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Interpretation of Resistance, Capacitance, Defect Density, and Activation Energy Levels in Single-Crystalline MAPbl₃

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the interpretation of resistance, capacitance, defect density, and activation energy levels in single-crystalline methylammonium lead iodide (MAPbI₃). In particular, the impedance response of the MAPbI₃ crystal as a function of applied bias and temperature (under both increasing and decreasing temperature cycles) is studied for the first time. From the detailed bias- and temperature-dependent studies, we found that the low-frequency capacitance values are influenced by ion density and mobility. Consequently, single-crystalline MAPbI₃ depicts an activation energy of 0.53-0.54 eV with an exceptionally low electronic trap density of $0.96 \times$



 10^{10} cm⁻³. The present study illustrates that the net electrochemical impedance spectra are due to ionic capacitance coupled to a resistance. The associated resistance is related to the conductivity of the perovskite crystal. These findings are helpful to understand the fundamental electrical properties of the MAPbI₃ single crystal, which could be useful for the further advancement of perovskite single-crystal-based applications.

1. INTRODUCTION

In the last 10 years, hybrid inorganic–organic lead halide perovskites have become the center of research interest as one of the most promising candidates for the development of nextgeneration photovoltaic¹⁻⁴ and optoelectronic devices.⁵⁻⁷ These semiconductors exhibit outstanding electrical and optical properties with the scopes of cost-effectiveness and ease of thinfilm fabrication processing.^{8,9} However, low operational stability¹⁰ and polarization at low frequencies/long time scales affect the charge transport and recombination during the operation of devices.^{11,12} It has been reported that issues mainly governed by ions and vacancy conductions need to be addressed to make perovskite solar cells (PSCs) commercially available.^{13,14}

Recently, extensive efforts have been devoted to study the growth methods and physicochemical characterizations of single-crystalline metal halide perovskites.¹⁵ Methylammonium lead iodide (MAPbI₃) is a prototypical photoabsorber in PSCs,^{16,17} and its single-crystalline form exhibits higher carrier diffusion length and carrier lifetime and a lower trap state density than the corresponding polycrystalline perovskite films.^{18,19} A number of studies involving other metal halide perovskites have also univocally shown that the optoelectronic properties of the single-crystalline perovskites are much better compared to those of pellets or thin-film counterparts due to the less amount of

grain boundaries and defects.^{18,20–22} Therefore, the utilization of perovskite single crystals for optoelectronic application has emerged as a new direction for boosting the efficiency and moisture stability of PSCs.^{23–25}

It is well established that migration of ionic species in a perovskite leads to instability in PSCs and has also been proposed as a possible cause of the hysteresis effect.²⁶ The activation energy of migrating ions in MAPbI₃ has been estimated theoretically and experimentally and covered a wide range of 0.2-0.6 eV.^{20,26-33} Due to this discrepancy on the energy barrier that determines the ion transport, it is still desirable to shed more light on the mechanism of ion conduction in perovskite materials.³⁴ The activation energy and electronic defect density of perovskite single crystals found in the literature are summarized in Table 1.

In this work, we report on the bias- and temperaturedependent behavior of single-crystalline $MAPbI_3$ using electrochemical impedance spectroscopy (EIS) and capacitance spectroscopy. Our results reveal that the low-frequency

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Table 1. Calculated Values of Activation Energy andElectronic Defect Density for Perovskite Single Crystals

activation energy and t	rap density of MAPbI ₃	single crystals
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activation energy	measurements	refs	
0.624 eV (I ⁻) (single crystal)	temperature-dependent ion conductivity	29	
0.45 eV (I ⁻) 0.70 eV (MA ⁺) (single crystal)	polarization-induced current—voltage hysteresis	30	
0.58 eV (I ⁻)	first principles	28	
2.31 eV (Pb ²⁺)			
0.84 eV (MA ⁺)			
0.82 eV (thin film)	temperature-dependent ion conductivity	31	
0.119 eV (MA ⁺)	Quasi-elastic neutron scattering measurements and first-principles analysis	32	
0.53 eV (I ⁻)	temperature-dependent electrochemical impedance spectroscopy (EIS)	26	
0.51 eV (I ⁻)	temperature-dependent ion conductivity from Warburg impedance	26	
1.05 eV (single crystal)	temperature-dependent ion conductivity	27	
0.5 eV (thin film)			
0.1 (I ⁻)	NMR spectra	20	
0.43 eV (I ⁻) (pellets)	temperature-dependent ion conductivity	33	
trap density of MAPbX ₃ (X = I, Br, Cl) single crystal using space-charge- limited current (SCLC) regime			
single crystals	trap density (cm ⁻³)	refs	
MAPbI ₃	5.8×10^{9}	35	
MAPbI ₃	3.3×10^{10}	18	
MAPbCl ₃	3.1×10^{10}	33	
MAPbI ₃	1.4×10^{10}	19	
MAPbBr ₃	3×10^{10}	19	

