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Rotational dynamics of diammonium cations in lead bromide composites investigated by quasi-elastic neutron scattering

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We report results from quasielastic neutron scattering (QENS) measurements of the dynamical nature of diammonium cations in the two-dimensional (2D) metal halide perovskites (MHPs) (1,3-PDA)PbBr₄ (PDA: phenylenediammonium), (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄ (XDA: xylylenediammonium), and in the zero-dimensional (0D) perovskitoid (1,3-XDA)₂PbBr₆. QENS spectra measured upon heating from 44 to 350 K reveal the onset of picosecond timescale dynamics of the respective organic cation at around 225 K for 1,3-PDA, 250 K for 1,4-PDA, 250 K for 1,3-XDA, and 350 K for 1,4-XDA. Analyses of the elastic incoherent structure factor of the materials suggest that the observed dynamics can be assigned to three-fold (C₃) and/or continuous rotational jump-diffusion dynamics of the terminal –NH₃ groups of the respective organic cation for all materials. An average, apparent, residence time of the jump-diffusion dynamics has been extracted from the QENS data and takes values of about 1 ps for (1,3-PDA)PbBr₄, 4–5 ps for (1,4-PDA)PbBr₄ and (1,3-XDA)Pb₂Br₆, and 10 ps for (1,4-XDA)PbBr₄ at 350 K. A comparison of the dynamical results with the length and symmetry of the organic cations suggests that a smaller organic cation (here PDA) and an asymmetric position of the two –NH₃ groups (here 1,3-PDA and 1,3-XDA) correlate with a lower onset temperature and faster dynamics. A comparison of the dynamics results with the photoluminescence (PL) spectra of the materials may indicate that slower –NH₃ dynamics correlates with a lower thermal stability of PL due to less dynamic disorder.

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1. Introduction

Layered, or 2D, metal halide perovskites (MHPs)—particularly those of the Ruddlesden–Popper type A₂A'_{n-1}M_nX_{3n+1} and the Dion–Jacobson type AA'_{n-1}M_nX_{3n+1}—are attracting considerable interest for optoelectronic applications.^{1–4} In these structures, A and A' are organic cations, and n denotes the number of stacked inorganic layers composed of octahedral MX₆ units, where M is a metal cation and X is a halide ion. Amongst the 2D MHPs developed so far, most are based on monoammonium cations, whereas diammonium cations have been less explored.^{5–9}

Recently, a new class of Dion–Jacobson type MHPs, with A = 1,3-phenylenediammonium (1,3-PDA, PDA = C₆H₅ (NH₃)₂²⁺), 1,4-phenylenediammonium (1,4-PDA), and 1,4-xylylenediammonium (1,4-XDA, XDA = C₆H₅ (CH₂NH₃)₂²⁺), M = Pb, and X = Br, as well as the chemically similar but structurally dissimilar material (1,3-XDA)₂PbBr₆, were reported.¹⁰ The latter

material exhibits isolated octahedral PbBr₆ building blocks and may therefore be described as a zero-dimensional (0D) perovskitoid; see Fig. 1 for the crystal structure of each material. While all four materials show PL in the visible wavelength region upon excitation at 300 nm,¹⁰ the nature of the PL varies markedly between the different materials. Specifically, the materials with the relatively short organic cations, (1,3-PDA)PbBr₄ and (1,4-PDA)PbBr₄, exhibit relatively broad PL in the wavelength range of 450–750 nm, whereas the material with the relatively long organic cation, (1,4-XDA)PbBr₄, exhibits a narrower PL in the wavelength range of 400–450 nm;¹⁰ the PL of (1,3-XDA)₂PbBr₆ has not yet been reported. Furthermore, the material with the more asymmetric organic cation, (1,3-PDA)PbBr₄, generally exhibits a weaker PL compared to (1,4-PDA)PbBr₄ and (1,4-XDA)PbBr₄, which have been correlated to the degree of distortion of the octahedral PbBr₆ building blocks.¹⁰

In addition to the correlation between the degree of octahedral distortion and PL properties, recent results based on QENS experiments on similar 2D MHPs, such as (nBA)₂PbBr₄, (ODA)PbBr₄ and (GABA)₂PbBr₄, where nBA = n-butylammonium, ODA = 1,8-diaminooctammonium, and GABA = 4-aminobutyric acid, suggest that the PL properties are affected by the reorientational motions of the organic cations.¹¹ In particular, it has been argued that the relatively smaller organic cation nBA is more

