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Structural properties of mixed conductor

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$Ba_{1-x}Gd_{1-y}La_{x+y}Co_2O_{6-\delta}$ (BGLC) compositions with large compositional ranges of Ba, Gd, and La have been characterised with respect to phase compositions, structure, and thermal and chemical expansion. The results show a system with large compositional flexibility, enabling tuning of functional properties and thermal and chemical expansion. We show anisotropic chemical expansion and detailed refinements of emerging phases as La is substituted for Ba and Gd. The dominating phase is the double perovskite structure *Pmmm*, which is A-site ordered along the *c*-axes and with O vacancy ordering along the *b*-axis in the Ln-layer. Phases emerging when substituting La for Ba are orthorhombic Ba-deficient *Pbnm* and cubic $LaCoO_3$ -based $R\bar{3}c$. When La is almost completely substituted for Gd, the material can be stabilised in *Pmmm*, or cubic *Pm\bar{3}m*, depending on thermal and atmospheric history. We list thermal expansion coefficients for $x = 0-0.3$, $y = 0.2$.

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Introduction

The double perovskite $Ba_{1-x}Gd_{1-y}La_{x+y}Co_2O_{6-\delta}$ (BGLC) has shown good performance as a positrode (oxygen electrode) for Proton Ceramic Electrochemical Cells (PCECs).^{1,2} The structural properties entail several crystallographic and chemical phases,³ depending on the stoichiometry factors x and y – both reflecting La, substituted for Ba (x) and Gd (y). Furthermore, the thermal and chemical expansion is vital for mechanical stability when this materials class is applied as PCEC electrodes and subjected to large variations in tempera-

ture and chemical environment. In this work, we utilise Powder X-ray Diffraction (PX), Synchrotron Radiation Powder X-ray Diffraction (SR-PXD) and Powder Neutron Diffraction (PND) to determine the development of BGLC's constituent phases, with variations in x (0–0.5) and y (0–1). A tetragonal or orthorhombic double perovskite unit cell with ordering between Ba and the lanthanides along the *c*-axis can be expressed by the general formula $A_{1-x}^I A_x^{I*} A_{1-y}^{II} A_y^{II*} Co_2 O_{6-\delta}$, where A^I and A^{II} are ordered A-sites, and * denotes substitutions on the respective A-sites.

This A-site cation ordering can be seen for other materials in the same family^{4–9} often also accompanied by ordering of the oxygen vacancies located in the Ln layers, as δ varies between 0 and 1, resulting in a doubled *b*-axis and stabilising an orthorhombic unit cell.^{4,10} The Co-based double perovskites generally exhibit high electronic and ionic conductivity,¹¹ and introducing a mix between di- and trivalent A-site cations opens the possibility of tuning both functional properties as well as chemical and structural stability by adjusting average valence states (x) and ionic radius ($x + y$) of the A-site cations. The overall aim is to influence hydration thermodynamics, electrochemical performance and chemical stability. The system has three end members: $BaLaCo_2O_{6-\delta}$ ($x = 0$, $y = 1$, BLC), $BaGdCo_2O_{6-\delta}$ ($x = 0$, $y = 0$, BGC), and $Ba_{0.5}Gd_{0.5}La_{0.7}Co_2O_{6-\delta}$ ($x = 0.5$, $y = 0.2$, BGLC587), where 50% of the Ba on A^I is nominally substituted with La. In BLC, the A-site cations can take ordered ($BaLaCo_2O_{6-\delta}$) or disordered ($Ba_{0.5}La_{0.5}CoO_3$) configurations.^{4,12,13} The reason for the tendency

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of A-site order with decreasing y is the difference in ionic radii between the relatively large Ba^{2+} (ionic radius 1.61 Å), the smaller La^{3+} (ionic radius 1.36 Å),¹⁴ and the even smaller Gd^{3+} (ionic radius 1.27 Å).¹⁴ The size difference at $y = 1$ is not big enough to unambiguously favour the ordered state. By reduction (lower Co valence and oxygen content) or substituting Gd for La, the structure will gradually stabilise in the A-site ordered double perovskite configuration.

In BGC, the size difference between Ba and Gd is too large to enable an A-site disordered cubic polymorph in the dry state. BGC is reported with two phase transitions upon cooling: at 350 K the structure changes from tetragonal $P4/mmm$ (space group 123) to orthorhombic $Pmmm$ (space group 47). By further cooling, there is a second order phase transition starting at 247 K from $Pmmm$ to $Pmma$, (space group 51), where Co^{2+} and Co^{3+} order.⁵ The differentiation between $P4/mmm$ and $Pmmm$ is rooted in the ordering of oxygen vacancies in the Ln–O layer, and is thus dependent on the oxygen non-stoichiometry (δ), which changes with temperature and $p\text{O}_2$. The identification of orthorhombic distortion can be subtle, and is sometimes only seen in high-resolution XRD data.¹⁵

In BGLC587, 50% of the Ba is substituted with La, and the phase identification is more complicated. The aim of this work is to show how the phase compositions and unit cells change as the compositions vary in the x and y ranges. Moreover, we investigate the thermal expansion when x is varied between 0 and 0.3.

Experimental

Powders for High Temperature PXD – BGLC ($x = 0, 0.1, 0.2,$ and $0.3, y = 0.2$) – were prepared by combustion reaction of a nitrate solution. 40 g citric acid was dissolved in water in a large beaker on a hot plate. BaCO_3 was slowly added until fully dissolved. La, Gd, and Co nitrates were subsequently dissolved in stoichiometric ratios. The water was evaporated on a hot plate with magnetic stir until a gel was formed, and the beaker was put in a heating cabinet at 250 °C for the combustion reaction to happen. The citric acid remains were burned off at 400 °C for 1 hour. The ashes were crushed thoroughly in a mortar and calcined at 1100 °C for five hours in an alumina crucible to obtain the desired BGLC phase.

For SR-PXD on compositions with $x = 0, 0.1, 0.2, 0.3, 0.4,$ and 0.5 and $y = 0.2$ we used commercial powders as described in a previous work.¹¹ A second series for SR-PXD and PND ($x = 0$ and $y = 0, 0.2, 0.3, 0.5, 0.7, 0.8,$ and 1) was prepared by a solid-state reaction as described in a previous study.⁴

HTPXRD data were collected on a Bruker D8 Advance diffractometer using Mo radiation with a focussing Goebel mirror and LynxEye XE high energy detector. 2.5° Soller slits were used to reduce beam divergence. Data were collected over a range of 10–45° 2θ with a step size of 0.02° and a count time of 1 s per step. The high temperature stage used was an Anton Paar HTK1200. Samples were measured in flat plate mode using

standard 0.3 mm deep corundum crucibles. The samples were heated to 850 °C and cooled again to room temperature, first in air, then under argon (after evacuating the chamber and filling with Ar 3 times).

