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Lattice interconversion of 1D ferrocene-based perovskite induced by solvent molecules for selective photocatalytic toluene oxidation†

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Metal halide perovskites (MHPs) have attracted enormous attention for potential applications in the fields of optoelectronics and photocatalysis due to their excellent optoelectronic characteristics, such as broad light absorption range, high charge mobility and long carrier diffusion length. However, using MHPs as photocatalysts for photocatalytic toluene oxidation is rare, and the underlying mechanisms affecting toluene oxidation are still unclear. In this study, we constructed two novel one dimensional (1D) ferrocene-based perovskite catalysts, $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$, of which $(C_{13}H_{17}FeNH)PbI_3$ can photocatalyze the oxidation of toluene to benzaldehyde effectively. It is interesting that the crystal lattices of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ can be converted to each other through the gain and loss of solvent molecules, which not only regulates the electronic band structure but also increases the separation efficiency of the photogenerated carriers. These results were confirmed by steady-state photoluminescence (PL) and time-resolved photoluminescence (TRPL) spectra, transient photocurrent response measurements and density functional theory (DFT) calculations. In addition, the lower exciton binding energy (43.2 meV) of $(C_{13}H_{17}FeNH)PbI_3$ further demonstrated its effective carrier separation efficiency. Furthermore, the selective adsorption of $(C_{13}H_{17}FeNH)PbI_3$ on toluene and benzaldehyde provided a prerequisite for the efficient selective oxidation of toluene. Finally, $(C_{13}H_{17}FeNH)PbI_3$ exhibited excellent catalytic activity for the photocatalytic oxidation of toluene to benzaldehyde with a conversion of 28.5% and selectivity (95.3%) towards benzaldehyde.

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Introduction

Selective oxidation of toluene to benzaldehyde for the production of highly value-added chemicals is important in the chemical industry.^{1–4} Benzaldehyde, as an aromatic carbonyl product, is a versatile stepping stone in the manufacturing of perfumes, solvents, dyes, plasticizers, flame retardants, and pharmaceuticals.^{5–9} However, to activate the thermodynamically strong and kinetically inert C(sp³)–H bonds, thermochemical processes are most frequently employed with the use of expensive metal catalysts or hazardous oxidants under

harsh reaction conditions (high temperature and/or high pressure).^{10,11} Such processes significantly suffer from the difficulty of controlling product selectivity and the simultaneous emission of large amount of waste, making their operation costly, energy-intensive, and environmentally unfriendly. Recently, the photocatalytic oxidation of hydrocarbons using clean and inexhaustible solar energy has garnered wide attention.^{12–16} It is well known that the active radicals (*e.g.*, $\cdot O_2^-$ and $\cdot OH$) generated by photocatalysis are extremely reactive with a high redox ability, which enables some challenging reactions to take place under green and mild conditions.¹⁷ Thus, several photocatalytic systems for the activation of C(sp³)–H bonds under solar or visible light illumination have been reported.

Over the past decade, metal halide perovskites (MHPs) have caused interdisciplinary interest and revolutionized optoelectronic fields due to their outstanding properties, such as excellent light absorption capability, large charge mobility, long carrier diffusion length, and low-cost solution processing.^{18–23} Inspired by these merits, in recent years, the utilization of MHPs for solar-to-chemical energy conversion has also received widespread attention, especially in the field

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of photocatalysis.^{24–27} More importantly, MHPs exhibit excellent catalytic performance as photocatalysts since they have high extinction coefficients, narrow band gaps with highly negative conduction band minimums (CBMs), low exciton binding energies (E_b), and fast charge transport.^{28,29} In recent years, some MHPs have become emerging catalysts for the photocatalytic oxidation of toluene. For instance, in 2020, Xu *et al.* employed $A_3Sb_2Br_9$ perovskite nanoparticles (NPs) as a new photocatalyst in the visible-light driven photocatalytic oxidation of toluene, indicating that the photocatalytic activity is strongly related to the distortion of the $[SbBr_6]$ octahedron affected by A-site cations.³⁰ In 2023, Chen *et al.* reported a layered all-inorganic lead-free MHP, $Cs_4ZnSb_2Cl_{12}$, which exhibits excellent photocatalytic performance in the transformation of toluene into benzaldehyde with a conversion rate of $1893 \mu\text{mol g}^{-1} \text{h}^{-1}$ and 95% selectivity.³¹ Although some progress has been made in the photocatalytic oxidation of toluene by MHPs, the underlying mechanisms affecting toluene oxidation are still unclear. In addition, lattice changes in MHPs may affect their semiconductor properties, such as photogenerated carrier dynamics, thereby affecting their photocatalytic activity. However, due to the lack of suitable systems, exploring the effect of lattice changes of MHPs on photocatalytic toluene oxidation has not been reported and remains a huge challenge.