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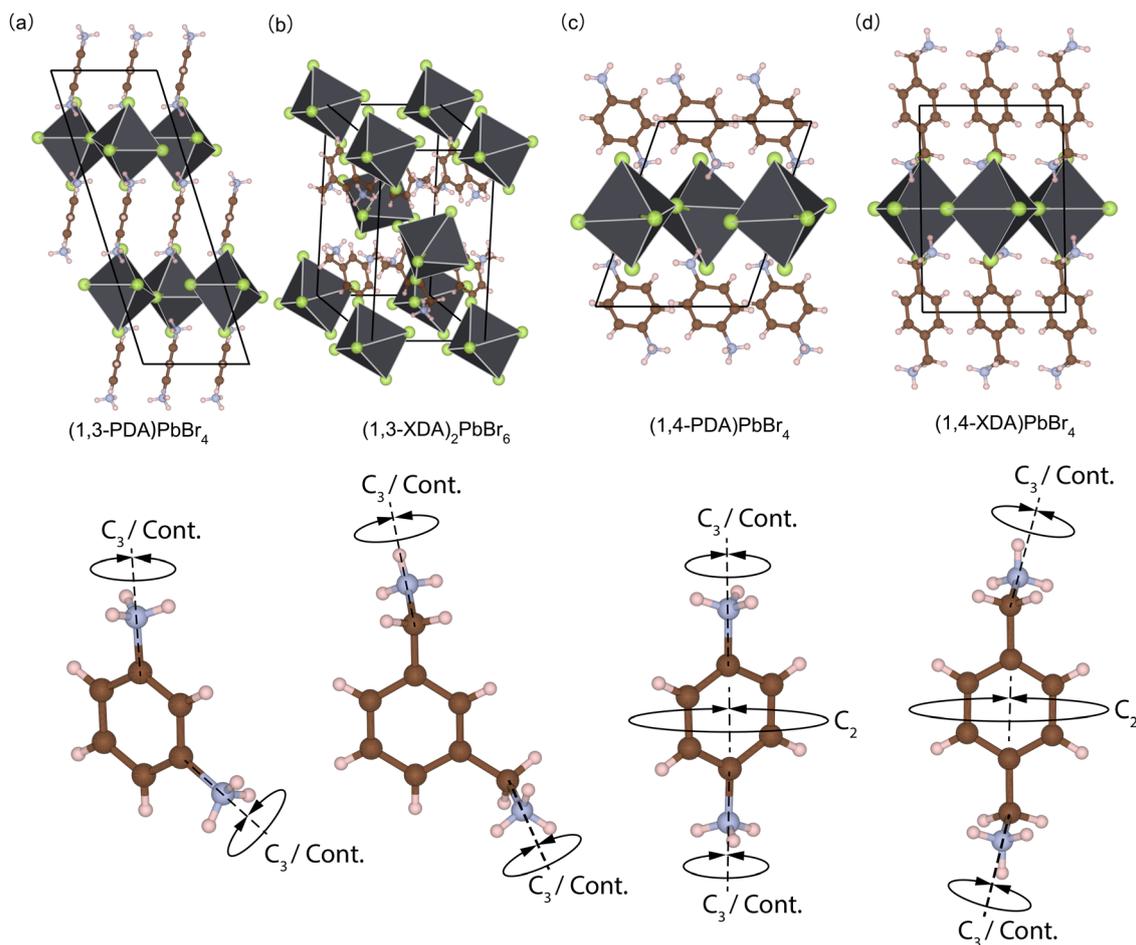



Fig. 1 Illustration of the crystal structures and organic cations of (a) (1,3-PDA)PbBr₄ (monoclinic crystal structure with space group $C2/c$), (b) (1,3-XDA)₂PbBr₆ (monoclinic crystal structure with space group $P2_1/c$), (c) (1,4-PDA)PbBr₄ (triclinic crystal structure with space group $P\bar{1}$), and (d) (1,4-XDA)PbBr₄ (monoclinic crystal structure with space group $P2_1/c$), according to ref. 10. The black lines refer to a unit cell. The lower part of the figure illustrates the C₃ and continuous (Cont.) rotational diffusion of the terminal –NH₃ groups, as well as C₂ rotational diffusion of the phenylene group (–C₆H₄–) along its long molecular axis for (1,4-PDA)PbBr₄ and (1,4-XDA)PbBr₄ (see the Results section). Br atoms are indicated as green spheres, C atoms as brown spheres, N atoms as light blue spheres, and H atoms as light pink spheres. The Pb atoms located inside the PbBr₆ octahedra are omitted. The crystal structures were plotted using VESTA.¹²

dynamic inside the inorganic sublattice, which gives rise to a larger “dynamic radius”, which yields a closer to ideal A-site geometry, preventing local structural distortions and broadband PL.¹¹

In this work, we investigate the reorientational dynamics of the organic cations in the novel 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and in the 0D perovskitoid (1,3-XDA)₂PbBr₆ using QENS. The aim of this study is to determine the onset temperature and geometry of the dynamics, how it correlates with the materials’ PL properties, and how it compares with the organic cation dynamics in other, related, MHPs.