Iodometric titration, SR-PXD, and PND were performed as described in previous work.⁴ Hydration of BGLC82 was done by heating the commercial BGLC82 powder in air at 300 °C for 12 hours while bubbling the gas through a saturated solution of KBr in H_2O to control the humidity at 2.6%. After 12 hours, the powder was quenched to room temperature and sealed.

Results

The BGLC series of compositions contain a range of crystallographic phases with compositional flexibility. With $x = 0$, the main phase is orthorhombic $Pmmm$, where Ba and La/Gd is ordered along the c -axis. BGLC's with $x = 0.2$ has previously been studied by powder XRD and reported with $P4/mmm$ space group.³ Our results shows a doubling of the b -axis caused by ordering of the oxygen vacancies located in the Ln layer. The orthorhombic distortion is subtle in PXD diffractograms, and can only be seen in the high-resolution diffractograms, as the one presented for isotope enriched $\text{Ba}^{160}\text{GdCo}_2\text{O}_{6-\delta}$ (B^{160}GC) in Fig. 1 taken with TOF-PND.

Phase compositions in BGLC ($x = 0-0.5, y = 0.2$). Refinements of SR-PXD diffractograms for the series of compositions with $x = 0-0.5$ and $y = 0.2$ showed formation of three main phases, depending on La substitution. For $x = 0$, the main phase is the orthorhombic, ordered double perovskite $Pmmm$ (80 wt%), while there is an 11 wt% fraction of La-substituted GdCoO_3 $Pbnm$ (space group 62) and smaller fractions of BaCO_3 and Co_3O_4 . The $Pbnm$ phase reflects a Ba deficiency – possibly due to BaCO_3 formation during ambient air storage.

By increasing x , the $Pmmm$ becomes indistinguishable from $P4/mmm$, and this phase gradually diminishes while a fraction of GdCoO_3 $Pbnm$ emerges. From $x = 0.2$, a third phase – LaCoO_3 $R\bar{3}c$ (space group 167) – emerges and increases, leaving the end member, BGLC587 ($x = 0.5$) as a three-phase composite with 23 wt% of the original BGLC $P4/mmm$ phase, 50 wt% LaCoO_3 , and 22 wt% GdCoO_3 . Both the LaCoO_3 and GdCoO_3 phases are substituted with La and Ba, giving a good fit for all composition when the refinements were constrained to nominal overall cation compositions (Fig. 2).

All compositions of x have minor amounts of at least two among the phases BaCO_3 , Gd_2O_3 and Co_3O_4 , as shown in Table 1 where all crystalline phases are shown with tentative compositions and fractions of total.

The evolvments of unit cell parameters with Ba/La fraction allowed an approximate assignment of substitutions in LaCoO_3 and GdCoO_3 . The unit cell parameters in the initial $P4/mmm$ phase were unaffected by La substitution, indicating that the Gd/La ratio of 8/2 was the most stable configuration with respect to formation of secondary phases. In Table 2: unit





Fig. 1 PND diffractogram of BGC with Rietveld refinement to orthorhombic $Pmmm$.

cell volumes of the three BGLC phases with $x = 0-0.5$, $y = 0.2$ ($x = 0$ refined to $P4/mmm$). We give unit cell volumes of the three main phases with increasing x . For simplicity and comparison we have here increased the symmetry from $Pmmm$ to $P4/mmm$ for $x = 0$.

As can be seen from Table 2, there is no trend in cell volumes for the three phases, indicating that they stabilise each other by compositionally dependent symmetry changes, also reflecting the uncertainty in compositional determination based on Rietveld refinements alone.

We have previously shown that the chemical stability of BGLC in high steam pressures increases with x when $y = 0.2$,² and also with y when $x = 0$.¹⁶ The increased stability for $x = 0.5$ can be suggested to stem from the ability of the secondary phases to incorporate Ba and La, and thus stabilising the main $P4/mmm$ phase from decomposing to binary hydroxides. In Fig. 3 we show phase fractions in BGLC ($x = 0-0.5$, $y = 0.2$).

Phase compositions in BGLC ($x = 0$, $y = 0-1$). Refinements of SR-PXD and PND data show that $Pmmm$ is the dominating phase for $x = 0$, $y = 0-1$. The only composition with a secondary phase with different chemical composition than the main phase is $x = 0$, $y = 0.2$ (BGLC82), where there are fractions of La-substituted $GdCoO_3$ ($Pbnm$, 11.2 wt%) and $BaCO_3$ (7 wt%). All other compositions are refined to single-phase $Pmmm$. The highest values for $y = 0.8$ and 1 can be stabilised both in A-site ordered $Pmmm$, and cubic, disordered $Pm\bar{3}m$ (space group 221), as illustrated in the refined SR-XRD diffractogram

of reduced BLC, which contains 30 wt% cubic $Pm\bar{3}m$ and 70 wt% A-site ordered $Pmmm$ (Fig. 4).

In Fig. 5, we present unit cell volume and cell parameters for BGLC compositions where $x = 0$ and y varies from 0 to 1, representing the solid solution of BGC and BLC.

The increased cell volume is anisotropic along the c -axis, reflecting mainly the increased average Ln^{3+} radius as La replace Gd. The a - and b -axes are not affected by increased Ln^{3+} radius, as they are mainly fixed by the larger Ba^{2+} radius in the ab plane. The anisotropic evolution of lattice parameters with increasing y is in accordance with previous results in the $0.2 \leq y \leq 0.8$ range.³ The substitution of La for Gd also entails an increase in Co oxidation state as oxygen vacancies are gradually filled with increasing y . This is shown in Fig. 6, where we display the average Co oxidation state obtained by iodometric titration and accompanying oxygen non-stoichiometry as a function of y for BGLC ($x = 0$).

To investigate the effect of O occupancy vs. Co–O bond angle, we have refined the three highest compositions of x to a $P4/mmm$ unit cell (Fig. 7, right) where O-occupancies on the O2 position were also refined, acknowledging that the refinement of occupancy on O2 is close to 1 and thus within uncertainty. The resulting correlation between O occupancies and O–Co–O bond angle is presented in Fig. 7.

