



Deposited via The University of Sheffield.

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/id/eprint/166168/>

Version: Accepted Version

Article:

Mostaed, A., Walkley, B., Hatnean, M.C. et al. (2021) Characterizing oxygen atoms in perovskite and pyrochlore oxides using ADF-STEM at a resolution of a few tens of picometers. *Acta Materialia*, 208. 116717. ISSN: 1359-6454

<https://doi.org/10.1016/j.actamat.2021.116717>

Article available under the terms of the CC-BY-NC-ND licence
(<https://creativecommons.org/licenses/by-nc-nd/4.0/>).

Reuse

This article is distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs (CC BY-NC-ND) licence. This licence only allows you to download this work and share it with others as long as you credit the authors, but you can't change the article in any way or use it commercially. More information and the full terms of the licence here: <https://creativecommons.org/licenses/>

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.

Characterizing oxygen atoms in perovskite and pyrochlore oxides using ADF-STEM at a resolution of a few tens of picometers

Ali Mostaed^{1*}, Brant Walkley^{2,3}, Monica Ciomaga Hatnean⁴, Geetha Balakrishnan⁴, Martin R. Lees⁴,
Richard Beanland⁴, Derek C. Sinclair² and Ian M. Reaney²

¹ *Department of Materials, University of Oxford, Parks Road, Oxford OX1 3PH, UK*

² *Department of Materials Science and Engineering, University of Sheffield, Sheffield S1 3JD, UK*

³ *Department of Chemical and Biological Engineering, University of Sheffield, Sheffield S1 3JD, UK*

⁴ *Department of Physics, University of Warwick, Gibbet Hill Road, Coventry CV4 7AL, UK*

Abstract

We present an aberration corrected scanning transmission electron microscopy (ac-STEM) analysis of perovskite (LaFeO_3) and pyrochlore ($\text{Yb}_2\text{Ti}_2\text{O}_7$ and $\text{Pr}_2\text{Zr}_2\text{O}_7$) oxides and demonstrate that both the shape and contrast of visible atomic columns in annular dark-field (ADF) images are sensitive to the presence of nearby atoms of low atomic number (e.g. oxygen). We show that point defects (e.g. oxygen vacancies), which are invisible – or difficult to observe due to limited sensitivity – in x-ray and neutron diffraction measurements, are the origin of the complex magnetic ground state of pyrochlore oxides. In addition, we present a method by which light atoms can be resolved in the quantitative ADF-STEM images. Using this method, we resolved oxygen atoms in perovskite and pyrochlore oxides and propose this method to be suitable for other materials containing both light and heavy elements.

Keywords: *STEM; Vacancies; Point defects; Perovskite; Pyrochlore*

* Corresponding author: ali.mostaed@materials.ox.ac.uk

1. Introduction

Ceramic oxides with perovskite (ABO_3) or pyrochlore ($A_2B_2O_7$) structure, where A and B are cations with very different sizes, exhibit interesting physical and electrical phenomena including magnetic [1, 2], ferroelectric [3-6], dielectric [7-9], pyroelectric [4, 10], piezoelectric [4, 11] and superconducting [12, 13] properties. The functionality of these oxides is often dependent on defects, in particular oxygen displacements or oxygen vacancies [14-19]. For example, local structural defects (e.g. vacancies) in the oxygen lattice can induce small changes in the Pr^{3+} ion network, and thus affect the magnetic properties of praseodymium zirconate ($Pr_2Zr_2O_7$). It has been demonstrated in x-ray diffraction and neutron scattering measurements that off-centring of Pr^{3+} ions from the ideal crystallographic position, which can be due to presence of oxygen vacancies, is at the origin of the complex magnetic ground state of single-crystal $Pr_2Zr_2O_7$ [20-23]. However, oxygen vacancies themselves remain undetected [20-23]. Although x-ray and neutron diffraction techniques have been widely used to study the presence and position of oxygen in oxide materials [24-28], these techniques only provide information regarding the average structure, as the information is collected from a large volume of a specimen. Transmission electron microscopy (TEM), on the other hand, offers the opportunity to probe their local structure. For instance, aberration corrected transmission electron microscopy (ac-TEM) can be exploited to obtain information about the structure of materials with sub-angstrom resolution. Nevertheless, detecting oxygen, which is a light element, in the presence of heavier elements in oxides remains challenging. Urban et al. [29] showed that it is feasible to make light atoms like oxygen visible in TEM images under negative spherical aberration conditions. However, exploiting this technique is not straight forward since the imaging area should be ultra-thin (about a few nonometers) and a comprehensive knowledge about the microscope's aberrations is required.

In scanning transmission electron microscopy (STEM), bright-field (BF) and annular bright-field (ABF) imaging techniques have been widely used to reveal the position of oxygen in oxide materials [30-35]; however, as intensities in ABF images are not related in a straightforward way to atomic number, they are not reliable to study the oxygen occupancy. On the other hand, annular dark-field (ADF) imaging provides quantitative information regarding the atomic structure of materials as the contrast of visible atom columns in ADF images is proportional to the atomic number (Z) of the atoms in those columns, i.e. ADF is a Z -contrast imaging technique [35-38]. The contribution of each single atom (Z^n ; where n is a constant with a value between 1.5 to 2 [39, 40]) in the specimen to the ADF image is

proportional to the atomic number of that atom as well as the electron probe intensity at that atom, $J(r)$, which can be simply written as a differential contribution (dI) as follows:

$$dI(\mathbf{r}_p, z) = Z^n J(\mathbf{r}_a - \mathbf{r}_p, z), \quad (1)$$

where, \mathbf{r}_p is the probe position, \mathbf{r}_a and z are the lateral position and depth of the atom [41]. When the probe is at or very close to an atom column containing heavy atoms, there is sufficient attractive Coulombic force from the heavy atoms to trap or focus the majority of the electron beam along the atom column and as a consequence avoid the probe spreading out from the column [42]. Thus, the contribution of heavy atom columns to the ADF intensity is significant, as not only do they contain heavy atoms (high Z^n in Eq. 1) but also the majority of the probe intensity distribution is on-column (high $J(r)$ in Eq. 1). On the other hand, as the attractive Coulombic potentials for the atom columns containing light atoms are too weak to trap the electron beam, probe de-channelling can occur when the probe is at or very close to these columns and, therefore, a fraction of the probe intensity can spread on their nearby atom columns, i.e. low $J(r)$ [41, 42]. Thus, columns with light atoms contribute far less than columns with heavy atoms to the intensity of ADF images as both their Z^n and $J(r)$ values are significantly smaller for columns with the lighter atoms. This is the reason why light elements, especially in the presence of relatively heavier elements, e.g. oxygen atoms in oxides, are invisible in ADF images. In contrast to the aforementioned reasons for invisible oxygen atoms in ADF images, Mostaed et al. [15] showed that the detailed intensity distribution around the visible cation columns in ADF-STEM images obtained from the pyrochlore $\text{Yb}_2\text{Ti}_2\text{O}_7$ is sensitive to the presence of their nearby invisible light atoms. In the current work, we explain how light atoms such as oxygen can contribute sufficiently to the intensity of ADF images to be detected.

Here we use ac-STEM **in conjunction with electron energy loss spectroscopy (EELS)** to study the atomic structure of pyrochlore ($\text{Yb}_2\text{Ti}_2\text{O}_7$ and $\text{Pr}_2\text{Zr}_2\text{O}_7$) and perovskite (LaFeO_3) oxides and demonstrate that the intensity as well as shape of visible cation columns in ADF-STEM images are sensitive to the presence of their nearby invisible oxygen atoms. We propose a novel method by which the invisible oxygen columns can be resolved in the ADF-STEM images. Using this method we, for the first time, resolve the position of oxygen columns in ADF-STEM images obtained from pyrochlore ($\text{Yb}_2\text{Ti}_2\text{O}_7$) as well as perovskite (LaFeO_3) oxides.

