2D nanosheets of layered double perovskites: synthesis, photostable bright orange emission and photoluminescence blinking†

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Lead (Pb)-free layered double perovskites (LDPs) with exciting optical properties and environmental stability have sparked attention in optoelectronics, but their high photoluminescence (PL) quantum yield and understanding of the PL blinking phenomenon at the single particle level are still elusive. Herein, we not only demonstrate a hot-injection route for the synthesis of two-dimensional (2D) ~2–3 layer thick nanosheets (NSs) of LDP, Cs₂CdBi₂Cl₁₂ (pristine), and its partially Mn-substituted analogue [i.e., Cs₂Cdₐ₂Mₐ₀.₄Bi₂Cl₁₂ (Mn-substituted)], but also present a solvent-free mechanochemical synthesis of these samples as bulk powders. Bright and intense orange emission has been perceived for partially Mn-substituted 2D NSs with a relatively high PL quantum yield (PLQY) of ~21%. The PL and lifetime measurements both at cryogenic (77 K) and room temperatures were employed to understand the de-excitation pathways of charge carriers. With the implementation of super-resolved fluorescence microscopy and time-resolved single particle tracking, we identified the occurrence of metastable non-radiative recombination channels in a single NS. In contrast to the rapid photo-bleaching that resulted in a PL blinking-like nature of the controlled pristine NS, the 2D NS of the Mn-substituted sample displayed negligible photo-bleaching with suppression of PL fluctuation under continuous illumination. The blinking-like nature in pristine NSs appeared due to a dynamic equilibrium flanked by the active and inactive states of metastable non-radiative channels. However, the partial substitution of Mn²⁺ stabilized the inactive state of the non-radiative channels, which increased the PLQY and suppressed PL fluctuation and photo-bleaching events in Mn-substituted NSs.

Introduction

Metal halide perovskite nanostructures have received significant attention owing to their outstanding optoelectronic properties. In spite of their unique properties, the foremost shortcoming is the presence of lead (Pb), which obstructs them from further applications. Hence, the search for environmentally-friendly replacements that maintain the remarkable optical properties of Pb-based perovskite halides has become an increasingly significant research area. Promising approaches included isovalent substitution of Pb²⁺ by less- or non-toxic metal ions, like Sn²⁺ and Ge²⁺. However, facile oxidation of these cations under ambient conditions leads to the inferior stability of crystal structures and impedes their applications. Subsequently, strategies for employing cations with an oxidation state other than +2 in place of Pb²⁺ in perovskites are being extensively explored. The lower dimensional A₂Mₙ⁺⁺Xₙ⁻ type perovskites, and paired monovalent-trivalent cation based double perovskites, A₂M⁺Mₙ⁺⁺Xₙ⁻ [where M(1) and M(III) correspond to Ag⁺, Cu⁺, Na⁺, etc. and Bi³⁺, Sb³⁺, In³⁺, etc., respectively], are being actively studied, and have shown promising properties.

Recently, dimensional reduction and heterovalent substitution resulted in layered double perovskite (LDP) halides, which can be reckoned as the two-dimensional (2D) variety of double perovskites or the double-metal version of layered perovskites. Different from the conventional double perovskites, LDPS integrate the divalent and trivalent cations into 2D structures, which further led to the chemical and structural diversity. Additionally, the dimensional reduction can engender modifications in the electronic structure. Solis-Ibarra and co-workers initially synthesized a mixed-metal 〈111〉-oriented layered double perovskite with the general...
formula of \( A_{n+1}[B(n)B'(n)]_nX_{3n+3} \), where \( n > 2 \) (i.e., \( \text{Cs}_4\text{Cu}_3\text{Sb}_2\text{Cl}_{12} \)). Subsequently, few LDPs are available, which are either computationally proposed or experimentally synthesized.\(^{3,8}\) Till now, the bulk form of LDPs (e.g., \( \text{A}_n\text{M}^{++}\text{M}^{+1}\text{X}_{12} \)) has been synthesized mainly by acid precipitation methods.\(^{3,8}\) Despite the great promise of this family of materials, very few reports have explored the synthesis of LDP nanostructures by the solution-based colloidal method.\(^9\) Recently, colloidal synthesis of \( \text{Cs}_4\text{M}^{n}[\text{B}_n\text{Cl}_{3n+1}] \) \( (\text{M} = \text{Cd}, \text{Mn}) \) nanocrystals (NCs) was reported using the hot-injection method; however, a maximum photoluminescence quantum yield (PLQY) of only 4.6% was achieved for the \( \text{Cs}_4\text{Cd}_1\text{–}_x\text{Mn}_x\text{Bi}_2\text{Cl}_12 \) NCs.\(^{3,8}\) Therefore, facile synthesis strategies for both the bulk form and the nanostructures (e.g., 2D nanolayers or nanosheets) of such LDPs should be developed to improve the PLQY and to understand the PL properties at the single particle level.

