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Strain-affected ferroelastic domain walls in RbMnFe charge-transfer materials undergoing collective Jahn–Teller distortion

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Many rubidium manganese hexacyanoferrate materials, with the general formula $\text{Rb}_x\text{Mn}[\text{Fe}(\text{CN})_6]_{(x+2)/3} \cdot z\text{H}_2\text{O}$, exhibit diverse charge-transfer-based functionalities due to the bistability between a high temperature $\text{Mn}^{\text{II}}(S = 5/2)\text{Fe}^{\text{III}}(S = 1/2)$ cubic phase and a low-temperature $\text{Mn}^{\text{III}}(S = 2)\text{Fe}^{\text{II}}(S = 0)$ tetragonal phase. The collective Jahn–Teller distortion on the Mn sites is responsible for the cubic-to-tetragonal ferroelastic phase transition, which is associated with the appearance of ferroelastic domains. In this study, we use X-ray diffraction to reveal the coexistence of 3 types of ferroelastic tetragonal domains and estimate the spatial extension of the strain around the domain walls, which represents about 30% of the volume of the crystal.

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Introduction

Many materials exhibit a ferroelastic phase transition,^{1–3} where the symmetry-breaking between different crystalline systems induces a spontaneous strain in the low symmetry phase. Ferroelastic materials show rich properties, with important applications in memory, multifunctional and novel controllable devices. In this respect, the cubic–tetragonal ferroelastic distortion was deeply investigated in many systems,^{4–9} with detailed analysis of the associated volume and tetragonal distortion strains. When such a symmetry breaking occurs, ferroelastic domains usually form,¹⁰ separated by a domain wall. The domain wall represents a twin interface where a multitude of properties can emerge.¹¹

There are many switchable magnetic and molecular-based materials, which exhibit ferroelastic phase transitions coupled to an electronic bistability, such as spin-crossover (SCO) or

charge-transfer (CT) in cyanide-bridge heterobimetallic materials.^{12–20} The effect of this coupling between both phenomena was rationalized within the frame of Collet's approach of the Landau theory of phase transition,²¹ which successfully explained the simultaneous or the sequential occurrence of electronic state switching and symmetry-breaking phenomena, the occurrence of hysteretic behaviour, and the emergence of various types of functions,^{22–33} including magnetoelectric (ME) effects and magnetic-field-induced spin state trapping (MIESST).^{34–40} The role of the ferroelastic domain walls and their dynamics was also scrutinized in Mn-based spin-crossover materials.^{12–15} It is now well understood that local properties can change around the deformed domain walls, giving rise for example to an electric polarization in an otherwise centrosymmetric system. The concept of functional domain walls opens new potential avenues for device designs.¹¹

Rubidium manganese hexacyanoferrate materials (Fig. 1a), with composition $\text{Rb}_x\text{Mn}[\text{Fe}(\text{CN})_6]_{(x+2)/3} \cdot z\text{H}_2\text{O}$, exhibit CT phase transitions, between $\text{Mn}^{\text{II}}(S = 5/2)\text{Fe}^{\text{III}}(S = 1/2)$ high temperature (HT) cubic phase and $\text{Mn}^{\text{III}}(S = 0)\text{Fe}^{\text{II}}(S = 0)$ low-temperature (LT) tetragonal phase, characterized by a wide thermal hysteresis.^{19,20,41,42} A Landau theory study revealed that this phase transition results from the coupling to the volume strain of the CT instability and the ferroelastic cubic–tetragonal phase transition associated with the collective Jahn–Teller distortion around the Mn^{III} sites.⁴³ Crystallographic studies investigated the important lattice changes between the HT cubic phase ($F\bar{4}3m$, $a_c \approx 10.5$ Å) and the LT phase described in the non-conventional $F\bar{4}2m$ tetragonal space group, $a_t \approx 10.0$ Å, $c_t \approx 10.5$ Å).^{43,44} The cubic \rightarrow tetragonal ferroelastic distortion

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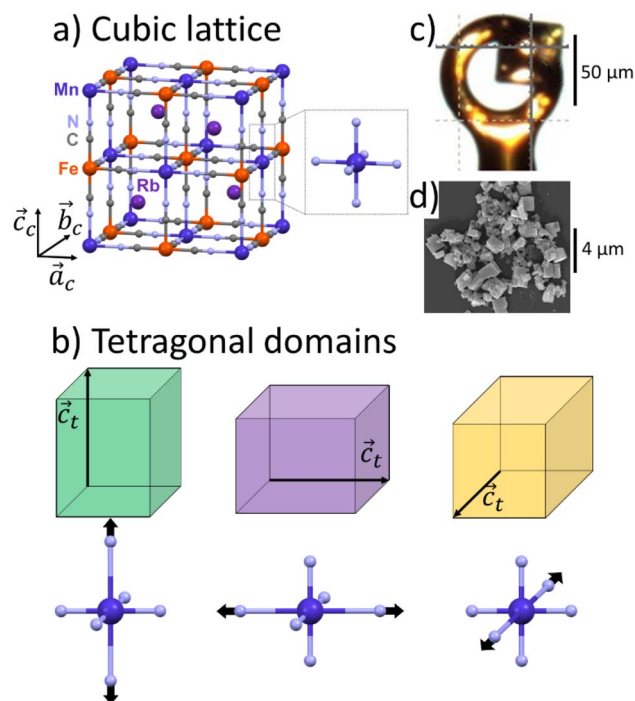


Fig. 1 (a) Structure of $\text{RbMn}[\text{Fe}(\text{CN})_6]$ in HT $\text{Mn}^{\text{II}}\text{Fe}^{\text{III}}$ cubic lattice ($a_c = b_c = c_c$), with MnN_6 sites of O_h symmetry. (b) Tetragonal domains in $\text{Mn}^{\text{III}}\text{Fe}^{\text{II}}$ LT phase ($a_t = b_t \neq c_t$), shown in green, purple and orange, with 3 possible orientations of the tetragonal axis c_t along a_c , b_c or c_c due to the Jahn–Teller distortion around the $\text{Mn}^{\text{III}}\text{N}_6$ sites with D_{4h} symmetry. (c) Image of the single crystal. (d) Scanning Electron Microscopy image of the powder.

is associated with the loss of the three-fold rotational symmetry of the cubic lattice. The structural instability occurs at the Γ point of the Brillouin zone and the Jahn–Teller symmetry-breaking order parameter belongs to the bidimensional E representation of the $F43m$ point group. Fig. 1b shows the cubic \rightarrow tetragonal distortion, associated with the Jahn–Teller deformation of the $\text{Mn}^{\text{III}}\text{N}_6$ core along the c_t axis in the tetragonal phase. This collective distortion can equally occur along the principal directions a_c , b_c or c_c of the initially cubic lattice.^{6,9,10,45} Therefore, 3 equivalent ferroelastic tetragonal domains can form in the LT phase.