2. Experimental methods

$\text{La}_{1-x}\text{FeO}_{3-\delta}$ ceramics were prepared via a solid-state reaction method using solid precursors La_2O_3 (Sigma Aldrich 99.99 wt.%) and Fe_2O_3 (Sigma Aldrich 99 wt.%). Stoichiometric amounts of the solid precursors were weighed and mixed by ball milling in isopropanol with yttria-stabilised zirconia (YSZ) media for 12h. The mixed powders were dried at 80 °C overnight and subsequently sieved and calcined in air at 1100 °C for 6 hours (5 °C/min ramp on heating and cooling) in an alumina crucible. Calcined powders were then sieved and ball milled in isopropanol with YSZ media for 12h, and subsequently dried at 80 °C for 12h. Pellets were then prepared from the dried calcined powders using a uniaxial press, and subsequently sintered in air for 6 hours at 1200 °C (5 °C ramp rate on heating and cooling in each case) in an alumina crucible.

$\text{Pr}_2\text{Zr}_2\text{O}_7$ samples were prepared in polycrystalline form by the conventional solid-state method. Powders of Pr_6O_{11} (99.9 %) and ZrO_2 (99 %) were weighed in stoichiometric amounts, mixed together and heat treated in air for several days (in four steps) at 1400-1450 °C, with intermediate grindings, see Ref. [43] for more information. The resulting sample (sample A) was brown in colour. A second praseodymium zirconate sample (sample B) was prepared under similar conditions, with an additional annealing step where the sample was heat treated in a reducing atmosphere ($\text{Ar} + 3\% \text{H}_2$) at 1200 °C for 2 days. The resulting powder was bright green coloured. Sample A is expected to be of the nominal composition $\text{Pr}_2\text{Zr}_2\text{O}_7$, whereas the reduced sample, B, is expected to be of composition $\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$.

The powder were blended in Al powder and then rolled into thin sheets. The TEM specimens from the thin sheets were prepared using standard routes, i.e. mechanical grinding and polishing followed by ion milling to electron transparency using Ar^+ initially at 6 keV and finally at 0.5 keV. A doubly-corrected JEOL ARM200F microscope, operating at 200 keV, was employed to study the atomic structure of samples. ADF-STEM images were obtained from thin ($t/\lambda < 0.5$, where t is the sample thickness and λ is the inelastic mean free path of electron in the samples), flat, defect-free and uniform regions of interest of the samples using an ADF detector where the inner angle of the ADF detector was $\sim 4.6\alpha$ (α , the convergence semi-angle of the electron probe, was ~ 15 or ~ 30 mrad). Up to sixty ADF images, all with a dwell time of 10 $\mu\text{s}/\text{pixel}$, were collected sequentially from a region of interest and subsequently the images were aligned using both normalized cross-correlation (using in-house code) and non-rigid image registration (using Smart Align software [44]) and then

summed to obtain high quality ADF-STEM images with a good (> 30) signal to noise ratio (SNR). The DetectColumns [15] program was used to calculate the position and intensity of atom columns in the atomic resolution ADF-STEM images. Multi-slice frozen phonon (>30 frozen phonons) image simulations were performed with Prismatic [45, 46] and Dr. Probe [47] software to compare with experimental data (see Section I in Supplemental Material for more information).

3. Results

3.1 Annular dark-field STEM

A representative ADF-STEM image from sample A ($\text{Pr}_2\text{Zr}_2\text{O}_7$) obtained along the $[2\ 1\ 1]$ direction is shown in Figs. 1(a). There are three types of cation columns in this projection as shown in Fig. 1(b); those containing only *A*-site cations (i.e. Pr), those containing only *B* site cations (i.e. Zr), and mixed (*M*) columns containing 50% *A* and 50% *B*. The *M* columns can be divided into two different types, *M1* and *M2*, according to the number and position of their nearby oxygen columns. There are twice as many oxygen atoms around *M1* columns in comparison to *M2* columns. The oxygen atoms around *M2* columns are closer to the centre of the cation columns. As a result, in the ADF-STEM image the *M2* columns have a higher peak intensity while *M1* columns have a higher radial intensity at larger radii (> 70 pm) as displayed in Fig. 1(c). These results, consistent with mean radial intensity profiles obtained from another pyrochlore $\text{Yb}_2\text{Ti}_2\text{O}_7$ [15] (Figs. 1(d)), indicate that invisible light atoms like oxygen have an influence on the contrast of their nearby visible cation columns with a relatively high atomic number in ADF images.

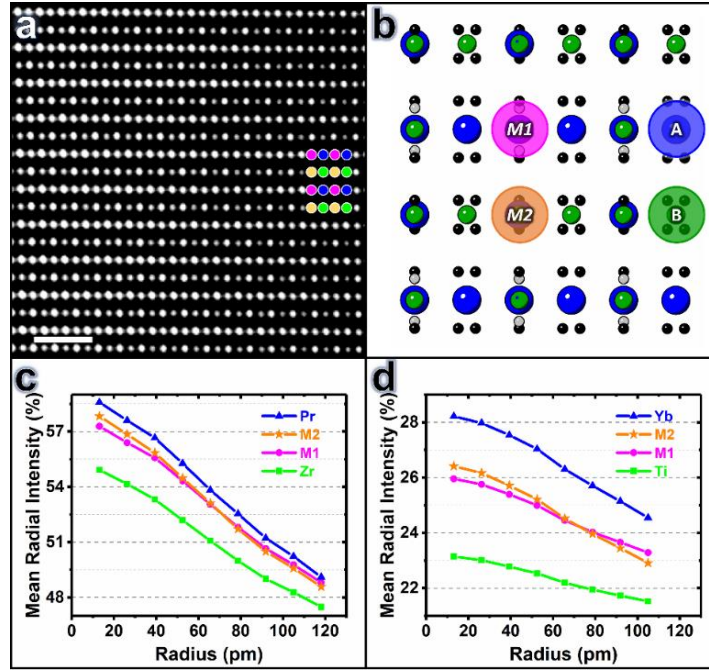


Fig. 1. (a) ADF-STEM image taken with an ADF detector inner angle of 2.4α from sample A ($\text{Pr}_2\text{Zr}_2\text{O}_7$) viewed along $[2\ 1\ 1]$ (scale bar is 1 nm). The four visible atom columns in this image, i.e. Pr, Zr, $M1$ and $M2$ columns, are shown in blue, green, pink and orange at the right of the image. (b) $[2\ 1\ 1]$ projection of the crystal structure of pyrochlore $A_2B_2O_7$ showing the four different types of atom columns. Here, A, B, $O(48f)$ and $O(8b)$ atoms are shown in blue, green, black and grey, respectively. (c) Mean radial intensity profiles for the four atom columns, i.e. Pr, Zr, $M1$ and $M2$, in (a) calculated using DetectColumns [15] program. (d) Mean radial intensity profiles for Yb, Ti, $M1$ and $M2$ atom columns in the ADF image of pyrochlore $\text{Yb}_2\text{Ti}_2\text{O}_7$ (Sample 1 in Ref. [15]) adapted from Ref. [15].

As illustrated in Fig. 2, we also investigated the intensity distribution around visible atom columns in ADF images obtained from sample B, which is expected to be oxygen deficient ($\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$). Interestingly, here we observe that the radial intensity of $M2$ columns is always more than that of $M1$, in other words, the radial intensity of $M1$ never exceeds that of $M2$. This could be due to the presence of oxygen vacancies, as Mostaed et al. [15] have already reported that oxygen vacancies alter the radial intensity distribution around M columns in ytterbium titanate. However, in the case of oxygen-deficient ytterbium titanate ($\text{Yb}_2\text{Ti}_2\text{O}_{7-\gamma}$), the intensity of $M1$ columns is always greater than that of $M2$ (Fig. 2(c)) whereas the intensity of $M2$ columns is always greater than that of $M1$ in $\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$ (Fig. 2(b)). This could be explained by oxygen vacancies in $\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$ being in different Wyckoff positions in comparison to those in $\text{Yb}_2\text{Ti}_2\text{O}_{7-\gamma}$. We will consider this further in the Discussion.

