

## RESEARCH ARTICLE

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# Reversible “on–off” conversion and ultra-high temperature sensitivity of a zero-dimensional lead-free $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$ perovskite†

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Zero-dimensional (0D) lead-free perovskites have garnered significant attention due to their unique opto-electronic 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  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  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  $\text{H}_2\text{O}$  molecules played a crucial role in the color tuning, enabling reversible “on–off” switching modes with rapid modulation rates. Moreover, the 0D lead-free halide perovskite  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  also exhibited excellent temperature sensitivity, with the relative sensitivity ( $S_R$ ) reaching up to  $9.39\% \text{ K}^{-1}$ . 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.

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## 1. Introduction

Recently, lead halide perovskites have shown excellent opto-electronic properties, with growing applications in various fields.<sup>1–8</sup> However, the toxicity of the lead element severely limits the practical application.<sup>9–15</sup> 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.<sup>16–22</sup> 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, light-emitting diodes, and lasers, bringing new development opportunities for modern optoelectronic technology.<sup>23–26</sup> However, the innovative synthesis route and optical characteristics of the zero-dimensional (0D) indium-based halide perovskite

$\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ , as well as its intrinsic physicochemical properties, still require further investigation.<sup>27</sup>

Currently, the incorporation of dopants is recognized as an effective strategy for improving the luminescence efficiency of perovskites.<sup>28</sup> Substantial advancements have been achieved in the study of the photoluminescence (PL) properties of indium-based halide perovskites doped with antimony ( $\text{Sb}^{3+}$ ). For instance,  $\text{Sb}^{3+}$ -doping in 0D  $\text{Cs}_2\text{SnCl}_6$  exhibits yellow-red emission.<sup>29,30</sup> Compared to the undoped counterparts, the photoluminescence quantum yield (PLQY) of  $\text{Sb}^{3+}$ -doped  $\text{A}_2\text{InCl}_5(\text{H}_2\text{O})$  ( $\text{A} = \text{Rb}, \text{Cs}$ ) has significantly improved, increasing from below 2% to a range of 85–95%.<sup>31</sup> In the family of  $\text{Cs}_2\text{InX}_5(\text{H}_2\text{O})$  ( $\text{X} = \text{Cl}^-, \text{Br}^-$  and  $\text{I}^-$ ), the octahedral units exhibit pronounced electron–phonon coupling, resulting in a substantial Stokes shift observed in these 0D perovskite compounds.<sup>32</sup> Substitution of the halide from  $\text{Cl}^-$  to  $\text{Br}^-$  and  $\text{I}^-$  also leads to redshifts in the photoluminescence (PL) and photoluminescence excitation (PLE) spectra of  $\text{Sb}^{3+}$ -doped  $\text{Cs}_2\text{InX}_5(\text{H}_2\text{O})$  compounds.<sup>33</sup> Additionally, applying high pressure enables precise modulation of the PL properties of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ , culminating in the observation of anomalous anti-Stokes emission.<sup>34</sup> In  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ , photogenerated charge carriers are spatially confined within isolated octahedra, enhancing the radiative recombination probability.<sup>35–38</sup> A-site cations have a slight effect on PL,<sup>39</sup> while water mole-

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cules, through their induced distortion of  $\text{In}^{3+}$  octahedra, result in a broad emission and exert a more pronounced influence on the self-trapped excitons (STEs).<sup>40</sup>

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.<sup>41–46</sup> Fortunately, the novel lead-free 0D halide perovskite  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  stands out for its exceptional stability and tunable crystal structure. Interestingly, its isolated octahedral structure incorporates a water molecule ( $\text{H}_2\text{O}$ ). As the temperature increases, the water molecule is released, modulating its luminescence from red to yellow.<sup>47,48</sup> 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.<sup>49</sup>