Unexpected spectroscopic changes also occur in the LT phase. IR spectroscopy revealed a broad IR band ($\approx 50 \text{ cm}^{-1}$) around 2100 cm^{-1} , attributed to the splitting of the C–N stretching mode, while a sharp ($\approx 10 \text{ cm}^{-1}$) band is observed in the HT phase.^{46,47} Optical spectroscopy revealed a broad and intense band around 500 nm ,^{48–50} identified as a Mn-centred d–d transition, which is supposed to be weak in the LT D_{4h} symmetry. This absorption band plays a key role for driving photoinduced phase transition in these materials.^{17,44,47,48} The origin of these unusual spectroscopic features was however not clearly explained so far. Hereafter we use single crystal and powder X-ray diffraction (XRD) to study the ferroelastic domains in the LT phase and estimate the spatial extension of the lattice deformation around the domain walls.

Experimental

Sample preparation

Single crystals of $\text{RbMn}[\text{Fe}(\text{CN})_6]$ (hereafter referred to as RbMnFe) were synthesized by gradually diffusing 2 mL of an aqueous solution of $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$ (1.0 mol dm^{-3}) and 2 mL of an aqueous solution of $\text{K}_3[\text{Fe}(\text{CN})_6]$ (1.0 mol dm^{-3}) into 100 mL of an aqueous solution of RbCl (0.5 mol dm^{-3}) over one week at 40°C . RbMnFe is known to exhibit a wide thermal hysteresis between $\approx 230 \text{ K}$ and $\approx 300 \text{ K}$.^{51,52} $\text{Rb}_{0.94}\text{Co}_{0.06}\text{Mn}_{0.94}[\text{Fe}(\text{CN})_6]_{0.98} \cdot 0.2\text{H}_2\text{O}$, referred to hereafter as $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$, was synthesized according to the procedure reported in previous works,¹⁷ where additional characterization data can be found. Magnetic measurements on the powder indicate a broad bistability regime around room temperature ($T_\downarrow = 253 \text{ K}$, $T_\uparrow = 328 \text{ K}$), between the LT and HT phases. The powder sample was either cooled below T_\downarrow to generate the LT phase, or warmed above T_\uparrow to generate the HT phase. It was then investigated at room temperature (293 K), where both phases are stable. Two identical batches of powder sample were prepared and analysed separately using two different powder X-ray diffraction techniques.

Single crystal X-ray diffraction

We used single-crystal X-ray diffraction to study the RbMnFe sample. The data were collected in the HT phase at 300 K and in the LT phase at 100 K . Single-crystal X-ray diffraction was conducted using a Rigaku XtaLAB Synergy-R/DW with monochromatic Mo K_α radiation ($\lambda = 0.70930 \text{ \AA}$). We used paratone N oil to mount the single crystal ($50 \mu\text{m}$ large) on a Micro MountsTM holder (Fig. 1c). The data analysis and the reciprocal space reconstruction was performed with CrisAlisPro software of Rigaku.

Conventional powder X-ray diffraction of $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$

Conventional powder X-ray diffraction data of the $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$ sample were collected using a Rigaku Ultima-IV diffractometer with Cu K_α radiation ($\lambda = 1.5418 \text{ \AA}$, scanning range: $10\text{--}70^\circ$, scanning speed: 1° min^{-1} , sampling width: 0.02°). The powder sample consists of plate-shaped crystals, with an average size of $0.9 \pm 0.3 \mu\text{m}$ (Fig. 1d). The LT phase was prepared by cooling the powder in HT phase with liquid nitrogen. After the cooling, the LT phase was sustained at room temperature due to the existence of the thermal hysteresis over room temperature. The attribution of the Bragg peaks was based on previous analysis using Rietveld refinement.¹⁷

Streaming powder X-ray diffraction of $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$

In the second approach, XRD measurements were performed at the XPP beamline of the LCLS X-ray Free-Electron Laser (X-FEL), using monochromatic X-rays ($\lambda = 1.890 \text{ \AA}$, $\Delta E/E \sim 10^{-4}$).^{53,54} We used the streaming powder technique, which makes it possible to study ultra-fast and persistent photoinduced phase transitions.^{17,23} The $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$ crystals were dispersed in ethanol with a 1:90 crystal-solvent weight ratio, and the



solution was streamed through a 75 μm free-flowing jet to interact with the X-ray beam. The diffracted intensity was collected shot-to-shot using an epix10k2M detector⁵⁵ in the transmission geometry, at the repetition rate of the LCLS X-FEL (120 Hz). Each 2D image was azimuthally integrated using pyFAI⁵⁶ and normalized on the X-ray intensity. A typical number of 10 000 shots are subsequently averaged, over an acquisition time of about 100 s. The resulting pattern extends in the $[0.5 \text{ \AA}^{-1}; 3.6 \text{ \AA}^{-1}]$ Q -range. It contains a broad scattering contribution due to the solvent, which is removed by subtracting the scattering pattern of pure ethanol measured in the same conditions. The shape of the diffraction peak(s) is then analysed following the same procedure as for static powder XRD.

Results and discussion

Fig. 2 shows single crystal XRD data, characteristic of the cubic \rightarrow tetragonal phase transition of $\text{RbMn}[\text{Fe}(\text{CN})_6]$. The phase transition is associated with a symmetry breaking from HT cubic phase with $F\bar{4}3m$ space group ($a_{\text{HT}} \approx 10.5 \text{ \AA}$) to LT non-conventional $F\bar{4}2m$ space group ($a_{\text{LT}} \approx 10.0 \text{ \AA}$, $c_{\text{LT}} \approx 10.5 \text{ \AA}$). In the cubic phase (Fig. 2a), the diffracted intensity in the $(h0l)$ reciprocal plane, with cubic reciprocal lattice vectors $a_c^* = c_c^*$, corresponds to sharp Bragg peaks on the node where the h , k

and l indices have same parity (F cell). The diffracted patterns are the same in the $(0kl)$ and $(hk0)$ planes due to cubic symmetry. In the LT phase (Fig. 2b), the lengths of the lattice vectors differ due to the tetragonal distortion ($a_t^* = b_t^* \neq c_t^*$). We can see in Fig. 2b a splitting of the Bragg peaks due to the formation of the ferroelastic domains, for which the tetragonal distortion defining the c_t^* axis can equally occur along the HT axes a_c^* , b_c^* or c_c^* . The diffraction pattern in LT phase (Fig. 2b), corresponding to the $(h0l)$ plane of HT phase (Fig. 2a), is then characteristic of the superposition of the diffracted signal from the 3 tetragonal domains with different orientations of the c_t^* axis in the LT phase: a square (a_t^* , b_t^*) lattice (yellow, with c_t^* perpendicular to the figure), and two rectangular (b_t^* , c_t^*) and (a_t^* , c_t^*) lattices with c_t^* vertical (green) or horizontal (purple). Consequently, the $(h0h)$ Bragg peaks split in 3 peaks. We can also observe an anisotropic broadening of the Bragg peaks in the LT phase, with some diffuse scattering in-between the peaks of the different domains. This is particularly clear for the (606) Bragg peaks for example. This peak broadening is characteristic of non-uniform lattice strains.^{57,58}

Fig. 3 shows the XRD powder diffraction patterns along the scattering vector Q of $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$, measured at room temperature in the HT and LT phases with conventional powder XRD. The most important changes in the diffraction pattern, due to the ferroelastic distortion in the LT phase, correspond to the splitting along scattering vector Q of the Bragg peaks $(h00)$ and $(00h)$, and of the Bragg peaks $(hh0)$ and $(h0h)$, which are initially equivalent in the HT phase. For instance, Fig. 2b shows that the (200) and (002) Bragg peaks are no longer located on the same diffraction ring defined by $|Q|$ in the LT phase since $|Q(200)| \neq |Q(002)|$. Compared to previous studies,^{42,59} the splitting of the (002) and (200) peaks, indexed the LT $F\bar{4}2m$ space group, corresponds to the splitting of the (200) and (110) peaks indexed the LT $I\bar{4}m2$ space group. A similar splitting is observed for different Bragg peaks. In addition, due to lattice contraction, the peaks shift to higher Q in the LT phase. Fig. 3 shows asymmetric Bragg peaks in the LT phase, compared to HT, with diffuse scattering in-between the peaks. As mentioned above, this is due to the lattice strains, also observed in single crystal XRD with the diffuse scattering around the Bragg peaks.

