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As a light absorber in photovoltaic applications, hybrid organic–inorganic halide perovskites should have longer and balanced diffusion lengths for both the separated electrons and holes before recombination, which necessitates high carrier mobility. In polar semiconductors, the room-temperature carrier mobility is often limited by the scattering between carriers and the lowest-frequency optical phonon modes. Using terahertz time-domain spectroscopy, we examine the temperature evolution of these phonon modes in CH$_3$NH$_3$PbBr$_3$ and obtained high carrier mobility values using Feynman’s polaron theory. This method allows us to estimate the upper limit of carrier mobilities without the need to create photogenerated free carriers, and can be applied to other heteropolar semiconductor systems with large polarons. Published by AIP Publishing.

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The lead halide perovskite family, CH$_3$NH$_3$PbI$_3$ (MAPbI$_3$), MAPbBr$_3$ and relatives, have attracted much interest as candidates for photo-absorbers in next-generation photovoltaic devices. The highest power conversion efficiency of laboratory-scale devices has reached 22.1%, as a result of broadband absorption in the solar spectrum, long-range balanced electron and hole diffusion lengths, and high carrier mobility.\textsuperscript{5–7}

The MAPbX$_3$ (X = I, Br) compounds consist of two sub-lattices comprising an inorganic PbX$_3$ octahedral framework and MA\textsuperscript{+} organic cations. First-principles calculations show that, in contrast to all-inorganic perovskites, hybrid organic–inorganic perovskites possess several unique properties, viz., that the interband transition near the band edge is dominated by the electronic states of the inorganic framework, while symmetry breaking associated with the organic cation produces disorder in the material.\textsuperscript{8–10}

Hybrid perovskites have been characterized by various experimental techniques including optical spectroscopy,\textsuperscript{6,11,12} X-ray diffraction,\textsuperscript{13} and neutron scattering.\textsuperscript{14} Many optical techniques, such as photoluminescence (PL) spectroscopy and transient-absorption spectroscopy, study processes such as carrier recombination, carrier relaxation, and electron-phonon coupling indirectly, based on interband transitions on electron-volt energy scales. In contrast, as a direct way to elucidate the evolution of free carriers, recent terahertz (THz) spectroscopic studies illustrate that both free carriers and phonon modes coexist in the THz frequency range.\textsuperscript{15,16} It is therefore important to study the lowest-lying phonon modes and their evolution with temperature, in order to characterise the transport and recombination processes in these materials.

In MAPbX$_3$, the orthorhombic-to-tetragonal phase transition is accompanied by the order-disorder behavior of the MA\textsuperscript{+} cation orientation.\textsuperscript{10,17} The THz response of MAPbI$_3$ has been studied both theoretically and experimentally,\textsuperscript{16–18} whereas the bromide analogue MAPbBr$_3$ has received relatively less attention. The latter has been widely employed in mixed-cation and mixed-halide perovskites,\textsuperscript{19} to improve the stability of all-perovskite tandem cells.

As a hybrid organic–inorganic crystal, the disorder of the organic MA\textsuperscript{+} cation plays a key role in the phase transitions. The orientation of the C–N axis is fixed in the orthorhombic phase, while the dipole has various possible orientations in the tetragonal and cubic phases. Thus, the transition from orthorhombic to tetragonal is not only related to the change in the crystal structure and unit-cell volume but is also linked to the order-disorder transition. The structural phase transitions in single-crystal MAPbBr$_3$ are reported to take place at\textsuperscript{10}
In this paper, we study the temperature evolution of the phonon modes in MAPbBr$_3$ thin films from 20 to 300 K in the region of 0.3–2.8 THz. We observe that two peaks in the tetragonal and cubic phases split into three peaks in the orthorhombic phase, with the additional mode at 1.8 THz gradually appearing near the phase-transition temperature. Comparing our results with first-principles density-functional theory (DFT) calculations, we attribute the phonon-mode splitting to the orthorhombic-to-tetragonal phase transition.

The perovskite precursor solution was prepared by dissolving MABr and PbBr$_3$ in a stoichiometric ratio in dimethylformamide/dimethyl sulfoxide (4:1 v/v, 1 M) at 80°C. The precursor solution was spin-coated onto a z-cut quartz substrate (10 × 10 × 1 mm) at 1000 rpm for 10 s and 4000 rpm for 30 s, with 250 μL of chlorobenzene being dripped onto the sample surface after 15 s during the second stage of the spin coating. Subsequently, the perovskite layer was annealed at 80°C for 5 min and 100°C for 10 min. The thickness of the polycrystalline MAPbBr$_3$ film was (932 ± 11) nm. The sample was kept under nitrogen before being mounted onto our THz-TDS system under vacuum (10$^{-7}$ mbar), which was maintained throughout the measurements to minimize exposure of the sample to air.

