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# PAPER

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# Phase transitions and optical properties of the trigonal perovskite $(CH_3NH_3)_2TeCl_6$ <sup>†</sup>

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Vacancy-ordered double perovskites  $A_2BX_6$  are attracting attention due to the isolated octahedra making them an ideal playground for studying perovskite structure-property relationships.  $MA_2TeCl_6$  single crystals have been successfully grown from solution. X-ray structure experiments as a function of temperature have demonstrated two structural phase transitions: from cubic  $Fm\bar{3}m$  (high temperature, phase I) to trigonal  $P\bar{3}m1$  (intermediate temperature, phase II) at 460 K and then to trigonal  $P\bar{3}1c$  (low temperature, phase III) at 200 K. Intriguingly, we observe a negative thermal expansion for the *c*-axis in phase II believed to caused by dynamic tilting of the C–N bond in the MA<sup>+</sup> ion. We have modified and extended the calculation of the Brown radii ratio in the perovskites to include hybrid  $A_2BX_6$  compounds by selecting the shortest A–X distance for the ratio equation. We find that for ratios between 0.89 and 1.17, typically the  $Fm\bar{3}m$  cubic structure is observed, while for ratios above 1.17, mostly distorted structures are found.

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## Introduction

Perovskite materials have received a great amount of attention in the photovoltaic field; however, chemical instability and toxicity issues are driving the demand for stable lead-free alternatives to the commonly used lead containing formulations. Vacancy-ordered double-perovskites  $A_2BX_6$  with B equal to a Group 16 metal offer a new avenue to replace hybrid lead iodide perovskites in photovoltaic applications due to improved air and moisture stability whilst retaining similar properties to the perovskite ABX<sub>3</sub> compounds. The double-perovskite structure is obtained by replacing half of the B-site cations with vacancies in an ordered fashion, which leads to doubling of the lattice parameter. This extended unit cell allows for greater flexibility in compositional modification.<sup>1-3</sup> Vibrations of the organic cations and octahedra contribute significantly to the lattice

dynamics, which, in turn, play an essential role in the material properties.<sup>4</sup> Hence, it is interesting to study the influence of the organic cations on the structure–property relationships to further understand these materials.<sup>2,5</sup>

Recently, tellurium-based double perovskites have been identified as potential lead-free solar cell materials with good electronic attributes and chemical stability.<sup>6</sup> In this study, we have added to this new field by investigating the crystal structure of (CH<sub>3</sub>NH<sub>3</sub>)<sub>2</sub>TeCl<sub>6</sub> (or MA<sub>2</sub>TeCl<sub>6</sub>) from 120 K to 500 K by variable temperature powder and single-crystal X-ray diffraction and neutron powder diffraction. Three phases were observed across the temperature range for MA2TeCl6; this contrasts with some other tellurium halides, for example Cs<sub>2</sub>TeCl<sub>6</sub> where no phase transitions were previously reported. We report a complete structure solution for each crystallographic phase. The cation dynamics of MA<sub>2</sub>TeCl<sub>6</sub> are notably different from other compounds in the series such as MA2SnCl6, which drives the necessity to study its structure-property relationship. The crystal structures analysed in our work, combined with other experiments to identify the phase-transition mechanism, will be helpful for further property investigations.<sup>7</sup> Finally, we have extended the Brown radiiratio calculation for perovskites to include the hybrid A2BX6 perovskites and have demonstrated an improved method to describe the lattice distortion that uses the shortest distance between the A-site and X-site atoms to calculate the radius ratio.

# Experimental

Tellurium oxide and aqueous methylamine were separately reacted with 12 M hydrochloric acid to form  ${\rm TeCl}_4$  and

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<sup>†</sup> Electronic supplementary information (ESI) available: Additional experimental detail, crystallographic tables, and additional supporting figures are supplied. CIF files have been deposited at the Cambridge Crystallographic Data Centre with REFCODES 2142618 (90 K), 2142619 (120 K), 2142620 (160 K), 2142621 (200 K), 2142622 (250 K) and 2142623 (295 K). Neutron powder diffraction data is available with https://doi.org/10.5286/ISIS.E.RB2090007-1. See DOI: https://doi.org/10.1039/d2tc02372e

methylammonium chloride (MACl) solutions. The acidic TeCl<sub>4</sub> and MACl solutions were mixed at room temperature and the resulting suspension was then filtered and dried to obtain MA<sub>2</sub>TeCl<sub>6</sub> as a microcrystalline powder. MA<sub>2</sub>TeCl<sub>6</sub> single crystals were grown by redissolving this MA<sub>2</sub>TeCl<sub>6</sub> powder in 12 M HCl at 60 °C to obtain a saturated clear MA<sub>2</sub>TeCl<sub>6</sub> solution, which was then slowly cooled to room temperature over 5 h to encourage crystal growth.