2. Experimental

2.1. Sample synthesis

Single crystals of (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, (1,3-XDA)₂PbBr₆ and (1,4-XDA)PbBr₄ were grown by dissolving lead(II) acetate powder in a large excess of 48% w/w aqueous

HBr and 50% w/w aqueous H₃PO₂, and heating the mixture to 140 °C. After solid dissolution, a stoichiometric amount of solid diamine (liquid for 1,3-XDA) was added. Subsequently, crystals formed upon slow cooling to room temperature. X-ray diffraction data, obtained using a Bruker D8 Discover with a Cu source, and energy-integrated neutron scattering data both confirm phase purity and that the samples adopt the crystal structures previously reported for these materials; see Fig. S1 in the SI.

2.2. Quasielastic neutron scattering

The QENS experiment was performed on the direct geometry time-of-flight spectrometer FOCUS at the continuous spallation neutron source SINQ at the Paul Scherrer Institute in Switzerland.^{13,14} The samples, approximately 1 g per composition, were contained in an aluminium foil rolled cylinder with an expanded size of 5 cm × 4 cm, which was sealed in an aluminium cylinder can with a lead wire. The instrument was



set up using an incident neutron wavelength ($\lambda_{\text{incident}}$) of 4 Å. By using this setting, the energy resolution (determined as the full-width at half maximum (FWHM) of the elastic line) was 0.2 meV, the accessible momentum transfer (Q) range was 0.4–2.5 Å⁻¹, and the observation time window of the dynamics was ~0.2–13 ps. Measurements were performed at temperatures (T) of 44 K, 200 K, 225 K, 250 K, 300 K, and 350 K, in that order. The measurement time at each temperature was approximately 2 h.

The data reduction was performed using the DAVE software.¹⁵ Data from the measurement of an empty sample cell at the same temperatures as those used for the samples were subtracted from the data for the samples, and all data were divided by the data from the measurement of a vanadium standard at 300 K to correct for variations in the efficiency of the different detectors. The measured scattering intensity is proportional to the dynamical structure factor $S(Q, E)$, which results from the sum of coherent and incoherent scattering contributions. The $S(Q, E)$ data were sliced into Q regions of 0.15 Å⁻¹ width.

3. Results

3.1. Dynamical structure factors

Fig. 2 shows the dynamical structure factors, $S(Q, E)$, as integrated over the measured Q -range, for temperatures between 200 and 350 K, together with the instrumental resolution function, $R(E)$. The latter is approximated with the spectrum measured at 44 K, where QENS studies of similar materials have shown that the diffusion dynamics of organic cations on the picosecond timescale are frozen-in.^{11,17} Note that, for (1,3-PDA)PbBr₄ there was a problem in the data collection at 44 K and, therefore, for the analysis of the neutron scattering data for this sample, the 44 K data for (1,3-XDA)₂PbBr₆ were used.

Our general observation is that the spectra for $T > 200$ K are markedly different from the spectra of the resolution function.

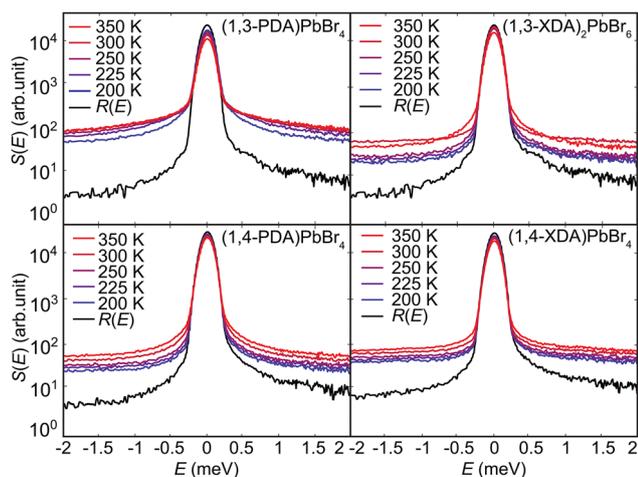


Fig. 2 Variable temperature Q -integrated dynamical structure factors, $S(E)$, for the 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and the OD perovskitoid (1,3-XDA)₂PbBr₆.

