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Elucidating the Structure and Photophysics of Layered Perovskites through Cation Fluorination

Eelco K. Tekelenburg, Simon Kahmann, Machteld E. Kamminga, Graeme R. Blake, and Maria A. Loi*

1. Introduction

Metal halide perovskites have become a popular class of materials due to their outstanding properties for optoelectronic applications. Layered lead iodide perovskites are denoted by the structural formula A$_2$PbI$_4$, where A is a large organic cation.

Optoelectronic devices based on layered perovskites containing fluorinated cations display a well-documented improved stability and enhanced performance over non-fluorinated cations. The effect of fluorination on the crystal structure and photophysics, however, has received limited attention until now. Here, 3-fluorophenethylnammonium lead iodide ((3-FPEA)$_2$PbI$_4$) single crystals are investigated and their properties to the non-fluorinated ((PEA)$_2$PbI$_4$) variant are compared. The bulkier 3-FPEA cation increases the distortion of the inorganic layers, resulting in a blue-shifted absorbance and photoluminescence. Temperature-dependent photoluminescence spectroscopy reveals an intricate exciton substructure in both cases. The fluorinated variant shows hot-exciton resonances separated by 12 to 15 meV, values that are much smaller than the 40 to 46 meV found for (PEA)$_2$PbI$_4$. In addition, high-resolution spectra show that the emission at lower energies consists of a substructure, previously thought to be a single line. With the analysis on the resolved photoluminescence, a vibronic progression is excluded as the origin of the emission at lower energies. Instead, part of the excitonic substructure is proposed to originate from bound excitons. This work furthers the understanding of the photophysics of layered perovskites that has been heavily debated lately.

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The incorporation of such cations results in the formation of inorganic layers sandwiched between the organic cations. Compared to purely 3D perovskites, mixtures of 2D and 3D perovskites have, for example, been incorporated into solar cells and light emitting diodes to improve their ambient and thermal stability as well as their overall performance.[1-4]

It is important to note that improvements in the stability are prerequisites for the use of metal halide perovskites on an industrial scale. The use of layered perovskites in photovoltaic devices has shown important stabilization effects, and in addition, these low dimensional systems are also particularly promising for light-emitting applications. Layered perovskites can be seen as natural quantum wells, where the charge carriers are confined to the inorganic layers. One of the consequences of the confinement is the formation of excitons with binding energies of hundreds of meV.[5,6] The excitonic nature of these materials results in bright and narrow emission, even at room temperature, which enables highly efficient and color-pure LEDs.[7]

In 2019, several reports showed that the use of fluorinated organic cations can further enhance the photovoltaic device performance and stability compared to non-fluorinated cations.[8–11] In addition, halogen-substituted organic cations have shown interesting photophysics, however, more studies are needed to fully understand their physical properties.[12,13]

Several studies on phenethylammonium lead iodide ((PEA)$_2$PbI$_4$) reported a detailed exciton substructure with multiple photoluminescence (PL) and absorption peaks at cryogenic temperatures.[14,15] The proposed origins vary and include emission from bound excitons,[16,17] vibronic progressions,[13,14] or the formation of distinct exciton polarons.[15,18] The even spacing of the peaks by 40 to 46 meV in both the absorbance and PL spectra was attributed to a single vibronic progression governed by a phonon mode of the organic cation.[14] The energy separation of the peaks could be reduced by substituting different halogens on the phenethylammonium cation, suggesting the importance of a low-energy vibrational mode of the heavier organic cation on the PL structure.[13] However, Thouin et al. reported two distinct exciton polaron peaks, associated with different sets of Raman modes, which excluded the existence of a single vibronic progression.[18] In addition, it was observed that
the spacing of the peaks, $35 \pm 5$ meV in their case, was identical in both butylammonium and phenethylammonium layered perovskites, suggesting an overarching mechanism in this class of materials.\cite{15} These recent reports show that no consensus has been reached regarding the origin of these evenly spaced PL peaks.\cite{13,18} With this work we want to contribute to the scientific debate by studying an unexplored fluorinated derivative of phenethylammonium with high spectral resolution.

In this study, the structural and optical properties of 3-fluorophenethylammonium lead iodide ((3-FPEA)$_2$PbI$_4$) single crystals are investigated and compared to the properties of the prototypical non-fluorinated variant ((PEA)$_2$PbI$_4$). In both cases, temperature-dependent PL spectra reveal that the emission spectra are dominated by free excitons whose luminescence is broadened by the same effective phonon mode associated with the inorganic cage. High-energy emission from hot-exciton resonances, however, exhibits a smaller separation for the fluorinated variant compared to the prototypical compound. In addition, high-resolution PL spectra at 5.4 K reveal a previously unreported substructure. The diverse spectral shifts and the time-resolved behavior of these PL resonances prove that the peaks do not originate from a single vibronic progression, while excitons bound to defects are their proposed origin.