In this work, we designed and synthesized two novel one dimensional (1D) ferrocene-based perovskites, $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$, of which $(C_{13}H_{17}FeNH)PbI_3$ can selectively and effectively photocatalyze the oxidation of toluene to benzaldehyde. Of note, the crystal lattices of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ can be converted to each other through the gain and loss of DEF molecules, which not only regulates the electronic band structure, but also increases the separation efficiency of photogenerated carriers. These results were confirmed by steady-state photoluminescence (PL) and time-resolved photoluminescence (TRPL) spectra, transient photocurrent response measurements and density functional theory (DFT) calculations. In addition, the E_b of $(C_{13}H_{17}FeNH)PbI_3$ was calculated to be 43.2 meV, much lower than that of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ (76.8 meV), further suggesting the effective carrier separation efficiency of $(C_{13}H_{17}FeNH)PbI_3$. Furthermore, the selective adsorption of $(C_{13}H_{17}FeNH)PbI_3$ on toluene and benzaldehyde provides a prerequisite for the efficient selective oxidation of toluene. Finally, $(C_{13}H_{17}FeNH)PbI_3$ exhibited excellent catalytic activity for the photocatalytic oxidation of toluene to benzaldehyde with a conversion of 28.5% and selectivity (95.3%) towards benzaldehyde.

Results and discussion

Crystal structural analysis of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$

Although $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ exhibit similar topological structures, their synthetic methods

are vastly different. The *N,N*-dimethylaminomethylferrocene-based 1D perovskite single crystal with DEF solvents, $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$, was grown by a conventional solvent evaporation method, while $(C_{13}H_{17}FeNH)PbI_3$ was obtained using an anti-solvent process. Single-crystal X-ray diffraction analysis indicated that $(C_{13}H_{17}FeNH)PbI_3$ crystallizes in the monoclinic $P112_1$ space group, featuring a 1D chain-based three-dimensional (3D) supramolecular structure. As shown in Fig. S1,† the basic structural motif of $(C_{13}H_{17}FeNH)PbI_3$ is composed of one $[PbI_3]^-$ and one protonated *N,N*-dimethylaminomethylferrocene ($C_{13}H_{17}FeNH$). Adjacent $[PbI_3]^-$ groups are connected together through coordination bonds between I^- ions and Pb^{2+} ions, forming a 1D anionic chain (Fig. 1a). $C_{13}H_{17}FeNH$ is arranged neatly around this chain as a counter cation to balance the charge. Finally, one 1D chain-based 3D supramolecular structure, $(C_{13}H_{17}FeNH)PbI_3$, is obtained through supramolecular interactions between adjacent 1D chains (Fig. 1b). Of note, as shown in Fig. 1b, the distance of adjacent 1D anionic chains at the (002) crystal planes is 13.2 Å. Interestingly, the C–N bond of $C_{13}H_{17}FeNH$ can rotate around the C atom as the center (Fig. 1c), which provides a prerequisite for the lattice change of $(C_{13}H_{17}FeNH)PbI_3$.

As shown in Fig. 1d, DEF is embedded into the lattice of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ in the form of molecules through hydrogen bonding interactions (Fig. S2†), which affects the rearrangement of $C_{13}H_{17}FeNH$ in the lattice, resulting in the number of layers of $C_{13}H_{17}FeNH$ in the lattice of these two perovskites changing from a single layer of $(C_{13}H_{17}FeNH)PbI_3$ to double layers of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$. This leads to a longer distance between the adjacent 1D anionic chains at the (002) crystal planes (17.3 Å) of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ compared to $(C_{13}H_{17}FeNH)PbI_3$. This lattice change induced by solvent molecules may cause differences in the semiconductor properties of these two perovskites, thereby affecting the photocatalytic activity. Pertinent crystal data and relevant refinement parameters for $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ are listed in Table S1.†

Lattice interconversion of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$

Firstly, powder X-ray diffraction (PXRD) was performed to confirm the successful synthesis of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$. As shown in Fig. 2a, the PXRD patterns of the synthesized $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ match well with their simulated ones from the simulated single crystal data, illustrating their good phase purities. It should be noted that the diffraction peak position of the (002) crystal plane of $(C_{13}H_{17}FeNH)PbI_3$ (7.2°) is obviously greater than that of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ (5.3°), which further indicates that the lattice of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ is larger than that of $(C_{13}H_{17}FeNH)PbI_3$, consistent with their crystallographic data. The (002) crystal planes of the two perovskites represented in the crystal structure can be found in Fig. S3.† Due to the weak interaction between the DEF molecule and $C_{13}H_{17}FeNH$ in the lattice of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$, we found that $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ can be con-