The onset of quasielastic scattering is observed at approximately 225 K for (1,3-PDA)PbBr₄, 250 K for (1,4-PDA)PbBr₄, 250 K for (1,3-XDA)PbBr₆, and 350 K for (1,4-XDA)PbBr₄. In this context, we note that H contributes approximately 90% for (1,3-PDA)PbBr₄, 90% for (1,4-PDA)PbBr₄, 93% for (1,3-XDA)₂PbBr₆, and 92% for (1,4-XDA)PbBr₄ to the total scattering cross section of the investigated materials, and H scatters mainly incoherently (*cf.* 80.3 barns for the incoherent scattering cross section and 1.8 barns for the coherent scattering cross section),¹⁶ which suggests that the quasielastic signal originates from the dynamics of the H of the organic cations. As reported in some other QENS studies of MHPs, strongly damped or overdamped vibrational dynamics of the octahedral units of the inorganic sublattice, here PbBr₆, can also give rise to significant quasielastic scattering.^{18–21} However, such dynamics are usually observed in the vicinity of displacive phase transitions. Given the absence of such phase transitions in the materials studied here, combined with the much smaller contribution from PbBr₆ to the total scattering cross section, it is reasonable to assume that the quasielastic signal can be approximated with the self-dynamics of the H of the organic cations.

In our more detailed analysis, $S(Q, E)$ was fitted according to the following function:

$$S(Q, E) = [I_{\text{el}} \cdot \delta(E) + I_{\text{qe}} \cdot \mathcal{L}(E)] \otimes R(Q, E) + \text{bkg}(Q),$$

$$\mathcal{L}(E) = \frac{1}{\pi} \frac{\gamma}{\gamma^2 + E^2} \quad (1)$$

Here, $I_{\text{el}} \cdot \delta(E)$ is the elastic scattering intensity, which accounts for the intensity at zero energy transfer ($E = 0$). In our case, elastic scattering originates from H atoms that can be considered immobile within the observation time window. $I_{\text{qe}} \cdot \mathcal{L}(E)$ is a Lorentzian function of amplitude I_{qe} and half-width at half maximum (HWHM) γ . \otimes denotes the convolution operation, $R(Q, E)$ is the instrumental resolution function, and $\text{bkg}(Q)$ is an E -independent background. The background relates to (in part) the scattering from phonons, which are centered at energy transfers ≈ 2 meV (see Fig. 3, which shows $S(E)$ for an extended E -range). In comparison, the quasielastic signal typically has a FWHM of less than 1.5 meV, thus not strongly overlapping with the phonon peaks.

Fig. 4 shows, as an example, the fits for $Q = 1.5$ Å⁻¹ and $T = 350$ K for all investigated materials, whereas the fits for all temperatures and selected Q -values are shown in Fig. S2–S5.

The quasielastic FWHM Γ ($\Gamma = 2\gamma$) is (within error) practically Q -independent for all materials (Fig. 5), which suggests that the dynamics associated with the quasielastic signal related to organic cation dynamics are localized in nature, in full agreement with other QENS studies of MHPs performed under similar experimental conditions.^{17,22} A characteristic timescale (τ) of the dynamics has been derived by taking $\tau = 2\hbar/\Gamma$,²³ where $\bar{\Gamma}$ refers to the Q -averaged value of Γ for each temperature and material. The obtained τ values are in the range of 1–10 ps and vary significantly between the four samples (Table 1), with the timescale of the dynamics being fastest for (1,3-PDA)PbBr₄ (1.2 ps at $T = 350$ K) and slowest for