Herein, we report a simple, solvent-free, scalable, and environment friendly all-solid-state mechanochemical synthesis of the bulk powders of LDPs, \( \text{Cs}_4\text{CdBi}_2\text{Cl}_{12} \) (pristine) and a partially Mn-substituted analogue \( \text{Cs}_4\text{Cd}_{0.6}\text{Mn}_{0.4}\text{Bi}_2\text{Cl}_{12} \). More importantly, we have synthesized the 2–3 layer thick 2D nanosheets (NSs) of the pristine and Mn-substituted analogue by the one-pot solution-based hot-injection method using benzoyl chloride as a halide precursor. These materials showed high thermal and environmental stability. The 2D NSs of the Mn-substituted analogue showed a relatively high PLQY of \( \sim 21\% \) at room temperature. A deeper understanding of the charge-carrier dynamics in these 2D NSs was achieved through time-resolved PL measurements both at room and cryogenic (77 K) temperatures. Super-resolved fluorescence microscopy and time-resolved single particle tracking were employed to investigate the PL blinking event in 2D NSs of the pristine and Mn-substituted analogue at the single particle level (herein, NS) level, which is a key process to control the emission efficiency. Photo-bleaching and temporal intermittency (blinking) behaviours have been perceived in controlled pristine NSs, whereas the single NS of the Mn-substituted analogue displayed relatively suppressed PL fluctuation characteristics. An enriched PL intensity with a high photon count was evidenced in Mn-substituted NSs via structured illumination microscopy (SIM) accompanied by negligible photo-bleaching behaviour under continuous light illumination.

**Results and discussion**

Here, we first discuss the solvent-free mechanochemical synthesis, stability, and optical properties of bulk polycrystalline powders, and subsequently, we present the colloidal synthesis of 2D nanosheets (NSs) of \( \text{Cs}_4\text{CdBi}_2\text{Cl}_{12} \) (pristine), and \( \text{Cs}_4\text{Cd}_{0.6}\text{Mn}_{0.4}\text{Bi}_2\text{Cl}_{12} \) (Mn-substituted analogue); and super-resolved fluorescence microscopy results to understand the PL fluctuation behaviour at the single NS level.

To synthesize the bulk powders of the pristine and Mn-substituted analogue, we have used \( \text{CsCl} \), \( \text{CdCl}_2 \), \( \text{MnCl}_2 \), and \( \text{BiCl}_3 \) as precursors in the appropriate stoichiometric ratio. Typically, a stoichiometric amount of the metal chloride precursors was mixed and ground mechanically in a mortar and pestle for 2 hours in an ambient laboratory environment (Schemes 1 and 2). Previously, such bulk powders have been synthesized by the precipitation method using concentrated hydrochloric acid.\(^{3,8}\) In contrast, our all-solid state mechanochemical approach is solvent-free, and scalable to \( \sim 1 \) g of pure crystalline product.