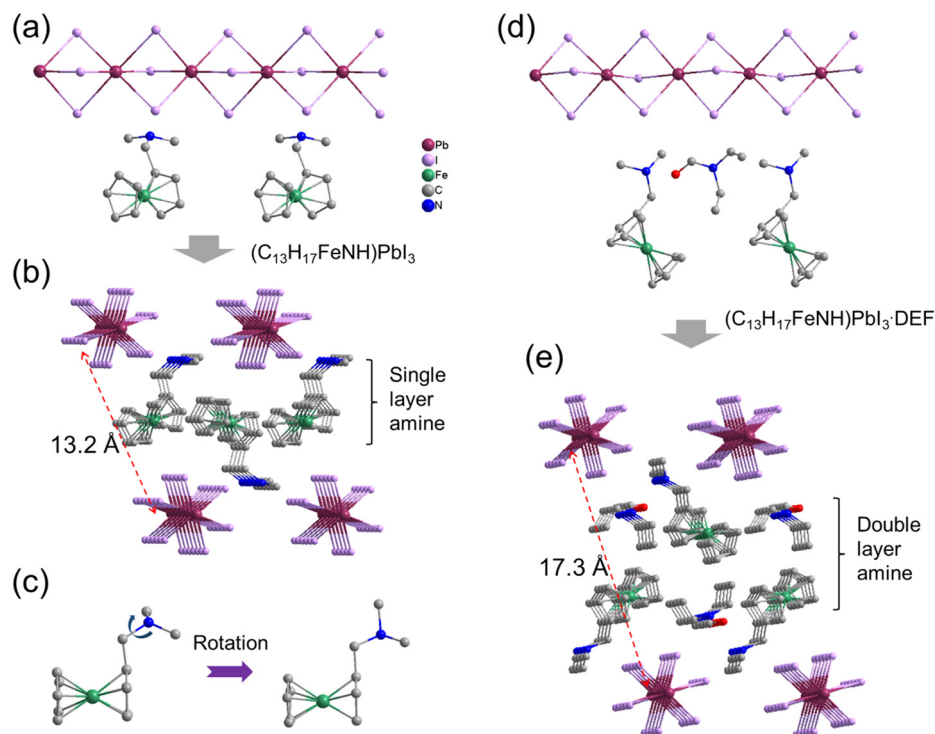


Fig. 1 1D chain structure of $(C_{13}H_{17}FeNH)PbI_3$ (a); 1D chain-based 3D supramolecular structure of $(C_{13}H_{17}FeNH)PbI_3$ (b); rotation diagram of the C-N bond of $C_{13}H_{17}FeNH$ (c); 1D chain structure of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ (d); 3D supramolecular structure of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ (e).

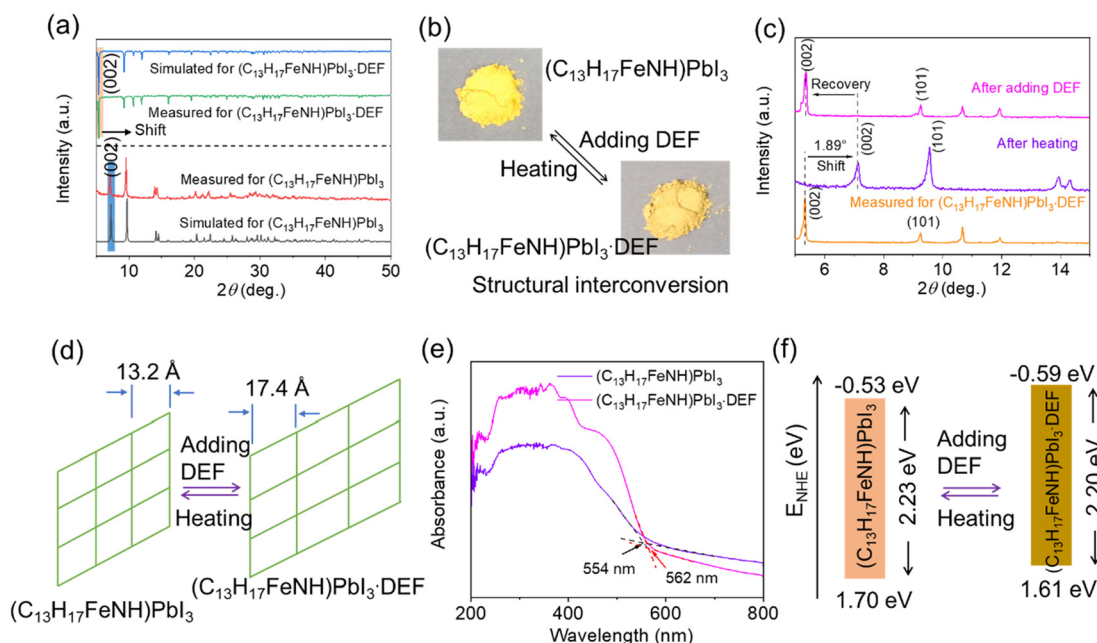


Fig. 2 As-synthesized and simulated PXRD patterns of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ single crystal powders (a); photos of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ powders and schematic of their structural interconversion (b); PXRD pattern changes of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ after heating and adding DEF (c); schematic of lattice transformation between $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ (d); UV-vis diffuse reflectance spectra of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ samples (e); electronic band structure diagram of $(C_{13}H_{17}FeNH)PbI_3$ and $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ (f).

verted to each other through the gain and loss of solvent molecules. As illustrated in Fig. 2b, $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ can be transformed into $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ by adding DEF dropwise, while, by heating, $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ can be converted into $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ gradually. This structural interconversion was also revealed by changes in the PXRD patterns. As shown in Fig. 2c, when $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ was heated at 80 °C for 20 min, the diffraction peak position of the (002) crystal plane after heating shifted towards a larger angle by 1.89°. It's interesting that the diffraction peak position of the (002) crystal plane after adding DEF returned to the original position of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$. This also demonstrates that the lattice interconversion between $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ can be achieved by controlling the gain and loss of solvent molecules. As shown in Fig. 2d, this lattice interconversion may lead to differences in the photoelectric properties of these two perovskites, further affecting their photocatalytic activities. The effect of DEF molecules on the thermal stabilities of these two perovskites was investigated by thermogravimetric analysis (TGA). As show in Fig. S4,† for $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$, the weight loss of about 10.4% at 25–85 °C corresponds to DEF molecules. Then, its framework starts to decompose at approximately 180 °C. $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$, due to the lack of DEF molecules in the crystal lattice, does not possess a weight loss process at 25–85 °C and maintains excellent thermal stability until 180 °C.