capacitances are influenced by the ion density and mobility. We demonstrate that the electrical parameters such as capacitance, resonance time, and impedance are temperature- and voltagedependent. Understanding the behavior of these parameters can provide a way to design and optimize perovskite crystals employed for various optoelectronic applications. In this regard, the conventional electrical direct current (DC) measurements still remain limited. The activation energy corresponding to ion migration was calculated to be in the range of 0.53-0.54 eV. Moreover, we find that the MAPbI₃ crystal depicts an exceptionally low trap density of 0.96×10^{10} cm⁻³. This study also shows that the EIS response of the MAPbI₃ single crystal exhibits different behavior from that observed in PSCs. While the EIS spectra of PSCs are defined by two semicircles, a single semicircle is observed for perovskite single crystal. The present study illustrates that the net electrochemical impedance spectra

are due to ionic capacitance coupled to a resistance. The associated resistance is related to the conductivity of the perovskite crystal. These results are helpful to understand the fundamental electrical properties of single-crystalline MAPbI₃, which could be useful for the further advancement of perovskite single-crystal-based applications.

2. EXPERIMENTAL SECTION

2.1. Synthesis of MAPbl₃ Single Crystals. The perovskite crystals were grown briefly using a 1.0 M MAPbI₃ solution prepared by dissolving equimolar amounts of precursors in γ -butyrolactone at 60 °C overnight. Before crystallization, the solution was filtered using a 0.2 μ m pore size PTFE filter. Next, 2 mL of the filtered precursor solution was kept at 160 °C for 30 min in an oil bath, which resulted in the growth of very small seed crystals. The seed crystal solution was further heated at 120 °C for 3 h. Finally, the remaining solution was discarded and the crystals were washed with acetone 2–3 times and then dried.

2.2. Characterization of Perovskite Single Crystals. Temperature-dependent powder X-ray diffraction (XRD) patterns were recorded with an Empyrean diffractometer (PANalytical) equipped with a copper lamp (40 kV, 40 mA). The samples were mounted inside the Anton Paar TTK-450 sample chamber for nonambient X-ray diffraction experiments. For the measurements, Ni-filtered Cu K α (λ = 0.154 nm) radiation was used and was detected with an X'Celerator 1D detector in the Bragg–Brentano θ – θ geometry. The XRD patterns were recorded over a 2θ range of 10–50° without rotating the sample. EIS measurements were performed using a potentiostat Autolab equipped with a frequency response analyzer. EIS measurements were performed as a function of applied bias and temperature.

3. RESULTS AND DISCUSSION

3.1. Effect of the Applied Bias. MAPbI₃ single crystals were grown using a recently reported procedure (for details, see the Section 2).³⁶ The electrical properties of a MAPbI₃ single crystal were studied using alternating current (AC) impedance spectroscopy in the frequency range of 1 Hz to 1 MHz under the dark condition in the temperature range of 30-90 °C. Additionally, a variable direct current (DC) voltage is applied on the sample during the electrochemical impedance spectroscopy (EIS) measurement. The impedance curves of the MAPbI₃ single crystal are found to be strongly affected by the measurement conditions. To perform temperature- and bias-dependent EIS measurements, thin layers of silver (Ag) were deposited on both sides of a MAPbI₃ single crystal with



Figure 1. EIS data of MAPbI₃ single crystal at ambient temperature (303 K) as (a) a function of applied bias measured under the dark condition and (b) capacitance $\left(\frac{\text{Im}(1/Z)}{\omega}\right)$ plot as a function of applied bias in the frequency range of 1 MHz to 1 Hz.

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Figure 2. Dark EIS measurement of the MAPbI₃ single crystal at 0 V DC bias in the frequency range of 1 MHz to 1 Hz as a function of temperature (313–363 K). Nyquist plots of the crystal (indicating approximately semicircles), by (a) increasing and (b) decreasing temperature. Apparent capacitance $\left(\frac{\text{Im}(1/Z)}{\omega}\right)$ of the crystal vs frequency curves, by (c) increasing and (d) decreasing temperature (the value of capacitance is increased with increasing temperature in the low-frequency region <1000 Hz).

dimensions of $2.26 \times 2.07 \times 0.6$ mm³. To confirm the validity of results, we repeated each measurement several times under the same conditions. All experiments were conducted on different crystals from the same batch. We observed that the crystals have different sizes even when crystallized from the same batch. Therefore, in all further experiments, we use the measuring units without any conversion factor.