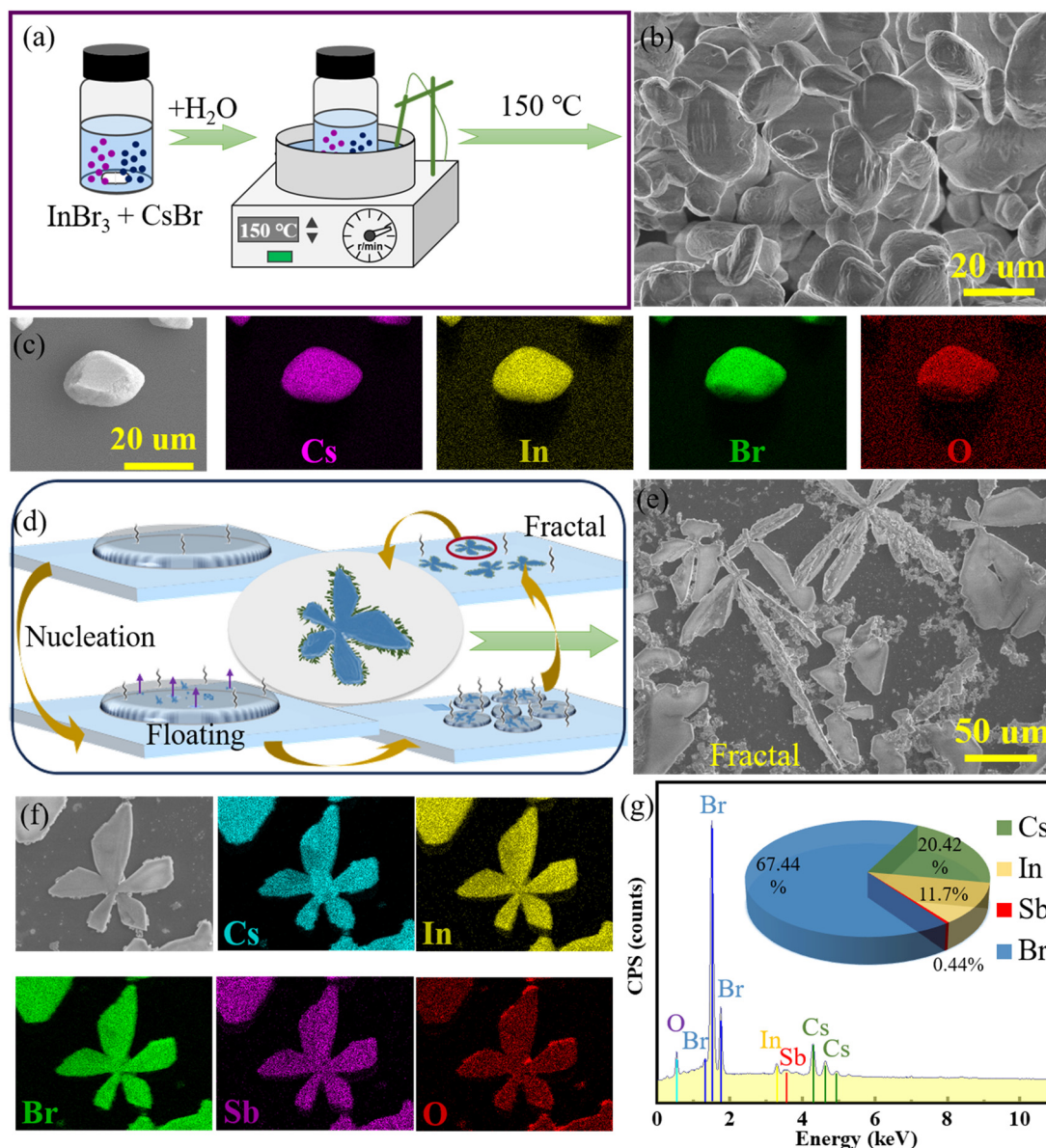
High-performance functional compounds that are reversible and responsive to stimuli such as temperature or humidity are highly desirable for sophisticated anti-counterfeiting technologies. However, many existing anti-counterfeiting functional compounds exhibit limited temperature responsiveness and necessitate further development to enhance their reversibility. In this study, we report the successful synthesis of a 0D halide perovskite,  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$ , 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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  are of significant importance. We observed that the water molecule in the  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  molecular structure acts as a switch, facilitating an “on-off-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  $3.55\% \text{ K}^{-1}$  and  $S_R$  reaching a peak of  $9.39\% \text{ K}^{-1}$ , indicating extremely high sensitivity. These research findings demonstrate the high sensitivity and responsiveness of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  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.