The powder X-ray diffraction (PXRD) pattern in Fig. 1b for the mechanochemically synthesized bulk powders was perfectly matched with the rhombohedral phase of pristine (space group \( \text{R}3\text{m} \)), which indicates the phase purity of the obtained product. A right shift of XRD peaks in 2\( \theta \) was evidenced for the bulk Mn-substituted analogue, which corresponds to the smaller ionic radius of Mn\(^{2+} \) (83 pm) as compared to that of Cd\(^{2+} \) (95 pm). \( \text{Cs}_4\text{Cd}_{1-x}\text{Mn}_x\text{Bi}_2\text{Cl}_{12} \) (where, \( x = 0 \) and 0.4) exhibits a layered double perovskite (LDP) crystal structure, which can be derived from the conventional cubic ABX\(_3\) perovskite structure. \( \text{Cs}_4\text{Cd}_{1-x}\text{Mn}_x\text{Bi}_2\text{Cl}_{12} \) possesses a unique structure with a layer of divalent metal halide octahedra (i.e., \( [\text{M}(n)\text{X}_{6}]^{4-} \)), \( \text{M} = \text{Cd}, \text{Mn} \) and/or Mn), sandwiched between the two layers of trivalent metal halide octahedra (i.e., \( [\text{M}(n)\text{X}_{6}]^{3-} \)), \( \text{M} = \text{Bi} \). The overall tri-layer motifs are separated by top and bottom vacancy (V) layers [an alternating M(n–M(n–)M(n–)M(n–)V pattern along the [001] crystal direction (the original [111] direction of the cubic lattice) of the LDP, as shown in Fig. 1a.\(^{3,8}\)

The visual appearances of as-synthesized bulk powders are demonstrated in Fig. 1c and d. The Mn-substituted powders showed intense orange emission, whereas pristine powders displayed low intense purple colour under UV light. To investigate the environmental stability, the PXRD patterns of bulk pristine and Mn-substituted powders were monitored after 30 days of exposure to the ambient conditions (Fig. S1a and b, ESI†), which confirmed the absence of degradation. The thermal stabilities of these mechanochemically synthesized bulk powders were investigated by thermogravimetric analysis (TGA), as shown in Fig. S2a (ESI†). Both the samples showed stability up to 400 °C, after which approximately 19% and 23% weight losses occurred at \( \sim 450 \) °C for pristine and Mn-substituted analogue, respectively, and further decomposition has been observed thereafter. The characteristic Raman active vibrational modes arising from the different metal octahedra in pristine and Mn-substituted samples were ascertained from the implementation of Raman spectroscopy (Fig. S2b, ESI†). For the pristine sample, the Raman active vibrational bands at 118, 268, and 291 cm\(^{-1} \) appeared due to the
Mn-substituted analogue, respectively. Extrapolating the linear region, which appeared to be
were estimated from the onset of the absorption edge by
presented in Fig. 2a. The optical band gaps of both samples

Electronic d transitions in the high spin Mn 2+ ion, respectively. The
X-ray photoelectron spectroscopy (XPS) measurement
symmetric stretching of the Mn
Cl bond.

Peaks in the visible region at
490 and 430 nm, whereas these
peaks are absent in the pristine sample (Fig. S4, ESI†).

Moreover, the emission
peak maxima in PL spectra of the Mn-substituted sample were
found to be independent of the excitation wavelength as shown
in Fig. 2d, which designates a single radiative decay pathway.†,10 Furthermore, the PL excitation (PLE) spectrum of
both samples monitored at λmax matched well with the
absorption spectrum (Fig. 2b).† Moreover, the broadband and
highly Stokes shifted PL emission of the Mn-substituted sample
could be due to the energy transfer from the intrinsic dark
self-trap excitonic (STE) state to the Mn 2+ luminescent centre.†

We have employed the solution-based hot-injection method
for the colloidal synthesis of pristine and Mn-substituted
nanostructured samples. In a typical synthesis of
Cs4Cd0.6Mn0.4Bi2Cl12, an appropriate proportion of Cs4CoOx
Cd(OAc)2·2H2O and/or Mn(OAc)2 and Bi(OAc)3 was dissolved in
1-octadecene (ODE), following which oleic acid (OA) and isopropanol. The detailed synthesis strategy is presented in
Scheme S1 (ESI†). The PXRD patterns of the as-synthesized
samples. In a typical synthesis of
Cs4Cd, Mn, Bi and Cl) in the Mn-substituted sample (Fig. S3a–e, ESI†).