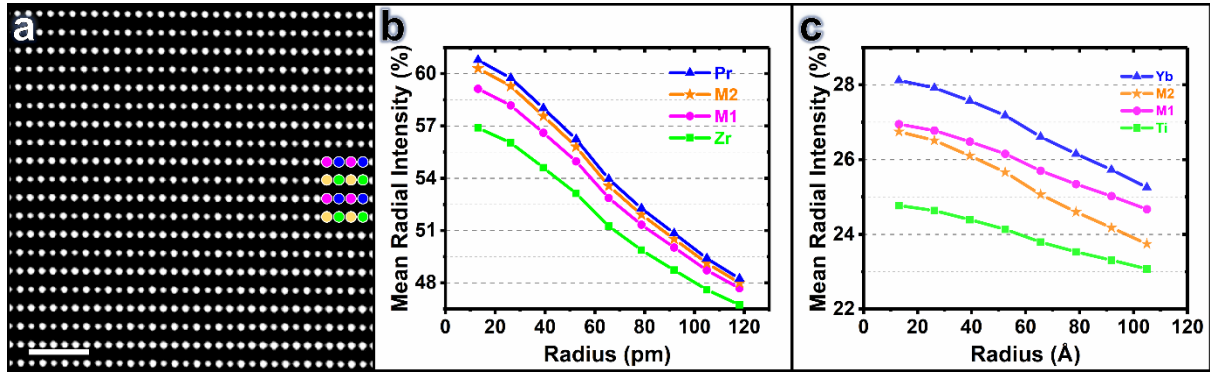


Fig. 2. (a) ADF-STEM image taken with an ADF detector inner angle of 2.4α from sample B ($\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$) viewed along $[2\ 1\ 1]$ (scale bar is 1 nm). The four visible atom columns in this image, i.e. Pr, Zr, *M1* and *M2* columns, are shown in blue, green, pink and orange at the top right of the image, respectively. (b) Mean radial intensity profiles for the four atom columns, i.e. Pr, Zr, *M1* and *M2*, in (a) calculated using DetectColumns [15] program. (c) Mean radial intensity profiles for Yb, Ti, *M1* and *M2* atom columns in the ADF image of pyrochlore $\text{Yb}_2\text{Ti}_2\text{O}_{7-\gamma}$ (Sample 2 in Ref. [15]) adapted from Ref. [15].

We also investigate the effect of oxygen atoms on the shape of their nearby cation columns in LaFeO_3 with a perovskite structure. Fig. 3(b) shows an ADF-STEM image viewed along $[110]$ direction obtained from LaFeO_3 . As schematically shown in Fig. 3(a), there are two types of cation columns in this projection; those containing only La (displayed in blue) and those containing only Fe (displayed in pink and yellow). The Fe columns can be divided into two different columns, *Fe1* and *Fe2*, according to the position of their nearby oxygen columns. As shown in Fig. 3(a), the oxygen atoms around *Fe1* columns are close to the vertical (y) axis of the image while those around *Fe2* columns are close to the horizontal (x) axis. As the number of oxygen atoms and their distance from the centre of cation columns are the same for both the *Fe1* and *Fe2* columns, the intensity of these atom columns in the ADF-STEM images should be similar. The intensities of atom columns shown in Fig. 3(b) are plotted as histograms in Fig. 3(c) and, as expected, there is no difference between the intensity of the *Fe1* and *Fe2* columns. Nevertheless, the width of the *Fe1* and *Fe2* columns are not the same. The calculated standard deviations obtained from two-dimensional (2D) Gaussian fitting on Fe columns (i.e. width of Fe columns) in ADF images obtained from LaFeO_3 samples are illustrated in Figs. 3(d) and 3(e). We observed that the *Fe2* columns are broader than the *Fe1* columns along the x direction (Fig. 3(d)) while the *Fe1* columns are broader along the y direction (Fig. 3(e)). These results are consistent with the position of oxygen atoms along x and y around these columns. In order to prove that the observed difference in the shape of the *Fe1* and *Fe2* columns is due to the difference in the position of their nearby oxygen atoms, we performed ADF-STEM image simulations using the Prismatic [45, 46] software. The simulation results (Fig. 3(f)), consistent

with the experimental results (Fig. 3(d) and 3(e)), indicate the *Fe2* columns are broader along the x direction whereas the *Fe1* columns are broader along the y direction; i.e. the oxygen atoms change the shape of their nearby cation columns.

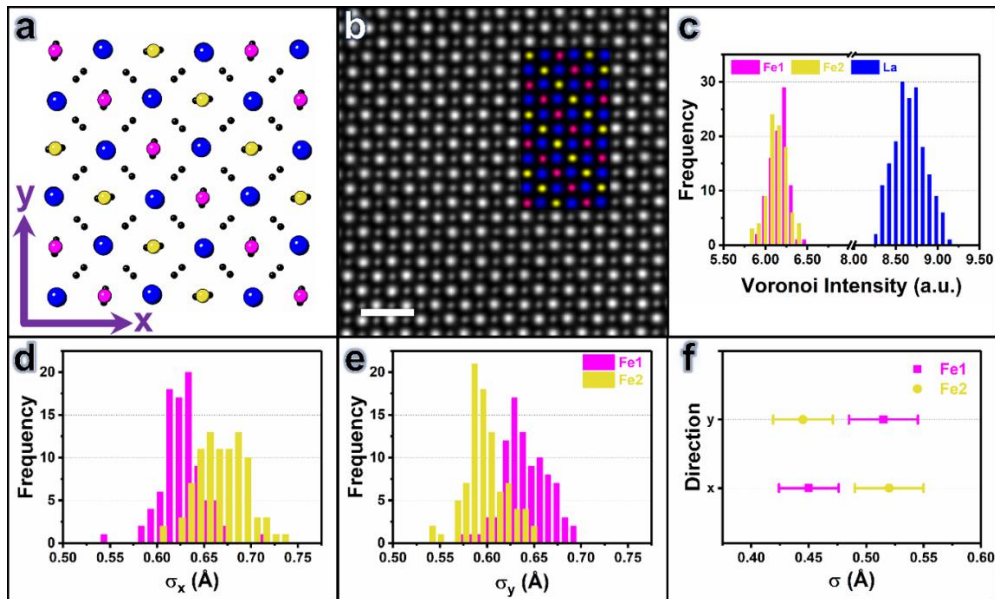


Fig. 3. (a) $[1\ 1\ 0]$ projection of the crystal structure of LaFeO_3 . Here, La and O atoms are displayed in blue and black, respectively. The Fe atoms, whose nature depends on the position of their nearby oxygen atoms, are shown in pink (the *Fe1* columns) and yellow (the *Fe2* columns). (b) ADF-STEM image taken with an ADF detector inner angle of 4.6α from the perovskite LaFeO_3 viewed along $[1\ 1\ 0]$ (scale bar is 1 nm). The 2D Gaussian fitting image on the three visible atom columns in this image is overlaid in which the La, *Fe1* and *Fe2* columns are shown in blue, pink and yellow, respectively. (c) Histograms of the extracted mean integrated intensities from (b) using Voronoi cells centred on each atomic column. (d) and (e) display the histograms of the standard deviations of the Fe columns along the horizontal (x) and vertical (y) axes obtained from the 2D Gaussian fitting on those columns shown in (b), respectively. (f) Standard deviations of the simulated Fe columns along the x and y axes.

3.2 Electron energy loss spectroscopy

The change in colour of praseodymium zirconate from brown ($\text{Pr}_2\text{Zr}_2\text{O}_7$) to green ($\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$) after reduction in $\text{Ar} + 3\%\text{H}_2$ atmosphere is due to partial reduction of Pr^{4+} cations to Pr^{3+} [20, 48]. As electron energy loss near-edge structure (ELNES) spectra can be used to obtain information on the Pr valence state in praseodymium oxides [49, 50], we examine the oxidation state of Pr in our praseodymium zirconate samples using Pr- $M_{5,4}$ ELNES spectra. Pr shows two white lines M_5 and M_4 , due to electron transitions from $3d_{5/2}$ and $3d_{3/2}$ subshells to the outer unoccupied $4f$ states, respectively. In praseodymium oxides, the number and position of peaks in the Pr- $M_{5,4}$ ELNES spectrum depends on the valence state of the Pr atoms. For instance, the Pr^{3+} - M_4 white line shows a shoulder on its low energy side that is absent in Pr^{4+} - M_4 [50]. In addition, a decrease in the Pr valence state leads to (a) a shift in the Pr- $M_{5,4}$ peaks to lower

energies and (b) a decrease in the intensity ratio I_{M_5}/I_{M_4} which is called the branching ratio [50].