Our THz spectrometer setup and data analysis method have been reported elsewhere. The THz conductivity $\sigma(\omega)$ is obtained from the relation $\sigma(\omega) = i\omega\epsilon(\omega)$, where $\epsilon(\omega)$ is the complex dielectric constant.

Above 150 K, the spectra are well fit by two Lorentzian oscillators, which is near the reported single-crystal-to-single-crystal tetragonal-to-orthorhombic phase transition temperature. Below 150 K, a shoulder feature at 1.8 THz (mode 3) gradually becomes a dominant peak at the lowest temperatures, necessitating the addition of a third Lorentzian function. This feature clearly indicates that the structural phase transition induces a phonon-mode splitting. A similar shoulder also appears in the mid-infrared spectra in the orthorhombic phase and is attributed to the organic cations at ~28 THz. Although weak, three more modes at 0.7, 1.0, and 2.6 THz can only be identified below 150 K.
Although room-temperature THz data on MAPbBr₃ have been reported earlier, we show here temperature-dependent THz measurements, which can show the fingerprints of structural phase transitions. To explore the possibility that the appearance of the shoulder/peak feature below 150 K is due to the tetragonal-to-orthorhombic phase transition, we performed DFT-based lattice-dynamics simulations of the THz spectra following a similar procedure to that in our previous work. Data for cubic MAPbBr₃ was taken from our previous study. An initial model of the orthorhombic phase was built from the optimized structure of the iodide analogue, which was then fully optimized using the VASP code with care taken to converge the electronic wavefunctions and forces (10⁻⁸ eV and 10⁻⁵ eVÅ⁻¹, respectively). We used the PBEsol exchange-correlation functional with projector-augmented-wave (PAW) pseudopotentials with the outermost s, p, and Pb d electrons in the valence shell. A plane-wave cut-off of 800 eV was employed with a regular 3 × 2 × 3 Γ-centered Monkhorst-Pack k-point mesh. The precision of the charge-density grids was set automatically to ensure that it is mode 2 in the tetraII phase that splits into modes 2 and 3 in the orthorhombic phase.

From Fig. 2(a), we see that, in the orthorhombic phase, both the experimental and simulated spectra show two major peaks (labelled 1 and 2) and one shoulder (labelled 3). The shoulder results from the two weak IR-active phonon modes in the 1.5–2 THz range in the orthorhombic phase, which are absent in the cubic phase, in agreement with our measurements. The lattice dynamic calculations therefore support our observation that the shoulder feature is present only in the orthorhombic phase. We note that both sets of simulated spectra are blue-shifted to higher energies compared to the experimental measurements. Similar phenomena have been observed by Ahmed et al. in MAPbI₃. More recently, high-quality Raman spectra, obtained from single-crystal MAPbI₃ and MAPbBr₃ samples, also found that the calculated frequencies of the lowest-lying phonon modes were higher than those obtained from the measured Raman spectra. This may be due in part to the fact that the spectra are simulated from the athermal 0 K structures—at finite temperature, thermal expansion of the lattice can result in a softening of the phonon frequencies and an increase in anharmonicity. It may also be partly due to the cation being frozen in one particular orientation in the model of the cubic phase, although this should not be an issue for the orthorhombic phase.

As noted earlier, we model the measured spectra, \( \tilde{\sigma}(\omega) \), using a set of Lorentz oscillators, with two oscillators for each phonon mode. The calculated loss function is shown as a black solid line in Fig. 3(a). The combined oscillator strength of modes 2 and 3 [shown as a blue dashed line in Fig. 3(a)] is conserved below 140 K, suggesting that it is mode 2 in the tetraII phase that splits into modes 2 and 3 in the orthorhombic phase.