The optical properties were investigated for both pristine
and Mn-substituted bulk powders by diffuse reflectance and
PL spectroscopic techniques in the solid state. Electronic
absorption spectra of pristine and Mn-substituted samples are
presented in Fig. 2a. The optical band gaps of both samples
were estimated from the onset of the absorption edge by
extrapolating the linear region, which appeared to be ∼372 nm
(∼3.33 eV) nm and ∼408 nm (∼3.08 eV) for pristine and
Mn-substituted analogue, respectively.† The absorption spectrum of the Mn-substituted sample involves two extra weaker
peaks in the visible region at ∼490 and ∼430 nm, whereas these
peaks are absent in the pristine sample (Fig. S4, ESI†).

The additional peaks in the Mn-substituted sample can be assigned
to the spin-forbidden 6A1g(S) → 1T2g(G) and 6A1g(S) → 1T1g(G) d–d transitions in the high spin Mn2+ ion, respectively. The
solid-state room-temperature PL spectra of the bulk pristine
sample displayed a PL band centred at 601 nm (Fig. 2c), while
the bulk Mn-substituted sample showed highly intense broad
PL centred at 595 nm with an emission line-width of 60 nm
(Fig. 2c). The representative orange PL emission of the
Mn-substituted sample is ascribed to the spin and parity
forbidden 1T1g (G) → 6A1g (S) transition within the 3d shell of
the octahedrally coordinated Mn2+.† Moreover, the emission
peak maxima in PL spectra of the Mn-substituted sample were
found to be independent of the excitation wavelength as shown
in Fig. 2d, which designates a single radiative decay pathway.†,10 Furthermore, the PL excitation (PLE) spectrum of
both samples monitored at λmax matched well with the
absorption spectrum (Fig. 2b).† However, the broadband and
highly Stokes shifted PL emission of the Mn-substituted sample
could be due to the energy transfer from the intrinsic dark
self-trap excitonic (STE) state to the Mn2+ luminescent centre.†

We have employed the solution-based hot-injection method
for the colloidal synthesis of pristine and Mn-substituted
nanostructured samples. In a typical synthesis of
Cs4Cd, Mn, Bi and Cl) in the Mn-substituted sample (Fig. S3a–e, ESI†).

The detailed structural analysis was carried out
Mn0.4Bi2Cl12 (Mn-substituted) nanostructured samples. (b) and (c) TEM images of pristine and Mn-substituted nanosheet (NS) samples, respectively. (d) and (f) depict the HRTEM images, and (e) and (g) represent the SAED patterns of the respective NS samples. (h) and (i) represent AFM images of pristine and Mn-substituted NS samples, respectively.

using the composition obtained from ICP-AES analysis (i.e., Cs4Cd0.67Mn0.33Bi2Cl12), and the pristine sample by employing Rietveld refinement [Fig. S5 and Tables S2–S3 (ESI†)]. The PXRD pattern of both the nanostructured samples after 30 days of exposure to ambient conditions suggests decent environmental stability (Fig. S6, ESI†). The final products were dispersed in toluene and checked under ambient light and UV light illumination. The toluene-dispersed Mn-substituted nanostructured sample exhibited strong orange emission as compared to the low-intense purple emission of the pristine sample under UV light, as presented in Fig. S7a (ESI†).