## 2. 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  $\text{InBr}_3$  (1 mmol, 99%, Sigma-Aldrich),  $\text{CsBr}$  (2 mmol, 99.9%, Sigma-Aldrich), and an appropriate amount of  $\text{SbBr}_3$  (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  $150^\circ\text{C}$  for 2 min, clearly displaying granular grains with sizes around  $20 \mu\text{m}$ . 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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ , thereby validating the successful synthesis of the  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  compound.

Figure 1d demonstrates crystallization near room temperature (around  $30^\circ\text{C}$ ), 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.<sup>50</sup> 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  $\text{Cs}:\text{In}:\text{Br} = 2:1:6$ , 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 as-prepared 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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ , with an  $R_{\text{wp}}$  value of 6.44% and a goodness-of-fit factor of 1.52, signifying a high degree of fitting accuracy. The crystallographic para-

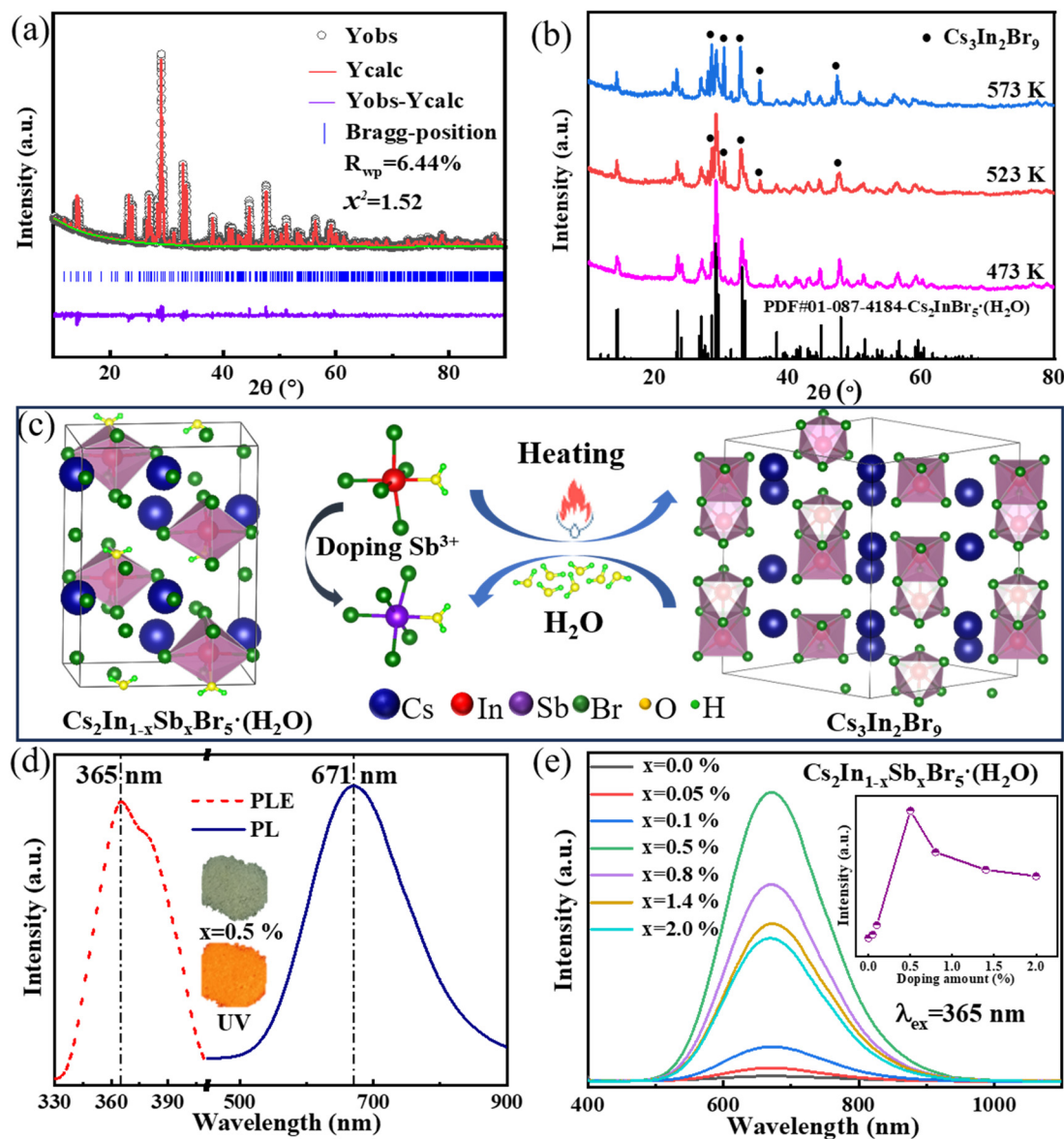


**Fig. 1** (a) Preparation process of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ ; (b) SEM image of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  prepared at  $150\text{ }^\circ\text{C}$ ; (c) EDX mapping of Cs, In, Br, and O elements; (d) fractal structure of  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}$ .

parameters derived were  $a = 14.8763\text{ \AA}$ ,  $b = 10.8116\text{ \AA}$ , and  $c = 7.6437\text{ \AA}$ , consistent with the  $pnma$  space group. Additionally, we observed a color change in the luminescence of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  gradually decreased, accompanied by the appearance of diffraction peaks corresponding to the  $\text{Cs}_3\text{In}_2\text{Br}_9$  phase. This fact indicates that with increasing temperature, water molecules are progressively released from