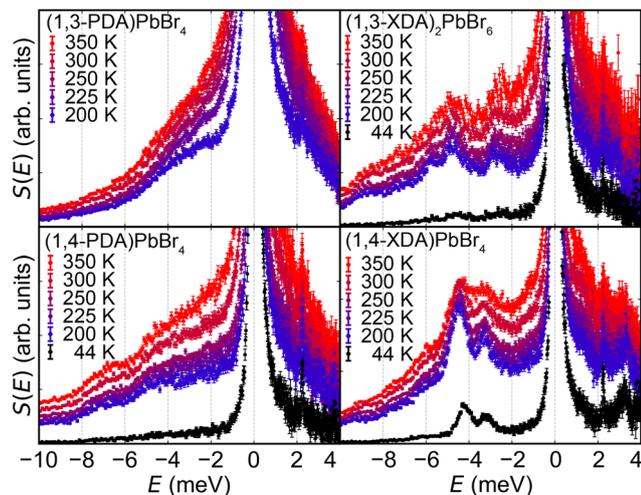


Fig. 3 Variable temperature Q -integrated dynamical structure factors, $S(E)$, for the 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and the 0D perovskitoid (1,3-XDA)₂PbBr₆, showing the quasi-elastic and inelastic regions.

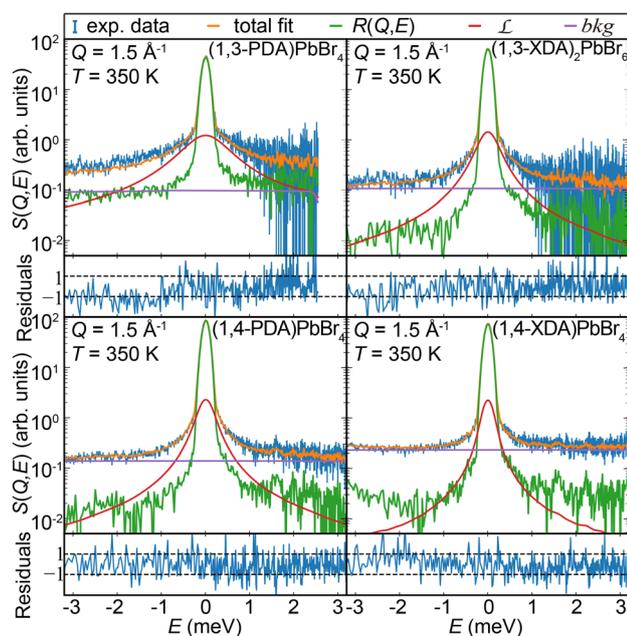


Fig. 4 $S(Q, E)$ for the 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and the 0D perovskitoid (1,3-XDA)₂PbBr₆ at $Q = 1.5 \text{ \AA}^{-1}$ and $T = 350 \text{ K}$.

(1,4-XDA)PbBr₄ (9.5 ps at $T = 350 \text{ K}$). Note that the quasi-elastic signals at $T < 250 \text{ K}$ for (1,4-PDA)PbBr₄ and (1,3-XDA)₂PbBr₆, and at $T < 350 \text{ K}$ for (1,4-XDA)PbBr₄ were too weak to allow a reliable fit according to eqn (1).

3.2. Elastic incoherent structure factors

Information about the spatial geometry of the dynamics can be obtained from the analysis of the elastic incoherent structure factor (EISF), which is defined as $\text{EISF} = I_{\text{el}} / (I_{\text{el}} + I_{\text{qe}})$. Fig. 6

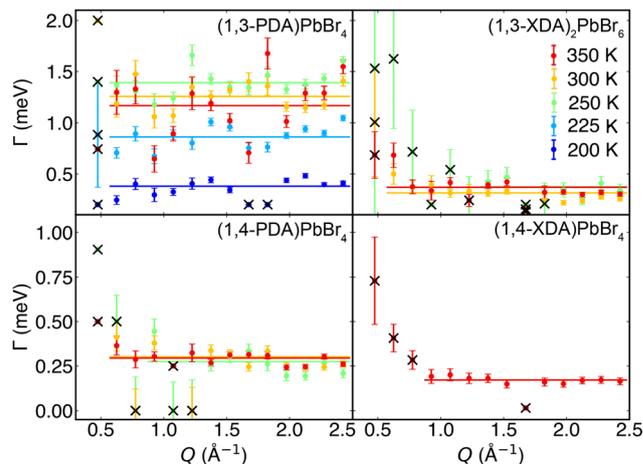


Fig. 5 Quasi-elastic linewidth Γ for the 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and the 0D perovskitoid (1,3-XDA)₂PbBr₆. Lines represent fits to a constant at each temperature. The points marked by cross marks are deemed unreliable outliers and are thus excluded from the fits.