To obtain information on the presence of added ligands (OA and OAm) on the surface of nanostructures, Fourier transform infrared spectroscopy (FTIR) was employed. Both samples showed similar FTIR spectra, as shown in Fig. S7b (ESI†). The FTIR confirmed the characteristic modes of the oleyl group [CH3(CH2)7–CH=CH–(CH2)8–], which exists in both OA and OAm. The peaks at ∼2854 and 2920 cm⁻¹ correspond to the symmetric and asymmetric stretching modes in –CH2, respectively. The νc=c stretching mode is also evident as a less intense peak at 1635 cm⁻¹. As expected, the 1710 cm⁻¹ peak appeared from the νc=o of the carboxylic group of OA. The 3175 cm⁻¹ and 1517 cm⁻¹ modes arose from the stretching and scissoring of NH2 groups of OAm, respectively. Moreover, the distinct 1460 cm⁻¹ and 1378 cm⁻¹ peaks appeared from the C–H bending vibration of the hydrocarbon chain of OA/OAm.⁴⁶ Therefore, the FTIR study confirms the existence of the ligands on the surface of nanostructured samples.

Transmission electron microscopy (TEM) revealed the 2D nanosheet (NS) morphology for both pristine and Mn-substituted samples (see Fig. 3b and c, respectively). High-resolution TEM (HRTEM) analysis revealed the d-spacing values from lattice fringes to be 0.385 nm and 0.277 nm (Fig. 3d and f), which correspond to the (110) and (119) crystal planes of pristine and Mn-substituted NSs, respectively. The selected area electron diffraction (SAED) patterns of the pristine and Mn-substituted NSs further confirm the single crystalline nature, with the diffraction spots corresponding to the [1–12] and [111] zone axis of the rhombohedral phase, respectively (Fig. 3e and g). Additionally, the atomic force microscopy (AFM) analysis was executed to investigate the thickness of NSs (Fig. 3h and i), in which the height profiles were found to be 7.33 and 11.35 nm in pristine and Mn-substituted NSs, respectively. These observations suggested that the NSs are two to three-layers thick (thickness of one layer corresponds to 3.35 nm). The lateral dimension of the NS ranges from 300 to 700 nm. The field-emission scanning electron microscopy (FESEM) images and EDAX elemental mapping for the two samples further confirmed the uniform distribution of all elements (Cs, Cd and/or Mn, Bi, and Cl) in both samples (Fig. S8 and S9, ESI†). Subsequently, the Mn-substituted NS sample was characterized by the electron paramagnetic resonance (EPR) measurement, as illustrated in Fig. S10 (ESI†). The single broad EPR resonance peak without any hyperfine splitting might be ascribed to the stronger magnetic interactions. The obtained g value of ~1.99 suggests the presence of Mn²⁺ (electronic configuration: [Ar]3d⁸, g ~ 2.0) in the corresponding NS sample.⁶⁶