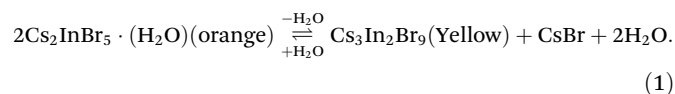
$\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ , leading to the partial transformation of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  into  $\text{Cs}_3\text{In}_2\text{Br}_9$ . Notably, this process is reversible, as the structure of the sample can be regenerated upon exposure to a humid environment.

The octahedral structure in the  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  perovskite is composed of one indium (In) atom, five bromine (Br) atoms, and a single  $\text{H}_2\text{O}$  molecule, with indium atoms being partially substituted by Sb atoms, as illustrated in the crystal structure in Fig. 2c. Two  $\text{Cs}^+$  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  $\text{Cs}_3\text{In}_2\text{Br}_9$ . Interestingly, the reverse transformation from



**Fig. 2** (a) Rietveld refinement results of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  samples; (b) XRD patterns after annealing at different temperatures; (c) schematic representation of the reversible transformation between  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  and  $\text{Cs}_3\text{In}_2\text{Br}_9$  crystal structures; (d) excitation (red line) and emission (blue line) spectra of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  (inset: photograph of the 0.5%  $\text{Sb}^{3+}$  sample under ambient light and 365 nm UV light); (e) photoluminescence spectra of  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  ( $x = 0\text{--}2.0\%$ ), the inset shows the changes in  $\text{Sb}^{3+}$  doping concentration and PL intensity.