The evolution of the fitted oscillator strengths of modes 2 and 3 with temperature is shown in Fig. 3(a). The combined oscillator strength of modes 2 and 3 [shown as a blue dashed line in Fig. 3(a)] is conserved below 140 K, suggesting that it is mode 2 in the tetraII phase that splits into modes 2 and 3 in the orthorhombic phase. Note that peaks in \( \sigma(\omega) \) give the frequencies (\( \omega_{TO} \)) and oscillator strengths of the TO phonon modes. Although distortions in the inorganic cages lead to local breaking of the space-group symmetry, which gives these TO phonon modes both TO and longitudinal optical (LO) character, we proceeded to plot the loss function \( -\text{Im}[1/\tilde{\epsilon}(\omega)] \) [which is the negative imaginary part of the complex response function \( 1/\tilde{\epsilon}(\omega) \)], whose peaks more accurately determine the LO phonon mode frequency \( \omega_{LO} \). Figure 3(b) shows the 300 K loss function data in red circles. Due to the frequency window of our setup, only one peak can be identified at 1.59 THz. However, since the fits to both the real and imaginary parts of \( \sigma(\omega) \) have been obtained, we use the fitting parameters to compute the loss function up to 5 THz. The calculated loss function is shown as a black solid line in Fig. 3(b), showing two peaks at \( \omega_{LO,1}/2\pi = 1.59 \) THz and \( \omega_{LO,2}/2\pi = 3.55 \) THz. Note that the positions of these two peaks are consistent with the Cochran-Cowley relation:

\[
\frac{\epsilon_{\text{static}}}{\epsilon_{\infty}} = \prod_{i=1}^{2} \frac{\omega_{LO,i}^2}{\omega_{TO,i}^2},
\]

where \( \epsilon_{\text{static}} \) is the static dielectric constant, thus giving an independent confirmation of the accuracy of these two values (see the supplementary material for details). The second LO-phonon peak was observed at 5 THz by Sendner et al., which differs from our 3.55 THz value. In the supplementary material, the possible reasons for the discrepancy are described in detail, such as the uncertainty in the value of \( \Delta L \).
TABLE I. Calculated 300 K carrier mobility based on carrier-LO phonon scattering in MAPbBr3. The LO phonon mode frequencies are taken from the peaks in the loss function $-\text{Im}[1/\varepsilon(\omega)]$. The effective LO phonon mode frequency is obtained by the sum of weighted LO modes. Mobilities are calculated using the effective LO phonon mode.

<table>
<thead>
<tr>
<th>Material</th>
<th>$\epsilon_{\infty}$</th>
<th>$\epsilon_{\text{static}}$</th>
<th>$\omega_{\text{LO,eff}}/2\pi$</th>
<th>Carrier</th>
<th>$m_0/m_h$</th>
<th>$\alpha$</th>
<th>$w$</th>
<th>$v$</th>
<th>$\mu$ (cm$^2$ V$^{-1}$ s$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MAPbBr$_3$</td>
<td>6.7</td>
<td>25</td>
<td>3.46 THz</td>
<td>$e$</td>
<td>0.25 (Ref. 30)</td>
<td>1.68</td>
<td>11.4</td>
<td>13.0</td>
<td>83</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>$h$</td>
<td>0.12 (Ref. 30)</td>
<td>1.17</td>
<td>11.5</td>
<td>12.6</td>
<td>265</td>
</tr>
</tbody>
</table>

and the determination of LO phonon mode frequency outside the experimental frequency window.

In systems with multiple LO-phonon modes, Hellwarth et al. derived an effective LO mode frequency $\omega_{\text{LO,eff}}$ from the frequencies and weights of the individual modes. This value of $\omega_{\text{LO,eff}}$ is then used to calculate the carrier mobility according to Hellwarth’s$^{36}$ and Feynman’s$^{37}$ theories. In our case, we use two Lorentz oscillators to fit the complex response function to obtain the weight $W_i^2$ for each LO mode $\omega_{\text{LO},i}$. We then obtain $\omega_{\text{LO,eff}}$ from

$$\omega_{\text{LO,eff}}^2 = \sum_i W_{\text{LO},i}^2 \sum_j \left( \frac{W_{\text{LO},j}^2}{\omega_{\text{LO},j}^2} \right),$$

(4)

To finally obtain $\omega_{\text{LO,eff}} = 3.46$ THz in MAPbBr$_3$.

In photovoltaic applications, it is always desirable to have high photogenerated carrier mobility at room temperature. The reported values vary from 10 to 100 cm$^2$ V$^{-1}$ s$^{-1}$, depending on the sample preparation method, measurement approaches, and corresponding assumptions.$^{38-40}$ One key question is that of whether there is an upper limit to the carrier mobility in hybrid perovskites.