The experimental Pr- $M_{5,4}$ ELNES spectra obtained from samples A and B are shown in Fig. 4. Peak energies, extracted by fitting the data to three Lorentzian curves, as well as intensity ratios are listed in Table 1. As shown in Fig. 4 and Table 1, the energy losses as well as the branching ratio for Pr- $M_{5,4}$ decrease from the brown sample to the green one. In other words, the Pr valence state reduced from $\text{Pr}_2\text{Zr}_2\text{O}_7$ to $\text{Pr}_2\text{Zr}_2\text{O}_{7-\delta}$, i.e. Pr-O bonds in praseodymium zirconate are affected by reduction in Ar + 3% H_2 atmosphere.

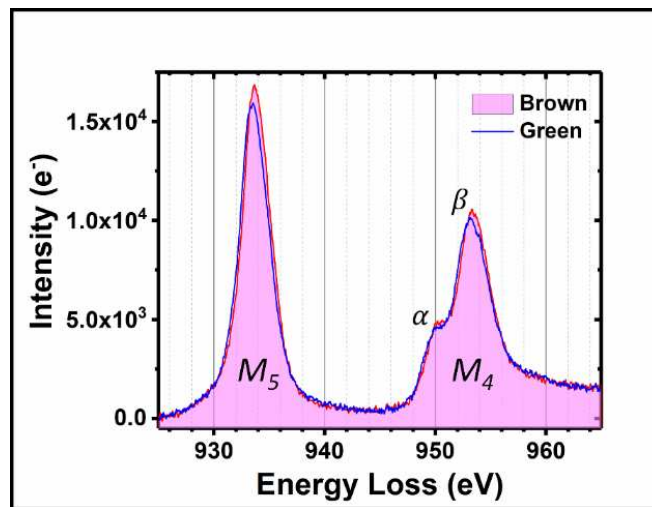


Fig. 4. Experimental EEL Pr- $M_{5,4}$ spectra of sample A and sample B ($t/\lambda = 0.35$).

Table 1. Pr ELNES energies (eV) and branching ratios for the Pr pyrochlore samples.

		Sample A	Sample B
Pr	E_{M_5}	933.74 ± 0.01	933.56 ± 0.01
	$E_{M_4}(\alpha)$	950.11 ± 0.03	949.83 ± 0.03
	$E_{M_4}(\beta)$	953.37 ± 0.02	953.22 ± 0.02
	I_{M_5}/I_{M_4}	1.30 ± 0.02	1.23 ± 0.02

4. Discussion

The ADF results obtained from the oxygen-deficient pyrochlores praseodymium zirconate and ytterbium titanate indicate that oxygen vacancies change the intensity distribution around $M1$ and $M2$ columns in both materials, but in different ways. According to the configuration of oxygen atoms around the M columns shown in Fig. 1(b), we expect a decrease in the occupancy of oxygen at $48f$ sites to reduce the ADF intensity of both the $M1$ and $M2$ columns. However,

as the distance between O-48*f* atoms and the centre of the *M2* columns (~50 pm) is much smaller than their distance from the centre of the *M1* columns (~100 pm), the reduction in the radial intensity of the *M2* columns caused by O-48*f* vacancies should mainly occur at small radii. Thus, the decrease in intensity for *M2* columns at small radii for Yb₂Ti₂O_{7-γ}, in comparison with stoichiometric Yb₂Ti₂O₇, can be attributed to O vacancies at 48*f* sites. In a pyrochlore structure (A₂B₂O₇), O-48*f* are located in tetrahedral voids created by two A-site and two B-site atoms (Fig. 5(a)) whereas O-8*b* are surrounded by only A-site cations (Fig. 5(b)). This means that B-site cation reduction can only be caused by oxygen vacancies at 48*f* sites. The EELS data presented in Ref. [15] showed that Ti exhibits a lower oxidation state in Yb₂Ti₂O_{7-γ} (Sample 2 in Ref. [15]) compared to Yb₂Ti₂O₇ (Sample 1 in Ref. [15]) which means that the oxygen vacancies are at 48*f* sites in Yb₂Ti₂O_{7-γ}. This is consistent with our radial intensity measurements.

Following the same reasoning, we expect a reduction in the ADF intensity only of *M1* columns for a decreased oxygen occupancy at 8*b* sites. Thus, the lower radial intensity of the *M1* columns, even at large radii, compared to *M2* columns in the ADF image obtained from Pr₂Zr₂O_{7-δ} (Fig. 2) can be explained by the presence of oxygen vacancies at 8*b* sites in this material. O-8*b* are located in tetrahedral voids created by four A-site atoms in a pyrochlore structure (Fig. 5(b)) and therefore O-8*b* vacancies only affect the oxidation state of A-site atoms. Our EELS results showed that Pr exhibits a lower oxidation state in Pr₂Zr₂O_{7-δ} compared to Pr₂Zr₂O₇, which means that the oxygen vacancies are at 8*b* positions in Pr₂Zr₂O_{7-δ}. This is consistent with our radial intensity measurements indicating that the oxygen vacancies in Pr₂Zr₂O_{7-δ} are at 8*b* positions. Thus, the off-centring of Pr³⁺ ions from their ideal crystallographic positions which has been demonstrated as the origin of the complex magnetic ground state of Pr₂Zr₂O₇ [20-23]) is due to Coulomb repulsion between the O-8*b* vacancies and the Pr cations. Our STEM technique allows us to observe the local defects in Pr₂Zr₂O_{7-δ} which were undetected in x-ray diffraction and neutron scattering measurements reported earlier [20-23].

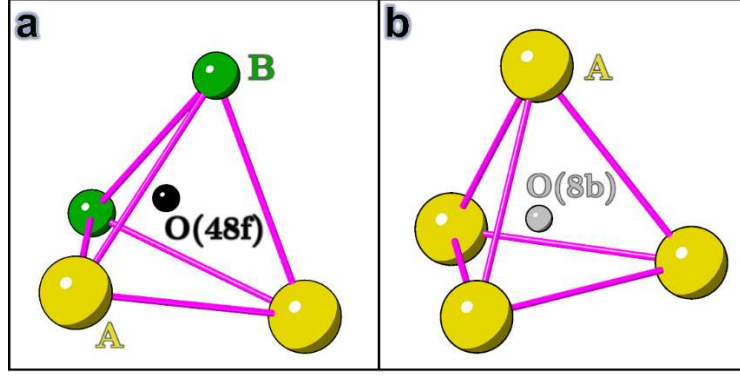


Fig. 5. Atom configuration around (a) O(48f) and (b) O(8b) in a pyrochlore $A_2B_2O_7$ structure.

The above results show that, while oxygen columns are invisible in ADF images, they make measurable contributions to the intensity of nearby cation columns. Careful comparisons of cation column intensity distribution for different samples can reveal changes in position, as well as occupancy, of oxygen atoms. The contribution of invisible light atoms to the contrast of ADF images can be direct or indirect, and we will return to this point later.