Single-crystal X-ray diffraction (SXD) measurements were made on  $MA_2TeCl_6$  using an Agilent SuperNova diffractometer between 90 K and 295 K. Full spheres of data were measured with Mo K $\alpha$  radiation in around 1 to 2 hours depending on temperature.

A room temperature powder X-ray diffraction (PXRD) pattern was obtained using a Stoe Stadi-P X-ray diffractometer in thinfoil transmission mode with Mo radiation. Subsequently, variable temperature powder X-ray diffraction (VT-PXRD) patterns were obtained using a Stoe Stadi-P X-ray diffractometer in capillary mode with Cu radiation. An Oxford Instruments CryojetHT was used to cool the sample to 120 K and then heat it in increments of 20 K to 500 K with PXRD patterns obtained at each temperature.

Room temperature neutron powder diffraction (NPD) was used to provide additional information on the light atom positions as carbon and nitrogen can be hard to distinguish reliably by X-rays in the presence of heavy atoms. Time-of-flight data were collected from  $MA_2TeCl_6$  using the Polaris at the ISIS Muon and Neutron Spallation Source, U.K.

Variable temperature optical absorption spectra were collected using an Agilent Cary 4000 UV Vis Spectrometer over the wavelength range 175 to 900 nm from 175 K to 500 K in increments of 25 K.

Further experimental details are available in the ESI.†

## Results and discussion

#### Crystal structure

A light vellow hexagonal single crystal was selected for SXD measurements, as shown in Fig. S1 (ESI<sup>+</sup>); the sample has a plate-like morphology with its width around ten times greater than its thickness. The crystal structures of the different phases of MA<sub>2</sub>TeCl<sub>6</sub> are shown in Fig. 1. From the SXD analysis, the crystal system of MA<sub>2</sub>TeCl<sub>6</sub> at room temperature (295 K) is trigonal, space group symmetry  $P\bar{3}m1$  with lattice parameters a = 7.3565(4) Å, c = 7.0812(7) Å, and Z = 1, which is in agreement with the literature.8 We denote this as phase II where the  $[TeCl_6]^{2-}$  octahedra occupy the corners of the hexagonal unit cell while the MA<sup>+</sup> cations lie between the octahedral units. Thus, the material consists of layers of  $[TeCl_6]^{2-} - MA^+ - MA^+ - MA^+$  $[\text{TeCl}_6]^{2-}$  stacked along the *c* axis with one formula unit per unit cell and with the axis of the MA<sup>+</sup> cations aligned parallel to c. The  $MA^+$  cations are orientated such the  $NH_3^+$  ionic head group points towards the Cl of three [TeCl<sub>6</sub>]<sup>2-</sup> anions with the non-polar -CH<sub>3</sub> groups lying in a plane formed by opposing  $MA^+$  cations.

As the temperature is lowered from room temperature to 200 K, jump rotations of the cation about its long axis are reduced in frequency, and a second-order transition to phase III occurs with frozen-in orientations of the MA<sup>+</sup> cation. Phase



**Fig. 1** The crystal structure of  $MA_2TeCl_6$  in (a) phase I at 500 K seen down the *a*-axis shown (upper) and obliquely (lower) in which a single N atom represents the whole  $MA^+$  cation due to the dynamical free rotation; (b) phase II at 295 K seen along the *c*-axis (upper) and down the *a*-axis (lower); and (c) phase III at 200 K seen along the *c*-axis (upper) and down the *a*-axis (lower).