2. Results

Two types of A$_2$PbI$_4$ single crystals, namely 3-fluorophenethylammonium and phenethylammonium lead iodide, were synthesized following the procedure described in our previous work.\cite{19} Figure 1a displays the crystal structure of (3-FPEA)$_2$PbI$_4$ solved in the monoclinic $I2/m$ space group from single crystal X-ray diffraction data collected at 100 K; full details of the structure refinement are given in Table S1, Supporting Information. This structure shows the typical arrangement of alternating organic and inorganic layers found in layered perovskites. The equatorial iodine atoms in the PbI$_6$ octahedra, depicted by the two-tone white and purple spheres in Figure 1a, show partial occupancies over two positions, which indicate disorder or an unresolved superstructure. The Pb atom also shows partial occupancies over two positions. These occupancies stem from two possible ways that the ammonium group on the 3-FPEA molecule forms hydrogen bonds to the inorganic layer, namely a “pinched-in” or “pinched-out” conformation in the $ab$-plane as previously reported for (3-FPEA)$_2$SnI$_4$.\cite{20} Furthermore, the mirror plane perpendicular to the $b$-axis results in two partially occupied orientations of the 3-FPEA molecule. We observe evidence for a superstructure with a doubled $b$-axis, as indicated by the weak and rather poorly defined diffraction spots shown in Figure S1, Supporting Information, but this structural model could not be refined in a satisfactory manner. The superstructure might be the result of local ordering of the organic cations, which was reported to be of long-range nature both in the (3-FPEA)$_2$PbI$_4$ and the (3-FPEA)$_2$SnI$_4$ crystals.\cite{8,20}

The structure of (PEA)$_2$PbI$_4$ at 100 K was solved in the triclinic $P\bar{T}$ space group and exhibits fully ordered octahedral rotations and disordered PEA molecular orientations (Figure 1b and Figure S2, Supporting Information). This results in a basal $ab$-plane that is enlarged by a factor of $a\sqrt{2} \times b\sqrt{2}$ compared to that of (3-FPEA)$_2$PbI$_4$. The unit cell contains a single inorganic layer and PEA bilayer. This is different to the monoclinic $P21/c$ structure that we previously determined for crystals grown using the antisolvent vapor-assisted method, where both the in-plane octahedral rotations and the orientations of the PEA molecules showed disorder over two positions.\cite{21} It is also different to the monoclinic $C2/m$ structure with disordered rotations reported by Calabrese et al. for crystals grown by a simple solvent evaporation method.\cite{22} Disordered rotations were also reported by Du et al. for crystals grown by a layered-solution method, although with a triclinic $P\bar{T}$ structure with a doubled $c$-axis at room temperature.\cite{23}

The $ab$-plane of (PEA)$_2$PbI$_4$ is displayed in Figure 1b with the in-plane Pb-I-Pb angle indicated by $\theta$. $\theta$ affects the band gap and the photophysics of layered perovskites and is influenced by the choice of the organic cation.\cite{13,24} The bulkier fluorinated cation introduces a larger distortion, decreasing the average $\theta$ from 152.24° for (PEA)$_2$PbI$_4$ to 150.23° for (3-FPEA)$_2$PbI$_4$. The tilting of the axial iodides is negligible in both materials, indicated by a small distortion up to 1.8°. In addition, the larger 3-FPEA cation introduces a longer interlayer distance, 16.7 Å versus 16.2 Å, in the direction perpendicular to the inorganic layers.

Whereas grain boundaries can influence the properties of multicrystalline thin films, single crystals allow for studying the intrinsic properties of the material. The single crystals were exfoliated in order to obtain clean surfaces. Figure 1c shows the PL map of a representative (3-FPEA)$_2$PbI$_4$ crystal flake obtained by confocal laser scanning microscopy. As expected from the layered crystal structure, clear step edges are evident as a result of the exfoliation.