We can capture the essence of the behaviour in a very simple model by taking the two-dimensional ADF intensity distribution of a mixed atom column $I_V(x, y)$ to be the sum of scattering from a column of heavy cations $I_C(x, y)$ and adjacent light atoms $I_L(x, y)$, plus a second order term $I_{(C,L,\vec{d})}(x, y)$ due to the enhanced scattering that results from increased electron channelling when atom columns are very close to each other [51-53]. The latter depends on not only the type of atom in both columns but also their relative displacement (\vec{d}). So, the ADF intensity distribution of a visible cation column in an ADF image can be written as:

$$I_V = I_C + I_L + I_{(C,L,\vec{d})}. \quad (2)$$

For example, the intensity of the *Fe1* and *Fe2* columns in LaFeO_3 can be written as:

$$I_1 = I_{Fe_1} + I_{O_1} + I_{(Fe_1,O_1,\vec{d}_1)}, \quad (3)$$

$$I_2 = I_{Fe_2} + I_{O_2} + I_{(Fe_2,O_2,\vec{d}_2)}. \quad (4)$$

In the [001] projection of Fig. 3 the number of iron and oxygen atoms, and the distance between them, is identical for both *Fe1* and *Fe2* columns in a specimen of fixed thickness. Taking bonding effects to be unimportant we may assume radially-symmetric scattering for I_{Fe} which means the only difference in I_1 and I_2 is the relative displacement of the Fe and O atom columns in the second and third terms of each equation. If we subtract the intensity distribution of *Fe2* columns from that of *Fe1*, the remaining intensity is

$$I_1 - I_2 = I_{O_1} - I_{O_2} + I_{(Fe,O,\vec{d}_1)} - I_{(Fe,O,\vec{d}_2)}. \quad (5)$$

That is, scattering from Fe columns cancels, leaving only an intensity distribution that is dominated by the oxygen columns. Accordingly, for Fig. 3 we calculated the average intensity distribution of all *Fe1* columns (i.e. \bar{I}_1), all *Fe2* columns (i.e. \bar{I}_2) and their difference (i.e. $\bar{I}_1 - \bar{I}_2$) which are displayed in Figs. 6(a-c), respectively. Very careful examination of Figs. 3(a) and 3(b) reveals that both the *Fe1* and *Fe2* columns have an oval shape, with their long axes matching the position of nearby oxygen atoms as shown on the left of Fig. 6(a) and Fig. 6(b), i.e. vertical and horizontal respectively. In Fig. 6(c), according to Eq. (5), we expect positive values to correspond to O_1 and negative values to O_2 . We may separate the two by taking only positive values (Fig. 6(d)) or negative values (Fig. 6(e)). Using this method, the oxygen columns in $\text{Yb}_2\text{Ti}_2\text{O}_7$ with the pyrochlore structure were also resolved (see Section II in Supplemental Material).

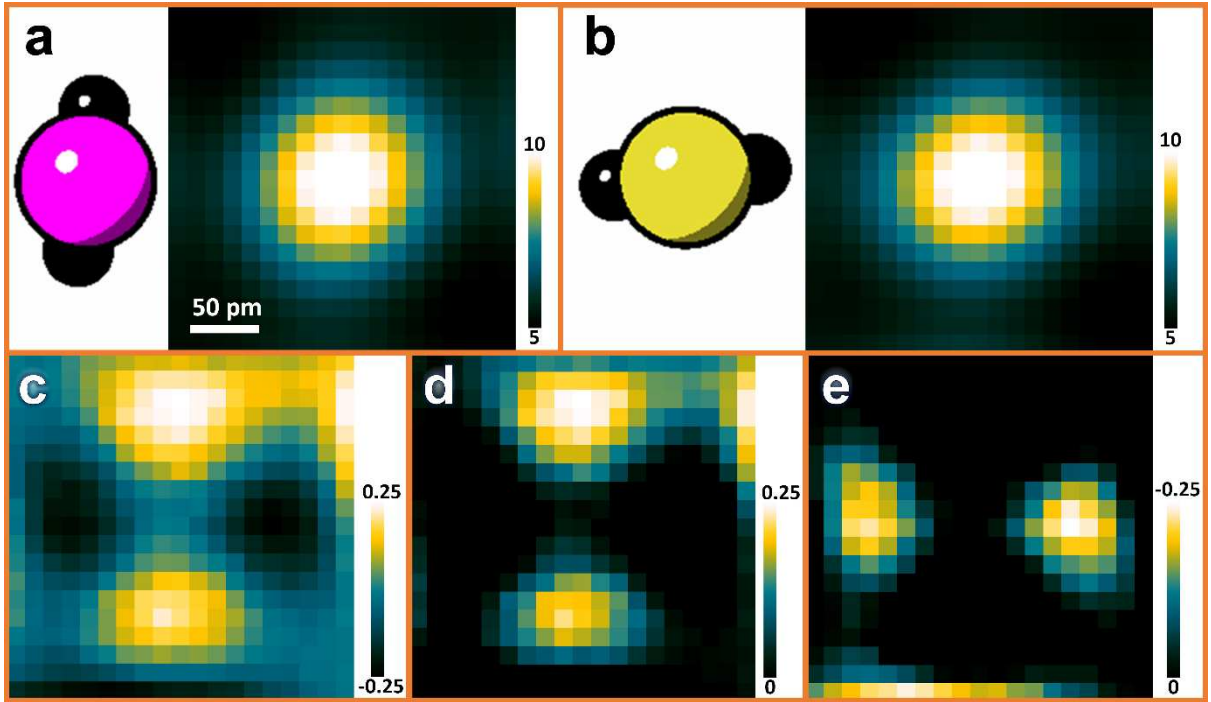


Figure 6: (a) Left: Schematic of the oxygen atom positions (black) around the Fe atoms (magenta) for the *Fe1* columns; Right: average ADF-STEM images obtained from all the *Fe1* columns shown in Fig. 3(b). (b) Left: Schematic of the oxygen atom positions (black) around the Fe atoms (yellow) for the *Fe2* columns; Right: average ADF-STEM images obtained from all the *Fe2* columns shown in Fig. 3(b). (c) Image of subtracted intensity for the average image of the *Fe2* columns from that of the *Fe1* columns, i.e. subtraction of (b) from (a). (d) Image of the positive values in (c) where the bright regions are consistent with the position of the oxygen atoms around the *Fe1* columns (with a distance of ~ 60 pm). (e) Image of the negative values in (c) where the bright regions are consistent with the position of the oxygen atoms around the *Fe2* columns (with a distance of ~ 60 pm).

The contrast of atom columns in ADF images is generated when a fraction of the incident electron beam is scattered to high angles by atoms in the specimen. As denoted in Eq. 1, the contribution of an oxygen column containing m oxygen atoms to the intensity of an ADF image would be proportional to mZ^n ($Z_O = 8$) is significantly lower than that of a column containing heavy cations with the same number of atoms. In the case of LaFeO_3 with a thickness of ~ 40 nm (Fig. 3), regardless the effects of dynamic scattering such as the probe channelling and dechannelling, the contribution of an oxygen column to the intensity of the ADF image would be less than 2% and 7% of the contribution of a La and Fe column, respectively (where $m_{La} = m_{Fe} = 100$, $m_O = 50$ and $n = 1.7$). In reality, the contribution of oxygen atoms to the intensity of ADF image is much lower even than this, because the probe intensity ($J(r)$ in Eq. 1) at the position of oxygen atom columns is much lower than that at heavy cation atom columns. In fact, when the probe is at the oxygen column positions, the probe is affected both by dechannelling away from oxygen columns, and channelling along heavy cation columns [41, 42]. This effect is shown in the simulation of Fig. 7 for different incident probe positions. When the probe is at the oxygen column positions (probe positions: $P1$ and $P3$ in Fig. 7(b)) or close to them (e.g. $P4$), the probe intensity spreads considerably on, and oscillates around, their nearby iron column. This will be the case even if there is no oxygen column around the Fe column (see Section IV in Supplemental Material) due to the attractive Coulombic potential of the heavy iron atoms. However, as shown in Figs. 7(c-e) the beam intensity at the iron column is significantly higher in the presence of oxygen columns. For instance, as illustrated in Fig. 7(d), when the probe is at the $P3$ position, the total beam intensity at the iron column in the presence of oxygen columns (A_{Fe}) is ~ 3.34 which is significantly higher ($\sim 27\%$) than that value in the absence of oxygen columns ($A_{Fe'} \approx 2.63$). Therefore, when the probe is at oxygen columns in the vicinity of heavy cation columns, the light oxygen atoms enhance spreading of the electron beam onto their nearby iron columns where heavy iron atoms, according to Eq. 1, can significantly contribute to the contrast of ADF images (see also Section V in Supplemental Material).