The electronic band structures of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ were then investigated.

The UV-vis diffuse reflectance spectra (DRS) of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ powder samples were obtained first to evaluate their bandgaps (E_g) (Fig. 2e). E_g can be defined as the intersection point between the energy axis and the line extrapolated from the linear portion of the adsorption edge in the plot of the Kubelka-Munk function F against E . $F = (1 - R)^2/2R$ was transformed from the measured diffuse reflectance data, in which R presents the reflectance of an infinitely thick layer at a given wavelength.^{32,33} As shown in Fig. S5a and b,† the E_g of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ were calculated to be 2.23 eV and 2.20 eV, respectively. Then, we calculated the conduction band minima (CBM) of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ through Mott-Schottky plots. As illustrated in Fig. S6,† the flat band potential (E_{fb}) of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ was calculated to be -0.67 eV vs. SCE, corresponding to -0.43 eV vs. NHE. Considering the CBM could be 0–0.1 eV more negative than the E_{fb} , the CBM of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ equals -0.53 eV vs. NHE.³⁴ As shown in Fig. S7,† the CBM of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ was calculated to be -0.59 eV vs. NHE. The position of the valence band maxima (VBM) of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ were calculated to be 1.70 eV and 1.61 eV, respectively, based on the E_g and CBM. Thus, the electronic band structure diagrams of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ are shown in Fig. 2f. Due to the lattice change induced by DEF molecules, the electronic band structures between $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ can be effectively regulated.

Low exciton binding energy and excellent photoelectrochemical properties

For photocatalysts, the separation efficiency of photogenerated carriers is a key factor affecting the photocatalytic performance. The PL and TRPL spectra were thus investigated. As shown in Fig. 3a, the relatively lower PL emission for $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ compared to that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ is due to the recombination of the photogenerated electron-hole being effectively suppressed, indicating low exciton binding energy and favorable charge transport. As shown in Fig. 3b, the TRPL curves of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ can be well fitted by a double-exponential function³⁵

$$I(t) = A_1 \exp\left(-\frac{t}{\tau_1}\right) + A_2 \exp\left(-\frac{t}{\tau_2}\right) \quad (1)$$

where τ_1 and τ_2 are the decay times of the first and second decay components, respectively, and A_1 and A_2 are the amplitudes of the first and second decay components, respectively. The amplitudes of the decay components reflect the total contribution of each lifetime component toward the average lifetime (τ_{ave}), which can be calculated as follows.^{36,37}

$$\tau_{\text{ave}} = \frac{A_1 \tau_1^2 + A_2 \tau_2^2}{A_1 \tau_1 + A_2 \tau_2} \quad (2)$$

All fitting parameters are tabulated in Table S2,† and τ_{ave} (0.352 ns) of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is shorter than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (0.541 ns), which further illustrates the higher separation efficiency and transfer ability of photogenerated charges of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$. Temperature-dependent photoluminescence was employed to explore the exciton binding energies of both perovskites. Fig. 3c and Fig. S8† show the temperature-dependent PL intensity plots of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$, respectively. The PL intensity decreases with the increase of temperature. The exciton binding energy is calculated by fitting the relation curve between PL integral intensity and temperature through the following equation^{38,39}

$$\frac{1}{I(T)} = A + B \exp\left(-\frac{E_b}{k_b T}\right) \quad (3)$$

where I is PL intensity, A and B are the fitting parameters, E_b represents exciton binding energy, k_b is the Boltzmann constant, and T is temperature. From Fig. 3d, the E_b of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ was calculated to be 43.2 meV, which is much lower than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (76.8 meV), further indicating the greater separation efficiency of photogenerated carriers of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$. Transient photocurrent response measurements can directly reflect the separation efficiency of photogenerated carriers. As shown in Fig. 3e, the photocurrent density of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ can reach as high as 9.1 $\mu\text{A cm}^{-2}$, about 3.3 times more than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (2.7 $\mu\text{A cm}^{-2}$). The higher photocurrent density of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ illustrates its higher separation efficiency of photogenerated carriers.