Representative PL and absorbance spectra at room temperature are shown in Figure 2a for (3-FPEA)$_2$PbI$_4$ (top) and (PEA)$_2$PbI$_4$ (bottom). Absorption peaks are located at 2.420 and 2.387 eV for (3-FPEA)$_2$PbI$_4$ and (PEA)$_2$PbI$_4$, respectively. The PL for both materials consists of two emission bands: a
narrow emission with peaks at 2.378 and 2.351 eV, and a broad emission with peaks at 1.732 and 1.765 eV for (3-FPEA)2PbI4 and (PEA)2PbI4, respectively. Thus, a small Stokes shift of 42 and 36 meV is observed for the narrow emission, and a Stokes shift of hundreds of meV for the broad emission. We previously showed that the broad emission is due to bulk defect states in the band gap, characterized by a longer PL decay compared to the narrow emission; also see Figure S3, Supporting Information. In this report, we aim to provide a deeper understanding of the narrow emission, which is commonly attributed to radiative recombination of free excitons.

A blue shift of 33 meV of the absorption peak is observed with the use of the fluorinated cation compared to the non-fluorinated cation. This shift is attributed predominantly to the larger in-plane distortion in (3-FPEA)2PbI4, as the larger distortion raises the bottom of the conduction band and lowers the top of the valence band, increasing both the electronic and optical band gap.

The asymmetric shape of the narrow emission toward lower energies suggests additional contributions from emitting states, as highlighted in Figure S4, Supporting Information. To verify this assumption, PL spectra were measured as a function of temperature, as shown in Figure 2b. Upon cooling, the narrow emission shows an initial blue shift followed by a red shift starting at 200 K (Figure S5, Supporting Information). This observation is explained by two competing effects: the blue shift is caused by a reduced exciton–phonon interaction, whilst the red shift is caused by the lattice contraction. The lattice contraction increases the overlap of anti-bonding hybridization of Pb 6s and I 5p orbitals of the valence band maximum and the Pb 6p and I 5p orbitals of the conduction band, which results in a smaller optical band gap. Both materials exhibit a similar behavior of the peak position, suggesting that the difference in distortion and crystal structure does not fundamentally change the behavior of the emission with temperature.

In addition to the shift of the peak position, a narrowing of the PL is observed upon cooling. The main PL peak is fitted with a Lorentzian function, and the full width at half maximum (FWHM) versus temperature is plotted in Figure 2c. Analysis of the PL linewidth can reveal different scattering mechanisms. The total PL linewidth is typically described by the summation of an inhomogeneous broadening term $\Gamma_0$, the scattering by longitudinal optical (LO) and acoustic phonons, and the scattering from ionized impurities:

\[
\Gamma(T) = \Gamma_0 + \frac{\gamma_{LO}}{e^{\frac{E_{LO}}{kT}} - 1} + \frac{\gamma_{imp}}{e^{\frac{E_{imp}}{kT}} - 1} + \frac{\gamma_{LO}}{e^{\frac{E_{LO}}{kT}} - 1} + \frac{\gamma_{imp}}{e^{\frac{E_{imp}}{kT}} - 1}
\]

where $\gamma_{LO}$, $\gamma_{imp}$, and $\gamma_{LO}$ are the effective coupling strengths of scattering by acoustic phonons, optical phonons, and ionized impurities, respectively. $E_{LO}$ is the effective optical phonon energy, and $E_{imp}$ is the effective binding energy of ionized impurities. Each of the terms in Equation (1) gives a distinct trend of the linewidth. Figure 2c shows that an excellent fit to the temperature-dependent broadening of the linewidth can be achieved by exclusively considering contributions from LO phonons. Thus, there is negligible contribution of acoustic phonons and ionized impurities to the FWHM, which is consistent with other metal halide perovskites. Therefore, we attribute the dominant broadening to scattering by LO phonons, and their values are displayed in Figure 2c.

Both materials show virtually identical values for the effective phonon energy (17 meV) and phonon coupling strength...
To verify this correlation in our materials, we compared the Raman modes of PbI2 with those of (3-FPEA)2PbI4 and (PEA)2PbI4 (Figure S7, Supporting Information). We observed Raman modes up to 20.6 meV (166 cm⁻¹) in PbI2. This suggests that the low-energy modes up to 17.4 meV are dominated by the lead iodide cage, whilst higher energy modes include Raman contributions of the organic cations. The Raman modes that are observed at room temperature play an important role also at low temperature, as shown in Figure S8, Supporting Information.