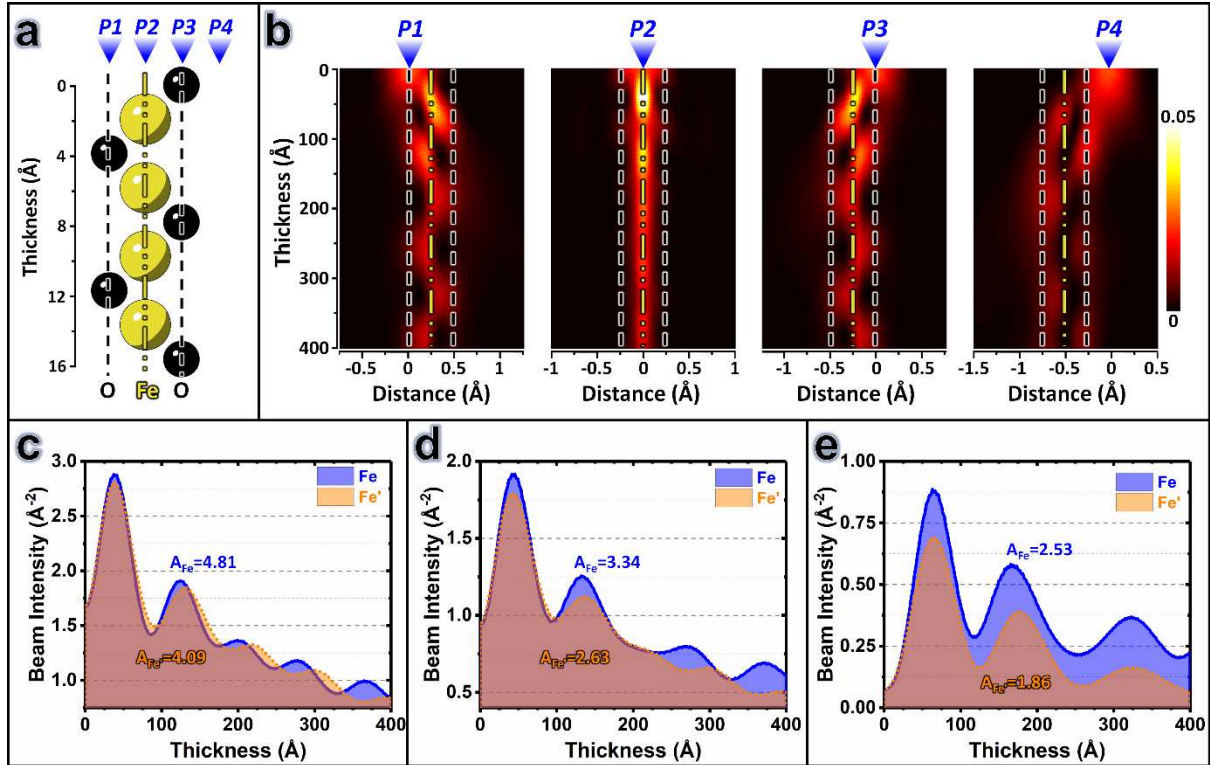


Figure 7: (a) Schematic of the atomic positions along an iron column and its nearby oxygen columns. (b) Simulated 2D beam intensity propagation profiles using Dr. Probe software [47] when the probe was at the $P1$, $P2$, $P3$ and $P4$ positions displayed in (a). Beam intensity depth profiles along the iron column in the presence of oxygen columns (blue lines) and in the absence of oxygen columns (orange dashed lines) when the probe is located at (c) $P2$, (d) $P3$ and (e) $P4$ positions. Here, A_{Fe} and $A_{Fe'}$ show the integration of the beam intensity over 40 nm along iron columns in the presence of oxygen columns and in the absence of oxygen columns, respectively.

The contribution of oxygen atoms to a typical ADF image is thus so small that oxygen columns are effectively invisible. Only by subtraction of images, in which the heavy atom contribution cancels, can the contribution of oxygen columns be observed as shown in our experimental and simulation results (Figs. 1–3 and 6). The result of this procedure is critically dependent upon the noise level, whose intensity is uncorrelated in the two images being subtracted and thus remains in the difference image. In our experimental ADF-STEM data, a sum of many individual images, the mean signal to noise ratio is between 30:1 and 60:1, allowing the oxygen atom signal to easily be detected. We can also produce noisier data by summing fewer images, and find that while oxygen columns can also be traced in ADF images with lower SNRs (see Fig. S3 in Section III of Supplemental Material) a minimum SNR of roughly 10:1 is needed.

In summary, using the methods described here, it is possible to obtain (quantitative) information about light atoms (e.g. O) in materials containing both light and relatively heavy elements (e.g. oxides) using ADF-STEM provided that we use certain viewing directions (zone

axes) for imaging in which the oxygen atoms are very close to atom columns containing heavy atoms. We expect the contrast of the resolved oxygen columns in the ADF images to contain quantitative information regarding the occupancy of oxygen sites. The methods can readily be used to investigate oxygen displacement/occupancy in other functional ceramic oxides with perovskite (e.g. BaTiO₃ [54, 55], LaMnO₃ [56], Na_{0.5}Bi_{0.5}TiO₃ [6], BiFeO₃ [57], etc.) or pyrochlore (e.g. Dy₂Ti₂O₇ [58], Tb₂Hf₂O₇ [59], etc.) structure where oxygen plays a key role in their physical and electrical properties. Furthermore, these methods can also be applied to characterize other invisible light atoms in the ADF images obtained from materials containing light and heavy elements, such as Li in lithium-ion battery materials [27, 60, 61].

5. Conclusions

We have investigated the atomic structure of three oxides with either a pyrochlore (praseodymium zirconate and ytterbium titanate) or a perovskite (lanthanum ferrite) structure using aberration corrected STEM. The intensity of visible atom columns in ADF images is sensitive to the number and position of their nearby invisible light atoms (i.e. oxygen). In addition, by examining the radial intensity profile of the cation columns, oxygen vacancies are observed at *8b* and *48f* Wyckoff sites in oxygen-deficient praseodymium zirconate (Pr₂Zr₂O_{7- δ}) and ytterbium titanate (Yb₂Ti₂O_{7- γ}), respectively. According to the atom configuration in pyrochlore *A₂B₂O₇*, oxygen vacancies at *8b* sites in Pr₂Zr₂O_{7- δ} or *48f* sites in Yb₂Ti₂O_{7- γ} would be compensated by a reduction in the oxidation state of Pr or either Yb or Ti cations, respectively. EELS data is consistent with those radial intensity measurements and showed that Pr in Pr₂Zr₂O_{7- δ} and Ti in Yb₂Ti₂O_{7- γ} have lower oxidation states in comparison to the stoichiometric material. We also devised a simple technique by which invisible light atoms like O can be resolved in the ADF images. Using this technique we have resolved the invisible oxygen columns in ADF-STEM images obtained from perovskite and pyrochlore structures. We expect this approach to be applicable for any other material comprised of atoms with widely differing atomic number.

6. Acknowledgements

The authors acknowledge the Engineering and Physical Sciences Research Council (EPSRC) UK grants EP/N032233/1, EP/L017563/1 and EP/T005963/1.