shows a comparison of the EISF extracted from the experimental data and the EISF model curves for different geometrically and, presumably, physically realistic reorientational motions of the organic cations. For all materials and temperatures, we observe an excellent agreement between the experimental data and the model for three-fold (C_3) and continuous (Cont.) rotational diffusion, respectively, of the $-\text{NH}_3$ group. These models are mathematically defined as:

$$\text{EISF}_{C_3} = c + (1 - c) \cdot \frac{1}{n_H} \left[n_H - 6 + 6 \cdot \frac{1}{3} (1 + 2 \cdot j_0(Qd)) \right] \quad (2)$$

$$\text{EISF}_{\text{Cont.}} = c + (1 - c) \cdot \frac{1}{n_H} \left[n_H - 6 + 6 \cdot \frac{1}{N_{\text{max}}} \sum_{N=0}^{N_{\text{max}}} j_0(Qd_N) \right] \quad (3)$$

Here, n_H is the number of hydrogens in an organic cation, $j_0(x)$ is the zeroth-order spherical Bessel function, d is the jump distance between two hydrogens in an $-\text{NH}_3$ group, N_{max} is the number of uniformly distributed jump locations on a circle (theoretically, $N_{\text{max}} \rightarrow +\infty$, but here we approximate with $N_{\text{max}} = 20$), and $d_N = D \sin(N\pi/N_{\text{max}})$ is the jump distance between position N and the original position on a circle with diameter D . The constant c accounts for additional elastic scattering, probably mostly due to H that experiences motions too slow to be resolved by the instrument resolution, but also to coherent scattering from the materials manifested as peaks in the energy-integrated neutron scattering data (Fig. S1). As can be seen, both models fit equally well to the experimental data and, hence, cannot be distinguished from each other in the relatively limited Q -range explored here. Other, geometrically feasible models, such as two-fold (C_2) rotation of the whole organic cation around its short molecular axis, or a more complex model that assumes C_3 rotation of the $-\text{NH}_3$ group together with C_2 rotation of the phenylene ring, were also considered; however, all



Table 1 Compilation of the characteristic timescale, τ , and additional elastic scattering, c , values extracted from the QENS data for 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and the OD perovskitoid (1,3-XDA)₂PbBr₆

T (K)	(1,3-PDA)PbBr ₄		(1,3-XDA) ₂ PbBr ₆		(1,4-PDA)PbBr ₄		(1,4-XDA)PbBr ₄	
	τ (ps)	c	τ (ps)	c	τ (ps)	c	τ (ps)	c
200	3.3(2)	0.66(1)	—	—	—	—	—	—
225	1.5(1)	0.55(1)	—	—	—	—	—	—
250	0.9(1)	0.47(1)	3.7(4)	0.91(1)	5.4(5)	0.91(1)	—	—
300	1.1(1)	0.39(1)	5.0(3)	0.71(1)	4.8(3)	0.82(1)	—	—
350	1.2(1)	0.25(1)	4.0(2)	0.54(1)	4.7(2)	0.71(1)	9.5(10)	0.63(1)

of these models showed a worse fit to the data and, hence, were not further considered here (see Fig. S6).

The additional elastic scattering shows a general decrease with increasing temperature, in a quite systematic manner for all materials, which we interpret as an increase in the proportion of organic cations that undergo rotational diffusion of the terminal $-\text{NH}_3$ groups within the observation time window; however, it must be noted that for (1,4-PDA)PbBr₄ we have data at only one temperature (350 K). Furthermore, a somewhat larger fraction of the organic cations seems mobile in (1,3-XDA)₂PbBr₆ and (1,3-PDA)PbBr₄, as compared to (1,4-PDA)PbBr₄ and (1,4-XDA)PbBr₄.

4. Discussion

An important result is the observation of picosecond timescale localized dynamics that can be assigned to C₃ and/or continuous rotational diffusion of the terminal $-\text{NH}_3$ groups of the respective organic cation for all the studied materials. In effect, this means that the spatial geometry of the observed organic cation dynamics is insensitive to the length (here the number of C atoms) and symmetry (here the mutual position of the two $-\text{NH}_3$ groups) of the organic cation, as well as to the dimensionality (2D or 0D) of the surrounding inorganic sublattice, for the materials studied here. In contrast, the onset temperature

and timescale of the observed dynamics vary significantly amongst the four materials. Firstly, when comparing organic cations of the same length, we find that the asymmetric cations exhibit faster dynamics (*cf.* 1 ps for (1,3-PDA)PbBr₄ vs. 5 ps for (1,4-PDA)PbBr₄, and 4 ps for (1,3-XDA)₂PbBr₆ vs. 10 ps (1,4-XDA)PbBr₄). Secondly, when comparing organic cations of the same symmetry, we find that the shorter cations exhibit a faster timescale of dynamics (*cf.* 1 ps for (1,3-PDA)PbBr₄ vs. 4 ps for (1,3-XDA)₂PbBr₆, and 4 ps for (1,4-PDA)PbBr₄ vs. 10 ps for (1,4-XDA)PbBr₄). Taken together, this suggests that, at a given temperature, a shorter organic cation (here PDA) and an asymmetric position of the two $-\text{NH}_3$ groups (here 1,3-PDA and 1,3-XDA) correlate with a lower onset temperature and faster dynamics.