The low-temperature PL spectra of (PEA)2PbI4 also shows high-energy peaks (Figure S12, Supporting Information), peaks HXe2 and HXe1 that are in accordance with previous reports[14,15] 3-FPEA opposes the view that the energy spacing of 12 to 15 meV is an overarching observation in these materials.[15,36] Strikingly, the interpeak separation is 40 to 46 meV for (3-FPEA)2PbI4, therefore much larger than the 12 to 15 meV separation for (3-FPEA)2PbI4. The smaller energy spacing of 12 to 15 meV in (3-FPEA)2PbI4 opposes the view that the energy spacing of 35 ± 5 meV is an overarching observation in these materials.[15,36] Similar to a previous report, we observe that the heavier mass of the halogen reduces the spacing of the hot-exciton emissions.[13] Assuming the hot-exciton peaks originate from a vibronic progression, it is remarkable that there is practically no change of modes observable in the Raman spectra (Figures S6 and S8, Supporting Information). Since Raman spectroscopy only probes the electronic ground state (GS), one explanation would be strikingly different modes in the electronic excited state governed by the organic cation. This observation warrants further studies to explain the impact of fluorination on the vibrational structure of the electronic excited state.

**Table 1. Summary of the energetic positions of the PL peaks.**

<table>
<thead>
<tr>
<th></th>
<th>HX2</th>
<th>HX1</th>
<th>X</th>
<th>LX1</th>
<th>LX2</th>
<th>LX3</th>
<th>LX4</th>
</tr>
</thead>
<tbody>
<tr>
<td>3-FPEA</td>
<td>2.411 eV</td>
<td>2.399 eV</td>
<td>2.384 eV</td>
<td>2.376 eV</td>
<td>2.371 eV</td>
<td>2.344 eV</td>
<td>2.340 eV</td>
</tr>
<tr>
<td>PEA</td>
<td>2.435 eV</td>
<td>2.389 eV</td>
<td>2.349 eV</td>
<td>2.347 eV</td>
<td>2.309 eV</td>
<td>2.305 eV</td>
<td>–</td>
</tr>
</tbody>
</table>

The time-resolved PL of the peaks HXe2, HXe1, and Xe are displayed in Figure 3b. Here, the peaks HXe2 and HXe1 show a fast decay with a lifetime of ≈2 to 3 ps. Peak Xe shows a much slower decay of 43 ps. In addition, peak Xe shows a delayed onset of approximately 10 ps, indicating that relaxation occurs from higher energy states toward the bottom of the electronic excited state (ES). The fast decay of peaks HXe2 and HXe1 together with their energy spacing coinciding with perovskite phonon modes leads us to ascribe these emissions to the radiative recombination of hot excitons as part of a vibronic progression; this is schematically depicted in Figure 3c.
Focusing on the low-energy side of the free exciton, Figure 4 shows a similar substructure in both compounds: $L_X F_3$ and $L_X F_4$ for $(3\text{-FPEA})_2\text{PbI}_4$; $L_X P_2$ and $L_X P_3$ for $(\text{PEA})_2\text{PbI}_4$. Analogously to the room-temperature PL, the PL at 5.4 K of $(3\text{-FPEA})_2\text{PbI}_4$ is shifted to higher energy compared to the PL of $(\text{PEA})_2\text{PbI}_4$ due to the increased distortion of the Pb-I-Pb angle. The substructures are only observed using a high-resolution grating. In addition, these high-resolution spectra allow to observe important differences near the main peak, that is, two individual peaks $L_X F_1$ and $L_X F_2$ for $(3\text{-FPEA})_2\text{PbI}_4$ and $L_X P_1$ for $(\text{PEA})_2\text{PbI}_4$. Mostly, these differences appear from a larger broadening of the PL of $(\text{PEA})_2\text{PbI}_4$, which could be determined by a form of long-range disorder characterizing this system. The small rotations of the inorganic layers around the stacking direction observed for this system (see Figure S1, Supporting Information), could be the source of such a disorder. The shoulder $L_X P_1$ could indicate a fine-structure splitting of the free exciton.[21,37] The plateau between $L_X F_2$ and $L_X F_3$ in $(3\text{-FPEA})_2\text{PbI}_4$ and between $L_X P_1$ and $L_X P_2$ in $(\text{PEA})_2\text{PbI}_4$ indicates emission from a large number of unresolved peaks.

