffraction peaks of Cs₂InBr₅(H₂O) gradually decreased, accompanied by the appearance of diffraction peaks corresponding to the Cs₃In₂Br₉ phase. This fact indicates that with increasing temperature, water molecules are progressively released from

Cs₂InBr₅(H₂O), leading to the partial transformation of Cs₂InBr₅(H₂O) into Cs₃In₂Br₉. Notably, this process is reversible, as the structure of the sample can be regenerated upon exposure to a humid environment.

The octahedral structure in the Cs₂InBr₅(H₂O):Sb³⁺ perovskite is composed of one indium (In) atom, five bromine (Br) atoms, and a single H₂O molecule, with indium atoms being partially substituted by Sb atoms, as illustrated in the crystal structure in Fig. 2c. Two Cs⁺ ions are spatially separated, forming a 0D configuration. Upon exposure to thermal annealing, dehydration of the octahedral units occurs, prompting the reorganization of adjacent In-Br bonds and the formation of Cs₃In₂Br₉. Interestingly, the reverse transformation from

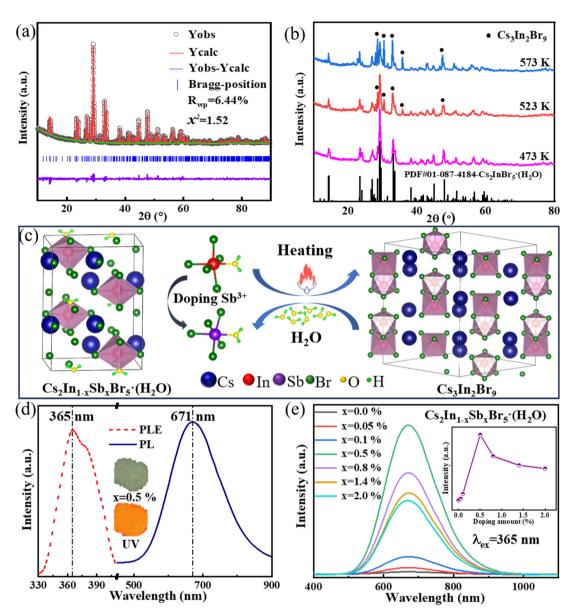


Fig. 2 (a) Rietveld refinement results of Cs₂InBr₅(H₂O) samples; (b) XRD patterns after annealing at different temperatures; (c) schematic representation of the reversible transformation between Cs₂InBr₅(H₂O) and Cs₃In₂Br₉ crystal structures; (d) excitation (red line) and emission (blue line) spectra of $Cs_2 ln Br_5(H_2O)$: Sb^{3+} (inset: photograph of the 0.5% Sb^{3+} sample under ambient light and 365 nm UV light); (e) photoluminescence spectra of $Cs_2In_{1-x}Sb_xBr_5(H_2O)$ (x = 0-2.0%), the inset shows the changes in Sb^{3+} doping concentration and PL intensity.

Cs₃In₂Br₉ back to Cs₂InBr₅(H₂O) is feasible upon rehydration with water molecules. This reversible hydration-dehydration process can be represented by the formula:

$$2Cs_2InBr_5 \cdot (H_2O)(orange) \mathop{\rightleftharpoons}_{+H_2O}^{-H_2O} Cs_3In_2Br_9(Yellow) + CsBr + 2H_2O. \tag{4}$$

The successful utilization of an in situ solution crystallization method to prepare a 0D lead-free Cs₂InBr₅(H₂O):Sb³⁺ halide perovskite is demonstrated. The structural transformation of the material is intimately associated with alterations in its color properties. The XRD patterns corresponding to

 $Cs_2In_{1-x}Sb_xBr_5(H_2O)$ compounds (x = 0-2.0%) are depicted in Fig. S2a.† These patterns reveal a close match between the diffraction peaks and those of the standard PDF card, indicating that introduction of Sb³⁺ as a dopant has a negligible impact on the overall crystal structure. Further examination of the XRD pattern at higher magnification, as shown in Fig. S2b,† unveils a noteworthy shift in diffraction peaks towards higher angles with increasing Sb3+ content. This shift can be attributed to the smaller ionic radius of Sb³⁺ (0.76 Å) compared to In³⁺ (0.80 Å), thereby suggesting that the doping concentration of Sb3+ exerts a subtle modulating effect on the structure.