It can be seen from Fig. 7 that the oxygen in the Co–O layer (O2) is positioned closer to the Ln layer with increasing occupancy, and that it is also positioned closer to the Ln layer with



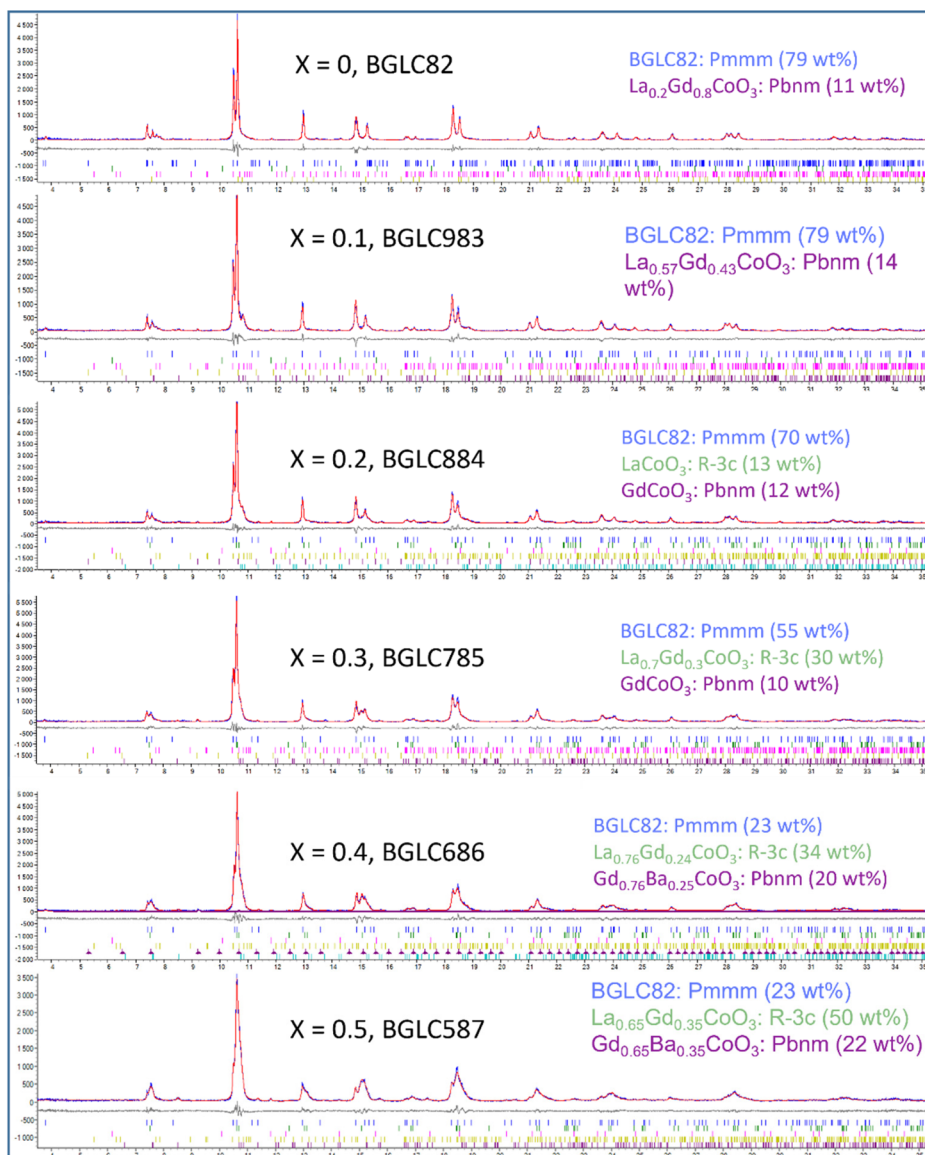


Fig. 2 SR-PXD rietveld refinements (red line) of BGLC ($x = 0-0.5$, $y = 0.2$). Minority phases Co_3O_4 , Gd_2O_3 and BaCO_3 are refined, but omitted in the figure legends.

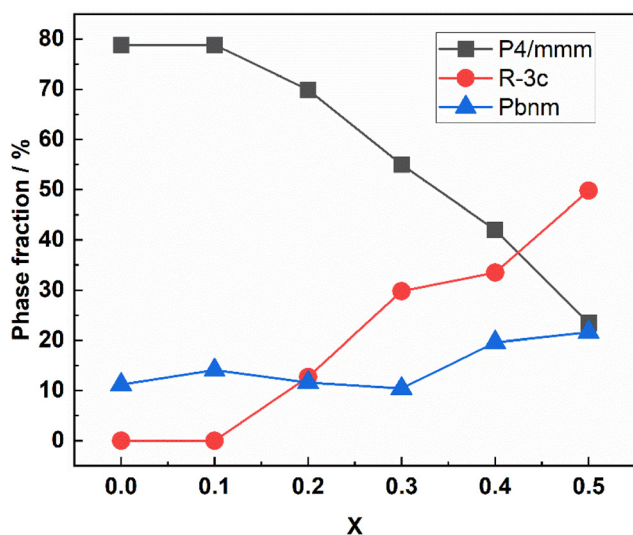
Table 1 Phase compositions in BGLC ($x = 0-0.5$, $y = 0.2$)

Sample	<i>Pmmm</i> BGLC wt% (composition)	<i>R</i> $\bar{3}$ <i>c</i> LaCoO ₃ wt% (composition)	<i>Pbnm</i> GdCoO ₃ wt% (composition)	Co ₃ O ₄ wt%	Gd ₂ O ₃ wt%	BaCO ₃ wt%
$x = 0$	78.8(4) BaGd _{0.8} La _{0.2} Co ₂ O _{6-δ}	—	11.2(5) La _{0.2} Gd _{0.8} CoO ₃	2.0(2)	—	7.2(2)
$x = 0.1$	78.8(4) BaGd _{0.8} La _{0.2} Co ₂ O _{6-δ}	—	14.1(2) La _{0.57} Gd _{0.43} CoO ₃	1.7(1)	0.58(4)	4.9(4)
$x = 0.2$	69.9(5) BaGd _{0.8} La _{0.2} Co ₂ O _{6-δ}	12.7(4) LaCoO ₃	11.6(4) La _{0.05} Gd _{0.88} Ba _{0.07} CoO ₃	1.54(8)	0.42(4)	3.8(2)
$x = 0.3$	55.0(7) BaGd _{0.8} La _{0.2} Co ₂ O _{6-δ}	29.8(8) La _{0.7} Gd _{0.3} CoO ₃	10.4(5) GdCoO ₃	—	0.73(4)	4.1(2)
$x = 0.4$	42.0(4) BaGd _{0.8} La _{0.2} Co ₂ O _{6-δ}	33.5(4) La _{0.76} Gd _{0.24} CoO ₃	19.6(2) Gd _{0.75} Ba _{0.25} CoO ₃	1.11(7)	0.64(4)	3.2(1)
$x = 0.5$	23.4(3) BaGd _{0.8} La _{0.2} Co ₂ O _{6-δ}	49.8(3) La _{0.65} Gd _{0.35} CoO ₃	21.6(3) Gd _{0.65} Ba _{0.35} CoO ₃	1.76(8)	—	3.4(2)



Table 2 Unit cell volumes of the three BGLC phases with $x = 0-0.5$, $y = 0.2$ ($x = 0$ refined to $P4/mmm$)

	BGLC82 $P4/mmm$	LaCoO ₃ $R\bar{3}c$	GdCoO ₃ $Pbnm$
$x = 0$	114.7(1)	—	215.80
$x = 0.1$	114.762(5)	—	214.12(8)
$x = 0.2$	114.803(5)	112.50(6)	214.92(7)
$x = 0.3$	114.668(8)	113.02(4)	217.64(8)
$x = 0.4$	114.600(9)	112.70(2)	216.26(6)
$x = 0.5$	114.64(2)	112.0(1)	215.99(6)

**Fig. 3** Phase fractions (wt%) vs. x in BGLC ($y = 0.2$).

decreasing occupancy on O3. We will therefore expect anisotropic chemical expansion as red-ox mainly occurs at O3. We will come back to this phenomenon later.