The optical properties of pristine and Mn-substituted NS samples were examined in the solution phase (dispersed in toluene). The UV-vis absorption spectra in Fig. S11a (ESI†) display a distinct and sharp absorption band maximum at ~335 nm for both samples, which probably signifies the presence of an excitonic peak, a representative of 2D layered materials.⁷ The absorption maximum appeared due to the electronically spin-forbidden 1S0 → 3P1 transition of Bi³⁺ ion.⁴⁶ While previous reports showed no noticeable PL emission at room temperature for Cs4CdBi2Cl12 NCs,⁷ we have obtained room temperature weak PL emission for the pristine NS sample, which is centred at 605 nm irrespective of the excitation wavelength (Fig. 4a). The peak possibly appears from the 3Eg → 1A₁g electronic transition of the Cd²⁺ ion in the [CdCl₄]²⁻ octahedral unit.⁹ Conversely, an intense PL emission centred at 601 nm was observed for the Mn-substituted NS sample, which also remains unaffected by the different excitation wavelengths (Fig. 4a). This intense orange emission can be ascribed to the 1G₄ (G) → 1A₁₈ (S) electronic transition of Mn²⁺ ions.⁶⁸ The PL emission of Mn-substituted NSs revealed a broad and large Stokes shifted emission without any overlap between its excitation and emission spectra. The large Stokes shifted broad PL emission of the pristine sample indicates that the emission originates from a characteristic STE state, whereas the broadband and Stokes-shifted PL emission of Mn-substituted NSs appears because of the proficient energy transfer from the STE state to the Mn²⁺ centre.⁷⁵ Fig. S11b and c (ESI†) present the PLE spectra (monitored near PL peak emission) of both samples, which displayed nearly overlapped profiles with the corresponding absorption spectra. The quantum yield (QY) of
both NS samples (dispersed in toluene) was measured at room temperature. The QY of pristine NSs was found to be 0.27%, whereas a relatively high quantum yield of 20.56% was recorded for the Mn-substituted NS sample. To the best of our knowledge, this is the highest PLQY of any of the 2D NS samples of Pb-free LDPS. Previously, Yang et al. reported a QY of 4.6% for Cs\textsubscript{4}[Cd\textsubscript{1–x}Mn\textsubscript{x}]Bi\textsubscript{2}Cl\textsubscript{12} NCs. The PLQY values of other earlier reported LDPS, such as Cs\textsubscript{4}Ba\textsubscript{0.12}Ca\textsubscript{0.88}SbCl\textsubscript{12} NCs, Cs\textsubscript{4}CuIn\textsubscript{2}Cl\textsubscript{12} NCs, and Cs\textsubscript{4}MnBi\textsubscript{2}Cl\textsubscript{12} NCs are found to be 3.90%, 1.70%, and <1%, respectively. Additionally, the PLQY values of earlier reported perovskite halide NS [or nanoplate (NPL)] samples were compared with the PLQY values of our NS sample along with their thickness values (Table S5, ESI†).

To examine the temperature stability of the NS sample, we performed temperature-dependent PL measurement of Mn-substituted NSs dispersed in toluene in the 25–85 °C temperature range. The PL spectra at 85 °C revealed a relative loss of only 26% of its initial intensity (Fig. S12a, ESI†), which can also be witnessed from the appearance of the sample under UV illumination (Fig. S12a,† inset). Furthermore, we added up to 120 μl of water into 3 ml of the NS sample in toluene and monitored the PL with gradual addition of water (Fig. S12b, ESI†). The PL spectra and visual appearance shown in Fig. S12b (ESI†) indicate a 57% relative loss of its initial intensity and a weak orange emission, respectively. These observations again lead to the relatively high temperature- and reasonable water stabilities of the as-synthesized Mn-substituted NS sample and show that this can be a promising candidate for optoelectronic application, comparable to the conventional Pb-based 3D perovskite NCs.

For better understanding of the charge carrier dynamics in these NS samples, PL measurement was performed at cryogenic temperature (77 K). Fig. 4b displays the low-temperature PL spectra for both the pristine and Mn-substituted NSs, respectively. An enhanced PL intensity and red-shifted emission towards the near-infrared (NIR) region were observed for both the samples at 77 K in comparison to room temperature PL. Similar trends in PL with temperature were reported earlier for Cs\textsubscript{2}BiBr\textsubscript{4} NCs. The enhanced PL intensity at 77 K for both samples is attributed to the suppression of the thermally activated non-radiative recombination pathways. The red-shifted PL events for Mn-substituted NSs were originated from the enhanced crystal field strength owing to the lattice contraction at low temperature, which resulted in reduced energy of the \( ^{4}T_{1g} \rightarrow ^{6}A_{1g} \) transition and a subsequent, red-shifted energy peak. The suppression of electron–phonon coupling at low temperature might be responsible for the emission line-width narrowing in the Mn-substituted sample.

In contrast, a broader PL signature was witnessed for pristine NSs at 77 K, which have been fitted with a Gaussian line-shape function into two contributions (Fig. S13, ESI†). The low-energy PL contribution is ascribed to the additional recombination from intragap localized states (i.e., shallow-defect states) that cause the unusual spectral broadening. Such observations were previously reported for Pb-based perovskites.