III has space group symmetry  $P\overline{3}1c$  with a = 12.6147(4) Å, c =14.1897(6) Å, and Z = 6. It is related to phase II by a doubling of the unit along c and a  $\sqrt{3}$  increase in a (Fig. S2, ESI<sup>†</sup>). As illustrated in Fig. 1(c),  $[TeCl_6]^{2-}$  octahedra still occupy the corners of the unit cell, but there is a second crystallographic [TeCl<sub>6</sub>]<sup>2-</sup> unit within the enlarged unit cell. The loss of mirror symmetry in going from space group symmetry  $P\bar{3}m1$  in phase II to  $P\bar{3}1c$  in phase III provides a greater freedom to the structure. Thus, to accommodate the MA<sup>+</sup> cations, the [TeCl<sub>6</sub>]<sup>2-</sup> octahedra rotate slightly about the *c*-axis with respect to the layers above and below (Fig. 2(a)). At 90 K, there is a noticeable rotation of *ca.*  $8.45^{\circ}$  for the Te(1) octahedron in contrast to the Te(2) octahedron where the rotation is only  $2.96^{\circ}$  (Fig. S3, ESI<sup>†</sup>). This difference leads to a reduction in the cavity size surrounding the MA<sup>+</sup> ion. Consequently, as supported by NMR results, a minor tilting of the C-N axis is observed in phase III (Fig. 2(b)).<sup>9</sup> Thus, the combination of anion and cation tilts improves the coordination of the  $-NH_3^+$  group by the Cl<sup>-</sup> of  $[TeCl_6]^{2-}$ . As shown in Fig. 2(c), in addition to rotational movement, there is a slight positional change of the octahedra in each layer, which are no longer constrained to lie by symmetry within the same ab plane.

Curiously, calorimetric studies suggest additional low temperature phases of  $MA_2TeCl_6$ ,<sup>7,10</sup> but no new phases were observed by SXD in the range 90 K to 295 K. The origin of this discrepancy is unclear and might be because calorimetry detects more subtle transitions (*e.g.* ferroelectric transitions), which are not easily observed in our SXD experiments. To confirm our observation of just two phases by SXD in the range 90 K to 295 K, VT-PXRD was performed. Initially, the purity of the  $MA_2TeCl_6$  powder at room temperature was verified by

(b)

**Fig. 2** Schematic of MA<sub>2</sub>TeCl<sub>6</sub> (III) showing (a) the rotational twist about the *c*-axis of the  $[Te(1)Cl_6]^{2-}$  anions from one octahedral layer to the next layer; (b) the minor tilting of the C–N axis of the MA<sup>+</sup> cation relative to the *c*-axis (shown as dashed red line); and (c) the slight positional offset of the octahedra in each layer, where red dashed lines represent *ab* planes at c = 0 and c = 1/2. The green colour shades are used to distinguish the Cl atoms belonging to different octahedra.

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**Fig. 3** Background subtracted VT-PXRD data collected on MA<sub>2</sub>TeCl<sub>6</sub> over the temperature range 120–500 K displayed as a surface plot. Diffraction intensity is represented using a red (high) to white (low) colour scale. Dotted lines indicate the temperatures of the two phase transitions. The transition from phase III to phase II is not obvious from this plot given the low intensity of the additional peaks that characterise phase III (see Fig. S5, ESI†).

Rietveld refinement using GSAS II<sup>11</sup> as shown in Fig. S4 (ESI†); all of the observed peaks in the data are accounted for in our structural model demonstrating the high quality of the powder. The CIF file, obtained from the SXD results, was used as the starting point for the refinement and  $R_{\rm wp}$  = 9.19% was achieved.

Our VT-PXRD data as a function of temperature is shown in Fig. 3. Two phase transitions were detected in MA<sub>2</sub>TeCl<sub>6</sub>, a firstorder phase transition on heating between 440 K and 460 K (phase II to phase I) and a second-order phase transition on heating between 200 K and 220 K (phase III to phase II). Focussing on the phase I structure at high temperature, it has been reported previously that the high-temperature phases of  $MA_2TeX_6$  (X = Br, I) are cubic with space group symmetry  $Fm\bar{3}m$ . The high-temperature phase I of MA<sub>2</sub>TeCl<sub>6</sub> seen in a calorimetric study was assumed to be cubic too, but this was not confirmed by X-ray diffraction.<sup>7,10</sup> From our data, we can conclusively show that MA2TeCl6 is isomorphic to MA2TeX6 (X = Br, I) above 460 K, in agreement with the isotropic result from polarised light microscopy.<sup>10</sup> We note that, at 460 K, the VT-PXRD pattern shows the coexistence of phases I and II; this is likely due to the relatively short wait time between changing temperatures and starting the next PXRD measurement in our study.

Next, we consider the low temperature phases II and III of  $MA_2TeCl_6$  seen in Fig. 3. Consistent with a second-order transition, the position of the peaks changes smoothly as a function of temperature down to our lowest temperature of 120 K. Below the transition temperature to phase III, we observe the appearance of several additional weak peaks (see Fig. S5, ESI†) congruous with a phase transition to a lower symmetry trigonal structure with a larger unit cell. This transition is in agreement with the calorimetric study.<sup>10</sup> As the temperature is increased from the base temperature (120 K), most of the peaks

(a)



**Fig. 4** Plot showing the variation (compared with T = 120 K) in lattice parameter and volume as a function of temperature obtained from fits to the data using GSAS II.<sup>11</sup> Values are given in Table S1 (ESI†). Lines joining the data points are a guide to the eye.

shift to lower angles consistent with thermal expansion but, in phase II, several peaks shift to higher angles, this is due to a shortening of the *c*-axis.