While conventional Rietveld refinement can be used to characterize uniform strain in powder patterns, it is usually inadequate to describe highly-anisotropic strains, that give asymmetric Bragg peaks.⁶⁰ Instead, we used an approach similar to the one presented by Daniels *et al.* to characterize the domain structures from diffraction profiles in tetragonal ferroelastic ceramics.⁶¹ We analysed the shape of each peak or group of peaks separately by individual fitting, using a combination of split pseudo-Voigt peak profiles with a polynomial background:

$$S_{\text{fit}}(Q) = S_{\text{bkg}}(Q) + \sum_{i=1}^N \text{SPV}_i(Q)$$

where $S_{\text{bkg}}(Q)$ is a second-order polynomial, and N is the number of fitted peaks ($N \in [1; 3]$). $S_{\text{bkg}}(Q)$ was fitted outside of the Bragg peak(s), and then fixed during the fit of the peak(s). The split pseudo-Voigt profile $\text{SPV}(Q)$ is defined as an

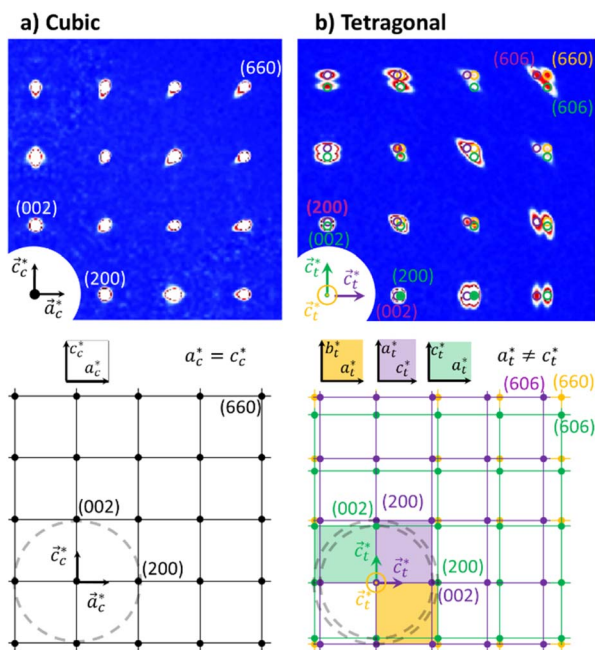


Fig. 2 Single crystal X-ray diffraction data of $\text{RbMn}[\text{Fe}(\text{CN})_6]$ (top) with a schematic drawing of the reciprocal lattice (bottom). (a) Diffracted intensity in the (a_c^*, c_c^*) plane with $k = 0$ for the cubic HT phase measured at 300 K, where the sharp Bragg peaks are located on the nodes of the cubic reciprocal lattice. (b) Diffracted intensity in the same reciprocal plane for the tetragonal LT phase (100 K), where the Bragg peaks are located on the nodes of the 3 superposing tetragonal reciprocal lattices of the 3 different domains (shown in orange, purple and green), characterized by different orientations of the tetragonal axis c_t^* . Diffuse scattering appears around the broader Bragg peaks due to lattice distortions around the domain walls.



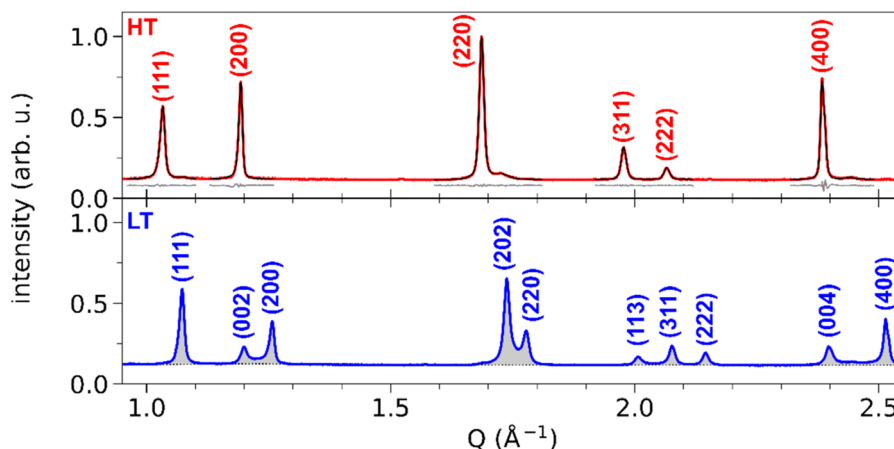


Fig. 3 Conventional X-ray powder diffraction patterns of $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$ in HT (red) and LT (blue) phases. (hkl) Miller indices correspond to the $F43m$ space group for the HT phase and to the $F42m$ space group for the LT phase. For the HT phase, the least-square fit of each peak is shown (black curve), together with the residuals (grey curve, shifted by +0.08). For the LT phase, only the fitted background is shown (black dotted line), highlighting the peak area (grey-shaded).

asymmetric peak shape, where the left (L) and right (R) sides around the peak centre Q_0 are described as two distinct pseudo-Voigt functions:

$$\text{SPV}(Q) = \begin{cases} A_L \cdot \text{PV}_L(Q), & \text{for } Q < Q_0 \\ A_R \cdot \text{PV}_R(Q), & \text{for } Q \geq Q_0 \end{cases}$$

where $\text{PV}_X(Q)$ ($X = L, R$) is the usual combination of a Lorentzian and a Gaussian profile with a common half width at half maximum σ_X and a mixing fraction $\alpha_X \in [0, 1]$:

$$\text{PV}_X(Q) = \alpha_X \frac{1}{\pi} \frac{\sigma_X}{(Q - Q_0)^2 + \sigma_X^2} + (1 - \alpha_X) \frac{\sqrt{\ln 2}}{\sigma_X \sqrt{\pi}} \exp\left(-\ln 2 \frac{(Q - Q_0)^2}{\sigma_X^2}\right)$$

Finally, ensuring continuity at the peak centre Q_0 imposes:

$$A_R = \frac{\text{PV}_L(Q_0)}{\text{PV}_R(Q_0)} A_L$$

As such, SPV profiles enable the analysis of asymmetry in the peak shape profiles, by extracting separately the half widths and the peak shape on both sides of the peaks. In the following, the reported values of the parameters correspond to the best fit of the data, and the associated uncertainties are the standard deviations of the least-square fit. In the case of (111), (220) and (400) Bragg peaks, we had to consider a weak additional signal due to an impurity, the passive $\text{Rb}_2\text{Mn}^{\text{II}}[\text{Fe}^{\text{II}}(\text{CN})_6] \cdot 3.5\text{H}_2\text{O}$ crystals ($Fm\bar{3}m$ $a = 10.185(6)$ Å), known to form during the synthesis.¹⁷ This impurity does not exhibit CT nor ferroelastic phase transition and we used the same contribution from this impurity in the analysis of the signal of HT and LT states at 293 K.