Additionally, we observed that Sb³⁺-doped Cs₂InBr₅(H₂O) samples exhibit bright luminescence properties, characterized

by a visually perceptible "orange peel" color. To quantify this luminescence behavior, photoluminescence (PL) measurements were conducted. The photoluminescence excitation (PLE) spectrum in Fig. 2d presents a peak at 365 nm. Consequently, we selected 365 nm as the excitation source and observed an emission peak at 671 nm. Under identical conditions, the PL spectra of Cs2In1-xSbxBr5(H2O) shown in Fig. 2e exhibit intense orange-red emissions for all samples. At a doping concentration of 0.5% Sb³⁺, the emission intensity reaches its maximum. Therefore, the optimal doping concentration for $Cs_2In_{1-x}Sb_xBr_5(H_2O)$ is determined to be x = 0.5%, designated as the sample group with the best performance for subsequent temperature and humidity responsiveness assessments. Many studies indicate that self-trapped excitons (STEs) are the primary mechanism underlying the broad spectral emission in the 0D Cs₂InBr₅(H₂O):Sb³⁺ perovskite.^{38,51} As shown in Fig. S3,† the luminescence mechanism of the Cs₂InBr₅(H₂O):Sb³⁺ perovskite involves the STE process.³³ Through coupling with polar lattice vibrations, excitons reduce the total energy of the system due to transient lattice deformations, thereby resulting in broad spectral emissions and larger Stokes shifts. Therefore, the luminescence of Sb3+ ions in Cs₂InBr₅(H₂O):Sb³⁺ is attributed to the formation of STE. The emission peaks originate from the structural distortion of Cs₂InBr₅(H₂O):Sb³⁺ crystals doped with Sb³⁺, which promotes the emission of both singlet and triplet STEs.

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We conducted a comprehensive study of the synthesized compound, Cs₂InBr₅(H₂O):Sb³⁺, to assess its responsiveness to environmental changes by subjecting it to various temperature and humidity conditions. Fig. 3a shows the inscription of "SUST" on the sample after grinding the as-prepared powder

with ethanol. Upon exposure to UV lamp irradiation at room temperature, the emitted light from "SUST" appeared yellow, but it changed to orange-red under humid conditions. The intensity of the emitted light decreased with increasing temperatures, resulting in a state of fluorescence thermal quenching. As depicted in Fig. 2b, elevating the sample temperature to high degrees resulted in a structural transformation from the 0D perovskite Cs₂InBr₅(H₂O) to the Cs₃In₂Br₉-type perovskite. *In situ* PL measurements under certain humidity conditions (as depicted in the inset of Fig. 3b) yielded the intensity-normalized spectrum. The PL emission peak migrated from 589 nm (red) to 671 nm (orange-red) over a period of 270 s, indicating a transition in color from yellow to orange-red light starting from 0 s after exposure to the humid environment. Furthermore, the response time for the transition from yellow to red light varies at different humidity levels. This is attributed to the formation of a distorted octahedral structure in Cs₃In₂Br₉, where the In-Br bonds form a regular octahedron. In contrast, Cs₂InBr₅(H₂O):Sb³⁺ contains five Br ions and one H₂O, forming five In-Br bonds and one In-O bond, leading to a distorted octahedral structure. This distortion breaks the symmetry of Cs₂InBr₅(H₂O):Sb³⁺ and results in a larger Stokes shift. Therefore, humidity levels during the reaction process are crucial for reversible processes.