In polar semiconductors, carrier mobility is limited by scattering with LO phonons. We estimate the upper limit of the carrier mobility using Feynman’s polaron theory.$^{36,37}$ This theory has been applied to various semiconductor systems, such as halide perovskites,$^{16,41}$ InSb,$^{42}$ Ca$_1-x$La$_x$MnO$_3$,$^{43}$ TiO$_2$,$^{44}$ and Bi$_{12}$SiO$_{20}$.$^{45}$

The carrier mobility, derived for large (Fröhlich) polarons, follows the expression$^{36,37}$

$$\mu = \frac{w^3 3e}{v^3 m_b \omega_{\text{LO,eff}}^2 2b \beta^2} \left( \frac{\beta}{K(a, b)} \right),$$

(5)

where $e$ is the elementary charge, $m_b$ is the band effective mass of the electrons or holes, $\alpha$ is the carrier-phonon coupling constant, and $\beta = h\omega_{\text{LO,eff}}/k_B T$, with $h$ the Planck constant, $k_B$ the Boltzmann constant, and $T$ the temperature. The factor $K(a, b)$ is a function of $w$ and $v$: $K(a, b) = \int_0^\infty du [\alpha^2 + \beta^2 - b \cos(\beta u)]^{-3/2} \cos(\beta u), \quad \beta = (\beta/2)^2 + R \beta \coth(\beta/2), \quad R = (v^2 - w^2)\beta v$.

The variational parameters $w$ and $v$ are obtained when the Osaka free energy achieves a minimum for a LO phonon at a particular temperature. We obtained the values of $w$ and $v$ for $\omega_{\text{LO,eff}}$ at 300 K.$^{36,37}$ The carrier-phonon coupling constants are calculated via $\alpha = \sqrt{m_b e^4 / 2\hbar^2 c^2 E_{\text{LO}}}, \quad 1/\epsilon_s = 1/\epsilon_{\text{static}}, \quad E_{\text{LO}} = \hbar\omega_{\text{LO,eff}}$, and are listed in Table I. The electron-phonon coupling constant $\chi_{e-ph} = 1.68$ agrees well with the value of 1.69 obtained from DFT calculations.$^{41}$ The small values of $\alpha$ (1–2) indicate that the system is in the large-polaron regime ($\alpha < 6$), and so justifies the use of Eq. (5) in our calculation of carrier mobility.

We summarize the parameters and calculated mobilities for electrons and holes in Table I. Mobilities of 10–100 cm$^2$ V$^{-1}$ s$^{-1}$ obtained from measurements on single-crystals,$^{36,39}$ especially the high hole-mobility ($\sim 210$ cm$^2$ V$^{-1}$ s$^{-1}$), agree well with our values. Note that Sendner et al.$^{22}$ performed the same analysis on MAPbBr$_3$ to obtain $\alpha = 1.69$. Also they used the reduced exciton mass in the mobility calculation to obtain $\mu = 158$ cm$^2$ V$^{-1}$ s$^{-1}$, whose value lies between our values of electron and hole mobilities.

From Eqs. (4) and (5), we see that the low-frequency THz modes make the largest contribution to the loss function, and therefore determine the effective frequency and, hence, upper limit to the carrier mobility. In real materials, carriers are also scattered by grain boundaries, defects, disor-
der, and ion impurities,$^{47}$ and so the observed mobility will likely be smaller than our calculated upper limit. Note that our previous work on MAPbI$_3$ used the low-temperature expression from Feynman’s theory, which calculated the low-temperature mobility, and therefore obtained values larger than those in this work.$^{16}$

In conclusion, we have studied the THz phonon modes in the hybrid halide perovskite MAPbBr$_3$. Phonon mode splitting is observed at 150 K, which we attribute to the tetra-
gonal–orthorhombic phase transition. We use Feynman’s polaron theory to estimate the upper limit of the carrier mobility based on scattering with LO phonons, and obtain values that agree well with other measurements. Our method gives an upper limit to the carrier mobility from the knowledge of the low-frequency phonon modes, without the need to generate carriers via photoexcitation. This method can be applied to other polar semiconductors where LO phonon frequencies, carrier-phonon coupling constant, and band-effective mass are known.

See supplementary material for the discussions on the uncertainty in the calculated loss function, consistency with the Cochran-Cowley relation, why the lowest phonon modes dominate the carrier mobility, and the real part of the dielectric function at 300 K.

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