To obtain insights into the excitonic recombination dynamics, time-resolved PL studies were carried out for both NS samples at 298 K and 77 K, as depicted in Fig. 4c and d. The PL decays of pristine and Mn-substituted NS samples are fitted with the biexponential function. The decay curve can be represented by using the following equation:

\[
A(t) = A_1 + A_2 \exp\left(-\frac{t}{\tau_1}\right) + A_2 \exp\left(-\frac{t}{\tau_2}\right)
\]

where \(A\) denotes the emission intensity, and \(A_1\) and \(A_2\) are the fitting parameters describing the relative amplitude for varied lifetime components, while \(\tau_1\) and \(\tau_2\) are the respective lifetime components. The details of the fitted lifetime parameters with their relative proportions are summarized in Table S5 (ESI†), in which the lifetimes fall in the order of the microsecond time scale due to the forbidden nature of the involved transitions. Notably, the PL decay plots for both NS samples in Fig. 4c and d suggested slower decay dynamics and a prolonged average lifetime (\(\tau_{avg}\)) at 77 K than that at 298 K (Table S5, ESI†), which might be due to the higher probability of radiative recombination, and subsequent elimination of non-radiative recombination channels. Furthermore, to verify the presence of the trap-state contribution at 77 K for pristine NSs, we have measured the lifetime by monitoring at two deconvoluted PL contributions, and a much shorter \(\tau_{avg}\) was achieved when monitored at low energy PL contribution. Previously reported Cs\textsubscript{2}Mn\textsubscript{0.12}Bi\textsubscript{1–x}Cl\textsubscript{12} NC samples also exhibited similar enhancement in the average lifetime at cryogenic temperature compared to that at 300 K.

In order to investigate the PL properties of individual single crystalline NSs, we acquired fluorescence images via super-resolved structured illumination microscopy (SIM) after depositing a diluted solution of NSs onto a glass coverslip. Fig. 5a and d show the bright-field optical images of the NSs, where spatially separated NSs can be clearly observed on the
Cs4Cd0.6Mn0.4Bi2Cl12 (Mn-substituted) nanosheet (NS) samples where determined by using the (FWHM) from the Gaussian size distribution profile (Fig. S15, ESI†) of the individual NS. The full width at half maximum (FWHM) from the Gaussian fit of this cross-sectional profile was reported as the size of the NSs. In an unbiased fashion, we have picked ~40 particles from the imaging window and measured the FWHM for determining the size distribution of the NSs. In this respect it should be noted that the typical achievable lateral (xy) resolution of SIM is around 100–120 nm. On the other hand, from our image analysis we found that even the smaller size NSs were >200 nm (Fig. 5c). This indicates that the size of NSs was well above the resolution limit of the SIM methods of imaging. Therefore, we believe that in this case, SIM images are capable of providing near accurate size estimation and it is comparable to the data obtained from the TEM analysis (Fig. 3b and c). Importantly, given that the average size from fluorescence images (∼400 nm) matches with that from other techniques (e.g., TEM), it ruled out the possibility of analysis being performed on aggregated NSs. In addition, a quantitative estimation from the fluorescence intensity plot in Fig. 5f clearly reflects a considerably higher average intensity from the Mn-substituted NSs compared to the pristine NSs. Moreover, we have also calculated the PL intensity vs. particle size data for a large number of NSs of the pristine and Mn-substituted samples. These data have been represented in a scatter plot in Fig S16,† which also indicated a considerably large fluorescence intensity in the case of Mn-substituted NSs with similar size particles.