Fig. 3c highlights the significant impact of environmental humidity on the luminescence properties of the material, with a notable emission peak shift observed at around 20 s from 597 nm to approximately 664 nm, accompanied by a gradual color change from yellow to orange-red, as depicted in the inset of Fig. 3c. Fig. S4a and b† show that an increase in dis-

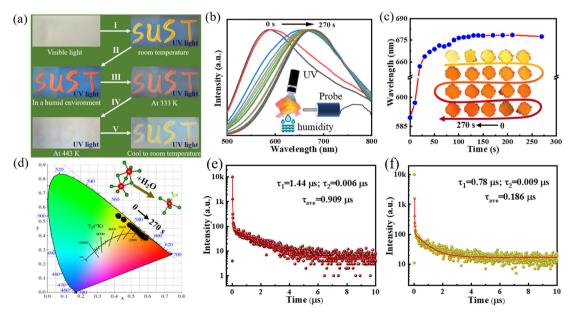


Fig. 3 Sensitivity of Cs₂InBr₅(H₂O):Sb³⁺ to water molecules. (a) Digital photograph of "SUST" under various conditions. (b) PL spectra variation in a humid environment; the illustration depicts the in situ testing process of PL. (c) Peak position as a function of exposure time to a humid environment (inset showing the color change of the sample's emission from 0 to 270 s). (d) CIE chromaticity diagram of the sample in a humid environment. The time-resolved PL decay curve for (e) Cs₂lnBr₅(H₂O):Sb³⁺ and (f) Cs₃ln₂Br₉.

tilled water content resulted in a redshift of the emission peak. The color alteration of Cs₂InBr₅(H₂O):Sb³⁺ from green light to red light is shown in the digital photographs in Fig. S4c† after the addition of 0.05 mL of distilled water. This change further verifies the high sensitivity of Cs₂InBr₅(H₂O):Sb³⁺ to humidity levels,³¹ enabling precise control over the material's luminescence color, as shown in Fig. 3d. We characterized the fluorescence lifetimes of Cs₂InBr₅(H₂O):Sb³⁺ (red light) and Cs₃In₂Br₉ (yellow light) to probe their photophysical properties. As shown in Fig. 3e and f, we observed a biexponential decay with an average lifetime of 0.909 µs for the 0D Cs₂InBr₅(H₂O):Sb³⁺ perovskite, attributed to characteristic singlet and triplet transitions of STEs. The average fluorescence lifetime of Cs₃In₂Br₉ decreased to 0.186 μs. Furthermore, in Fig. S5,† by employing the integrating sphere method, we measured the excitation and emission peaks, calculated the difference in their areas, and determined the photoluminescence quantum yields (PLOYs) of Cs₂InBr₅(H₂O) and Cs₃In₂Br₉ to be 87.8% and 22.8%, respectively. This significant contrast suggests that water molecules (H2O) may exert a notable influence on the PLQY of Cs₃In₂Br₉. Therefore, modulating the structure of Cs₃In₂Br₉ or doping specific ions could potentially enhance its luminescence performance, thereby increasing its practical utility.