In Fig. 4, we show the temperature dependence of the lattice parameters and unit cell volume. For comparative purposes, the unit cell volume *V* is normalised by the number of formula units *Z* in each phase; for phase III, the lattice constant *a* is divided by  $\sqrt{3}$  and *c* is divided by 2. The sudden change in volume between phase II and phase I is evidence of first-order behaviour. In contrast, no discontinuous change in the lattice parameters at the phase II–III boundary is observed, consistent with the second-order transition reported previously.<sup>10</sup>

SXD may struggle to accurately determine the positions of low atomic number atoms in a crystal structure containing heavy atoms. Hence, to complete the characterisation of the room temperature structure (phase II), a complementary neutron powder diffraction (NPD) study was performed to confirm the orientation of the MA<sup>+</sup> cation. Fig. S6 (ESI<sup>†</sup>) shows the refinement results without and with the MA+ cation included in the structure. From previous studies, <sup>1</sup>H and <sup>14</sup>N NMR data indicate that there is a continuous rotation of the methyl and ammonium groups about a non-static C-N axis spreading around a cone with a half angle of 22  $\pm$  1° at 293 K,<sup>7</sup> while the <sup>35</sup>Cl NQR relaxation time indicates a hindered rotation of the [TeCl<sub>6</sub>]<sup>2-</sup> octahedra at room temperature.<sup>9</sup> It should be noted that our model of phase II has the MA<sup>+</sup> cation aligned along the 3-fold symmetry axis: the d-spacing resolution in both SXD and NPD experiments do not allow us to distinguish between atoms lying on the axis and just off the axis with sixfold disorder (see Fig. S7, ESI<sup>†</sup>). The model with the MA<sup>+</sup> cation aligned on the axis uses fewer parameters but results in an apparently shortened C-N bond length consistent with rotation about a cone. For comparison in MAPbI<sub>3</sub>, the MA<sup>+</sup> cations are fully ordered below 100 K, have fourfold disorder in the tetragonal structure at room temperature, and become isotropically disordered in the cubic phase above 352 K.<sup>12</sup>

The bandgap of  $MA_2TeCl_6$  as a function of temperature, as determined from the Tauc plot of the absorption spectra, is



Fig. 5 Optical bandgap of  $MA_2TeCl_6$  calculated as a function of temperature from the Tauc plot shown in Fig. S8 (ESI†).

shown in Fig. 5. As the temperature increases, a red shift is induced such that the bandgap decreases linearly within phases III and II (with a temperature dependence of  $-8.7(1) \times 10^{-4}$  eV K<sup>-1</sup>). The band gap shows an obvious drop after the transition to phase I. The data demonstrates that the cubic phase perovskite has a lower bandgap than trigonal structure, despite the larger volume of the cubic phase. Secondly, the first-order phase transition has a marked effect on the bandgap. The bandgap of MA<sub>2</sub>TeCl<sub>6</sub> decreases as a function of temperature to a larger extent than that seen in Cs<sub>2</sub>AgBiBr<sub>6</sub>,<sup>13</sup> which may due to differences in the crystal structure, but the temperature dependence has a similar value to that observed for other semiconductors.<sup>14</sup> This thermochromic effect may lead to new applications for MA<sub>2</sub>TeCl<sub>6</sub>.

#### Comparison of MA2TeCl6 and MA2SnCl6

The structure of MA<sub>2</sub>TeCl<sub>6</sub> is notably different from other MA<sub>2</sub>BCl<sub>6</sub> (B = Sn, Pt, Se, Pd and Pb) compounds, all of which exhibit rhombohedral structures at room temperature with space-group symmetry  $R\bar{3}m$ . These other compounds all undergo a structural phase transition to space group  $R\bar{3}$  at temperatures,  $T_{t}$ , between 100 and 200 K.<sup>10,15</sup> For example, the crystal structure of MA<sub>2</sub>SnCl<sub>6</sub>, shown in Fig. S9 (ESI†) has symmetry  $R\bar{3}m$  at room temperature, and transforms to  $R\bar{3}$  below 156 K.<sup>15</sup> Similar to the MA<sub>2</sub>TeCl<sub>6</sub> room temperature phase, the octahedral [SnCl<sub>6</sub>]<sup>2–</sup> anions occupy the corners of a trigonal unit cell, while MA<sup>+</sup> cations separate the layers of [SnCl<sub>6</sub>]<sup>2–</sup> anions. However, along the *c*-axis direction in MA<sub>2</sub>TeCl<sub>6</sub> planes of [TeCl<sub>6</sub>]<sup>2–</sup> anions lie directly above each other, whereas in MA<sub>2</sub>SnCl<sub>6</sub> the layer at  $z = \frac{1}{3}$  is displaced relative to the layer at z = 0 by 2/3 and 1/3 in *a* and *b*, respectively (see Fig. S9, ESI†).