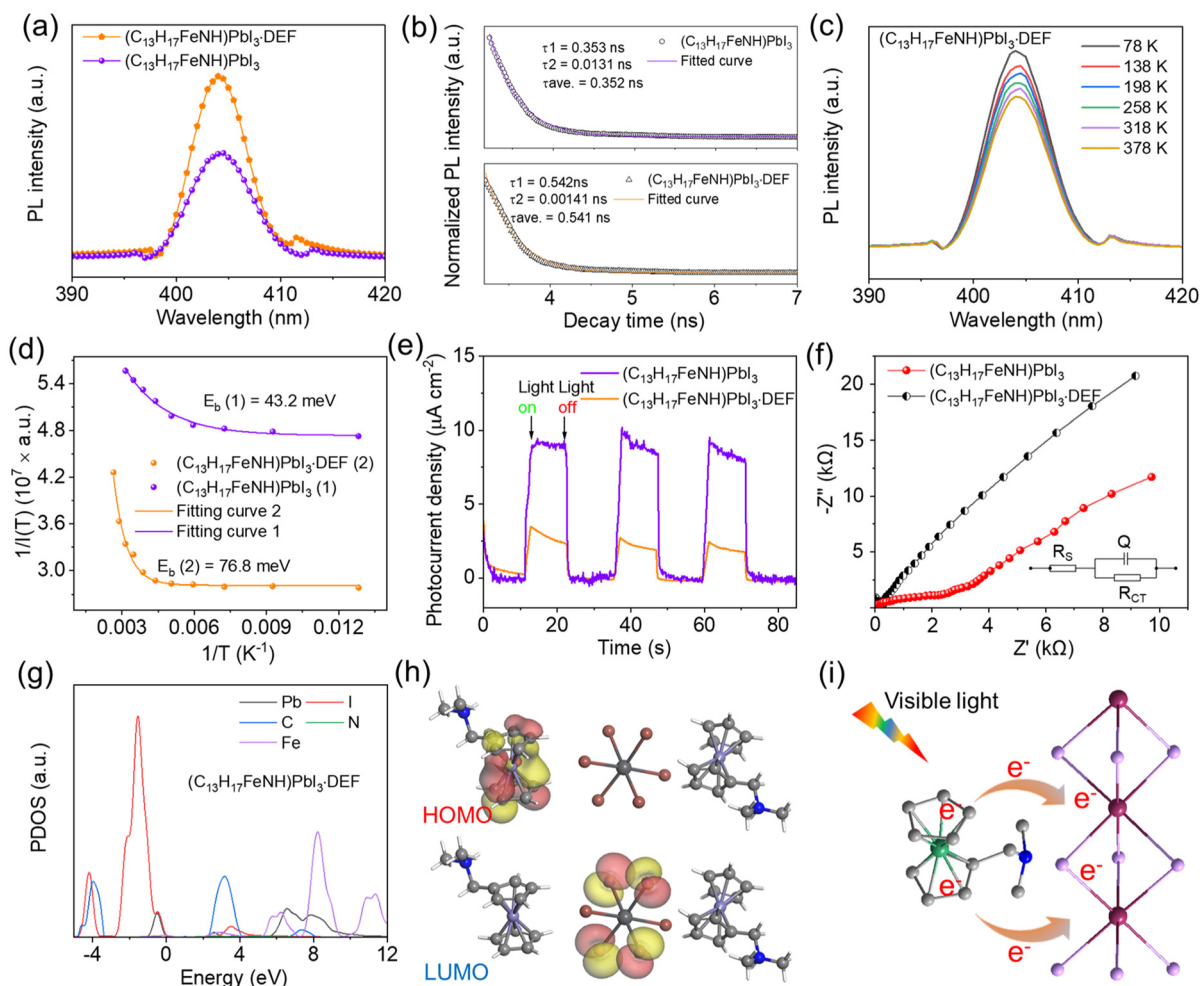


Fig. 3 PL (a) and TRPL (b) spectra of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$, respectively; temperature-dependent PL intensity plots of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (c); exciton binding energies from fitting the integrated PL intensity to temperature (d); transient photocurrent responses of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$ under 0.5 V bias voltage (e); electrochemical impedance spectroscopy of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$ (f); PDOS plots of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (g); HOMO and LUMO of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (h); schematic of charge transition process of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (i).

The electrochemical impedance spectroscopy (EIS) Nyquist plots of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$ were measured to illustrate their capacities of charge separation and transfer (Fig. 3f). The equivalent circuit diagram is inserted. From fitting the EIS data, the charge transfer resistance values (R_{CT}) of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$ electrodes were found to be $4.58 \times 10^4 \Omega$ and $7.82 \times 10^4 \Omega$, respectively. In general, the smaller the R_{CT} , the smaller the radius of the impedance spectrum, which means a higher efficiency of charge separation and transfer. Thus, the charge transfer capacity of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is larger than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$. To further reveal the contributions of different atoms of these two perovskites to the LUMO and HOMO orbitals, as well as the differences of the charge transfer process of these two perovskites, DFT calculations were carried out. As shown in Fig. 3g, the electron density of the LUMO orbital of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ mainly lies on the I 5p and Pb 6s orbitals of the $[\text{PbI}_3]^-$ group. The HOMO orbital of

$(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is mostly concentrated on the C 2p and Fe 3d orbitals of the $\text{C}_{13}\text{H}_{17}\text{FeNH}$ group. These results show that the electrons of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ primarily transition from the C 2p and Fe 3d orbitals of $\text{C}_{13}\text{H}_{17}\text{FeNH}$ to the I 5p and Pb 6s orbitals of the $[\text{PbI}_3]^-$ group (Fig. 3h and i), indicating the efficient separation of photogenerated carriers. However, in contrast to $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$, the HOMO and LLUMO orbitals of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\text{-DEF}$ are both contributed by the $\text{C}_{13}\text{H}_{17}\text{FeNH}$ groups (Fig. S9[†]), illustrating that the presence of DEF molecules in the crystal lattice is not conducive to the separation of photogenerated carriers, which is consistent with the PL, TRPL, and transient photocurrent response measurements.