Hydration of BGLC: A-site order-disorder. We have previously reported the hydration of BGLC82,¹ BGC and BLC.⁴ In all of these ordered double perovskites, hydration induces a slow, irreversible weight gain, tentatively ascribed to increasing A-site cation disorder with accompanying oxidation upon hydration. SR-PXD shows no evolution of the unit cell reflecting this process. The diffractograms may however be refined

both to a reduced, ordered orthorhombic $Pmmm$ with unit cell parameters $a2b2c$, and to an oxidised, A-site disordered $Pmmm$ with unit cell parameters $2a3b3c$, both giving the same Bragg reflections and intensities. In Fig. 8 we give an example of fitting the SR-PXD diffractograms of dry BGLC82 to A-site ordered $Pmmm$ $a2b2c$, and hydrated BGLC82 to A-site disordered $Pmmm$ $2a3b3c$. Both refinements give equally good fit.

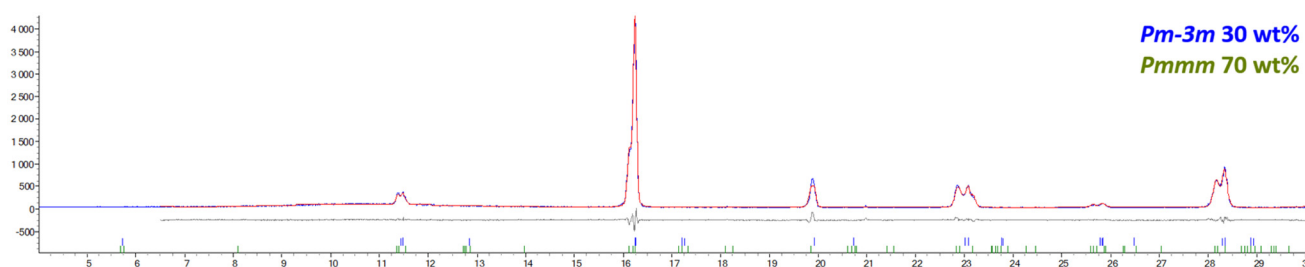
The refinements yield a suggested phase transition as shown in Fig. 9.

TEM/STEM. TEM analysis of BGLC82 ($x = 0$, $y = 0.2$) was performed after dehydration in dry air at 400 °C for 129 hours. The Selected Area Electron Diffraction (SAED) and Fast Fourier Transform images (Fig. 10) show a mix of domains with reflections from a larger unit cell in addition to the main phase. These phases are not compliant with $Pbnm$, and suggest that a deeper analysis must be performed, including also STEM-EDS on dry and hydrated samples to control for A-site order-disorder in the various domains.

STEM-EDS atomic mapping was performed on the same sample in dry state, and the results show how Ba and Gd order along the c -axis (Fig. 11). The La signal is not strong enough for this composition to distinguish A-site ordering of Ba and La from background signals.

HT-XRD. High temperature XRD reveals that BGLC maintains the tetragonal $P4/mmm$ unit cell from RT to 850 °C both in air and Ar. As can be seen in Fig. 12, chemical expansion is anisotropic, showing higher chemical expansion in the ab plane than along the c -axis at high temperatures for all compositions ($x = 0-0.3$, $y = 0.2$). There is a clear shift in thermal expansion for a and c from 300 °C and upwards, and as seen from our previous work,¹¹ 300 °C is the onset temperature for oxygen loss. Due to the high chemical expansion in the ab plane and the relative contraction in the c -direction upon oxygen loss, total volume expansion is almost linear over the temperature range, shifting from $19 \times 10^{-6} \text{ K}^{-1}$ from RT to 300, to $22 \times 10^{-6} \text{ K}^{-1}$ from 300 to 800 °C for $x = 0$ and 0.1. For $x = 0.2$ and 0.3, there is a clear shift also in volume expansion below and above 300 °C. The temperature behaviour of the lattice parameters and cell volume are given in Fig. 12a-d for BGLC ($x = 0, 0.1, 0.2$, and 0.3).

The volume expansion of BGLC shows a linear dependence on temperature for $x = 0$ and 0.1 despite substantial oxygen loss upon heating above 300 °C. The absence of a net chemical

**Fig. 4** SR-PXD with rietveld refinement of reduced, partly A-site ordered BLC.

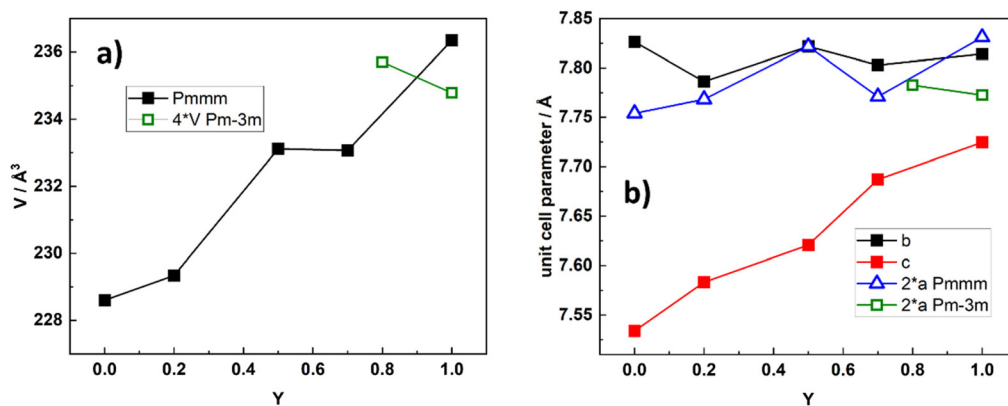


Fig. 5 Unit cell volume (a) and cell parameters (b) for BGLC ($x = 0, y = 0-1$).