In MA<sub>2</sub>SnCl<sub>6</sub>, it has been demonstrated that the MA<sup>+</sup> cation rotates freely about the C–N axis far below  $T_t$ , indicating that the hydrogen bonding between cations and anions is too weak to drive the phase transition.<sup>15,16</sup> Furthermore, NMR studies indicate that the motion of the MA<sup>+</sup> cations makes a smaller contribution to the phase transition compared to the rotation of the octahedra about the *c*-axis; also, NQR suggests that the phase transition may be driven by a soft rotary mode of the [SnCl<sub>6</sub>]<sup>2–</sup> anion.<sup>17</sup> In the high temperature  $R\bar{3}m$  phase,

the anions perform a hindered rotation, but their motion is restricted in the low-temperature  $R\bar{3}$  phase and they can only vibrate. Consequently, the loss of mirror symmetry and slight rotation of the octahedra are expected to drive the phase transition in MA<sub>2</sub>SnCl<sub>6</sub>. Intriguingly, X-rays experiments did not detect this rotation. The softening of an acoustic mode at the zone centre may be considered the cause of transition, as supported by the non-observation of a soft mode in Raman spectroscopy.<sup>15,16,18-20</sup> The situation concerning the anions in MA2TeCl6 is expected to be similar as there is also a loss of mirror symmetry in going from  $P\bar{3}m1$  in phase II to  $P\bar{3}1c$  in phase III. However, although the MA<sup>+</sup> cation has 3*m* symmetry in both the  $R\bar{3}m$  phase of MA<sub>2</sub>SnCl<sub>6</sub> and the  $P\bar{3}m1$  phase of MA<sub>2</sub>TeCl<sub>6</sub>, there is a distinct difference in the low temperature phase of each: the threefold symmetry of the MA<sup>+</sup> cation is retained in MA<sub>2</sub>SnCl<sub>6</sub>. In contrast, the symmetry of the cation is lost entirely in MA2TeCl6 due to the tilt ordering of the C-N axis. We, therefore, expect the phase transition in MA<sub>2</sub>TeCl<sub>6</sub> to be strongly affected by the order-disorder change of the cation.

Further insight into the nature of the structure and the phase transition can be gleaned from a comparison of the coefficients of thermal expansion in MA<sub>2</sub>SnCl<sub>6</sub> and MA<sub>2</sub>TeCl<sub>6</sub>. As illustrated in Fig. 6, the *c*-axis length of MA<sub>2</sub>SnCl<sub>6</sub> remains almost unchanged below 180 K; above this temperature, it increases with a thermal expansion coefficient of about  $0.6 \times 10^{-4} \text{ K}^{-1}$ , while *a* expands steadily with a coefficient of  $0.7 \times 10^{-4} \text{ K}^{-1}$  in both phases.<sup>21</sup> Similarly, based on the PXRD data, MA<sub>2</sub>TeCl<sub>6</sub> exhibits a steady increase in *a* with increasing temperature (Fig. 6) with a thermal expansion coefficient of  $0.9 \times 10^{-4} \text{ K}^{-1}$ , which is greater than for the tin compound. As in the case of MA<sub>2</sub>SnCl<sub>6</sub>, the *c* parameter of MA<sub>2</sub>TeCl<sub>6</sub> remains unchanged in phase III but then *decreases* in phase II with a negative thermal expansion coefficient of  $-0.4 \times 10^{-4} \text{ K}^{-1}$ .



**Fig. 6** Variation of the lattice parameters (a and c) and unit cell volume (V) of  $MA_2TeCl_6$  and  $MA_2SnCl_6$  as a function of temperature relative to 120 K highlighting the difference in the behaviour of c for the two compounds. The vertical lines shown in red and black mark the phase transition boundary  $T_t$  for the lower temperature phases of  $MA_2TeCl_6$  and  $MA_2SnCl_6$ , respectively.

These results demonstrate that temperature has a more dramatic effect on the *c* parameter compared with the *a* parameter.