Photocatalytic oxidation of toluene

To explore the influence of lattice changes induced by DEF on photocatalytic activities, the photocatalytic $\text{C}(\text{sp}^3)\text{-H}$ bond activation in toluene molecules was investigated under visible

light irradiation ($\lambda > 420$ nm) using these two perovskites as catalysts at room temperature (Fig. 4a). As shown in Fig. 4b, both $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ can photocatalyze the oxidation of toluene, but the yield of benzaldehyde over $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is higher than the yield using $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ as catalyst within all reaction times. Using $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ as photocatalyst, the amount of benzaldehyde after 4 h of reaction can reach as high as 251 μmol , which is about 1.8 times higher than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (140 μmol). As shown in Fig. 4c, the selectivity of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ towards BA is as high as 95.3%, which is larger than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (73%). In addition, both $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ have very low selectivity for BY production, with 3.2% and 12%, respectively. Of note, the conversion rate of toluene using $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ as catalyst can reach 28.5%, which is also larger than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (26%). These results indicate that the DEF molecules in the crystal lattice of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ are not conducive to its photocatalytic activity for toluene oxidation. We further compared the catalytic performance of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ with those of some other reported catalysts, as shown in Table S3.[†] The results indicate that $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ possesses excellent catalytic performance for photocatalytic toluene oxidation. As shown in Fig. 4d, the apparent quantum efficiency (AQE) value for the oxidation of toluene to benzaldehyde on $(\text{C}_{13}\text{H}_{17}\text{FeNH})$

$\text{PbI}_3\cdot\text{DEF}$ is 2.5%. $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ presents the significantly increased AQE of 4.3%, further demonstrating the excellent photocatalytic performances of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$. Reusability and stability are important figures of merit to evaluate a catalyst. As shown in Fig. 4e, only a slight decrease in the yield of benzaldehyde (from 251 μmol to 223 μmol) was found after five cycling experiments, indicating that the structural framework of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is basically stable after five cycles.

For the purpose of further exploring and verifying the catalytic potential of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$, the reactivities of various toluene derivatives were examined under the same conditions (Table 1). When *p*-xylene as a substrate was added into this reaction system, a substrate conversion rate of 37.3% was achieved with a corresponding aldehyde selectivity of 88.4%. In addition, *o*-xylene and *m*-xylene as substrates also present higher conversion than toluene, and the corresponding aldehyde selectivities are around 80%. Several electron-withdrawing *para*-position-substituted toluenes, including -F, -Cl and -Br, were subjected to the optimal conditions, and all can also convert into their corresponding aldehydes. However, compared to toluene, electron-withdrawing substituted toluenes show slightly lower conversion, and electron-donating substituted toluenes show relatively higher conversion. These results indicate that $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ can serve as an effective photocatalyst for photocatalytic oxidation of toluene derivatives.

Photocatalytic mechanism analysis

To clarify the reaction mechanism of photocatalytic toluene oxidation over $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$, a series of free-radical-capture experiments were performed to reveal the key reactive species induced by light (Fig. 5a). The introduction of tetramethylpiperidine *N*-oxide (TEMPO) as the quenching agent for all radicals in the reaction completely suppresses the conversion of toluene, indicating the key role of the radicals for toluene oxidation. The addition of *tert*-butanol (TBA) as an $\cdot\text{OH}$ scavenger has no noticeable effect on toluene conversion, indicating that $\cdot\text{OH}$ is not involved in the reaction. The addition of butylated hydroxytoluene (BHT) as the quenching

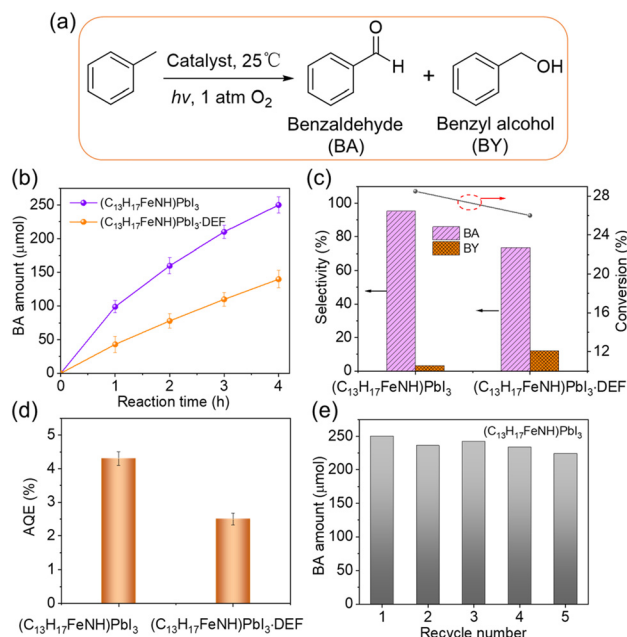


Fig. 4 Photocatalytic selective oxidation of toluene; conditions: photocatalyst (50 mg), toluene (1 mmol), MeCN (3 mL), O_2 (1 atm), $\lambda > 420$ nm, and irradiation time (4 h) (a); effect of reaction time on BA amount over $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ catalysts (b); selectivity of BA and BY as well as the conversion of toluene over $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ catalysts (c); AQE of toluene oxidation for $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ ($\lambda = 365$ nm) (d); photocatalytic cycling experiment for toluene oxidation (e).