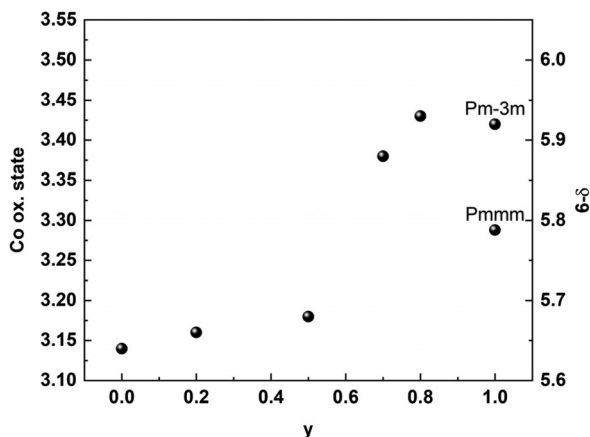


Fig. 6 Cobalt oxidation state and oxygen non-stoichiometry vs. y in BGLC ($x = 0, y = 0-1$).

volume expansion is seen also in BGCO and stems from anisotropic chemical expansion. This can be seen by comparing cell parameters as a function of temperature in air and Ar. The formation of oxygen vacancies has an expanding effect in the a - b

plane and a contracting effect in the c direction, as exemplified in Fig. 13 for $x = 0.1$. The same effect of air and Ar can be seen for $x = 0.2$ and 0.3 , while for $x = 0$, there is no change in a - or c -axis upon air vs. Ar atmosphere (Fig. 14).

It can also be seen from Fig. 13 and 14 that as oxygen loss starts at ~ 300 °C, the thermal expansion increases in the ab -plane and decreases in the c -direction, both in Ar and Air atmospheres.

This anisotropic change in a/b - and c -axes can be seen as a result of reduction, and hence a higher oxygen non-stoichiometry with increasing temperature. It has previously been shown that the formation of oxygen vacancies in $\text{LnBaCo}_2\text{O}_{6-\delta}$ double perovskites induces a contraction in the c -axis and an expansion in the ab -plane.¹⁷ Two main reasons may be suggested for this expansion behaviour; coulombic forces, and red-ox of the B-site cation giving changes in the ionic radius of Co. When a new oxygen vacancy is formed, the radius of the vacancy is smaller than for the host oxide ion,¹⁸ but it still causes coulombic repulsion between the surrounding cations.¹⁹ This repulsion can be counteracted by an inward attraction felt by the surrounding oxide ions towards the oxygen vacancy (Fig. 15).

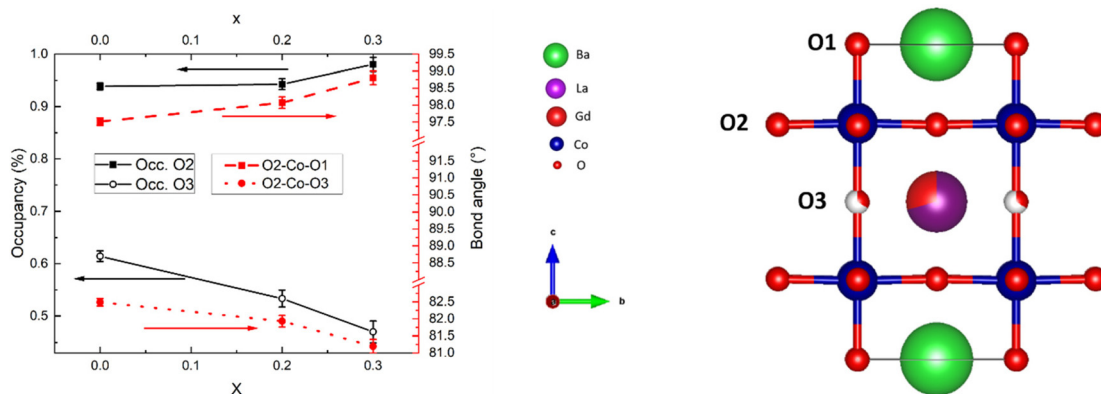


Fig. 7 SR-PXD data at room temperature showing occupancies on O2 and O3 sites (left axis) and shift in O2-Co-O1 and O2-Co-O3 bond angles (right axis) with increasing La substitution x .



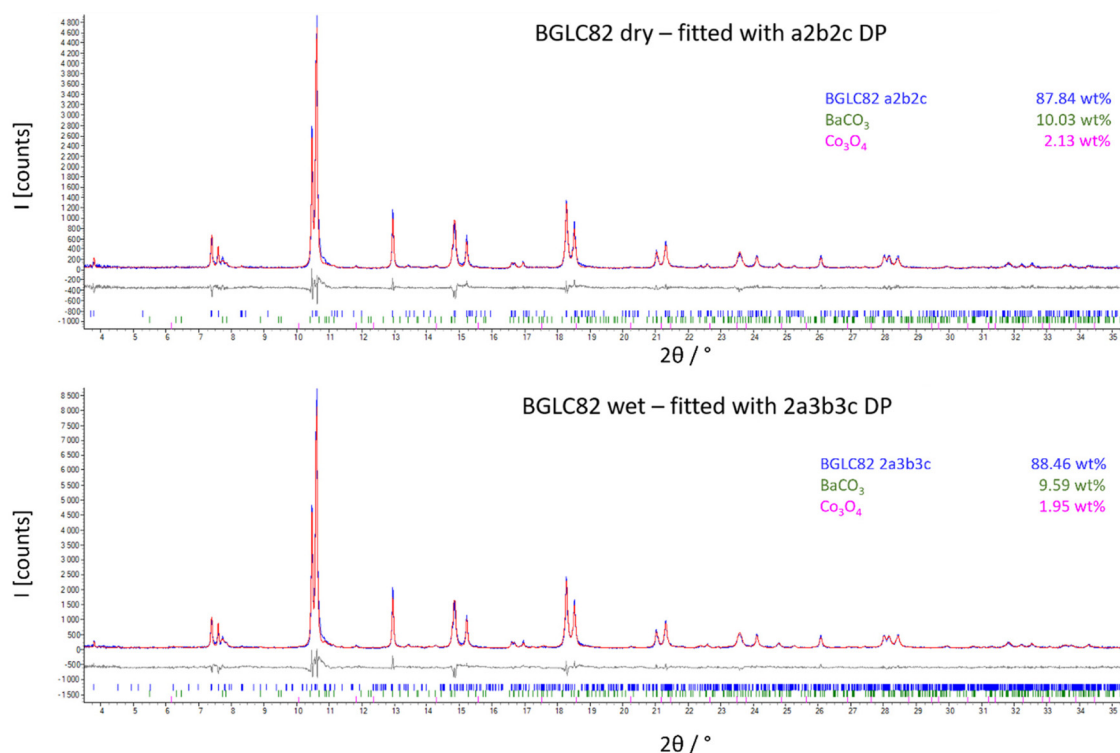


Fig. 8 SR-PXD of BGLC82 treated in dry (upper) and wet (lower) atmosphere at 300 °C. The spectra are fitted to a *Pmmm* a2b2c (dry), and a *Pmmm* 2a3b3c (wet) unit cell.