The thermal expansion below  $T_t$  can be understood by considering the librational motion of the molecular ions. Previous work has discussed the effect of librational motion of both the MA<sup>+</sup> and [SnCl<sub>6</sub>]<sup>2-</sup> ions. With librational motion about the *c*-axis, then with increasing temperature, the molecular ions are forced apart in the *ab*-plane, which has the effect of increasing a, but leaving c essentially unchanged. The behaviour of the a and c lattice parameters in phase III of  $MA_2TeCl_6$  is seen to be similar. Vibrations of the ions would typically result in expansion along c, but this may be offset by an increased off-axis rotation of the MA<sup>+</sup> cations with increasing temperature, effectively shortening the projection of the C-N bond length on the c-axis: 1.468(7) Å at 90 K versus 1.407(8) Å at 295 K. This apparent shortening was also noted in the X-ray study on MA<sub>2</sub>SnCl<sub>6</sub> (1.522(25) Å at 120 K versus 1.487(30) Å at 298 K) and was also seen in the later neutron diffraction study on a deuterated sample.<sup>21,22</sup>

The distinct difference in the behaviour of the lattice parameter c above  $T_t$  in MA<sub>2</sub>TeCl<sub>6</sub> versus MA<sub>2</sub>SnCl<sub>6</sub> may be due to the different stacking sequences of the  $MA^+ - [Cl_3BCl_3]^{2-}$  -MA<sup>+</sup> layers. While the immediate coordination environment of the MA<sup>+</sup> cations is similar in both cases, *i.e.* they each have 12 Cl neighbours, these 12 Cl ions originate from different numbers of  $[BCl_6]^{2-}$  octahedral units: the MA<sup>+</sup> cations are coordinated to 4  $[SnCl_6]^{2-}$  anions in the tin compound, but to 6  $[TeCl_6]^{2-}$  anions in the tellurium one (Fig. 7). The larger size of a 6-coordinate cavity permits a greater degree of motion of the MA<sup>+</sup> cations. Further, in the low temperature phase transition of MA<sub>2</sub>TeCl<sub>6</sub>, the MA<sup>+</sup> cations tilt with respect to the *c*-axis (see Fig. 2) in contrast to MA<sub>2</sub>SnCl<sub>6</sub>, where the cations remain colinear with respect to the *c*-axis. On heating MA<sub>2</sub>TeCl<sub>6</sub> through the III-II phase transition, it is likely that the increase in temperature results in the MA<sup>+</sup> cations having an increased tilt (despite our model, see Fig. S7, ESI<sup>+</sup>): thus, the dynamicallydisordered C-N axis forms a cone with an increased cone angle about the threefold axis and, consequently, a decrease in the lattice parameter *c* is observed.



**Fig. 7** Schematic diagram of the MA<sup>+</sup> cations located in (a) a cavity coordinated by 6  $[TeCl_6]^{2-}$  anions in MA<sub>2</sub>TeCl<sub>6</sub> and (b) the cavity formed by only 4  $[SnCl_6]^{2-}$  anions in MA<sub>2</sub>SnCl<sub>6</sub>. For clarity, hydrogen atoms are not shown.

#### Paper

To understand the reasons for the constant *c*-axis below  $T_t$ , the magnitude of *c* can be broken down into the component distances between cation and anion layers, *i.e.*, the C–N bond length, the 'height' of the octahedron (Cl–Cl), the intermole-cular distance between chlorine ions from different octahedra (Cl···Cl), *etc.*, as shown schematically in Fig. S10 (ESI†). These intermolecular distances and bond lengths are calculated from the atomic coordinates derived from the variable temperature SXD results and are summarised in Fig. 8 and Table S2 (ESI†), where, for comparative purposes, changes in both *a* and *c* are included.

The chloride anion distances (as projected along *a*) both inside and between octahedra seem to contribute similarly to *a* in that both distances increase with increasing temperature, which is due to thermal expansion and the small rotation of the octahedra about their 3-fold axes. However, along the *c* direction, although the intermolecular distances, as projected on *c*, for C···C and Cl···Cl increase, the contraction of the projection of the C–N bond length along *c* and the reduction in octahedra 'height' (Cl–Cl) due to librational motion offset the lattice expansion; thus the length of *c*-axis stays approximately constant in phase III. The apparent shortening of C–N bond length has been discussed already.