Table 1 Substrate scope on $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ ^a

Entry	R	Conv. (%)	Sel. (%)	
			Aldehyde	Alcohol
1	H	28.5	95.3	3.2
2	<i>o</i> -CH ₃	33.7	81.5	8.7
3	<i>m</i> -CH ₃	32.1	83.2	9.5
4	<i>p</i> -CH ₃	37.3	88.4	5.2
5	<i>p</i> -F	17.7	82.1	9.4
6	<i>p</i> -Cl	21.2	89.6	6.6
7	<i>p</i> -Br	24.8	93.0	4.6

^a Reaction conditions: substrate (1 mmol), MeCN (3 mL), O_2 (1 atm), photocatalyst (50 mg), $\lambda > 420$ nm, and reaction time (4 h).

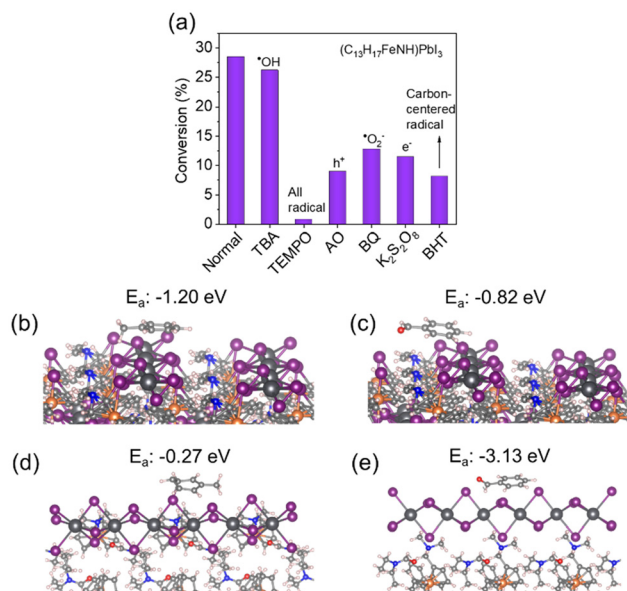


Fig. 5 Control experiments of radical quenching for the photocatalytic oxidation of toluene (a); adsorption models for toluene on $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (b) and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (d); adsorption models for benzaldehyde on $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (c) and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (e).

agent of carbon-centered radicals in the reaction produces an apparent decline in toluene conversion, indicating that the carbon-centered radicals are important intermediates for the oxidation of toluene to benzaldehyde. In Fig. 5a, it can be seen that the conversion of toluene was significantly blocked after ammonium oxalate (AO) was added as a hole (h^+) scavenger, suggesting that holes play a key role for toluene conversion. Furthermore, there was a significant decline in toluene conversion when either benzoquinone (BQ) or $\text{K}_2\text{S}_2\text{O}_8$ was added, which are scavengers for superoxide radicals ($\cdot\text{O}_2^-$) and photogenerated electrons (e^-), respectively. Since the formation of $\cdot\text{O}_2^-$ is through the combination of molecular oxygen with photogenerated electrons, the quenching of photogenerated electrons would result in the decline of $\cdot\text{O}_2^-$ formation. The results strongly suggest that the $\cdot\text{O}_2^-$ play a critical role in the photocatalytic toluene oxidation.

DFT calculations were further employed to investigate the difference in adsorption capacities of these two perovskites for toluene and benzaldehyde. As shown in Fig. 5b, the calculated adsorption energy (E_{ad}) of toluene molecules on $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ was -1.20 eV, which is substantially lower than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ (-0.27 eV), confirming that the adsorption of toluene on $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is energy favorable. This is a prerequisite for $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ to exhibit a higher conversion rate to toluene. In addition, considering that a large adsorption energy of the catalyst to the substrate is not conducive to the removal of the substrate from the catalyst, which will affect the selectivity of the catalyst towards the production of substrate, we further calculated the E_{ad} of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ for benzaldehyde. As shown in Fig. 5e, the E_{ad} of benzaldehyde on $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ is

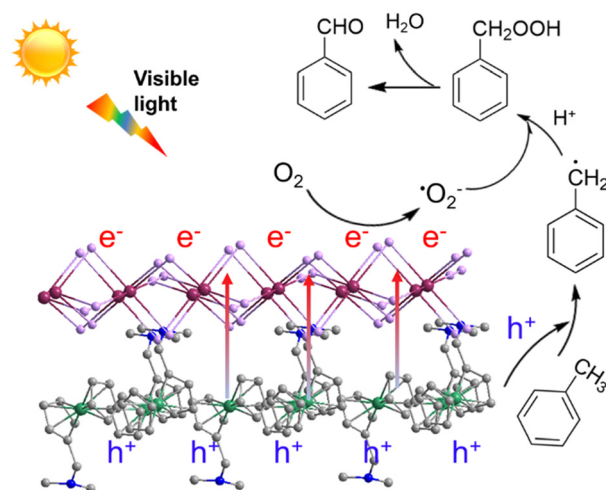


Fig. 6 Possible photocatalytic reaction mechanism of toluene oxidation over $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$.

as low as -3.13 eV, which is much lower than that of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ (-0.82 eV). This indicates that benzaldehyde is more easily removed from $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$, hindering its further oxidation and improving the selectivity of benzaldehyde.