Intuitively, the onset and timescale of the dynamics should be correlated to the strength of the hydrogen-bonding interactions between the $-\text{NH}_3$ groups and the surrounding Br atoms of the PbBr₆ octahedra, but results from density functional theory (DFT) calculations revealed no systematic differences in the electronic distribution of the investigated materials that would promote the formation of either stronger or weaker hydrogen bonds.¹⁰ Instead, we note that (1,3-PDA)PbBr₄ and (1,3-XDA)₂PbBr₆, *i.e.* the materials exhibiting the fastest dynamics, least “immobile” species, and lowest onset temperature of the dynamics, exhibit the largest level of distortion of the PbBr₆ octahedral units, whereas the materials exhibiting the slowest dynamics and highest fraction of “immobile” species, (1,4-PDA)PbBr₄ and (1,4-XDA)PbBr₄, correlate with less octahedral distortion.¹⁰ Additionally, we note that materials having a longer organic cation (XDA) generally exhibit less and slower dynamics compared to their shorter-cation (PDA) counterparts. This feature may be attributed to the additional methylene group ($-\text{CH}_2-$) in the longer cations. More specifically, we speculate that the dynamics of the $-\text{CH}_2-$ group could take up some thermal energy, which could, possibly, explain that the onset temperature of the dynamics is higher for (1,4-XDA)PbBr₄ compared to (1,4-PDA)PbBr₄, and for (1,3-XDA)₂PbBr₆ compared to (1,3-PDA)PbBr₄. Additionally, we note that C₂ rotational diffusion of the phenylene group ($-\text{C}_6\text{H}_4-$) around the long molecular axis may be more readily happening for the more symmetric molecules, 1,4-PDA and 1,4-XDA (see Fig. 1). This may modify the dynamical characteristics of the terminal $-\text{NH}_3$ groups, such that some thermal energy is stored in $-\text{C}_6\text{H}_4-$ dynamics rather than in $-\text{NH}_3$ dynamics, thus further increasing its onset temperature.

In comparison to the literature, the observation of C₃ and/or continuous rotational dynamics of the $-\text{NH}_3$ groups on the picosecond timescale, as observed here, is in agreement with

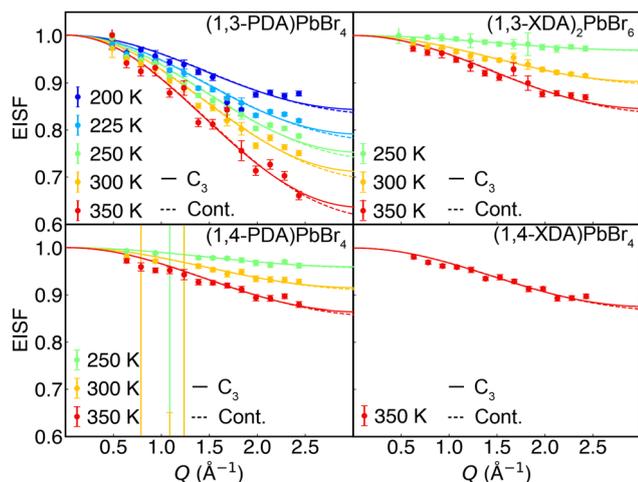


Fig. 6 Experimental EISF (markers) for 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, and (1,4-XDA)PbBr₄, and the OD perovskitoid (1,3-XDA)₂PbBr₆. The solid lines are fits to a C₃ jump-diffusion model, and the dashed lines are fits to a model describing continuous diffusion on a circle.



other QENS studies of MHPs, such as MAPbX₃ (MA = methylammonium, CH₃NH₃; X = I, Br, Cl),^{24–27} and 2D (OA)₂PbI₄ (OA = C₈H₁₇NH₃).¹⁷ However, it is faster than the reported same type of molecular dynamics observed in the 2D MHPs (*n*BA)₂PbBr₄, (ODA)PbBr₄ and (GABA)₂PbBr₄, for which C₃ rotational diffusion of the –NH₃ groups was found to occur on a timescale of 30–230 ps at 275–300 K.¹¹ The slower timescale dynamics compared to our study may be attributed to differences in the observation time window limited by the *E* range because different instruments were used.