The top panel of Fig. 3 shows the least-square fit of all the Bragg peaks for the HT phase, where the SPV fit yields good results with quite symmetric profiles. For example, the fit of the (200) Bragg peak provides quite similar values for L/R sides: σ_L

$= 4.90(0.04) \times 10^{-3} \text{ Å}^{-1}$ vs. $\sigma_R = 3.88(0.03) \times 10^{-3} \text{ Å}^{-1}$ and $\alpha_L = 0.69(0.02)$ vs. $\alpha_R = 0.45(0.02)$. The same fitting procedure was applied to the LT phase. We focus first our attention on the (002) and (200) peaks. The top panel of Fig. 4a shows the fit with 2 SPVs, which is in good agreement with the XRD data. In this LT phase, the peak profiles are however strongly asymmetric, as characterized by the fitting parameters.

For the (002) Bragg peak, we obtained $\sigma_L = 6.7(0.3) \times 10^{-3} \text{ Å}^{-1}$ vs. $\sigma_R = 10.3(0.3) \times 10^{-3} \text{ Å}^{-1}$ and $\alpha_L = 0.59(0.07)$ vs. $\alpha_R = 1$ and for the (200) Bragg peak $\sigma_L = 7.51(0.08) \times 10^{-3} \text{ Å}^{-1}$ vs. $\sigma_R =$

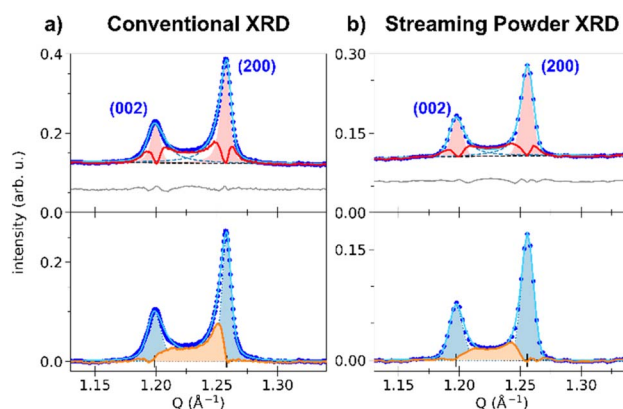


Fig. 4 (a) Zoom of the LT conventional powder diffraction pattern on the (002) and (200) peaks (blue dots). (Top panel): the least-square fit is shown (light blue curve), together with the two SPV components describing each peak (blue dashed lines), the polynomial background (black dashed line) and the residuals (grey line, shifted by +0.06). For both peaks, the HT peak model is shown as the red-shaded area, and the associated differential between the LT pattern and the HT peak model is displayed as the red curve (see text). (Bottom panel): background-subtracted pattern, together with the fit (light blue curve), the symmetrized peak shapes (blue-shaded areas and blue dotted lines), and the strain-affected intensity (orange area). (b) Same analysis for monochromatic X-FEL diffraction, in the streaming powder configuration.



$4.39(0.08) \times 10^{-3} \text{ \AA}^{-1}$ and $\alpha_L = 1$ vs. $\alpha_R = 0.68(0.03)$. Compared to the HT phase, both LT peaks are broader ($\sigma_{L,R}^{LT} > \sigma_{L,R}^{HT}$), and display a significant L/R asymmetry: the width of the peaks is larger in between the two Bragg peaks compared to their outer part ($\sigma_R^{(002)} > \sigma_L^{(002)}$ and $\sigma_L^{(200)} > \sigma_R^{(200)}$). The inner part of the two peaks also have a more pronounced Lorentzian character, as α_x reaches its maximum value (1). This fitting procedure thus maps the non-negligible intensity observed between the two peaks in the LT phase, in agreement with Fig. 2b. This asymmetry of the powder diffraction peaks is well understood from the single crystal data in Fig. 2b, as diffuse scattering is observed at $|Q|$ in between $|Q(200)|$ and $|Q(002)|$. The present fit shows that this intensity cannot be explained by a simple isotropic broadening of the peaks as observed in other ferroelastic materials.

We can highlight the asymmetry of the LT (002) and (200) peaks by comparing their profiles with the more symmetric one of the HT (200) peak. The LT peaks were fitted with the SPV profile of the HT phase, with widths $\sigma_{L,R}$ and fractions $\alpha_{L,R}$ fixed to the HT values and refined amplitudes and centres. The result is shown in Fig. 4a, where the red-shaded areas corresponds to fit of the peaks with the HT profiles. The differential between the experimental XRD pattern and the result from this fit is shown by the red curve. It is clear that the parameters of the sharp HT Bragg peaks poorly reproduce the LT data, as only the maximum of each peak is well described. The differential curve between the fit and the experimental data highlights the strong XRD signal between the (002) and (200) Bragg peaks, their asymmetry and broadening. Interestingly, a differential signal is also observed on the outer part of the two Braggs (*i.e.*, below 1.20 \AA^{-1} and above 1.26 \AA^{-1}).

Thus, the LT Bragg peaks are globally broadened compared to the HT phase and the signal observed in between (002) and (200) is characteristic of lattice strains in the LT phase, due to the formation of ferroelastic domains walls in the crystal.

Indeed, the strong asymmetry of the (200) and (002) Bragg peaks is due to the fact that tetragonal crystals have 90° {101}-type ferroelastic domain walls.^{62–65} There is an important strain due to the difference between the LT lattice parameters, with a_t

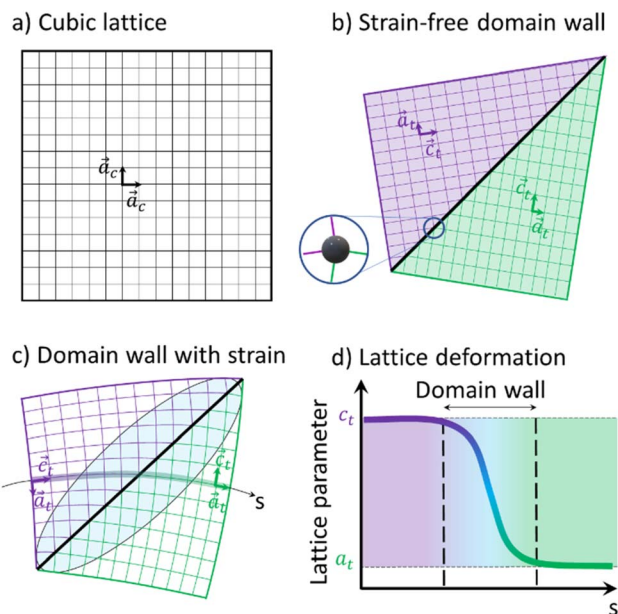


Fig. 6 Schematic representations of the cubic lattice (a) in the HT phase, and two tetragonal domains in the LT phase (b) with the tetragonal axis c_t horizontal (purple) or vertical (green) for a strain-free domain wall. The inset shows the distortion of the metallic sites on the wall. (c) Domain wall with strained region represented by the blue shaded elliptical area. (d) Evolution of the lattice parameter through the domain wall.