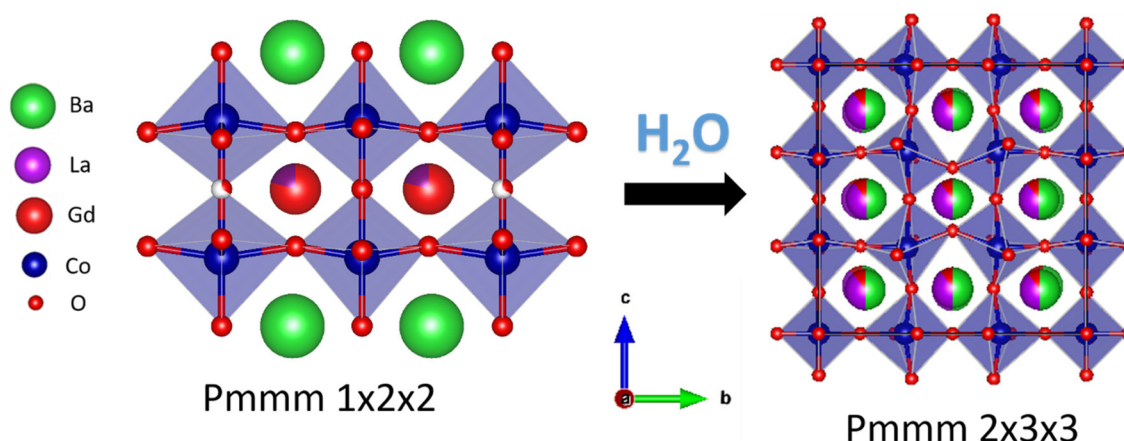


Fig. 9 Proposed order–disorder transition of BGLC82 upon hydration.

The tendencies of O2 to shift position towards the Ln layer with increased concentrations of O vacancies located there can be seen from the relation between O3 occupancies and O2–Co–O3 bond angles in Fig. 7.

The HT-XRD analysis (Fig. 12) shows different temperature behaviour for the two compositions with higher A^I substitution ($x = 0.2$ and 0.3) than for the two with lower ($x = 0$ and 0.1). Looking at the *a*-axis, it can be seen that the two lower substituted compositions ($x = 0$ and 0.1) go from low TEC (17 and $18 \times 10^{-6} \text{ K}^{-1}$) in the temperature interval from RT to 300 °C to

higher values (25 and $26 \times 10^{-6} \text{ K}^{-1}$) in the temperature interval from 350 to 850 °C. In the high temperature interval, the structure releases oxygen, and the increased TEC can thus be suggested to stem from an added expansion in the *a*–*b* plane due to formation of oxygen vacancies. If we compare with the *c*-axis, the trend is opposite. The TEC for the two same compositions is lowered in the high temperature interval, suggesting that chemical expansion has a negative contribution to the overall TEC, and that the *c*-axis thus contracts upon formation of oxygen vacancies.



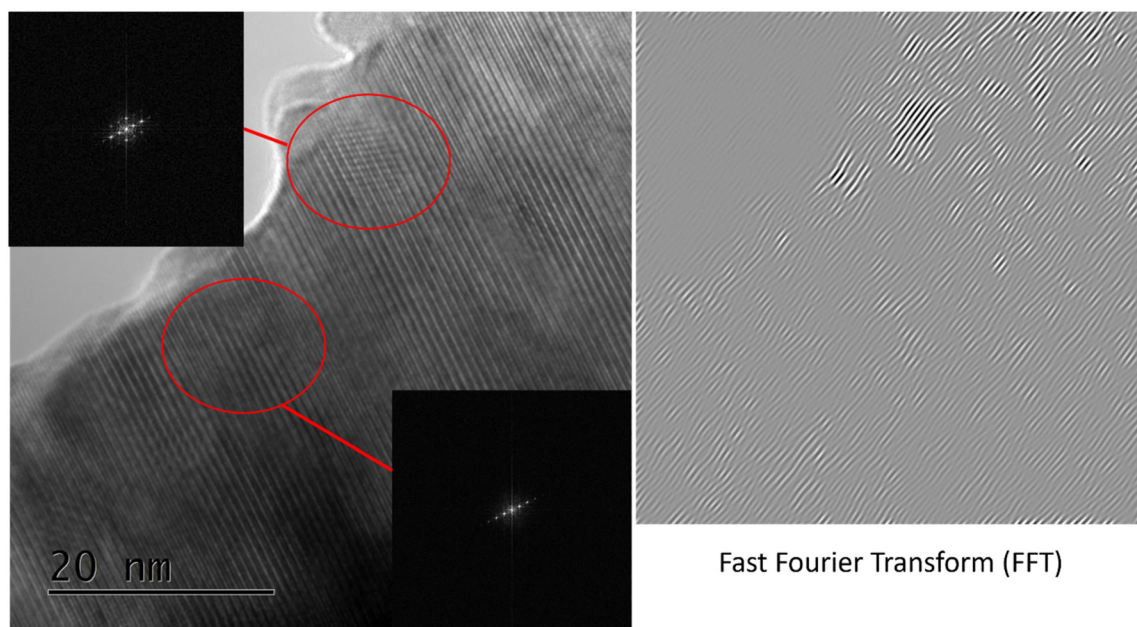


Fig. 10 SAED and FFT images of BGLC 82, $x = 0$, $y = 0.2$.