$\text{Cs}_3\text{In}_2\text{Br}_9$  back to  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  is feasible upon rehydration with water molecules. This reversible hydration–dehydration process can be represented by the formula:



The successful utilization of an *in situ* solution crystallization method to prepare a 0D lead-free  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  halide perovskite is demonstrated. The structural transformation of the material is intimately associated with alterations in its color properties. The XRD patterns corresponding to

$\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  compounds ( $x = 0\text{--}2.0\%$ ) 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  $\text{Sb}^{3+}$  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  $\text{Sb}^{3+}$  content. This shift can be attributed to the smaller ionic radius of  $\text{Sb}^{3+}$  (0.76 Å) compared to  $\text{In}^{3+}$  (0.80 Å), thereby suggesting that the doping concentration of  $\text{Sb}^{3+}$  exerts a subtle modulating effect on the structure.

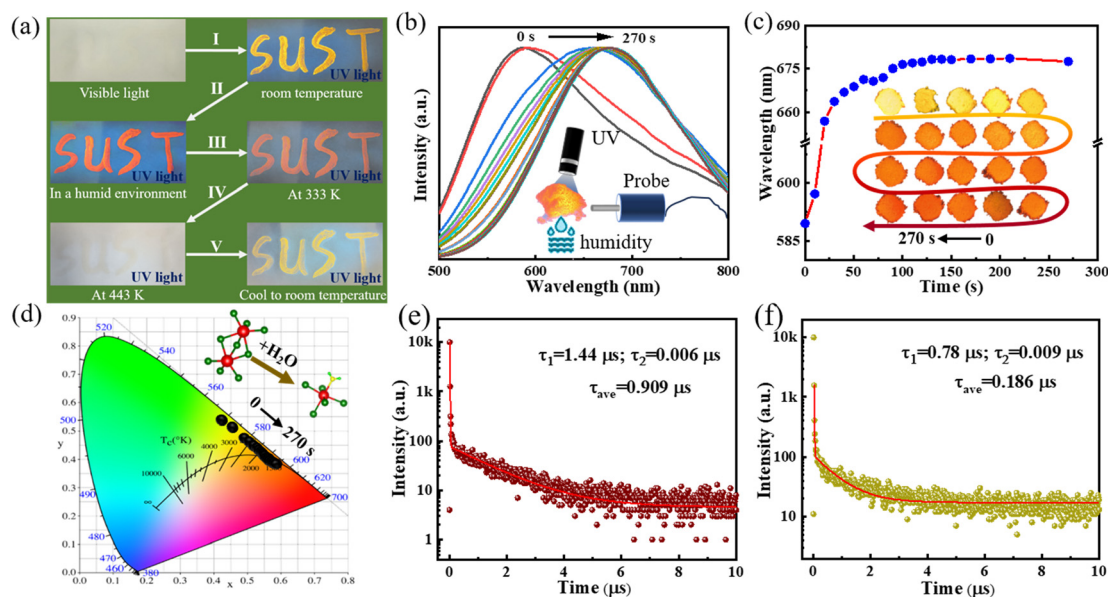
Additionally, we observed that  $\text{Sb}^{3+}$ -doped  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  samples exhibit bright luminescence properties, characterized

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  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  shown in Fig. 2e exhibit intense orange-red emissions for all samples. At a doping concentration of 0.5%  $\text{Sb}^{3+}$ , the emission intensity reaches its maximum. Therefore, the optimal doping concentration for  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  perovskite.<sup>38,51</sup> As shown in Fig. S3,† the luminescence mechanism of the  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  perovskite involves the STE process.<sup>33</sup> 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  $\text{Sb}^{3+}$  ions in  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  is attributed to the formation of STE. The emission peaks originate from the structural distortion of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  crystals doped with  $\text{Sb}^{3+}$ , which promotes the emission of both singlet and triplet STEs.

We conducted a comprehensive study of the synthesized compound,  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$ , 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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  to the  $\text{Cs}_3\text{In}_2\text{Br}_9$ -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  $\text{Cs}_3\text{In}_2\text{Br}_9$ , where the In–Br bonds form a regular octahedron. In contrast,  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  contains five Br ions and one  $\text{H}_2\text{O}$ , forming five In–Br bonds and one In–O bond, leading to a distorted octahedral structure. This distortion breaks the symmetry of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  and results in a larger Stokes shift. Therefore, humidity levels during the reaction process are crucial for reversible processes.