For the same reason, the increased librational motion of the  $[\text{TeCl}_6]^{2-}$  octahedra results in an apparent contraction of the Te–Cl bond and a reduction in its size. 'N···Cl' represents the interleaved distance between MA<sup>+</sup> and octahedron layers, decreasing with increasing temperature. However, because of the overlap between layers, the decrease of distance will positively affect *c*, which is the reason for a positive slope in Fig. 8. The lack of SXD results makes it challenging to analyse MA<sub>2</sub>SnCl<sub>6</sub> similarly. However, based on the observation in MA<sub>2</sub>TeCl<sub>6</sub>, the constant *c* in MA<sub>2</sub>SnCl<sub>6</sub> in its low temperature



**Fig. 8** Percentage interatomic distance changes,  $d/d_{90 \text{ K}}$ , as projected in *a* and *c* for a variety of atoms in MA<sub>2</sub>TeCl<sub>6</sub> as a function of temperature relative to the structure at 90 K; the values are derived from SXD. Although there is a significant percentage increase in N···Cl with respect to *c*, in absolute terms, this contributes only a small to expansion in *c*. The error bar is smaller than the symbol; all error values can be found in Table S2 and S3 (ESI†).

phase is likely to be caused by the offset from increased thermal vibration of molecules. Further evidence is needed to prove the impact of corresponding interatomic distances.

#### The cation cavity

Geometric structure stability conditions in lattices are helpful to chemists as it allows them to quantify lattice-class stability and distortions easily. A classic example is the cation radius ratios and tolerance factors in perovskites.<sup>23</sup> A similar approach was proposed for cubic  $A_2BX_6$  halide perovskites by Brown in 1964, who used the radius ratio (*R*) of the cation size ( $r_A$ ) to the cavity size ( $r_{cavity}$ ), given in eqn (1), to describe the distortion of the structure in  $A_2BX_6$  crystals.<sup>24</sup>

$$R = \frac{r_{\rm A}}{r_{\rm cavity}} \tag{1}$$

It has been suggested that the distortion of high-symmetry crystal structures are driven by the packing of large anions with small cations.<sup>25</sup> Hence, the definition of the cavity is crucial, for example, in MA<sub>2</sub>TeCl<sub>6</sub> where the MA<sup>+</sup> cation is surrounded by 12 chloride ions as shown in Fig. 7(a), with the 12 chloride ions provided by 6 octahedra. Three of the 6 octahedra are face-coordinated, with the remaining three being vertex coordinated. As the temperature decreases, the rotation of the octahedra leads to a distortion of the MA<sup>+</sup> cavity, which drives the observed phase transition. However, the precise size of the cavity is difficult to define. Brown applied the halogen–halogen contacts as the size of the cavity eqn (2), where  $d_{X-X}$  is defined as the average value of interionic and intraionic halogen–halogen distances and  $r_X$  is the radius of halogen anion.

$$r_{\rm cavity} = d_{\rm X-X} - r_{\rm X} \tag{2}$$

However, the corresponding radius ratio is insufficient to distinguish the  $Fm\bar{3}m$  cubic structure from other structures. This might be because eqn (2) is most applicable to the K<sub>2</sub>PtCl<sub>6</sub> cubic structure type, where the distance between the spherical A<sup>+</sup> cation and halogen roughly equals the halogen–halogen contact distance. However, the distance between cation and halogen can vary significantly for a distorted structure. We can improve the estimate of the cavity size by calculating the *average* distance from the centre of mass of the cation to the coordinate halogen atom  $d_{A-x}$  as given in eqn (3). Yet this does not wholly capture the variations from a distorted structure either due to the wide variation in contact distances.

$$r_{\rm cavity} = d_{\rm A-X} - r_{\rm X} \tag{3}$$

Further improvement can be made to the calculation by only considering the smallest  $r_{\text{cavity}}$  length, which is less sensitive to the spread of the contact distances but still captures the magnitude of the distortion. This approach makes more logical sense because in hybrid compounds, the A site cavity contains an organic cation, which are generally not only much larger than the inorganic cations but also behave non-spherically at low temperatures. Using the shortest  $d_{A-X}$  for the radius ratio calculations, a larger variation in *R* occurs, as shown in Fig. 9,

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**Fig. 9** Radius ratio for  $A_2$ TeX<sub>6</sub> and  $A_2$ SnCl<sub>6</sub> compounds calculated from the shortest  $d_{A-X}$  values. The MA<sub>2</sub>TeCl<sub>6</sub> phases determined in this work are highlighted in red. *Fm*3m cubic structures are found in the range 0.89–1.17 as picked out in grey. (Square, diamond, and triangle symbols represent the *Fm*3m cubic structure, other cubic structures, and non-cubic structures respectively; solid symbols represent organic cation structures and hollow symbols represent inorganic ones).

in contrast to using the shortest  $d_{X-X}$  distance (see Table S4 and Fig. S11, ESI<sup>†</sup>). Our radius ratio range is larger than Brown's work (0.89 < R < 0.98), which did not consider hybrid compounds. Essentially, octahedral tilting in the hybrid compounds leads to lower symmetry structures with larger R values, which we capture in our modified equation allowing a more diverse range of materials to be compared.