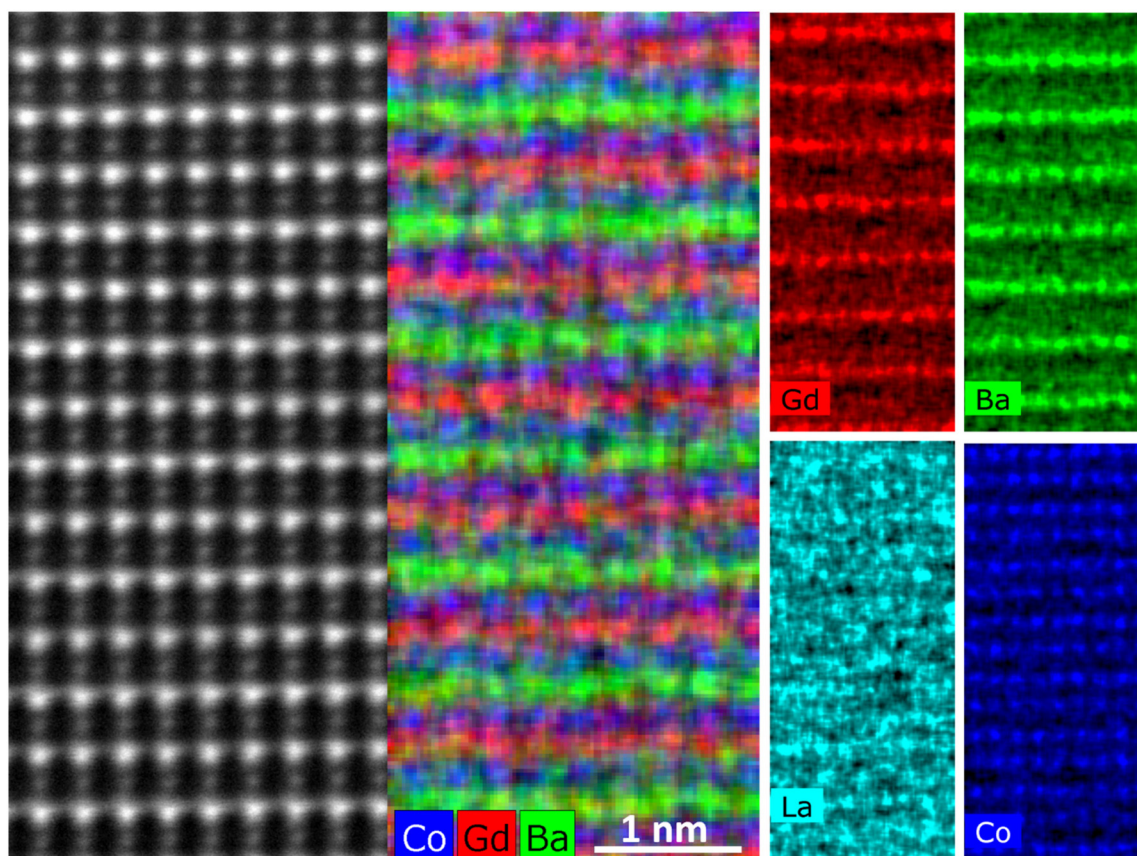


Fig. 11 STEM dark-field image (left) and EDS elemental mapping (middle and right) of BGLC82 in dry state.

For the two compositions with 20 and 30% La substitution for Ba on A^1 ($x = 0.2$ and 0.3), the trends are significantly altered. High TECs ($33\text{--}36 \times 10^{-6} \text{ K}^{-1}$) for both a - and c -axes in

the low temperature interval are lowered to $13 \times 10^{-6} \text{ K}^{-1}$ for the a -axis and $9 \times 10^{-6} \text{ K}^{-1}$ for the c -axis above $350 \text{ }^\circ\text{C}$ through an apparent phase transition.





Fig. 12 Thermal and chemical expansion of *a*- and *b*-axes and volume expansion in air for $x = 0$ (a), $x = 0.1$ (b), $x = 0.2$ (c), and $x = 0.3$ (d).



Fig. 13 Cell parameters for *a*-axis (left) and *c*-axis (right) vs. temperature for $x = 0.1$ in air (closed squares) and Ar (open squares) atmospheres.

The *z*-coordinate of the B-site Co in the structure varies slightly between the four compositions, albeit shifted towards the Ln layer for all compositions, indicating that the layered structure is not interrupted. Upon heating, the Co *z*-coordinate moves from an off-centred position, shifted towards the Ln layer to a symmetric position of 0.25 above 600 °C, indicating that some A-site ordering may be lost above this temperature. The positioning of the Co *z*-coordinate vs. temperature is given for BGLC ($x = 0$, $y = 0.2$) in air and inert atmosphere in Fig. 16.

Table 3 gives the extracted thermal expansion coefficients – including chemical expansion – for each axis and the volumetric thermal expansion coefficients at high and low temperatures and in air and Ar for all compositions.

Conclusions

We have shown the development of crystalline phases in double perovskite $\text{Ba}_{1-x}\text{Gd}_{1-y}\text{La}_{x+y}\text{Co}_2\text{O}_{6-\delta}$ (BGLC) in the inter-





Fig. 14 Cell parameters for *a*-axis (left) and *c*-axis (right) vs. temperature for $x = 0$ in air (closed squares) and Ar (open squares) atmospheres.



Fig. 15 Coulombic forces around an oxygen vacancy in the Ln layer.

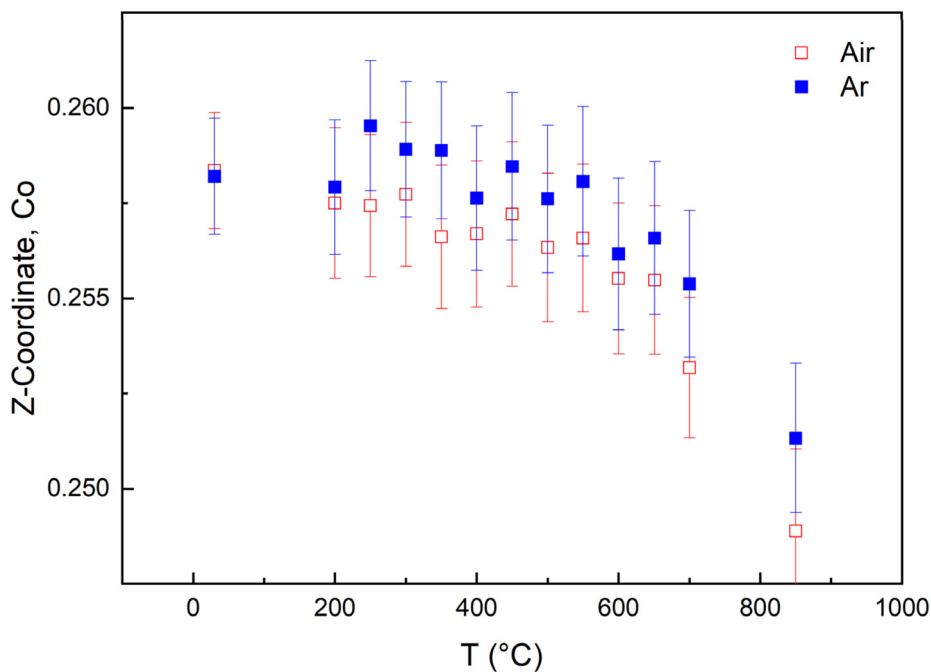


Fig. 16 Z-coordinate of Co in BGLC ($x = 0, y = 0.2$) upon heating from RT to 850 °C in air and Ar atmosphere.