Variation of the excitation fluence can help in discriminating recombination processes from each other. The PL intensity of these processes can be described by $I \propto P^k$ with $P$ the excitation fluence.[38,39] Monomolecular recombination processes, for example, exciton recombination, typically show an exponent $k$ of one. Exponents equal to or lower than one can be linked to free-to-bound and donor–acceptor pair transitions, whereas both a linear and superlinear dependence are associated with bound excitons.[38] A superlinear dependence has also been attributed to the emission of biexcitons in layered perovskites.[21,28,40]

The PL intensities versus increasing excitation fluence are presented in Figure S9, Supporting Information for $(3\text{-FPEA})_2\text{PbI}_4$ and Figure S10, Supporting Information for $(\text{PEA})_2\text{PbI}_4$. The main peaks X rise linearly with increasing excitation fluence, corroborating our assignment of free exciton emission. Interestingly, peaks $L_X P_3$ and $L_X P_4$ in $(3\text{-FPEA})_2\text{PbI}_4$ show a linear dependence opposed to the superlinear dependence of peaks $L_X F_2$ and $L_X F_3$ in $(\text{PEA})_2\text{PbI}_4$. This suggests either the absence of biexcitons in $(3\text{-FPEA})_2\text{PbI}_4$ or the misattribution of biexcitons in $(\text{PEA})_2\text{PbI}_4$, since a superlinear dependence points both to the emission of bound excitons and biexcitons.[21,38]

We extracted the time-resolved traces of the free exciton and the low-energy peaks for both materials, as shown in Figure 4b,c. The traces of the low-energy peaks are combined as these could not be displayed separately; see Figure S13, Supporting Information for the streak camera images. The PL for both materials shows a biexponential decay, where the free exciton decays faster than the low-energy peaks, as summarized in Table 2. Additionally, the low-energy traces exhibit a delayed onset up to tens of picoseconds, suggesting a transfer process from the free exciton. The similarity of PL decays of $L_X F_1 + L_X F_2$ and $L_X F_3 + L_X F_4$ suggests a similar origin of these emissions. We note that a long PL tail of $L_X F_1 + L_X F_2$ continues well beyond the displayed time window and persists for hundreds of ns into the μs range (Figure S14, Supporting Information). It has been argued that the long decay stems from weakly allowed emission from a dark state due to spin-orbit coupling and intermixing of a triplet state with a singlet state.[21,44,46] This leads us to assume the presence of a dark state slightly below the main peak of $(3\text{-FPEA})_2\text{PbI}_4$ as observed in $(\text{PEA})_2\text{PbI}_4$; see for additional discussion Note S1, Supporting Information.[21]

As mentioned earlier, the assignment of the low-energy PL peaks in layered perovskites based on PEA has been a source...
of contradicting reports.\textsuperscript{[13–18,43]} Explanations include excitons bound to defects,\textsuperscript{[16,17]} vibronic progressions,\textsuperscript{[13,14,43]} and exciton polarons.\textsuperscript{[15,18]}

Our data, namely, the intricate substructure, the PL intensity dependence on the excitation fluence, and the slower PL decay of the low-energy peaks suggests the presence of bound excitons. This explanation is further corroborated by the marked change in PL intensity upon temperature variation, an observation only expected for bound excitons (Figure S15, Supporting Information).\textsuperscript{[17]} It is clear that, as discussed above, the crystal structure influences the emission energy of the of the bound excitons. The larger emission linewidth of $X_F$ compared to $X_P$ could indicate an increased disorder in (PEA)$_2$PbI$_4$; an example could be the small rotation of the inorganic planes around the stacking direction.

Furthermore, for the emission $L_X3$ and $L_X4$, we exclude that phonon replicas could be their origin as the required modes in the Raman spectra are absent. In addition, we have no clear evidence of the energetic separation of 35 meV to confirm the proposed model of exciton polarons being important for this class of materials.

### 3. Conclusion

To conclude, in this study we address a current debate in the literature and contribute to the understanding of the emission properties of layered perovskites. This is achieved by investigating the structural and optical properties of (3-FPEA)$_2$PbI$_4$ and (PEA)$_2$PbI$_4$ single crystals. Fluorination of the organic cation increases the distortion of the octahedral layer, resulting in a shift to higher energy of the absorbance and PL spectrum. Interestingly, the PL spectra exhibit an intricate substructure at 5.4 K, where the interpeak separation of two hot-exciton peaks is reduced from 40 to 46 meV for (PEA)$_2$PbI$_4$ to 12 to 15 meV for (3-FPEA)$_2$PbI$_4$. In addition, high-resolution PL spectra show a previously unreported substructure in the low-energy peaks. Analysis of these spectra reveals that the emission at lower energies originates from bound excitons, in contrast to the commonly reported phonon replicas.

### Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

### Acknowledgements

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### Conflict of Interest

The authors declare no conflict of interest.

### Data Availability Statement

Research data are not shared.

### Keywords

crystal substructure, fluorinated cations, layered perovskites, luminescence spectroscopy, photophysics, Ruddlesden–Popper phase, single crystals

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