$\approx 10.0 \text{ \AA}$ and $c_t \approx 10.5 \text{ \AA}$. The different possible orientations of the {101}-type domain walls are shown in Fig. 5.

Fig. 6 shows a 2D representation of the formation of two ferroelastic domains. Compared to the regular HT cubic lattice (Fig. 6a), tetragonal domains can form in the LT phase, separated by a wall. The geometrically strain-free twinning angle only depends on the c_t/a_t ratio (Fig. 6b). However, due to constraints during the cubic-to-tetragonal phase transition and the development of the domains, this ideal twinning angle is not achieved. This is especially true in this polymeric crystal as the metallic sites, including the ones on the wall, are stabilized in D_{4h} symmetry (Fig. 1b). The sites on the wall for the strain-free twinning strongly deviates from D_{4h} symmetry as shown by the inset in Fig. 6b, which costs too much energy. Therefore, the lattice has to deform as shown in Fig. 6c, which results in strain fields in the vicinity of domain walls. The lattice constant changes gradually from c_t to a_t as we cross the domain wall along the path s shown in Fig. 6c, which represents the Mn–N–C–Fe bridges of the lattice. This change of lattice parameter is schematically shown in Fig. 6d. These strain fields around the wall, corresponding to a gradient of lattice parameter, result in the broadening of the Bragg peaks and diffuse scattering around some of them. The volume of material close to the domain wall that is affected by this strain field depends on the elastic modulus of the material. In the powder pattern, the signal between (002) and (200) is a direct signature of this strain field and the shape and amplitude of this inner signal reflects the strains extending around the domain walls.

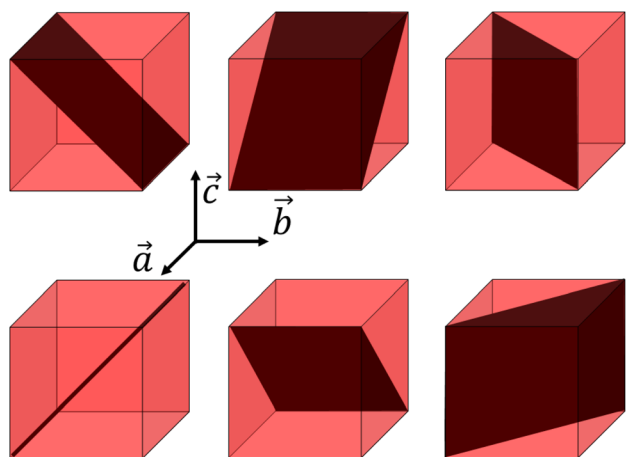


Fig. 5 Domain walls in a cubic lattice with tetragonal distortion, corresponding to {101} planes or symmetry-equivalent.



To further characterize the gradient of lattice parameters around the domain walls, we performed a strain analysis, with an approach similar to the one proposed by Daniels *et al.*⁶¹ In the hypothetical absence of gradient, both (002) and (200) peaks would display symmetric profiles. Thus, we artificially constructed symmetrized peak profiles $S_{\text{symm}}(Q)$, using the fit parameters of the outer parts of the two peaks, which is not affected by the gradient:

$$S_{\text{symm}}^{(002)}(Q) = A_L^{(002)} \cdot \text{PV}_L^{(002)}(Q)$$

$$S_{\text{symm}}^{(200)}(Q) = A_R^{(200)} \cdot \text{PV}_R^{(200)}(Q)$$

The X-ray diffraction signal affected by the strain of the gradient can then be calculated by subtracting the symmetrized peak profiles:

$$S_{\text{strain}}(Q) = S(Q) - [S_{\text{symm}}^{(002)}(Q) + S_{\text{symm}}^{(200)}(Q)]$$

The resulting signal is shown as the orange curve in Fig. 4a. As can be seen, the signal spreads over the whole Q range between the two peaks. Its shape is similar to the expected strain distribution around domain walls, as discussed in the literature for other ferroelastic phase transitions.⁶¹ A maximum is observed on the upper Q limit, that can be attributed to difference in the peak multiplicity between ($h00$) and ($00h$) Bragg peaks in the tetragonal lattice. Furthermore, this analysis enables to estimate the total volume of the crystals that is affected by the strain, by integrating $S_{\text{strain}}(Q)$ and comparing it to the total integrated signal. In the present case, our fit indicates that the strain-affected volume represents $\sim 30\%$ of the crystalline volume. This means that the formation of the ferroelastic domain walls induces important strains, which extend over about one third of each crystal.

Additionally, the width of the outer part of (002) and (200) peaks brings additional signature of the presence of ferroelastic domains. The outer parts of these peaks are indeed not affected by the gradient of the domain walls, and their broadening compared to the HT model cannot be related to lattice strain from c_t to a_t . Instead, the larger width is due to the small size of the ferroelastic domains forming during the HT-to-LT phase transition. Indeed, as domains are created in the LT phase, the coherence length of the diffraction signal decreases from the crystal size in the HT phase to the domain size in the LT phase. The formation of small domains increases the width of the peaks, according to the Scherrer equation:

$$\Delta\text{FWHM}_\theta = \frac{K \cdot \lambda}{L_C \cdot \cos \theta}$$

where ΔFWHM_θ is the full width at half maximum (FWHM) of the peak, corrected from the instrument response function (IRF), K is the shape factor of the order of unity, λ is the X-ray wavelength, L_C is the coherence length, and θ the diffraction angle of the Bragg peak.

We can therefore estimate the domain size from the broadening of (002) and (200) Bragg peaks relative to the HT phase, by considering that the width of HT (200) peak is mainly due to the IRF and that only the small size of the domains is responsible for the peak broadening in the LT phase. In this way, the increase of width of the LT (002) and (200) peaks, compared to HT (200) peak, gives $\Delta\text{FWHM}_\theta^{(002)}$ and $\Delta\text{FWHM}_\theta^{(200)}$. Given that we considered pseudo-Voigt peak profiles, ΔFWHM_θ was estimated as a mixture of Gaussian- and Lorentzian-corrected FWHMs:

$$\Delta\text{FWHM}_\theta^X = \alpha_X (\sigma_X^{\text{LT}} - \sigma_X^{\text{HT}}) + (1 - \alpha_X) \cdot \sqrt{(\sigma_X^{\text{LT}})^2 - (\sigma_X^{\text{HT}})^2}$$

In this way, we find that L_C is within the range of 200–600 nm. This means that the size of the ferroelastic domains is of the order of several hundreds of nanometers. Since the average size