References

- [1] J.S. Gardner, M.J.P. Gingras, J.E. Greedan, Magnetic pyrochlore oxides, *Reviews of Modern Physics*, 82 (2010) 53-107.
- [2] R.J.D. Tilley, *Perovskites: Structure-Property Relationships*, John Wiley & Sons, Ltd, Chichester, 2016.
- [3] N.W. Thomas, Crystal structure–physical property relationships in perovskites, *Acta Crystallographica Section B-Structural Science*, 45 (1989) 337-344.
- [4] Y. Bai, T. Siponkoski, J. Perantie, H. Jantunen, J. Juuti, Ferroelectric, pyroelectric, and piezoelectric properties of a photovoltaic perovskite oxide, *Applied Physics Letters*, 110 (2017).
- [5] I.A. Sergienko, V. Keppens, M. McGuire, R. Jin, J. He, S.H. Curnoe, B.C. Sales, P. Blaha, D.J. Singh, K. Schwarz, D. Mandrus, Metallic "ferroelectricity" in the pyrochlore $\text{Cd}_2\text{Re}_2\text{O}_7$, *Physical Review Letters*, 92 (2004).
- [6] M. Li, M.J. Pietrowski, R.A. De Souza, H.R. Zhang, I.M. Reaney, S.N. Cook, J.A. Kilner, D.C. Sinclair, A family of oxide ion conductors based on the ferroelectric perovskite $\text{Na}_{0.5}\text{Bi}_{0.5}\text{TiO}_3$, *Nature Materials*, 13 (2014) 31-35.
- [7] J. Chun, P.G. Reuvekamp, D. Chen, C. Lin, R.K. Kremer, Promising high-k dielectric permittivity of pyrochlore-type crystals of $\text{Nd}_2\text{Hf}_2\text{O}_7$, *Journal of Materials Chemistry C*, 3 (2015) 491-494.
- [8] G. Wang, Z.L. Lu, H.J. Yang, H.F. Ji, A. Mostaed, L.H. Li, Y.Q. Wei, A. Feteira, S.K. Sun, D.C. Sinclair, D.W. Wang, I.M. Reaney, Fatigue resistant lead-free multilayer ceramic capacitors with ultrahigh energy density, *Journal of Materials Chemistry A*, 8 (2020) 11414-11423.
- [9] Z. Lu, G. Wang, W. Bao, J. Li, L. Li, A. Mostaed, H. Yang, H. Ji, D. Li, A. Feteira, F. Xu, D.C. Sinclair, D. Wang, S.-Y. Liu, I.M. Reaney, Superior energy density through tailored dopant strategies in multilayer ceramic capacitors, *Energy & Environmental Science*, 13 (2020) 2938-2948.
- [10] S.L. Swartz, C.A. Randall, A.S. Bhalla, Dielectric and Pyroelectric Properties of Cadmium Niobate Ceramics, *Journal of the American Ceramic Society*, 72 (1989) 637-641.
- [11] S. Murakami, D.W. Wang, A. Mostaed, A. Khesro, A. Feteira, D.C. Sinclair, Z.M. Fan, X.L. Tan, I.M. Reaney, High strain (0.4%) $\text{Bi}(\text{Mg}_{2/3}\text{Nb}_{1/3})\text{O}_3\text{-BaTiO}_3\text{-BiFeO}_3$ lead-free piezoelectric ceramics and multilayers, *Journal of the American Ceramic Society*, 101 (2018) 5428-5442.
- [12] C.A. Hancock, J.M. Porras-Vazquez, P.J. Keenan, P.R. Slater, Oxyanions in perovskites: from superconductors to solid oxide fuel cells, *Dalton Transactions*, 44 (2015) 10559-10569.
- [13] R. Jin, J. He, S. McCall, C.S. Alexander, F. Drymiotis, D. Mandrus, Superconductivity in the correlated pyrochlore $\text{Cd}_2\text{Re}_2\text{O}_7$, *Physical Review B*, 64 (2001).
- [14] J. Lim, J. Yu, Role of oxygen vacancy in the spin-state change and magnetic ordering in $\text{SrCoO}_{3-\delta}$, *Physical Review B*, 98 (2018).
- [15] A. Mostaed, G. Balakrishnan, M.R. Lees, Y. Yasui, L.J. Chang, R. Beanland, Atomic structure study of the pyrochlore $\text{Yb}_2\text{Ti}_2\text{O}_7$ and its relationship with low-temperature magnetic order, *Physical Review B*, 95 (2017).
- [16] D.D. Cuong, B. Lee, K.M. Choi, H.S. Ahn, S. Han, J. Lee, Oxygen vacancy clustering and electron localization in oxygen-deficient SrTiO_3 : LDA+U study, *Physical Review Letters*, 98 (2007).
- [17] J. He, A. Borisevich, S.V. Kalinin, S.J. Pennycook, S.T. Pantelides, Control of Octahedral Tilts and Magnetic Properties of Perovskite Oxide Heterostructures by Substrate Symmetry, *Physical Review Letters*, 105 (2010).
- [18] G. Sala, M.J. Gutmann, D. Prabhakaran, D. Pomaranski, C. Mitchelitis, J.B. Kycia, D.G. Porter, C. Castelnovo, J.P. Goff, Vacancy defects and monopole dynamics in oxygen-deficient pyrochlores, *Nature Materials*, 13 (2014) 488-493.
- [19] A. Mostaed, G. Balakrishnan, M.R. Lees, R. Beanland, Electron-irradiation induced defects in $\text{Yb}_2\text{Ti}_2\text{O}_7$, *Acta Materialia*, 143 (2018) 291-297.

- [20] S.M. Koohpayeh, J.J. Wen, B.A. Trump, C.L. Broholm, T.M. McQueen, Synthesis, floating zone crystal growth and characterization of the quantum spin ice $\text{Pr}_2\text{Zr}_2\text{O}_7$ pyrochlore, *Journal of Crystal Growth*, 402 (2014) 291-298.
- [21] N. Martin, P. Bonville, E. Lhotel, S. Guitteny, A. Wildes, C. Decorse, M.C. Hatnean, G. Balakrishnan, I. Mirebeau, S. Petit, Disorder and Quantum Spin Ice, *Physical Review X*, 7 (2017).
- [22] S. Petit, E. Lhotel, S. Guitteny, O. Florea, J. Robert, P. Bonville, I. Mirebeau, J. Ollivier, H. Mutka, E. Ressouche, C. Decorse, M.C. Hatnean, G. Balakrishnan, Antiferroquadrupolar correlations in the quantum spin ice candidate $\text{Pr}_2\text{Zr}_2\text{O}_7$, *Physical Review B*, 94 (2016).
- [23] J.J. Wen, S.M. Koohpayeh, K.A. Ross, B.A. Trump, T.M. McQueen, K. Kimura, S. Nakatsuji, Y. Qiu, D.M. Pajerowski, J.R.D. Copley, C.L. Broholm, Disordered Route to the Coulomb Quantum Spin Liquid: Random Transverse Fields on Spin Ice in $\text{Pr}_2\text{Zr}_2\text{O}_7$, *Physical Review Letters*, 118 (2017).
- [24] J.A. Alonso, C. Cascales, I. Rasines, Oxygen Vacancy Ordering in the Defect Pyrochlore $\text{Pb}_2[\text{TiSb}]\text{O}_{6.5}$: A Rietveld Refinement of Neutron Powder Diffraction Data, *Acta Crystallographica Section C-Crystal Structure Communications*, 45 (1989) 3-7.
- [25] J. Blasco, J. Garcia, J.M. deTeresa, M.R. Ibarra, J. Perez, P.A. Algarabel, C. Marquina, C. Ritter, Structural, magnetic, and transport properties of the giant magnetoresistive perovskites $\text{La}_{2/3}\text{Ca}_{1/3}\text{Mn}_{1-x}\text{Al}_x\text{O}_{3-\delta}$, *Physical Review B*, 55 (1997) 8905-8910.
- [26] R.I. Dass, J.B. Goodenough, Multiple magnetic phases of $\text{La}_2\text{CoMnO}_{6-\delta}$ ($0 \leq \delta \leq 0.05$), *Physical Review B*, 67 (2003).
- [27] C.R. Fell, D.N. Qian, K.J. Carroll, M.F. Chi, J.L. Jones, Y.S. Meng, Correlation Between Oxygen Vacancy, Microstrain, and Cation Distribution in Lithium-Excess Layered Oxides During the First Electrochemical Cycle, *Chemistry of Materials*, 25 (2013) 1621-1629.
- [28] S. McIntosh, J.F. Vente, W.G. Haije, D.H.A. Blank, H.J.M. Bouwmeester, Oxygen stoichiometry and chemical expansion of $\text{Ba}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.8}\text{Fe}_{0.2}\text{O}_{3-\delta}$ measured by in situ neutron diffraction, *Chemistry of Materials*, 18 (2006) 2187-2193.
- [29] K.W. Urban, C.L. Jia, L. Houben, M. Lentzen, S.B. Mi, K. Tillmann, Negative spherical aberration ultrahigh-resolution imaging in corrected transmission electron microscopy, *Philosophical Transactions of the Royal Society a-Mathematical Physical and Engineering Sciences*, 367 (2009) 3735-3753.
- [30] S.D. Findlay, N. Shibata, H. Sawada, E. Okunishi, Y. Kondo, Y. Ikuhara, Dynamics of annular bright field imaging in scanning transmission electron microscopy, *Ultramicroscopy*, 110 (2010) 903-923.
- [31] X. He, L. Gu, S.D. Guo, Z.L. Liu, R.C. Yu, Z.X. Mei, X.L. Du, B.G. Liu, Y. Ikuhara, X.F. Duan, Oxygen polarity and interfacial atomic arrangement in an $\text{Mg}_x\text{Zn}_{1-x}\text{O}/\text{C-MgO}/\text{sapphire}$ heterostructure, *Journal of Physics D-Applied Physics*, 46 (2013).
- [32] E. Okunishi, H. Sawada, Y. Kondo, Experimental study of annular bright field (ABF) imaging using aberration-corrected scanning transmission electron microscopy (STEM), *Micron*, 43 (2012) 538-544.
- [33] S.J. Zheng, C.A.J. Fisher, T. Kato, Y. Nagao, H. Ohta, Y. Ikuhara, Domain formation in anatase TiO_2 thin films on LaAlO_3 substrates, *Applied Physics Letters*, 101 (2012).
- [34] Y.M. Kim, S.J. Pennycook, A.Y. Borisevich, Quantitative comparison of bright field and annular bright field imaging modes for characterization of oxygen octahedral tilts, *Ultramicroscopy*, 181 (2017) 1-7.
- [35] I. MacLaren, Q.M. Ramasse, Aberration-corrected scanning transmission electron microscopy for atomic-resolution studies of functional oxides, *International Materials Reviews*, 59 (2014) 115-131.
- [36] P.D. Nellist, S.J. Pennycook, The principles and interpretation of annular dark-field Z-contrast imaging, *Advances in Imaging and Electron Physics*, Vol 113, 113 (2000) 147-203.
- [37] S.J. Pennycook, Atomic-scale imaging of materials by Z-contrast scanning transmission electron microscopy, *Analytical Chemistry*, 64 (1992) A263-A272.
- [38] S.J. Pennycook, B. Rafferty, P.D. Nellist, Z-contrast imaging in an aberration-corrected scanning transmission electron microscope, *Microscopy and Microanalysis*, 6 (2000) 343-352.
- [39] P. Hawkes, J.C.H. Spence, *Science of Microscopy*, Springer-Verlag New York 2007.