First, we conduct the EIS measurement as a function of applied DC bias at ambient temperature, as shown in Figure 1a. The Nyquist plots of MAPbI₃ crystals show a suppressed single semicircle. The distortion in the semicircle is due to the involvement of the physical process associated closely with the low and high frequencies. To test the reproducibility of our results, we measured the EIS response of other MAPbI₃ crystals and found similar behavior (see Figure S1). A decrease in the radii of the semicircles was observed with an increase in the applied bias from 0 to 1 V, which suggests that the applied potential has a significant impact on the charge characteristics.²⁶ In our previous works, we demonstrated that the EIS spectra (Nyquist plots) of PSCs show two well-separated semicircles in the dark and under illumination.^{37,38} These semicircles were associated with the low- and high-frequency responses of the devices. The high-frequency response is due to the recombination resistance and geometrical capacitance, whereas the lowfrequency response is due to the coupled response of ions and charges.³⁹ In the present case, we believe that the single semicircle is due to symmetrical contacts and the single-crystal nature of MAPbI₃. In PSCs, the recombination takes place at interfaces between the perovskite absorber layer and contacts and at grain boundaries. In EIS, these features are observed in the high-frequency region $(10^5 - 10^6 \text{ Hz})$ and analyzed by the real part of high-frequency impedance.^{37,38,40} However, no such response is observed for MAPbI₃ single crystals. It is possible that the absence of grain boundaries, use of symmetrical

contacts, and dominant ionic response in the low-frequency capacitance $(<10^3 \text{ Hz})$ have suppressed the high-frequency features of MAPbI3 crystals. To further test the occurrence of Nyquist spectra with a central frequency in the range of $< 10^3$ Hz, MAPbBr₃ single crystals were synthesized and measured. Figure S2 shows the single semicircle Nyquist spectra for the MAPbBr₃ crystal, confirming that the EIS spectra of the single crystal differ from those found in PSCs. Figure S3 shows the resonance time constant (au) as a function of applied bias. The resonance time (τ) is obtained using the expression $\tau = 1/\omega_{(\min)}$, where $_{(\min)}$ is the angular frequency, corresponding to the minimum of imaginary of complex impedance. We found that τ is affected by the applied bias. Note that the obtained magnitude of τ in the range of ms is too high in magnitude to explain the electronic charge recombination, which generally occurs in the time scale of μ s. The origin of the τ in MAPbI₃ single crystals is another topic, which requires further discussion.

Figure 1b shows the capacitance vs frequency (C-F) plots as a function of applied bias. Two distinguishing features of capacitance corresponding to low- and high-frequency regions are observed. In general, the capacitance at low frequency ($<10^3$ Hz) is associated with ion and charge accumulation or electrode polarization, whereas the high-frequency capacitance is due to geometrical capacitance.⁴¹ With the increase in the applied bias, the low-frequency capacitance increases and also shifts to high frequency. Neukom et al. analytically and experimentally investigated the role of ion density and mobility in the C-Fplot of perovskites.⁴² It was concluded that a variation in the ion mobility changes the transition frequency; the low-frequency capacitance value increased with increasing ion density. Another study revealed that a higher bulk conductivity leads to a higher capacitance, which further increases with the applied bias and illumination.⁴³ Therefore, the obtained capacitance response at

the frequency ($<10^3$ Hz) of MAPbI₃ crystals is said to be influenced by the ion density and mobility.