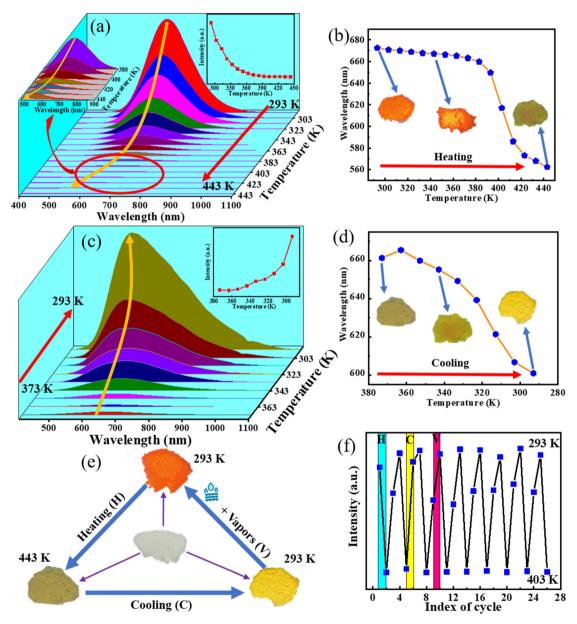


Fig. 4 (a) PL-temperature spectra of $Cs_2In_{0.995}Sb_{0.005}Br_5(H_2O)$ (inset: intensity *versus* temperature curve); (b) variation of the emission peak position during the heating process; (c) PL spectra during the temperature cooling process (inset: intensity versus temperature); (d) variation of the emission peak position during the cooling process; (e) color change of the sample during the cycling process; (f) changes in PL intensity during cycles of heating (H), cooling (C), and vapors (V).

In response to the temperature-induced alterations observed in Cs₂InBr₅(H₂O):Sb³⁺, as depicted in Fig. 3a, we undertook an examination of the temperature-dependent photoluminescence (PL) spectra of Cs₂In_{0.995}Sb_{0.005}Br₅(H₂O), spanning a temperature range of 293-443 K. As shown in Fig. 4a, the fluorescence intensity gradually decreases with increasing temperature. Concurrently, during the heating process, the emission peak gradually blue-shifts (Fig. 4b), and the full width at half maximum (FWHM) initially broadens and then narrows as illustrated in Fig. S6.† Fig. 4c depicts the cooling process, where the fluorescence intensity gradually increases as the temperature decreases, accompanied by a blue-shift in the emission peak (Fig. 4d), with the FWHM initially narrowing and then broadening. Fig. 4b and d insets illustrate that the sample emits orange-red light at room temperature. Upon heating to 443 K, faint yellow light is observed, and cooling back to room temperature emits bright yellow light. The cyclic process of heating (H), cooling (C), and vapors (V) of the Cs₂In_{0.995}Sb_{0.005}Br₅(H₂O) sample is represented in Fig. 4e. The color change is attributed to the evaporation of water molecules from the molecular structure, leading to yellow light emission as the temperature increases. This process demonstrates reversibility, as samples emitting yellow light in a humid environment reverted to orange-red after a storage period. The temperature-dependent spectra in cycles of

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heating and cooling demonstrate good stability, as shown in Fig. S7.† This observation highlights the high sensitivity of Cs₂In_{0.995}Sb_{0.005}Br₅(H₂O) to both temperature and humidity. The evaporation of water molecules causes the color changes. Water molecules play a crucial role in the color-changing process, allowing for repeatable and reversible color tuning. Fig. 4 shows the dynamic interaction between temperature, humidity, and molecular structure, highlighting the potential for further exploration in the manipulation and control of water molecules in these materials.

The compound's temperature sensing performance was determined using the fluorescence intensity variation shown in Fig. 4a. Given the material's broad single-peak emission profile, the conventional fluorescence intensity ratio (FIR) temperature calculation formula could not be utilized. Fig. 5a shows the fluorescence intensity values of the material at different temperatures, decreasing continuously with temperature elevation within the range of 293-413 K. By employing a nonlinear regression analysis of the data, the derived formula is:

$$I = A \exp(-T/B) + C. \tag{2}$$

The values for A, B, and C are 5.64×10^9 , 28.55, and -3.41×10^9 10³. These values can be utilized to analyze absolute sensitivity

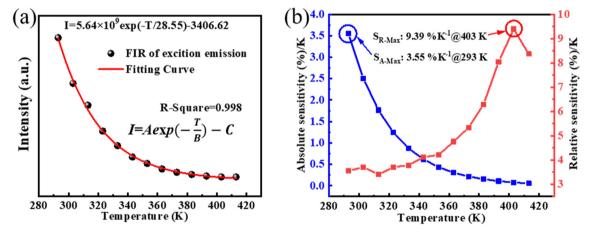


Fig. 5 (a) Plot of intensity at different temperatures with nonlinear fitting curves; (b) S_A curve and S_R curve as a function of temperature.