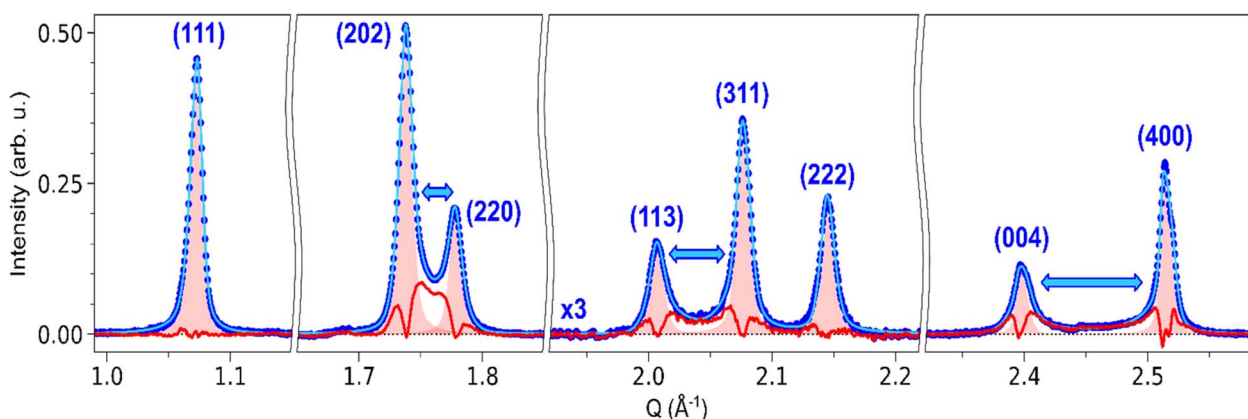


Fig. 7 Fit of (111), (202)/(220), (113)/(311)/(222) and (004)/(400) Bragg peaks in the LT phase: background-subtracted pattern (blue dots), least-square fit with SPV functions (light blue curve), HT peak models corresponding to each peak (red-shaded area), and the associated differential between the LT pattern and the HT peak model (red curve). For the (113)/(311)/(222), the signal is magnified by a factor of 3 for visibility. Blue arrows indicate the peaks that are split in the LT phase due to the phase transition. In the specific case of the fit of (004)/(400), SPV functions are replaced with split Pearson VII distributions, to account for the large splitting between the two peaks.



of the crystals in the powder is typically 900 nm,¹⁷ crystals are composed of a few to tens of domains with few 100 nm size.

These signatures of ferroelastic domains were further confirmed by streaming powder X-ray diffraction measurements on $\text{RbMn}_{0.94}\text{Co}_{0.06}\text{Fe}$ at the XPP beamline at LCLS (Fig. 4b). Similar trends are measured, with asymmetric peak shapes and peak broadening. Slight differences are visible in the shape of the strain-affected signal, whose maximum is less pronounced, and with a strain-affected volume of $\sim 20\%$. This difference, compared to Fig. 4a, may be due to the batch used or to the different thermal cycling of the samples during the experiments. Nonetheless, the overall signatures of the strain around the ferroelastic domain walls are clearly observed.

A similar influence of the strain around the domain walls is also expected for other Bragg peaks. Fig. 7 shows the fit of (111), (202)/(220), (113)/(311)/(222) and (004)/(400) peaks, and their comparison with the corresponding HT model. The profile of the (111) peak matches the HT model, as indicated by a negligible differential. This is also true for the (222) peak. On the other hand, (202)/(220), (113)/(311) and (004)/(400) all display a behaviour similar to (002)/(200): strong differential scattering intensity is observed between the peaks, as well as broadening on the outer part of the peaks. These three groups of peaks indeed correspond to peak splitting due to the ferroelastic phase transition, as (002)/(200). The gradient imposed by the domain walls during the transition thus affects the corresponding lattice constant. However, compared to the seminal case of (002)/(200) splitting, further strain analysis is difficult for these peaks, since their broadening can be affected on both left and right sides by the strain around the domain walls. For instance, (202) is affected by the $a_{\text{LT}} \rightarrow c_{\text{LT}}$ gradient on the left side, and by the $c_{\text{LT}} \rightarrow a_{\text{LT}}$ gradient on the right side. In the case of (111) and (222), the good fit with the HT model shows that the strain field is weaker along these crystalline directions, as expected for {101}-type ferroelastic domain walls.

Conclusions

This X-ray diffraction study of the ferroelastic phase transition from the cubic $\text{Mn}^{\text{II}}\text{Fe}^{\text{III}}$ HT to the tetragonal $\text{Mn}^{\text{III}}\text{Fe}^{\text{II}}$ LT phase of RbMnFe materials reveals the formation of ferroelastic domains walls during the cooperative Jahn–Teller distortion around the Mn sites. Three types of ferroelastic tetragonal domains, with c_t axis pointing in all three possible directions, coexist in the LT phase, as revealed by single-crystal X-ray diffraction. Our analysis shows that the spatial extension of the strain around the domain walls represents $\approx 30\%$ of the volume. Because of these distortions of the polymeric Mn–N–C–Fe network, the local symmetry of the ligand field of the metal ions around the wall deviates from D_{4h} , as shown in Fig. 6c. In this figure, a single domain wall is represented. However, as explained in Fig. 5, different domain walls with different orientations can form. Along the path s in Fig. 6c, the c_t axis points up, while the a_t axis points down. However, around a symmetry-equivalent wall the c_t axis will point down, while the a_t axis will point up. In average, c_t and a_t axes are horizontal. The angular distribution of these axes broadens transversally

the Bragg peaks (Fig. 2b). Our analysis, evidences that a large fraction of the sample ($\approx 30\%$ of the volume) is affected by the lattice strain around the domain wall. This translates through the appearance of scattering signal in-between Bragg peaks, such as (002)/(200) for example. Such a diffuse signal was observed in previous experiments, but strain analysis was never performed so far. Our strain analysis makes it possible to understand the unexpected spectroscopic changes occurring in the LT phase. On the one hand, a broad IR band ($\approx 50 \text{ cm}^{-1}$) of the C–N stretching mode is observed in the LT phase, compared to the sharp IR band ($\approx 10 \text{ cm}^{-1}$) in the HT phase.⁴⁷ This broadening results from both a splitting of this IR band due to the symmetry-breaking in the tetragonal phase and a broadening of these bands due to the strain field around the wall. On the other hand, a broad and intense optical band, which has a strong Mn-centred d–d character, appears around 500 nm in the LT phase,⁴⁸ which is not expected for Mn sites in D_{4h} symmetry. Our results show that about 30% of the Mn sites are in a strong strain field. Consequently, the associated lattice strain, bending the Mn–N–C–Fe bridges, activates this optical transition. The strain field around domain walls plays therefore an important role for the optical control of the magnetic properties of RbMnFe materials, as the Mn-centred d–d transition is also used to drive photomagnetic properties.^{17,44,47,48} This highlights the important consequences of the formation of a domain wall. The present strain analysis is thus complementary to the structural information obtained though, *e.g.*, Rietveld refinements. Our next goal is to extend this study to the time domain, to understand the evolution of lattice strains during ultrafast and persistent photoinduced phase transition.¹⁷ This family of multifunctional materials¹⁹ has good chemical stability and durability, and does not contain any banned elements, making it a promising candidate for practical applications, such as barocaloric effects,⁶⁶ photomagnetism,^{20,41,67} or Ferroelectricity.⁶⁸ It was also shown that the application of an electric field above a threshold value leads to a transition from the high- to the low-temperature phase⁶⁹ and consequently changes in their magnetic, optical, and electronic properties. This can open the way for novel electro-optical devices, and here again the domain walls may also play an important role in the process.

Data availability

The powder X-ray diffraction data and their fit are available at Zenodo <https://doi.org/10.5281/zenodo.13366596>.