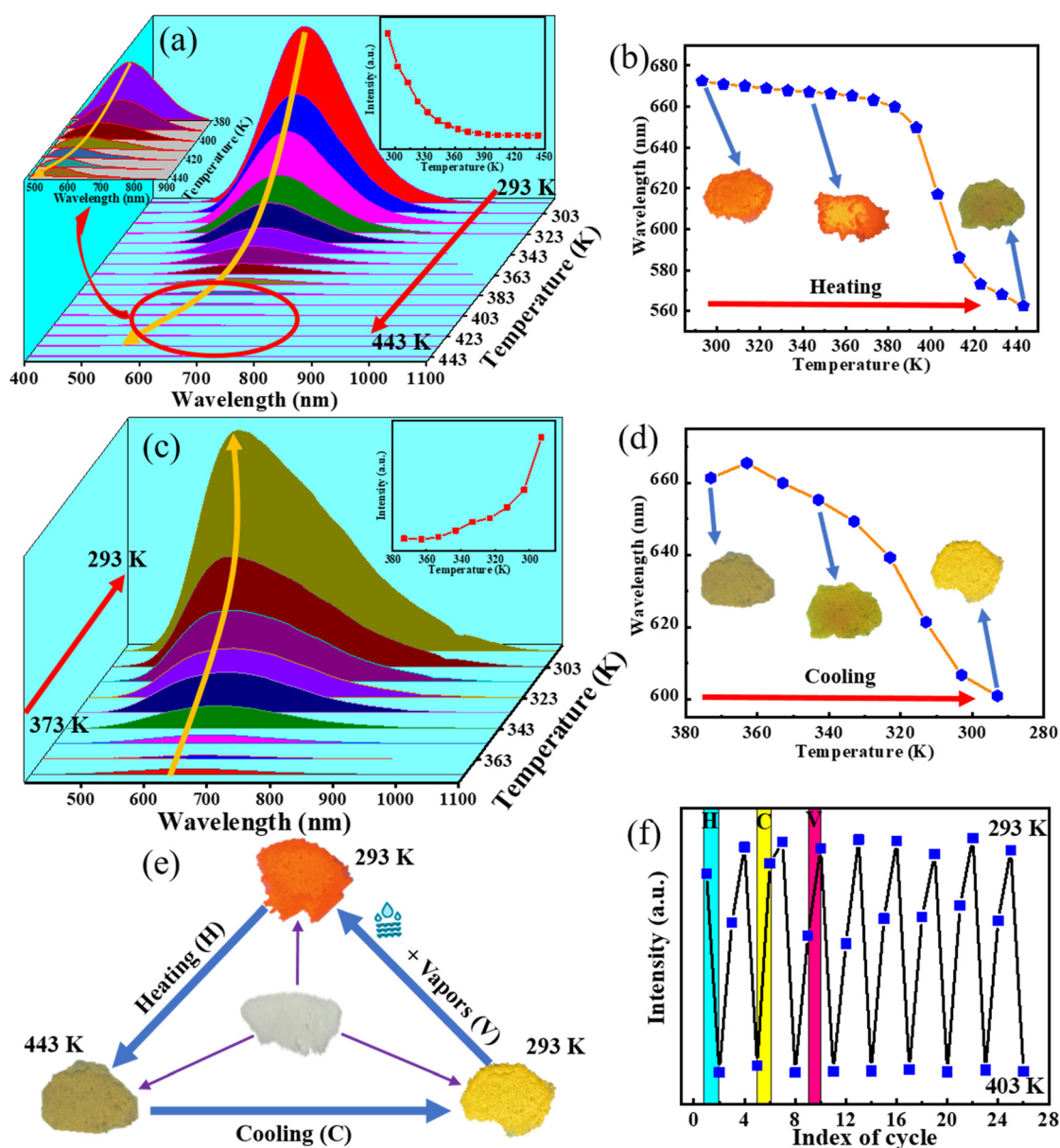
Fig. 3c highlights the significant impact of environmental humidity on the luminescence properties of the material, with a notable emission peak shift observed at around 20 s from 597 nm to approximately 664 nm, accompanied by a gradual color change from yellow to orange-red, as depicted in the inset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