Table 1 Temperature sensitivity performance of other luminescent materials

Samples	Temperature range (K)	$S_{\text{R-Max}}$ temperature (K)	$S_{\text{R-Max}} \left(\% \text{ K}^{-1} \right)$	Ref.
Bi ₂ Ti ₂ O ₇ :Yb ³⁺ /Er ³⁺	298-540	298	1.53	54
$Cs_3Cu_2I_5:Mn^{2+}$	298-498	498	0.547	55
CsPbCl ₂ Br:Eu ³⁺	80-440	400	3.10	56
NaYbF ₄ :Er ³⁺	175-475	175	3.46	57
CaTiO ₃ :Yb ³⁺ /Er ³⁺	303-523	303	1.17	58
$Cs_2Ag_{0.6}Na_{0.4}In_{0.9}Bi_{0.1}Cl_6$	300-470	345	1.05	59
Cs ₂ Ag _{0.6} Na _{0.4} In _{0.9} Bi _{0.1} Cl ₆ Cs ₂ NaYbCl ₆ :Er ³⁺ /Sb ³⁺	80-310	310	11.21	60
Cs ₄ PbBr ₆ :Sm ³⁺	303-423	303	3.83	61
$Cs_2InCl_5(H_2O)$: Te^{4+}	80-380	320	6.2	62
$Cs_2InBr_5(H_2O):Sb^{3+}$	293-443	403	9.39	This worl

 (S_A) and relative sensitivity (S_B) using the following formulas:52,53

$$S_{\rm A} = \frac{{\rm d}(I(T)/I(0))}{{\rm d}T} \quad S_{\rm R} = S_{\rm A} \frac{I(0)}{I(T)}.$$
 (3)

The initial fluorescence intensity and the intensity corresponding to temperature T are represented by the symbols I(0)and I(T), respectively. S_A and S_R of the samples at different temperatures are shown in Fig. 5b, where S_{A-Max} is 3.55% K^{-1} @293 K and S_{R-Max} is 9.39% K^{-1} @403 K. The S_{R-Max} achieved in this study is notably higher than those of other perovskites, as indicated by a comparison with literature values for other temperature sensing materials (refer to Table 1). This discrepancy suggests the potential importance of the Cs₂InBr₅(H₂O):Sb³⁺ compound in temperature sensing applications, presenting new opportunities for the advancement of higher-performance temperature sensors and other optoelectronic devices due to its enhanced temperature sensitivity.

Conclusions 3.

In our study, we synthesized a 0D halide perovskite using an environmentally friendly fabrication method, which revealed fractal structure growth through an in situ solution crystallization approach. Upon exposure to 365 nm UV excitation, the material exhibited an emission peak that could be tuned from 589 nm (yellow) to 671 nm (orange-red). Our confirm of the reversible structural and photoluminescence transitions triggered by water molecules or thermal reactions in the Sb3+doped Cs₂InBr₅(H₂O). We found that heating the 0D Cs₂In_{1-x}Sb_xBr₅(H₂O) perovskite leads to dehydration and transformation into the Cs₃In₂Br₉ structure. Notably, brief exposure to defined humidity levels (30 s) under specific relative humidity (RH) conditions reversed this transformation, reinstating the compound's original 0D configuration. By precisely manipulating temperature and humidity, we induced multiple color transitions in the material, revealing its versatile photoluminescence properties. This reversible luminescence behavior of the 0D lead-free halide perovskite sheds light on the structural and luminescence characteristics of the perovskite family, thereby broadening their prospective integration into reusable smart material systems. Furthermore, Cs₂InBr₅(H₂O):Sb³⁺ exhibited remarkable temperature sensitivity, revealing a relative sensitivity (S_R) of 9.39% K⁻¹@403 K. The high sensitivity highlights the material's broad potential for temperature sensing applications, showcasing its adaptability in humidity- and temperature-sensitive systems.

Data availability

All relevant data are within the manuscript and its additional files. The data that support the findings of this study are available from the corresponding author upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors gratefully acknowledge the funding from the National Natural Science Foundation of China (52073165).

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