To further investigate the change in the conductivity of MAPbI₃ crystals due to the applied bias, the value of resistance as a function of applied bias is plotted (Figure S4). The value of resistance was obtained from the real part of the impedance spectra. We observe a gradual decrease in resistance with increasing bias. This phenomenon is consistent with the recent study by Moia et al., in which this behavior was described as the "ionic-to-current amplification", and mobile ions open the door for the electronic charge transport.⁴³ Figure S5a shows the current-voltage (I-V) characteristic of the MAPbI₃ crystal measured under forward and reverse directions and the dark condition at a scan rate of 100 mV s⁻¹. The measured I-Vresponse exhibits the hysteresis feature. In perovskite solar cells, the existence of hysteresis is very common and generally assigned to the ionic nature of the perovskite semiconductor. The difference in the current magnitude during I-V measurements in forward and reverse directions (see Figure S5a) confirms that the net electronic current is influenced by the ions. Figure S5b shows the I-V plot of the MAPbI₃ crystal as a function of the voltage sweep rate. It was found that iodine ions are responsible for hysteresis in PSCs due to its 4 orders higher diffusion coefficient $(10^{-12} \text{ cm}^2 \text{ s}^{-1})$ compared to MA⁺ ions $(10^{-16} \text{ cm}^2 \text{ s}^{-1})$.^{28,44} In the present case, the increase in hysteresis is observed at a low scan rate. At a low scan rate, the iodine ions have sufficient time to move through the perovskite structure. Moreover, it is well established that the hysteresis index is proportional to the magnitude of low-frequency capacitance.^{45,46} From the observation of C-F and I-Vanalyses, we conclude that the low-frequency capacitance and hysteresis in the I-V curve are due to mobile ions.

3.2. Effect of Applied Temperature at Fixed Bias. To study the effect of temperature on the EIS response of the MAPbI₃ crystal, the EIS spectra were collected at a range of different temperatures (303-363 K) at 0 V bias under the dark condition. Before the measurement, powder X-ray diffraction of the MAPbI₃ crystal was measured to ensure the structural stability at applied temperatures. The temperature-dependent powder X-ray diffraction (pXRD) analysis confirms the structural stability of the MAPbI₃ crystal after heating up to 90 °C (Figure S6). Moreover, we observed the tetragonal–cubic phase transition between 323 and 333 K.⁴⁷

The impedance responses in the form of Nyquist and C-F plots as a function of temperature are shown in Figure 2. The collected Nyquist plots exhibit a similar trend upon increasing and decreasing temperatures, which indicates that the electronic properties of the investigated crystal are stable under probed temperature ranges. We observe a decrease in the semicircle radius of the EIS spectra with increasing temperature (Figure 2a). Moreover, after plotting the Bode curves shown in Figure S7, there is a shift in resonance frequency toward the higher-frequency range with increasing temperature. The observed shift suggests that there is a temperature-activated process. Comparing our results with other reported works on EIS studies of PSCs under different temperatures, the following key observations are concluded:

1. In PSCs, the high-frequency resonance peak is generally observed in the frequency range of 10^5-10^6 Hz and mainly assigned to the recombination resistance and geometrical capacitance.^{38,40} In the present study on the MAPbI₃ crystal, the frequency resonance peak was

observed in the range of 10^2-10^3 Hz and there is no peak in the frequency range of 10^5-10^6 Hz.

2. According to the recent report by Garcia-Rodriguez et al. on the role of bromide content in iodine migration, the temperature has a least impact on the high-frequency spectra (10^5-10^6 Hz) .⁴⁰ In turn, the maximum shift at low- and mid-frequency (10^2-10^3 Hz) spectra is observed with change in temperature. Consistent with the finding of the mid-frequency (10^2-10^3 Hz) spectra of PSCs, the MAPbI₃ crystal also shows a similar temperaturedependent behavior in the same frequency range. The shift in the probed frequency range is seen by plotting the variation of τ against temperature, as shown in Figure S8.

These results suggest that the EIS response of PSCs is different from that found in single crystals, where the net capacitance response in the frequency range ($<10^3$ Hz) is mainly governed by the mobile ions in the later case. Note that parameters such as capacitance, resonance time, and impedance are found to be temperature- and voltage-dependent. Understanding the effect of these parameters can provide a way to design and optimize perovskite crystals, which are employed for various optoelectronic applications, while the conventional electrical DC measurements, remain limited in this regard. After establishing the fact that the EIS response of crystals is due to ionic capacitance coupled to a resistance, in the next step, we calculate the activation energy of ions and electronic density of defects.