Based on experimental and DFT calculation results, a plausible reaction mechanism for the selective photocatalytic oxidation of toluene to benzaldehyde over $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ can be proposed as follows (Fig. 6): (1) $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ is excited to generate photoinduced electrons and holes under visible light irradiation; (2) the accumulated holes at the VBM of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ oxidize toluene to yield benzyl radicals and the accumulated electrons at the CBM of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ react with O_2 to generate $\cdot\text{O}_2^-$; (3) benzyl radicals couple with $\cdot\text{O}_2^-$ and a proton to form the hydroperoxide intermediate; (4) the hydroperoxide species can be dehydrated to form the benzaldehyde species, which desorbs from $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ quickly due to its low adsorption energy to produce benzaldehyde, avoiding the overoxidation process.

Conclusions

In summary, two novel 1D ferrocene-based perovskites, $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$, were constructed for the selective photocatalytic oxidation of toluene to benzaldehyde. Impressively, we found that the crystal lattices of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ and $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3\cdot\text{DEF}$ can be converted to each other through the gain and loss of DEF molecules, which not only regulates the electronic band structure, but also increases the separation efficiency of photogenerated carriers. These results were confirmed by PL, TRPL, and transient photocurrent response measurements. In addition, the significantly lower exciton binding energy of $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ further demonstrates its favorable charge separation. Finally, $(\text{C}_{13}\text{H}_{17}\text{FeNH})\text{PbI}_3$ exhibited excellent catalytic activity for the

photocatalytic oxidation of toluene to benzaldehyde with a conversion of 28.5% and selectivity (95.3%) towards benzaldehyde. This work reveals the influence of perovskite lattice changes on their optoelectronic properties and photocatalytic performances, which may provide new ideas and insights for designing perovskite-based photocatalysts with highly efficient separation efficiencies of photogenerated carriers and selectivity towards substrates.

Experimental

Materials and characterization

All reagents utilized in this study were of analytical purity and used without additional processing. Additional information regarding the reagents and instrument characterizations can be found in the ESI.†

Synthesis of $C_{13}H_{17}FeNH$

6.8 mL of HI was dissolved in 15 mL ethyl alcohol, and then 10 mL of $C_{13}H_{17}FeN$ was slowly added dropwise to the above solution. After the mixed solution was stirred in an ice bath for 2 h, $C_{13}H_{17}FeNH$ powder was obtained using a rotary evaporator and purified by washing with dichloromethane. The $C_{13}H_{17}FeNH$ powder was dried in a vacuum oven overnight at 55 °C before use.

Synthesis of $(C_{13}H_{17}FeNH)PbI_3$

A mixture of $C_{13}H_{17}FeNH$ (0.01 mol, 3.71 g) and PbI_2 (0.01 mmol, 4.61 g) was dissolved in 3 mL of DEF in a 10 mL beaker at 50 °C. After stirring at 100 °C for 2 h, the solution was immediately filtered through a 0.45 μm poly(tetrafluoroethylene) filter to obtain a clear perovskite precursor solution. Then, the $(C_{13}H_{17}FeNH)PbI_3$ single crystal can be obtained by placing above solution in a closed container with a dichloromethane atmosphere for 7 days.

Synthesis of $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$

A mixture of $C_{13}H_{17}FeNH$ (0.01 mol, 3.71 g) and PbI_2 (0.01 mmol, 4.61 g) was dissolved in 3 mL of DEF in a 10 mL beaker at 50 °C. After stirring at 100 °C for 2 h, the solution was immediately filtered through a 0.45 μm poly(tetrafluoroethylene) filter to obtain a clear perovskite precursor solution. Then, the $(C_{13}H_{17}FeNH)PbI_3 \cdot DEF$ single crystal was obtained after the solution slowly evaporated at 55 °C for 3 days.

Photocatalytic oxidation of toluene experiments

The photocatalytic oxidation of toluene to benzaldehyde was performed in a 10 mL quartz reactor with water-cooling system under visible light irradiation ($\lambda > 420$ nm, 100 mW cm⁻²). A 300 W xenon lamp (Perfect Light Company, Beijing) was chosen as the light source. For each photocatalytic reaction, 1 mmol of toluene, 50 mg of photocatalyst and 3 mL of acetonitrile (CH_3CN) were uniformly mixed in the quartz reactor. O_2 was bubbled through the mixture at a rate of 1.0 mL min⁻¹ for 1 h. Then, the quartz reactor was sealed with a balloon to keep

the oxygen gas at one atmosphere pressure. The suspension was stirred in the dark for 30 min before illumination. Then, the reaction was carried out under visible light irradiation. After the reaction, the catalyst was removed by centrifugation and the products were directly analyzed with a gas chromatograph (Agilent GC 7820A). The conversion (con.) of toluene and selectivity (sel.) of the products were calculated according to the following equations.^{40,41}

$$\text{Con. (\%)} = \frac{\text{moles of toluene reacted}}{\text{moles of initial toluene}} \times 100\% \quad (4)$$

$$\text{Sel. (\%)} = \frac{\text{moles of corresponding product}}{\text{moles of toluene reacted}} \times 100\% \quad (5)$$

Data availability

The data supporting the findings of this study are available from the corresponding authors upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

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