- [40] E.J. Kirkland, *Advanced Computing in Electron Microscopy*, 2 ed., Springer US2010.
- [41] P.M. Voyles, J.L. Grazul, D.A. Muller, Imaging individual atoms inside crystals with ADF-STEM, *Ultramicroscopy*, 96 (2003) 251-273.
- [42] R.J. Wu, A. Mittal, M.L. Odlyzko, K.A. Mkhoyan, Simplifying Electron Beam Channeling in Scanning Transmission Electron Microscopy (STEM), *Microscopy and Microanalysis*, 23 (2017) 794-808.
- [43] M. Ciomaga Hatnean, C. Decorse, M.R. Lees, O.A. Petrenko, G. Balakrishnan, Zirconate Pyrochlore Frustrated Magnets: Crystal Growth by the Floating Zone Technique, *Crystals*, 6 (2016).
- [44] L. Jones, H. Yang, T.J. Pennycook, M.S.J. Marshall, S. Van Aert, N.D. Browning, M.R. Castell, P.D. Nellist, Smart Align-a new tool for robust non-rigid registration of scanning microscope data, *Advanced Structural and Chemical Imaging*, 1 (2015).
- [45] C. Ophus, A fast image simulation algorithm for scanning transmission electron microscopy, *Advanced Structural and Chemical Imaging*, 3 (2017).
- [46] A. Pryor, C. Ophus, J.W. Miao, A streaming multi-GPU implementation of image simulation algorithms for scanning transmission electron microscopy, *Advanced Structural and Chemical Imaging*, 3 (2017).
- [47] J. Barthel, Dr. Probe: A software for high-resolution STEM image simulation, *Ultramicroscopy*, 193 (2018) 1-11.
- [48] S. Ferro, Physicochemical and Electrical Properties of Praseodymium Oxides, *International Journal of Electrochemistry*, 2011 (2011).
- [49] S.C. Cheng, V.P. Dravid, T.J. Goodwin, R.N. Shelton, H.B. Radousky, Determination of the valence of Pr in $(\text{Eu}_{1.5-x}\text{Pr}_x\text{Ce}_{0.5})\text{Sr}_2\text{Cu}_2\text{NbO}_{10}$ superconducting compounds by electron-energy-loss spectroscopy, *Physical Review B*, 53 (1996) 11779-11783.
- [50] J. Richter, A. Braun, A.S. Harvey, P. Holtappels, T. Graule, L. Gauckler, Valence changes of manganese and praseodymium in $\text{Pr}_{1-x}\text{Sr}_x\text{Mn}_{1-y}\text{In}_y\text{O}_3$ -delta perovskites upon cation substitution as determined with XANES and ELNES, *Physica B-Condensed Matter*, 403 (2008) 87-94.
- [51] M. Haruta, H. Kurata, H. Komatsu, Y. Shimakawa, S. Isoda, Effects of Electron channeling in HAADF-STEM intensity in $\text{La}_2\text{CuSnO}_6$, *Ultramicroscopy*, 109 (2009) 361-367.
- [52] L.F. Kourkoutis, M.K. Parker, V. Vaithyanathan, D.G. Schlom, D.A. Muller, Direct measurement of electron channeling in a crystal using scanning transmission electron microscopy, *Physical Review B*, 84 (2011).
- [53] R.F. Loane, E.J. Kirkland, J. Silcox, Visibility of single heavy atoms on thin crystalline silicon in simulated annular dark-field STEM images, *Acta Crystallographica Section A*, 44 (1988) 912-927.
- [54] G.Y. Yang, E.C. Dickey, C.A. Randall, D.E. Barber, P. Pinceloup, M.A. Henderson, R.A. Hill, J.J. Beeson, D.J. Skamser, Oxygen nonstoichiometry and dielectric evolution of BaTiO_3 . Part I - improvement of insulation resistance with reoxidation, *Journal of Applied Physics*, 96 (2004) 7492-7499.
- [55] G.Y. Yang, G.D. Lian, E.C. Dickey, C.A. Randall, D.E. Barber, P. Pinceloup, M.A. Henderson, R.A. Hill, J.J. Beeson, D.J. Skamser, Oxygen nonstoichiometry and dielectric evolution of BaTiO_3 . Part II - insulation resistance degradation under applied dc bias, *Journal of Applied Physics*, 96 (2004) 7500-7508.
- [56] J.T. Mefford, W.G. Hardin, S. Dai, K.P. Johnston, K.J. Stevenson, Anion charge storage through oxygen intercalation in LaMnO_3 perovskite pseudocapacitor electrodes, *Nature Materials*, 13 (2014) 726-732.
- [57] J. Seidel, P. Maksymovych, Y. Batra, A. Katan, S.Y. Yang, Q. He, A.P. Baddorf, S.V. Kalinin, C.H. Yang, J.C. Yang, Y.H. Chu, E.K.H. Salje, H. Wormeester, M. Salmeron, R. Ramesh, Domain Wall Conductivity in La-Doped BiFeO_3 , *Physical Review Letters*, 105 (2010).
- [58] H.M. Revell, L.R. Yaraskavitch, J.D. Mason, K.A. Ross, H.M.L. Noad, H.A. Dabkowska, B.D. Gaulin, P. Henelius, J.B. Kycia, Evidence of impurity and boundary effects on magnetic monopole dynamics in spin ice, *Nature Physics*, 9 (2013) 34-37.
- [59] R. Sibille, E. Lhotel, M.C. Hatnean, G.J. Nilsen, G. Ehlers, A. Cervellino, E. Ressouche, M. Frontzek, O. Zaharko, V. Pomjakushin, U. Stuhr, H.C. Walker, D.T. Adroja, H. Luetkens, C.

- Baines, A. Amato, G. Balakrishnan, T. Fennell, M. Kenzelmann, Coulomb spin liquid in anion-disordered pyrochlore $Tb_2Hf_2O_7$, *Nature Communications*, 8 (2017).
- [60] X. Gao, C.A.J. Fisher, T. Kimura, Y.H. Ikuhara, H. Moriwake, A. Kuwabara, H. Oki, T. Tojigamori, R. Huang, Y. Ikuhara, Lithium Atom and A-Site Vacancy Distributions in Lanthanum Lithium Titanate, *Chemistry of Materials*, 25 (2013) 1607-1614.
- [61] R. Huang, Y. Ikuhara, STEM characterization for lithium-ion battery cathode materials, *Current Opinion in Solid State & Materials Science*, 16 (2012) 31-38.