Finally, we compare our dynamical results to the PL properties of the materials. In this context, (1,3-PDA)PbBr₄ and (1,4-PDA)PbBr₄ exhibit broadband PL (*ca.* 500–900 nm), whereas (1,4-XDA)PbBr₄ exhibits narrowband PL at 420 nm together with weaker, broadband PL between 500 and 800 nm.¹⁰ The broadband PL of (1,3-PDA)PbBr₄ and (1,4-PDA)PbBr₄ has been assigned to radiative relaxation of self-trapped excitons (STEs) due to their more distorted inorganic sublattice and shows a general, systematic decrease in intensity with increasing temperature.^{10,28} The broad nature of STE emission in MHPs may also be associated with low-energy, anharmonic phonons due to their generally soft lattices.²⁰ In this regard, we notice that the phonon peaks for (1,3-PDA)PbBr₄ and (1,4-PDA)PbBr₄ are broader and have a lower energy compared to (1,4-XDA)PbBr₄, for which the phonon peaks are more distinct and exhibit less contribution close to the elastic line (see Fig. 3). The difference in these phonon properties, as well as structural disorder, may thus correlate with the broadness of the PL of the studied materials.

The systematic decrease in the PL intensity with increasing temperature is generally believed to originate from enhanced electron–phonon processes, which depopulate the light-emitting levels in a non-radiative way.¹⁰ However, the softness of the perovskite lattice of MHPs implies that the lattice dynamically reacts to the organic cation dynamics and *vice versa*, and, therefore, the phonons of the inorganic sublattice and the organic cation dynamics are intricately coupled. We speculate that the increase in the portion of mobile –NH₃ species, as indicated by the decreasing immobile fraction (*cf.* Table 1) and the concomitant increase in local structural fluctuations (dynamic disorder) induced by transient hydrogen-bonding interactions between the –NH₃ groups and the inorganic sublattice, promotes thermal quenching of luminescence. Indeed, previous studies of the 3D MHPs ABX₃ (A = Cs, FA, or MA; B = Sn or Pb; X = I, Br, or Cl)^{29–31} and the 2D MHP (C₄H₉NH₃)₂PbBr₄³² have shown that dynamic disorder may lead to an increased tendency for non-radiative relaxation through luminescence killer centers. For (1,4-XDA)PbBr₄, complexities arising from reabsorption and weak photostability hinder a reliable quantitative analysis of the PL–dynamics relationship.¹⁰ Nevertheless, its comparatively suppressed thermal quenching of luminescence between 77 and 297 K may be related to the relatively slower rotational diffusion of the –NH₃ groups in this material compared to the other materials. Therefore, our results may indicate that slower dynamics yields less dynamic disorder and higher thermal stability of PL.

5. Conclusions

To conclude, our variable temperature (200–350 K) QENS study of the 2D MHPs (1,3-PDA)PbBr₄, (1,4-PDA)PbBr₄, (1,4-XDA)PbBr₄, and the 0D perovskitoid (1,3-XDA)₂PbBr₆ unravelled organic cation dynamics with a characteristic timescale in the range of 1–10 ps that can be attributed to C₃ and/or continuous rotational diffusion of the terminal –NH₃ groups. The proportion of mobile species generally increases with increasing temperature from 200 to 350 K, and a correlation of the dynamical results with the structural properties of the materials suggests that a smaller organic cation (here PDA) and an asymmetric position of the two –NH₃ groups (here 1,3-PDA and 1,3-XDA) correlate with a lower onset temperature and faster dynamics. A comparison of the dynamical results to the PL properties of the investigated materials may indicate a correlation between relatively slow dynamics and little dynamic disorder with high thermal stability of PL; however, it must be noted that the structure–dynamics–PL relationship is likely to be very complex.

Conflicts of interest

There are no conflicts to declare.

Data availability

Data for this article, including the neutron scattering data measured at the Paul-Scherrer Institute (PSI), are available in ref. 33.

Supplementary information (SI): energy integrated neutron scattering data, additional *S(Q, E)* data and fits, and additional EISF models and fits. See DOI: <https://doi.org/10.1039/d5cp04945h>.

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