**Fig. 3** Sensitivity of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  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)  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  and (f)  $\text{Cs}_3\text{In}_2\text{Br}_9$ .

titled water content resulted in a redshift of the emission peak. The color alteration of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  to humidity levels,<sup>31</sup> enabling precise control over the material's luminescence color, as shown in Fig. 3d. We characterized the fluorescence lifetimes of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  (red light) and  $\text{Cs}_3\text{In}_2\text{Br}_9$  (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  $\mu\text{s}$  for the 0D  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  perovskite, attributed to characteristic

singlet and triplet transitions of STEs. The average fluorescence lifetime of  $\text{Cs}_3\text{In}_2\text{Br}_9$  decreased to 0.186  $\mu\text{s}$ . 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 (PLQYs) of  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$  and  $\text{Cs}_3\text{In}_2\text{Br}_9$  to be 87.8% and 22.8%, respectively. This significant contrast suggests that water molecules ( $\text{H}_2\text{O}$ ) may exert a notable influence on the PLQY of  $\text{Cs}_3\text{In}_2\text{Br}_9$ . Therefore, modulating the structure of  $\text{Cs}_3\text{In}_2\text{Br}_9$  or doping specific ions could potentially enhance its luminescence performance, thereby increasing its practical utility.



**Fig. 4** (a) PL-temperature spectra of  $\text{Cs}_2\text{In}_{0.995}\text{Sb}_{0.005}\text{Br}_5(\text{H}_2\text{O})$  (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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$ , as depicted in Fig. 3a, we undertook an examination of the temperature-dependent photoluminescence (PL) spectra of  $\text{Cs}_2\text{In}_{0.995}\text{Sb}_{0.005}\text{Br}_5(\text{H}_2\text{O})$ , 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  $\text{Cs}_2\text{In}_{0.995}\text{Sb}_{0.005}\text{Br}_5(\text{H}_2\text{O})$  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

heating and cooling demonstrate good stability, as shown in Fig. S7.† This observation highlights the high sensitivity of  $\text{Cs}_2\text{In}_{0.995}\text{Sb}_{0.005}\text{Br}_5(\text{H}_2\text{O})$  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:

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

The values for  $A$ ,  $B$ , and  $C$  are  $5.64 \times 10^9$ , 28.55, and  $-3.41 \times 10^3$ . These values can be utilized to analyze absolute sensitivity