Table 3 Thermal expansion coefficient for *a*- and *c*-axes in air and Ar at high and low temperatures for all compositions. Volume thermal expansion coefficients in air and Ar at high and low temperatures for all compositions

	Atmosphere	<i>T</i> (°C)	TEC ($\times 10^{-6} \text{ K}^{-1}$)			
			<i>x</i> = 0	<i>x</i> = 0.1	<i>x</i> = 0.2	<i>x</i> = 0.3
<i>a</i> -Axis	Air	RT–300	17	18	33	34
		350–850	25	25	13	13
	Ar	RT–300	16	21	34	35
		350–850	24	26	14	14
<i>c</i> -Axis	Air	RT–300	25	23	36	34
		350–850	15	15	9	9
	Ar	RT–300	25	23	35	36
		350–850	15	15	8	9
Volume expansion	Air	RT–300	19	19	34	34
		350–850	22	22	12	12
	Ar	RT–300	19	22	34	36
		350–850	21	22	12	12

vals $x = 0-0.5$, $y = 0.2$ and $x = 0$, $y = 0-1$. The end members are BGLC587 ($x = 0.5$, $y = 0.2$), BGC ($x = 0$, $y = 0$), and BLC ($x = 0$, $y = 1$). The most studied and reported composition of BGLC with $x = 0$, $y = 0.2$ (BGLC82) is found to be a double perovskite with 79 wt% orthorhombic *Pmmm*, 11 wt% orthorhombic Ba-deficient *Pbnm*, and minor impurities of BaCO_3 and Co_3O_4 . Substitution of La for Ba on the *Pmmm* A¹ site gives formation of a cubic LaCoO_3 -based *R3c* phase when $x \geq 0.2$. The end member BGLC587 is a three-phase material composed of 50 wt% LaCoO_3 with space group *R3c*, 23 wt% $\text{BaGd}_{0.8}\text{La}_{0.2}\text{Co}_2\text{O}_{6-\delta}$ with space group *Pmmm* and 22 wt% GdCoO_3 with space group *Pbnm*, where Ba is distributed over *Pmmm* and *Pbnm*, and Gd is distributed over all three phases. For the compositions with $x = 0.1$ and 0, the dominating phase is always *Pmmm*. By increasing La-substitution for Gd (with $x = 0$), the orthorhombic phase dominates until $y = 0.9$ and 1, where the structure can be stabilised in *Pmmm* or cubic *Pm3m* depending on thermal and atmospheric history. The thermal expansion of BGLC is generally anisotropic, where reduction causes chemical expansion and contraction in the *a*-*b* plane and *c*-direction, respectively. Upon hydration, we propose the formation of an oxidised $2a3b3c$ *Pmmm* supercell formation with disordering of Ba and La, giving domains where Bragg-reflections overlap with the *Pmmm* $a2b2c$ unit cell.

Conflicts of interest

There are no conflicts to declare.

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References

- R. Strandbakke, V. A. Cherepanov, A. Y. Zuev, D. S. Tsvetkov, C. Argiris, G. Sourkouni, S. Prunte and T. Norby, *Solid State Ionics*, 2015, **278**, 120.
- E. Vøllestad, R. Strandbakke, M. Tarach, D. Catalán-Martínez, M.-L. Fontaine, D. Beeff, D. R. Clark, J. M. Serra and T. Norby, *Nat. Mater.*, 2019, **18**(7), 752.
- D. Malyshev, A. Novikov, I. Ivanov, V. Sereda, D. Tsvetkov and A. Zuev, *J. Alloys Compd.*, 2020, **845**, 156309.
- S. L. Wachowski, I. Szpunar, M. H. Sørby, A. Mielewczyk-Gryń, M. Balaguer, C. Ghica, M. C. Istrate, M. Gazda, A. E. Gunnæs, J. M. Serra, T. Norby and R. Strandbakke, *Acta Mater.*, 2020, **199**, 297.
- N. Ishizawa, T. Asaka, T. Kudo, K. Fukuda, A. Yasuhara, N. Abe and T.-h. Arima, *Chem. Mater.*, 2014, **26**(22), 6503.
- A. Grimaud, K. J. May, C. E. Carlton, Y.-L. Lee, M. Risch, W. T. Hong, J. Zhou and Y. Shao-Horn, *Nat. Commun.*, 2013, **4**(1), 2439.
- D. S. Tsvetkov, I. L. Ivanov, D. A. Malyshev and A. Y. Zuev, *Dalton Trans.*, 2014, **43**(31), 11862.
- A. A. Taskin, A. N. Lavrov and Y. Ando, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2005, **71**(13), 134414.
- C. Frontera, A. Caneiro, A. E. Carrillo, J. Oró-Solé and J. L. García-Muñoz, *Chem. Mater.*, 2005, **17**(22), 5439.
- A. Maignan, C. Martin, D. Pelloquin, N. Nguyen and B. Raveau, *J. Solid State Chem.*, 1999, **142**(2), 247.
- E. Vøllestad, M. Schrade, J. Segalini, R. Strandbakke and T. Norby, *J. Mater. Chem. A*, 2017, **5**(30), 15743.
- C. Bernuy-Lopez, K. Høydalsvik, M.-A. Einarsrud and T. Grande, *Materials*, 2016, **9**(3), 154.
- I. Szpunar, S. Wachowski, T. Miruszewski, K. Dzierzgowski, K. Górnicka, T. Klimczuk, M. H. Sørby, M. Balaguer, J. M. Serra, R. Strandbakke, M. Gazda and A. Mielewczyk-Gryń, *J. Am. Ceram. Soc.*, 2020, **103**(3), 1809.
- Y. Q. Jia, *J. Solid State Chem.*, 1991, **95**(1), 184.
- I. Szpunar, R. Strandbakke, M. H. Sørby, S. L. Wachowski, M. Balaguer, M. Tarach, J. M. Serra, A. Witkowska, E. Dzik, T. Norby, M. Gazda and A. Mielewczyk-Gryń, *Materials*, 2020, **13**(18), 4044.
- A. Mielewczyk-Gryń, S. Yang, M. Balaguer, R. Strandbakke, M. H. Sørby, I. Szpunar, A. Witkowska, S. Wachowski, J. M. Serra, A. Navrotsky and M. Gazda, under publication.
- D. S. Tsvetkov, I. L. Ivanov, D. A. Malyshev, A. L. Sednev, V. V. Sereda and A. Y. Zuev, *Pure Appl. Chem.*, 2019, **91**(6), 923.
- C. Chatziechristodoulou, P. Norby, P. V. Hendriksen and M. B. Mogensén, *J. Electroceram.*, 2015, **34**(1), 100.
- Y. Wang, C. Cai, L. Li, L. Yang, Y. Zhou and G. Zhou, *AIP Adv.*, 2016, **6**(9), 095113.