Finally, to acquire deeper insights into the PL characteristics of the pristine and Mn-substituted NSs, we investigated temporal instability of PL from individual single crystalline NSs under continuous excitation. In general, the nature of time-dependent PL intermittency (or “blinking”) is dependent on the metastable non-radiative recombination sites.23 This further results in a low-emissive dark (OFF) state or quenched PL state (active), whereas a smaller number of such traps provides an enhanced photo-stability in the intense high emissive (ON) state or bright PL state (in-active).23,24 These low-emissive states in blinking events are a major setback in wide-ranging applications, such as in LEDs and biological settings for tracking single biomolecules.24 Therefore, it is important to characterize the temporal instability of PL behaviour of the as-synthesized NSs under constant illumination. For the fluorescence time-trace analysis, we continuously collected emission (video) over 100 s for pristine and Mn-substituted NS samples individually under the total internal reflection fluorescence (TIRF) setting (videos S1 and S2, respectively, ESI†). Fig. 6a and d depict the representative NSs that were considered for the temporal PL instability study in the
was observed earlier for halide perovskite-semiconductor suppressed PL to arrest the dark OFF-state. Such a rare behaviour with absence of the non-zero OFF state and a distinctly suppressed result from the trap state contribution (Table S3, ESI†).

Interestingly, from the time-dependent PL traces up to 120 s in Fig. 6e and the recorded video (video S2, ESI†) for Mn-substituted NSs, no significant decrease in luminescence intensity from PL-versus-time traces was observed from a single particle. The PL intensity is found to slightly fluctuate between the high intensity ON states, which is well-above the background intensity level. A zoomed in plot within 50 s of the high intensity ON states, which is well-above the continuous laser radiation. This blinking-like nature can be presumably ascribed to the presence of trap states that results in the OFF state.† This is further supported by the time-resolved PL decay at room temperature, where nearly 50% contribution in the life-time originates from the faster component ($\tau_1$), which resulted from the trap state contribution (Table S3, ESI†).

The weak PL emission in pristine NSs is induced by the dynamic equilibrium between active and in-active states of the metastable non-radiative recombination channels, causing PL fluctuations or a blinking-like nature. However, in partially Mn-substituted NSs, the PLQY is much higher, which means the number of non-radiative centers is much smaller, but we still see some PL fluctuation like events. Similar defects are probably still present in Mn-substituted NSs; however, their number is much less, and these non-radiative centers are mostly in the passive state. Thereby, partial Mn$^{2+}$ substitution seems to stabilize the in-active state of the non-radiative centers, which leads to an intense PL emission, and seldom and short-lived off states.

Conclusions

Facile solvent-free mechanochemical synthesis and optical property investigations have been demonstrated for the bulk powder of pristine and its partially Mn-substituted analogue. Importantly, a modified hot-injection route was implemented to synthesize the 2D NSs of pristine and Mn-substituted analogue. A weak PL emission is perceived for pristine NSs at room temperature with a QY of < 1%. Conversely, the Mn-substituted NSs showed an intense orange emission at room temperature together with a QY of ~ 21%. The steady-state and time-resolved PL measurements at cryogenic temperature (77 K) revealed the involvement of trap states in weak PL emission of the pristine sample, which is substantially reduced by the addition of Mn$^{2+}$ in partially Mn-substituted NSs, resulting in an intense PL. Finally, super resolution fluorescence microscopy was implemented to understand the single nanosheet’s PL properties in the pristine NSs and the Mn-substituted analogue. The pristine NS exhibited illumination induced photo-bleaching and PL blinking-like nature, whereas the Mn-substituted NS analogue demonstrated suppression of PL fluctuations with negligible photo-bleaching behaviour. The PL intensity traces with time at the single NS level established that the introduction of Mn$^{2+}$ in pristine NSs can stabilize the in-active state of the metastable non-radiative channels. Characterized by an intense orange emission with suppressed PL fluctuations and negligible photo-bleaching, the Mn-substituted NSs demonstrate their unique optical properties at the single NS level, which might be utilized for further optoelectronic applications in the future.

Data availability

All data are available in the manuscript and in the ESI†.

Author contributions

K. B. conceived the idea and designed the study. A. B., K. K., P. A., J. P. and K. B. carried out the synthesis, structural and optical measurements, and other analyses. R. S., S. K. and S. S. A. studied the photoluminescence blinking properties. K. K. wrote the first draft and everyone contributed to editing the manuscript.

Conflicts of interest

The authors declare no conflict of interest.

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