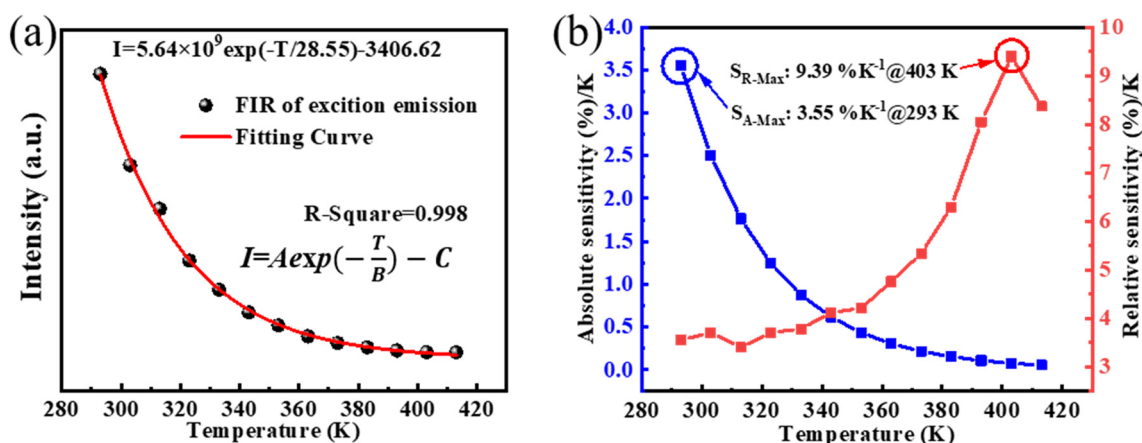


Fig. 5 (a) Plot of intensity at different temperatures with nonlinear fitting curves; (b)  $S_A$  curve and  $S_R$  curve as a function of temperature.

Table 1 Temperature sensitivity performance of other luminescent materials

| Samples  | Temperature range (K) | $S_{R-\text{Max}}$ temperature (K) | $S_{R-\text{Max}}$ (% $\text{K}^{-1}$ ) | Ref.      |
|--|-----------------------|------------------------------------|---|-----------|
| $\text{Bi}_2\text{Ti}_2\text{O}_7\text{:Yb}^{3+}/\text{Er}^{3+}$                     | 298–540               | 298                                | 1.53                                    | 54        |
| $\text{Cs}_3\text{Cu}_2\text{I}_5\text{:Mn}^{2+}$                                    | 298–498               | 498                                | 0.547                                   | 55        |
| $\text{CsPbCl}_2\text{Br:Eu}^{3+}$   | 80–440                | 400                                | 3.10                                    | 56        |
| $\text{NaYbF}_4\text{:Er}^{3+}$  | 175–475               | 175                                | 3.46                                    | 57        |
| $\text{CaTiO}_3\text{:Yb}^{3+}/\text{Er}^{3+}$                                       | 303–523               | 303                                | 1.17                                    | 58        |
| $\text{Cs}_2\text{Ag}_{0.6}\text{Na}_{0.4}\text{In}_{0.9}\text{Bi}_{0.1}\text{Cl}_6$ | 300–470               | 345                                | 1.05                                    | 59        |
| $\text{Cs}_2\text{NaYbCl}_6\text{:Er}^{3+}/\text{Sb}^{3+}$                           | 80–310                | 310                                | 11.21                                   | 60        |
| $\text{Cs}_4\text{PbBr}_6\text{:Sm}^{3+}$  | 303–423               | 303                                | 3.83                                    | 61        |
| $\text{Cs}_2\text{InCl}_5(\text{H}_2\text{O})\text{:Te}^{4+}$                        | 80–380                | 320                                | 6.2                                     | 62        |
| $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$                        | 293–443               | 403                                | 9.39                                    | This work |

( $S_A$ ) and relative sensitivity ( $S_R$ ) using the following formulas:<sup>52,53</sup>

$$S_A = \frac{d(I(T)/I(0))}{dT} \quad S_R = S_A \frac{I(0)}{I(T)}. \quad (3)$$

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  $S_{A-\text{Max}}$  is 3.55%  $\text{K}^{-1}$ @293 K and  $S_{R-\text{Max}}$  is 9.39%  $\text{K}^{-1}$ @403 K. The  $S_{R-\text{Max}}$  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  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  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.

### 3. Conclusions

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  $\text{Sb}^{3+}$ -doped  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})$ . We found that heating the 0D  $\text{Cs}_2\text{In}_{1-x}\text{Sb}_x\text{Br}_5(\text{H}_2\text{O})$  perovskite leads to dehydration and transformation into the  $\text{Cs}_3\text{In}_2\text{Br}_9$  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 photoluminescence 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,  $\text{Cs}_2\text{InBr}_5(\text{H}_2\text{O})\text{:Sb}^{3+}$  exhibited remarkable temperature sensitivity, revealing a relative sensitivity ( $S_R$ ) of 9.39%  $\text{K}^{-1}$ @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.

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