Author contributions

Marius Hervé: methodology, data curation, formal analysis, writing – original draft; Shintaro Akagi: resources, data curation, investigation, formal analysis; Laurent Guérin, formal analysis; Leland Gee: data curation; Ryan Ribson: data curation; Matthieu Chollet: data curation; Marco Cammarata: data curation, funding acquisition; Shuntaro Nagashima: data curation, investigation, formal analysis; Shin-ichi Ohkoshi: funding acquisition, supervision; Hiroko Tokoro: resource, data



curation, funding acquisition, supervision; Eric Collet: funding acquisition, data curation, investigation, conceptualization, supervision, writing – original draft, review & editing. All authors critically reviewed and revised the manuscript draft and approved the final version for submission.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 E. K. Salje, *Phase Transitions in Ferroelastic and Co-elastic Crystals*, Cambridge University Press, Cambridge, 1991.
- 2 E. K. H. Salje, *Annu. Rev. Mater. Res.*, 2012, **42**, 265–283.
- 3 M. A. Carpenter, E. K. H. Salje and A. Graeme-Barber, *Eur. J. Mineral.*, 1998, **10**, 621–691.
- 4 S. C. Tarantino, M. Giannini, M. A. Carpenter and M. Zema, *Iucrj*, 2016, **3**, 354–366.
- 5 M. A. Carpenter and E. K. Salje, *Eur. J. Mineral.*, 1998, **10**, 693.
- 6 V. Janovec, V. Dvořák and J. Petzelt, *Czech. J. Phys. B*, 1975, **25**, 1362–1396.
- 7 T. Yamanaka, T. Mine, S. Asogawa and Y. Nakamoto, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2009, **80**, 134120.
- 8 D. C. Palmer, E. K. H. Salje and W. W. Schmahl, *Phys. Chem. Miner.*, 1989, **16**, 714–719.
- 9 J. D. Axe and Y. Yamada, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1981, **24**, 2567–2569.
- 10 V. Janovec and J. Privratska, *International Tables for Crystallography*, 2006, D, 449–505.
- 11 E. K. H. Salje, *J. Appl. Phys.*, 2020, **128**, 164104.
- 12 V. B. Jakobsen, E. Trzop, E. Dobbelaar, L. C. Gavin, S. Chikara, X. Ding, M. Lee, K. Esien, H. Müller-Bunz, S. Felton, E. Collet, M. A. Carpenter, V. S. Zapf and G. G. Morgan, *J. Am. Chem. Soc.*, 2022, **144**, 195–211.
- 13 E. Dobbelaar, V. B. Jakobsen, E. Trzop, M. Lee, S. Chikara, X. Ding, H. Müller-Bunz, K. Esien, S. Felton, M. A. Carpenter, E. Collet, G. G. Morgan and V. S. Zapf, *Angew. Chem., Int. Ed.*, 2022, **61**, e202114021.
- 14 V. B. Jakobsen, S. Chikara, J.-X. Yu, E. Dobbelaar, C. T. Kelly, X. Ding, F. Weickert, E. Trzop, E. Collet, H.-P. Cheng, G. G. Morgan and V. S. Zapf, *Inorg. Chem.*, 2021, **60**, 6167–6175.
- 15 V. B. Jakobsen, E. Trzop, L. C. Gavin, E. Dobbelaar, S. Chikara, X. Ding, K. Esien, H. Müller-Bunz, S. Felton, V. S. Zapf, E. Collet, M. A. Carpenter and G. G. Morgan, *Angew. Chem., Int. Ed.*, 2020, **59**, 13305–13312.
- 16 S. Chikara, J. Gu, X. G. Zhang, H.-P. Cheng, N. Smythe, J. Singleton, B. Scott, E. Krenkel, J. Eckert and V. S. Zapf, *Nat. Commun.*, 2019, **10**, 4043.
- 17 M. Hervé, G. Privault, E. Trzop, S. Akagi, Y. Watier, S. Zerdane, I. Chaban, R. G. Torres Ramírez, C. Mariette, A. Volte, M. Cammarata, M. Levantino, H. Tokoro, S. Ohkoshi and E. Collet, *Nat. Commun.*, 2024, **15**, 267.
- 18 S. Ohkoshi, H. Tokoro and E. Collet, *Chimia Rep.*, 2019, **22**, 498–507.
- 19 H. Tokoro and S. Ohkoshi, *Bull. Chem. Soc. Jpn.*, 2015, **88**, 227–239.
- 20 S. Ohkoshi and H. Tokoro, *Acc. Chem. Res.*, 2012, **45**, 1749–1758.
- 21 E. Collet and G. Azzolina, *Phys. Rev. Mater.*, 2021, **5**, 044401.
- 22 E. Collet, G. Azzolina, J. Jeftić and M.-H. Lemée-Cailleau, *Adv. Phys.: X*, 2023, **8**, 2161936.
- 23 M. Reinhard, K. Kunnus, K. Ledbetter, E. Biasin, D. B. Zederkof, R. Alonso-Mori, T. B. van Driel, S. Nelson, M. Kozina, O. J. Borkiewicz, M. Lorenc, M. Cammarata, E. Collet, D. Sokaras, A. A. Cordones and K. J. Gaffney, *ACS Nano*, 2024, **18**, 15468–15476.
- 24 F. J. Valverde-Muñoz, R. G. Torres Ramírez, A. Ulhe, E. Trzop, M. Dutta, C. Das, P. Chakraborty and E. Collet, *CrystEngComm*, 2023, **25**, 3588–3597.
- 25 R. G. Torres Ramírez, E. Trzop and E. Collet, *Dalton Trans.*, 2024, **53**, 10159–10167.
- 26 K. W. Törnroos, M. Hostettler, D. Chernyshov, B. Vangdal and H.-B. Bürgi, *Chem.–Eur. J.*, 2006, **12**, 6207–6215.
- 27 S. Lakhroufi, P. Guionneau, M. H. Lemée-Cailleau, P. Rosa and J. F. Létard, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2010, **82**, 132104.
- 28 V. A. Money, J. Elhaik, I. Radosavljevic Evans, M. A. Halcrow and J. A. K. Howard, *Dalton Trans.*, 2004, 65–69, DOI: [10.1039/B311262B](https://doi.org/10.1039/B311262B).
- 29 G. A. Craig, J. Sánchez Costa, O. Roubeau, S. J. Teat and G. Aromí, *Chem.–Eur. J.*, 2011, **17**, 3120–3127.
- 30 A. Djemel, O. Stefanczyk, M. Marchivie, E. Trzop, E. Collet, C. Desplanches, R. Delimi and G. Chastanet, *Chem.–Eur. J.*, 2018, **24**, 14760–14767.
- 31 S. Floquet, N. Guillou, P. Négrier, E. Rivière and M.-L. Boillot, *New J. Chem.*, 2006, **30**, 1621–1627.
- 32 B. Weber, E. S. Kaps, J. Obel, K. Achterhold and F. G. Parak, *Inorg. Chem.*, 2008, **47**, 10779–10787.
- 33 Y.-Y. Zhu, C.-W. Liu, J. Yin, Z.-S. Meng, Q. Yang, J. Wang, T. Liu and S. Gao, *Dalton Trans.*, 2015, **44**, 20906–20912.