We have extended and updated the radius ratio calculation of Brown to include new hexahalotellurate and hexachlorostannate (IV) complexes obtained from the Inorganic Crystal Structure Database (ICSD)<sup>26,27</sup> as shown in Fig. 9 and Table S4 (ESI<sup>†</sup>). The coordination number of the A-site cation by halide ions remains 12 across all compounds. The ionic radii of halogen and metal are taken from Shannon radii,<sup>28</sup> and the radii of the complete isocharge density sphere are used as effective radii of organic cations.<sup>29</sup> The grey region picked out in Fig. 9 covers the ratio range 0.89 < R < 1.17, and this ratio R shows an improved ability to distinguish between high-symmetry ( $Fm\bar{3}m$ ) cubic phases (depicted as squares in Fig. 9) and other structures compared (diamonds and triangles in Fig. 9) to using the average  $d_{X-X}$  distance (see Table S4, ESI<sup>†</sup>). The highest temperature cubic phase of MA<sub>2</sub>TeCl<sub>6</sub> (500 K) is clearly located inside the 0.89 < R < 1.17 range, whilst the other two phases fall above this range.

Most of the surveyed compounds follow a similar trend as  $MA_2TeCl_6$ : the cubic structure is stable in the lower R range while the lower symmetry structures tend to fall in the higher R regime. However, there are clear outliers in Fig. 9, e.g., Rb<sub>2</sub>TeI<sub>6</sub> (P4/mnc), (NH<sub>4</sub>)<sub>2</sub>TeI<sub>6</sub> (P2<sub>1</sub>/n), K<sub>2</sub>TeBr<sub>6</sub> (P4/mnc), K<sub>2</sub>TeCl<sub>6</sub> (I2/m) and K<sub>2</sub>SnCl<sub>6</sub> (P4/mnc). These compounds have relatively small cations, so the cavity distorts more heavily to accommodate them, which suggests that our definitions could be further refined to include, for example, the shortest distance between the octahedra. However, this is beyond the scope of this work. K2TeBr6 and K2SnCl6 transform upon cooling from their high temperature cubic  $(Fm\bar{3}m)$  phase to tetragonal (P4/mnc) and then to a monoclinic (P2<sub>1</sub>/n) phase, unlike the large cation compound MA2TeBr6, which follows a similar trend to our material. Again, the reason for this behaviour is likely related to the A-site ion radius. Although only the room temperature phases of (NH<sub>4</sub>)<sub>2</sub>TeI<sub>6</sub> and K<sub>2</sub>TeCl<sub>6</sub> are included in the ICSD, their highest temperature structures were reported as pseudo-cubic or cubic.<sup>30,31</sup>

For those compounds with multiple phases as a function of temperature, the radius ratio increases as the temperature decreases, reflecting the well understood relationship of crystal symmetry and temperature. One source of error in this method may come from the calculated cation radii, which we have assumed to be temperature independent but will likely increase at elevated temperatures. Thus, as the cavity size is known, the

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higher temperature phase's radius ratio would be underestimated (with a larger ionic radii), whereas the lower temperature phase's radius ratio should be over-estimated (with a smaller ionic radii).

# Conclusions

Single crystals of MA<sub>2</sub>TeCl<sub>6</sub> have been successfully grown by a slow cooling solvent method in HCl. VT-PXRD clearly identifies three phases from 120 K to 500 K. We corroborate that phase I (high temperature) is cubic structure ( $Fm\bar{3}m$ ), while single crystal XRD has determined the low temperature phase III structure as  $P\bar{3}1c$  at 200 K. Lattice distortions upon cooling were analysed, and the increased C–N axis length, which is related to the softening of molecule libration, is believed to be the main driver of the constant length of *c*-axis below the second phase transition temperature. We propose a new radius ratio (*R*) beyond Brown's work calculated using the shortest distance between A-site cation and halogen atom, which captures the distortion of MA cavity more effectively. The calculation of radius ratio was extended to hybrid A<sub>2</sub>BX<sub>6</sub> compounds, and we found that  $Fm\bar{3}m$  cubic structure generally locates in the range of 0.89 < *R* < 1.17.

## Conflicts of interest

There are no conflicts to declare.

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