Figure 2c,d shows the frequency-dependent capacitance plots at different temperatures. No significant difference is observed between the C-F curves measured upon increasing and decreasing temperature cycles. As expected, the capacitance in the low-frequency region increases with increasing temperature. From the obtained C-F plot, a change in inflection frequencies (f_0) in the range of 10^1-10^2 Hz is observed. Figure 3 shows the



Figure 3. Arrhenius plots of the inflection frequencies vs 1000/T (ln (f_0) vs 1000/T) during increasing and decreasing the temperature of the MAPbI₃ single crystal. E_{act} is the activation energy for the traps.

plots of $\ln(f_0)$ vs 1/T. By taking the slope of these curves, activation energies of 0.53 ± 0.17 and 0.54 ± 0.14 eV during heating and cooling of the MAPbI₃ crystal were obtained, respectively. A small difference in the value of activation energy for the MAPbI₃ crystal was observed during the heating and cooling measurement. This might be due to the local lattice distortion connected with the tetragonal-cubic phase transition. In the literature, point defects due to the ion migration in MAPbI₃ have been related to the activation energy ranging from 0.1 to 0.6 eV, which depends on the applied characterization method and material crystallinity (Table 1).^{26,28,48,49} The

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Figure 4. Trap density measurement for the MAPbI₃ single crystal. Temperature-dependant dark current–voltage responses of the crystal by (a) increasing and (b) decreasing the temperature.

obtained activation energy corresponds to the ion migration, which further confirms our results that the associated capacitance in the low-frequency range ($<10^3$ Hz) is due to mobile ions. Very recently, Almora et al. have studied the temperature-activated low-frequency capacitance ($<10^1$ Hz) for PSCs and attained $\sim10^{21}$ eV⁻¹ cm⁻³ density of states (DOS).⁵⁰ It was stressed that such a high value of DOS is hardly connected to ionic or electronic defects levels.

To calculate the electronic trap states in the crystals, a wellestablished space-charge-limited current (SCLC) for perovskite crystals is used.^{18,19,22,35} To quantify the electronic defect density in MAPbI₃ crystals, dark current–voltage (*I–V*) measurements as a function of temperature were carried out. As shown in Figure 4, the *I–V* characteristic of the crystal exhibits different slopes at different bias. In the low applied bias, a linear region is observed that causes a sharp rise ($I\alpha V^{n>3}$) at a high applied bias. This voltage is called the kink point or trapfilled limit voltage (V_{TFL}).⁵¹ By considering a linear dependence between trap density (η_{traps}) and V_{TFL} , the defect density is calculated using the given expression

$$\eta_{\rm traps} = \frac{2\varepsilon\varepsilon_0 V_{\rm TFL}}{eL^2} \tag{1}$$

where e and ε_0 are the elementary charge (1.6 × 10⁻¹⁹ C) and vacuum permittivity (8.854 × 10⁻¹² F m⁻¹), respectively, and L and ε are the thickness of the crystal (~0.6 mm) and dielectric constant $(28.8)^{52}$ of the MAPbI₃ single crystal, respectively. We calculated V_{TFL} for each cycle and estimated the trap density. The average value of $\eta_{\rm traps}$ equals 0.96×10^{10} cm⁻³, which is in agreement with the previous study.¹⁸ Interestingly, the trap density is significantly lower than that found in the well-known inorganic semiconductor like single crystals of CIGS, ⁵³ CdTe, ⁵ or Si.⁵⁵ It should be noted here that the applied method is mainly associated with the electronic defect level. To elucidate the role of ions in the electronic trap states, we measure the I-Vhysteresis and calculate the trap density (see Figure S5c). From the obtained plot, hysteresis behavior confirms that the change in current is due to ions and has a negligible effect on the trap density. This signature requires further electrical or optical characterization to clearly elucidate the role of ions on trap defect density. Our results confirm that the MAPbI₃ single crystal has a very high quality with very low defect density, which is required for high-performance photoelectronic devices such as the photodetector.

4. CONCLUSIONS

In conclusion, we systematically studied the electrical properties of single-crystalline MAPbI₃, such as resistance, capacitance,

defect density, and activation energy levels. Detailed bias- and temperature-dependent studies revealed that the low-frequency capacitance values are influenced by the ion density and mobility. We found that electrical parameters such as capacitance, resonance time, and impedance are temperature- and voltage-dependent. The calculated activation energy values in the range of 0.53–0.54 eV correspond to ion migration. The MAPbI₃ crystal depicts an exceptionally low electronic trap density of 0.96 × 10¹⁰ cm⁻³. These findings are helpful to understand the fundamental electrical properties of single-crystalline MAPbI₃.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.jpcc.9b11343.

Nyquist plots of a MAPbI₃ and MAPbBr₃ single crystals; relaxation time constant; real impedance; I-V curves under forward and reverse bias; temperature-dependent powder X-ray diffraction (pXRD); phase-frequency Bode plot; and Arrhenius plots of time constant vs 1000/*T* of MAPbI₃ single crystal (PDF)

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Notes

The authors declare no competing financial interest.

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