- 34 Y. Otsuki, S. Kimura, S. Awaji and M. Nakano, *AIP Adv.*, 2019, **9**, 085219.
- 35 M. Nakano, G.-e. Matsubayashi and T. Matsuo, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2002, **66**, 212412.
- 36 J. Olguín, *Coord. Chem. Rev.*, 2020, **407**, 213148.
- 37 N. A. Spaldin and R. Ramesh, *Nat. Mater.*, 2019, **18**, 203–212.
- 38 S.-W. Cheong and M. Mostovoy, *Nat. Mater.*, 2007, **6**, 13–20.
- 39 N. A. Spaldin and M. Fiebig, *Science*, 2005, **309**, 391–392.
- 40 X. Zhang, W. H. Xu, W. Zheng, S. Q. Su, Y. B. Huang, Q. Shui, T. Ji, M. Uematsu, Q. Chen, M. Tokunaga, K. Gao, A. Okazawa, S. Kanegawa, S. Q. Wu and O. Sato, *J. Am. Chem. Soc.*, 2023, **145**, 15647–15651.
- 41 H. Tokoro, T. Matsuda, T. Nuida, Y. Moritomo, K. Ohoyama, E. D. L. Dangui, K. Boukheddaden and S. Ohkoshi, *Chem. Mater.*, 2008, **20**, 423–428.
- 42 H. Tokoro, K. Nakagawa, K. Imoto, F. Hakoe and S. Ohkoshi, *Chem. Mater.*, 2012, **24**, 1324–1330.
- 43 G. Azzolina, R. Bertoni, C. Ecolivet, H. Tokoro, S. Ohkoshi and E. Collet, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2020, **102**, 134104.
- 44 G. Azzolina, R. Bertoni, C. Mariette, M. Cammarata, E. Trzop, C. Ecolivet, M. Sander, M. Levantino, H. Tokoro, K. Imoto, M. Yoshikiyo, S. Ohkoshi and E. Collet, *J. Mater. Chem. C*, 2021, **9**, 6773–6780.
- 45 P. Toledano, M. M. Fejer and B. A. Auld, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1983, **27**, 5717–5746.
- 46 H. Tokoro, A. Namai, M. Yoshikiyo, R. Fujiwara, K. Chiba and S. Ohkoshi, *Sci. Rep.*, 2018, **8**, 63.
- 47 G. Privault, M. Hervé, N. Godin, R. Bertoni, S. Akagi, J. Kubicki, H. Tokoro, S. Ohkoshi, M. Lorenc and E. Collet, *Angew. Chem., Int. Ed.*, 2024, e202408284.
- 48 G. Azzolina, H. Tokoro, K. Imoto, M. Yoshikiyo, S. Ohkoshi and E. Collet, *Angew. Chem., Int. Ed.*, 2021, **60**, 23267–23273.
- 49 S. Ohkoshi, T. Nuida, T. Matsuda, H. Tokoro and K. Hashimoto, *J. Mater. Chem.*, 2005, **15**, 3291.
- 50 E. D. Loutete-Dangui, E. Codjovi, H. Tokoro, P. R. Dahoo, S. Ohkoshi and K. Boukheddaden, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **78**, 014303.
- 51 S. Ohkoshi, H. Tokoro, M. Utsunomiya, M. Mizuno, M. Abe and K. Hashimoto, *J. Phys. Chem. B*, 2002, **106**, 2423–2425.
- 52 H. Tokoro, S. Ohkoshi, T. Matsuda and K. Hashimoto, *Inorg. Chem.*, 2004, **43**, 5231–5236.
- 53 M. Chollet, R. Alonso-Mori, M. Cammarata, D. Damiani, J. Defever, J. T. Delor, Y. Feng, J. M. Glowina, J. B. Langton, S. Nelson, K. Ramsey, A. Robert, M. Sikorski, S. Song, D. Stefanescu, V. Srinivasan, D. Zhu, H. T. Lemke and D. M. Fritz, *J. Synchrotron Radiat.*, 2015, **22**, 503–507.
- 54 C. Antolini, V. Sosa Alfaro, M. Reinhard, G. Chatterjee, R. Ribson, D. Sokaras, L. Gee, T. Sato, P. L. Kramer, S. L. Raj, B. Hayes, P. Schleissner, A. T. Garcia-Esparza, J. Lim, J. T. Babicz, A. H. Follmer, S. Nelson, M. Chollet, R. Alonso-Mori and T. B. van Driel, *Molecules*, 2024, **29**, 2323.
- 55 T. B. van Driel, S. Nelson, R. Armenta, G. Blaj, S. Boo, S. Boutet, D. Doering, A. Dragone, P. Hart, G. Haller, C. Kenney, M. Kwaitowski, L. Manger, M. McKelvey, K. Nakahara, M. Oriunno, T. Sato and M. Weaver, *J. Synchrotron Radiat.*, 2020, **27**, 608–615.
- 56 G. Ashiotis, A. Deschildre, Z. Nawaz, J. P. Wright, D. Karkoulis, F. E. Picca and J. Kieffer, *J. Appl. Crystallogr.*, 2015, **48**, 510–519.
- 57 B. D. Cullity, *Elements of X-Ray Diffraction*, Addison-Wesley, Reading, MA, USA, 2nd edn, 1978.
- 58 H. Boysen, *Z. Kristallogr.*, 2005, **220**, 726–734.
- 59 G. Azzolina, E. Collet, C. Mariette, M. Cammarata, E. Trzop, M. Sander, M. Levantino, K. Nakagawa, H. Tokoro, S. Ohkoshi and R. Bertoni, *Eur. J. Inorg. Chem.*, 2019, **2019**, 3142–3147.
- 60 G. Tutuncu, M. Motahari, M. R. Daymond and E. Ustundag, *Acta Mater.*, 2012, **60**, 1494–1502.
- 61 J. E. Daniels, J. L. Jones and T. R. Finlayson, *J. Phys. D: Appl. Phys.*, 2006, **39**, 5294.
- 62 J. Sapriel, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1975, **12**, 5128–5140.
- 63 S. H. Curnoe and A. E. Jacobs, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2000, **62**, R11925–R11928.
- 64 T. J. Parker and J. C. Burfoot, *J. Phys. D: Appl. Phys.*, 1969, **2**, 1168.
- 65 G. Arlt and P. Sasko, *J. Appl. Phys.*, 1980, **51**, 4956–4960.
- 66 S. Ohkoshi, K. Nakagawa, M. Yoshikiyo, A. Namai, K. Imoto, Y. Nagane, F. Jia, O. Stefanczyk, H. Tokoro, J. Wang, T. Sugahara, K. Chiba, K. Motodohi, K. Isogai, K. Nishioka, T. Momiki and R. Hatano, *Nat. Commun.*, 2023, **14**, 8466.
- 67 H. Tokoro, T. Matsuda, K. Hashimoto and S. Ohkoshi, *J. Appl. Phys.*, 2005, **97**, 10M508.
- 68 S. Ohkoshi, H. Tokoro, T. Matsuda, H. Takahashi, H. Irie and K. Hashimoto, *Angew. Chem., Int. Ed.*, 2007, **46**, 3238–3241.
- 69 T. Mahfoud, G. Molnár, S. Bonhommeau, S. Cobo, L. Salmon, P. Demont, H. Tokoro, S.-I. Ohkoshi, K. Boukheddaden and A. Bousseksou, *J. Am. Chem. Soc.*, 2009, **131**, 15049–15054.

